



**UZ
LEUVEN**



LVAD and the failing right heart

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Introduction

- Right heart failure in LVAD patients

Pre-operative



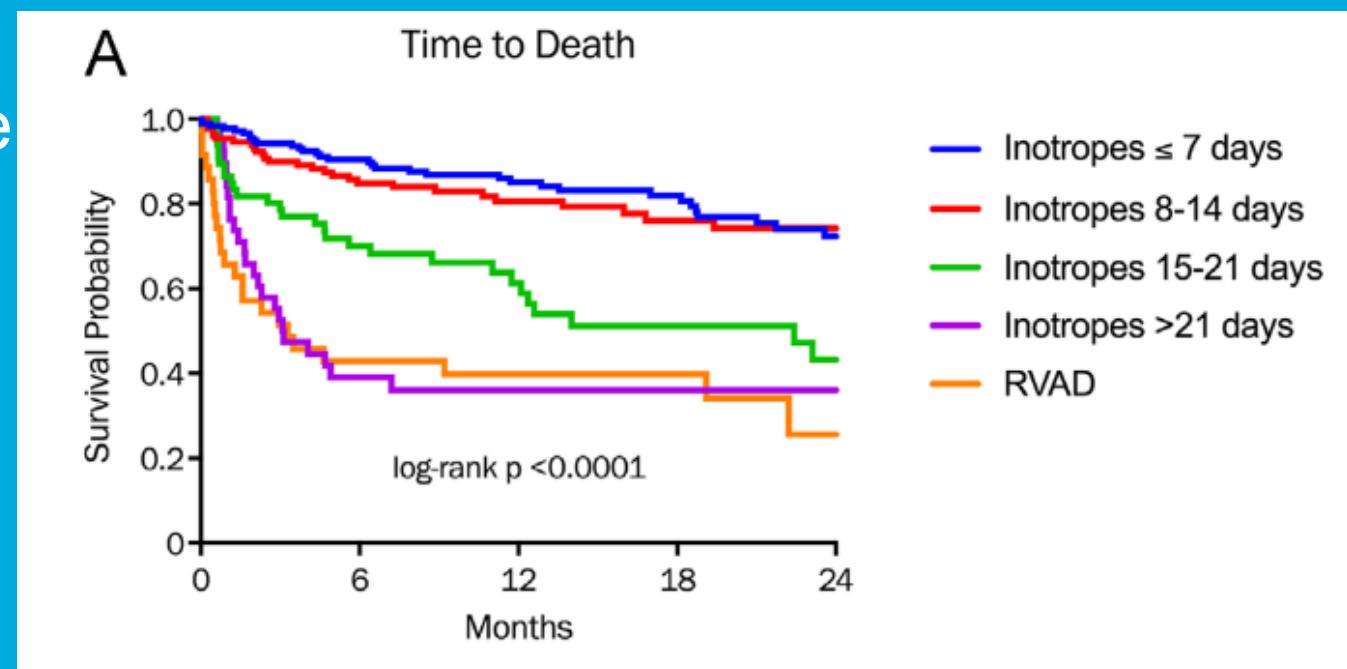
Peri-operative

Post-operative



Incidence of peri-operative RV failure

- Peri-operative: 5 – 46%
 - Mild
 - Moderate
 - Severe



LaRue S, et al. Clinical outcomes associated with INTERMACS-defined right heart failure after left ventricular assist device implantation J Heart Lung Transplant 2017; 36(4):475-477.

Definition of RHF after LVAD

TABLE 2 | 2014 INTERMACS definition for right heart failure.

Definition: Symptoms or findings of persistent right heart failure characterized by BOTH of the following:

1. Documentation of elevated central venous pressure by:
 - Direct measurement with right atrial pressure > 16 mmHg OR
 - Findings of significantly dilated inferior vena cava with absence of inspiratory variation by echocardiography OR
 - Clinical findings of elevated jugular distention at least halfway up the neck in an upright patient
2. Manifestations of elevated central venous pressure characterized by:
 - Clinical findings of peripheral edema (>2+ either new or unresolved) OR
 - Presence of ascites or palpable hepatomegaly on physical examination or diagnostic imaging OR
 - Laboratory evidence of worsening hepatic congestion (total bilirubin > 2.0 mg/dl) or renal dysfunction (creatinine > 2.0 mg/dl)
3. If the patient meets definition of both criteria above then a severity scale for right heart failure will be graded utilizing post implant inotropes, inhaled nitric oxide or intravenous vasodilators, need for right ventricular assist device and timing from surgery.

