The epidemiology and molecular characterization of methicillin-resistant staphylococci sampled from a healthy Jordanian population

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SUMMARY

The prevalence of natural carriage and molecular epidemiology of methicillin-resistant Staphylococcus aureus (MRSA) and methicillin-resistant coagulase-negative staphylococci (MR-CoNS) isolates in a Jordanian community were investigated. The MRSA nasal carriage rate in 227 healthy volunteers was 7.5% and the majority (81%) of MRSA harboured the resistance element SCCmec type IVe and were of a novel spa type t9519 (76%); other significant spa gene types were t223 (14.7%) and t044 (5.9%). All MRSA isolates were susceptible to other classes of antibiotics, and tested positive for at least three virulence factor encoding genes, but only two harboured the pvl gene. MR-CoNS carriage was 54.2% and these isolates were characterized by single, double and untypable SCCmec elements, with Staphylococcus epidermidis SCCmec type IVa predominating. Of eight subjects with nasal co-colonization of MR-CoNS + MRSA, three shared SCCmec type IV in both groups of organisms. This is the first report of methicillin-resistant staphylococci carriage in a Jordanian community and its findings are important for epidemiological study and infection control measures of these organisms.

Key words: Jordan, MR-CoNS, MRSA, SCCmec, spa.

INTRODUCTION

Since the 1990s, the epidemiology of methicillin-resistant Staphylococcus aureus (MRSA) infections has changed markedly owing to the emergence of community-associated MRSA (CA-MRSA) infections as a serious health problem in many parts of the world [1]. Initially CA-MRSA strains were identified as being responsible for the increase of staphylococcal infections in communities worldwide, but now they have expanded to also become a causative agent of nosocomial infections and thus the distinction between hospital- and community-acquired strains is becoming less clear [1]. Methicillin resistance in MRSA and coagulase-negative staphylococci (CoNS) is encoded by the mecA gene which is located within the staphylococcal cassette element (SCCmec) integrated in the bacterial chromosome [2–4]. Genotypic and phenotypic differences between CA-MRSA and hospital-acquired MRSA (HA-MRSA) strains are well
recognized and include the type of SCCmec element, their virulence factor profile, and wider antimicrobial susceptibility [5].

Nasal carriage of MRSA by hospital staff in Jordan was first reported in the early 1990s [6]. Subsequently, high rates of S. aureus (40%) and MRSA (19%) were recorded in young Jordanian adults in the community [7], which were somewhat higher than the rate (22.7%) found in a survey of 132 healthy students published 4 years earlier [8]. Recently, Khalil et al. [9] reported ST80-MRSA-IV as the dominant clonal type in hospitalized children in Jordan, although the extent of MRSA and MR-CoNS carriage in the healthy Jordanian community and the molecular characterization of these strains remain inadequately investigated.

A major risk factor for infections with methicillin-resistant staphylococci (MRS) is the carriage of these microorganisms at different body sites. Indeed their co-existence might facilitate the exchange of mobile resistance and other genetic elements and there is some evidence to suggest that MR-CoNS may act as a source of SCCmec for MRSA [4, 10, 11]. Although MR-CoNS colonization might be correlated with the emergence of MRSA, there are few epidemiological studies examining co-colonization in the healthy population for these two groups of organisms and it has been shown that there is no significant difference in terms of nasal carriage for S. aureus and MRSA [12] in pre-clinical and clinical university students; therefore university students are a suitable representative group on which to draw conclusions on the spread of MRSA in the wider community population [12, 13].

In the current study we aimed to determine the skin and nasal carriage rates of MR-CoNS and MRSA in healthy preclinical students and faculty staff in the Jordanian community. A secondary aim was to obtain baseline epidemiological data to inform the design and implementation of appropriate infection surveillance and control practices. Third, we investigated all isolates from the survey in detail with regard to their molecular genetic types and virulence gene profiles, and wider antimicrobial susceptibility.

