

XI Reunión. Estado del Arte en
INSUFICIENCIA CARDIACA

PRÁCTICA CLÍNICA Y MODELOS ORGANIZATIVOS

Sede: Hotel Meliá MaríaPita, A Coruña

A CORUÑA 27-28 SEPTIEMBRE 2024



XI Meeting. State of the Art in
HEART FAILURE

CLINICAL PRACTICE AND ORGANIZATIONAL MODELS

Venue: Hotel Meliá MaríaPita, A Coruña

#ACoruñaHF2024

A CORUÑA 27-28 SEPTEMBER 2024

Galicia CS code: Short term mechanical circulatory support

Guillermo Bastos Fernandez, MD

Interventional Cardiology Unit

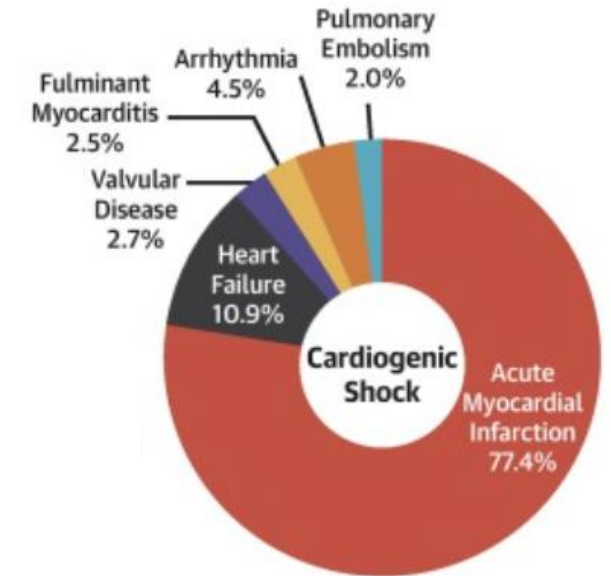
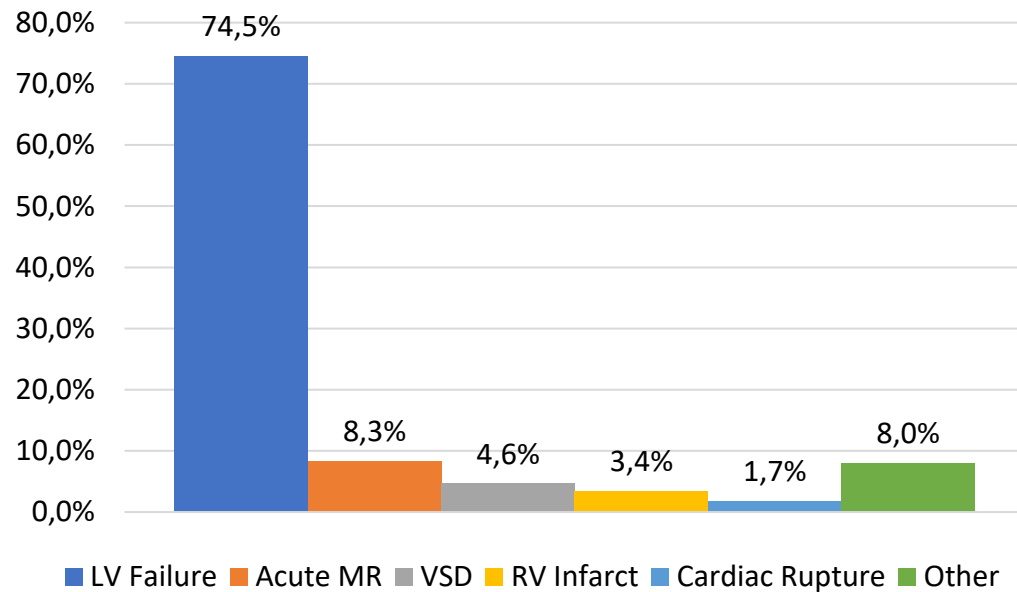
Alvaro Cunqueiro University Hospital - Vigo, Spain

Short term mechanical circulatory support

I do not have any potencial conflict of interst to declare

THE PROBLEM OF CS

- Acute coronary syndrom represent first cause of CS
 - ≈ 50% of all identifiable causes
- LV predominant fenotipe



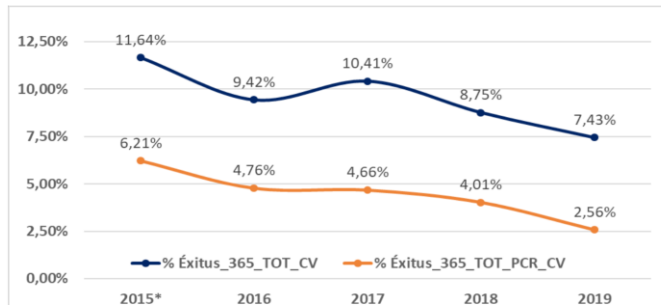
JACC Asia. 2022 Oct 31;3(1):122-134.

Hochman, J.S., et al. J Am Coll Cardiol, 2000. 36(3 Suppl A): p. 1063-70.

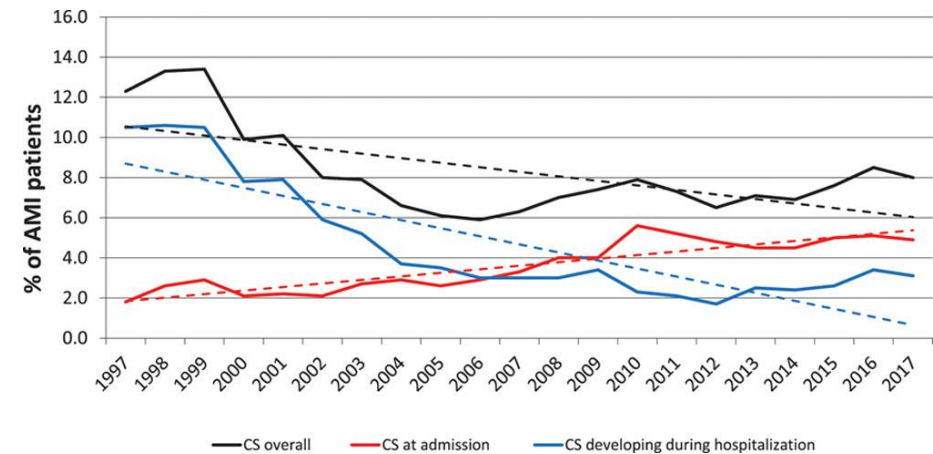
THE PROBLEM OF CS

- Rapid reperfusion strategies
 - Reduced complications and mortality <10%

Figura 8. Mortalidad cardiovascular (1 año). REGALIAM 2015-2019.



Regaliam 1y cardiovascular mortality

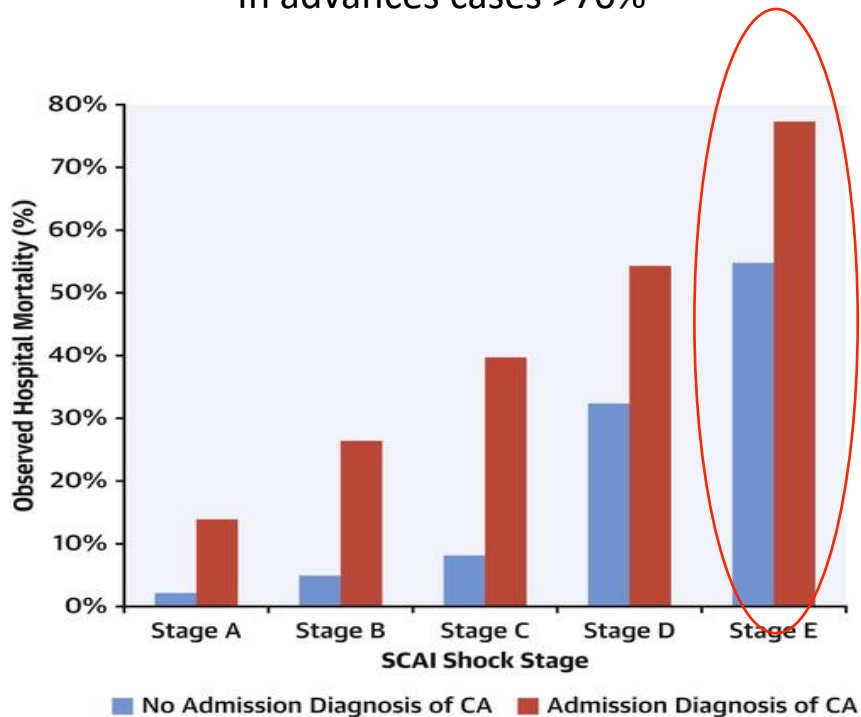


Circ Cardiovasc Interv. 2019 Apr;12(4):e007293.

- There is still and important AMI-CS subgroup (5-10%).

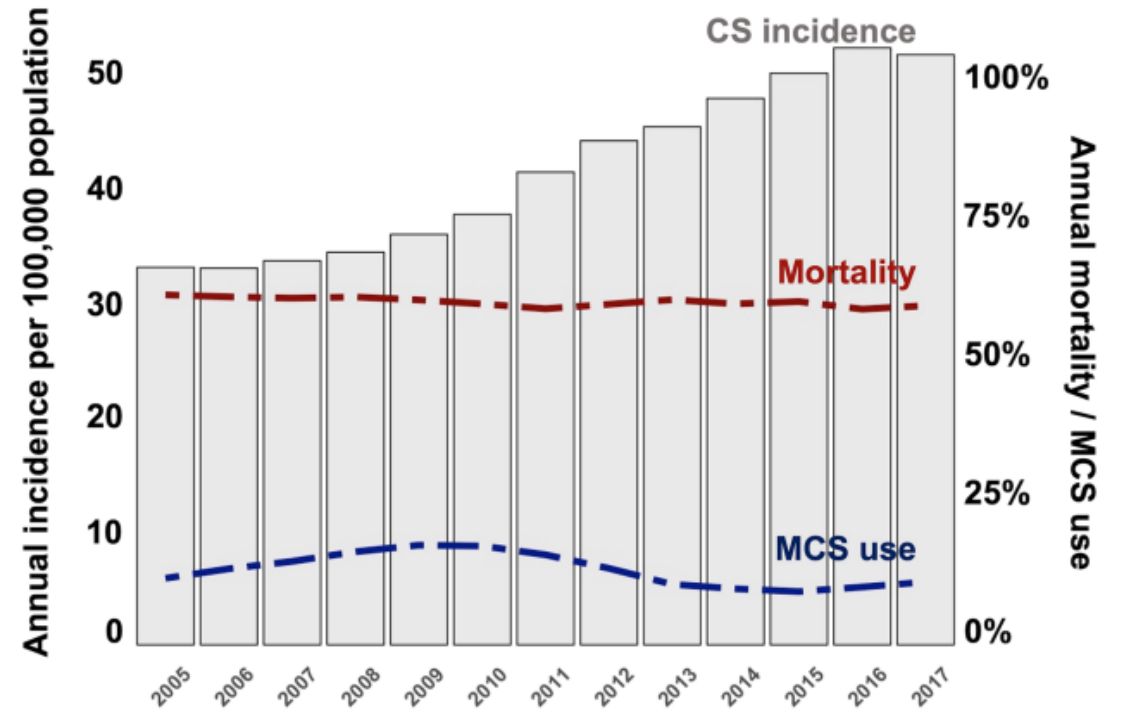
THE PROBLEM OF CS

- Very high mortality
 - Overall >50%
 - In advances cases >70%



Jacob C. Jentzer et al. JACC 2019; 74:2117-2128. Mayo Clinic CICU CS registry >10.000 pts

- Minimal improvement over time



ESC Heart Fail. 2021 Apr;8(2):1295-1303.

To start with... Nightmare case

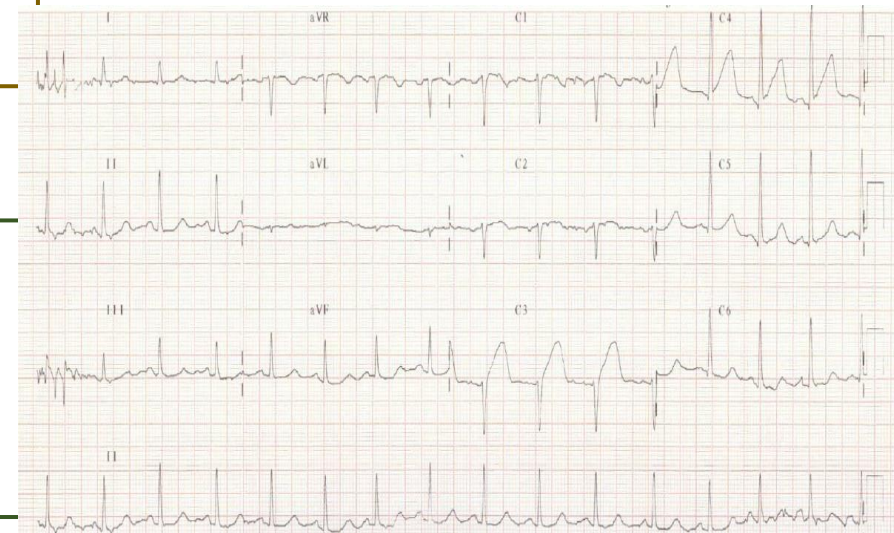
Female, 43 y.o no CV risk factors

Cardiac History

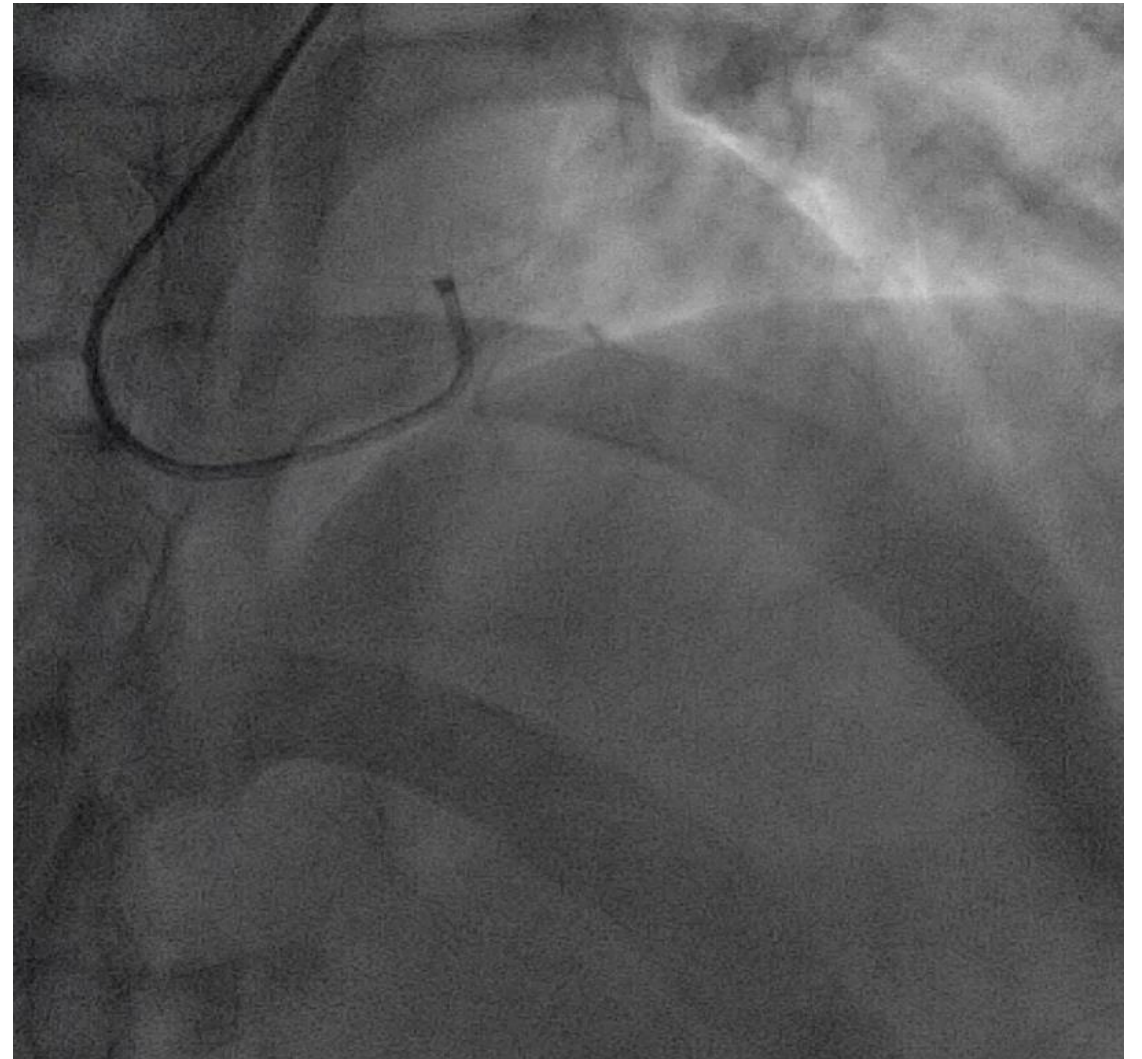
- **2014: Apical STEMI:** Spontaneous Coronary Artery Dissection (medical treatment). **TTE: Preserved LVEF**
- **Treatment:** Bisoprolol 2.5mg, AAS 100mg.

History of Present Illness

- **Chest pain with irradiation to left arm for the last 2h.**
- **ECG:** ST elevation in V3-V4
- **TpUs I:** 4441 ng/L
- **TTE:** Preserved LVEF, akinesia in LAD territory.



