

CARDIO RUN 2023



TAVI... INTRODUCTION...
RESULTATS des 200 Iers TAVI CSC



27 septembre 2023

Dr C. POUILLOT

TAVI

Trans-catheter Aortique Valve Implantation

Le rétrécissement aortique du sujet âgé à la Réunion

RAo valvulopathie la plus fréquente du sujet âgé : prévalence 2.8 à 4.6 %
> 75 ans (USA).

Longue latence (asymptomatique) puis progression rapide après 1ers
symptômes

(50 % patients décédés à 2 ans, 80 % à 3 ans)

France : 67,8 M habitants (INSEE 2021)

6 291 749 > 75 ans (9,2 %)

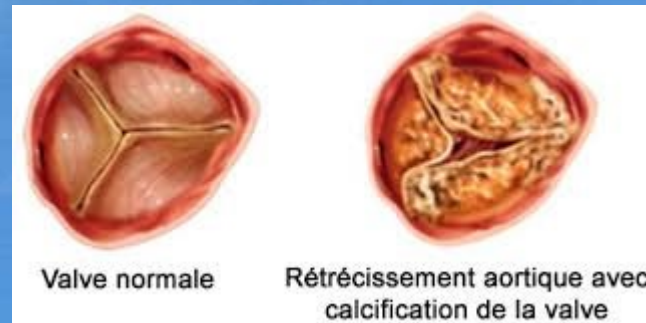
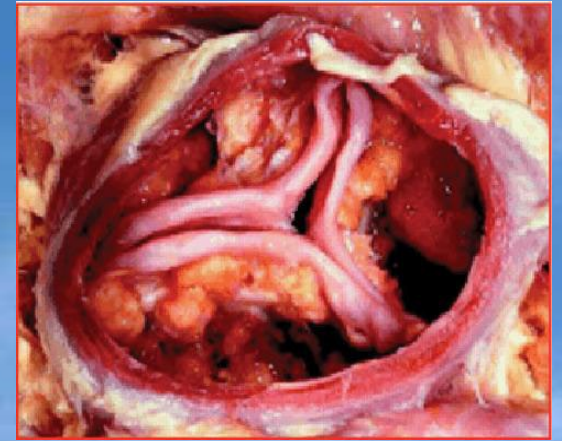
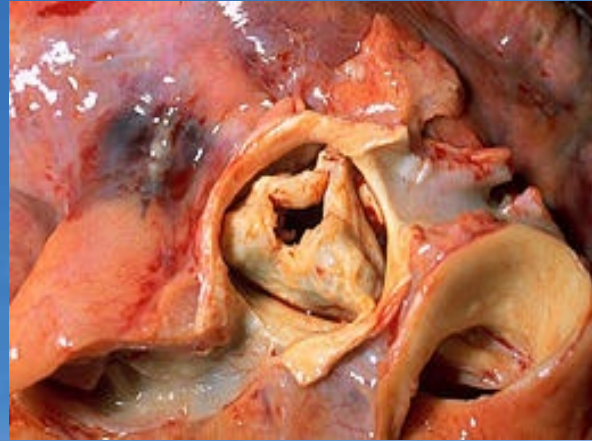
environ 230 000 patient RAo

Réunion : 858 450 habitants (INSEE 2021)

40 355 > 75ans (4,7%) 2018 VS 25 594 (3,2 %) en 2008 :↑ Rapide

environ 1500 patients RAo

Anatomo pathologie



TAVI : Bioprothèse aortique sur structure métallique (stent) implantée à l'intérieur de la valve aortique native



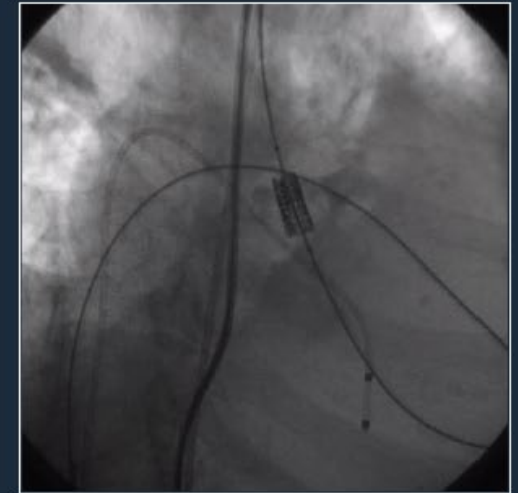
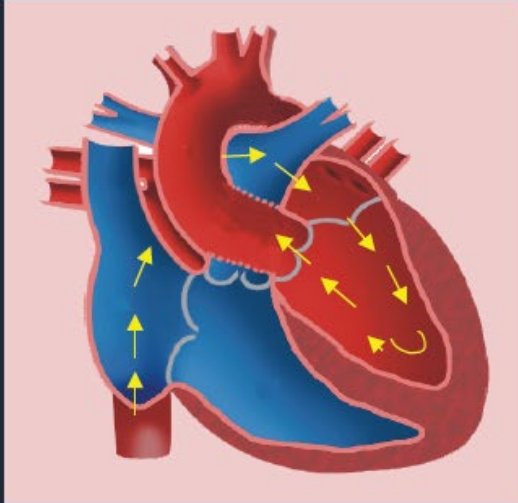
Bioprothèse aortique
Péricarde bovin
Stent cobalt chrome ballon expandable
Sapien 3 Edwards



Bioprothèse aortique
Péricarde porcin
Stent Nitinol self expandable
Core Valve evolute R Medtronic

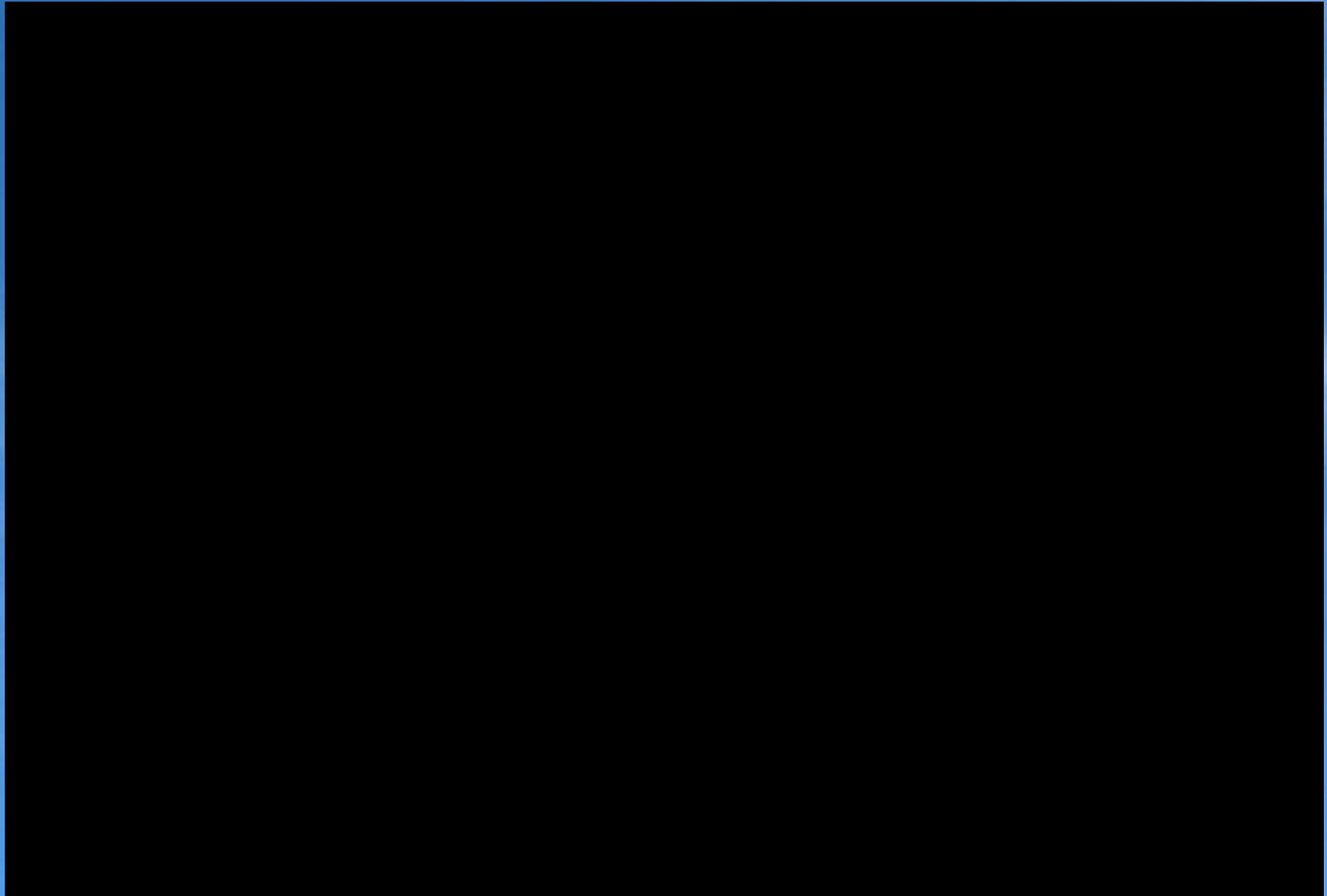
2002, Rouen, F.I.M –TAVI

Du rêve à la réalité



Marguerita H. 78 ans, OAP, RAo serré, V max 4.6 , gradient moyen 55 mm Hg, SAo 0.65 cm²

LES 22.6 %, STS 7.1 % IRC sévère C 9 ml/mn



TAVI CSC 25 octobre 2016 (30 cc Ultravist): bon résultat immédiat

**ESC**European Society
of Cardiology

European Heart Journal (2021) 00, 1–72

doi:10.1093/eurheartj/ehab395

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

Developed by the Task Force for the management of valvular heart disease of the European Society of Cardiology (ESC) and the European Association for Cardio-Thoracic Surgery (EACTS)

