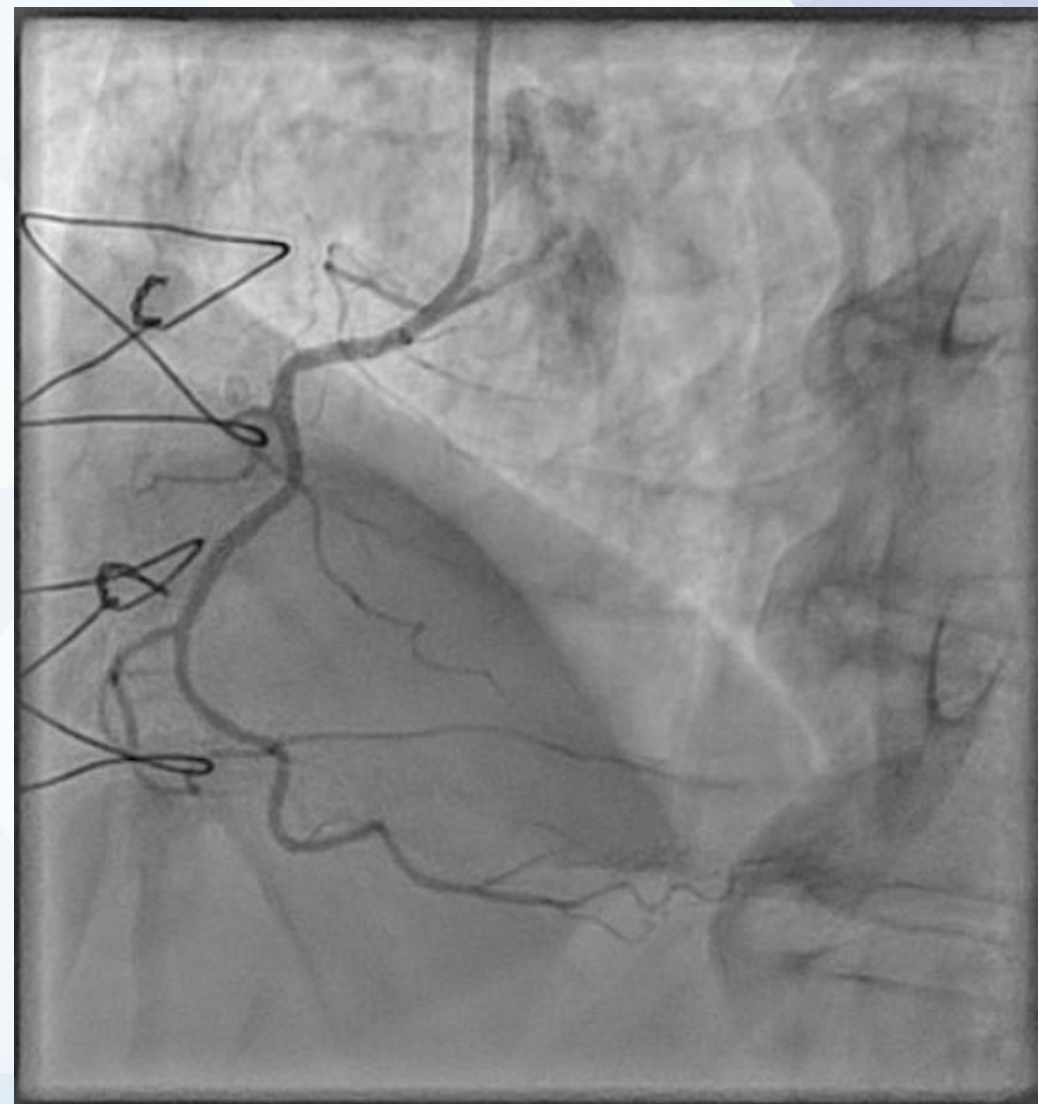
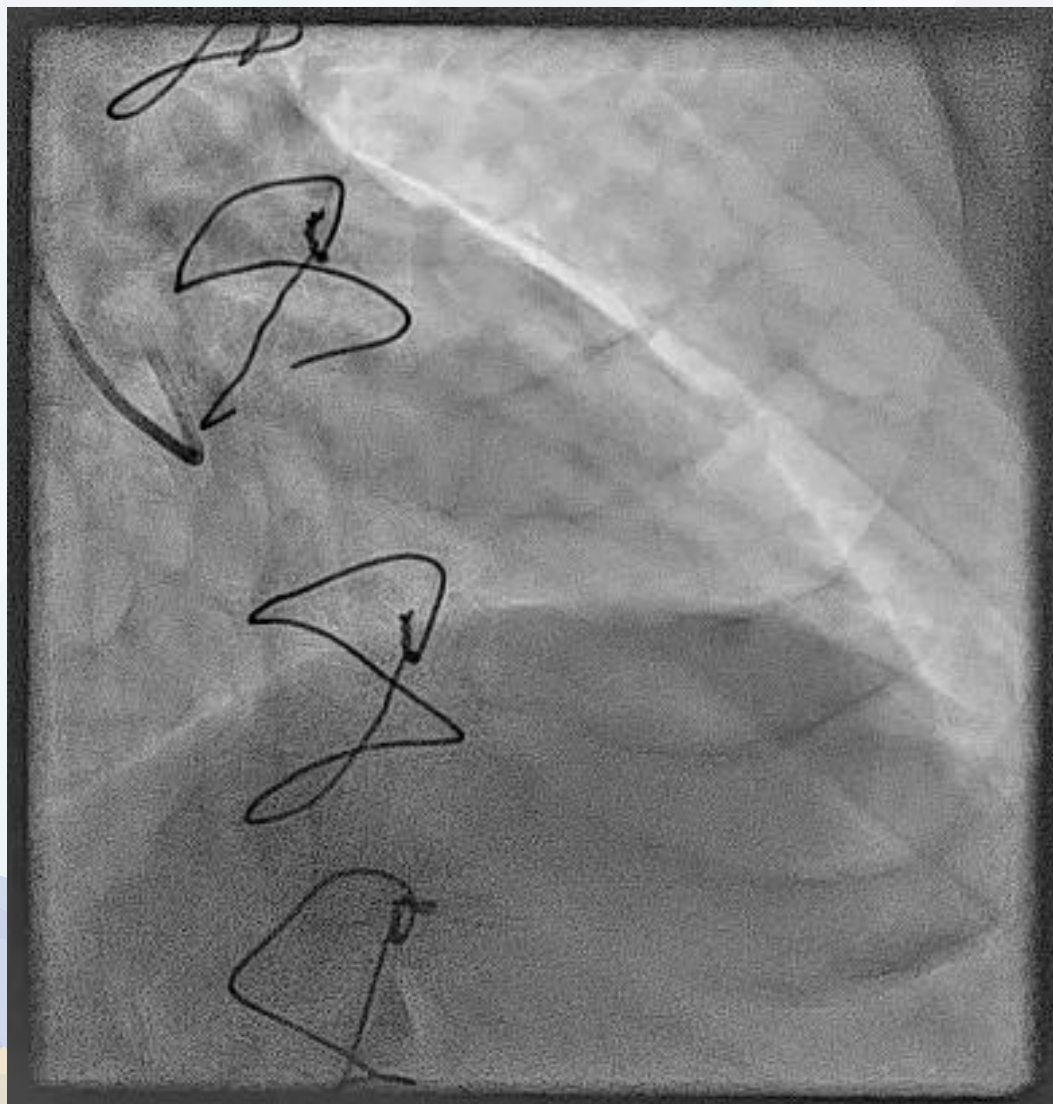


QUÉ HACER Y CÓMO RESOLVER

*IAM y shock
cardiogénico*

*Paula Vela Martín y Juan Francisco Oteo Domínguez
Hospital Universitario Puerta de Hierro*

Octubre 2024



Diciembre 2021

*Varón 55 años
Fumador*

*No HTA, no DM, no DL conocidas
No antecedentes médicos de interés*

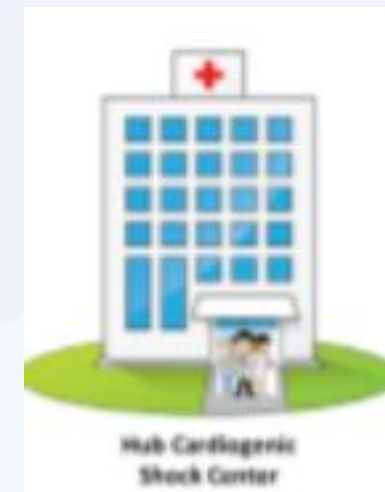
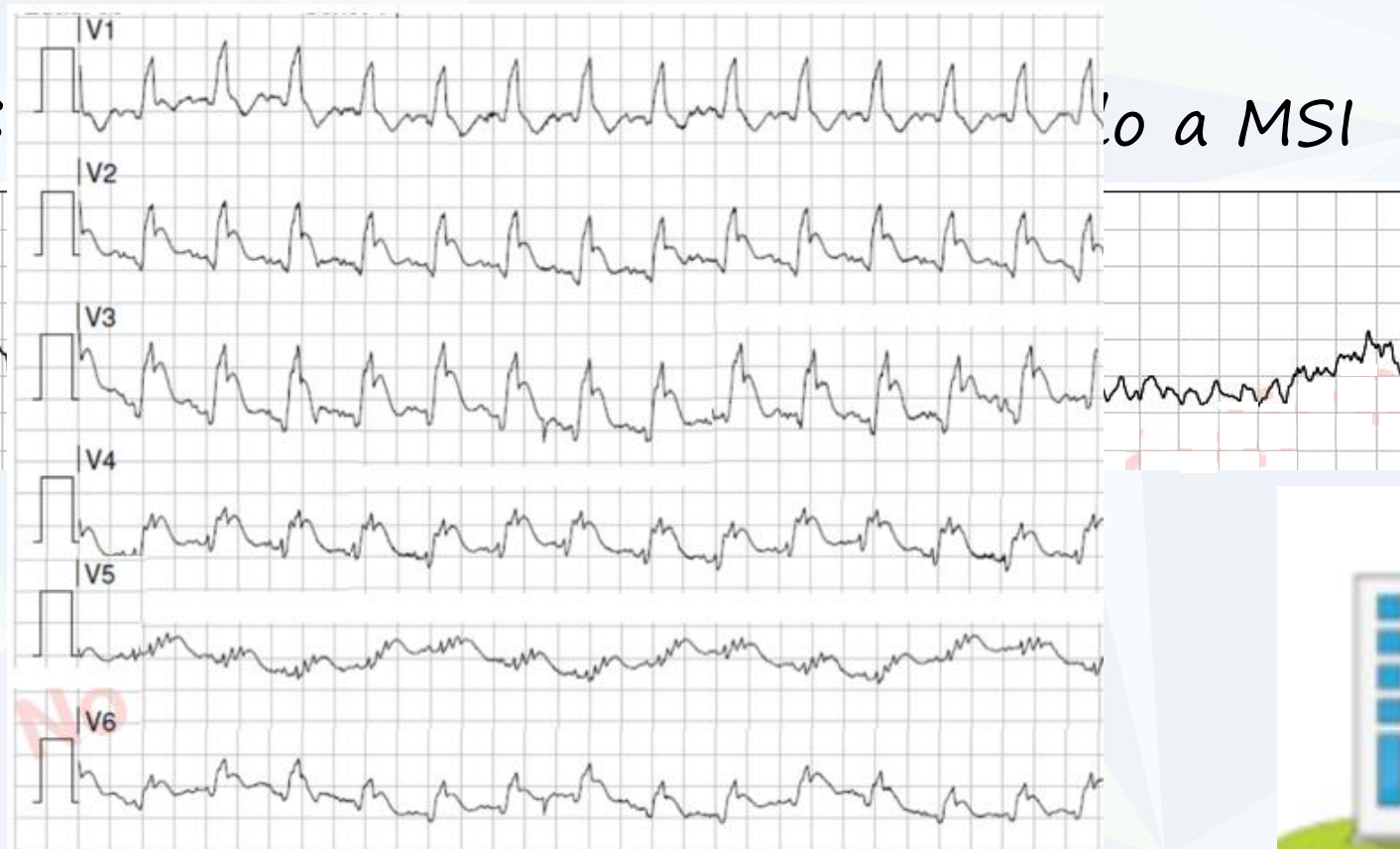
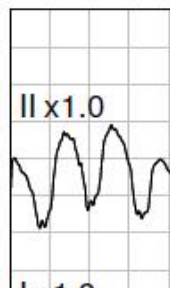
Padre fallecido a los 74 años por IAM

