



45th Annual International Congress of the
EGYPTIAN SOCIETY OF CARDIOLOGY
CardioEgypt 2018

**STEMI with Cardiogenic shock and
multivessel coronary artery disease**

ELSayed Farag MD ,FSCAI
Zagazig University

Clinical Data

- Female 65 yrs old diabetic
- Chest pain 24 hours before being transferred
- AWMi ...SK
- Bed side echoDoppler .. Severe HK in all segments with EF 39 %
- Shocked

Multivessel Disease in AMI

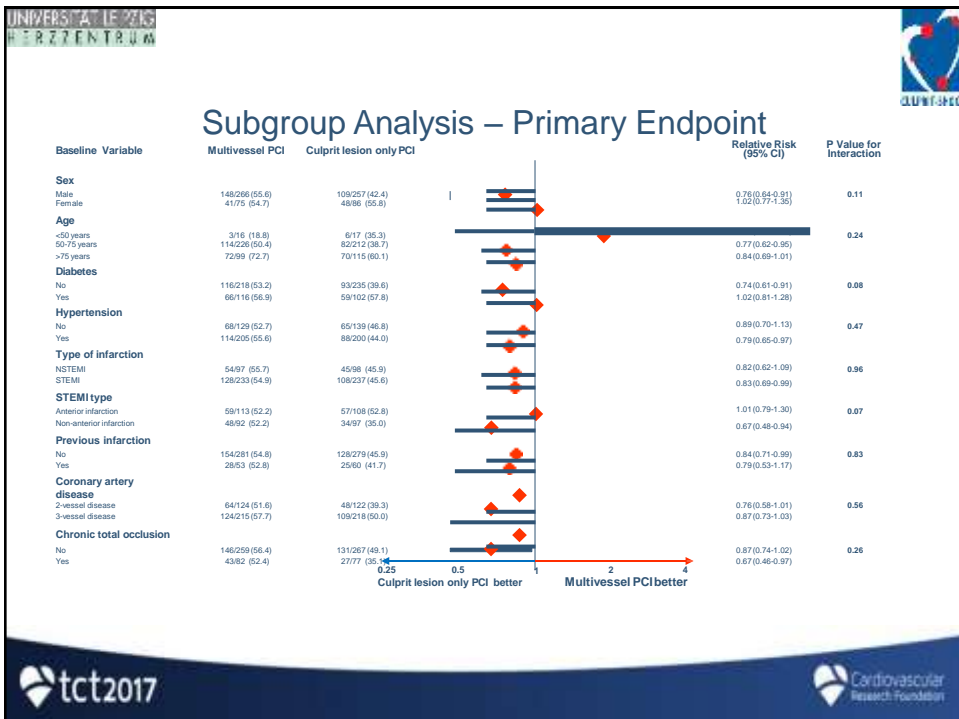
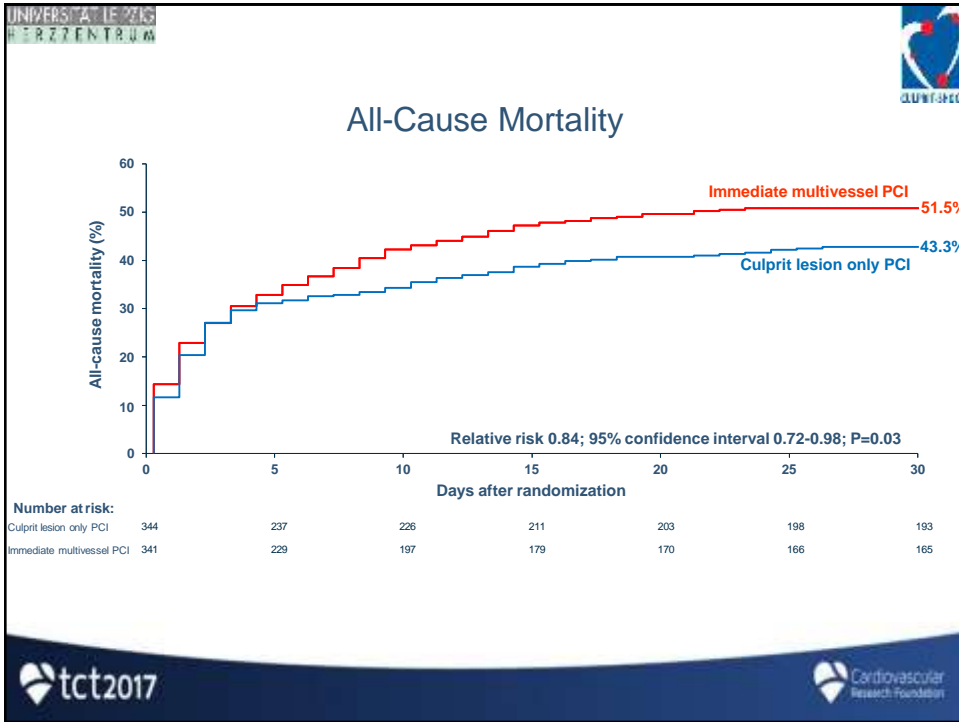
- ◆ **Multivessel disease occurs in 40-60% of patients with STEMI, and 70-80% of patients with shock**
- ◆ **It confers higher risk of death, reinfarction, stent thrombosis, lack of compensatory hyperkinesis of the non-infarct zone and development of shock**
- ◆ **Multiple culprits may be present due to a systemic inflammatory state**
- ◆ **Therefore, treatment of non-culprit vessels may be beneficial**

