

Al-Azhar International Medical Journal

Volume 5 | Issue 8

Article 20

8-31-2024 Section: Plastic surgery

Incidence of Methicillin Resistant Staphylococcus Aureus Acquisition in Burn Intensive Care Unit, A meta-analysis study

AbdElrahman Awadeen AbdElrahman Department of Plastic and Burn Surgery, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt

Mohammed Abdel Hamid Basiouny Khedr Department of Clinical Pathology, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt

Mohammed Abdelsamie El-Zoeiky Department of Plastic and Burn Surgery, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt

Ahmed Seliem Hussein Mohammed Department of Plastic and Burn Surgery, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt, slym02190@gmail.com

Follow this and additional works at: https://aimj.researchcommons.org/journal

Part of the Medical Sciences Commons, Obstetrics and Gynecology Commons, and the Surgery Commons

How to Cite This Article

AbdElrahman, AbdElrahman Awadeen; Khedr, Mohammed Abdel Hamid Basiouny; El-Zoeiky, Mohammed Abdelsamie; and Mohammed, Ahmed Seliem Hussein (2024) "Incidence of Methicillin Resistant Staphylococcus Aureus Acquisition in Burn Intensive Care Unit, A meta-analysis study," *Al-Azhar International Medical Journal*: Vol. 5: Iss. 8, Article 20. DOI: https://doi.org/10.58675/2682-339X.2596

This Meta Analysis is brought to you for free and open access by Al-Azhar International Medical Journal. It has been accepted for inclusion in Al-Azhar International Medical Journal by an authorized editor of Al-Azhar International Medical Journal. For more information, please contact dryasserhelmy@gmail.com.

META-ANALYSIS

Incidence of Methicillin Resistant Staphylococcus Aureus Acquisition in Burn Intensive Care Unit, A meta-analysis study

AbdElrahman A. AbdElrahman ^a, Mohammed A. B. Khedr ^b, Mohammed A. El Zoeiky ^a, Ahmed S. H. Mohammed ^{a,*}

^a Department of Plastic and Burn Surgery, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt

^b Department of Clinical Pathology, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt

Abstract

Background: The skin acts as the body's first line of defense against harmful germs.

Aim: To assess the incidence and outcomes of MRSA acquisition in admitted patients within burn ICU in the last ten years. Patients and methods: The current meta-analysis involved 16 articles with a total of 7623 individuals, the majority of these studies were retrospective and only four studies were prospective most of these studies were from USA and China, and no studies from Egypt were found, so more prospective studies were needed to be performed among ICU burn patients in Egypt.

Results: In all, four studies examined the effect of MRSA on mortality rate. No significant heterogeneity was determined. Therefore, a fixed effect model was utilized for analysis ($I^2 = 0\%$, P = .56). The combined OR & 95% CIs was 1.28 (0.69 to 1.40). The combined result suggested that MRSA-positive patients are not associated with a higher mortality rate (Z = 0.78, P = .44).

Conclusion: The study found 14.7% MRSA infections in burn patients admitted to burn ICUs, with 3.8% colonization and 8.6% acquisition. Risk factors include age, burn surface area, male gender, inhalational injury, and burn cause.

Keywords: Staphylococcus Aureus; Methicillin Resistant; Acquisition; burn intensive care unit

1. Introduction

 \frown he skin acts as the primary barrier,

protecting against harmful microbes. This is lost with burns, along with systemic and topical immunosuppression. With significant morbidity Mortality for burn patients, infection is thus one of the most dangerous side effects.¹

When it comes to burn patients, MRSA has become one of the biggest concerns. Burn wounds provide a breeding ground for bacteria that may cause a deadly chain reaction that includes septicemia, invasive infection, multiorgan failure, burn wound sepsis, and death.²

Surveillance and prophylactic treatment have parallel roles in the prevention of infection. Besides that, medical education of medical staff plays a role when dealing with carriers or cases such as hand wash, antiseptic use, wise use of antibiotic therapy, and isolation equipment. Surveillance is the key restriction during the use of antibiotics.³

The objective of this research was to evaluate the incidence and outcomes of MRSA acquisition in admitted individuals within burn ICUs in the last ten years.

