

Quelle Assistance pour le Cœur ?

Dr. Jean Beissel
INCCI



Assistance Circulatoire

Choc Cardiogénique

PCI à haut risque



Le Choc Cardiogénique

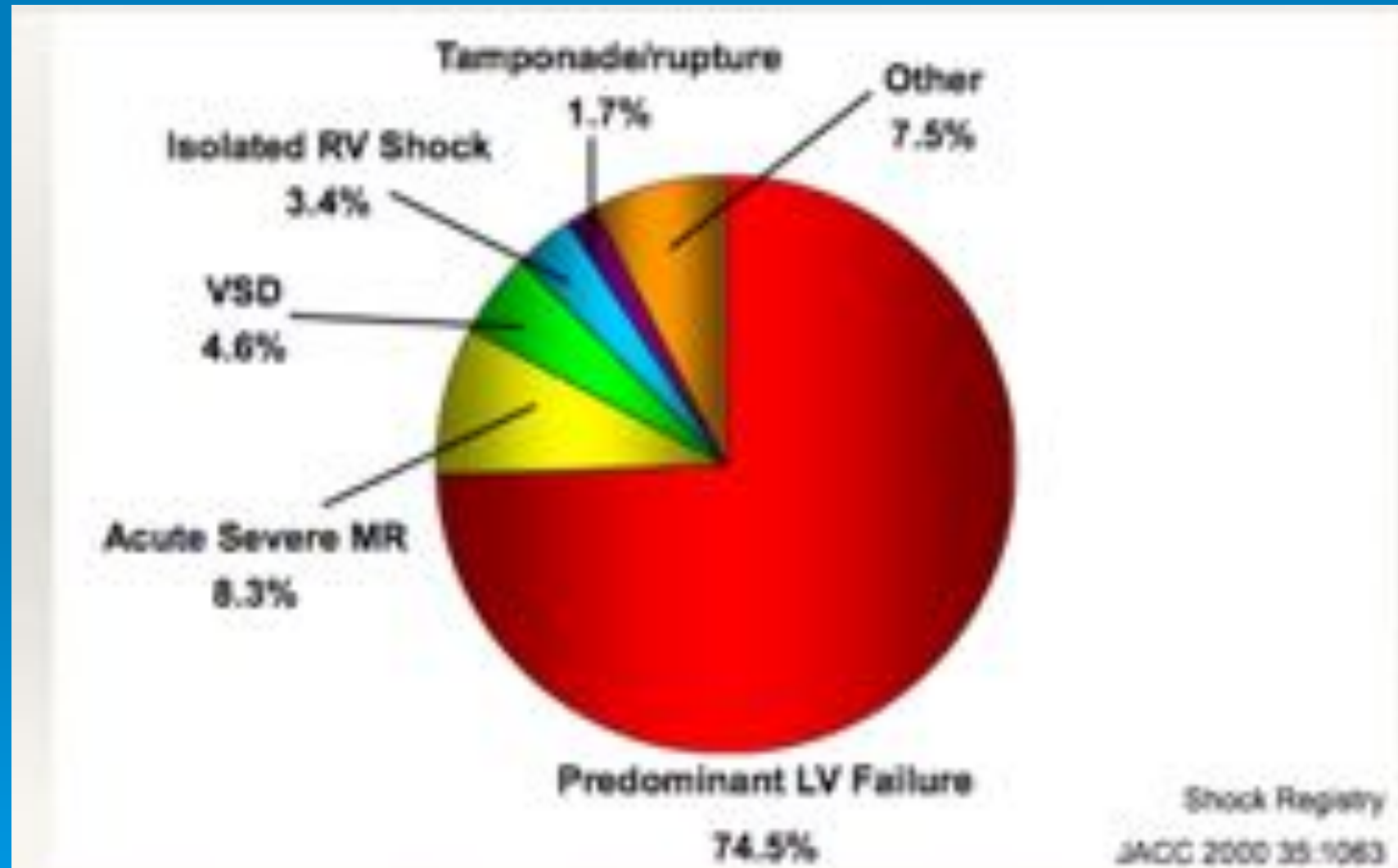
Le choc cardiogénique est une complication grave de l'infarctus dont la mortalité dépasse à l'heure actuelle toujours les 50% malgré les inotropes et les moyens d'assistance circulatoire

Assistance Circulatoire

Choc cardiogénique

- Etiologies multiples pas toujours secondaires à un SCA
- Cardiomyopathies
- Myocardites
- CP ischémique chronique
- Tr du rythme (Orage rythmique)
- CP valvulaires
- Intoxications
- ARCA

Etiologies du choc cardiogénique



CHOC CARADIOGENIQUE

- **Définition** : hypoperfusion tissulaire liée à une défaillance cardiaque après correction de la pré-charge.
 - PAS < 90 mm Hg ou chute de la PAS de 30 mm Hg
 - IC < 2.2 l/min/m²
 - Avec FC > 60 bpm
 - Pcap > 16
 - Diurèse < 0.5ml/kg/min
 - +/- signes de congestion.

TRAITEMENT

- Repos
- Oxygénothérapie / Support ventilatoire.
- Optimisation de la précharge
- Correction des troubles du rythme
- Inotropes :
 - Dobutamine
 - Inhibiteurs des phosphodiesterases
 - Vasopresseurs
 - Adrénaline si réfractaire
- Diurétiques.
- Etiologique : revascularisation...

ECHEC

Choc cardiogénique réfractaire

MORTALITE 75 à 90%

Doll Am Thor Surg 2004

Récupération, traitement étiologique, ou
greffe/assistance chronique envisagée.

