Methicillin Resistant Staphylococcus aureus and public fomites: a review

Ziad W Jaradat^a, Qutaiba O Ababneh^a, Sherin T Sha'aban^b, Ayesha A Alkofahi^b, Duaa Assaleh^b and Anan Al Shara^b

^aJordan University of Science and Technology, Irbid, 22110, Jordan; ^bDepartment of Applied Biological Sciences, Jordan University of Science and Technology, Irbid, 22110, Jordan

ABSTRACT

Staphylococcus genus is a Gram-positive coccus normally associated with skin and mucous membranes of warm-blooded animals. It is part of the commensal human microflora, or found in animals, or contaminating surfaces in the community and hospital settings. *Staphylococcus aureus* is the most pathogenic species belonging to this genus, as it possesses a collection of virulence factors that are expressed solely to evade the immune system. The increase in the misuse of antimicrobial agents predisposed *S. aureus* to develop antibiotic resistance, including the resistance to methicillin which led to the emergence of Methicillin-Resistant *S. aureus* (MRSA). MRSA is considered one of the most dangerous nosocomial pathogens causing many hard to treat infections in hospitals and was named as Hospital Associated MRSA (HA-MRSA). Over the past 20–25 years, MRSA was isolated from community settings and thus Community Associated MRSA (CA-MRSA) has emerged. Inside hospitals, MRSA has been isolated from fomites in contact with patients, as well as staff's protective and personal items. This review highlights the worldwide prevalence of MRSA on fomites within the contexts of hospital and community settings.

KEYWORDS

Methicillin resistance; Staphylococcus aureus; public fomites; Hospital Associated; Community Associated; review

Taylor & Francis

Check for updates

Taylor & Francis Group

Introduction

Staphylococcus species are Gram-positive, non-motile, non-spore forming microorganisms. *S. aureus* is the most pathogenic strain among this species with a potential to cause a wide range of diseases both in communities and hospitals [1].

As a commensal microbe, *S. aureus* mainly colonizes the nasal cavity of humans and many animals [2]. In addition, *S. aureus* is found on the skin, inside oral cavity, upper respiratory tract, lower urogenital tract, and gastrointestinal tract of humans [3]. In fact, 25–30% of people are permanently colonized with *S. aureus* and about 60% of the population are transiently colonized with this pathogen [4].

S. aureus possesses several virulence factors that allow the pathogen to thrive in diverse host environments and to survive extreme conditions [5]. *S. aureus* produces a set of cell-surface and secreted virulence factors such as enterotoxins and hemolysins which play a role in enhancing its pathogenicity [5,6]. Due to its wide distribution in the environment surrounding humans, *S. aureus* is considered one of the most harmful human pathogens implicated in several diseases ranging from mild skin-related infections to more severe life-threatening and systemic infections, such as bacteremia [3,7,8].

S. aureus was one of the first pathogens to develop resistance to penicillin shortly after its introduction [9].

In 1959, a new semi-synthetic β -lactam antibiotic, called methicillin was introduced to replace penicillin. However, only 2 years after its first use (1961), several cases of *S. aureus* strains resistant to methicillin were reported. These strains were also found to be resistant to other β -lactam antibiotics including oxacillin, cefoxitin, and other antibiotic families. Later, strains with this spectrum of resistance were given the name, methicillin-resistant *Staphylococcus aureus* (MRSA) [10,11].

Resistance to methicillin in *S. aureus* is primarily mediated by *mecA* gene carried on a mobile genetic element called staphylococcal chromosomal cassette (SCCmec) which is highly diverse and is classified into 13 types [12]. The *mecA* gene encodes a modified, low-affinity penicillin-binding protein (PBP2'), which confers cross-resistance to almost all β -lactam antibiotics [9,13].

Traditionally, MRSA has been considered a nosocomial pathogen responsible for health-careassociated (HA-MRSA) infections in patients, hospital personnel, and their associates [14]. However, the 1990s marked the emergence of MRSA within households as well as community facilities such as schools, daycares, geriatric homes, prisons, recreation centers, and many other institutions or settings [14,15].

Over the years, scientists have identified certain *S. aureus* lineages to be more prevalent in certain geographic areas. These linages are circulating all

CONTACT Ziad W Jaradat 🔯 jaradatz@just.edu.jo 🗈 Department of Biotechnology and Genetic Engineering, Jordan University of Science and Technology, P. O Box 3030, Irbid, Jordan

over the globe. Therefore, they were named so that they can easily be tracked. These lineages include the clonal complex 5 (CC5)/USA100, CC30/USA200, CC8/ USA300, CC1/USA400, and CC45/USA600 [16]. Each lineage is endemic in a certain area; for example, *S. aureus* USA 100 is prevalent in the USA, both in health and community settings. *S. aureus* USA 400 is community-associated and is predominant in southern Alaska [16]. Further, several other lineages were detected in other parts of the world; St 22 and St772 in India [17], St 80 in Europe and Middle East [18], St 59 in Taiwan [19] and St 72 in Korea [20].

In the United States, the USA 300 lineage is the most well-studied clonal group. It is associated with the production of high levels of cytotoxins such as α-toxin, arginine catabolic mobile element type 1 and a set of virulence genes including lukS-PV/lukF-PV, sek, and seq [16,21]. And thus, it is incriminated in most of the community-associated MRSA infections causing severe skin and soft-tissue infections [16]. USA300 has been classified as one of the highest pathogenic strains [22], as it became an increasingly common cause of health-care-associated MRSA infections as well [23]. Interestingly, a variant of the USA300 called USA300 LA is predominant in some countries in Latin America such as Columbia and Ecuador [24].

Community-Associated MRSA (CA-MRSA) strains are genetically distinct from HA-MRSA strains [25]. CA-MRSA is mainly associated with skin and soft-tissue infections [26]. For instance, Carrel et al. (2015) reported in their review that among isolates with any reported anatomic site of isolation, skin, and soft-tissue infections accounted for 62.6% of USA300 and 19.1% of USA100 isolates. In addition, these, PVL-positive CA-MRSA causes severe diseases such as necrotizing pneumonia, and bacteremia [23].

CA-MRSA possesses SCCmec type IV, producing the virulence factor Panton-Valentine leukocidin (PVL), but frequently exhibit a non-multi drug resistance profile [13,27,28].

Certainly, fomites play a role in the transmission of CA-MRSA infections where inanimate objects are considered the potential reservoir and the source of these infections [29]. Pathogens may be transferred directly through the surface to mouth contact, or indirectly by contaminating hands/fingers, which subsequently transfer the pathogen to mouth, eye, ear, nasal cavities, or genitals [30–33]. Body fluids from infected areas may serve as a source of the transmission of the pathogen to fomites again [15,34–36]. Several studies have shown that MRSA survives on fomites for several hours, days or even months depending on the number of cells deposited and on other conditions related to the microstructure of fomites surfaces and environmental conditions [37–39].

This review will focus on the different types of community and hospital fomites known to harbor MRSA and their role in the widespread transmission of the CA-MRSA lineages and HA-MRSA nosocomial infections. The mode of transmission of MRSA from fomites to humans, and the methods to control MRSA colonization and transmission as well as the risk factors will be also addressed.

Community-associated methicillin-resistant Staphylococcus aureus

Community-associated MRSA (CA-MRSA) is defined as MRSA strains isolated from individuals who have not been recently exposed to the health care from patients exhibiting MRSA infections or having an infection incubation period at the time of admission to a healthcare facility [40]. Two decades ago, MRSA infections were confined only to healthcare systems, and were dubbed Hospital associated-MRSA, until the emergence of CA-MRSA in 1990s [41]. Interestingly, CA-MRSA is not limited to causing infections in community settings, but rather, they have been found to cause infections in hospitals and other healthcare settings [42]. CA-MRSA is now considered a public health problem in some parts of the world due to the increase in the number of infections caused by this pathogen [29]. Unlike HA-MRSA, which is associated with major and lifethreatening infections, CA-MRSA usually causes minor skin and soft-tissue infections [26,43]. Nevertheless, CA-MRSA infections were reported to cause severe lifethreatening illnesses such as pneumonia, pelvic osteomyelitis, septic thrombophlebitis, ocular infections, and necrotizing fasciitis [44]. The following parts of this review will shed light on the prevalence of MRSA in different community settings with special emphasis on the risk factors associated with these settings.

Athletic communities

The prevalence of MRSA infections among athletes in different types of sports and fitness centers is reviewed.

MRSA plays a major role in causing skin infections among athletes due to multiple risk factors associated with sport communities and athletes [45]. Several risk factors, such as physical contact between athletes, poor hygiene following physical activity and contaminated surfaces of fitness centers and training facilities, have greatly contributed to this problem [46]. In addition, multi-user equipment, high-contact surfaces, skin-to-skin contact, carpeted areas, locker rooms, and training areas are also among the major risk factors [46]. Furthermore, some risk factors are associated with certain types of sports. In football for example, skin abrasions, turf burns, and player tactical positions all contribute to the risk of obtaining an MRSA infection [47]. Indeed, numerous reports documented such infections in the last two decades [47-53]. Several other outbreaks of *Staphylococcus* infections in athletes and athletic settings were recently reported as outlined in the Table 1.

Emergency medical services and ambulances

Emergency medical services (EMS) settings and its personnel have also been colonized with a battery of microorganisms including MRSA. EMS personnel are at high risk of acquiring pathogens in the community and transmitting them to patients during medical emergencies [64-66]. In EMS settings, patients and EMS personnel occupy the same small area within the cabin of ambulances, and therefore, increasing the risk of MRSA transmission between patient, EMS personnel, and ambulance interior environment. In addition, some EMS systems suffer from a high rate of patient turnover, which means that subsequent patients will occupy the same cabin within a short period of time, and therefore, leaving a limited time for cleaning of the ambulance's interiors and equipment before reporting to duty. Such practice will certainly increase the risk of MRSA transmission. Several studies reported the colonization of MRSA in samples collected from ambulance's cabin interiors and equipment. In a study conducted on ambulances in the Chicago metropolitan area, 5 out of 71 ambulances tested in early 2010 were found contaminated with MRSA [67]. Similarly, MRSA was isolated from 2 out of the 17 ambulances investigated in July 2012 in a Spanish study conducted in Bilbao city [68]. In contrast, Roline et al. (2007) and Brown et al. (2010) documented the presence of MRSA in almost 50% of ambulances investigated in the West Coast and Southern Maine, respectively [69,70]. A recent study from Egypt reported that 46.1% of the S. aureus strains isolated in early 2016 from 25 ambulance cabins were MRSA [71]. In all these studies, MRSA was detected in both the patient-contact sites and the non-contacted sites inside the ambulance cabin.

Other similar studies reported lower percentages of MRSA contamination, for example, one study from Germany conducted in 2009 detected MRSA contamination only on a patient stretcher in 8 out of the 89 ambulances after the transport of MRSA-colonized /infected patients [72]. Likewise in California, USA, Kei, and Richards (2011) tested 40 samples collected in the summer of 2006 from many surfaces and devices inside the emergency department for the presence of *S. aureus* and reported only one sample (2.5%) to be positive for MRSA [73].

Fire stations

The communal lifestyle inside fire stations and the potential contact with high-risk populations puts fire-fighters at a high risk of exposure to pathogens. A study conducted in 2010 on environmental surfaces

I and I and I and a difference	table 1. Janimary of the recently papiloned reports on million prevarince more animary national and million services.			
Country of report	Sample number	S. aureus isolated	MRSA percentage	Reference/Year
Ohio, USA	81 environmental samples from 9 High school training facilities and wrestling mats	ND	61.7%	[54]/(2010)
Ohio, USA	90 surface samples from 10 high school training facilities and related locker rooms	MD	42%	[55]/(2010)
Florida, USA	240 samples from university recreational facility, a high school gymnasium, and a private gymnasium	None	None	[56]/(2011)
Texas, USA	125 from various areas and equipment from athletic facility	38%	6%	[54]/(2012)
Virginia, USA	99 environmental samples from fitness center	10 samples	None	[57]/(2012)
Thohoyandou, South Africa	500 from hand touched surfaces inside 3 fitness center	36%	1.6%	[58]/(2046)
Tennessee, USA	32 from skin-contact surfaces inside 4 membership-based fitness centers	90.6%	37.5%	[59]/(2016)
Ohio, USA	288 swabs on fomites	38.2% of which 36.4% MDR	30%	[60]/(2019)
16 fitness centers				
Malaysia, various fitness centers	42 swabs on fomites	73.81%	ND	[61]/(2018)
USA	Multiple samples on surfaces	ND	24% of surfaces were MRSA and VRE +ve	
Ohio, USA	288 environmental swabs from 16 fitness facilities	38.2%	11.5%	[60]/2019
Ohio, USA	280 environmental samples from 10 playgrounds	31.8%	3.9%	[63]/2019

in fire stations and training sites in Arizona, USA revealed that the highest prevalence of MRSA was on couches (20%; 4/20); class desks (10%, 1/10); and commonly touched surfaces in offices (6.7% 2/30) [64]. In another study conducted in 2011, MRSA was isolated from 8% of the 653 samples swabbed from 33 different fire stations in Washington State, USA. Among the tested fire stations, 19 of 33 (58%) were positive for MRSA [74]. Likewise, another research group isolated MRSA from 44 of the 1,064 samples (4.1%) collected in 2011 from two fire stations in two independent districts in the northwestern United States [75]. In the last two studies, most of MRSA isolates were recovered from surfaces in the living rooms and garbage disposal areas.

Public restrooms

Restrooms or washrooms are considered shared public spaces with clear pathogen transmission potential [76]. Factors that are likely to contribute to the spread and persistence of pathogens inside restrooms include the presence of human feces, temperature, humidity, and the inappropriate use of disinfectants [77]. Several studies documented the isolation of MRSA from restroom floors, handles, toilet seats, doorknobs, and water faucets. Roberts et al. (2011) reported the isolation of MRSA from university dormitory bathroom floors, toilet flush handles, light switches, and doorknobs [78]. In a study conducted during the Muslim pilgrimage (Hajj) season in Saudi Arabia, swabs were collected from doorknob surfaces of 224 toilets serving hundreds of thousands of people. Out of the 42 S. aureus strains recovered, four (10%) were identified as MRSA [79]. In another study, 32 staphylococcal strains were isolated from 18 public washrooms in London, UK [77]. Three of these strains were identified as EMRSA-15 clone. In UK, the MRSA-15 clone is frequently isolated from patients with bacteremia caused by MRSA, which indicates that infection control measures failed to limit the spread of such clones in both hospital and community environments.

MRSA was also isolated from hospitals restrooms, especially the ones that might be shared between patients, staff, and visitors. In a study conducted in 2009 in a children's cancer hospital in Tennessee USA, daily samples were collected for 4 weeks from toilet seats in restrooms equipped with alcohol wipes and from restrooms lacking the alcohol wipes [80]. MRSA was isolated from 3.3% of samples collected from toilets lacking the alcohol wipes while no MRSA was isolated from the toilets equipped with wipes, indicating the significance of such simple measures in controlling the spread of this pathogen. A similar study conducted early 2012 in Japan investigated the presence of bacteria in bidet-type toilets at a universityaffiliated hospital [81]. Of the 292 bidet-toilet seats sampled, MRSA was only found in one water-jet nozzle and one toilet seat. The results of these studies indicate that hospital toilets are potential risk to patients who may acquire MRSA from colonized persons. In addition, they represent a potential reservoir for nosocomial spread and serve as foci for community-acquired, hospital-related strains.

Public beaches

Beaches are considered a potential source of community-associated S. aureus infection, which is evident by the correlation between gastrointestinal illness, the ear, skin, and eye infections among bathers, and the density of S. aureus and other Staphylococcus species [82]. Bathers can shed S. aureus into water and sand, and S. aureus concentrations are correlated with the density of bathers and are attributed to human activities such as eating, playing, and littering [83], as well as other sources including stormwater [84], wastewater [85], sea mammals and domestic pets [86]. Several studies documented the isolation of MRSA from beach water, wet sand, and dry sand (Table 2) [82,87–96]. The results from all these studies indicate that beaches are potential reservoir for transmission of MRSA to bathers, especially those with skin lesions.

Homeless shelters

Closed community settings are considered areas of high-risk for CA-MRSA transmission, colonization, and infection due to increased person-to-person contact [97]. Homeless shelters are a very good example of such high-risk community settings. Crowdedness, lack of hygiene, and the frequent use of the facilities makes homeless shelters potential reservoirs for MRSA as well as a source of other pathogenic microorganisms [46]. In agreement with this, a study conducted between July 2012–June 2014 on a homeless shelter in Kansas City showed that individuals in

this homeless shelter are at high risk of obtaining an MRSA infection. In addition, the risk of infections within such individuals was significantly higher than the general population with a prevalence of 9.8% in homeless groups compared to 1.8% in the general population [97]. Another study conducted in April, 2006 in Canada revealed a 4.5% prevalence of MRSA colonization among residents of three homeless shelters in Ottawa, Ontario [98]. Poor hygiene, lack of sanitation, and immune compromise due to HIV and malnutrition are the most important risk factors contributing to the spread of MRSA infections among residents of such facilities [99].

