

# Patient Selection for VA ECMO

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University of Western Ontario, London, Ontario, Canada

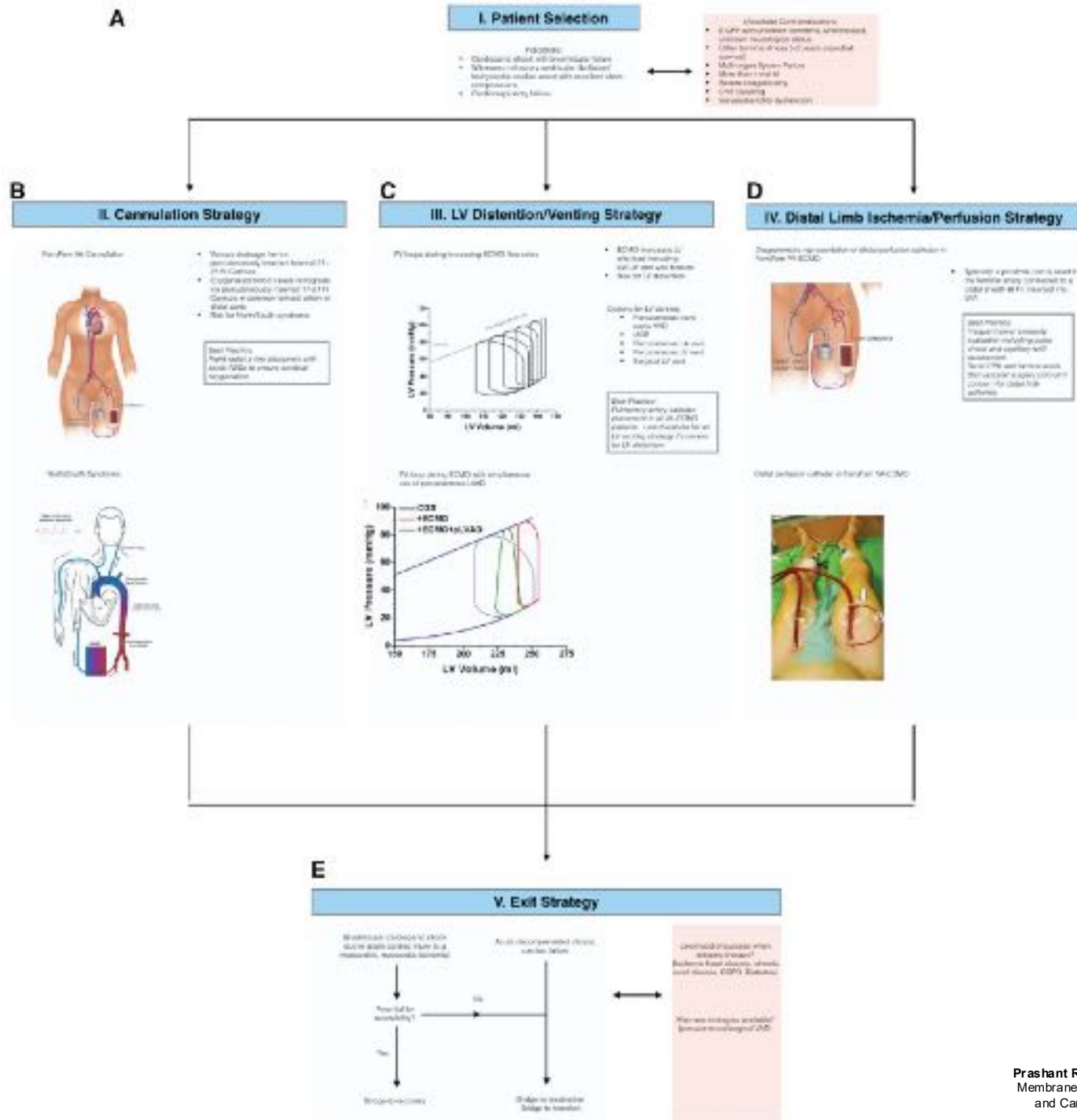
# Outline

Patient Selection

VA ECMO Initiation

VA ECMO Goals





- Absorbable drug molecules
- If CYP activity is inhibited, absorption is not affected
- Other forms of drug CYP activity are not affected
- Multidrug System P-glycoprotein
- More than 10000
- Resists excretion
- Drug transport
- Involves ATP dependent

- Continuous about all intermediate values
- Whence, all values and limits of functions
- Indispensable studies about all number class
- Continuous very famous

### Flowchart 10: Case resolution



**Best Tips:**  
Night vision is less accurate with acids/alkalis so ensure careful navigation.

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## 8

#### IV. Distal Limb Ischemia/Perfusion Strategy

Diagramm zur Darstellung der Ergebnisse der Analyse der Daten der Studie.



1990, March 10  
 "Super-Corn" is widely  
 available, including some  
 whole and partially-hull  
 products.  
 Super-Corn and Super-Corn  
 that is not a Super-Corn  
 product is a Super-Corn  
 product.

David Nelson, Cathode Interface SAC/NC



# Week 8

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# Outline

Patient Selection

VA ECMO Initiation

VA ECMO Goals



# Case Presentation

ID

- 26-year-old female

PMHx

- Hypothyroidism on replacement therapy

# Case Presentation

ID

- 26-year-old female

PMHx

- Hypothyroidism on replacement therapy

HPI

- Viral prodrome 5 days ago
- Feeling very weak 2 days ago
- Presented to peripheral hospital

# Case Presentation

O/E

- Cold clammy extremities
- BP: 78/49, HR: 136 (sinus), SpO<sub>2</sub> 89% on RA

Labs

- Creatinine 138 mmol/L, BUN 14
- Lactate: 6.2 mmol/L, pH 7.21, HCO<sub>3</sub> 15



# Case Presentation

## O/E

- Cold clammy extremities
- BP: 78/49, HR: 136 (sinus), SpO<sub>2</sub> 89% on RA

## Labs

- Creatinine 138 mmol/L, BUN 14
- Lactate: 6.2 mmol/L, pH 7.21, HCO<sub>3</sub> 15

## Next Steps

- Started on norepinephrine infusion
- POCUS

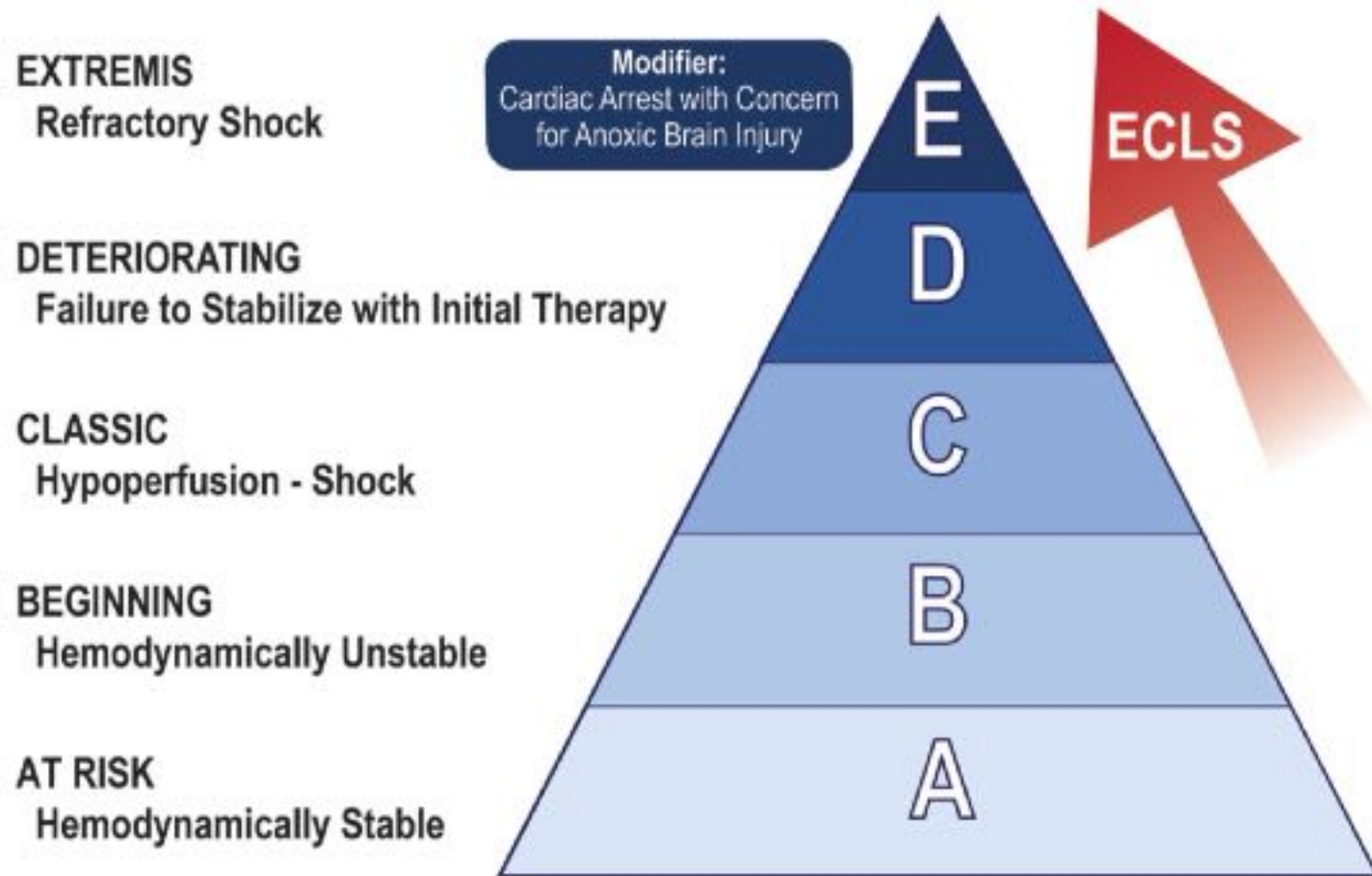


# Question 1

Which **SCAI** class of cardiogenic shock is she in?

- A. Class A
- B. Class B
- C. Class C
- D. Class D
- E. Class E

# Society of Cardiovascular Angiography & Interventions (SCAI) Staging for Cardiogenic Shock



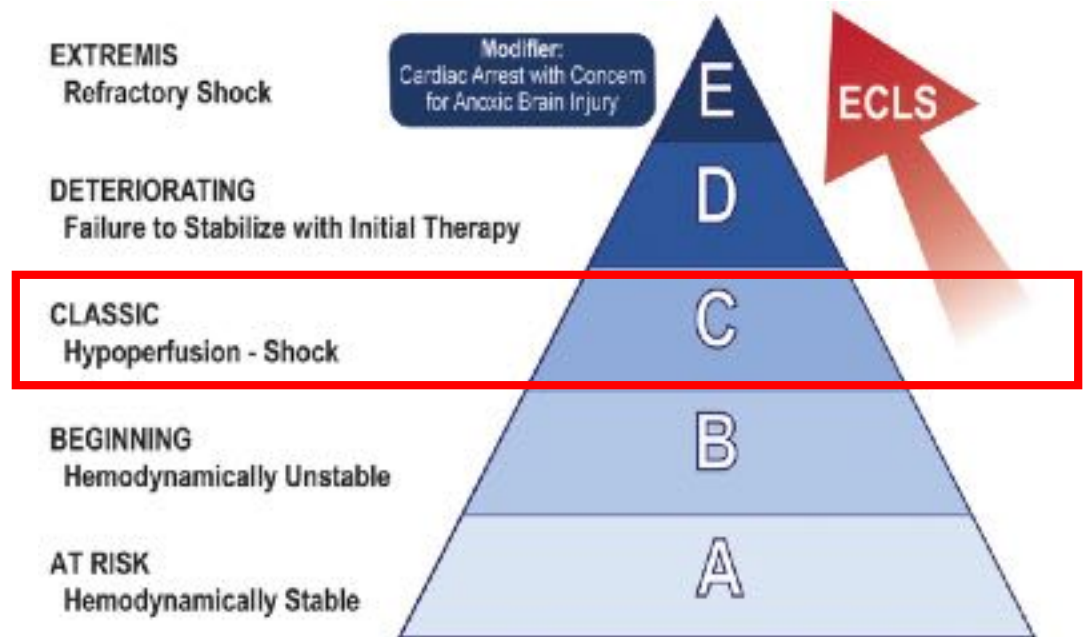
# Society of Cardiovascular Angiography & Interventions (SCAI) Staging for Cardiogenic Shock

		VA ECMO
E	A patient with refractory shock or actual/impeding circulatory collapse	Indicated ± eCPR
D	A patient who has clinical evidence of shock that worsens or fails to respond to medical therapy	Indicated
C	A patient who has clinical evidence of hypoperfusion that initially responds to medical therapy but requires <b>mechanical support</b> . Hypotension is usually present	Strongly consider based on clinical response.
B	A patient who has clinical evidence of hemodynamic instability (hypotension and/or abnormal systemic hemodynamics) without hypoperfusion	Early planning
A	A hemodynamically stable patient who is not experiencing signs or symptoms of shock or at risk for its development	Not indicated

# Question 1

Which **SCAI** class of cardiogenic shock is she in?