METHODS

Subjects and data collection

During June 2009 to December 2009, a total of 454 microbiological samples were collected from 227 apparently healthy volunteer pharmacy students and employees at the Faculty of Pharmacy, The University of Jordan. Most (89%) of the students had not been exposed to clinical training within the 2 months prior to sampling. Written consent was obtained prior to commencing the study which complied with ethical guidelines of experimental approval (number 14/2007–2008) obtained from the Scientific Research Council at the Deanship of Academic Research, The University of Jordan. The following data were collected: demographic characteristics, medical history including previous and recent hospitalization (within a year), antibiotic consumption, and family member being a healthcare worker.

Sampling and bacteriological identification

The dorsum of the forearm (1 cm²) and the anterior nares of each volunteer subject were sampled [14] using sterile dry cotton swabs (Deltalab, Spain) pre-moistened with brain heart infusion (BHI) broth (Oxoid, UK) containing 6.5% NaCl. Each swab was immersed in the same broth and incubated at 37 °C for 2 h. The broth was subcultured onto mannitol-salt-agar (Oxoid) supplemented with methicillin (10 mg/l), and incubated at 35 °C for 24–72 h. Each distinctive colony morphotype was selected, Gram-stained and biochemically identified which included catalase (Merck, Germany), tube coagulase (Remel, UK) and DNase tests (Oxoid). Presumptive MRSA isolates (n=37) and randomly selected MR-CoNS isolates (n=51) were analysed using MICROBACT™ 12S (Oxoid) according to the manufacturer’s recommendations. Isolates were stored in BHI broth supplemented with 10% glycerol, at −70 °C.

DNA extraction and PCR procedures

LB broth (Merck) overnight cultures of the MRSA and MR-CoNS isolates were prepared. Chromosomal DNA was extracted according to the manufacturer’s instructions using the Wizard Genomic DNA purification kit (Promega, USA), with lysostaphin (Sigma, USA) at 25 µg/ml for the lysis step. MRSA isolates (n=37) and MR-CoNS isolates (n=298) were analysed by PCR in a PTC-100 thermocycler (MJ Research, USA) for the presence of nuc gene [15] and tuf gene [16], respectively. The multiplex PCR method [17] was used for the determination of SCCmec types
for all methicillin-resistant isolates and subtyping of SCCmec type IV was performed according to Milheirico et al. [18]. All MRSA isolates were analysed for the presence of Panton–Valentine leukocidin (PVL) encoding genes (lukS-PV, lukF-PV), γ-haemolysin [19], toxic shock syndrome toxin (tst) and enterotoxin (sea-see, seg-sej, sem-seo) genes [20].

The following were used as reference strains, MRSA strains: COL, ANS46, MW2, 8/6-3P, Q2314, JCSC4469, HAR22, WIS, HDE288 [17, 18], Bk2464 [21].

spa typing of Staphylococcus aureus

The polymorphic X region of the spa gene was amplified according to Shopsin et al. [22]. The PCR products were purified using a PCR product purification kit (Qiagen, Germany) and DNA sequenced (Macrogen, Korea). The obtained sequences were analysed using RidomStaphType software version 2.2.1 (Ridom GmbH, Germany) and spa types were assigned according to the spa server database (http://spaserver.ridom.de). The relationship of the spa types was analysed using the BURP (based upon repeat patterns) algorithm with default parameters within the RidomStaphType software package.

Antibiotic susceptibility testing

Antibiotic susceptibility of isolates was determined using a disc diffusion method as described previously [23] with the following antibiotics (Oxoid): penicillin (10 IU), oxacillin (1 μg) cefoxitin (30 μg), vancomycin (30 μg), gentamicin (10 μg), erythromycin (15 μg), clindamycin (2 μg) norfloxacin (10 μg), tetracycline (30 μg) and linezolid (30 μg). MR-CoNS isolates, which were erythromycin resistant and appeared to be susceptible to clindamycin, were further tested for possible inducible clindamycin resistance as described previously [23].

