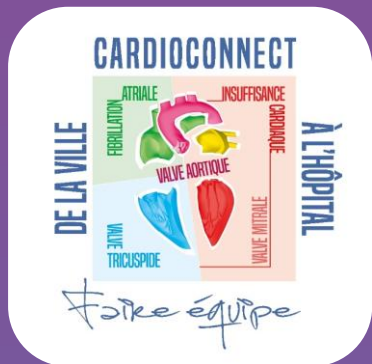


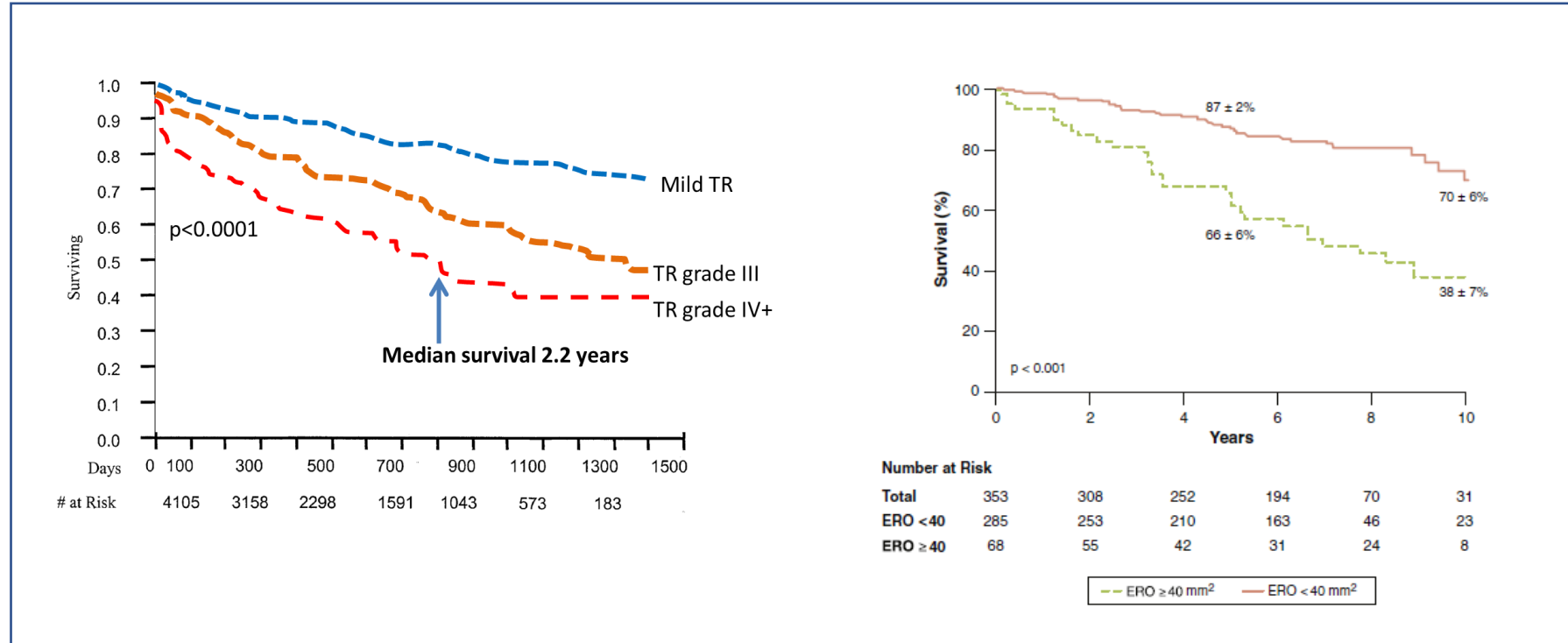
# Traitement percutané des valves tricuspides : réparation ou remplacement ?

Dr Romain GALLET, Pr Emmanuel TEIGER

CHU Henri Mondor, APHP, Créteil



# Pronostic en cas d'IT ?

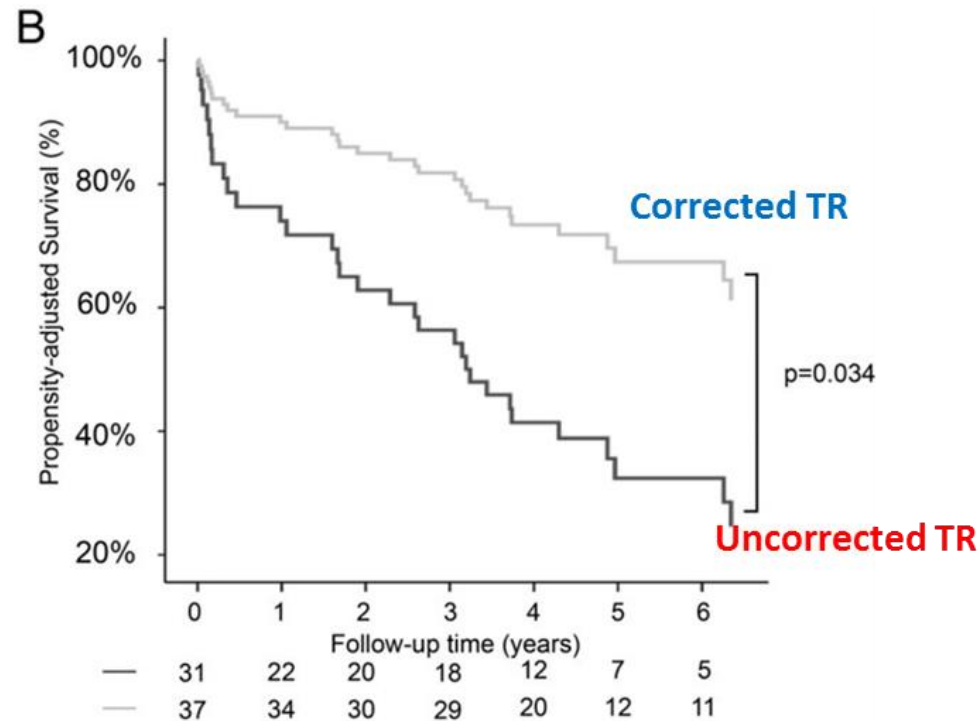


- ❑ Le pronostic de la fuite tricuspide sous traitement médical dépend de sa sévérité
- ❑ La  $\text{SOR} \geq 40 \text{ mm}^2$  a été validée comme un critère pronostic indépendant dans les IT isolées
- ❑ La médiane de survie est 2 à 6 ans selon l'étiologie de la fuite.

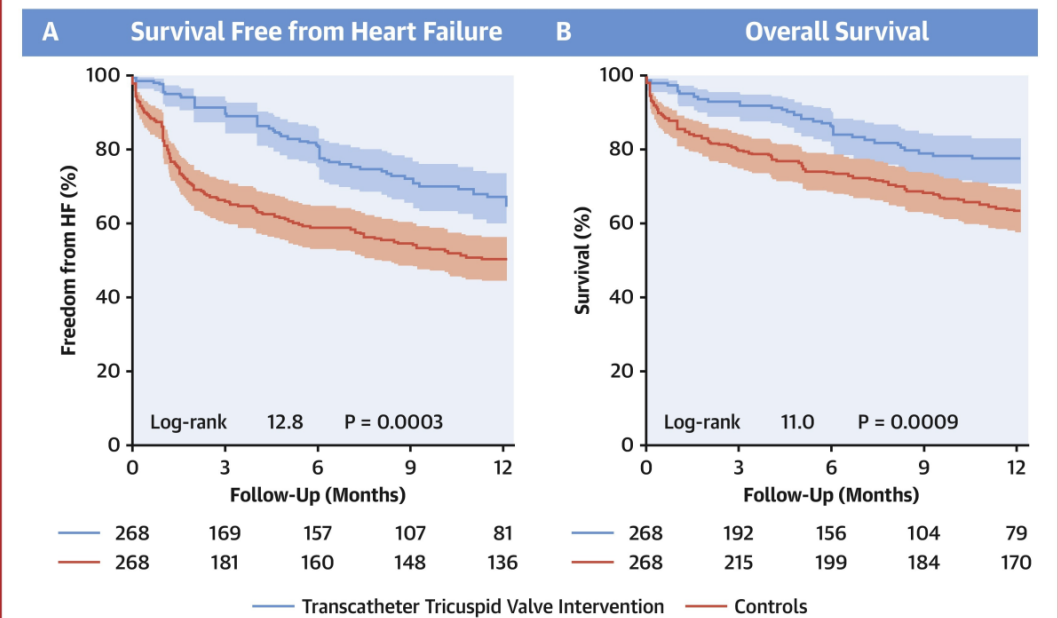
# Faut-il corriger les IT?