Authors/Task Force Members: Alec Vahanian * (ESC Chairperson) (France), Friedhelm Beyersdorf*¹ (EACTS Chairperson) (Germany), Fabien Praz (ESC Task Force Coordinator) (Switzerland), Milan Milojevic¹ (EACTS Task Force Coordinator) (Serbia), Stephan Baldus (Germany), Johann Bauersachs (Germany), Davide Capodanno (Italy), Lenard Conradi¹ (Germany), Michele De Bonis¹ (Italy), Ruggero De Paulis¹ (Italy), Victoria Delgado (Netherlands), Nick Freemantle¹ (United Kingdom), Martine Gilard (France), Kristina H. Haugaa (Norway), Anders Jeppsson¹ (Sweden), Peter Jüni (Canada), Luc Pierard (Belgium), Bernard D. Prendergast (United Kingdom), J. Rafael Sádaba¹ (Spain), Christophe Tribouilloy (France), Wojtek Wojakowski (Poland), ESC/EACTS Scientific Document Group

A) Symptomatic aortic stenosis	Class ^b	Level ^c
Intervention is recommended in symptomatic patients with severe, high-gradient aortic stenosis [mean gradient ≥ 40 mmHg, peak velocity ≥ 4.0 m/s, and valve area ≤ 1.0 cm ² (or ≤ 0.6 cm ² /m ²)]. ^{235,236}	I	B
Intervention is recommended in symptomatic patients with severe low-flow (SVi ≤ 35 mL/m ²), low-gradient (< 40 mmHg) aortic stenosis with reduced ejection fraction ($< 50\%$), and evidence of flow (contractile) reserve. ^{32,237}	I	B
Intervention should be considered in symptomatic patients with low-flow, low-gradient (< 40 mmHg) aortic stenosis with normal ejection fraction after careful confirmation that the aortic stenosis is severe ^d (Figure 3).	IIa	C
Intervention should be considered in symptomatic patients with low-flow, low-gradient severe aortic stenosis and reduced ejection fraction without flow (contractile) reserve, particularly when CCT calcium scoring confirms severe aortic stenosis.	IIa	C
Intervention is not recommended in patients with severe comorbidities when the intervention is unlikely to improve quality of life or prolong survival > 1 year.	III	C

B) Asymptomatic patients with severe aortic stenosis

Intervention is recommended in asymptomatic patients with severe aortic stenosis and systolic LV dysfunction (LVEF <50%) without another cause.^{9,238,239}

I

B

Intervention is recommended in asymptomatic patients with severe aortic stenosis and demonstrable symptoms on exercise testing.

I

C

Intervention should be considered in asymptomatic patients with severe aortic stenosis and systolic LV dysfunction (LVEF <55%) without another cause.^{9,240,241}

IIa

B

Intervention should be considered in asymptomatic patients with severe aortic stenosis and a sustained fall in BP (>20 mmHg) during exercise testing.

IIa

C

Intervention should be considered in asymptomatic patients with LVEF >55% and a normal exercise test if the procedural risk is low and one of the following parameters is present:

- Very severe aortic stenosis (mean gradient ≥ 60 mmHg or $V_{\max} > 5$ m/s).^{9,242}
- Severe valve calcification (ideally assessed by CCT) and V_{\max} progression ≥ 0.3 m/s/year.^{164,189,243}
- Markedly elevated BNP levels ($> 3 \times$ age- and sex-corrected normal range) confirmed by repeated measurements and without other explanation.^{163,171}

IIa

B

PARTNER Manuscripts in NEJM (October, 2010 – May, 2012)



The NEW ENGLAND JOURNAL of MEDICINE

ESTABLISHED IN 1812

OCTOBER 21, 2010

VOL. 363 NO. 17

Transcatheter Aortic-Valve Implantation for Aortic Stenosis in Patients Who Cannot Undergo Surgery

Martin B. Leon, M.D., Craig R. Smith, M.D., Michael Mack, M.D., D. Craig Miller, M.D., Jeffrey W. Moses, M.D., Lars G. Svensson, M.D., Ph.D., E. Murat Tuzcu, M.D., John G. Webb, M.D., Gregory P. Fontana, M.D., Raj R. Makkar, M.D., David L. Brown, M.D., Peter C. Block, M.D., Robert A. Guyton, M.D., Augusto D. Pichard, M.D., Joseph E. Bavaria, M.D., Howard C. Herrmann, M.D., Pamela S. Douglas, M.D., John L. Petersen, M.D., Jodi J. Akin, M.S., William N. Anderson, Ph.D., Duolao Wang, Ph.D., and Stuart Pocock, Ph.D., for the PARTNER Trial Investigators*

The NEW ENGLAND JOURNAL of MEDICINE

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JUNE 9, 2011

VOL. 364 NO. 23

Transcatheter and Surgical Aortic-Valve Replacement in High-Risk Patients

Craig R. Smith, M.D., Martin B. Leon, M.D., Michael J. Mack, M.D., D. Craig Miller, M.D., Jeffrey W. Moses, M.D., Lars G. Svensson, M.D., Ph.D., E. Murat Tuzcu, M.D., John G. Webb, M.D., Gregory P. Fontana, M.D., Raj R. Makkar, M.D., Mathew Williams, M.D., Todd Dewey, M.D., Samir Kapadia, M.D., Vasilis Babaliaros, M.D., Vinod H. Thourani, M.D., Paul Corso, M.D., Augusto D. Pichard, M.D., Joseph E. Bavaria, M.D., Howard C. Herrmann, M.D., Jodi J. Akin, M.S., William N. Anderson, Ph.D., Duolao Wang, Ph.D., and Stuart J. Pocock, Ph.D., for the PARTNER Trial Investigators*

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

Transcatheter Aortic-Valve Replacement for Inoperable Severe Aortic Stenosis

Raj R. Makkar, M.D., Gregory P. Fontana, M.D., Hasan Jilaihawi, M.D., Samir Kapadia, M.D., Augusto D. Pichard, M.D., Pamela S. Douglas, M.D., Vinod H. Thourani, M.D., Vasilis C. Babaliaros, M.D., John G. Webb, M.D., Howard C. Herrmann, M.D., Joseph E. Bavaria, M.D., Susheel Kodali, M.D., David L. Brown, M.D., Bruce Bowers, M.D., Todd M. Dewey, M.D., Lars G. Svensson, M.D., Ph.D., Murat Tuzcu, M.D., Jeffrey W. Moses, M.D., Matthew R. Williams, M.D., Robert J. Siegel, M.D., Jodi J. Akin, M.S., William N. Anderson, Ph.D., Stuart Pocock, Ph.D., Craig R. Smith, M.D., and Martin B. Leon, M.D., for the PARTNER Trial Investigators*

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

Two-Year Outcomes after Transcatheter or Surgical Aortic-Valve Replacement

Susheel K. Kodali, M.D., Mathew R. Williams, M.D., Craig R. Smith, M.D., Lars G. Svensson, M.D., Ph.D., John G. Webb, M.D., Raj R. Makkar, M.D., Gregory P. Fontana, M.D., Todd M. Dewey, M.D., Vinod H. Thourani, M.D., Augusto D. Pichard, M.D., Michael Fischbein, M.D., Wilson Y. Szeto, M.D., Scott Lim, M.D., Kevin L. Greason, M.D., Paul S. Teirstein, M.D., S. Chris Malaisrie, M.D., Pamela S. Douglas, M.D., Rebecca T. Hahn, M.D., Brian Whisenant, M.D., Alan Zajarias, M.D., Duolao Wang, Ph.D., Jodi J. Akin, M.S., William N. Anderson, Ph.D., and Martin B. Leon, M.D., for the PARTNER Trial Investigators*



The NEW ENGLAND JOURNAL of MEDICINE

Transcatheter Aortic-Valve Replacement with a Self-Expanding Prosthesis

David H. Adams, M.D., Jeffrey J. Popma, M.D., Michael J. Reardon, M.D.,
Steven J. Yakubov, M.D., Joseph S. Coselli, M.D., G. Michael Deeb, M.D.,
Thomas G. Gleason, M.D., Maurice Buchbinder, M.D., James Hermiller, Jr., M.D.,
Neal S. Kleiman, M.D., Stan Chetcuti, M.D., John Heiser, M.D., William Merhi, D.O.,
George Zorn, M.D., Peter Tadros, M.D., Newell Robinson, M.D.,
George Petrossian, M.D., G. Chad Hughes, M.D., J. Kevin Harrison, M.D.,
John Conte, M.D., Brijeshwar Maini, M.D., Mubashir Mumtaz, M.D.,
Sharla Chenoweth, M.S., and Jae K. Oh, M.D.,
for the **J.S. CoreValve Clinical Investigators***

High Risk

8 Mai 2014



CoreValve®

TRANSCATHETER AORTIC VALVE IMPLANTATION (TAVI) PLATFORM



The NEW ENGLAND JOURNAL of MEDICINE

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

Martin B. Leon, M.D., Craig R. Smith, M.D., Michael J. Mack, M.D., Raj R. Makkar, M.D., Lars G. Svensson, M.D., Ph.D., Susheel K. Kodali, M.D., Vinod H. Thourani, M.D., E. Murat Tuzcu, M.D., D. Craig Miller, M.D., Howard C. Herrmann, M.D., Darshan Doshi, M.D., David J. Cohen, M.D., Augusto D. Pichard, M.D., Samir Kapadia, M.D., Todd Dewey, M.D., Vasilis Babaliaros, M.D., Wilson Y. Szeto, M.D., Mathew R. Williams, M.D., Dean Kereiakes, M.D., Alan Zajarias, M.D., Kevin L. Greason, M.D., Brian K. Whisenant, M.D., Robert W. Hodson, M.D., Jeffrey W. Moses, M.D., Alfredo Trento, M.D., David L. Brown, M.D., William F. Fearon, M.D., Philippe Pibarot, D.V.M., Ph.D., Rebecca T. Hahn, M.D., Wael A. Jaber, M.D., William N. Anderson, Ph.D., Maria C. Alu, M.M., and John G. Webb, M.D., for the **PARTNER 2 Investigators**

