



# Agr Typing and Detection of Mupirocin Resistance in Multi-Drug Resistant Clinical Isolates of *Staphylococcus aureus* in Northern Iran

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## ABSTRACT

**Background:** Mupirocin is a topical antibiotic inhibiting most Gram-positive cocci. Shortly after taking mupirocin, drug resistance emerges. This study aimed to determine mupirocin resistance in *Staphylococcus aureus* strains isolated from clinical specimens in Rasht.

**Materials & Methods:** In this study, a total of 85 clinical isolates of *S. aureus* were collected. Biofilm formation ability and antibacterial resistance patterns of isolates were investigated. Disc diffusion method and MIC determination were used to determine the susceptibility of strains to mupirocin antibiotic. Agr types, the presence of *mupA*, and mutation in *ileS-1* were evaluated in mupirocin non-susceptible isolates by PCR and PCR sequencing, respectively.

**Findings:** Out of 85 tested strains, 57 (67%) isolates were recognized as biofilm producers, and all of which showed multidrug resistance phenotype. Agr type 1 was the most commonly detected type. Additionally, 12 mupirocin-resistant strains were identified in the disc diffusion and MIC tests. A total of four strains were *mupA* positive and showed high-level resistance. In sequencing and mutation evaluation of the *ileS-1* gene in eight low-level mupirocin-resistant strains, 12 types of silent mutation and one type of missense mutation were determined.

**Conclusion:** The study of mupirocin-resistant strains in this study showed the need to identify factors affecting the occurrence of resistance and to take control and prevention measures before mupirocin losses its efficacy.

**Keywords:** *Staphylococcus aureus*, Mupirocin resistance, *ileS-1*, *MupA*.

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## Introduction

*Staphylococcus aureus* is a microbial flora of the skin and nasal passages of some people and one of the most common pathogens responsible for a wide range of infections from superficial skin infections to many serious infections such as septicemia, endocarditis, and osteomyelitis in hospitalized patients. Drug resistance is more common in *S. aureus* strains than other bacteria [1-2]. Mupirocin is one of the most effective antibiotics against Gram-positive cocci, used as a topical antimicrobial agent [3]. Nowadays, this antibiotic is used to prevent or treat skin surface infections, such as yellow ulcers, wound, and burn infections, and remove *S. aureus* nasal carriers and control MRSA transmission in healthcare settings [1, 4]. Mupirocin is a polyketide antibiotic with broad-spectrum antibacterial effects. Mupirocin epoxide side chain has a structure similar to isoleucine and could be linked to a specific position of isoleucine in isoleucine-tRNA synthetase and inhibit this enzyme and prevent the synthesis of protein and RNA, leading to bacterial death [5]. However, increased use of mupirocin causes the development of mupirocin resistance among *S. aureus* isolates. The development of resistance to mupirocin could be of both chromosomal and plasmid origins and is phenotypically divided into low- and high-level mupirocin resistance [6-7]. Low-level resistance appears due to decreased sensitivity to mupirocin as a result of mutation in chromosomal *ileS-1* gene encoding isoleucine-tRNA synthetase [7]. This type of resistance is sustainable and could be transferred to the next generation [8]. High-level resistance, which is a major threat to the clinical use of mupirocin, occurs due to the acquisition of plasmid-borne resistance genes including *mupA* (also known as *ileS-2*) and *mupB*, both of which encode an alternative isoleucine-tRNA synthetase

[9-10]. Conjugation of transgenic plasmids expressing this protein has been proven in staphylococcal strains in vitro and in vivo. It is believed that this plasmid transfer could contribute to the development of mupirocin resistance. The *mupB* gene encoding plasmid is responsible for very high-level mupirocin resistance [7].

The accessory gene regulator (*agr*) is one of the major factors regulating and controlling cell surface proteins and virulence gene expression in *S. aureus* strains. The *agr* operon includes *agrA*, *agrB*, *agrC*, and *agrD* genes. *Agr* system is polymorphic and permits the classification of *S. aureus* strains into four groups (*agr* I, *agr* II, *agr* III, and *agr* IV) according to sequences diversity in variable regions, comprising the last third of *agrB* and *agrD* as well as the first half of *agrC* genes [11-12].

**Objectives:** The aim of this study was to evaluate *agr* types and mupirocin resistance in clinical isolates of *S. aureus* in Guilan province.

