



72° CONGRESSO
BRASILEIRO DE
CARDIOLOGIA

03 a 05
Novembro | 2017

SÃO PAULO EXPO
EXHIBITION & CONVENTION CENTER



Mesa-Redonda

Avaliação do Paciente com Estenose Aórtica de Moderado e Alto Risco Cirúrgico

Visão do Cirurgião Cardíaco

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Cirurgião Cardiovascular

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Declaração de Potencial Conflito de Interesse

Nome do Palestrante:

Renato A. K. Kalil

Título da Apresentação:

Avaliação do Paciente com Estenose Aórtica de Moderado e Alto
Risco Cirúrgico

Visão do Cirurgião Cardíaco

**Não possuo nenhum conflito de interesse relacionado a
esta apresentação**

Definição do Risco

Escores mais utilizados tem base em pacientes com cardiopatia isquêmica

STS score (<http://209.220.160.181/STSWebRiskCalc261/>)

Euroscore I e II (www.euroscore.org/calc.html)

Para valvulares:

Ambler: Ambler, G., Omar, R.Z., Royston, P. et al. Generic, simple risk stratification model for heart valve surgery. *Circulation*. 2005; 112: 224–231

Guaragna: Guaragna JC, Bodanese LC, Bueno FL, Goldani MA. *Arq Bras Cardiol*. 2010 Apr;94(4):541-8

Escores mais específicos, incluindo itens como fragilidade, agilidade e força, tem sido propostos

Validade dos Escores de Risco

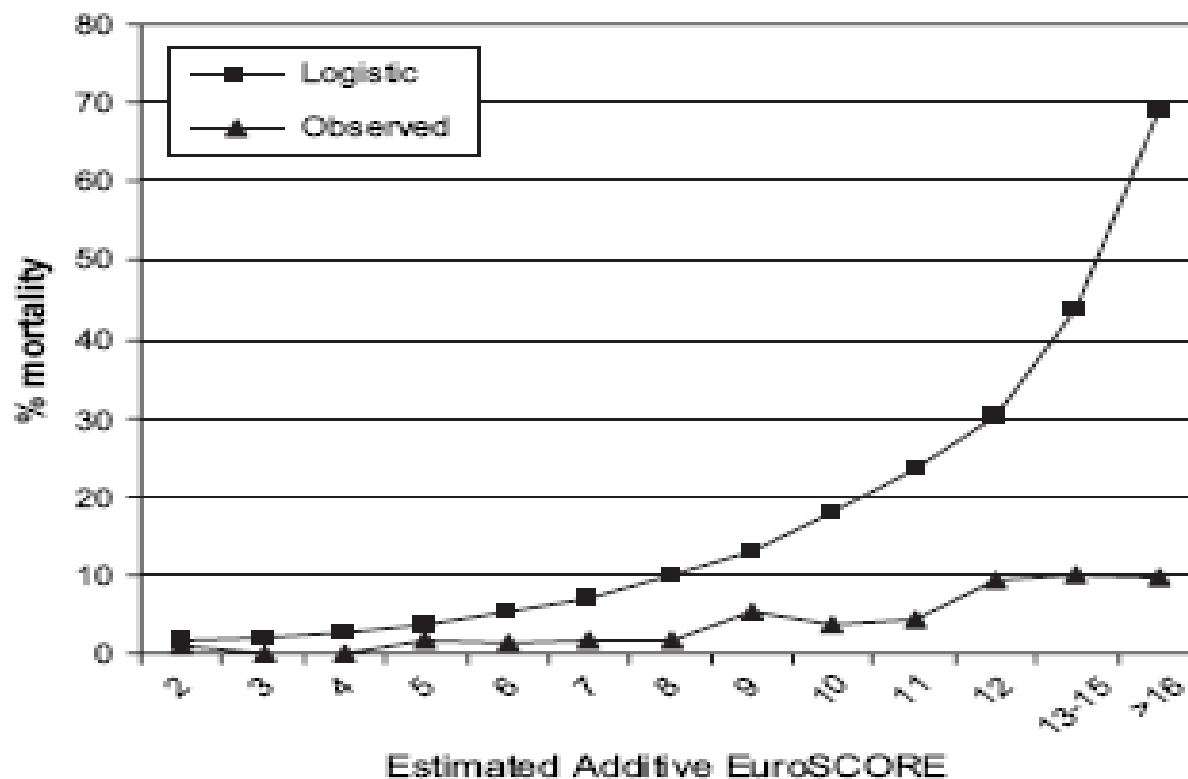
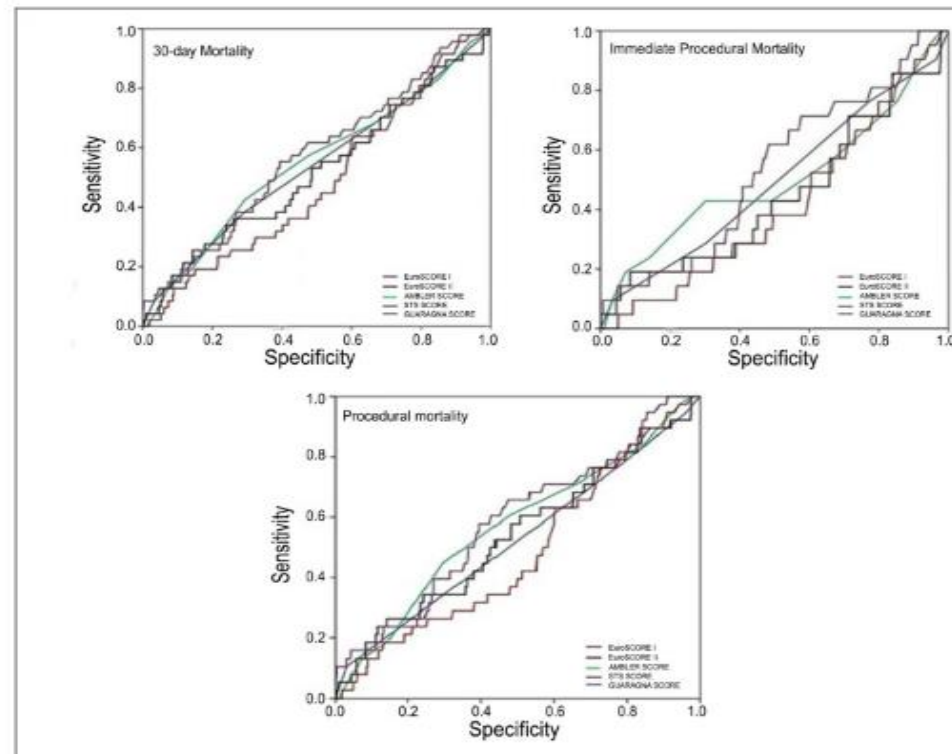


Figure 2. Actual and predicted mortality for the logistic EuroSCORE model. The mortality rate is on the Y axis, and patients are divided into groups on the basis of the predicted additive EuroSCOREs on the X axis.

Is the European System for Cardiac Operative Risk Evaluation model valid for estimating the operative risk of patients considered for percutaneous aortic valve replacement?

Performance of surgical risk scores to predict mortality after transcatheter aortic valve implantation.

Figure 1



ROC curves for the outcomes assessed using the different surgical risk scores.

CONCLUSIONS:

In this real world Brazilian registry, the surgical risk scores were inaccurate in predicting mortality after TAVI. Risk models specifically developed for TAVI are required.

Assessment of Commonly Used Frailty Markers for High- and Extreme-Risk Patients Undergoing Transcatheter Aortic Valve Replacement

4 itens:

albumina sérica
teste caminhada
força, *grip strenght*,
índice Katz de
independência
atividade diária

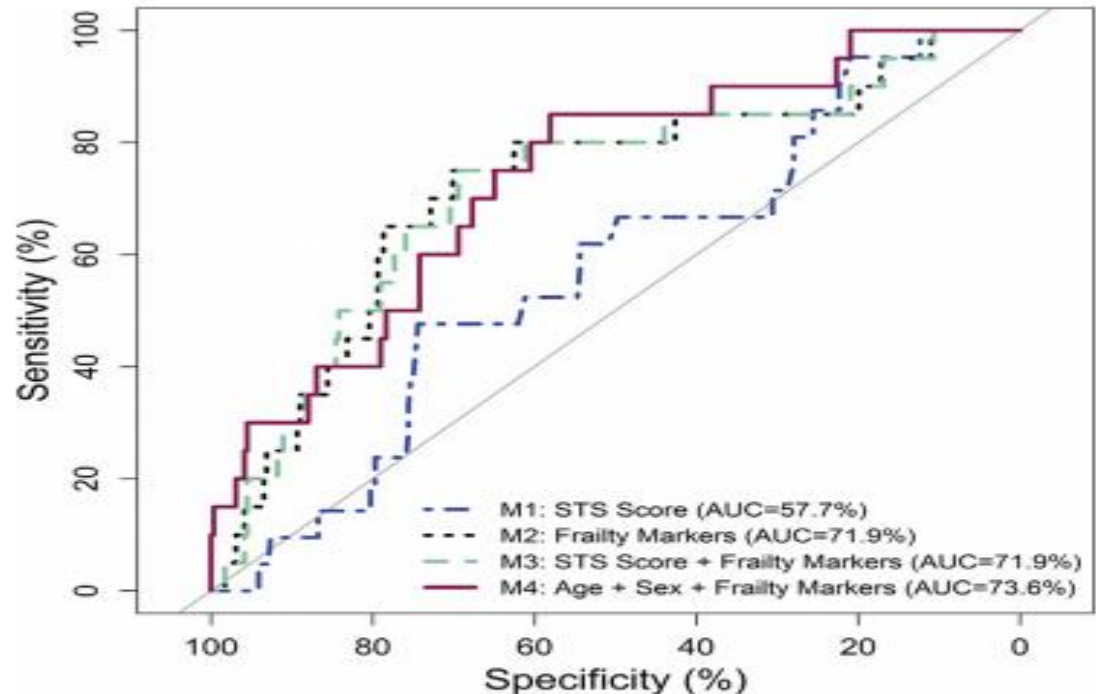


Fig 1. Receiver-operating characteristics curves for 30-day mortality (M): M1 (blue line) The Society of Thoracic Surgeons (STS) score (area under the curve [AUC] = 57.7%); M2 (black line) frailty markers (AUC = 71.9%); M3 (green line) STS score plus frailty markers (AUC = 71.9%); and M4 (purple line) age plus sex plus frailty markers (AUC = 73.6%).

Resultados da Cirurgia Convencional Aberta

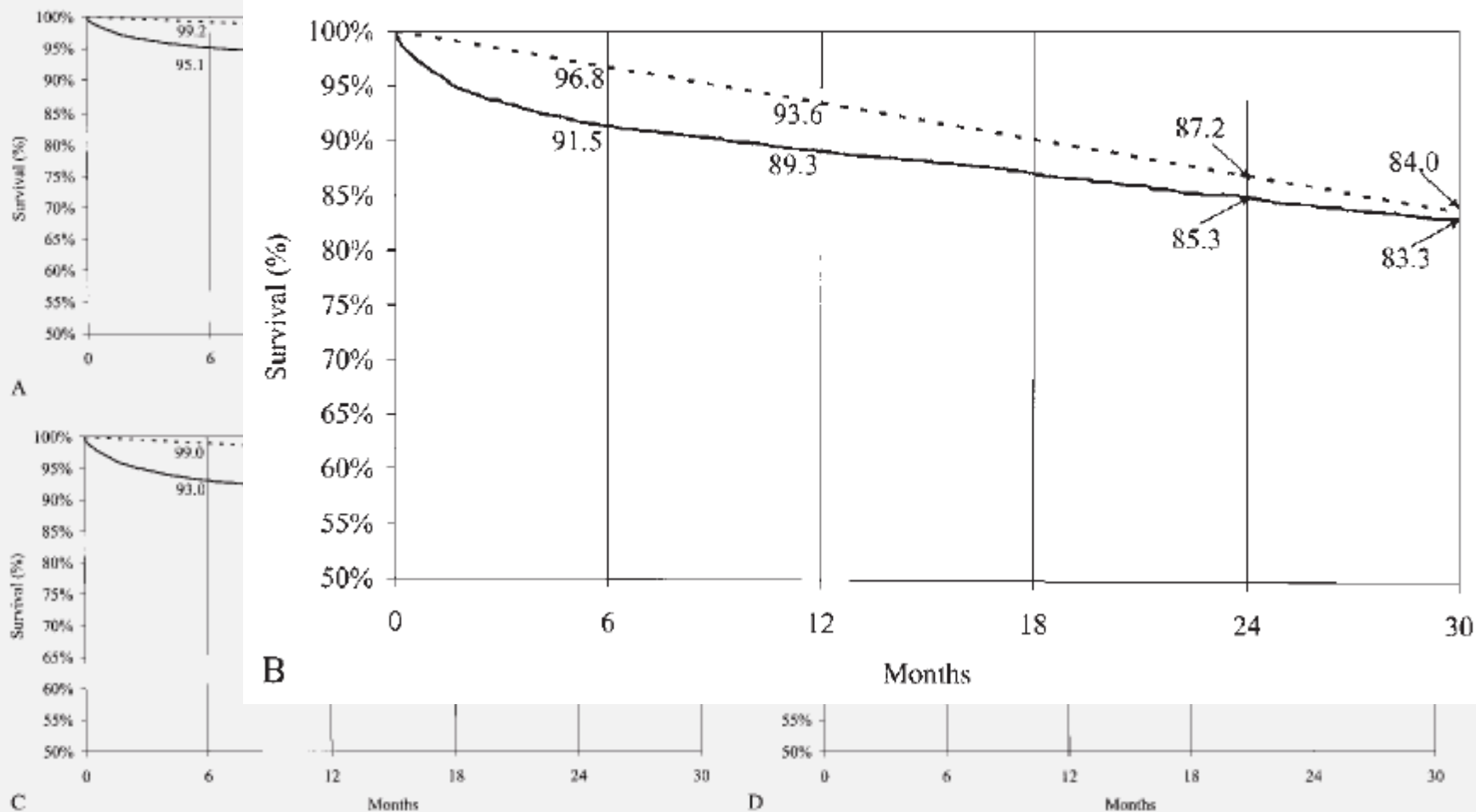


Fig 2. Survival after aortic valve replacement according to patient age. Dashed lines are survival for age- and sex-matched US population. Solid lines represent risk-adjusted survival in selected age and surgery subgroups. (A) Nonelderly patients (age < 75 years) with isolated aortic valve replacement. (B) Elderly patients (age > 75 years) with isolated aortic valve replacement. (C) Nonelderly patients undergoing aortic valve replacement with coronary artery bypass graft surgery. (D) Elderly patients undergoing aortic valve replacement with coronary artery bypass graft surgery.