N Engl J Med 2016; 374:1609-1620 | April 28, 2016 | DOI: 10.1056/NEJMoa1514616

SURTAVI



The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

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

Michael J. Reardon, M.D., Nicolas M. Van Mieghem, M.D., Ph.D., Jeffrey J. Popma, M.D., Neal S. Kleiman, M.D., Lars Søndergaard, M.D., Mubashir Mumtaz, M.D., David H. Adams, M.D., G. Michael Deeb, M.D., Brijeshwar Maini, M.D., Hemal Gada, M.D., Stanley Chetcuti, M.D., Thomas Gleason, M.D., John Heiser, M.D., Rüdiger Lange, M.D., Ph.D., William Merhi, D.O., Jae K. Oh, M.D., Peter S. Olsen, M.D., Nicolo Piazza, M.D., Ph.D., Mathew Williams, M.D., Stephan Windecker, M.D., Ph.D., Steven J. Yakubov, M.D., Eberhard Grube, M.D., Ph.D., Raj Makkar, M.D., Joon S. Lee, M.D., John Conte, M.D., Eric Vang, Ph.D., M.P.H., Hang Nguyen, B.S., Yanping Chang, M.S., Andrew S. Mugglin, Ph.D., Patrick W.J.C. Serruys, M.D., Ph.D., and Arie P. Kappetein, M.D., Ph.D., for the SURTAVI Investigators*

N Engl J Med 2017; 376:1321-1331 [April 6, 2017](#) DOI: 10.1056/NEJMoa1700456

The **NEW ENGLAND**
JOURNAL *of* **MEDICINE**

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MAY 2, 2019

VOL. 380 NO. 18

Transcatheter Aortic-Valve Replacement with a Balloon-
Expandable Valve in Low-Risk Patients

M.J. Mack, M.B. Leon, V.H. Thourani, R. Makkar, S.K. Kodali, M. Russo, S.R. Kapadia, S.C. Malaisrie, D.J. Cohen, P. Pibarot, J. Leipsic, R.T. Hahn, P. Blanke, M.R. Williams, J.M. McCabe, D.L. Brown, V. Babaliaros, S. Goldman, W.Y. Szeto, P. Genereux, A. Pershad, S.J. Pocock, M.C. Alu, J.G. Webb, and C.R. Smith,
for the PARTNER 3 Investigators*

ORIGINAL ARTICLE

Transcatheter Aortic-Valve Replacement with a Self-Expanding Valve in Low-Risk Patients

Jeffrey J. Popma, M.D., G. Michael Deeb, M.D., Steven J. Yakubov, M.D., Mubashir Mumtaz, M.D., Hemal Gada, M.D., Daniel O'Hair, M.D., Tanvir Bajwa, M.D., John C. Heiser, M.D., William Merhi, D.O., Neal S. Kleiman, M.D., Judah Askew, M.D., Paul Sorajja, M.D., Joshua Rovin, M.D., Stanley J. Chetcuti, M.D., David H. Adams, M.D., Paul S. Teirstein, M.D., George L. Zorn III, M.D., John K. Forrest, M.D., Didier Tchétché, M.D., Jon Resar, M.D., Antony Walton, M.D., Nicolo Piazza, M.D., Ph.D., Basel Ramlawi, M.D., Newell Robinson, M.D., George Petrossian, M.D., Thomas G. Gleason, M.D., Jae K. Oh, M.D., Michael J. Boulware, Ph.D., Hongyan Qiao, Ph.D., Andrew S. Mugglin, Ph.D., and Michael J. Reardon, M.D., for the Evolut Low Risk Trial Investigators*

Transcatheter aortic valve implantation vs. surgical aortic valve replacement for treatment of symptomatic severe aortic stenosis: an updated meta-analysis

George C.M. Siontis^{1†}, Pavel Overtchouk ^{1†}, Thomas J. Cahill^{2†}, Thomas Modine³, Bernard Prendergast ⁴, Fabien Praz ¹, Thomas Pilgrim ¹, Tatjana Petrinic⁵, Adriani Nikolakopoulou⁶, Georgia Salanti⁶, Lars Søndergaard⁷, Subodh Verma⁸, Peter Jüni⁹, and Stephan Windecker ^{1*}

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Aims

Owing to new evidence from randomized controlled trials (RCTs) in low-risk patients with severe aortic stenosis, we compared the collective safety and efficacy of transcatheter aortic valve implantation (TAVI) vs. surgical aortic valve replacement (SAVR) across the entire spectrum of surgical risk patients.

Methods and results

The meta-analysis is registered with PROSPERO (CRD42016037273). We identified RCTs comparing TAVI with SAVR in patients with severe aortic stenosis reporting at different follow-up periods. We extracted trial, patient, intervention, and outcome characteristics following predefined criteria. The primary outcome was all-cause mortality up to 2 years for the main analysis. Seven trials that randomly assigned 8020 participants to TAVI (4014 patients) and SAVR (4006 patients) were included. The combined mean STS score in the TAVI arm was 9.4%, 5.1%, and 2.0% for high-, intermediate-, and low surgical risk trials, respectively. Transcatheter aortic valve implantation was associated with a significant reduction of all-cause mortality compared to SAVR {hazard ratio [HR] 0.88 [95% confidence interval (CI) 0.78–0.99], $P=0.030$ }; an effect that was consistent across the entire spectrum of surgical risk (P -for-interaction = 0.410) and irrespective of type of transcatheter heart valve (THV) system (P -for-interaction = 0.674). Transcatheter aortic valve implantation resulted in lower risk of strokes [HR 0.81 (95% CI 0.68–0.98), $P=0.028$]. Surgical aortic valve replacement was associated with a lower risk of major vascular complications [HR 1.99 (95% CI 1.34–2.93), $P=0.001$] and permanent pacemaker implantations [HR 2.27 (95% CI 1.47–3.64), $P<0.001$] compared to TAVI.

Conclusion

Compared with SAVR, TAVI is associated with reduction in all-cause mortality and stroke up to 2 years irrespective of baseline surgical risk and type of THV system.

All-cause mortality

Trial	HR (95% CI)	<i>P</i>
PARTNER 1A	0.90 (0.71 - 1.15)	
US CoreValve high risk	0.79 (0.61 - 1.01)	
NOTION	0.72 (0.33 - 1.59)	
PARTNER 2A	0.92 (0.74 - 1.13)	
SURTAVI	0.98 (0.72 - 1.34)	
PARTNER 3	0.41 (0.14 - 1.17)	
Evolut low risk	0.83 (0.41 - 1.67)	
Overall	0.88 (0.78 - 0.99)	0.030

(Heterogeneity $\tau^2 < 0.001$, $p = 0.727$)

0.2

0.5

1

2

5

Favours TAVI

Favours SAVR

RESULTATS DES 200 PREMIERS TAVI À LA CSC



- ❖ 1^{er} cas de TAVI à La Réunion en octobre 2014
(Dr C.Pouillot)
- ❖ 200 TAVI à la CSC entre 10/2014 et 10/2022.
Drs C.Pouillot ,G.Rambaud et Y.Gadri..
- ❖ Clinique Sainte Clotilde en 2023 environ 60 TAVI trans femoraux et 6 TAVI chirurgicaux (trans aortique et trans apical) .

FRANCE TAVI REGISTRE MIS EN PLACE EN 2013 POUR 5 ANS.. ACTUELLEMENT TOUJOURS EN COURS..)



France TAVI



OBSERVATOIRE SUR LES BIOPROTHÈSES VALVULAIRES AORTIQUES IMPLANTABLES PAR CATHÉTER

Centre : Clinique Sainte Clotilde

Principal Investigateur : DR C. POUILLOT

Nombre de patients TAVI pris en charge : 33

Nombre d'inclusions dans l'observatoire : 29 (1 décès per procédure non inclus)