2. Patients and methods

The current meta-analysis involved 16 articles with a total of 7623 individuals; the majority of these studies were retrospective, and only four studies were prospective. Most of these studies were from the USA and China, and no studies from Egypt were found, so more prospective studies need to be performed among ICU burn patients in Egypt.

Accepted 21 August 2024. Available online 31 August 2024

* Corresponding author at: Plastic and Burn Surgery, Faculty of Medicine for Boys, Al-Azhar University, Cairo, Egypt. E-mail address: slym02190@gmail.com (A. S. H. Mohammed).

https://doi.org/10.58675/2682-339X.2596 2682-339X/© 2024 The author. Published by Al-Azhar University, Faculty of Medicine. This is an open access article under the CC BY-SA 4.0 license (https://creativecommons.org/licenses/by-sa/4.0/). Types of included studies: A literature search was conducted in PUBMED, EMBASE, ScienceDirect, Web of Science, Scopus, Ovid, Proquest, Google Scholar, and the Cochran Library. The search was limited to English articles published between 2012 and 2022. Databases were approached through the Egyptian Knowledge Bank. The search was conducted in December 2022.

Study eligibility criteria

Inclusion criteria: Retrospective, prospective, or cross-sectional studies, Patients with major burn, Patients admitted in burn ICU, Patients with MRSA positive (nasal or wound); you were able to report on at least one of the outcome measures, the data could be extracted from the entire articles.

Exclusion criteria: Every item that failed to satisfy the inclusion requirements was eliminated. Studies without control groups, case reports, meta-analyses, letters, expert opinions, abstracts, editorials, and reviews without original data were also disqualified. Research on languages other than English was not included.

Data collection and data items

Data from each study was extracted into a form with the following parameters according to a prespecified protocol: primary author, publication year, type of the research, number of patients, age of patients, TBSA, male gender, cause of burn, inhalational injury, DM, duration of hospital stay, duration of ICU stay, intubation, active infection, numbers of operation and Mortality.

Statistical analysis of the data and synthesis of results

Data were entered into the computer system for analysis utilizing version 20.100 of the MedCalc application. Statistical software significance was assigned to p-values that were less than or equal to 0.05, with a CI of ninety-five percent established. I2 (observed variance for heterogeneity) and Q (total variance for heterogeneity) were utilized to evaluate statistical heterogeneity. Qualitative data are presented as the total number of occurrences and the number of events, whereas quantitative data are presented as the mean standard deviation.

3. Results



Figure 1. PRISMA Flow Diagram of the Study Selection Process

A total of 16 examined were selected for the current analysis, including a total of 7623 individuals admitted to regional specialized burn units. The number of included patients ranged from 31 patients by Schweitzer et al.5 to 2036 patients by Cleland et al.¹⁹ (Table 1)

Table 1. Study Characteristics (N = 16 studies, 7623 patients)

1020 pui	criwj					
FIRST	YEAR	COUNTRY	DESIGN	STU	DY	SAMPLE
AUTHOR				PER		SIZE
				From	То	
KAISER ⁴	2011	USA	Retrospective	2007	2009	752
SCHWEITZER ⁵	2012	USA	Prospective	2009	2010	31
ALRAWI ⁶	2013	UK	Retrospective	NA	NA	139
RODRIGUES ⁷	2013	Brazil	Retrospective	2006	2009	367
ISSLER-	2015	Australia	Retrospective	2012	2013	357
FISHER ⁸						
JOHNSON ⁹	2015	USA	Retrospective	2011	2014	554
HAITH ¹⁰	2016	USA	Retrospective	2010	2011	826
DRUM ¹¹	2016	USA	Retrospective	2011	2016	725
TEJIRAM ¹²	2017	USA	Retrospective	2012	2014	601
PARK ¹³	2017	Australia	Retrospective	2013	2014	238
AMISSAH ¹⁴	2017	Ghana	Prospective	2014	2015	62
LIU ¹⁵	2018	China	Prospective	2012	2016	112
JIANG ^{`16}	2018	China	Prospective	2011	2016	177
PANGLI ¹⁷	2019	Canada	Retrospective	2012	2016	396
MATER ¹⁸	2020	Saudi	Retrospective	2016	2017	250
		Arabia				
CLELAND ¹⁹	2022	Australia	Retrospective	2014	2020	2036
		NA: DATA NO	OT AVAILABLE			

