

RESEARCH NOTE

INTERNATIONAL MICROBIOLOGY (2004) 7:63–66

INTERNATIONAL
MICROBIOLOGY

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Simultaneous PCR detection of *ica* cluster and methicillin and mupirocin resistance genes in catheter-isolated *Staphylococcus*

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Summary. Recent data show that more than 50% of catheter-associated bloodstream infections are caused by staphylococci. Staphylococcal infections produced by intercellular-adhesion cluster (*ica*) carriers can be even more problematic due to the presence of methicillin and mupirocin resistance genes. In the present study, a multiplex PCR protocol that allows the simultaneous identification of staphylococci and detection of both the *ica* and methicillin and/or mupirocin resistance genes was developed. Furthermore, the method allows differential detection of the *ica* locus from *Staphylococcus aureus* and *Staphylococcus epidermidis*. [*Int Microbiol* 2004; 7(1):63–66]

Key words: *Staphylococcus aureus* · *Staphylococcus epidermidis* · intercellular adhesion gene cluster · antibiotic resistance · multiplex PCR

Received 9 April 2003
 Accepted 13 May 2003

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Introduction

A major achievement within the medical field in the twentieth century was the invention and development of prosthetic surgical implants. However, such implants frequently become infected by bacteria [7,18,23]. This is a serious complication, especially when the infection is caused by multiresistant bacteria, which are difficult to eradicate from the prosthetic material. Additionally, the efficiency of some bacteria in their ability to colonize indwelling medical devices,

such as catheters [6,7], poses particular problems. These bacteria develop a highly consolidated structure: the biofilm [6,7,25]. In the biofilm, microbial populations reside within a matrix that facilitates cell-to-cell and/or cell-surface adherence, resulting in an inherent structural resistance towards antimicrobial agents, such as antibiotics, disinfectants, and germicides [7,8,19,24]. In the case of staphylococci, formation of the biofilm requires polysaccharidic intercellular adhesin (PIA), which is synthesized by enzymes encoded by the intercellular adhesion cluster (*ica*) [5,11,14].

Recent data show that 50% of pathogens isolated from

hospital-acquired bloodstream infections, normally associated with the use of central-venous catheters, are staphylococci [3,4,9]. Furthermore, many *Staphylococcus aureus* and *Staphylococcus epidermidis* strains carry the *ica* cluster. Staphylococcal infections produced by *ica* carriers can, in turn, be even more problematic due to the presence of methicillin and mupirocin resistance genes [10,15,20].

Nosocomial infections due to methicillin-resistant staphylococci have increased subsequent to the widespread use of β -lactamic antibiotics. The transmission of resistant strains among hospital departments, and the ability of bacteria to transmit resistance genes are well-known [18,22]. With respect to mupirocin, several recent studies indicate the utility of topical mupirocin application in order to avoid the colonization of indwelling medical devices. While mupirocin is not an antibiotic for treatment of infections once they occur, it is used prophylactically to prevent *S. aureus* strains from reaching the inner part of the device. However, this application has led to the selection of highly mupirocin-resistant staphylococci [2,4,16]. For example, in our hospital, the incidence of mupirocin resistance has increased sharply from 7.7% in 1998 to 19% in 2000 [22].

Since most *S. aureus* are *ica* carriers, the colonization of devices by isolates harboring both resistance genes and adhesion genes has become a serious problem. Rapid detection of the *ica* locus in hospital staphylococcal isolates, together with simultaneous detection of antibiotic-resistance genes, may allow the development of methods to prevent or reduce the ability of bacterial to invade indwelling medical devices. It should be noted that the presence of the *ica* locus does not guarantee its expression; thus, it does not directly reflect biofilm formation. However, anticipating and detecting the possibility of biofilm colonization of catheters before it actually occurs would be of great help in preventing infection. For this reason, we developed a method to simultaneously detect *ica* and methicillin- and mupirocin-resistance markers. The method described here is a rapid and easy means to verify that catheters are not being colonized by mupirocin-resistant, methicillin-resistant staphylococci harboring the *ica* cluster. The presence within the hospital of these staphylococci would not only make the preventive use of mupirocin useless, it could also lead to an increased risk of infections.

