Methicillin Resistance Transfer from *Staphyloccocus epidermidis* to Methicillin-Susceptible *Staphylococcus aureus* in a Patient during Antibiotic Therapy

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**Abstract**

**Background:** The *mecA* gene, encoding methicillin resistance in staphylococci, is located on a mobile genetic element called Staphyloccocal Cassette Chromosome *mec* (SCCmec). Horizontal, interspecies transfer of this element could be an important factor in the dissemination of methicillin-resistant *S. aureus* (MRSA). Previously, we reported the isolation of a closely related methicillin-susceptible *Staphyloccocus aureus* (MSSA), MRSA and potential SCCmec donor *Staphylococcus epidermidis* isolate from the same patient. Based on fingerprint techniques we hypothesized that the *S. epidermidis* had transferred SCCmec to the MSSA to become MRSA. The aim of this study was to show that these isolates form an isogenic pair and that interspecies horizontal SCCmec transfer occurred.

**Methodology/Results:** Whole genome sequencing of both isolates was performed and for the MSSA gaps were closed by conventional sequencing. The SCCmec of the *S. epidermidis* was also sequenced by conventional methods. The results show no difference in nucleotide sequence between the two isolates except for the presence of SCCmec in the MRSA. The SCCmec of the *S. epidermidis* and the MRSA are identical except for a single nucleotide in the *ccrB* gene, which results in a valine to alanine substitution. The main difference with the closely related EMRSA-16 is the presence of SaPI2 encoding toxic shock syndrome toxin and exfoliative toxin A in the MSSA-MRSA pair. No transfer of SCCmec from the *S. epidermidis* to the MSSA could be demonstrated in vitro.

**Conclusion:** The MSSA and MRSA form an isogenic pair except for SCCmec. This strongly supports our hypothesis that the MRSA was derived from the MSSA by interspecies horizontal transfer of SCCmec from *S. epidermidis*.

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**Introduction**

Methicillin-resistant *Staphyloccocus aureus* (MRSA) is a major cause of nosocomial and community-acquired infections. Resistance to methicillin is a result of the expression of penicillin-binding protein 2a (PBP2a), which causes a low affinity for β-lactam antibiotics. PBP2a is encoded by the *mecA* gene, which is located on a mobile genetic element called Staphyloccocal Cassette Chromosome *mec* (SCCmec) [1]. To date, several types of SCCmec have been described which were recently reclassified [2].

Horizontal transfer of mobile genetic elements contributes to the worldwide dissemination of (multi)-resistant pathogens. Leverstein-van Hall et al. described a high rate of interspecies transfer of resistance genes among multidrug resistant Enterobacteriaceae in patients [3]. Berglund et al. described the likely transfer of a type V SCCmec from methicillin resistant *Staphyloccocus haemolyticus* to *S. aureus* in a neonatal intensive care unit [4]. Although this interspecies transfer of SCCmec could contribute profoundly to the dissemination of MRSA, evidence is still inconclusive.

Previously, we described the isolation of a successive pair of *mecA* and *mecA*+ *S. aureus* and a methicillin resistant *Staphyloccocus epidermidis* isolate, which was likely the donor of the SCCmec, from a male infant suffering Pierre Robin syndrome [5]. The patient was intubated and mechanically ventilated 4 days after birth because of respiratory insufficiency. A suspected respiratory tract infection was treated with amoxicillin/clavulanic acid. As he became bacteraemic after 3 days the antibiotic regimen was changed to amoxicillin and cefotaxime. When his blood cultures grew *mecA*+ methicillin-susceptible *S. aureus* (MSSA) treatment was changed to flucloxacillin. He seemed to recover, but amoxicillin/clavulanic acid treatment was reinstated for 10 more days on day 32 after the respiratory tract infection had recurred. On day 56, routine cultures from nasal swabs unexpectedly showed a *mecA*+ MRSA. Several strains of *mecA*+ coagulase-negative staphyloccoci (CNS) were also identified. Pulsed-Field Gel Electrophoresis (PFGE) of the isolates using *Smal* showed a 40 kb shift of a single DNA fragment between the MSSA and MRSA. Southern hybridization showed that the larger DNA fragment contained the *mecA* gene. The MRSA and MSSA had identical ribotypes and...
Results