Further, two studies conducted in USA by Landers et al. (2009) and Leibler et al. (2017) investigated the prevalence of *S. aureus* and MRSA nasal colonization in non-hospitalized homeless individuals in 2009 and

		Reference, Time of sample
Sample source	Note	collection
Dry, wet and inundated sand from subtropical beach in Florida, USA	5 MRSA CFU per g of dry sand	[87](08/2009)
Sand samples were collected from 37 Californian beaches	One beach sample tested positive for MRSA	[92] (10/2009)
Sand from 2 beaches; Avalon and Doheny in California, USA.	MRSA was detected in 11 out of 155 sand samples collected over two years	[83] (08/2007)
Public intertidal beach sand and marine water from Washington State, USA	5 MRSA were isolated	[90] (02–09/2008)
Sand from 4 beaches (Avalon, Doheny, and Malibu Surf rider) in California, USA.	MRSA was detected in 10 samples (2.7%), out of 366.	[82] (05–09/2009)
Dry and wet sand from marine water beaches and freshwater beach in the Seattle WA	Dry and wet sand from marine water beaches and freshwater beach in the Seattle WA MRSA was isolated from one dry sand sample [1.9%; n = 53] and six from wet sand [14%; n = 43]	[91] (06–08/2010)
Sand from Marine Mammal Conservancy beaches, three recreational beaches, a residential beach in Florida. USA	MRSA detected in 7 sand samples out of 11 isolates from environmental samples	[93] (06–08/2011)
Sand samples from subtropical recreational beach from South Florida, USA.	3 out of 36 (8.3%) sand samples were positive for MRSA	[94] (07–08/2009)
Recreational marine sand and fresh water beaches from Seattle, USA.	Thirty-one (10.5%) of the 296 recreational beach samples were positive for MRSA	[88, 2010]
Intertidal beach sand in the Eastern Cape Province of South Africa	One out of 67 samples was positive for MRSA	[95] (04/2015-04/2016)
Water and sand from 10 freshwater recreational beaches in Northeast Ohio, USA	MRSA was detected in 15 (7.1%), of 210 samples	[96] (06–11/2014)

2015, respectively, and reported a range from 8.3% to 26.5%. This high prevalence of MRSA infections was attributed to medical risk factors including recent use of antibiotics, renal failure, endocarditis, and to behavioral factors, like alcoholism, smoking, and probably narcotics use [100,101].

Daycare centers

As discussed earlier, the prevalence of CA-MRSA is diverse, and multiple risk factors could contribute to the spread of infections. Children at daycare centers can also be reservoirs for MRSA and can further increase the spread of these pathogens in the community.

Several studies showed that children at daycare centers are at a low risk of obtaining MRSA infections. Nonetheless, restricted health care and hygiene strategies should be implemented to prevent the spread of MRSA infections among children and their families. Probably the shared use of toys and other objects by children might be the reason behind the incidence of MRSA among this group.

When surfaces in daycare centers were sampled, MRSA was isolated. In a study conducted between Feb 2009 and Feb 2010 on a child care center in Iowa, MRSA was isolated from only 3 out of the 214 surface samples collected [102]. In another study conducted by Ryan et al. (2013) in two daycare centers serving staff of two academic institutions in Florida, no MRSA was isolated out of the 87 surface swabs tested [103]. Indeed, the very low MRSA prevalence in these daycare centers reflect the implementation of high hygienic standards.

Prisons

Infections and outbreaks of CA-MRSA occur in correctional facilities, such as jails and prisons, due to poor hygienic practices, crowdedness, low educational status of prison inmates and the low hygienic quality of these facilities [104,105].

In a study conducted by Felkner et al. (2009) in Texas, USA, 132 swabs were collected from jail surfaces, from which 8 (6%) MRSA isolates were recovered [106].

A study conducted in 2004 aimed to examine the incidence of MRSA infections in a Texas prison population, and to identify the risk factors associated with these infections, showed an incidence rate of 12 MRSA infections per 1000 person out of a total of 299,179 inmates (both males and females) between 1999 and 2001 [107]. This study correlated elevated rates of MRSA infections between inmates and risk factors including the health of the inmates, circulatory diseases, cardiovascular diseases, diabetes, end-stage liver disease, human

immunodeficiency virus infection, or acquired immunodeficiency syndrome, and skin diseases [107]. Furthermore, the heavy usage of prison fomites and surfaces accompanied by low hygienic practices among prisoners certainly predisposes the users to infections including MRSA.

Unfortunately, the spread of MRSA infections through correctional facilities is difficult to control, nonetheless understanding the risk factors may help in the prevention of MRSA infections and to control its transmission, before a chronic problem that disseminates MRSA clones to the outside community happens [107].

Hotels

Hotel rooms and amenities are potential sources of community-associated infections due to the high number of quests passing through such settings, with the consequence of acquiring or transmitting of infectious agents [108]. In addition to that, hotels cleaning protocols are based on visual rather than microbial assessments [109]. Foodborne and waterborne infections and outbreaks in hotels have been documented by numerous reports. However, very few studies investigated the contamination of contact surfaces for the presence of infectious agents within hotel rooms and amenities [110]. Although no MRSA was isolated in their study, Xu and colleagues (2015) isolated multidrug-resistant Staphylococcus species from samples collected from inanimate objects inside rooms of three large hotels in London, UK. The authors of this study concluded that these Staphylococcus species represent potential reservoirs for MRSA [111]. In contrast, a study done in Canada in summer 2012 reported the presence of MRSA on high-contact surfaces inside 54 hotel rooms from 6 different hotel chains [108]. MRSA was isolated from the comforter, TV remote, bedside lamp, telephone, bathroom countertop, faucet, and toilet seat.

Academic institutions

School and university environments contain many high-contact surfaces frequently touched by thousands of students and staff on a daily basis. In such environments, close physical contact, common shared spaces, and variable hygiene habits among students are some factors likely to contribute to the transmission of infectious agents. MRSA have been isolated from a wide range of high-touch surfaces within schools and university environments, such as class-rooms [112,113], restrooms [78,88,112,113], lockers [78,88], elevators [78,88], athletic training facilities [114,115], dormitory fomites [78,88,116] as well as public computers [117,118]. In addition, student's personal items have been found contaminated with MRSA including cellphones [119] and door keys [120]. Details of the MRSA prevalence in these places can be reviewed in the cited references.

Nursing homes

Elderly individuals in nursing care facilities are at an increased risk of acquiring MRSA due to their premorbid conditions, frequent hospitalization, shared rooms, and common areas [111,121]. Several studies reported the colonization of MRSA in nursing care facilities and residents as well [111,121-128]. MRSA have been also isolated from inanimate objects inside nursing facilities. For example, in Oct/Nov 2006, MRSA was isolated from 2 (1.2%) of the 163 environmental specimens collected from fomites in private and common rooms inside an elderly long-term facility located in USA [127]. These MRSA strains also colonized the residents in the contaminated rooms. In another study conducted in 36 residential elderly care homes in Hong Kong in the year 2010, more than 400 environmental samples were collected [121]. MRSA was isolated from bedside tabletops (19.4%), hand covers (15.6%), commodes (11.4%), sofas (8.6%), armchair handrails (8.3%), soap dispensers (5.7%) and wheelchairs (5.6%). In the same study, 20.4% of nasal swabs collected from 2776 residents tested positive for MRSA, which explains the presence of the MRSA on fomites and other surfaces in the facility.

Mobile phones and pagers

Mobile phones are increasingly becoming an inherent part of human's life. A major problem of using such devices is that they are rarely cleaned by the majority of their users, and thus, collect many types of bacteria and could act as reservoirs for the transmission of many pathogens [129–130]. Interestingly, Morris et al. (2012) conducted a one-day microbiological survey of hospital doctors fingerprints before and after using their mobile phones [131]. Among the 20 doctors enrolled in the study, the fingers of five doctors were contaminated with *Staphylococcus* species including MRSA. This indicates that doctors themselves can spread MRSA to patients.

Many researchers in different countries conducted studies on the prevalence of MRSA on mobile phones and pagers used by many health-care personnel and reported varying degrees of microbial contamination including MRSA. Table 3 lists a number of such studies. These studies indeed indicate the potential danger that mobile phones pose on the health of hospitalized patients and certainly shed light on the possible reasons behind the widespread of nosocomial infections inside hospitals. These studies also signify the difficulties encountered by hospital infection control personnel to eradicate nosocomial infections as mobile phones have

	No. of phones contaminated	No. of phones contaminated		Country of	
Phone belongs to	with bacteria (%)	with MRSA (%)	Facility	study	Reference
Hospital doctors and staff (132), college faculty and staff (54), medical students (100) and control group (100)	316/386 (81.8%)	16 (4%)	Hospital	India	[130]
Nurses, 32 (17.5%) from laboratory workers, and 57 (31.1%) from health care staff	179/183 (97.8%)	17 (9.5%)	Hospital	Turkey	[129]
HCWs ¹	157/213 (73.7 %)	1.4%	Intensive care units (ICUs), pediatric intensive care units (PICUs), and neonatal care units (NCUs)	Kuwait	[134]
Health care personnel	65/120 (54.2%)	11(16.4%)	General	India	[135]
Hands and mobile phones of HCWs		16/27 (59%)	General	India	[136]
Mobile phones of HCWs	13/101 (13%)	4/101 (4%)	General	Korea	[137]
Mobile phones with HCWs	144/200 (72%)	26/200 (13%)	General	India	[138]
Pagers of HCWs	12/100 (12%)	3 (3%)	General	USA	[139]
Students	107/309 (34.6%)	ND^{2}	University	Japan	[140]
Patients, patients' companions, visitors,	121/133 (90%)	ND	Hospital	Turkey	[141]
HCW	58/67 (85.6%)				
Patients & HCW	40/40 (100%)	21 (53%)	Hospital	Egypt	[142]
Veterinary students, technicians, residents/interns and clinical faculty	3/123 (1%)	1/123 (0.8%)	Veterinary teaching hospitals	Canada	[143]
¹ HCW; health-care workers ² ND: not determined					

Table 3. Summary of the prevalence of pathogens on mobile phones.

become so indispensable, subjected to no rules for their use, and used freely within the health-care facilities.

MRSA prevalence in paper currency and coins

Paper currency and coins are frequently handled by many people with varying health and hygienic standards. Therefore, paper currency and coins can pose a public health risk especially when they are handled by individuals with direct contact with food, such as in restaurants and bakeries. MRSA can survive for a long time on paper currency and coins surfaces, making these surfaces potential reservoirs of infection [132].

In a study conducted by Gedik and coworkers (2013), the survival of MRSA and other bacteria on selected paper currencies from different countries was tested. Interestingly, it appeared that the types of fabric used to make paper currencies play a major role in supporting the survival of bacteria. For instance, some paper currencies like the Romanian Leo, US, and Canadian dollars supported confluent growth while some currencies like Indian Rupees, Euros, Morocco Dirhams, and Croatian Kunas did not support any bacterial growth or survival. In addition, some types of the currency fabric promoted the transmission of bacteria to handlers while other fabric types did not promote any transmission. The Romanian Leo was the most effective in transmitting S. aureus from the currency to volunteers followed by US dollar, while the Euro did not lead to any transmission [133]. Similarly, Tolba et al. (2007) tested the survival of several MRSA strains on metal coins and reported that no epidemic nosocomial or community-associated MRSA survived on coins lightly heat treated. It was concluded from this study that when no organic protection is offered, no MRSA will survive; however, bacteria may survive well when soil, pus, and blood is present on metal coins, therefore offering protection from drying and other environmental conditions [144]. In the same token, several researchers [39,145] reported that many Gram-positive and Gram-negative bacteria (see references for list of bacteria) can survive for months on surfaces that contain some organic matter, which might serve as a shield that helps in their survival. In a study from Nigeria conducted in Feb/Mar 2012, 128 paper currencies were collected from meat sellers and 36 (28%) of the tested bank notes were found contaminated with MRSA, while none of the very new currency notes contained MRSA [146]. In a similar study conducted by the same researcher (Neel, 2013), of the 205 different currency notes collected from a market place (hotels and restaurants), 53 (26%) and 6 (3%) were found contaminated with S. aureus and MRSA, respectively [147]. However, a meta-study conducted by Angelakis et al. (2014) reported the presence of multiple types of bacteria (MRSA was not specified) on paper currency and coins issued by many countries [132].

ATMs

Automatic teller machine (ATM) surfaces are likely to be contaminated with bacteria due to their direct contact with hands from multiple users. People with different socioeconomic backgrounds and hygienic statuses use ATMs on a daily basis. The point of contact is the user's fingers touching the keypad and/or screen surfaces. Several studies reported the isolation of low levels of MRSA from ATM surfaces [78,148–150]. Other studies reported a high prevalence of colonization of ATMs with coagulase-negative *S. aureus* [151,152].

MRSA and public transportation

Millions of people use public ground transportation on a daily basis. Thus, ground transportation serves as one of the favorable routes of microbial transmission. Users touch and contaminate objects within transportation vehicles such as door handles, seats, windows and they may even drop contaminated items on the floor such as tissue. This certainly creates a potential reservoir for multiple types of microbes, some of which might be very pathogenic, while others might be opportunistic or nonpathogenic [153]. Table 4 summarizes the prevalence of MRSA in public transportation vehicles and waiting stations reported by several studies.

Hospital associated MRSA and fomites

Nosocomial or healthcare-associated infections (HAI) are infections acquired while receiving medical care at a health facility but were not present at the time of admission. In the year 2011, The World Health Organization reported that on average 7–15% of the population in developed and underdeveloped countries, respectively, suffer from HAI at any given time with a mortality rate at nearly 10% [165].

These infections have severe consequences on the life status of patients and are considered a high financial burden for health-care systems [166–169]. The discussion of HAI is often associated with MRSA as a major causative agent for the problem [170–172]. MRSA is one of the most prevalent pathogens among HAIs and it has a severe impact on health, especially among immunocompromised patients such as neonates and ICU patients [173–176]. Inside hospitals, MRSA can be transmitted by air, droplets, and through direct (skin-to-skin contact) or indirect contact (via fomites) [177,178]. Any inanimate object at a health facility is a potential fomite, even healthcare personnel themselves could be a transmission source through their contaminated apparel [179–182].

MRSA survival on different hospital surfaces

A critical factor that allows the transmission of MRSA from a person to the environment and then to other

people, is the pathogen's ability to survive on different types of surfaces under low humidity conditions, and its persistence on these surfaces for extended periods [39,183]. It is noteworthy that the antibiotic-resistance trait of MRSA does not affect the length of the survival period on fomites compared to, for example, Methicillinsensitive S. aureus (MSSA). Rather, MRSA survival time on fomites is affected by inoculum concentration [184]. This is consistent with the phenomenon of cryptic growth; where cells in a nutrient-limited environment will live on the remains of surrounding dying cells, and thus, an increased inoculum concentration will provide more dying cells for longer periods of time to sustain the lives of the remaining bacteria [185]. However, others reported contradictory data on the differences in the survival ability/length of MRSA vs MSSA. For instance, Wagenvoort and Penders (1997) reported the survival of an MRSA linage for 175 days in hospital dust while an MSSA linage survived only for 4 weeks under the same conditions [186]. Zarpellon et al. (2015) reported the survival of MRSA and vancomycin-resistant S. aureus (VRSA) on vinyl floors and formica for 40-45 days while on latex for only 2 days while MSSA survived on latex for only one day (Table 5) [187].

In a study aimed at determining the survival time of different types of bacteria on common hospital materials, MRSA was shown to persist on polyester for the longest time, reaching up to 56 days. However, on cotton, the bacterial survival was far less than that on polyester. For instance, MRSA lasts for one week on pure cotton and two weeks on cotton terry, while it lasts on polyester-cotton blend for less than one week [184]. These observations have significant infection control implications. For example, polyester is commonly used in making privacy drapes that are frequently touched by patients and staff. The frequent use of the polyester drapes, alongside the increased survival time of MRSA on them, makes them a high-risk source for hosting and transmitting MRSA. Even though MRSA survives for the least period on polyester-cotton blends, it does survive for at least a day. Polyester-cotton blends are most commonly used in making different kinds of clothes including healthcare workers' (HCWs) lab coats and scrubs. Thus, even within one day, the movement of the HCWs between different patients could promote transmission of MRSA from one patient to another [184]. Apparently, the length of MRSA survival period on surfaces and textiles differs with the type of surface or textile it occupies as well as its initial inoculum size and other environmental conditions local to each tested place.

