Intra-aortic balloon counterpulsation and infarct size in patients with acute anterior myocardial infarction without shock: The CRISP AMI Randomized Trial



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# Background



- Despite improvements in STEMI care
  - The 6 month mortality remains high ~10%¹
- Intra-aortic balloon counterpulsation
  - — 
     †Diastolic arterial pressure (coronary perfusion pressure)
  - – ↓Simultaneously decrease afterload and left ventricular end diastolic pressure (LVEDP) both work to decrease oxygen consumption
  - Decreases infarct expansion when placed prior to reperfusion in animal studies <sup>2,3</sup>

<sup>&</sup>lt;sup>1</sup>Heart disease and stroke statistics--2009 update. Circulation 2009;119:e21-181.

<sup>&</sup>lt;sup>2</sup>LeDoux JF et. al.. Catheterization & Cardiovascular Interventions 2008;72:513-21.

<sup>&</sup>lt;sup>3</sup>Azevedo CF et. al. European Heart Journal 2005;26:1235-41.

# **Primary Objective**



To determine whether routine initiation of intraaortic balloon counterpulsation (IABC) before mechanical reperfusion compared to standard of care (SOC) primary PCI decreases infarct size in patients with anterior ST-segment elevation myocardial infarction (STEMI) without cardiogenic shock

# Study Design



# Anterior STEMI without Shock

### **Inclusion Criteria**

- Anterior STEMI
  - 2 mm in 2 contiguous leads or at least 4 mm in the anterior leads
- •Planned Primary PCI within 6 hrs
- Adult able to consent

Intra-aortic Balloon Counterpulsation prior to PCI

Randomize Open Label (n ~ 300)

Standard of Care Primary PCI

At least 12 hours of IABC post PCI

Routine Post PCI care

Cardiac MRI performed day 3-5 post PCI

Primary Endpoint: Infarct Size on CMR

- 1. All Patients with CMR data
- 2. Patients with Prox LAD occlusion TIMI 0/1 flow

Clinical Events – 6 months

### **Exclusion Criteria**



- Known Contraindication to MRI
- Prior Thrombolytic Therapy for STEMI
- Cardiogenic Shock
- Prior MI, CABG, or ESRD
- Contraindications to IABC
  - Known Severe AI, AAA, or severe peripheral artery disease
  - ->400 lbs of < 4 feet

# Statistical Methodology



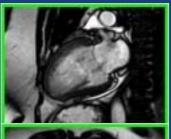
- Sample Size
  - Estimated Infarct size
    - All patients (25.3 -26.6% LV)<sup>1,2</sup> and (19.9 28.8% LV)<sup>1,2</sup> prox. LAD TIMI 0/1
  - 25% reduction (270 patients) 10% CMR data missing
  - >80% power, Type 1 error 0.025 (2-sided)
  - − ~ 300 patients
- Primary Endpoint Evaluation: Infarct Size on CMR
  - Modified ITT all patients with CMR data
  - All CMR patients with proximal LAD occlusion TIMI 0/1
- Primary Safety Evaluation: Major vascular complications and Major bleeding
- Clinical Outcomes: 6-month rate all cause mortality, MACE

<sup>&</sup>lt;sup>1</sup> Patel et al. Jacc: Cardiovascular Imaging 2010;3:52-60

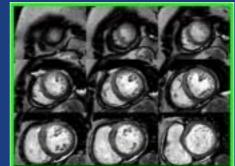
<sup>&</sup>lt;sup>2</sup> Thiele et al. Circulation 2008 Jul 1;118(1):49-57 Epub 2008 Jun 16

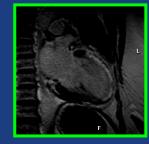
### **CMR Protocol**

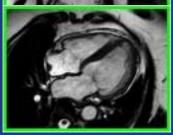


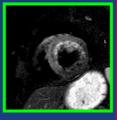


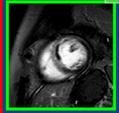
Contrastinjection 0.15 mmol/kg/KG Bolus Gadovist i.v.



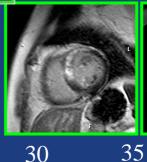


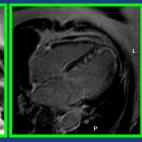






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Survey

0

Function
4-chamber
2-chamber

5

**Edema** 3 short axes

10

Early enhancement Short axes Apex-Base Function
Short axes
Apex-Base

20

Delayed enhancement Short axes Apex-Base Delayed enhancement 4-chamber 2-chamber

40

SSFP sequence (TR/TE/flip = 3.2ms/1.2ms/60°)

T2 STIR sequence (TR/TE/flip = 2 heart beats/80ms/90°) slice thickness: 8-10 mm

Inversion recovery gradient echo sequence (TR/TE/flip 2.8ms/1.1ms/15°) slice thickness: 8-10 mm, no gap SSFP sequence (TR/TE/flip = 3.2ms/1.2ms/60°) slice thickness: 8-10 mm, no gap