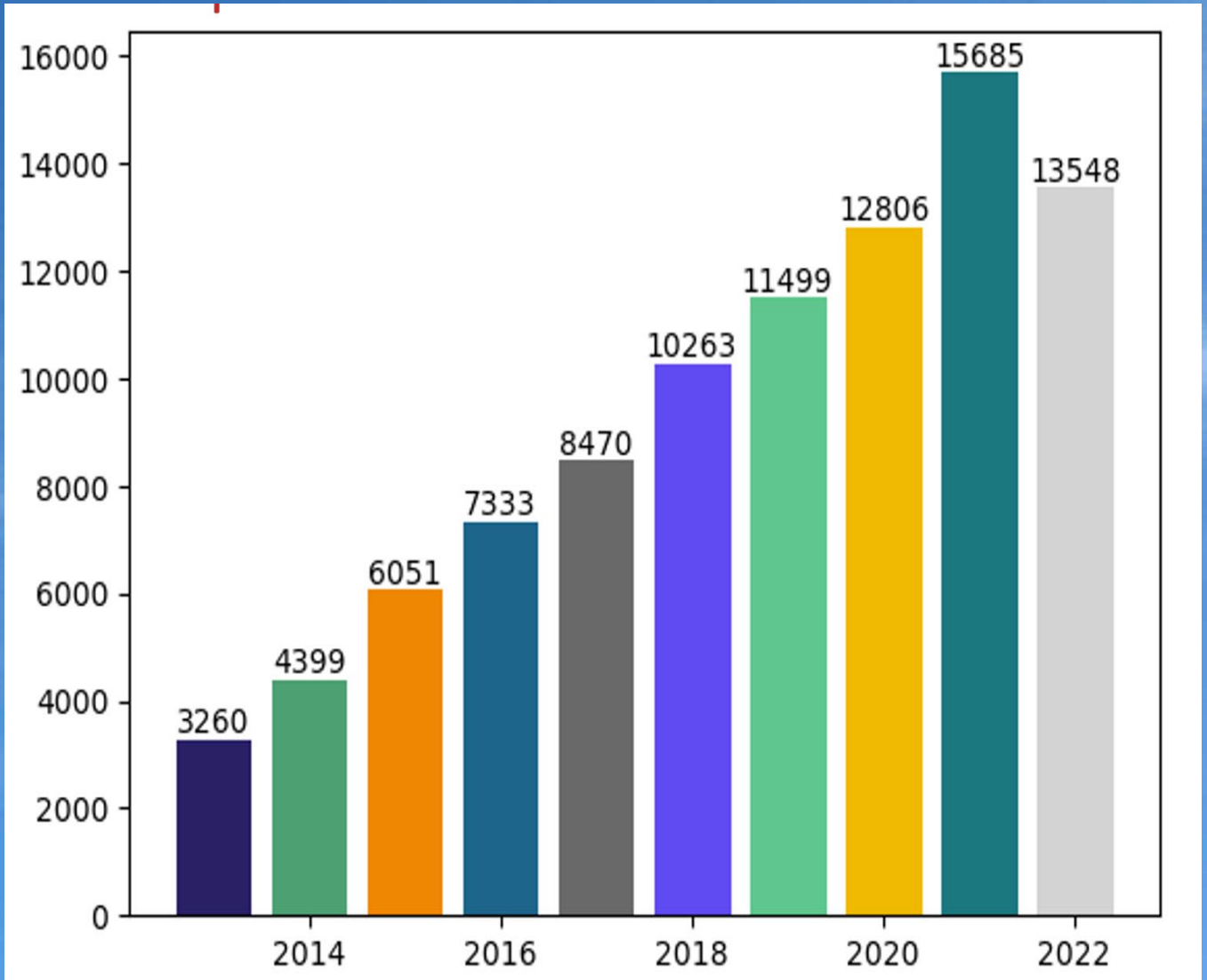
Tous centres : 10264 patients (17/11/2015)

La CSC a intégré France TAVI des 2015

AMIENS	HOPITAL SUD - CHU AMIENS	1	MULHOUSE	C.H. DU MOENCHSBERG - EMILE MULLER	27
ANGERS	C.H.U. D' ANGERS	2	NANTES	C.H.U	28
BESANCON	C.H.U	3	NEUILLY	CLINIQUE AMBROISE PARE	29
BOIS-BERNARD	HOPITAL PRIVE DE BOIS-BERNARD-LENS	4	PARIS	INSTITUT MUTUALISTE MONTSOURIS	30
BORDEAUX	CLINIQUE SAINT-AUGUSTIN	5	PARIS	GROUPE HOSPITALIER PITIE SALPETRIERE (AP-HP)	31
BREST	CHU HOPITAL CAVALE BLANCHE	6	PARIS	HOPITAL EUROPEEN GEORGES POMPIDOU	32
BRON	Hôpital Cardiologique Louis Pradel	7	PARIS	GROUPE HOSPITALIER BICHAT CLAUDE BERNARD	33
CAEN	CENTRE HOSPITALIER PRIVE ST MARTIN	8	PESSAC	CHU HOPITAUX DE BORDEAUX	34
CAEN	CHU COTE DE NACRE - CAEN	9	POITIERS	CHU - LA MILETRIE	35
CALUIRE ET CUIRE	INFIRMERIE PROTESTANTE DE LYON	10	PRINGY	CENTRE HOSPITALIER REGION ANNECIENNE	36
CLERMONT-FERRAND	C.H.U. - HOPITAL G. MONTPIED	11	REIMS	HOPITAL ROBERT DEBRE CHR REIMS	37
CRETEIL	GROUPE HOSPITALIER HENRI MONDOR - ALBERT CHENEVIER	12	RENNES	C.H.U DE RENNES - HOPITAL PONTCHAILLOU	38
DIJON	CHU HOPITAL DU BOCAGE	13	ROUEN	HOPITAL CHARLES NICOLLE - CHU ROUEN	39
GRENOBLE	CHU GRENOBLE - MICHALLON	14	SAINT-DENIS	CENTRE CARDIOLOGIQUE DU NORD	40
LE CHESNAY	CENTRE MEDICO CHIRURGICAL DE PARLY II	15	SAINT-ETIENNE	CHU SAINT ETIENNE - HOPITAL NORD	41
LE PLESSIS-ROBINSON	CENTRE CHIRURGICAL MARIE LANNELONGUE	16	SAINT-LAURENT-DU-VAR	INSTITUT ARNAULT TZANCK	42
LILLE	CHRU de Lille - Hôpital Cardiologique	17	STRASBOURG	CHU DE STRASBOURG - HOPITAL CIVIL	43
LILLE	POLYCLINIQUE DU BOIS	18	TOULOUSE	HOPITAL DE RANGUEIL - CHU TOULOUSE	44
LIMOGES	CHU DUPUYTREN LIMOGES	19	TOULOUSE	CLINIQUE PASTEUR	45
LYON	CLINIQUE DE LA SAUVEGARDE	20	TOURS	CLINIQUE SAINT GATIEN	46
MARSEILLE	HOPITAL PRIVE CLAIRVAL	21	TOURS	HOPITAL TROUSSEAU - CHU	47
MARSEILLE	HOPITAL LA TIMONE - CHU	22	VANDOEUVRE-LES-NANCY	HOPITAL DE BRABOIS - CHU DE NANCY	48
MARSEILLE	HOPITAL SAINT-JOSEPH	23	VILLEURBANNE	CLINIQUE DU TONKIN	49
MASSY	INSTITUT HOSPITALIER JACQUES CARTIER - ICPS	24	LA REUNION	CLINIQUE SAINTE CLOTILDE	50
METZ	HOPITAL-CLINIQUE CLAUDE BERNARD	25			
MONTPELLIER	CHU MONTPELLIER	26			