## Transcatheter Versus Medical Treatment of Patients With Symptomatic Severe Tricuspid Regurgitation

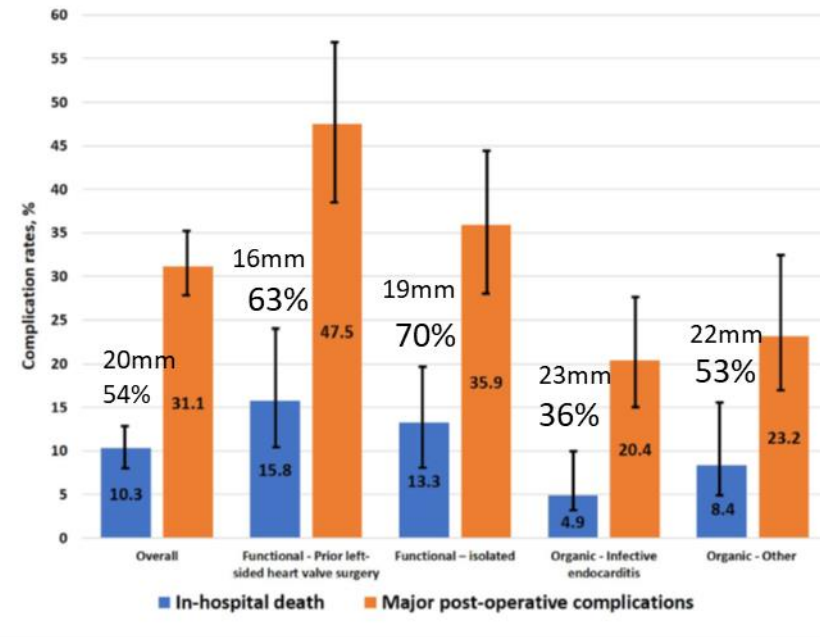
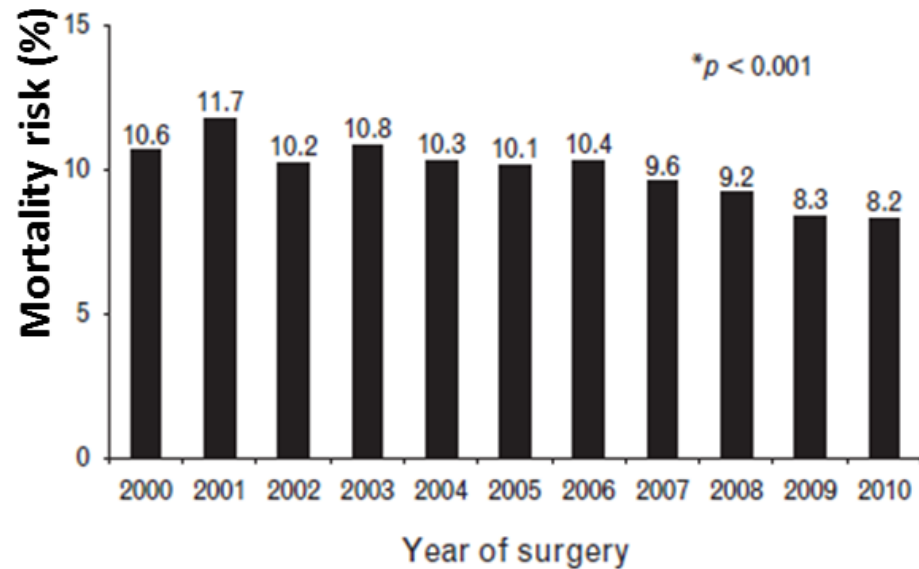


### CENTRAL ILLUSTRATION: Transcatheter Treatment of Severe Tricuspid Regurgitation: Primary and Secondary Endpoints



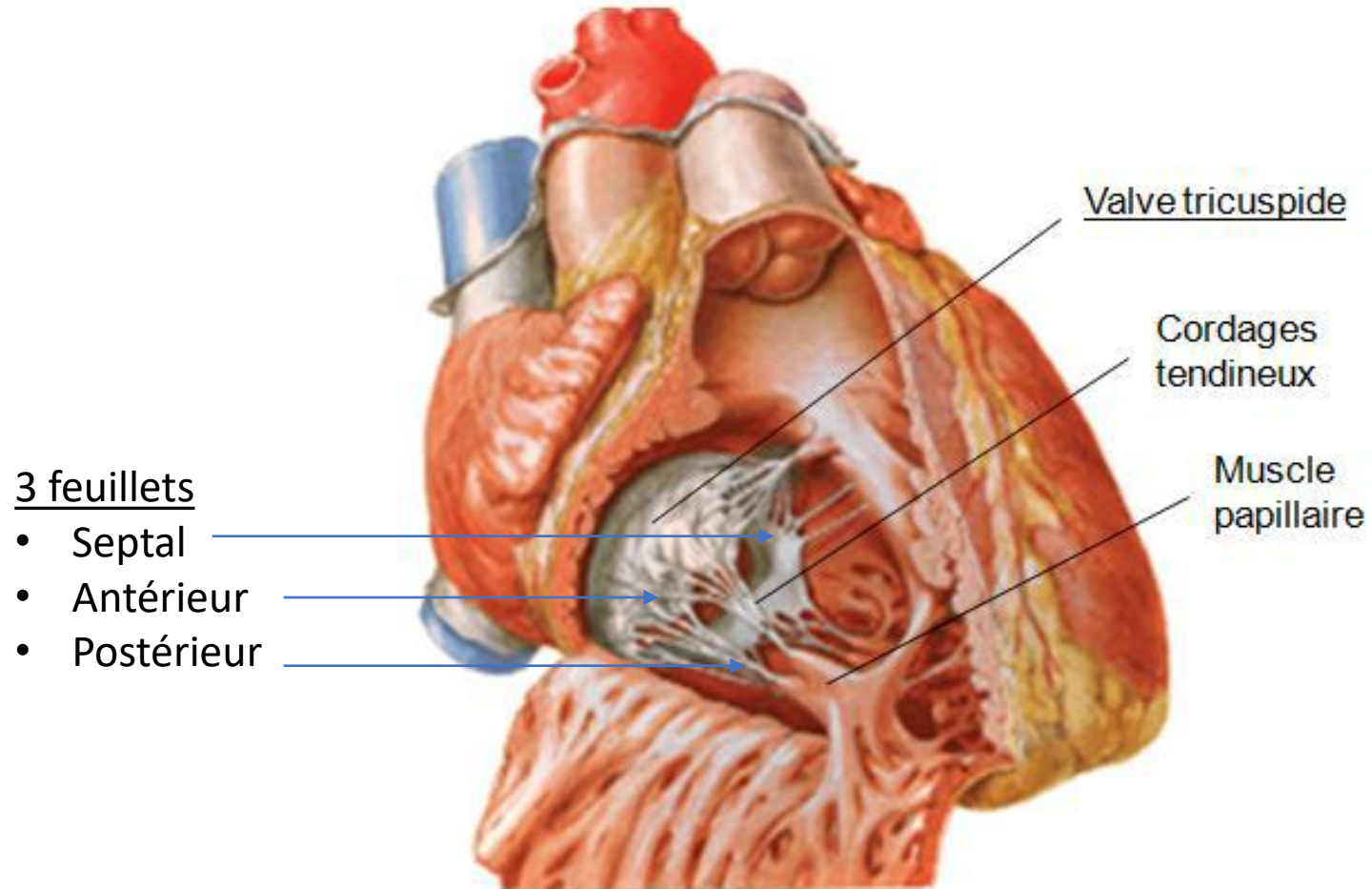
Taramasso, M. et al. J Am Coll Cardiol. 2019;74(24):2998-3008.

# Risque opératoire



- ❑ La mortalité est d'environ 10% lorsque l'on inclut les fuites organiques et fonctionnelles
- ❑ Dans les fuites fonctionnelles pures, la mortalité est plutôt de 13 à 16%

# Réparer ou remplacer?

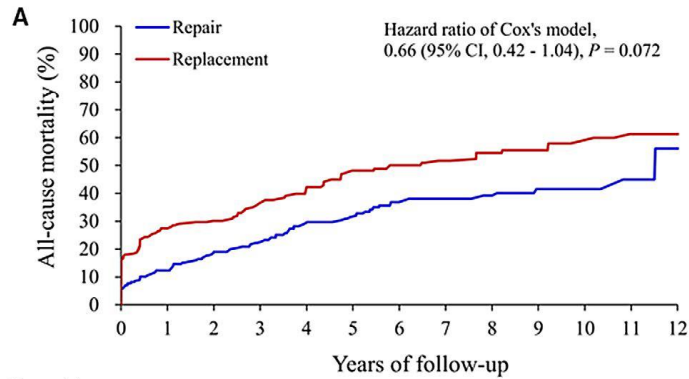


3 feuillets

Nombreux cordages

→ complique potentiellement la réparation

## Isolated tricuspid surgery

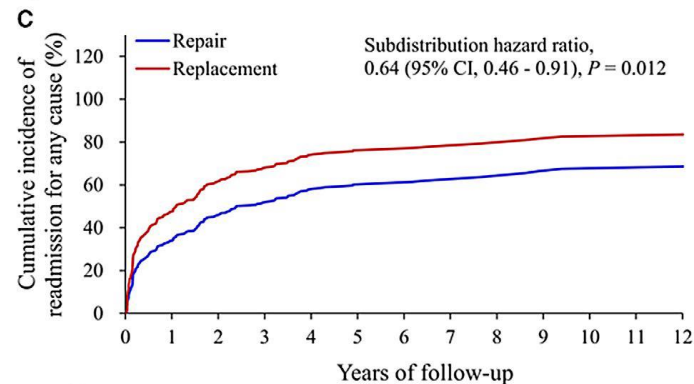


No. at risk:

Repair	330	223	162	107	75	47	22
Replacement	340	194	148	103	73	49	31

No. at risk:

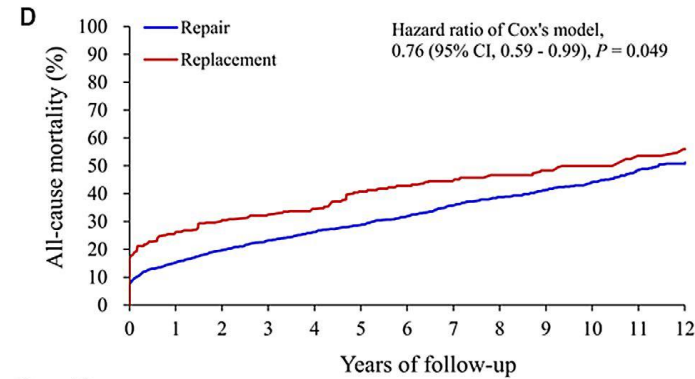
Repair	311	200	147	94	65	36	13
Replacement	293	151	87	73	55	37	29