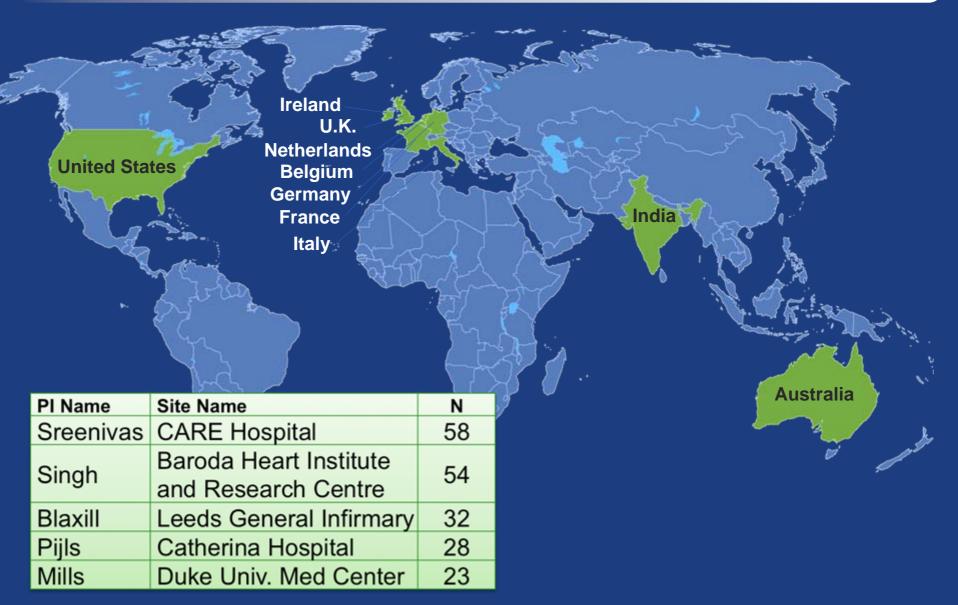
25

Inversion recovery gradient echo sequence (TR/TE/flip 2.8ms/1.1ms/15°) slice thickness: 8-10 mm, no gap Time (min)

### **Enrollment**

9 countries, 30 sites, 337 patients





# Study Conduct



Randomized\* N=337

		IABC N=161	SOC N=176		
Received intervention		153 (95.03%)	161 (91.48%)		
Withdrew		4	2		
Lost to follow-up		Crossing over to IABC			
MRI not performed	Sustained hypotension/Cardiogenic shock To prevent event post vessel dissection Failed PCI of IR vessel Continued chest pain				
Died					
Unstable					
Metallic contraindica	cation 3 1		1		
Unable to tolerate	11 18		18		
Other		6	0		
MRI performed, not ev	aluable	5	7		

# Baseline Demographics



	AII (N=337)	IABC (N=161)	SOC (N=176)
Age, median (25th, 75th), yrs	56.6 (48.4, 65.6)	56.1 (48.3, 64.3)	57.7 (48.6, 66.4)
Male, %	81.9	82.0	81.8
Race, %			
White	47.8	50.3	45.5
Asian	45.1	46.6	43.8
Black or African American	4.7	1.9	7.4
Other	2.1	1.2	2.8
Medical history, %			
Hypertension on drug tx.	29.4	24.2	34.1
Current nicotine use	31.8	33.1	30.7
Dyslipidemia on drug tx.	12.5	12.5	12.5
Diabetes mellitus	18.7	16.8	20.5

# Baseline Demographics (cont.)



	AII (N=337)	IABC (N=161)	SOC (N=176)
SBP, median (25th, 75th), mm Hg	131.0 (118.0, 150.0)	130.0 (113.0, 150.0)	135.0 (120.0, 151.0)
DBP, median (25th, 75th), mm Hg	80.0 (70.0, 92.0)	80.0 (70.0, 92.0)	80.0 (71.5, 92.0)
HR, median (25th, 75th), bpm	81.0 (71.0, 94.0)	81.0 (71.0, 93.0)	80.0 (70.0, 94.0)
ST ↑ in anterior leads, no. (%)			
0–<2 mm	0 (0.0)	0 (0.0)	0 (0.0)
2–<4 mm	1 (0.3)	0 (0.0)	1 (0.6)
4–<6 mm	135 (40.1)	61 (37.9)	74 (42.0)
≥6 mm	201 (59.6)	100 (62.1)	101 (57.4)