Discuter l'ASSISTANCE
CIRCULATOIRE

Indications for percutaneous ventricular support

- High risk PCI
 - High risk anatomy (left main, last vessel...)
 - Severe LV dysfunction (EF < 30%)
- Cardiogenic shock
 - Acute MI
- Bridge to LVAD/heart transplant

History:

1962 Animal studies
Moulapoulos et al. Am Heart J 1962;63:669-679

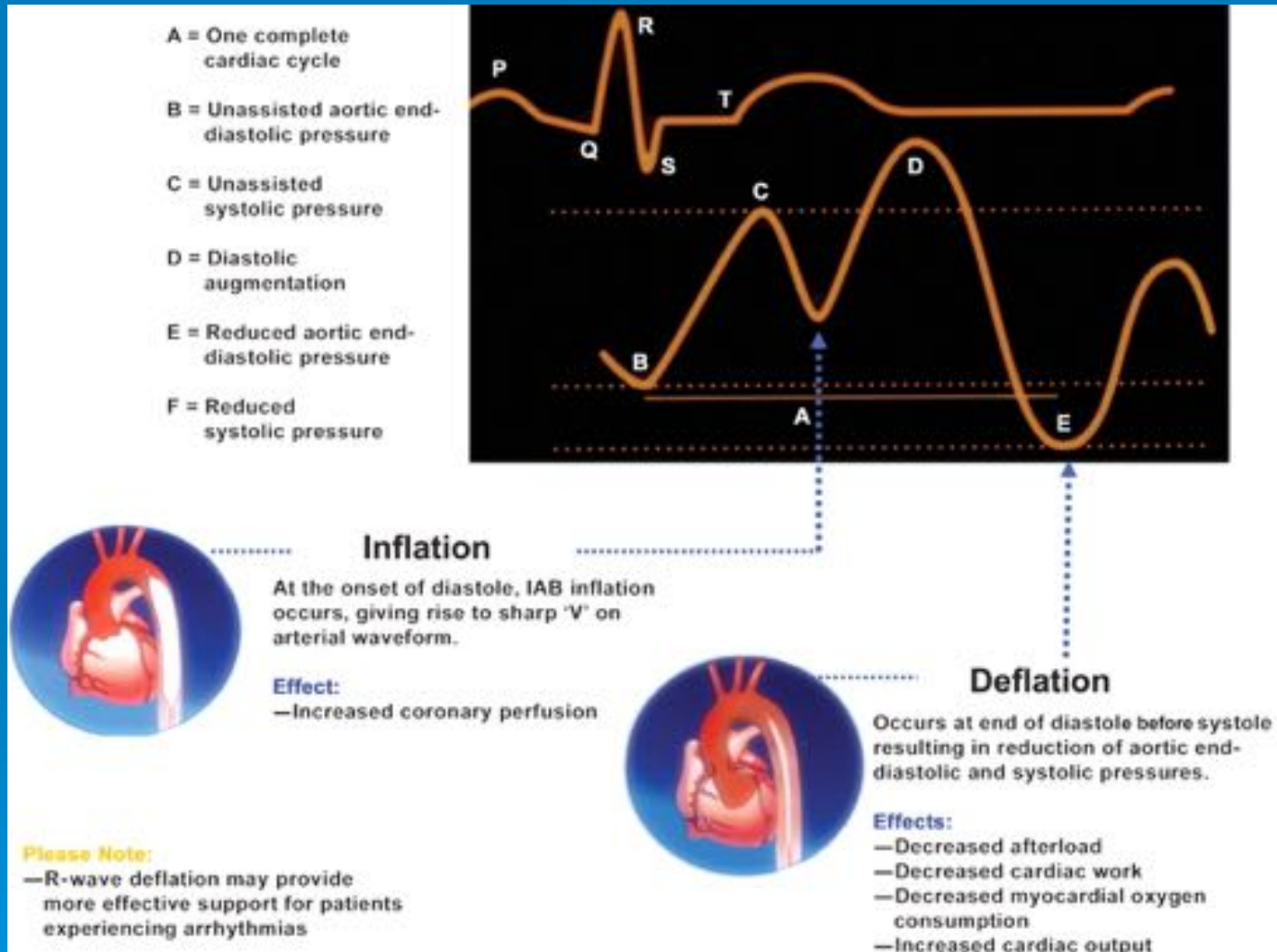
1968 First clinical description in shock
Kantrowitz et al. JAMA 1968;203:138-140

1973 Hemodynamic effects in shock,
Mortality unchanged
Scheidt et al. NEJM 1973;288:979-984

> 40 years > 1 Million patients treated, low complication rate,
Benchmark registry
Ferguson et al. JACC 2001;38:1456-1462



