

# Assessment of the SPECIFIC REVEAL® Rapid AST System in the determination of the sensitivity of *Pseudomonas aeruginosa* to antibiotics

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## Introduction

In *P. aeruginosa* bacteremias, reducing the time to obtain an antibiogram is believed to be a key element in improving the survival rate of patients infected with multi-drug resistant strains. Where appropriate, it also helps to limit the use of last resort antibiotics, accordance with the Antibiotic Stewardship Program.

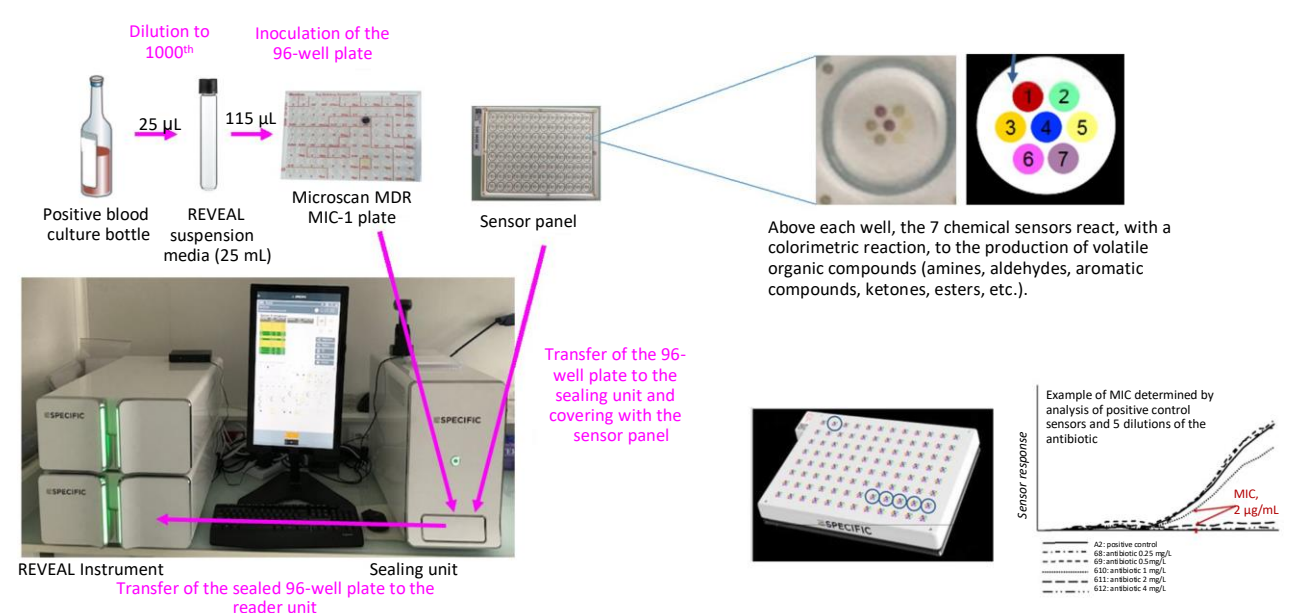
## Goal

The aim of the study was to assess the performance of Specific Diagnostics' SPECIFIC REVEAL® Rapi AST System (SPECIFIC REVEAL) capable of determining bacterial sensitivity to antibiotics (MIC) by microdilution from blood culture bottles with the detection of volatile organic compounds.