Materials and methods

Bacterial isolates, identification, susceptibility testing, and DNA extraction method. *S. epidermidis* strains 1855-25, 9295-79, and 9951-79 were isolated from catheters obtained from the Oncology Service of our hospital (Nuestra Señora de Candelaria University Hospital, Santa Cruz de Tenerife, Spain). *S. aureus* strain 229 was isolated from the sputum of a patient with sepsis. All strains were biochemically identified, and

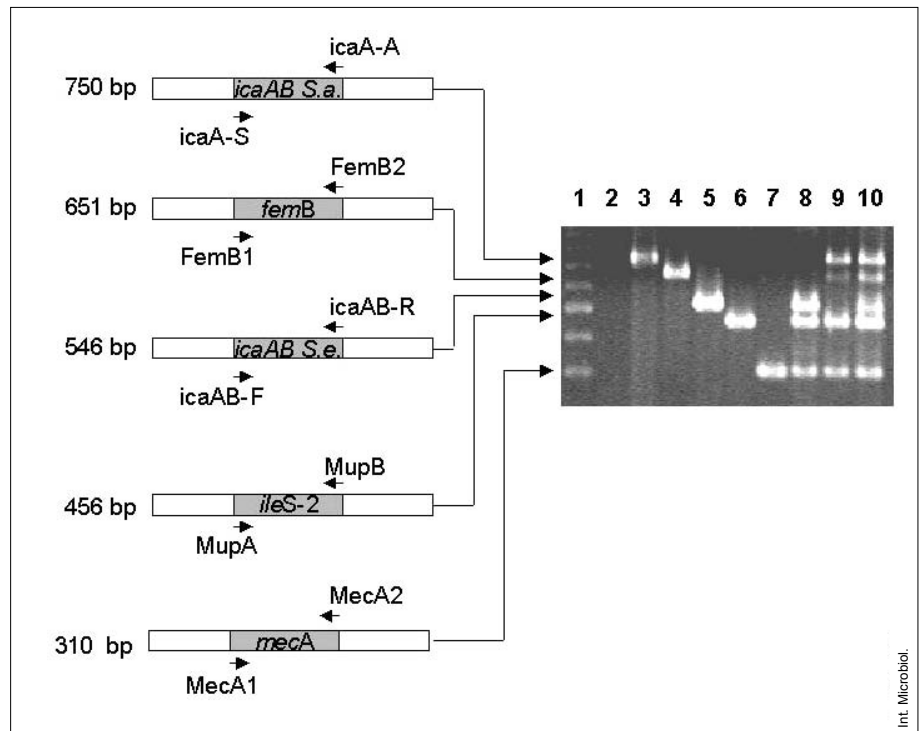
antibiotic-resistance patterns were determined according to the standard laboratory criteria of the Microbiology Service of our hospital. *S. aureus* 229 and *S. epidermidis* 1855-25 and 9295-79 were methicillin and mupirocin resistant, whereas *S. epidermidis* 9295-79 was mupirocin and methicillin sensitive.

The DNA extraction method has been described previously [20,21]. After overnight culture on brain-heart infusion agar plates, one or two colonies (one from a *S. aureus* isolate and one from a *S. epidermidis* isolate) were suspended in 20 ml of sterile distilled water, and the suspension was then heated at 100°C for 20 min. From this suspension, a 5-ml aliquot was directly used as template for PCR amplification.