9 de diciembre 2021

Avisa al S

o a MSI

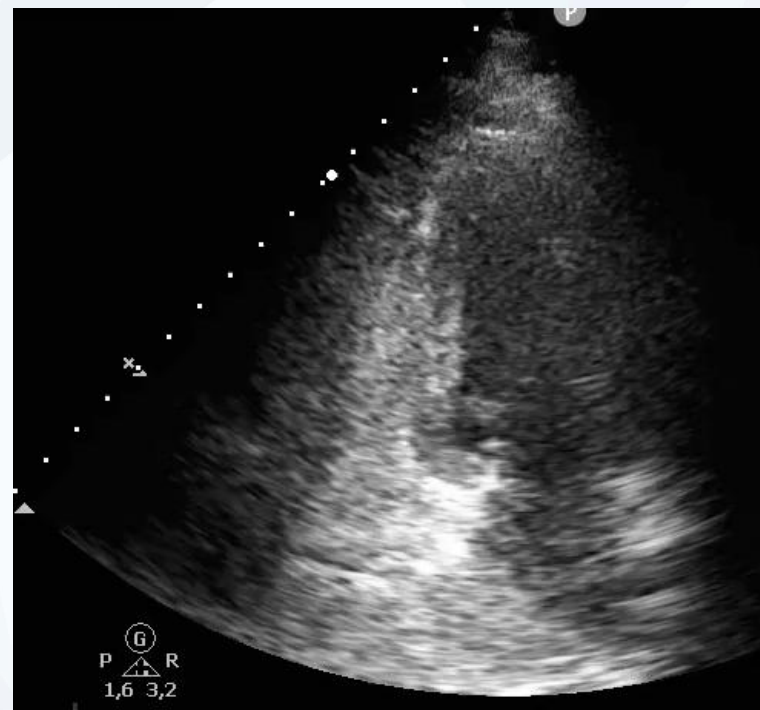
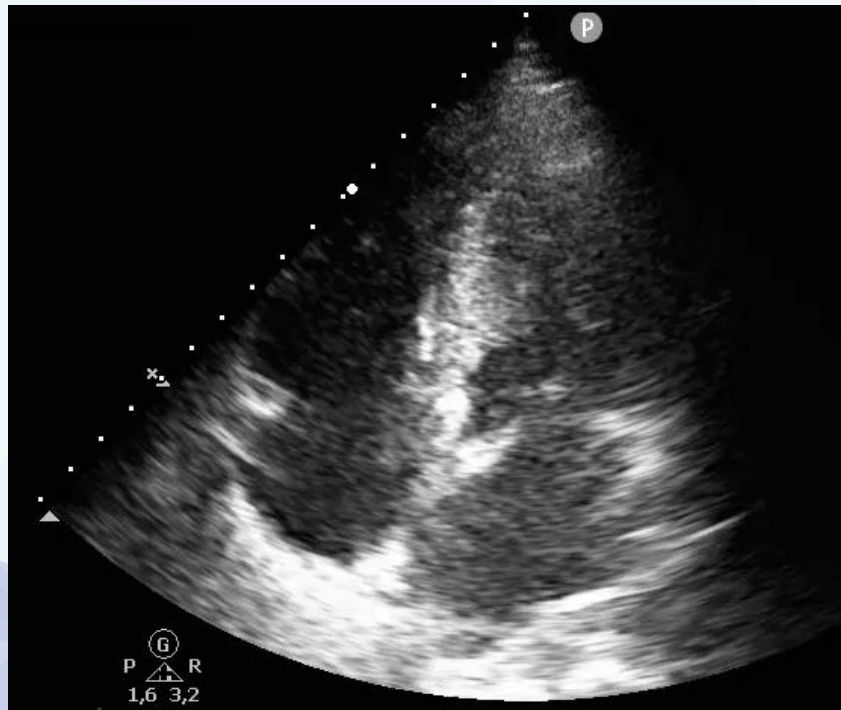
SUMMA:



CÓDIGO IAM

9 de diciembre 2021

A su llegada: TA 80/40 mmHg. FC 100 lpm. SatO2 98% basal
MEG, frialdad cutánea. Eupneico en reposo.



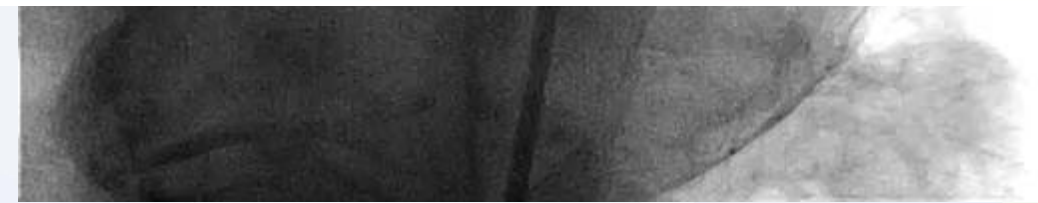
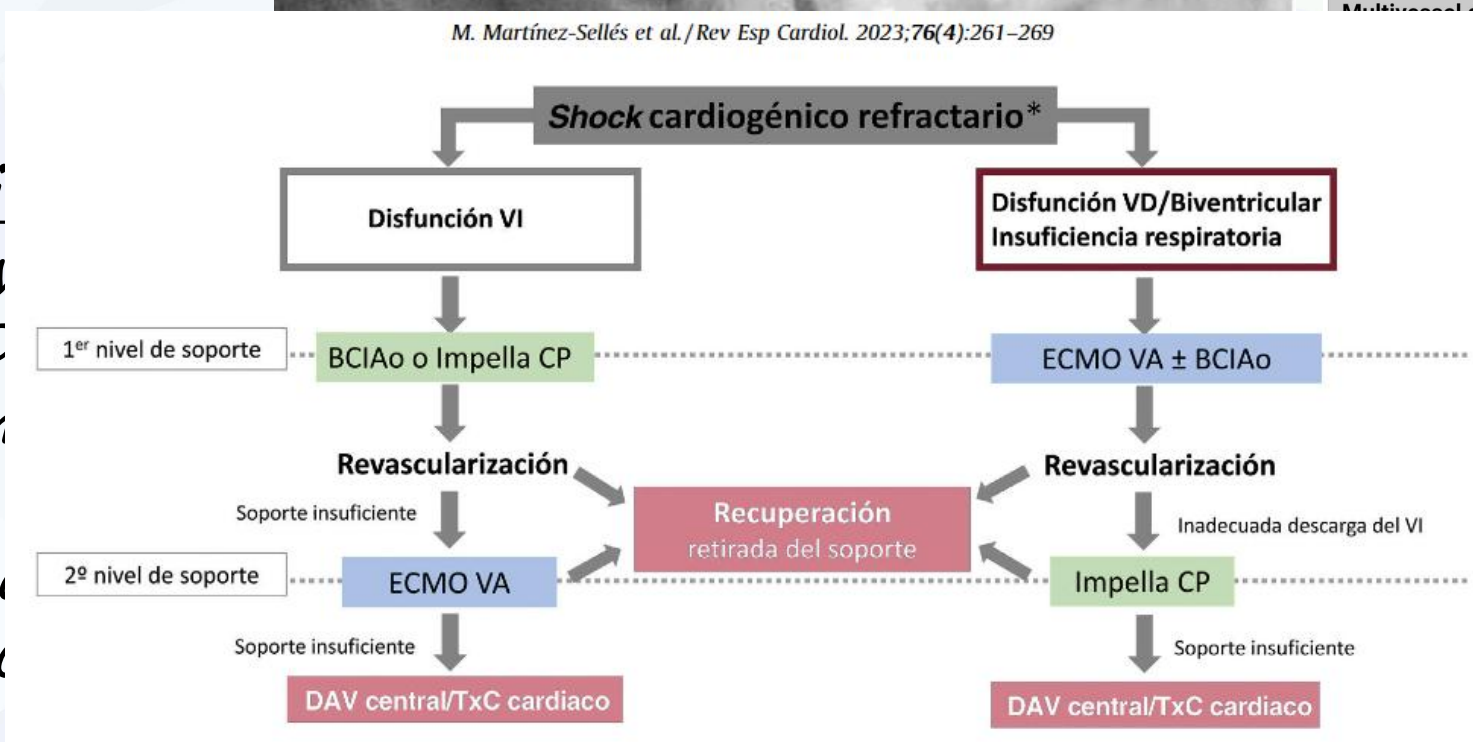
9 de diciembre 2021

SCACEST anterior
Disfunción VI severa
Inestabilidad HD
NA 1 mcg/Kg/v

- Oclusión aguda
- Estenosis crítica
- OCT de CD



M. Martínez-Sellés et al. / Rev Esp Cardiol. 2023;76(4):261-269

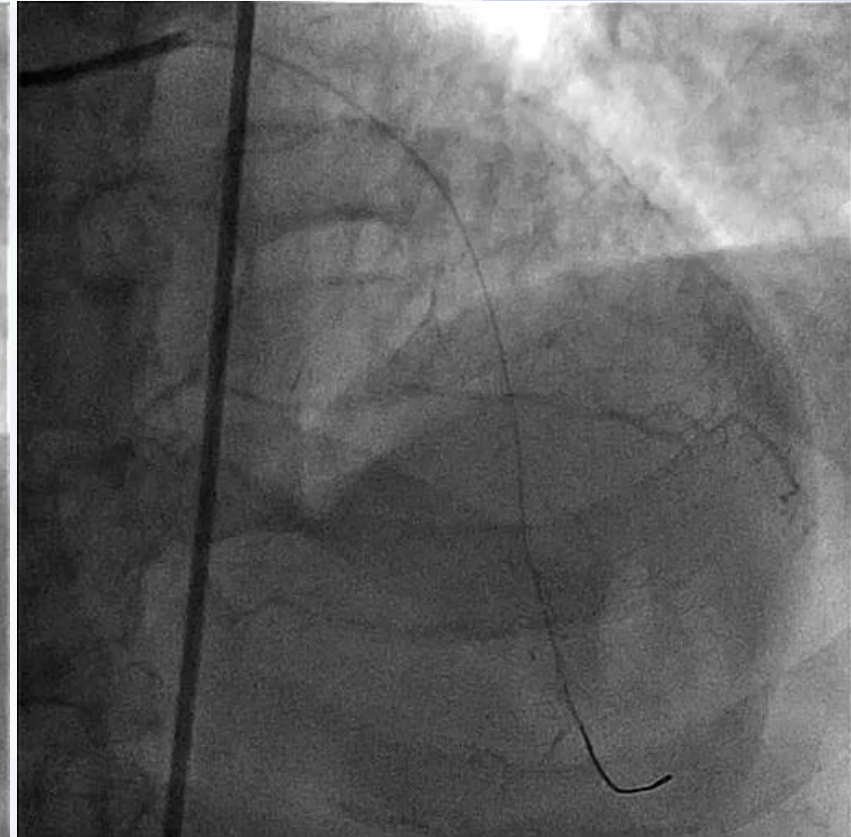
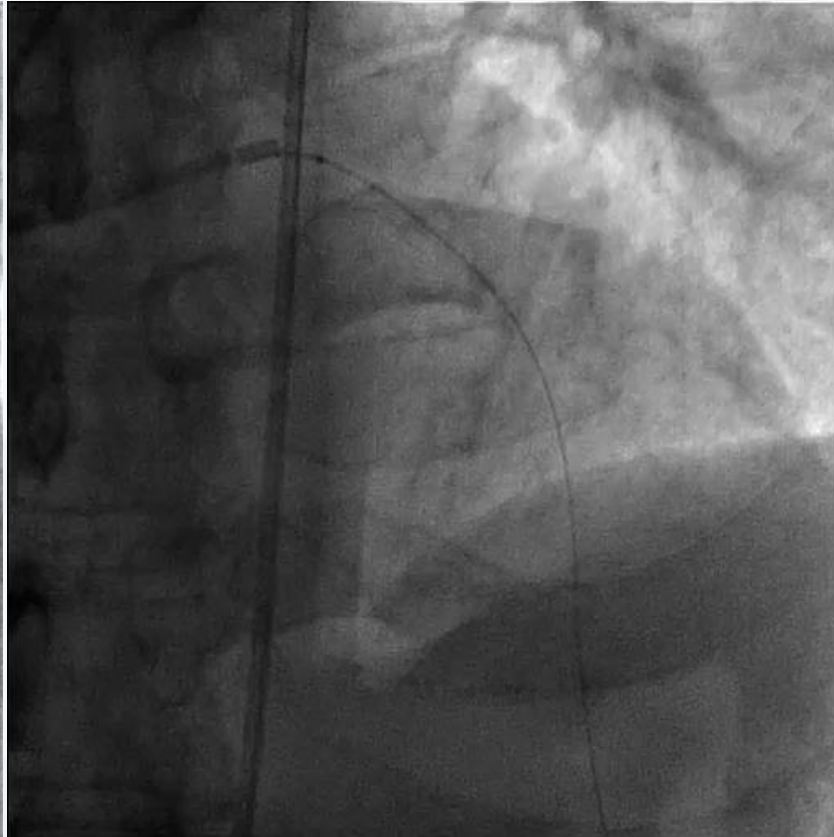
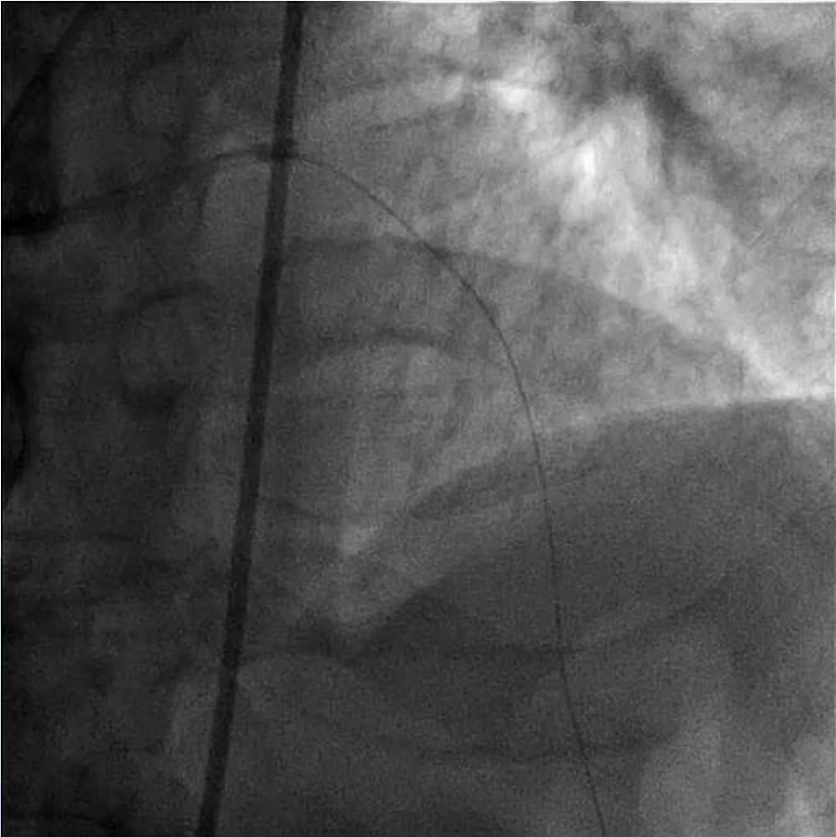


