Virulence genes profile and biofilm formation of Methicillin-resistant and Methicillin-susceptible *Staphylococcus aureus* isolates from diabetic foot infections

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ABSTRACT

Aims: Diabetic foot infections (DFIs) represent one of the most important risk factors for lower extremity amputation. One of the major infection agents that causes DFIs is *Staphylococcus aureus*. *Staphylococcus aureus* is an important human pathogen causing variety of clinical manifestations which can lead to invasive infections, sepsis and even death. Outcomes of antibiotic treatment of diabetic foot infections may depend not only on the antimicrobial susceptibility of the etiological agents, but also their ability to produce diverse virulence factors. This study was aimed to investigate biofilm production and the presence of various virulence genes among Methicillin-resistant *Staphylococcus aureus* (MRSA) and Methicillin-susceptible *Staphylococcus aureus* (MSSA) isolates obtained from patients with DFIs.

Methodology and results: A total of 48 clinical MRSA and MSSA isolates obtained from diabetic foot patients were studied for their biofilm formation and the presence of 29 known virulence genes. The biofilm formation was observed, analyzed and quantified using the microtiter plate method. Biofilm production was observed as 95.50% and 92.00% in the MRSA and MSSA isolates, respectively. Among the 29 virulence genes tested on the 48 clinical isolates, 19 virulence genes were detected. It was found that *aap* (62.50%), *etd* (54.17%), *icaD* (50.00%), *aae* (50.00%), *seh* (31.25%) and *icaADB* (22.92%) were the most prevalent genes. A total of 10 virulence genes (*etb, gehD, icaB, icaC, seb, hla_haem, hld_epid, altE, fbe* and *sesI*) were absent in all the isolates used.

Conclusion, significance and impact of study: Virulence genes play important role in clinical infections. Our results showed the presence rates of biofilm formation and accumulation-associated factors that are high among MRSA as well as MSSA isolates from DFIs. These results confirmed the importance of biofilm formation as regarded for DFIs.

Keywords: MRSA (Methicillin-resistant *Staphylococcus aureus*), MSSA (Methicillin-susceptible *Staphylococcus aureus*), Diabetic foot infections (DFIs), virulence factors

INTRODUCTION

Diabetic foot infections (DFIs) are soft tissue or bone infections below the malleoli, representing one of the most important risk factors for lower-extremity amputation, substantial morbidity and resulting in high treatment cost (Lipsky *et al.*, 2016). The pathophysiology of DFIs is complicated, relating to both host-disorders like neuropathy, arteriopathy and pathogen-related factors like virulence and antibiotic resistance (Dunyach-Remy *et al.*, 2016). Bacteria causing DFIs normally differ based on the geographical region in which they are allocated. The aerobic Gram-positive cocci like *Staphylococcus aureus* are usually found in the North America and Europe, while in Africa and Asia are usually aerobic Gram-negative bacilli such as *Pseudomonas aeruginosa* (Hatipoglu *et al.*, 2014). *Staphylococcus aureus* is among the first

pathogens to be described in the clinical manifestation and it is still considered as one of the deadliest pathogens of humans. The exceptional level of *S. aureus* to possess various resistance against multiple antibiotic classes contributes to clinical complications of this pathogen, leading to difficulties in choosing appropriate antibiotics (Diekema *et al.*, 2001; Schito, 2006).

Semisynthetic antibiotic methicillin was discovered in the 1950s, with methicillin-resistant *S. aureus* (MRSA) clinically identified in the 1960s (Bergdoll *et al.*, 1981). MRSA strains produce penicillin-binding protein (PBP) with decreased affinity for most semisynthetic penicillin. This protein is encoded by *mecA* gene, which is carried on a mobile genetic element (MGE), designated *Staphylococcal cassette chromosome mec* (*SCCmec*) (Freer and Arbuthnott, 1982). Infections associated with MRSA strains results in high mortality rate than infections

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caused by methicillin-susceptible *S. aureus* (MSSA) strains (Liu *et al.*, 2015). Among the multiple virulence factors produced by *S. aureus* is the production of cytotoxins, which includes hemolysins, staphylococcal enterotoxins, toxic-shock syndrome toxin-1 (*tsst-1*) and Panton-valentine leukocidin (Mariutti *et al.*, 2017).