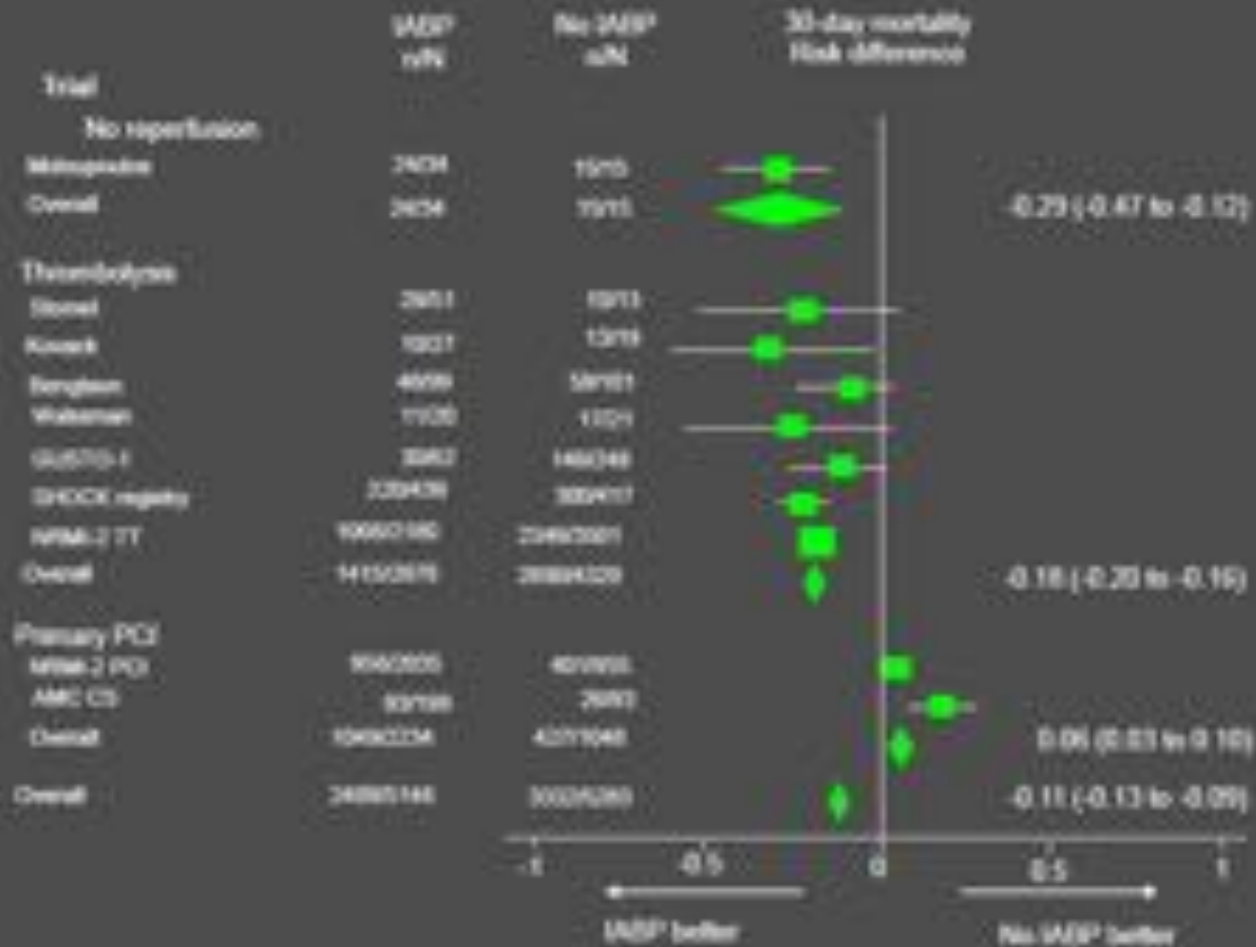
One complete cardiac cycle and the corresponding waveform of the IABP during inflation and deflation.



Krishna M , and Zacharowski K Contin Educ Anaesth Crit Care Pain 2009;9:24-28

Background

Mortality IABP vs no IABP - Metaanalysis



Sjauw et al. Eur Heart J 2009;30:459-468

**Randomized comparison of
intraaortic balloon counterpulsation
versus
optimal medical therapy in addition to early
revascularization in acute myocardial infarction
complicated by cardiogenic shock**

Holger Thiele, MD

Uwe Zeymer, MD; Franz-Josef Neumann, MD; Miroslaw Ferenc,
MD; Hans-Georg Olbrich, MD; Jörg Hausleiter, MD; Gert Richardt, MD;
Marcus Hennersdorf, MD; Klaus Empen, MD; Georg Fuernau, MD; Steffen Desch, MD;
Ingo Eitel, MD; Rainer Hambrecht, MD; Jörg Fuhrmann, MD; Michael Böhm, MD;
Henning Ebel, MD; Steffen Schneider, PhD;
Gerhard Schuler, MD; Karl Werdan, MD

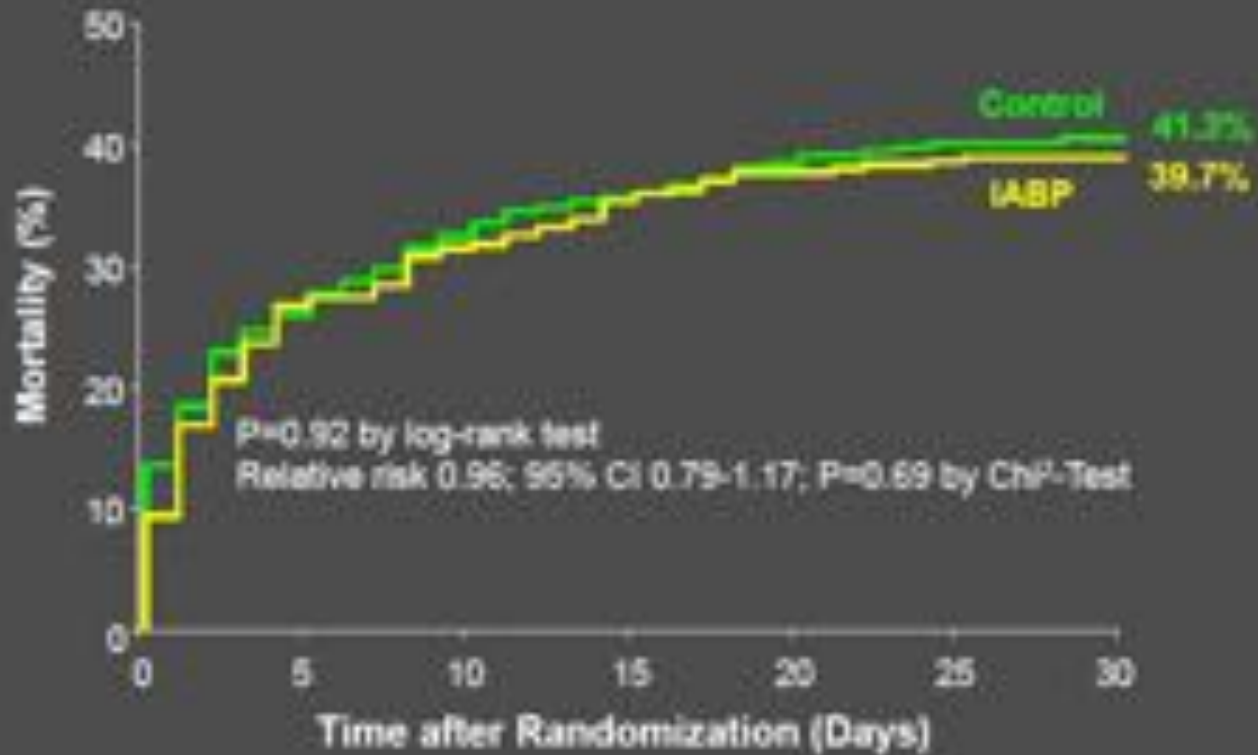
on behalf of the **IABP-SHOCK II Trial** Investigators