Definition of RHF after LVAD

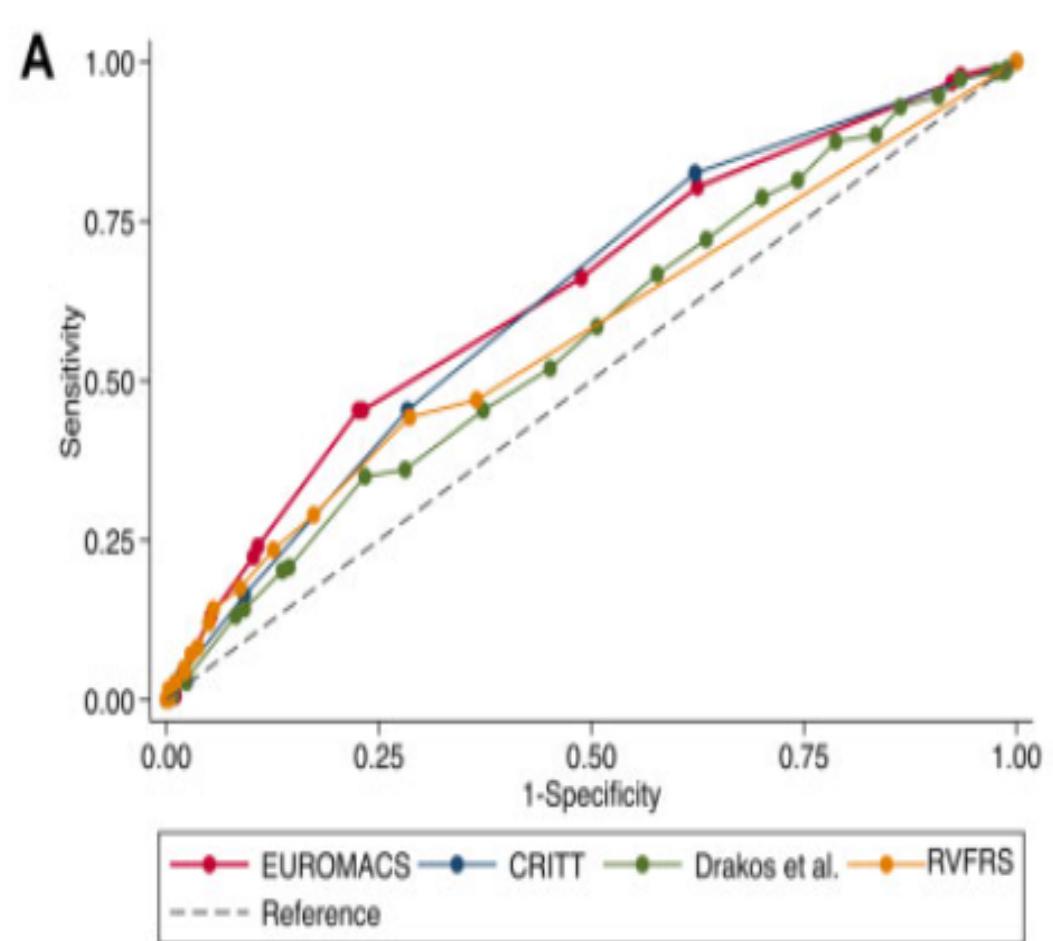
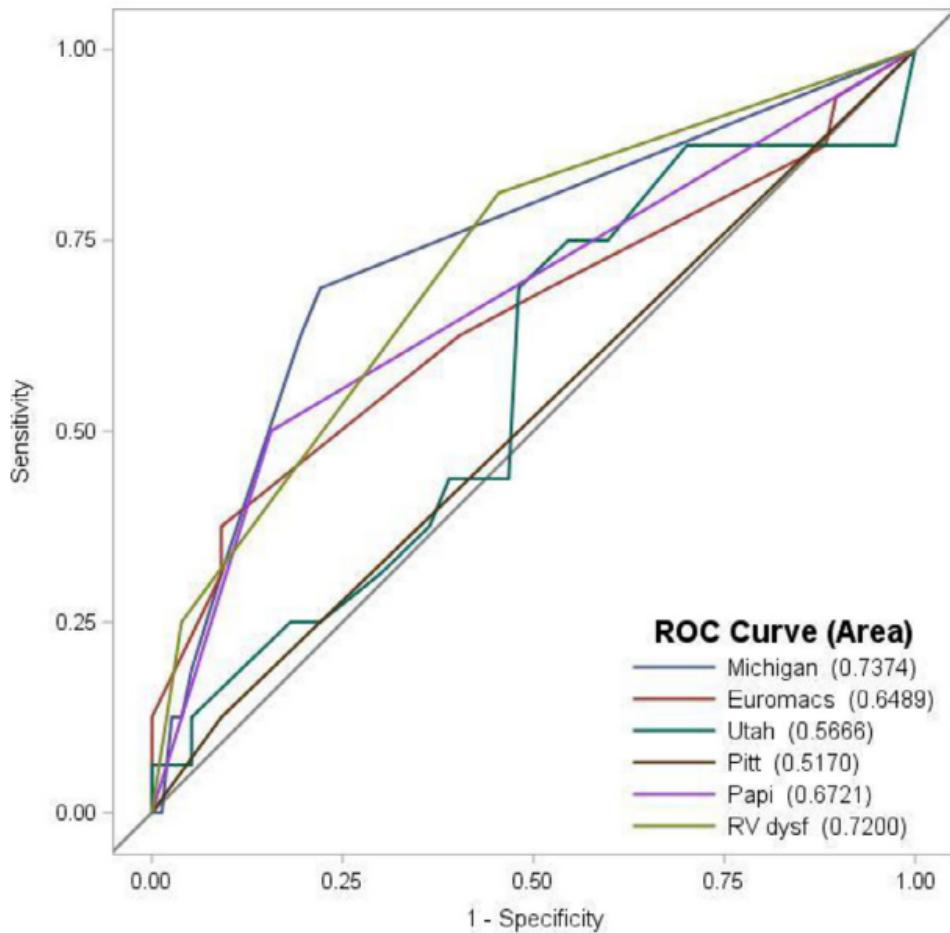
TABLE 3 | 2020 Academic Research Consortium definition for right heart failure.

Early acute right heart failure	Early post-implant right heart failure
Need for implantation of right ventricular assist device at time of left ventricular assist device implantation	<p>(A) Need for implantation of right ventricular assist device <30 days of left ventricular assist device implantation OR (*B) Failure to wean from inotropic support or inhaled nitric oxide within 14 days following LVAD implantation or having to initiate this support within 30 days of implant for a duration of at least 14 days OR (C) Death occurring in patients within 14 days of LVAD implant who have not received an RVAD but who remain on inotropes or vasopressors at the time of death and meet criteria for the diagnosis of Right Heart Failure on the basis of the above clinical findings</p> <p>* For Criteria B: At least two of the following must be present:</p> <ul style="list-style-type: none"> - Ascites - Peripheral edema (>2+) - Elevated central venous pressure (>16 mmHg) - Elevated jugular venous pressure atleast half way up the neck in an upright patient <p>OR</p> <p>At least one of the following must be present:</p> <ul style="list-style-type: none"> - Renal failure with creatinine > 2 × baseline value - Liver injury with at least 2× upper limit normal in AST/ALT - Total bilirubin > 2.0 - SVO₂ < 50% - Cardiac index < 2.2 liter/min/m² - Elevated lactate > 3.0 mmol/liter - Reduction in pump flow of >30% from previous baseline in absence of cardiac tamponade, tension pneumothorax or other mechanical causes.