Statistical analysis

Categorical variables between groups were compared by means of Pearson’s χ² test [exact significance (two-sided)] using SPSS version 16 (SPSS Inc., USA) to evaluate the relationship between risk factors with the carriage rate of MRS, MRSA, MR-CoNS and MRSA+MR-CoNS co-colonization. A P value of <0.05 was defined as significant.

RESULTS

Population characteristics

The basic demographics of the volunteers screened by nasal and skin swabs are presented in Table 1.

<table>
<thead>
<tr>
<th></th>
<th>MRS (n=130)</th>
<th>MR-CoNS only (n=111)</th>
<th>MRSA only (n=7)</th>
<th>MR-CoNS + MRSA (n=12)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Females</td>
<td>99 (76.2)</td>
<td>83 (74.8)</td>
<td>5 (71.4)</td>
<td>11 (91.7)</td>
</tr>
<tr>
<td>Males</td>
<td>31 (23.8)</td>
<td>28 (25.2)</td>
<td>2 (28.6)</td>
<td>1 (8.3)</td>
</tr>
<tr>
<td>Hospitalized</td>
<td>14 (10.8)</td>
<td>12 (10.8)</td>
<td>0 (0.0)</td>
<td>2 (16.7)</td>
</tr>
<tr>
<td>Contact with HCW</td>
<td>36 (27.7)</td>
<td>29 (26.1)</td>
<td>3 (42.9)</td>
<td>4 (33.3)</td>
</tr>
<tr>
<td>Antibiotic exposure</td>
<td>41 (31.5)</td>
<td>36 (32.4)</td>
<td>2 (28.6)</td>
<td>3 (25)</td>
</tr>
</tbody>
</table>

MRS, Methicillin-resistant staphylococci; MR-CoNS, methicillin-resistant coagulase-negative staphylococci; MRSA, methicillin-resistant Staphylococcus aureus; HCW, healthcare worker.

* Statistically significant trend (*P < 0.05) calculated by Pearson’s χ² test.

<table>
<thead>
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<td>2 (28.6)</td>
<td>3 (25)</td>
</tr>
</tbody>
</table>

Table 1. Demographics of 227 healthy volunteer subjects and distribution of MRS, MR-CoNS only, MRSA only and MR-CoNS+MRSA isolates among these subjects

Carriage rates of community MRS

Initial biochemical identification of the colonies grown from 454 samples resulted in the isolation of
335 MRS from 130/227 (57.2%) subjects (Table 1). Of these, 37 isolates were MRSA and were recovered from 16 females and three males (Table 2). The prevalence of MRSA nasal carriage was 7.5% (17/227) and 8.4% (19/227) for both nasal and extranasal colonization (Table 2). The remaining isolates (n = 298) were identified as MR-CoNS and were isolated from 123 participants (carriage rate 54.2%), predominantly from nasal samples (Table 2). Of the 12 individuals with MRSA and MR-CoNS carriage, nasal co-colonization was detected in eight. None of the studied risk factors was significantly correlated with carriage in either group of staphylococci (Table 1).

The species-specific tuf gene was identified in 152/298 MR-CoNS isolates from 87 subjects and 51 of the tuf-positive isolates were additionally identified by MICROBACT. Eighty percent were MR-S. epidermidis (MRSE) with the remainder being S. capitis (7.8%), S. chromogenes (3.9%), S. hominis (3.9%), S. cohnii (2%) and S. haemolyticus (2%).

**Antimicrobial susceptibility**

All MRSA isolates were resistant to all β-lactam antibiotics, but susceptible to all other agents tested. The majority of tuf-positive MR-CoNS isolates were susceptible to gentamicin (97.4%), tetracycline (90.8%), norfloxacin (84.9%), and clindamycin (79.6%). Furthermore, 52.6% showed resistance to erythromycin. Inducible clindamycin resistance was detected in 11 (7.2%) of 54 clindamycin-sensitive/erythromycin-resistant MR-CoNS isolates. All MR-CoNS were susceptible to vancomycin and linezolid.