No. at risk:

Repair	311	140	75	54	34	19	8
Replacement	293	71	32	19	12	3	3

## Concomitant tricuspid surgery

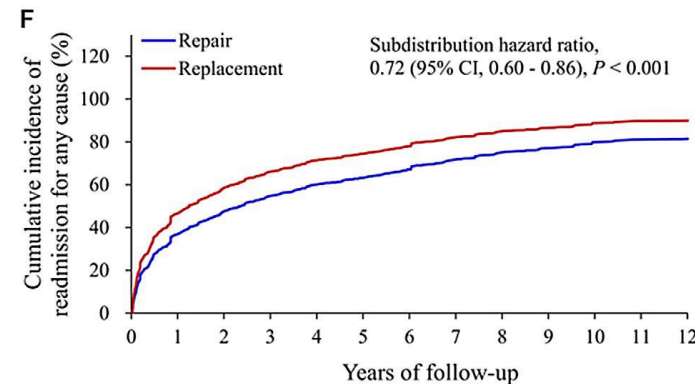


No. at risk:

Repair	2312	1586	1186	888	602	380	161
Replacement	2281	1348	1038	643	420	255	118

No. at risk:

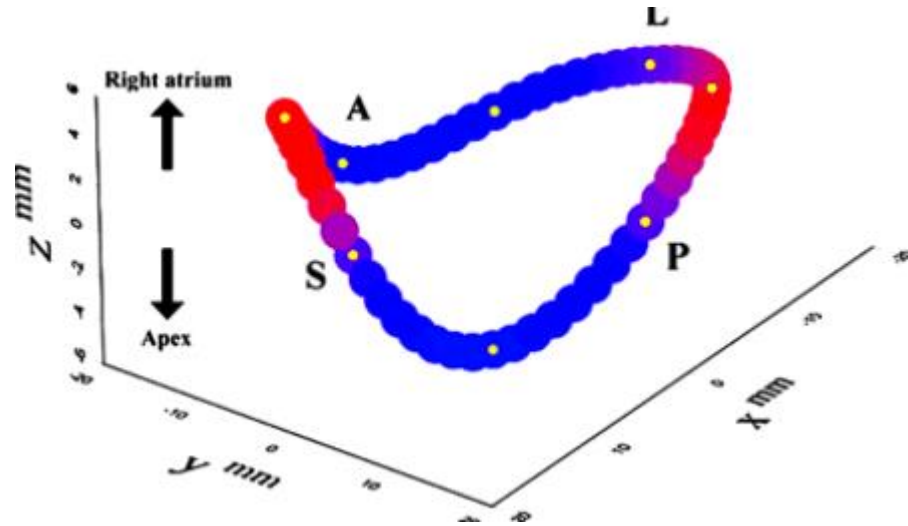
Repair	2144	1412	988	705	460	275	109
Replacement	1910	991	723	455	274	129	65



No. at risk:

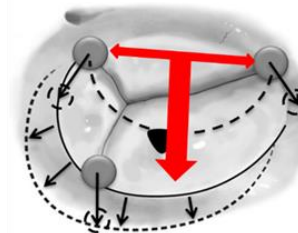
Repair	2144	931	516	321	168	83	30
Replacement	1910	604	401	199	84	38	26

# Réparer ou remplacer?

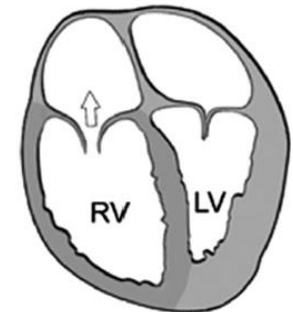


Diamètre <40mm ou <21mm/m<sup>2</sup>  
Normal=28±5mm<sup>2</sup>  
Surface =8cm<sup>2</sup>  
Variation de surface 30%

1- Dilatation de l'anneau



2- Dilatation du VD

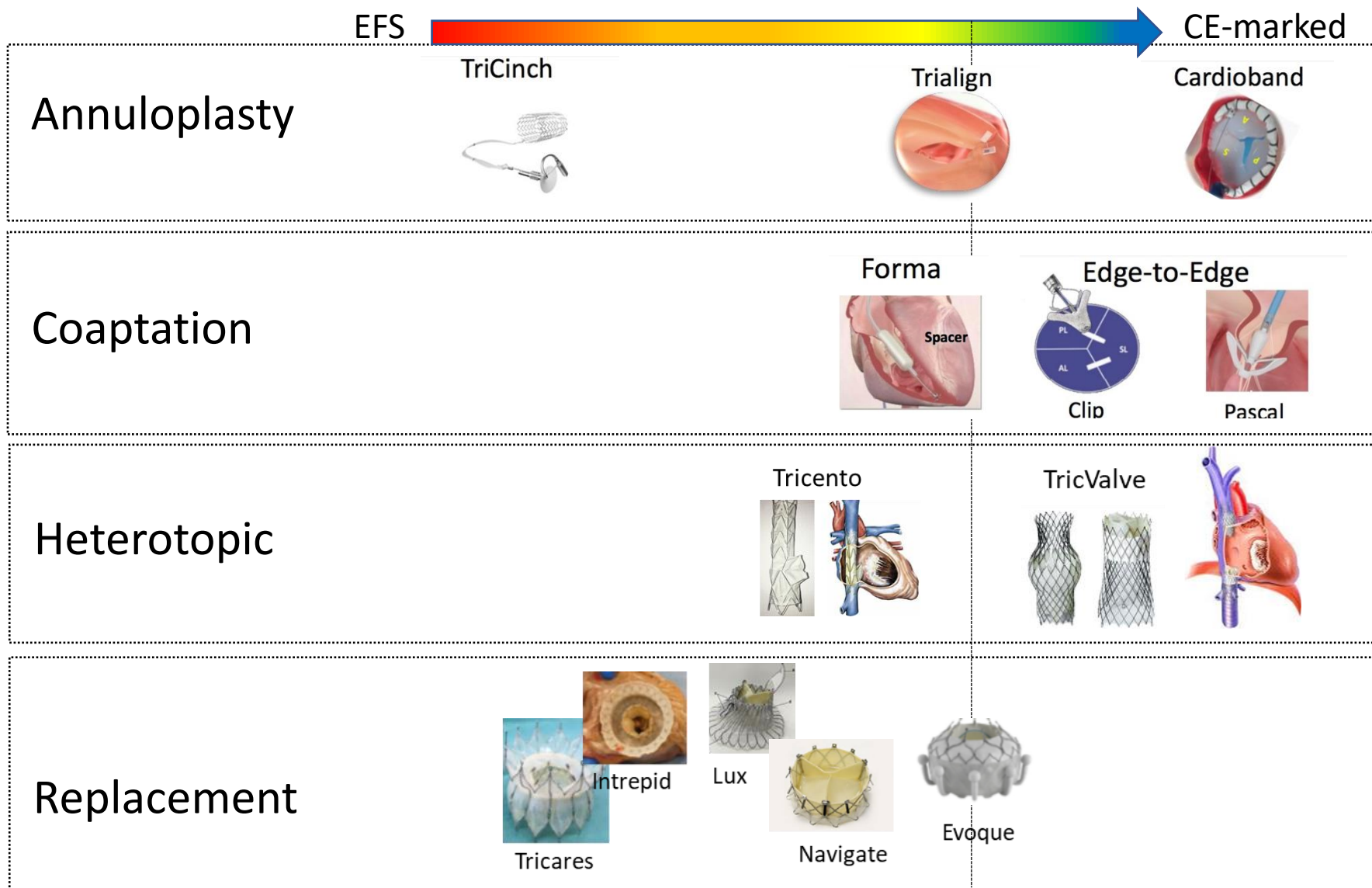


- Anneau en forme de selle de cheval
- Variation au cours du cycle cardiaque
- Dilatation importante anneau et VD
- Pas de « landing zone évidente, pas de calcifications

