

Antimicrobial resistance, virulence genes profiling and molecular relatedness of methicillin-resistant *Staphylococcus aureus* strains isolated from hospitalized patients in Guangdong Province, China

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Purpose: The main objective of this study was to decipher the prevalence, antimicrobial resistance, major virulence genes and the molecular characteristics of methicillin-resistant *Staphylococcus aureus* (MRSA) isolated from different clinical sources in southern China.

Materials and methods: The present study was performed on 187 non-duplicate *S. aureus* clinical isolates collected from three tertiary hospitals in Guangdong Province, China, 2010–2016. Antimicrobial susceptibility testing was performed by the disk diffusion method and by measuring the minimum inhibitory concentration. Screening for resistance and virulence genes was performed. Clonal relatedness was determined using various molecular typing methods such as multilocus sequence typing, *spa* and staphylococcal chromosomal cassette *mec* (SCC*mec*) typing. Whole genome sequencing was performed for three selected isolates.

Results: Out of 187 isolates, 103 (55%) were identified as MRSA. The highest prevalence rate was found among the skin and soft tissue infection (SSTI) samples (58/103), followed by sputum samples (25/103), blood stream infection samples (15/103) and others (5/103). Antimicrobial susceptibility results revealed high resistance rates for erythromycin (64.1%), clindamycin (48.5%), gentamicin (36.9%) and ciprofloxacin (33.98%). All isolates were susceptible to vancomycin. Resistance genes and mutation detected were as follows: *aac(6′)-aph(2′′)* (24.3%), *dfpG* (10.7%), *rpoB* (21.4%), *efr* (0%), *fexA* (1.94%), *gyrA* (35.92%), *gyrB* (0.97%), *glaA* (20.4%), *glaB* (10.68%), *ermA* (21.4%), *ermB* (18.44%), *ermC* (21.4%) and *lnuA* (18.44%). Profiling of virulence genes revealed the following: *sea* (11.7%), *seb* (21.4%), *sec* (0.97%), *sed* (0.97%), *hla* (86.41%), *hly* (17.48%), *hlg* (10.68%), *hld* (53.4%), *Tsst-1* (3.9%) and *pvl* (27.2%). Clonal relatedness showed that ST239-SCC*mecA* III-t37 clone was the most prevalent clone.

Conclusion: Our study elucidated the prevalence, antibiotic resistance, pathogenicity and molecular characteristics of MRSA isolated from various clinical sources in Guangdong, China. We found that the infectious rate of MRSA was higher among SSTI than other sources. The most predominant genotype was ST239-SCC*mecA* III-t37 clone, indicating that ST239-t30 clone which was previously predominant had been replaced by a new clone.

Keywords: MRSA, antibiotic resistance, resistance genes, virulence factors, molecular typing, whole genome sequencing

Introduction

Staphylococcus aureus is a highly virulent opportunistic pathogen capable of causing a variety of infections, including skin and soft tissue infection (SSTI), blood stream infection (BSI) and toxin-mediated syndromes as well as life-threatening diseases.^{1–3} *S. aureus*, as one of the “ESKAPE” organisms, is a growing threat worldwide, as it can cause a variety of serious nosocomial infections.⁴ Infections caused by *S. aureus* are divided into two types: methicillin-sensitive *S. aureus* (MSSA) and methicillin-resistant *Staphylococcus aureus* (MRSA), which is also known as oxacillin-resistant *S. aureus*.⁵

Recently, MRSA strains have become responsible for about 25%–50% of clinical infections caused by *S. aureus* in many countries.⁶ Commonly, the MSSA becomes MRSA via acquisition of large and potentially transmissible genomic islands called staphylococcal chromosomal cassette *mec* (SCC*mec*), which harbor *mecA/mecC* genes. Awareness of the different antibiotic resistance patterns and the molecular characteristics of MRSA can effectively help in the treatment of MRSA infections as well as slowing the epidemic spread of it. MRSA has acquired multiple resistance to a wide range of antibiotics including aminoglycosides. Practically, MRSA was found to be resistant to all available β -lactam antibiotics.^{3,7}

Several reports about the prevalence, molecular characterization and antimicrobial resistance pattern of MRSA in China are available since China represents a country of high prevalence rate of MRSA, especially in Guangdong Province, one of the largest southern coastal provinces which has a large population and is adjoining Hong Kong and Taiwan. However, a large proportion of these reports are mainly focusing on a single source of MRSA infection, ignoring the possibility of MRSA infection from multiple sources. Therefore, our study aims to focus on MRSA isolates collected from different clinical sources.^{8–16}

Materials and methods

Clinical isolates

This cross-sectional study was conducted on a total of 187 non-repetitive *S. aureus* isolates collected from different clinical specimens in three tertiary hospitals in Guangdong Province, China, denoted as H1, H2 and H3, in the year 2010–2016. All the MRSA isolates were collected from the routine laboratory work of these hospitals. Specimens were collected from different wards including medical ward, surgical ward, intensive care unit and pediatrics department and were recovered from different clinical sources including SSTI, blood, sputum and others (inflamed joint fluid and

ascitic fluid). The isolates were identified as *S. aureus* by using the BD Phoenix™100 microbial identification system (bioMérieux, Marcy l’Etoile, France).

For detection of MRSA, all the identified *S. aureus* were tested for their antimicrobial susceptibility by disk diffusion technique using cefoxitin (FOX; 30 μ g) according to the Clinical and Laboratory Standards Institute (CLSI, 2017) guidelines. Then *mecA* gene was detected by PCR as previously described.¹⁷ The samples positive for the *mecA* gene and resistant to FOX were identified as MRSA.