MRSA transmission to- and from- inanimate objects

Fomites or inanimate objects, when contaminated with microbes, can serve as reservoirs for the

Table 4. Summary of the published studies on the presence of MRSA inside transportation vehicles and stations.	esence of MRSA inside transportatic	on vehicles and stations.		
Type of swab- sample collected	Type of transportation vehicle/station (number)	Type of transportation vehicle/station No. (%) of samples/vehicles positive for MRSA out of the (number)	Country of study	References, Time of sample collection
Handrails (1400 swabs) Handrails Straps and handrails	Trolleybuses, trams and buses (n = 55) 0% of samples Buses (n = 85) 26% of buses Subway trains (n = 349) 2.3% of trains	0% of samples 26% of buses 2.3% of trains	Belgrade, Serbia Oporto, Portugal Tokyo and Niigata, Japan	[154] (NI) * [155] (05–2009/02-2010) [156] (2008–2010)
Handrails, ticket machines and hand-touched surfaces Handrails, seat rails, handgrips, stop buttons and tickets machine	buses and stations Buses (n = 199)	0% 36.2% of buses	Lyon, France Lisbon, Portugal	[157] (NI) [158] (05–2011/05-2012)
Pooled samples from handrails, seats, seat rails, driver's area handrails and stop buttons Hand rails, seats, stanchions, Ticket Vending Machines (TVMs)	Buses (n = 40) Buses (n = 112) stations and carriages	63 % of buses 16.1% of buses 2.5% of stations	Ohio, USA Porto, Portugal Guangzhou, China	[159] (07–2010/10-2010) [160] (10–2010/05-2011) [161] (11–2013)
and escatators. Toilet door handles Grab rail, armrest and vinyl seat	Airport (n = 136) Buses (n = 15)	0.7% of airports 12 isolates from 45 samples	Worldwide Chittagong City, Bangladesh	[162] (12–2012/11-2015) [163] (NI)
380 surface samples Handle and seat surfaces	railway stations and coach stations 1.58% of the 149 5 Public transportation vehicles (n = 28) 31.4 % of vehicles	1.58% of the 149 Staphylococcus species 31.4 % of vehicles	China Kathmandu valley, Nepal	[164] (12–2013-01-2014) [153] (06–2017-08-2017)
* NI; not identified				

Tab	le !	5. MRSA	survival	rate	on	different	hospita	l surfaces.
-----	------	---------	----------	------	----	-----------	---------	-------------

Reference	Type of fomite	Survival (days)
Fabrics		
[184]	Smooth cotton	4 to 21
	Cotton terry	2 to 14
	Cotton-polyester blend	1 to 3
	Polyester	1 to 40
[188]	Cotton	37
	Cotton-polyester blend	37
	Wool	41
	Silk	37
[183]	Polyester cloth curtain	9
Plastics		
[183]	Plastic charts	11
	Plastic laminated bedside table	> 12
[184]	Polypropylene plastic	40 to > 51
Other surfaces		
[189]	Glass	41
	Tile	45
	Countertop	≥ 60
[190]	Dry mops	56
[187]	Vinyl floors and formica	40
	Latex	2

dissemination of disease-causing agents to human hosts. Each of these inanimate objects can host an entire community of bacteria, viruses, fungi, or even metabolic products such as toxins that can lead to illnesses in humans [191]. The transmission of CA-MRSA from fomites to humans certainly plays an important role in the spread of MRSA in communities as well as in hospitals.

Patients colonized or infected with MRSA are considered the main contributors to MRSA contamination of their surrounding environment. Sexton and his colleagues (2006) have demonstrated that in hospital rooms where MRSA patients are isolated, more than half of the cultured surfaces tested positive for the pathogen, and the isolated strains were similar to those found in samples from patients occupying the same rooms as determined by pulsed-field gel electrophoresis (PFGE) analysis [192]. The frequency of contamination is influenced by the number and type of culture-positive body sites; patients with an MRSAcontaminated active wound, in the urine, or having diarrhea, shed more of the pathogen than patients who are only colonized, or just harbor the pathogen in other body sites [193,194]. Furthermore, another study has shown that in rooms occupied with patients who have MRSA colonizing their groin, 31% (75/240) of the surfaces were contaminated, compared to only 3.6% (27/760) contaminated surfaces in rooms occupied with MRSA-positive patients whose groins are MRSA-free [195]. Not surprisingly, it is more likely to find contaminated surfaces in the patient's surrounding environment if MRSA is isolated from the palm of the patient [196]. Nonetheless, MRSA could be found in the vicinity of the immediate environment of MRSAnegative patients, which could be transmitted via visitors or HCWs. Villamaria et al. (2015) collected samples from surfaces of 100 hospital rooms and were able to isolate 202 and 1,830 MRSA isolates from 32 MRSA-

Table 6. MRSA contamination rate on different fomites in patient's surroundings.

	Source (room)	Type of fomite	MRSA Contamination rate (number of contaminate fomites/total sample)
Tables			
216]	MRSA-positive patients ^a	Over bed table	25% (6/24)
196]	MRSA-positive patients	Over bed table	22.4 (19/85)
194]	Gastrointestinal MRSA-positive patients with diarrhea	Table	62.5% (5/8)
	MRSA-negative patients ^b	Table	16.7% (1/6)
193]	Wound or urine MRSA-positive patients	Over bed table	44.4% (12/27)
217]	MRSA-positive patients	Over bed table	3.8% (1/26)
	MRSA-negative patients	Over bed table	0% (0/9)
203]	ICU	Bedside table	40% (4/10)
218]	Burn unit	Bedside table	6.67% (2/30)
eds			
216]	MRSA-positive patients	Bedside rails	31.6% (6/19)
196]	MRSA-positive patients	Bed linens	40.2% (41/102)
100]	with positive patients	Bed side rails	20.9 (18/86)
194]	Gastrointestinal MRSA-positive patients with	Bedside rails	100% (8/8)
194]	diarrhea		
	MRSA-negative patients	Bedside rail	66.7% (4/6)
193]	Wound or urine MRSA-positive patients	Bed linen	55.6% (15/27)
		Bedside rails	29.6% (8/27)
217]	MRSA-negative patients	End of bed	3.8% (1/26)
-	MRSA-positive patients	End of bed	0% (0/10)
203]	ICU	Bed rails	50% (10/20)
		Bed crank	40% (4/10)
2101	Male surgical ward	Bed rails	
210]	Male surgical ward		4.7% (6/128)
218,219] urtains	Burn unit	Bed rails	6.67% (2/30)
216]	MRSA-positive patient	Curtains	0% (0/24)
217]	MRSA-positive patients	Privacy curtain	12.5% (1/8)
217]	MRSA-negative patients	Privacy curtain	0% (0/26)
oor hand	÷ .		070 (0/20)
196]	MRSA-positive patient	Inner side room door handle	2.7% (2/74)
[90]			
10.41		Outer side room door handle	4.1% (3/74)
194]	Rooms of gastrointestinal MRSA-positive patients with diarrhea	Room door handles	37.5% (3/8)
	MRSA-negative patients	Room door handles	16.7% (1/6)
220]	MRSA-positive patients	Room door handle	19% (4/21)
	MRSA-negative patients	Room door handle	7.4% (13/175)
193]	Wound or urine MRSA-positive patients	Bath door handle	22.2% (6/27)
-		Room door handle	7.4 (2/27)
210]	Male surgical ward	Room door handle	10.7% (3/28)
73]	Urban Emergency department	Door handle of ambulance bay	2.5% (1/40)
[0]	orban Emergency department	door key pad	2.5% (1/40)
utton dev	vices		
193]	Gastrointestinal MRSA-positive patients with diarrhea	TV remote	75% (6/8)
	MRSA-negative patients	TV remote	16.7% (1/6)
	Gastrointestinal MRSA-positive patients with	Nurse call button	37.5% (3/8)
	diarrhea	N. II.I	
	MRSA-negative patients	Nurse call button	33.3% (2/6)
221]	Colorectal surgical unit	Bed-control handsets	12.9% (9/70)
222]	Randomly chosen hospital rooms	Bed handset	0.87% (1/115)
	Male surgical ward	Nurse call button	7.7% (2/26)
210]			
oilets	-	Toilet seat	
oilets	Gastrointestinal MRSA-positive patients with	Toilet seat Toilet rail	62.5% (5/8)
oilets 194]	Gastrointestinal MRSA-positive patients with diarrhea	Toilet rail	62.5% (5/8) 50% (4/8)
oilets 194]	Gastrointestinal MRSA-positive patients with	Toilet rail Toilet seat	62.5% (5/8) 50% (4/8) 33.3% (2/6)
oilets 194]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients	Toilet rail Toilet seat Toilet rail	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6)
'oilets 194] 218,219]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit	Toilet rail Toilet seat	62.5% (5/8) 50% (4/8) 33.3% (2/6)
218,219]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with	Toilet rail Toilet seat Toilet rail	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6)
oilets 194] 218,219] Other furn	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea	Toilet rail Toilet seat Toilet rail Toilet floor Dresser	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8)
oilets 194] 218,219] Other furn 194]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients	Toilet rail Toilet seat Toilet rail Toilet floor Dresser Dresser	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6)
oilets 194] 218,219] Other furn 194] 193]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients	Toilet rail Toilet seat Toilet rail Toilet floor Dresser Dresser Floor	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27)
oilets 194] 218,219] Other furn 194] 193]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients	Toilet rail Toilet seat Toilet rail Toilet floor Dresser Dresser Floor Bulletin board	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6)
oilets 194] 218,219] Other furn 194] 193]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients	Toilet rail Toilet seat Toilet rail Toilet floor Dresser Dresser Floor	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27)
oilets 194] 218,219] Other furn 194] 193]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients	Toilet rail Toilet seat Toilet rail Toilet floor Dresser Dresser Floor Bulletin board	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27) 0% (0/23)
oilets 194] 218,219] Other furn 194]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients MRSA-negative patients	Toilet rail Toilet seat Toilet seat Toilet floor Dresser Dresser Floor Bulletin board Chair back Television	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27) 0% (0/23) 16% (4/25) 0% (0/23)
Toilets 194] 218,219] Dther furn 194]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients	Toilet rail Toilet seat Toilet seat Toilet floor Dresser Dresser Floor Bulletin board Chair back Television Bulletin board	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27) 0% (0/23) 16% (4/25) 0% (0/23) 0% (0/7)
Toilets 194] 218,219] Other furn 194] 193]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients MRSA-negative patients	Toilet rail Toilet seat Toilet seat Toilet floor Dresser Dresser Floor Bulletin board Chair back Television Bulletin board Chair back	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27) 0% (0/23) 16% (4/25) 0% (0/23) 0% (0/7) 0% (0/7) 0% (0/10)
oilets 194] 218,219] Other furn 194] 193]	Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Burn unit iture and floor Gastrointestinal MRSA-positive patients with diarrhea MRSA-negative patients Wound or urine MRSA-positive patients MRSA-negative patients	Toilet rail Toilet seat Toilet seat Toilet floor Dresser Dresser Floor Bulletin board Chair back Television Bulletin board	62.5% (5/8) 50% (4/8) 33.3% (2/6) 16.7% (1/6) 10% (3/30) 50% (4/8) 16.7% (1/6) 59.3% (16/27) 0% (0/23) 16% (4/25) 0% (0/23) 0% (0/7)

(Continued)

Table 6. (Continued).

Reference	Source (room)	Type of fomite	MRSA Contamination rate (number of contaminated fomites/total sample)
[223]	ED in tertiary care hospital	Treatment room desk	1.4% (1/69)
Others			
[210]	Male surgical ward	Ventilation duct/grill Radiator	8.3% (4/48) 36.4% (16/44)
[224]	Hospital wards	Multiple environmental locations	24.7% (174/705)
[218]	Burn unit	Chairs and nurse station	6.67% (2/30)
[225]	Tertiary care hospital	Nurses coats	2% (9/436)

a MRSA-positive patients are patients infected or colonized with MRSA

b MRSA-negative patients are patients who are MRSA-free

noncontact rooms, and 68 MRSA-contact rooms, respectively. One can speculate that the presence of MRSA in rooms occupied by MRSA-free patients can be attributed to residual contamination from previous occupants, visitors, or HCWs with hands or gloves contaminated with MRSA. Gloved hand contamination is just as likely to occur after contacting commonly examined skin sites in MRSA patients, and after contacting the patient's surrounding environment [197]. In one experiment, Desai et al, (2011) assessed the survival and transmission of CA-MRSA strain USA 300-0114 from 9 different fomites that are used by humans [198]. Experimentally, CA-MRSA was transmissible from many fomites to skin with the non-porous fomites exhibiting transmissibility for many weeks, while porous fomites transmit pathogens fast but for short times. In another study, Moore et al. (2013) evaluated the MRSA transmission between different types of gloves worn by HCWs and fomite surfaces [199]. In this study, they have found that the bacteria transfer occurs at a very low rate ranging from 0% to 20%, with the difference is anticipated due to variations in the type and hydrophobicity of the materials used to make glove. Surface characteristics certainly influence microbial survival and the rates of transfer to and from humans, in addition to the type of use and the frequent use of gloves by the end users.

MRSA-contaminated hospital surfaces and instruments, and their role in MRSA outbreaks

Several studies have reported the presence of MRSA contamination on different fomites in the patient's surrounding environment, highlighting the potential role of these inanimate objects in transmitting the pathogen [200,201]. Using a quantitative approach based on the frequency of patient and HCWs contact with different objects, Huslage et al. (2012) defined 'high-touch surfaces' as the bed surface, bed rails, overbed tables, intravenous pumps, and supply carts [202]. In rooms occupied with patients having heavy gastro-intestinal MRSA colonization, a study conducted in England between June 2005 and Feb 2006 showed that the pathogen had colonized bedside rails, blood

pressure cuffs, television remote controls, and toilet seats (Tables 6 and Tables 7) [194]. In addition, one study conducted in Brazil in Oct 2008, reported MRSA colonization of 46% (29/63) of cultured surfaces close to ICU patients. These surfaces were bedside rails, bed cranks, bedside tables, infusion pump buttons, and cotton gowns [203]. Other instruments and accessories that are used frequently in the hospital were also tested and found to be contaminated with MRSA and other bacteria. For instance, X-ray facilities tested positive for MRSA as well as chairs in nursing stations and land phones and others.

