

Asian Journal of Biotechnology and Bioresource Technology

Volume 9, Issue 3, Page 31-40, 2023; Article no.AJB2T.107926 ISSN: 2457-0125

Isolation and Identification of Air Microflora in Clifford University Medical Center, Ihie Campus, Owerrinta, Abia State, Nigeria

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/AJB2T/2023/v9i3186

Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <u>https://www.sdiarticle5.com/review-history/107926</u>

Original Research Article

Received: 22/08/2023 Accepted: 30/10/2023 Published: 07/11/2023

ABSTRACT

The settling plate method was used to examine the air microflora in 6 units at the Clifford University Medical Centre, Ihie campus, Owerrinta, Abia state, Nigeria. The male ward, male restroom, female ward, female restroom, theater room, and environment were among the 6 units. At each of the sample locations, the culture plates containing potato dextrose agar and nutrient agar media were exposed to the atmosphere for about 15 minutes. A total count of 11 CFU and 295 CFU fungal and bacterial isolates respectively were identified from the medical center but 5 fungal isolates and 29 bacterial isolates were obtained as pure cultures. These fungi were *Aspergillus flavus, Aspergillus niger, Trichophyton* sp, *Penicillium* sp, and white mold. The bacterial genera were *Staphylococcus*

Asian J. Biotechnol. Bioresour. Technol., vol. 9, no. 3, pp. 31-40, 2023

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sp, *Streptococcus* sp, *Micrococcus* sp, and *Bacillus* sp. *Staphylococcus* sp and *Aspergillus* sp. were the two most prevalent microorganisms in this investigation. It can be said that the Clifford University Medical Centre employed for this study featured numerous types of Gram-positive bacteria, a high prevalence of pathogenic bacteria, and a high prevalence of pathogenic fungi in both the indoor and outdoor air. The majority of the fungal isolates are molds, whereas certain bacterial species are commensals that live on human skin, the fungal isolates are of environmental origin. The bacterial and fungal isolates contained a few non-pathogenic species. The study was carried out to determine if air microfloral affects the infection rate in the University Medical Centre.

Keywords: Air Microflora; university medical centre; fungal and bacterial isolates; pathogenic; nonpathogenic organisms.

1. INTRODUCTION

The study of living microorganisms (bioaerosols) and toxins that are suspended in the atmosphere is known as Aeromicrobiology [1]. Although the quantity of microorganisms in the air is far lower than in the soil and water, they are nevertheless numerous enough to have an impact on the atmosphere. These aerosols have ecological significance because they are linked to illnesses in people, animals, and plants. [2,3]. The majority of the particles that end up suspended in the air come from terrestrial and aquatic environments, and air turbulence. In addition to being largely transported by wind, bioaerosols can also be deposited by a number of different mechanisms, such as gravity, contact with surfaces, or mixing with rain to bring the dust back down to the earth's surface [4,5]. Bioaerosols can spread infections. endotoxins. and allergens that sensitive people are allergic to. The types of bacteria or toxins, the types of particles they are connected with, such as dust or mist, and the gases in which the bioaerosols are suspended, are just a few of the variables that affect the content and size of bioaerosols. Fungi, bacteria, viruses, and pollen are examples of bioaerosols. Bioaerosol survival rates are influenced by a variety of biotic and abiotic parameters, such as weather patterns, ultraviolet (UV) light, temperature, and humidity, as well as the availability of resources in clouds or dust [6]. Bacteria make up the majority of bioaerosols found in marine areas, whilst those found in terrestrial ecosystems are abundant in bacteria, fungi, and pollen [7].

