

Performance of the LifeScale Rapid AST vs. Clinical Standard of Care



P1454

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Background

Rapid ASTs (rASTs) must meet the challenge of reducing time to results without compromising accuracy. Six hospitals evaluated the LifeScale AST system (Affinity Biosensors), an FDA-cleared rAST that provides phenotypic AST results for Gram-negative organisms direct from blood cultures. Its performance was compared to each site's standard-of-care (SOC) automated AST platform.

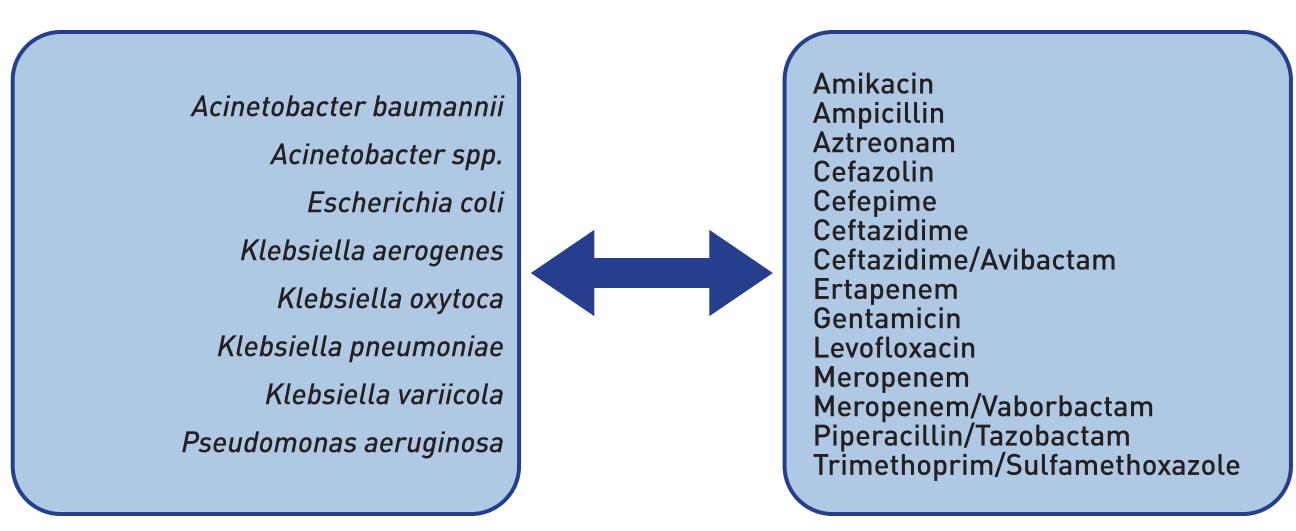
Introduction

Bloodstream infections are a significant cause of death and a major burden on hospitals. Furthermore, treatment using unnecessary broad-spectrum or combination regimens are associated with adverse effects and select for antimicrobial resistance. For these reasons, there is a strong need for shorter turnaround times required to perform AST. LifeScale AST system is an FDA cleared rapid AST shown to produce accurate results within 5 hours¹. The systems accuracy, ease-of-use and time to results were evaluated at the following institutions:

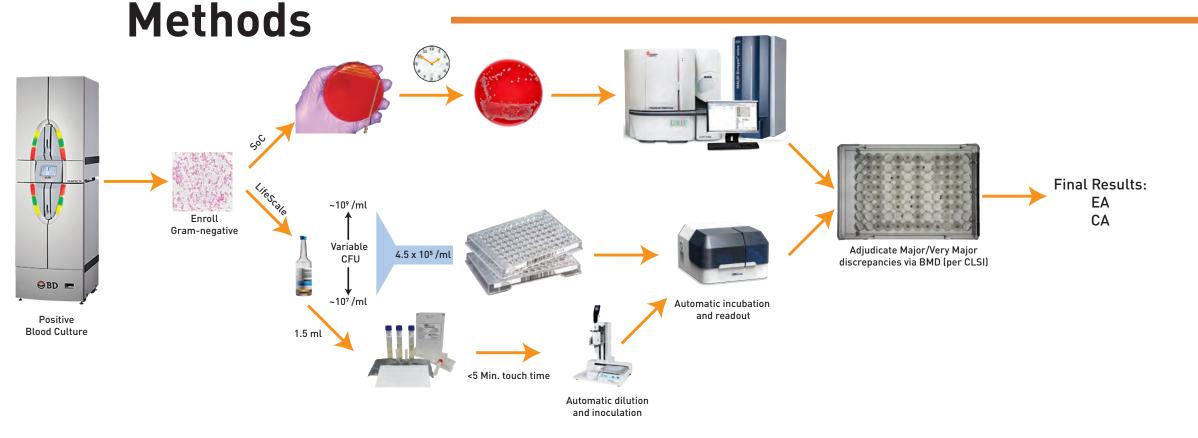
- University of Louisville, Louisville KY
- Baylor Scott and White, Temple, TX
- New York-Presbyterian Columbia University Irving Medical Center, New York, NY
- Associated Regional and University Pathologists (ARUP), Salt Lake City, UT
- Spartanburg Medical Center, Spartanburg, SC
- Northwestern Memorial Hospital, Chicago, IL

Microfluidic sensor detects individual microbes The mass of each microbe is measured: Population Profile Population Profile AI PREDICTOR MICROFILE AI PREDICTOR

The LifeScale AST system counts and weighs the mass of individual microbes. This allows for an accurate representation of the microbes' response to differing concentrations of antibiotics within a shorter incubation period.