Table 2 summarizes the data related to screening techniques adopted by each study as regards site of sampling, method of MRSA testing, and timing of sampling.

Table 2. Screening Techniques (N = 16 studies, 7623 patients)

FIRST	SIT OF	METHOD	TIMING (OF SAMPLING
AUTHOR	SAMPLING	OF	Initial	SUBSEQUENT
		TESTING	Sample	SAMPLES
KAISER ⁴	Nose	Culture	Before 72	AFTER 72 HR
			hr	
SCHWEITZER ⁵	Nose	Culture	On	NA
			Admission	
ALRAWI ⁶	Wound	Culture	On	EACH
			Admission	DRESSING
RODRIGUES ⁷	Nose/Axilla/Throat/	Culture	On	WEEKLY
	Perineum/Wound		Admission	
ISSLER-	Nose/Groin/Wound	Culture	Before 48	AFTER 48 HR
FISHER ⁸			hr	
JOHNSON ⁹	Nose/Wound	Culture	On	WEEKLY
			Admission	
$HAITH^{10}$	Nose	PCR	On	WEEKLY
			Admission	
DRUM ¹¹	Nose/Groin/Axilla/	Culture	On	NA
	Wound		Admission	
TEJIRAM ¹²	Nose	Culture	Before 48	NA
			hr	
PARK ¹³	Wound	Culture	Before 48	NA
			hr	
AMISSAH ¹⁴	Nose/Wound	PCR/Culture	Before 24	AFTER 24 HR
			hr	
LIU^{15}	Blood	Culture	NA	NA
JIANG ¹⁶	Blood/Wound	PCR/Culture	NA	NA
PANGLI ¹⁷	Nose/Perineum/	Culture	Before 48	AFTER 48 HR
	Wound		hr	
MATER ¹⁸	Blood/Wound	Culture	NA	NA
CLELAND ¹⁹	Nose/Wound	Culture	Before 48	AFTER 48 HR
			hr	

NA: DATA NOT AVAILABLE

Table 3 summarizes the colonization rate, acquistion rate, and prevalence of MRSA amongst burn individuals in included studies.

Table 3 MRSA Prevalence (N = 7623 patients)

				(
STUDY	SAMPLE	MRSA		M	RSA	MRSA		
	SIZE	COLONIZATION		ACQU	ISITION	PREVALENCE		
		No.	%	No.	%	No.	%	
KAISER ⁴	752	30	4.0	40	5.3	70	9.3	
SCHWEITZER ⁵	31	4	12.9	NA	NA	4	12.9	
ALRAWI ⁶	139	1	0.7	19	13.7	20	14.4	
RODRIGUES ⁷	367	8	2.2	88	24.0	96	26.2	
ISSLER-	357	10	2.8	47	13.2	57	16.0	
FISHER ⁸								
JOHNSON9	554	37	6.7	51	9.2	88	15.9	
HAITH ¹⁰	826	59	7.1	18	2.2	77	9.3	
DRUM ¹¹	725	60	8.3	NA	NA	60	8.3	
TEJIRAM ¹²	601	24	4.0	NA	NA	24	4.0	
PARK ¹³	238	2	0.8	NA	NA	2	0.8	
AMISSAH ¹⁴	62	7	11.3	5	8.1	12	19.4	
LIU ¹⁵	112	NA	NA	NA	NA	52	46.4	
JIANG ¹⁶	177	NA	NA	NA	NA	159	89.8	
PANGLI ¹⁷	396	10	2.5	71	17.9	81	20.5	
MATER ¹⁸	250	NA	NA	NA	NA	33	13.2	
CLELAND ¹⁹	2036	15	0.7	58	2.8	73	3.6	