- A. Class A
- B. Class B
- C. **Class C**
- D. Class D
- E. Class E



# Cardiogenic Shock Diagnosis

<b>Symptoms/Signs</b>	Altered mental status, confusion, chest pain or pressure, cold and clammy extremities, rapid pulse, low pulse pressure (<25% of SBP), elevated jugular venous pressure, crackles, rales, orthopnea, paroxysmal nocturnal dyspnea, lower extremity edema
<b>Urine output</b>	Oliguria or anuria, <30 mL/h (<0.5 mL/[kg·h])
<b>Sustained hypotension</b>	SBP <90 mm Hg, MAP <65 mm Hg for >30 min or a >30-mm Hg decrease from baseline, or the need for pharmacological or mechanical support to maintain SBP >90 mm Hg
<b>Perfusion</b>	Evaluate markers of end-organ malperfusion, including lactic acid >2 mmol/L, ALT >200 U/L or >3× upper limit of normal, creatinine ≥2× upper limit of normal, pH <7.2, metabolic acidosis without another known cause



S  
U  
S  
P  
E  
C  
T

# Cardiogenic Shock Diagnosis

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<b>ECG/Echocardiogram</b>	Evaluate acute ischemia, including ECG and sonographic evidence of STEMI (regional wall motion abnormalities); evidence of LV or RV dilation and systolic dysfunction; valvular pathology
<b>Congestion</b>	Presence or absence of congestion based on physical signs and hemodynamics; elucidation of ventricular involvement (LV vs RV vs BiV)



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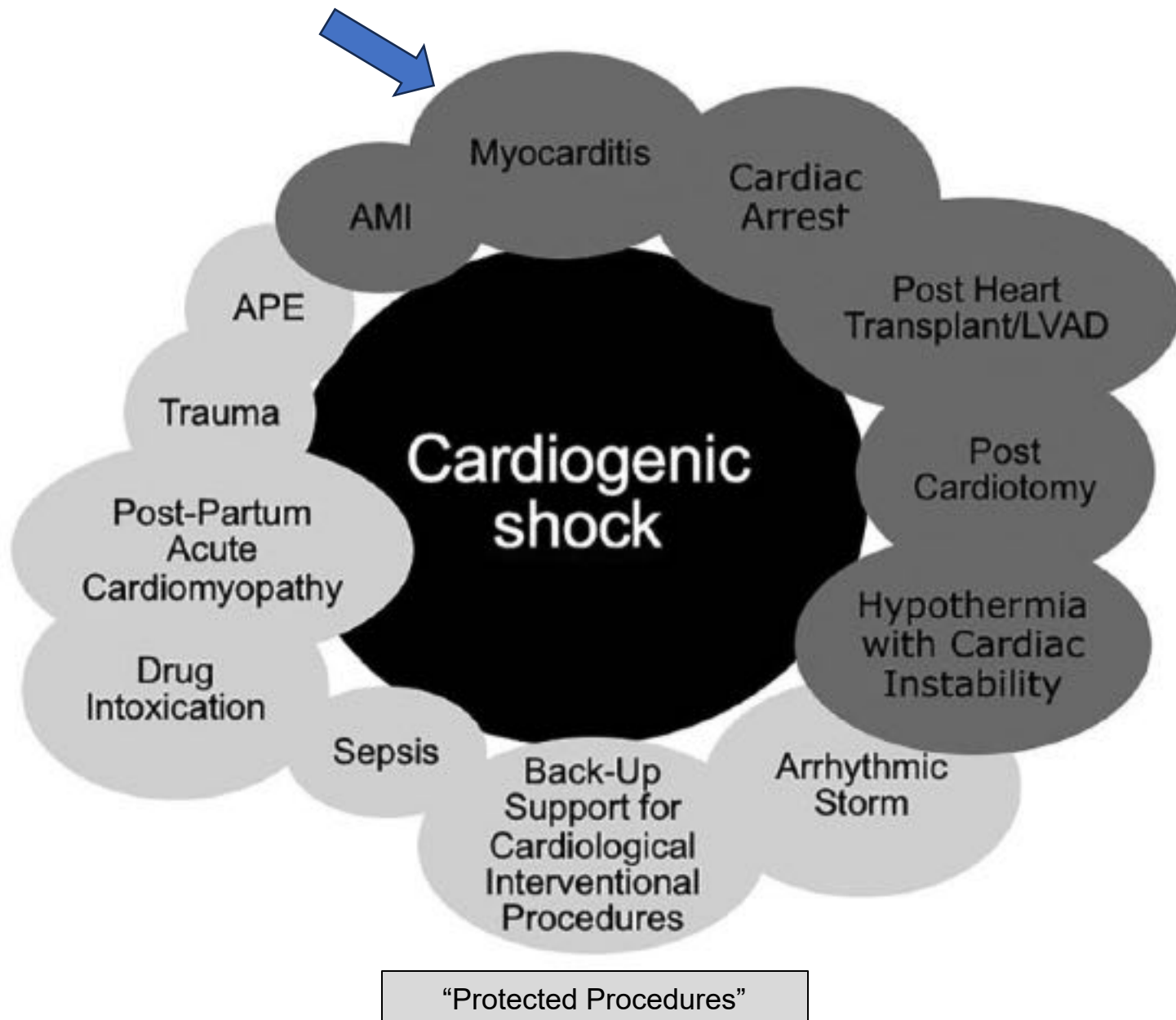


# Cardiogenic Shock Diagnosis



S  
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<b>Congestion</b>	Presence or absence of congestion based on physical signs and hemodynamics; elucidation of ventricular involvement (LV vs RV vs BiV)
<b>Triage</b>	Appropriate triage/shock team activation or possible transfer to a higher level of care



# Case Presentation

## Steps

- Central line & art line inserted
- Norepi escalated to 0.3 mcg/kg/min
- Dobutamine started at 5 mcg/kg/min

## Vitals

- New vitals:  
BP 92/51, HR 141, SpO<sub>2</sub> 92% on FMO<sub>2</sub> 6 L/m

## Labs

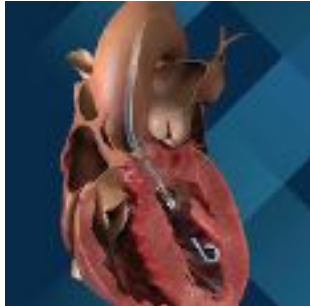
- Creatinine 138 mmol/L, BUN 14
- Lactate 6.9 mmol/L (from 6.2 mmol/L)
- pH 7.20, HCO<sub>3</sub> 14

You call your CS Team  
(or ECMO Team)

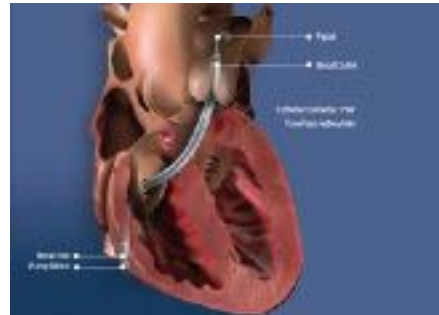
# MCS Choices



IABP



Impella



Impella RP



Protek Duo



VA ECMO



Durable LVAD  
(Heart Mate 3)



## Question 2

What would be the most appropriate MCS device choice for this patient?

- A. Intra-aortic balloon pump
- B. Impella CP
- C. Protek Duo
- D. VA ECMO
- E. Durable LVAD

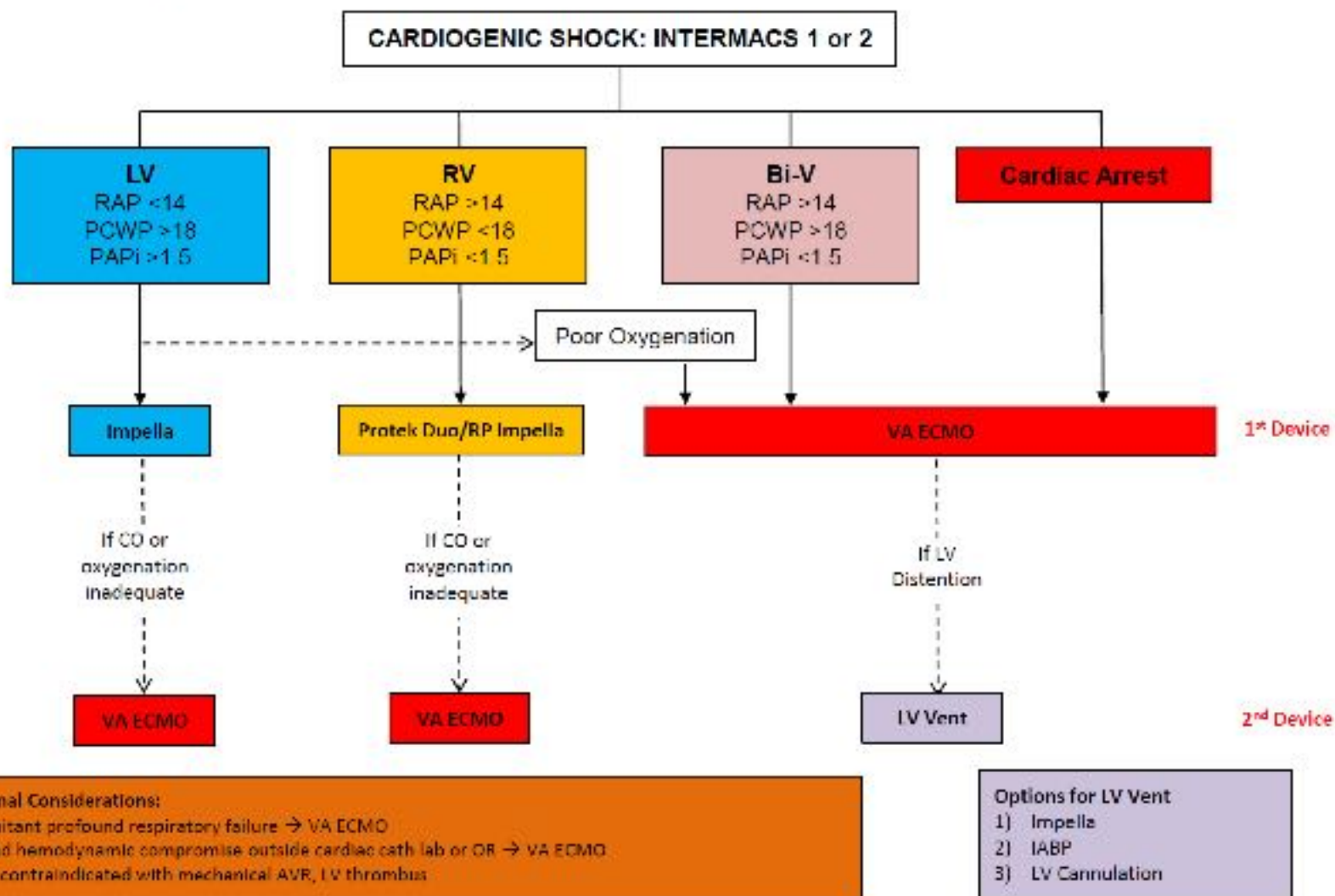
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# Short-Term MCS Decision Algorithm