Healthcare facilities present an extra, particular issue in addition to the infectious threats faced by all office and commercial buildings, a high density of potentially infectious and immunecompromised individuals. Ventilation is necessary in hospitals and other healthcare facilities because they are complicated

environments that must manage hazardous emissions while providing comfort for patients [8]. Furthermore, patients may spread dangerous microbes to staff, hospital visitors, and other patients, making the biological quality of the air in hospital environments a special issue [9]. Despite the fact that hospitalization and medical operations are intended to treat illnesses, they occasionally unintentionally can introduce dangerous bacteria into the body and start a Nosocomial Infection (NI), the infected patient is the main source of airborne germs inside the hospital [10]. Airborne transmission occurs when pathogenic microorganisms are tran smitted from an infected person to a vulnerable person through the air. The primary process that makes infections airborne, is the production of aerosol droplets from coughing or sneezing, along with the subsequent loss of water that permits them to float in the air for long periods of time and over great distances [11]. Biological aerosols include spores of veast, mold, bacteria, other microorganisms, viruses. and skin lesions may potentially be a source of airborne particles in specific clinical situations [10]. Some microorganisms, especially Gram-positive ones like Staphylococcus aureus and Streptococcus pneumoniae, can persist for several months on dust particles. Viruses and fungus spores can both endure for longer lengths of time. Since many modern buildings are enclosed feature self-contained circulating and air systems for temperature management, the prevalence of airborne infections has increased [12].

Bioaerosol properties are complex and diverse, and have a direct impact on the environment, climate, and human health, to accurately obtain the atmospheric chemical characteristics of bioaerosols, their effective identification in the atmosphere is very significant [13]. Airborne bacteria and fungi may cause several adverse effects, especially infectious, allergenic, and immunotoxic disorders [13]. Microbiological air quality is an important criterion that must be into consideration when taken indoor workplaces are being designed to provide a safe environment. [14] More people are dying every year from hospital infections. Although, many pathogens can cause hospital infection those that are able to survive in the hospital environment for long periods and resist also disinfections are particularly important in this respect [15]. This study was designed to identify both pathogenic and non-pathogenic air microflora and determine if they affect the infection rate in the hospital environment, bacterial and fungal aerosols were collected in this study using the settling plate technique and isolated in the microbiology laboratory.

2. MATERIALS AND METHODS

2.1 Study Site

Clifford University Medical Centre was used as a case study for this research project. Air samples were collected at the following units of the hospital: Male ward, Female ward, Male toilet, Female toilet, Theatre room, and External environment and investigated by settling plate technique.

2.2 Sample Collection and Isolation of Organisms

2.2.1 Collection of sample

The 12 culture plates containing the Nutrient agar media and the 12 culture plates containing Potato dextrose agar were exposed to the Medical Centres' atmosphere for a period of 15 minutes where 2 culture plates were exposed at each of the sampling units. After which, the culture plates were carefully covered back and wrapped in Santana nylons and placed in a cool, dry, and clean carton, and transported to the Microbiology laboratory at 25 °C for isolation of organisms.

2.2.2 Isolation of organisms

After 24 hours of incubation at 37 °C, 29 bacterial pure cultures were isolated in the Microbiology laboratory while after 3-5 days of incubation at 25°C, 5 fungal pure cultures were isolated [16,17].

2.3 Identification of Organisms

Various biochemical tests were carried out in this study after the isolation for the identification of the bacteria isolated from the Medical Centre while fungal staining and microscopy were carried out for the identification of fungi isolated from the Medical Centre. The biochemical tests are catalase, coagulase, citrate utilization, oxidase, hydrogen sulphide production, sugar fermentation, and indole tests [16,17].

3. RESULTS

The bacterial counts and fungal counts of each unit of the medical centre ranged from 19 CFU to 115 CFU and 1 CFU to 4 CFU respectively. The microbial counts (CFU) are presented in Table 1, the biochemical characteristics and identification of bacterial isolates are presented in Table 2 while Table 3 is the cultural characteristics of fungal isolates.

The bacterial isolates identified in the various units of Clifford University Medical Centre were *Micrococcus* sp., *Staphylococcus* sp., *Streptococcus* sp., and *Bacillus* sp. The occurrence of the bacterial isolates is presented in Table 2.