NA: DATA NOT AVAILABLE

Nine studies reported on the MRSA acquisition among burn patients. A random effect model was used as substantial heterogeneity was determined $(I^2 = 97\%, P=.000)$. The overall acquisition rate of MRSA ranged from 0.022 to 0.240, with the pooled estimate being 0.086 (95% CI: 0.048 - 0.150). (Figure 2)



Figure 2. MRSA Acquisition

In all, four studies examined the effect of MRSA on ICU stay. We analyzed the data using the random-effects model after identifying substantial heterogeneity ($I^2 = 91\%$, P <.001). The combined MD & 95% CIs was 2.79 (-2.21 to 7.8). The combined result suggested that MRSA positive patients are not associated with longer ICU stay (Z = 1.09, P =.27). (Figure 3)

	MRSA Positive		MRSA	MRSA Negative			Mean Difference	Mean Difference	
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% Cl	IV, Random, 95% Cl
Issler-Fisher 2015	5	4	31	2	2	78	27.5%	3.00 [1.52, 4.48]	•
Tejiram 2017	7	7.1	24	12.1	18.6	577	25.3%	-5.10 [-8.32, -1.88]	+
Liu 2018	25	11	52	22	6	60	25.0%	3.00 [-0.35, 6.35]	-
Pangli 2019	12	17	46	0.7	1.5	310	22.2%	11.30 [6.38, 16.22]	-
Total (95% CI)			153			1025	100.0%	2.79 [-2.21, 7.80]	•
Heterogeneity: Tau ² = 23.13; Chi ² = 34.18, df = 3 (P < 0.00001); P = 91%								to te to	
Test for overall effect: Z = 1.09 (P = 0.27)								-50 -25 U 25 50 MRSA Positive MRSA Negative	

Figure 3. Forest plot of ICU stay demonstrates no statistically substantial variance amongst groups

In all, four studies examined the effect of MRSA on mortality rate. No significant heterogeneity was detected. Therefore, a fixed effect model was utilised for analysis ($I^2 = 0\%$, P =.56). The combined OR & 95% CIs was 1.28 (0.69 to 1.40). The combined result suggested that MRSA positive patients are not associated with higher mortality rate (Z = 0.78, P = .44). (Figure 4)

	MRSA Pos	itive	MRSA Negative		Odds Ratio		Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl	M-H, Fixed, 95% Cl
Kaiser 2011	3	70	11	682	11.6%	2.73 [0.74, 10.03]	
Issler-Fisher 2015	0	57	6	300	12.3%	0.39 [0.02, 7.09]	
Tejiram 2017	0	24	6	577	3.1%	1.79 [0.10, 32.77]	
Liu 2018	22	52	23	60	72.9%	1.18 [0.55, 2.52]	
Total (95% CI)		203		1619	100.0%	1.28 [0.69, 2.40]	+
Total events	25		46				
Heterogeneity: Chi [#] = 2.04, df = 3 (P = 0.56); I [#] = 0%							2004 dt 1 40 400
Test for overall effect: Z = 0.78 (P = 0.44)							MRSA Positive MRSA Negative

Figure (4): Forest plot of mortality rate shows no statistically substantial variance amongst collections

Patient isolation was carried out in two studies, including Kaiser et al.⁴ and Johnson et al.⁹ Contact precautions were described in three studies, including Kaiseret al.,⁴ Fisheret al.,⁸ and Johnson et al.⁹ Weekly surveillance was reported in six studies, including Kaiseret al.,⁴ Rodrigueset al.,⁷ Issler-Fisheret al.,⁸ Johnsonet al.,⁹ Haithet al.,¹⁰ and Pangli et al.¹⁷ (Table 4)

Table (4): MRSA Control Policies (N = 16 Studies)