Toxins play important role in the pathogenicity of *S. aureus* by producing lesion directly or indirectly to the membrane of the targeted cells (Amagai *et al.*, 2002). Staphylococcal enterotoxins (SE) are globular water-soluble proteins, rich in lysine, aspartic acid and glutamic acid with a molecular weight of 26–29 kDa. Enterotoxins are relatively resistant to heat and the presence of its proteolytic enzyme allows it to breach via gastro-intestinal tract during infections (Wolk *et al.*, 2009). Biofilm serve as a key barrier interfering antimicrobial agents and host immune system during staphylococcal infections. Biofilm is a fundamental factor for chronic bacterial infections, which is composed of a cell-multilayer embedded in a slime matrix. Cell aggregation and biofilm accumulation are mediated by the composition of *icaADB* and ica*C* genes, which is responsible for encoding essential proteins for the production of polysaccharide intercellular adhesion (PIA) and capsular polysaccharide/adhesion in *Staphylococcus* spp. (Katayama *et al.*, 2000). Another important toxin is the Panton-valentine leukocidin (PVL), a pore-forming cytotoxin that causes destruction of human and rabbit mononuclear and polymorph nuclear cells (Bergdoll *et al.*, 1971). Exfoliative toxins are also known as the epidermolytic toxins. They are specific serine proteases produced by *S. aureus*, which recognize and hydrolyze desmosome cadherin in the superficial layers of epidermis in humans. It causes cleavage of keratinocytes junction and cell-to-cell adhesion in the epidermis of the host cell, leading to peeling and blistering of the skin (Cramton *et al.*, 1999; Prévost *et al.*, 2001).

The evaluation and correlation between specific virulence genotypic markers in *S. aureus* isolates from DFIs has being studied on various dimensions. It is reported that Staphylococci is the virulent pathogen that is most frequent identified in clinical isolates in DFIs (Lipsky *et al.*, 2016). Recently, due to the increase in multi-drug resistant (MDR) isolates, the complications of DFIs have increased rapidly which makes it difficult to differentiate true infectious pathogens from colonizers (Spichler *et al.*, 2015). The clinical importance of bacterial virulence factors is related to whether or not they are expressed, and if expression leads to increase virulence. In this regard, several molecular studies have been performed to detect virulence factors responsible for Staphylococcal pathogenicity in relation to wound adherence and colonization (Sotto *et al.*, 2008; Liesse Iyamba *et al.*, 2011; Víquez-Molina *et al.*, 2018). In this study, we aimed to investigate the ability of biofilm production and the presence of various virulence factor genes among MRSA and MSSA isolates cultured from patients with DFIs in Turkey.

MATERIALS AND METHODS

Bacterial isolates

A total of 48 *S. aureus* isolates from diabetic foot infections were included in the present study. The isolates were obtained from routine clinical microbiology laboratory collection of Aydın Adnan Menderes University Hospital, Turkey. All samples were from the debridement material of diabetic foot wound infections. MRSA and MSSA represent 23 and 25 of the total 48 isolates, respectively. The isolates were grown aerobically on Tryptic Soy Agar (TSA) for 18–24 hours at 37 °C and stored in 2× skimmed milk at –20 °C.

For PCR setup, *S. aureus* G3959 was served as positive control to identify *hld_epid, gehD, icaA, icaB, icaC, icaD, hlb_epid, atlE, fbe, aae, aap, gehC* and *sesl* genes. While *S. aureus* G4005 used to identify *she* and *hla_haem* genes. These strains were received from clinical microbiology laboratory collection, Aydın Adnan Menderes University Hospital, Turkey.

Observation of Hemolysis

To investigate the ability of the isolates to perform hemolysis, all the isolates were streaked on blood agar medium composed of Luria Bertani agar: 1.0% tryptone (Merck Laboratory), 0.5% yeast extract (Merck Laboratory), 1.0% NaCl (Merck Laboratory), supplemented with pre-warmed (37 °C) 5% sheep erythrocytes (Biomerieux, Ca. No. 55822). The plates were incubated at 37 °C overnight and the presence or absence of hemolysis was accessed based on the hemolytic zones produced (Figure 1).

Figure 1: Hemolysis on blood agar. Hemolytic reaction of four isolates tested on sheep blood agar after 24 hours. Hemolysis; positive for 3 isolates (B, C and D) and negative for 1 isolate (A).