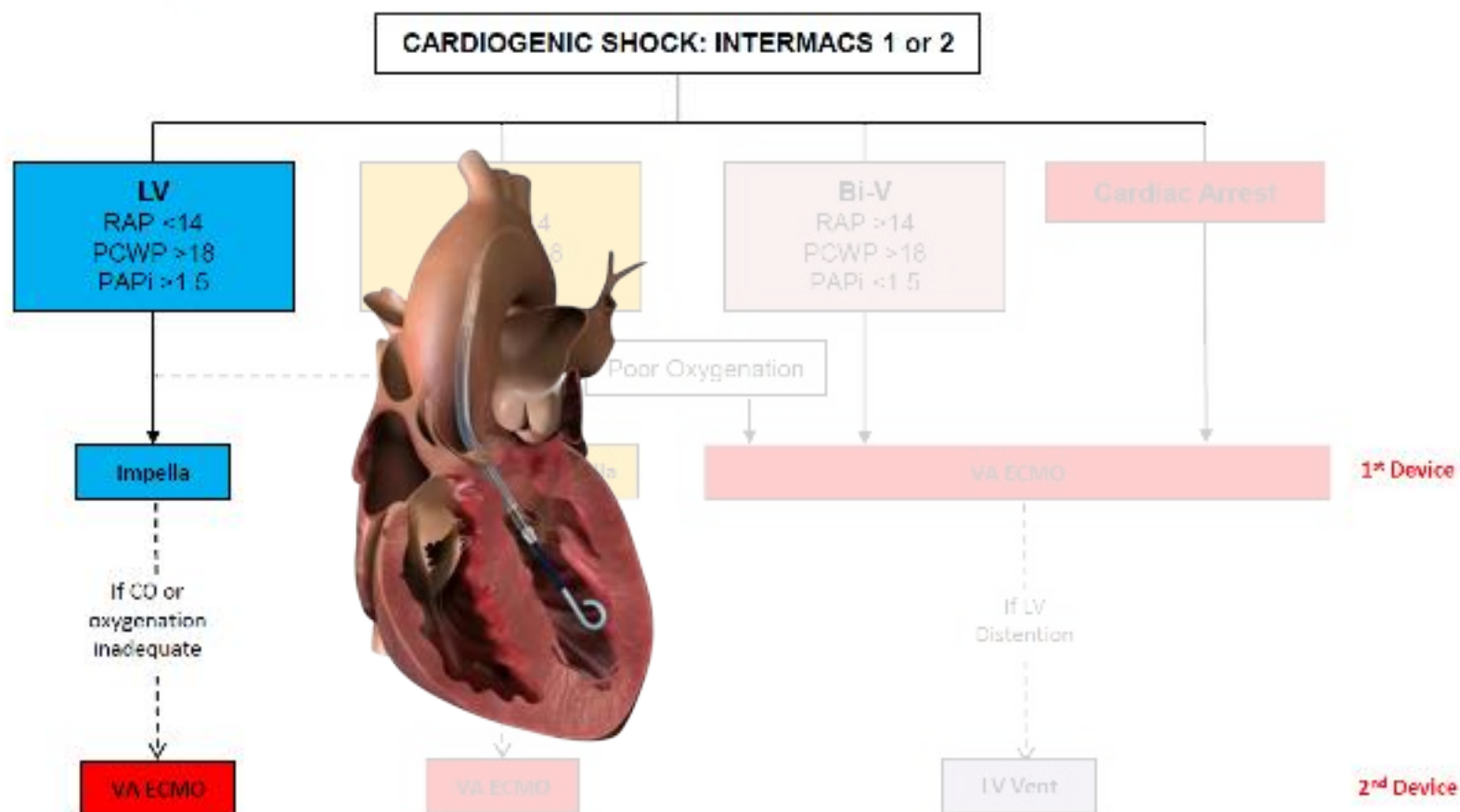




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$$PAPi = (PASP - PADP) / RAP$$

## Short-Term MCS Decision Algorithm



### Additional Considerations:

Concomitant profound respiratory failure → VA ECMO  
Profound hemodynamic compromise outside cardiac cath lab or OR → VA ECMO  
Impella contraindicated with mechanical AVR, LV thrombus

### Options for LV Vent

- 1) Impella
- 2) IABP
- 3) LV Cannulation



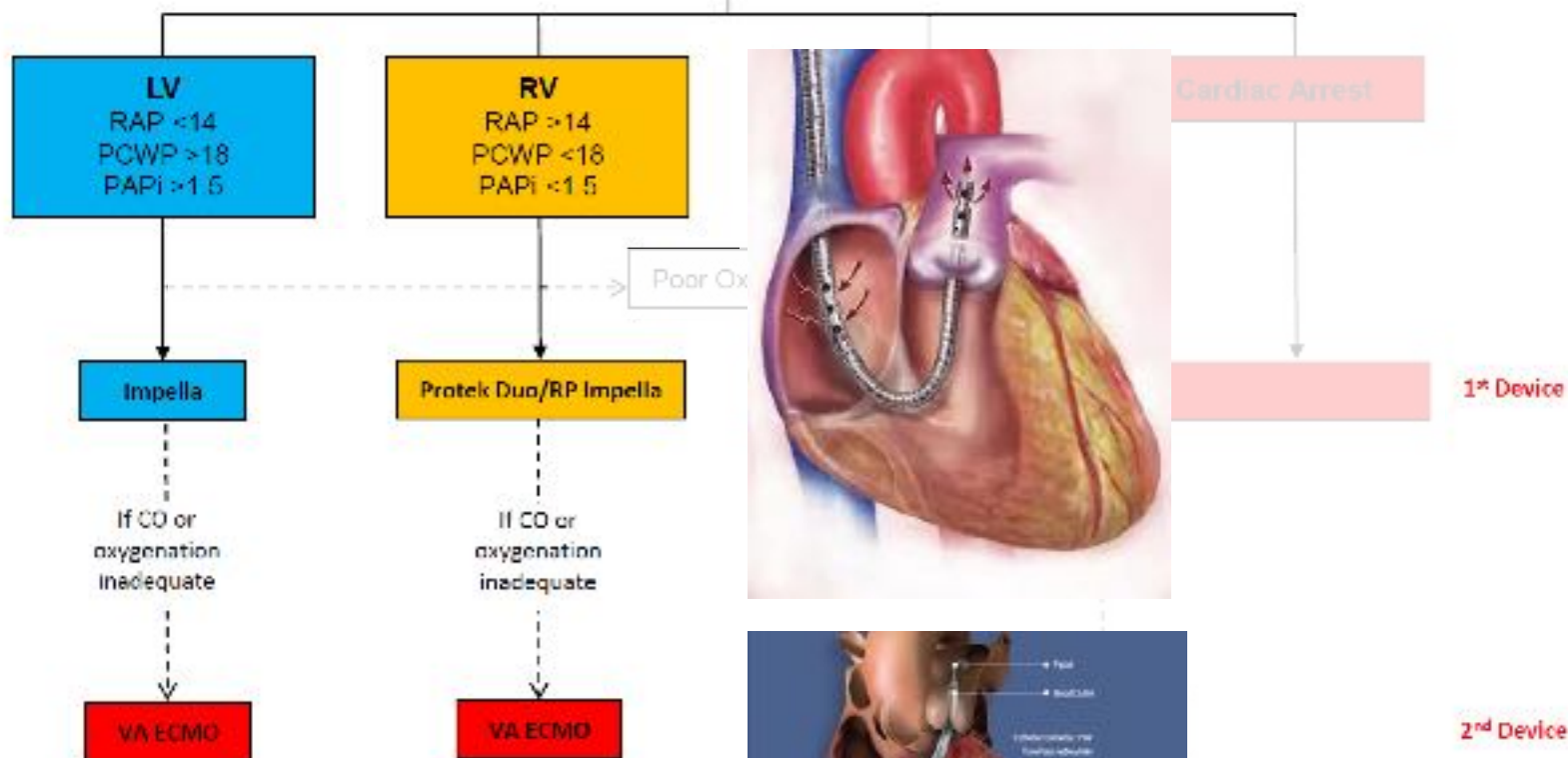


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$$PAPi = (PASP - PADP) / RAP$$

## Short-Term MCS Decision Algorithm

**CARDIOGENIC SHOCK: INTERMACS 1 or 2**



### Additional Considerations:

Concomitant profound respiratory failure → VA ECMO

Profound hemodynamic compromise outside cardiac cath lab or OR → VA ECMO

Impella contraindicated with mechanical AVR, LV thrombus

### Options for LV Vent

Impella

IABP

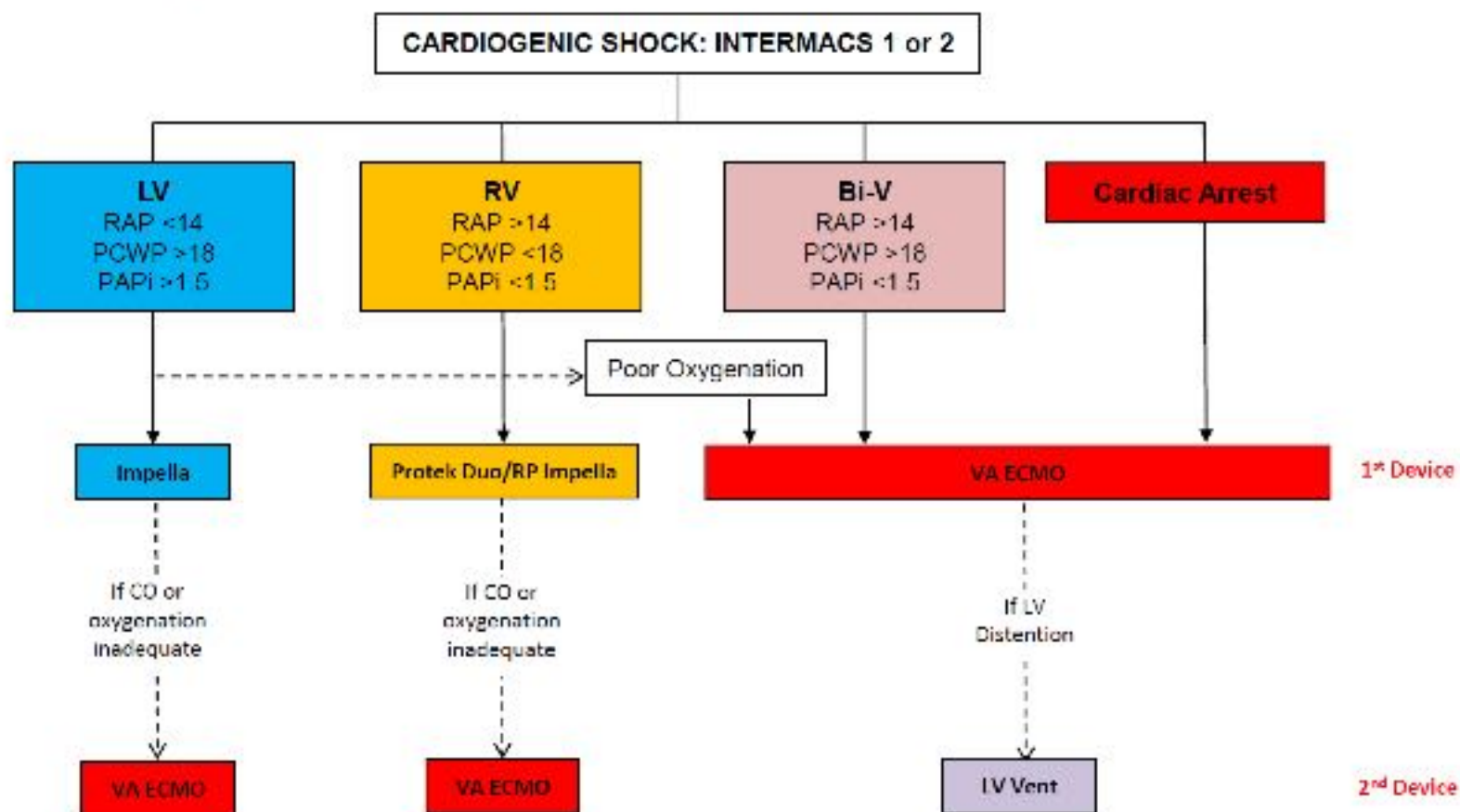
LV Cannulation



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## Short-Term MCS Decision Algorithm



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### Options for LV Vent

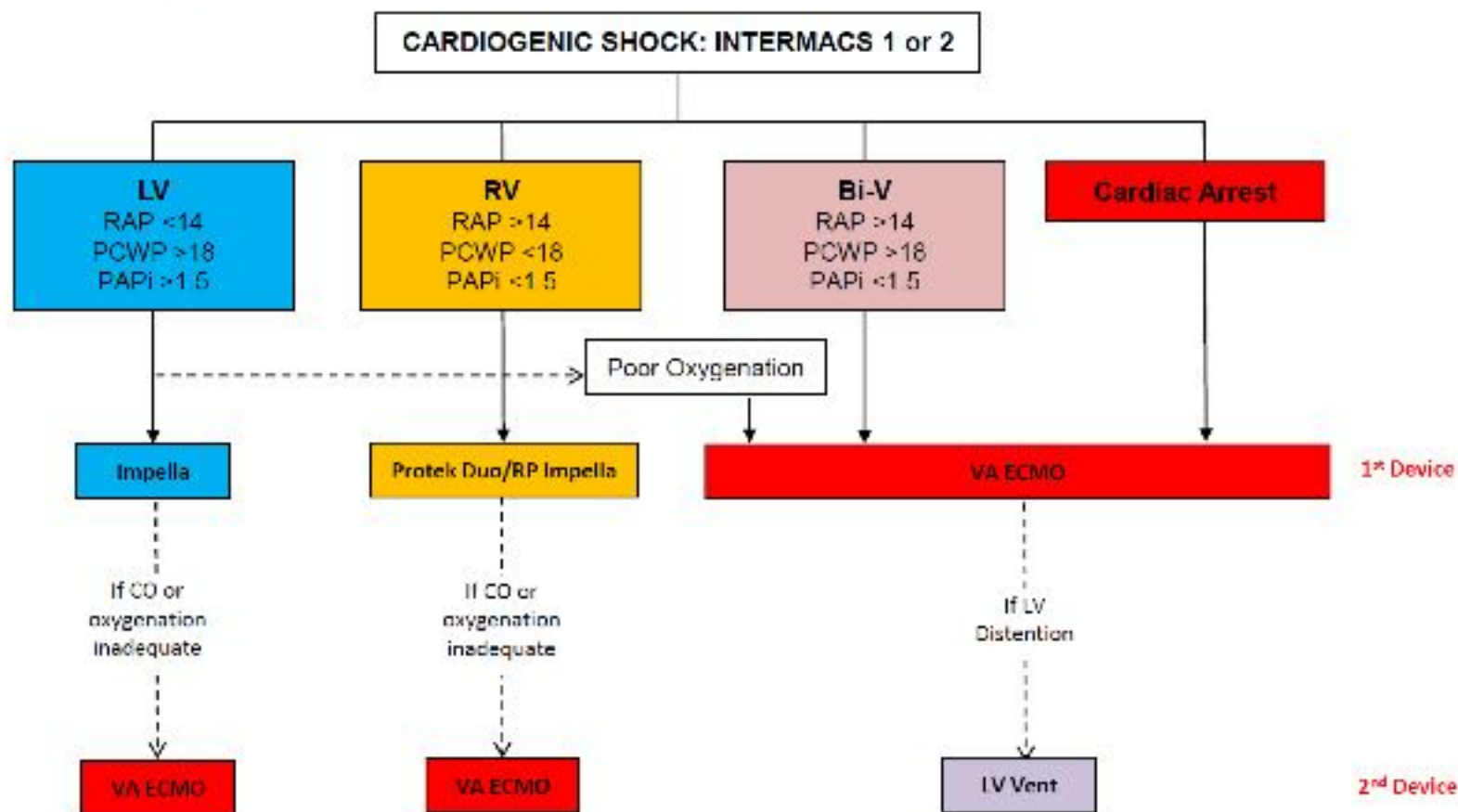
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$$PAPi = (PASP - PADP) / RAP$$