	DECOLONIZATION	PATIENT	CONTACT	SURVEILLANCE					
	PROTOCOL	ISOLATION	PRECAUTIONS	TIMELINE					
KAISER ⁴	NA	Yes	Yes	WEEKLY					
SCHWEITZER ⁵	NA	NA	NA	NA					
ALRAWI ⁶	NA	NA	NA	NA					
RODRIGUES ⁷	No	NA	NA	WEEKLY					
ISSLER- FISHER ⁸	No	NA	Yes	WEEKLY					
JOHNSON9	Nasal mupirocin for 5 days and chlorhexidine baths	Yes	Yes	WEEKLY					
HAITH ¹⁰	Nasal mupirocin for 5 days	NA	NA	WEEKLY					
DRUM ¹¹	NA	NA	NA	NA					
TEJIRAM ¹²	NA	NA	NA	NA					
PARK ¹³	NA	NA	NA	NA					
AMISSAH ¹⁴	NA	NA	NA	NA					
LIU ¹⁵	NA	NA	NA	NA					
JIANG ¹⁶	NA	NA	NA	NA					
PANGLI ¹⁷	NA	NA	NA	WEEKLY					
MATER ¹⁸	NA	NA	NA	NA					
CLELAND ¹⁹	NA	NA	NA	NA					
	NA: DATA NOT AVAILABLE								

4. Discussion

Prolonged MRSA epidemics in multiple hospitals have led to individuals with burns developing complications, including bacteremia , sepsis, and pneumonia. As a consequence, the burn intensive care unit has emerged as a significant reservoir for MRSA in the hospital environment.²⁰

The current meta-analysis involved 16 articles with a total of 7623 individuals; the majority of these studies were retrospective, and only four studies were prospective. Most of these studies were from the USA and China, and no studies from Egypt were found, so more prospective studies need to be performed among ICU burn patients in Egypt.

The majority of studies used nasal and burn wound sampling, all studies used culture, and a small proportion used PCR, and some studies used both.

All studies reported on the overall incidence of MRSA among burn patients. Given the presence of high heterogeneity (I2 = 97%, P =.000), a random effect model was used to analyze the data. The meta-analysis demonstrated that the overall incidence of MRSA ranged from 0.8% to 89.8%, with the pooled estimate being 14.7% (95% CI: 0.092 - 0.226).

The lowest incidence (0.8%) was reported in an Australian study by Park et al.¹³ reflecting the higher hygiene level in burn ICU, while the highest incidence (89.8%) was reported in a Chinese study by Jiang et al.¹⁶ reflecting lower hygiene practices, may be due to the higher burden on medical service higher due to

population density.

Beyond the scope of the most recent metaanalysis, Khan et al. assessed the occurrence of MRSA, which comprised 21 publications and a total of 481 patients with illness. The metaanalysis was conducted on burned individuals who were brought to the intensive care unit. A meta-analysis was conducted, and the findings showed that the probability of MRSA isolates in the burn intensive care unit was 55.0% higher (odds ratio = 0.55, 95% confidence interval = 0.32-0.94). According to the authors, the high prevalence of MRSA isolates was explained by the fact that the majority of the studies that were included were from South Asia.²¹

For the purpose of determining the prevalence of Methicillin-Resistant Staphylococcus aureus that was isolated from burned individuals in Iran, Emaneini et al. conducted yet another metaanalysis. This review covered thirteen different research. Among culture-positive cases, the incidence of MRSA infections in burn patients was found to be 77.9%, with a 95% confidence interval (CI) ranging from 70.2-84. This was determined by the meta-analyses.²²

However, in contrast the present to investigation, another meta-analysis conducted by Kalligeros et al. evaluated the incidence of MRSA colonization upon admission as well as the occurrence of MRSA acquisition inside burn units; the meta-analysis included 16 articles that showed that MRSA incidence on admission was 4.1% (95% CI 2.7%–5.7%). The lower incidence is because most of the included studies were conducted in North America and Europe, reflecting the higher level of medical care.²³

Thirteen studies in this thesis reported on MRSA colonization among burn patients. A random effect model was used as significant heterogeneity was detected (I² = 90%, P =.000). The overall colonization rate of MRSA ranged from 0.7% to 12.9%, with the pooled estimate being 3.8% (95% CI: 0.025 – 0.057).