University of Leipzig – Heart Center

Results

Primary Study Endpoint (30-Day Mortality)

IABP
SHOCK II



ECMO

L'ECMO est flexible veino veineuse ou veino arterielle.

Elle donne 4,5 -5 l/min Le patient doit être bien rempli pour « nourrir » le Cathéter veineux.

Le cathéter veineux est placé dans la VCI près de l'oreillette Le sang veineux est oxygéné et reconduit vers l'Aorte descendante en court-circuitant le cœur et les poumons ce qui entraîne une élévation de la post-charge au niveau du VG.

L'augmentation de la post-charge au niveau du VG a pour effet d'aggraver la défaillance VG en aggravant le risque d'OAP et d'augmenter la consommation en O₂ très défavorable pour ces cœurs déjà ischémiques.

Parfois on peut traiter la surcharge du VG en ponctionnant le septum interauriculaire pour décharger le cœur gauche et la circulation

Extra - Corporeal Membrane Oxygenation

- ⊗ Pompe centrifuge. Flux actif
- ⊗ 4 - 4,5L /min
- ⊗ Oxygénation associée
- ⊗ Double abord vasculaire avec veine 18-31F et art 15-22F
- ⊗ 15 min insertion

- ⊗ Risques: ↑ PTDVG
thrombose VG



Cardiohelp



Problems with ECMO

- Small series of patients from high volume LVAD centers (A. Combes, Paris)
- No LV unloading
- Increased LV afterload
- More personnel (cardiac surgeon, perfusionist) and more difficult to organize rapidly in acute setting
- More pronounced systemic inflammatory response

