# Assessing the clinical impact of rapid pathogen identification and antimicrobial susceptibility testing provided by the Accelerate Pheno<sup>™</sup> System at Hampshire Hospitals NHS Trust

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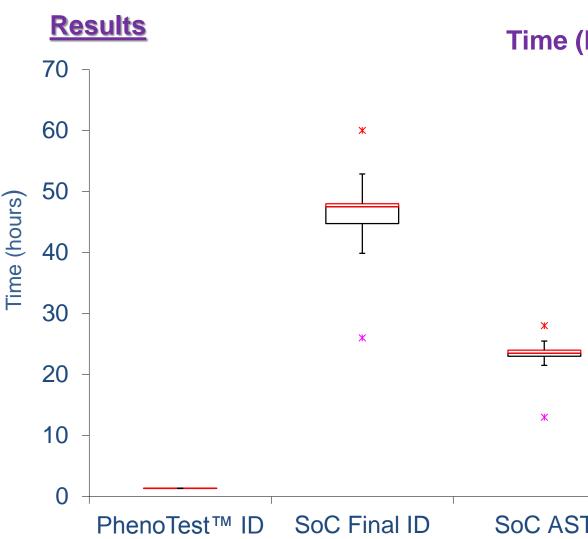
#### Background

Rapid identification (ID) and antimicrobial susceptibility testing (AST) of microorganisms cultured from blood is essential for optimising management of patients with sepsis. Clinical outcomes are influenced by timeliness of appropriate antimicrobials and investigations for source control. Timely de-escalation and targeted therapy are a vital component of antimicrobial stewardship

#### **Materials/Methods**

A prospective study was designed to investigate the clinical impact of the rapid ID and AST generated by the Accelerate Pheno<sup>™</sup> System versus our standard of care (SoC) of direct disc susceptibility and VITEK2. We processed signal positive blood cultures from 172 septic patients in ITU and on admission to ED and determined the clinical impact of these early interventions. 170 eligible assays were reported totalling 179 isolates (9 dual infections). AST testing from both the US and European panels PhenoTest<sup>™</sup> were performed, only the more relevant new European panel is reported here.

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## **Performance vs SoC (n=179)**

	Gram-negative	Gram-positive	Yeast	Overall
Sensitivity	96.3% [93.1-99.5%]	87.2% [76.7-97.7%]	100% [100-100%]	94.3% [90.8-97.7%]
Specificity	99.8% [99.6-100%]	99.9% [99.7-100%]	100% [100-100%]	99.9% [99.8-100%]
				[95% C

#### PhenoTest<sup>™</sup> BC (European Abx Panel) performance vs SoC (n= 55)

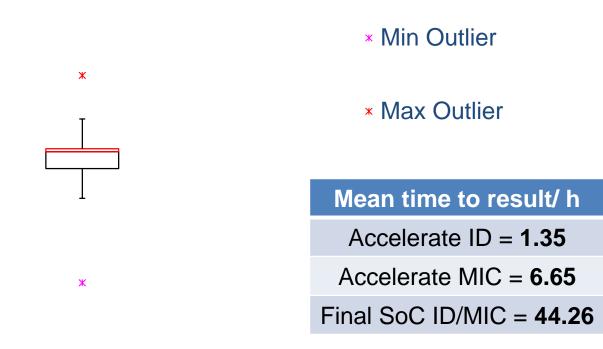
Essential Agreement (MIC)	Categorical Agreement (S, I, R)	Very Major Error (VME)*	Major Error (ME)	Minor Error (MiE)
97% [95.5-98.6%]	95.4% [93.5-97.3%]	5.3%	1.1%	2.3%
				[95% CI]

- Clinical action **1.57** days earlier than SoC.
- Lower sensitivity for Gram-positives reported due to favouring Gramnegatives in this study meaning less were assayed.
- VME<sup>\*</sup> due to Meropenem, 2x Tobramycin and 2x Cefuroxime (4x E. coli and P. aeruginosa.

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### Time (h) to ID & AST



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Т	SoC MIC	PhenoTest™ AST/MIC	T	

Clinical Impact	*106 blood cultures (BCs)
Optimisation of treatment	31
eduction of overall antibiotic use	19
larrowing of antibiotic spectrum	16
IV to oral switch	9
Infection control	12
	*analysis on going

analysis on-going

# **Hampshire Hospitals NHS Foundation Trust**

Annualised projections for our Trust based on 106 signal positive BCs

~5,500 doses saved/annum of Tazocin

~2,500 doses saved/annum of total antibiotic usage

1278 patient exposure days to multidrug resistant organisms saved

Clinical actions a mean of 1.57 days earlier compared to SoC

# Conclusion

The Accelerate Pheno<sup>™</sup> System represents an exciting innovative platform with potential for significantly decreasing the interval to antimicrobial optimisation in blood stream infection. The potential clinical impact is greatest in pathogens with unpredictable antibiograms like those we encounter locally in our Gram-negatives.

rapid clinical These demonstrable interventions can deliver significant benefits for individual patients and healthcare organisations in terms of quality of care, patient safety, antimicrobial stewardship and prevention infection measures, notwithstanding the potential associated financial savings.