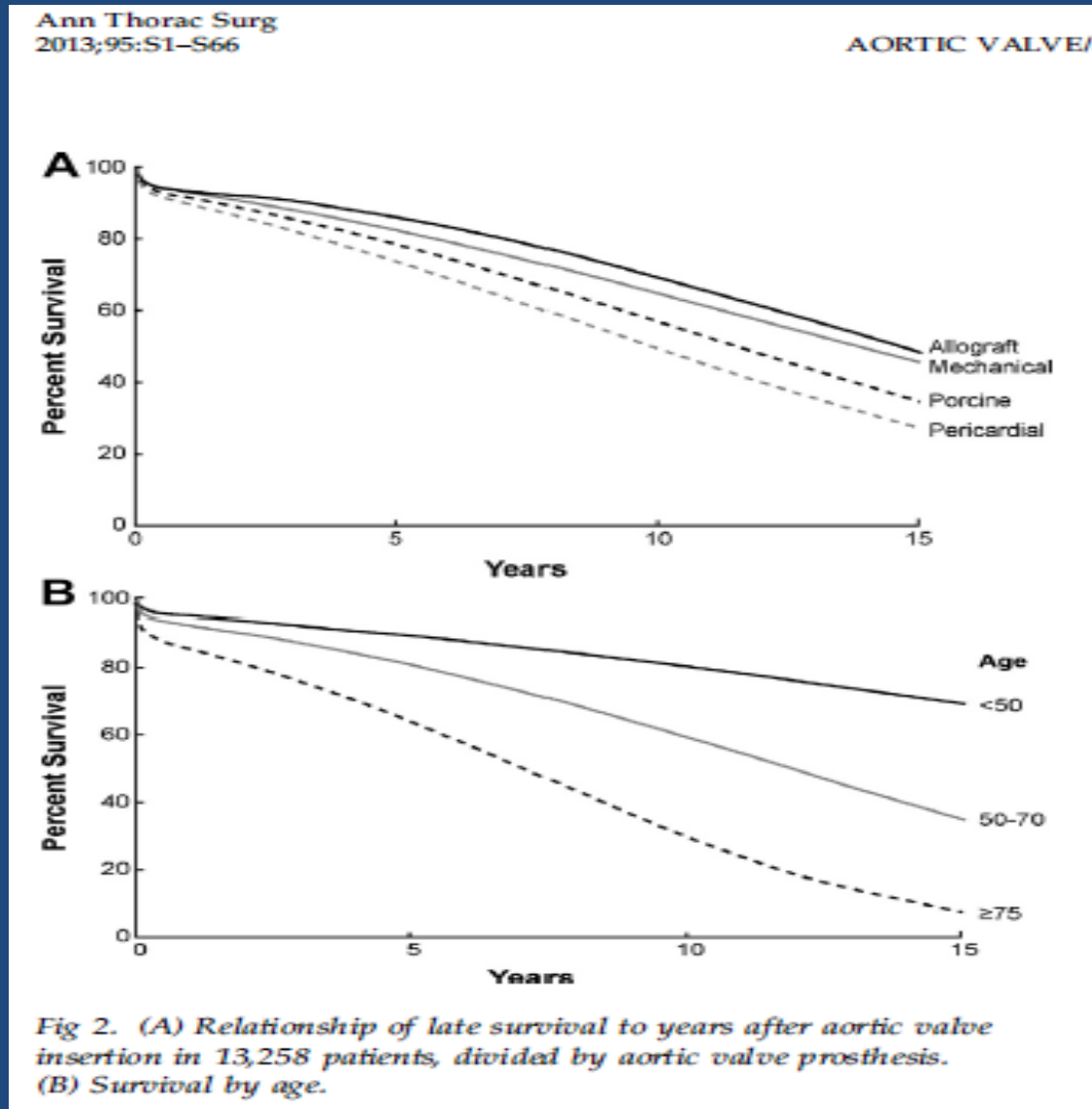
- **Table 3 Multivariable Predictors for 30-Month Survival in Patients With Severe Aortic Stenosis Undergoing Aortic Valve Replacement Surgery in New York State From 2003 to 2005**

- | Predictor | Prevalence | Coefficient | p Value | HazardRatio |
|-----------|------------|-------------|---------|-------------|
| Age <65 y | 17.57 | Reference | 1.00 | |
| 65–74 | 28.75 | 0.453 | 0.003 | 1.57 |
| 75–84 | 44.36 | 0.7807 | <0.0001 | 2.18 |
| >85 | 9.33 | 1.3767 | <0.0001 | 3.96 |

Mortalidade hospitalar média 3,97% para Ao isolada

Hannan, Ann Thorac Surg 2009

Relação Sobrevida X Idade

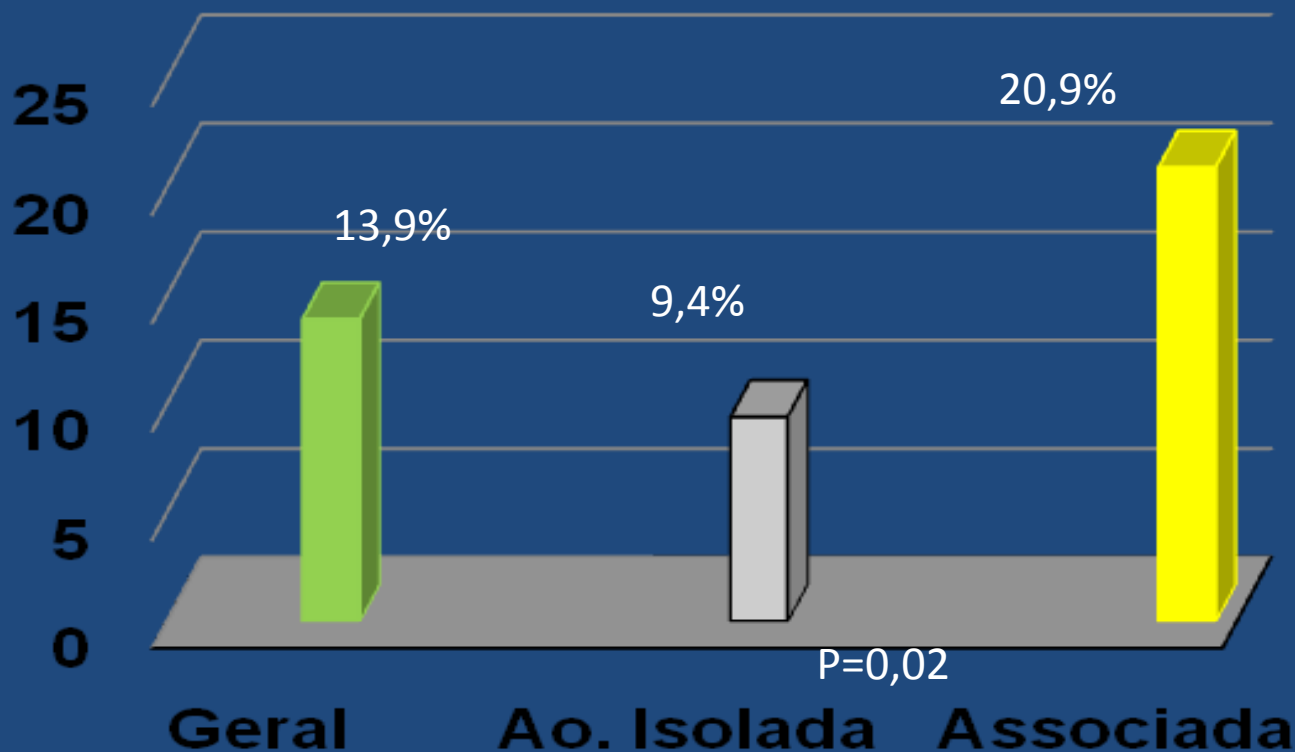


Cirurgia EAo >75anos no IC/FUC

Período 2002-2007

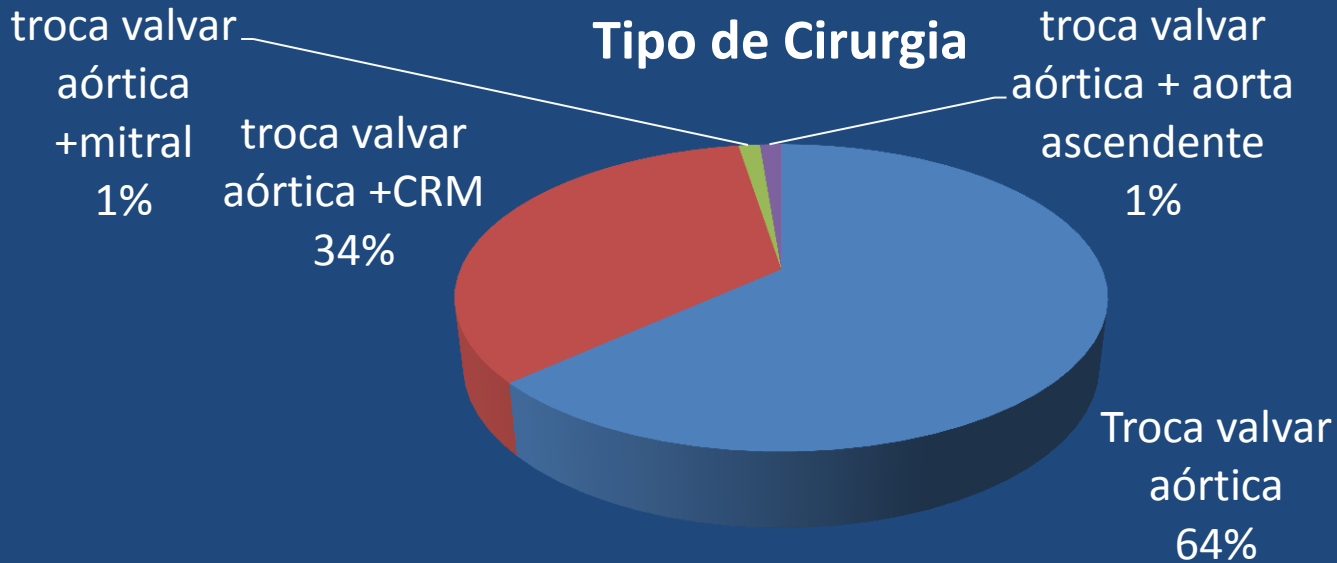
$n = 230$

Mortalidade hospitalar



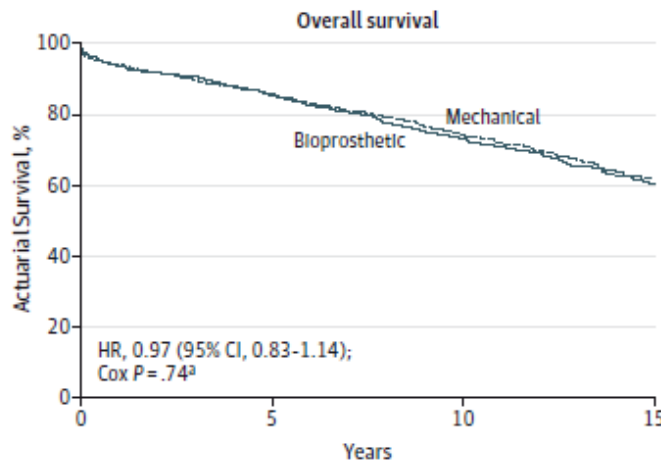
Pacientes acima de 70 anos submetidos a troca valvar aórtica por estenose no IC-FUC entre 2015-2016

n=79



| | | |
|--|---|------------------------------------|
| Mortalidade Geral: | 8 (10,1%) | |
| Mortalidade por tipo de cirurgia: | Troca valvar aórtica isolada: 5 de 49 (10,2%) | Cirurgia combinada: 3 de 30(10,0%) |
| Mortalidade por grupo etário: | 70-75 anos: 1 (1,2%) | >75 anos: 7 (8,8%) |
| Causa do óbito: | Cardiovasc: 3 (3,79%) | Não Cardiovasc: 5 (6,3%) |
| | | |

Figure 1. Overall Survival Among Propensity-Matched Patients Aged 50 to 69 Years After Bioprosthetic vs Mechanical Aortic Valve Replacement



| No. at risk | 0 | 5 | 10 | 15 |
|---------------|------|-----|-----|----|
| Bioprosthetic | 1001 | 860 | 589 | 91 |
| Mechanical | 1001 | 856 | 611 | 89 |

There were 322 all-cause deaths in the bioprosthesi group vs 318 in the mechanical prosthesis group.

^a P value calculated using a marginal Cox model with a robust sandwich variance estimator.

Survival and Long-term Outcomes Following Bioprosthetic vs Mechanical Aortic Valve Replacement in Patients Aged 50 to 69 Years

DESIGN, SETTING, AND PARTICIPANTS Retrospective cohort analysis of 4253 patients aged 50 to 69 years who underwent primary isolated aortic valve replacement using bioprosthetic vs mechanical valves in New York State from 1997 through 2004, identified using the Statewide Planning and Research Cooperative System. Median follow-up time was 10.8 years (range, 0 to 16.9 years); the last follow-up date for mortality was November 30, 2013. Propensity matching yielded 1001 patient pairs.

Based on NY Registry

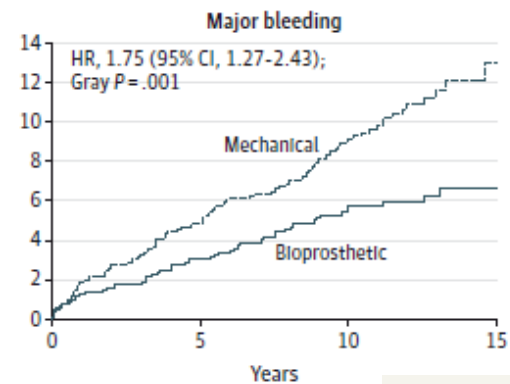
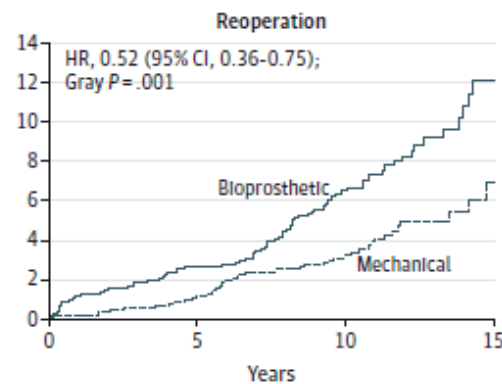
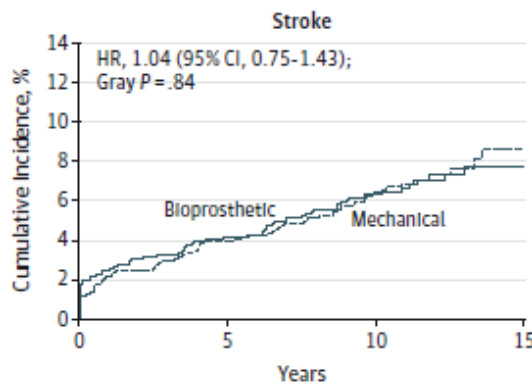
Mortality after complications:

18,7% after stroke

9,0% after reoperation

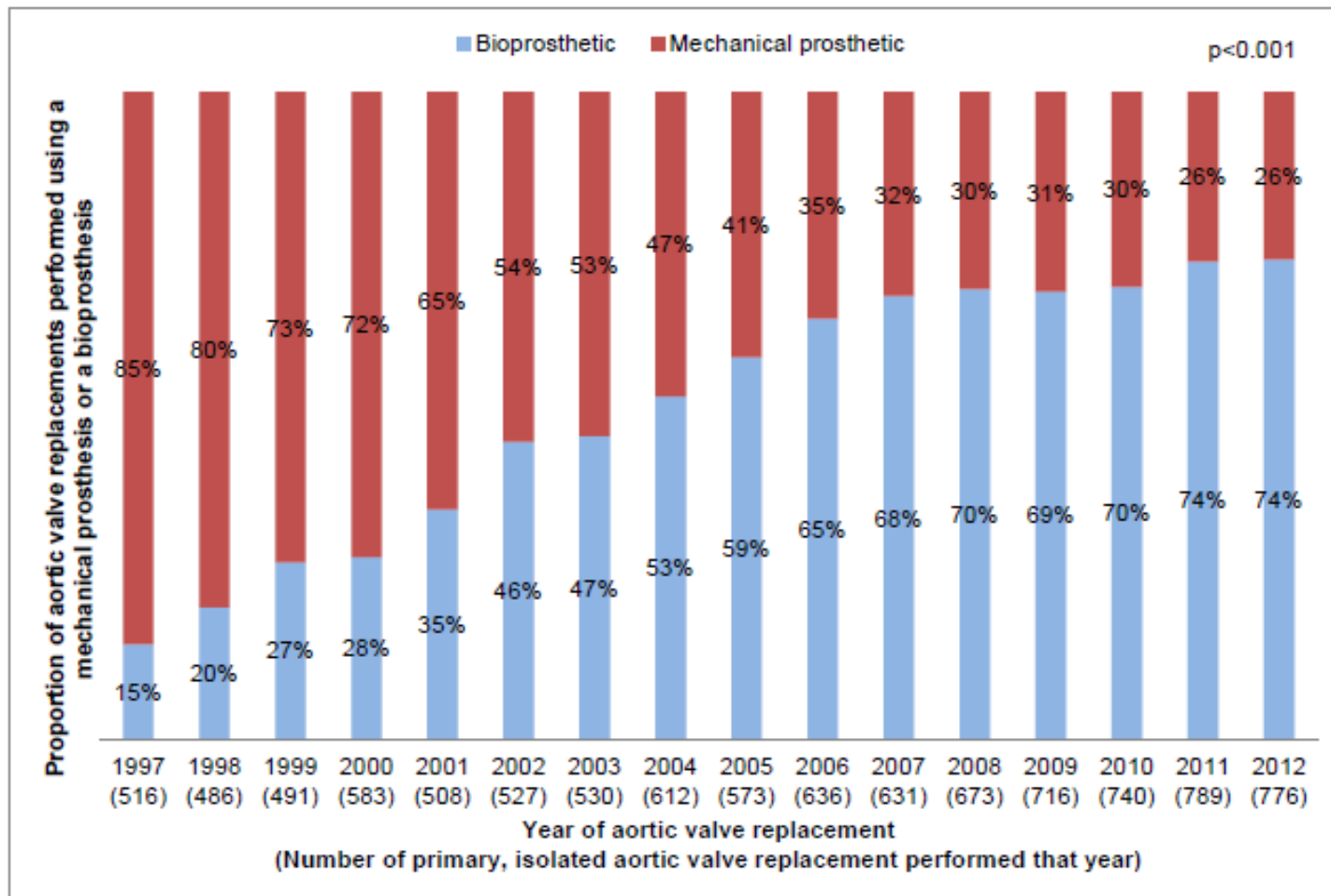
13,2% after major bleeding

Figure 2. Cumulative Incidence of Major Morbidity (Stroke, Reoperation, Major Bleeding) Among Propensity-Matched Patients Aged 50 to 69 Years After Bioprosthetic vs Mechanical Aortic Valve Replacement



