



How do we use the rapid LAMP-PCR information of Enterobacterales ESBL producers causing bacteremia to guide empirical treatment?

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Inadequate empirical antimicrobial therapy (EAT) for bacteremia is associated with an increase in the mortality rate. The early detection of resistance mechanisms will imply a more suitable empiric therapy, reducing the time to obtain the complete antimicrobial susceptibility testing (AST).

The **aim** of this study is to evaluate whether the information generated from a rapid molecular technique for ESBL detection modifies the treatment of bacteremia caused by Gram-negative bacilli (GNB) before the definitive AST report is available.

Methods

A multicenter 3 year-intervention study was performed on Enterobacterales monobacterial bacteremia were included. The study was conducted in two phases:

Pre-intervention phase using a standard work flow routine without performing the molecuar test.

Intervention phase with a modified routine in order to detect resistance mechanisms directly from positive blood cultures by LAMP-PCR (Genie, eazyplex SuperBug CRE, Menarini)

All clinical and microbiological relevant information were recorded in RedCAP database. EAT evaluation was performed *a posteriori* (optimal, appropriate, inappropriate, inadequate) according the AST result at three points: (1) no microbiology information available; (2) the blood cultured tested positive and identification or partial AST was available; (3) antibiogram-guided treatment. An EAT rating scale (-2 to +2) was applied to evaluate goodness of EAT compared with the targeted antibiotic indicated after microbiology lab information was available (Gram stain/MALDI-TOF in pre-intervention and LAMP-PCR in intervention phase).

Results

A total of 163 bacteremia were included. Clinical information of the patients included in both periods is shown in Table 1. Evaluation of empirical treatment modification with and without LAMP-PCR information is shown in Figure 1.

Pre-intervention and intervention empirical treatment rating score were 0.86 and 1.2, respectively.

Pre-intervention	Intervention
02	80
83	80
13,30%	18,80%
52,9%	45,0%
47%	55%
72 уо	75 уо
57,8%	62,5%
28,9%	27,5%
13,3%	10,0%
	Pre-Intervention 83 13,30% 52,9% 47% 72 γο 57,8% 28,9% 13,3%



OPTIMAL

The most accurate or favorable antibiotic treatment according to the AST as well as clinical condition of the patient.

APPROPRIATE

Appropriate according to the AST but the spectrum could be reduced. A reduction in the spectrum is not possible because of the patient's clinical condition.

INAPPROPRIATE

Unnecessarily broad-spectrum antibiotic therapy. The

Urinary	65 <i>,</i> 0%	62,5%
Abdominal	9,6%	6,3%
Biliar	15,7%	22,5%
Other	9,7%	8,7%
Previous	5,9%	10,0%
colonization		
Preescriptor		
Service		
Emergency	69,4%	65,0%

treatment is not justified based on the clinical condition of the patient.

INADEQUATE

Not covering the cultured microorganism based on the in vitro AST or no antibiotic therapy at all when antibiotic therapy was required,

Table 1. Clinical information of the patients included in both periods.

Figure 1. Evaluation of EAT when microbiology reports partial information of the result.

Conclusions

- Optimal and appropiate EATs are improved when molecular resistance is available.
- Early knowledge of the resistance mechanism helps to reduce the number of inappropriate and inadequate EATs.
- Early reporting of the resistance mechanisms to the clinician improves the appropriateness of the EATs in Enterobacterales bacteremias.

