

Results from 2015 Antimicrobial Susceptibility Testing External Quality Assessment (EQA) Exercise Organised for EARS-Net Participants

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Introduction

The United Kingdom National External Quality Assessment Service (UK NEQAS) for Microbiology has provided external quality (EQA) for antimicrobial assessment to the European testing susceptibility Surveillance Resistance Antimicrobial Network (EARS-Net - formerly EARSS) since 2000. The aim is to assess and monitor the comparability of results between laboratories and countries and thus justify the pooling and routinely comparison of collected antimicrobial susceptibility test data across Europe.

analysis was carried out on the An performance of participants in the quality assessment exercise. Participation was invited from 974 laboratories in 30 countries and results were returned by 902 laboratories. The organisms distributed were: Enterococcus faecalis, Klebsiella pneumoniae, Staphylococcus aureus, Streptococcus pneumoniae, Escherichia coli and Pseudomonas aeruginosa. Participants' results for identification and antimicrobial susceptibility testing were collated and assessed.

Methods

• to assess the ability of participating laboratories to identify antimicrobial resistance of clinical and public health importance • to determine the accuracy of antimicrobial susceptibility test results reported by individual

- laboratories
- education



Specimen 3083 - Klebsiella pneumoniae This organism produces an **OXA-48** carbapenamase conferring reduced susceptibility to carbapenems.

MICs of third generation of cephalosporins were within the susceptible category but were borderline, and this is reflected in the high discrepancy rates for these agents (Chart 1).



S



In the presence of OXA-48 the MICs of carbapenems are commonly raised, often resulting in resistance to ertapenem, while the effect on other carbapenems is less marked. A total of 99.2% of participants correctly reported the resistance to ertapenem, (MIC 8mg/L).

Susceptibility to imipenem (MIC 4-8 mg/L) was intermediate-resistant with EUCAST breakpoints and resistant with CLSI breakpoints. Susceptibility to meropenem (MIC 2-4 mg/L) was susceptible-intermediate with EUCAST breakpoints and intermediate-

resistant with CLSI breakpoints. The differences in breakpoints and borderline susceptibility were reflected in the variable reporting, (Chart 2).





Specimen 3084 - Staphylococcus aureus

This organism exhibits low level resistance to vancomycin and teicoplanin (VISA). It is the same strain of VISA that was distributed in the EARS-Net EQA exercise in 2014 and there is little evidence of change in performance of participants (table 1).

Results continued

		Table 1
	2014 (n=819)	2015 (n=877)
S	394 (48.1)	431 (49.2)
I	80 (9.8)	81 (9.2)
R	345 (42.1)	365 (41.6)

Isolates of S. aureus with vancomycin MICs of 4-8 mg/L were originally termed "vancomycin intermediate S. aureus" (VISA) because the level of resistance is low and is distinguishable from the high-level resistance displayed by S. aureus carrying the mecA gene (MICs >8 mg/L). While CLSI have maintained this distinction EUCAST does not have an intermediate category because VISA strains are clinically resistant.

Reports of susceptible were less frequent among 738 participants following EUCAST/EUCASTrelated guidelines (48% susceptible) than among the 139 following CLSI guidelines (55.4% susceptible). In line with breakpoints, most nonsusceptible reports with CLSI guidelines were in the intermediate category while with EUCAST guidelines most were in the resistant category (chart 3).

The objective of the EQA exercise:

• to assess the comparability of results between laboratories and countries and thus justify the pooling and comparison of routinely collected antimicrobial susceptibility test data across Europe

Results

With some notable exceptions the level of performance was high (overall concordance with the intended results ranged from 27% to 100%).

Data are presented for some of the • general findings and a few of the





EQA is a valuable tool in the quality assurance of antimicrobial susceptibility testing in the diagnostic laboratory and demonstrates the validity of comparing collated data between laboratories. In this exercise overall concordance between participating laboratories was high except where there was borderline susceptibility or different guidelines used.

Acknowledgement

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