DNA extraction

All the samples were cultured on Mueller–Hinton agar plates at 33°C overnight. Then, a single bacterial colony was suspended onto 3 mL of sterile lysogeny broth medium (Oxoid, Hampshire, UK) and subsequently incubated at 33°C with vigorous shaking for 8 hours. Then, DNA was extracted using Hipure bacterial DNA kit (Magen, Guangzhou, China) according to the manufacturer’s instructions.

Antimicrobial susceptibility and minimum inhibitory concentration (MIC) determination

In vitro antimicrobial susceptibility of MRSA isolates was determined by modified Kirby–Bauer disk diffusion method according to the CLSI 2017 guidelines. Antimicrobial drugs tested included: penicillin (P), FOX, rifampicin (RFD), quinupristin/dalfopristin (Q/D) and trimethoprim–sulfamethoxazole (SXT). MIC was detected by broth microdilution method and interpreted according to the CLSI 2017 guidelines for erythromycin (ERY), clindamycin (CLI), gentamicin (GEN), ciprofloxacin (CIP), chloramphenicol (CHL), nitrofurantoin (NIT) and vancomycin (VAN). Tigecycline (TGC) was evaluated using MICs according to the European Committee on Susceptibility Testing guidelines (version 1.3, 2010; Basel, Switzerland).

Molecular detection of antibiotic resistance genes

Isolates were screened for the production of the common resistance genes and mutation for different antimicrobial classes by PCR as follows: RFD (*rpoB*), CHL (*cfp* and *flexA*), aminoglycosides (*acc(6′)-aph(2′′)*), macrolides (*ermA*, *ermB* and *ermC*), lincosamides (*lnuA*), SXT (*dfpG*) and fluoroquinolones (*gyrA*, *gyrB*, *grlA* and *grlB*).^{14,18–22} Purified positive PCR products were sequenced using BigDye terminator chemistry on an automated ABI 3130 sequencer (PE Applied Biosystems, Foster City, CA, USA) based on Sanger’s sequencing method. Gene sequences were corrected

using CodonCode Aligner version 7.1.2. sequencing analysis software and then analyzed using BlastN and BlastP against the National Center for Biotechnology Information database (www.ncbi.nlm.nih.gov).

Detection of virulence factors

Detection of virulence genes was performed among MRSA isolates. Ten exotoxins coding gene, including four enterotoxins (*sea*, *seb*, *sec* and *sed*), four hemolysin toxins (*hla*, *hlb*, *hld* and *hlg*), toxic shock syndrome toxin (*tsst-1*) and Panton–Valentine leukocidin (*pvl*) were amplified by multiplex PCR as previously described.²

Molecular typing of MRSA

All the MRSA strains were typed by three different molecular characterization methods as follows: SCCmec, *spa* type and multilocus sequence typing (MLST).

In the SCCmec typing, MRSA strains were identified as SCCmec type I, II, III, IV or V by performing PCR using nine pairs of primers and the primers for the *mecA* gene as previously described.¹⁷ The SCCmec types were determined on the basis of the band pattern obtained.

Spa typing was performed according to the website <https://spa.ridom.de/>. The isolates were assigned to the specific *spa* types according to the guidelines described by the publicly available Center of Genomic Epidemiology (<https://cge.cbs.dtu.dk/services/spaTyper-1.0/>).²³

MLST was performed based on seven housekeeping genes (*arcC*, *aroE*, *glpF*, *gmK*, *pta*, *tpiA* and *yqiL*) obtained from the MLST database as previously described.²⁴ Isolates were assigned sequence types (STs) according to the MLST database (<http://saureus.mlst.net>).

Whole genome sequencing (WGS)

WGS was performed for three selected isolates, QY152, L2316 and L2317. QY152 was selected for its relatedness to ST398, which is not common in China, but common in Europe. Both L2316 and L2317 were selected for their TGC resistance (the only two TGC-resistant isolates) as well as their relatedness to ST9. The ST398 isolate (SO385, GenBank: AM990992) was used as a reference strain for comparative genomics.

The resulting draft genomes were described. DNA libraries were constructed with 350 bp paired-end fragments. The isolate reads were produced by Illumina HiSeq2000 platform. Reads were assembled using SPAdes. The genome annotation was performed using PROKKA, E-value setting as $1e^{-5}$ ²⁵ and Rapid Annotations Subsystems Technology. Bacte-

riophages detection was performed using PHAge Search Tool.²⁶ Antimicrobial resistance genes were identified using the Comprehensive Antibiotic Resistance Database (<https://card.mcmaster.ca/>) and the virulence factors were screened using the Virulence Factors Database (<http://www.mgc.ac.cn/VFs/>). This Whole Genome Shotgun project of the three selected isolates has been deposited at GenBank under the following accession numbers: RDQU000000, RDQT000000 and RDQS000000, respectively.

Statistical analysis

Statistical analysis for resistance and virulence genes as well as for the resistance rates to the tested antibiotics among our MRSA was performed using chi-squared test (SPSS version 20). *P*-values <0.01 were considered statistically significant.

Results

MRSA identification

Among 187 identified *S. aureus* samples, 103 (55.1%) were confirmed as MRSA depending on *mecA* gene and resistance to FOX. The majority of these MRSA (58/103, 56.31%) were recovered from SSTI, followed by sputum (25/103, 24.3%), BSI (15/103, 14.6%) and others (5/103, 4.9%), as shown in Table 1.

