

Specific detection of coagulase positive methicillin-resistant *Staphylococcus aureus* by multiplex PCR method

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Background and Aims: This study conducted to implement, for the first time in our laboratory, a triplex PCR technique for detection of genes encoding resistance to methicillin (*mecA* and *femA*) in *Staphylococcus aureus* from clinical samples.

Methods: After sample collection from three University hospitals in Zanjan province, antimicrobial susceptibilities were determined by Kirby-Bauer disk diffusion according to the NCCLS recommendation. DNA was extracted by phenol – chloroform standard method. By using multiplex PCR strategy two regions of *mecA* and *femA* genes were co amplified. A third *Staphylococcal* genomic region was used as internal control and PCR products were analyzed by electrophoresis.

Results: 50 methicillin-resistant *S. aureus* identified by antibiotic susceptibility testing from hospitals of Zanjan University of Medical Sciences. The *mecA* and *femA* were found in 100% of coagulase positive *S. aureus*.

Conclusions: These results suggest that the multiplex PCR method mentioned above can be used to provide a specific, rapid, simple, and highly sensitive detection of coagulase positive *S. aureus* in clinical samples.

Keywords: *S. aureus*; Multiplex PCR; *mecA*; *femA*