



# Evaluation of a Rapid Blood Culture Assay for Phenotypic Antimicrobial Susceptibility Testing of Gram-negative Bacteria on Antimicrobial Use in Children



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## Introduction

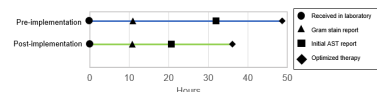
- Rapid identification and antimicrobial susceptibility testing (AST) from positive blood cultures can decrease the time to optimal therapy and reduce the use of broad-spectrum agents.
- The Accelerate Pheno Blood Culture panel (Pheno) provides AST of select on-panel Gram-negative organisms directly from positive blood cultures.
- We sought to determine the performance and the clinical impact of Pheno at our pediatric hospital compared to the BD Phoenix AST system (reference).

## Results

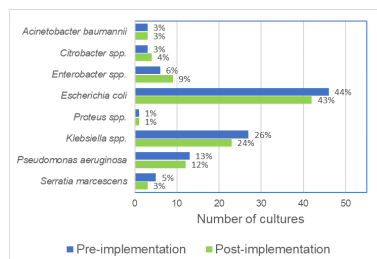
**Table 1.** Demographics and clinical outcomes

Demographics	Pre-implementation (n = 90)	Post-implementation (n = 93)	p value
Median age	6.4	2.3	0.16
Female	40 (44.4)	40 (43.0)	0.88
Immunocompetent	6 (6.7)	17 (18.3)	0.02
<b>Chart review (n = 100)</b>			
Mean length of stay	17.0 days	14.0 days	
30 day mortality	4 (4.0)	7 (7.2)	0.37
CVAD line removal	28 (28.0)	22 (22.7)	0.42
Community-onset	56 (56.0)	63 (64.9)	0.24
Hospital-onset	44 (44.0)	34 (35.1)	0.19
<b>Antimicrobial duration (n = 100)</b>			
Meropenem	47.7 hours	25.2 hours	<0.01

**Figure 1.** Median time to AST and optimization of therapy from time of receipt in laboratory

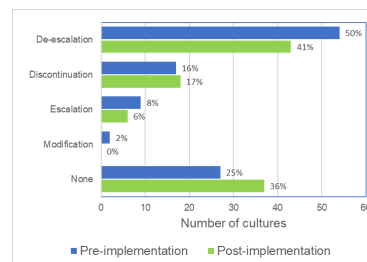


**Figure 2.** Organisms identified in blood cultures



## Results

**Figure 3.** Category of antimicrobial change after susceptibility testing results



**Table 2.** Antimicrobial susceptibility agreement between Pheno and reference method

Antibiotic	Minor errors (%)	Major errors (%)	Very major errors (%)
Amikacin	0	0	0
Ampicillin-sulbactam	20.7	1.7	1.7
Aztreonam	2.6	0	0
Cefazolin	16.1	5.4	0
Cefepime	6.8	0	0
Ceftazidime	16.1	2.3	0
Ceftriaxone	0	1.3	0
Ciprofloxacin	3.4	0	0
Ertapenem	1.4	0	0
Gentamicin	4.6	2.3	0
Meropenem	4.6	0	0
Minocycline	0	0	0
Piperacillin-tazobactam	14.9	0	0
Tobramycin	4.6	0	0
<b>Total</b>	<b>7.56</b>	<b>0.94</b>	<b>0.10</b>

## Results

Differences in categorical agreement

- 72 minor errors
  - Overcalling
    - 26.4% (R) when reference was (I)
    - 61.1% (I) when reference was (S)
  - Undercalling
    - 5.6% (I) when reference was (R)
    - 6.9% (S) when reference was (I)
- 9 major errors
  - 1 ampicillin-sulbactam, 3 cefazolin, 2 ceftazidime, 1 ceftriaxone, 2 gentamicin
- 1 very major error (ampicillin-sulbactam in Klebsiella pneumoniae)
- 9 of 12 ampicillin-sulbactam minor errors were due to overcalling resistance in *Escherichia coli* when the reference method was intermediate

## Methods

- We conducted chart review on a total of 100 cases tested by conventional AST directly from positive blood culture cell pellet during the period of May 2018 - April 2019 and a total of 97 cases tested by Pheno during May 2019 - March 2021. A total of 183 patients were tested.
- Pheno results in the test group were compared to the BD Phoenix AST system in the reference group.
- Duration of therapy, time to optimal therapy, and length of stay were calculated.

## Conclusions

- Pheno had accurate performance compared to the reference method. The majority of the minor errors were due to overcalling intermediate resistance when the reference was susceptible.
- The median time to initial AST report and optimal therapy decreased significantly after Pheno implementation.
- There was no significant impact on clinical outcomes such as 30-day mortality or central venous access device removal.
- There were significantly more immunocompetent patients in the post-implementation group, potentially impacting these results.
- The median duration on broad-spectrum meropenem decreased by 22.5 hours after Pheno implementation (P<0.01).

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