Can we predict the RV failure?

- Risk scores:
 - Utah RVF
 - Pittsburgh decision tree
 - Michigan RVF
 - CRITT
 - Penn
 - EUROMACS right-sided heart failure risk score
 - ...

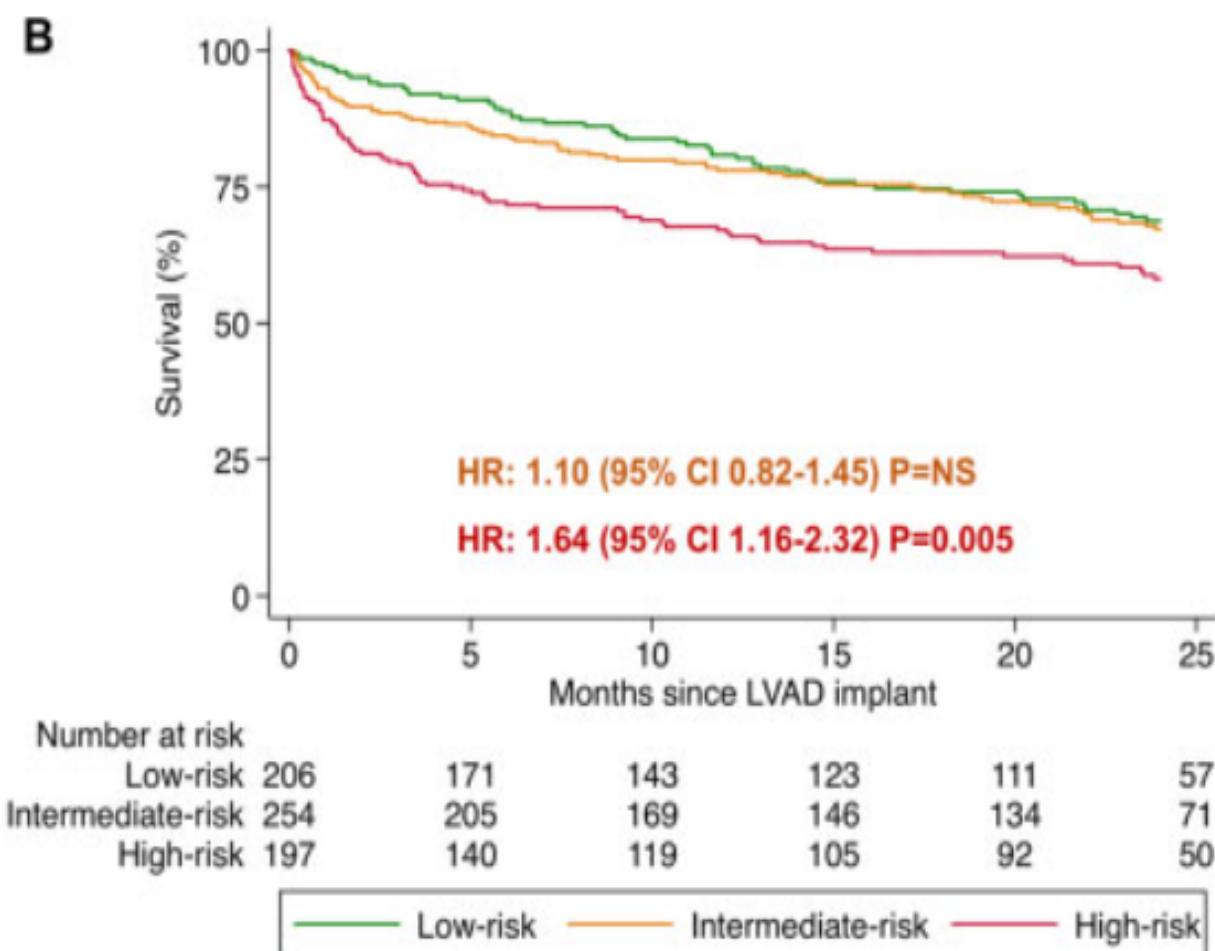
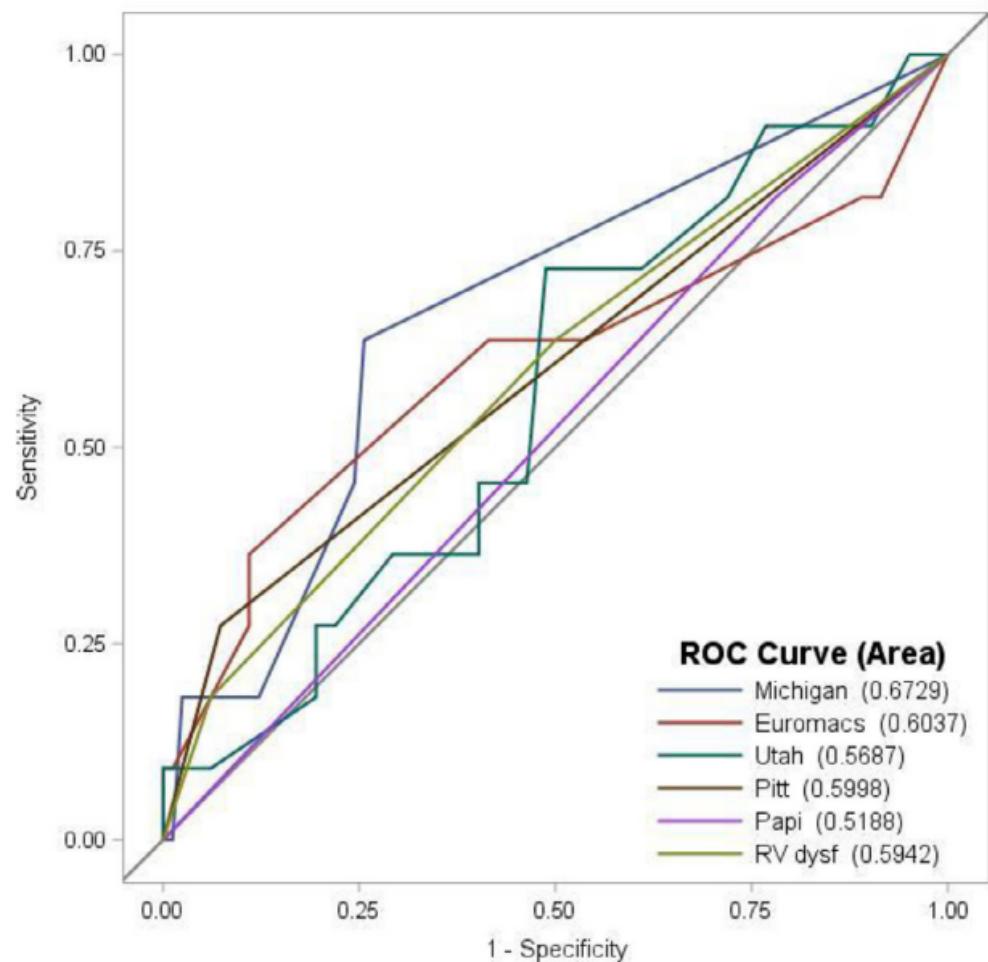
Can we predict the RV failure?