→ complique potentiellement le remplacement



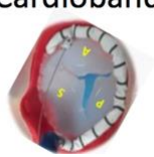









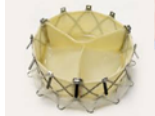

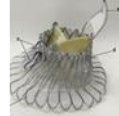




# Différentes possibilités



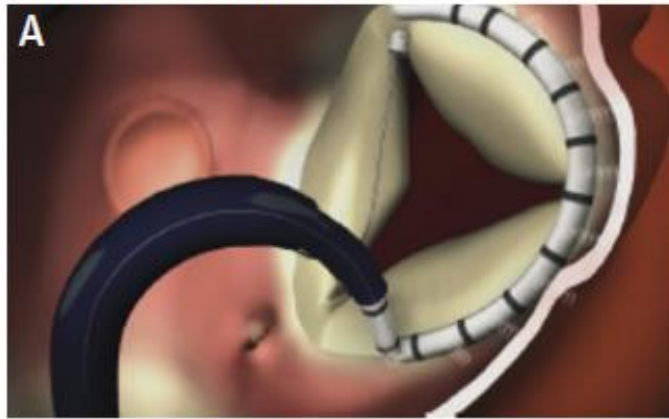


# Différentes possibilités

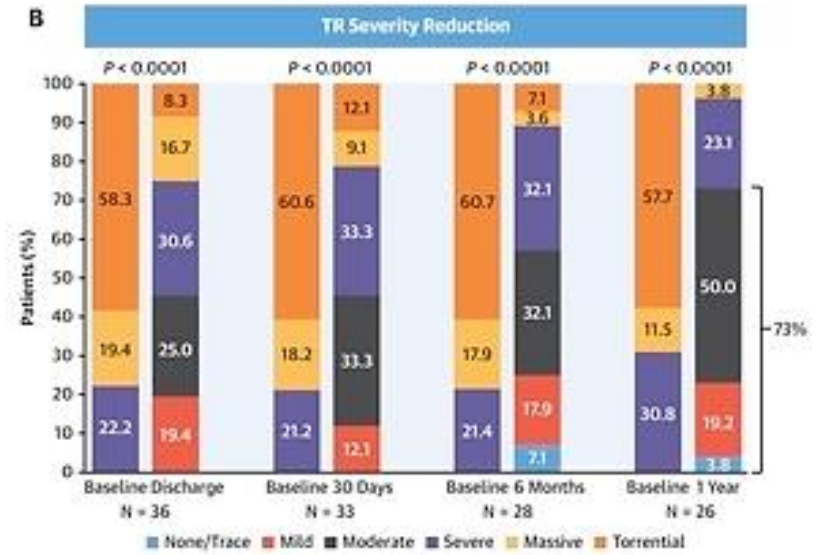
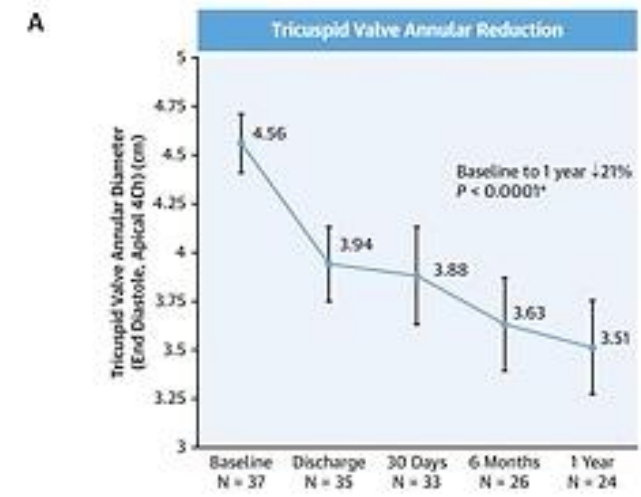
	EFS	CE-marked	
Annuloplasty	TriCinch 	Trialign 	Cardioband 
Coaptation		Forma 	Edge-to-Edge  Clip  Pascal 
Heterotopic		Tricento  	TricValve   
Replacement	Navigate 	Intrepid 	Lux 
		Tricares 	Evoque 

# Annuloplasty: cardioband

Cardioband Anchor Deployment



Reduction in Tricuspid Valve Annular Diameter and TR Severity at 1 Year With Cardioband System

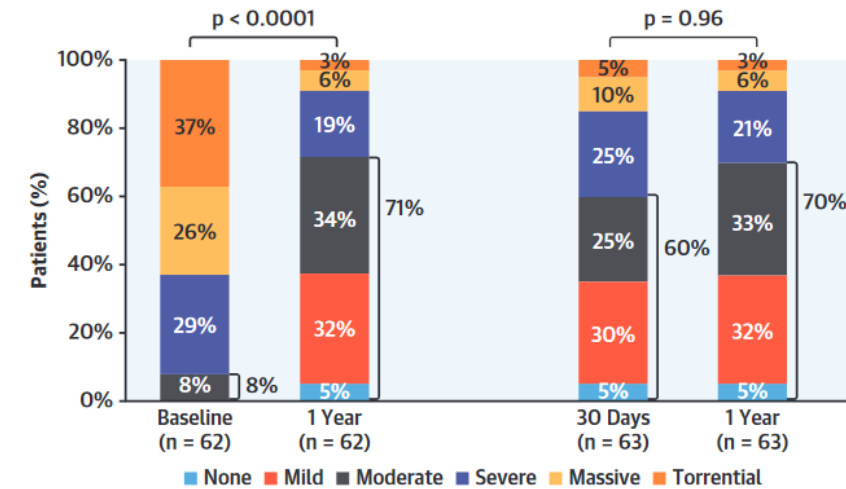
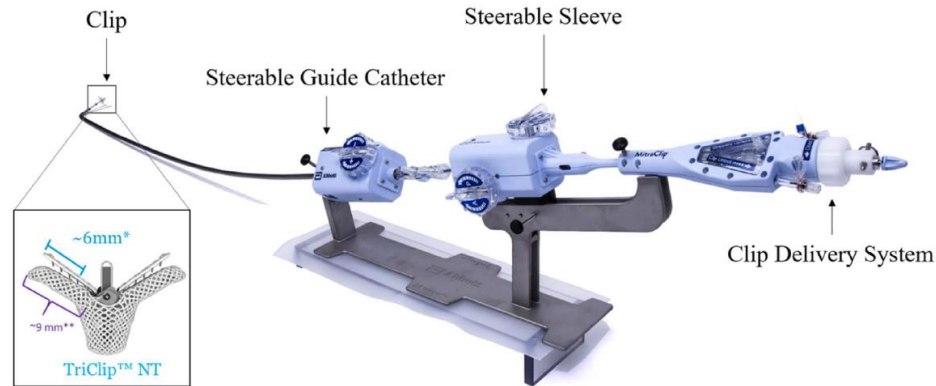


**Population:** IT fonctionnelle sévère (EROA=0.79±0.51mm<sup>2</sup>)

**Résultats:** Succès d'implantation 100%, mortalité péri-procédure: 6,7%  
Amélioration clinique (82% vs. 17% en classe NYHA II à 2 ans)

**Limites**  
Durée de procédure : **4,2±1,5 heures** (16 vis en moyenne) - **ETO+++**  
Efficacité modérée sur la fuite [**24% grade ≤II**, à 1 an]

# Réparation bord à bord: triclip



**Population:** N=85 patients IT fonctionnelle modérée à sévère (EROA=0.65±0.29 mm<sup>2</sup>)

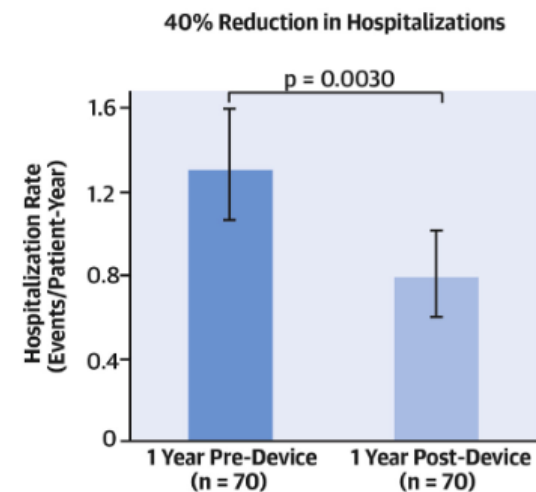
**Résultats:** 5% de mortalité à 1 mois (non liée à la procédure) 90% Succès implantation –  
Amélioration clinique soutenue (80% vs. 25% en classe NYHA I-II à 1 mois et un an)

## Limites

**Durée de procédure :** 2,6±1 heures (67% 1-2 clips -77% sur la commissure AS)

Qualité imagerie ETO +++

Efficacité modérée sur la fuite [37% grade ≤II, 70% ≤3]

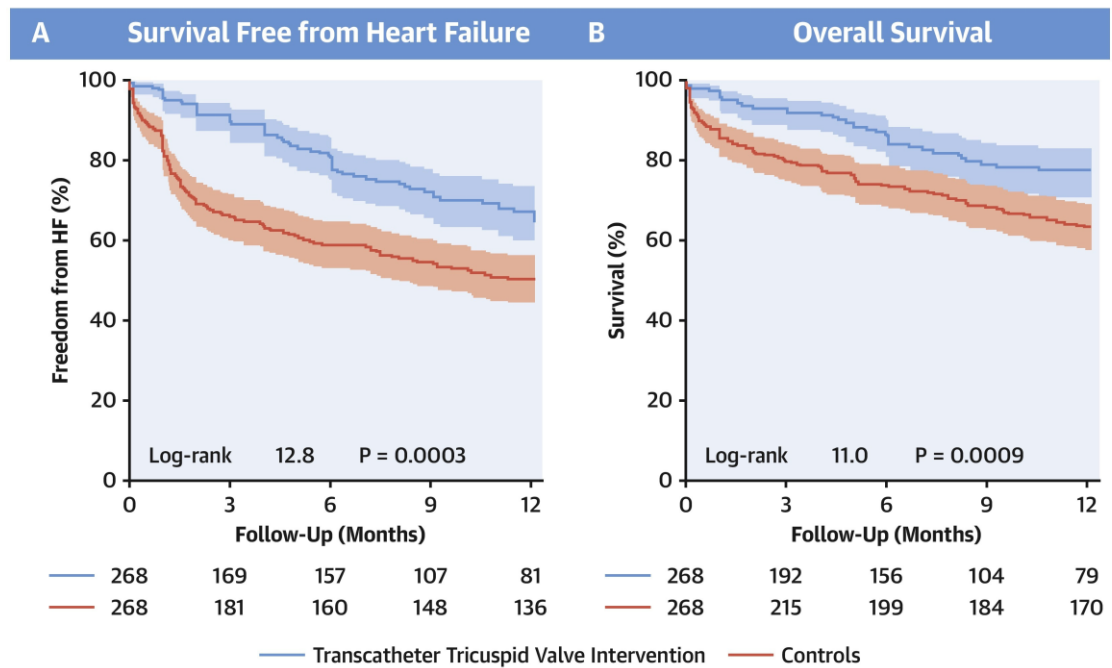


# Réparation bord à bord: clip



## Transcatheter Versus Medical Treatment of Patients With Symptomatic Severe Tricuspid Regurgitation

### CENTRAL ILLUSTRATION: Transcatheter Treatment of Severe Tricuspid Regurgitation: Primary and Secondary Endpoints



Taramasso, M. et al. J Am Coll Cardiol. 2019;74(24):2998-3008.