UNIVERSITÄT LEIPZIG
HERZ ZENTRUM

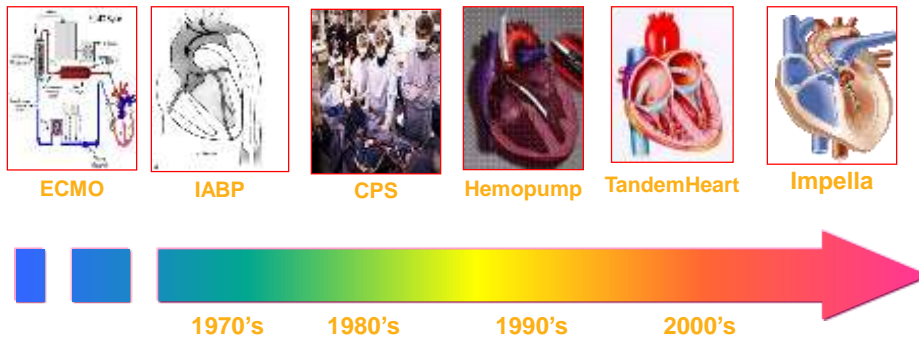


CULPRIT-SHOCK: A Randomized Trial of Multivessel PCI in Cardiogenic Shock

Holger Thiele, MD
on behalf of the CULPRIT-SHOCK Investigators



Historical Perspectives of Cardiac Support Devices



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

Study Sites and Organisation



DSMB:

Steering committee:

Support +
Patronage:



Summary + Conclusions

- IABP support in cardiogenic shock is safe without significant inherent complications.
- However, IABP support did not reduce 30-day mortality in this large, randomized, multicenter trial in cardiogenic shock patients complicating myocardial infarction undergoing early revascularization.
- The primary study endpoint results are supported by a lack of benefit in secondary endpoints.

Guidelines

ESC



Class IC

ACC/AHA



Class IB

ESC STEMI Guidelines 2017

CHANGE IN RECOMMENDATIONS 2012		2017 NEW RECOMMENDATIONS
Radial access* BASKET™		<ul style="list-style-type: none"> Additional lipid lowering therapy (LDL >1.8 mmol/L or 70 mg/dL) with statin PROVE-IT COURAGE™ Complete revascularization during index primary PCI in STEMI patients in which lesion severe Congestive HF: 1 medication have not been given (QUINQUE) Switch to potent P2Y₁₂ inhibitors 48 hours after fibrinolysis (don't repeat) Excess fibrinogen up to 30 months in high-risk patients (SCASTIN-UP) Use of polytail to increase adherence (SCOR) Reduce use of deferred imaging (CLASSIC)
DES over BMS COMPELL™, COMPELL™, COMPELL™, COMPELL™		
Complete Revascularization* REACT™, CANARY™, REBEL™, CRYSTAL™, Coronary-ACAD™		
Thrombus Aspiration* ASAP™, ASAP™		
Bivalirudin MATERION™, RECOVER™		
Enoxaparin VENEL™, New-onset™		
Early Hospital Discharge* Small study & observational study		
<p>Class I: Class IIa: Class IIb: Class III:</p> <p>Dose (X) Thrombolysis same in all patients</p> <p>STRAT™</p> <p>Dose (X) Thrombolysis half in Pt >75 years</p>		
2017 NEW / REVISED CONCEPTS		
MINOCA AND QUALITY INDICATORS: <ul style="list-style-type: none"> New chapters dedicated to these topics. 	TIME LIMITS FOR ROUTINE OPENING OF AN IBA: <ul style="list-style-type: none"> 0-2h (Class I); 1.5-4h (Class IIa); >4h (Class III) 	
STRATEGY SELECTION AND TIME DELAYS: <ul style="list-style-type: none"> Clear definition of first medical contact (FMC) Definition of "time 0" to chosen reperfusion strategy (i.e. the strategy clock starts at the time of STEMI diagnosis) Selection of PCI over fibrinolysis when procedural delay from "STEMI diagnosis" to wire crossing is <30 min. Maximum delay since from "STEMI diagnosis" to initiation of fibrinolysis agent is set to 30 min. "Door-to-Balloon" term eliminated from guidelines. 	ELECTROCARDIOGRAM AT PRESENTATION: <ul style="list-style-type: none"> Left and right bundle branch block considered equal for recommending urgent angiography if ischemic symptoms. 	
	TIME TO ANGIOGRAPHY AFTER FIBRINOLYSIS: <ul style="list-style-type: none"> Timeliness is set to >2.5h after successful fibrinolysis. 	
	PATIENTS TAKING ANTICOAGULANTS: <ul style="list-style-type: none"> Acute and chronic management presented. 	