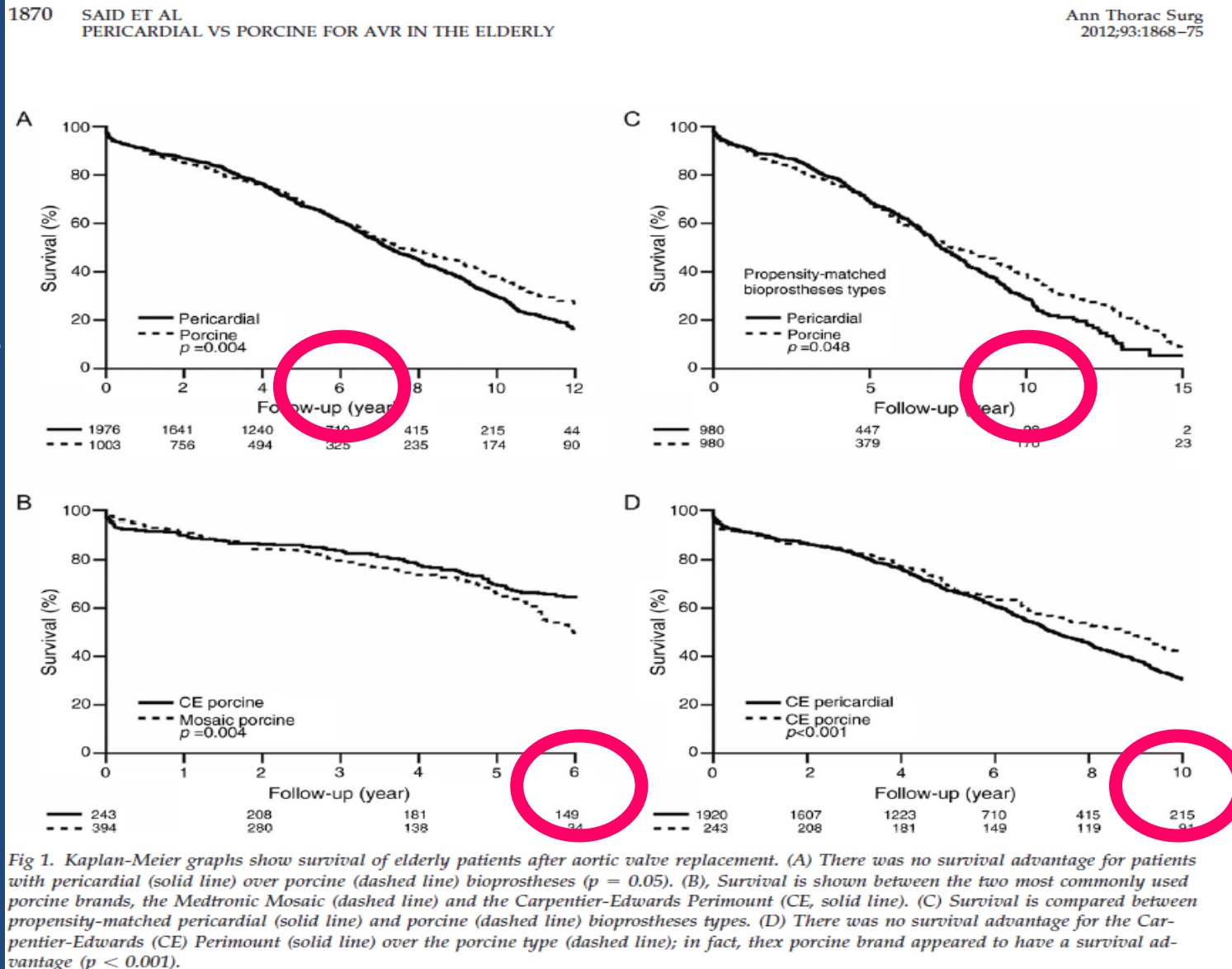
Survival and Long-term Outcomes Following Bioprosthetic vs Mechanical Aortic Valve Replacement in Patients Aged 50 to 69 Years

eFigure 2. Trend in Mechanical versus Bioprosthetic Valve Usage for Aortic Valve Replacement in Patients Aged 50 to 69 in New York State^a



Há tipos de biopróteses de maior durabilidade?

n=2979
 >65anos
 período
 1993-2007,
 Mayo,
 Mass Gen e
 Brigham



Long-Term Survival After Bovine Pericardial Versus Porcine Stented Bioprosthetic Aortic Valve Replacement: Does Valve Choice Matter?

Table 1. Stented Bioprosthetic Aortic Valves Included in Study

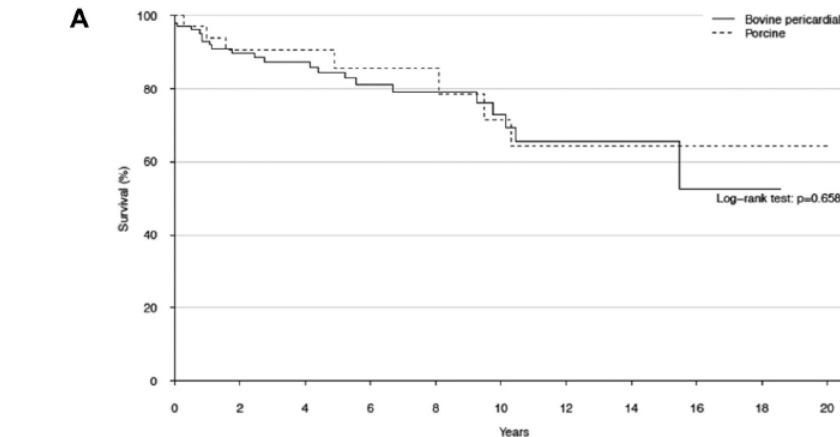
| Valves | Total (No.) | Isolated | |
|--|----------------|--------------|-------------------|
| | | AVR (No.) | AVR+CABG (No.) |
| Bovine pericardial | 1,411 | | |
| Carpentier-Edwards Perimount ^a | 1,273 | 734 | 539 |
| Sorin Mitroflow ^b | 26 | 16 | 10 |
| St. Jude Trifecta ^c | 112 | 51 | 61 |
| Porcine | 599 | | |
| St. Jude Biocor ^c | 128 | 46 | 82 |
| Carpentier-Edwards Porcine ^a | 210 | 111 | 99 |
| Medtronic Hancock ^d | 105 | 44 | 61 |
| Medtronic Mosaic ^d | 156 | 140 | 16 |

^a Edwards Lifesciences, Irvine, California. ^b Sorin Group Inc, Arvada, Colorado. ^c St. Jude Medical Inc, St. Paul, Minnesota. ^d Medtronic, Minneapolis, Minnesota.

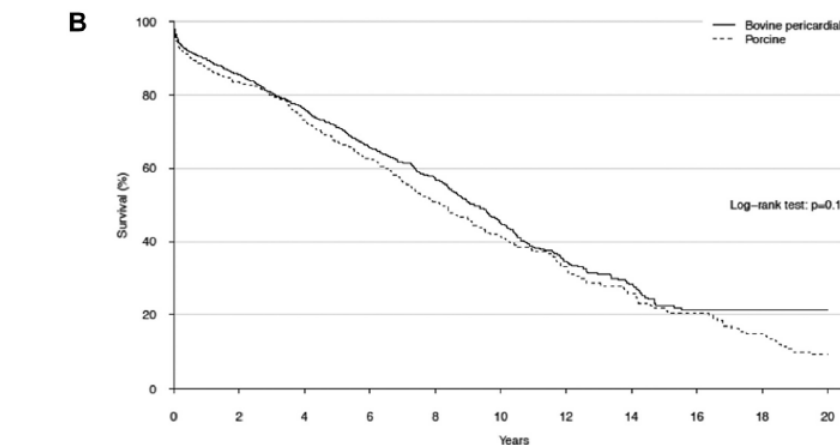
AVR = aortic valve replacement; CABG = coronary artery bypass grafting.

Long-Term Survival After Bovine Pericardial Versus Porcine Stented Bioprosthetic Aortic Valve Replacement: Does Valve Choice Matter?

Fig 4. Overall survival analysis in patients with bovine pericardial (solid line) and porcine (dashed line) valves according patient age (A) 18 to 55 years and (B) age older than 55 years at aortic valve replacement.



| | | | | | | | | | | |
|--------------------|-----|----|----|----|----|----|----|----|---|---|
| Bovine pericardial | 104 | 78 | 64 | 42 | 32 | 21 | 15 | 11 | 4 | 1 |
| Porcine | 34 | 27 | 18 | 17 | 13 | 10 | 8 | 7 | 7 | 7 |



| | | | | | | | | | | |
|--------------------|------|-----|-----|-----|-----|-----|-----|----|----|----|
| Bovine pericardial | 1307 | 917 | 725 | 472 | 318 | 173 | 109 | 64 | 23 | 10 |
| Porcine | 565 | 432 | 237 | 194 | 133 | 106 | 65 | 38 | 30 | 21 |

In conclusion, for patients undergoing AVR with a stented bioprosthetic valve, with or without CABG, the choice of a porcine vs bovine pericardial bioprosthesis does not appear to affect long-term survival or the need for reoperation, regardless of valve size or patient age. As such, stented bioprosthetic valves would appear to be fungible, and therefore, valve choice should be driven by local market factors similar to other commodities.

Sutureless Aortic Valves



PERCEVAL, Sorin



INTUITY, Edwards



Fig. 1. Photo of the ATS 3f Enable® Aortic Bioprosthesis Model 6000.

3f ENABLE, Medtronic Inc.



? Inovare, Braile

Resultados a curto e longo prazo de
TAVI em moderado e alto risco
cirúrgico

Transcatheter or Surgical Aortic Valve Replacement in
Intermediate Risk Patients
with Aortic Stenosis:
Final Results from the PARTNER 2A Trial

Craig R. Smith, MD

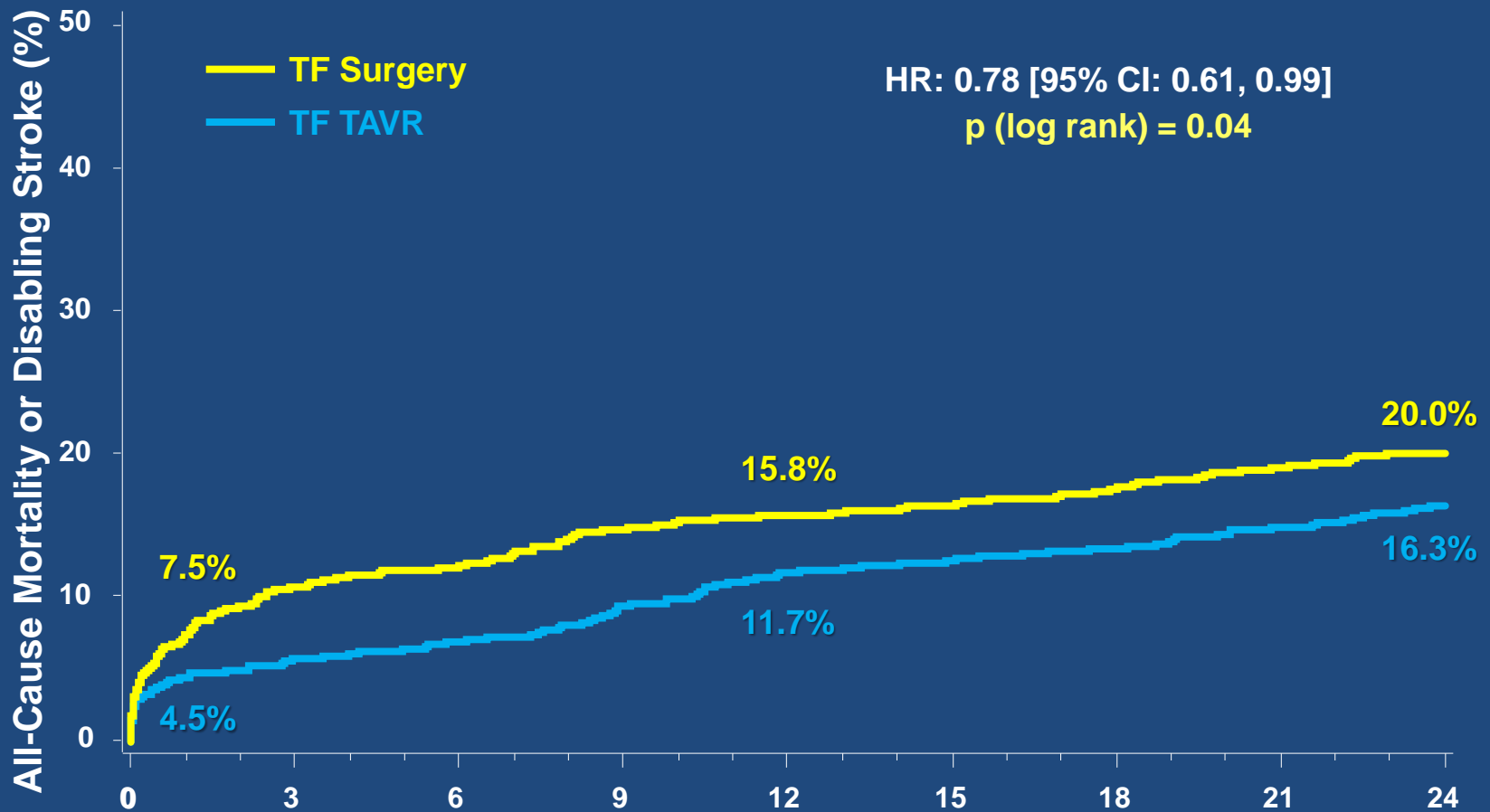
on behalf of the PARTNER Trial Investigators

ACC 2016 | Chicago | April 2, 2016



TF Primary Endpoint (AT)