Registre TTVR vs. contrôles appariés  
229/268 mitraclip (85%)

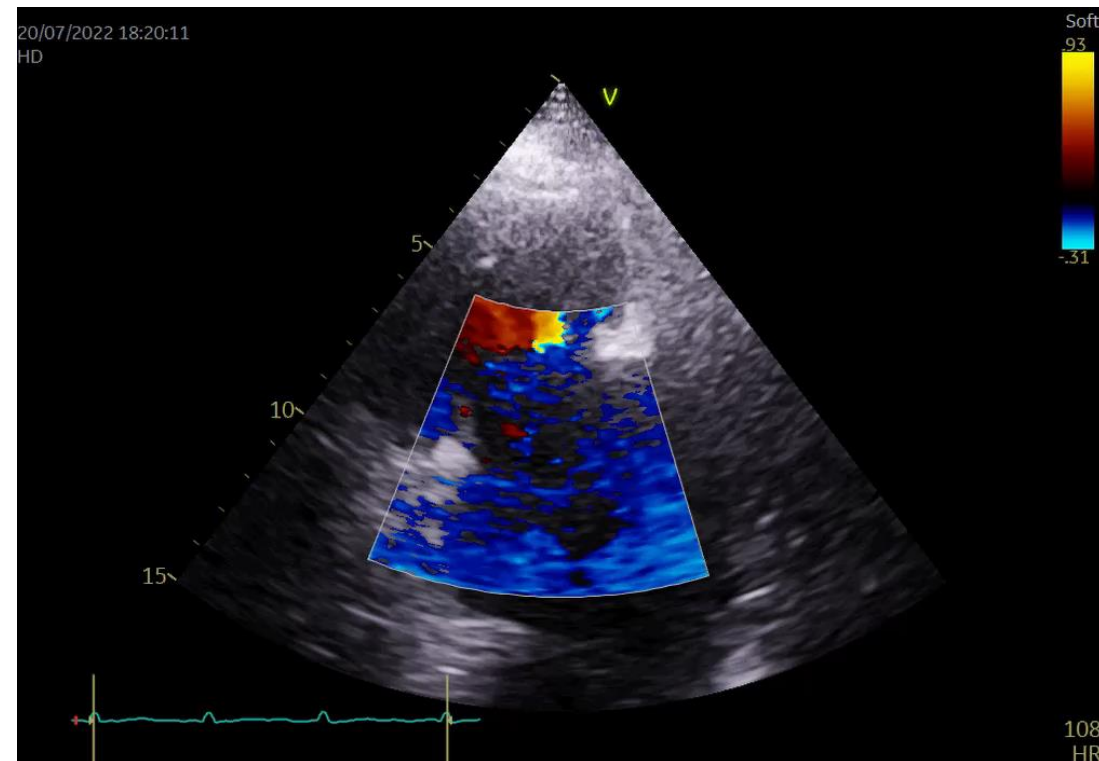
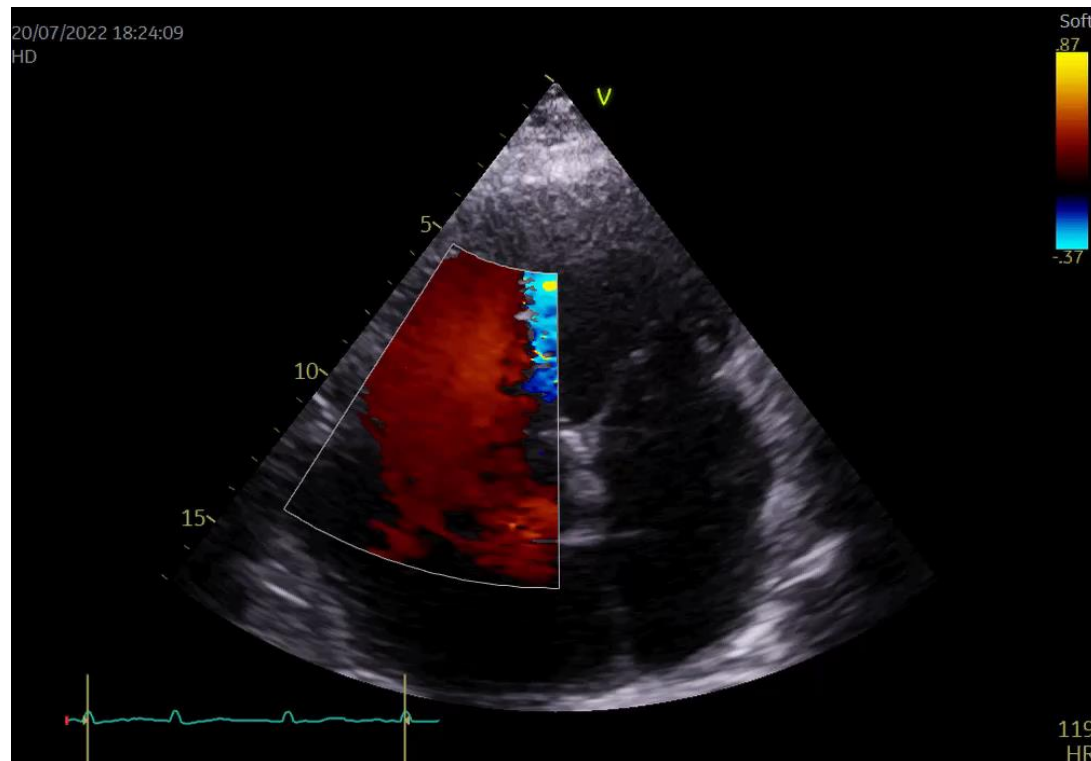
# Réparation bord à bord: clip

Femme 75 ans

FA permanente depuis 2010

Cardiopathie rythmique avec dilatation majeure de l'OD et fuite tricuspide torrentielle

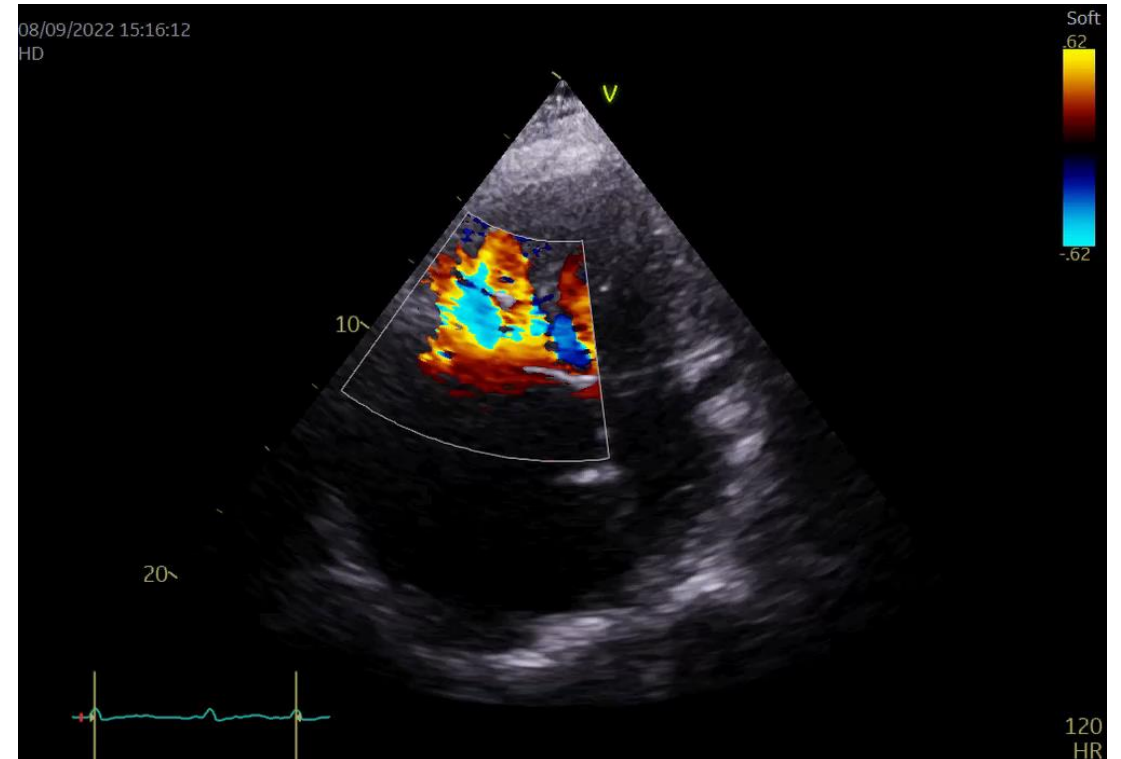
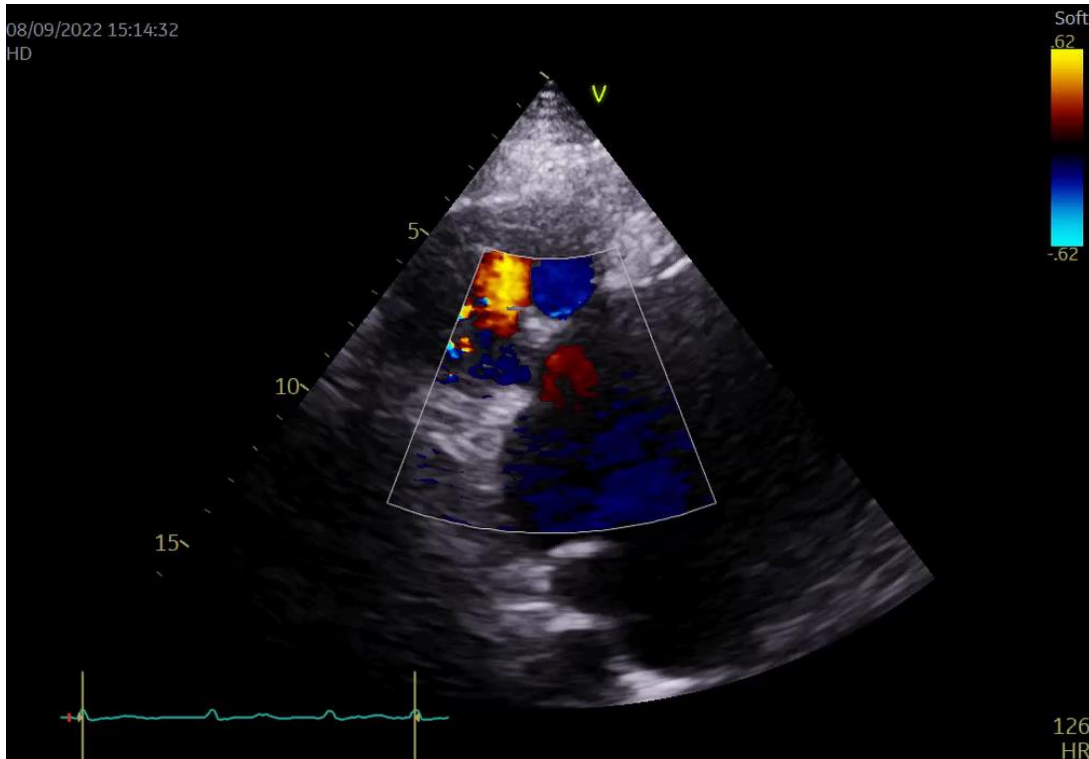
Décision de réparation percutanée par Mitraclip