Cardiac Catheterization

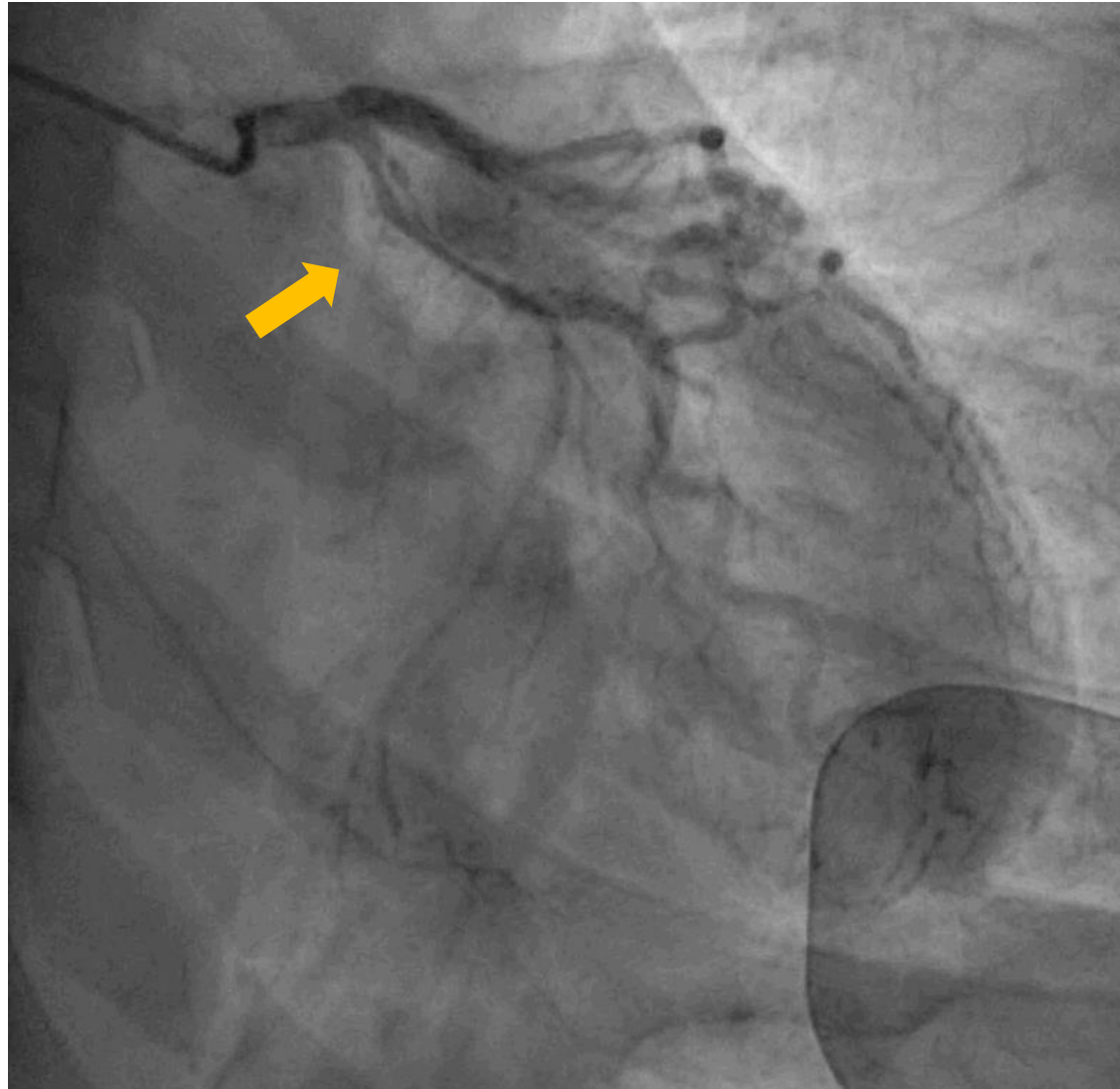


Cardiac Catheterization

Spontaneous
coronary dissection



TIMI III culprit vessel



Cardiac Catheterization

Cardiac Arrest due to VF
DEF x5 and CPR 10 min

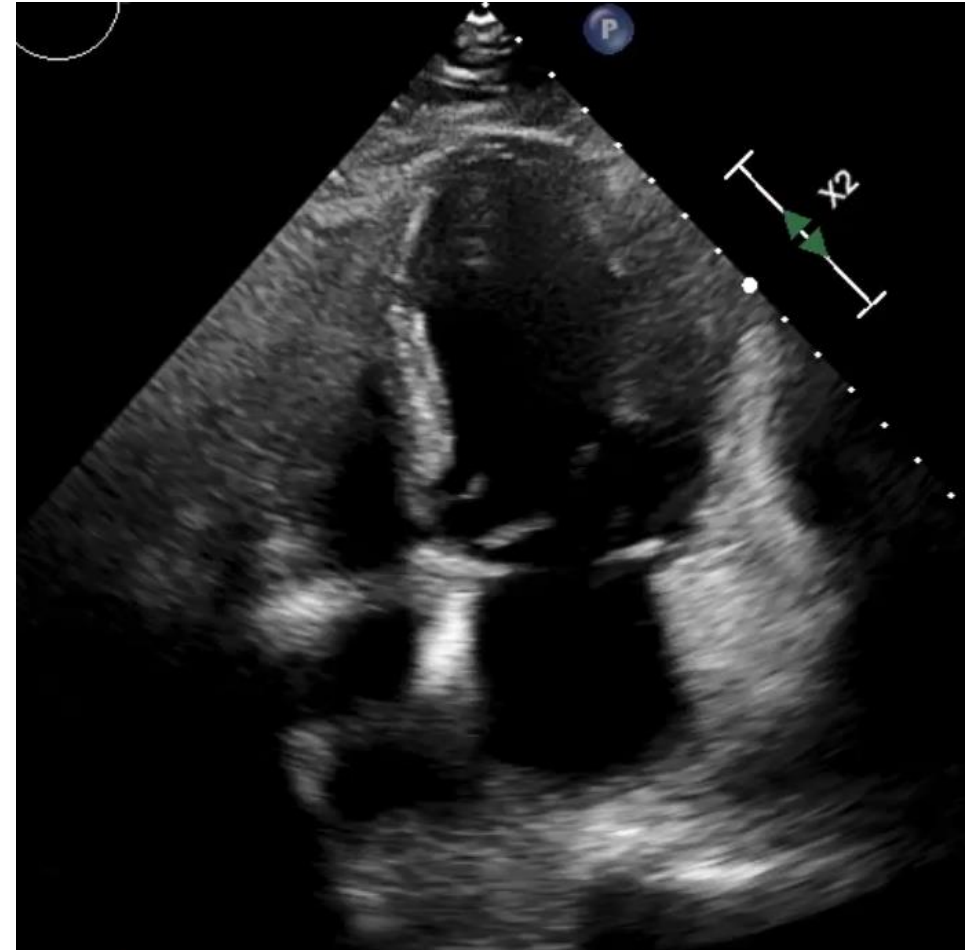
ROSC with ST elevation (anterior leads)

Mechanical Ventilation

Lactate 7mmol/L

BP 80/55mmHg (NEP 0.2 mcg/kg/min)

+



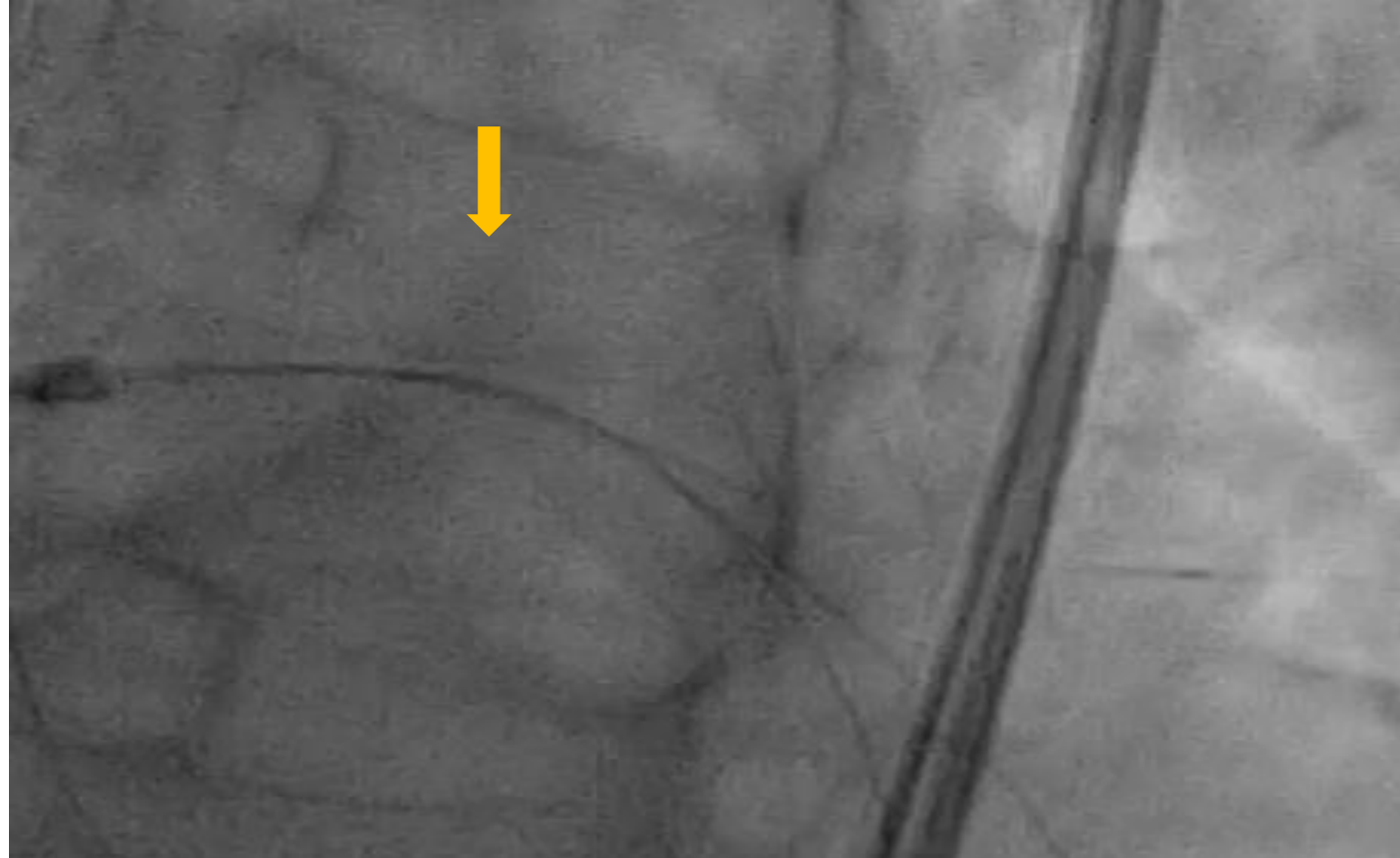
Severely decreased LVEF (25%)

Cardiac Catheterization (*LM-LAD*)

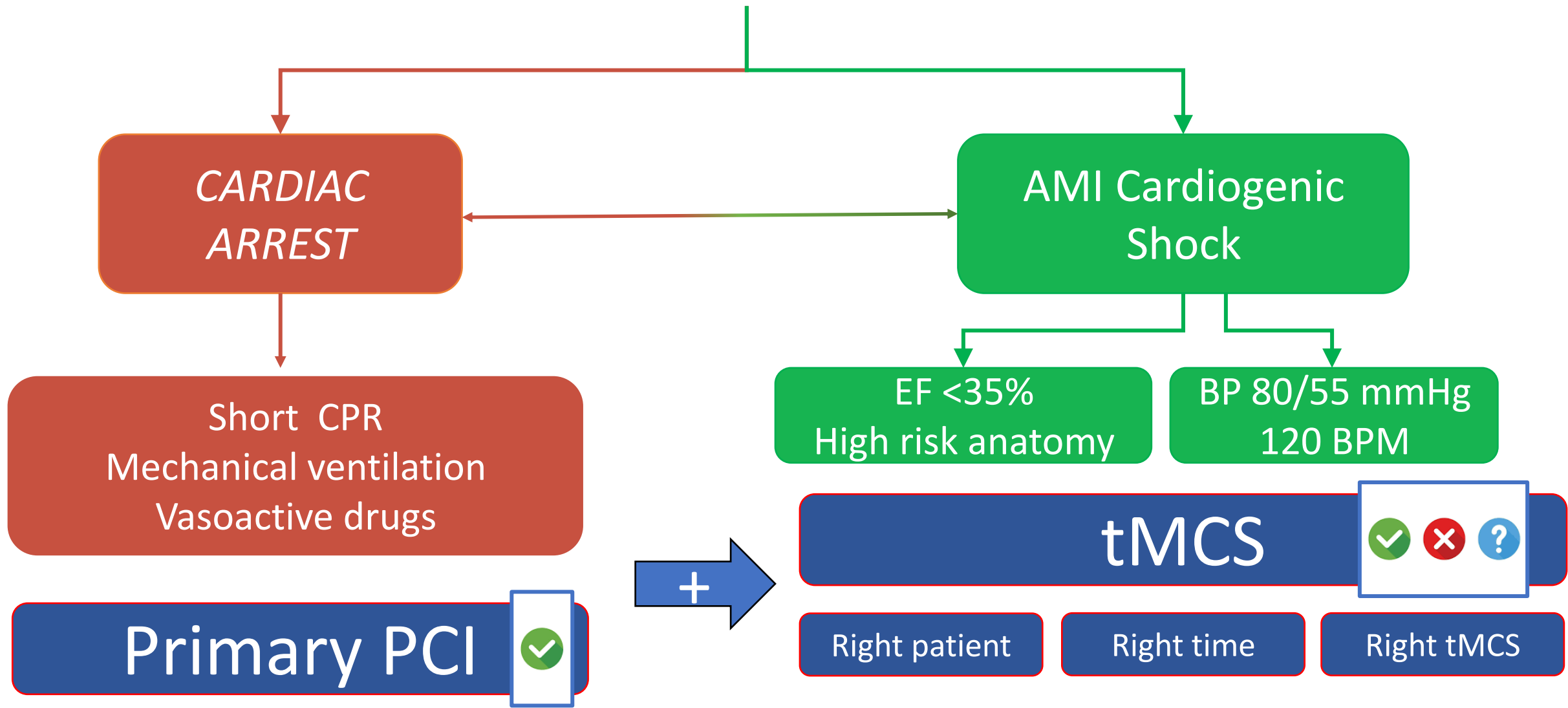
Coronary
dissection progression



LM hematoma
TIMI II flow



Heart Team Evaluation



Objetives of tMCS

- To improve hemodynamic stability before and during PCI
- To „buy time“ for recovery of stunned myocardium after PCI
- To bridge to LVAD/Tx if no myocardial recovery



Guidelines CS and tMCS



ESC
European Society
of Cardiology
European Journal of Heart Failure (2022) 24, 4–131
doi:10.1002/ejhf.2333

ESC GUIDELINES

2021 ESC Guidelines for the diagnosis and treatment of acute and chronic heart failure

Recommendations for the use of short-term mechanical circulatory support in patients with cardiogenic shock

Recommendations	Class ^a	Level ^b
Short-term MCS should be considered in patients with cardiogenic shock as a BTR, BTD, BTB. Further indications include treatment of the cause of cardiogenic shock or long-term MCS or transplantation.	IIa	C

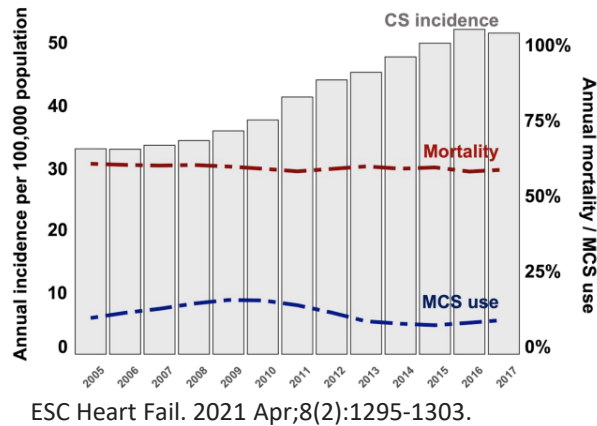
VS

Inotropic agents		
Inotropic agents may be considered in patients with SBP <90 mmHg and evidence of hypoperfusion who do not respond to standard treatment, including fluid challenge, to improve peripheral perfusion and maintain end-organ function. ³⁸⁸	IIb	C
Inotropic agents are not recommended routinely, due to safety concerns, unless the patient has symptomatic hypotension and evidence of hypoperfusion. ^{388,468,479}	III	C
Vasopressors		
A vasopressor, preferably norepinephrine, may be considered in patients with cardiogenic shock to increase blood pressure and vital organ perfusion. ^{486–488}	IIb	B

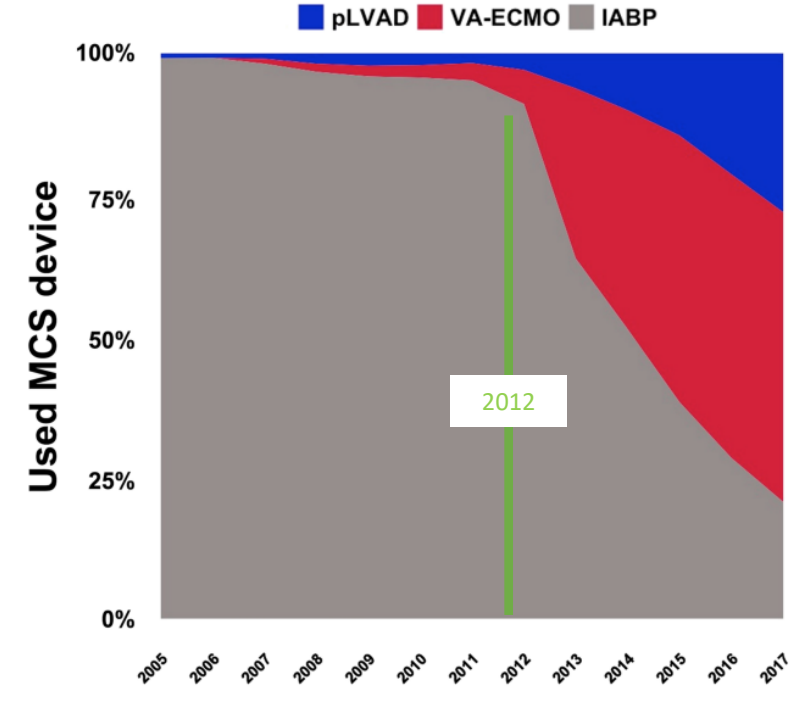
Daily practice tMCS

- Low MCS use: Only 10-15% of the CS

IABP treated	pLVAD treated	VA-ECMO treated	No device
(N = 42 636; 10%)	(N = 4063; 1%)	(N = 9825; 2%)	(N = 385 172; 87%)