## Materials and Methods

**Test bacteria:** *S. aureus* isolates were collected from hospitalized patients in Rasht during 2018. To isolate test bacteria, samples were cultured on mannitol salt agar and blood agar (Merck, Germany). Different biochemical characteristics including coagulase, catalase, and DNase production were investigated in the isolates. Subsequently, the identification of *S. aureus* isolates was confirmed by a pair of 23SrRNA specific primers via PCR as described previously [13].

**Biofilm formation assay:** This assay was performed in a microtiter plate. Overnight cultures of *S. aureus* isolates ( $1.5 \times 10^8$  CFU/mL) were diluted (1:100) in brain heart infusion broth supplemented with 1% glucose. Then 200  $\mu$ L of each bacterial suspension was transferred into each well

of a 96-well flat-bottom polystyrene plate and incubated at 37 °C for 48 hours. After incubation, the plank tonic bacteria were removed, wells were gently rinsed three times with sterile physiological saline and fixed by methanol for 20 min. Subsequently, the cells attaching to the surface were stained with crystal violet and rewashed. Then the assay was followed by destaining of biofilm-associated crystal violet using 1 mL of ethanol-acetone (95:5, vol/vol) solution. Subsequently, the optical density of the mixed solution was measured at 570 nm. The isolates were divided into three categories according their ability to produce biofilm as follows: non-biofilm producer ( $OD_{570} < 0.2$ ), weak-biofilm producer ( $0.2 < OD_{570} < 1.0$ ) and strong-biofilm producer ( $OD_{570} > 1.0$ )<sup>[14]</sup>.

**Antibacterial resistance:** Antibacterial resistance of *S. aureus* isolates was investigated using the disc diffusion method according to the CLSI guideline. The antibiotic disks used in this study (High Media-India) included clindamycin (2 µg), gentamicin (10 µg), teicoplanin (30 µg), linezolid (30 µg), mupirocin (20 µg), co-trimoxazole (23.75 µg), ciprofloxacin (5 µg), erythromycin (15 µg), cephalixin (30 µg), cephalothin (30 µg), methicillin (5 µg), amoxicillin (25 µg), amoxiclav (30 µg), and penicillin G. The tests were repeated twice, and bacterial resistance or susceptibility to antibiotics was determined by measuring the inhibition zone diameter according to the CLSI guideline. In addition, susceptibility to vancomycin was assessed by MIC determination based on the CLSI guideline (CLSI, 2018). The standard strain of *S. aureus* ATCC 33591 was used as control. *S. aureus* isolates resistant to at least three antimicrobial classes were considered as multidrug resistant (MDR)<sup>[15]</sup>.

**Agr typing:** *Agr* gene types were determined by *agr* group-specific multiplex PCR using primers specific to four types of this gene as described previously<sup>[12]</sup>. PCR reaction

was performed in a total volume of 25 µL, including 0.5 µL of dNTPs (10 µM), 5 µL of enzyme buffer (10X), 1 µL of each (8 µL in total) reverse and forward primers (10 pM), 2 µL of template DNA (2 µg), 0.5 µL of *Taq* polymerase enzyme (2.5 units) (Bioneer, South Korea), and 9 µL of distilled water. Thermocycler thermal conditions consisted of an initial denaturation step at 94 °C for 5 min, followed by 30 cycles of denaturation at 94 °C for 30 s, 55 °C for 30 s, and 72 °C for 60 s. Then a final extension step at 72 °C for 10 min was included. PCR products were detected by electrophoresis using a 1% agarose gel and confirmed by sequencing.

**Identification of mupirocin resistant strains:** Mupirocin resistance was determined using a standard disc diffusion method and determination of MIC by broth microdilution method (CLSI, 2018).

**Amplification of *ileS-1*, *mupA*, and *mupB* genes:** To amplify *ileS-1* and *mupA* genes, two pairs of primers specific to these genes were used. Table 1 shows the nucleotide sequences of these primers. The PCR reaction was carried out in a total volume of 25 µL consisting of 0.5 µL of dNTPS (10 µM), 5 µL of enzyme buffer (10x), 1.5 µL of each reverse and forward primers (10 pM), 2 µL of template DNA (2 µg), 0.5 µL of *Pfu* enzyme (2.5 units) (Bioneer, South Korea), and 14 µL of distilled water. Thermocycler thermal conditions consisted of an initial denaturation step at 94 °C for 5 min, followed by 30 cycles of denaturation at 94 °C for 45 s, annealing according to table 1 for 45 s and 72 °C for 60 s. Then a final extension step at 72 °C for 10 min was included. PCR products were detected by electrophoresis using a 1% agarose gel, and one PCR product from each type was confirmed by sequencing<sup>[10, 17]</sup>.