Peters A. et al Comparative Analysis of Established Risk Scores and Novel Hemodynamic Metrics in Predicting Right Ventricular Failure in Left Ventricular Assist Device Patients *J Card Fail* 2019; 25(8):620-628.

Rivas-Lasarte M. et al Prediction of right heart failure after left ventricular assist implantation: external validation of the EUROMACS right-sided heart failure risk score *Eur Heart J Acute Cardiovasc Care* 2021; 10(7):723-732.

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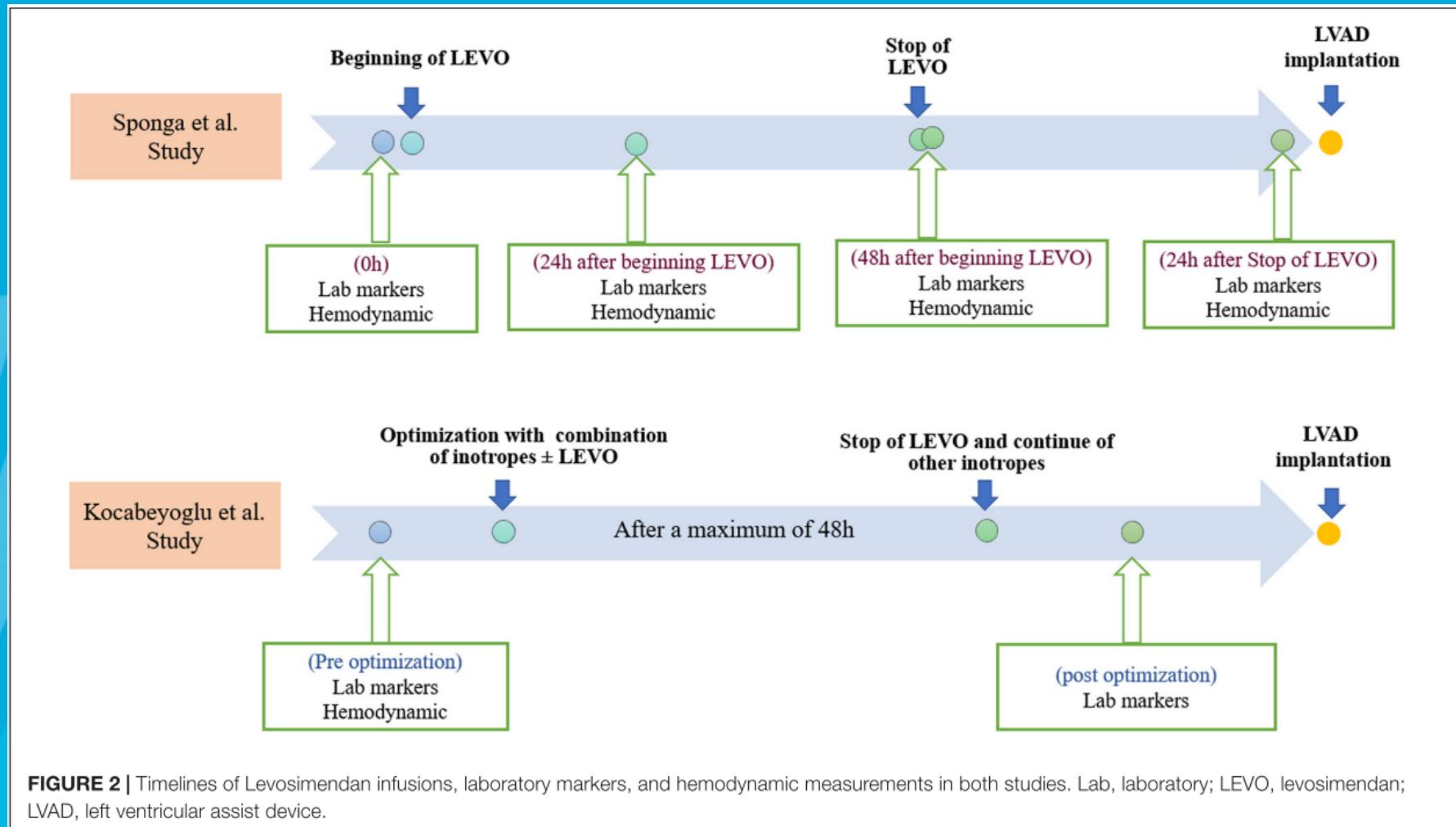
Are there risk factors?