# PCI Procedure

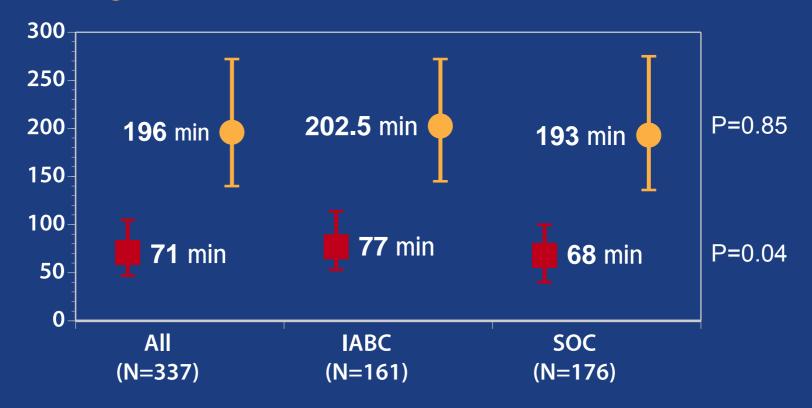


	All N=337	IABC N=161	SOC N=176
PCI			
PCI performed, %	94.3	96.3	92.6
Infarct-related artery			
Left anterior descending, %	97.6	99.4	96.0
Infarct-related artery stenosis location			
Proximal, %	62.9	64.8	61.2
Infarct-related artery TIMI flow pre-intervention	n		
Grade 0, %	65.3	66.0	64.7
Grade 1, %	10.3	11.3	9.4
Infarct-related artery final TIMI flow post-inter			
Grade 3, %	94.2	92.9	95.3

### Time to Treatment



- First medical contact to first device\*
- ——— Symptom onset to 1st device



# Primary outcome



	AII (N=337)	IABC (N=161)	SOC (N=176)	P Value
Primary endpoint				
Infarct size (% LV), modified ITT all patients with CMR data				
N	275	133	142	
Mean	39.8	42.1	37.5	
Median	38.8	42.8	36.2	
Infarct size (% LV), modified ITT patients prox. LAD and TIMI flow 0/1				
N	192	93	99	
Mean	44.4	46.7	42.3	
Median	42.1	45.1	38.6	

Co-primary endpoint: 2-sided p=0.025

# 30-day Clinical Events

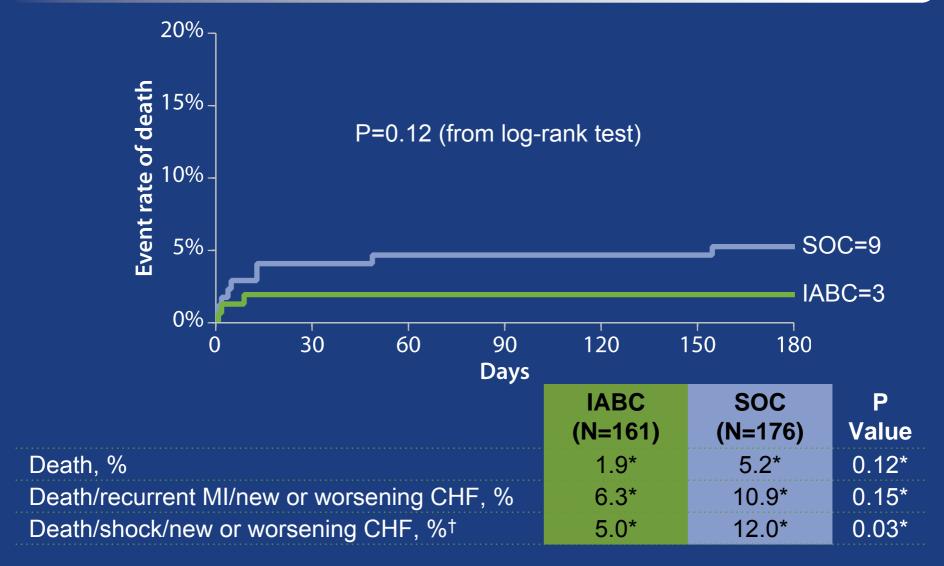


	IABC (N=161)	SOC (N=176)	P Value
Death, %	1.9*	4.0*	0.26*
Stroke, %	1.9	0.6	0.35
Major bleed per GUSTO 1 definition or transfusion, %	3.1	1.7	0.49
Vascular complications, (n) %	7(4.3)	2 (1.1)	0.09
Major limb ischemia requiring operative intervention (n)	0	0	
Distal embolization (n)	0	0	
Major dissection (n)	2	0	
Pseudoaneurysm or AV fistula (n)	3	2	
Hematoma >5 cm (n)	3	0	

<sup>\*</sup>From KM curves and log-rank test.

### All Cause Death – 6 months





<sup>\*</sup>From KM curves and log-rank test. †Exploratory analysis.

### Conclusion



Among Patients with Acute Anterior STEMI without cardiogenic shock use of Intra-aortic counterpulsation prior to PCI compared to standard of care PCI:

- 1.Does not reduce infarct size
- 2.All cause mortality at 6 months was not different
- 3. Exploratory composite clinical endpoint favored of IABC

### Lessons for Current and Future Care



- These findings do not support the <u>routine</u> use of IABC prior to PCI in Anterior STEMI patients without cardiogenic shock,
- Clinicians should continue to be vigilant about identifying patients who are at risk for rapid deterioration or hypotension that may benefit from support, as seen with the cross-over in this trial (8.5%)
- Acute STEMI studies are feasible without significant increases in door-to-device times

# Acknowledgements



### **CRISP Steering Committee**

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# CRISTO Counterpulsation reduces infarct size pre pci in acute myocardial infarction