The lowest MRSA colonization (0.7%) was reported by an Australian study by Cleland et al.¹⁹ and a study from USA by Kaiser et al.⁴ reflecting the higher hygiene and infection control level in burn ICU, while the highest prevalence (89.8%) was reported in USA study Schweitzer et al.,⁵ may be due to the higher severity of the studied patients.

Another meta-analysis conducted by Kalligeros et al. indicated that the total pooled occurrence of MRSA colonization during the first 72 hours after admission (also known as colonization on admission) to the burn unit was 4.1 percent (95% confidence interval: 2.7%–5.7%). This finding is comparable to the most recent research.²³

It was revealed that 9 studies reported on the MRSA acquisition among burn patients. A

random effect model was used as substantial heterogeneity was determined ($I^2 = 97\%$, P =.000). The overall acquisition rate of MRSA ranged from

2.2% to 24%, with the pooled estimate being 8.6% (95% CI: 0.048 – 0.150).

The lowest MRSA colonization (2.2%) was reported in a study from the USA by Haith et al.¹⁰ reflecting the higher hygiene and infection control level in burn ICU, while the highest prevalence (89.8%) was reported in a Brazilian study by Rodrigues et al.⁷ reflecting lower hygiene practices, may be due to the higher overcrowding of patients.

Kalligeros et al. conducted a meta-analysis, which revealed that the acquisition of MRSA in trials that did not include a decontamination process was 21.2% (95% confidence interval: 13.2%-30.5%), with a statistically significant decrease trend over the course of recent years. The MRSA acquisition incidence rate was found to be 4.5 percent in studies that utilized a decontamination approach, with а 95% confidence interval ranging from 0.9% to 10.6%. MRSA acquisition was shown to be greater amongst individuals who had suffered from inhalation injuries (odds ratio: 3.96, 95% confidence interval: 2.51-6.23), flame burns (odds ratio: 1.85, 95% confidence interval: 1.25-2.73), or admission to the intensive care unit (odds ratio: 3.12, 95% confidence interval: 2.18-4.47).²³

MRSA acquisition in the ICU is associated with untrained medical staff, contaminated items and equipment, and unplanned measures when dealing with cases and carriers, so early detection of the acquisition is the most important issue in decreasing and hindering the spread of infection.

Many factors determine outcomes of MRSA, such as length of hospital stay, length of ICU stay, intubation, duration of intubation, number of operations, onset of active infection, and Mortality.

Regarding length of hospital stay, the current meta-analysis found five studies that examined the effect of MRSA on hospital stay. Following the identification of considerable heterogeneity (I2 = 89%, P <.001), we opted to conduct the analysis using the random-effect model. The value of the combined mean 95% confidence interval was 22.6 (13.7 to 31.6). The combined result suggested that MRSA-positive patients are associated with longer hospital stays (Z = 4.97, P <.001).

Emaneini et al. said in their meta-analysis that individuals with burns who are infected with MRSA had a higher risk of morbidity, Mortality, duration of hospital stay, and expenses. This finding is in accordance with what they said.²² These data show a reciprocal association between MRSA hospital stays. Hospital stay is associated with an increased chance of MRSA infection, and MRSA infection is associated with an increased duration of hospital stay. A severe sickness, the use of medical devices, and chronic and continuous exposure to healthcare professionals and/or individuals who may be colonized or infected with MRSA are all factors that may lead to infection in individuals who have been hospitalized for an extended period of time.

Regarding the length of ICU stay, the current meta-analysis found four studies that examined the effect of MRSA on ICU stay. Following the identification of considerable heterogeneity (I2 = 91%, P <.001), we opted to conduct the analysis using the random effect model. A combined mean 95% confidence interval was found to be 2.79 (-2.21 to 7.8). The combined result suggested that MRSA-positive patients are not associated with longer ICU stays (Z = 1.09, P =.27).