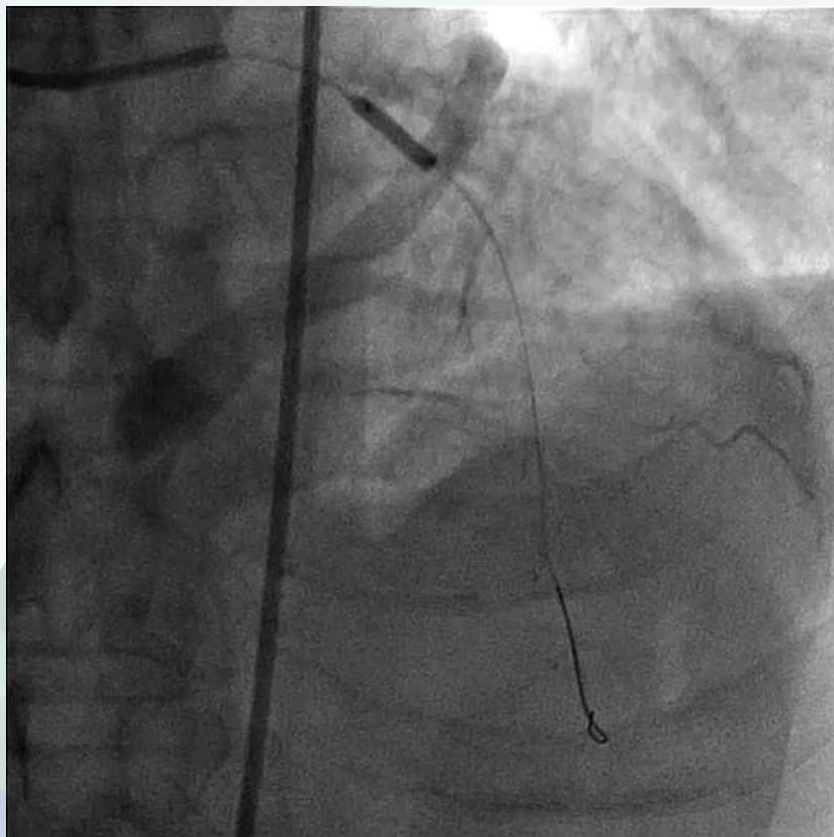
Multisystem disease in ACS patients presenting in car-

ring the index commended.	I	B
on-IRA should	Ila	C

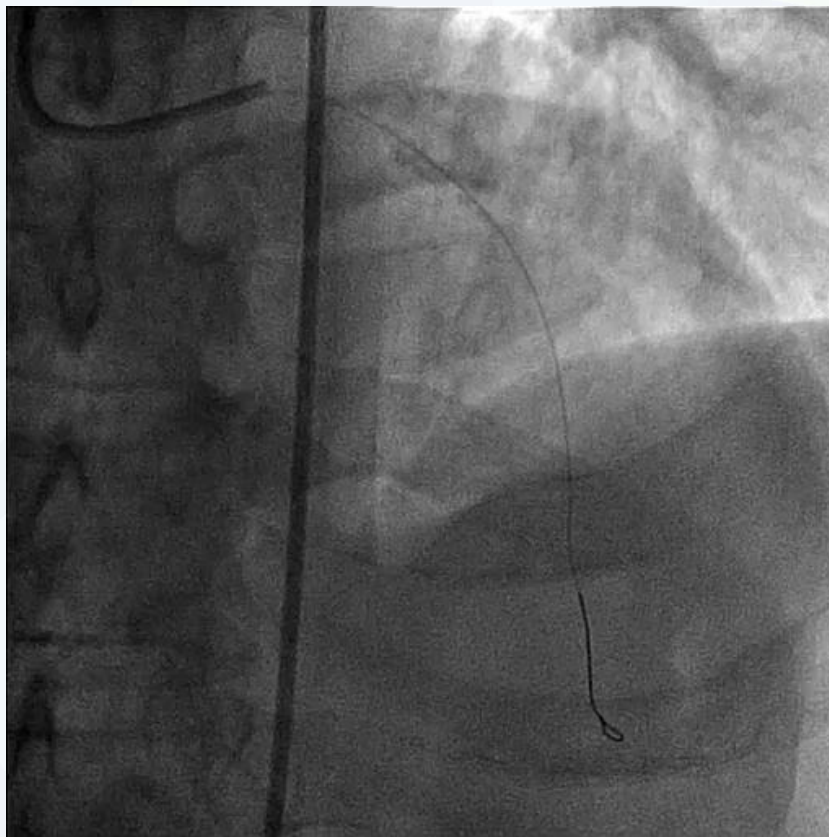


- ICP
- ICP + SMC
- SMC + ICP

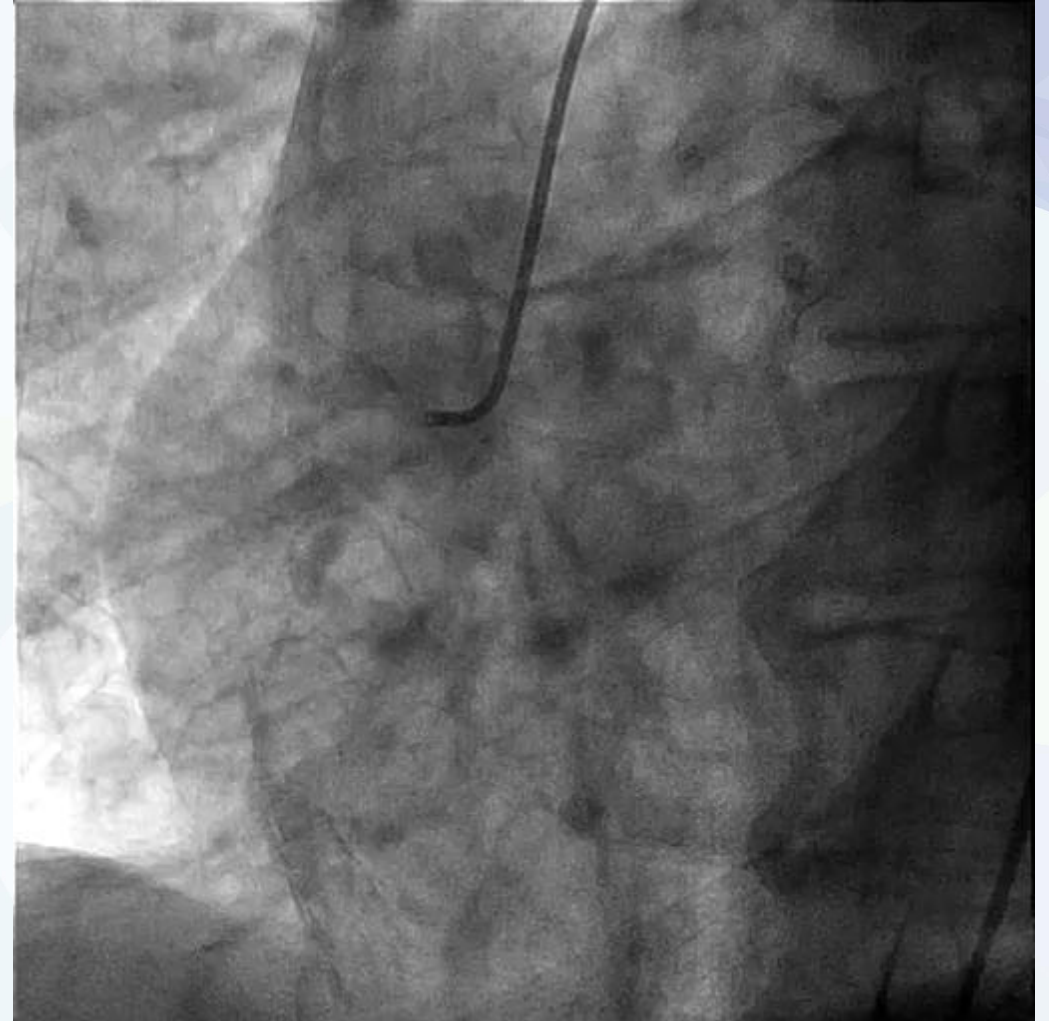
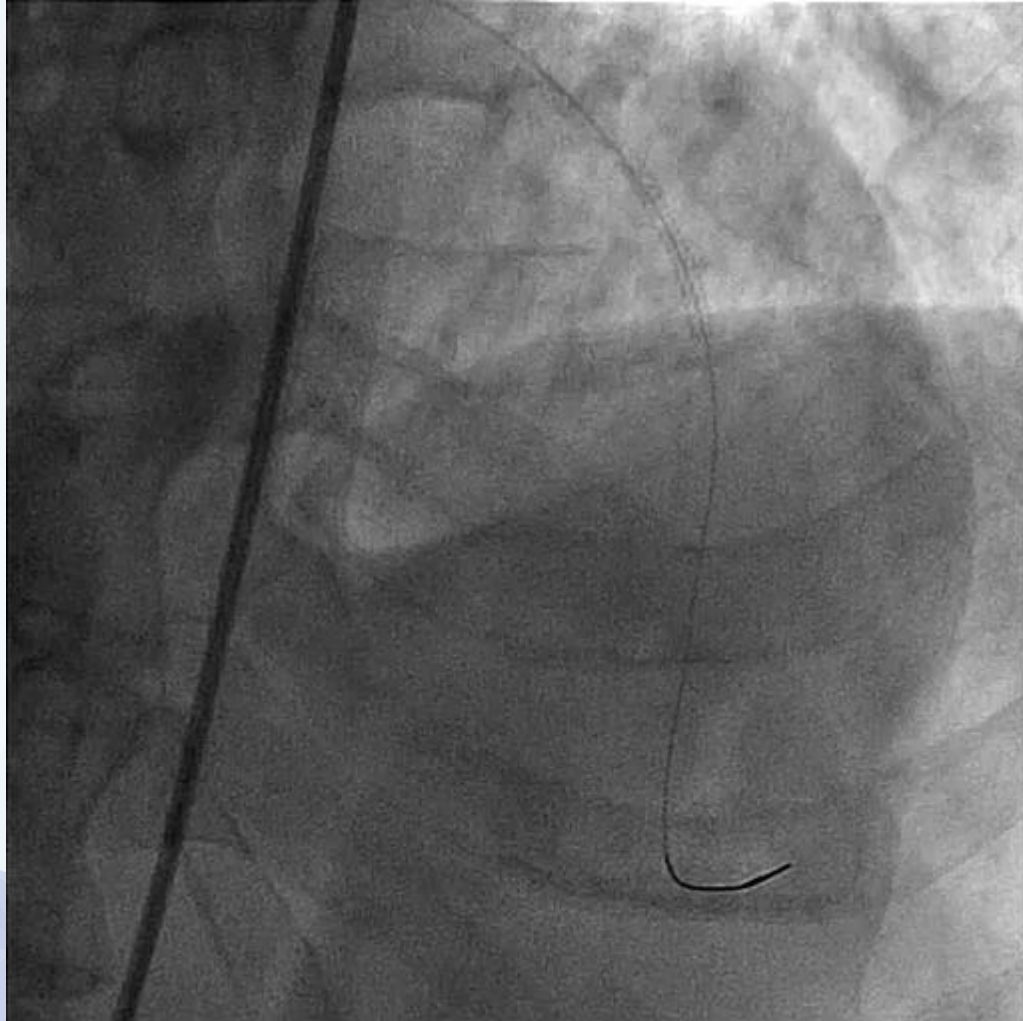




DES 2.75/12mm



DES 2.5/22mm



Acute coronary syndrome with unstable presentation

In patients with ACS and severe/refractory CS, short-term mechanical circulatory support may be considered.

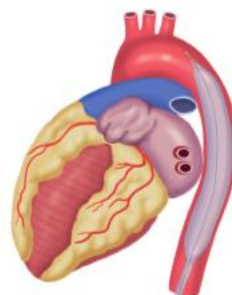
IIb

C

The routine use of an IABP in ACS patients with CS and without mechanical complications is not recommended.

III

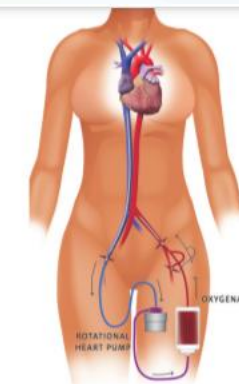
B



BCIAo



Impella®



ECMO VA

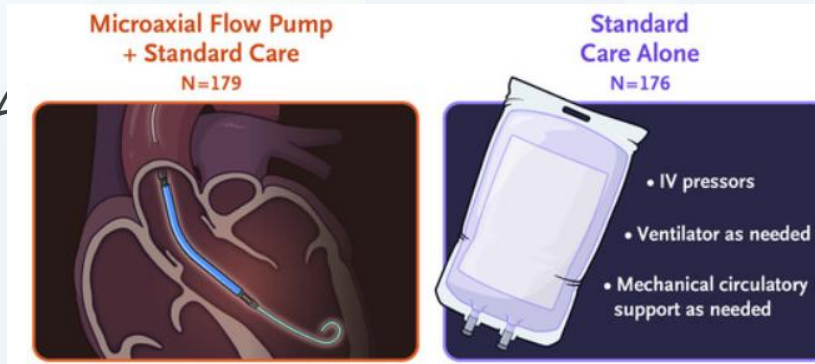
Soporte circulatorio	-	++/+++	++++
Descarga VI	+	+++	-
Ventajas	+ Accesible Facil de implantar	Facil de implantar Ensayos clínicos +	Soporte circulatorio y respiratorio
Contras	Efecto hemodinámico	Hemólisis Complicaciones vasculares	Aumenta postcarga Complicaciones vasculares

ORIGINAL ARTICLE

DanGer Shock trial

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

- Randomizado, multicéntrico (Alemania, Dinamarca, Reino Unido)
- Shock cardiogénico + IA