# Réparation bord à bord: clip

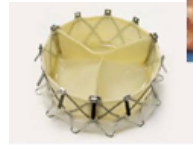
Après 2 mitraclips implantés en position tricuspide entre les feuillets antérieur et septal





# Différentes possibilités

Remplacement



Navigate



Intrepid



Lux

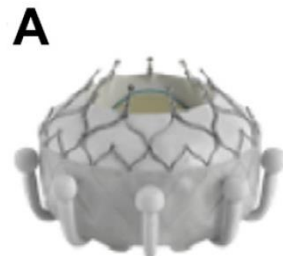


Tricare



Evoque

# EVOQUE (EDWARDS)



A  
Deux tailles 44mm (32%) 48 (64%) mm



B  
28F – femoral access

**Résultats:** 92% Succès implantation – **0% de mortalité péri-procédure**  
 Amélioration clinique (76% vs.4% en classe NYHA I-II à 1 mois)  
**Efficacité sur la fuite [92% grade ≤I]**

**Limites**

Durée de procédure : 2,3±2,5 heures

8% de BAV III

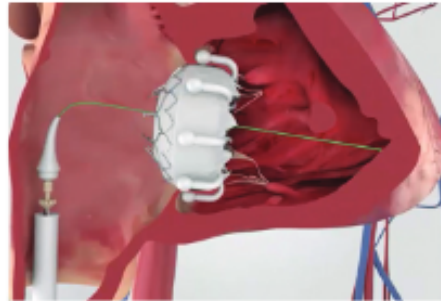
**Taille des prothèses**

**TABLE 4 Clinical Events Committee-Adjudicated Safety Events at 30 Days (N = 56)**

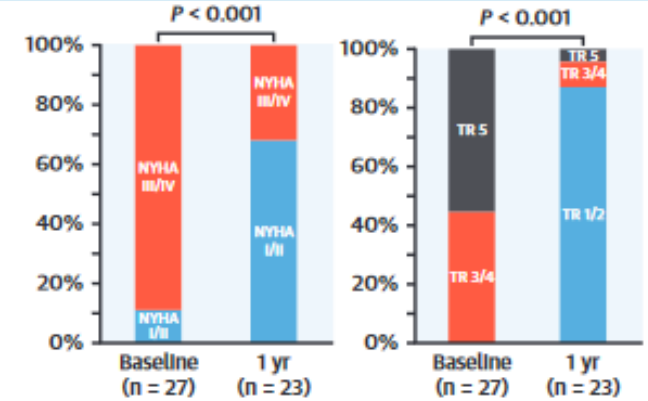
Cardiovascular mortality	1 (1.8) <sup>a</sup>
Myocardial infarction	0
Stroke	0
Renal complications requiring dialysis or renal replacement therapy	0
New need for renal replacement therapy	0
Severe bleeding <sup>b</sup>	15 (26.8)
Fatal	0
Life-threatening	0
Extensive	7 (12.5)
Major	8 (14.3)
Nonelective tricuspid valve reintervention	2 (3.6)
Major access site and vascular complications requiring intervention	1 (1.8)
Major cardiac structural complications	0
Device-related pulmonary embolism	0
Composite MAE rate	15 (26.8)

# EVOQUE

## EVOQUE Transfemoral Tricuspid Replacement 1-Year Clinical and Echocardiographic Outcomes



1-Year Follow-Up

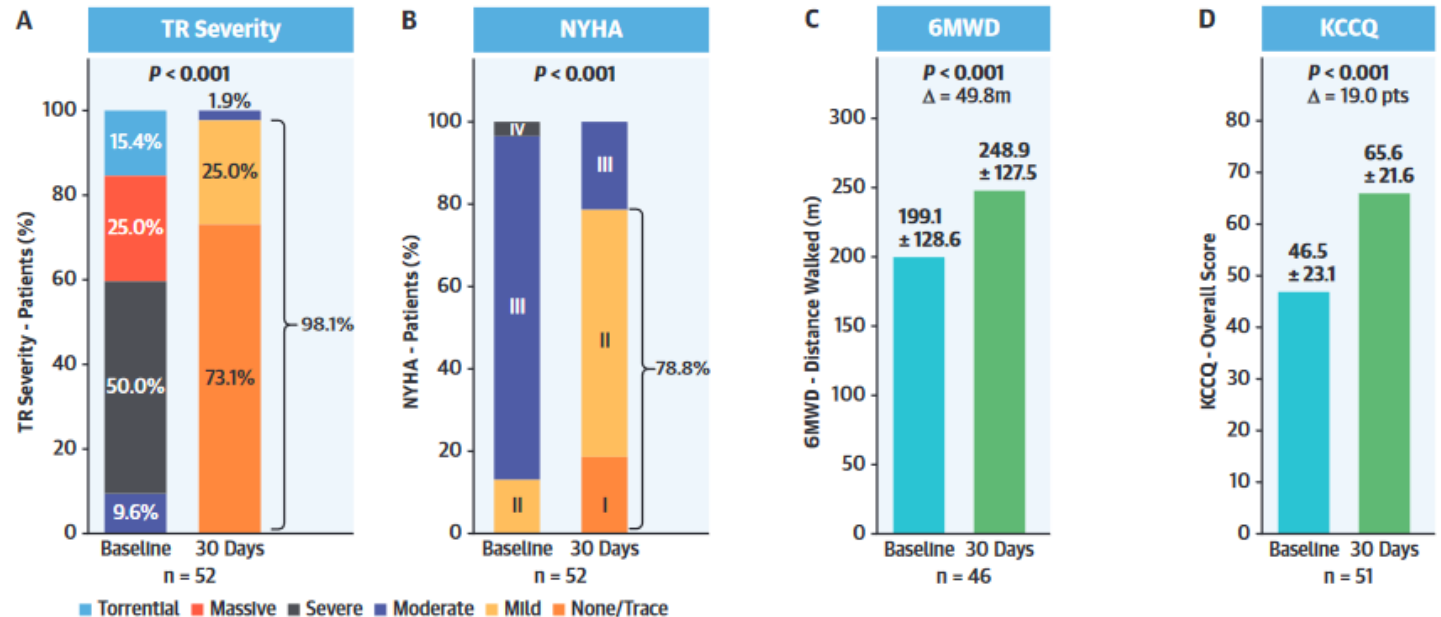


27 patients with severe TR treated with the EVOQUE system  
7 sites (Canada, Europe, U.S.)  
May 2019 to July 2020

All-cause mortality: 7%  
HF hospitalization: 7%  
New pacemaker: 7% within 30 days,  
4% beyond 30 days

Sustained Improvement in NYHA functional class as well as Improvement in TR degree suggesting that the EVOQUE System is a promising treatment option for this population

