# Results From The Minimizing Adverse Haemorrhagic Events By Transradial Access Site And Systemic Implementation of Angiox-MATRIX Access and Anti-thrombin Programs



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#### Background

- Previous studies have come to differing conclusions with regards to the role of radial access and bivalirudin in reducing adverse outcomes in patients with ACS
- It remains unclear whether avoiding access site bleeding and vascular complications through routine transradial intervention or routine use of bivalirudin improves outcomes in unselected patients with ACS undergoing invasive management

#### **MATRIX** Program

NCT01433627

**NSTEACS** or **STEMI** with invasive management

Aspirin+P2Y12 blocker

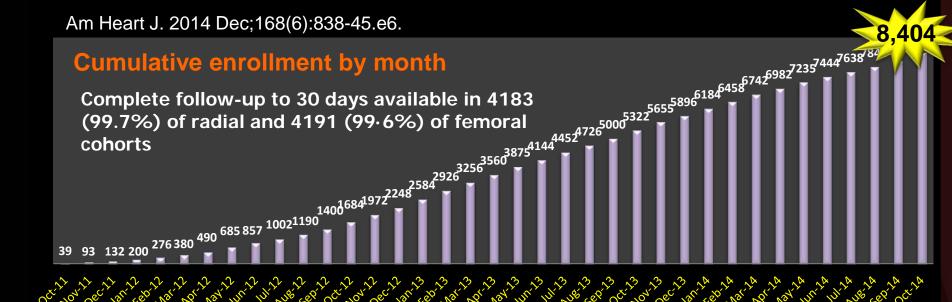
1:1**Trans-Radial Trans-Femoral ACCESS** Access Access 1:1 Bivalirudin **Unfractionated** Heparin with planned or **Bailout GPI** bailout GPI **ANTITHROMBIN** 



#### MATRIX Access

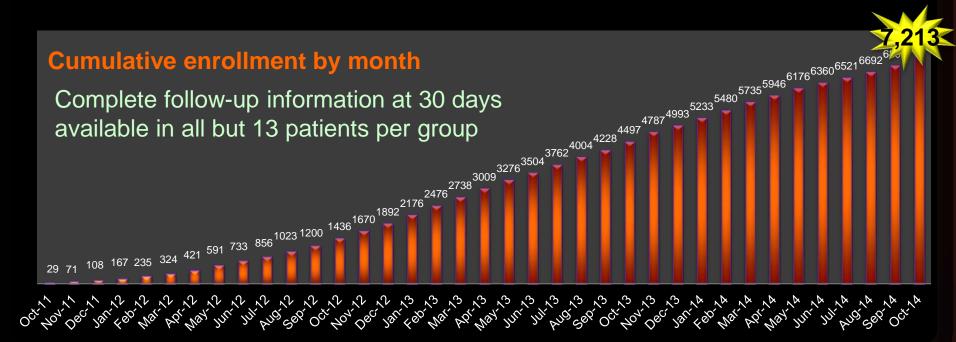


- 8,404 patients with ACS undergoing coronary angiography ± PCI from 11<sup>th</sup> Oct 2011 to 7<sup>th</sup> Nov 2014
- Operator Eligibility Criteria: Interventional cardiologist expertise in TRI and TFI including at least 75 transradial coronary interventions and at least 50% of interventions performed via radial route in the year preceding site initiation



#### MATRIX Recruiting timelines: Anti-thrombin program

First Recruited patient: 11<sup>th</sup> Oct 2011 Last Recruited patient: 7<sup>th</sup> Nov 2014





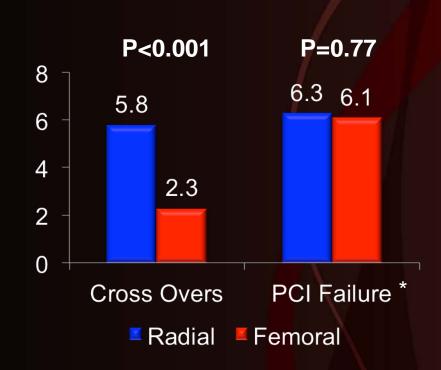
## **Cross Over and Procedural Success Rates**