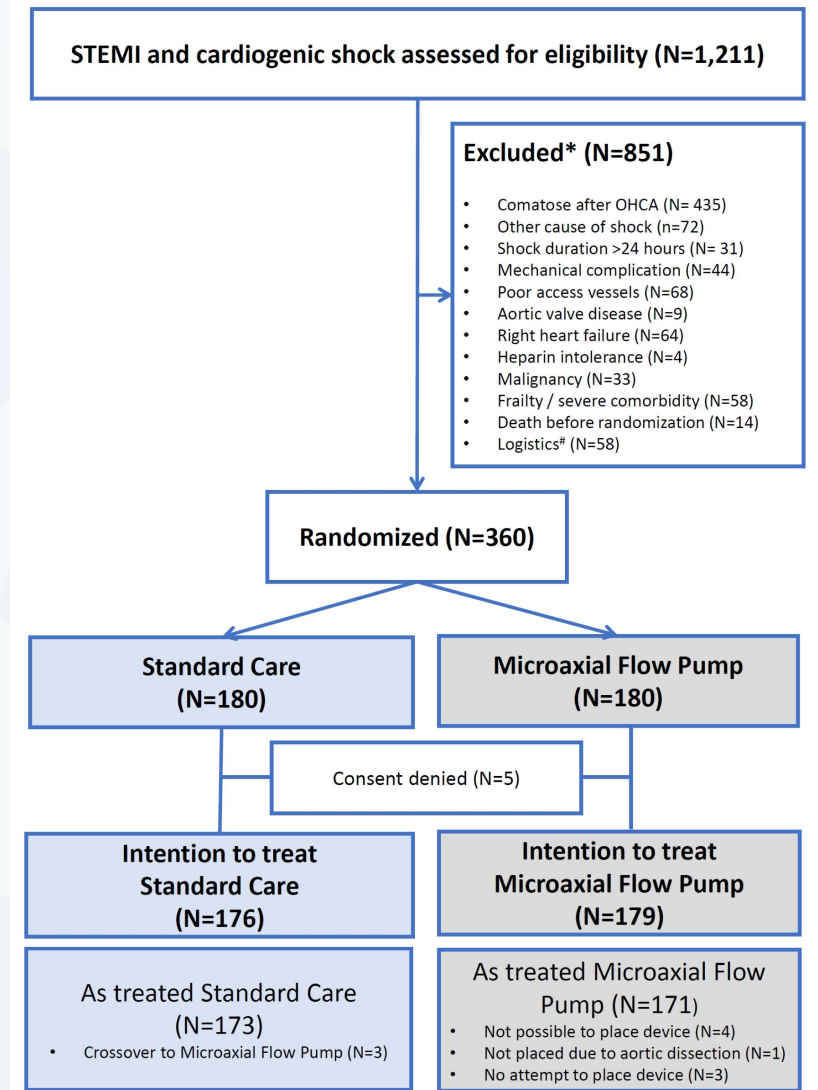


Endpoint primario: muerte por cualquier causa a los 180 d.

Endpoints secundarios: sangrados, isquemia en MMII, hemólisis, fallo dispositivo, IAo

Criterios de exclusión:

- Pacientes recuperados de PCR extrahospitalaria y que permanecieran con bajo nivel de conciencia a su llegada a la sala de hemodinámica.
- Fallo VD.



Characteristic	Microaxial Flow Pump plus Standard Care (N = 179)	Standard Care Alone (N = 176)
Median age (IQR) — yr	67 (58–76)	69 (61–76)
Male sex — no. (%)	142 (79.3)	139 (79.0)
Medical history — no. (%)		
Hypertension	89 (49.7)	94 (53.4)
Diabetes	33 (18.4)	47 (26.7)
Myocardial infarction	29 (16.2)	28 (15.9)
Heart failure	16 (8.9)	17 (9.7)
Chronic kidney disease	17 (9.5)	18 (10.2)
Median systolic blood pressure (IQR) — mm Hg	84 (72–91)	82 (72–91)
Median of the mean arterial blood pressure (IQR) — mm Hg	63 (55–72)	64 (55–73)
Median heart rate (IQR) — beats/min	94 (77–110)	95 (76–111)
Median arterial lactate level (IQR) — mmol/liter	4.6 (3.4–7.1)	4.5 (3.2–6.9)
Median left ventricular ejection fraction (IQR) — %	25 (20–31)	25 (15–30)
Resuscitation before randomization — no. (%)	39 (21.8)	33 (18.8)
Intubation before randomization — no. (%)	35 (19.6)	28 (15.9)
Transfer from outside hospital — no. (%)	51 (28.5)	48 (27.3)
Anterior myocardial infarction — no. (%)	126 (70.4)	129 (73.3)
SCAI–CSWG stage at admission — no. (%) [†]		
C	100 (55.9)	97 (55.1)
D	51 (28.5)	50 (28.4)
E	28 (15.6)	29 (16.5)
No. of diseased vessels on coronary angiography — no. (%)		
0	1 (0.6)	0
1	52 (29.1)	47 (26.7)
2	70 (39.1)	64 (36.4)
3	56 (31.3)	65 (36.9)
Timing of randomization		
Median time from symptom onset to randomization (IQR) — hr	4.8 (2.4–12.8)	3.8 (2.2–9.4)
Randomization performed before revascularization — no. (%)	99 (55.3)	102 (58.0)
Randomization performed in the catheterization laboratory but after revascularization — no. (%)	48 (26.8)	48 (27.3)
Randomization performed ≤12 hr after departure from the catheterization laboratory — no. (%)	32 (17.9)	26 (14.8)

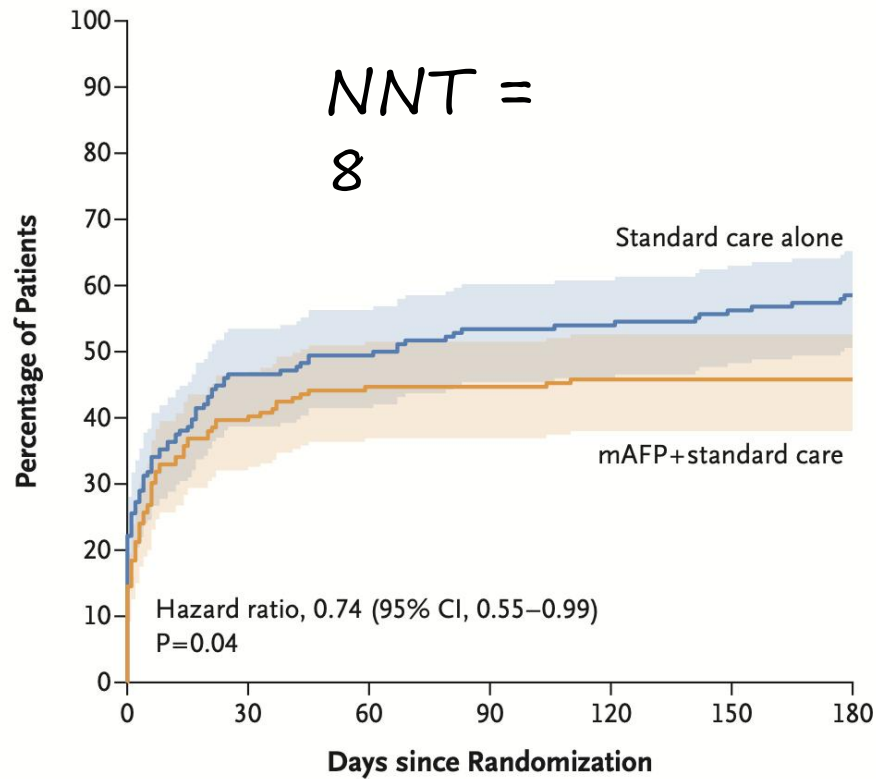
Escalation to additional mechanical circulatory support

Placement of Impella 5.0 device — no. (%)	7 (3.9)	5 (2.8)
Placement of Impella CP for venting during venoarterial ECMO therapy — no. (%)	0	4 (2.3)
Placement of Impella 2.5 device — no. (%)	0	1 (0.6)
Placement of Impella RP device — no. (%)	0	0
Venoarterial ECMO — no. (%)	21 (11.7)	33 (18.8)
Median time from randomization to placement of venoarterial ECMO (IQR) — hr	14 (4–54)	2 (1–5)
Placement of permanent LVAD — no. (%)	10 (5.6)	4 (2.3)
Any escalation to additional mechanical circulatory support — no. (%)	28 (15.6)§	37 (21.0)¶