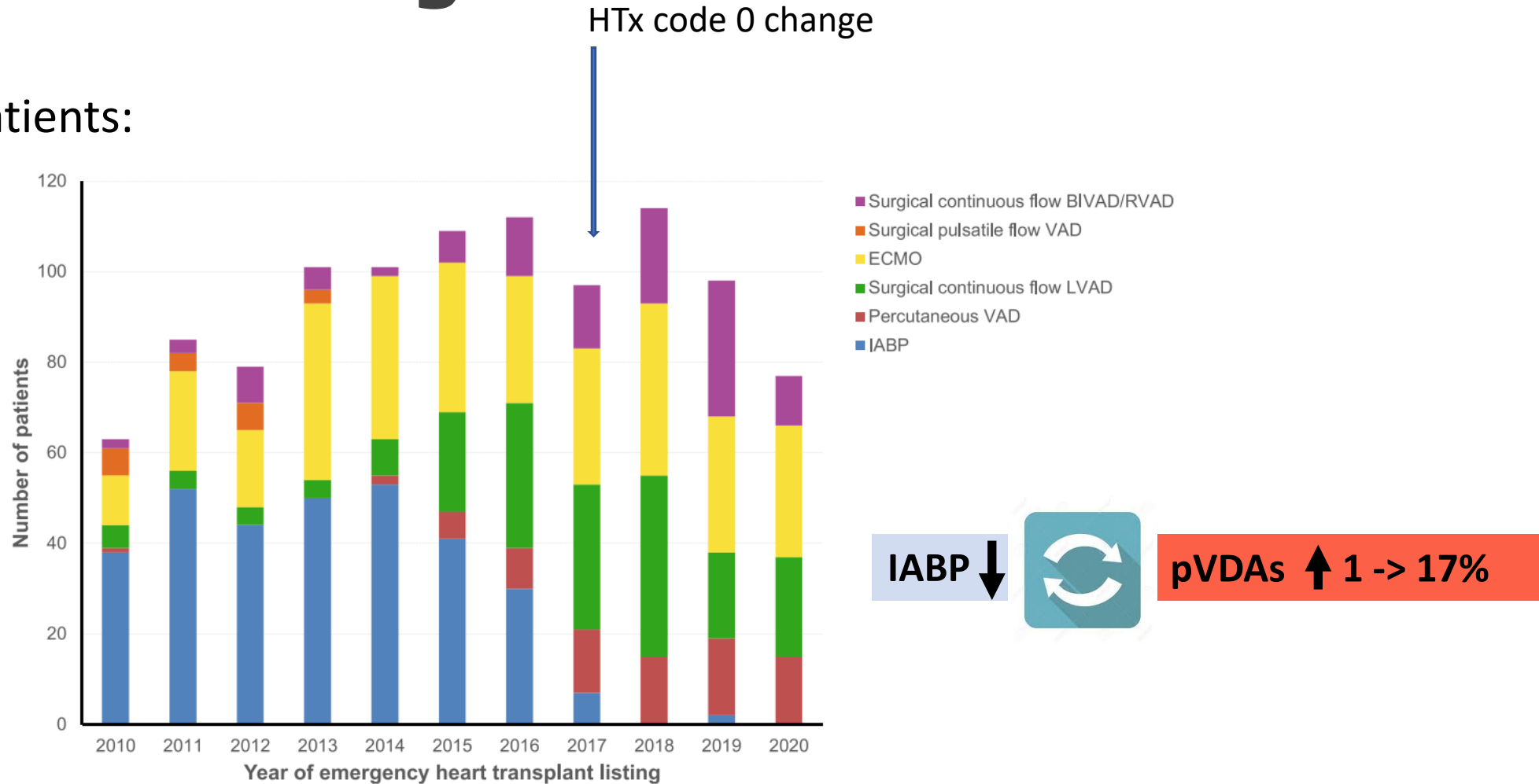
- Remarkable change after the IABP II-shock trial in AMI-CS patients



MCS preHTx bridge

- Also in preTx patients:

Asis-TC registry
Spanish
Emergency HTx



J Heart Lung Transplant. 2023 Apr;42(4):488-502.

Types of percutaneous tMCS

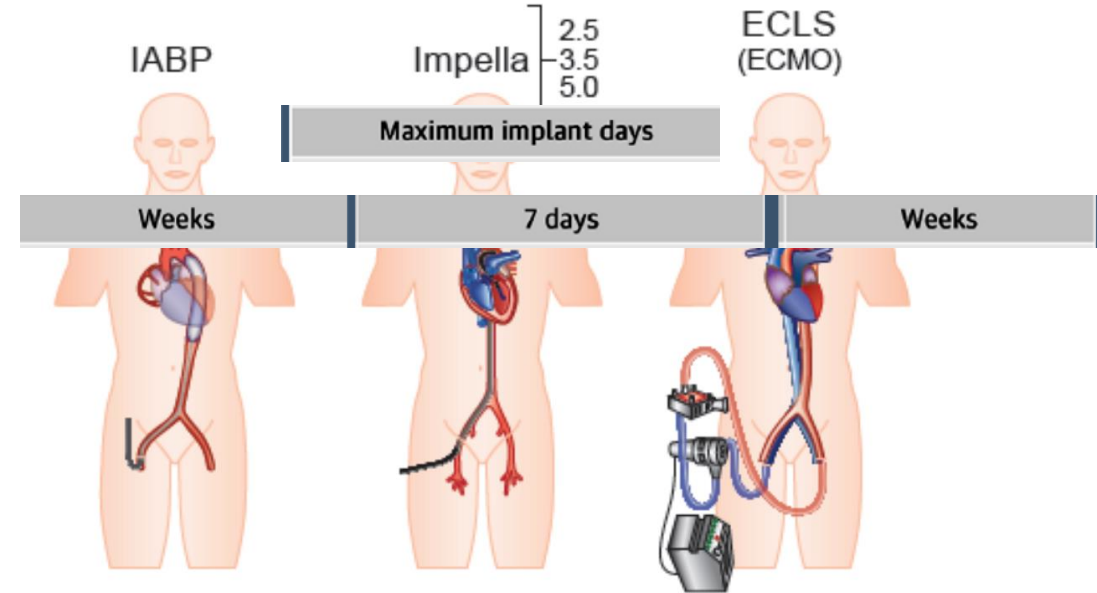
Disadvantage

Advantage

"Not bad"



	VA-ECMO	IABP	Impella (2.5, CP, 5.0, 5.5)
Flow	max 7.0 l/min	0.5 l/min	2.5 - 5.5 l/min
Pump Speed	max 5000 rpm	NA	max 51,000 rpm
Mechanism	Centrifugal flow continuous pump (RA-to-AO)	Balloon inflation-deflation (AO)	Axial flow continuous pump (LV-to-AO)
Cannula Size	14-19 F arterial 17-21 F venous	7-8 F arterial	13-21 F arterial
Insertion/Placement	Femoral vein Femoral artery	Femoral artery Axillary artery	Femoral artery Axillary artery
LV Unloading	-	+	+to+++
RV Unloading	++	-	-
Cardiac Power	↑↑	↑	↑↑
Afterload	↑↑	↓	↓↓
Coronary Perfusion	-	↑	↑

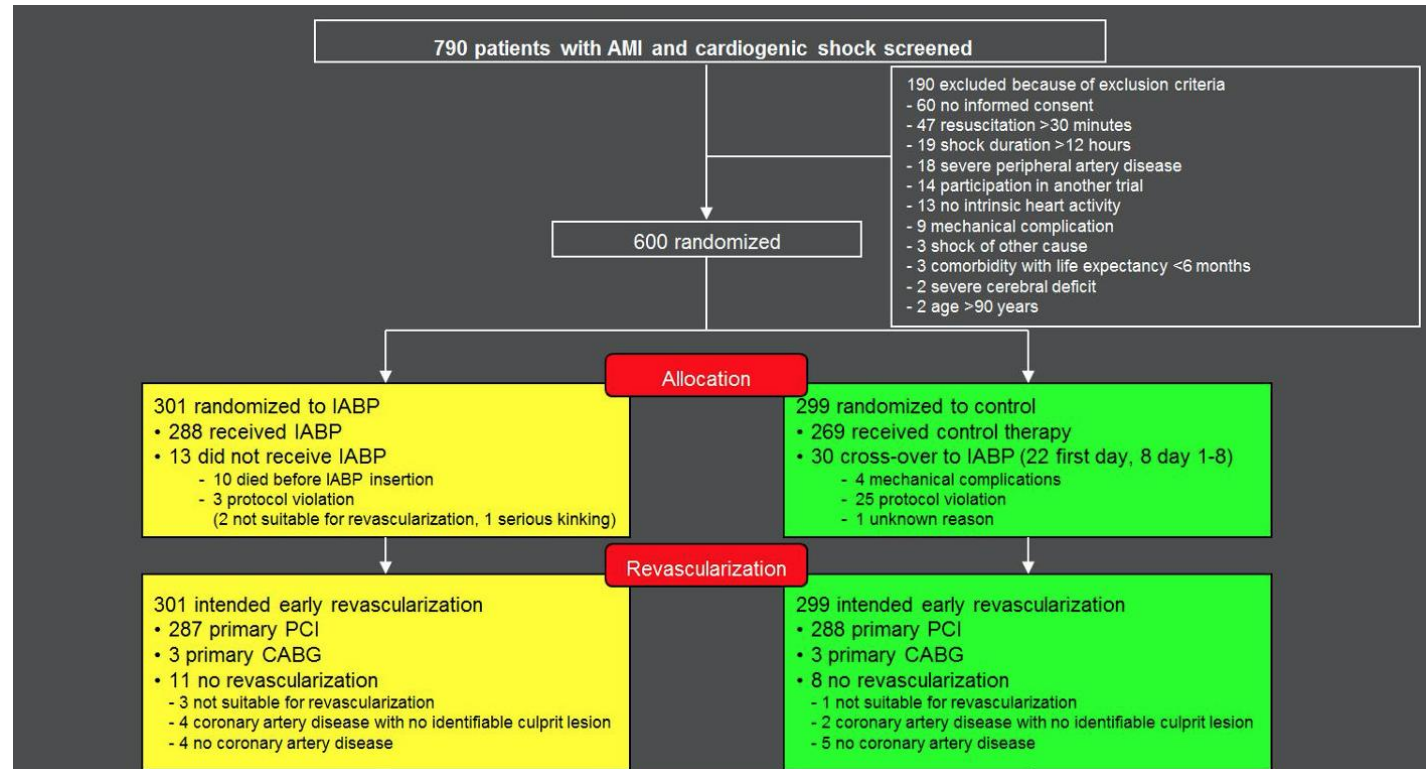
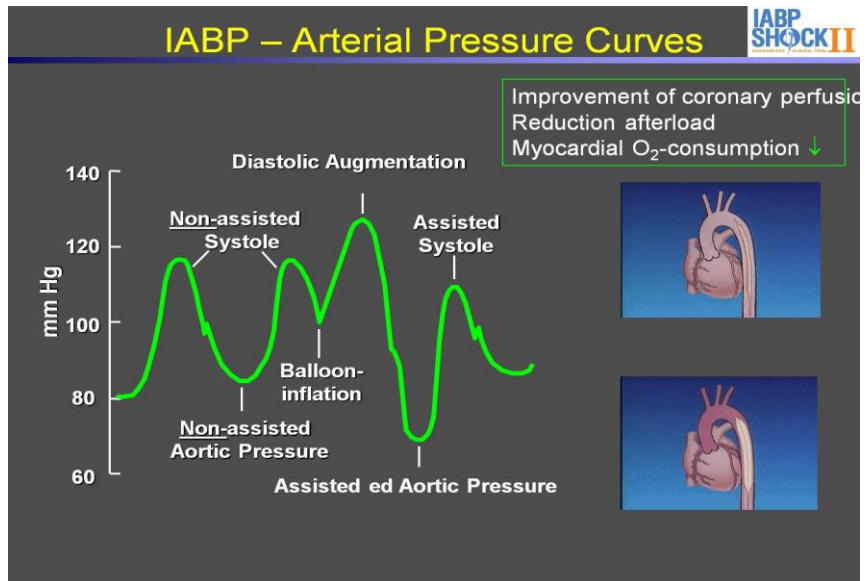
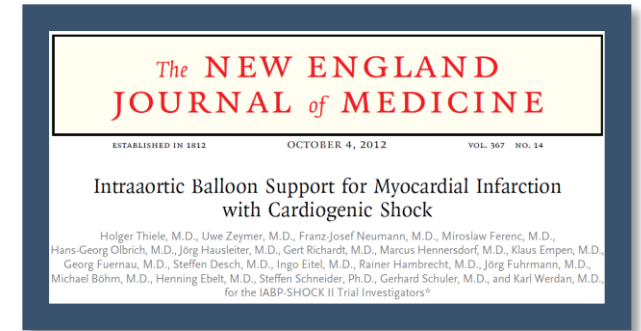


Eur J Heart Fail. 2018 Jan;20(1):178-186

Respiratory support

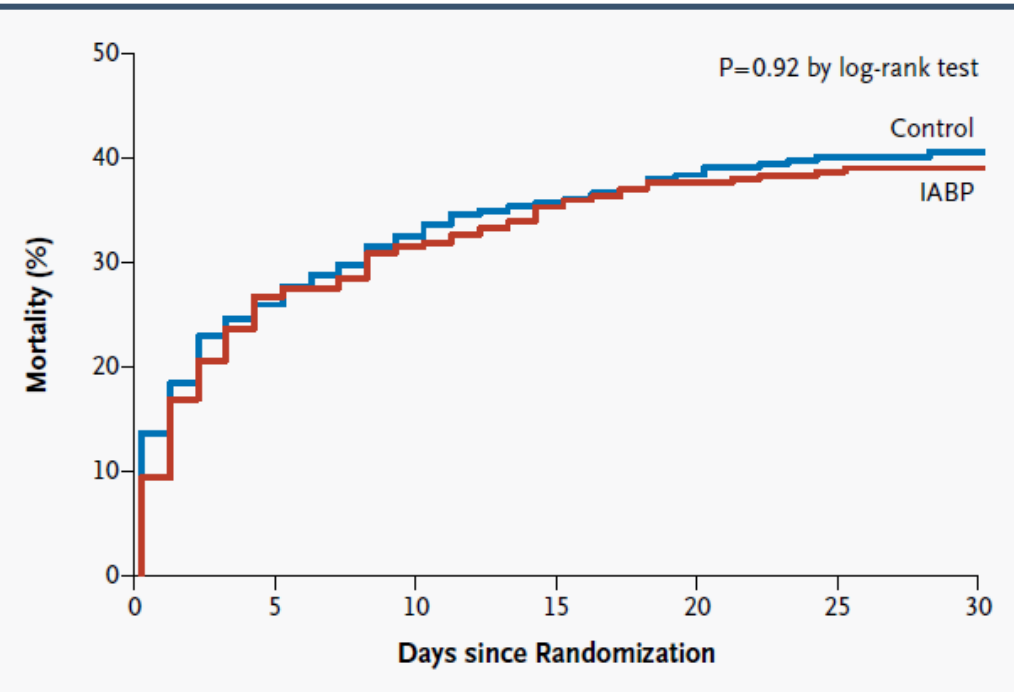


IABP in CS

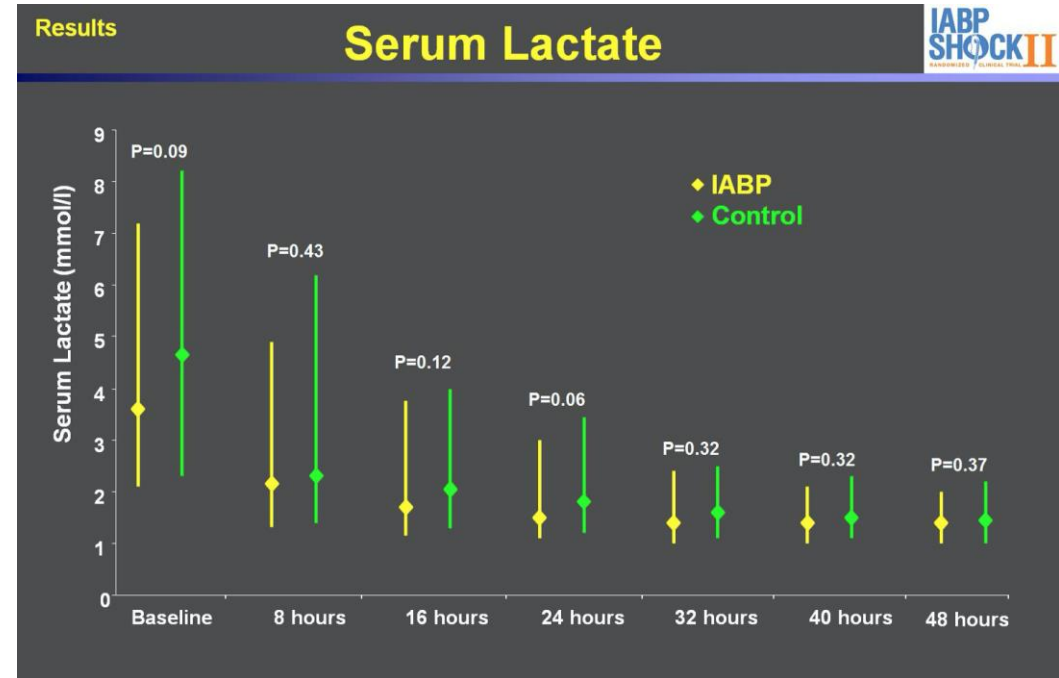


Thiele, H., et al. *N Engl J Med*, 2012. 367(14): p. 1287-96.

IABP in CS no clinical benefit



The use of **IABP did not significantly reduce 30-day mortality** in patients with CS complicating AMI for whom an early revascularization strategy was planned.



No effect on distal perfusion markers .