**SCCmec types for MRSA and MR-CoNS**

All MRSA isolates harboured the SCCmec type IV element of subtype e (81%) and subtype c. By contrast, MR-CoNS displayed great variability in SCCmec types (Table 3). Sixty-six of 152 isolates were of a single type comprising type IV (n = 59, of which 53 were MRSE), type V (five MRSE) and type VI (two MRSE). Four isolates harboured two types, IV + I (three MRSE), and IV + V (one MRSE). The SCCmec element in 82 of MR-CoNS (78 MRSE) could not be identified by PCR. The identified SCCmec IV subtypes in MR-CoNS were: subtype a (56%), subtype c (5%), subtype d (1.7%), subtype e (5%), subtype g (5%) or untypable (27.1%). For non-S. epidermidis species only MR-CoNS-IV or MR-CoNS-NT were recognized. Significantly, 3/8 subjects colonized with both groups of staphylococci yielded isolates harbouring SCCmec type IV. The other five subjects were co-colonized by MRSA-IV either with MRSE and/or MR-CoNS of SCCmec types V and IV + I.

**spa types of MRSA and virulence factors**

The spa type of the 37 MRSA isolates and their associated virulence factors are presented in Table 4. A single novel spa type, designated t9519, accounted for 76% of the isolates and only two other spa types were identified, i.e. t223 (14.7%) and t044 (5.9%). Two isolates were positive for PVL encoding genes lukS-PV-lukF-PV and belonged to SCCmec IV subtype c, spa type t044. All isolates harboured γ-haemolysin and tst genes, and were enterotoxin gene positive with the most common being seb (97.3%), seo (94.6%) and sei (91.9%). All isolates were negative for other enterotoxin genes tested: sea, sec, sed, see, seh, and sej. The PVL-positive strain carried only the seb gene.

**DISCUSSION**

Jordan had been reported as being among the countries with hyperendemic antimicrobial resistant bacterial strains. Previously, the Antibiotic Resistance in the southeastern Mediterranean (ARMed) project, reported Jordan as the country with the highest prevalence of significant clinical MRSA infections among the Mediterranean countries [24, 25]. This is expected to be correlated to antimicrobial misuse and overuse [26, 27] and probably also to a high carriage.
rate of this microorganism. Surprisingly, we were unable to identify epidemiological factors in this study population, including antibiotic exposure or close contact with a healthcare worker, that were associated significantly with carriage of either MRSA or MR-CoNS. Of interest, the reported nasal carriage rate of MRSA found here remains unchanged from that reported previously (7.6%) in non-medical Jordanian university students [7], whereas the geographically adjacent countries of Lebanon and Saudi Arabia have reported markedly lower carriage rates of 1.6% and 1.3%, respectively [28, 29].

To the best of our knowledge, the present study is the first to comprehensively assess the genotypic and phenotypic characteristics of MRSA isolated from healthy Jordanians in the community. Genotypically, they belonged to SCC\textit{mec} type IV, a type that is often associated with CA-MRSA infections, and is also associated with a minority of hospital strains including some reported from Jordan [8, 9]. This might imply that some strains are circulating and disseminating between the hospital and community.

We report here the identification of a novel \textit{spa} type (t9519) of MRSA with the methicillin resistance element SCC\textit{mec} IVe as the predominant type carried by healthy Jordanians. BURP analysis revealed a close relationship to t012 which is a common \textit{spa} type in Europe (spaserver.ridom.de). Interestingly, t044,