Amines



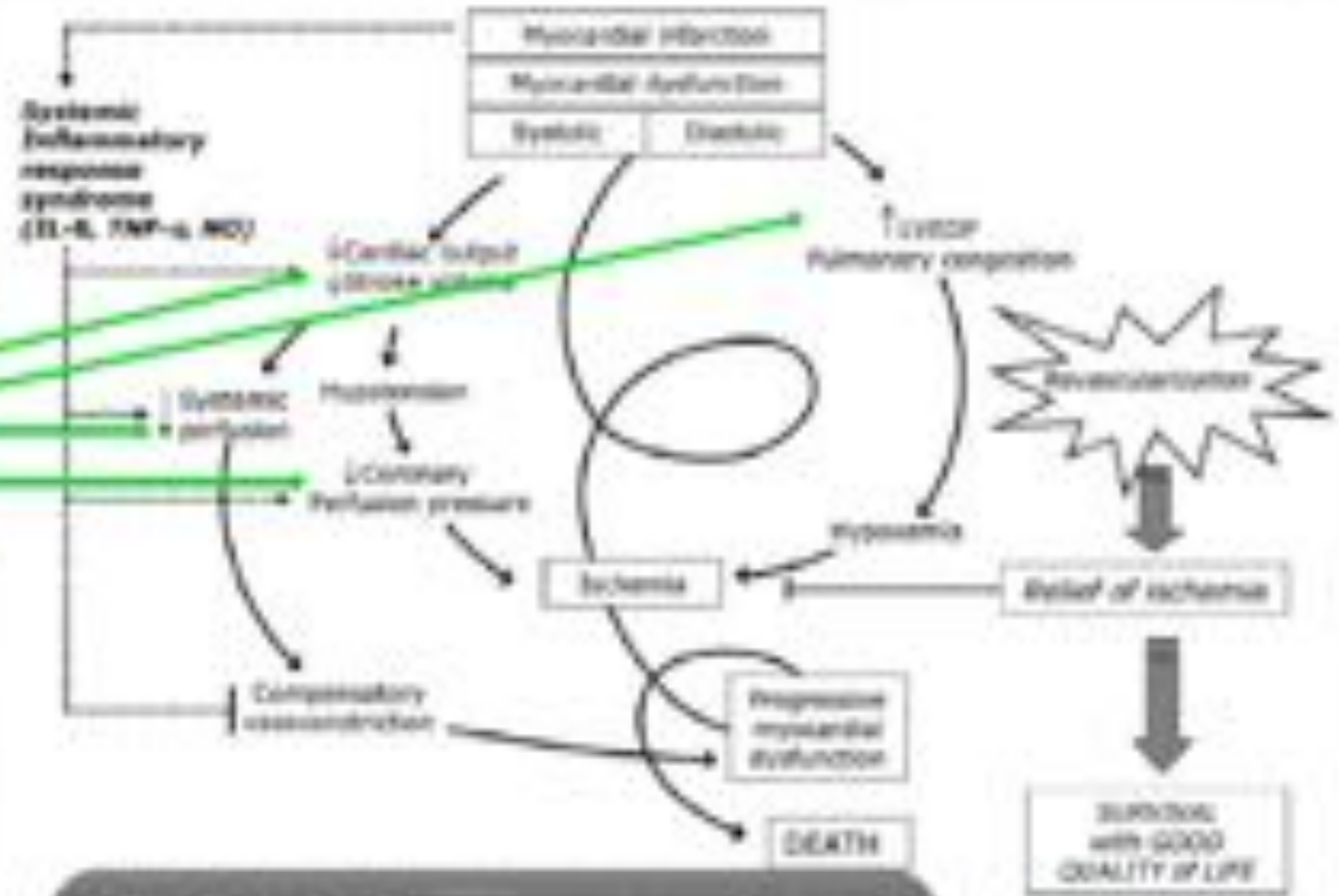
Assistance cardiaque
Antérograde

Assistance circu
rétrograde



Assistance Anterograde

Assistance
Cardiaque
Antérograde



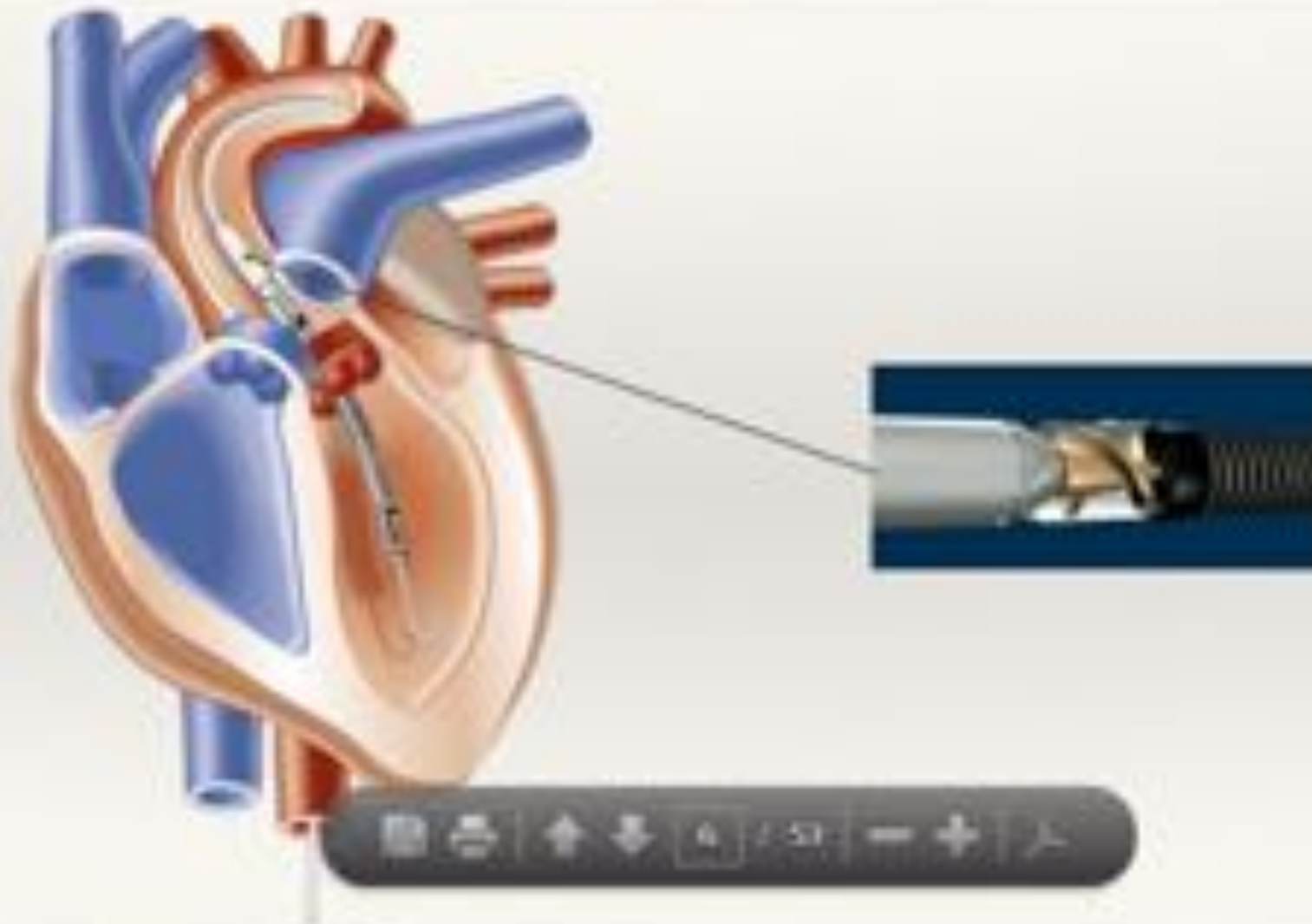
Impella: LV-aortic axial flow pump



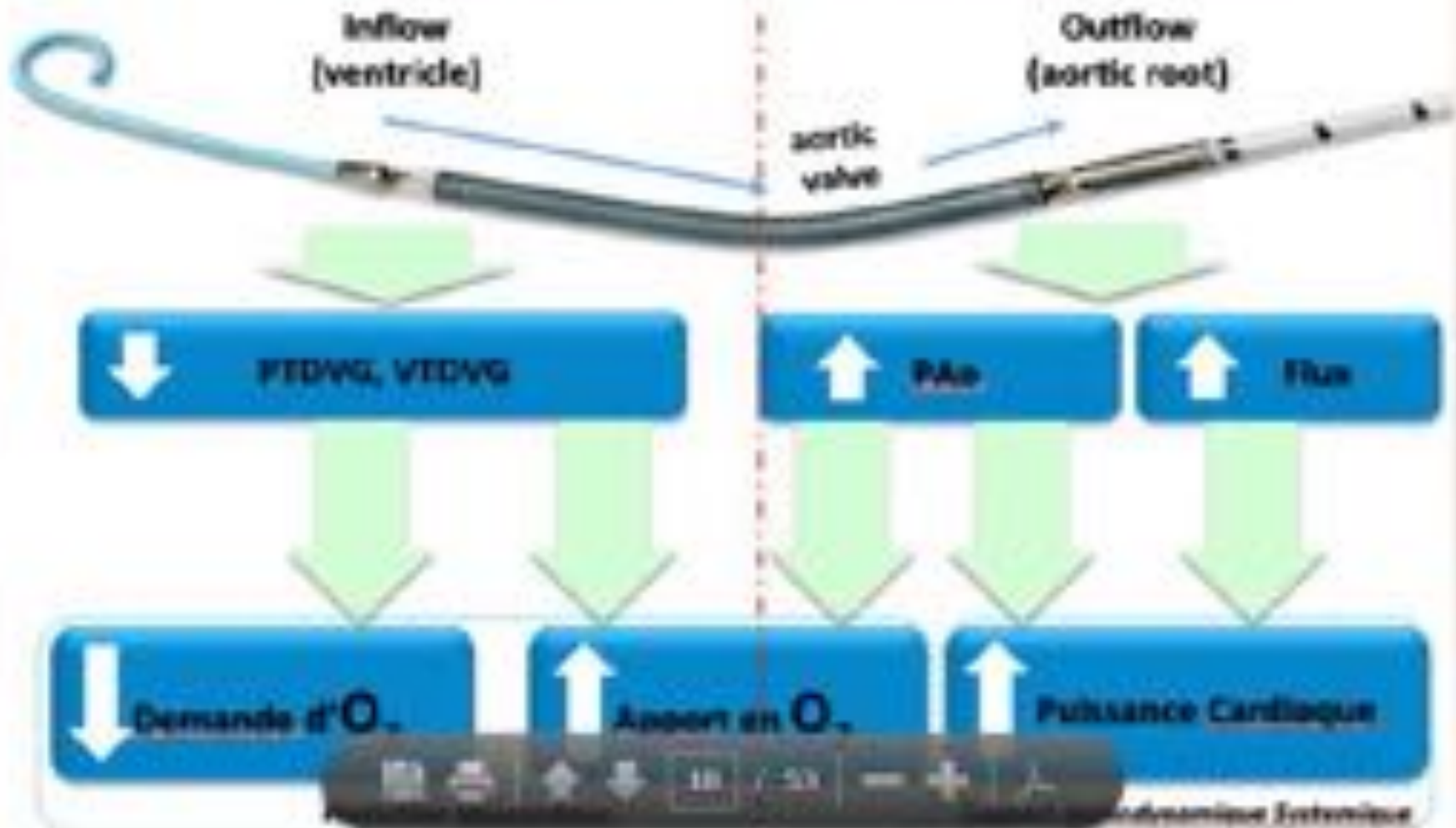
- Miniaturized 12-13F pump
- Single Femoral access
- Placement with 0.018" wire
- Actively unloads the LV
- Forward Flow up to 4.0L/min
- Low anticoagulation

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IMPELLA



Impella Principe d'Action



Impella Percutaneous Circulatory Support System

Key Design Differences

	Impella® 2.5	Impella CP™
Flow Rate (L/min, max)	2.5	Up to 4.0
Catheter Size	9 Fr	9 Fr
Pump Size	12 Fr	14 Fr
Insertion Method	Percutaneous via 13 Fr Introducer Sheath	Percutaneous via 14 Fr Introducer Sheath
Guidewire	0.018" Silicone Wire	0.018" PTFE Wire
Placement Measurement	Fluid-filled Pressure Lumen	Fluid-filled Pressure Lumen
Cannula Geometry	Curved, Pigtail	Curved, Pigtail



Impella Physiopathologie

Etudes Impella	Augmente Le Débit Cardiaque	Apports en O2 Augmentés (Flux coronaire)	Demande en O2 diminuée (Décharge VG)
Burzotta et al., J Cardiovasc Med. 2008	✓		
Saunen et al., Artif Organs. 2007		✓	✓
Valgimigli et al., Cath Cardiovasc Interv. 2005	✓		✓
Meyns et al., Thorac Cardio Surg. 2003	✓		✓
Meyns et al., J Am Coll Cardio. 2003			✓
Remmelink et al., Cath Cardio Interv. 2007		✓	
Reesink et al., CHEST 2004	✓		✓
Dixon et al., J Am Coll Card Interv. 2009	✓		
Seyfarth et al., J Am Coll Card			

Danish Cardiogenic Shock Trial (DanShock)

- In 2012 a more powerful Impella has been introduced that is able to deliver 3.5l/min (approximately 75% of a normal cardiac output).
- The hypothesis of the current study is to reduce mortality and morbidity of patients with cardiogenic shock using the Impella cVAD.
- The study will be carried out as a randomized multicenter study where eligible patients will be randomized to receive conventional circulatory support or support with the Impella device and inotropic support if needed.
- A total of 360 patients are planned to be enrolled, and the primary endpoint will be death.