Microtiter Plate Assay for Biofilm Formation

The microtiter assay was used to evaluate the biofilm formation of MRSA and MSSA isolates as described previously (Stepanović *et al.*, 2007). Briefly, bacteria were inoculated into TSB containing 0.25% glucose and incubated at 37 °C overnight. The overnight cultures were then diluted in 1:100 TSB with 0.25% glucose. Then, 200 µL of the overnight culture was transferred to U-shaped 96 well micro plates and incubated for 48 hours at 37 °C. For negative control, fresh TSB medium was used. Following incubation, the wells were discarded and gently washed three times with 200 µL sterile phosphate buffered saline (PBS; pH 7.4). Biofilms were fixed dried at 60 °C for 45 min. Subsequently, plates were stained by adding 125 µL of 1% crystal violet and incubated at room temperature for 15 min. Excess crystal violet was rinsed under running water and left to ambient temperature to be dried. To dissolve any remaining stain, 150 µL of 80% (v/v) ethanol with 20% (v/v) acetone was added.

The optical density (OD) of each well was measured spectrophotometrically at 595nm (A_{595}) using the microplate reader (Thermo Multiscan Spectro). The negative control containing uninoculated TSB was used to determine the background OD. Formation of biofilms by MRSA and MSSA were analyzed and categorized depending on the absorbance of slime cells attached to crystal violet as described previously (Ohadian *et al.*, 2014). The OD_{595} values less than 0.125 were grouped as non-biofilm producers, those between 0.125–0.250 (2 × 0.125) as weak biofilm producers; between 0.250–0.5 (4 × 0.125) as moderate biofilm producers, and values higher than 1.0 (8 × 0.125) were considered as strong biofilm producers.

Extraction of DNA for Polymerase Chain Reaction (PCR)

All isolates were cultured aerobically on Tryptic Soy Agar (TSA) at 37 °C overnight. The DNA was extracted using DNA4PCR kit (R Tech Aydin, Turkey) protocol as recommended by the manufacturer. Briefly, 1 or 2 bacterial colonies were homogenized in 1 mL distillated water and centrifuged for 5 min at 13000 rpm. The supernatant was discarded, and pellet was resuspended in 100 µL of DNA4PCR solution. The mixture was incubated at 56 °C for 20 min, vortexed and incubated at 100 °C for 10 min. After vortexing for 15 sec and centrifugation at 4 °C, 13000 rpm for 5 min, 1 µL of the supernatant was used for PCR. The purified DNAs were stored at −20 °C until use.

Detection of Virulence Genes by PCR

The presence of virulence genes: *eta, etb, etd, etx, tst, luk- pvl, hla_ haem, hlb_epid, hld_ epid, icaA, icaB, icaC, icaD, icaADB, gehD, sea, seb, sec, sed, see, seg, seh, sei, atlE, fbe, aae, aap, gehC* and *sesI* from the isolates of MRSA and MSSA were determined using PCR with the oligonucleotide primers as given in Table 1. The PCR was performed in a final volume of 50 µL reaction mixture containing: 1× Taq buffer with Mg^{2+}, 0.2 mM dNTP, 0.4 pmol forward primer and 0.4 pmol reverse primer, 1.5 U Taq polymerase and 1.0 µL purified DNA. PCR program was performed with an initial denaturing step at 95 °C for 3 min, followed by 35 cycles of denaturing at 95 °C for 30 sec, annealing at 50 °C for 30 sec, extension at 72 °C for 1 min and the reaction was finalized with an extension step at 72 °C for 5 min. The PCR product (5 µL) were mixed with 1 µL of loading dye solution and carefully loaded into the 2% agarose gel well. The lambda PstI marker (Thermo fisher Scientific, USA) was used to approximate the size of the amplicons. The gel was then electrophoresed at 100 V for 30 min and visualized under UV trans-illuminator device (Vilber Lourmat 08-200229).

RESULTS AND DISCUSSION

Despite the development of diverse control measures, bacterial infections remain among the leading causes of death among wound patients. Challenges in the treatment of staphylococcal infections are accompanied by several mechanisms of biofilm formation relating to this bacterium (Ohadian *et al.*, 2014).