PATIENTS INCLUS DANS FRANCE-TAVI

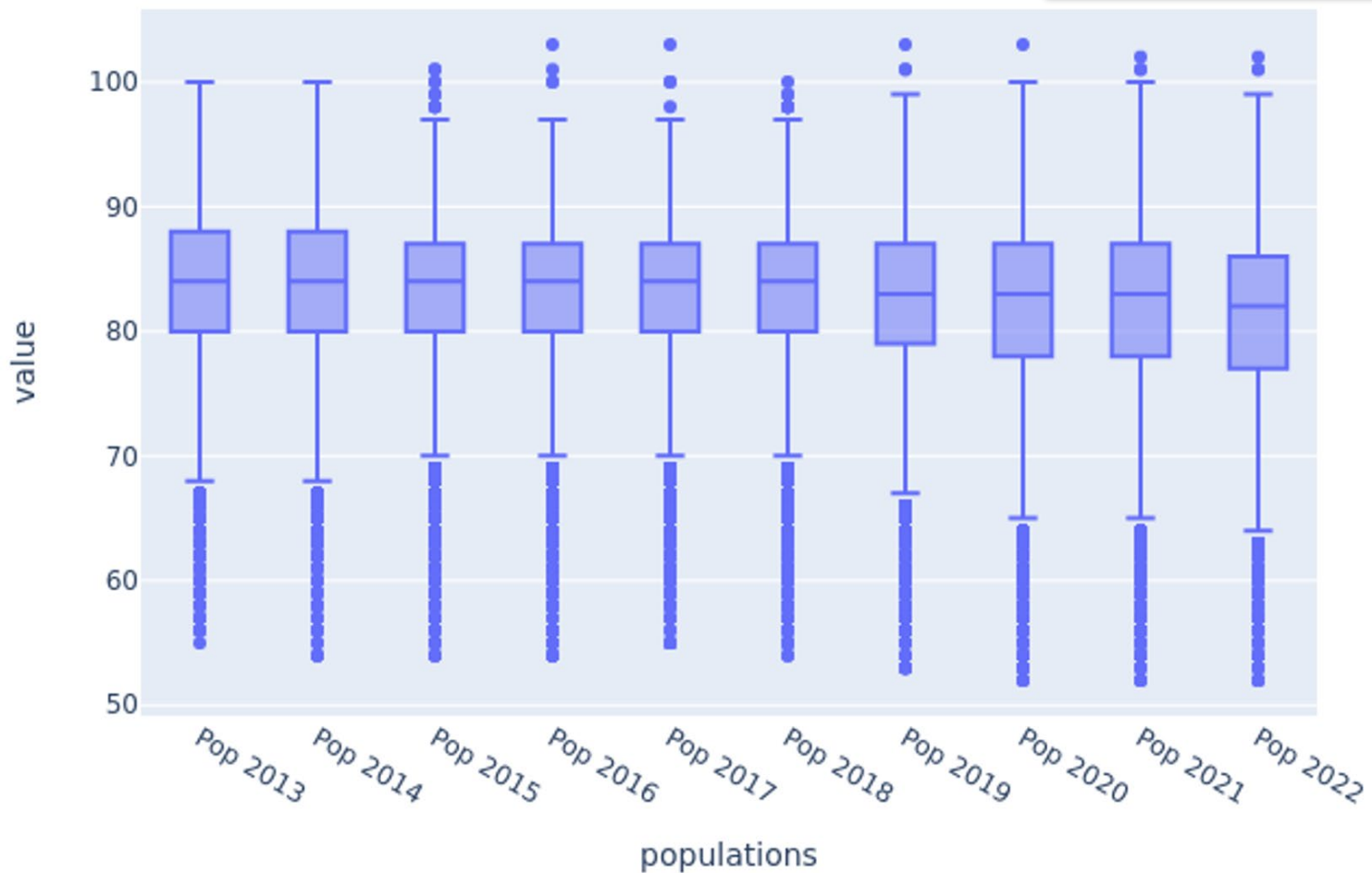
Année 2013-2022



AGES DES PATIENTS

Evolution 2013-2022

Age moy > 80 ans

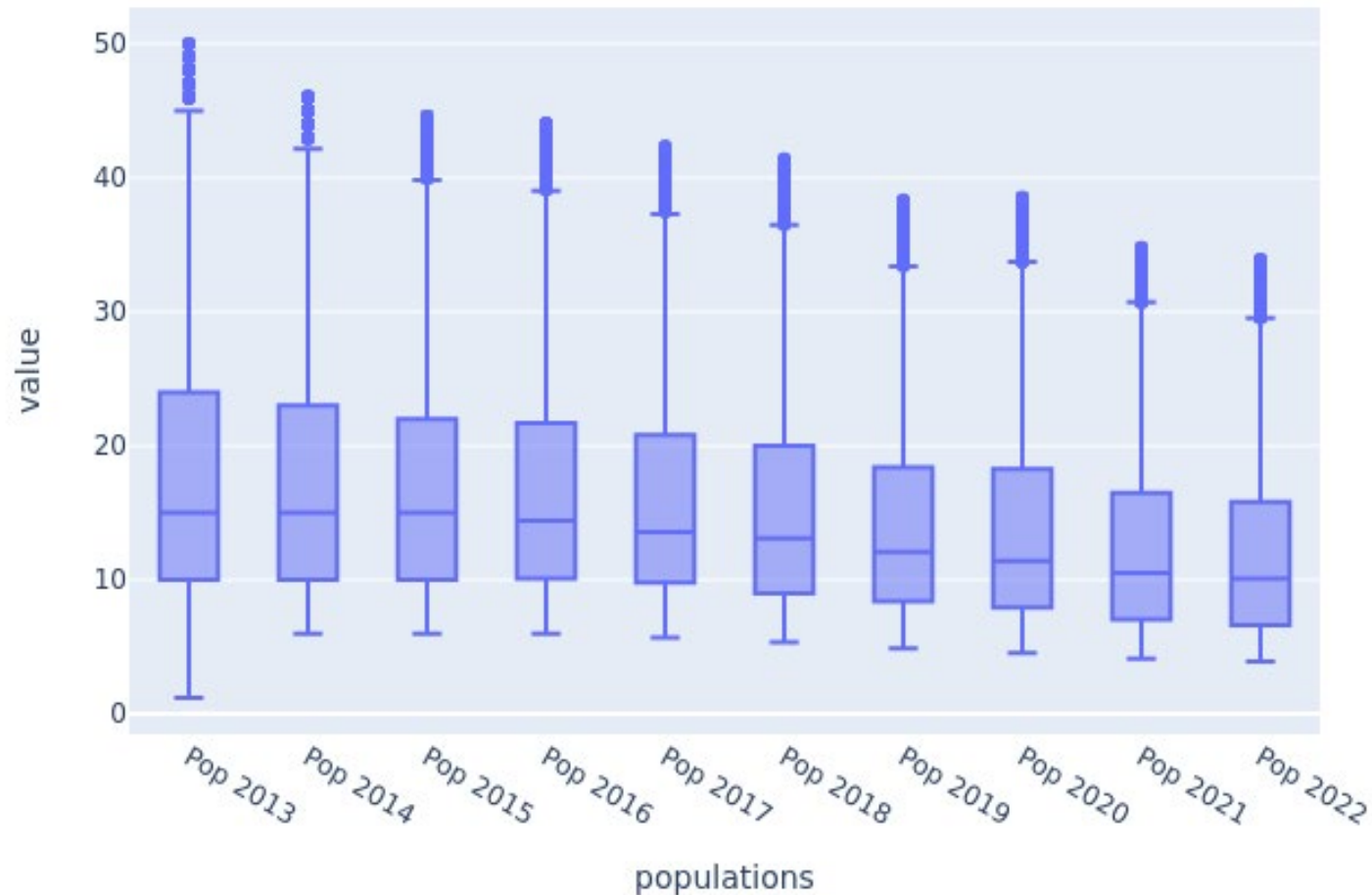


France TAVI

LOGISTIC EUROSCORE

Evolution 2013-2022

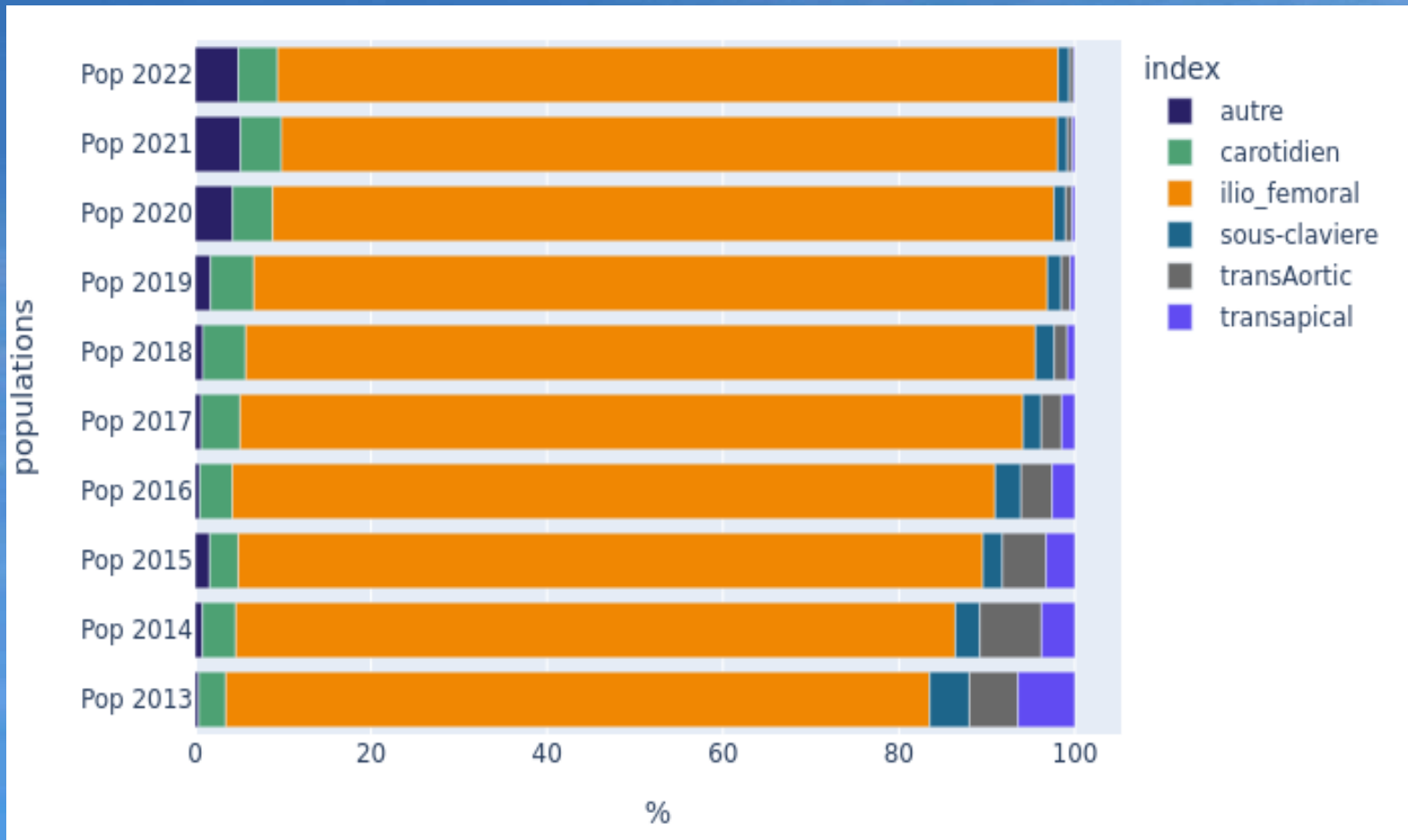
Euroscore > 10



France TAVI

VOIE D'ABORD

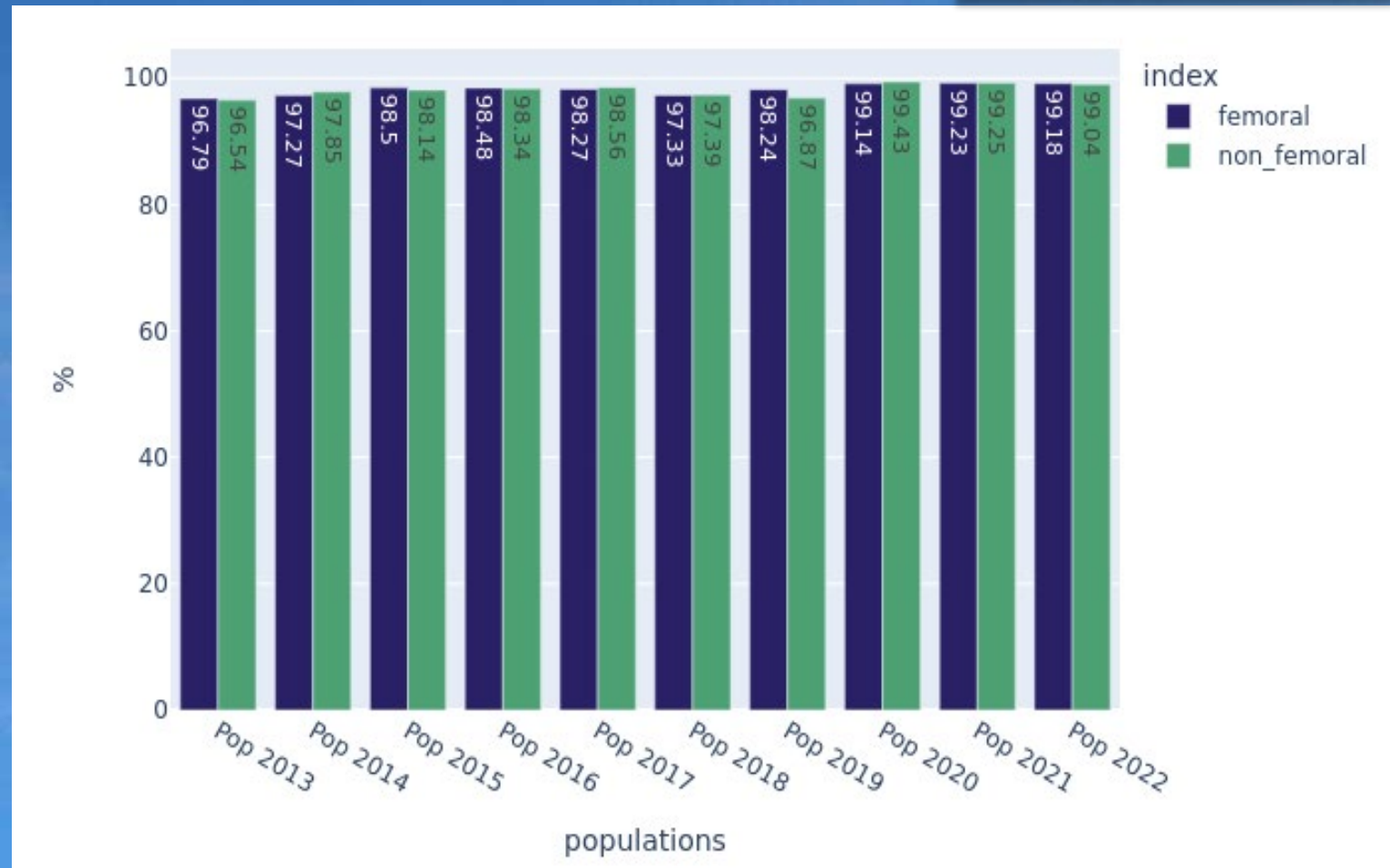
Evolution 2013-2022



TAUX DE SUCCÈS IMMÉDIAT

Evolution 2013-2022

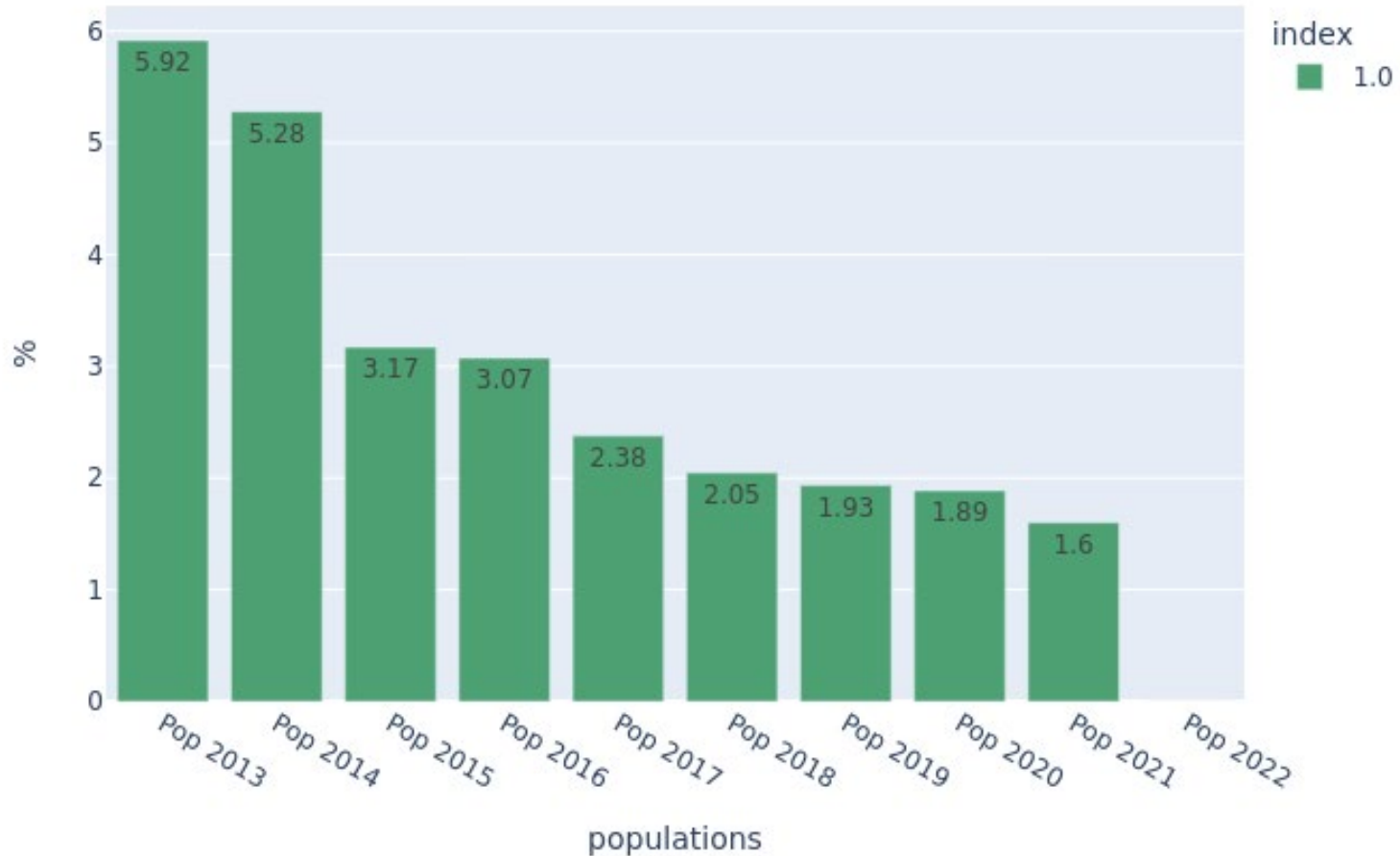
99.2 % de succès
Par voie Fém.



MORTALITÉ HOSPITALIÈRE

Evolution 2013-2022

Mortalité Hosp. 1,6 %



Les résultats CSC TAVI vs France TAVI

	Clinique Sainte Clotilde		France TAVI 2021		Reunion
	n =200	%	n =13081	%	
Genre					
Hommes	82	41%		55,3%	
Femmes	118	59%		44,7%	+ de femmes
Age					
Minimum	54		-		
Moyenne +/- écart-type	81	+/- 7,19	81,79		≈Un an + jeune
Maximum	101		103		

INDICATION ET CALCUL SCORES

Indication principale	Clinique Sainte Clotilde		France TAVI 2021		Run
	n	%		%	
Chirurgie contre indiquée	21	10,5%		3,27%	
Haut risque opératoire	51	25.5%		22,01%	
Fragilité	27	13.5%		17,82%	
Autre (refus RVAo)	8	4%		0,64%	+ pts refus RVAo
Symptomatique	191	90%		87,87%	
Logistic Euroscore					
Minimum					
Moyenne +/- écart-type	18.72		13.84		< 0.01 patients plus sévères
Maximum					

HISTOIRE MÉDICALE ET ANTÉCÉDENTS

	Clinique Sainte Clotilde		France TAVI 2021		Run
		%		%	
Diabète					
Oui	91	45,5		33.54%	+ de diabétiques
Insuffisance rénale chronique					
Non	105	52,5		67.11%	+ IRC
Modérée	59	29,5		23.31%	