All-Cause Mortality or Disabling Stroke



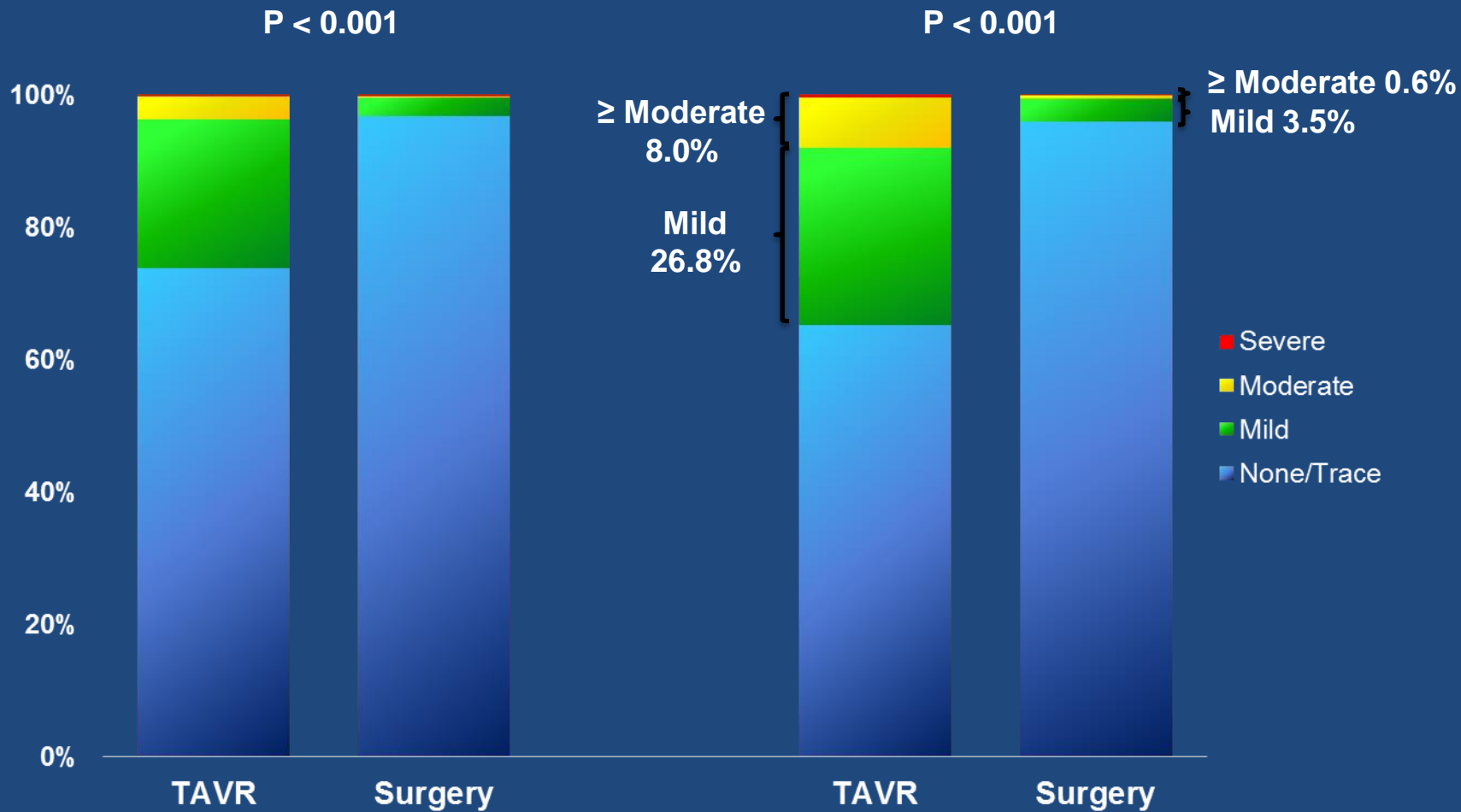
Number at risk:

| | 0 | 3 | 6 | 9 | 12 | 15 | 18 | 21 | 24 |
|------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| TF Surgery | 722 | 636 | 624 | 600 | 591 | 573 | 565 | 555 | 537 |
| TF TAVR | 762 | 717 | 708 | 685 | 663 | 652 | 644 | 634 | 612 |

Months from Procedure

Paravalvular Regurgitation (VI)

3-Class Grading Scheme



No. of echos

30 Days

2 Years

TAVR

872

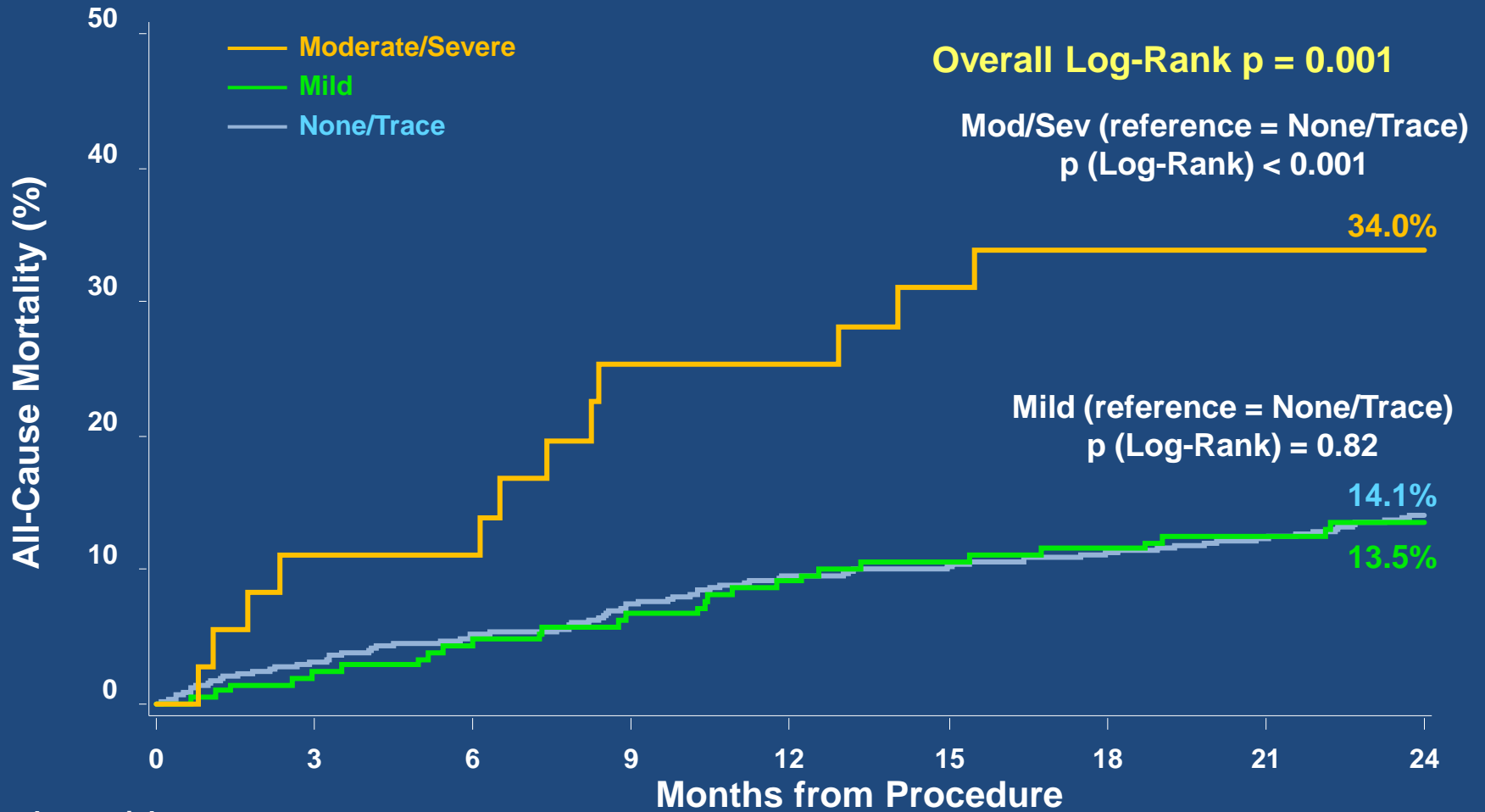
600

Surgery

757

514

Severity of PVR at 30 Days and All-cause Mortality at 2 Years (VI)

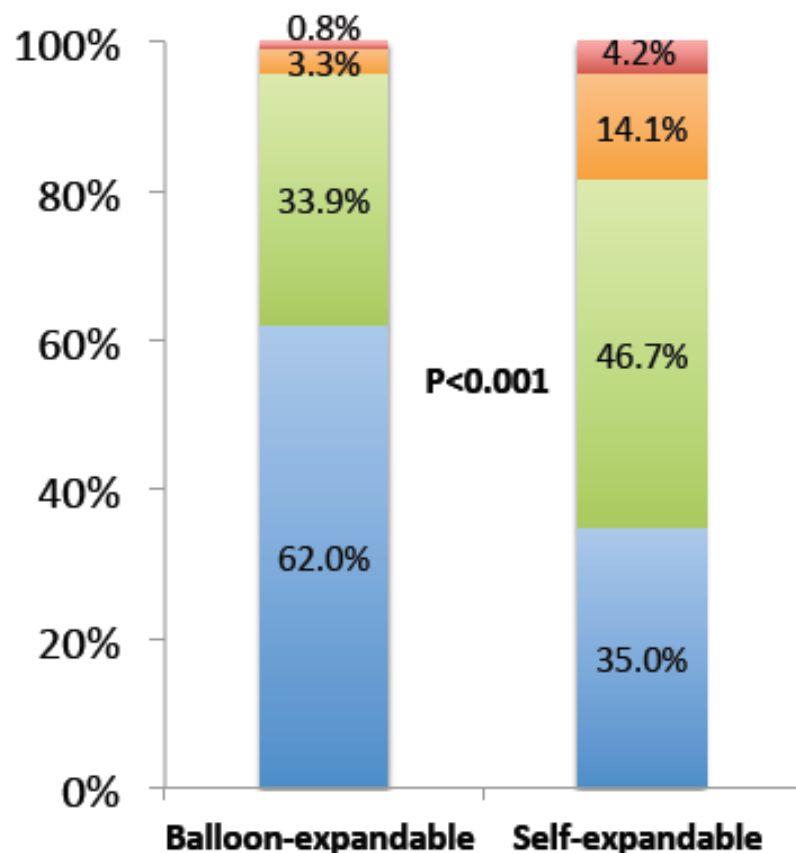


Number at risk:

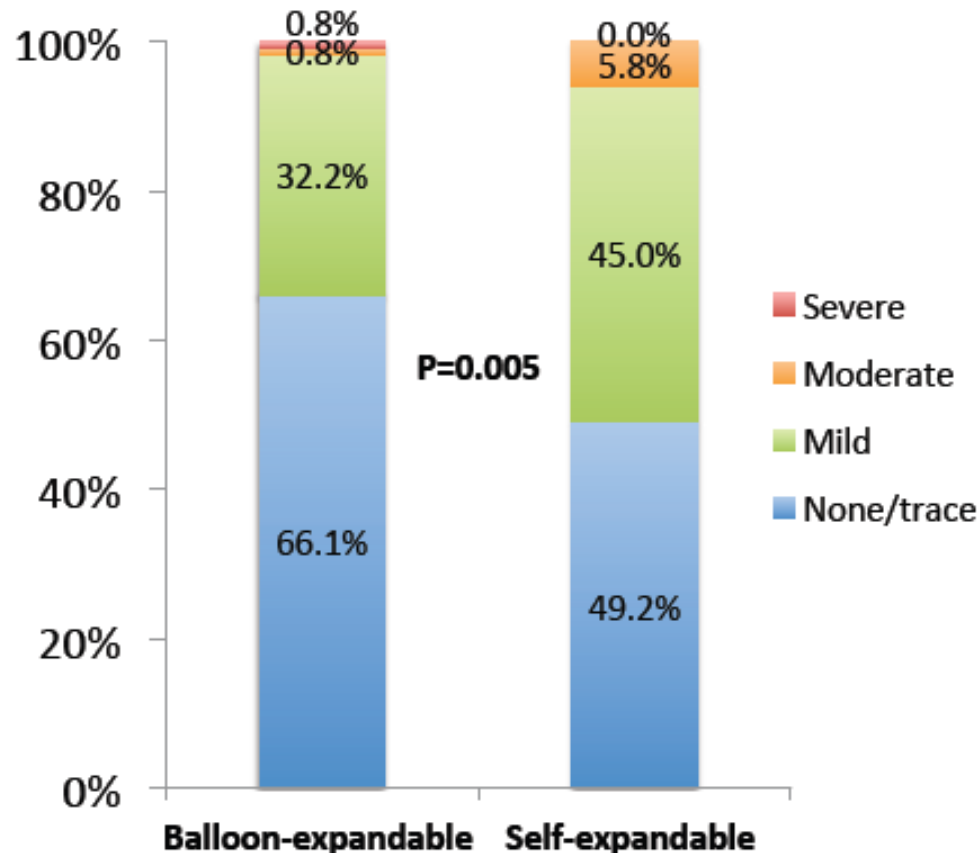
| | | | | | | | | | |
|--------------|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| Moderate/Sev | 36 | 32 | 32 | 26 | 26 | 24 | 22 | 22 | 21 |
| Mild | 210 | 204 | 199 | 194 | 188 | 184 | 182 | 180 | 175 |
| None/Trace | 701 | 678 | 664 | 647 | 628 | 621 | 612 | 605 | 585 |

Post-Procedural Aortic Regurgitation

AR by Angiography



AR by Echocardiography



| | Balloon-expandable (n=116) | Self-expandable (n=114) | p-value |
|------------------------|----------------------------|-------------------------|---------|
| Dimensionless AR Index | 29.0±7.1 | 27.3±7.2 | 0.08 |

The PARTNER 2A Trial

Clinical Implications

- *The results from PARTNER 2A support the use of TAVR as an alternative to surgery in intermediate risk patients, similar to those included in this trial.*
- In patients who are candidates for transfemoral access, TAVR may result in additional clinical advantages.
- Long-term durability assessments of transcatheter bioprosthetic valves are still lacking and extrapolation of these findings to low-risk patients requires further clinical trial validation.

Vieses dos ensaios clínicos

De metodologia e de interpretação
dos resultados

ORIGINAL ARTICLE

Transcatheter Aortic-Valve Replacement with a Self-Expanding Prosthesis

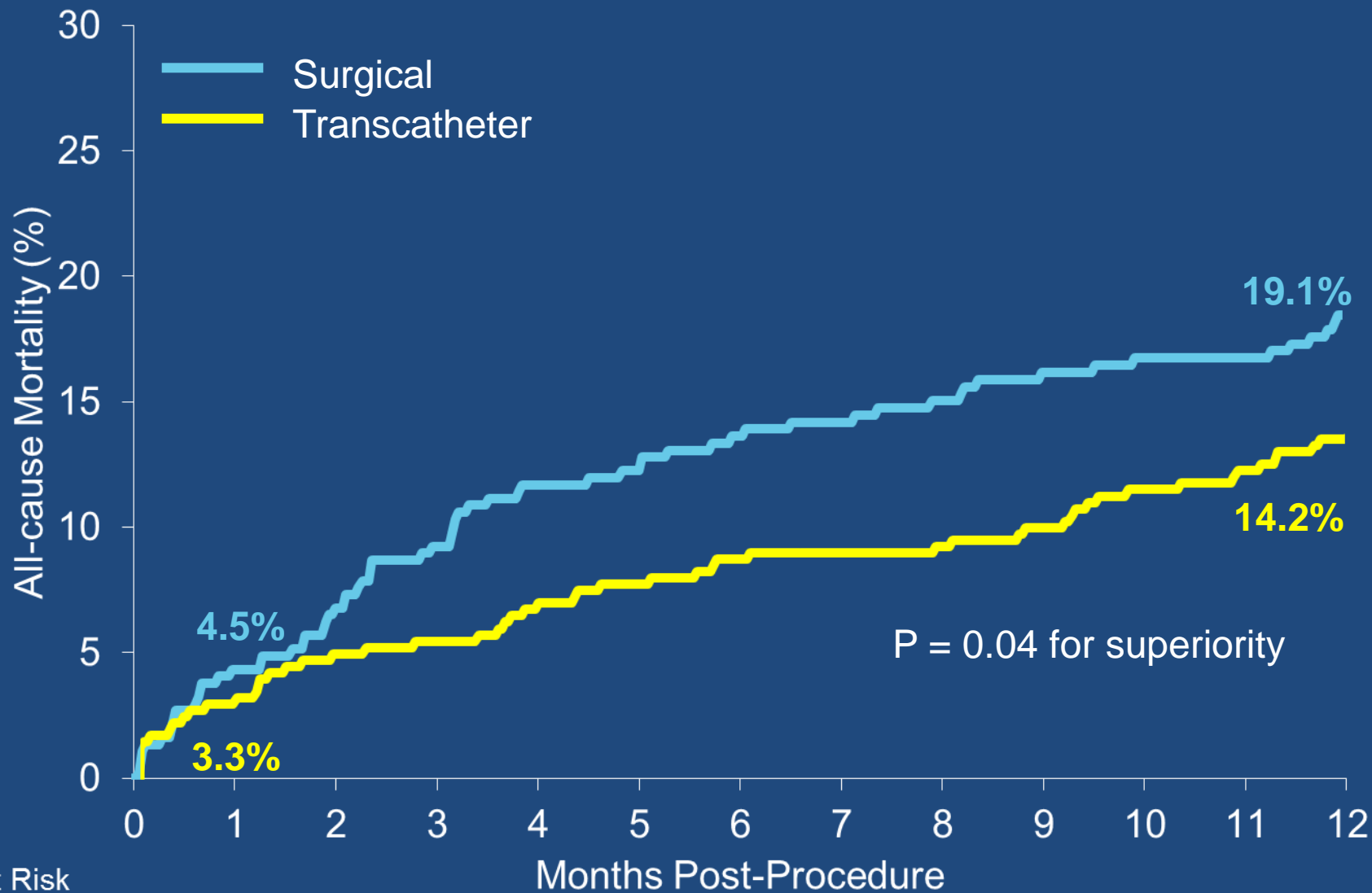
David H. Adams, M.D., Jeffrey J. Popma, M.D., Michael J. Reardon, M.D.,

CONCLUSIONS

In patients with severe aortic stenosis who are at increased surgical risk, TAVR with a self-expanding transcatheter aortic-valve bioprosthesis was associated with a significantly higher rate of survival at 1 year than surgical aortic-valve replacement. (Funded by Medtronic; U.S. CoreValve High Risk Study ClinicalTrials.gov number, NCT01240902.)