A total of 48 *S. aureus* isolates from DFIs among which 23 (48%) MRSA and 25 (52%) MSSA were studied. The ability of bacteria to produce biofilm was tested using the microtiter plate assay. Based on our results (Table 2), 16 (69.5%) MRSA isolates were examined as moderate biofilm producer. Non-biofilm was formed by 1 (4.3%) of MRSA isolates, 2 (8.6%) of the MRSA isolates formed weak biofilm and 4 (17.4%) of the MRSA isolates were considered as strong biofilm producers. On the other hand, among the 25 MSSA isolates, 15 (60.0%) isolates were examined as moderate biofilm producer. Non-biofilm producer was recorded by 2 (8.0%) of the MSSA isolates, 1 (4.0%) of MSSA isolates as weak biofilm producer, whilst 7 (28.0%) of the MSSA isolates formed strong biofilms. Smith *et al.* (2008) examined the biofilm formation of 763 (53.8%) MRSA and 209 (43.5%) MSSA isolates and they found that 20.5% of the MRSA and 28.0% of the MSSA isolates established fully established biofilm. O'Neil *et al.* (2007) documented similar study of biofilm formation in 114 clinical isolates of MRSA and found that only 9% had the ability to form fully established biofilms (strong biofilm producers). These differences in biofilm formation may be due to the variation in isolation site or the geographical differences in the isolated genotypes of the strains included in the study.

Virulence genes investigation among the isolates of MRSA and MSSA is presented in Table 3. From the 29 virulence genes tested, our results showed the presence of 19 virulence genes and absence of 10 virulence genes (*etb, gehD, icaB, icaC, seb, hla_haem, hld_epid, altE, fbe* and *sesI*) in all the isolates with the primers used. The virulence genes detected were *eta, etd, etx, sea, sec, sed, see, seg, seh, sei, icaA, icaD, icaADB, aae, aap, luk_pvl, tst, hlb_epid* and *gehC*. The most prevalent

Table 1: Primer sequences used in the PCR assays for detecting virulence genes among the MRSA and MSSA isolates.

Gene	Primer	Nucleotide sequence (5'–3')	Product size (bp)	References
icaA (intercellular adhesion A)	forward	ACAGTCGCTACGAAAAGAAA	103	(Arciola *et al.*, 2005)
	reverse	GGAAATGCCATAATGAGAAC		
icaB (intercellular adhesion B)	forward	CTGATCAAGAATTTAAATCACAAA	302	(Arciola *et al.*, 2005)
	reverse	AAAGTCCCATAAGCATATGTTT		
icaC (intercellular adhesion C)	forward	TAACTTTAGGCGCATATGTTT	400	(Arciola *et al.*, 2005)
	reverse	TTCCAGTTAGGCTGGTATTG		
icaD (intercellular adhesion D)	forward	ATGGTCAAGCCCAGAGAG	198	(Arciola *et al.*, 2005)
	reverse	CGTGTTT TCAACATTTAATGCAA		
icaADB (intercellular adhesion ADB)	forward	TTATCAATGCCGCAGTTGTC	546	(Arciola *et al.*, 2005)
	reverse	GTTTAACGCGAGTGCGCTAT		
aap (accumulation-associated protein)	forward	ATACAACTGGTGCAGTGGTTG	400	(Vandecasteele *et al.*, 2003)
	reverse	GTAGCCGTCCAAGTTTTACCAG		
aae (surface associated protein and autolysin)	forward	GAGGAGGATTTTAAAGTGC	858	(Heilmann *et al.*, 2003)
	reverse	AACATGACCATAGTAACC		
sea (enterotoxin A)	forward	TTGGAAACGGTTAAAACGAA	120	(Johnson *et al.*, 1991)
	reverse	GAACCTTCCCATCAAAAACA		
seb (enterotoxin B)	forward	TCGCATCAAACTGACAAACG	478	(Johnson *et al.*, 1991)
	reverse	GCAGGTACTCTATAAGTGCC		
sec (enterotoxin C)	forward	GACATAAAAGCTAGGAATTT	257	(Johnson *et al.*, 1991)
	reverse	AAATCGGGATTAACATTATCC		
sed (enterotoxin D)	forward	CTAGTTTGGTAATATCTCCT	317	(Johnson *et al.*, 1991)
	reverse	TAATGCTATATCTTATAGGG		
see (enterotoxin E)	forward	CAAAGAAATGCTTTAAGCAATCTTAGGCCAC	170	(Mehrotra *et al.*, 2000)
	reverse	CTTACCGCCAAAGCTG		
seg (enterotoxin G)	forward	AATTATGTTGAATGCTCAACCCGATC	642	(Omoe *et al.*, 2002)
	reverse	AAACTTATATGGAACAAAAGGTACTAGTTC		
seh (enterotoxin H)	forward	CAATCACATCATATGCGAAAGCAG	376	(Omoe *et al.*, 2002)
	reverse	CATCTACCCAAACATTAGCACC		
sei (enterotoxin I)	forward	CTCAAGGTGATATTGGTGTAGG	576	(Omoe *et al.*, 2002)