<table>
<thead>
<tr>
<th>SCC\textit{mec} type</th>
<th>Total</th>
<th>\textit{S. epidermidis (MICROBACT + tuf)}</th>
<th>\textit{S. epidermidis (tuf)}</th>
<th>\textit{S. capitis}</th>
<th>\textit{S. chromogenes}</th>
<th>\textit{S. cohnii}</th>
<th>\textit{S. haemolyticus}</th>
<th>\textit{S. hominis}</th>
</tr>
</thead>
<tbody>
<tr>
<td>IV</td>
<td>59</td>
<td>35 (59.3)</td>
<td>18 (30.5)</td>
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<td>2 (3.4)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>2 (3.4)</td>
</tr>
<tr>
<td>IVa</td>
<td>33</td>
<td>20 (60.6)</td>
<td>11 (33.3)</td>
<td>2 (6.1)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>IVc</td>
<td>3</td>
<td>2 (66.7)</td>
<td>1 (33.3)</td>
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<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
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</tr>
<tr>
<td>Iv</td>
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<td>1 (100)</td>
<td>0 (0)</td>
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<td>0 (0)</td>
<td>0 (0)</td>
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<td>0 (0)</td>
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<tr>
<td>IVe</td>
<td>3</td>
<td>1 (33.3)</td>
<td>2 (66.7)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>IVg</td>
<td>3</td>
<td>2 (66.6)</td>
<td>1 (33.3)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
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<td>0 (0)</td>
</tr>
<tr>
<td>NT*</td>
<td>16</td>
<td>8 (50)</td>
<td>4 (25)</td>
<td>2 (12.5)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>2 (12.5)</td>
</tr>
<tr>
<td>V</td>
<td>5</td>
<td>1 (20)</td>
<td>4 (80)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>VI</td>
<td>2</td>
<td>0 (0)</td>
<td>2 (100)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
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<tr>
<td>IV + I</td>
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<td>3 (100)</td>
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<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>IV + V</td>
<td>1</td>
<td>1 (100)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>NT†</td>
<td>82</td>
<td>4 (4.9)</td>
<td>74 (90.2)</td>
<td>2 (2.4)</td>
<td>0 (0)</td>
<td>1 (1.2)</td>
<td>1 (1.2)</td>
<td>0 (0)</td>
</tr>
<tr>
<td>Total</td>
<td>152</td>
<td>101</td>
<td>4</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>2</td>
<td></td>
</tr>
</tbody>
</table>

Values given are n (%).
* NT, Untypable by Milheirico et al. [18].
† NT, Untypable by Milheirico et al. [17].

Table 4. Resistance element, \textit{spa} type and virulence gene profile patterns of MRSA carriage isolates

<table>
<thead>
<tr>
<th>\textit{spa} type</th>
<th>Isolate no.</th>
<th>SCC\textit{mec} type</th>
<th>Presence (+) or absence (−) of gene</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>plv</td>
</tr>
<tr>
<td>t223</td>
<td>3</td>
<td>IV†</td>
<td>−</td>
</tr>
<tr>
<td>T223</td>
<td>1</td>
<td>IV†</td>
<td>−</td>
</tr>
<tr>
<td>T223</td>
<td>1</td>
<td>IV†</td>
<td>−</td>
</tr>
<tr>
<td>T044</td>
<td>2</td>
<td>IV e</td>
<td>+</td>
</tr>
<tr>
<td>T9519</td>
<td>25</td>
<td>IV e</td>
<td>−</td>
</tr>
<tr>
<td>T9519</td>
<td>1</td>
<td>IV e</td>
<td>−</td>
</tr>
<tr>
<td>T9519</td>
<td>1</td>
<td>IV e</td>
<td>−</td>
</tr>
<tr>
<td>NT*</td>
<td>2</td>
<td>IV e</td>
<td>+</td>
</tr>
<tr>
<td>NT*</td>
<td>1</td>
<td>IV e</td>
<td>+</td>
</tr>
</tbody>
</table>

* Unable to identify the \textit{spa} type.
† Untypable by Milheirico et al. [18].
which is a widely distributed *spa* lineage of HA-
MRSA in European [5] and some Middle Eastern
countries, including Jordan [9, 30], was found in
the present study in two healthy individuals.