## Short-Term MCS Decision Algorithm



### Additional Considerations:

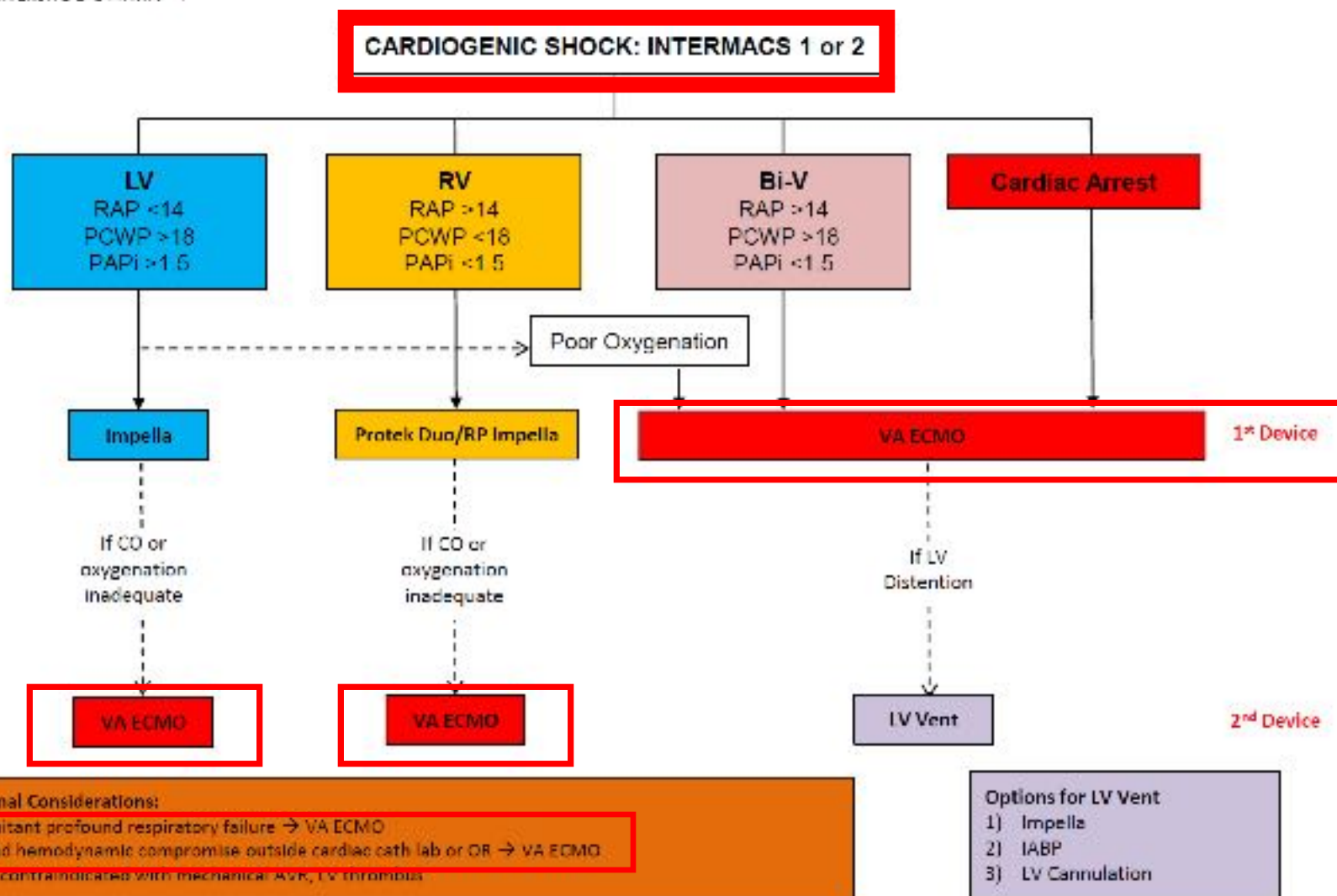
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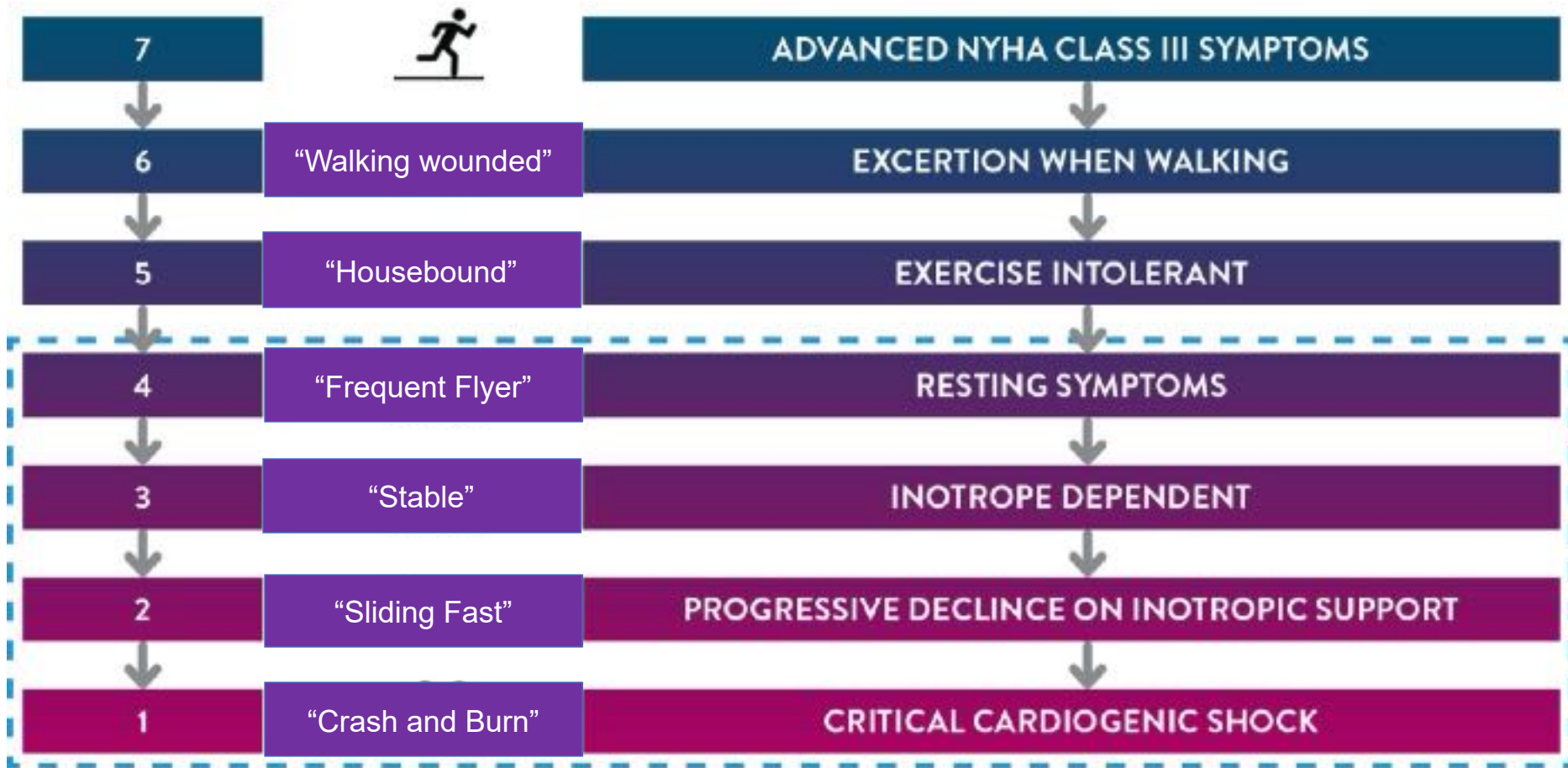
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# Short-Term MCS Decision Algorithm

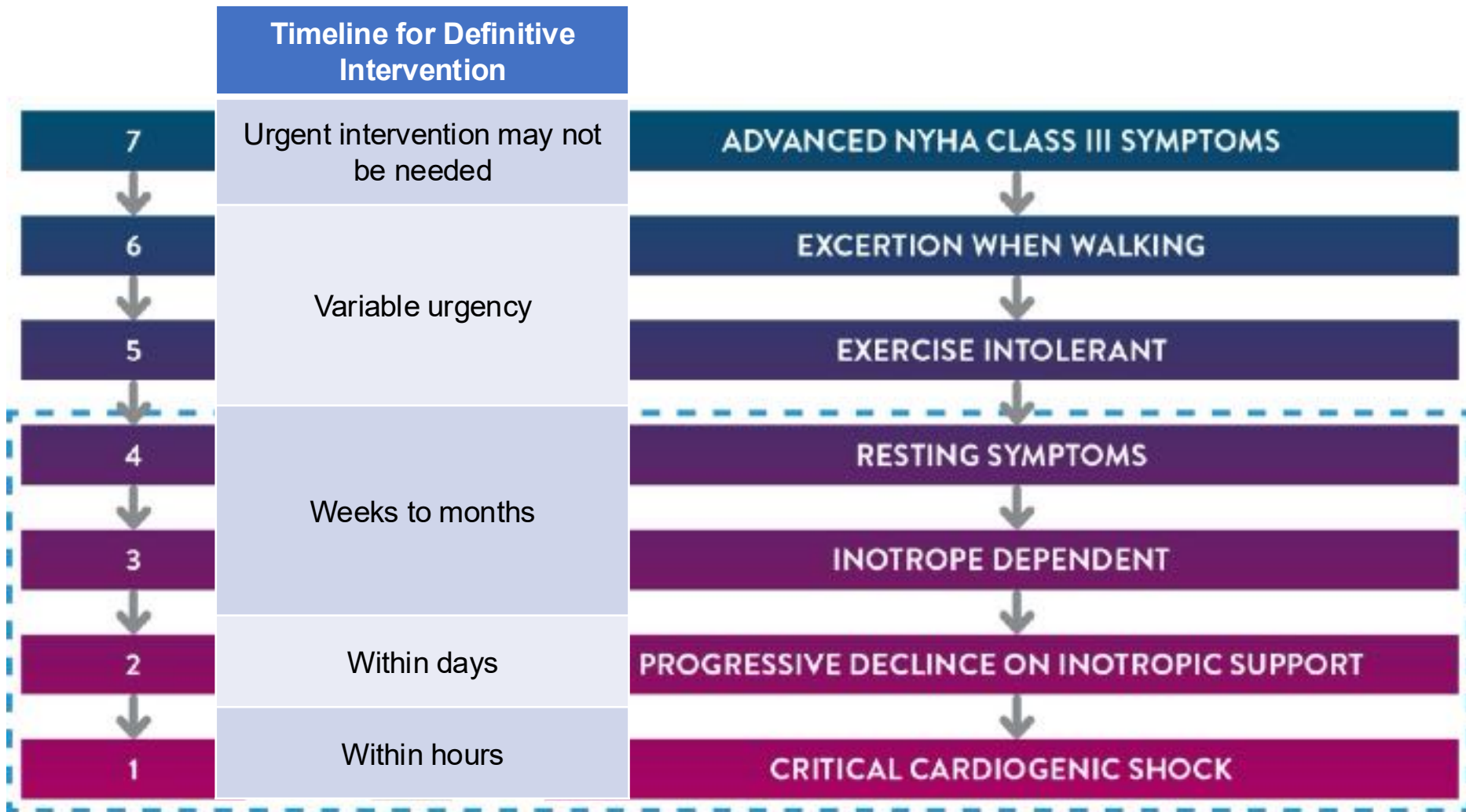


# Intermacros Classification





# Intermacros Classification



# Case Presentation

## Vitals

- Tachypneic, increased work of breathing → Intubated, good SpO<sub>2</sub> on FiO<sub>2</sub> 0.4, AC/PC: PIP/PEEP 20/5

## Steps

- Post-intubation vitals: BP 90/50, HR 141
- Norepi 0.5 mcg/kg/min, Dobutamine 5 mcg/kg/min

## Labs

- Creatinine 138 mmol/L, BUN 14
- Lactate 7.2 mmol/L (from 6.9 mmol/L), pH 7.20, HCO<sub>3</sub> 14

You are still on the phone with  
your CS Team (or ECMO Team)

## Question 3

What is our “time window” for intervention?