(Continued)

	reverse	AAAAAACTTACAGGCAGTCCATCTC		
tsst-1 (toxic shock syndrome toxin-1)	forward	AGCATCTACAAACGATAATATAAAGG	481	(Seier-Petersen *et al.*, 2015)
	reverse	CATTGTTATTTTCCAATAACCACCCG		
eta (exfoliative toxin A)	forward	CTATTTACTGTAGGAGCTCTCTAT	741	(Amagai *et al.*, 2002)
	reverse	ATTTATTTGATGCTCTCTAT		
etb (exfoliative toxin B)	forward	ATACACACATTACGGATAAT	629	(Amagai *et al.*, 2002)
	reverse	CAAAGTGTCTCCAAAAGTAT		
etd (exfoliative toxin D)	forward	AACTATCATGTATCAAGG	376	(Amagai *et al.*, 2002)
	reverse	CAGAATTTCCCGACTCAG		
etx (exfoliative toxin X)	forward	TACTCATACTGTGGGAACTTCGATACAAGC	403	(Amagai *et al.*, 2002)
	reverse	CTCATCTCCCATAACTGCACTATAATTTCG		
luk_pvl (pore forming toxin, Panton valentine leukocidin)	forward	ATCATTAGGTAAAATGTCTGGACATGATCCA	433	(Lina *et al.*, 1999)
	reverse	GCATCAATGTATTGGATAGCAAAAGC		
atlE (attachment to polystyrene and vitronectin)	forward	CAACTGCTCAACCGAGAACA	682	(Vandecasteele *et al.*, 2003)
	reverse	TTTGTAGATGTTGTGCCCA		
gehC (Lipase binding to collagen)	forward	CAAAAGATAGCCAATCAACAG	700	(Bowden *et al.*, 2002)
	reverse	GCGTACAATCGCTTGTTACC		
gehD (Triacylglycerol lipase D)	forward	TTTGAATTCTGCAAGCTCAATATAA	1179	(Bowden *et al.*, 2002)
	Reverse	TTTGCGGCCGCTATCGCTACTTACGTGTAA		
sesI (surface binding protein)	forward	GCTGATTATGTAAATGACTCAAAT	408	(Johnson *et al.*, 1991)
	Reverse	AGCTTTTGTTGTTTGAGCTTC		
hla_haem (alpha hemolysin)	forward	TGGGCCATAAACTTCAATCGC	72	(Rosec and Gigaud, 2002)
	reverse	ACGCCACCTACATGCAGATTT		
hlb_epid (beta hemolysin)	forward	TGGTGGCGTTGGTAT	335	(Rosec and Gigaud, 2002)
	reverse	ACCCCAAGATTCAC		
hld_epid (delta hemolysin)	forward	TGGTGGCGTTGGTATTGTGA	541	(Rosec and Gigaud, 2002)
	reverse	ACCCCAAGATTTCACGGACC		
fbe (fibrinogen binding protein)	forward	TAAACACCGACGATAATAACCAAA	495	(Arciola *et al.*, 2004)
	reverse	GGTCTAGCCTTATTTTCATATTCA		

Table 2: Results of biofilm formation of MRSA and MSSA isolates in the presence of 0.25% glucose using the microtiter plate technique.

	Biofilm formation			
	Non-biofilm producer[A] (ODs ≤ 0.125) N (%)	Weak biofilm producer[B] (0.125 < ODs ≤ 0.25) N (%)	Moderate biofilm producer[C] (0.25 < ODs ≤ 0.50) N (%)	Strong biofilm producer[D] (1.0 < ODs) N (%)
MRSA (n=23)	1 (4.3)	2 (8.6)	16 (69.5)	4 (17.4)
MSSA (n=25)	2 (8.0)	1 (4.0)	15 (60.0)	7 (28.0)