A yearlong study (March 2009-Feb 2010) performed by Balen et al. (2016) in USA showed that the hospital's environmental surfaces that are exposed to general public contact could also be colonized by MRSA. In this study, hand-rails, coffee machines, and elevators were the most commonly contaminated general public surfaces, and to lesser extent, hand sanitizer dispensers and land phones were shown to be contaminated at certain times [204]. The results from the abovementioned study highlight the fact that different hospital surfaces are colonized with MRSA, but they do not provide a direct link for MRSA transmission from fomites to patients. Such links could be provided by outbreak reports, in which, the causative agent is attributed to contaminated inanimate objects. Molecular typing methods such as PFGE, Multi-locus sequence typing (MLST), variable number tandem repeats (VNTR), restriction fragment length polymorphism (RFLP) and whole-genome sequencing provide the means for linking pathogens isolated from different places and patients [205-207]. For example, in an MRSA outbreak in a postnatal ward in UK, which lasted for almost a year (Nov 89-Oct. 90) and affected 82 mothers and 28 babies, the bed mattresses were found contaminated with the same MRSA isolate that was identified as the causative agent for the outbreak. The mattresses were covered with old covers that were permeable to water. This outbreak was contained after the rooms were thoroughly cleaned, and the old furniture and mattresses were incinerated [208]. In another MRSA outbreak inside a plastic surgery/burn unit that occurred in 1996 in Manitoba, Canada,

ReferenceSource[226]Teaching hospital, Italy[223]ED in Tertiary care hospital Baltimore, USA[204]Two wards in an academic medical center,[204]Two wards in an academic medical center,[204]Two wards in an academic medical center,[204]Two wards in an academic medical center,[180]Radiology department, Korea[227]Ceneral hospital, Japan[229]General hospital, Japan[215]Tertiary hospital, Taiwan	Source Teaching hospital, Italy ED in Tertiary care hospital Baltimore, USA Two wards in an academic medical center, USA (a year-long study) Radiology department, Korea	Fomite Telephone handsets Computer keyboards Nursing station keyboard,Nursing station phone Chart holders	Contamination rate
	tal Baltimore, USA nic medical center, USA (a year-long study) corea	Telephone handsets Computer keyboards Nursing station keyboard,Nursing station phone Chart holders	18.9% (7/37)
	tal Baltimore, USA iic medical center, USA (a year-long study) corea	Computer keyboards Nursing station keyboard,Nursing station phone Chart holders	
	tal Baltimore, USA nic medical center, USA (a year-long study) corea	Nursing station keyboard,Nursing station phone Chart holders	22.2% (6/2/)
	ic medical center, USA (a year-long study) corea	Chart holders	4.3% (3/69)
	in metrod center, oon ta year forg study) (orea		(60/1) (1/0/1/2) (1/0/
	Órea		(F2/C) 0/C:/C plip (F2/LL) 0/C/C
	Órea	Medicine caris (drawers)	(10/24) Alla 41.1 % (10/24) (10/24) (10/24)
	Órea	Medicine cart laptops	12.5% (3/24) and 16.7% (4/24)
	Orea	Copy machine	16.7% (2/12) and 8.3% (1/12)
	Orea	Medicine room	50% (6/12)
	(orea	Computers	8.3% (3/36) and 25% (3/12)
	(orea	Access doors to office's halls of faculty and health care professionals	33.3% (4/12)
	órea	Markers at main nurses' board	8.3% (1/12)
	lorea lorea Lorea lorea lore lorea lorea lore lorea lorea	General use laptops	14.3% (3/21) and 15% (3/20)
		X-ray cassettes	16.2% (6/37)
		MRI	0.8% (1/25)
	di Arabia	ICU patients' files	6.8% (7/102)
		Surgery ward patients' files	1.1% (1/89)
		Working tables in ward staff centers	30.4% (17/56)
		Kevhoard	0.7% (2/282)
		Mouse	0.4% (1/282)
		Sporial unite' madical charts	0 3% (10/107)
		General wints incurate that's	(18/455)
Madian Janian			
ical devices			
[194] Gastrointestinal MRSA-po	Gastrointestinal MRSA-positive patients with diarrhea	Blood pressure cuffs	87.5% (7/8)
MRSA-negative patients		IV pump	25% (2/8)
		Blood pressure cuffs	16.7% (1/6)
		IV pump	0% (0/6)
[193] Wound or urine MRSA-positive patients	ositive patients	Blood pressure cuff	33.3% (9/27)
		Infusion pump button	18.5% (5/27)
[231] General intensive care unit	nit	Blood pressure cuffs	5% (5/100)
[232] Hospital wards		Blood pressure cuffs	8% (2/24)
[233] Critical care unites		Blood pressure cuffs	66.6% (33/50)
[234] Different places in a teaching hospital	ching hospital	Blood pressure cuffs	20% (10/50)
	Staff	Tourniquets	29% (7/24)
[236] Used by hospital staff pul	Used by hospital staff public and private sector hospitals	Tourniquets	17.4% (9/51)
[203] ICU		Infusion pump button	30% (3/10)
		Surgical gallons	46% (6/13)
[210] Male surgical ward		Medical equipment	13.2% (16/121)

Table 7. MRSA contamination rate in HCW contact objects.

Table 8. Summary of the range of MRSA contamination rate on HCW attire and devices, (adapted from [179]).

Sample type	MRSA contamination rate	Number of studies
Stethoscopes	0–42%	20
Digital devices	0-50%	23
White coats	0-79%	5
Neckties	3–32%	3
Pens	0-25%	4
Uniforms	4–20 %	4
ID badges	0–5 %	2

a hand-held shower and a shower stretcher in the hydrotherapy room tested positive for the outbreak strain, and were implicated as the source of contamination [209]. Moreover, in an outbreak in a male surgical ward that lasted from Jan 98 tell May 99 and affected 69 patients in UK, the MRSA strain isolated from the ward environment was the same one isolated from all patients [210]. Further, MRSA-contaminated ultra-sonic nebulizers inside a university tertiary care hospital in the Netherlands, was identified as a potential source of an outbreak that lasted for almost 6 months (May–Nov 2000) and spread to two ICU units. Such outbreaks demonstrate the role of fomites in aerosol transmission of MRSA, a route of transmission that causes a wider spread of the pathogen and more challenging to be contained [211]. Table 6 lists all the different studies reporting MRSA in hospital environments and fomites.

In a very recent study, Phoon et al. (2018) tested environmental samples from a tertiary hospital in Malaysia and isolated only one MRSA clone that was a multi-drug resistant [212].

HCWs contribution to environmental MRSA contamination

Hospital regulations force HCWs who are in direct contact with MRSA-colonized patients to wear gloves and gowns. However, even with those enforced guidelines, studies have shown that after the removal of gloves and gowns, many HCWs acquire the pathogen on their hands [213,214]. Unfortunately, not all health-care personnel are adherent with hand hygiene guidelines, which is a necessity for the prevention of MRSA transmission between HCWs and patients. For example, in a study aimed at examining MRSA contamination on medical charts, 4% and 9.3% of the medical charts examined in general wards and special units, respectively, were contaminated. Such results could be explained by the lack of handwashing by HCWs between patient rounds [215]. Beside medical charts, various medical objects and devices that are in direct contact with HCWs have been shown to be contaminated with MRSA (Table 7). Moreover, several reports have demonstrated the presence of MRSA contamination on HCWs attire and personal devices. A recent systematic study has shown that white coats, neckties, pens, and stethoscopes as well as other digital devices are commonly contaminated with MRSA. The results of this study are summarized in Table 8 [179]. The results from these studies could be utilized to redefine practices and guidelines aimed at preventing and controlling MRSA infections inside healthcare settings.

MRSA infection control strategies

MRSA is one of the most important nosocomial infections in hospitals that also causes infections in the community. MRSA infections created so much burden on the health systems and led to thousands of deaths annually all over the world. Several countries have launched aggressive measures to curb this pathogen and decrease the incidences of MRSA outbreaks. Different measures were taken including the availability of bedside hand sanitizer, isolation of MRSA patients in single rooms, use of barrier precautions inwards containing MRSA patients, decolonization of MRSA patients using mupirocin, and routine antiseptic washing of MRSA patients with chlorohexidine [237]. These procedures were implemented in both surgical wards and ICU units but with more heavy use in ICU units. It appeared that countries with decreasing MRSA proportions showed a strict implementation of various prevention measures while those with lower decreasing proportions did not follow such strict measures.

The environment may have a minor role in propagating MRSA's spread overall, but this role is certainly important [238]. Decontaminating the hospital environment, therefore, is vital for controlling MRSA spread among patients and different hospital wards. All the instruments used by doctors and nurses, linen, curtains, floors, attires, beds, tables, and many other instruments are considered parts of the environment that should be decontaminated.

Are MRSA incidences increasing or decreasing?

There is a plethora of studies summarizing the dynamics of MRSA incidences over the years from all over the world. However, it is very hard to get a consensus on a trend of whether MRSA-related infections are increasing or decreasing globally.

This part summarizes data from studies reported from scattered parts of the world showing that MRSA infections are generally decreasing. For instance, from 2005 to 2010, MRSA bacteremia cases dropped in 10 European countries; however, these years also witnessed the appearance of the community-associated MRSA strains [239]. In contrast, Otter and French (2010) have stated that generally in Europe MRSA incidence rates are low compared to other regions, but actually are still increasing in some parts of Europe [240]. In a recent study, McGough et al. (2018) reported a decreasing rate of both *S. aureus* and MRSA in 28 European countries between 2000 and 2016 [241].

In Germany, the proportion of MRSA among *S. aureus* isolates decreased continuously from 16% in 2010 to 10% in 2015 and the decrease was seen for all types of care and for the majority of sample types [242]. Further, a decreasing trend from 64.7% to 36.4%, in prevalence of MRSA bloodstream infections was observed in Greece between the years 2000–2015 [243].

results concerning Contrasting communityassociated MRSA infections were reported from Ontario, Canada. A significant increase in communityassociated MRSA from 23.6% in 2010 to 43.0% was observed in 2016 [244]. In the same token, in another report from Canada, the proportion of CA-MRSA genotypes increased significantly from 20.8% in 2007 to 56.3% in 2016 while HA-MRSA genotypes decreased from 79.2% to 43.8% throughout the study period [245]. Kavanagh et al. (2017) conducted a thorough review of several studies to analyze the trend of the MRSA infections in the USA [246]. They have reported that the incidences of MRSA infections varied widely depending upon the type of population studied, the types of infections captured, and the definitions and terminology used to describe the results in these studies. Kavanagh et al. (2017) concluded that they were unable to identify a firm evidence that there is a significant decrease in the total number of HA-MRSA infections in the USA. In contrast, the Military veteran's health-care system in the USA reported a decrease of 87% MRSA infections in ICUs and about 80% reduction in other wards between Oct 2007 and Oct 2015. However, in a CDC report published in 2019 observing MRSA trend from 2005 to 2012, they have concluded that rates of hospital-onset MRSA bloodstream infection decreased by 17.1% annually, but the decline slowed during 2013-2016 [247]. Further, community-onset MRSA declined but less markedly with a 6.9% annual rate during 2005–2016 and relating the decline to a general decline in health care-associated infections [247].

In a study conducted in Shanghai, China to evaluate the trend of the MRSA cases over ten year period, Dai et al. (2018) reported that the proportion of MRSA markedly decreased from 83.5% to 54.2%, in the years between 2008 and 2017 [248]. Similarly, there was a 64.6% decrease in the number of isolates of methicillin-resistant *S. aureus*, in Taiwan and Korea over the study period which lasted from 2008 to 2015 [249].

However, it is very hard to obtain a consensus on whether the MRSA incidence is decreasing or increasing in the world. Nonetheless, the author's impression is that the infection control measures are helping in decreasing MRSA and other nosocomial infections in places where the control measures are strictly adhered. A similar effect was not observed in places where such control measures were not applied.

Emerging antibiotic-resistance infections other than MRSA

MRSA is not the only multidrug-resistant bacteria (MDR), there are other microbes that have developed multidrug resistance to many antibiotics. Vancomycin-Resistant *Enterococcus* (VRE), Carbapenem-Resistant *Enterobacteriaceae* (CRE) and the Extended Spectrum Beta Lactamase (ESBL) producers *Enterobaceriaceae* are examples of MDR organisms that are causing major problems for the health systems worldwide. Nevertheless, it appears that MRSA is still leading the way globally in the high number of cases [237,250–252].

Conclusions

Fomites, whether in community or hospital settings, can be contaminated with multiple types of pathogenic bacteria. Among these, S. aureus are frequently isolated from all objects that are tested for contamination. Hospital fomites represent a major source of nosocomial infections, S. aureus for instance, was isolated from every tested object. Further, MRSA was isolated almost from all tested objects too, but to a lesser extent than MSSA. Interestingly, fomites in public community environments such as transportation, daycare, athletic facilities, airports, and other high-traffic areas were also considered reservoirs for many pathogenic bacteria including S. aureus, and particularly MRSA, which indicates the lack of proper hygiene practices among individuals who work or participate in activities associated with these environments. This certainly poses a risk of spreading the infectious diseases that are challenging to manage from public health perspectives.

Furthermore, some of the studies highlighted in this review described the differences among the various types of fabrics and fomites used in hospitals in supporting the growth of multiple pathogenic bacteria. The results from these studies should have an impact on the healthcare policy decision-makers when purchasing the textiles, clothing, and construction materials for hospital settings and personnel.

Acknowledgments

We acknowledge the Jordan University of Science and Technology for continuous support and we greatly acknowledge Dr. Ben Davis Tall from the Virulence Mechanisms Branch, US FDA, (Maryland, USA) for reviewing the manuscript. Thanks are also extended to Aseel Ziad Jaradat who is majoring in English, for reviewing the final version of the manuscript.

Disclosure statement

No potential conflict of interest was reported by the authors.

References

- Otto M. Staphylococcus aureus toxins. Curr Opin Microbiol [Internet]. 2014 Feb [cited 2019 Aug 26];17:32–37. Available from: http://www.ncbi. nlm.nih.gov/pubmed/24581690
- [2] Warnke P, Harnack T, Ottl P, et al. Nasal screening for Staphylococcus aureus - daily routine with improvement potentials. PLoS One. 2014;9(2):1–7.
- [3] Todar K Staphylococcus (page 5) ©. 2003;(5).
- [4] Fournier B, Philpott D. Reconocimiento de carácteristicas de la degradación de pasturas en el Rancho San Luis - Morelia - Caquetá - Colombia. Rev Fac Ciencias Agropecu. 2010;18(3):3–12.
- [5] Liu GY. Molecular pathogenesis of *Staphylococcus aureus* Infection. Pediatr Res. 2009;65:71–77.
- [6] Pengov A. *Staphylococcus aureus* do we really have to live with it?. Slov Vet Res. 2006;43(1):41–46.
- [7] Lowy FD. Staphylococcus aureus Infections. N Engl J Med [Internet]. 1998 Aug 20;339(8):520–532. Available from.
- [8] Roberts S, Chambers S. Diagnosis and management of Staphylococcus aureus infections of the skin and soft tissue. Intern Med J. 2005;35(SUPPL. 2):S97–S105.
- [9] Lowy F. Antimicrobial resistance: the example of Staphylococcus aureus. J Clin Invest [Internet]. 2003;111(9):1265–1273. Available from: http://www. pubmedcentral.nih.gov/articlerender.fcgi?artid= 154455&tool=pmcentrez&rendertype=abstract% 5Cnhttp://www.jci.org/cgi/content/abstract/111/9/ 1265.
- [10] Guignard B, Entenza J, Moreillon P. β-lactams against methicillin-resistant *Staphylococcus aureus*. Curr Opin Pharmacol [Internet]. 2005 Oct 1 [cited 2017 Oct 24];5 (5):479–489. Available from: http://www.sciencedirect. com/science/article/pii/S1471489205001153?via% 3Dihub
- [11] Chambers HF. The changing epidemiology of Staphylococcus aureus? Emerg Infect Dis [Internet]. 2001;7(2):178–182. Available from: http://www.ncbi. nlm.nih.gov/pmc/articles/PMC2631711/.
- [12] Urushibara N, Aung MS, Kawaguchiya M, et al. Novel staphylococcal cassette chromosome mec (SCCmec) type XIV (5A) and a truncated SCCmec element in SCC composite islands carrying speG in ST5 MRSA in Japan. J Antimicrob Chemother [Internet]. 2020 Jan 1 [cited 2020 Sep 3];75(1):46–50. Available from: https:// pubmed.ncbi.nlm.nih.gov/31617906/
- [13] Deurenberg RH, Vink C, Kalenic S, et al. The molecular evolution of methicillin-resistant *Staphylococcus aureus*. Clin Microbiol Infect [Internet]. 2006;13 (3):222–235. Available from:.
- [14] LG M, Diep BAA. Colonization, fomites, and virulence: rethinking the pathogenesis of community-associated methicillin-resistant *Staphylococcus aureus* Infection. Clin Pract. 2008;46(5):752–760.

- [15] Lopez GU, Gerba CP, Tamimi AH, et al. Transfer efficiency of bacteria and viruses from porous and nonporous fomites to fingers under different relative humidity conditions. Appl Environ Microbiol. 2013;79 (18):5728–5734.
- [16] King JM, Kulhankova K, Stach CS, et al. Phenotypes and virulence among *Staphylococcus aureus* USA100, USA200, USA300, USA400, and USA600 clonal lineages. mSphere [Internet]. 2016 [cited 2020 Apr 20];1(3). Available from: http://www.ncbi.nlm.nih. gov/pubmed/27303750
- [17] Balaji V, Yamuna DB, Francis YI, et al. Molecular characterization of panton-valentine leukocidin (PVL) toxin-encoding phages from South India [Internet]. New Microbes New Infect [cited 2020 Sep 2]. 2017;20:34–38. Available from: /pmc/articles/ PMC5682882/?report=abstract
- [18] Mairi A, Touati A, Lavigne JP. Methicillin-resistant Staphylococcus aureus ST80 clone: A systematic review [Internet]. Toxins (Basel) [cited 2020 Sep 2]. 2020;12:119. Available from: /pmc/articles/ PMC7076798/?report=abstract
- [19] Liu CY, Lai CC, Chiang HT, et al. Predominance of methicillin-resistant *Staphylococcus aureus* in the residents and environments of long-term care facilities in Taiwan. J Microbiol Immunol Infect [Internet]. 2019 Feb 1 [cited 2020 Sep 3];52:62–74. Available from: https://pubmed.ncbi.nlm.nih.gov/29530709/
- [20] Kang S, Lee J, Kim M. The association between Staphylococcus aureus nasal colonization and symptomatic infection in children in Korea where ST72 is the major genotype. Med (United States) [Internet]. 2017 Aug 1 [cited 2020 Sep 3];96(34). Available from: https://pubmed.ncbi.nlm.nih.gov/28834892/
- [21] Glaser P, Martins-Simões P, Villain A, et al. Demography and intercontinental spread of the USA300 community-acquired methicillin-resistant *Staphylococcus aureus* lineage. MBio. 2016 Feb 16;7 (1):e02183–15.
- [22] Yokomori R, Tsurukiri J, Moriya M, et al. First report of fatal infection caused by community-acquired methicillin-resistant *Staphylococcus aureus* USA300 clone in a collegiate athlete. Jma J. 2020;3(1):78–82.
- [23] Carrel M, Perencevich EN, David MZ. USA300 methicillin-resistant *Staphylococcus aureus*, United States, 2000–2013. Emerg Infect Dis. 2015 Nov 1;21 (11):1973–1980.
- [24] Arias CA, Reyes J, Carvajal LP, et al. A prospective cohort multicenter study of molecular epidemiology and phylogenomics of *Staphylococcus aureus* bacteremia in nine Latin American countries. Antimicrob Agents Chemother [Internet]. 2017 [cited 2020 Apr 20];61(10). Available from: http://www.ncbi.nlm. nih.gov/pubmed/28760895
- [25] Mediavilla JR, Chen L, Mathema B, et al. Global epidemiology of community-associated methicillin resistant *Staphylococcus aureus* (CA-MRSA). Curr Opin Microbiol. 2012;15(5):588–595.
- [26] Johnson JK, Khoie T, Shurland S, et al. Skin and soft tissue infections caused by methicillin-resistant *Staphylococcus aureus* USA300 Clone. Emerg Infect Dis [Internet]. 2007 Aug [cited 2019 Sep 2];13 (8):1195–1200. Available from: http://www.ncbi.nlm. nih.gov/pubmed/17953091
- [27] Davis SL, Perri MB, Donabedian SM, et al. Epidemiology and outcomes of community-associated

methicillin-resistant *Staphylococcus aureus* infection. J Clin Microbiol. 2007;45(6):1705–1711.