IABP in CS low complication rate

Outcome	IABP (N=300) <i>number (percent)</i>	Control (N=298) <i>number (percent)</i>	P Value	Relative Risk with IABP (95% CI)
Primary end point: all-cause mortality at 30 days	119 (39.7)	123 (41.3)	0.69	0.96 (0.79–1.17)
Reinfarction in hospital	9 (3.0)	4 (1.3)	0.16	2.24 (0.70–7.18)
Stent thrombosis in hospital	4 (1.3)	3 (1.0)	0.71	1.32 (0.30–5.87)
Stroke in hospital	2 (0.7)	5 (1.7)	0.28	0.40 (0.08–2.03)
Ischemic	2 (0.7)	4 (1.3)	0.45	0.49 (0.09–2.71)
Hemorrhagic	0	1 (0.3)	0.50	—
Peripheral ischemic complications requiring intervention in hospital	13 (4.3)	10 (3.4)	0.53	1.29 (0.58–2.90)
Bleeding in hospital*				
Life-threatening or severe	10 (3.3)	13 (4.4)	0.51	0.76 (0.34–1.72)
Moderate	52 (17.3)	49 (16.4)	0.77	1.05 (0.74–1.50)
Sepsis in hospital	47 (15.7)	61 (20.5)	0.15	0.77 (0.54–1.08)

IABP in the Guidelines

ESC Guidelines AMI routine use IABP

2012 IABP-SHOCK II Trial

2019

Class IC

Class IIb B

Class III



ESC
European Society
of Cardiology
European Heart Journal (2021) 42, 3599–3726
doi:10.1093/eurheartj/ehab368

ESC GUIDELINES

2021 ESC Guidelines for the diagnosis and
treatment of acute and chronic heart failure

Class IIa C

Mechanical complications, as a bridge to more advanced supports

Class IIb C

Refractory shock not due to myocardial infarction

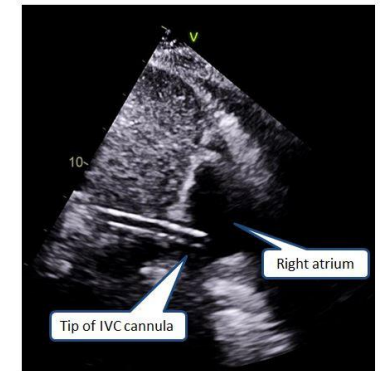
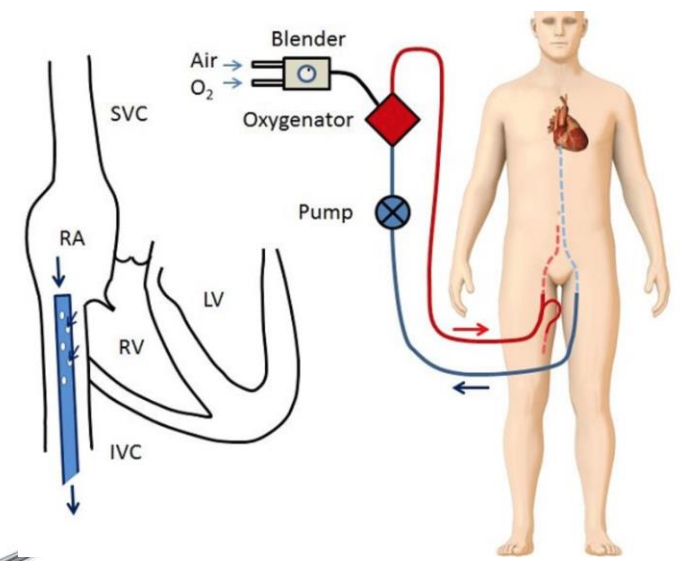
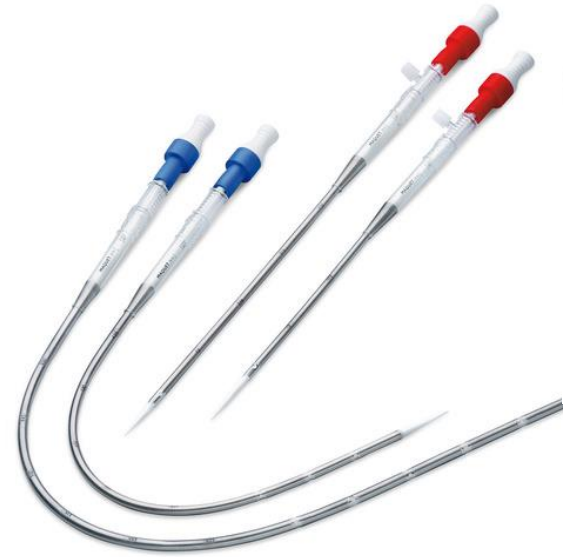
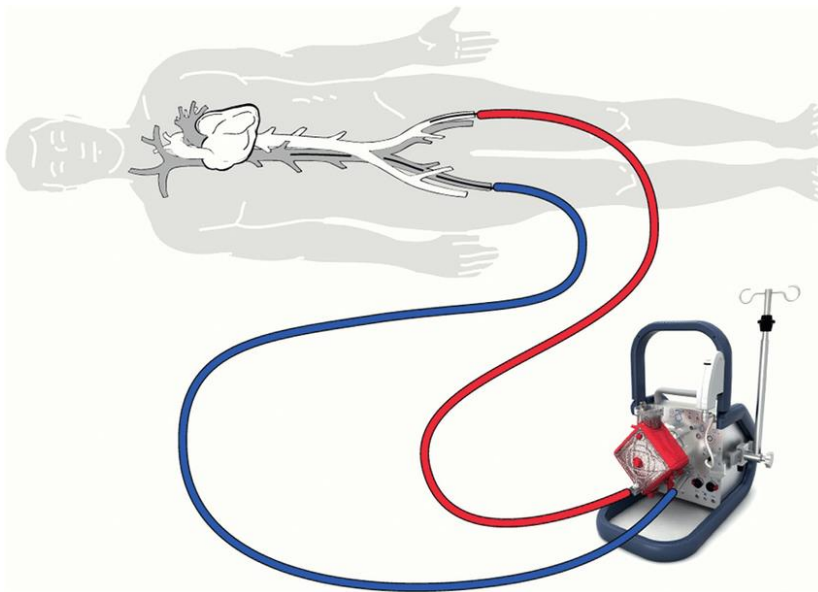
No based in any RCTs

Holger Thiele
@thiele_holger
This is where IABP console may have its place if you follow evidence 🤔



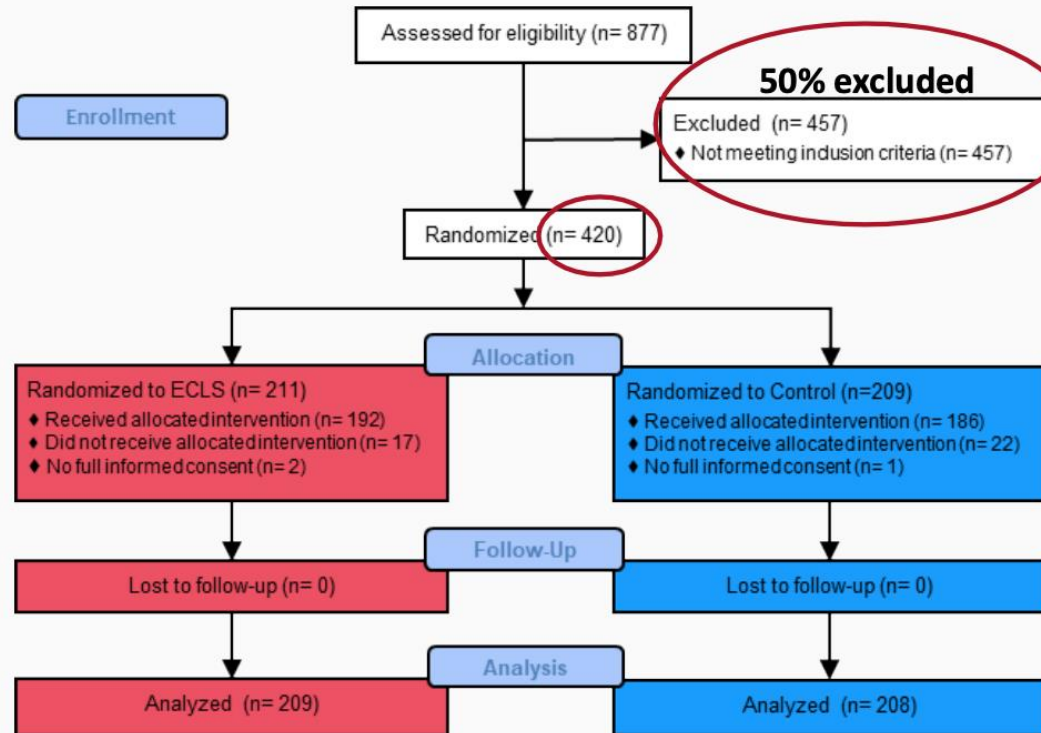
ECMO VA

VA-ECMO
max 7.0 l/min
max 5000 rpm
Centrifugal flow continuous pump (RA-to-AO)
14-19 F arterial 17-21 F venous
Femoral vein Femoral artery
-
++
↑↑
↑↑
-



ECMO VA in CS ECLS SHOCK trial

420 pts in 44 centers in 3 years: nearly 3 patient per year/center



The NEW ENGLAND JOURNAL of MEDICINE


ORIGINAL ARTICLE

Extracorporeal Life Support in Infarct-Related Cardiogenic Shock

H. Thiele, U. Zeymer, I. Akin, M. Behnes, T. Rassaf, A.A. Mahabadi, R. Lehmann, I. Eitel, T. Graf, T. Seidler, A. Schuster, C. Skurk, D. Duerschmied, P. Clemmensen, M. Hennersdorf, S. Fichtlscherer, I. Voigt, M. Seyfarth, S. John, S. Ewen, A. Linke, E. Tigges, P. Nordbeck, L. Bruch, C. Jung, J. Franz, P. Lauten, T. Goslar, H.-J. Feistritz, J. Pöss, E. Kirchhof, T. Ouarrak, S. Schneider, S. Desch, and A. Freund, for the ECLS-SHOCK Investigators*

Inclusion Criteria	Exclusion Criteria
<ul style="list-style-type: none"> Cardiogenic shock complicating AMI (STEMI or NSTEMI) plus obligatory: SCAI C-D-E 1. Planned revascularization 2. SBP <90 mmHg >30 min or catecholamines required to maintain SBP >90 mmHg 3. Signs of impaired organ perfusion with at least one of the following criteria: <ul style="list-style-type: none"> ➢ Altered mental status ➢ Cold, clammy skin and extremities ➢ Oliguria with urine output <30 ml/h 4. Arterial lactate >3 mmol/l * Informed consent 	<ul style="list-style-type: none"> Resuscitation >45 minutes Mechanical cause of cardiogenic shock Onset of shock >12 h Severe peripheral artery disease with impossibility to insert ECLS cannulae Age <18 years or >80 years Shock of other cause (bradycardia, sepsis, hypovolemia, etc.) Other severe concomitant disease with limited life expectancy <6 months Pregnancy Participation in another trial

ECMO VA in CS ECLS SHOCH trial

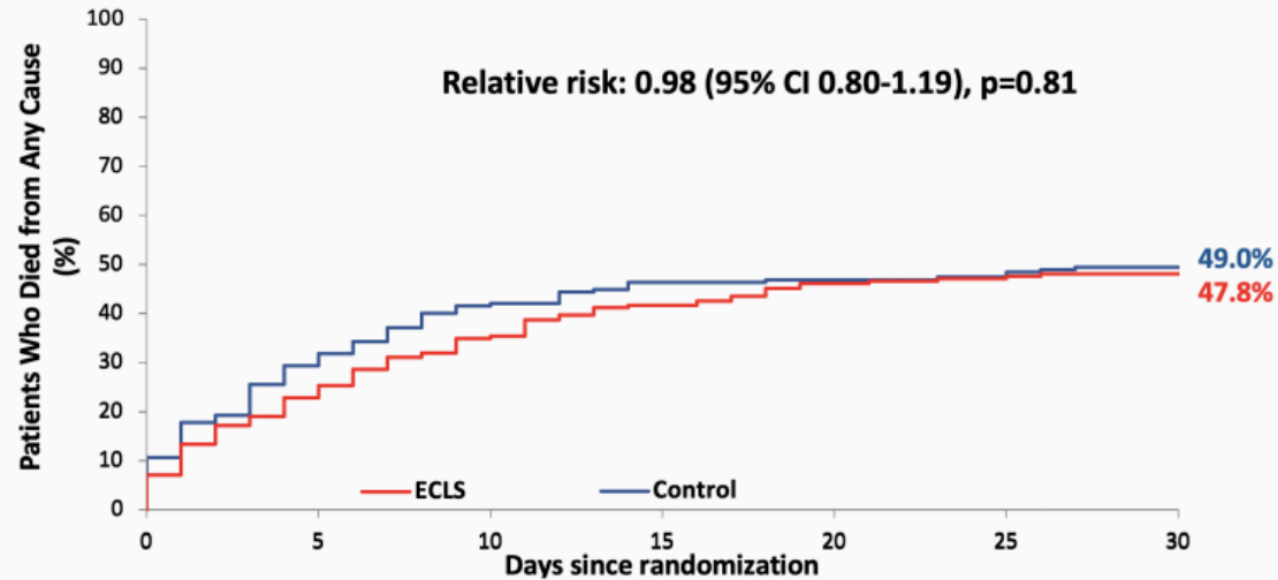
Results	Baseline Characteristics		
	ECLS (n=209)	Control (n=208)	
Age (years); median (IQR)	62 (56 - 69)	63 (57 - 71)	
Male sex; n/total (%)	170/209 (81.3)	169/208 (81.3)	
Mean blood pressure (mmHg); median (IQR)	71 (61 - 87)	72 (60 - 88)	
STEMI*; n/total (%)	135/204 (66.2)	141/207 (68.1)	
* 46.8 % vs 48.5%. LAD as IRA			
Resuscitation before randomization; n/total (%)	162/209 (77.5)	162/208 (77.9)	
Laboratory values on admission			
pH; median (IQR)	7.2 (7.1 - 7.3)	7.2 (7.1 - 7.3)	
Lactate (mmol/L); median (IQR)	6.8 (4.5 - 9.6)	6.9 (4.6 - 10.0)	
SCAI Shock classification; n/total (%)			
C	104/209 (49.8)	111/208 (53.4)	
D	38/209 (18.2)	18/208 (8.7)	
E	67/209 (32.1)	79/208 (38.0)	
Median time until return of spontaneous circulation 20 minutes in both groups			

N Engl J Med. 2023 Oct 5;389(14):1286-1297

ECMO VA in CS ECLS SHOCK trial

Results

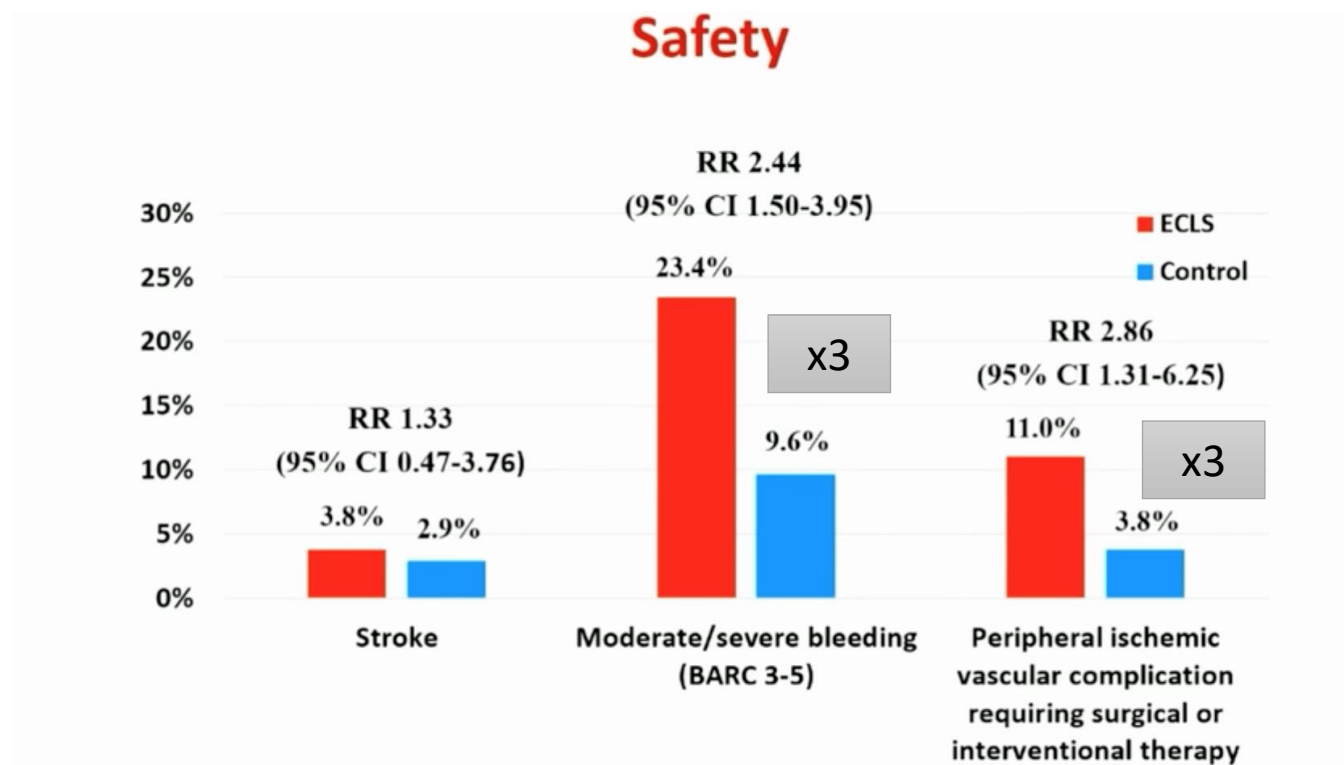
Primary Endpoint – 30-Day All-Cause Mortality



No. at Risk

ECLS	209	161	136	119	109	107	105
Control	208	146	120	109	105	104	100

ECMO VA in CS ECLS SHOCH trial



Microaxial Flow Pump in CS

	Impella (2.5, CP, 5.0, 5.5)
Flow	2.5 - 5.5 l/min
Pump Speed	max 51,000 rpm
Mechanism	Axial flow continuous pump (LV-to-AO)
Cannula Size	13-21 F arterial
Insertion/Placement	Femoral artery Axillary artery
LV Unloading	+ to +++
RV Unloading	-
Cardiac Power	↑↑
Afterload	↓↓
Coronary Perfusion	↑



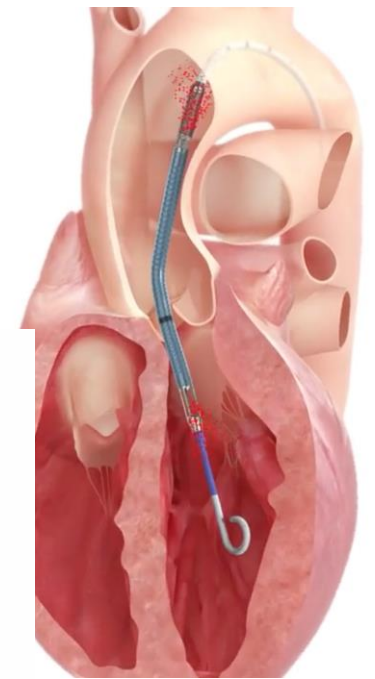
Impella CP® with SmartAssist®
 Percutaneous insertion, increased flow and repositioning without imaging



Impella 5.5® with SmartAssist®
 Designed for surgeons; delivers full forward flow from the ventricle



Impella RP® with SmartAssist®
 The first percutaneous, single access pump designed for right heart support



Automated Impella Controller™
 The primary user control interface for the Impella Platform

NCSI early tMCS registry

Journal of the American Heart Association

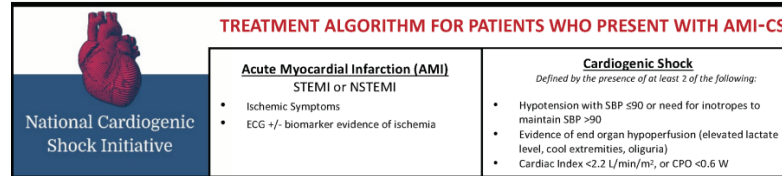
Volume 12, Issue 23, 5 December 2023
<https://doi.org/10.1161/JAHA.123.031401>



ORIGINAL RESEARCH

Early Utilization of Mechanical Circulatory Support in Acute Myocardial Infarction Complicated by Cardiogenic Shock: The National Cardiogenic Shock Initiative

NATIONAL CSI: CLINICAL SITES & ENROLLMENT



ACTIVATE CATH LAB

VASCULAR ACCESS

- Obtain femoral arterial access (via direct visualization with use of ultrasound, fluoroscopy & micropuncture)
- Obtain venous access (femoral or internal jugular)

CONFIRMATION OF CLINICAL DIAGNOSIS

- Clear AMICS diagnosis = Place MCS
- Unclear AMICS diagnosis = Perform RHC/Echo as needed

MECHANICAL CIRCULATORY SUPPORT
 Once AMICS diagnosis confirmed, place MCS

PCI
 Perform culprit PCI with the goal of establishing TIMI III flow, careful selective MV-PCI can be considered. Avoid CTO PCI

CPO (Cardiac Power Output)
 $\frac{MAP \times CO}{451}$

PAPI (Pulmonary Artery Pulsatility Index)
 $\frac{sPAP - dPAP}{RA}$

WEANING OF INOTROPES / ESCALATION OF SUPPORT

- For patients requiring ≥ 2 inotropes, operators should wean inotropes in the Cath Lab and reassess hemodynamics to determine if patient would benefit from early MCS escalation.
- If CPO remains ≤ 0.6 (requiring inotropes), operators should consider escalation of MCS in the Cath Lab (**estimated survival <50%**):
 - If PAPI is ≤ 0.9 & RA pressure >12 , consider escalation of right-sided MCS
 - If PAPI >0.9 , escalate left-sided MCS
- If CPO is >0.6 without inotropes (or low-moderate doses of a single inotrope), the patient should be transferred to the ICU (**estimated survival >70%**)

VASCULAR ASSESSMENT

- Prior to transfer from the Cath Lab, a detailed vascular exam should be performed including femoral angiogram, physical examination, and Doppler assessment of the affected limb.
- If indicated, external bypass should be performed.