**Investigation of mutation of *ileS-1* gene:** After assuring the production of the desired product, it was sent to Bioneer Company (South Korea) for sequencing. Nucleotide

**Table 1)** Oligonucleotide primers used for the amplification of particular sequences of *S. aureus mupA*, *mupB*, and *ileS-1* genes

Gene	Primer Sequence	Amplicon Size (bp)	Annealing Tem. (°C)	Ref.
<i>Agr1-F</i>	5'-ATG CAC ATG GTG CAC ATG C-3'	441	56	16
<i>Agr1-R</i>	5'-GTC ACA AGT ACT ATA AGC GCT GAT-3'			
<i>Agr2-F</i>	5'-ATG CAC ATG GTG CAC ATG C-3'	575	56	16
<i>Agr2-R</i>	5'-TAT TAC TAA TTG AAA AGT GGC CAT AGC-3'			
<i>Agr3-F</i>	5'-ATG CAC ATG GTG CAC ATG C-3'	323	56	16
<i>Agr3-R</i>	5'-GTA ATG TAA TAG CTT GTA TAA TAA TAC CCA G-3'			
<i>Agr4-F</i>	5'-ATG CAC ATG GTG CAC ATG C-3'	659	56	16
<i>Agr4-R</i>	5'-CGA TAA TGC CGT AAT CG-3'			
<i>mupA-F</i>	5'-TAAGGAAGGGAGAATGGGAA -3'	456	60	17
<i>mupA-R</i>	5'-CTGAGCAAACGGCATAGAGC -3'			
<i>mupB-F</i>	5'-CTAGAAGTCGATTTTGGAGTAG-3'	674	60	10
<i>mupB-R</i>	5'-AGTGTCTAAAATGATAAGACGATC-3'			
<i>iles-1-F</i>	5'-CAGTTGCTACAAGAGGAGTGTCCACC-3'	915	65	17
<i>iles-1-R</i>	5'-CACCATGTTTCATAAGCTGTTGCC -3'			

sequences of the isolated strains *ileS-1* gene were compared to that of standard strain of *S. aureus* in gene bank, and changes in base and amino acid sequences were determined using online software, such as BLAST, Chromas (Ver. 1.45), and CLC Main Workbench (Ver.3.5).

## Findings

**Identification of *S. aureus* and biofilm formation ability:** Totally, 85 *S. aureus* strains were isolated. Distribution of *S. aureus* isolates according to the type of samples was as follows: 45 isolates from urine and 40 isolates from wound and abscess. According to amplification of the *23s rRNA* gene, all 85 isolates were confirmed as *S. aureus*. In phenotypic assays, 57 isolates (67%) were recognized as biofilm producers: 19 strong and 38 weak biofilm producers; the others (28 isolates) did not form biofilm.

**Antibacterial resistance pattern:** All the tested isolates showed multidrug resistance phenotype. Resistance rate of *S. aureus* isolates to 14 selected antibiotics was as follows: clindamycin, 6 (7.05%); gentamicin, 8 (9.41%); vancomycin, 3 (3.53%); teicoplanin, 4 (4.7%); linezolid, 5 (5.9%); methicillin, 53 (62.35%); co-trimoxazole, 32 (37.64%); ciprofloxacin, 38 (44.7%); erythromycin, 44 (51.76%); cephalixin, 49 (57.64%); cephalothin, 35 (41.17%); amoxicillin, 51 (50.59%); amoxiclav, 26 (30.59%); and penicillin G, 80 (94.12%).  
**Agr typing:** Among the 85 isolates studied, *agr* gene was identified in 81 *S. aureus* isolates using primers specific to four types of this gene. A total of 65 isolates (76.5%) were identified as *agr* type 1, eight isolates (9.4%) as *agr* type 2, five isolates (5.9%) as *agr* type 3, and three isolates (3.5%) as *agr* type 4. None of the *agr* types were detected in four isolates.

