Genome Reports



Complete Genome Sequence of a Methicillin-Resistant *Staphylococcus aureus* Sequence Type 72 Strain SA520 Isolated from Korean Hospital

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Once known as a major community-genotype, sequence type (ST) 72 methicillin-resistant *Staphylococcus aureus* (MRSA) strains have been increasingly identified in hospital-associated infections in Korea. Here, we report the complete genome sequence of SA520 isolated from a patient in Korea.

Keywords: Methicillin-resistant Staphylococcus aureus, sequence type 72, SA520

Community-associated methicillin-resistant *Staph*ylococcus aureus (CA-MRSA) usually carrying type IV or V staphylococcal cassette chromosome *mec* (SCC*mec*) has emerged in global communities [1]. Although distinctive lineages of CA-MRSA have been found in different countries [2], a unique sequence type (ST) 72 MRSA-SCC*mec* IV has become a major cause of infection in both community and hospital settings in Korea [3, 4]. We report here the complete genome sequence of a ST72 MRSA-SCC*mec* IV strain, SA520, isolated in Korea.

S. aureus SA520 was isolated from a blood sample form a patient with bacteremia hospitalized in Seoul National University Bundang Hospital in Korea [5], and identified using 16S ribosomal RNA gene sequencing (Bionics, Seoul, Korea). For whole genome sequencing, genomic DNA was extracted by using a Genmed DNA kit (Korea). Sequencing libraries were constructed by hybrid sequencing using an Oxford Nanopore MinION

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(Oxford Nanopore Technologies, UK) and Illumina iSeq platforms (Illumina, USA). *De novo* assembly of Unicycler v0.4.8 was employed to generate high-quality complete genome. The assembled genome was annotated using both Prokka v1.14.6 and Rapid Annotation using Subsystem Technology (RAST) v.2.0.

The total length of the completed genome was 2,765,395 bp with $607 \times$ genome coverages, which comprised of one large circular chromosome of 2,762,063 bp (G+C content, 32.84%) and a plasmid (pSA520) of 3,332 bp (G+C content, 29.53%). The complete genome sequences of *S. aureus* SA520 and a 3-kb plasmid pSA520 have been deposited in the GenBank under the accession numbers CP101312 and CP101313, respectively. The sequence data were compared with a previously reported sequences of ST72 CA-MRSA CN1 strain resulting in a 99.93% average nucleotide identity (ANI) value [6].

In silico genotyping was performed using Center for Genomic Epidemiology (CGE) software for multilocus sequence typing, as well as SCC*mec*, *spa*, and *agr* typing. Antimicrobial resistance genes (ARGs) and virulence-

Strain	SA520	CN1ª
Genome size (bp)	2,765,395	2,757,070
G+C content (%)	32.8	32.79
Coverage	607	40-60
Plasmid	pSA520	pHL1 and pHL2
MLST	ST72	ST72
SCCmec	IV	IV
spa type	t13921	t324
<i>agr</i> type	I	I
ARGs	mecA, blaZ	aadD, blaZ, bleO, mecA
Virulence genes	Gamma-hemolysin (<i>hlgA, hlgB, hlgC</i>), Leukocidins (<i>lukD, lukE</i>), IEC (<i>sak, chp, scn</i>), SEs (<i>seg, sei, sem, sen, seo, seu</i>), Aureolysin (<i>aur</i>), Serine protease (<i>splA, splB</i>)	Gamma-hemolysin (<i>hlgA, hlgB, hlgC</i>), Leukocidins (<i>lukD, lukE</i>), IEC (<i>sak, chp</i>), SEs (<i>seg, sei, sem, sen, seo, seu</i>), Aureolysin (<i>aur</i>), Serine protease (<i>splA, splB, splC, splD</i>)

Table 1. Genetic characteristics of ST72 CA-MRSA SA520 and CN1 strains.

IEC, Immune evasion cluster; SE, Staphylococcal enterotoxin ^aGenBank accession number: CP003979.1

associated genes were identified by ResFinder (https:// cge.food.dtu.dk/services/ResFinder/) of CGE. Standard antimicrobial susceptibility assay [7] revealed that SA520 strain displayed resistance to β -lactams (ampicillin, cefoxitin, and penicillin). SA520 MRSA strain was identified as ST72 clonal lineage carrying SCCmec IV with spa type t13921 and agr type I. ResFinder analysis indicated two ARGs corresponding to the resistance phenotypes: mecA, and blaZ. Moreover, virulence genes encoding gamma-hemolysin components, leukocidins, immune evasion clusters (IECs), staphylococcal enterotoxins (SEs), aureolysin, and serine protease were identified in SA520 (Table 1).

Acknowledgments

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Conflict of Interest

The authors have no financial conflicts of interest to declare.

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