[A]ODs ≤ ODc; [B]ODc < ODs ≤ 2x ODc; [C]ODc < ODs ≤ 4x ODc; [D]1.0 < ODs
ODc = Optical density at 595 nm of negative control
ODs = Optical density at 595 nm of samples

genes among the isolates were *aap* (62.50%), followed by *etd* (54.17%), *icaD* (50.00%), *aae* (50.00%), *seh* (31.25%), *icaADB* (22.92%), *sei* (18.75%), *gehC* (16.67%), *see* (10.42%), *icaA* (8.33%), *sea* (6.25%), *seg* (6.25%), *sec* (4.17%), *sed* (4.17%), *eta* (4.17%), *etx* (4.17%), *luk_pvl* (4.17%), *hlb_epid* (2.08%) and *tst-1* (2.08%). We also found that among the MRSA isolates; *aae, aap* and *icaD* genes were significant while *etd, icaADB* and *gehC* were significantly frequent in the MSSA isolates. These results indicate the presence of distinct virulence genes among MRSA and MSSA strains.

Staphylococci devote significant amount of energy to the production of virulence factors to protect the bacteria from host immune surveillance, and also promote the bacterial survive in hostile environment (Richard *et al.*, 2011). Clinical outcome of *S. aureus* infections is influenced by the presence of various virulence genes. In this study, we found that MRSA and MSSA strains isolated from DFIs were characterized by a high prevalence of biofilm formation (*icaD*, *icaADB*), biofilm production (*aap*), autolysin (*aae*) and exfoliative (*etd*) genes, whereas low frequency of other toxin genes (*tsst-1, luk_pvl* and *hlb_epid*).

Biofilm formation contributes to the attachment of microbial cells to surfaces contributing to the initial stage of infections. It is considered as a characteristic virulence factor used by microbial organisms in pathogenesis enabling it to survive adverse environments within the host (Mashouf *et al.*, 2015). Among these virulence genes responsible for biofilm is the *icaD* gene which encodes the N-acetylglucosaminyl transferase; an enzyme involved in the synthesis of N-acetylglucosamine oligomers from UDP-N-acetylglucosamine (Costerton *et al.*, 1999). The prevalence of *icaD* gene in our study was 50.0% from the total isolates. MRSA and MSSA isolates displayed 60.0% and 40.0%, respectively. This result is in accordance to the study of Ghasemian *et al.* (2016), which reported the presence of *icaD* gene with a frequency of (69.0%) MRSA and (54.0%) MSSA in 209 clinical isolates tested. Aggarwal *et al.* (2019) also documented the presence of *icaD* gene in 109 isolates from wound infections with a higher frequency in both MRSA (81.3%) and MSSA (84.4%). The activation of accumulation associated protein (*aap*) is controlled by

bacterial and host-accompanied proteases (Conlon *et al.*, 2014). In our study, the frequency of *aap* gene was 62.5%, which MRSA and MSSA accounted for 69.5% and 56.0%, respectively. These results emphasized the relationship between bio-film formations and adhesive mechanism during infections. The relationship between biofilm formation and *aap* gene had been studied largely in *S. epidermidis* (Schaeffer *et al.*, 2015; Salgueiro *et al.*, 2017; Guo *et al.*, 2019), but not in *S. aureus* isolates.

Superantigens play essential role in the inflammations that occur in DFIs. Staphylococcal enterotoxins are important superantigens causing non-specific activation of the immune system especially the T cells. However, they do not all involved in food poisoning. The important food poisoning toxins are encoded by *sea, seb, sec, sed* and *see* genes. More recently, it was found enterotoxin genes show superantigenic effect but not emetic effect (Omoe *et al.*, 2002), except *seh* gene which is associated with food poisoning outbreak with raw milk (Jørgensen *et al.*, 2005). In our study, *seh* gene was more frequently identified in (39.10%) MRSA isolates, whereas *sei* gene was in the (32.00%) MSSA isolates. In contrast, Sila *et al.* (2009) reported the absence of *seh* gene in all the MRSA and MSSA isolates from the 200 clinical samples (mostly skin and wound) tested. The frequency of *seb* gene reported in Gorgan, Iran (Kamarehei *et al.*, 2013), Canada (Mehrotra *et al.*, 2000) and Tehran (Norouzi *et al.*, 2012) from the clinical samples were 61.3%, 15.8% and 73.6%, respectively. However, none of the MRSA and MSSA isolates harbored the *seb* gene from our study. These differences may be due to the existence of different genotypes among the strains and the toxin profile of specific isolates which are affected by the origins of their geographical location (Moore and Lindsay, 2001).