Danish (DanShock) Trial

Acute MI (STEMI < 36 h)

Shock

Primary Endpoint:
Total Mortality

Inclusion criteria

- Lactate ≥ 2.5 mmol/l
- Hemodynamics
SBP < 100 mmhg or
vasopressors
- LV-EF < 35%

PCI (CABG)

Randomization (N=360)

IABP pre PCI (n=180)

Impella cVAD pre PCI (n=180)

The Current Use of Impella 2.5 in Acute Myocardial Infarction Complicated by Cardiogenic Shock: Results from the USpella Registry

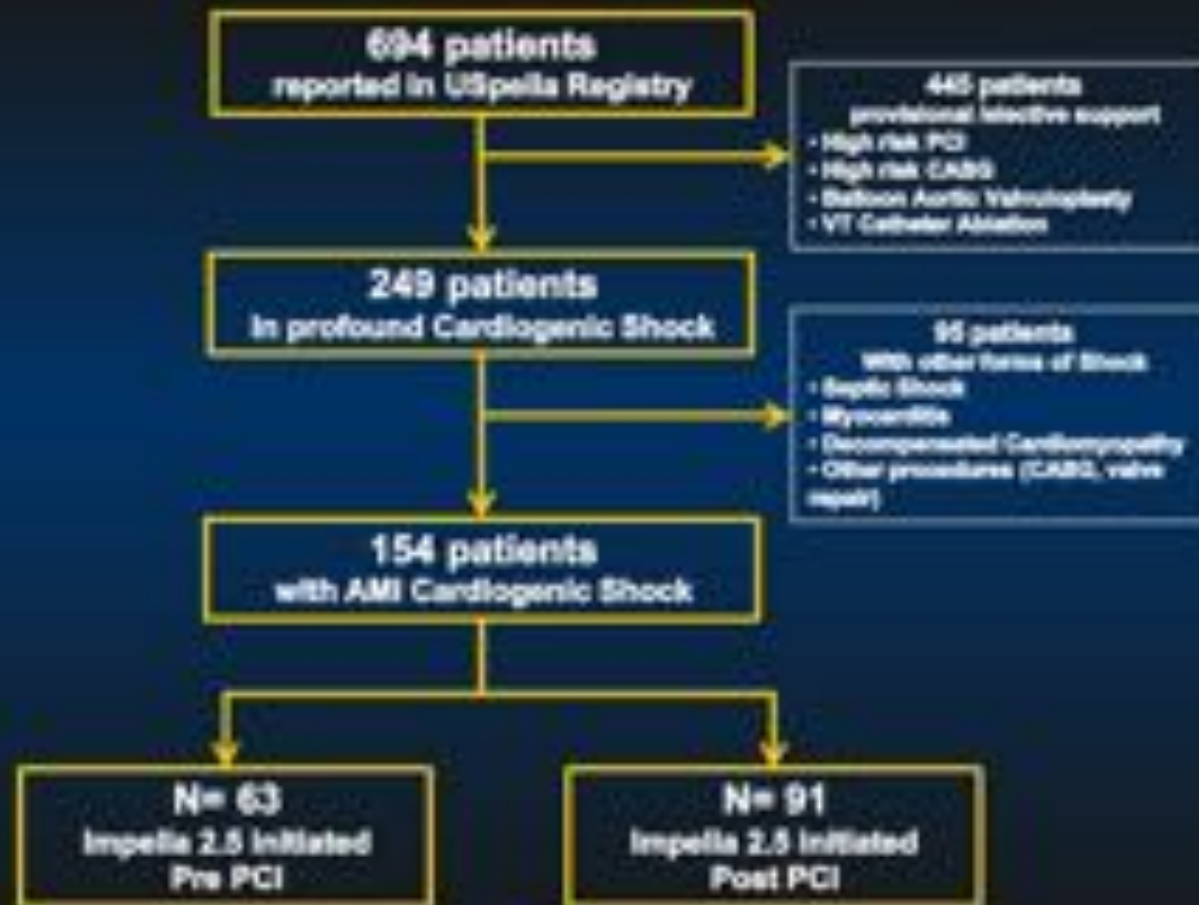
William W. O'Neill, MD¹; Theodore Schreiber, MD²; David H. W. Wahna, MD³; Charanjit Rihal, MD⁴;
Brijhari S. Naidu, MD⁵; Andrew B. Civitello, MD⁶; Simon R. Dixon, MChB⁷; Joseph M. Massaro,
PhD⁸; Brijeshwar Maini, MD⁹; E. Magnus Ohman, MD¹⁰

From the ¹Henry Ford Hospital, Detroit, MI, USA; ²Detroit Medical Center, Detroit, MI, USA; ³Spectrum Health, Grand Rapids, MI, USA; ⁴Mayo Clinic, Rochester, MN, USA; ⁵Winthrop University Hospital, Mineola, NY, USA; ⁶Texas Heart Institute, Houston, TX, USA;

⁷Beaumont Hospital, Royal Oak, MI, USA; ⁸Harvard Research Institute, Boston, MA, USA;
⁹Pinnacle Health Medical Center, Womersleyburg, PA, USA; ¹⁰Duke University Medical Center, Durham, NC, USA.

Study Flow Chart

(06/08-05/12)



O'Neill et al, J Intervent Cardiol 2013;9999:1-11

USpella Cardiogenic Shock

Hemodynamics All Patients

N = 154 (mean \pm SD or %)

	Pre-support	On Support	P-Value
SBP, mmHg	85.4 \pm 25.6 (143)	126.7 \pm 31.4 (144)	<0.0001
DBP, mmHG	50.8 \pm 18.6 (143)	78.7 \pm 21.1 (143)	<0.0001
MAP, mmHg	62.7 \pm 19.2 (143)	94.4 \pm 23.1 (143)	<0.0001
PCWP, mmHg	31.9 \pm 11.0 (25)	19.2 \pm 9.7 (25)	<0.0001
Cardiac Output, L/min	3.4 \pm 1.3 (23)	5.3 \pm 1.7 (23)	<0.0001
Cardiac index, L/min/m ²	1.9 \pm 0.7 (23)	2.7 \pm 0.7 (23)	<0.0001
Cardiac power output, W	0.48 \pm 0.17 (23)	1.06 \pm 0.48 (23)	<0.0001