Antimicrobial susceptibility testing

All MRSA isolates were found to be resistant to penicillin and FOX, while all were susceptible to VAN and NIT. Some of the tested antimicrobials revealed high resistance pattern, including ERY (66/103, 64.1%), CLI (50/103, 48.5%), GEN (38/103, 36.9%) and CIP (35/103, 33.98%), while others presented relatively low resistance rates, including RFD (19/103, 18.5%), CHL (12/103, 11.65%), SXT (11/103, 10.7%), Q/D (9/103, 8.74%) and TGC (2/103, 1.94%). All the isolates were found to be resistant to at least two antibiotics, and 68/103 (66.2%) were multidrug resistant (MDR), exhibiting resistance to three or more antibiotic classes, where 44.12% (30/68) of MDR isolates were recovered from SSTI, 35.3% (24/68) from sputum, 13.2% (9/68) from BSI and 7.4% (5/68) from other sites. The resistance rates of GEN, ERY, CIP, RFD and Q/D were found to be higher in sputum than in SSTI or BSI; however, statistically differences were only found in GEN and CIP ($P < 0.01$, Table 1; Figure 1).

PCR amplification and sequencing of antibiotic resistance genes

Screening for fluoroquinolone resistance-encoding genes revealed the presence of *gyrA* (37/103, 35.92%), *gyrB* (1/103,

Table 1 Prevalence and resistance pattern of MRSA isolates

Clinical sample (N=103)	Hospitals			Resistance pattern (%)			
	H1	H2	H3	P	FOX	DA	SXT
SSTI (58, 56.3%)	11, 18.97%	0	47, 81.03%	58, 100%	5, 100%	5, 8.62%	4, 6.9%
Sputum (25, 24.3%)	24, 96%	1, 4%	0	25, 100%	25, 100%	3, 12%	4, 16%
BSI (15, 14.6%)	2, 13.33%	1, 6.67%	12, 80%	15, 100%	15, 100%	1, 6.67%	3, 20%
Others (5, 4.9%)	3, 60%	0	2, 40%	5, 100%	5, 100%	0	0%
Total	40, 38.83%	2, 1.94%	61, 59.22%	103, 100%	103, 100%	9, 8.74%	11, 10.68%
P-value				–	–	0.927	0.287

Note: ^{a,b}Each subscript letter denotes a subset of group categories whose column proportions do not differ significantly from each other at the 0.01 level.

Abbreviations: BSI, blood stream infection; CHL, chloramphenicol; CIP, ciprofloxacin; CLI, clindamycin; ERY, erythromycin; FOX, ceftioxin; GEN, gentamicin; MDR, multidrug resistance; MRSA, methicillin-resistant *Staphylococcus aureus*; NIT, nitrofurantoin; RFD, rifampicin; SSTI, skin and soft tissue infection; SXT, trimethoprim-sulfamethoxazole; TGC, tigecycline; VAN, vancomycin.

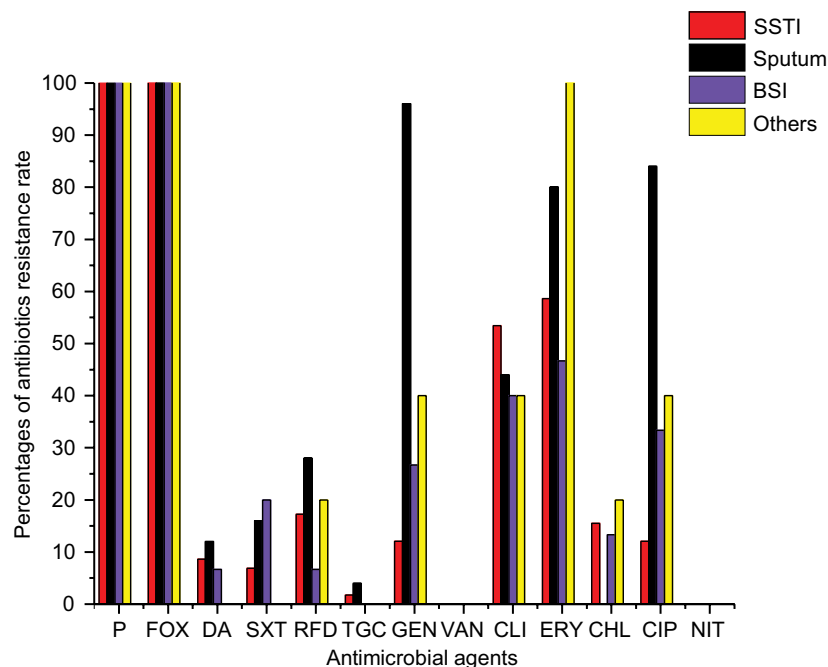


Figure 1 Percentages of antibiotic resistance rates of MRSA isolated from various clinical sources.

Abbreviations: BSI, blood stream infection; CHL, chloramphenicol; CIP, ciprofloxacin; CLI, clindamycin; ERY, erythromycin; FOX, ceftioxin; GEN, gentamicin; MRSA, methicillin-resistant *Staphylococcus aureus*; NIT, nitrofurantoin; P, penicillin; RFD, rifampicin; SSTI, skin and soft tissue infection; SXT, trimethoprim-sulfamethoxazole; TGC, tigecycline; VAN, vancomycin.