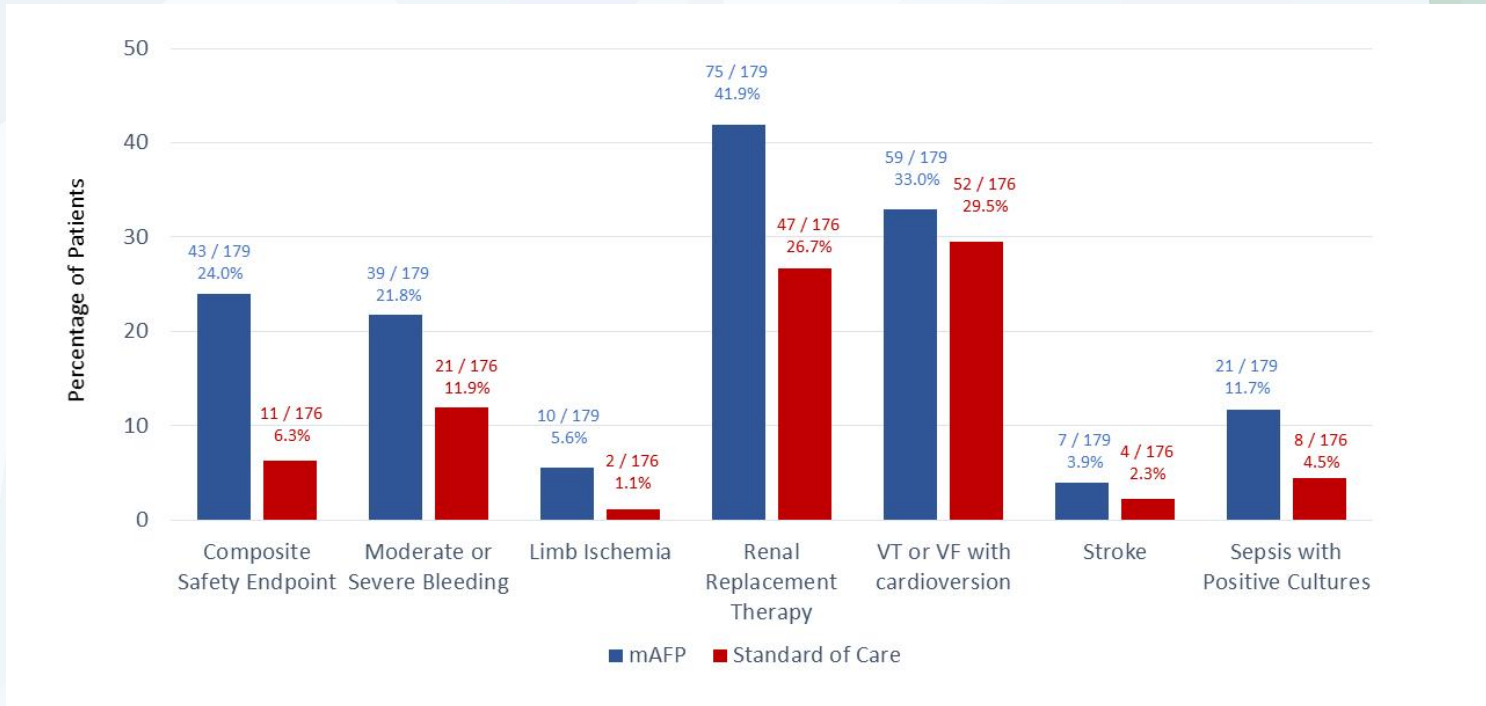
ORIGINAL ARTICLE

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

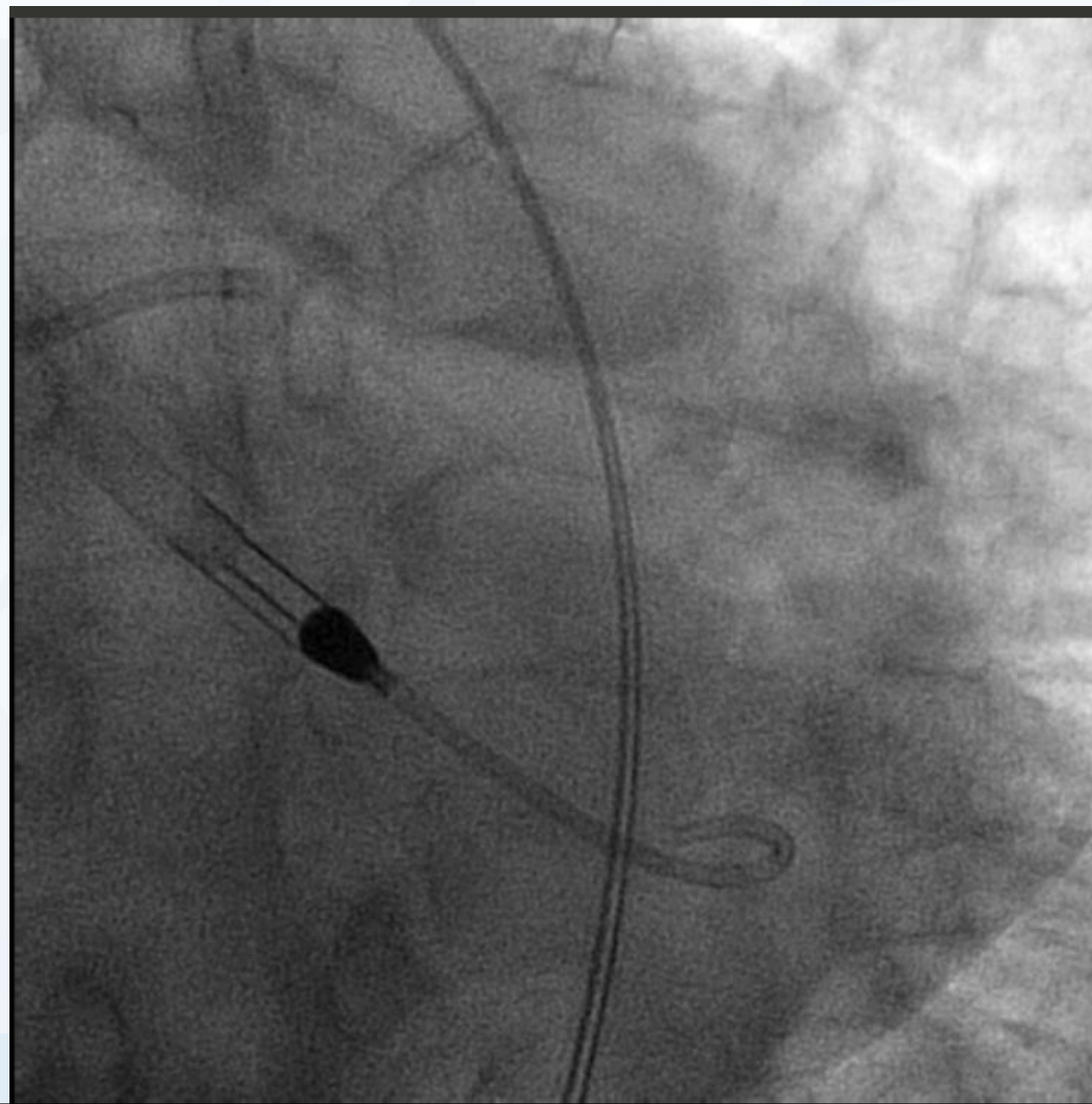
A Death from Any Cause



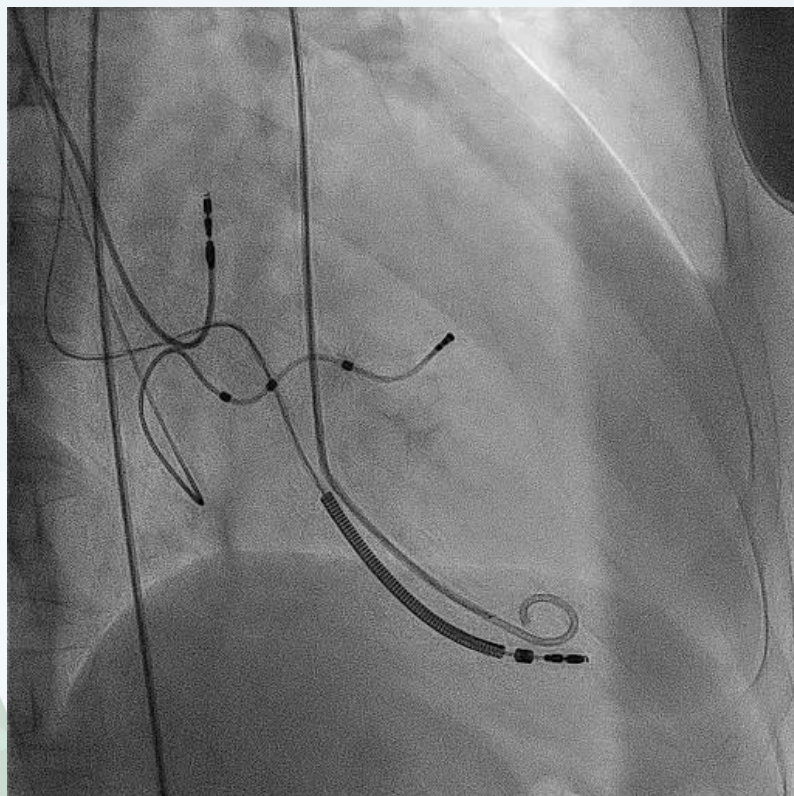
No. at Risk	0	30	60	90	120	150	180
Standard care	176	94	89	82	81	77	72
mAFP+standard care	179	108	99	99	97	97	97



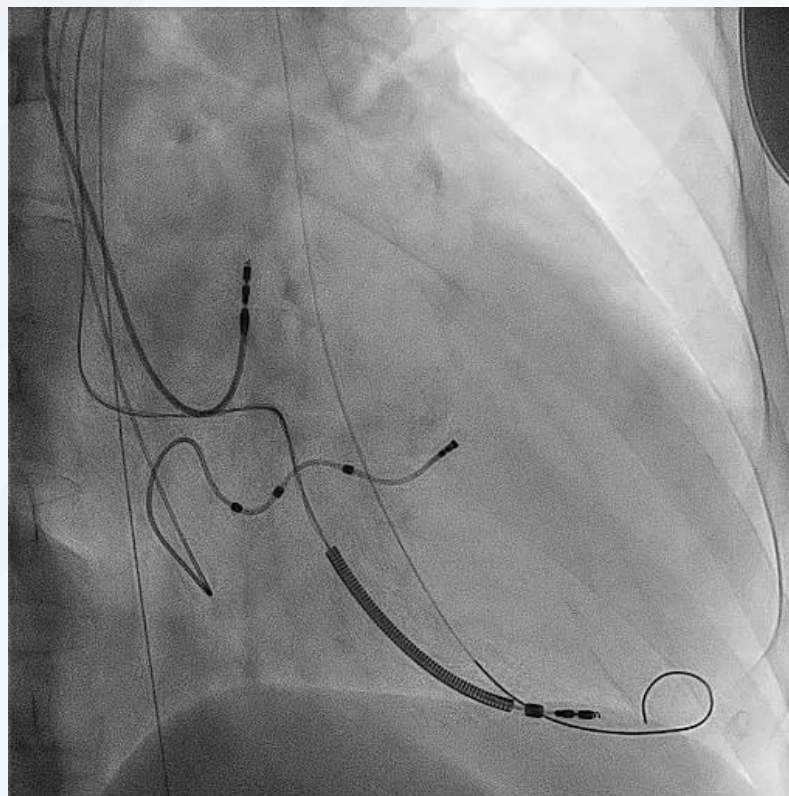
Implante de IMPELLA CP



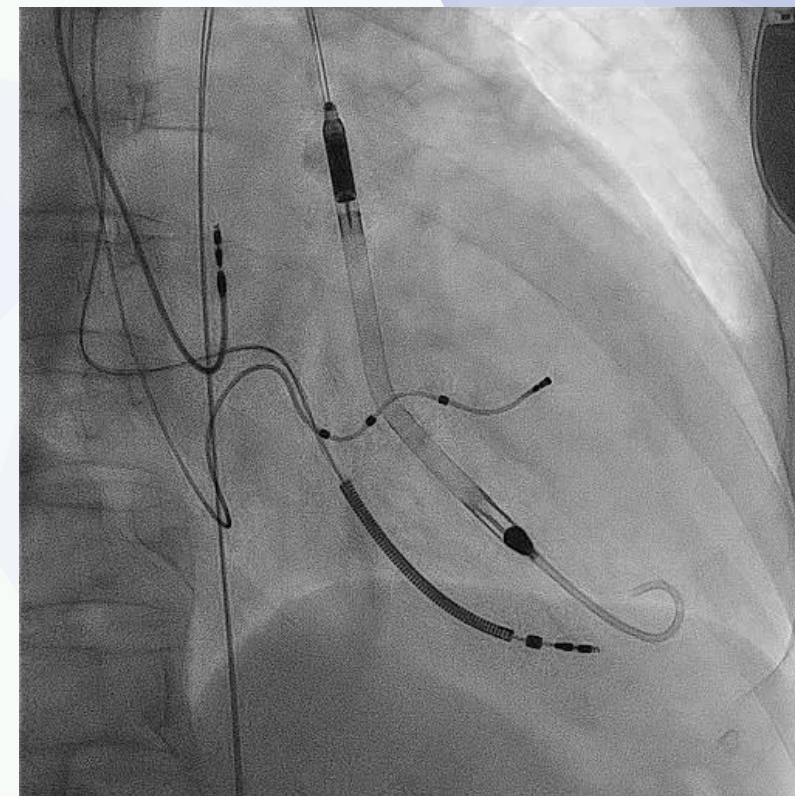
Implante Impella



Pigtail hasta la punta del VI

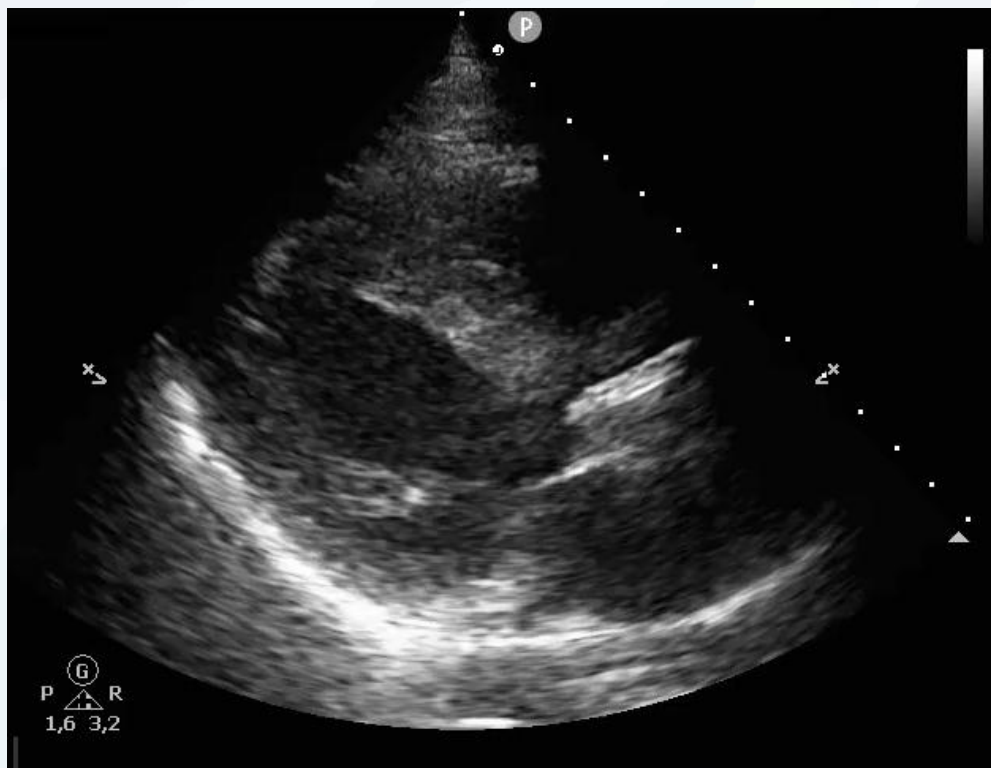


Guía de alto soporte (0.018")



Se avanza Impella y se retira guía

Comprobar posición en ETT y curvas



Evolución

9/1



ICP DA
IMPELLA CP



- NRL: consciente y orientado con Remifentanilo 1.2 mcg/Kg/min
- RESP: eupneico, SatO₂ 98% con GN 2L
- HD: TAM 70mmHg con **NA 0.5 mcg/Kg/min + DBT 2.3 mcg/Kg/min**
Impella P8, 3.5L
***Láctico 4.5 → 3.8 mmol/l**
- NEF: FRA (cr 1.09 → 1.6) y oligoanuria a pesar de Furosemida iv
- Elevación de transaminasas: GPT 214, GOT 953. Troponina (Tnlhs) 558000 ng/L (<72)

Evolución

9/1



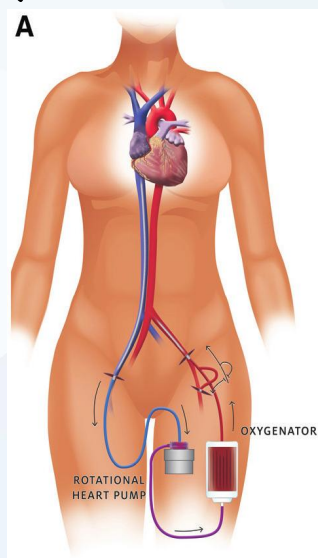
ICP DA
IMPELLA CP



10/



ECMO
VA
periféric



- NRL: consciente y orientado con Dexdor
- RESP: eupneico, SatO2 98% con GN 4L y ECMO FiO2 0.8
- HD: TAM 75mmHg con **Nicardipino 3ml/h + DBT 1.5 mcg/Kg/min + LVS 15ml/h**

Impella P5, 3L

ECMO 3600rpm, 3.2L

***Láctico 0.6**

- NEF: diuresis ok con furosemida iv. Cr 1.2

Evolución

9/1



ICP DA
IMPELLA CP

10/

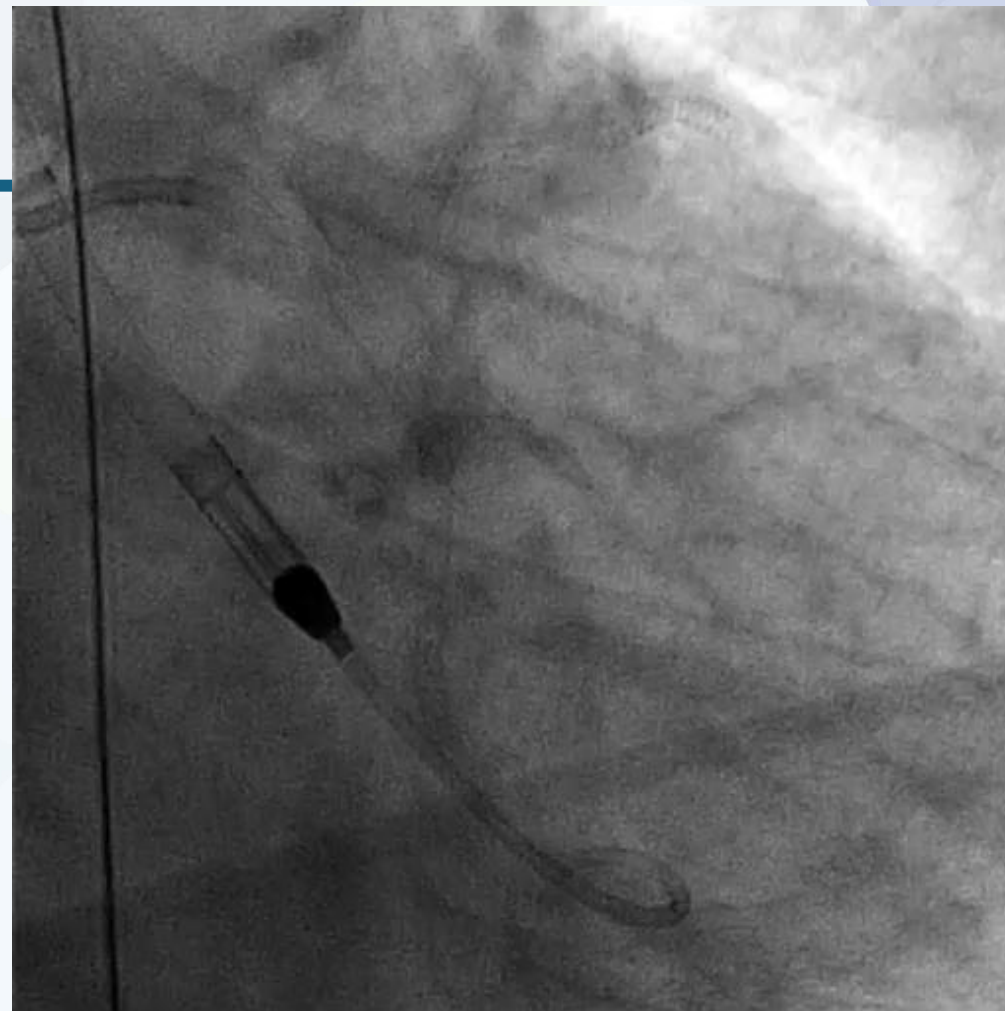
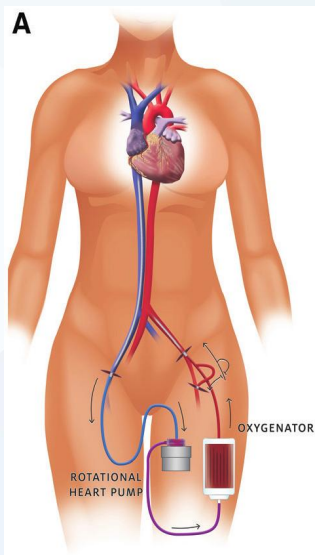


