

the speed of glutaraldehyde at their respective minimum effective concentrations.

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PCR and Conventional Tests Used for MRSA Detection

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Conventional and molecular techniques are being used in the detection of methicillin resistance in *Staphylococcus aureus* (MRSA) but they do not always show concordant results. Araj and coinvestigators from the Department of Pathology and Laboratory Medicine, American University of Beirut Medical Center, Lebanon, compared a *mecA* polymerase chain reaction (PCR)-based amplification with the 1 µg oxacillin disk-diffusion test and the Epsilon meter test (E-test) for detection of minimum inhibitory concentrations (MICs). Among 31 isolates initially character-

ized as MRSA by the disk-diffusion test, *mecA* was detected in only 13 isolates (42%). The E-test showed a wide range of oxacillin MICs (0.5- >256 µg/mL) among these 31 MRSA isolates: 7 isolates had an MIC of >256 µg/mL, 1 had 64 µg/mL, 2 had 4 µg/mL, 2 had 3 µg/mL, 1 had 2.5 µg/mL, 9 had 2 µg/mL, 3 had 1.5 µg/mL, 5 had 1 µg/mL, and 1 had 0.5 µg/mL.

Comparing the *mecA* PCR results with the E-test oxacillin MIC findings revealed that *mecA* was detected in 7 of 8 isolates (87.5%) with an MIC of ≥64 µg/mL, in 3 of 14 isolates (21.4%) with an MIC of 2 to 4 µg/mL, and in 3 of 9 isolates (33.3%) with an MIC of <2 µg/mL. B-Lactamase production was positive in 28 of 31 isolates (90.3%).

Because of this variation between tests, and because several resistance mechanisms are known to mediate methicillin resistance in *S aureus*, the reliable detection of MRSA cannot be based solely on detection of *mecA* gene in *S aureus*.

At this stage, and until new guidelines are introduced by an official body such as NCCLS, a combination of conventional methods alone or together with a molecular method should be used every time *S aureus* is tested for detection of methicillin resistance.

FROM: Araj GF, Talhouk RS, Simaan CJ, Maasad MJ. Discrepancies between *mecA* PCR and conventional tests used for detection of methicillin resistant *Staphylococcus aureus*. *Int J Antimicrob Agents* 1999;11:47-52.