BEST PRACTICES:

- MCS Pre-PCI
- Door to Support **<90 minutes**
- Establish TIMI III Flow
- RHC Utilization
- Wean Inotropes
- Maintain CPO **>0.6W**
- Survival to Hospital Discharge Goal **$\geq 80\%$**

Early tMCS

Door-to-support time in STEMI, min*	78 [41–237]	80 [41–238]	70 [43–226]	0.60
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Characteristics	All (N=406)	Stage C/D (N=295)	Stage E (N=111)	P value
Impella insertion				
Pre-PCI	70% (285)	69% (201)	76% (84)	0.38
Intraprocedural	9% (35)	9% (27)	8% (7)	
Post-PCI	21% (84)	22% (65)	17% (19)	

J Am Heart Assoc. 2023 Dec 5;12(23):e031401.



NCSI early tMCS registry

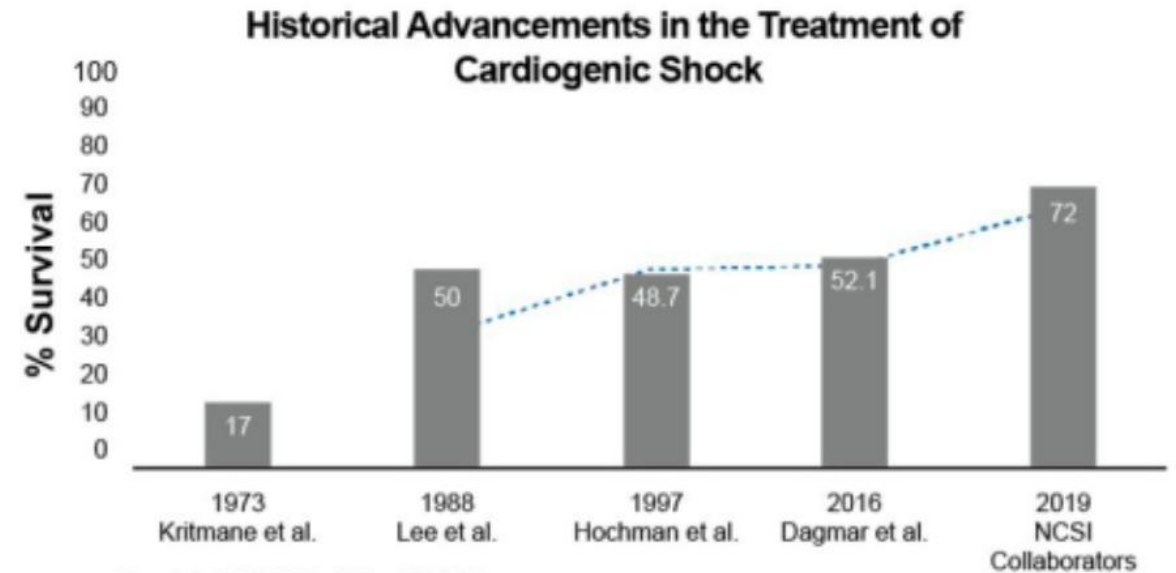
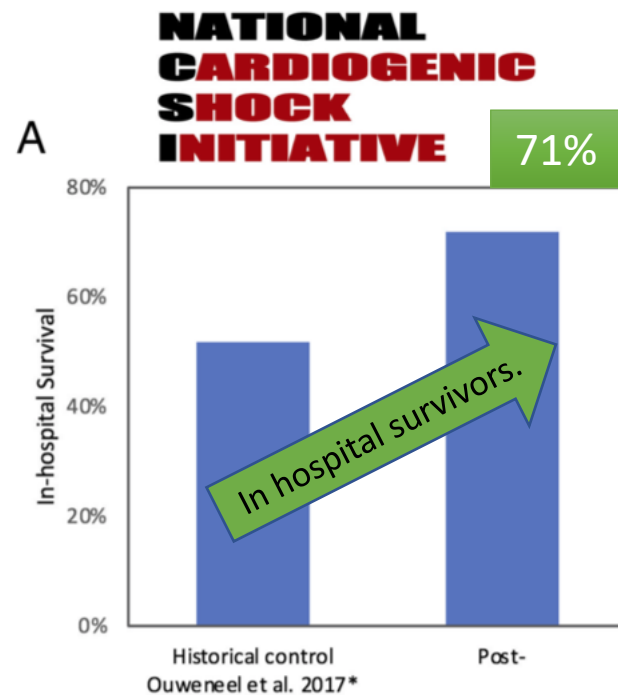
Journal of the American Heart Association

Volume 12, Issue 23, 5 December 2023
<https://doi.org/10.1161/JAHA.123.031401>



ORIGINAL RESEARCH

Early Utilization of Mechanical Circulatory Support in Acute Myocardial Infarction Complicated by Cardiogenic Shock: The National Cardiogenic Shock Initiative



J Am Heart Assoc. 2023 Dec 5;12(23):e031401.

Danger SHOCK Trial

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

Microaxial Flow Pump or Standard Care
in Infarct-Related Cardiogenic Shock

Study Design

Multicenter, double-blinded, randomized trial

STEMI and Cardiogenic Shock
N = 360

Impella CP
N = 179

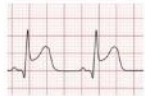
BMS
N = 176

Primary Endpoint: Death from any cause at 180
days

Danger SHOCK Trial

Inclusion

Lactate >2.5 mmol/L



STEMI



Hypotension and hypoperfusion



LVEF < 45%

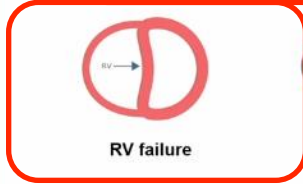


Randomization when shock was diagnosed

Exclusion



Comatose OHCA

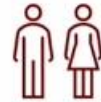


RV failure



Mechanical complication

Patients characteristics – N=355



Median 67 years
79% male



Median lactate 4.5 mmol/L



72% LAD or LM culprit
72% Multi vessel disease



Median 4 hrs from onset of STEMI symptoms to randomization

84% randomized in cath lab



Median LVEF 25%



55% SCAI class C
45% SCAI class D or E



Median systolic BP 82 mmHg

Apply 20% all CS

Very selected population

Acute STEMI: symptoms <5h

LV predominant CS

Low CPR 20% rate
(only short, no comatose)

10y to complete reclutation (2013-2023)

N Engl J Med 2024;390:1382-1393

Danger SHOCK Trial

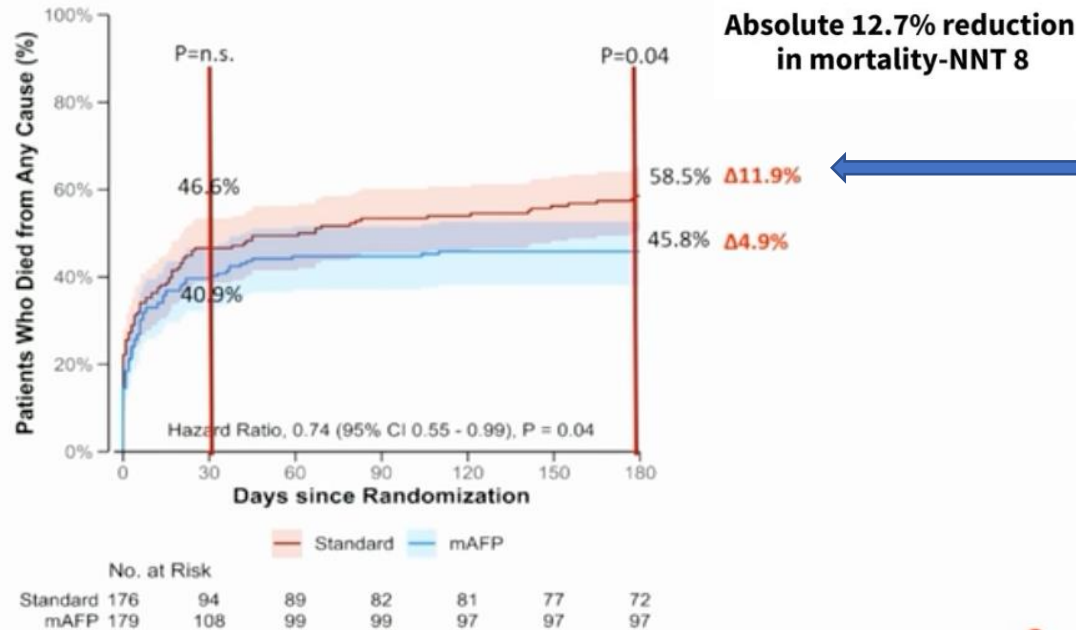
Primary Outcome Measures

Time Frame

All-cause mortality

180 days

Primary Endpoint - 6-Month Mortality



Conclusion

- The routine use of a mAFP on top of standard care reduced death from any cause in patients with STEMI and cardiogenic shock.

Very high mortality in control group from 30d-6M

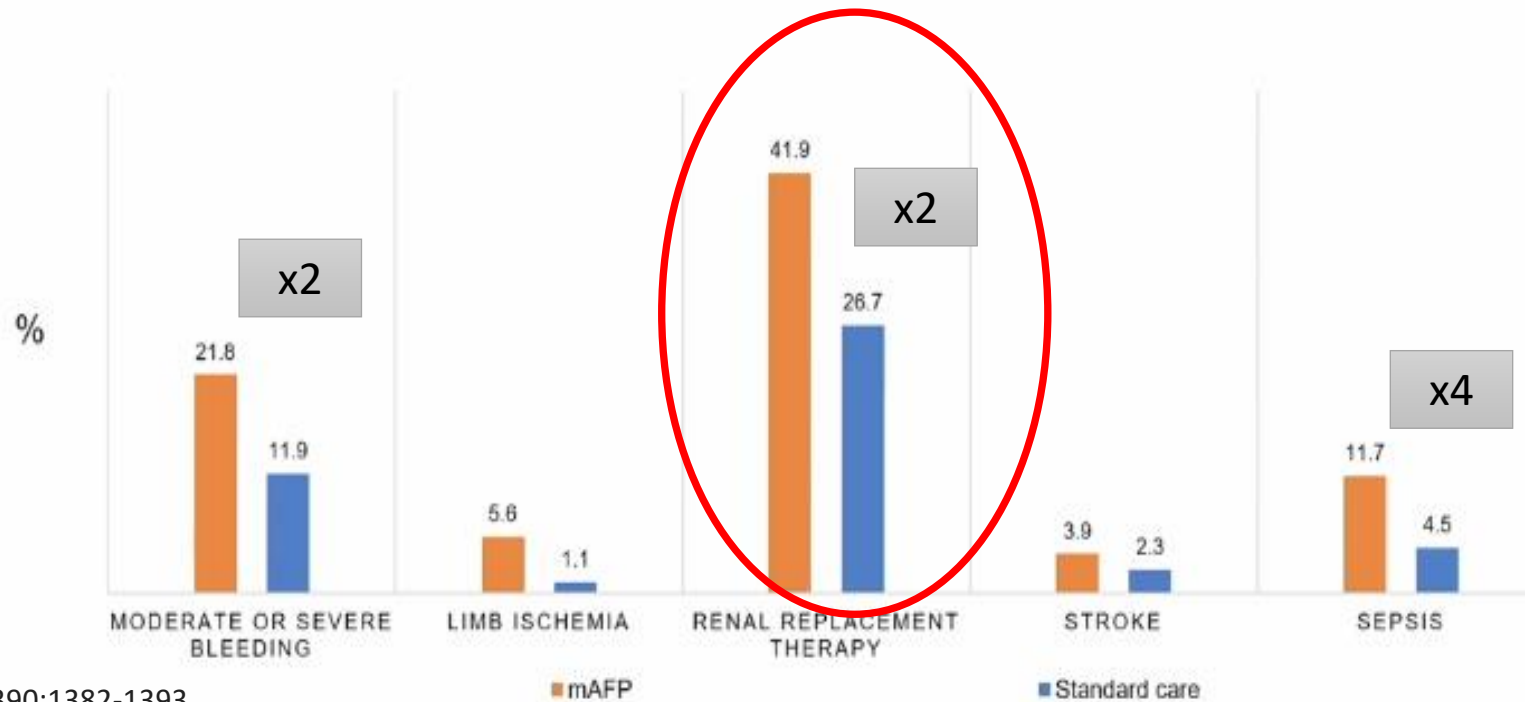
N Engl J Med 2024;390:1382-1393

Danger SHOCK Trial

Microaxial Flow Pump or Standard Care in Infarct-Related Cardiogenic Shock

Møller JE et al. DOI: 10.1056/NEJMoa2312572

Adverse events



N Engl J Med 2024;390:1382-1393

Danger SHOCK Trial

Microaxial Flow Pump or Standard Care in Infarct-Related Cardiogenic Shock

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Conclusion

- The routine use of a mAFP on top of standard care reduced death from any cause in patients with STEMI and cardiogenic shock.

NSTEMI ≈ 25%

n (%)	No CS (n=47 752)	CS (n=4090)	P Value
Age, y; mean (SD)	65.8 (13.2)	69.6 (12.5)	<0.001
Sex: female (%)	12 631/47 752 (26.5)	1263/4090 (30.9)	<0.001
Resuscitation prior admission (%)	1569/47 549 (3.3)	1223/4059 (30.1)	<0.001
Delay, min (symptoms on admission); median (IQR)	230 (115–660)	165 (84–483)	<0.001
Vital signs at admission, mean (SD)			
Systolic blood pressure, mm Hg	138.4 (27.1)	111.7 (31.0)	<0.001
Diastolic blood pressure, mm Hg	80.6 (16.8)	67.4 (21.5)	<0.001
Heart rate, bpm	78.2 (19.2)	89.5 (28.2)	<0.001
STEMI (%)	27 466/47 752 (57.5)	3027/4090 (74.0)	<0.001
NSTEMI (%)	20 286/47 752 (42.5)	1063/4090 (26.0)	<0.001

Circ Cardiovasc Interv. 2019 Apr;12(4):e007293.

≈ 20-30% AMI
CARDIOGENIC SHOCK

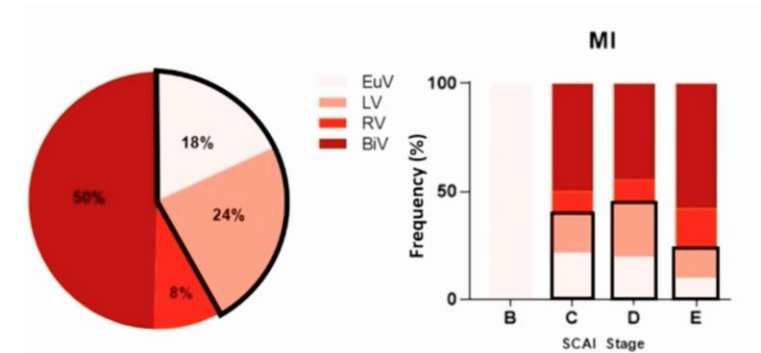
OHCA in CS ≈ 5%

Table 1 – Incidence of OHCA within patients treated for STEMI by year (95% confidence interval in parenthesis).

	OHCA+	OHCA-	OHCA + incidence (%)
2014	263	4284	5.78 [5.12–6.50]
2015	306	5253	5.50 [4.92–6.14]
2016	288	5733	4.78 [4.26–5.35]
2017	342	5894	5.48 [4.93–6.08]
2018	336	5384	5.87 [5.38–6.52]
Total	1535	26548	5.47 [5.20–5.74]

Resusc Plus 2021 Mar 31:6:100113.

BiV / VD STEMI SHOCK ≈ 50%



~40% of MI-CS 21

ECLS SHOCK Vs Danger SHOCK trial

No mortality reduction

How Can these Differences Be Explained?



LV Loading

STEMI + NSTEMI
77% Resuscitation
-> 20% hypoxic brain injury
Lactate 6.9 mmol/l
Ventilation: 90%
Catecholamines: 90%

30 days

RRT 8.1% vs 13.1%
ICU time: 10 vs 8 days
MCS duration: 2.7 days

Device?

Patient selection?

Follow-up?

Treatment Bias?