ECMO
VA
periférico

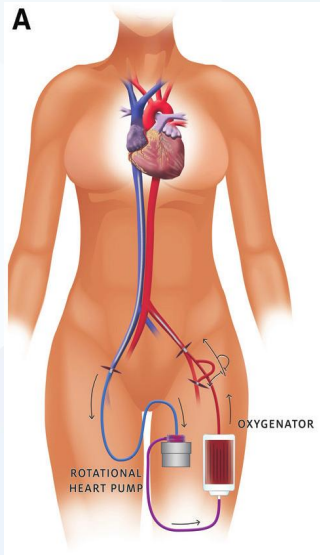
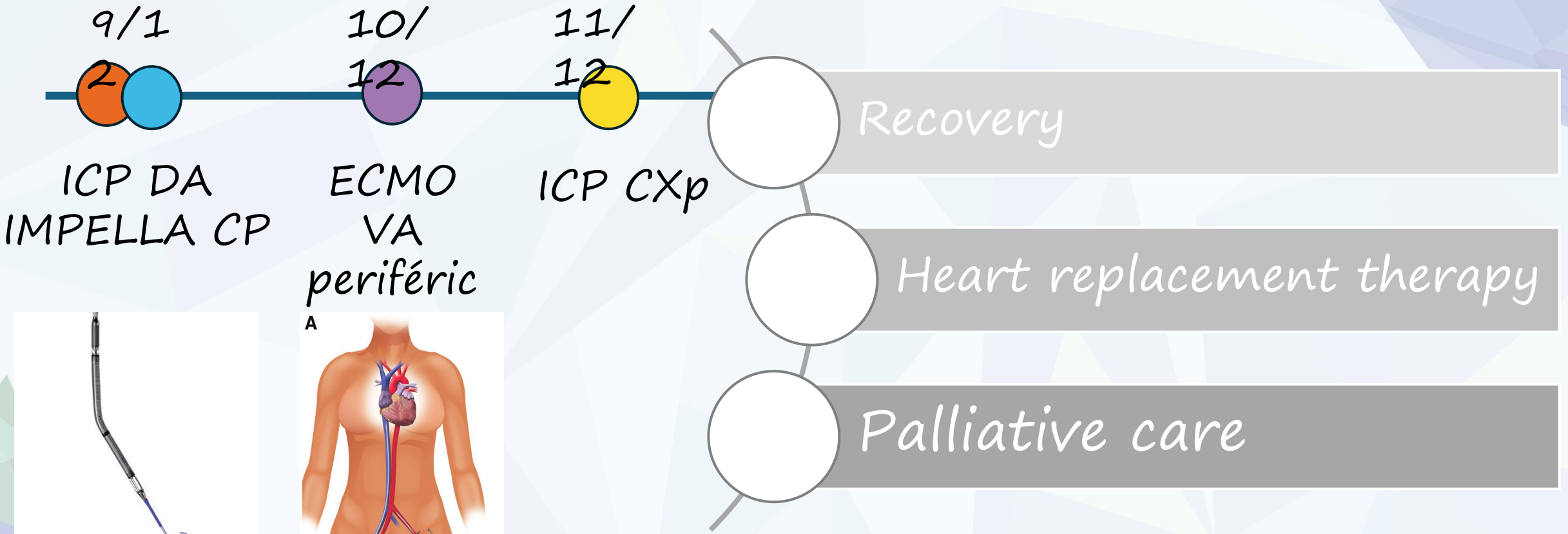
11/



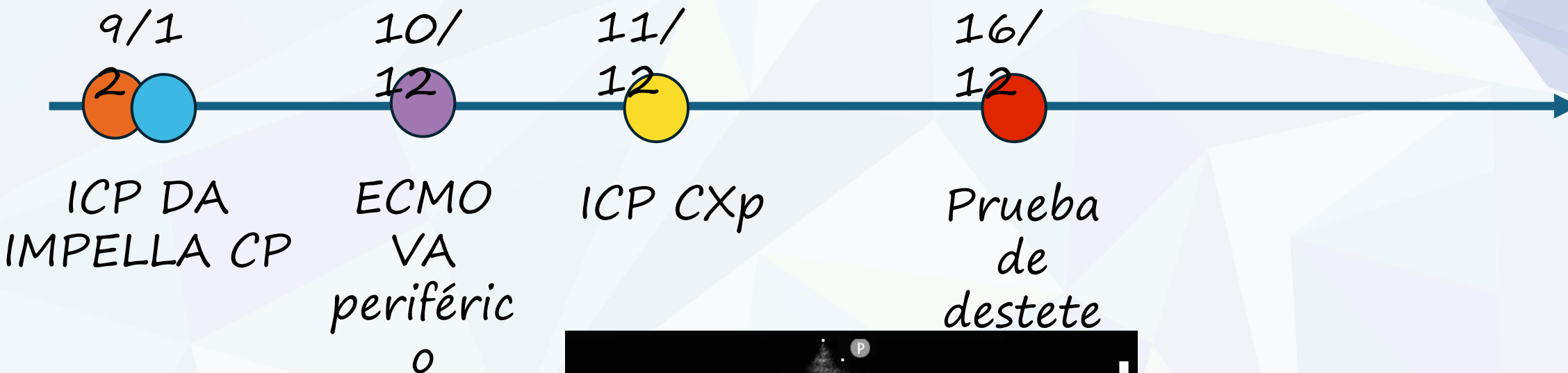
ICP CXp



Evolución



Evolución



Evolución

9/1



ICP DA
IMPELLA CP

10/



ECMO
VA
periféric
o

11/



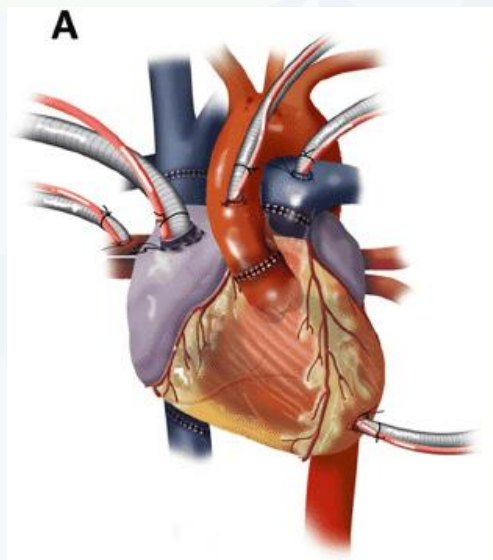
ICP CXp

17/

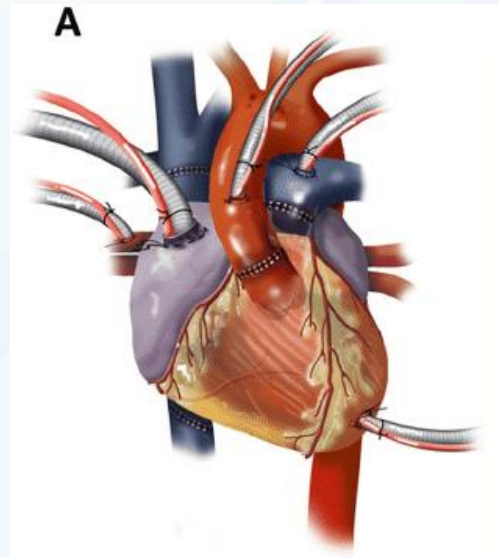
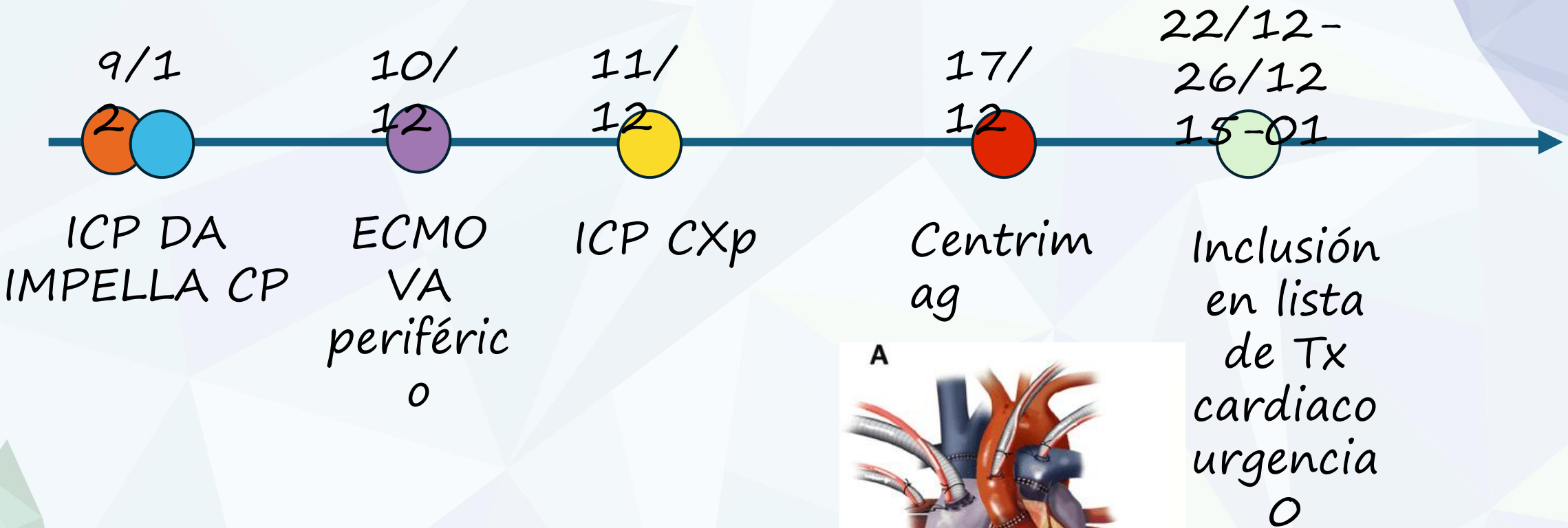


Centrim
ag

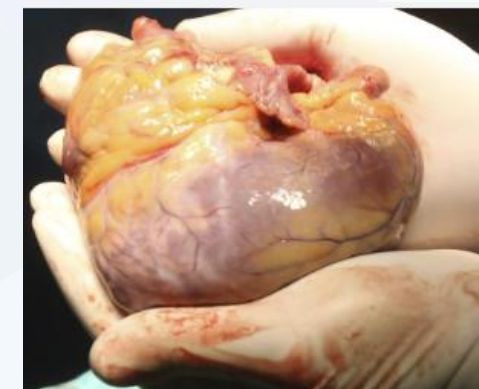
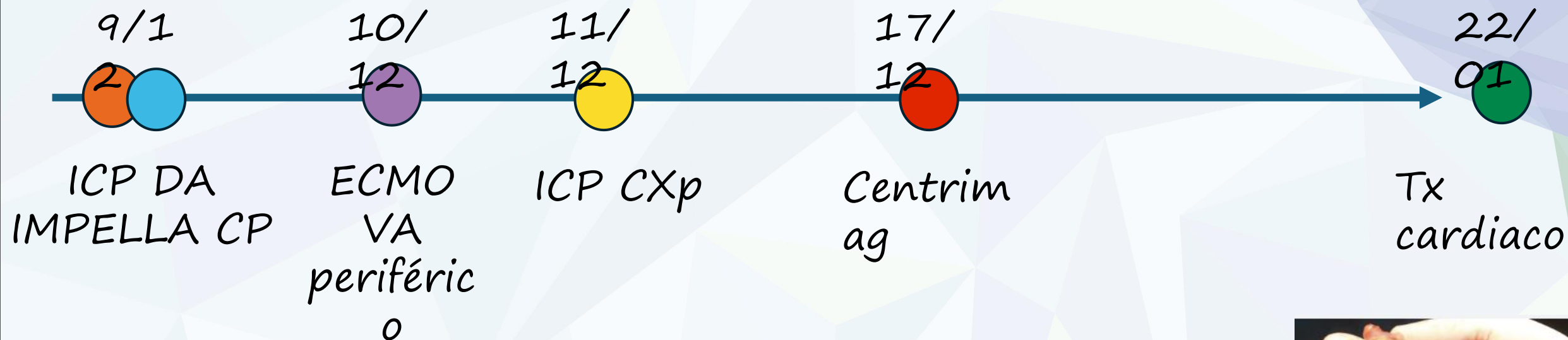
A