Sévère	25	12,5%		7,89%	+ IRC sévères
Dialyse	11	5,5%		1.69%	+ hémodialyse

BILAN ECHO PRÉ IMPLANTATION

	Clinique Sainte Clotilde	France TAVI 2021		Run
	%		%	
Echographie : Gradient moyen (mm Hg)				
Moyenne	45,76	48,0		Gdt moyen -2
Echographie : Surface valvulaire (cm²)				
Moyenne +/- écart-type	0,69	0,75		SAo - 0.06
Echographie : FEVG (%)				
Moyenne +/- écart-type	58.85 %	52.92 %		FE + 5%
HTAP				
Non	13,5%		35.04 %	
Modérée (30-55)	65%		56.88 %	+ HTAP
Sévère (>55)	12,5%		8.08 %	+ HTAP sévère

CARACTÉRISTIQUES PROCÉDURES

	Clinique Sainte Clotilde	France TAVI 2022	Run
	%	%	
Abord			
Ilio-fémoral	100%	# 90%	<0.001
Anesthésie			
Locale et/ou sédation	100%	88,3%	minimal invasive TAVI

TYPES DE VALVES

	Clinique Sainte Clotilde		France TAVI 2021	Run
	n	%	%	
Type de la valve 1				
Corevalve Evolut R Evolut Pro Evolut Pro +	13	6,5%	28.33 %	
Edwards Sapien 3 + XT	185	92.5%	64.98 %	Sapien 3
Diamètre de la valve 1				
23	108	50,2%	25,8%	Plus petites valves
26	73	34%	39,8%	
29	24	11,2%	26,9%	
20	4	1,9%		

PROCÉDURES

Succès procédure (définition VARC)	CSC		France TAVI 2022		Run
	Oui	197	98,5%	12139	99,2%

COMPLICATIONS PER PROCEDURES

Complications	CSC		France TAVI 2021	Run
	Nombre	%	%	
AVC	2	1%	2.09 %	- AVC
Tamponnade	4	2%	< 1 %	+ tamponnade
Troubles conductifs avec implantation d'un pacemaker (durant hospitalisation)	28	14%	16.21 %	Un peu - PM
Rupture d'anneau	1	0,5%	< 1 %	id
Occlusion coronaire	0	0%	< 1 %	
Migration de valve	1	0,5%	< 1 %	id
Complication Abord fémoral (stent, réparation chir...)	1		2.63 %	Pas +...