For MSSA WKZ-1 a total of 45 contigs larger than 1000 nucleotides was obtained with an average read size of 244 nucleotides and a 23.5-fold coverage, whereas for WKZ-2 a total of 136 contigs larger than 1000 nucleotides were obtained with an average read size of 187 nucleotides and a 9.9-fold coverage. The isolates belong to Sequence Type (ST) 30. MRSA252, which is an EMRSA-16 and has ST36 and belongs to the same Clonal Complex (CC30) was the most closely related strain from which the whole genome was sequenced. Therefore, the sequence of this strain was used as a scaffold to determine the order and orientation of the contigs for further DNA sequencing of the gaps between the contigs. Except for SCCmec no difference in the mobile genetic elements of WKZ-1 and WKZ-2 was found.

The WKZ isolates differ from MRSA252 in a number of mobile genetic elements. The WKZ-2 MRSA contains a SCCmec type IV, whereas MRSA252 carries a SCCmec type II of 58.8 kb which includes pUB110 encoding bloemycin and kanamycin resistance and Tn554 encoding resistance to erythromycin and streptomycin. At the position of the Staphylococcus aureus Pathogenicity Island 4 (SaPI4) in MRSA252 the WKZ isolates carries a slightly different SaPI (bp 356,242-372,489). It shows differences at the presence or absence of open reading frames (orfs) and similarity between homologous genes (Fig. 1). The function of these proteins is unknown, but a BLAST against the GenBank database shows homology with pathogenicity island proteins. In addition, SaPI2 is present in the WKZ isolates (bp 2,116,959-2,131,693) which encodes toxic shock syndrome toxin and exfoliative toxin A. The genomic island vSaX is identical for the WKZ isolates and MRSA252 with two exceptions. One exotoxin gene shows 3 nucleotide deletions leading to a slightly different C-terminal amino acid sequence for this protein. The second dissimilarity, which is in a protein part of a restriction modification system, contains 20 point-mutations which lead to 5 amino acid changes. In addition, the restriction modification system present in genomic island vSaβ of MRSA252 is not functional in the WKZ isolates as the gene encoding the methylase is truncated. Finally, an additional gene for a SD-repeat protein (sdrD) is present in the WKZ isolates but not in MRSA252. Both MRSA252 and the WKZ isolates contain the sdrC and bhp gene, which are also SD-repeat proteins. The latter gene encodes bone sialoprotein binding protein.

The WKZ isolates differs in 423 single nucleotide polymorphisms or SNPs from MRSA252 (181 missense mutations, 97 silent mutations 85 mutations at intergenic regions, and 60 indels) in the core genome of S. aureus. The number of SNPs in the variable part of the S. aureus shared by the 2 isolates was much higher with 1001 SNPs (558 missense mutations, 344 silent mutations 22 mutations at intergenic regions, and 77 indels). Based on these data the dN/dS ratio for the core is approximately 0.6. The close relationship between the WKZ and MRSA252 isolates is further confirmed by a phylogenetic analysis based on the sequence of WKZ-1 and 17 other published complete genomes. In this analysis WKZ-1 and MRSA252 cluster together (Fig. 2).

No differences were observed in the sequences of WKZ-1 and WKZ-2, except for the presence of SCCmec in the latter isolate. The isolates can therefore be considered isogenic. The SCCmec of the proposed donor S. epidermidis isolate O7.1 belonged to type IVa. However, the SCCmec of WKZ-2 contains an additional sequence that is homologous to the 6987 bp cassette chromosome(SCC)-like element of strain ATCC25923 [6]. This element was already present in WKZ-1. The SCCmec IVa sequence of S. epidermidis O7.1 is identical to that of the WKZ-2 isolate except for a single point-mutation that resulted in a valine to alanine amino acid substitution in ccrB, which encodes a recombinase involved in the excision and integration of SCCmec. The lack of differences between WKZ-1 and WKZ-2 and the presence of only one point-mutation between the SCCmec elements of the proposed donor S. epidermidis and WKZ-2 strongly support our hypothesis that WKZ-2 was derived from WKZ-1 by transfer of SCCmec from S. epidermidis O7.1.