Table 2 Resistance pattern and virulence genes of MRSA isolates from different clinical sources

Clinical sample (N=103)	Resistance genes and mutation (%)										
	<i>aac6-aph2</i>	<i>dfrG</i>	<i>rpoB</i>	<i>cfr</i>	<i>fexA</i>	<i>gyrA</i>	<i>gyrB</i>	<i>griA</i>	<i>griB</i>	<i>ermA</i>	<i>ermB</i>
SSTI (58, 56.3%)	4, 6.9% ^a	4, 6.9%	11, 8.97%	0	1, 1.7%	8, 13.8% ^a	0, 0 ^a	5, 8.6% ^a	2, 3.44% ^a	5, 8.6% ^a	13, 22.4% ^{a,b}
Sputum (25, 24.3%)	18, 72% ^b	5, 20%	9, 36%	0	1, 4%	21, 84% ^b	1, 4% ^b	14, 56% ^b	7, 28% ^b	14, 56% ^b	0 ^b
BSI (15, 14.6%)	3, 20% ^a	2, 13.3%	1, 6.7%	0	0	6, 40% ^a	0, 0% ^a	2, 13.3% ^a	2, 13.3% ^{a,b}	2, 13.3% ^a	4, 26.7% ^a
Others (5, 4.9%)	0 ^a	0	1, 20%	0	0	2, 40% ^{a,b}	0, 0% ^{a,b}	0, 0% ^{a,b}	0 ^{a,b}	1, 20% ^{a,b}	2, 40% ^a
Total	25, 24.3%	11, 10.7%	22, 21.4%	0	2, 1.94%	37, 35.92%	1, 0.97%	21, 20.4%	11, 10.68%	22, 21.4%	19, 18.5%
P-value	<0.01	0.275	0.151	–	0.685	<0.01	0.437	<0.01	<0.01	<0.01	0.0102

Note: ^{a,b,c}Each subscript letter denotes a subset of group categories whose column proportions do not differ significantly from each other at the 0.01 level.

Abbreviations: BSI, blood stream infection; MRSA, methicillin-resistant *Staphylococcus aureus*; SSTI, skin and soft tissue infection.

RFD	TGC	GEN	VAN	CLI	ERY	CHL	CIP	NIT	MDR
10, 17.24%	1, 1.72%	8, 12.07% ^a	0	31, 53.45%	34, 58.62%	9, 15.51%	7, 12.07% ^a	0	30, 51.72% ^a
7, 28%	1, 4%	24, 96% ^b	0	11, 44%	20, 80%	0%	21, 84% ^b	0	24, 96% ^b
1, 6.67%	0%	4, 26.67% ^a	0	6, 40%	7, 46.67%	2, 13.33%	5, 33.33% ^a	0	9, 60% ^a
1, 20%	0%	2, 40% ^a	0	2, 40%	5, 100%	1, 20%	2, 40% ^{a,b}	0	5, 100% ^{a,b}
19, 18.45%	2, 1.94%	38, 36.89%	0	50, 48.54%	66, 64.07%	12, 11.65%	35, 33.98%	0	68, 66.02%
0.371	0.685	<0.01	–	0.734	0.039	0.731	<0.01	–	<0.01

1.94%), *grlA* (21/103, 20.4%) and *grlB* (11/103, 10.68%), as shown in Table S1. Macrolide-encoding resistance genes detected were as follows: *ermA* (22/103, 21.4%), *ermB* (19/103, 18.44%) and *ermC* (22/103, 21.4%). *Aac(6′)-aph(2′′)* gene which is correlated with GEN resistance was detected in 25/103 (24.3%) of MRSA. Regarding CHL, *fexA* (2/103, 1.94%) was detected, while no *cfr* was found, indicating the low antimicrobial resistance rate of MRSA to CHL. The *rpoB* gene was detected in 22/103 (21.4%) isolates, *lnuA* gene in 19/103 (18.44%) isolates, while *dfpG* was detected in 11/103 (10.7%) of MRSA isolates (Table 2).

Virulence genes profiling

Multiplex PCR revealed that 97/103 (94.2%) MRSA isolates were harboring virulence genes, including four enterotoxin genes, *sea* (12/103, 11.7%), *seb* (22/103, 21.4%), *sec* (1/103, 0.97%) and *sed* (1/103, 0.97%), and six exotoxin genes, whereas for hemolysin toxins (*hla*, *hly*, *hlg* and *hld*), results revealed that *hla* gene was the highest detected gene that was harbored by 89/103 isolates (86.41%), followed by *hly* (55/103, 53.4%), *hly* (18/103, 17.48%) and *hlg* (11/103, 10.68%). The majority of *hla*-positive strains were recovered from SSTI (44/89, 49.4%) followed by sputum (25/89, 28.1%), BSI (15/89, 16.85%) and others (5/89, 5.62%). Concerning *tsst-1* gene, it was harbored by 4/103 isolates

(3.9%). PCR assay for detection of *pvl* gene showed that 28/103 (27.2%) MRSA were *pvl* positive and the majority of them were isolated from SSTI (25/28, 89.3%; Table 2).

Molecular typing of MRSA

Results of SCCmec typing indicated that five types of SCCmec elements were detected. Type III was the most predominant (39/103, 37.86%) followed by type IV (17/103, 16.5%), wherein the majority of isolates belonged to type IVa (15/17 88.2%) then type IVb (2/17, 11.8%). The prevalence of SCCmec types II and V was very low (2/103, 1.94% and 1/103, 0.97%, respectively), while no isolates belonging to type I were found. Out of 103 MRSA isolates, 44 (42.7%) were classified as non-typable MRSA. The majority of the isolates harboring *hla* gene (37/89, 41.57%) were found to be belonging to SCCmec type III, followed by type IV (12/89, 13.48%), type II (2/89, 2.24%) and type V (1/89, 1.12%), while 36/89 (40.44%) of them were found to be associated with non-typable MRSA. *Tsst-1* gene was associated with SCCmec type II (2/4, 50%). Regarding PVL-positive isolates, 16/28 (57.1%) isolates belonged to the predominant SCCmec type III.