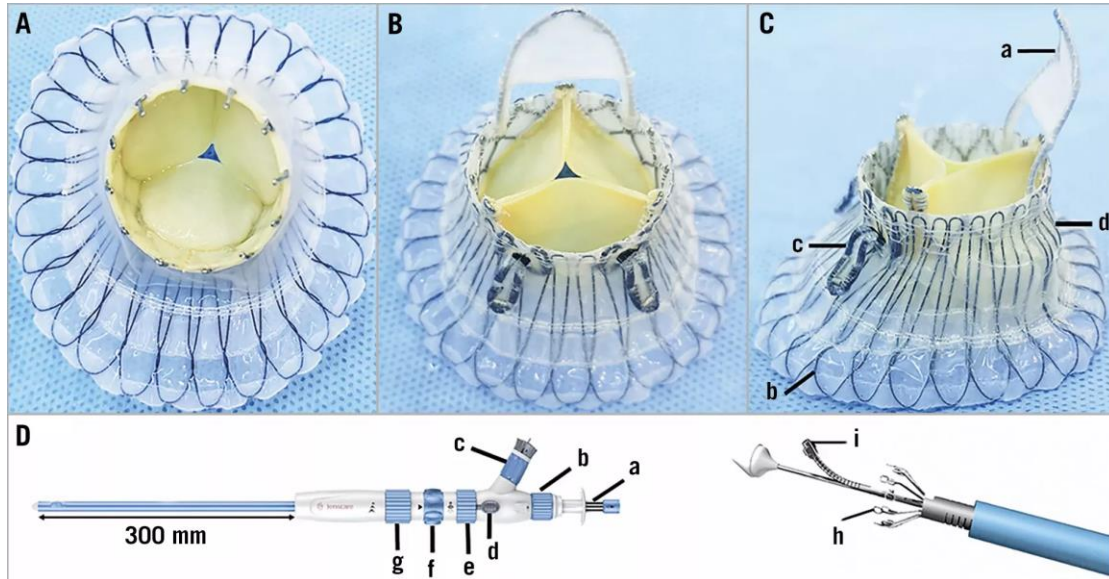
Webb, J.G. et al. J Am Coll Cardiol Intv. 2022;15(5):481-491.



Kodali et al. TRISCEND Study, JACC CI 2022



# LuX-Valve: 1-month



**Table 2. Clinical outcome at 30-day follow-up (n=6).**

Mortality	0 (0)
Myocardial infarction	0 (0)
Stroke	0 (0)
Convention to surgery	0 (0)
Heart failure hospitalisation	0 (0)
Major bleeding	1 (17%)
Conduction abnormality requiring pacemaker	0 (0)
Dialysis	0 (0)

# LuX-Valve: 12-months

**Table 3. Baseline and follow-up echocardiographic parameters.**

Baseline echo parameters		Patient #1	Patient #2	Patient #3	Patient #4	Patient #5	Patient #6
LAESVI, ml/s <sup>2</sup>		1,068.9	401.3	159.7	94.9	89.3	306.3
LVEDVI, ml/s <sup>2</sup>		97.1	56.6	45.6	83.2	56.4	68.0
LVEF, %		60	60	64	56	70	55
RAESVI, ml/s <sup>2</sup>		167.3	145.7	75.1	126.0	319.5	330.5
RVEDAI, cm <sup>2</sup> /s <sup>2</sup>		14.3	16.5	13.7	16.1	24.6	28.9
TAPSE, cm		1.5	1.8	2.0	1.2	1.3	1.1
RV fractional area change, %		25	45	47	23	22	48
RV global longitudinal strain, %		-13.7	-10.2	-12.6	-15.7	-14.9	-12.8
Mean transvalvular gradient, mmHg		2.2	4.6	3.7	3.3	6.6	4.5
TR defining parameters	EROA, cm <sup>2</sup>	1.0	0.7	1.1	0.9	1.2	0.8
	VC width, mm	11	8	13	10	16	9
	IVC, mm	28	24	29	28	30	32
TR grade		Torrential	Massive	Torrential	Torrential	Torrential	Torrential
TV annular diameter (4Ch, end-diastole), cm		4.0	4.1	4.1	4.6	4.7	4.4
Minimum/maximum diameters of TV annular (3D TEE, end-diastolic), cm		3.8x4.1	4.0x4.3	4.1x4.2	4.1x4.6	4.1x4.7	4.0x4.5
Baseline CT annular measurements (end-diastolic)	Minimum/maximum diameters, cm	3.9x4.3	3.9x4.4	4.0x4.2	4.2x4.8	4.3x5.0	4.1x4.7
	LuX-Valve model implanted	JS/TTVI-28-50	JS/TTVI-28-50	JS/TTVI-28-50	JS/TTVI-28-55	JS/TTVI-28-55	JS/TTVI-28-55
Release time of the guiding sheath, min		10.7	6.2	8.5	9.8	13.6	9.1
Intraprocedural, post-device TEE	Tricuspid regurgitation, total	Moderate	No	No	Mild	Mild	Mild
	Central	No	No	No	No	No	No
	Paravalvular	Moderate	No	No	Mild	Mild	Mild
One-year follow-up, post-device TTE	RAESVI, ml/s <sup>2</sup>	/	85.9	56.4	93.3	105.4	152.0
	RVEDAI, cm <sup>2</sup> /s <sup>2</sup>	/	9.7	8.3	11.5	19.8	17.7
	TAPSE, cm	/	2.0	1.8	1.7	1.5	1.9
	RV fractional area change, %	/	47	39	32	26	40
	RV global longitudinal strain, %	/	-17.5	-19.3	-22.1	-20.4	-18.2
	Mean transvalvular gradient, mmHg	/	3.1	2.4	1.8	4.1	2.2
	Tricuspid regurgitation, total	/	No	No	Mild	Mild	Mild
	Central	/	No	No	No	No	No
	Paravalvular	/	No	No	Mild	Mild	Mild

CT: computed tomography; EROA: effective regurgitant orifice area; IVC: inferior vena cava; LAESVI: left atrium end-systolic volume index; LVEDVI: left ventricular end-diastolic volume index; LVEF: left ventricular ejection fraction; RAESVI: right atrium end-systolic volume index; RV: right ventricular; RVEDAI: right ventricular end-diastolic area index; TAPSE: tricuspid annular plane systolic excursion; TEE: transoesophageal echocardiography; TR: tricuspid regurgitation; TV: tricuspid valve

Release time of the guiding sheath, min		10.7	6.2	8.5	9.8	13.6	9.1
Intraprocedural, post-device TEE	Tricuspid regurgitation, total	Moderate	No	No	Mild	Mild	Mild
	Central	No	No	No	No	No	No
	Paravalvular	Moderate	No	No	Mild	Mild	Mild
One-year follow-up, post-device TTE	RAESVI, ml/s <sup>2</sup>	/	85.9	56.4	93.3	105.4	152.0
	RVEDAI, cm <sup>2</sup> /s <sup>2</sup>	/	9.7	8.3	11.5	19.8	17.7
	TAPSE, cm	/	2.0	1.8	1.7	1.5	1.9
	RV fractional area change, %	/	47	39	32	26	40
	RV global longitudinal strain, %	/	-17.5	-19.3	-22.1	-20.4	-18.2
	Mean transvalvular gradient, mmHg	/	3.1	2.4	1.8	4.1	2.2
	Tricuspid regurgitation, total	/	No	No	Mild	Mild	Mild
	Central	/	No	No	No	No	No
	Paravalvular	/	No	No	Mild	Mild	Mild

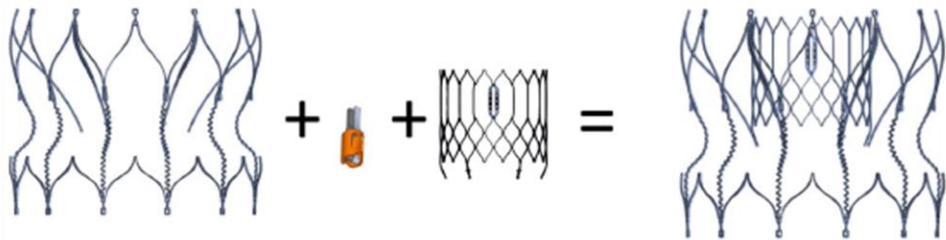
CT: computed tomography; EROA: effective regurgitant orifice area; IVC: inferior vena cava; LAESVI: left atrium end-systolic volume index; LVEDVI: left ventricular end-diastolic volume index; LVEF: left ventricular ejection fraction; RAESVI: right atrium end-systolic volume index; RV: right ventricular; RVEDAI: right ventricular end-diastolic area index; TAPSE: tricuspid annular plane systolic excursion; TEE: transoesophageal echocardiography; TR: tricuspid regurgitation; TV: tricuspid valve

**Twelve-month outcomes of the LuX-Valve for transcatheter treatment of severe tricuspid regurgitation**

EuroIntervention 2021;17:818-826



# TRICARES

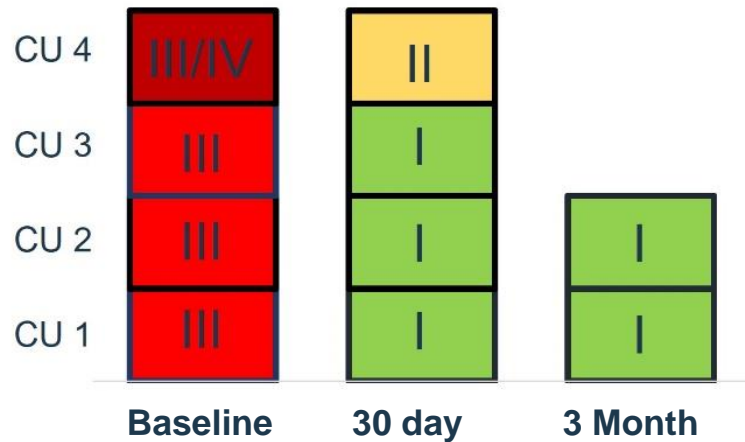


- stent très souples permettant de suivre les variations géométriques du VD
- Dé-corrélation de la fonction ancrage et valvulaire