Primary Endpoint: 1 Year All-cause Mortality

ACC 2014



No. at Risk

| | | | | |
|---------------|-----|-----|-----|-----|
| Surgical | 357 | 341 | 297 | 274 |
| Transcatheter | 390 | 377 | 353 | 329 |

Redefining Optimal TAVR Outcomes

The CoreValve TAVR System has high long-term survival rates. In a randomized trial comparing CoreValve TAVR to surgical aortic valve replacement (SAVR), CoreValve TAVR significantly outperforms surgical valve replacement at one year.



Performance
That Matters

Learn more at
CoreValve.com

Supplementary Appendix

This appendix has been provided by the authors to give readers additional information about their work.

Supplement to: Adams DH, Popma JJ, Reardon MJ, et al. Transcatheter aortic-valve replacement with a self-expanding prosthesis. *N Engl J Med*. DOI: 10.1056/NEJMoa1400590

Inclusion Criteria

1. Subject must have co-morbidities such that one cardiologist and two cardiac surgeons agree predicted risk of operative mortality is $\geq 15\%$ (and predicted operative mortality or serious, irreversible morbidity risk of $< 50\%$) at 30 days.

Non-cardiovascular death:

- Any death not covered by the above definitions, such as death caused by infection, malignancy, sepsis, pulmonary causes, accident, suicide, or trauma.

Table S9: Clinical Outcomes After 30 Days and 1 Year in the As-Treated Population

| Outcome | 30 Days | | | 1 Year | | |
|---|------------------------------------|-------------------------------|---------|------------------------------------|-------------------------------|---------|
| | Transcatheter Replacement N=390 | Surgical Replacement N=357 | P Value | Transcatheter Replacement N=390 | Surgical Replacement N=357 | P Value |
| Death | | | | | | |
| All-cause – no. (%) | 13 (3.3) | 16 (4.5) | 0.43 | 55 (14.2) | 67 (19.1) | 0.04* |
| Cardiovascular – no. (%) | 12 (3.1) | 16 (4.5) | 0.32 | 40 (10.4) | 44 (12.8) | 0.31 |
| Stroke – no. (%) | 19 (4.9) | 22 (6.2) | 0.46 | 33 (8.8) | 42 (12.6) | 0.10 |
| Major – no. (%) | 15 (3.9) | 11 (3.1) | 0.55 | 22 (5.8) | 23 (7.0) | 0.59 |
| Minor – no. (%) | 4 (1.0) | 12 (3.4) | 0.03 | 11 (3.0) | 20 (6.0) | 0.05 |
| Transient ischemic attack – no. (%) | 3 (0.8) | 1 (0.3) | 0.36 | 6 (1.6) | 5 (1.6) | 0.93 |
| All-cause mortality or major stroke – no. (%) | 23 (5.9) | 24 (6.7) | 0.68 | 63 (16.3) | 79 (22.5) | 0.03 |
| MACCE – no. (%) | 30 (7.7) | 37 (10.4) | 0.22 | 79 (20.4) | 96 (27.3) | 0.03 |
| Myocardial infarction – no. (%) | 3 (0.8) | 3 (0.8) | 0.92 | 7 (1.9) | 5 (1.5) | 0.70 |
| Reintervention – no. (%) | 3 (0.8) | 0 (0.0) | 0.10 | 7 (1.9) | 0 (0.0) | 0.01 |

Supplementary Appendix

This appendix has been provided by the authors to give readers additional information about their work.

Supplement to: Adams DH, Popma JJ, Reardon MJ, et al. Transcatheter aortic-valve replacement with a self-expanding prosthesis. *N Engl J Med*. DOI: 10.1056/NEJMoa1400590

Table S10: Serial Echocardiography by Treatment Group

| | Baseline | | Discharge | | 1 Month | | 1 Year | |
|---|-------------|-------------|------------|---------------|------------|---------------|------------|---------------|
| | TAVR | SAVR | TAVR | SAVR | TAVR | SAVR | TAVR | SAVR |
| Patients – no. | 389 | 353 | 370 | 328 | 365 | 317 | 299 | 228 |
| Mean aortic gradient, mm Hg | 387 | 350 | 356 | 317 | 356 | 311 | 291 | 224 |
| | 48.27±15.31 | 47.65±13.85 | 9.85±4.41 | 13.20±6.20*** | 8.88±3.87 | 11.71±5.71*** | 9.07±3.49 | 12.40±7.38*** |
| Effective orifice area, cm ² | 349 | 306 | 328 | 256 | 344 | 280 | 274 | 206 |
| | 0.72±0.23 | 0.73±0.24 | 1.94±0.58 | 1.58±0.50*** | 1.95±0.56 | 1.60±0.51*** | 1.91±0.51 | 1.57±0.49*** |
| Total aortic regurgitation, N | 385 | 346 | 363 | 306 | 359 | 308 | 297 | 223 |
| None – no. (%) | 58 (15.1) | 51 (14.7) | 63 (17.4) | 197 (64.4) | 44 (12.3) | 201 (65.3) | 85 (28.6) | 152 (68.2) |
| Trace – no. (%) | 146 (37.9) | 115 (33.2) | 134 (36.9) | 81 (26.5) | 148 (41.2) | 70 (22.7) | 105 (35.4) | 48 (21.5) |
| Mild – no. (%) | 161 (41.8) | 159 (46.0) | 133 (36.6) | 25 (8.2) | 131 (36.5) | 33 (10.7) | 86 (29.0) | 20 (9.0) |
| Moderate – no. (%) | 20 (5.2) | 19 (5.5) | 28 (7.7) | 1 (0.3) | 29 (8.1) | 4 (1.3) | 20 (6.7) | 2 (0.9) |
| Severe – no. (%) | 0 (0.0) | 2 (0.6) | 5 (1.4) | 2 (0.7) | 7 (1.9) | 0 (0.0) | 1 (0.3) | 1 (0.4) |
| Moderate or Severe – no. (%) | 20 (5.2) | 21 (6.1) | 33 (9.1) | 3 (1.0)*** | 36 (10.0) | 4 (1.3)*** | 21 (7.0) | 3 (1.3)** |
| Paravalvular regurgitation, N | | | 359 | 302 | 356 | 307 | 295 | 221 |
| None – no. (%) | NA | NA | 85 (23.7) | 255 (84.4) | 60 (16.9) | 263 (85.7) | 113 (38.3) | 190 (86.0) |
| Trace – no. (%) | NA | NA | 123 (34.3) | 37 (12.3) | 137 (38.5) | 31 (10.1) | 88 (29.8) | 20 (9.0) |
| Mild – no. (%) | NA | NA | 123 (34.3) | 9 (3.0) | 127 (35.7) | 10 (3.3) | 76 (25.8) | 10 (4.5) |
| Moderate – no. (%) | NA | NA | 24 (6.7) | 1 (0.3) | 26 (7.3) | 3 (1.0) | 17 (5.8) | 1 (0.5) |
| Severe – no. (%) | NA | NA | 4 (1.1) | 0 (0.0) | 6 (1.7) | 0 (0.0) | 1 (0.3) | 0 (0.0) |
| Moderate or severe – no. (%) | NA | NA | 28 (7.8) | 1 (0.3)*** | 32 (9.0) | 3 (1.0)*** | 18 (6.1) | 1 (0.5)*** |

TAVR = transcatheter group; SAVR = surgical group; *P<0.05; **P<0.01; ***P<0.001.

Conclusion

- We assessed the safety and effectiveness of TAVR with the CoreValve prosthesis compared to surgical valve replacement in symptomatic patients with severe aortic stenosis at increased surgical risk
- The rate of death from any cause at 1 year was significantly reduced with TAVR performed with the CoreValve prosthesis

Adams D, ACC 2014

Non-cardiovascular death:

- Any death not covered by the above definitions, such as death caused by infection, malignancy, sepsis, pulmonary causes, accident, suicide, or trauma.

Clinical Outcomes After Transcatheter Aortic Valve Replacement Using Valve Academic Research Consortium Definitions

A Weighted Meta-Analysis of 3,519 Patients From 16 Studies

Philippe Généreux, MD,*† Stuart J. Head, MSc,‡ Nicolas M. Van Mieghem, MD,§

30-day event rates after TAVR

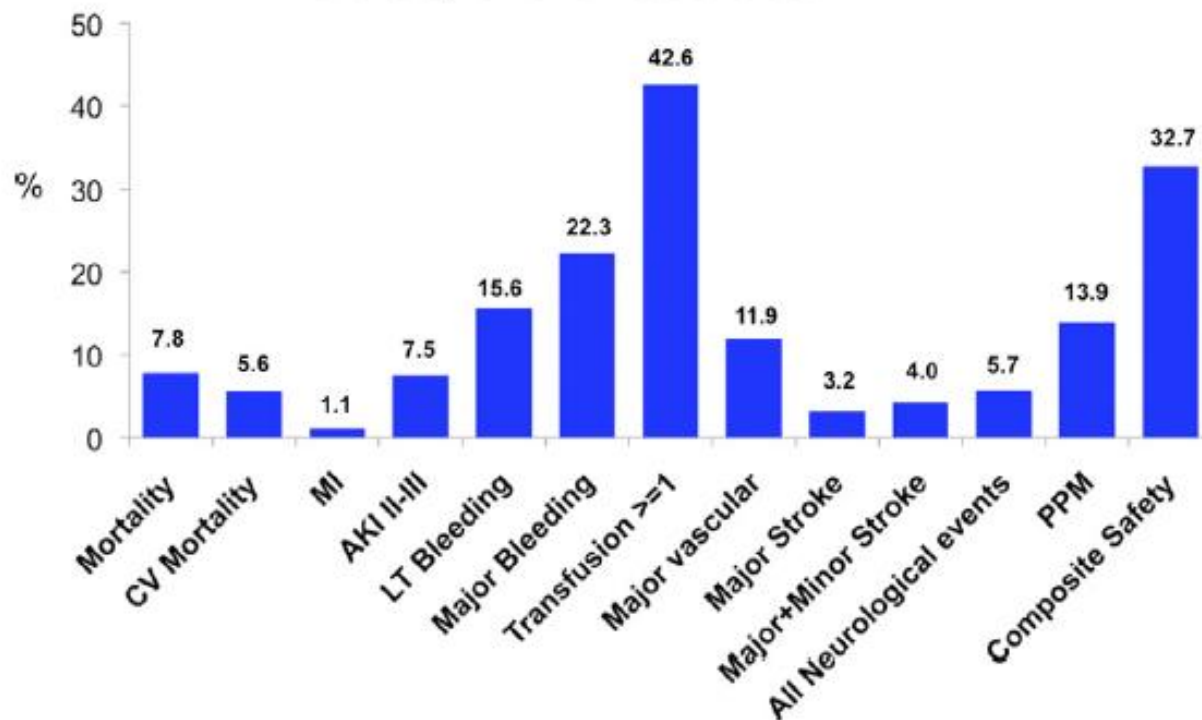


Figure 2 30-Day Event Rates of Major VARC-Related Outcomes

AKI = acute kidney injury; CV = cardiovascular; LT = life-threatening; MI = myocardial infarction ≤ 72 h after procedure; PPM = permanent pacemaker; VARC = Valve Academic Research Consortium.