Spa typing revealed a diverse range of *spa* types, wherein 25 types were detected among MRSA. t437 (22/103, 21.4%) was the most predominant type, followed by t37 (18/103,

		Virulence genes (%)									
<i>ermC</i>	<i>lnuA</i>	<i>sea</i>	<i>seb</i>	<i>sec</i>	<i>sed</i>	<i>hla</i>	<i>hly</i>	<i>hld</i>	<i>hlg</i>	<i>Tsst-1</i>	<i>Pvl</i>
12, 20.7%	12, 20.7% ^{a,b}	7, 12.1%	18, 31.03% ^a	0	1, 1.7%	44, 75.9%	9, 15.5%	26, 44.8%	5, 8.6%	2, 3.44%	25, 43.10% ^c
8, 32%	0 ^b	3, 12%	0, 0% ^b	0	0	25, 100%	4, 16%	23, 92%	6, 24%	1, 4%	0 ^a
0	5, 33.3% ^a	1, 6.7%	4, 26.7% ^a	0	0	15, 100%	3, 20%	3, 20%	0	0	1, 6.7% ^{a,b}
2, 40%	2, 40% ^a	1, 20%	0, 0% ^{a,b}	1, 20%	0	5, 100%	2, 40%	3, 60%	0	1, 4%	2, 40% ^{b,c}
22, 21.4%	19, 18.5%	12, 11.7%	22, 21.4%	1, 0.97%	1, 0.97%	89, 86.4%	18, 17.5%	55, 53.4%	11, 10.7%	4, 3.9%	28, 27.2%
0.042	<0.01	0.797	<0.01	0.194	1	<0.01	0.51	<0.01	0.095	0.31	<0.01

17.5%). The remaining *spa* types were distributed homogeneously among MRSA isolates. Also, 17/103 (16.5%) of MRSA isolates were not identified in the *spa* database (Table S2).

According to the MLST typing, MRSA isolates were assigned to 23 different STs; also, 18 novel STs were detected and deposited in the database of PubMLST (<https://pubmlst.org/saureus/>), as shown in Table S2. ST239 was the most prevalent among our isolates (24/103, 23.3%), followed by ST338 (18/103, 17.5%). ST239 was mainly distributed in H1, while ST338 was circulating in H3. It is noteworthy that 65 different clones have been detected where ST239-SCCmecA III-t37 clone was found to be the most predominant genotype circulating in H1 and H2 (9/103, 8.7%), followed by ST338-SCCmecA III-t437 (6/103, 5.83%), which was found to be circulating mainly in H3.

Isolates harboring *hla* gene were related to different clones, while the majority were associated with ST338 (13/89, 14.61%) and t437 (16/89, 17.97%). Of the 28/103 isolates harboring *pvl* gene, 7/28 (25%) isolates belonged to ST338-SCCmecA III-t437 clone. The *tsst-1* gene (4/103) was associated with ST1, ST72, ST641 and ST863 (Figure 2).

Whole genome sequencing

WGS was performed for three selected isolates (QY152, L2316 and L2317). QY152 was found to be harboring six resistance genes and two common virulence factors: hemolysin and adhesin. The *mepA* and *mepR* genes were detected among the TGC-resistant isolates (Table 3).

Discussion

As a clinically important drug-resistance pathogen, MRSA can cause a variety of infections, including SSTI, respiratory tract infection, BSI and others. Herein, we investigated the prevalence, antibiotic resistance, virulence factors and the molecular characteristics of MRSA isolated from different clinical sources as shown in Table 1. The prevalence of MRSA in the present study was high. However, the prevalence among SSTI was much lower than that reported from a burn center in southeastern China in 2014–2015,¹⁶ but was consistent with that reported in a study from Taiwan in 2008.²⁷ The prevalence of our MRSA isolates in BSI was lower than those reported from China in 2016 and Latin America in 2011–2014.^{28,29} The detection rate of MRSA among our sputum samples was much higher than that reported from Dongguan and Shenzhen, two cities of Guangdong Province,^{12,30} but much lower than that reported in a multicenter study in China from 2007 to 2009.³¹

Among our isolates, high drug resistance was observed. More than half of the isolates were MDR, which mostly came from sputum and SSTI and were found related to ST239 (23/24) and ST1016 (7/8). The high MDR problem of ST239-SCCmecA III-t37/t30 clone is still extremely serious. The MDR rate in our study was a little lower than that reported in a previous study in Beijing Children's Hospital, China.³²

We believe that different clinical samples show differences in developing drug resistance, which is attributed to the different selection processes. However, in contrast to our hypothesis, the antimicrobial resistance rate among our MRSA isolates did not show a statistical difference, except for CIP and GEN, where *acc(6')-aph(2'')*, *gyrA*, *gyrB*, *griA* and *griB* were found to be more prevalent among sputum samples ($P < 0.01$). We hypothesized that the reason was that fluoroquinolones were often used in pneumonia, which may increase the resistance rate.^{33,34}