LV Unloading

STEMI only
20% Resuscitation - only witnessed CA
-> 2% hypoxic brain injury
Lactate 4.5 mmol/l
Ventilation: 70%
Catecholamines: 85%

6 months

RRT 41.9% vs 26.7%
ICU time: 6 vs 3 days
MCS duration: 2.5 days

Absolute 12.7% reduction in mortality- NNT 8

tMCS Meta-analysis

Temporary mechanical circulatory support in infarct-related cardiogenic shock: an individual patient data meta-analysis of randomised trials with 6-month follow-up

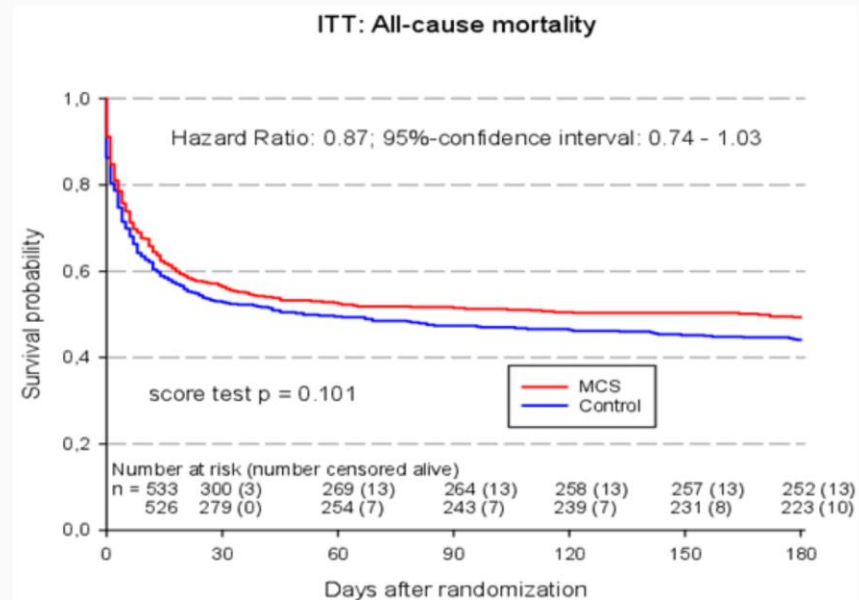


Holger Thiele*, Jacob E Møller*, Jose P S Henriques, Margriet Bogerd, Melchior Seyfarth, Daniel Burkhoff, Petr Ostadal, Richard Rokaya, Jan Belohlavek, Steffen Massberg, Marcus Flather, Matthias Hochadel, Steffen Schneider, Steffen Desch, Anne Freund, Hans Eiskjær, Norman Mangner, Janine Pöss, Amin Polzin, P Christian Schulze, Carsten Skurk, Uwe Zeymer†, Christian Hassager†, on behalf of the MCS Collaborator Scientific Group†

>1000 pts

9 RCTs included

6-Month Mortality - MCS versus no MCS



Lancet 2024 Sep 14;404(10457):1019-1028.

tMCS Meta-analysis

Temporary mechanical circulatory support in infarct-related cardiogenic shock: an individual patient data meta-analysis of randomised trials with 6-month follow-up



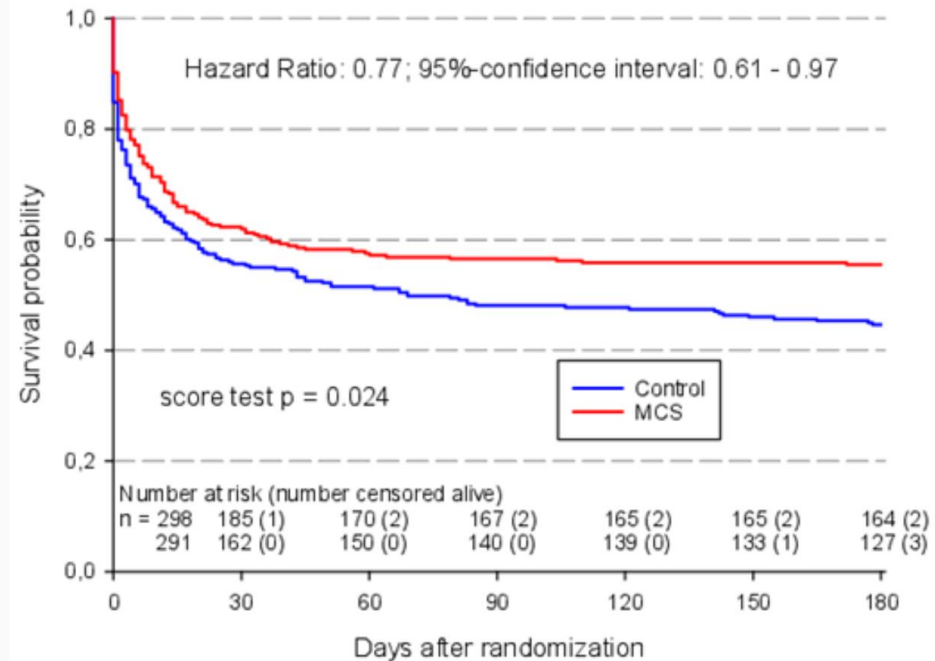
>1000 pts

9 RCTs included

Holger Thiele*, Jacob E Møller*, Jose P S Henriques, Margriet Bogerd, Melchior Seyfarth, Daniel Burkhoff, Petr Ostadal, Richard Rokytka, Jan Belohlavek, Steffen Massberg, Marcus Flather, Matthias Hochadel, Steffen Schneider, Steffen Desch, Anne Freund, Hans Eiskjær, Norman Mangner, Janine Pöss, Amin Polzin, P Christian Schulze, Carsten Skurk, Uwe Zeymer†, Christian Hassager†, on behalf of the MCS Collaborator Scientific Group†

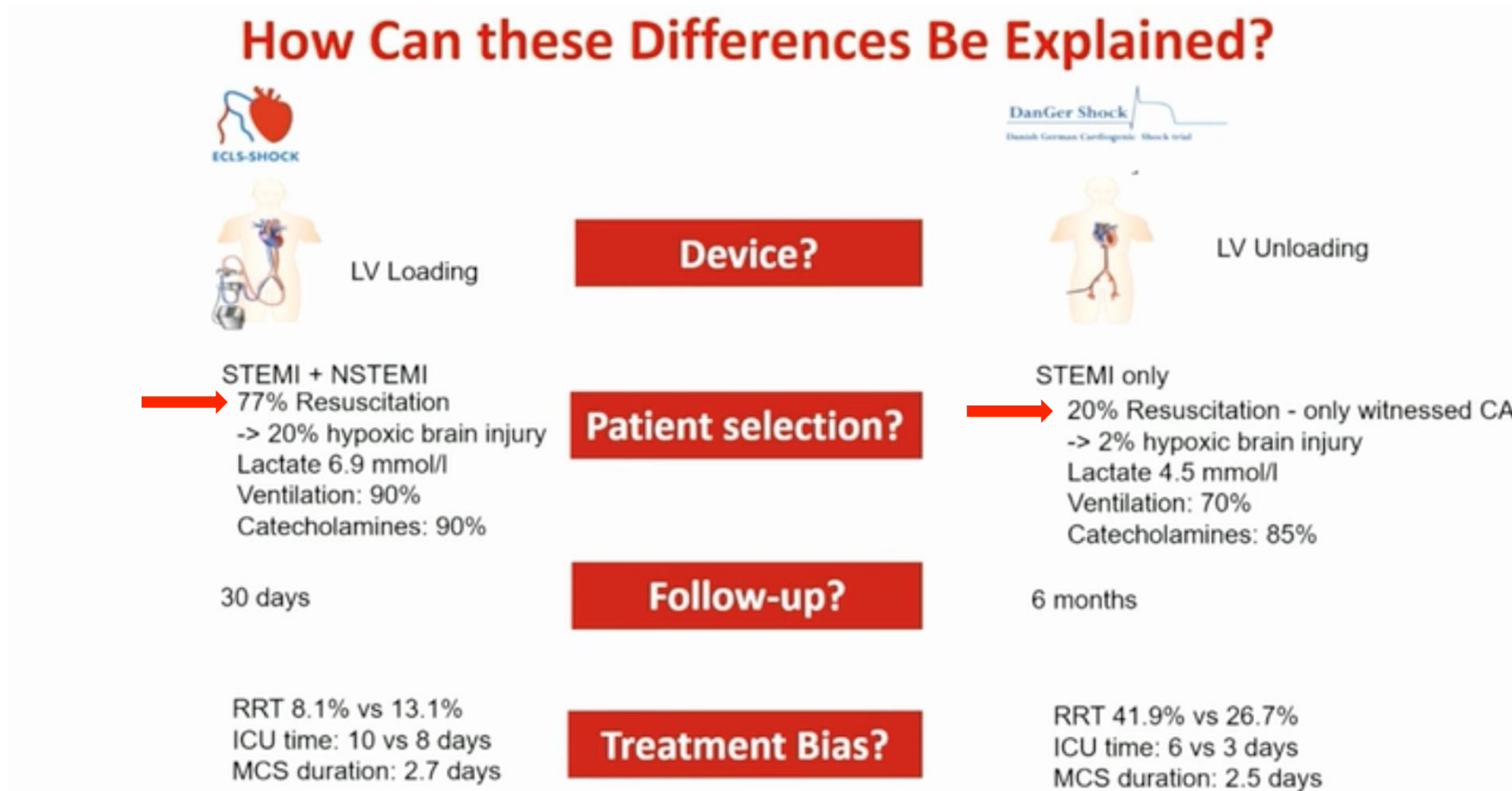
6-Month Mortality – Patient Selection MCS versus no MCS

Patient selection to STEMI no risk of hypoxic brain injury: All-cause mortality



Lancet 2024 Sep 14;404(10457):1019-1028.

ECMO VA Vs Microaxial Flow Pump in CS

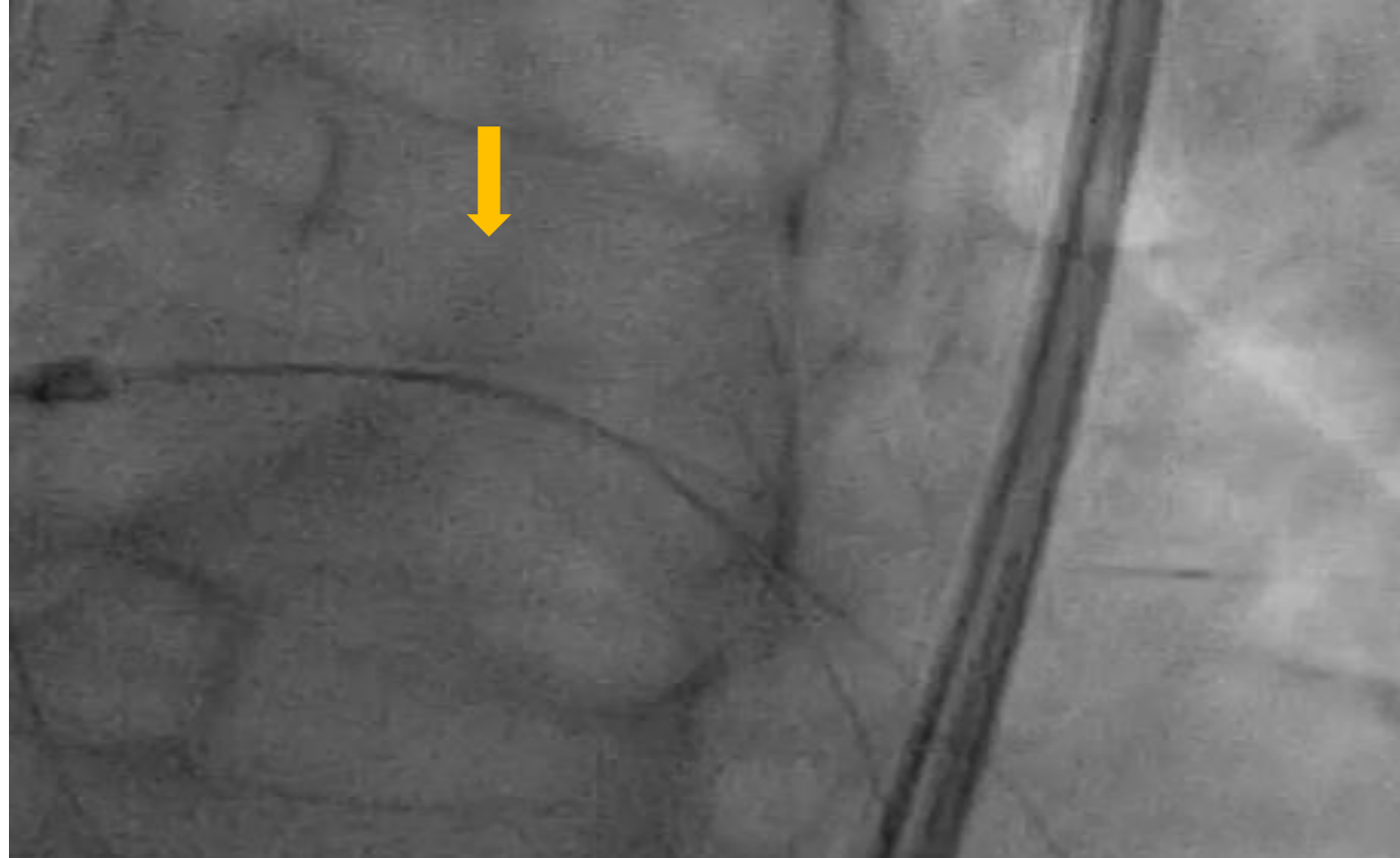


Cardiac Catheterization (*LM-LAD*)

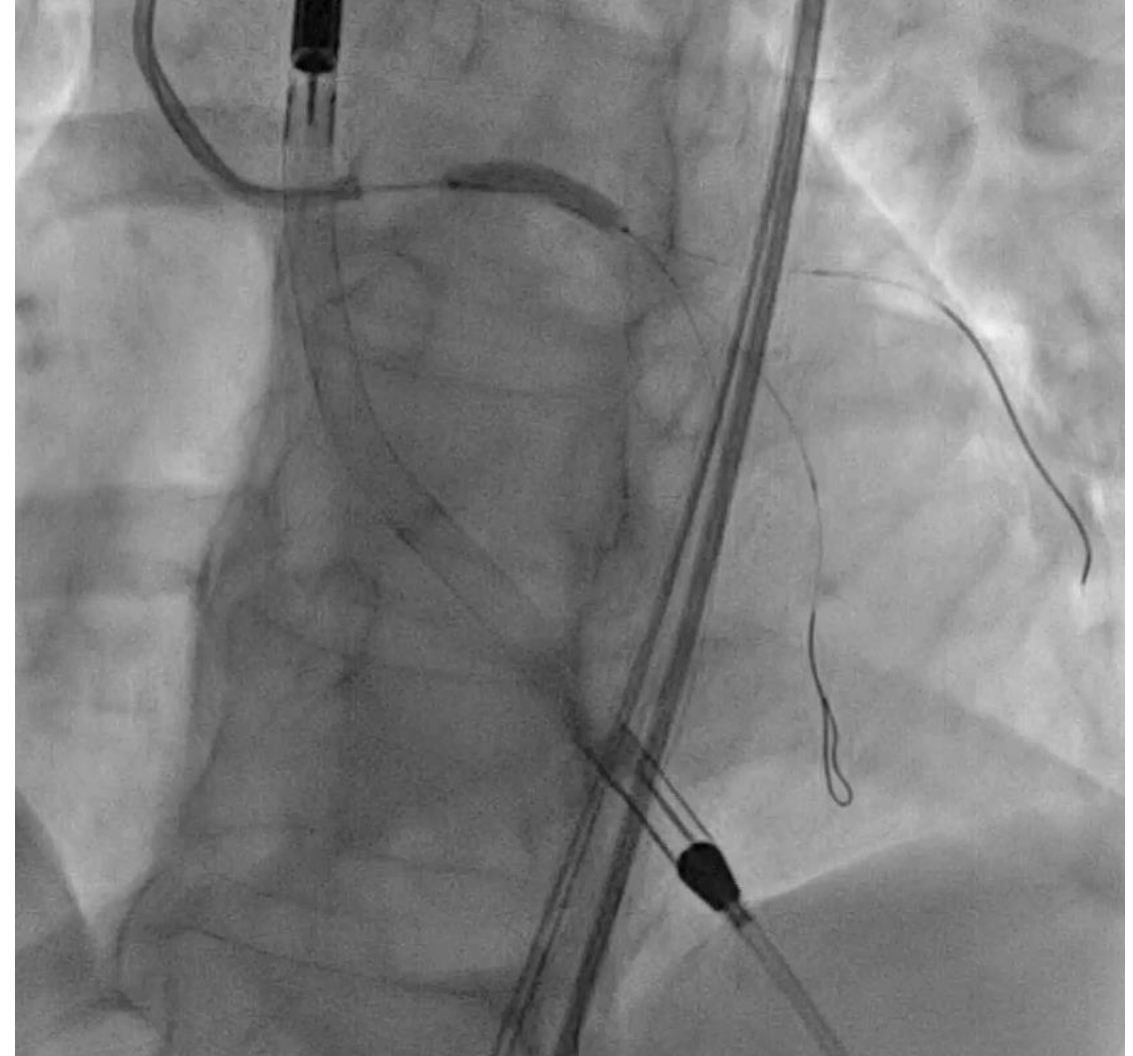
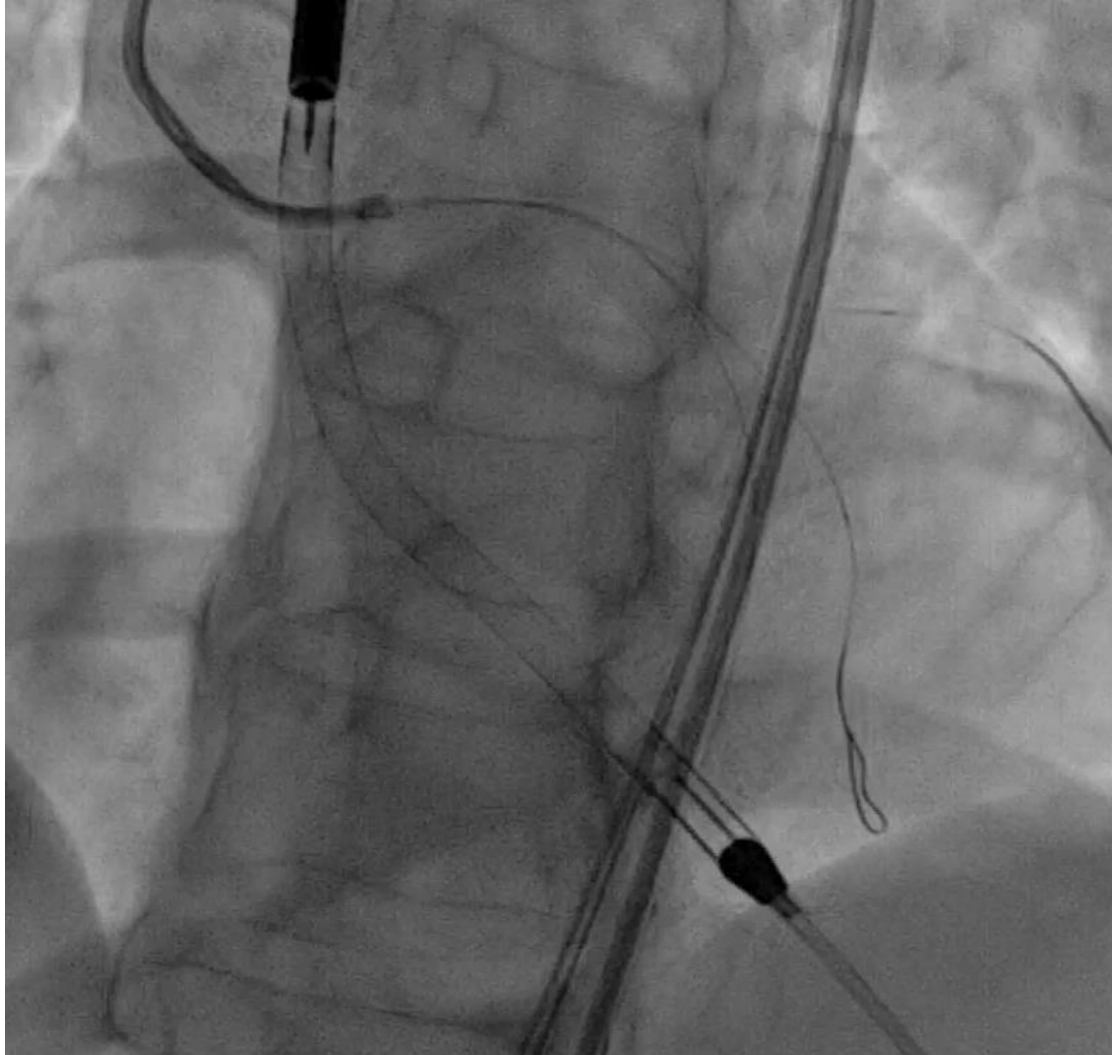
Coronary
dissection progression