Resistance to antibiotics was studied. S. epidermidis O7.1 was resistant to cefoxitin, ampicillin, penicillin, amoxicillin/clavulana- nate, and trimethoprim/sulfametoxazol. MSSA WKZ-1 was not resistant to any of the tested antibiotics. WKZ-2 MRSA was resistant only to oxacillin.
Inactivation of a restriction modification enzyme in possible cause for the ease of the acceptance of SCC elements is also be excised from the supposed alternative site was also noted previously [7]. The element could elements either independently or together, but excision at an occasion. This strain has the ability to excise these SCC encoded recombinase.

Discussion

We previously reported the isolation of a closely related MSSA and MRSA and potential SCCmec donor S. epidermidis isolate from the same patient. Based on fingerprint techniques we hypothesized that the S. epidermidis had transferred SCCmec to the WKZ-1 MSSA to become the WKZ-2 MRSA [5]. In the present study, we performed whole genome sequencing on both isolates. The results show no difference in nucleotide sequence between the two WKZ isolates except for the presence of SCCmec in WKZ-2. A previous study had shown that SCCmec of WKZ-2 was bordered by the SCC-like element of S. aureus strain ATCC25923, whereas this element was lacking in the S. epidermidis strain. The SCC-like element may encode 2 transposases and 3 hypothetical proteins. Sequencing of the SCCmec of the S. epidermidis showed that it was identical to the SCCmec of WKZ-2 except for a single point-mutation. The SCC-like element that is also present in WKZ-2 was also previously shown to be present in WKZ-1 [7]. In the current study we show that the sequences of the SCC-like element are identical in both isolates, except for a single point-mutation in the ccrB gene which leads to a valine to alanine substitution in the encoded recombinase.

Apparently, the WKZ strain was able to accept SCC elements on 2 occasions. This strain has the ability to excise these SCC elements either independently or together, but excision at an alternative site was also noted previously [7]. The element could also be excised from the supposed S. epidermidis donor [7]. A possible cause for the ease of the acceptance of SCC elements is the inactivation of a restriction modification enzyme in VSaβ by

The WKZ-2 is likely to be more pathogenic than MRSA252, which is a representative of EMRSA-16, as WKZ-2 contains a number of additional virulence factors. Most notably, the WKZ-2 strain contains SaPI2, which encodes both toxic shock syndrome toxin and exfoliative toxin. Especially toxic shock syndrome toxin has been associated with severe disease [11], whereas exfoliative toxin A, which acts as a specific protease has been associated with skin damage [12]. The acquisition of SaPI may also be the result of the impaired restriction modifications systems. So, impaired restriction modification may lead to the emergence of novel antibiotic-resistant strains, but also more pathogenic strains. In addition, the sbd gene is present in WKZ-2 which encodes a putative adhesion protein. This may play a role in tissue specificity of the strain. A difference in a VSaβ encoded exotoxin was also observed.

The relatedness of WKZ-1 (ST30) and MRSA252 (ST36) is demonstrated by the relatively low number of SNPs (423 in the S. aureus core genome and 1001 in the shared variable genome). In addition, the phylogenetic tree based on the complete genomes of 18 S. aureus strains, shows the close relatedness of MRSA252 and the WKZ isolates. The interpretation of the dN/dS ratio of 0.6 is not possible, because it may indicate negative or positive selection or no selection at all in intraspecies analysis of selective pressure [13]. To establish whether particular genes or operons were under selective pressure the presence of multiple mutations in single genes or operons were analyzed. Eight genes and one operon [14] (Table 1) were identified as having more than mutation had occurred. Only of one the genes had only silent mutations. The mutations in DNA topoisomerase I can be explained by differences in fluoroquinolone susceptibility. The very large surface anchored protein is most likely exposed on the outside of the bacterium and might be sensitive to immune pressure. The only operon is involved in the synthesis of the S. aureus pigment, a known virulence factor [15,16], how mutations affect the synthesis in different hosts is unknown. The mutations in the genes for DNA polymerase I, a possible formate dehydrogenase, and a natural resistance associated protein family member are not readily explained, but may represent chance mutations that have not been purified from the strain through selection [13].