Evolución

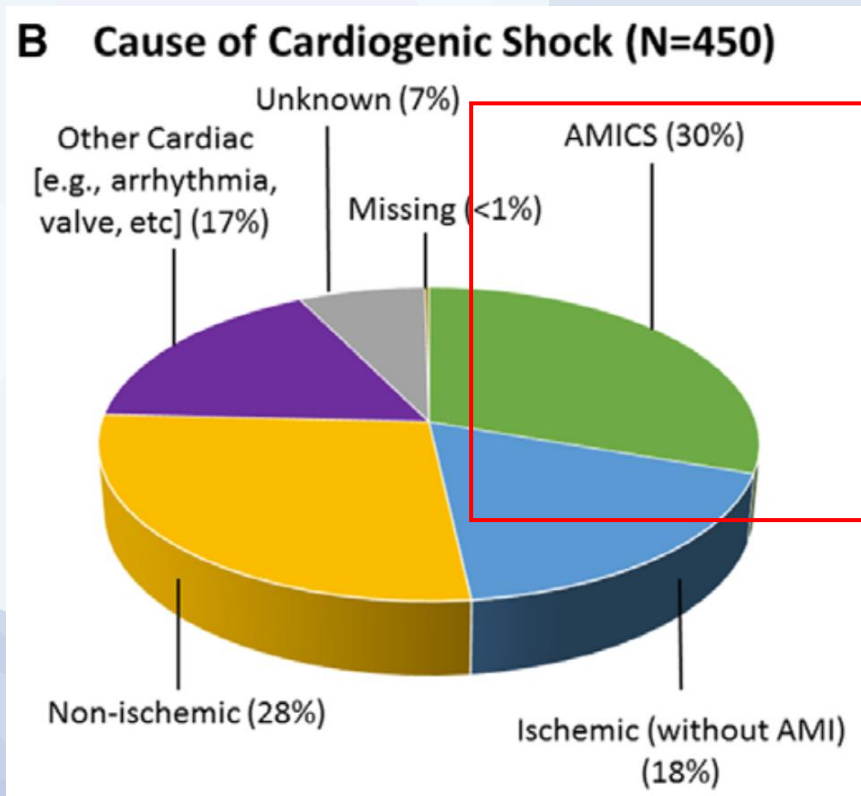


Evolución

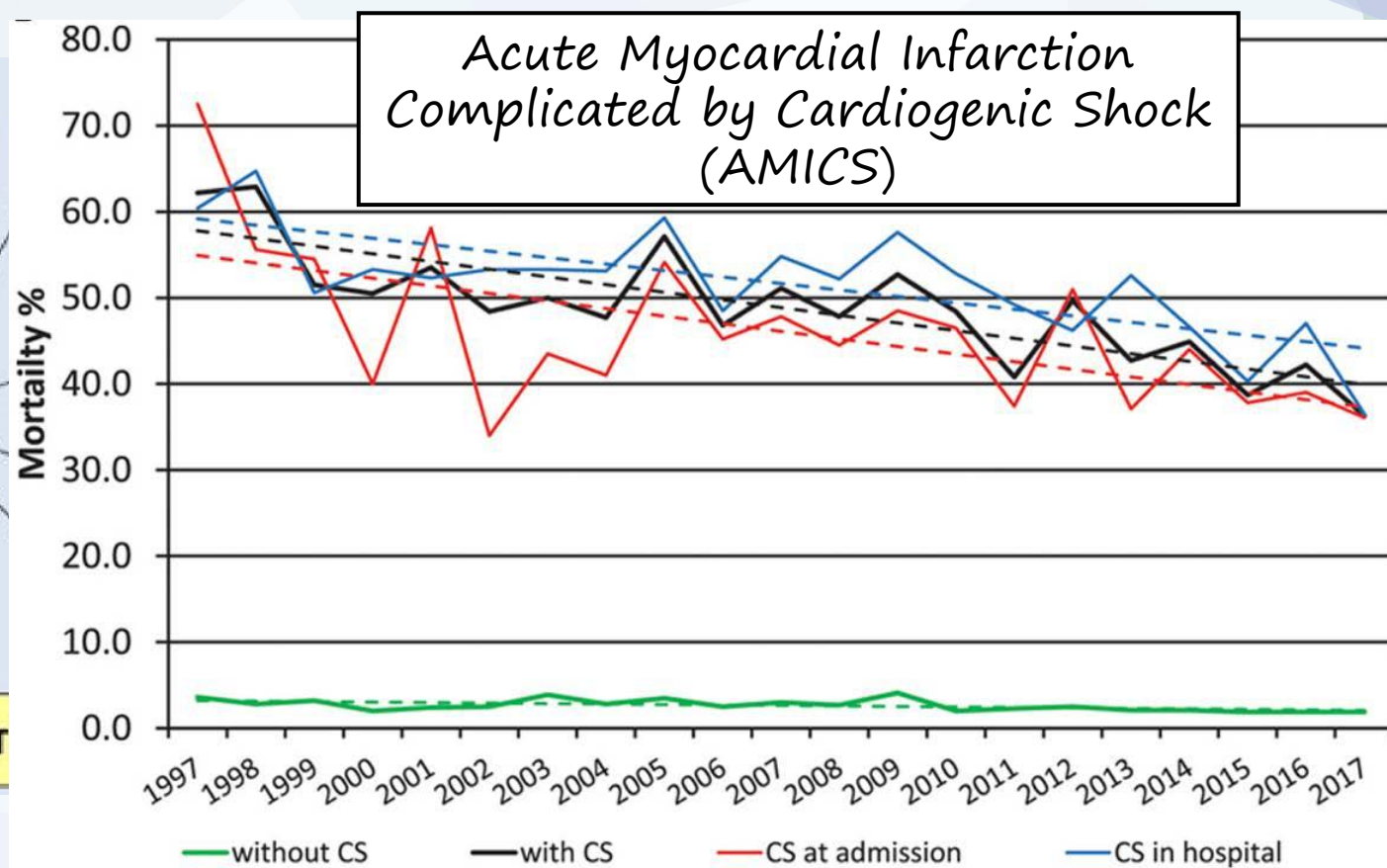


Shock cardiogénico

Condición aguda y potencialmente letal que consiste en la incapacidad para proporcionar una adecuada perfusión tisular que consiga satisfacer las demandas metabólicas y provoca un fallo multiorgánico

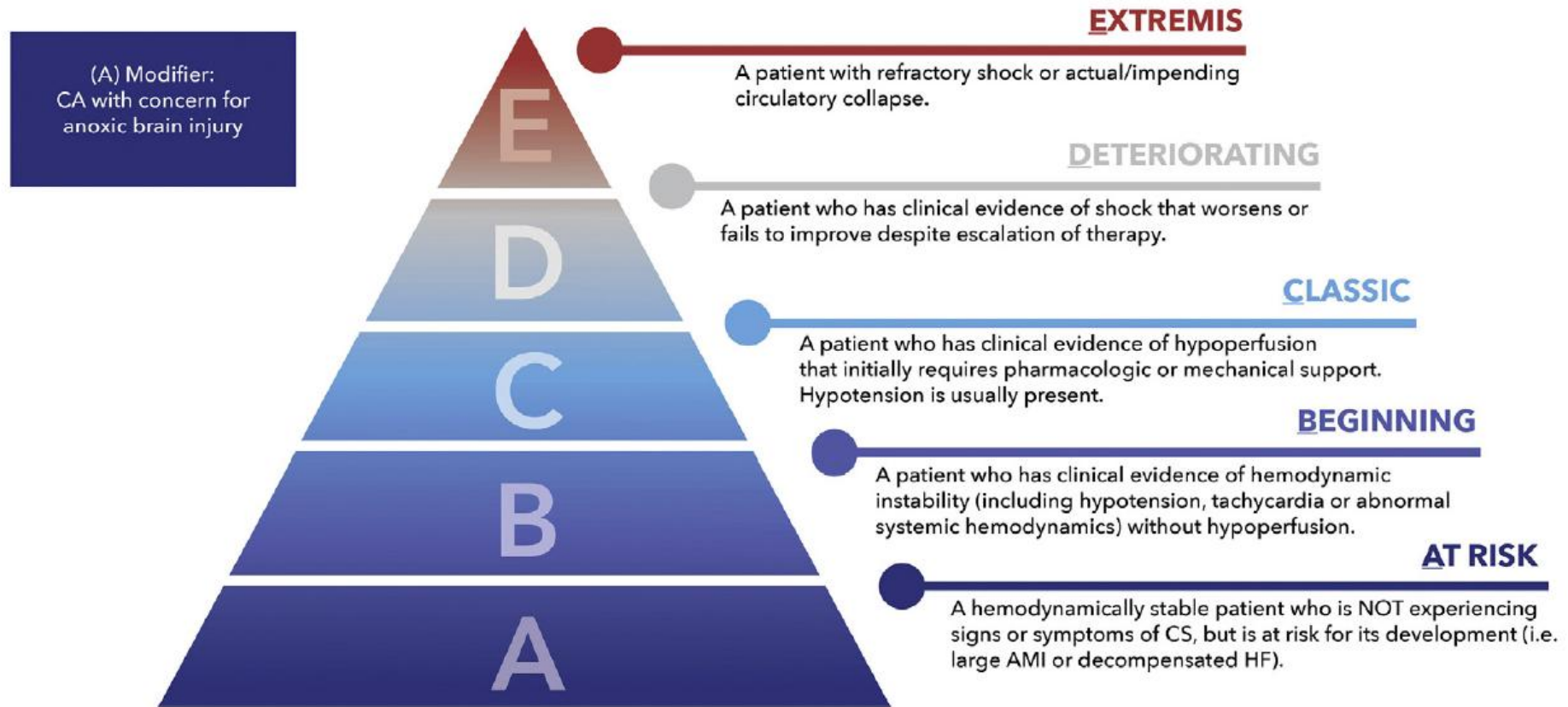


Berg DD et al. *Circ Cardiovasc Qual Outcomes*. 2019



Henry T et al. *Circulation*. 2021

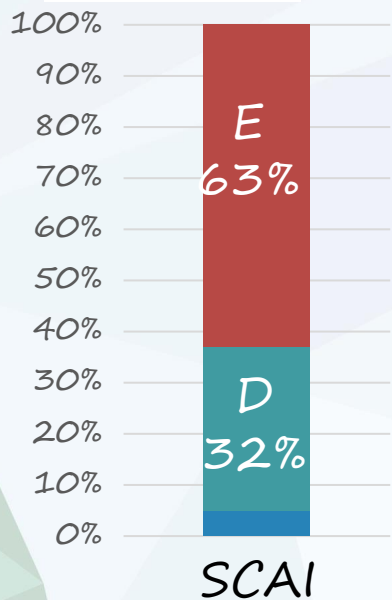
FIGURE 4 Updated SCAI SHOCK Classification Pyramid



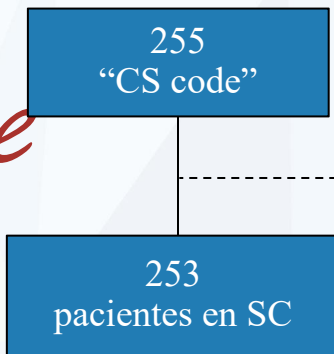
AMI = acute myocardial infarction; CS = cardiogenic shock; HF = heart failure; SCAI = Society for Cardiovascular Angiography and Interventions.

Puerta de Hierro Hub & Spoke

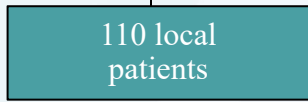
8 years
Sept 2014 - Aug 2022



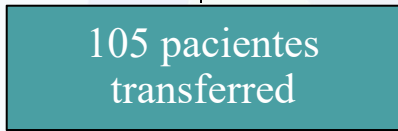
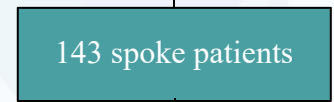
95%
SCAI
D-E



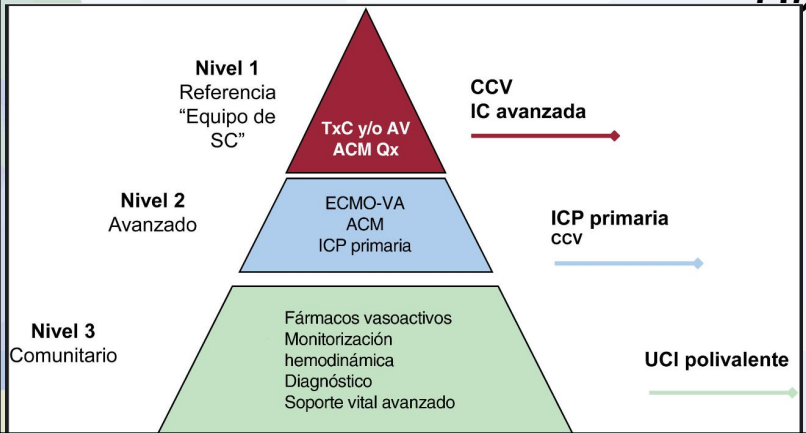
2 patients - 2 episodes of CS
AMI y PGF
Electrical storm & ADHF



Level 1
Hub



2 patients Not CS criteria
12 patients Futility
23 patients Collaborative
1 paciente Logistic