94.1% of radial and 97.4% of femoral cohorts received respective treatment as allocated

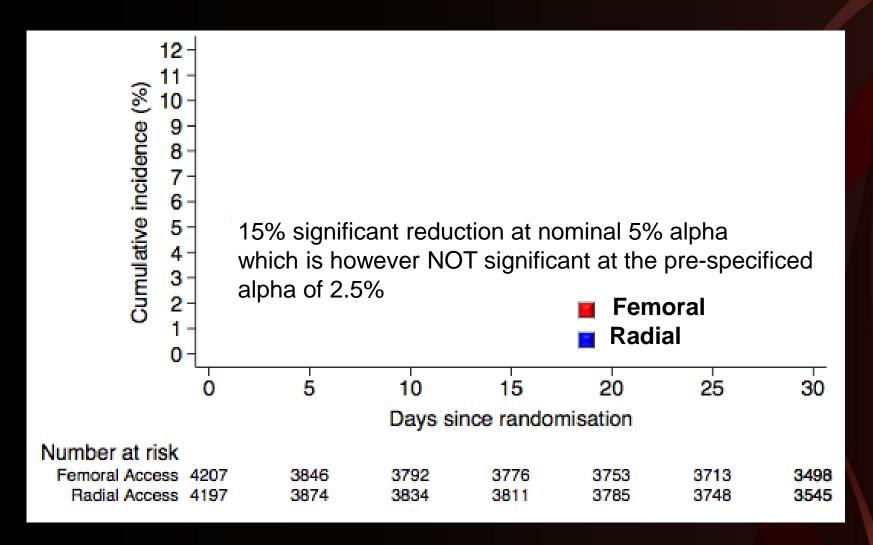
In 5.8% of radial and 2.3% of TF cohort the allocated access was attempted but failed.

In 3 (0.1%) in the radial and 13 (0.3%) patients in the femoral groups the allocated access was not attempted