- A. 24 hours
- B. 48 hours
- C. 72 hours
- D. 1 week





**ORIGINAL RESEARCH**

# Timing of Initiation of Extracorporeal Membrane Oxygenation Support and Outcomes Among Patients With Cardiogenic Shock

Jacob C. Jentzer , MD; Stavros G. Drakos , MD, PhD; Craig H. Selzman , MD; Clark Owyang, MD; Felipe Teran , MD; Joseph E. Tonna , MD

- ELSO Registry (8,619) patients in cardiogenic shock.
- ECMO initiated **> 24 hours** after admission had a higher risk of in-hospital death after multivariable adjustment (adjusted OR, 1.20 [95% CI, 1.06–1.36];  $P = 0.004$ ).
- Each **12-hour increase** in the time from admission to ECMO initiation was incrementally associated with higher adjusted in-hospital mortality rate (adjusted OR, 1.06 [95% CI, 1.03–1.10];  $P < 0.001$ ).



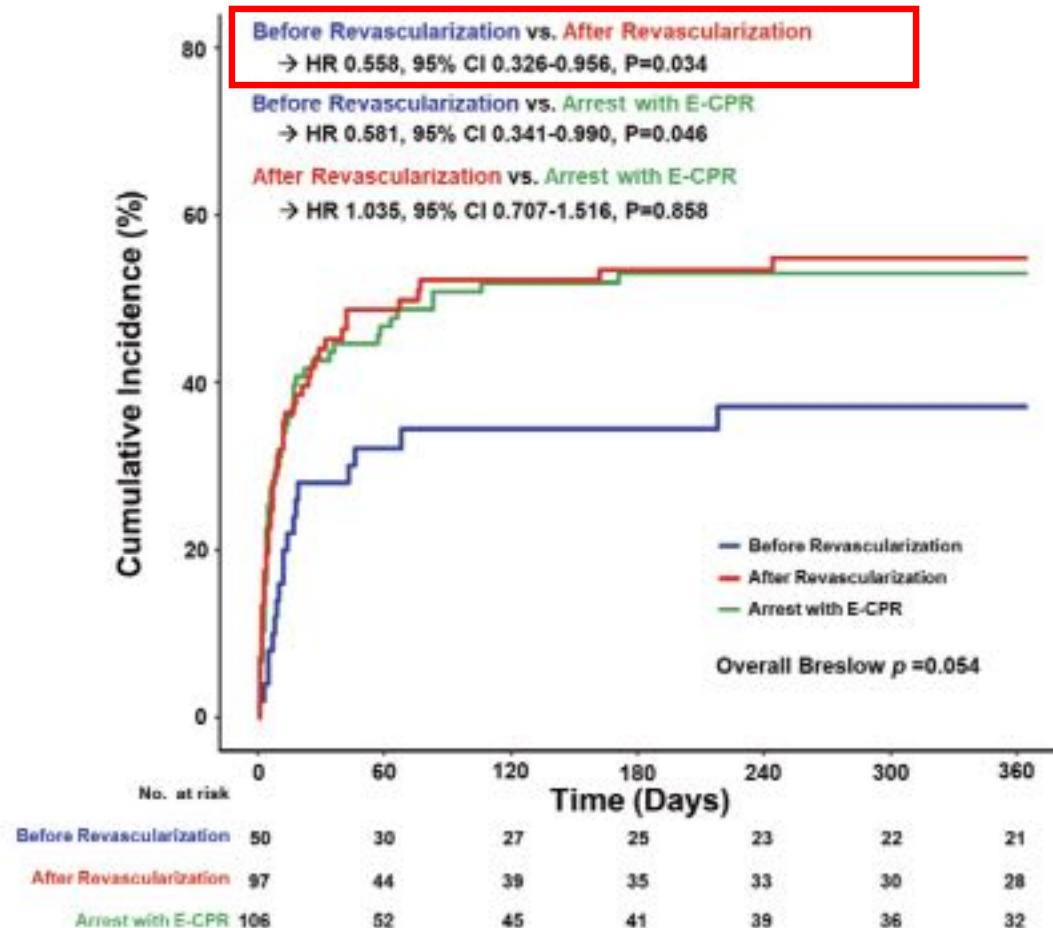
## Optimal Timing of Venoarterial-Extracorporeal Membrane Oxygenation in Acute Myocardial Infarction Patients Suffering From Refractory Cardiogenic Shock

Ki Hong Choi, MD; Jeong Hoon Yang, MD; David Hong, MD; Taek Kyu Park, MD; Joo Myung Lee, MD; Young Bin Song, MD; Joo-Yong Hahn, MD; Seung-Hyuk Choi, MD; Jin-Ho Choi, MD; Su Ryeun Chung, MD; Yang Hyun Cho, MD, PhD; Dong Seop Jeong, MD; Kiick Sung, MD; Wook Sung Kim, MD; Young Tak Lee, MD, PhD; Hyeon-Cheol Gwon, MD

- AMI patients in cardiogenic shock.
- VA ECMO before revascularization (n=50) versus VA ECMO after revascularization (n=97).
- Primary outcome (composite of in-hospital mortality, LVAD implantation, heart transplantation) **significantly lower in VA-ECMO before revascularization** than in VA-ECMO after revascularization (32.0% vs. 49.5%, OR 0.480, 95% CI 0.235-0.982, P=0.045).

# Optimal Timing of VA ECMO in AMI Patients with Refractory CS

## A. All-cause Death, LVAD Implantation or Heart Transplantation



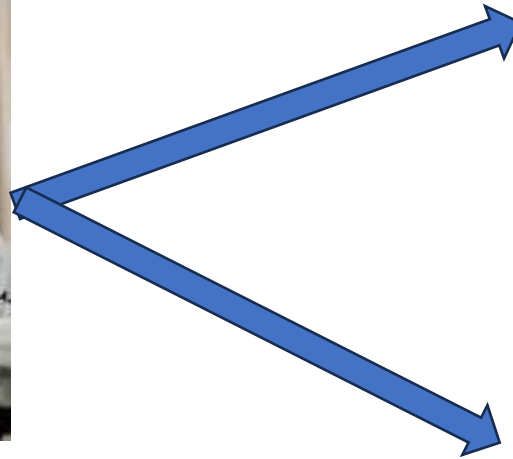
## Question 3

What is our “time window” for intervention?

- A. 24 hours
- B. 48 hours
- C. 72 hours
- D. 1 week



# ECMO Triage



VV ECMO



VA ECMO



## Question 4

Which comorbidity is an **absolute contraindication** to VA ECMO?

- A. Hypothyroidism
- B. Obesity
- C. Moderate aortic incompetence
- D. Mild NASH cirrhosis (non-alcoholic steatohepatitis)

# Patient Selection



Bridge-to-Decision

- Recovery
- Heart Transplant
- Long-term MCS
- Withdrawal of Life Support

# Contraindications

- **Cardiac recovery unlikely** and not a candidate for heart transplant or durable LVAD.
- Non-recoverable **advanced comorbidity** with poor life expectancy (end-stage peripheral-organ diseases, malignant tumor, massive pulmonary embolisms in cancer patients, chemotherapy-induced chronic cardiomyopathy, etc.)
- **Severe neurologic impairment** (i.e., prolonged anoxic brain damage, extensive trauma and bleeding)
- Liver cirrhosis (Child-Pugh class B and C)

VA ECMO will not be of benefit



# Contraindications

- **Moderate to severe aortic valve regurgitation**
- **Acute Type A or B aortic dissection** with extensive aortic branches (ascending, supra-aortic and femoral) involvement
- Severe immunologic disease with marked blood and coagulation disorders
- Severe vascular disease with extensive aortic and peripheral vessel involvement (calcification, stenosis, and closure), including axillary arteries

VA ECMO may make the patient worse  
Or will be too challenging to insert

## Question 4

Which comorbidity is an **absolute contraindication** to VA ECMO?

A. Hypothyroidism

B. Obesity

C. Moderate aortic incompetence

D. Mild NASH cirrhosis (non-alcoholic steatohepatitis)



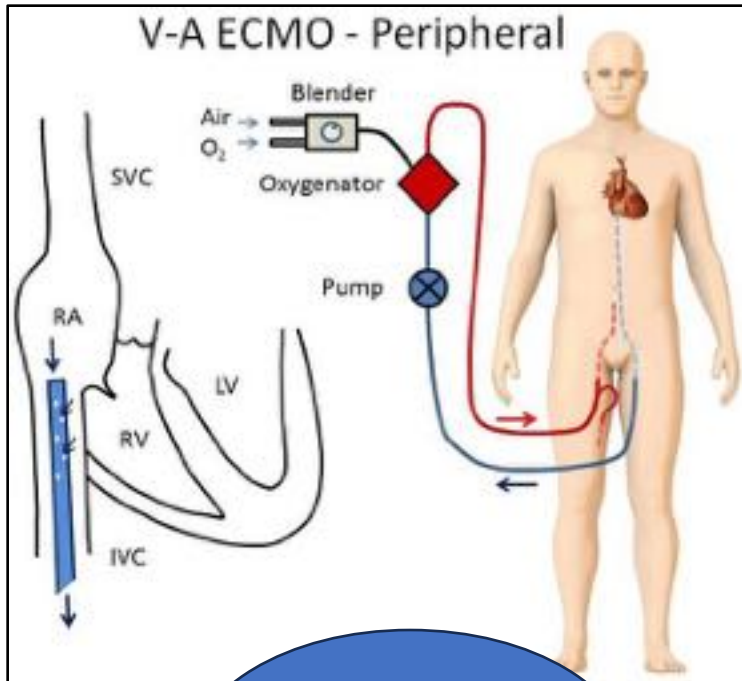
Luckily, our patient has  
no contraindications

## Question 5

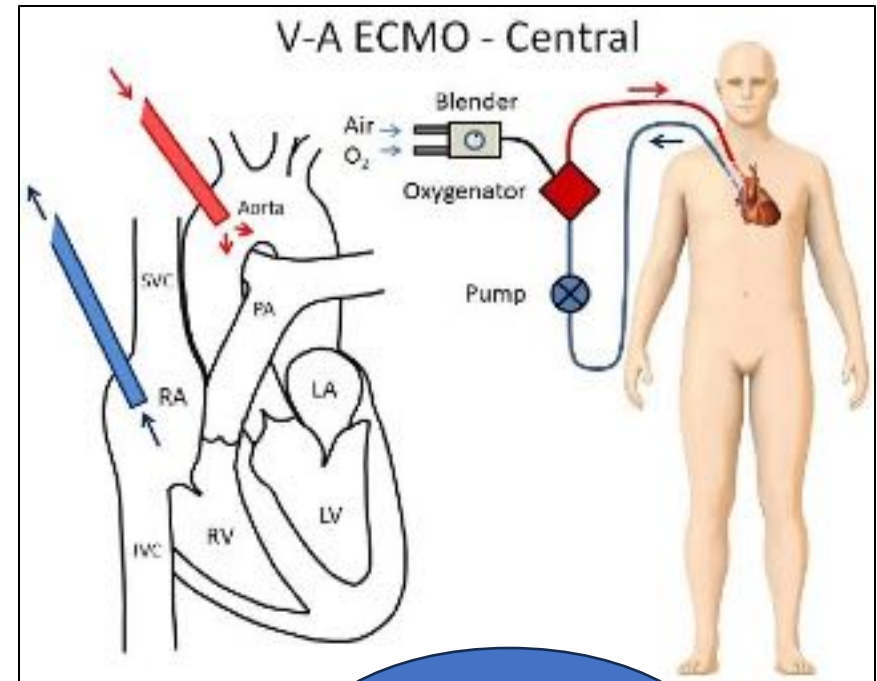
What is the best ECMO approach for this patient?