	Model						
	Michigan (RVFRS)	EUROMACS	Penn (Fitzpatrick)	Utah (Drakos)	CRITT	Kormos (Heartmate II)	Pittsburgh Decision Tree
Model variables							
Demographic				Obesity			Age
Preoperative status	Vasopressor requirement	≥3 inotropes	Previous cardiac surgery	Inotrope dependency	Preoperative intubation	Need for ventilatory support	No. of inotropes
		INTERMACS class 1–3		IABP			
Hemodynamic		RAP/PCWP ≥0.54	Cln ≤2.2 L/min per m ²	PVR	CVP >15 mm Hg	CVP/PCWP >0.63	RAP
			RVSWI ≤0.25 mm Hg L/m ²		HR >100 bpm		TPG
			SBP ≤96 mm Hg				HR
Echocardiographic		Severe RV dysfunction	Severe RV dysfunction		Severe RV dysfunction		
					Severe TR		
Laboratory	AST ≥80 UI/L	Hb ≤10 g/dL	Cr ≥1.9 mg/dL			BUN >39 mg/dL	INR
	Bili ≥2 mg/dL						WBC
	Cr ≥2.3 mg/dL						ALT
Device indication				DT			
Medication				ACE inhibitor/ARB			
				β-Blocker			

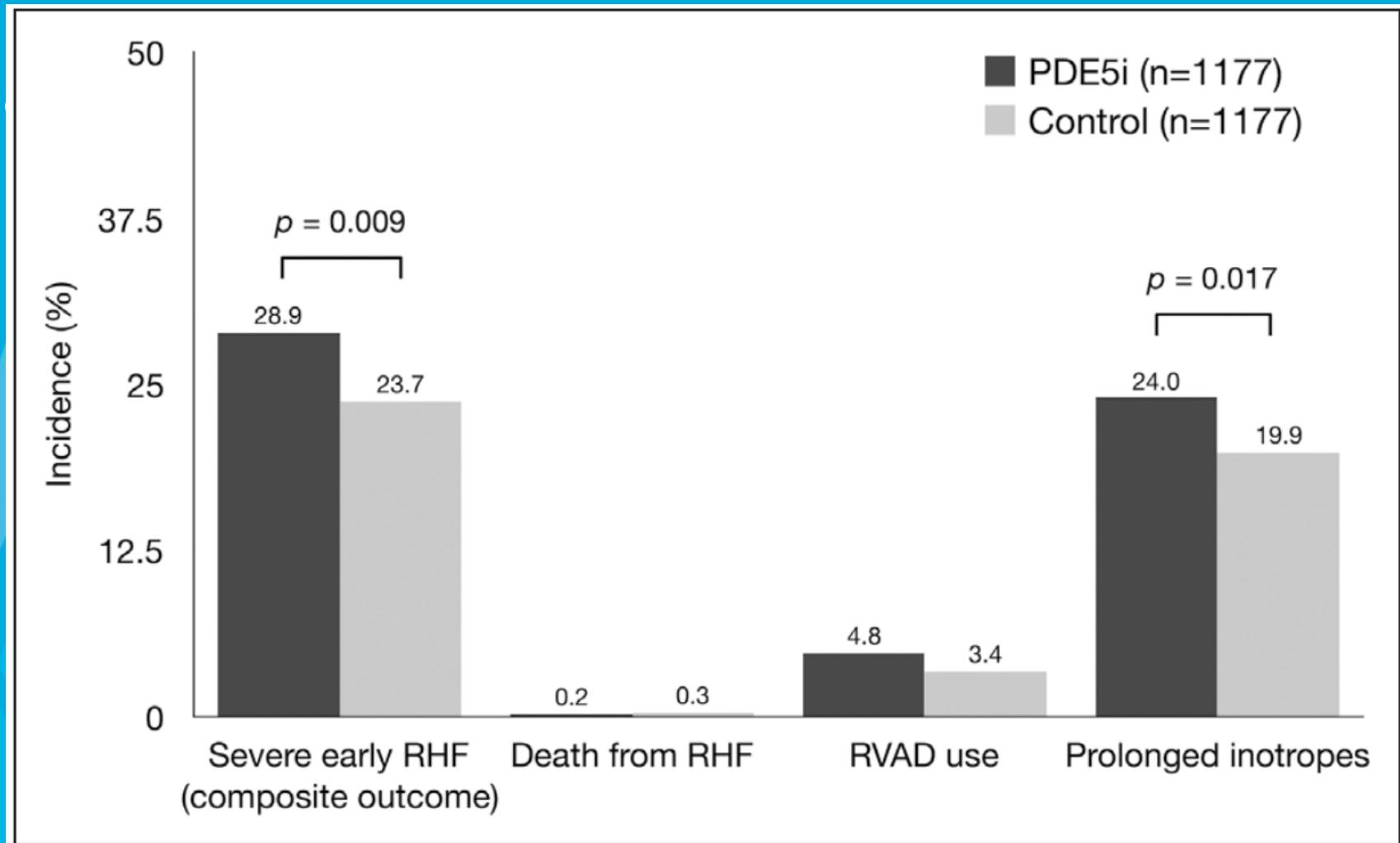
Are there risk factors?