Multiplex PCR amplification. The oligonucleotide primers have all been previously described [1,10,13,17,20]. *ica*AB-F: 5'-TTATCAATGC-CGCAGTTGTC-3' and *ica*AB-R: 5'-GTTTAACGCGAGTGCCGCTAT-3' from partial *icaAB* genes of *S. epidermidis*, *ica*A-S: 5'-AAACTTGGTGCG-GTTACAGG-3' and *ica*A-E: 5'-TCT GGGCTTGACCATGTTG-3' from partial *icaAB* genes of *S. aureus*, FemB1: 5'-TTACAGAGTAACTGT-TACC-3' and FemB2: 5'-ATACAAATCCAGCACGCTCT-3' from *femB*, MupA: 5'-TATATTATGCGATGGAAGGTTGG-3' and MupB: 5'-AATAAAATCAGCTGGAAAGTGTG-3' from *ileS*-2 and MecA1: 5'-GTAGAAATGACTGAACGTCCGATAA-3' and MecA2: 5'-CCAATTC-CACATTGTTTCGGTCTAA-3' from *mecA*. Multiplex PCR assays were all carried out directly using the bacterial suspension obtained after the rapid DNA extraction method described above. A 5-ml aliquot of this suspension was added to 45 ml of PCR mixture consisting of 1 \times reaction buffer [16 mM (NH₄)₂SO₄, 67 mM Tris-HCl (pH 8.8)], 0.2 mM of each of the four dNTPs (Promega, Madison, WI, USA), 3.5 mM MgCl₂, 3.6 μ M of each *femB* primer, 0.2 μ M of each *ileS*-2 primer, 0.1 μ M of each *mecA* primer, and 0.8 μ M of each *icaAB* primer and 1.25 U of Taq DNA polymerase (Bioline, UK). For each sample, one reaction was done with the *femB* pair of primers to identify *S. aureus* strains, with the *mecA* and *ileS*-2 pairs of primers to detect both resistance markers, and with the two *icaAB* pairs of primers to detect the *ica* cluster from *S. aureus*, *S. epidermidis*, or simultaneously from both. In order to reduce the formation of nonspecific extension products, a "hot-start" PCR protocol was used. All multiple PCR assays were carried out with a negative control containing all of the reagents without DNA template. DNA was amplified in a GeneAmp PCR system 9700 thermocycler (PE, Applied Biosystems, CA, USA) with the following thermal cycling profile: an initial denaturation step at 94°C for 5 min was followed by 10 cycles of amplification (denaturation at 94°C for 30 s, annealing at 66°C for 45 s, and extension at 72°C for 60 s); 5 cycles of amplification (denaturation at 94°C for 45 s, annealing at 64°C for 45 s, and extension at 72°C for 60 s); and 25 cycles of amplification (denaturation at 94°C for 45 s, annealing at 50°C for 45 s, and extension at 72°C for 60 s), ending with a final extension step at 72°C for 10 min. After PCR amplification, 5 μ l were removed and subjected to 2% agarose gel electrophoresis to estimate the size of the amplification products by comparison with a 100-bp molecular size standard ladder (Roche Diagnostics, Mannheim, Germany). The gel was stained with ethidium bromide and the amplicons were visualized using a UV light.

The reaction conditions for the multiplex PCR assay were optimized to ensure that all of the target gene sequences were satisfactorily amplified. The primers used in this study differ in annealing temperatures, which increased the possibility of occurrence of unwanted bands originating from non-specific amplification. Therefore, both a "hot-start" and three rounds of amplification with different annealing temperatures were carried out. In addition, there is evidence indicating that multiplex PCR with targets that differ widely in size often favors amplification of the shorter target over the longer one, resulting in different amounts of amplified products [21]. For this reason, in order to optimize the conditions for the multiplex PCR analysis, different primer concentrations, template DNA preparations, and MgCl₂ concentrations were assayed. As described above, the final quantities generated optimal results were 3.5 mM MgCl₂, 180 pmol of each *femB* primer, 10 pmol of each *ileS*-2 primer, 5 pmol of each *mecA* primer, 40 pmol of each *icaAB* primer and 5 ml of the DNA solution obtained with our DNA extraction method.

Fig. 1. Agarose gel electrophoresis patterns showing single (lanes 3–7) and multiplex (lanes 8–10) PCR results: lane 3, *icaAB* amplicon from *Staphylococcus aureus* isolate 229; lane 4, *femB* amplicon from *S. aureus* isolate 229; lane 5, *icaAB* product from *Staphylococcus epidermidis* isolate 9951-79; lane 6 *ileS-2* amplicon from *S. epidermidis* isolate 9295-79; lane 7, *mecA* amplicon from *S. epidermidis* isolate 9295-79; lane 8, *icaAB*, *ileS-2* and *mecA* amplicons from a *S. epidermidis* isolate 1855-25; lane 9, *femB*, *icaAB*, *ileS-2* and *mecA* amplicons from *S. aureus* isolate 229; lane 10, *femB*, *icaAB*, *ileS-2*, *mecA* and both *icaAB* from a mixture of *S. aureus* and *S. epidermidis* isolates 229 and 9951-79, respectively, simultaneously amplified. Lane 1, DNA molecular size marker (100-bp ladder). No bands were present in the negative control (lane 2). A schematic representation of the fragments amplified is shown on the left.