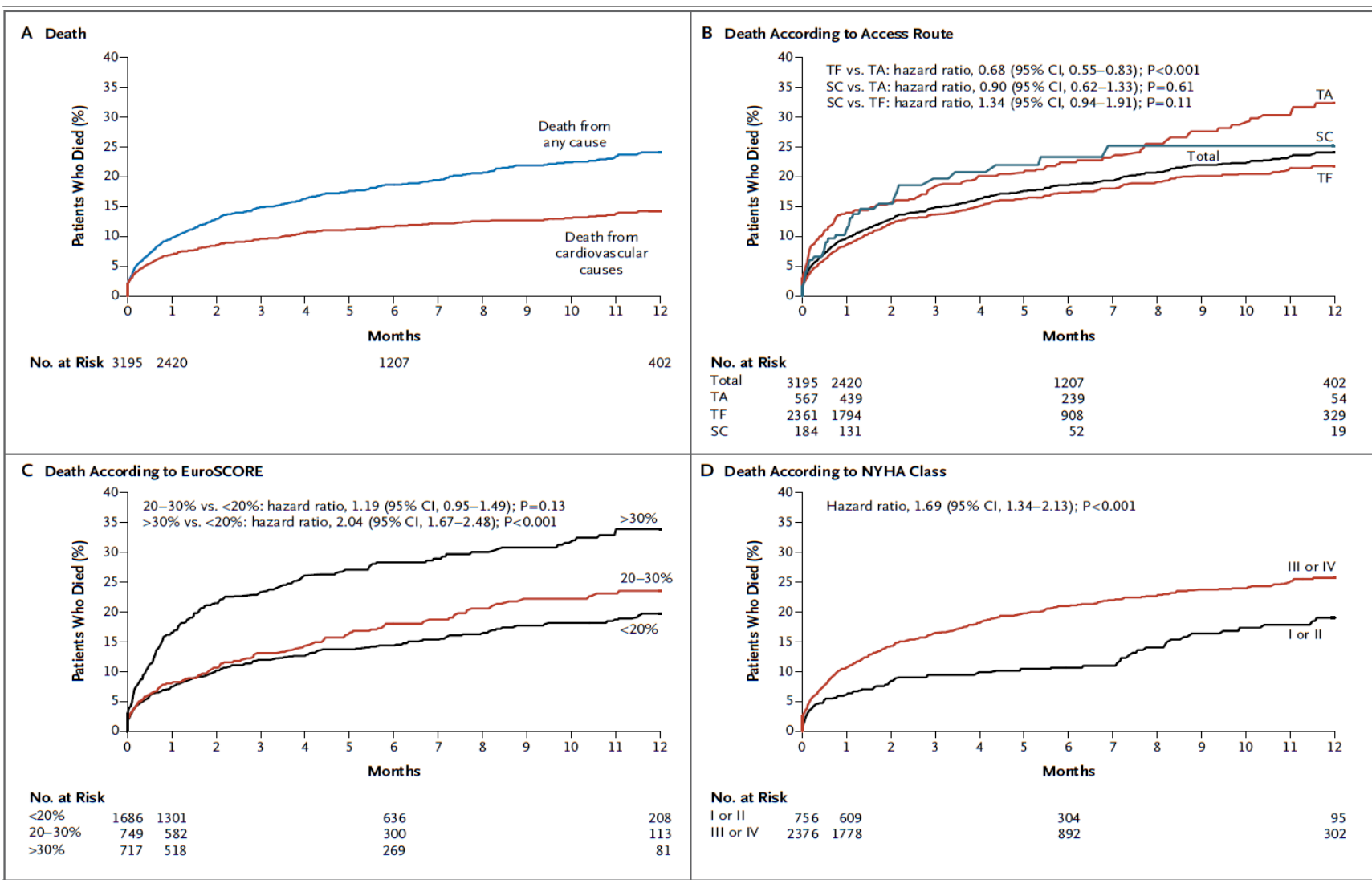
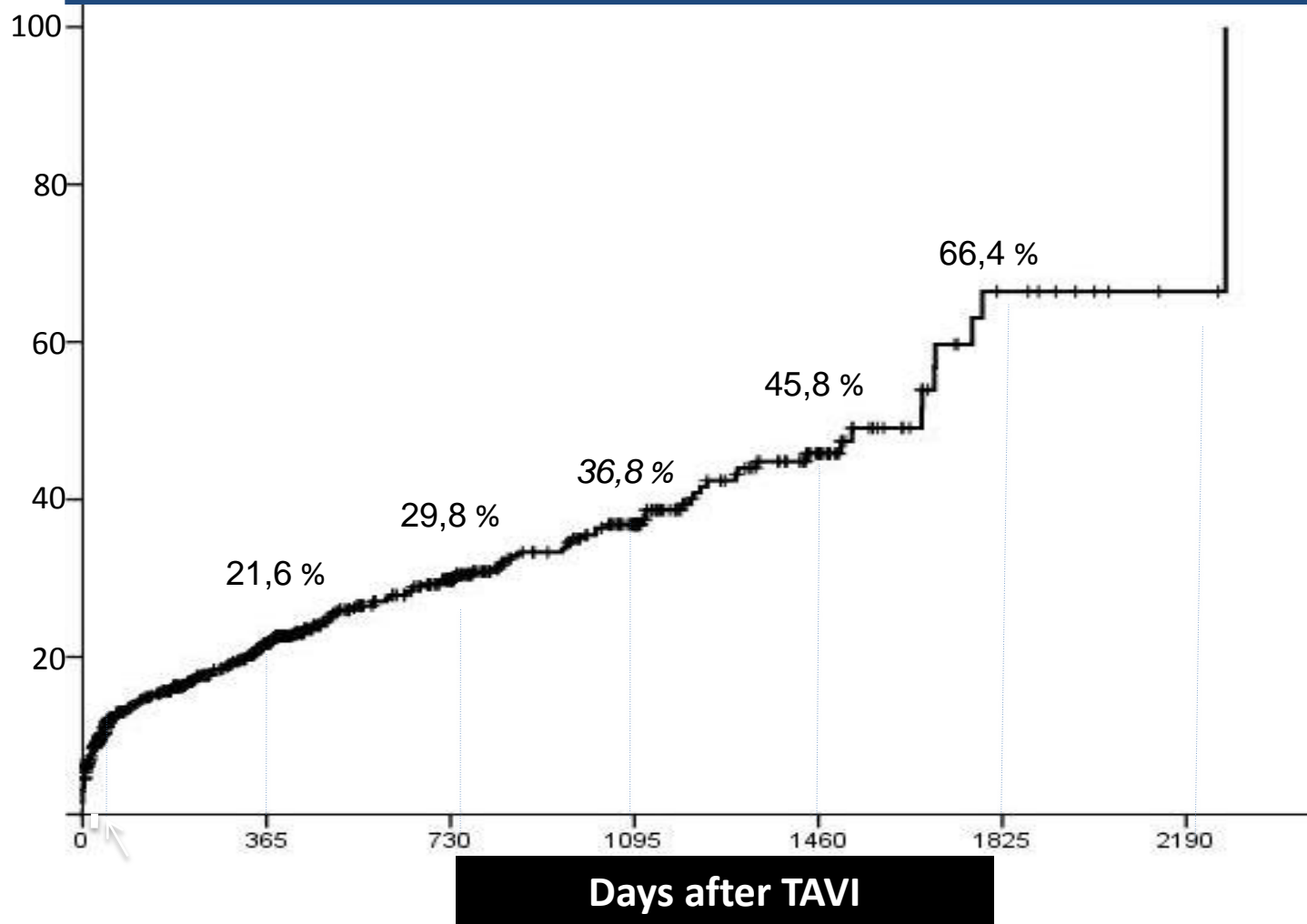


Figure 1. Time-to-Event Curves for the Primary End Point and Other Selected End Points.
 Panel A shows the rate of death from any cause (the primary end point) and from cardiovascular causes among patients in the French national transcatheter aortic-valve implantation (TAVI) registry, FRANCE 2. Panel B shows rates of death from any cause according to the TAVI access route: transapical (TA), transfemoral (TF), or subclavian (SC). Panel C shows rates of death from any cause according to the logistic EuroSCORE (with a score of >20% indicating very high surgical risk). Panel D shows rates of death from any cause according to New York Heart Association (NYHA) class. Event rates were calculated with the use of Kaplan–Meier methods and were compared by means of the log-rank test. Deaths from unknown causes were assumed to be from cardiovascular causes.

Brazilian TAVI Registry All-Cause Mortality



N. at risk
total: 819

713

428

228

123

47

9

2

Procedure Failure / Complications

| | (n = 819) |
|---|-----------------|
| Device Failure, n (%) | 185 (22.6%) |
| PVR Moderate / Severe, n (%) | 55 / 729 (7.5%) |
| Mean Gradient \geq 20 mmHg, n (%) | 30 / 587 (5.1%) |
| 2nd Prosthesis, n (%) | 41 (5.0%) |
| Surgical Conversion, n (%) | 14 (1.7%) |
| Device Malpositioning, n (%) | 43 (5.3%) |
| Device Emboization, n (%) | 29 (3.5%) |
| Coronary Occlusion, n (%) | 3 (0.4%) |
| Mitral Damage / Dysfunction, n (%) | 2 (0.2%) |
| Annulus Rupture / Ventricular Septal Perforation, n (%) | 2 (0.2%) |
| Tamponade, n (%) | 29 (3.5%) |
| LV Perforation, n (%) | 15 (1.8%) |



The German Aortic Valve Registry: 1-year results from 13 680 patients with aortic valve disease[†]

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Hans Reiner Figulla^e, Jochen Cremer^f, Karl-Heinz Kuck^g, Rüdiger Lange^h, Ralf Zahnⁱ, Stefan Sack^j,
Gerhard Schuler^k, Thomas Walther^k, Friedhelm Beyersdorf^l, Michael Böhm^m, Gerd Heuschⁿ,
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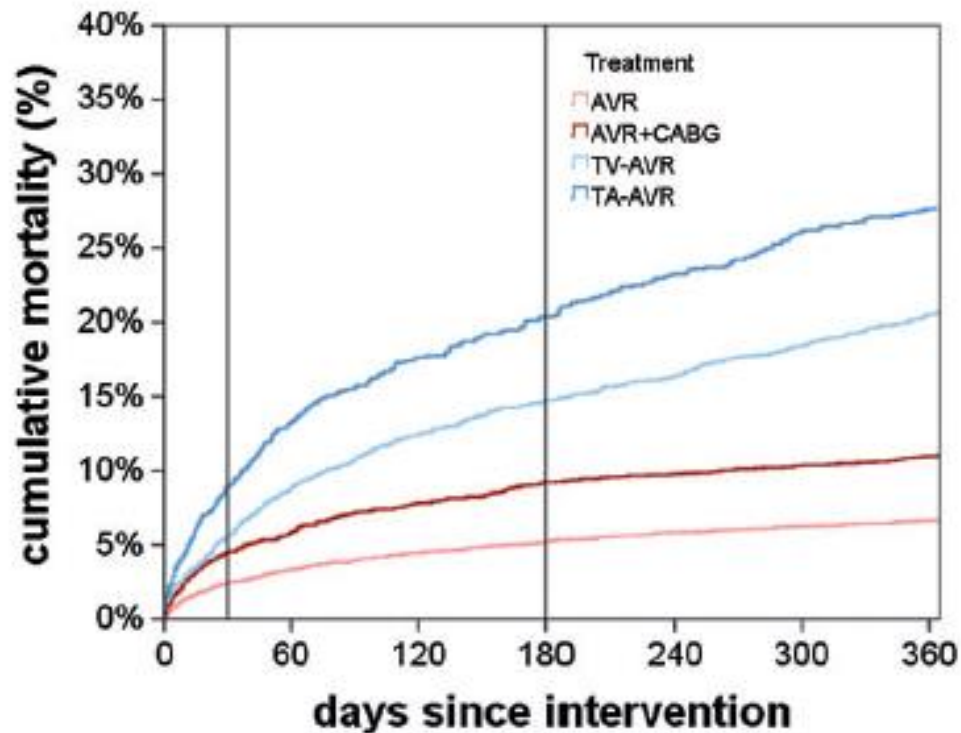
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Received 16 April 2014; received in revised form 27 May 2014; accepted 17 June 2014

N= 13860

AVR= 6523, AVR+CABG=3462, TF-TAVI=2694, TA-TAVI= 1181



| # at Risk – day | 0 | 30 | 180 | 365 |
|-----------------|------|------|------|------|
| AVR | 6523 | 6346 | 6069 | 5962 |
| AVR+CABG | 3462 | 3293 | 3079 | 3016 |
| TV-AVR | 2694 | 2533 | 2235 | 2073 |
| TA-AVR | 1181 | 1065 | 912 | 822 |

Figure 1: Overall death rates within the first year. Pairwise tests: for multiple comparison to correct by Bonferroni-Holm-Shaffer (6-3-3-3-2-1 rule). AVR: aortic valve replacement; CABG: coronary artery bypass grafting; TA: transapical; TV: transvascular.

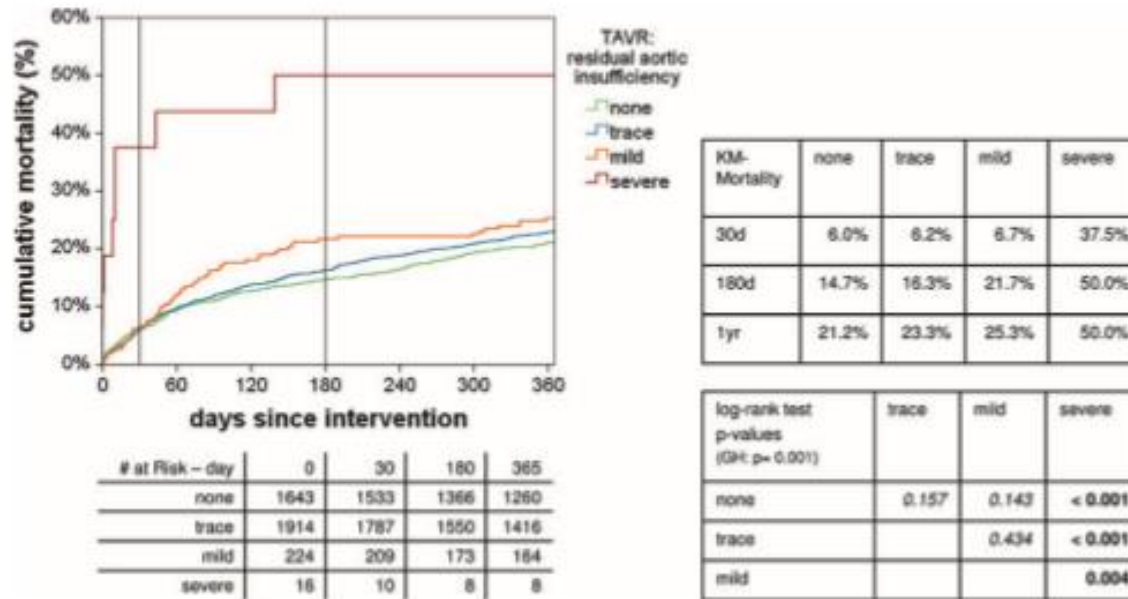


Figure 4: Influence of residual aortic regurgitation on survival. KM: Kaplan-Meier; GH: global hypothesis; TAVR: transcatheter AVR.

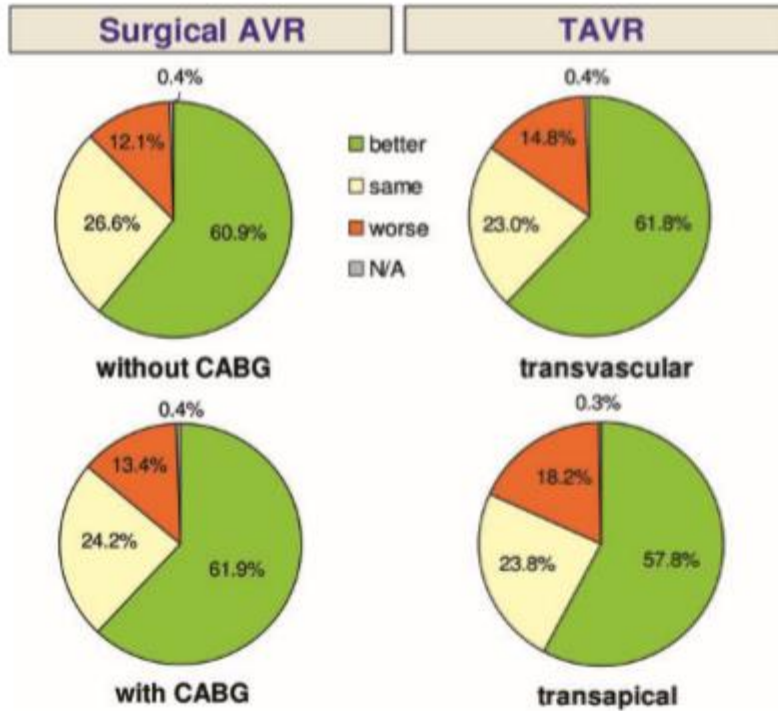


Figure 5: One-year follow-up: subjective rating of general health condition when compared with condition prior to the intervention. KM: Kaplan-Meier; GH: global hypothesis; TAVR: transcatheter AVR.

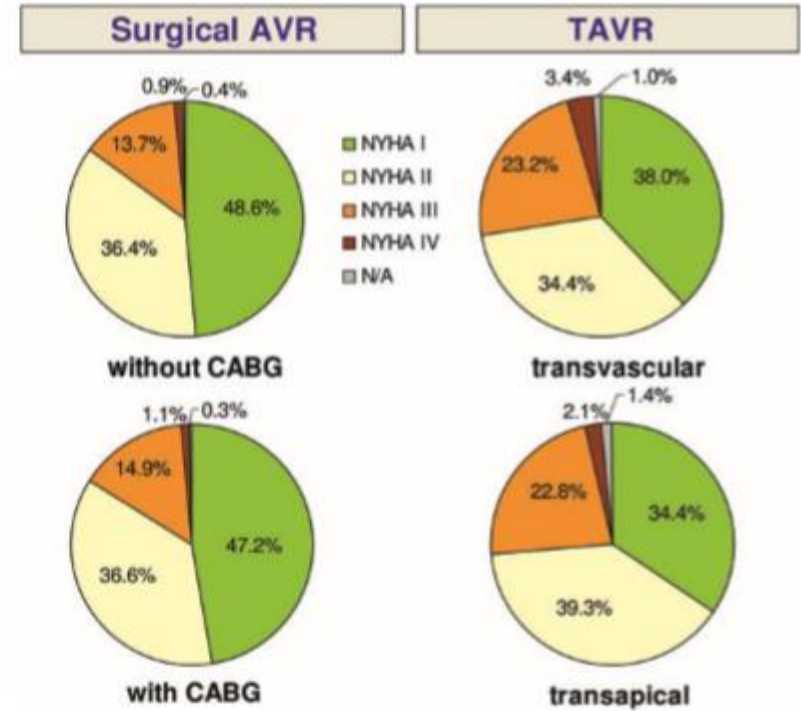


Figure 7: Heart failure symptom rating (NYHA) at 1 year post-intervention.