In conclusion, the data presented strongly support our hypothesis that interspecies transfer of a SCCmec took place, in a patient undergoing antibiotic therapy, from S. epidermidis O7.1 to MSSA WKZ-1, to become MRSA WKZ-2. Our hypothesis is that MSSA WKZ-1 is receptive for mobile genetic elements, due to an
Table 1. Genes and operons with multiple SNPs between WKZ-1 and MRSA252.

| potential function                        | type of mutation | bp position in MRSA252 |
|-------------------------------------------|------------------|------------------------|
| natural resistance associated             | N                | 1,125,728              |
| macrophage protein family protein         | N                | 1,125,731              |
|                                            | N                | 1,125,792              |
|                                            | N                | 1,125,806              |
| hypothetical protein                       | S                | 1,126,585              |
|                                            | N                | 1,126,662              |
| DNA topoisomerase I                        | S                | 1,279,226              |
| very large surface anchored protein        | N                | 1,503,515              |
|                                            | N                | 1,505,090              |
|                                            | S                | 1,508,470              |
|                                            | S                | 1,514,431              |
|                                            | N                | 1,521,422              |
|                                            | S                | 1,529,950              |
| DNA polymerase I                           | S                | 1,832,748              |
|                                            | N                | 1,834,661              |
| putative formate dehydrogenase             | S                | 2,462,696              |
|                                            | N                | 2,463,997              |
| squalene synthetase                        | N                | 2,731,700              |
| squalene desaturase                        | S                | 2,732,542              |
| putative glycosyl transeferase             | N                | 2,733,561              |

N is non-synomous, S is synomous.

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Materials and Methods

Ethics statement

An ethics statement is not required for this work.

DNA sequencing

The MSSA and MRSA isolates WKZ-1 and WKZ-2 were sequenced by KeyGene (Wageningen, The Netherlands) using 454 sequencing technology. For WKZ-1 the gaps between the contigs with the exception of the rRNA operons were closed by PCR amplification of the missing fragments followed by conventional DNA sequencing. MRSA252 was used as a scaffold. Annotation was performed by JCVI. The sequencing of WKZ-1 has been reported in GenBank under project ID 40253. The SCMc mec sequence is accessible under no. GQ918137.

Single-nucleotide polymorphism (SNP) analysis was performed using NUCMER [http://mummer.sourceforge.net/] followed by confirmation using PCR amplification and conventional DNA sequencing.

The SCMc mec of the S. epidermidis was sequenced by conventional sequencing of PCR products. PCR amplifications were based on the sequence of the SCMc mec of WKZ-2. Conventional sequencing was performed by BaseClear (Leiden, The Netherlands).

In vitro SCMc mec transfer

To select S. aureus that may have obtained SCMc mec from the S. epidermidis donor strains during in vitro mating experiments the MSSA was made resistant to linezolid by serial transfer, because the S. epidermidis isolate was resistant to commonly used antibiotics. Bacteria were mated on nitrocellulose filters. Mating experiments without antibiotics were performed using TSA 5% sheep blood plates (BD Diagnostic Systems, USA). Mating experiments under antibiotic pressure were performed using cloxacillin at concentrations of 0.5, 0.25 and 0.125 μg/mL and cefotaxime at concentrations of 1.0, 0.5, 0.25 μg/mL in LB agar. Conjugants were selected using both linezolid and cloxacillin at 4.0 μg/mL each.

Antibiograms

To assess antimicrobial resistance of the WKZ-1 MSSA and the S. epidermidis 07.1 isolates, the minimal inhibitory concentration (MIC) were determined for a number of clinically relevant antibiotics.

Author Contributions

Conceived and designed the experiments: ALAB ACF. Performed the experiments: ECB. Analyzed the data: ALAB ECB ACF. Wrote the paper: ALAB ACF.

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