Another important gene observed in our study was the exfoliative genes. Four staphylococcal exfoliative toxins, *eta*, *etb*, *etd* and *etx* were tested in this study. Our result showed a high frequency rate of *etd* gene in MSSA (84.00%) isolates. In contrary, the *etd* gene was not found in any of the MSSA clinical isolates tested by (Wu *et al.*, 2011). The *etb* gene was not detected in our study. Similarly, Peacock *et al.* (2002) did not detect *etb* gene in all the isolates tested. Moreover, the frequency of *eta* gene (4.17%) was low in our study. Our results coincide

Table 3: Number of virulence genes of MRSA (n=23) and MSSA (n=25) isolates from diabetic foot infections (DFIs).

| Gene group	Virulence genes	Total positive	Positive MRSA	Positive MSSA	Positive MRSA (%)	Positive MSSA (%)	Total positive (%)
Biofilm formation	*icaA*	4	0	4	0	16.00	8.33
	icaB	0	0	0	0	0	0
	icaC	0	0	0	0	0	0
	icaD	24	14	10	60.00	40.00	50.00
	icaADB	11	1	10	4.35	40.00	22.92
Biofilm production	*aap*	30	16	14	69.50	56.00	62.50
Enterotoxins	*sea*	3	3	0	13.00	0	6.25
	seb	0	0	0	0	0	0
	sec	2	1	1	4.35	4.00	4.17
	sed	2	1	1	4.35	4.00	4.17
	see	5	4	1	17.40	4.00	10.42
	seg	3	0	3	0	12.00	6.25
	seh	15	9	6	39.10	24.00	31.25
	sei	9	1	8	4.35	32.00	18.75
Hemolysin toxins	*hla_haem*	0	0	0	0	0	0
	hlb_epid	1	0	1	0	4.35	2.08
	hld_epid	0	0	0	0	0	0
Fibrinogen Binding	*fbe*	0	0	0	0	0	0
Lipase binding to collagen	*gehC*	8	0	8	0	32.00	16.67
	gehD	0	0	0	0	0	0
Autolysin	*aae*	24	15	9	65.20	36.00	50.00
	altE	0	0	0	0	0	0
Surface protein I	*sesI*	0	0	0	0	0	0
Exfoliative toxins	*eta*	2	1	1	4.35	4.00	4.17
	etb	0	0	0	0	0	0
	etd	26	5	21	21.70	84.00	54.17
	etx	2	1	1	4.35	4.00	4.17
β-pore-forming toxins syndrome toxin-1	*luk_pvl*	2	2	0	8.70	0	4.17
Toxic-shock syndrome toxin	*tsst-1*	1	0	1	4.35	0	2.08
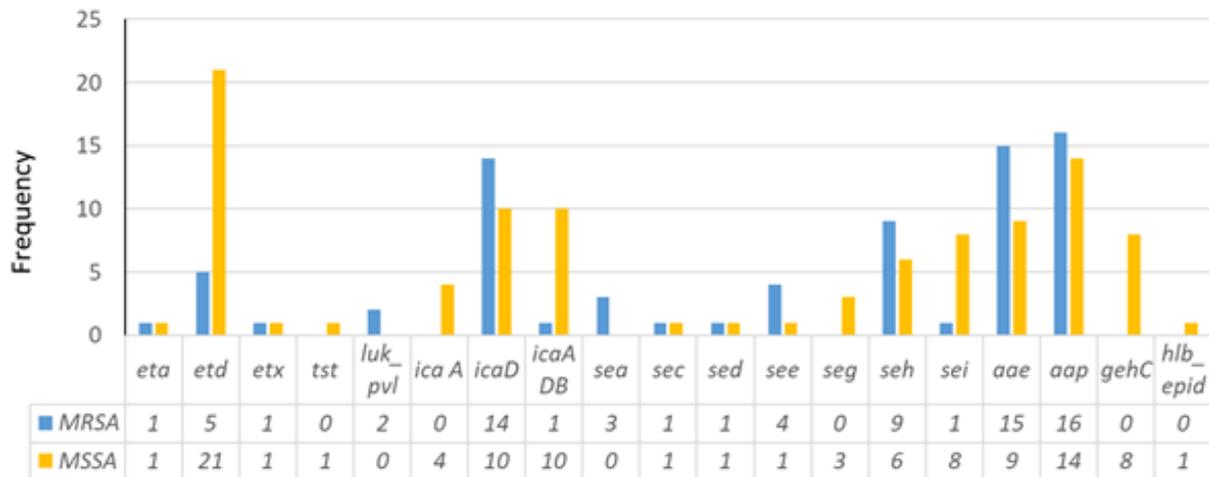
Figure 2: Frequency of virulence genes among MRSA and MSSA isolates. *eta*, exfoliative toxin A; *etd*, exfoliative toxin D; *etx*, exfoliative toxin *X*; *tsst-1*, toxic shock syndrome toxin-1; *luk_pvl*, Panton-valentine leukocidin; *icaA*, intercellular adhesion *A*; *icaD*, intercellular adhesion D; *icaADB*, intercellular adhesion ADB; *sea*, staphylococcal enterotoxin A; *sec*, staphylococcal enterotoxin C; *sed*, staphylococcal enterotoxin D; *see*, staphylococcal enterotoxin E; *seg*, staphylococcal enterotoxin G; *seh*, staphylococcal enterotoxin H; *sei*, staphylococcal enterotoxin I; *aae*, autolysin *A*; *aap*, accumulation-association proteins; *gehC*, triacylglycerol lipase C; *hlb_epid*, beta hemolysin.