FUITES AORTIQUES

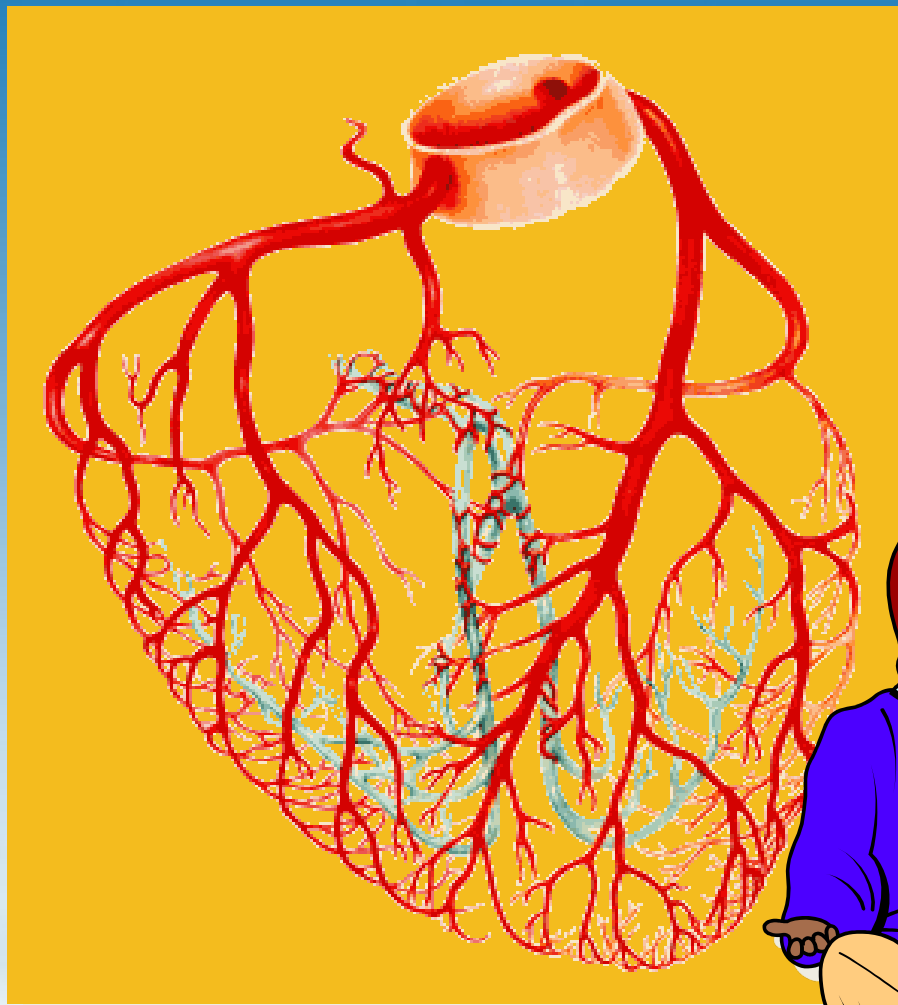
Fuite aortique ≥ 2 en fin de procédure

Oui	8	4%	458	4,46 %	id
Non	167	96%	5378	95.54 %	

MORTALITE

CSC			France TAVI 2020 - 2021	Run
Décès	Nombre	Pourcentage	Pourcentage	Pourcentage
Hospitalier	3	1.5 %	1.45 %	id
Décès < J30	9	4,5%		
Décès < 1 an	46	23 % Dans ¾ cas non cardiaque..	11.02 %	+ décès à 1 an (dans 3/4 cas non cardiaques)

Merci



**COMMISSION NATIONALE D'ÉVALUATION
DES DISPOSITIFS MÉDICAUX ET DES TECHNOLOGIES DE SANTÉ**

AVIS DE LA CNEDiMTS

16 mars 2021

Faisant suite à l'examen du 19/01/2021, la CNEDiMTS a adopté un projet d'avis le 02/02/2021.

Ce projet d'avis a fait l'objet d'une phase contradictoire le 16/03/2021. La CNEDiMTS a adopté l'avis le 16/03/2021.

CONCLUSIONS

EDWARDS SAPIEN 3, bioprothèse valvulaire aortique implantée par voie transfémorale (système COMMANDER)

Demandeur : EDWARDS LIFESCIENCES SAS (France)

Fabricant : EDWARDS LIFESCIENCES LLC (Etats-Unis)

Les modèles et références retenus sont ceux proposés par le demandeur (cf. page 5)

Indications retenues :	<p>Patients avec sténose aortique native sévère symptomatique (SVAoi < 0,5 cm²/m²). L'indication doit être posée lors d'une réunion multidisciplinaire en prenant en compte les scores de risque et les comorbidités associées. Pour les patients opérables avec un score STS < 4%, l'indication est limitée aux patients de plus de 65 ans, avec un orifice tricuspide, ne pas avoir d'indication de chirurgie valvulaire mitrale ou coronaire (tronc commun et/ou SYNTAX > 32) associée et avec une anatomie favorable à la voie transfémorale.</p> <p>Les patients ayant une espérance de vie inférieure à 1 an compte tenu de facteurs extracardiaques (comorbidités) ou ayant des calcifications importantes dans la chambre de chasse sous aortique ne sont pas éligibles à la technique (non-indication). Il est rappelé la nécessité du respect des contre-indications figurant au marquage CE du dispositif EDWARDS SAPIEN 3.</p>
Service Attendu / Rendu (SA/SR) :	Suffisant.
Comparateurs retenus :	<p>Pour les patients avec contre-indication à la chirurgie de remplacement valvulaire aortique ou les patients opérables avec un score STS ≥ 4% : aux autres bioprothèses valvulaires aortiques implantées par voie transcathéter inscrites sur la LPPR dans les indications retenues.</p> <p>Pour les patients opérables avec un score STS < 4% : chirurgie de remplacement valvulaire aortique.</p>

Heart team : qui fait quoi ?

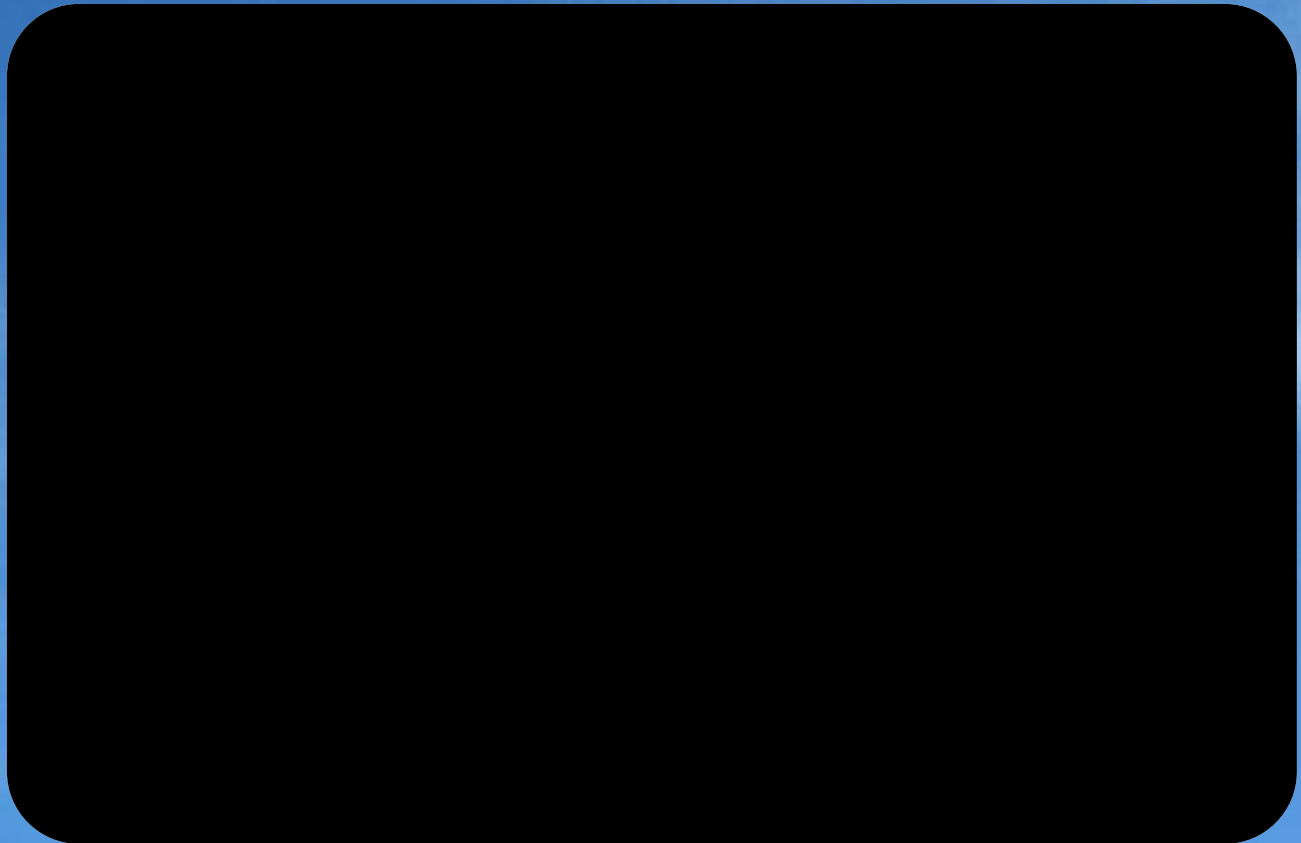
Indications du TAVI	QUI évalue ?
Rétrécissement aortique serré symptomatique	→ Cardiologue
Non opérabilité	→ Heart Team
Espérance de vie > 1 an	→ Score pronostic cardiologique (STS, EuroSCORE) → Cardiologue → Éval. Fragilité et co-morbidités → Gériatre
Possibilité d'amélioration de la qualité de vie	→ Gériatre

RVAo ? TAVI ?