## Methods

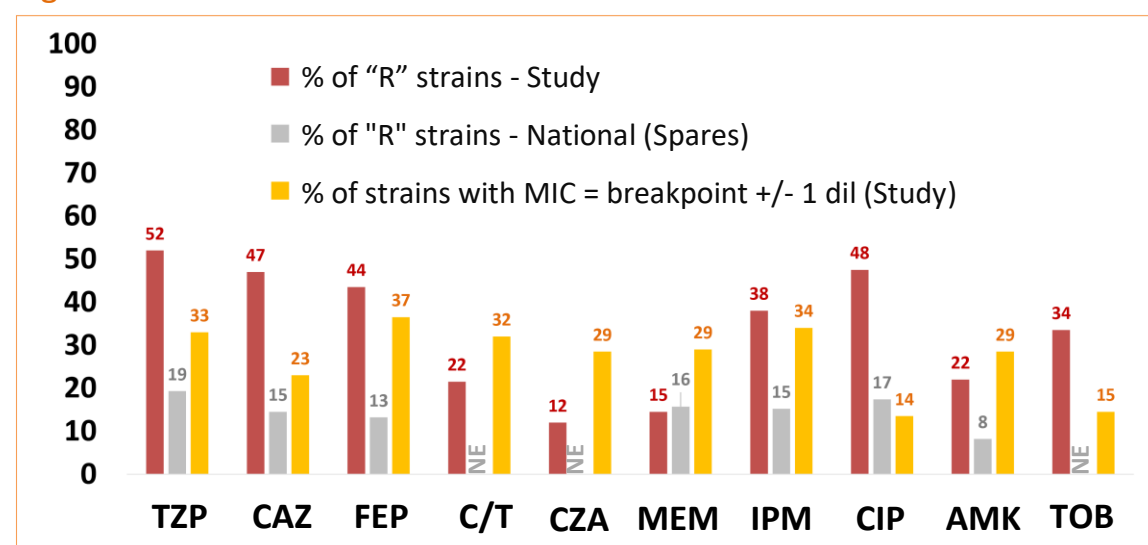
- 200 strains of previously genotyped *P. aeruginosa* were selected on the basis of their antibiotic sensitivity profiles.
- BACTEC blood culture bottles were inoculated with 10<sup>2</sup> CFU of each strain and incubated until bacterial growth.
- The broths from the positive bottles were then withdrawn and diluted to inoculate the microplates for the SPECIFIC REVEAL BC-AST Assay (Figure 1).
- A broth microdilution method (Sensititre® GN6F) was used as the reference.
- The results were interpreted according to CA-SFM 2021.
- The performances have been calculated according to the ISO 20776-2:2007 standard.

Figure 1. SPECIFIC REVEAL - Analytical Process



## Results

Figure 2. Characteristics of the studied strains



Spares, Mission Spares 2019 surveillance data (May 2021); NE, not assessed

Table 1. Performance of the SPECIFIC REVEAL

	TZP	CAZ	FEP	C/T	CZA	MEM	IPM	CIP	LVX	AMK	TOB	Total
<b>Render time (average in h)</b>	7:35	6:12	5:56	6:23	5:56	6:31	6:28	5:59	6:31	6:23	6:03	<b>6:22</b>
<b>% uninterpretable results</b>	0.0	0.0	0.0	0.0	0.0	0.5	0.5	0.0	0.0	0.0	0.0	<b>0.1</b>
<b>% CA</b>	96.5	94.5	89.5	99.0	98.5	92.5	92.5	98.0	98.5	97.5	100.0	<b>96.1</b>
<b>% VME</b>	1.0	3.2	0.0	4.5	0.0	0.0	0.0	2.1	3.1	2.3	0.0	<b>1.6</b>
<b>% ME</b>	6.3	7.5	18.6	0.0	1.7	0.0	11	1.9	0.0	2.6	0.0	<b>4.2</b>
<b>% Me</b>	-	-	-	-	-	7.0	-	-	-	-	-	<b>0.6</b>

ISO 20776-2:2007 acceptance criteria: category agreement (CA) ≥ 90%; very major errors (VME) ≤ 3% and major errors (ME) ≤ 3%

- **Collection studied.** The strains analyzed were more resistant than those normally isolated routinely (Spares 2019 data). For 13% to 36% of the strains, the MICs were located within +/- 1 dilution of the CA-SFM breakpoint depending on the molecules.
- **Practicability.** The antibiogram preparation was simple and quick (<5 min).
- **Time to result.** Results were available in an average of 6 hours 22 minutes, and were released after the species information was entered in to the SPECIFIC REVEAL.
- **Performance.** In view of the collection studied and compared to the reference technique, the performance of the SPECIFIC REVEAL was very satisfactory (%CA > 90%, %VME < 3%). However, the %ME was > 3% due to a relatively high percentage of strains erroneously made resistant to cefepime (FEP) and imipenem (IPM).

## Conclusions

The SPECIFIC REVEAL can reliably determine the sensitivity of *P. aeruginosa* strains in antibiotics from blood culture bottles, within an average of 6.5 hours. Its impact on the early management of patients with bacteremia must now be evaluated on a routine basis.

The poster is available on our website