71 patients
68%



Level 2
CVS

F.J. Hernández-Pérez, mayo 2023



Level 3
ICU

34 patients
32%



Transfer: stabilization

- SCAI
- ECAI
- SCAI C



Level 1
Hub



Patients transferred from level 2 centers arrive more stable



Level 2
CVS

Lactate $3,8 \pm 5,4$

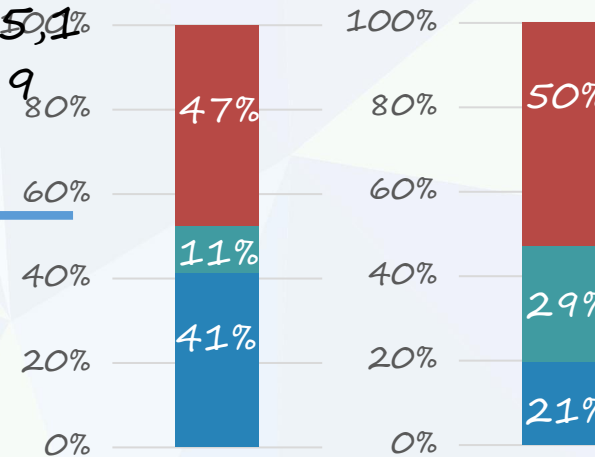
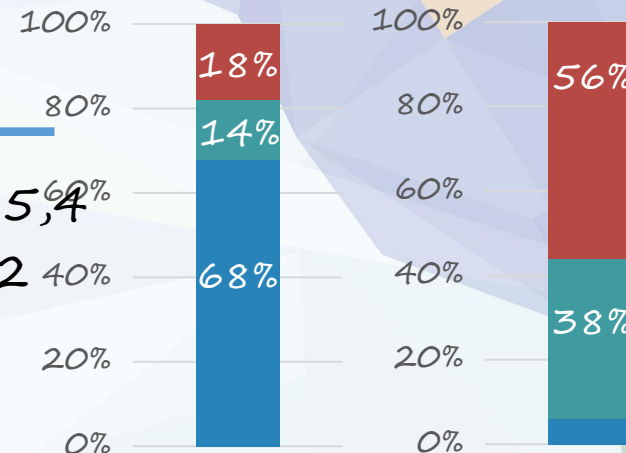
VIS 27 ± 32

$p < 0,01$

Lactate $5,1 \pm 5,1$

VIS 35 ± 39

Level 3 ICU





Transfer: stabilization

- SCAI
- ECAI
- SCAI C



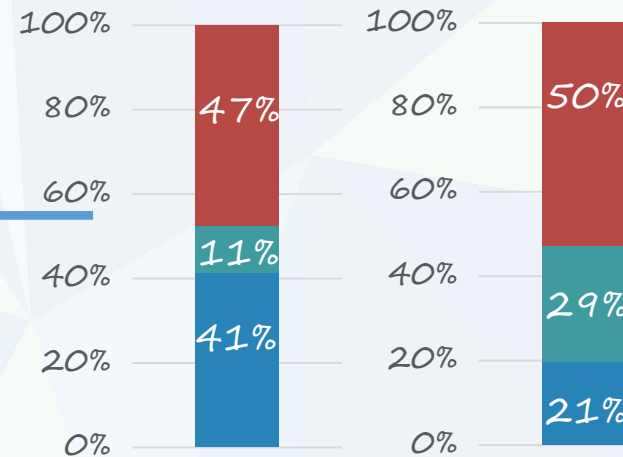
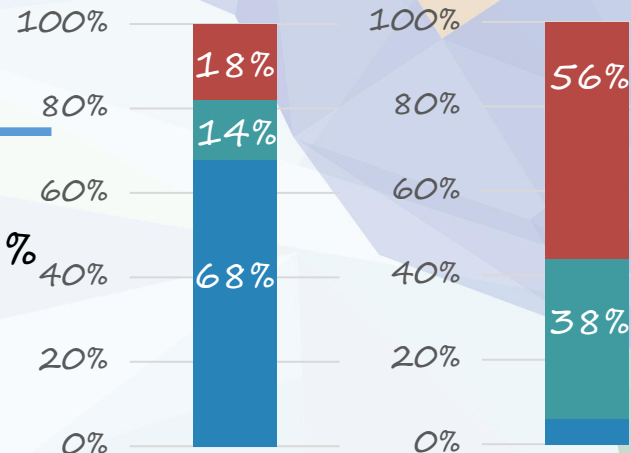
Level 1
Hub

Level 2
CVS

IABP 25%
ECMO 37%
T-VAD 16%
MCS 78%

IABP 35%

Level 3 ICU

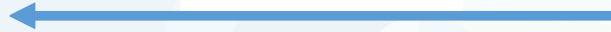


★ Stability is associated with increased use of MCS

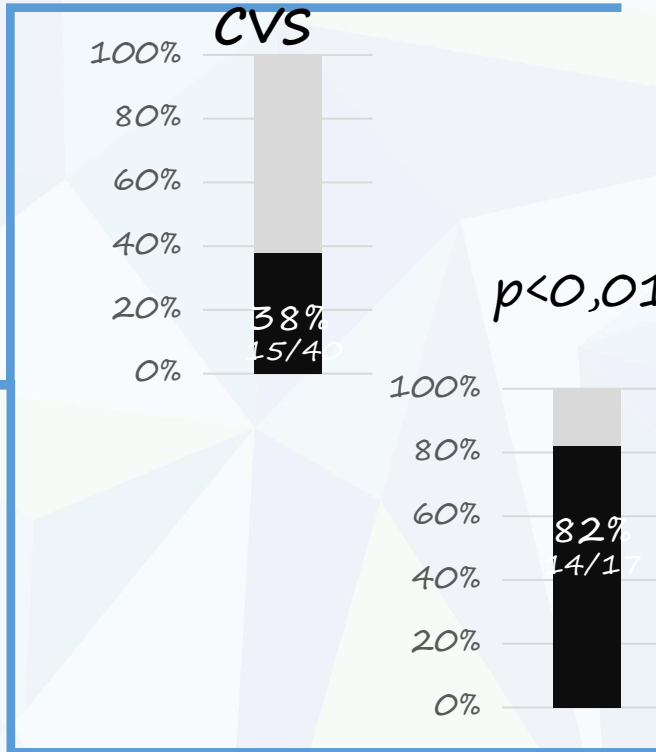
- SCAI
- E SCAI
- B SCAI C



Level 1
Hub



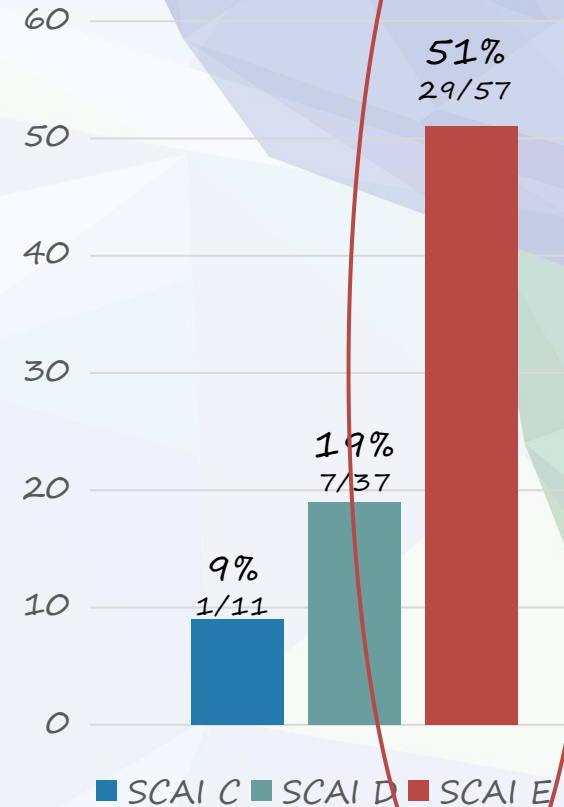
Level 2
CVS



Level 3 ICU



Mortality dx SCAI

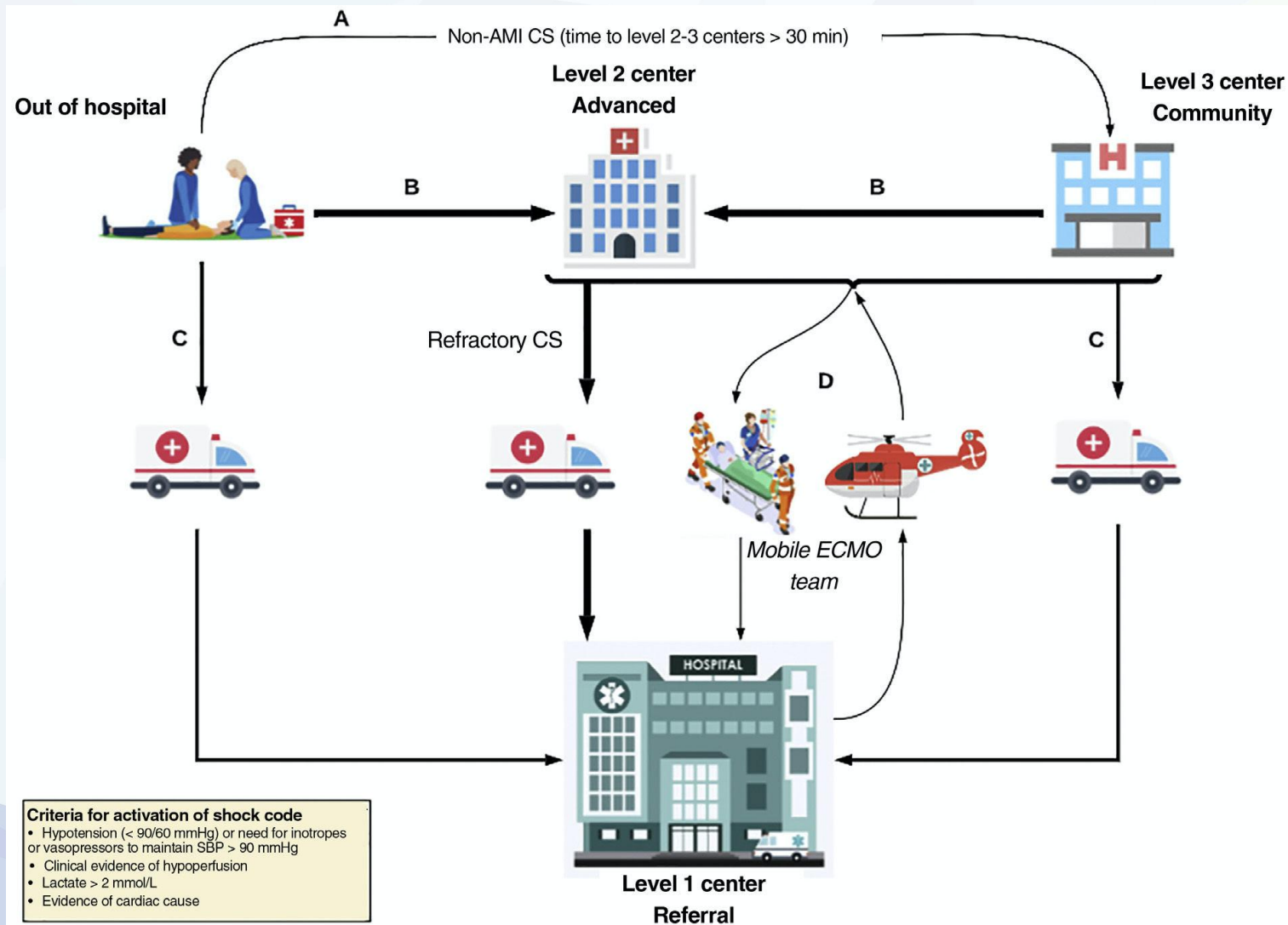


■ SCAI C ■ SCAI D ■ SCAI E

★ Transferring **SCAI E** patients from level 3 patients is associated with a prohibitive mortality



Propuesta código shock cardiogénico

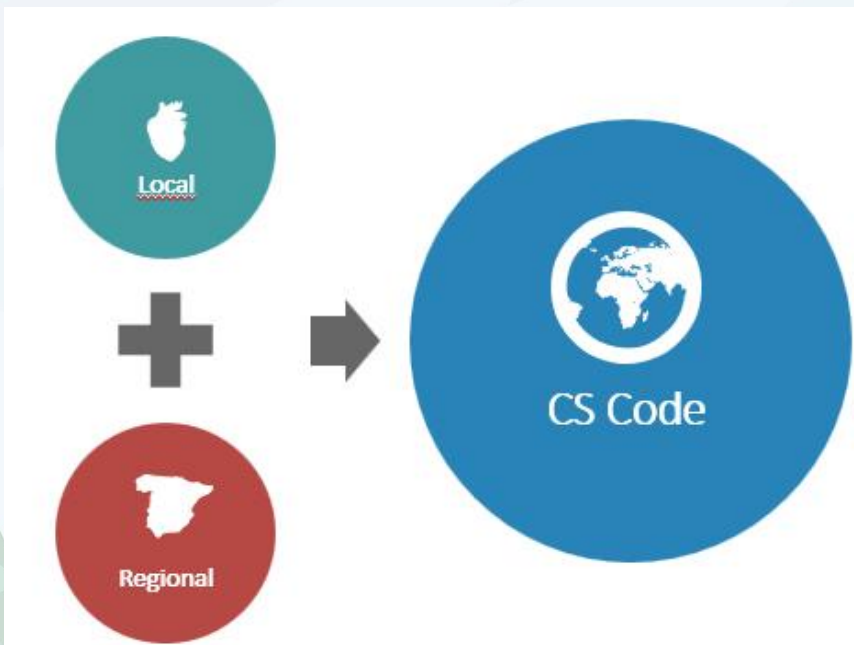


Mail goals

1. Early detection
2. Expedited transfers to level 1-2 centers
3. Exit strategies



Shock team



1. Early detection & characterization
2. Revascularization and MCS
3. Exit strategies



<p>Advanced HF</p> <p>Coordination HT & LVADs ONT</p>	<p>CICU</p> <p>Diagnosis & triage Medical treatment Invasive hemodynamics MCS Management</p>
<p>Interventional Cardiology</p> <p>Primary angioplasty 24/7 pMCS</p>	<p>Cardiac Surgery</p> <p>ECMO Temporary-VAD LVAD HT</p>



Adapt Doll et al. CCI 2016:88;424-433

Puerta de Hierro

Take home messages

- En los casos de shock cardiogénico en contexto de un IAM, la ICP primaria puede no ser suficiente, debemos disponer de dispositivos de soporte mecánico circulatorio en las salas de hemodinámica adheridas al programa de código IAM.
- Impella ha demostrado en estos pacientes reducir la mortalidad vs el tratamiento médico convencional.
- Sin embargo, la mortalidad de estos pacientes sigue siendo elevada.
- La coordinación mediante el CÓDIGO SHOCK de los distintos centros sanitarios puede mejorar la supervivencia de los pacientes en shock cardiogénico.