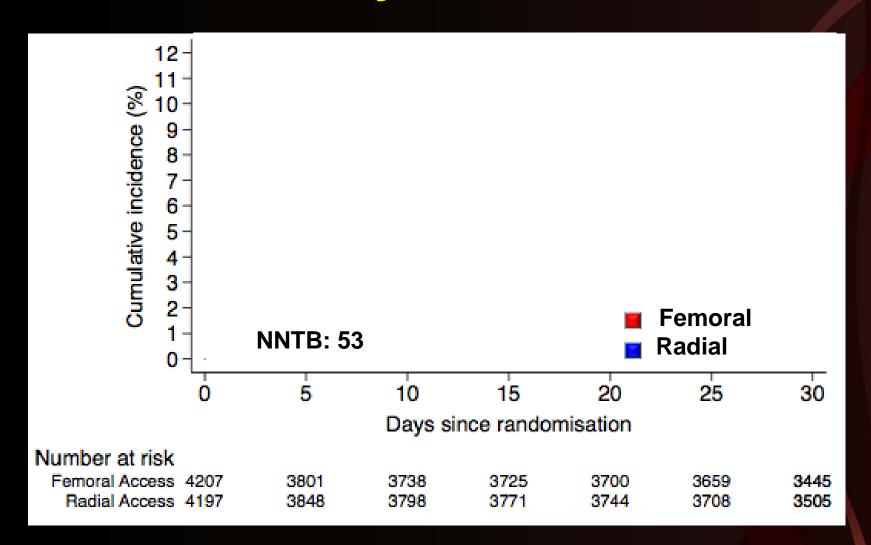


#### Primary EP: MACE





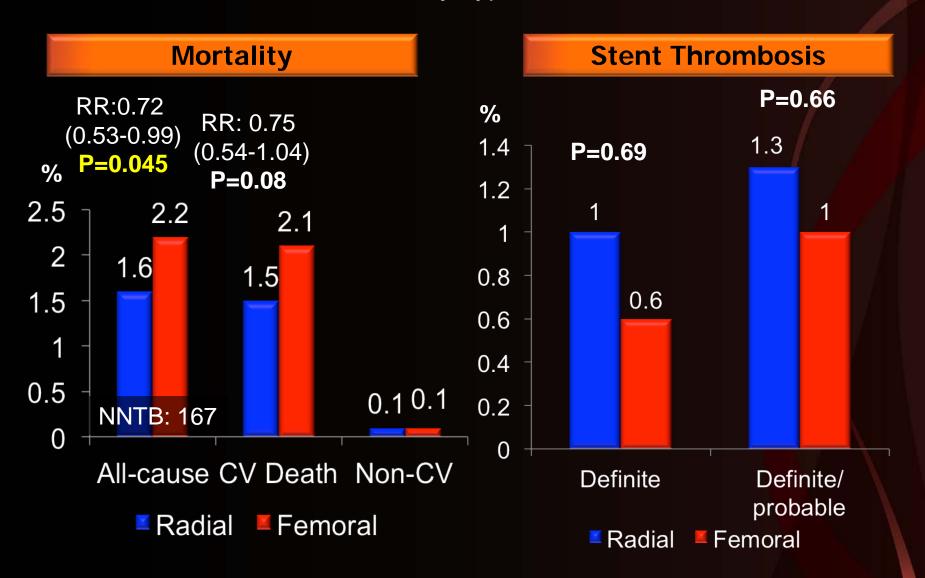
#### Primary EP: NACE



#### Fatal and ST EPs:



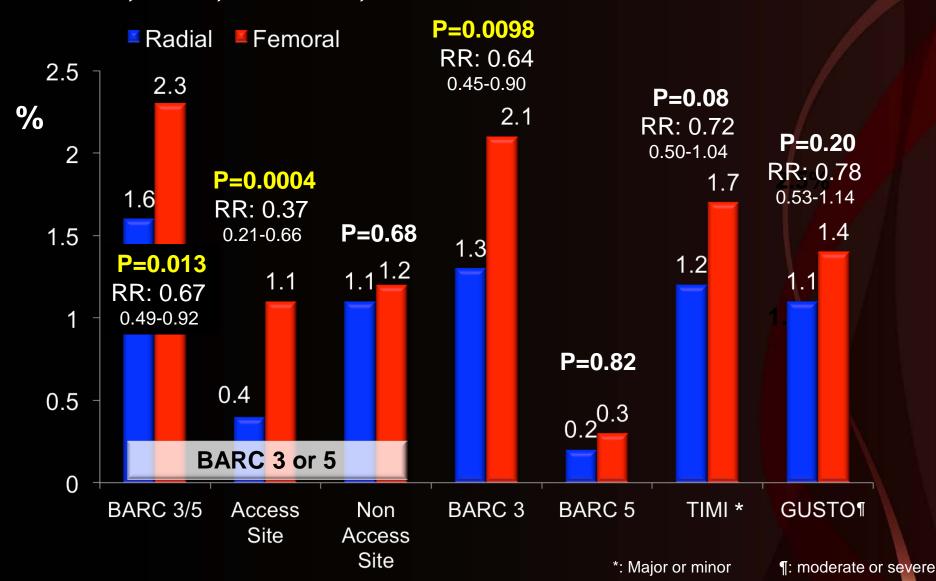
All-Cause, Cardiac, non-CV mortality, type of stent thrombosis



#### Bleeding endpoints:



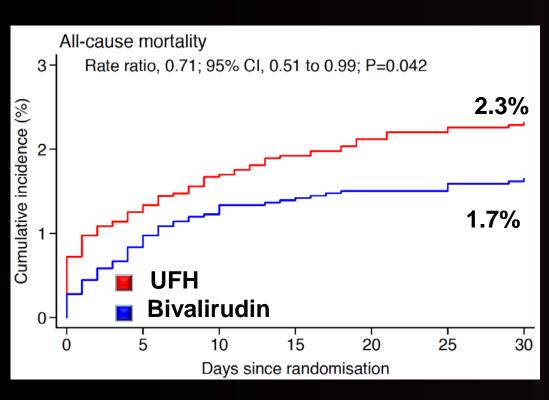
BARC, TIMI, GUSTO, access vs non-access related