In the meta-analysis that Kalligeros et al. conducted, they found that the acquisition of MRSA was greater among individuals who had been admitted to the intensive care unit (OR 3.12, 95% CI: 2.18–4.47). However, they did not investigate the impact that MRSA acquisition had on the duration of stay.²³

However, the duration of stay at ICU, if regular swabs from the wound and nose, hygiene precautions, isolation of new cases, and regular policy of antibiotic use are done, will decrease risks of infections, duration of stay, and costs for treatments and operations.

Regarding Mortality, four studies examined the effect of MRSA on mortality rate. No significant heterogeneity was detected. Therefore, a fixed effect model was used for analysis ($I^2 = 0\%$, P =.56). The total odds ratio 95% confidence intervals came out at 1.28 (0.69 to 1.40). Based on the compilation of findings, it was determined the individuals who tested positive for MRSA did not have a higher death rate (Z = 0.78, P = .44). The integrity of this barrier is compromised as a consequence of burns, which also lead to immunosuppression on both the local & systemic levels. Concurrently, burn wounds may serve as a microenvironment for bacteria due to the presence of necrotic tissue, which has significant levels of protein and does not have any vascularization. The individual who has suffered a burn is at a significant risk of experiencing significant morbidity & death as a result of infection, which is one of the most damaging consequences.²³

This comes in agreement with Emaneini et al., who, in their meta-analysis, stated that Individuals with burns who are infected with MRSA have a higher risk of morbidity and death.22 Also, according to the systematic review by Shuping et al., burn patients infected with MRSA have a significantly higher rate of Mortality due to many factors (as the leading cause of Mortality is deaths caused by burns, ICU is multi-organ failure).²⁴

4. Conclusion

The study found 14.7% MRSA infections in burn patients admitted to burn ICUs, with 3.8% colonization and 8.6% acquisition. Risk factors include age, burn surface area, male gender, inhalational injury, and burn cause.

Disclosure

The authors have no financial interest to declare in relation to the content of this article.

Authorship

All authors have a substantial contribution to the article

Funding

No Funds : Yes

Conflicts of interest

There are no conflicts of interest.

References

- Li X, Luck ME, Hammer AM, Cannon AR, Choudhry MA. 6-Formylindolo (3, 2-b) Carbazole (FICZ)-mediated protection of gut barrier is dependent on T cells in a mouse model of alcohol combined with burn injury. Biochim Biophys Acta Mol Basis Dis. 2020; 1866(11):165901.
- Norbury W, Herndon DN, Tanksley J, Jeschke MG, Finnerty CC. Infection in Burns. Surg Infect (Larchmt). 2016;17(2):250-255.
- 3. van Duin D, Paterson DL. Multidrug-Resistant Bacteria in the Community: An Update. Infect Dis Clin North Am. 2020;34(4):709-722.
- Kaiser ML, Thompson DJ, Malinoski D, Lane C, Cinat ME. Epidemiology and risk factors for hospital-acquired methicillin-resistant Staphylococcus aureus among burn patients. J Burn Care Res. 2011;32(3):429-434.
- Schweizer M, Ward M, Cobb S. The epidemiology of methicillin-resistant Staphylococcus aureus on a burn trauma unit. Infect Control Hosp Epidemiol. 2012;33(11):1118-1125.
- Alrawi M, Crowley TP, Pape SA. Bacterial colonisation of the burn wound: a UK experience. J Wound Care. 2014;23(5):274-277.
- 7. Rodrigues MV, Fortaleza CM, Riboli DF, Rocha RS, Rocha C, da Cunha Mde L. Molecular epidemiology of methicillin-resistant Staphylococcus aureus in a burn unit from Brazil. Burns. 2013;39(6):1242-1249.
- Issler-Fisher AC, McKew G, Fisher OM, Harish V, Gottlieb T, Maitz PK. Risk factors for, and the effect of MRSA colonization on the clinical outcomes of severely burnt patients. Burns. 2015;41(6):1212-1220.