- A. Central VA ECMO
- B. Peripheral VA ECMO with a femoral-femoral approach
- C. Peripheral VA ECMO with an internal jugular-subclavian approach
- D. Peripheral V-AV ECMO with femoral-femoral/internal jugular approach

# VA ECMO Configurations



Fem-fem  
cannulation



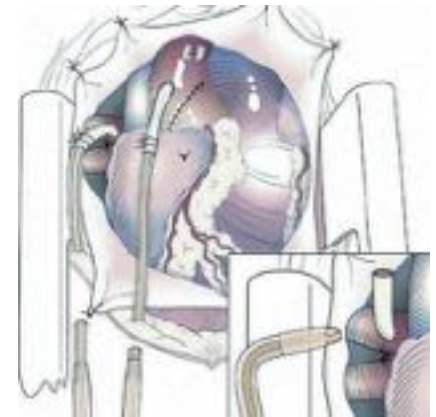
Atrio-aortic  
cannulation

# Central VA ECMO

- Drain RA directly and return to ascending aorta

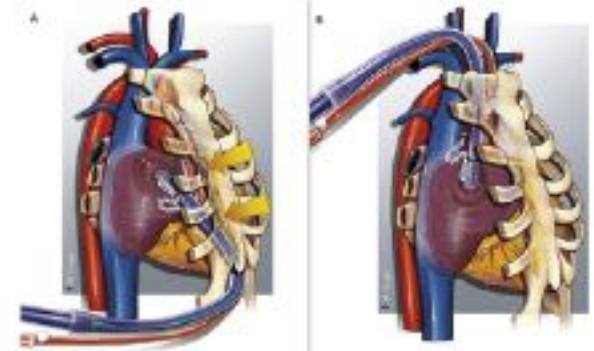
## Advantages:

- Same CPB cannulas used for ECMO (may bring out below sternum to close the chest).
- Avoids femoral artery puncture (often in a fully heparinized patient), avoids lower limb ischemia, differential hypoxia.
- Higher flows
- More options for venting.

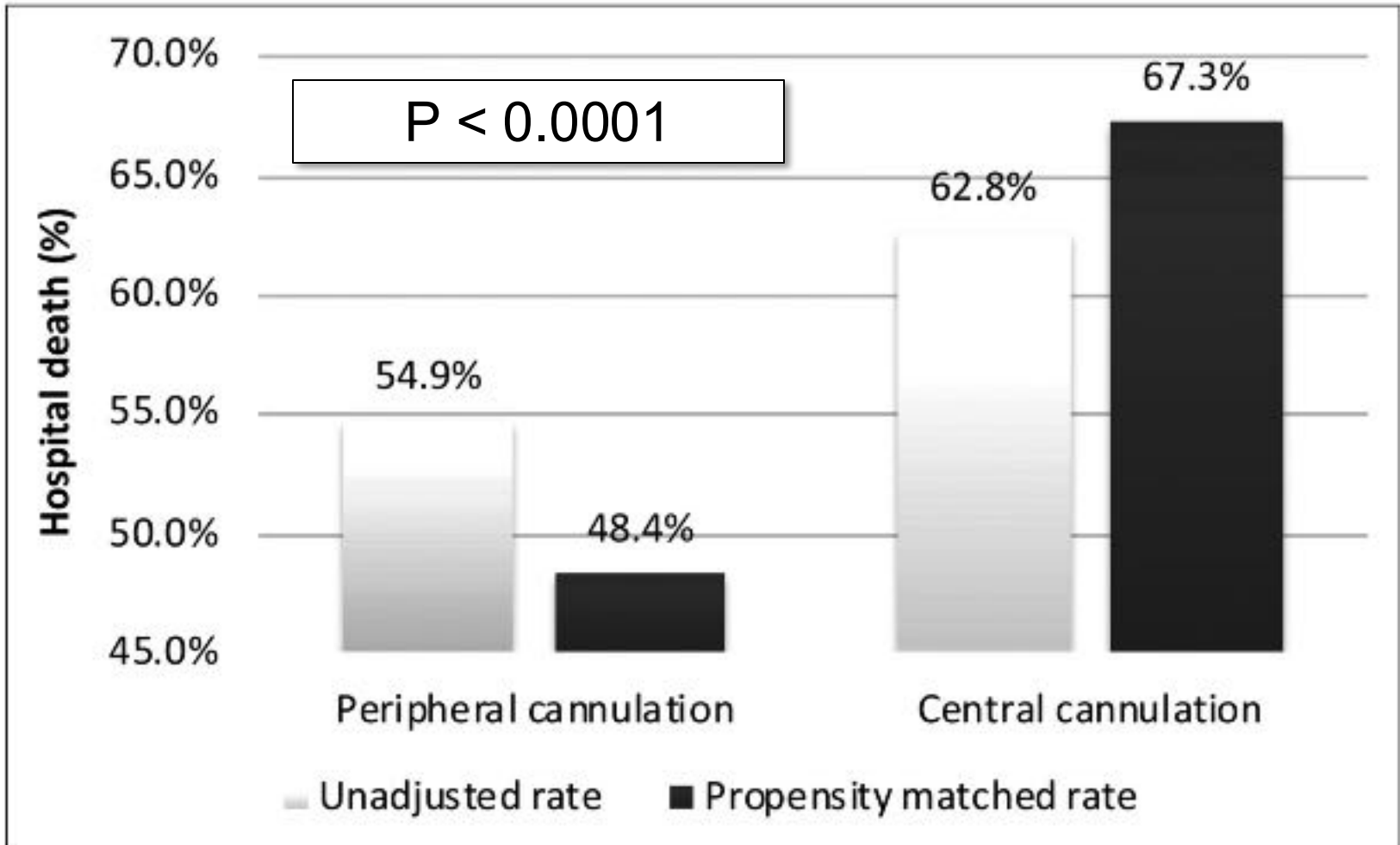


## Disadvantages

- Needs reopening for decannulation.
- Higher risk of mediastinitis, bleeding.
- Higher risk of strokes.
- Generally, carries a higher mortality



# Postcardiotomy ECLS



# Peripheral VA ECMO

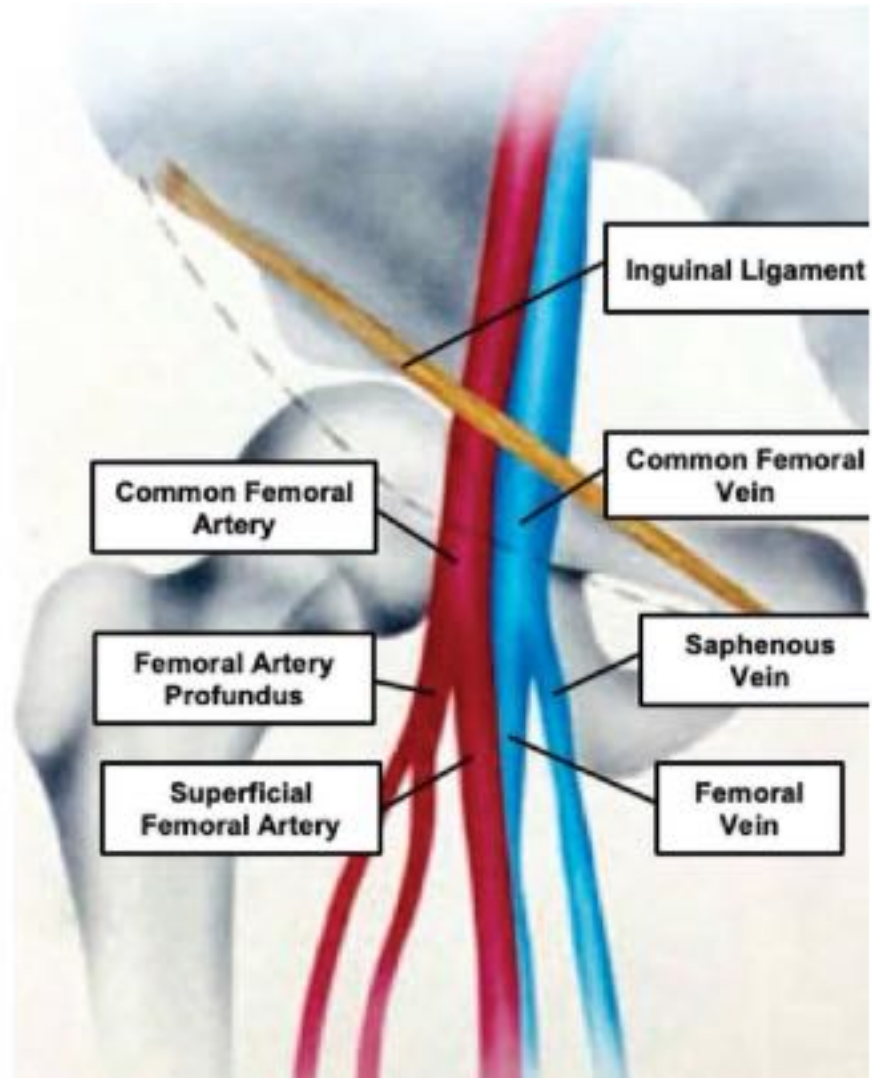
## Puncture points:

- Common femoral vein
- Common femoral artery (just below the inguinal ligament and above their respective bifurcations)

## Sequence:

- Start with venous (drainage cannula)
- Heparinize after guidewire in vein (target ACT 180-220)

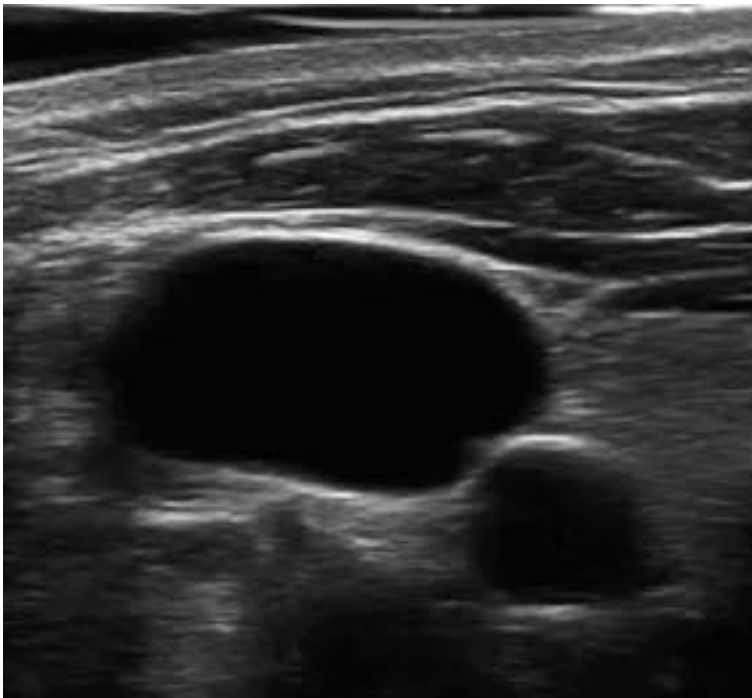
**Drainage**  
21 – 25 Fr



# Peripheral VA ECMO

Percutaneous

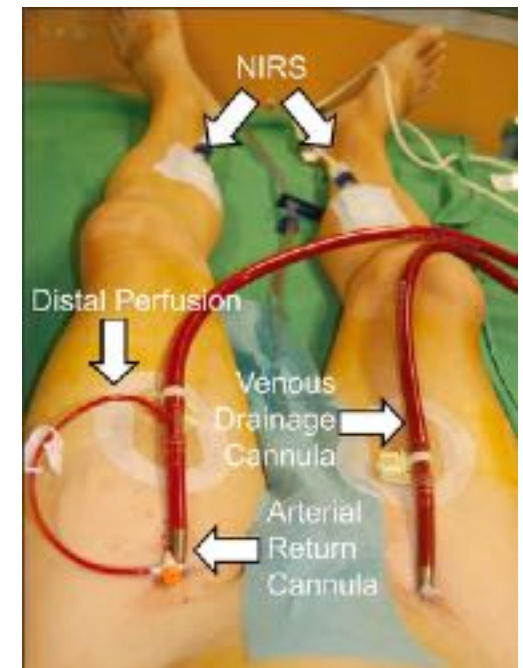
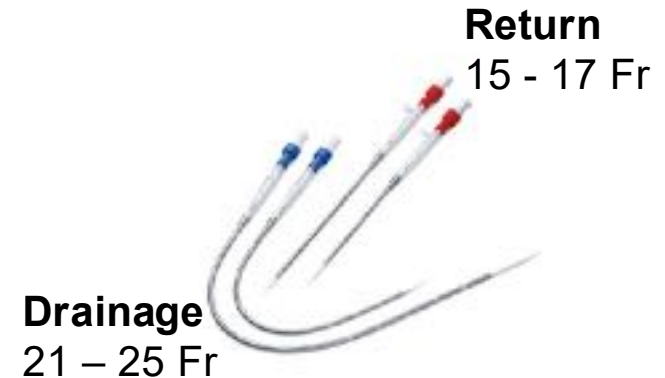
Open