La discussion Heart team

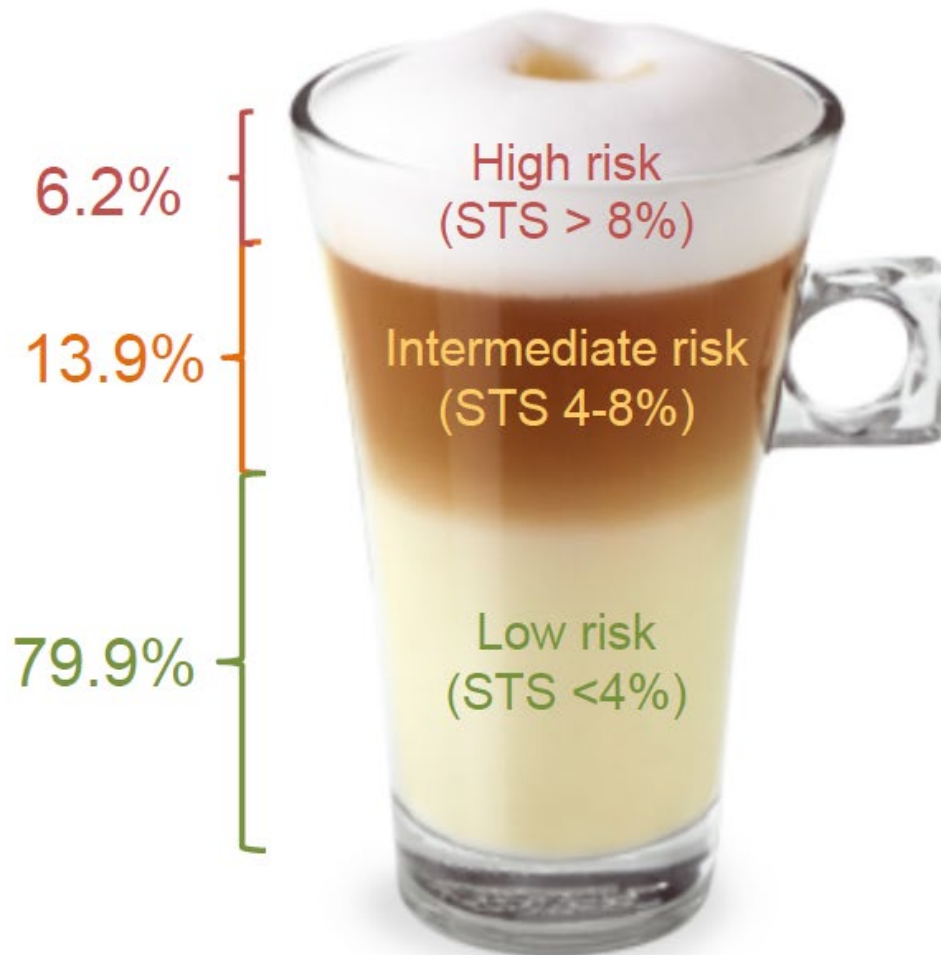
- Le cardiologue interventionnel
- Le (la) cardiologue correspondant (e), (non invasif)
- Le chirurgien cardiaque
- L' anesthésiste
- La gériatre

C'est quoi le TAVI ?

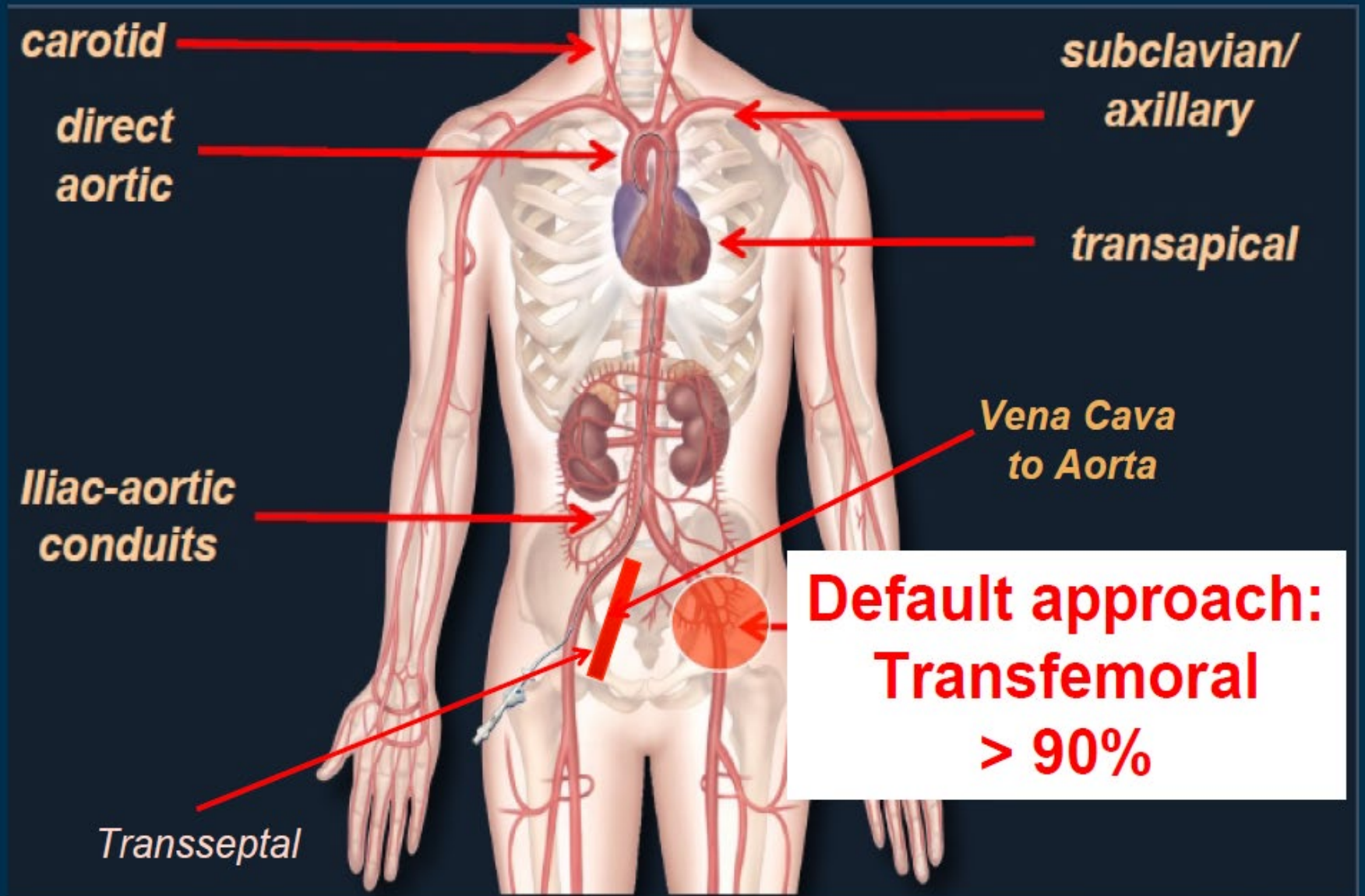


STS database 2002-2010

n=141,905 patients

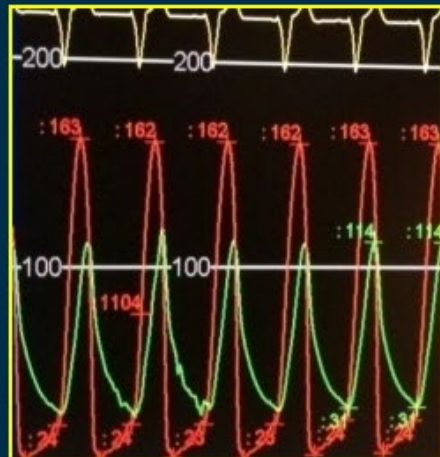


Multiple access sites for improved sa

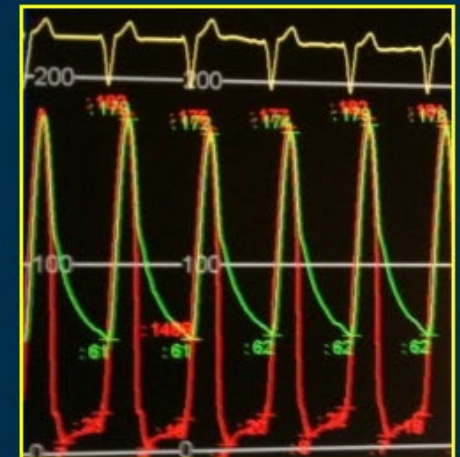


Valve – in – Valve for degenerated bioprosthesis

Dvir et al-VIR multicentric Registry, JAMA 2014 : 459 pts



- Importance of Pt selection (type of valve, internal diameter)
- ViV application



The 3MM strategy in Vancouver

- Multidisciplinary
- Multimodality
- Minimalist
 - ✓ TF access
 - ✓ next day discharge

THE VANCOUVER SUN



Dr David Wood from St Paul Hospital: Same day discharge after TAVR!



A 97 years old women discharged the same day after a TAVR procedure stretching while waiting for the bus to get home... read more on page 3.

PROPORTION DE FEMMES

Evolution 2013-2022

44.7 % de femmes