**Table 2)** Drug resistance patterns of mupirocin non-susceptible isolates

Bacterial Isolate	Mupirocin Resistance Level	Drug Resistance Pattern
1	Low	E, TET, CP, SXT, FOX, CFX, PEN, MET, AMX, AMC
2	Low	CC, GM, E, TET, CP, CFX, PEN, MET
3	Low	E, LZD, TET, CP, SXT, FOX, CFX, PEN, MET, AMX, AMC
4	Low	E, TET, CP, SXT, FOX, CFX, PEN, MET, AMX, AMC
5	Low	GM, TEK, TET, CFX, PEN, V, MET, AMX
6	Low	E, CP, CFX, PEN, MET, AMX, AMC
7	Low	E, TET, SXT, FOX, CFX, PEN, MET, AMX, AMC
8	Low	GM, E, TET, SXT, FOX, CFX, PEN, MET, AMX, AMC
9	High	CC, E, TET, CP, SXT, CFX, PEN, MET, AMX
10	High	E, LZD, TEK, TET, CP, FOX, CFX, PEN, V, MET, AMX
11	High	CC, GM, E, LZD, TET, CP, SXT, FOX, CFX, PEN, MET
12	High	CC, GM, E, TEK, TET, SXT, FOX, CFX, PEN, V, MET, AMX, AMC

CC: clindamycin, GM: gentamicin, E: erythromycin, LZD: linezolid, TEK: teicoplanin, TET: Tetracyclin, CP: ciprofloxacin, SXT: trimethoprim-sulfamethoxazole, FOX: ceftiofloxacin, CFX: cephalexin, PEN: penicillin, V: vancomycin, MET: methicillin, AMX: amoxicillin, and AMC: amoxiclav.

**Detection of mupirocin resistant strains:**

In the phenotypic assay, eight isolates with a mupirocin MIC of 8-128 µg/mL were identified as low-level mupirocin resistant strains, and four isolates with MIC of 512 µg/mL were recognized as high-level resistant strains. All of the mupirocin non-susceptible isolates were methicillin resistant and belonged to *agr* type 1. Linezolid was the most potent antibiotic against mupirocin resistant strains. Table 2 presents antibacterial drug resistance patterns in mupirocin non-susceptible *S. aureus* isolates.

**Detection of *mupA* and *mupB* genes:**

Using *mupA* gene specific primers in PCR, a PCR product with an approximate length of 456 bp was obtained in four isolates (5.9%) with a MIC of 512 µg/mL. All of low-level mupirocin-resistant strains were negative for *mupA* gene. All of *mupA* positive strains were biofilm producer in the phenotypic

assay. None of the tested isolates harbored the *mupB* gene.

**Investigation of mutation of *ileS-1* gene:**

PCR amplification of the *ileS-1* gene in eight low-level mupirocin-resistant strains resulted in 915 bp amplicons. Comparing the nucleoside sequences of the isolates *ileS-1* gene with that of *S. aureus* standard strain in the gene bank, one type of mutation was detected in three isolates. This mutation resulted in replacing E707 with G, leading to the replacement of glutamic acid with glycine. Furthermore, 12 different silent mutations were identified in this investigation. Table 3 presents the obtained results for each isolate.

**Discussion**

Mupirocin was first introduced in 1985 in the United Kingdom; in 1986, it was used in hospitals (SFGH) to eradicate methicillin resistant *S. aureus* (MRSA) and intrinsic

**Table 3)** MIC of mupirocin, biofilm formation ability, and mutations in the *ileS-1* gene in low-level mupirocin-resistant *S. aureus* isolates

Bacterial Isolate	Sample Type	Biofilm Formation	Mupirocin MIC ( $\mu\text{g/mL}$ )	Missense Mutation	Number of Silent Mutations
1	Wound	-	8	-	7
2	Abscess	+	16	-	6
3	Urine	-	32	-	8
4	Wound	+	64	E707G	9
5	Abscess	+	128	-	8
6	Wound	+	128	-	7
7	Urine	+	128	E707G	7
8	Wound	+	128	E707G	8

methicillin sensitive *S. aureus* (MSSA) strains. Two years after introducing mupirocin, the first case of mupirocin resistance was reported, and since then, various types of resistance have been reported [18].