- 9. Johnson AT, Nygaard RM, Cohen EM, Fey RM, Wagner AL. The Impact of a Universal Decolonization Protocol on Hospital-Acquired Methicillin-Resistant Staphylococcus aureus in a Burn Population. J Burn Care Res. 2016;37(6):e525-e530.
- 10.Haith LR Jr, Patton ML, Guilday RE. Antimicrobial Selection in the Face of Nasal Methicillin-Resistant Staphylococcus aureus-Polymerase Chain Recombination Testing in Thermal Injury Patients. J Burn Care Res. 2016;37(2):115-121.
- 11.Drum BE, Collinsworth K, Arnoldo BD, Sreeramoju PV. Hospital-Onset Bloodstream Infection Rates After Discontinuing Active Surveillance Cultures for Methicillin-Resistant Staphylococcus aureus in a Regional Burn Center. Infect Control Hosp Epidemiol. 2017;38(3):371-372.
- 12. Tejiram S, Johnson LS, Mete M. Screening nasal swabs for methicillin resistant Staphylococcus aureus: A regional burn center's experience. Burns. 2017;43(4):771-779. doi:10.1016/j.burns.2017.01.009
- 13.Park KH, Chong YP, Kim SH. Community-associated MRSA strain ST72-SCCmecIV causing bloodstream infections: clinical outcomes and bacterial virulence factors. J Antimicrob Chemother. 2015;70(4):1185-1192.
- 14.Amissah NA, Buultjens AH, Ablordey A. Methicillin Resistant Staphylococcus aureus Transmission in a Ghanaian Burn Unit: The Importance of Active Surveillance in Resource-Limited Settings. Front Microbiol. 2017;8:1906.
- 15.Liu Y, Du FL, Liu PP. Molecular Epidemiology and Virulence Features of Staphylococcus aureus Bloodstream Isolates in a Regional Burn Center in China, 2012-2016. Microb Drug Resist. 2018;24(9):1354-1360.
- 16.Jiang B, Yin S, You B. Antimicrobial resistance and virulence genes profiling of methicillin-resistant Staphylococcus aureus isolates in a burn center: A 5-year study. Microb Pathog. 2018;114:176-179.
- 17.Pangli H, Papp A. The relation between positive screening results and MRSA infections in burn patients. Burns. 2019;45(7):1585-1592.
- 18.Mater ME, Yamani AE, Aljuffri AA, Binladen SA. Epidemiology of burn-related infections in the largest burn unit in Saudi Arabia. Saudi Med J. 2020;41(7):726-732.
- 19.Cleland H, Fernando DT, Gabbe BJ. Trends in Victorian burn injuries 2008-2017. Burns. 2022;48(3):703-712.
- 20.Norbury W, Herndon DN, Tanksley J, Jeschke MG, Finnerty CC. Infection in Burns. Surg Infect (Larchmt). 2016;17(2):250-255.
- 21.Khan TM, Kok YL, Bukhsh A, Lee LH, Chan KG, Goh BH. Incidence of methicillin resistant Staphylococcus aureus (MRSA) in burn intensive care unit: a systematic review. Germs. 2018;8(3):113-125.
- 22.Emaneini M, Beigverdi R, van Leeuwen WB. Prevalence of methicillin-resistant Staphylococcus aureus isolated from burn patients in Iran: A systematic review and metaanalysis. J Glob Antimicrob Resist. 2018;12:202-206.
- 23.Kalligeros M, Shehadeh F, Karageorgos SA, Zacharioudakis IM, Mylonakis E. MRSA colonization and acquisition in the burn unit: A systematic review and meta-analysis. Burns. 2019;45(7):1528-1536.
- 24.Shuping LL, Kuonza L, Musekiwa A, Iyaloo S, Perovic O. Hospital-associated methicillin-resistant Staphylococcus aureus: A cross-sectional analysis of risk factors in South African tertiary public hospitals. PLoS One. 2017;12(11):e0188216.