PROTECT II



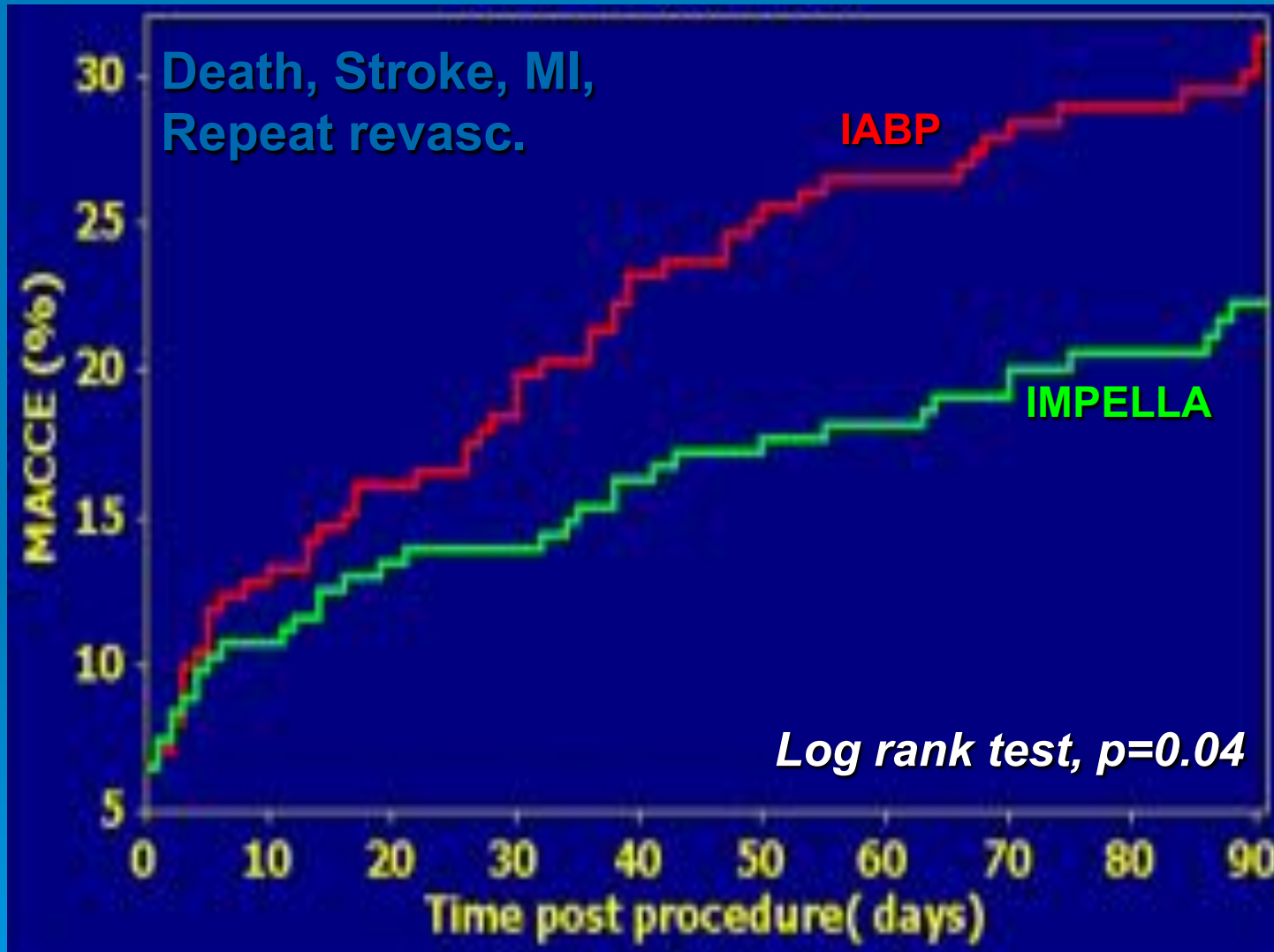
*Prospective Multicenter Randomized Trial
Comparing IMPELLA to IABP in High Risk PCI:
90 Day Results*

William O'Neill, Neal Kleiman, Jose
Henriques, Simon Dixon, Joseph Massaro,
~~Ioana Ghiu, Brijeshwar Maini, Suresh~~
Mulukutla, Vladimir Dzavik, James
Revenaugh, Hadley Wilson, Karim Benali,
Magnus Ohman

On behalf of all PROTECT II Investigators

PROTECT II MACCE**

Per Protocol Population, N=426



**Using $\times 8$ ULN threshold for biomarkers or Q-wave for Peri-procedural MI (Stone et al Circulation 2001;104:642-647) and $2\times$ ULN threshold for biomarkers for Spontaneous MI (Universal MI definition)

Impella or ECMO in Refractory Cardiogenic Shock?

- Absence of randomized data compared to routine care for both devices
- No randomized comparison between ECMO and Impella
- Advantages of Impella:
 - 2.5 and CP: percutaneous insertion, smaller sheaths
 - Left ventricular unloading

American Guidelines for Impella

- 2011 ACCF/AHA/SCAI Guideline for Percutaneous Coronary Intervention. *JACC* 2011
 - High risk patients: Class IIb
 - PCI and Cardiogenic Shock: Class I
- 2013 ACCF/AHA Guideline for the Management of ST-Elevation Myocardial Infarction. *Circulation* 2012
 - STEMI and Cardiogenic Shock: Class IIb
 - STEMI and urgent CABG: Class IIa
- Use of Mechanical Circulatory Support: American Heart Association. *Circulation* 2012
 - Acutely decompensated heart failure patients: Class IIa
- 2013 International Society for Heart and Lung Transplantation Guidelines for Mechanical Circulatory Support. *The Journal of Heart and Lung Transplantation*, 2013
 - Temporary mechanical support for patients with multi-organ failure: Class I
- 2013 ACCF/AHA Guideline for the Management of Heart Failure, *Journal of American College of Cardiology* 2013
 - "Bridge to Recovery" or "Bridge to Decision" for patients with acute, profound hemodynamic compromise: Class IIa

Conclusion: INCCI algorithm for shock?