- Higher risk in patients with:
 - Mechanical ventilation, CRRT
 - Higher NT-proBNP, INR, WBC
 - Lower RVSWI, higher CVP

Can we prevent RV failure?

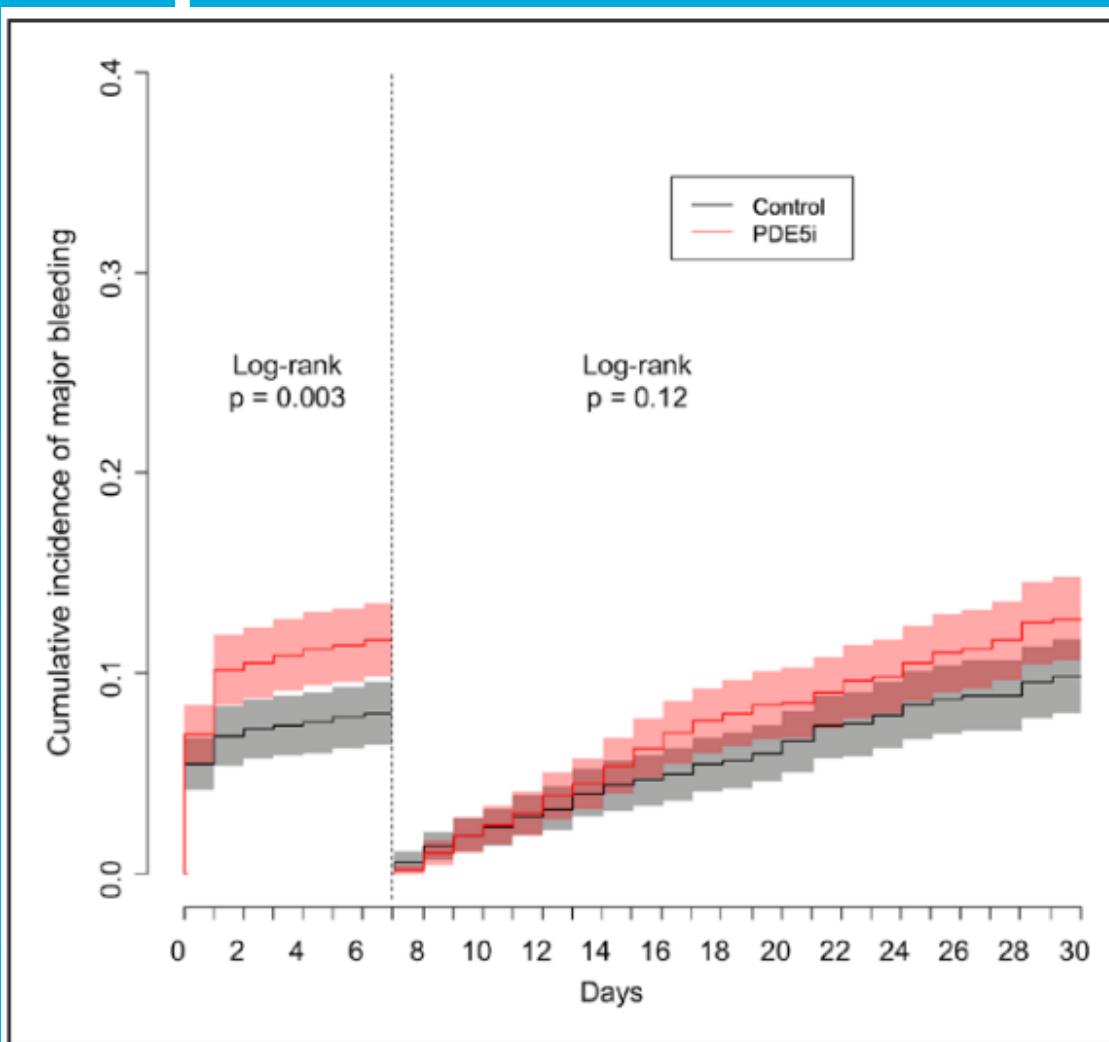


Can we prevent RV failure?



Gulati G. et al Preimplant phosphodiesterase-5 inhibitor use is associated with higher rates of severe early right heart failure after left ventricular assist device implantation an INTERMACS analysis *Circ Heart Fail* 2019 Jun; 12(6):e005537.

Can we prevent RV failure?



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Surgical points of attention

- Use of carbon dioxide
- Carefull de-airing (Trendelenburg)

Surgical points of attention

- Start low, go slow
- Filling state!