LM hematoma
TIMI II flow

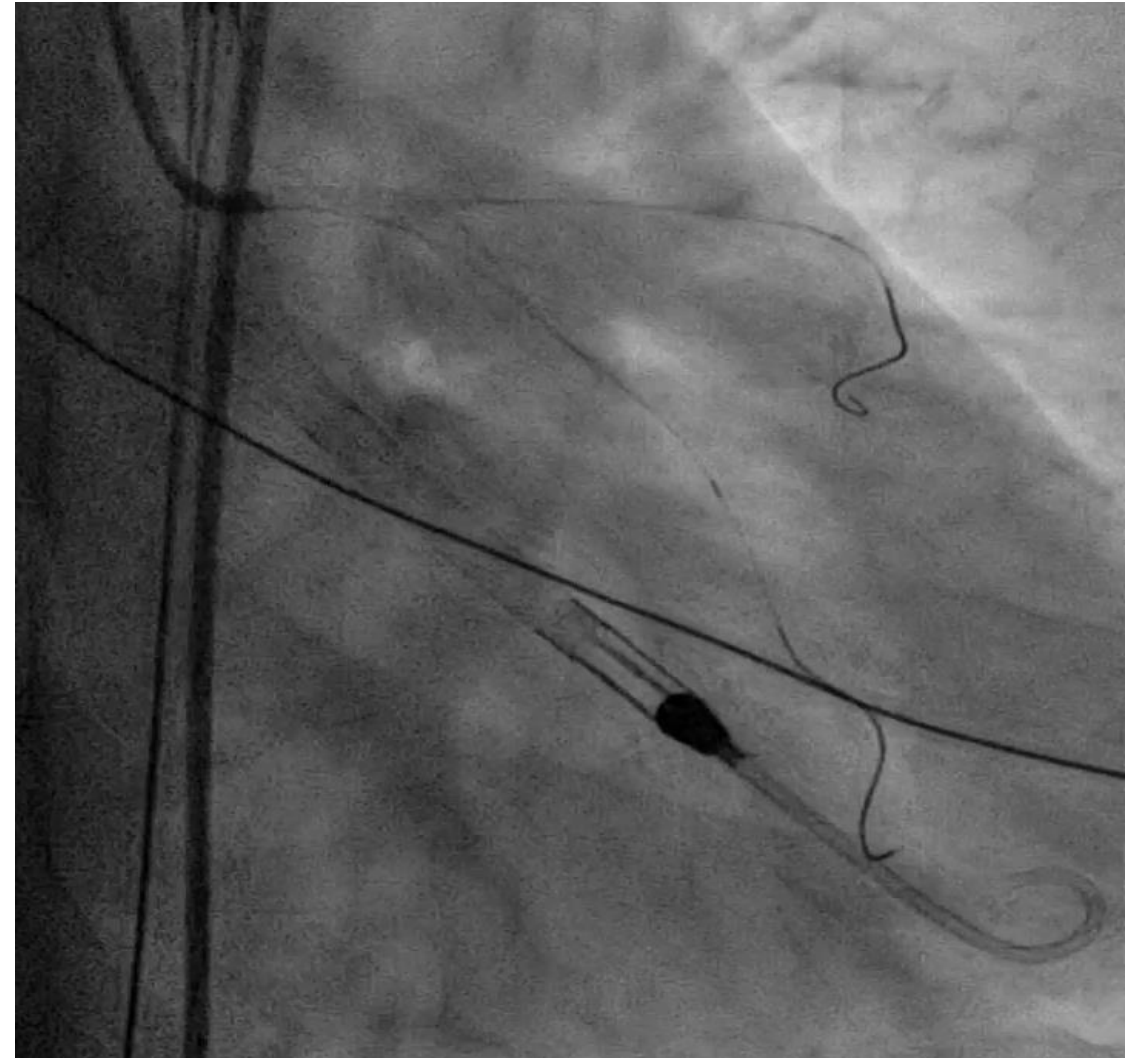
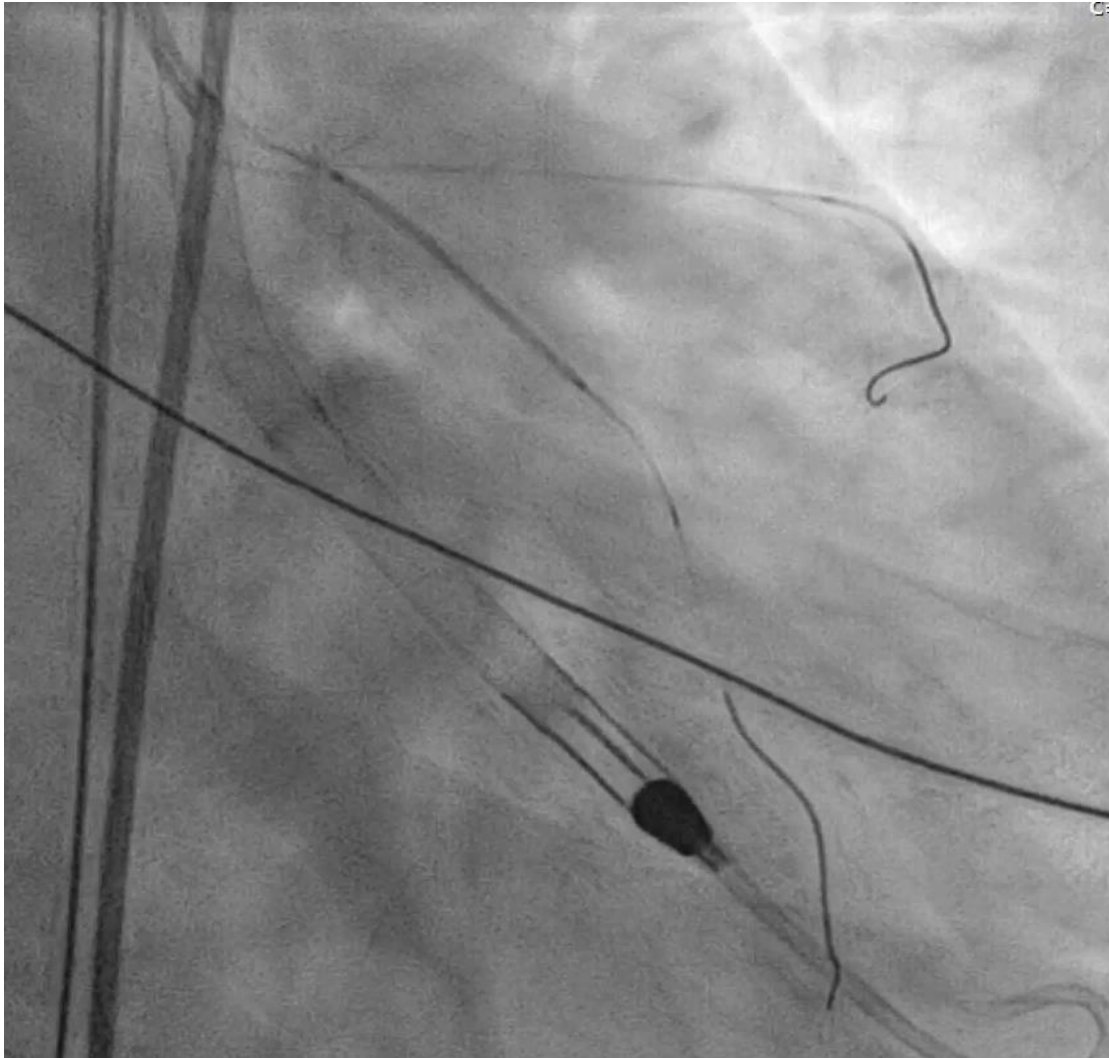


Cardiac Catheterization (*TCI-LAD*)

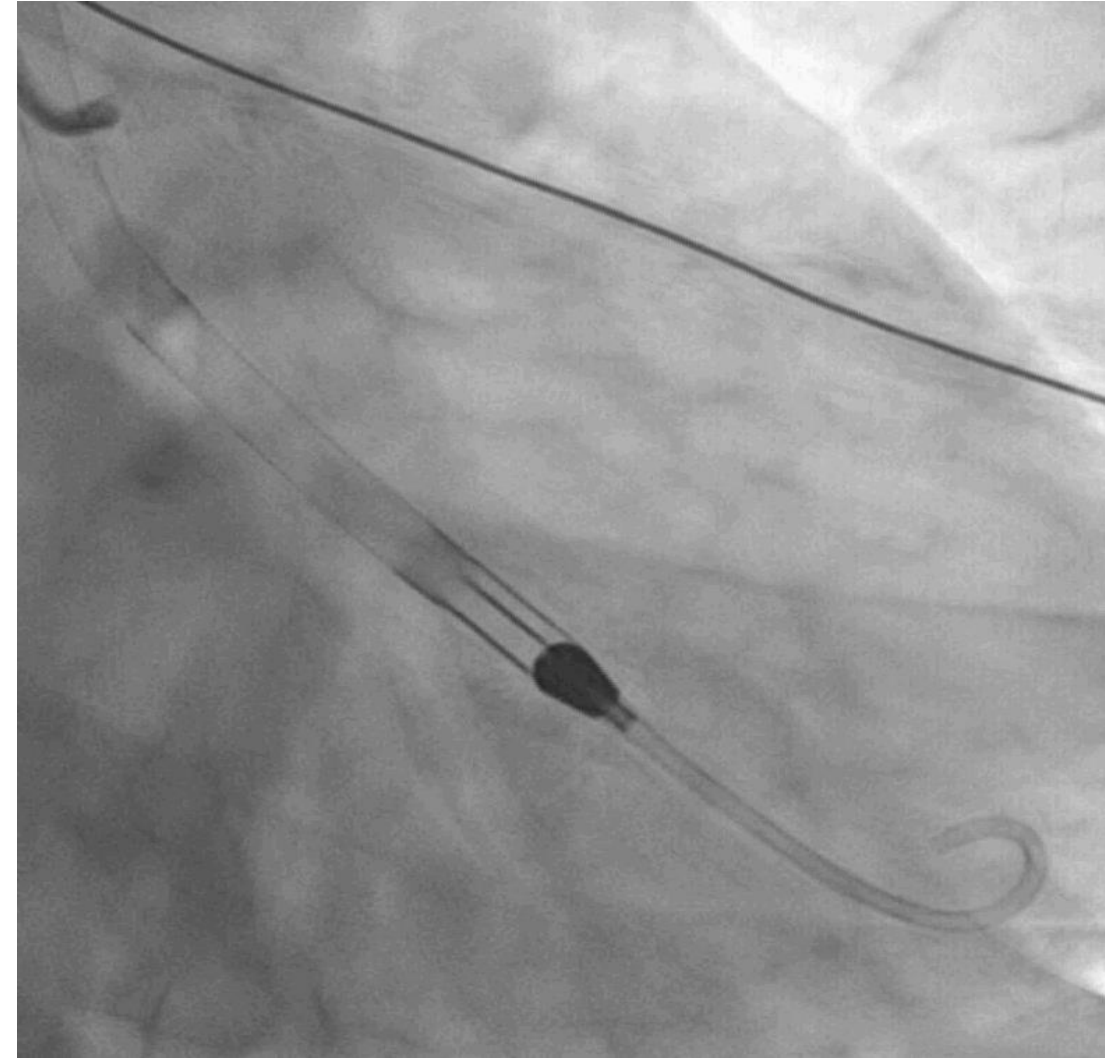
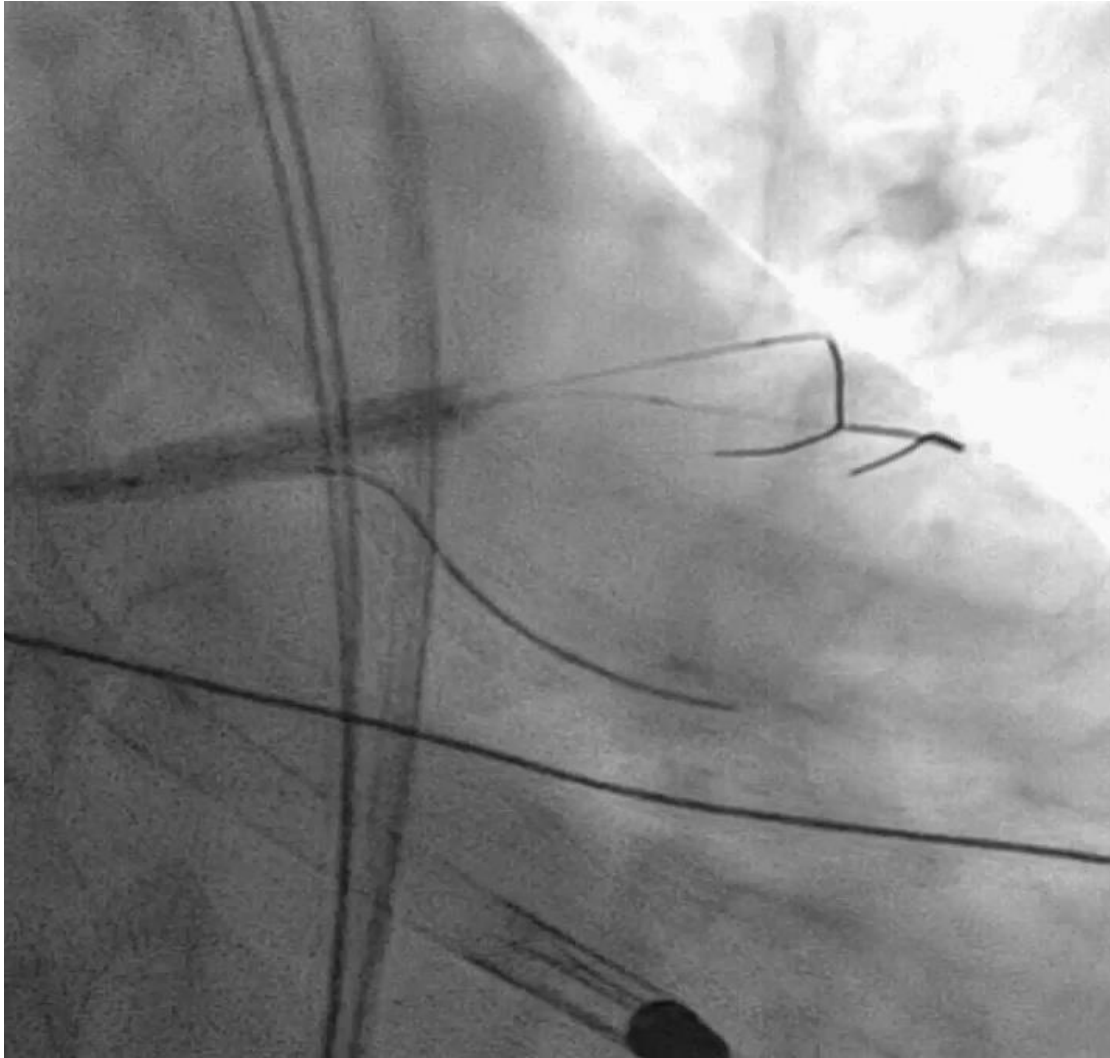


Synergy 3/20mm

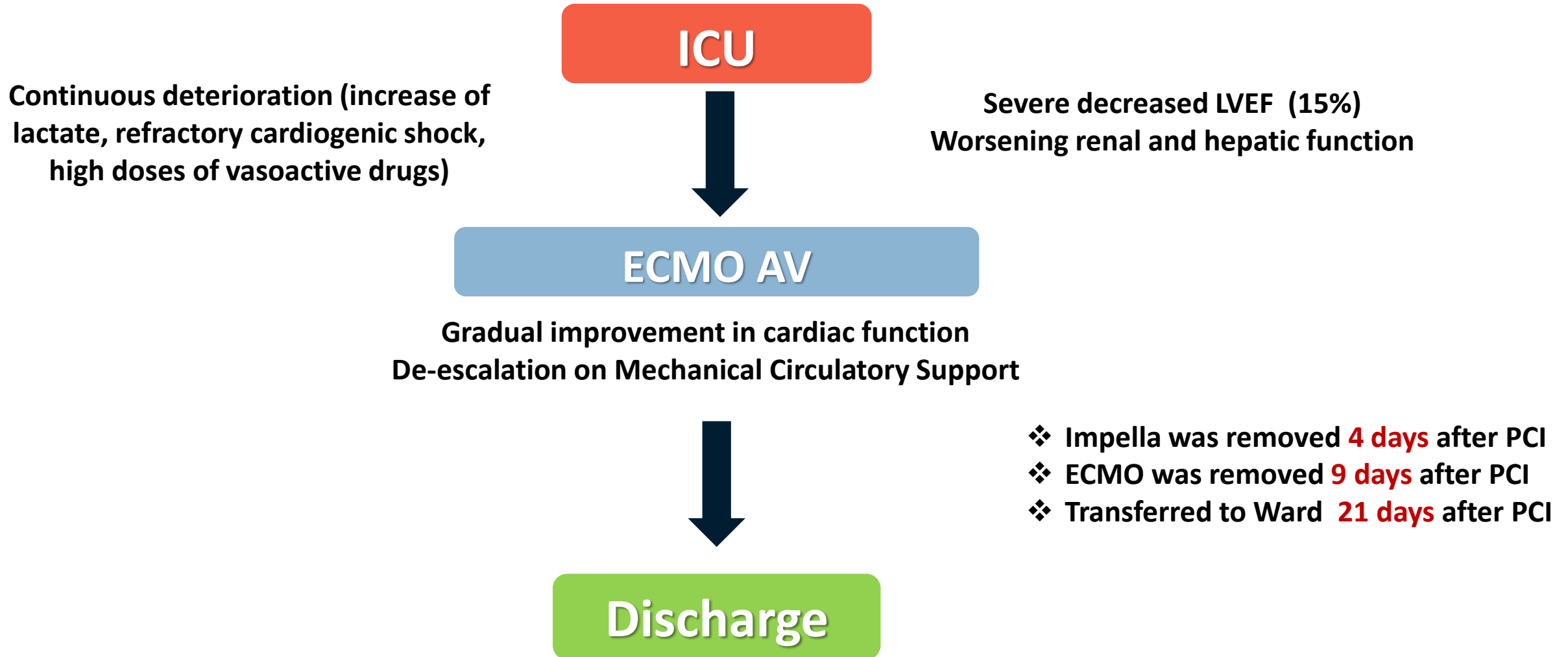
Cardiac Catheterization (*TCl-Cx*)