Table 1. Counts of Indoor and Outdoor Air Microflora of	of Clifford University Medical Centre.
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Sample Source	Total Bacterial Counts Per Unit (CFU)	Total Fungal Counts Per Unit (CFU)
Male Toilets	25	2
Male Wards	115	2
Female Toilets	37	1
Female Wards	19	4
External Environment	28	1
Theatre room	71	1
Total Microbial Counts	295	11

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Sample source	Cultural characteristics	Shape	Gram reaction	Catalase	Coagulase	Oxidase	Indole	Citrate	S	В	G	H₂S	Possible Bacterial Species (sp)
MT1A	Orange, flat, punctiform, smooth.	Cocci	Positive	+	-	-	-	-	+	+	-	-	Staphylococcus sp.
MT1B	Milky, flat, punctiform, smooth.	Cocci	Positive	+	-	+	-	+	+	+	-	-	Staphylococcus sp.
MT2	Milky, flat, smooth, punctiform.	Cocci	Positive	+	-	+	-	+	+	+	-	-	Staphylococcus sp.
MW1A	Milky, flat, smooth, punctiform.	Cocci	Positive	+	-	+	-	+	-	+	-	-	<i>Micrococcus</i> sp.
MW1B	Milky, round, smooth, flat.	Cocci	Positive	+	-	+	-	+	+	+	-	-	Staphylococcus sp.
MW2A	Milky, round, flat, smooth.	Short rod	Positive	+	-	+	-	+	-	+	+	+	<i>Bacillus</i> sp.
MW2B	Creamy, flat, round, smooth.	Long rod	Positive	+	-	+	-	+	-	+	-	-	<i>Bacillus</i> sp.
FT1A	Milky, flat, round, smooth.	Cocci	Positive	+	-	+	-	+	-	-	+	-	<i>Micrococcus</i> sp.
FT1B	Milky, flat, punctiform, smooth.	Cocci	Positive	+	-	+	-	-	+	+	-	-	<i>Micrococcus</i> sp.
FT2A	Milky, flat, punctiform, smooth.	Cocci	Positive	+	-	+	-	+	+	+	+	-	Staphylococcus sp.
FT2B	Milky, opaque, round, smooth.	Cocci in chain	Positive	+	-	+	-	-	-	-	-	-	Micrococcus sp.
FW1A	Milky, rhizoid, hilly, rough.	Short rod	Positive	+	-	+	-	-	+	+	+	-	<i>Bacillus</i> sp.
FW1B	Milky, opaque, round, smooth.	Cocci in chain	Positive	+	-	+	-	-	+	+	-	-	Staphylococcus sp.
FW1C	Milky, smooth, punctiform.	Short rod	Positive	+	-	-	-	+	-	-	-	-	<i>Bacillus</i> sp.
FW2A	Milky, smooth, dark- centered, round.	Cocci	Positive	+	-	+	-	+	+	+	-	-	Staphylococcus sp.

Table 2. Biochemical characteristics and identification of bacterial isolates from different units of Clifford University Medical Centre.