The susceptibility profiles of the MRSA isolates
recovered in the present study are also consistent
with a community origin being susceptible to all
antimicrobials with the exception of the β-lactams
tested [1]. It has been suggested that antimicrobial
susceptibility-based classification of CA-MRSA lacks
sensitivity and the widely used marker of cipro-
floxacin susceptibility may miss approximately
one-third of these strains [31]. Nevertheless, PVL, a
common marker of CA-MRSA, was quite rare in our
isolates in accord with reports from both Japan and
Ireland [32, 33]. The carriage of at least three viru-
ence factor encoding genes suggests that the majority
of our MRSA isolates have the potential to cause in-
fections as the combinatorial effect of virulence factors
is strongly linked to their potential involvement in
severe infections [34].

We present, for the first time, data on the preva-
ience and molecular epidemiology of MR-CoNS in
healthy Jordanians. Consistent with previous reports,
the SCC*mec* elements in MR-CoNS exhibited genetic
diversity where some isolates harboured two SCC*mec*
types and others were untypable [3, 35]. SCC*mec* type
IV, a relatively common type in MR-CoNS of com-
munity origin [4, 35], prevailed among our *tuf*-positive
MR-CoNS isolates which proved to be mostly MRSE
[36]. In addition other MR species commonly isolated
from human clinical specimens, such as *S. capitis*,
*S. cohnii*, *S. haemolyticus* and *S. hominis*, were identi-
cified as nasal and skin colonizers. Of note, in our
community pool of MR-CoNS isolates, MRSE-IV
and MRSE-NT prevailed, and co-harbouring of
different SCC*mec* elements (MRSE-IV+V, MRSE-
IV+I) was observed, a finding consistent with MRSE
genome plasticity in the community [37]. In agree-
ment with Garza-Gonzalez *et al.* [3], antibiotic sus-
ceptibility patterns in MR-CoNS isolates are diverse
and this is clearly associated with the heterogeneity of
SCC*mec* types present in these strains.

Resistance to β-lactams in MR-CoNS is encoded
by SCC*mec* elements which have high homology with
certain types of SCC*mec* of MRSA [4]. Such high
homology might suggest a common origin and indi-
cate a probable horizontal cross-transmission of re-
sistance genes upon co-colonization [10, 11]. Thus,
the sharing of the same SCC*mec* type between most
of the co-colonized MRSA and MR-CoNS, might
suggest *in vivo* horizontal gene transfer but this would
need to be confirmed by further genetic characteriza-
tion not only of these groups but also of co-resistant
methicillin-sensitive *S. aureus*.

This study has some limitations. We were not able
to estimate the true prevalence of CA-MRSA infec-
tion in the general Jordanian population and wider
based surveillance studies in different sections of local
and regional geographical sectors are warranted to
achieve this. Furthermore, the socio-demographic
characteristics and the risk factors associated with
MRSA acquisition were not analysed or compared
with non-colonized subjects. Due to the spread of
various MRSA clones between the community and
hospital [38], and across national boundaries [38], a
full molecular characterization of strain lineages by
the internationally validated MLST system in order to
define them in a global epidemiological context is of
importance in both the clinical and infection control
setting [39].

As successfully demonstrated in countries or re-
regions with low, decreasing or stabilized MRSA
infection rates [40], it is of utmost importance to de-
velop comprehensive surveillance and prevention
strategies that will lead to effective and robust control
not only of MRSA but also other multidrug-resistant
infections. Major elements of such strategies include
implementation of active screening procedures adap-
ted to the specific endemic situation, adherence to
basic infection control practices, introduction and
control of microorganism-specific hygiene measures,
and antibiotic control programmes [41].

In conclusion, this study is the first to demon-
strate a high incidence of MRS nasal and skin carriage
in healthy Jordanians, a finding consistent with a com-
munity origin for these organisms. We found a high
prevalence of MRSA in the community compared to
other countries in the region and that most isolates
are of the same genotype and share the same β-lactam
antimicrobial resistance mechanism. In addition we
identified nasal co-colonization with both MRSA and
MR-CoNS with highly similar resistance genes in a
small number of subjects which raises the possibility
of horizontal resistance gene transfer between the
different groups of staphylococci.

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DECLARATION OF INTEREST
None.

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