# Peripheral VA ECMO

- Arterial cannula: 15 – 17 Fr adequate. When higher flow needed (e.g. septic patient), 19 -21 Fr may be used
- Larger cannulas associated with more vascular complications e.g. limb ischemia
- Opinion:
  - arterial and venous cannulas in separate limbs to reduce vascular complications and to facilitate decannulation.
  - If feasible, the venous cannula should be placed in right femoral vein (more direct path to IVC and right atrium)



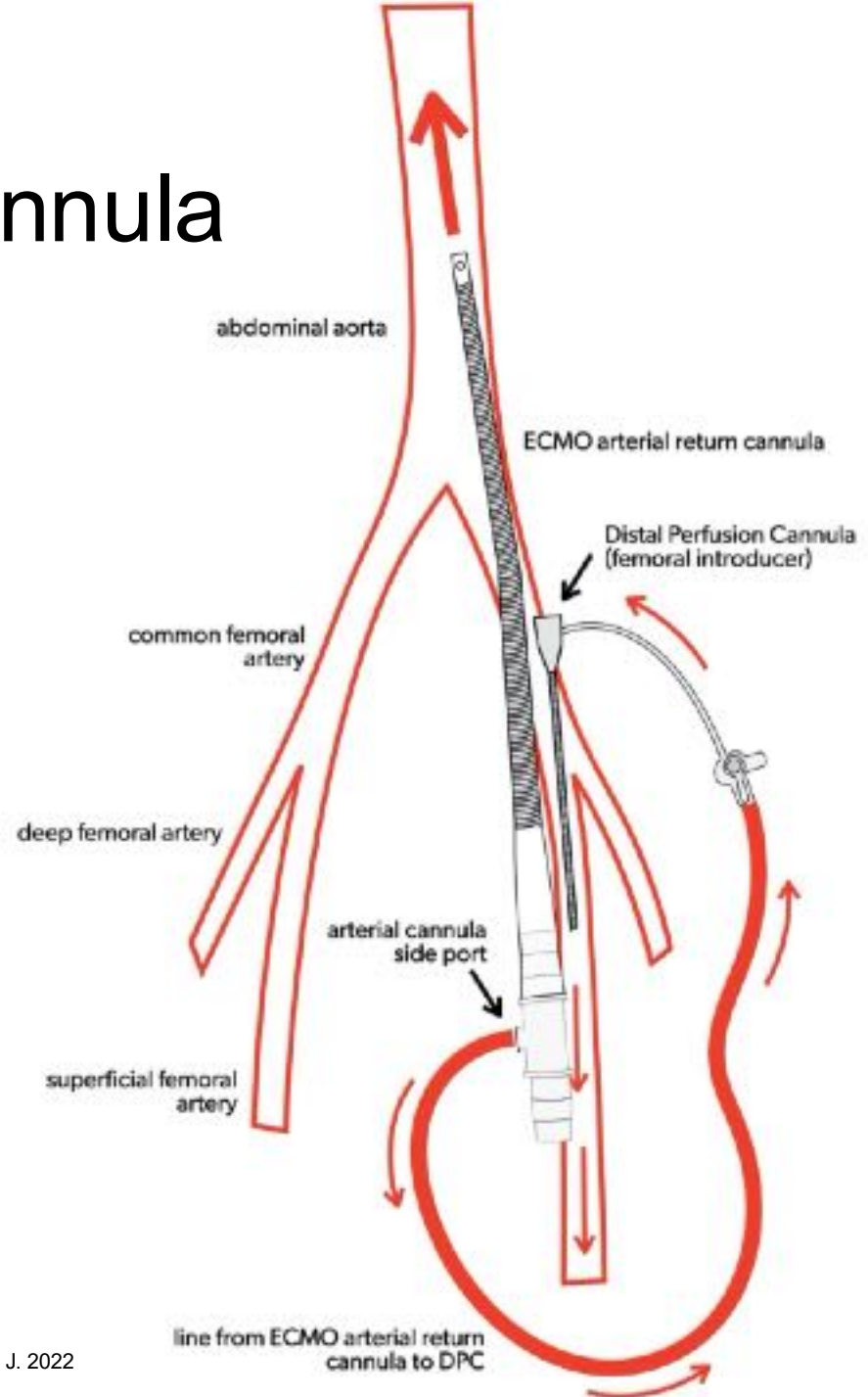
# Limb Ischemia – Prevention

## Distal Perfusion Cannula (DPC)

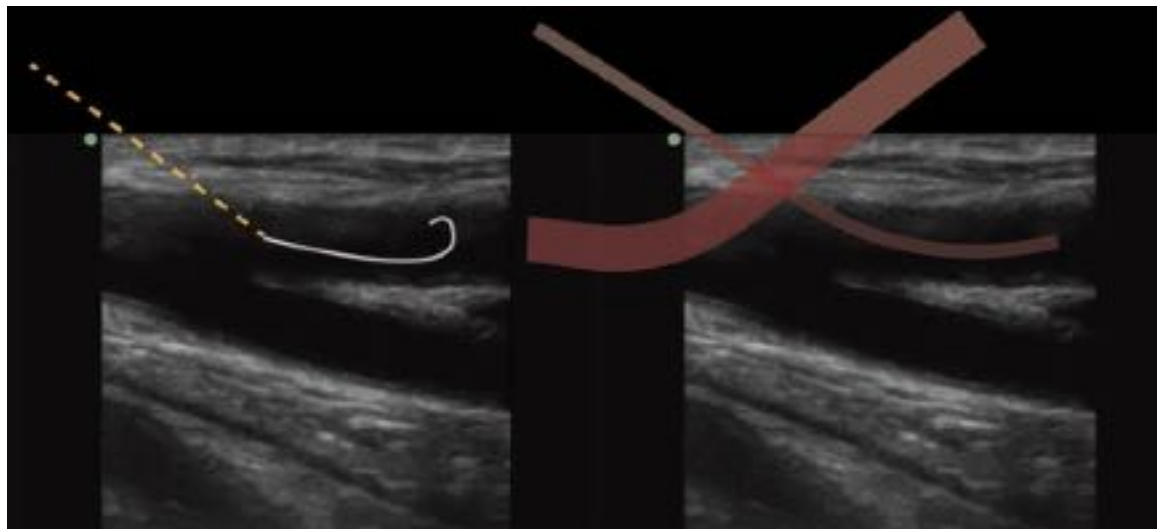
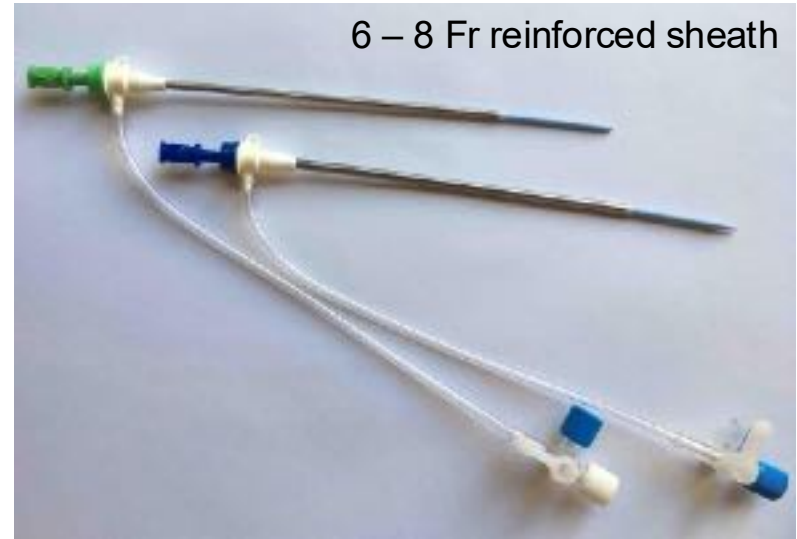
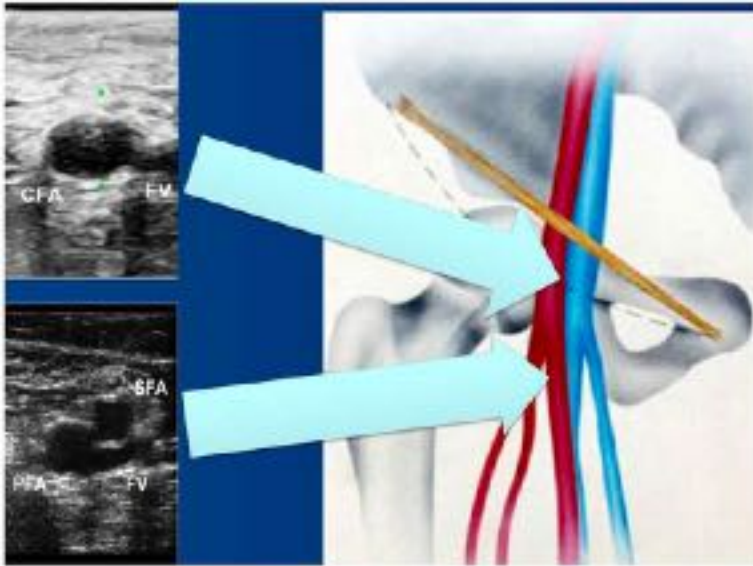
Set-up	Incidence of Lower Extremity Ischemia
Without DPC	7%
With DPC	0-3%

# Distal Perfusion Cannula

- Placed under US guidance or direct vision into the **superficial femoral artery**.
- Puncture point maybe common femoral artery (above bifurcation), then directed to superficial femoral.
- May confirm guide wire in superficial femoral artery by US or fluoroscopy
- Confirm flow in popliteal artery by ultrasound and continuous flow Doppler.

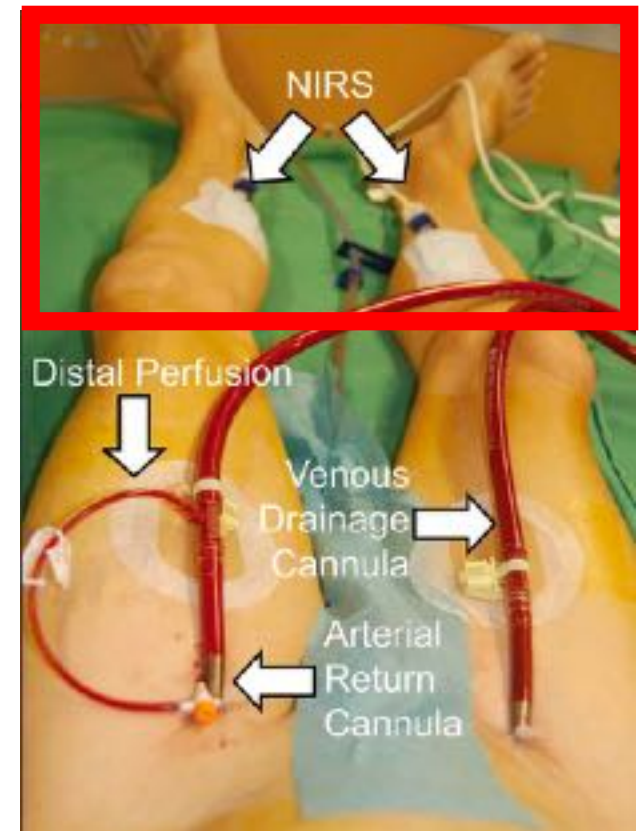


# Distal Perfusion Cannula



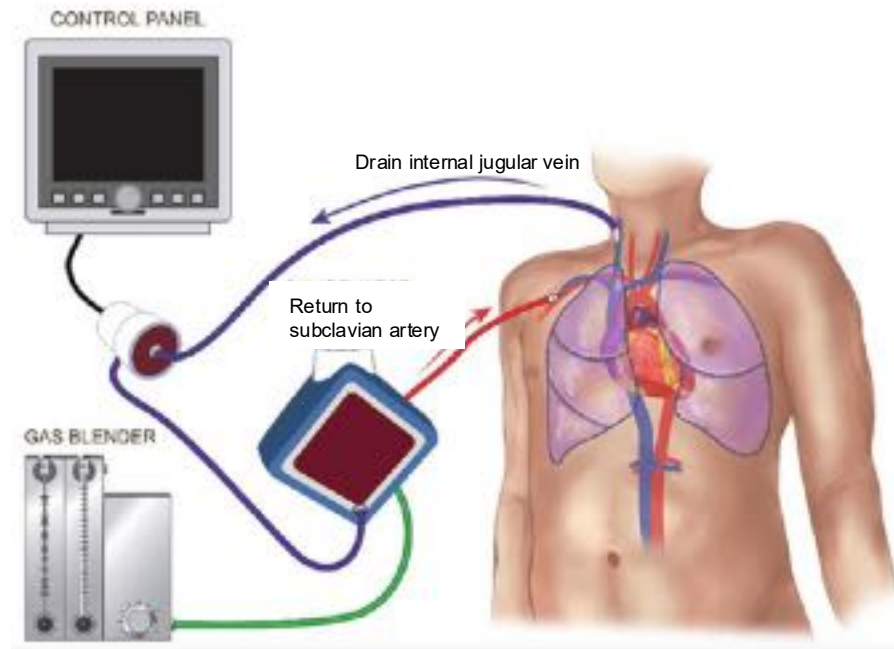
# Distal Perfusion Cannula

- With small return cannulas (15 Fr & 17 Fr), distal perfusion not always needed.
- Decision making may be informed by Near-Infrared Spectroscopy (NIRS).
- DPC needed with NIRS values:
  - < 50% (preferably < 60%)
  - > 20% difference between extremities
- Insertion timing:
  - On ECMO institution
  - Salvage intervention



# Alternative Peripheral Cannulation Approach

- Drain RA via internal jugular
- Return to subclavian or axillary artery via surgical cut-down
- **Advantages:**
  1. May be of use in patients with severe peripheral vascular disease
  2. Avoid complications of femoral artery site
  3. Minimize differential oxygenation
  4. Facilitate patient mobilization in case of predicted long run
- **Disadvantages:**
  1. Smaller size cannulas, less flow
  2. Return cannula by cut-down



## Question 5

What is the best ECMO approach for this patient?