- [28] Kluytmans J, Struelens M Meticillin resistant Staphylococcus aureus in the hospital. BMJ [Internet]. 2009 Feb 12;338. Available from: http://www.bmj. com/content/338/bmj.b364.abstract
- [29] Uhlemann AC, Knox J, Miller M, et al. The environment as an unrecognized reservoir for community-associated methicillin resistant *Staphylococcus aureus* USA300: A Case-Control study. PLoS One. 2011;6(7):1–9.
- [30] Coughenour C, Stevens V, Stetzenbach LD. An evaluation of methicillin-resistant *Staphylococcus aureus* survival on five environmental surfaces. Microb Drug Resist [Internet]. 2011;17(3):457–461. Available from: http://www.liebertonline.com/doi/abs/10.1089/mdr. 2011.0007.
- [31] Nwankwo E. Isolation of pathogenic bacteria from fomites in the operating rooms of a specialist hospital in Kano, North-western Nigeria. Pan Afr Med J. 2012;12:90.
- [32] Cookson B, Bonten MJM, MacKenzie FM, et al. Meticillin-resistant Staphylococcus aureus (MRSA): screening and decolonisation. Int J Antimicrob Agents [Internet]. 2011;37(3):195–201. Available from:.
- [33] Atkinson MP, Wein LM. Spatial queueing analysis of an interdiction system to protect cities from a nuclear terrorist attack. Oper Res [Internet]. 2008;56 (1):247–254. Available from: http://www.jstor.org/ stable/25147180.
- [34] Haas BK. Clarification and integration of similar quality of life concepts. J Nurs Scholarsh [Internet]. 1999 Sep 1 [cited 2017 Oct 23];31(3):215–220. Available from: http://onlinelibrary.wiley.com/doi/10.1111/j.1547-5069.1999.tb00483.x/abstract/#.We57Nkb_160. mendeley
- [35] Hall C, Douglas R. Modes of transmission of respiratory syncytial virus. J Pediatr. 1981;99(1):100–102.
- [36] Hendley JO. Epidemic Keratoconjunctivitis and hand washing. N Engl J Med [Internet]. 1973 Dec 20;289 (25):1368–1369. Available from.
- [37] Abad-Franch F, Paucar CA, Carpio CC, et al. Biogeography of triatominae (Hemiptera: reduviidae) in Ecuador: implications for the design of control strategies. Mem Inst Oswaldo Cruz. 2001;96 (5):611–620.
- [38] Boone SA, Gerba CP. Significance of fomites in the spread of respiratory and enteric viral disease. Appl Environ Microbiol. 2007;73(6):1687–1696.
- [39] Kramer A, Schwebke I, Kampf G. How long do nosocomial pathogens persist on inanimate surfaces? A systematic review. BMC Infect Dis. 2006;6 (1):130.
- [40] Barton M, Hawkes M, Moore D, et al. Guidelines for the prevention and management of community-associated methicillin-resistant *Staphylococcus aureus*: A perspective for Canadian health care practitioners. Can J Infect Dis Med Microbiol. 2006;17(SupplC):4C–24C.
- [41] Skov RL, Jensen KS. Community-associated meticillin-resistant *Staphylococcus aureus* as a cause of hospital-acquired infections. J Hosp Infect [Internet]. 2009;73(4):364–370.
- [42] Otter JA, French GL. Community-associated meticillin-resistant *Staphylococcus aureus*: the case for a genotypic definition. J Hosp Infect. 2012;81 (3):143–148.
- [43] Nurjadi D, Klein S, Zimmermann S, et al. Transmission of ST8-USA300 latin american variant

methicillin-resistant *Staphylococcus aureus* on a neonatal intensive care unit: recurrent skin and soft- tissue infections as a marker for epidemic community-associated-MRSA colonization. Infect Control Hosp Epidemiol [Internet]. 2017 Jul 18 [cited 2019 Sep 2];38(7):883–885. Available from: https:// www.cambridge.org/core/product/identifier/ S0899823X17000885/type/journal_article

- [44] Shilo N, Quach C. Pulmonary Infections and Community Associated Methicillin Resistant Staphylococcus aureus: a dangerous mix? Paediatr Respir Rev [Internet]. 2011;12(3):182–189. Available from
- [45] Kirkland EB, Adams BB. Methicillin-resistant *Staphylococcus aureus* and athletes. J Am Acad Dermatol. 2008;59(3):494–502.
- [46] Friedman L, Wallar L, Papadopoulos A. Environmental risk factors for psychosis. Clin Res. 2015;7(1):69–80.
- [47] Kazakova SV, Hageman JC, Matava M, et al. A clone of methicillin-resistant *Staphylococcus aureus* among professional football players. N Engl J Med [Internet]. 2005 Feb 3 [cited 2019 Sep 2];352:468–475. Available from: http://www.nejm.org/doi/abs/10.1056/ NEJMoa042859
- [48] Romano R, Lu D, Holtom P. Outbreak of community-acquired methicillin-resistant *Staphylococcus aureus* skin infections among a collegiate football team. J Athl Train. 2006;41 (2):141–145.
- [49] Begier EM, Frenette K, Barrett NL, et al. A high-morbidity outbreak of methicillin-resistant *Staphylococcus aureus* among players on a college football team, facilitated by cosmetic body shaving and turf burns. ClinInfectDis. 2004;39 (1537–6591):1446–1453.
- [50] Nguyen DM, Mascola L, Bancroft E. Recurring methicillin-resistant *Staphylococcus aureus* infections in a football team. Emerg Infect Dis. 2005;11 (4):526–532.
- [51] Stacey AR, Endersby KE, Chan PC, et al. An outbreak of methicillin resistant *Staphylococcus aureus* infection in a rugby football team. Br J Sports Med [Internet]. 1998 Jun 1 [cited 2019 Sep 2];32(2):153–154. Available from: http://www.ncbi.nlm.nih.gov/pubmed/9631224
- [52] Lindenmayer JM, Schoenfeld S, O'Grady R, et al. Methicillin-resistant Staphylococcus aureus in a high school wrestling team and the surrounding community. Arch Intern Med [Internet]. 1998 Apr 27 [cited 2019 Sep 2];158(8):895. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/9570176
- [53] Huijsdens XW, van Lier AMC, van Kregten E, et al. Methicillin-resistant Staphylococcus aureus in Dutch soccer team. Emerg Infect Dis [Internet]. 2006 Oct [cited 2019 Sep 2];12(10):1584–1586. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 17176578
- [54] Stanforth B, Krause A, Starkey C, et al. Prevalence of community-associated methicillin-resistant *Staphylococcus aureus* in high school wrestling environments. J Environ Health [Internet]. [cited 2019 Sep 2].2010;72(6):12–16. Available from: http://www. ncbi.nlm.nih.gov/pubmed/20104828
- [55] Montgomery K, Ryan TJ, Krause A, et al. Assessment of athletic health care facility surfaces for MRSA in the secondary school setting. J Environ Health. 2010 Jan [cited 2019 Sep 2];72(6). Available from: http://www. ncbi.nlm.nih.gov/pubmed/20104827.

- [56] Ryan KA, Ifantides C, Bucciarelli C, et al. Are gymnasium equipment surfaces a source of staphylococcal infections in the community? Am J Infect Control [Internet]. 2011 Mar [cited 2019 Sep 19];39 (2):148–150. Available from: http://www.ncbi.nlm.nih. gov/pubmed/21356432
- [57] Markley JD, Edmond MB, Major Y, et al. Are gym surfaces reservoirs for *Staphylococcus aureus*? A point prevalence survey. Am J Infect Control [Internet]. 2012 Dec [cited 2019 Sep 2];40(10):1008–1009. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/22622513
- [58] Mashau TP Antibiogram and molecular characterization of *Staphylococcus aureus* isolated from gym equipment in public fitness centres in Thohoyandou, Vhembe District, Limpopo Province [Internet]. University of Venda; 2016 [cited 2019 Sep 2]. Available from: http://univendspace.univen.ac.za/han dle/11602/792
- [59] Mukherjee N, Sulaiman IM, Banerjee P. Characterization of methicillin-resistant *Staphylococcus aureus* isolates from fitness centers in the Memphis metropolitan area, Tennessee. Am J Infect Control [Internet]. 2016 Dec 1 [cited 2019 Sep 2];44(12):1681–1683. Available from: http://www.ncbi.nlm.nih.gov/pubmed/27658767
- [60] Dalman M, Bhatta S, Nagajothi N, et al. Characterizing the molecular epidemiology of *Staphylococcus aureus* across and within fitness facility types. BMC Infect Dis [Internet]. 2019 Dec 18 [cited 2019 Sep 19];19:69. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/30658587
- [61] Maurice Bilung L, Tahar AS, Kira R, et al. High occurrence of *Staphylococcus aureus* isolated from fitness equipment from selected gymnasiums. La Torre G, editor. J Environ Public Health [Internet]. 2018;2018:4592830. Available from
- [62] LaBelle MW, Knapik DM, Arbogast JW, et al. Infection risk reduction program on pathogens in high school and collegiate athletic training rooms. Sports Health [Internet]. 2020 Jan 1 [cited 2020 Apr 20];12:51–57. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/31660785
- [63] Thapaliya D, Kadariya J, Capuano M, et al. Prevalence and molecular characterization of *Staphylococcus aureus* and methicillin-resistant *S.aureus* on children's playgrounds. Pediatr Infect Dis J [Internet]. 2019 Mar [cited 2019 Sep 19];38(3):e43–7. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/29746375
- [64] Sexton JD, Reynolds KA. Exposure of emergency medical responders to methicillin-resistant *Staphylococcus aureus*. Am J Infect Control [Internet]. 2010 Jun [cited 2019 Sep 2];38(5):368–373. Available from: https://linkinghub. elsevier.com/retrieve/pii/S0196655310001483
- [65] Al Amiry A, Bissell RA, Maguire BJ, et al. Methicillinresistant Staphylococcus aureus nasal colonization prevalence among emergency medical services personnel. Prehosp Disaster Med [Internet]. 2013 Aug 24 [cited 2019 Sep 2];28(4):348–352. Available from: https://www.cambridge.org/core/product/identifier/ S1049023X13003476/type/journal_article
- [66] Hudson AJ, Glaister GD, Wieden H-J. The emergency medical service microbiome. Drake HL, editor. Appl Environ Microbiol [Internet]. 2017 Dec 8 [cited 2019 Sep 2];84(5). Available from: http://www.ncbi.nlm.nih. gov/pubmed/29222105
- [67] J V R, Buhs LK, Makarovaite V, et al. Detection and analysis of *Staphylococcus aureus* isolates found in

ambulances in the Chicago metropolitan area. Am J Infect Control [Internet]. 2012 Apr [cited 2019 Sep 2];40(3):201–205. Available from: https://linkin ghub.elsevier.com/retrieve/pii/S0196655311011758

- [68] Varona-Barquin A, Ballesteros-Peña S, Lorrio-Palomino S, et al. Detection and characterization of surface microbial contamination in emergency ambulances. Am J Infect Control. 2017;45:69–71.
- [69] Roline CE, Crumpecker C, Dunn TM. Can methicillin-resistant *Staphylococcus aureus* be found in an ambulance fleet? Prehosp Emerg Care [Internet]. 2007 Jan 2 [cited 2019 Sep 2];11 (2):241–244. Available from: http://www.tandfon line.com/doi/full/10.1080/10903120701205125
- [70] Brown R, Minnon J, Schneider S, et al. Prevalence of methicillin-resistant *Staphylococcus aureus* in ambulances in southern Maine. Prehosp Emerg Care [Internet]. 2010 Jun 3 [cited 2019 Sep 2];14
 (2):176–181. Available from: http://www.tandfonline. com/doi/full/10.3109/10903120903564480
- [71] El-Mokhtar MA, Hetta HF. Ambulance vehicles as source of multidrug-resistant infections: а a multicenter study in Assiut City, Egypt. Infect Drug Resist [Internet]. 2018 Apr [cited 2019 Sep 2];11:587-594. Available from: https://www.dove press.com/ambulance-vehicles-as-a-source-ofmultidrug-resistant-infections-a-mul-peer-reviewedarticle-IDR
- [72] Eibicht SJ, Meticillin-resistant VU. Staphylococcus aureus (MRSA) contamination of ambulance cars after short term transport of MRSA-colonised patients is restricted to the stretcher. J Hosp Infect [Internet]. 2011 Jul [cited 2019 Sep 2];78(3):221–225. Available from: https://linkinghub.elsevier.com/retrieve/pii/ S0195670111000351
- [73] Kei J, Richards JR. The prevalence of methicillin-resistant *Staphylococcus aureus* on inanimate objects in an urban emergency department. J Emerg Med [Internet]. 2011 Aug [cited 2019 Sep 2];41(2):124–127. Available from: http://www. ncbi.nlm.nih.gov/pubmed/19111424
- [74] Roberts MC, No DB. Environment surface sampling in 33 Washington State fire stations for methicillin-resistant and methicillin-susceptible *Staphylococcus aureus*. Am J Infect Control [Internet]. 2014;42(6):591–596. Available from:.
- [75] Roberts MC, Soge OO, No D, et al. Isolation and characterization of methicillin-resistant *Staphylococcus aureus* from fire stations in two northwest fire districts. Am J Infect Control. 2011;39 (5):382–389.
- [76] Flores GE, Bates ST, Knights D, et al. Microbial biogeography of public restroom surfaces. Liles MR, editor. PLoS One [Internet]. 2011 Nov 23 [cited 2019 Sep 2];6: e28132. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/22132229
- [77] Mkrtchyan HV, Xu Z, Cutler RR. Diversity of SCCmec elements in Staphylococci isolated from public washrooms. BMC Microbiol [Internet]. 2015 Dec 14 [cited 2019 Sep 2];15(1):120. Available from: http://www.biomedcentral.com/1471-2180/ 15/120
- [78] Roberts MC, Soge OO, No D, et al. Characterization of methicillin-resistant *Staphylococcus aureus* isolated from public surfaces on a university campus, student homes and local community. J Appl Microbiol. 2011;110(6):1531–1537.