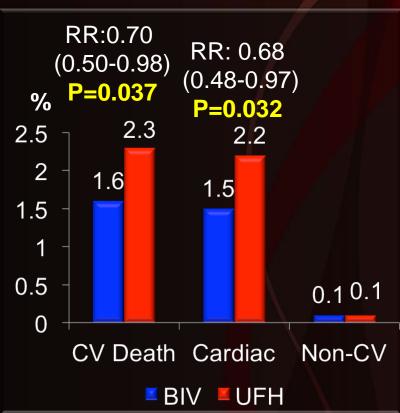


#### **Mortality:**



All-Cause, Cardiac, Vascular and non-CV mortality

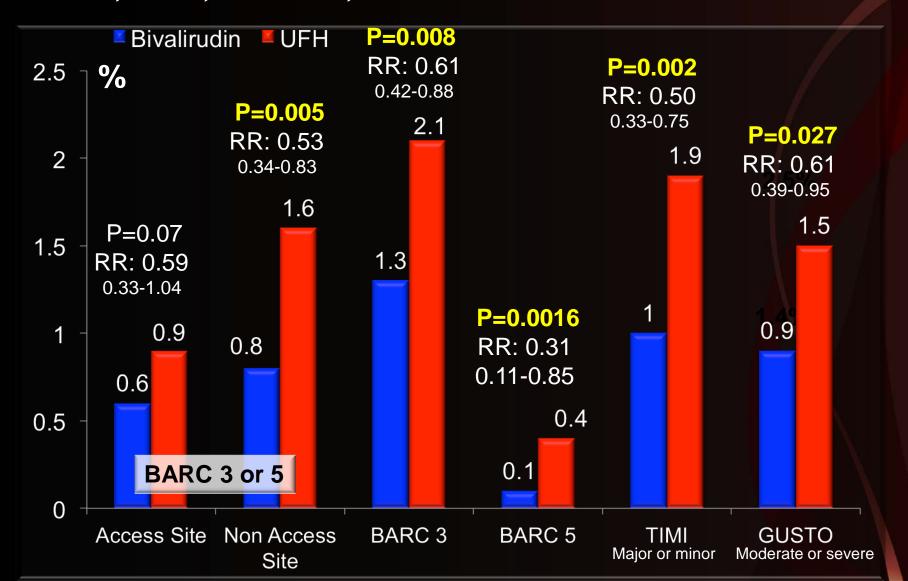




#### **Bleeding endpoints:**



BARC, TIMI, GUSTO, access vs non-access related



### Subgroup Analysis



MACE	HAZARD RATIO (95% CI)	HAZARD RATIO (95% CI)	P-VALUES	
			Superiority	Interaction
Femoral Radial	<b>-≡</b> -	0.94 (0.77-1.15)	0.56	0.98
	- <u>■</u> -	0.95 (0.76-1.17)	0.62	0.30
NACE				
Femoral	- <u>m</u> -	0.87 (0.72-1.05) 0.15	0.15	0.66
Radial	<b>-M</b> -	0.93 (0.75-1.14)	0.46	
Death		,		
Femoral		0.66 (0.42-1.03)	0.07	
Radial		0.77 (0.47-1.28)	0.31	0.64
Bleeding				
Femoral Radial	—×-	0.50 (0.32-0.79)	0.0024	0.51
		0.64 (0.37-1.10)	0.10	
			3113	
	2 1 0.25	<b>→</b>		
	<b>UFH Better</b> Bivalirudin I	Better		

#### Summary

- Radial as compared to femoral access lowered the risk of NACE, driven by a significant reduction of mortality and bleeding with no difference for myocardial infarction or stent thrombosis. This effect was independent from the anticoagulant used during PCI but expertise in TRI was a treatment modifier.
- Bivalirudin, when compared to UFH plus provisional GPI (used in 25% of patients) did not reduce the composite EP of MACE or NACE. however, the use of bivalirudin was associated to reduction of allcause and CV death and bleeding including fatal ones. This effect was independent from the access site used during the procedure.