According to the US Food and Drug Administration black box, TGC was a last-line agent for MRSA infections until September 2010.³⁶ In the present study, TGC resistance was very low (2/103, 1.94%), similar to that reported in a study from Zhejiang Province. The rate of TGC resistance in China is still low, but the number of MRSA isolates resistant to TGC is actually growing.³⁵ Here, WGS revealed the occurrence of the efflux protein coding gene *mepA* among the whole genome analyzed sequences, which was found to be related to decreasing the susceptibility to TGC. *MepA* is rarely reported in China, while it is more prevalent in African countries.^{37,38}

Regarding the virulence genes, our study showed relatively similar findings for *sea*, *seb*, *sec*, *sed*, *tsst-1*, *pvl* and *hla* with those reported of a burn center in southeastern China.¹⁶

Isolates recovered from SSTI were associated with >15 types of MLST and *spa* types, which indicated high diversity and complexity in their genetic background. Our results were consistent with those reported in a multicenter study of SSTI in China³⁹ where the predominant *spa* type was t437. In the present study, ST338-t437 was the predominant clone, which is inconsistent with other studies which revealed that ST59-t437 clone was predominant.²⁸ However, it has been reported that both ST338-t437 and ST59-t437 belonged to the same clonal complex (CC59), where ST59 was the predominant ST, while single-locus variants belonging to ST338 were obtained.⁴⁰ Therefore, we could probably suppose that ST239-t30 clone which was predominant in 2013 had been gradually replaced by ST338-t437 or ST59-t437.

Our results were similar to those of a study conducted in Wenzhou, China from 2012 to 2013 in which all the isolates

Main molecular types

- SCCmecA III
- SCCmecA IV
- ST239
- ST338
- ST59
- ST1016
- New ST
- t37
- t437
- t1081

- Antibiotic resistance
- ◁ Resistance pattern
- ☆ Virulence genes

Antibiotic resistance	Resistance pattern	Virulence genes
T R G C E C C D S	r aac l e e e f g g g g d s s s s h h h h t P	
G F E L R H I A X	p (6)-n r r r l y y r r f e e e e l l l l s V	
C D N I Y L P T	B o p h u m m e r r l l r a b c d a b d g s L	
	(2) A A B C x A B A B G	t
	A	-
]



Figure 2 Phylogenetic tree of 103 MRSA isolates.

Notes: A maximum likelihood phylogeny was used of every MRSA isolate. Strain names are color-coded based on the hospital: blue is hospital 1, black is hospital 2 and red is hospital 3. The main molecular types are shown in various colors as shown in the legend. The inner bar indicates the main SCCmecA and the outer bar indicates the main spa types. Main STs are indicated by the middle bar.

Abbreviations: MRSA, methicillin-resistant *Staphylococcus aureus*; SCCmec, staphylococcal chromosomal cassette mec; ST, sequence type.

Table 3 General characteristics of the genomes of QY152, L2316 and L2317 in comparison with SO385

Elements and characteristics	SO385 ^a	QY152	L2316	L2317
Chromosome				
Size (bp)	2,872,582	2,715,979	2,799,472	2,792,378
G + C content (%)	32.9	32.8	32.7	32.7
tRNA gene	59	57	57	57
rRNA operon	6	8	7	6
Protein-coding genes	2,699	2,502	2,659	2,592
Insertion sequences	12	6	5	5
Bacteriophages	2	1	3	2
Plasmids	3	0	0	0
Resistance genes	<i>copA</i> , <i>tet(M)</i> , <i>tet(K)</i> , <i>str</i>	<i>blaZ</i> , <i>ermC</i> , <i>norA</i> , <i>mepA</i> , <i>tet(38)</i> , <i>sav1866</i>	<i>blaZ</i> , <i>ermC</i> , <i>norA</i> , <i>mepA</i> , <i>tet(38)</i> , <i>sav1866</i> , <i>tet(L)</i> , <i>ant(4')-Ib</i> , <i>aac(6')-aph(2'')</i> , <i>fosB</i> , <i>dfgG</i> , <i>fexA</i>	<i>blaZ</i> , <i>ermC</i> , <i>norA</i> , <i>mepA</i> , <i>tet(38)</i> , <i>sav1866</i> , <i>tet(L)</i> , <i>ant(4')-Ib</i> , <i>aac(6')-aph(2'')</i> , <i>fosB</i> , <i>dfgG</i> , <i>fexA</i>
Virulence factors	vSaβ, NWMN0280	Hemolysin, adhesin, staphostatin B and others	Hemolysin, adhesin, aur and others	Hemolysin, adhesin, aur and others

Note: ^aSO385 livestock-associated MRSA (ST398 isolate) from the Netherlands. GenBank: AM990992.

Abbreviation: MRSA, methicillin-resistant *Staphylococcus aureus*.

belonging to CC59 were susceptible to NIT, Q/D and SXT; however, 81.8% of them were resistant to ERY and CLI.⁴¹

Noteworthy, two of our isolates recovered from SSTI at the same hospital were found to be belonging to ST398, which has been known as a livestock-associated MRSA (LA-MRSA) clone.⁴² In previous studies, it was reported that most of the LA-MRSA isolates in Asia belonged to ST9, while ST398-related isolates were mainly reported in European and North American countries.^{43,44} Generally, due to the lack of several common virulence factors, the ST398 MRSA in Europe did not cause the frequent invasive diseases in humans.⁴⁵ However, a recent report indicated that the Chinese LA-MRSA ST398 isolates are far different from those of other continents, because they likely evolved from human-adapted MSSA.⁴⁶ Recently, a report about the emergence of ST398 in a pig farm in Guangdong, China was published,¹³ which indicated the transmission of this clone between different continents. Therefore, we supposed that the two isolates belonging to ST398 may be transmitted via livestock, which is in accordance with other reports that showed the possibility of transmission of ST398 from the animal reservoir to humans.^{28,47,48} Also, since the source of infection was SSTI, we can conclude that the possible way of transmission of MRSA was via skin contact with the livestock, which requires certain precautions.