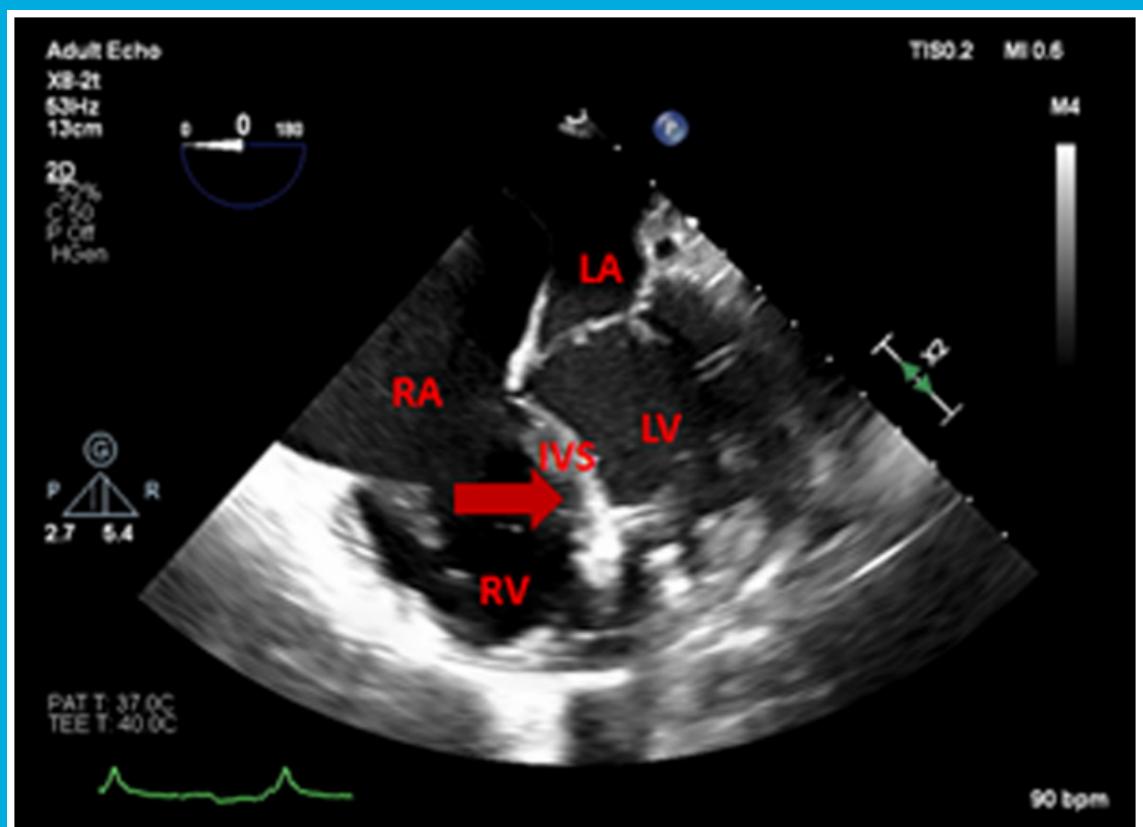


Table 5 Primary and Secondary Outcome Measures in the Intent-to-Treat Population

Outcome Measure	iNO	Placebo	p-value
Patients meeting RVD criteria ≤48 hours			0.330
No. of total (%)	7/73 (9.6)	12/77 (15.6)	
95% CI	2.8–16.3	7.5–23.7	
Males, No. (%)	7/64 (10.9)	7/65 (10.8)	>0.99
Females, No. (%)	0/9 (0.0)	5/12 (41.7)	0.045
PVRI <270.5 dyne/sec/cm ⁻⁵	6/51 (11.8)	6/48 (12.5)	>0.99
PVRI ≥270.5 dyne/sec/cm ⁻⁵	1/7 (14.3)	5/7 (71.4)	0.103
Days on mechanical ventilation ^a	70	67	0.077
Mean (SD)	5.37 (7.72)	11.10 (24.81)	
Median (range)	2.0 (1–30)	3.0 (0–160)	
No. of ICU days ^b	60	58	0.630
Mean (SD)	20.52 (32.31)	19.90 (24.38)	
Median (range)	11.0 (3–194)	9.0 (3–115)	
No. of total hospital days ^c	58	58	0.979
Mean (SD)	40.57 (32.19)	40.76 (29.41)	
Median (range)	32.0 (11–194)	31.5 (10–156)	
Quantity of blood products used	73	77	
Mean, ml (SD)	4,232 (4675)	4,885 (7760)	0.226
Patients requiring RRT, No. (%) ^d	10/71 (14.1)	8/70 (11.4)	0.637
Non-survival at Day 28, No. (%)	8/71 (11.3)	8/70 (11.4)	0.924
Patients needing RVAD by Day 28, No. (%)	4/71 (5.6)	7/70 (10.0)	0.468

ICU, intensive care unit; iNO, inhaled nitric oxide; PVRI, pulmonary vascular resistance index; RRT, renal replacement therapy; RVAD, right ventricular assist device; RVD, right ventricular dysfunction; SD, standard deviation.

^aIntent-to-treat population, extubated before study end (Day 28).

^bIntent-to-treat population, discharged from ICU before study end (Day 28).

^cIntent-to-treat population, discharged from hospital before study end (Day 28).

^dIncludes those patients in the intent-to-treat population for whom RRT information was available.

Can we treat RV failure?