- [79] Ahmed OB, Sirag B. Microbial contamination of doorknobs in public toilets during Hajj. Asian J Sci Technol [Internet]. 2016 [cited 2019 Sep 2];7(10):3676–3679. Available from: https://www.researchgate.net/publica tion/309547998_Microbial_contamination_of_door knobs_in_public_toilets_during_Hajj
- [80] Giannini MA, Nance D, McCullers JA. Are toilet seats a vector for transmission of methicillin-resistant *Staphylococcus aureus*? Am J Infect Control [Internet]. 2009 Aug [cited 2019 Sep 2];37(6):505–506. Available from: https://linkinghub.elsevier.com/retrieve/pii/ S0196655309000042
- [81] Kanayama Katsuse A, Takahashi H, Yoshizawa S, et al. Public health and healthcare-associated risk of electric, warm-water bidet toilets. J Hosp Infect [Internet]. 2017 Nov [cited 2019 Sep 2];97(3):296–300. Available from: http://www.ncbi.nlm.nih.gov/pubmed/28756169
- [82] Goodwin KD, McNay M, Cao Y, et al. A multi-beach study of *Staphylococcus aureus*, MRSA, and enterococci in seawater and beach sand. Water Res [Internet]. 2012 Sep 1 [cited 2019 Sep 2];46(13):4195–4207. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 22652414
- [83] Goodwin KD, Pobuda M. Performance of CHROMagarTM Staph aureus and CHROMagarTM MRSA for detection of *Staphylococcus aureus* in seawater and beach sand – comparison of culture, agglutination, and molecular analyses. Water Res [Internet]. 2009 Nov [cited 2019 Sep 2];43(19):4802–4811. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/19577788
- [84] Selvakumar A, Borst M. Variation of microorganism concentrations in urban stormwater runoff with land use and seasons. J Water Health [Internet]. 2006 Mar [cited 2019 Sep 2];4(1):109–124. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/16604843
- [85] Börjesson S, Matussek A, Melin S, et al. Methicillinresistant Staphylococcus aureus (MRSA) in municipal wastewater: an uncharted threat? J Appl Microbiol [Internet]. 2010 Apr [cited 2019 Sep 2];108 (4):1244–1251. Available from: http://www.ncbi.nlm. nih.gov/pubmed/19735317
- [86] Baptiste KE, Williams K, Willams NJ, et al. Methicillinresistant Staphylococci in companion animals. Emerg Infect Dis [Internet]. 2005 Dec [cited 2019 Sep 2];11 (12):1942–1944. Available from: http://www.ncbi.nlm. nih.gov/pubmed/16485485
- [87] Shah AH, Abdelzaher AM, Phillips M, et al. Indicator microbes correlate with pathogenic bacteria, yeasts and helminthes in sand at a subtropical recreational beach site. J Appl Microbiol [Internet]. 2011 Jun [cited 2019 Sep 2];110(6):1571–1583. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/21447014
- [88] Roberts MC, Soge OO, No D, Comparison of multi-drug resistant environmental methicillin-resistant *Staphylococcus aureus* isolated from recreational beaches and high touch surfaces in built environments. Front Microbiol [Internet]. 2013 [cited 2019 Sep 2];4:74. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/23577006
- [89] Goodwin KD, Pobuda M. Performance of CHROMagar Staphylococcus aureus and CHROMagar MRSA for detection of *Staphylococcus aureus* in seawater and beach sand-comparison of culture, agglutination, and molecular analyses. Water Res [Internet]. 2009 Nov [cited 2019 Sep 2];43(19):4802–4811. Available

from: https://linkinghub.elsevier.com/retrieve/pii/ S0043135409003893

- [90] Soge OO, Meschke JS, No DB, et al. Characterization of methicillin-resistant Staphylococcus aureus and methicillin-resistant coagulase-negative *Staphylococcus* spp. isolated from US West Coast public marine beaches. J Antimicrob Chemother [Internet].
 2009 Dec 1 [cited 2019 Sep 2];64(6):1148–1155. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/19837712
- [91] Levin-Edens E, Soge OO, No D, et al. Methicillinresistant Staphylococcus aureus from Northwest marine and freshwater recreational beaches. FEMS Microbiol Ecol [Internet]. 2012 Feb [cited 2019 Sep 2];79(2):412–420. Available from: http://www. ncbi.nlm.nih.gov/pubmed/22092827
- [92] Yamahara KM, Sassoubre LM, Goodwin KD, et al. Occurrence and persistence of bacterial pathogens and indicator organisms in beach sand along the california coast. Appl Environ Microbiol. 2012 Mar 15 [cited 2019 Sep 2];78(6): 1733–1745. Available from: http://www.ncbi.nlm.nih.gov/pubmed/22247142
- [93] Hower S, Phillips MC, Brodsky M, et al. Clonally related methicillin-resistant *Staphylococcus aureus* isolated from short-finned pilot whales (globicephala macrorhynchus), human volunteers, and a bayfront cetacean rehabilitation facility. Microb Ecol [Internet]. 2013 May 19 [cited 2019 Sep 2];65:1024–1038. Available from: http://www.ncbi.nlm.nih.gov/pubmed/23508733
- [94] Plano LRW, Shibata T, Garza AC, et al. Humanassociated methicillin-resistant *Staphylococcus aureus* from a subtropical recreational marine beach. Microb Ecol [Internet]. 2013 May 4 [cited 2019 Sep 2];65:1039–1051. Available from: http://www. ncbi.nlm.nih.gov/pubmed/23553001
- [95] Akanbi OE, Njom HA, Fri J, et al. Antimicrobial susceptibility of *Staphylococcus aureus* isolated from recreational waters and beach sand in eastern cape province of south africa. Int J Environ Res Public Health [Internet]. 2017 Sep 1 [cited 2019 Sep 2];14(9):1001. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/28862669
- [96] Thapaliya D, Hellwig EJ, Kadariya J, et al. Prevalence and Characterization of *Staphylococcus aureus* and methicillin-resistant *Staphylococcus aureus* on Public recreational beaches in Northeast Ohio. GeoHealth [Internet]. 2017 Dec 1 [cited 2019 Sep 2];1 (10):320–332. Available from:
- [97] Ottomeyer M, Graham CD, Legg AD, et al. Prevalence of nasal colonization by methicillin-resistant *Staphylococcus aureus* in persons using a homeless shelter in Kansas City. Front Public Health [Internet]. 2016;4(October):234. Available from: http://ovidsp. ovid.com/ovidweb.cgi?T=JS&PAGE=reference&D= prem&NEWS=N&AN=27826546
- [98] Szakacs TA, Toye B, Turnbull JM, et al. Prevalence of methicillin-resistant *Staphylococcus aureus* in a Canadian inner-city shelter. Can J Infect Dis Med Microbiol = J Can Des Mal Infect La Microbiol Medicale [Internet]. 2007 Jul [cited 2019 Sep 2];18 (4):249–252. Available from: http://www.ncbi.nlm.nih. gov/pubmed/18923739
- [99] Moffa M, Cronk R, Fejfar D, et al. A systematic scoping review of environmental health conditions and hygiene behaviors in homeless shelters. Int J Hyg Environ Health [Internet]. 2019 Apr [cited 2019

Sep 2];222(3):335–346. Available from: http://www. ncbi.nlm.nih.gov/pubmed/30583994

- [100] Landers TF, Harris RE, Wittum TE, et al. Colonization with Staphylococcus aureus and Methicillin-Resistant S. aureus among a Sample of Homeless Individuals, Ohio. Infect Control Hosp Epidemiol [Internet]. 2009 Aug 2 [cited 2019 Sep 2];30(8):801–803. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 19591582
- [101] Leibler JH, León C, Cardoso LJP, et al. Prevalence and risk factors for MRSA nasal colonization among persons experiencing homelessness in Boston, MA. J Med Microbiol. 2017 Aug;66(8):1183–1188.
- [102] Moritz ED, Hanson BM, Kates AE, et al. Molecular characteristics of *Staphylococcus aureus* isolated from employees, children, and environmental surfaces in lowa child daycare facilities. Am J Infect Control [Internet]. 2015 May 1 [cited 2019 Sep 19];43(5):482–488. Available from: https:// www.sciencedirect.com/science/article/abs/pii/ S0196655315000644
- [103] Ryan K, Black E, Saliba H, et al. Is methicillin-resistant Staphylococcus aureus colonization changing? A study of academic health center daycare facilities. Infect Control Hosp Epidemiol [Internet]. 2013 Oct 2 [cited 2019 Sep 19];34:1124–1126. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/24018938
- [104] Main CL, Jayaratne P, Haley A, et al. Outbreaks of Infection caused by community-acquired methicillin-resistant *Staphylococcus aureus* in a Canadian correctional facility. Can J Infect Dis Med Microbiol [Internet]. 2005 Nov [cited 2019 Sep 2];16 (6):343–348. Available from: http://www.ncbi.nlm.nih. gov/pubmed/18159517
- [105] Maree CL, Eells SJ, Tan J, et al. Risk factors for infection and colonization with community-associated methicillin-resistant *Staphylococcus aureus* in the los angeles county jail: a case-control study. Clin Infect Dis. 2010;51(11):1248–1257.
- [106] Felkner M, Andrews K, Field LH, et al. Detection of Staphylococcus aureus Including MRSA on Environmental Surfaces in a Jail Setting. J Correct Heal Care [Internet]. 2009 Oct 21 [cited 2019 Sep 19];15:310–317. Available from: http://www.ncbi. nlm.nih.gov/pubmed/19622846
- [107] Baillargeon J, Kelley MF, Leach CT, et al. Methicillin-Resistant Staphylococcus aureus Infection in the Texas Prison System. Clin Infect Dis [Internet]. 2004 May 1 [cited 2018 Aug 8];38(9):e92–5. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/15127360
- [108] Xu C, Weese SJ, Namvar A, et al. Sanitary status and incidence of methicillin-resistant *Staphylococcus aureus* and Clostridium difficile within Canadian hotel rooms. J Environ Health [Internet]. 2015 Apr [cited 2019 Sep 2];77(8):8–15. Available from: http://www. ncbi.nlm.nih.gov/pubmed/25876260
- [109] Almanza BA, Kirsch K, Kline SF, et al. How clean are hotel rooms? Part II: examining the concept of cleanliness standards. J Environ Health [Internet]. [cited 2019 Sep 2]. 2015;7z8(1):14–18. Available from: http://www. ncbi.nlm.nih.gov/pubmed/26427263
- [110] Xu Z, Mkrtchyan HV, Cutler RR. Antibiotic resistance and mecA characterization of coagulase-negative staphylococci isolated from three hotels in London, UK. Front Microbiol [Internet]. 2015 Sep 9 [cited 2019 Sep 2];6:947. Available from: http://www.ncbi.nlm.nih. gov/pubmed/26441881

- [111] Reynolds C, Quan V, Kim D, et al. Methicillin-Resistant Staphylococcus aureus (MRSA) Carriage in 10 nursing homes in Orange County, California. Infect Control Hosp Epidemiol [Internet]. 2011 Jan 2 [cited 2019 Sep 2];32:91–93. Available from: https://www.cam bridge.org/core/product/identifier/ S0195941700039710/type/journal_article
- [112] Mbogori C, Muigai A, Kariuki S. Detection and Characterization of methicillin resistant Staphylococcus aureus from toilet and classroom door handles in selected secondary schools in Nairobi county. Open J Med Microbiol [Internet]. 2013 [cited 2019 Sep 2];03(4):248–252. Available from:
- [113] Thapaliya D, Taha M, Dalman MR, et al. Environmental contamination with *Staphylococcus aureus* at a large, Midwestern university campus. Sci Total Environ [Internet]. 2017 Dec [cited 2019 Sep 2];599–600:1363–1368. Available from: https://linkin ghub.elsevier.com/retrieve/pii/S0048969717311804
- [114] Montgomery K, Ryan TJ, Krause A, et al. Assessment of athletic health care facility surfaces for MRSA in the secondary school setting. J Environ Health [Internet]. 2010 [cited 2019 Sep 2];72(6):8–11;quiz 66. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 20104827
- [115] Oller AR, Province L, Curless B. Staphylococcus aureus recovery from environmental and human locations in 2 collegiate athletic teams. J Athl Train [Internet]. 2010 May [cited 2019 Sep 2];45(3):222–229. Available from: http://www.ncbi.nlm.nih.gov/pubmed/20446834
- [116] Tonn K, Ryan TJ. Community-associated methicillin-resistant Staphylococcus aureus in college residential halls. J Environ Health [Internet]. 2013 [cited 2019 Sep 2];75(6):44–49. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/23397649
- [117] Kassem II, Sigler V, Esseili MA. Public computer surfaces are reservoirs for methicillin-resistant staphylococci. Isme J [Internet]. 2007 Jul 31 [cited 2019 Sep 2];1(3):265–268. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/18043636
- [118] Boa TT, Rahube TO, Fremaux B, et al. Prevalence of methicillin-resistant staphylococci species isolated from computer keyboards located in secondary and postsecondary schools. J Environ Health [Internet]. 2013 [cited 2019 Sep 2];75(6):50–58. Available from: http://www.ncbi.nlm.nih.gov/pubmed/23397650
- [119] Amini R, As A, Chung C, et al. Circulation and transmission of methicillin-resistant Staphylococcus aureus among college students in Malaysia (cell phones as reservoir). Asian Biomed [Internet]. 2012 [cited 2019 Sep 2];6(5):659–673. Available from: http://eprints.umsha.ac.ir/1080/1/%5BAsian Biomedicine%5D Circulation and transmission of methicillin-resistant Staphylococcus aureus among college students in Malaysia %28cell phones as reservoir%29.pdf
- [120] Amini R. Isolation and identification of methicillin-resistant *Staphylococcus aureus* from keys of college students using different detection methods. Br Biotechnol J [Internet]. 2012 Jan 10 [cited 2019 Sep 2];2(1):13–25. Available from: http://www.science domain.org/abstract.php?iid=93&id=11&aid=331
- [121] Chuang VW, Tsang IH, Keung JP, et al. Infection control intervention on meticillin resistant *Staphylococcus aureus* transmission in residential care homes for the elderly. J Infect Prev [Internet]. 2015 Mar 27 [cited 2019 Sep 2];16:58–66. Available from: http://www. ncbi.nlm.nih.gov/pubmed/28989403

- [122] Goettsch W, Geubbels E, Wannet W, et al. MRSA in nursing homes in the Netherlands 1989 to 1998: a developing reservoir? Eurosurveillance [Internet].
 2000 Mar 1 [cited 2019 Sep 2];5(3):28–31. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 12631868
- [123] Baum H, Von, Schmidt C, Svoboda D, et al. Risk factors for methicillin-resistant *Staphylococcus aureus* carriage in residents of german nursing homes. Infect Control Hosp Epidemiol [Internet]. 2002 Sep 2 [cited 2019 Sep 2];23(9):511–515. Available from: http://www. ncbi.nlm.nih.gov/pubmed/12269448
- [124] Kerttula A-M, Lyytikainen O, Vuopio-Varkila J, et al. Molecular epidemiology of an outbreak caused by methicillin-resistant *Staphylococcus aureus* in a health care ward and associated nursing home. J Clin Microbiol [Internet]. 2005 Dec 1 [cited 2019 Sep 2];43:6161–6163. Available from: http://www. ncbi.nlm.nih.gov/pubmed/16333120
- [125] Vovko P, Retelj M, Cretnik TZ, et al. Risk factors for colonization with methicillin-resistant *Staphylococcus aureus* in a long-term-care facility in Slovenia. Infect Control Hosp Epidemiol [Internet]. 2005 Feb 21 [cited 2019 Sep 2];26:191–195. Available from: http://www. ncbi.nlm.nih.gov/pubmed/15756891
- [126] Ho P-L, Wang TKF, Ching P, et al. Epidemiology and genetic diversity of methicillin-resistant *Staphylococcus aureus* strains in residential care homes for elderly persons in Hong Kong. Infect Control Hosp Epidemiol [Internet]. 2007 Jun 2 [cited 2019 Sep 2];28:671–678. Available from: http://www. ncbi.nlm.nih.gov/pubmed/17520539
- [127] O'Fallon E, Schreiber R, Kandel R, et al. Multidrugresistant gram-negative bacteria at a long-term care facility: assessment of residents, healthcare workers, and Inanimate surfaces. Infect Control Hosp Epidemiol [Internet]. 2009 Dec 2 [cited 2019 Sep 2];30(12):1172–1179. Available from: http://www. ncbi.nlm.nih.gov/pubmed/19835474
- [128] LUDDEN C, BRENNAN G, MORRIS D, et al. Characterization of methicillin-resistant *Staphylococcus aureus* from residents and the environment in a long-term care facility. Epidemiol Infect [Internet]. 2015 Oct 2 [cited 2019 Sep 2];143(14):2985–2988. Available from: http://www.ncbi.nlm.nih.gov/pubmed/25640407
- [129] Ustun C, Cihangiroglu M. Health care workers' mobile phones: A potential cause of microbial crosscontamination between hospitals and community. J Occup Environ Hyg. 2012;9(9):538–542.
- [130] Pal S, Juyal D, Adekhandi S, et al. Mobile phones: reservoirs for the transmission of nosocomial pathogens. Adv Biomed Res [Internet]. 2015 Jul 27;4:144. Available from: http://www.ncbi.nlm.nih. gov/pmc/articles/PMC4549928/
- [131] Morris TC, Moore LS, Shaunak S. Doctors taking a pulse using their mobile phone can spread MRSA. BMJ [Internet]. 2012;344(January):e412. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/22252697.
- [132] Angelakis E, Azhar El, Bibi F, et al. Paper money and coins as potential vectors of transmissible disease. Future Microbiol [Internet]. 2014;9(2):249–261. Available from:
- [133] Gedik H, Voss TA, Voss A. Money and transmission of bacteria. Antimicrob Resist Infect Control [Internet]. 2013;2(22):1–3. Available from: http://www.aricjour nal.com/content/2/1/22.