The present study determined *agr* types, biofilm formation ability, antibacterial resistance pattern, and mupirocin resistance mechanisms in *S. aureus* strains isolated from clinical specimens collected from Rasht hospitals, northern Iran. In this assay, all of the tested strains were multidrug resistant. The highest resistance rate was against penicillin G, followed by methicillin and cephalexin. Vancomycin, linezolid, and ticoplanin were the most potent antibiotics. Moreover, 67% of these isolates were biofilm producers, and *agr* type 1 was the most common type in these isolates. More than 75% of the isolates belonged to this type. Predominance of *agr* type 1 is in accordance with previous data from Iran [11-12]. In the phenotypic assay, 12 of 85 isolates tested were mupirocin resistant. All of the mupirocin-resistant strains were methicillin resistant and biofilm producer while belonging to *agr* type 1; among which, eight isolates were identified as low-level mupirocin resistant. The MIC of mupirocin in these isolates ranged from 8-128 mcg/mL. In 3 of them, E707G single point amino acid substitution was identified. All of these

isolates had silent mutations, and totally, 12 different types of silent mutations were recognized. In most cases, the number of mutations in *ileS-1* gene was positively correlated with MIC value. E707G amino acid substitution identified in *ileS-1* has not been previously reported. In a study conducted by Fujimora et al. (2003), A637G, G1762T, G1891T, and A2412T amino acids substitution in *ileS-1* was reported in mupirocin-resistant *S. aureus* strains [19]. In a study by Antonio et al. (2002), 22 silent mutations and 9 types of missense mutations, including A110C, T635C, A637G, A769G, G808T, G1891T, T1948A, A2330G, and G1762T, were reported in the *ileS-1* gene in *S. aureus* isolates with decreased sensitivity to mupirocin [20].

In addition, Yang et al. (2006) reported that all mupirocin resistant isolates showed mutation at V588F. Furthermore, they reported P187F, K226T, F227L, Q612H, and V767D mutations [21].

Moreover, in the present study, four high-level mupirocin resistance phenotypes were identified, all of which were positive for *mupA* gene. MIC of mupirocin was 512  $\mu\text{g/mL}$  in these isolates. In similar studies, Hesami et al. (2013) identified 11 mupirocin-resistant *mupA* positive isolates [22]. Contrary to the present study, highly resistant isolates had

a higher prevalence than low-level resistant strains. On the other hand, in a study by Daskalaki et al. (2009) in Spain, out of 23 (11.3%) mupirocin-resistant isolates, 18 (78.3%) isolates had high-level resistance, and five isolates (21.7%) had low-level resistance. All high-level mupirocin-resistant isolates were positive for the presence of the *ileS-2* gene in PCR reaction, while the isolates with low-level resistance were negative for the presence of this gene [23]. In a study by Chaturvedi et al. (2014), out of 15 mupirocin-resistant strains, eight ones (53.3%) showed high-level resistance, and seven (46.7%) isolates exhibited low-level resistance [24].

Among mupirocin non-susceptible isolate, four (5.9%) isolates were identified as high-level mupirocin resistant with a MIC of  $\geq 512$   $\mu\text{g/mL}$ . This frequency was high in comparison with that reported by Mahmoudi et al. (2019), identifying mupirocin resistance in 3% of clinical isolates of *S. aureus* in Tehran, Iran. In their study, the *mupA* gene was detected in two out of four high-level mupirocin resistant isolates, and the *mupB* gene was not detected in any isolates, which is in accordance with the present study results [10]. In another study performed in Tehran, the prevalence of low-level and high-level mupirocin-resistant isolates as well as the *mupA* gene was 3.5, 1, and, 5.8%, respectively [25]. A study in Arak reported that the frequency of high-level mupirocin resistance and the *mupA* gene in isolates was 7.3 and 6%, respectively [22]. Also, in 2018, 12.5% of *S. aureus* clinical isolates from two major hospitals in Shiraz were high-level mupirocin-resistant, and 55 (45.8%) isolates carried the *mupA* gene [26].

### Conclusion

Overall, the present study and different studies worldwide indicate the emergence and spread of mupirocin resistance in *S.*

*aureus* isolates. Identification of mupirocin-resistant strains highlights the need to identify factors affecting the occurrence of these resistance and to take control and prevention measures before mupirocin losses its efficacy. Therefore, regular surveillance of *S. aureus* strains resistance to this antimicrobial and establishment of adequate infection control measures against resistant isolates are suggested.

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