Multivessel Percutaneous Coronary Intervention in Patients With ST-Segment Elevation Myocardial Infarction With Cardiogenic Shock

ABSTRACT

BACKGROUND Recent trials demonstrated a benefit of multivessel percutaneous coronary intervention (PCI) for noninfarct-related artery (non-IRA) stenosis over IRA-only PCI in patients with ST-segment elevation myocardial infarction (STEMI) multivessel disease. However, evidence is limited in patients with cardiogenic shock.

OBJECTIVES This study investigated the prognostic impact of multivessel PCI in patients with STEMI multivessel disease presenting with cardiogenic shock, using the rationale, multicenter, prospective KAMIR-NHI (Korea Acute Myocardial Infarction-National Institutes of Health) registry.

METHODS Among 13,104 consecutive patients enrolled in the KAMIR-NHI registry, we selected patients with STEMI with multivessel disease presenting with cardiogenic shock and who underwent primary PCI. Primary outcome was 1-year all-cause death, and secondary outcomes included patient-oriented composite outcome (a composite of all-cause death, any myocardial infarction, and any repeat revascularization) and its individual components.

RESULTS A total of 659 patients were treated by multivessel PCI (n = 260) or IRA-only PCI (n = 399) strategy. The risk of all-cause death and non-IRA repeat revascularization was significantly lower in the multivessel PCI group than in the IRA-only PCI group (21.3% vs. 31.7%; hazard ratio: 0.59; 95% confidence intervals: 0.49 to 0.82; $p = 0.001$; and 6.7% vs. 8.2%; hazard ratio: 0.39; 95% confidence intervals: 0.17 to 0.90; $p = 0.028$, respectively). Results were consistent after multivariable regression, propensity-score matching, and inverse probability weighting to adjust for baseline differences. In a multivariable model, multivessel PCI was independently associated with reduced risk of 1-year all-cause death and patient-oriented composite outcome.

CONCLUSIONS Of patients with STEMI and multivessel disease with cardiogenic shock, multivessel PCI was associated with a significantly lower risk of all-cause death and non-IRA repeat revascularization. Our data suggest that multivessel PCI for complete revascularization is a reasonable strategy to improve outcomes in patients with STEMI with cardiogenic shock. *J Am Coll Cardiol* 2018;71:844-56 | © 2018 by the American College of Cardiology Foundation.

Lee, J.M. et al. *J Am Coll Cardiol*. 2018;71(8):844-56.

STEMI with Cardiogenic Shock

Adjusted HR (95% CI) for All-Cause Death

Strategy	Adjusted HR (95% CI)
Unadjusted	0.58 (0.49-0.82)
HR-Adjusted	0.47 (0.41-0.70)
PS-Matched	0.59 (0.41-0.84)
IPW-Adjusted	0.60 (0.44-0.82)
IPW-Matching	0.64 (0.39-0.90)

Events: Multivessel PCI (n=260) vs. IRA-Only PCI (n=399) as a Reference

45th Annual International Congress of the
EGYPTIAN SOCIETY OF CARDIOLOGY
CardioEgypt 2018

Take Home Message

- Support and support ...
- Ventilate if mandate ...
- Cardiac arrest setup
- Minimal amount of dye...
- Be simple...
- Try to be complete ... to complete you job !