- [134] Heyba M, Ismaiel M, Alotaibi A, et al. Microbiological contamination of mobile phones of clinicians in intensive care units and neonatal care units in public hospitals in Kuwait. BMC Infect Dis [Internet]. 2015;15 (1):434. Available from: http://bmcinfectdis.biomed central.com/articles/10.1186/s12879-015-1172-9
- [135] Tambe NN, Pai CA. Study of microbial flora and mrsa harboured by mobile phones of health care personnel [Internet]. Int J Recent Trend SciTechnol [cited 2019 Sep 2]. 2012;4. Available from: https://statperson.com/ Journal/ScienceAndTechnology/Article/ Volume4lssue1/A Study1.pdf
- [136] Angadi K, Gupta U, Misra R, et al. Study of the role of mobile phones in the transmission of Hospital acquired infections. Med J Dr DY Patil Univ [Internet].
 2014 [cited 2019 Sep 2];7(4):435. Available from: http://www.mjdrdypu.org/text.asp?2014/7/4/435/ 135256
- [137] Kim J, Kwon O, Song W, et al. Isolation of healthcare-associated pathogens from cellular phones used by medical personnel. Korean J Nosocom Infect Control. 2010;15(1):36–40.
- [138] Datta P, Rani H, Chander J, et al. Bacterial contamination of mobile phones of health care workers. Indian J Med Microbiol [Internet]. 2009 [cited 2019 Sep 19];27
 (3):279. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/19584520
- [139] Singh D, Kaur H, Gardner WG, et al. Bacterial contamination of hospital pagers. Infect Control Hosp Epidemiol [Internet]. 2002 May 2 [cited 2019 Sep 2];23(5):274–276. Available from: http://www. ncbi.nlm.nih.gov/pubmed/12026153
- [140] Furuhata K, Ishizaki N, Isolation SK. Identification and antibacterial susceptibility of *Staphylococcus* spp. Associated with the mobile phones of university students. Biocontrol Sci. 2016.21(2):91–8.
- [141] Tekerekoğlu MS, Duman Y, Serindağ A, et al. Do mobile phones of patients, companions and visitors carry multidrug-resistant hospital pathogens? Am J Infect Control. 2011;39(5):379–381.
- [142] Selim HS, Abaza AF. Microbial contamination of mobile phones in a health care setting in Alexandria, Egypt. GMS Hyg Infect Control [Internet]. 2015;10: Doc03. Available from: http://www.pubmedcentral. nih.gov/articlerender.fcgi?artid=4332273&tool= pmcentrez&rendertype=abstract
- [143] Julian T, Singh A, Rousseau J, et al. Methicillin-resistant staphylococcal contamination of cellular phones of personnel in a veterinary teaching hospital. BMC Res Notes [Internet]. 2012;5:193. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/22533923% 5Cnhttp://www.pubmedcentral.nih.gov/articlerender. fcgi?artid=PMC3393609
- [144] Tolba O, Loughrey A, Goldsmith CE, et al. Survival of epidemic strains of nosocomialand community-acquired methicillin-resistant *Staphylococcus aureus* on coins. Am J Infect Control. 2007;35(5):342–346.
- [145] Taylor J, Davies M, Canales M, et al. The persistence of flood-borne pathogens on building surfaces under drying conditions. Int J Hyg Environ Health [Internet]. 2013;216(1):91–99. Available from: http://www.science direct.com/science/article/pii/S1438463912000491.
- [146] Neel R. Multidrug resistance of isolates of miticilin resistant *Staphylococcus aureus* (MRSA) in paper currency notes from meat sellers in Tanga, Tanzania. Int J LifeSc Bt Pharm Res [Internet]. 2012 [cited 2019

Sep 19];1(4):8–14. Available from: http://www.ijlbpr. com/uploadfile/2015/0414/20150414033401317.pdf

- [147] Neel R. Multidrug resistance of isolates of methicillin resistant Staphylococcus aureus (MRSA) in paper currency notes from restaurants and hotels in Lusaka in Zambia. Int J Pharm Pharm Sci [Internet]. 2013 [cited 2019 Sep 19];5(1):363–366. Available from: https:// www.researchgate.net/publication/282726219_ Multidrug_resistance_of_isolates_of_Methicillin_resis tant_Staphylococcus_aureus_MRSA_in_paper_cur rency_notes_from_restaurants_and_hotels_in_ Lusaka_in_Zambia
- [148] Nagajothi JD, Vigneshwaran S, Kumar RP, et al. Study of prevalence of microbial contamination with its antibiotic resistance pattern in automated teller machine in and around Puducherry, India. J Earth, Environ Heal Sci [Internet]. 2015 [cited 2019 Sep 2];1(1):27. Available from: http://www.ijeehs.org/article.asp?=2423-7752; year=2015;volume=1;issue=1;spage=27;epage=31; aulast=Nagajothi
- [149] Tekerekoğlu MS, Yakupogullari Y, Otlu B, et al. Bacteria found on banks automated teller machines (ATMs). African J Microbiol Res [Internet]. 2013 Apr 16 [cited 2019 Sep 2];7(16):1619–1621. Available from: http:// academicjournals.org/journal/AJMR/article-abstract /F5E8A0813069
- [150] Zhang M, O'Donoghue M, Boost MV. Characterization of staphylococci contaminating automated teller machines in Hong Kong. Epidemiol Infect [Internet]. 2012 Aug 19 [cited 2019 Sep 2];140(8):1366–1371. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/22011397
- [151] Chukwudozie Onuoha S, Fatokun K. Bacterial contamination and public health risk associated with the use of banks' automated teller machines (atms) in Ebonyi State, Nigeria. Am J Public Heal Res [Internet]. 2014 Mar 9 [cited 2019 Sep 2];2(2):46–50. Available from: http://pubs.sciepub.com/ajphr/2/2/2/index.html
- [152] Sribenjalux P, Palarach W, Wannarat A, et al. Bacterial contamination on automatic teller machine keypad in Khon Kaen University. J Med Technol Assoc Thail [Internet]. 2011 Apr 1 [cited 2019 Sep 2];39(1). Available from: http://jmt-amtt.com/?journal=jmtamtt&page=article_view&path%5B%5D=93
- [153] Angbuhang KB, Neupane M, Adhikari A, et al. Detection of methicillin resistant *Staphylococcus aureus* in public transportation of Kathmandu Valley, Nepal. Tribhuvan Univ J Microbiol [Internet]. 2018 Sep 26 [cited 2019 Sep 2];5:51–56. Available from: https://www.nepjol.info/index.php/tujm/article/view/ 22312
- [154] Stepanović S, Ćirković I, Djukić S, et al. Public transport as a reservoir of methicillin-resistant staphylococci. Lett Appl Microbiol [Internet]. 2008 Oct 1 [cited 2019 Sep 2];47(4):339–341. Available from:
- [155] Simões RR, Aires-de-Sousa M, Conceição T, et al. High prevalence of EMRSA-15 in Portuguese Public Buses: A worrisome finding. Otto M, editor. PLoS One [Internet]. 2011 Mar 2 [cited 2019 Sep 2];6(3):e17630. Available from: https://dx.plos.org/10.1371/journal. pone.0017630
- [156] Iwao Y, Yabe S, Takano T, et al., Isolation and molecular characterization of methicillin-resistant *Staphylococcus aureus* from public transport. Microbiol Immunol [Internet]. 2012 [cited 2019 Sep 2];56:76–82. Available from: https://onlinelibrary.wiley.com/doi/pdf/10.1111/j. 1348-0421.2011.00397.x

- [157] Gaymard A, Pichon M, Degaud M, et al. Staphylococcus aureus in the environment of public transport: data from the metropolitan network in Lyon, France. Int J Antimicrob Agents [Internet]. 2016 Oct [cited 2019 Sep 2];48(4):459–462. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/27575729
- [158] Conceição T, Diamantino F, Coelho C, et al. Contamination of public buses with MRSA in Lisbon, Portugal: A possible transmission route of major MRSA clones within the community. PLoS One. 2013;8 (11):1–6.
- [159] Lutz JK, van Balen J, Mac CJ, et al. Methicillin-resistant Staphylococcus aureus in public transportation vehicles (buses): another piece to the epidemiologic puzzle. Am J Infect Control [Internet]. 2014 Dec [cited 2019 Sep 2];42(12):1285–1290. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/25465258
- [160] Â M, Martins da Costa P, Rego D, et al. Contamination of public transports by *Staphylococcus aureus* and its carriage by biomedical students: point-prevalence, related risk factors and molecular characterization of methicillin-resistant strains. Public Health [Internet]. 2015 Aug [cited 2019 Sep 2];129(8):1125–1131. Available from: https://linkinghub.elsevier.com/ retrieve/pii/S0033350615002085
- [161] Peng Y, Ou Q, Lin D, et al. Metro system in Guangzhou as a hazardous reservoir of methicillin-resistant Staphylococci: findings from a point-prevalence molecular epidemiologic study. Sci Rep [Internet]. 2015 Dec 29 [cited 2019 Sep 2];5:16087. Available from: http:// www.nature.com/articles/srep16087
- [162] Schaumburg F, Köck R, Leendertz FH, et al. Airport door handles and the global spread of antimicrobial-resistant bacteria: across sectional study. Clin Microbiol Infect. 2016;22(12):1010–1011.
- [163] Chowdhury T, Mahmud A, Barua A, et al. Bacterial contamination on hand touch surfaces of public buses in Chittagong City, Bangladesh. J Environ Sci Toxicol Food Technol [Internet]. 2016 [cited 2019 Sep 2];10(4):48–55. Available from: https://www. researchgate.net/publication/303273428_Bacterial_ Contamination_On_Hand_Touch_Surfaces_Of_ Public_Buses_in_Chittagong_City_Bangladesh
- [164] Lin JL, Peng Y, Ou QT, et al. A molecular epidemiological study of methicillin-resistant Staphylococci environmental contamination in railway stations and coach stations in Guangzhou of China. Lett Appl Microbiol. 2017 Feb 1;64(2):131–137.
- [165] Storr J, Twyman A, Zingg W, et al. Core components for effective infection prevention and control programmes: new WHO evidence-based recommendations. Antimicrob Resist Infect Control [Internet]. 2017 Jan 10 [cited 2020 Sep 2];6:1–18. Available from: www.who.int/
- [166] Kurutkan MN, Kara O, Eraslan İH. An implementation on the social cost of hospital acquired infections. Int J Clin Exp Med. 2015;8(3):4433–4445.
- [167] Stone PW. Economic burden of healthcare-associated infections: an American perspective. Rev Pharmacoecon Outcomes Res. 2009;9(5):417–422.
- [168] Graves N, Nicholls TM, Morris AJ. Modeling the Costs of Hospital-Acquired Infections in New Zealand. Infect Control Hosp Epidemiol [Internet]. 2003 Mar 2 [cited 2018 Aug 8];24(3):214–223. Available from: http:// www.ncbi.nlm.nih.gov/pubmed/12683515
- [169] Klevens RM, Edwards JR, Richards CL, et al. Estimating Health Care-Associated Infections and Deaths in. 433021. 2007;122(April):160–166.

- [170] Lobdell KW, Stamou S, Sanchez JA. H o s p i t a l A c q u i re d Infections. Surgical Clinics of North America. 2012;92:65–77.
- [171] Cardoso T, Arag I, Costa-pereira A. Differences in Microbiological Profile between and Hospital-Acquired Infections. Acta Med Port. 2013;26 (4):377–384.
- [172] Durai R, Ng PCH, Hoque H. *Staphylococcus aureus*: an update. AORN J. 2010;91(5):599–609.
- [173] Nelson MU, Gallagher PG. Methicillin-resistant Staphylococcus aureus in the neonatal intensive care unit. Semin Perinatol. 2013;36(6):424–430.
- [174] Usukura Y, Igarshi T. Examination of severe, hospital acquired infections affecting extremely low birthweight (ELBW) infants. Pediatr Int. 2003;46 (January):230–232.
- [175] Sista RR, Oda G, Methicillin-resistant BJ. *Staphylococcus aureus* infections in ICU patients. Anesthesiology Clinics of North America. 2004;22:405–435.
- [176] Achermann Y, Seidl K, Kuster SP, et al. Epidemiology of Methicillin-Susceptible Staphylococcus aureus in a Neonatology Ward. Infect Control Hosp Epidemiol [Internet]. 2015;36(11):1305–1312. Available from: https://www.cambridge.org/core/product/identifier/ S0899823X15001841/type/journal_article
- [177] Saadoun I, Jaradat ZW, Al TIA, et al. Airborne methicillin-resistant *Staphylococcus aureus* in the indoor environment of King Abdullah University Hospital, Jordan. Indoor Built Environ [Internet]. 2015 May 19 [cited 2019 Sep 2];24(3):315–323. Available from: http://journals.sagepub.com/doi/10.1177/ 1420326X14526604
- [178] Siegel JD, Rhinehart E, Jackson M, et al. 2007 guideline for isolation precautions : preventing transmission of infectious agents in health care settings. Am J Infect Control. 2007;65:S65–S164.
- [179] Haun N, Hooper-lane C, Safdar N. Healthcare Personnel Attire and Devices as Fomites : A Systematic Review. Infect Control Hosp Epidemiol. 2016.37 (11):1367–1373.
- [180] Kim JS, Kim HS, Park JY, et al. Contamination of X-ray cassettes with methicillin-resistant *Staphylococcus aureus* and methicillin-resistant *Staphylococcus haemolyticus* in a radiology department. Ann Lab Med. 2012;32 (3):206–209.
- [181] Campos-Murguía A, León-Lara X, Muñoz JM, et al. Stethoscopes as potential intrahospital carriers of pathogenic microorganisms. Am J Infect Control. 2014;42(1):82–83.
- [182] Hogue MH, Heilmann KP, Callaghan JJ. Wearing ID badges in the operating room environment: is reconsideration warranted? J Arthroplasty. 2017 Jul;32 (7):2231–2233.
- [183] Huang R, Mehta S, Weed D, et al. Methicillin-resistant *Staphylococcus aureus* survival on hospital fomites. Infect Control Hosp Epidemiol. 2006;27(11):1267–1269.
- [184] Neely AN, Maley MP. Survival of enterococci and Staphylococci on hospital fabrics and plastics. J Clin Microbiol. 2000;38(2):724–726.
- [185] Watson SP, Clements MO, Foster SJ. Characterization of the starvation-survival response of *Staphylococcus aureus*. J Bacteriol. 1998;180(7):1750–1758.
- [186] Wagenvoort JH, Penders RJ. Long-term in-vitro survival of an epidemic MRSA phage-group III-29 strain. J Hosp Infect [Internet]. 1997 Apr [cited 2019 Sep 2];35(4):322–325. Available from: http://www. ncbi.nlm.nih.gov/pubmed/9152828

- [187] Zarpellon MN, Gales AC, Sasaki AL, et al. Survival of vancomycin-intermediate *Staphylococcus aureus* on hospital surfaces. J Hosp Infect [Internet]. 2015 Aug [cited 2019 Sep 2];90(4):347–350. Available from: http://www.ncbi.nlm.nih.gov/pubmed/25986166
- [188] Koca O, Altoparlak U, Ayyildiz A, et al. Persistence of nosocomial pathogens on various fabrics. Eurasian J Med. 2012;44(1):28–31.
- [189] Williams C, Davis DL. Methicillin-resistant Staphylococcus aureus fomite survival. Clin Lab Sci. 2009;22(1):34.
- [190] Oie S, Kamiya A. Survival of methicillin-resistant *Staphylococcus aureus* (MRSA) on naturally contaminated dry mops. J Hosp Infect. 1996;34(2):145–149.
- [191] Stephens B, Azimi P, Thoemmes MS, et al. Microbial exchange via fomites and implications for human health. Curr Pollut Reports [Internet]. 2019;5 (4):198–213. Available from:.
- [192] Sexton T, Clarke P, Neill EO, et al. Environmental reservoirs of methicillin-resistant *Staphylococcus aureus* in isolation rooms : correlation with patient isolates and implications for hospital hygiene. J Hosp Infect. 2006;62:187–194.
- [193] Boyce JM, Potter-bynoe G, Chenevert C, et al. Enironmental contamination due to methiclillin-resistant *Staphlylococcus auresus*: Possible infection control implications. Infect Control Hosp Epidemiol. 1997;18 (9):622–627.
- [194] Boyce JM, Havill NL, Otter JA, et al. Widespread environmental contamination associated with patients with diarrhea and methicillin-resistant *Staphylococcus aureus* colonization of the gastrointestinal tract. Infect Control Hosp Epidemiol. 2007.28(10):1142–7.
- [195] Rohr U, Kaminski A, Wilhelm M, et al. Colonization of patients and contamination of the patients ' environment by MRSA under conditions of single-room isolation. Int J Hyg Environ Health. 2009;212:209–215.
- [196] Oie S, Suenaga S, Sawa A, et al. Association between isolation sites of methicillin-resistant *Staphylococcus aureus* (MRSA) In patients with MRSA-positive body sites and mrsa contamination in their surrounding environmental surfaces. Jpn J Infect Health. 2007;367–369.
- [197] Stiefel U, Cadnum JL, Eckstein BC, et al. Contamination of hands with methicillin-resistant *Staphylococcus aureus* after contact with environmental surfaces and after contact with the skin of colonized patients. Infect Control Hosp Epidemiol. 2014;32(2):7–10.
- [198] Desai R, Pannaraj PS, Agopian J, et al. Survival and transmission of community-associated methicillin-resistant *Staphylococcus aureus* from fomites. Am J Infect Control. 2011 Apr;39(3):219–225.
- [199] Moore G, Dunnill CW, Wilson APR. The effect of glove material upon the transfer of methicillin-resistant *Staphylococcus aureus* to and from a gloved hand. Am J Infect Control [Internet]. 2013 Jan [cited 2020 Apr 20];41(1):19–23. Available from: http://www.ncbi. nlm.nih.gov/pubmed/22981164
- [200] Saba CKS, Amenyona JK, Kpordze SW. Prevalence and pattern of antibiotic resistance of *Staphylococcus aureus* isolated from door handles and other points of contact in public hospitals in Ghana. Antimicrob Resist Infect Control [Internet]. 2017 Dec 10 [cited 2018 Aug 8];6(1):44. Available from: http://aricjournal.bio medcentral.com/articles/10.1186/s13756-017-0203-2
- [201] Torkar KG, lvić S. Surveillance of bacterial colonisation on contact surfaces in different medical wards. Arch