Among the sputum clinical samples, 6/25 (24%) of MRSA belonged to t1081 type, which is supposed to be more prevalent in long-term care facilities (LTCFs).⁴⁹ Several reports indicated that t1081 had spread from the Netherlands to Hong Kong and Taiwan.^{50–54} The present study was done in Guangdong, which is a province located near Taiwan and

Hong Kong; therefore, presumably, t1081 might not only be introduced from LTCF residents to hospitalized patients, but also from Hong Kong and Taiwan to Mainland China.

Conclusion

In conclusion, our study elucidated and gave insight on the high prevalence of MRSA in southern China, especially from SSTI and sputum clinical samples. Notably, GEN and CIP should be used carefully in respiratory tract infections with MRSA and rational use of antibiotics should be undertaken to slow the epidemic spread of MDR in China. Our study also showed that ST239-SCCmecA III-t37 clone was the most prevalent among our isolates, indicating that ST239-t30 clone, which was previously predominant worldwide, had been replaced by a new clone, providing confirmation that clonal replacement is persistent.

Acknowledgment

This work was supported by the National Natural Science Foundation of China (grant numbers 81471988 and 81722030).

Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

Table S1 Mutation of *gyrA*, *gyrB*, *grlA* and *grlB* among MRSA isolates from different sources

No.	<i>gyrA</i>	<i>gyrB</i>	<i>grlA</i>	<i>grlB</i>
QY3	C251T (Ser84Leu)	/	/	/
QY35	C251T (Ser84Leu)	/	/	/
QY56	G317A (Phe106Tyr)	/	/	/
QY110	C251T (Ser84Leu)	/	/	/
L2301	C251T (Ser84Leu)	/	/	/
L2303	C251T (Ser84Leu), T290A (Met90Lys)	/	/	/
L2307	C251T (Ser84Leu)	/	/	/
L2308	C251T (Ser84Leu)	/	/	/
L2309	C251T (Ser84Leu)	/	G253A (Arg85Gln)	/
L2311	C251T (Ser84Leu)	/	C242T (Ser81Phe)	/
L2312	C251T (Ser84Leu)	/	T138G (Ile46Met)	A1266T (Glu422Asp)
L2313	C251T (Ser84Leu)	/	C242T (Ser81Phe), G253A (Arg85Gln)	/
L2314	C251T (Ser84Leu)	/	/	/
L2315	C251T (Ser84Leu)	/	/	/
L2316	T250G (Ser84Ala)	/	C242T (Ser81Phe)	/
L2317	T250G (Ser84Ala)	/	C242T (Ser81Phe)	/
L2318	C251T (Ser84Leu)	/	G253A (Arg85Gln)	/
L2319	C251T (Ser84Leu)	/	/	G1294A (Asp432Asn)
L2320	C251T (Ser84Leu)	/	/	C1351T (Pro451Leu)
L2321	C251T (Ser84Leu)	/	C242T (Ser81Phe)	/
L2322	C251T (Ser84Leu), G317A (Phe106Tyr)	/	C242T (Ser81Phe)	G1294A (Asp432Asn)
L2323	C251T (Ser84Leu)	/	C242T (Ser81Phe)	/
L2325	C251T (Ser84Leu)	/	G253A (Arg85Gln)	/
L2326	C251T (Ser84Leu)	/	C242T (Ser81Phe)	C1351T (Pro451Leu)
L2327	C251T (Ser84Leu), G317A (Phe106Tyr)	/	G253A (Arg85Gln)	/
L2328	C251T (Ser84Leu)	/	G253A (Arg85Gln)	/
L2330	C251T (Ser84Leu)	/	/	/
L2331	C251T (Ser84Leu), T253C (Ser85Pro)	/	/	/
L2332	C251T (Ser84Leu)	/	G253A (Arg85Gln)	/
L2335	C251T (Ser84Leu)	/	/	/
L2336	C251T (Ser84Leu)	/	C242T (Ser81Phe)	A1266T (Glu422Asp), C1387A, T1389A (Gly463Arg)
L2337	C251T (Ser84Leu)	/	C242T (Ser81Phe)	C1351T (Pro451Leu)
L2338	C251T (Ser84Leu)	/	C242T (Ser81Phe)	G1294A (Asp432Asn)
L2343	C251T (Ser84Leu), G317A (Phe106Tyr)	/	C242T (Ser81Phe)	G1294A (Asp432Asn)
L2344	C251T (Ser84Leu)	/	G253A (Arg85Gln)	/
L2503	C251T (Ser84Leu), A263T (Glu88Asp), G286A, A287C, T288A (Asp96Thy)	/	/	C1282T (Ser428Phe), C1330A, C1332A (Leu444Ile), A1346T (Tyr449Phe), T1358A (Val453Asp), C1387A, T1389A (Gly463Arg)
L2508	C251T (Ser84Leu)	G1390A (Val464Ile)	C242T (Ser81Phe)	G1294A (Asp432Asn)

Note: Reference: GenBank: AASB02000026.1 (*gyrA/gyrB*), GenBank: AKYW15.1 (*grlA/grlB*).

Abbreviation: MRSA, methicillin-resistant *Staphylococcus aureus*.