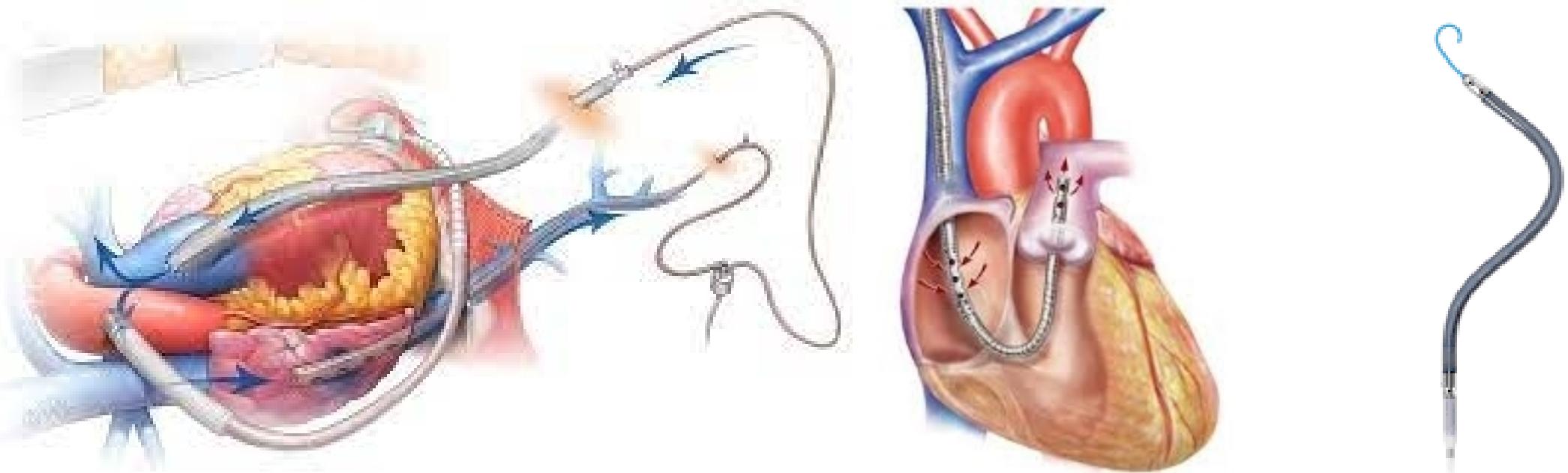
- Drug therapy

Inotrope and vasopressor support

Norepinephrine should be considered as a first-line vasopressor in case of postoperative hypotension or shock.	IIa	B	[9, 328, 329]
Dopamine may be considered in case of postoperative hypotension or shock.	IIb	B	[9, 328, 329]
The combination of norepinephrine and dobutamine should be considered instead of epinephrine in case of postoperative hypotension and low cardiac output syndrome with RV failure.	IIa	C	[9, 71, 330, 331]
Epinephrine may be considered in case of postoperative hypotension and low cardiac output syndrome with RV failure.	IIb	C	
Phosphodiesterase 3 inhibitors may be considered in patients with long-term mechanical circulatory support with postoperative low cardiac output syndrome and RV failure.	IIb	C	[332, 333]
The use of levosimendan in case of postoperative low cardiac output syndrome may be considered.	IIb	A	[334, 335]

— Milrinone

Mechanical RV support



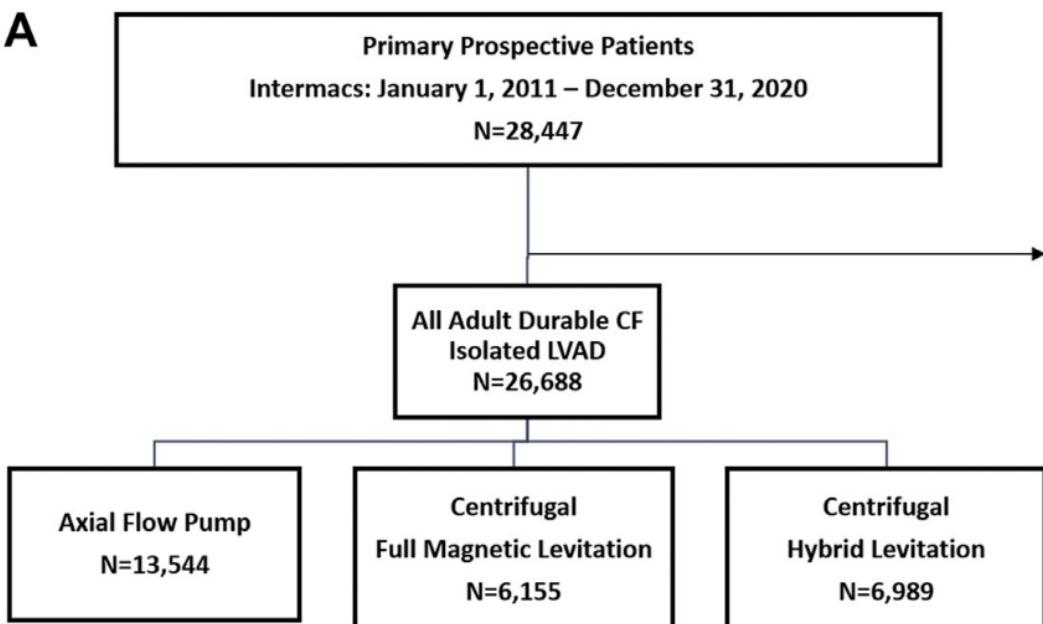
- Long term:
 - BiVAD
 - HTX



Chronic RVAD and LVAD

12th INTERMACS report

A



Exclude:	n
Isolated PF RVAD	19
Isolated CF RVAD	10
TAH	426
Isolated PF LVAD	30
PF BiVAD	138
CF BiVAD	1,136
Total	1,759

PF = pulsatile flow pump
 CF = continuous flow pump
 RVAD: right ventricular assist device
 LVAD: left ventricular assist device
 TAH: total artificial heart
 BiVAD: biventricular assist device

3th EUROMACS report

A total of 6192 implants in
5735 patients

- Before 2011: 473 implants
- Age <18: 397 implants
- Center with <60% follow up: 295
- Excluded centers: 193

A total of 4834 implants in
4486 patients

- | |
|-------------------|
| LVAD: 4304 |
| LVAD + RVAD*: 195 |
| BiVAD: 68 |
| RVAD: 142 |
| SVAD: 3 |
| TAH: 74 |
| Unknown*: 48 |

Leuven experience with RV failure

- BIVAD in Leuven
 - Long term: 0
 - Temporary RVAD: 14/252 (5,6%)
 - Urgent HTX: 4
 - Weaned: 9
 - Death: 1
 - In hospital mortality
 - HTX: 2
 - Weaned: 3

Conclusions

- RV failure is common after LVAD implantation
- RV failure is hard to predict
- A struggling RV benefits from:
 - Careful fluid and LVAD management
 - iNO
 - Vasopressors / inotropes
 - Temporary RV support