Results and Discussion

Prior to optimizing the multiple PCR, the PCR protocol was evaluated in order to ensure that it was adequate for the individual amplification of all five DNA fragments. Each individual amplification yielded a fragment of the expected size, i.e. 750-, 651-, 546-, 456-, and 310-bp, respectively. Figure 1 shows an agarose gel that illustrates results typically obtained with the optimized PCR assay. Amplification of *icaAB*, *femB*, *ileS-2* and *mecA* targets produced distinct, easily recognizable bands corresponding to their respective molecular size (lanes 3–7). The *icaAB* fragments were clearly differentiated when amplified from *S. aureus* or from *S. epidermidis* (lanes 8–10); *femB* was always amplified in the case of *S. aureus* strains (lane 9) and never in the case of *S. epidermidis* (lane 8). Results obtained for the *mecA* and *ileS-2* fragments were in accordance with the resistance pattern of the isolates: methicillin- and mupirocin-resistant isolates *S. epidermidis* 1855-25 and *S. aureus* 229 presented the *mecA* fragment and showed the *ileS-2* marker (lanes 8–10). PCR from mixed *S. aureus* 229 and *S. epidermidis* 9951-79 colonies permitted the simultaneous detection of the five different targets (lane 10).

This protocol, including the rapid extraction of DNA from single colonies and electrophoretic analysis of the amplified products on an agarose gel, was completed in

less than 6 h. Moreover, it can be used for *S. aureus* and *S. epidermidis* and for a mixture of both staphylococci.

Nowadays, since few antibiotics, including mupirocin, constitute the last resort against MRSA, and due to the increasing incidence of catheter-associated staphylococcal bloodstream infections, a fast, sensitive laboratory method to immediately detect multiple-resistant staphylococci harboring the *ica* cluster is urgently needed. Although most *S. aureus* carry the *ica* cluster, confirmation of its presence in a particular isolate is a necessary step in preventing colonization by potentially biofilm-forming strains. The method described herein is highly sensitive and specific, fast and feasible, and thus provides a new tool for controlling catheter-borne infections.

Acknowledgements. This work was supported by grants 2001/020 from the *Consejería de Educación, Cultura y Deportes*, Canarian Autonomous Government, FIS PI 01/3150 from the Health Spanish Ministry, and Bio 2002/00953 from the Spanish Ministry for Science and Technology. S.M.A. was supported by FIS contract 99/3060 (Fondo de Investigación Sanitaria, Spain).

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Detección simultánea del cluster *ica* y de los genes de resistencia a meticilina y mupiricina en *Staphylococcus* aislados de catéteres

Resumen. Estudios recientes han demostrado que más del 50% de las septicemias asociadas al uso de catéteres están causadas por estafilococos. Las infecciones estafilocólicas producidas por cepas portadoras del operón de adhesión intercelular (*ica*) pueden agravarse por la presencia de genes de resistencia a la meticilina y/o a la mupirocina. Hemos desarrollado un protocolo de PCR múltiple que permite, simultáneamente, identificar estafilococos y detectar la presencia de *ica* y de los genes de resistencia a la meticilina y/o a la mupirocina. Además, dicho método permite la detección diferencial del locus *ica* de *Staphylococcus aureus* y/o *S. epidermidis*. [*Int Microbiol* 2004; 7(1):63–66]

Palabras clave: *Staphylococcus aureus* · *Staphylococcus epidermidis* · cluster génico de adhesión intercelular · resistencia a antibióticos · PCR múltiple

Detecção simultânea do cluster *ica* e dos genes de resistência a meticilina e mupiricina em *Staphylococcus* isolados de catéteres

Resumo. Estudos recentes têm demonstrado que mais de 50% das septicemias associadas ao uso de catéteres são causadas por estafilococos. As infecções estafilocólicas produzidas por cepas portadoras de operón de adesão intercelular (*ica*) podem agravar-se pela presença de genes de resistência a meticilina e/ou a mupirocina. Foi desenvolvido um protocolo de PCR múltiplo que permite, simultaneamente, identificar estafilococos e detectar a presença de *ica* e dos genes de resistência a meticilina e/ou a mupirocina. Este método permite a detecção diferencial do locus *ica* de *Staphylococcus aureus* e/ou de *S. epidermidis*. [*Int Microbiol* 2004; 7(1):63–66]

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