Final Result



Hospital Course



FOLLOW UP (3 weeks)



LVEF 64%



No valvulopathies

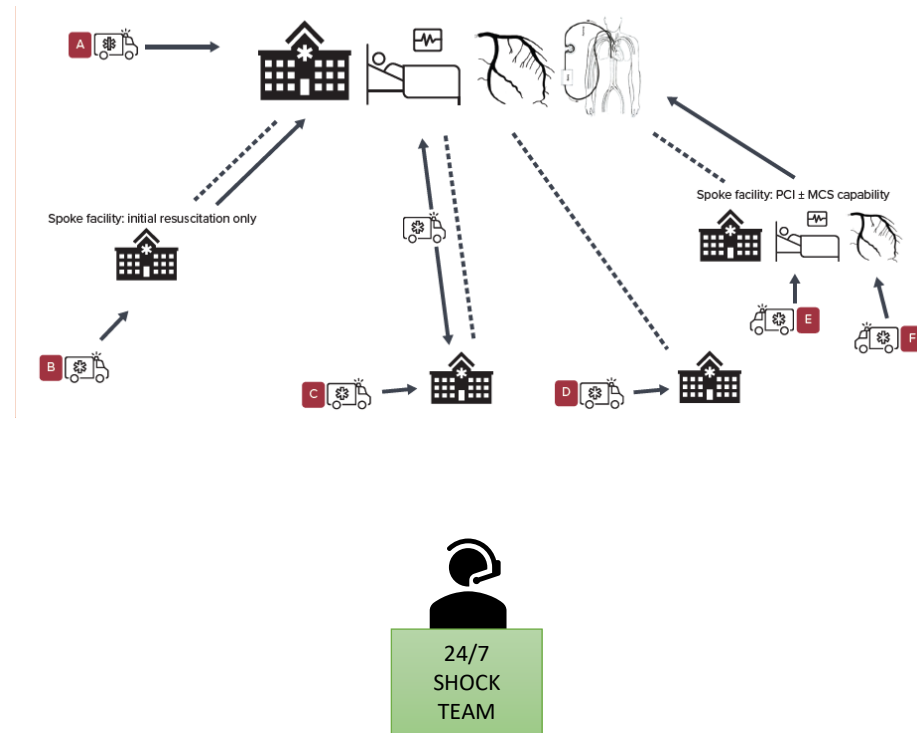
CÓDIGO SHOCK CARDIOGÉNICO

Atención al paciente en Shock Cardiogénico en el Servicio Gallego de Salud.



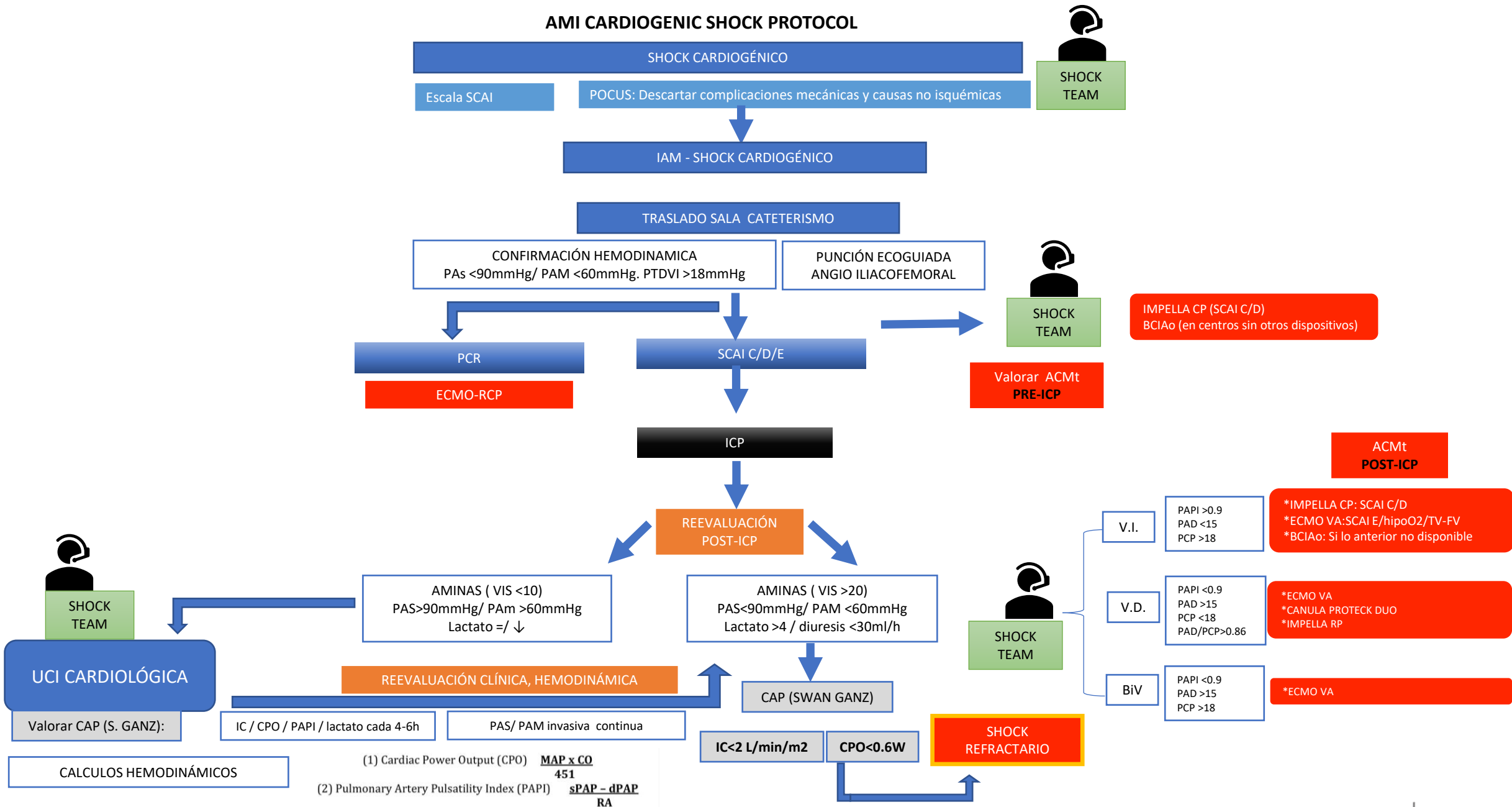
Grupo de trabajo Código Shock Cardiogénico.

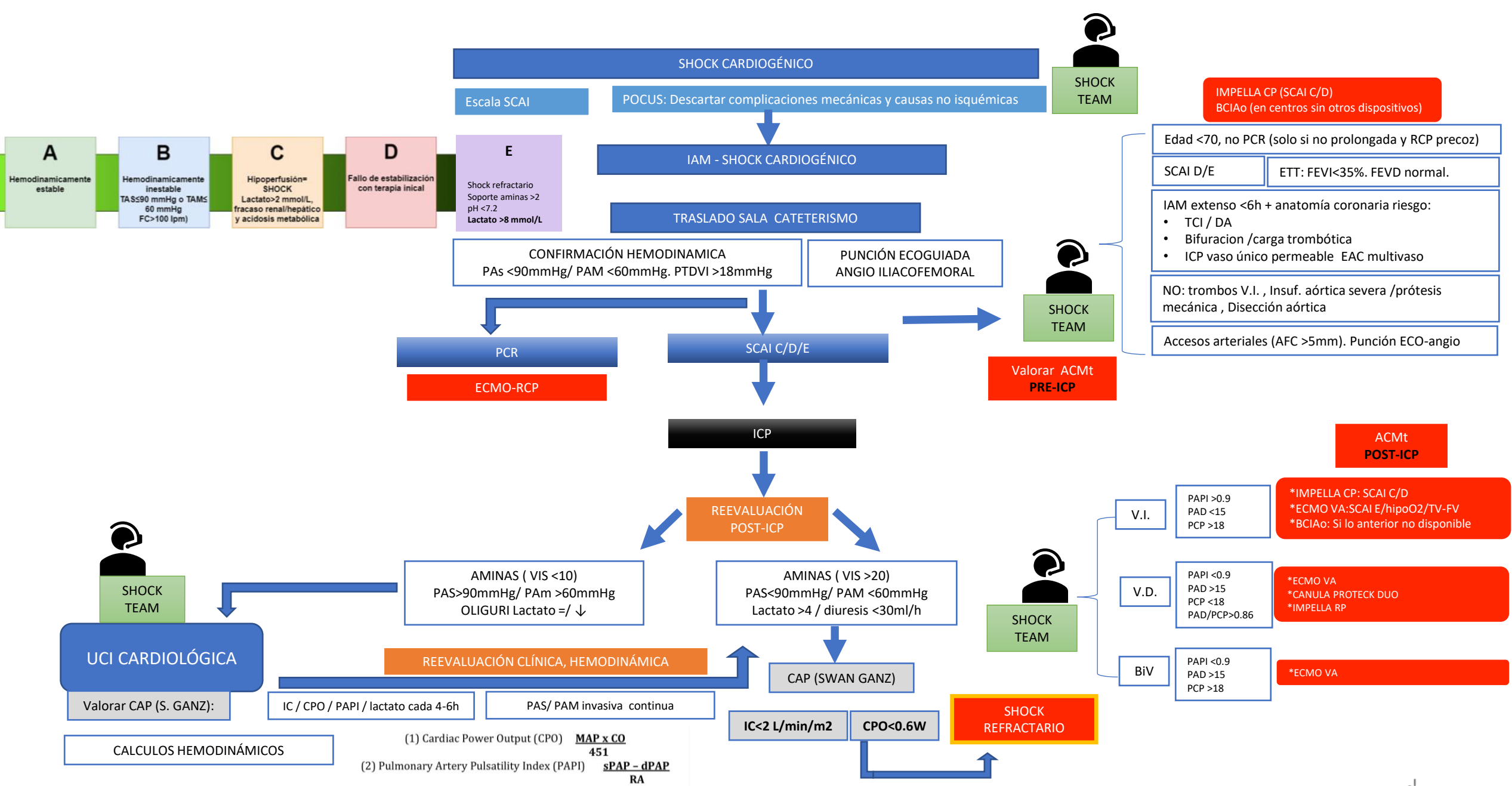
SHOCK CENTER
High-volumen
24/7 Cath Lab, Intensive Unit, on call SHOCK TEAM, Cardiac Surgery, Advanced MCS



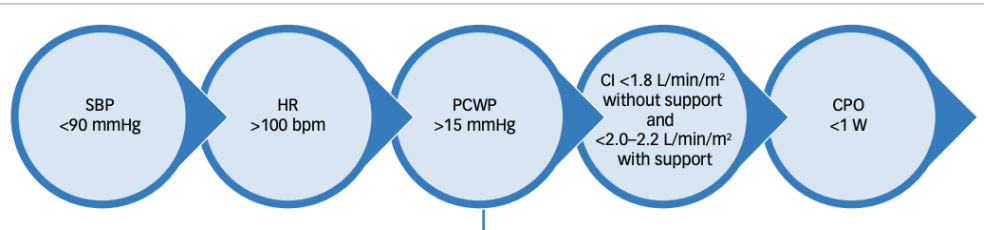
CARDIOGENIC SHOCK PROTOCOL

AMI CARDIOGENIC SHOCK PROTOCOL



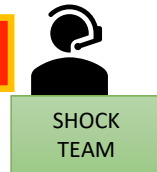


ACS REFRACTORY CS POST PCI



INICIATE or tMCS ESCALATION

HUB SHOCK CENTER TRANSFER



SHOCK TEAM

High-volumen

High dose inotropes (VIS >20)
pH <7.2 / Lactate >4 /Urine output <30ml/h

Bridge to decision/ recovery / LAVD-TX

<70 y/o

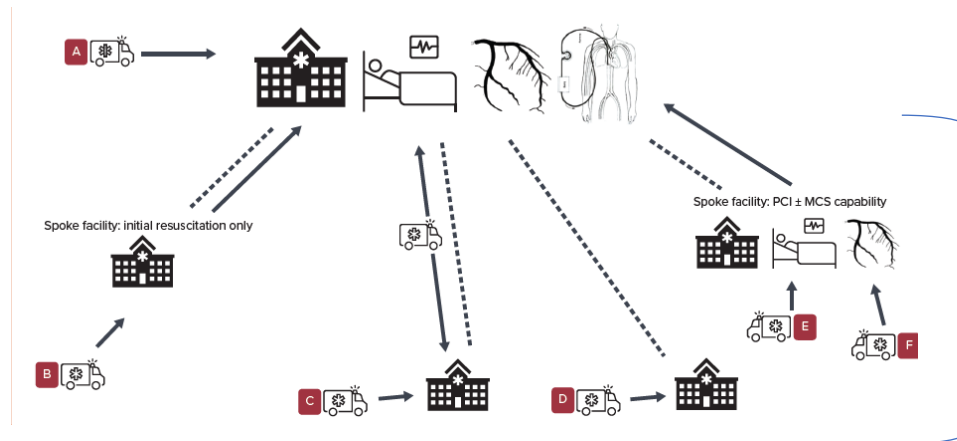
No comatose / short cardiac arrest

No contraindications for tMCS

E
Extremis

D
Deteriorating

24/7 Cath Lab, Intensive Unit, on call SHOCK TEAM, Cardiac Surgery, Advanced MCS



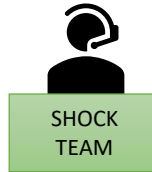
Spoke (low-volume) centers: triage, Culprit PCI, short MCS

ACS REFRACTORY CS



INICIATE
or
tMCS ESCALATION

HUB SHOCK CENTER
TRANSFER



The GOLDEN HOUR: "Time S2S"

E
Extremis

D
Deteriorating

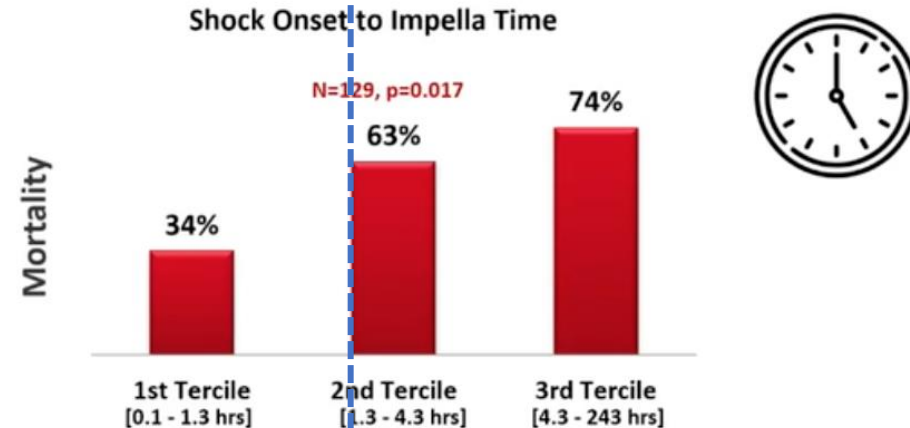
TIME DEPENDENT
DECISION

MORTALITY

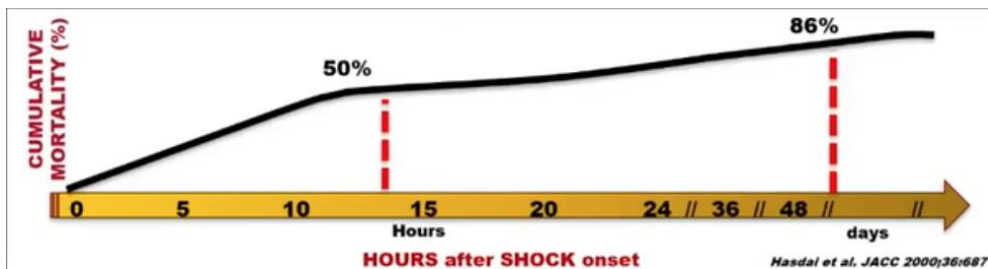
<90 min

>90 min: x2 mortality

Shock Onset to Impella Time



Basir et al. Am J Cardiol. 2017;119:845-851.





Henry T et al. *Circulation* 2023; 47:465–468.



TAKE HOME MESSAGE

- Is tMCS a routine option in CS ? Yes / **No**
- Does tMCS have any mortality benefit? **Yes** / No

But ...

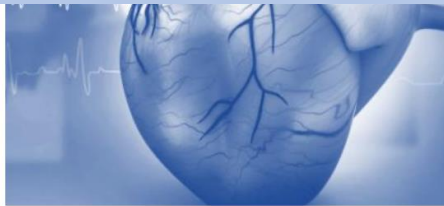
- Attending to RCTs (DANGER trial) and registries (NCSI) experience
 - Only CS very selected patients: Lactate >4, Extense STEMI, LV predominant, no prolonged CPR
 - Device selection according hemodynamics
 - High volumen centers: minimize complications
 - Time dependent therapy: “Time S2S” <90min

REGIONAL SC code programs

CÓDIGO SHOCK CARDIOGÉNICO

MULTIDISCIPLINARY SHOCK TEAMS

Essential for a good quality tMCS program...



Grupo de trabajo Código Shock Cardiogénico.

Mayo de 2024



¡GRACIAS!