A. Central VA ECMO

B. Peripheral VA ECMO with a femoral-femoral approach

C. Peripheral VA ECMO with an internal jugular-subclavian approach

D. Peripheral V-AV ECMO with femoral – femoral / internal jugular approach



## Question 6

6. She is now cannulated, and VA ECMO has been established.

The patient's younger brother is a medical student. He is asking what the likelihood is that she'll survive?

Her approximate average survival is:

- A. 80%
- B. 60%
- C. 40%
- D. 20%

### 26-year-old female

Viral myocarditis

64 kg

On norepi & dobutamine  
BP: 92/51 mmHg

Intubated 4 hours ago on AC PC  
PIP/PEEP: 20 / 5 cmH<sub>2</sub>O

Creatinine: 138 mmol/L

Lactate 6.9, pH 7.20, HCO<sub>3</sub>: 14 mmol/L

No liver failure (yet)



# Risk Scores for VA-ECMO

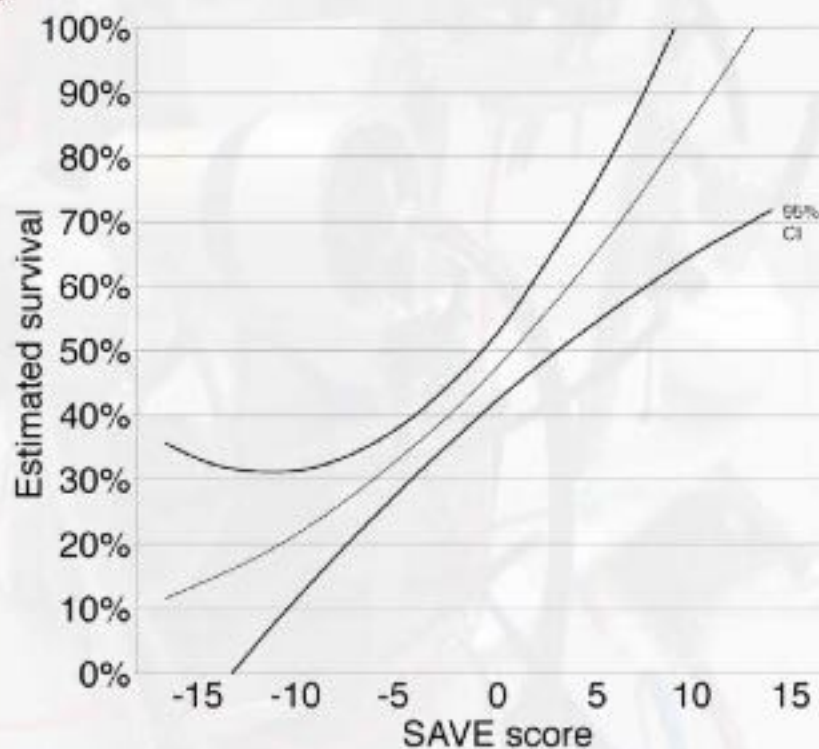
Risk Score Name <sup>Reference</sup>	VA ECMO Setting	Predictors
ENCOURAGE Score <sup>7</sup>	Cardiogenic shock (post AMI)	Age, sex, BMI, Glasgow Coma Scale, creatininemia, serum lactate value, prothrombin activity
REMEMBER Score <sup>8</sup>	Post cardiectomy cardiogenic shock (post CABG)	Older age, left main coronary artery disease, Inotropic score, CK-MB, serum creatinine, platelet count
CARDShock Score <sup>9</sup>	Cardiogenic shock	Older age, neurologic status, previous myocardial infarction or CABG, blood lactate value, acute coronary syndrome etiology, LV systolic dysfunction, estimated glomerular filtration

# SAVE

Survival After Veno-arterial ECMO

The SAVE Score has been developed by [ELSO](#) and [The Department of Intensive Care at The Alfred Hospital, Melbourne](#). It is designed to assist prediction of survival for adult patients undergoing Extra-Corporeal Membrane Oxygenation for refractory cardiogenic shock. It should not be considered a substitute for clinical assessment.

For more information see: [Predicting survival after ECMO for refractory cardiogenic shock: the survival after veno-arterial-ECMO \(SAVE\)-score](#)



The patient's SAVE Score is

Diagnosis: ☒

Myocarditis

Refractory VT/VF

Post heart or lung transplantation

Congenital heart disease

Other diagnoses

Age (years):

18-38 ☐

39-52 ☐

53-62 ☐

≥63 ☐

Weight (kg):

<65 ☐

65-89 ☐

≥90 ☐

Cardiac:

Pulse pressure pre ECMO ≤20 mmHg ☒

Diastolic BP pre ECMO ≥40 mmHg ☒

Pre-ECMO cardiac arrest

Respiratory:

Peak inspiratory pressure ≤20 cmH<sub>2</sub>O

Intubation duration pre ECMO (hrs)

≤10 ☐

11-29 ☐

≥30 ☐

Renal:

Acute renal failure ☒

Chronic renal failure ☐

HCO<sub>3</sub> pre ECMO ≤15 mmol/L ☒

Other organ failures pre ECMO:

Central nervous system dysfunction ☒

Liver failure ☒

☐

☐

☐

☐

☐

☐

# Question 5

The patient's SAVE Score is

5

## Diagnosis:

Myocarditis  
Refractory VT/VF  
Post heart or lung transplantation  
Congenital heart disease  
Other diagnoses

☒ Yes  
☐ No  
☐ No  
☐ No  
☐ No

## Age (years):

18-38 ☒  
39-52 ☐  
53-62 ☐  
≥63 ☐

## Weight (kg):

<65 ☒  
65-89 ☐  
≥90 ☐

## Cardiac:

Pulse pressure pre ECMO ≤20 mmHg  
Diastolic BP pre ECMO ≥40 mmHg  
Pre-ECMO cardiac arrest

☐ Yes  
☒ No  
☐ No

## Respiratory:

Peak inspiratory pressure ≤20 cmH<sub>2</sub>O  
Intubation duration pre ECMO (hrs)

☒ Yes  
☐ No  
≤10 ☒  
11-29 ☐  
≥30 ☐

## Renal:

Acute renal failure  
Chronic renal failure  
HCO<sub>3</sub> pre ECMO ≤15 mmol/L

☒ Yes  
☐ No  
☒ No

## Other organ failures pre ECMO:

Central nervous system dysfunction  
Liver failure

☐ Yes  
☐ No  
☒ No

26-year-old female

Viral myocarditis

64 kg

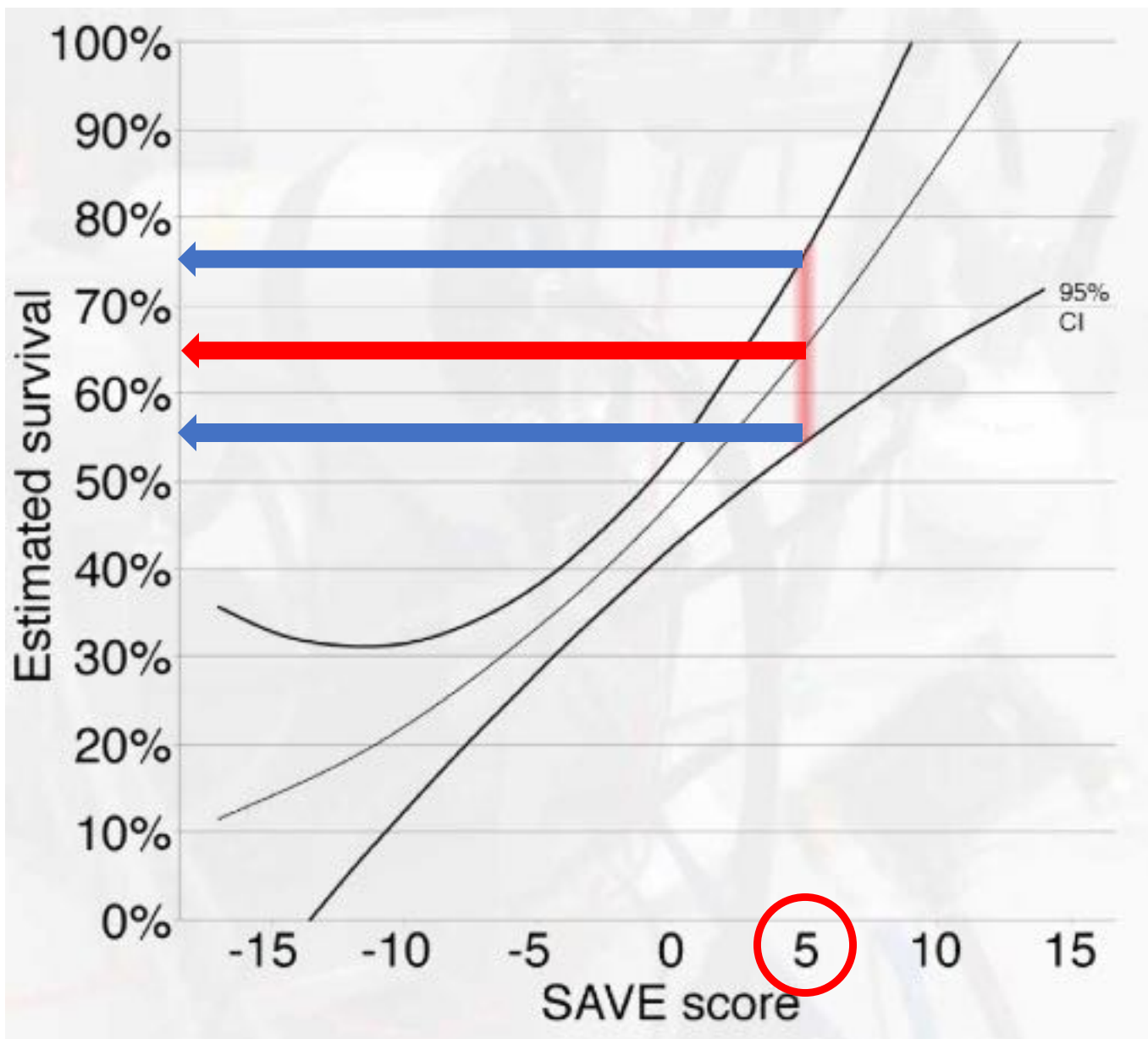
On norepi & dobutamine  
BP: 92/51 mmHg

Intubated 4 hours ago on AC PC  
PIP/PEEP: 20 / 5 cmH<sub>2</sub>O

Creatinine: 138 mmol/L

Lactate 6.9, pH 7.20, HCO<sub>3</sub>: 14 mmol/L

No liver failure (yet)



## Question 6

She is now cannulated, and VA ECMO is established. The patient's younger brother is a medical student. He is asking what is the likelihood she'll survive?

- A. Average survival 80%
- B. Average survival 60%**
- C. Average survival 40%
- D. Average survival 20%

### 26-year-old female

Viral myocarditis

64 kg

On norepi & dobutamine  
BP: 92/51 mmHg

Intubated 4 hours ago on AC PC  
PIP/PEEP: 20 / 5 cmH<sub>2</sub>O

Creatinine: 138 mmol/L

Lactate 6.9, pH 7.20, HCO<sub>3</sub>: 14 mmol/L

No liver failure (yet)

# Outline

Patient Selection

VA ECMO Initiation

VA ECMO Goals



# VA ECMO Goals

- Provides hemodynamic support
- Restores end-organ perfusion
- Buys time to heal

## **Rationale for VA ECMO:**

- Bedside cannulation
- Biventricular support
- Respiratory support

# VA ECMO Goals


## **Restore Organ Perfusion**

- Reduce inotropes
- Reduce vasopressors
  
- Assess for lactate normalization
- Improvement in urine output
- Resolution of shock liver



# Case Resolution

- 
- Decannulated on ECMO day 19

- 
- Discharged from ICU day 31
  - Discharged from hospital day 47

- 
- Came back to visit us in ICU!

# Outline

Patient Selection

VA ECMO Initiation

VA ECMO Goals



Thank you!