Ind Hyg Toxicol [Internet]. 2017 Jun 27 [cited 2018 Aug 8];68(2):116–126. Available from: http://www. ncbi.nlm.nih.gov/pubmed/28665797

- [202] Huslage K, Rutala WA, Sickbert E, et al. A quantitative approach to defining " high-touch " surfaces in hospitals. Infect Control Hosp Epidemiol. 2012;31 (8):850–853.
- [203] Ferreira AM, De AD, Rigotti MA, et al. Methicillinresistant *Staphylococcus aureus* on surfaces of an Intensive Care Unit. Acta Paul. enferm. 2011;24 (4):453–458.
- [204] van Balen J, Bottichio L, Stevenson K, et al. Understanding the introduction and circulation of environmental methicillin-resistant *Staphylococcus aureus* in a large academic medical center during a nonoutbreak, year-long period. Am J Infect Control. 2016;44(8):925–930.
- [205] Güven Gökmen T, Kalayci Y, Yaman A, et al. Molecular characterization of methicillin-resistant *Staphylococcus aureus* strains by spa typing and pulsed field gel electrophoresis methods. BMC Microbiol [Internet]. 2018 Dec 24 [cited 2019 Sep 2];18(1):155. Available from: https://bmcmicrobiol.biomedcentral.com/articles/10. 1186/s12866-018-1305-6
- [206] Bendary MM, Solyman SM, Azab MM, et al. Genetic diversity of multidrug resistant *Staphylococcus aureus* isolated from clinical and non clinical samples in Egypt. Cell Mol Biol (Noisy-le-grand) [Internet]. 2016 Aug 31 [cited 2019 Sep 2];62(10):55–61. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 27609475
- [207] Jaradat ZW, Ababneh QO, Saraireh S, et al. Analysis of genetic heterogeneity of *Staphylococcus aureus* strains isolated from food and clinical samples from northern Jordan using VNTR, toxin profiles and antibiograms. Malays J Microbiol [Internet]. 2016 Sep [cited 2019 Sep 19];12(3):254–264. Available from: http://mjm.usm. my/index.php?r=cms/entry/view&id=1007&slug= Analysis-of-genetic-heterogeneity-of-Sta phylo co ccus -aureus-strains-isolated-from-food-and-clinical-samples -from-northern-Jordan-using-VNTR%2C-toxin-profi lesand-antibiograms
- [208] Ndawula EM, Brown L. Mattresses as reservoirs of epidemic methicillin-resistant *Staphylococcus aureus*. Lancet. 1991;337:488.
- [209] Embil JM, Mcleod JA, Al-barrak AM, et al. An outbreak of methicillin resistant *Staphylococcus aureus* on a burn unit : potential role of contaminated hydrotherapy equipment. Burns. 2001;27:681–688.
- [210] Rampling A, Wiseman S, Davis L, et al. Evidence that hospital hygiene is important in the control of methicillin-resistant *Staphylococcus aureus*. J Hosp Infect. 2001;49:109–116.
- [211] Schultsz C, Meester HHMM, Kranenburg AMHH, et al. Ultra-sonic nebulizers as a potential source of methicillin-resistant *Staphylococcus aureus* causing an outbreak in a university tertiary care hospital. J Hosp Infect. 2003;55(4):269–275.
- [212] Phoon HYP, Hussin H, Hussain BM, et al. Distribution, genetic diversity and antimicrobial resistance of clinically important bacteria from the environment of a tertiary hospital in Malaysia. J Glob Antimicrob Resist [Internet]. 2018 Sep [cited 2019 Sep 19];14:132–140. Available from: http://www.ncbi. nlm.nih.gov/pubmed/29540306
- [213] Morgan DJ, Rogawski E, Thom KA, et al. Transfer of multidrug-resistant bacteria to healthcare workers'

gloves and gowns after patient contact increases with environmental contamination. Crit Care Med. 2012;40(4):1045–1051.

- [214] Snyder GM, Thom KA, Furuno JP, et al. Detection of methicillin-resistant *Staphylococcus aureus* and vancomycin-resistant enterococci by healthcare workers on infection control gown and gloves. Infect Control Hosp Epidemiol. 2008;29(7):583–589.
- [215] Chen K, Chen L, Wang Y. Contamination of medical charts : an important source of potential infection in hospitals. PLOS One. 2014;9(2):1–7.
- [216] Kurashige EJO, Oie S, Furukawa H. Contamination of environmental surfaces by methicillin-resistant *Staphylococcus aureus* (MRSA) in rooms of inpatients with MRSA-positive body sites. Brazilian J Microbiol. 2016;47(3):703–705.
- [217] Faires MC, Pearl DL, Berke O, et al. The identification and epidemiology of meticillin-resistant *Staphylococcus aureus* and Clostridium difficile in patient rooms and the ward environment. BMC Infectious Diseases. 2013;24:1–13.
- [218] Andrade C, Champagne S, Caruso D, et al. Staphylococcus aureus: an assessment of environmental contamination in a burn center. Am J Infect Control [Internet]. 2009 Aug [cited 2019 Sep 2];37(6):515–517. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 19643281
- [219] Andrade C, Champagne S, Caruso D, et al. Staphylococcus aureus: an assessment of environmental contamination in a burn center. Am J Infect Control [Internet]. 2009 Aug [cited 2019 Sep 2];37(6):515–517. Available from: https://linkinghub.elsevier.com/ retrieve/pii/S0196655308009024
- [220] Oie S, Hosokawa I, Kamiya A. Contamination of room door handles by methicillin-sensitive/methicillinresistant *Staphylococcus aureus*. J Hosp Infect. 2002;51(2):140–143.
- [221] Brady RRW, Kalima P, Damani NN, et al. Bacterial contamination of hospital bed control handsets in a surgical setting: A potential marker of contamination of the health care environment. Ann R Coll Surg Engl. 2007;89(7):656–660.
- [222] Young JM, Naqvi M, Richards L. Microbial contamination of hospital bed handsets. Am J Infect Control. 2002;33:170–174.
- [223] Otter JA, Passaretti CL, French GL, et al. Low frequency of environmental contamination with methicillin-resistant *Staphylococcus aureus* in an inner city emergency department and a human immunodeficiency virus outpatient clinic. Am J Infect Control [Internet]. 2011 Mar [cited 2019 Sep 2];39(2):151–153. Available from: https://linkin ghub.elsevier.com/retrieve/pii/S0196655310007364
- [224] Lemmen S, Häfner H, Zolldann D, et al. Distribution of multi-resistant Gram-negative versus Gram-positive bacteria in the hospital inanimate environment. J Hosp Infect [Internet]. 2004 Mar [cited 2019 Sep 2];56(3):191–197. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 15003666
- [225] Pandey A, Asthana A, Tiwari R, et al. Physician accessories: doctor, what you carry is every patient's worry? Indian J Pathol Microbiol [Internet]. 2010 [cited 2019 Sep 2];53(4):711. Available from: http://www.ncbi.nlm. nih.gov/pubmed/21045398
- [226] Messina G, Ceriale E, Lenzi D, et al. Environmental contaminants in hospital settings and progress in

disinfecting techniques. BioMed Research International. 2013;23:139.

- [227] Shelly MJ, Scanlon TG, Ruddy R, et al. Meticillinresistant Staphylococcus aureus (MRSA) environmental contamination in a radiology department. Clin Radiol [Internet]. 2011 Sep [cited 2019 Sep 2];66(9):861–864. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/21676384
- [228] Panhotra BR, Saxena AK, Al-Mulhim AS. Contamination of patients' files in intensive care units: an indication of strict handwashing after entering case notes. Am J Infect Control [Internet]. 2005 Sep [cited 2019 Sep 2];33(7):398–401. Available from: http://www. ncbi.nlm.nih.gov/pubmed/16153486
- [229] Omaki MO, Orioka KY, le SO, et al. *Staphylococcus aureus* contamination on the surface of working tables in ward staff centers and its preventive methods. Biol Pharm Bull. 2006;29(7):1508–1510.
- [230] Lu P-L, Siu LK, Chen T-C, et al. Methicillin-resistant Staphylococcus aureus and Acinetobacter baumannii on computer interface surfaces of hospital wards and association with clinical isolates. BMC Infect Dis. 2009;9:164.
- [231] Baruah J, Kumar S, Gratrix A. Blood pressure cuffs as a potential fomite for transmission of pathogenic micro-organisms: A prospective study in a university teaching hospital. Br J Infect Control [Internet]. 2008 [cited 2019 Sep 2];9(4). Available from: http://www
- [232] Walker N, Gupta R, Cheesbrough J. Blood pressure cuffs: friend or foe? J Hosp Infect [Internet]. 2006 Jun [cited 2019 Sep 2];63(2):167–169. Available from: http://www.ncbi.nlm.nih.gov/pubmed/16616799
- [233] Jain K, Chavan NS, Jain SM. Blood pressure cuff-as a fomite for cross infection [Internet]. Int.J.Curr. Microbiol.App.Sci. [cited 2019 Sep 2]. 2014;3. Available from: http://www.ijcmas.com
- [234] Jeyakumari D, Nagajothi S, Kumar RP, et al. Bacterial colonization of blood pressure cuff: A potential source of pathogenic organism: A prospective study in a teaching hospital. Med Sci J Com [Internet]. 2016;35–37. Available from: http://www.medicalscien cejournal.com/download/173/2-6-17-864.pdf
- [235] Berman DS, Schaefler S, Simberkoff MS, et al. Tourniquets and nosocomial methicillin-resistant *Staphylococcus aureus* infections. N Engl J Med. 1986;315(8):514–515.
- [236] Mehmood Z, Mubeen SM, Afzal MS, et al. Potential risk of cross-infection by tourniquets: a need for effective control practices in pakistan. Int J Prev Med [Internet]. 2014 Sep [cited 2019 Sep 2];5(9):1119–1124. Available from: http://www.ncbi.nlm.nih.gov/pubmed/ 25317294
- [237] Martin P, Abou Chakra CN, Williams V, et al. Prevalence of antibiotic-resistant organisms in Canadian hospitals. Comparison of point-prevalence survey results from 2010, 2012, and 2016. Infect Control Hosp Epidemiol [Internet]. 2019 Jan 1 [cited 2020 Apr 20];40:53–59. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/30394232
- [238] Cimolai N. MRSA and the environment: implications for comprehensive control measures. Eur J Clin Microbiol Infect Dis. 2008;27:481–493.

- [239] Köck R, Becker K, Cookson B, et al. Methicillin-resistant *Staphylococcus aureus* (MRSA): burden of disease and control challenges in Europe. Eurosurveillance. 2010;15:1–9.
- [240] Otter JA, French GL. Molecular epidemiology of community-associated meticillin-resistant *Staphylococcus aureus* in Europe [Internet]. Lancet Infect Dis [cited 2020 Apr 20]. 2010;10:227–239. Available from: http://www.ncbi.nlm.nih.gov/ pubmed/20334846
- [241] McGough SF, MacFadden DR, Hattab MW, et al. Rates of increase of antibiotic resistance and ambient temperature in Europe: a cross-national analysis of 28 countries between 2000-2016. bioRxiv 414920.2018.
- [242] Walter J, Noll I, Feig M, et al. Decline in the proportion of methicillin resistance among *Staphylococcus aureus* isolates from non-invasive samples and in outpatient settings, and changes in the co-resistance profiles: an analysis of data collected within the Antimicrobial Resistance Surveillance Network, Germany 2010 to 2015. BMC Infect Dis [Internet]. 2017 Feb 23 [cited 2020 Sep 3];17(1). Available from: /pmc/articles/ PMC5324250/?report=abstract
- [243] Nikolaras GP, Papaparaskevas J, Samarkos M, et al. Changes in the rates and population structure of methicillin-resistant *Staphylococcus aureus* (MRSA) from bloodstream infections: A single-centre experience (2000–2015). J Glob Antimicrob Resist. 2019 Jun 1;17:117–122.
- [244] Guthrie JL, Teatero S, Hirai S, et al. Genomic epidemiology of invasive methicillin-resistant *Staphylococcus aureus* infections among hospitalized individuals in Ontario, Canada. J Infect Dis [Internet]. 2020 May 20 [cited 2020 Sep 3]. Available from: https://academic. oup.com/jid/advance-article/doi/10.1093/infdis/ jiaa147/5841126
- [245] Nichol KA, Adam HJ, Golding GR, et al. Characterization of MRSA in Canada from 2007 to 2016. J Antimicrob Chemother [Internet]. 2019 Aug 1 [cited 2020 Sep 3];74:iv55–63. Available from: https://academic.oup.com/jac/article/74/ Supplement_4/iv55/5553082
- [246] Kavanagh KT, Abusalem S, Calderon LE. The incidence of MRSA infections in the United States: is a more comprehensive tracking system needed? Antimicrob Resist Infect Control [Internet]. 2017;6(1):34. Available from:.
- [247] Kourtis AP, Hatfield K, Baggs J, et al. Vital signs: epidemiology and recent trends in methicillin-resistant and in methicillin-susceptible Staphylococcus aureus bloodstream infections — united states. MMWR Morb Mortal Wkly Rep [Internet]. 2019 Mar 8 [cited 2020 Sep 3];68:214–219. Available from: http://www. cdc.gov/mmwr/volumes/68/wr/mm6809e1.htm?s_ cid=mm6809e1_w
- [248] Dai Y, Liu J, Guo W, et al. Decreasing methicillinresistant *Staphylococcus aureus* (MRSA) infections is attributable to the disappearance of predominant MRSA ST239 clones, Shanghai, 2008–2017. Emerg Microbes Infect [Internet]. 2019 Jan 1 [cited 2020 Sep 3];8:471–478. Available from: /pmc/articles/ PMC6455123/?report=abstract

450 😉 Z. W. JARADAT ET AL.

- [249] Chiang CH, Pan SC, Yang TS, et al. Healthcareassociated infections in intensive care units in Taiwan, South Korea, and Japan: recent trends based on national surveillance reports. Antimicrob Resist Infect Control [Internet]. 2018 Nov 7 [cited 2020 Sep 3];7(1). Available from: https://pubmed.ncbi.nlm. nih.gov/30455867/
- [250] Williams V, Simor AE, Kiss A, et al. Is the prevalence of antibiotic-resistant organisms changing in Canadian hospitals? Comparison of point-prevalence survey results in 2010 and 2012. Clin Microbiol Infect [Internet]. 2015 Jun 1 [cited 2020 Sep 4];21:553–559. Available from: https:// pubmed.ncbi.nlm.nih.gov/25677630/
- [251] Kang Cl, Song JH. Antimicrobial resistance in Asia: current epidemiology and clinical implications. Infect Chemother. 2013 Mar;45(1):22–31.
- [252] Kim D, Ahn JY, Lee CH, et al. Increasing resistance to extended-spectrum cephalosporins, fluoroquinolone, and carbapenem in gram-negative bacilli and the emergence of carbapenem non-susceptibility in *klebsiella pneumoniae*: analysis of korean antimicrobial resistance monitoring system (KARMS) data from 2013 to 2015 [Internet]. Ann Lab Med [cited 2020 Sep 4]. 2017;37:231–239. Available from: https://pubmed.ncbi.nlm.nih.gov/ 28224769/