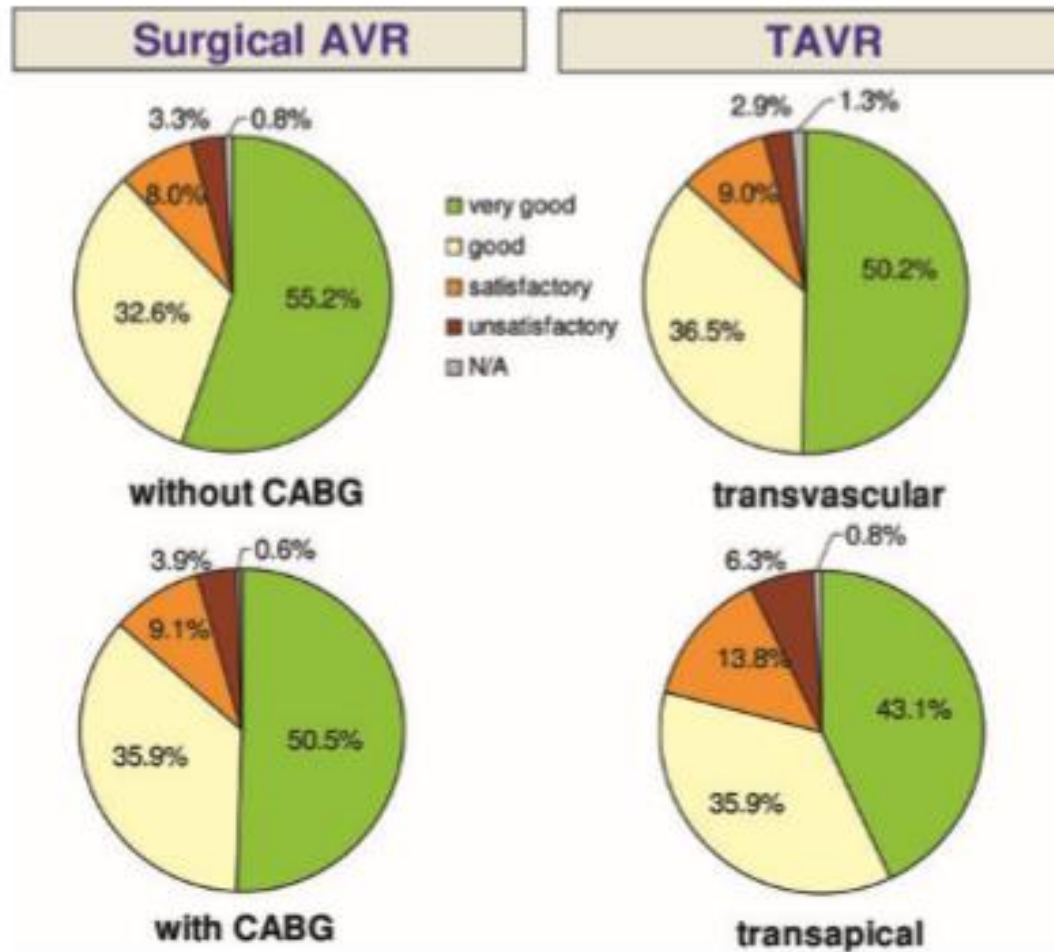


Figure 6: Satisfaction with outcome 1 year after the procedure.

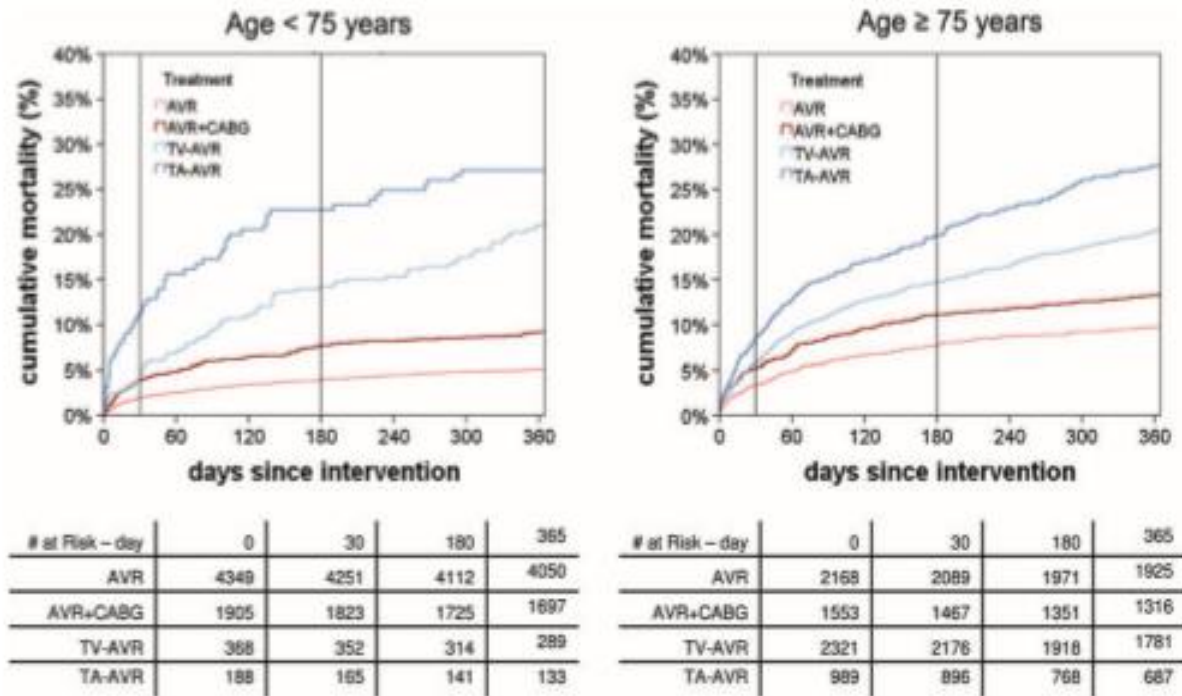


Figure 2: Time-to-event curves for death stratified by age. AVR: aortic valve replacement; CABG: coronary artery bypass grafting; TA: transapical; TV: transvascular.

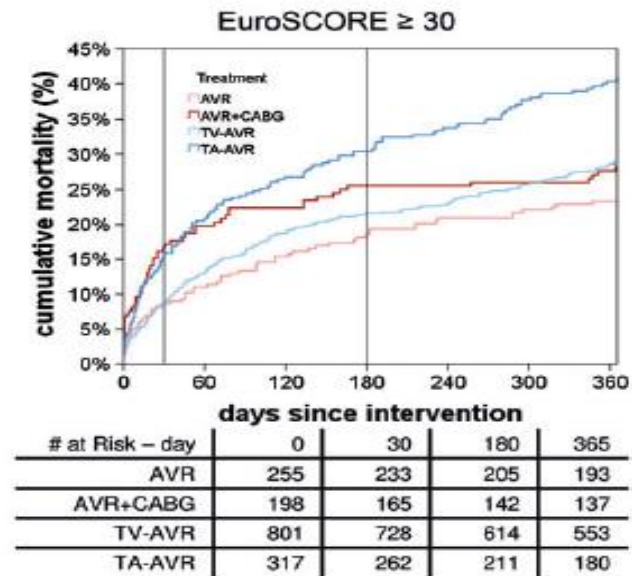
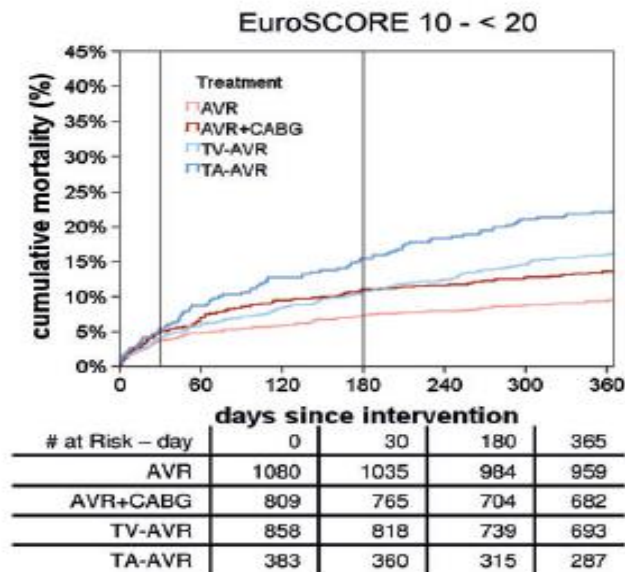
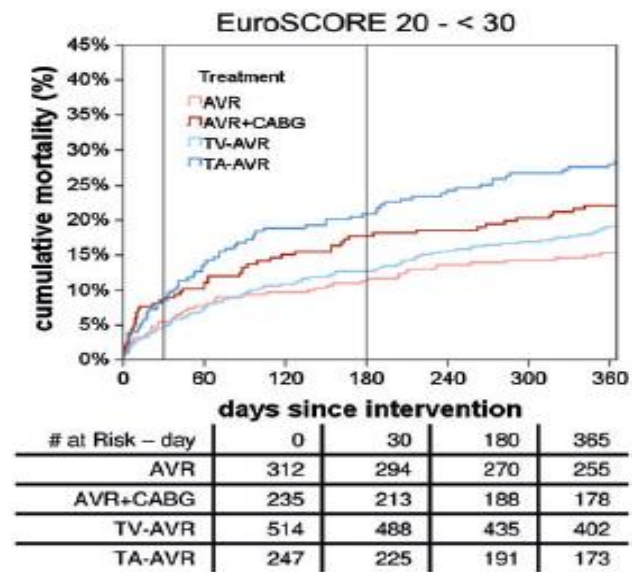
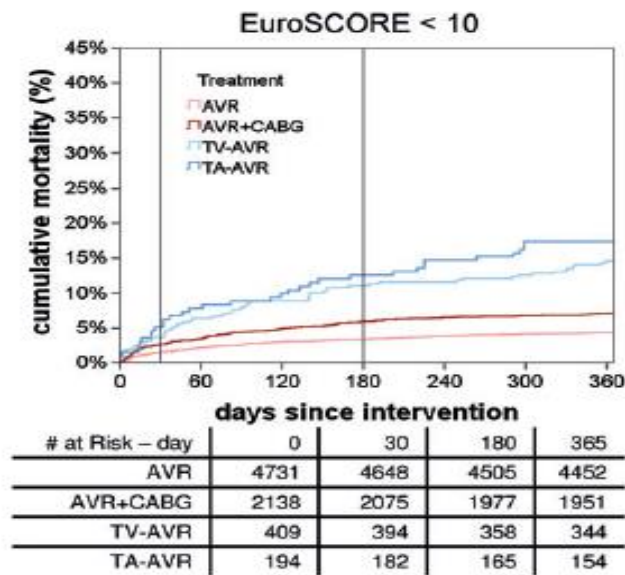


Figure 3: Time-to-event curves for death stratified by the logistic EuroSCORE. AVR: aortic valve replacement; CABG: coronary artery bypass grafting; TA: transapical; TV: transvascular.

TAVI a longo prazo



Centre for
Heart Valve Innovation
St. Paul's Hospital, Vancouver

2016 | euro
PCR

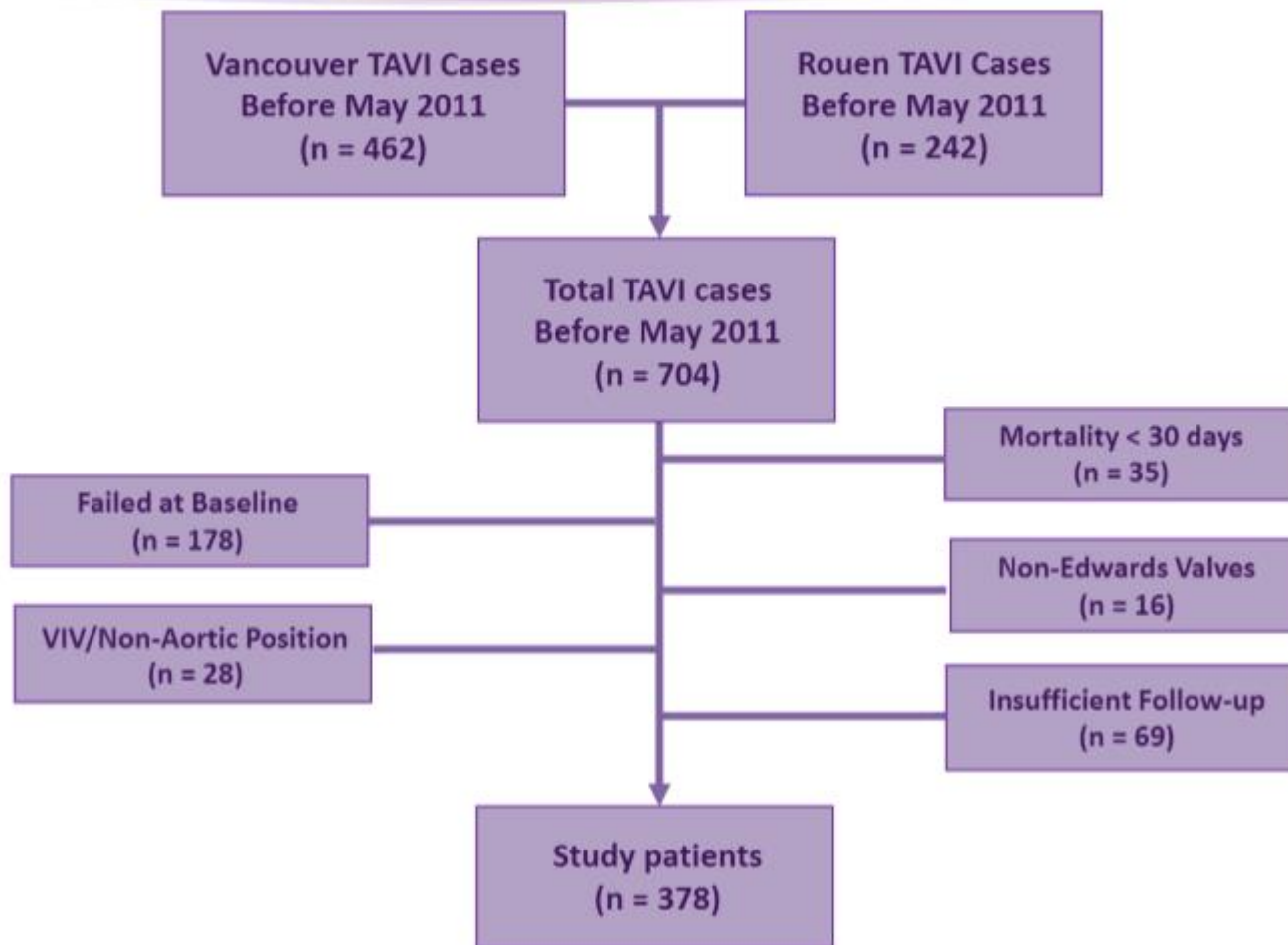
First look at long-term durability of transcatheter heart valves: Assessment of valve function up to 10-years after implantation

Danny Dvir, St. Paul's Hospital, Vancouver, Canada.