Table S2 MLST, SCCmec types and spa types of MRSA isolates from different sources

Molecular typing	Total n (%)	SSTI n (%)	Sputum n (%)	BSI n (%)	Others n (%)
SCCmec					
I	0	0	0	0	0
II	2 (1.94)	0	1 (4)	0	1 (20)
III	39 (37.86)	17 (29.31)	16 (64)	4 (26.67)	2 (40)
IV	17 (16.5)	14 (24.14)	1 (4)	2 (13.33)	0
a	15 (88.24)	13 (22.41)		2 (13.33)	0
b	2 (11.76)	1 (1.72)	1 (4)	0	0
c	0	0	0	0	0
d	0	0	0	0	0
V	1 (0.97)	0	1 (4)	0	0
No type	44 (42.72)	27 (26.21)	6 (24)	9 (60)	2 (40)
MLST					
239	24 (23.30)	7 (12.07)	14 (56)	2 (13.33)	1 (20)
338	18 (17.48)	16 (27.59)	0	1 (6.67)	1 (20)
59	8 (7.77)	5 (8.62)	0	3 (20)	0
1061	8 (7.77)	2 (3.44)	6 (24)	0	0
88	3 (2.91)	2 (3.44)	0	1 (6.67)	0
45	2 (1.94)	2 (3.44)	0	0	0
398	2 (1.94)	2 (3.44)	0	0	0
965	2 (1.94)	1 (1.72)	0	1 (6.67)	0
1	1 (0.97)	1 (1.72)	0	0	0
5	1 (0.97)	1 (1.72)	0	0	0
6	1 (0.97)	1 (1.72)	0	0	0
9	2 (1.94)	1 (1.72)	1 (4)	0	0
10	1 (0.97)	1 (1.72)	0	0	0
30	1 (0.97)	1 (1.72)	0	0	0
72	1 (0.97)	1 (1.72)	0	0	0
188	1 (0.97)	0	0	1 (6.67)	0
630	1 (0.97)	1 (1.72)	0	0	0
641	1 (0.97)	0	0	0	1 (20)
863	1 (0.97)	0	1 (4)	0	0
889	1 (0.97)	0	1 (4)	0	0
2148	1 (0.97)	0	0	1 (6.67)	0
3188	1 (0.97)	1 (1.72)	0	0	0
4860	1 (0.97)	1 (1.72)	0	0	0
4940	1 (0.97)	0	0	0	1 (20)
4942	1 (0.97)	1 (1.72)	0	0	0
4943	1 (0.97)	0	1 (4)	0	0
4944	1 (0.97)	0	1 (4)	0	0
4945	1 (0.97)	0	0	1 (6.67)	0
4946	1 (0.97)	0	0	1 (6.67)	0
4947	1 (0.97)	0	0	1 (6.67)	0
4948	2 (1.94)	1 (1.72)	0	0	0
4949	1 (0.97)	1 (1.72)	0	0	0
4957	1 (0.97)	0	0	1 (6.67)	0
4958	1 (0.97)	2 (3.44)	0	0	0
4959	1 (0.97)	1 (1.72)	0	0	0
4960	2 (1.94)	2 (1.72)	0	0	0
4961	1 (0.97)	0	0	0	1 (20)
4962	1 (0.97)	1 (1.72)	0	0	0
4963	1 (0.97)	1 (1.72)	0	0	0
4964	1 (0.97)	1 (1.72)	0	0	0
4965	1 (0.97)	0	0	1 (6.67)	0
spa					
t437	22 (21.36)	19 (32.76)	1 (4)	2 (13.33)	0
t37	18 (17.48)	6 (10.34)	9 (36)	2 (13.33)	1 (20)

(Continued)

Table S2 (Continued)

	Total n (%)	SSTI n (%)	Sputum n (%)	BSI n (%)	Others n (%)
t30	6 (5.83)	1 (1.72)	4 (16)	1 (6.67)	0
t1081	6 (5.83)	0	6 (24)	0	0
t189	4 (3.88)	2 (3.44)	0	2 (13.33)	0
t441	4 (3.88)	2 (3.44)	0	1 (6.67)	1 (20)
t62	3 (2.91)	1 (1.72)	0	1 (6.67)	1 (20)
t116	3 (2.91)	3 (5.17)	0	0	0
t34	2 (1.94)	2 (3.44)	0	0	0
t899	2 (1.94)	1 (1.72)	1 (4)	0	0
t14034	2 (1.94)	1 (1.72)	0	1 (6.67)	0
t2	1 (0.97)	0	0	0	1 (20)
t19	1 (0.97)	1 (1.72)	0	0	0
t114	1 (0.97)	1 (1.72)	0	0	0
t421	1 (0.97)	0	1 (4)	0	0
t701	1 (0.97)	1 (1.72)	0	0	0
t1212	1 (0.97)	1 (1.72)	0	0	0
t1244	1 (0.97)	1 (1.72)	0	0	0
t2358	1 (0.97)	1 (1.72)	0	0	0
t3523	1 (0.97)	1 (1.72)	0	0	0
t5554	1 (0.97)	1 (1.72)	0	0	0
t6497	1 (0.97)	1 (1.72)	0	0	0
t7707	1 (0.97)	0	0	1 (6.67)	0
t12011	1 (0.97)	0	0	0	0
t17403	1 (0.97)	0	1 (4)	0	0
Not available	17 (16.5)	10 (17.24)	2 (8)	4 (26.67)	1 (20)

Abbreviations: BSI, blood stream infection; MLST, multilocus sequence typing; MRSA, methicillin-resistant *Staphylococcus aureus*; SCCmec, staphylococcal chromosomal cassette mec; SSTI, skin and soft tissue infection.

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