The LifeScale AST System is FDA cleared for use on 93% of species/antibiotic combinations that are indicated for use, not intrinsically resistant and have FDA breakpoints available.



Positive blood culture samples detected positive by FDA approved continuous-monitoring blood culture system and confirmed as Gram-negative by Gram stain were enrolled in the study. Blood cultures were processed on the LifeScale AST system and FDA approved standard-of-care (SOC) AST system at each site (MicroScan WalkAway, VITEK 2, BD Phoenix). Very major and major discrepancies were adjudicated using BMD testing in accordance with CLSI guidelines, ^{2,3}.

Acknowledgements

We would like to thank the teams at University of Louisville, Baylor Scott and White, New York-Presbyterian - Columbia University Irving Medical Center, Associated Regional and University Pathologists, Spartanburg Medical Center and Northwestern Memorial Hospital for their collaboration on this work.

Links









Results

Table 1. LifeScale AST overall performance, Walkaway: 3; Phoenix: 2; Vitek: 1

Essential Agreement	Categorical Agreement	VMEs	MEs	mEs	Resistance
95.53%	94.74%	7/1410 (0.50%)	18/5019 (0.36%)	320/6582 (4.86%)	21.66%

Table 2. LifeScale error rate less than for SOC c.f rBMD

	LifeScale	SOCs
Very Major Errors	23/449 (5.12%)	38/449 (8.46%)
Major Errors	28/1024 (2.73%)	41/1013 (4.05%)

Table 3. Average time to results to report all antibiotics by genus

Genus	Species	Time to Result H:MM
Escherichia	coli	4:48
Klebsiella	species	4:57
Pseudomonas	aeruginosa	5:11
Acinetobacter	species	4:50
	Average	4:54
	Average (removing stacked delay)	4:32

Table 4. LifeScale AST performance by species.

Genus	No. Evaluated EA ¹	No. Evaluated CA	Essential Agreement	Categorical Agreement	Resistance
Acinetobacter	164	164	95.73%	90.85%	45.12%
Escherichia	3956	3952	96.76%	94.86%	23.56%
Klebsiella	1968	1978	95.12%	95.75%	18.10%
Pseudomonas	466	488	86.70%	90.98%	9.63%

Table 5. LifeScale AST performance by antibiotic.

	Essential Agreement ¹	Categorical Agreement	VMEs	MEs	mEs	Resistance
All	95.53%	94.74%	7/1410 (0.50%)	18/5019 (0.36%)	320/6582 (4.86%)	21.66%
Amikacin	91.13%	96.14%	0/19 (0.00%)	0/522 (0.00%)	21/544 (3.86%)	2.94%
Ampicillin	99.69%	99.08%	0/215 (0.00%)	0/112 (0.00%)	3/327 (0.92%)	65.75%
Aztreonam	95.01%	95.70%	0/121 (0.00%)	2/312 (0.64%)	17/442 (3.85%)	27.83%
Cefazolin	94.00%	82.53%	1/187 (0.53%)	2/264 (0.76%)	80/475 (16.84%)	38.95%
Cefepime	90.29%	95.47%	0/145 (0.00%)	1/439 (0.23%)	25/596 (4.19%)	25.00%
Ceftazidime	96.01%	92.64%	0/140 (0.00%)	2/457 (0.44%)	44/625 (7.04%)	23.36%
Ceftazidime/Avibactam	97.14%	99.04%	1/20 (5.00%)	1/189 (0.53%)	0/209 (0.00%)	10.53%
Ertapenem	98.75%	99.50%	0/38 (0.00%)	0/361 (0.00%)	2/401 (0.50%)	9.98%
Gentamicin	96.17%	97.67%	1/80 (1.25%)	1/514 (0.19%)	12/601 (2.00%)	13.31%
Levofloxacin	97.13%	89.65%	0/156 (0.00%)	1/435 (0.23%)	64/628 (10.19%)	25.32%
Meropenem	97.29%	97.93%	1/64 (1.56%)	0/561 (0.00%)	12/629 (1.91%)	10.65%
Meropenem/Vaborbactam	96.69%	97.77%	1/2 (50.00%)	0/174 (0.00%)	3/179 (1.68%)	1.12%
Piperacillin/Tazobactam	94.38%	92.08%	1/66 (1.52%)	5/453 (1.10%)	37/543 (6.81%)	11.79%
Trimethoprim/Sulfamethoxazole	98.17%	98.96%	1/157 (0.64%)	3/226 (1.33%)	0/383 (0.00%)	39.69%

Table 6. The studies tested 477 prospective and 179 seeded challenge samples comprising 7,188 organism/antibiotic combinations of which 21.4% were resistant.

Species	Number Enrolled	LifeScale%
baumannii/baumannii complex	34	
lwoffii	2	21.66%
radioresistens	2	4.18%
species	3	74.16%
ursingii	1	
coli	329	SOC%
aerogenes	21	300%
oxytoca	38	21.44%
pneumoniae	147	2.10%
variicola	1	76.46%
aeruginosa	78	
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¹ Essential agreement adjusted for differences in panel ranges.

Conclusions

LifeScale AST produced rapid results from positive blood cultures in under 5 hours that were in excellent agreement with three major standard-of-care ASTs. Notably, LifeScale's rate of Very Major and Major errors was lower than for the SOCs, demonstrating that its speed does not compromise accuracy, and that its rapid results can be relied upon to improve antimicrobial stewardship and patient outcomes.

References

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- 3. CLSI. Methods for Dilution Antimicrobial Susceptibility Tests for Bacteria That Grow Aerobically, 12th ed. CLSI standard M07. Clinical and Laboratory Standards Institute; 2024.