# Topaz TTVR - Early Compassionate Use Experience

NYHA Classification Over Time



Procedure Discharge	CU 1	CU 2	CU 3	CU4
Procedure Time*	18 min	12 min	20 min	48 min
NYHA Class Discharge	I	I	I	II
Tricuspid Regurgitation (Grade 1-5)	None (0)	None (0)	None (0)	Mild (1)
Discharged on POD	4	4	15	15

\* Definition: Time from Topaz delivery system in, followed by valve deployment to delivery system out

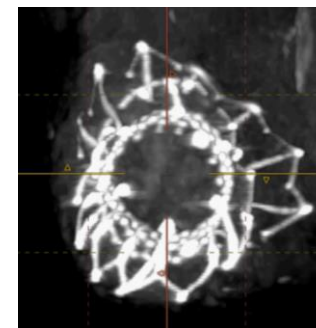
CU 1  
post deployment



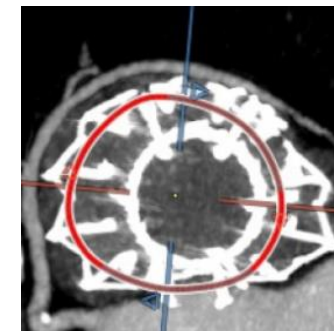
CU 2  
post deployment



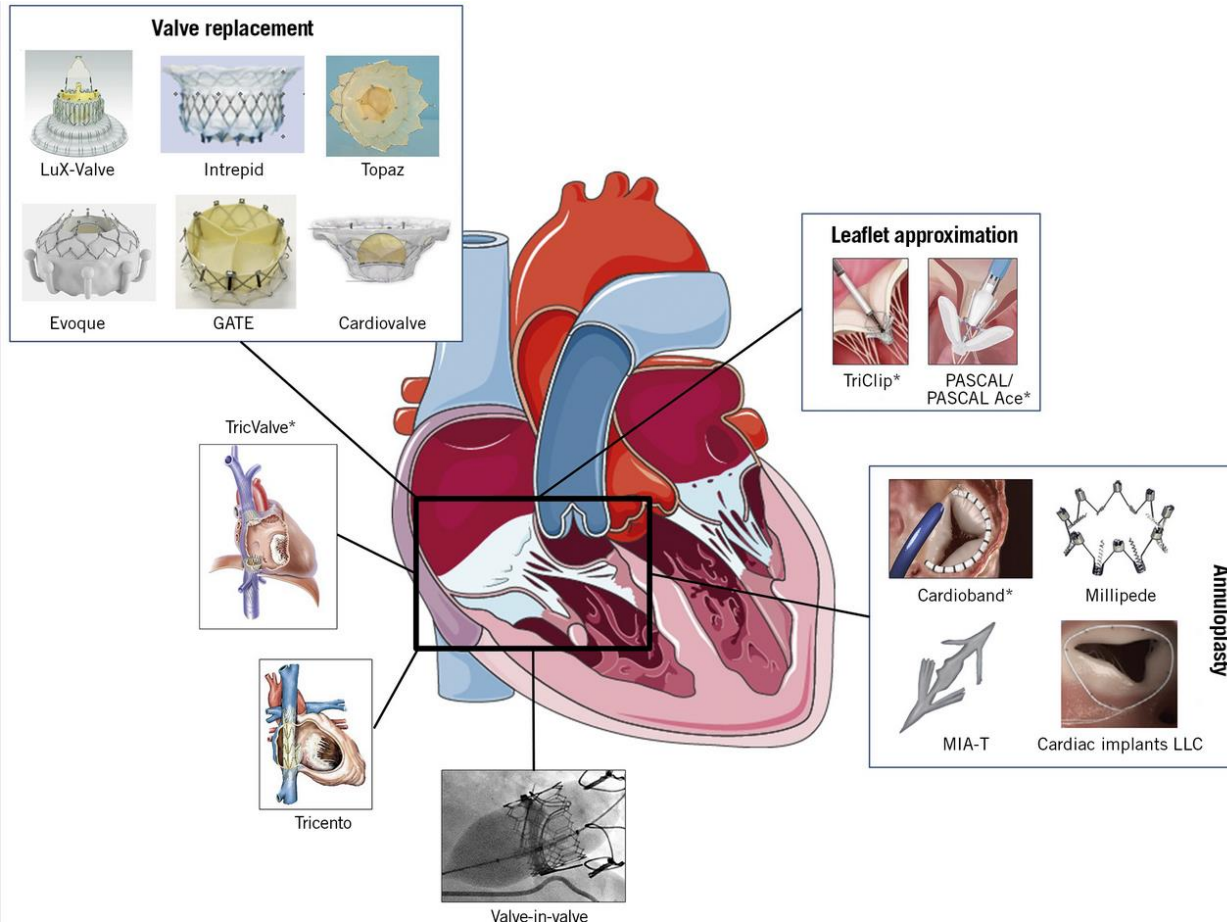
CU 1  
CT 3 Mon FUP



CU 2  
CT 3 Mon FUP



# Algorithme décisionnel



**Table 3. Anatomical criteria for device selection.**

Strategy	Favourable anatomy	Feasible anatomy	Unfavourable anatomy
Leaflet approximation	<ul style="list-style-type: none"> <li>Small septolateral gap <math>\leq 7</math> mm<sup>10</sup></li> <li>Anteroseptal jet location</li> <li>Confined prolapse or flail</li> <li>Trileaflet morphology</li> </ul>	<ul style="list-style-type: none"> <li>Septolateral coaptation gap <math>&gt; 7</math> but <math>\leq 8.5</math> mm<sup>65</sup></li> <li>Posteroseptal jet location</li> <li>Non-trileaflet morphology</li> <li>Incidental CIED RV lead (i.e., without leaflet impingement)</li> </ul>	<ul style="list-style-type: none"> <li>Large septolateral coaptation gap <math>&gt; 8.5</math> mm<sup>65</sup></li> <li>Leaflet thickening/shortening (rheumatic, carcinoid)/perforation</li> <li>Dense chordae with marked leaflet tethering</li> <li>Anteroposterior jet location</li> <li>Poor echocardiographic leaflet visualisation</li> <li>CIED RV lead leaflet impingement</li> <li>Unfavourable device angle of approach</li> </ul>
Annuloplasty	<ul style="list-style-type: none"> <li>Annular dilatation as primary mechanism of TR</li> <li>Mild tethering (tethering height <math>&lt; 0.76</math> cm, tenting area <math>&lt; 1.63</math> cm<sup>2</sup>, tenting volume [3D] <math>&lt; 2.3</math> mL)<sup>110,111</sup></li> <li>Central jet location</li> <li>Sufficient landing zone for anchoring</li> </ul>	<ul style="list-style-type: none"> <li>Moderate tethering (tethering height <math>\geq 0.76</math> cm but <math>&lt; 1.0</math> cm, tenting area <math>&gt; 1.63</math> but <math>&lt; 2.5</math> cm<sup>2</sup>, tenting volume [3D] <math>\geq 2.3</math> mL but <math>\leq 3.5</math> mL)<sup>110,111</sup></li> <li>Incidental CIED RV lead (i.e., without leaflet impingement)</li> </ul>	<ul style="list-style-type: none"> <li>Excessive annular dilatation (exceeding device size)</li> <li>Severe tethering (tethering height <math>&gt; 1.0</math> cm, tenting volume <math>&gt; 3.5</math> mL). Poor echocardiographic annular visualisation<sup>110,111</sup></li> <li>Annular proximity of RCA</li> <li>CIED RV lead leaflet impingement</li> </ul>
Orthotopic valve implantation	<ul style="list-style-type: none"> <li>Previous surgical repair or bioprosthetic valve replacement</li> <li>Leaflet thickening/shortening (rheumatic, carcinoid)</li> <li>Incidental CIED RV lead (i.e., without leaflet impingement)</li> <li>Any leaflet morphology</li> </ul>	<ul style="list-style-type: none"> <li>Large coaptation gap</li> <li>CIED RV lead leaflet impingement</li> </ul>	<ul style="list-style-type: none"> <li>Excessive annular dilatation (exceeding device size)</li> <li>Unfavourable device angle of approach</li> <li>Severe right ventricular dysfunction</li> </ul>
Heterotopic valve implantation	<ul style="list-style-type: none"> <li>Appropriate caval diameters (and intercaval distance)</li> <li>No option for direct valve treatment</li> </ul>		<ul style="list-style-type: none"> <li>Proximity of the RA to the orifice of the liver veins (<math>&lt; 10</math>-<math>12</math> mm)</li> <li>Severely increased pulmonary artery and RA pressures due to the risk of fracture of bicaval valved stents</li> </ul>

3D: three-dimensional; CIED: cardiac implantable electronic device; RA: right atrium; RCA: right coronary artery; RV: right ventricular; TR: tricuspid regurgitation

## State of the Art

[Transcatheter treatment for tricuspid valve disease](#) Praz F. et al EUROINTERVENTION

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# Conclusion



- Réparation: TRICLIP, PASCAL, Annuloplastie
  - Exclusions anatomiques: fréquentes
  - Durée de procédure: longue à très longue
  - Dépendance à l'imagerie per-procédure: importante
  - Résultats incomplets ?: à valider sur les études en cours
- Valves percutanées
  - Exclusion anatomique: moins fréquente (évolution taille des dispositifs)
  - Durée de procédure: courte
  - Dépendance à l'imagerie: faible
  - Résultats semble-t-il plus complets