with similar studies from Iran (Alfatemi *et al.*, 2014), Germany (Becker *et al.*, 1998) and Columbia (Smyth and Kahlmeter, 2005) which reported the frequency rates of *eta* gene as 0.68%, 2% and 3%, respectively.

The *tsst-1* gene was only detected in one MRSA isolates (2.08%) in our study. Our result contravenes with the study in Iran (Koosha *et al.*, 2016), which reported a high frequency of *tsst-1* gene among MRSA (69.8%) and MSSA (56.0%) in 172 wound isolates. The frequency of *luk_pvl* was reported in Tehran 24.2% (Norouzi *et al.*, 2012), Isfahan 75% (Ohadian-Moghadam *et al.*, 2012) and Columbia 73% (Ertas *et al.*, 2010). The frequency of *luk_pvl* gene in our study was 4.17%. The frequency obtained in our study is similar to the study in Iran, which reported the frequency of *pvl* gene to be 5.47% (Alfatemi *et al.*, 2014). These differences may be due to the geographical differences in the isolated genotypes of the MRSA and MSSA strains included in the study.

Hemolysins are important virulence factors contributing to Staphylococcal infections. Phenotypically, we accessed the presence or absence of hemolysis among the MRSA and MSSA isolates based on the hemolytic zone area produced. From the 23 MRSA isolates examined, 10 (43.5%) isolates showed hemolysis, whilst 13 (56.5%) did not show any hemolysis. As for MSSA isolates, 15 (60.0 %) isolates exhibited hemolysis, whilst 10 (40.0%) MSSA isolates did not produce any hemolysis. However, our results showed the absence of *hla_haem* and *hld_epid* genes in all the isolates tested. The absence of these genes is in contrary to the study conducted in Iran (Motamedi *et al.*, 2018), which reported the presence of hemolysin genes (*hla, hlb, hld* and *hlg*) among 389 clinical MRSA isolates as 9.19%, 8.04%, 13.79% and 3.44% respectively. Although we could not find any genetic determinant that we

investigated by PCR, our results may indicate the presence of other hemolysis gene(s) among these isolates (Figure 1).

Despite evaluation of limited number of strains from a single center, our study showed the importance of biofilm associated virulence genes (*icaD*, *icaADB*) in contributing to diabetic foot infections. The virulence genes were more diverse and abundant in MSSA than in MRSA strains as shown in Figure 2.

CONCLUSION

In conclusion, we found that the genes that encode for proteins related to the production and formation of biofilms (*aap*) are remarkably significant among MRSA and MSSA strains. Interestingly, an exfoliative gene (*etd*) was also found in high frequency among the diabetic isolates studied. This may be related to their pro-inflammatory effects. Despite the importance of hemolysins in *S. aureus* infections, our findings did not indicate the presence of *hla_haem* and *hld_epid* genes among the MRSA and MSSA isolates tested.

ETHICAL APPROVAL

As the testing used in this study is only a laboratory procedure done on specimens obtained for clinical purposes, there was no need for ethical approval.

INFORMED CONSENT

Patients were treated by their physicians according to standard protocols. As no additional specimens were collected for the virulence testing, and their identity was not linked in this publication to any identifying features,

we did not need to obtain informed consent for this testing.

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