On behalf of coauthors: Helene Eltchaninoff, Jian Ye, Arohumam Kan, Eric Durand, Anna Bizios, Anson Cheung, Mina Aziz, Matheus Simonato, Christophe Tron, Yaron Arbel, Robert Moss, Jonathon Leipsic, Hadas Ofek, Gidon Perlman, Marco Barbanti, Michael A. Seidman, Philippe Blanke, Robert Yao, Robert Boone, Sandra Lauck, Sam Lichtenstein, David Wood, Alain Cribier, John Webb



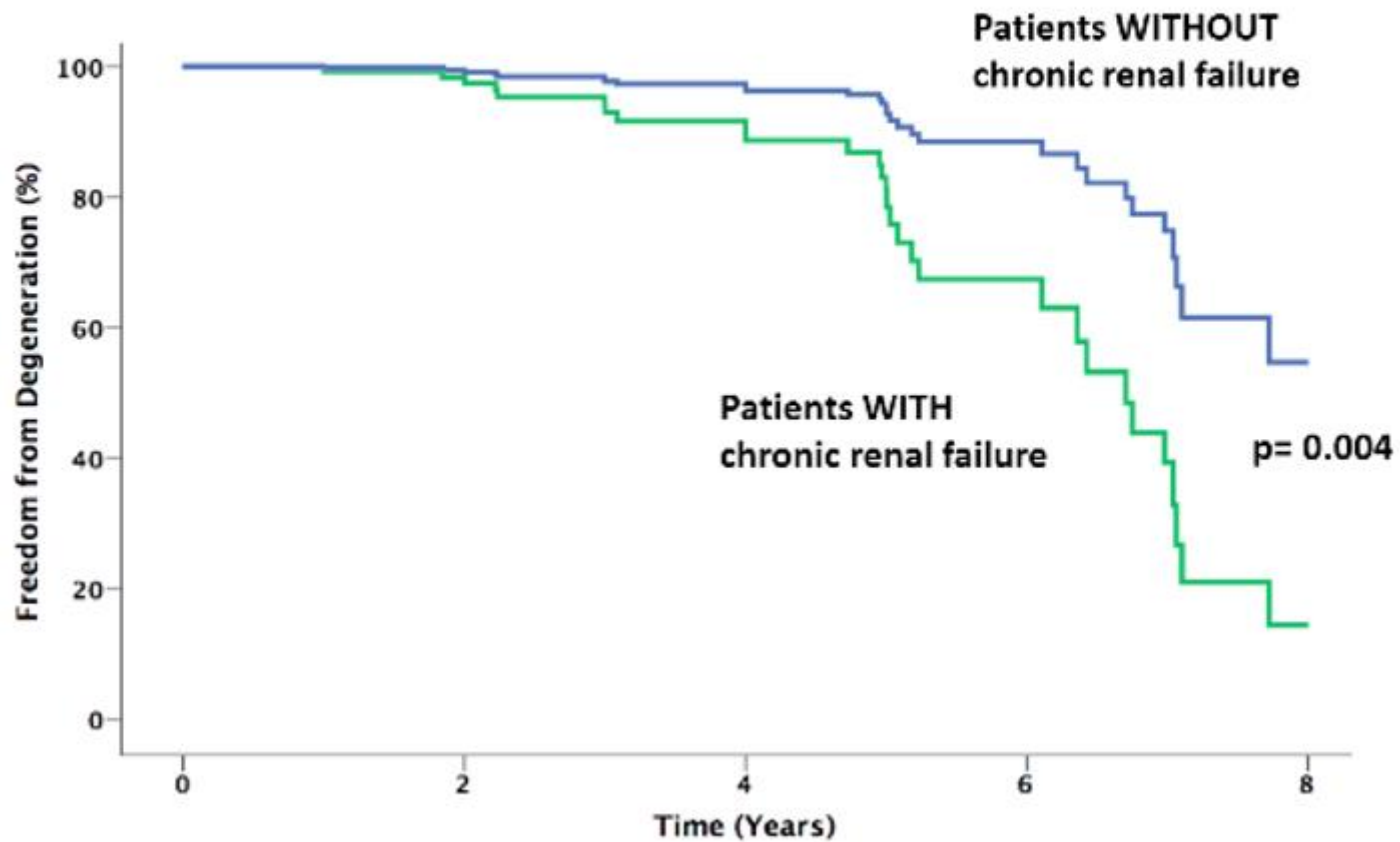
Methods



2016

euro
PCR

Freedom from THV degeneration

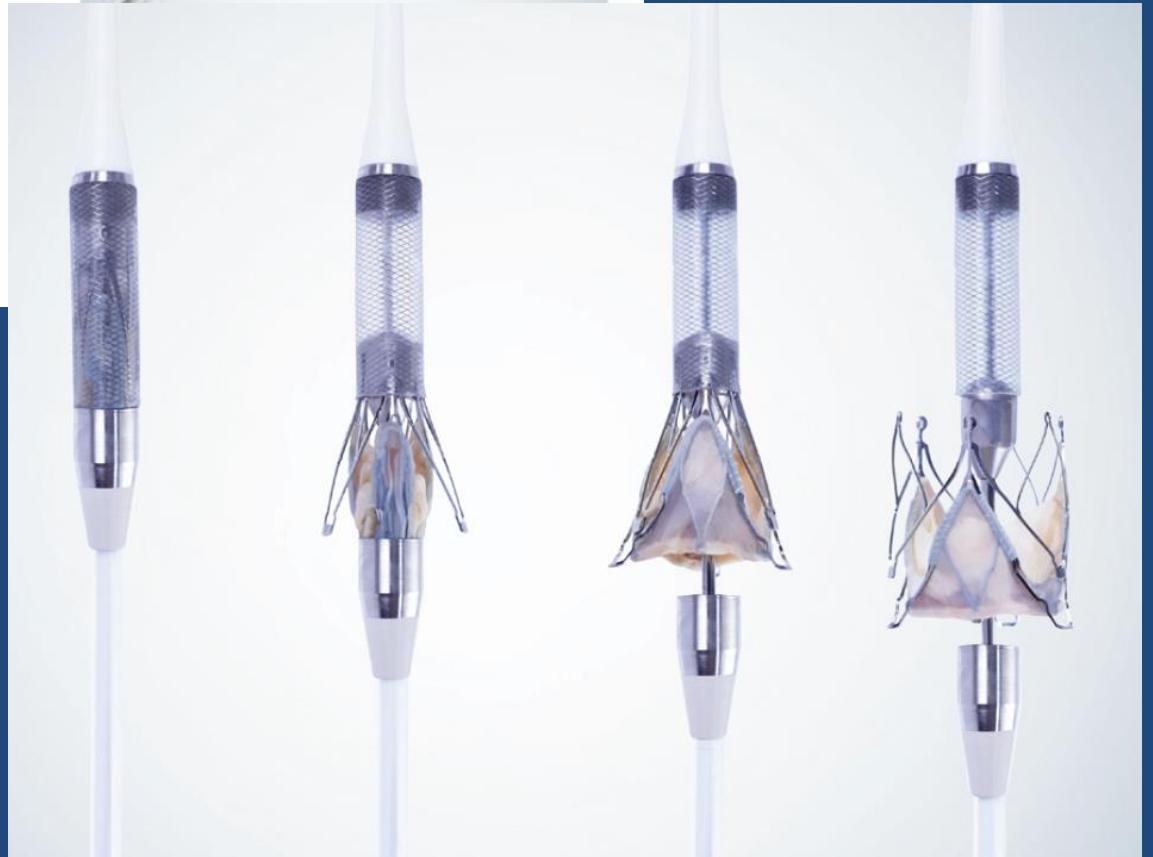


Summary

- The current analysis includes a first look at long-term durability after TAVI, evaluating cases performed 5-14 years ago with early-generation balloon-expandable THV devices.
- In this preliminary report, a significant increase in degeneration rate was observed between 5-7 years after TAVI.
- Estimate of THV degeneration (resulting in at least moderate stenosis AND/OR regurgitation) was ~50% within 8 years.
- Renal failure was the strongest correlate of THV degeneration.



Die JenaValve



A escolha do procedimento

Estenose Aórtica

Escolha do tipo de Intervenção

I

B-NR

See Online Data Supplements 5 and 9
(Updated From 2014 VHD
Guideline)

Surgical AR is recommended for symptomatic patients with severe AS (Stage D) and asymptomatic patients with severe AS (Stage C) who meet an indication for AVR when surgical risk is low or intermediate (42,43).

MODIFIED: LOE updated from A to B-NR. Prior recommendations for intervention choice did not specify patient symptoms. The patient population recommended for surgical AVR encompasses both symptomatic and asymptomatic patients who meet an indication for AVR with low-to-intermediate surgical risk. This is opposed to the patient population recommended for TAVR, in whom symptoms are required to be present. Thus, all recommendations for type of intervention now specify the symptomatic status of the patient.

Estágios da EAo

A – em risco de doença valvar

B – EAo assintomática progressiva

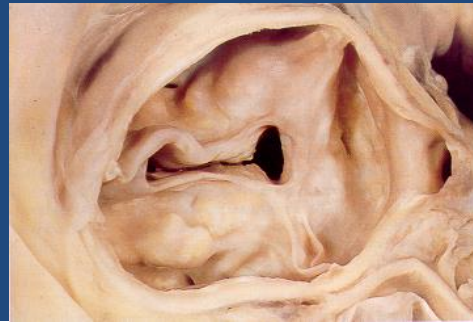
C – EAo assintomática, grave

D – EAo sintomática, grave



Estenose Aórtica

Escolha do tipo de Intervenção



See Online Data Supplement 9
(Updated From 2014 VHD
Guideline)

Surgical AVR or TAVR is recommended for symptomatic patients with severe AS (Stage D) and high risk for surgical AVR, depending on patient-specific procedural risks, values, and preferences (49-51).

MODIFIED: COR updated from IIa to I, LOE updated from B to A. Longer-term follow-up and additional RCTs have demonstrated that TAVR is equivalent to surgical AVR for severe symptomatic AS when surgical risk is high.



See Online Data Supplements 5 and 9
(Updated From 2014 VHD
Guideline)

TAVR is recommended for symptomatic patients with severe AS (Stage D) and a prohibitive risk for surgical AVR who have a predicted post-TAVR survival greater than 12 months (58-61).

MODIFIED: LOE updated from B to A. Longer-term follow-up from RCTs and additional observational studies has demonstrated the benefit of TAVR in patients with a prohibitive surgical risk.



See Online Data Supplements 5 and 9
(Updated From 2014 VHD
Guideline)

TAVR is a reasonable alternative to surgical AVR for symptomatic patients with severe AS (Stage D) and an intermediate surgical risk, depending on patient-specific procedural risks, values, and preferences (62-65).

NEW: New RCT showed noninferiority of TAVR to surgical AVR in symptomatic patients with severe AS at intermediate surgical risk.

Obs limited follow-up

Estenose Aórtica

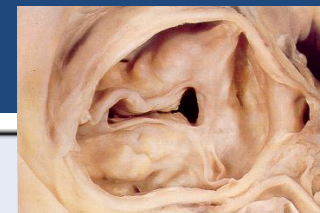
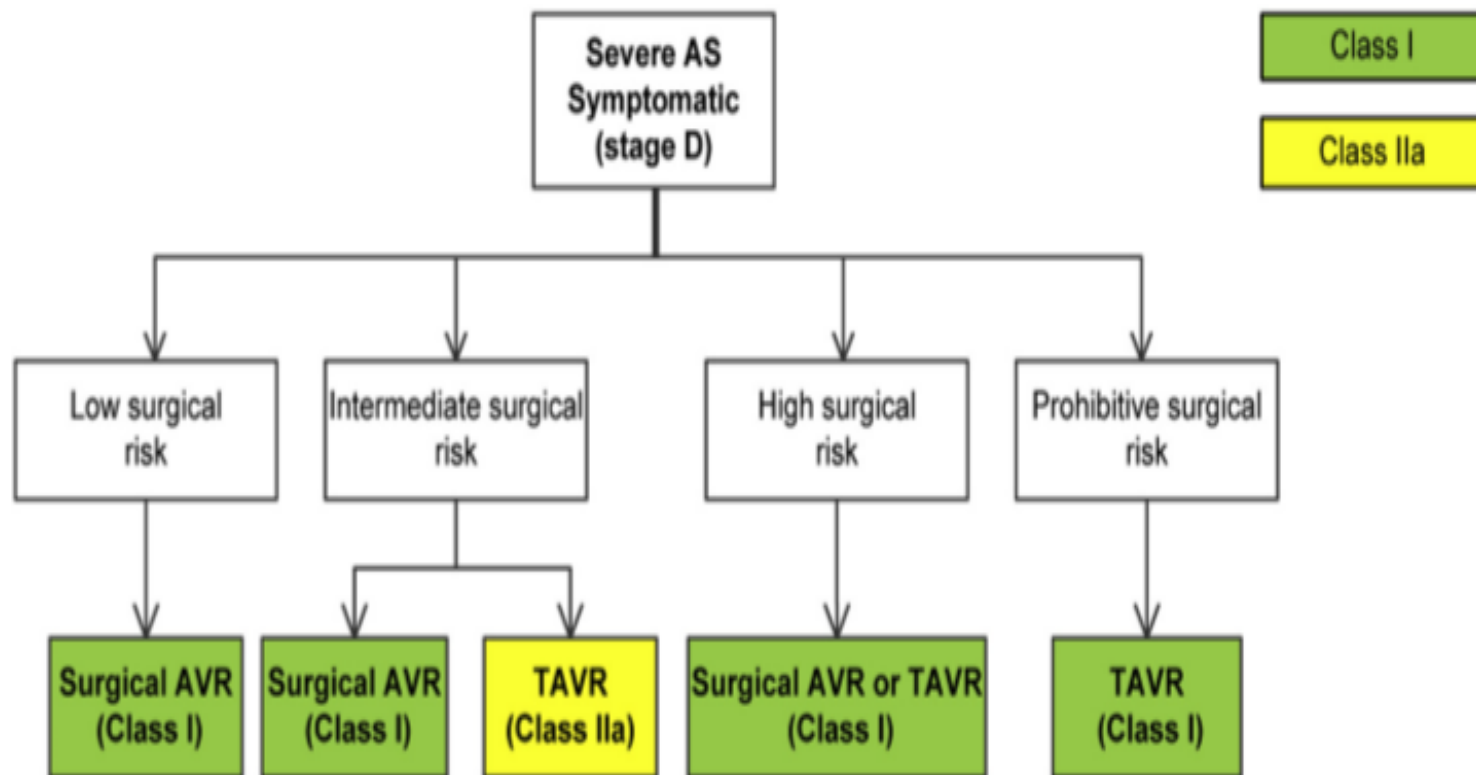


FIGURE 1 Choice of TAVR Versus Surgical AVR in the Patient With Severe Symptomatic AS



AS indicates aortic stenosis; AVR, aortic valve replacement; and TAVR, transcatheter aortic valve replacement.

Limitações

TAVI

Tamanho do anel valvar

Anatomia: bicúspide, angulações

Lesões associadas: valva mitral, ectasia e an.aorta

Acessos vasculares comprometidos

Tecido biológico: <60 anos, não

Mortalidade não desprezível

Morbidade idem

Longo prazo desconhecido

Não é “só um furinho na perna”

CIRURGIA CONVENCIONAL

Porte do procedimento

Recuperação imediata

Dor

Aceitação pelo paciente

Infecção

Mais transfusões,

insuficiência renal

insuficiência respiratória

fibrilação atrial

Maiores tempos de

procedimento, recuperação,
hospitalização

Conclusões

- Cirurgia convencional persiste como padrão-ouro por algum tempo: *indicada em todas as situações*
- Valvoplastias em IAo e EAo poderão ter incremento
- Biopróteses serão preferidas sobre próteses mecânicas
- TAVI terá incremento, dependente de avanço tecnológico, na EAo e *valve-in-valve*. *Em IAo?*
- Biopróteses *Sem sutura/Implante rápido* deverão buscar seu lugar
- Critérios de indicação e escolha do procedimento deverão seguir avaliações de resultados objetivas e isentas
- Custos deverão sempre ser considerados



72° CONGRESSO
BRASILEIRO DE
CARDIOLOGIA

03 a 05

Novembro | 2017

SÃO PAULO EXPO
EXHIBITION & CONVENTION CENTER



Mesa-Redonda

Avaliação do Paciente com Estenose Aórtica de Moderado e Alto Risco Cirúrgico

Visão do Cirurgião Cardíaco

Renato A. K. Kalil

Cirurgião Cardiovascular

Professor-Titular de Clínica Cirúrgica da UFCSPA

Professor Emérito do Programa de Pós-Graduação do IC/FUC

Membro Titular da Academia Sul-Rio-Grandense de Medicina

Pesquisador CNPq

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