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Sample source	Cultural characteristics	Shape	Gram reaction	Catalase	Coagulase	Oxidase	Indole	Citrate	S	В	G	H₂S	Possible Bacterial Species (sp)
FW2B	Milky, smooth, round.	Cocci	Positive	+	-	+	-	+	+	+	-	-	Staphylococcus sp.
FW2C	Milky, smooth, punctiform.	Cocci	Positive	+	-	+	-	+	+	+	+	-	Staphylococcus sp.
EE1A	White, rough, dark- centered, round.	Cocci	Positive	+	-	-	-	+	+	+	+	-	Staphylococcus sp.
EE1B	Creamy, rough, rough, rough, rhizoid.	Cocci	Positive	+	-	-	-	+	+	+	-	-	Staphylococcus sp.
EE1C	Orange, round, hilly, smooth.	Cocci	Positive	+	-	-	-	-	+	+	+	-	Staphylococcus sp.
EE1D	Milky, round, flat, smooth.	Cocci	Positive	+	-	-	-	-	+	+	-	-	Staphylococcus sp.
EE2A	White, rough, dark- centered, filamentous, rhizoid.	Cocci in chain	Positive	+	-	-	-	-	+	+	-	-	Staphylococcus sp.
EE2B	Milky, round, smooth, flat.	Cocci in chain	Positive	-	-	-	-	-	-	+	-	-	Streptococcus sp.
EE2C	Milky, smooth, punctiform, flat.	Long rod	Positive	+	+	-	-	-	+	+	+	-	<i>Bacillu</i> s sp.
T1	Milky, rhizoid, rough.	Cocci	Positive	-	-	+	-	-	-	+	-	-	Streptococcus sp.
T2	Orange, round, dark-centered, moist, flat	Cocci	Positive	+	-	-	-	+	+	+	-	-	Staphylococcus sp.
Т3	White, rough, dry, umbolate.	Short rod	Positive	-	-	+	-	-	+	-	-	-	<i>Bacillu</i> s sp <i>.</i>
Т4	Milky, flat, smooth, punctiform.	Cocci	Positive	+	-	+	-	-	+	+	-	-	Staphylococcus sp.
Т5	Milky, round, smooth, moist.	Short rod	Positive	+	-	+	-	-	-	+	+	-	<i>Bacillu</i> s sp.

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KEY: MT- Male Toilet, MW- Male Ward, FT- Female Toilet, FW- Female Ward, EE- External Environment, T- Theatre, SFT- Sugar Fermentation Test, S- Slant, G- Gas, B- Butt, H₂S- Hydrogen sulphide, + (positive), - (negative)

Sample Source	Organism	Cultural Characteristics
Theatre	White mold	White and filamentous
Environment	Trichophyton sp.	Orange and non-filamentous
Male Ward	Aspergillus sp. Penicillium sp.	Non-filamentous, velvet, circular and leave-green with white circumference. Non-filamentous, velvet, circular, greenish-blue with white circumference.
Male Toilet	Aspergillus sp. Trichophyton sp.	Non-filamentous, velvet, circular, dark-brown with white edges. Orange and non-filamentous
Female Toilet	White mold	White and filamentous
Female Ward	Aspergillus sp. Aspergillus sp.	Non-filamentous, velvet, circular, dark-brown with white edges. Non-filamentous, velvet, circular and leave-green with
	<i>Penicillium</i> sp.	white circumference. Non-filamentous, velvet, circular, greenish-blue with white circumference.
	White mold	White and filamentous

Table 3. Cultural Characteristics of Fungal Isolates from Different Units of Clifford University Medical Centre

4. DISCUSSION

The bacterial isolates from the different units of Clifford University Medical Centre: male ward, male toilet, female ward, female toilet, theatre, and the environment, were subcultured for 24 hours at 37 °C. Afterward, 29 pure cultures were isolated and grouped based on their different sources of the sample.

The results of the current study showed that the air at Clifford University Medical Centre included a great diversity of bacteria. Micrococcus. Staphylococcus, Streptococcus, and Bacillus genera were the bacterial isolated. predominated Staphylococcus in the environment. In a related study, Staphylococcus, Proteus, Streptococcus, and Micrococcus were isolated from indoor air of laboratories, while Micrococcus, Paracoccus, Staphylococcus, and Enhydrobacter were isolated in the study of indoor and outdoor environments of child care facilities [18,19]. Also, Sanaa & Amani [15] isolated S. aureus, E. coli, Klebsiella sp, P. aeruginosa, and Bacillus sp from hospital delivery and nursing rooms in Khartoum district Sudan. All of the bacteria found in this study were Gram-positive rods and cocci, just like the ones found in hospital lobbies' air [20]. Since Gram-positive bacteria have a thicker covering of peptidoglycan in their cell walls than Gram-negative bacteria, this is not surprising. As a result, they dominated the study's bacterial flora.

The organisms recovered in this study can be categorized as either commensals or pathogens depending on how they interact with their human hosts. Either humans or the soil environment is the source of these bacteria. The commensals are part of the typical human body flora and are present on the skin, nose, and mouth. They do not infect their host normally, with the exception of when the immune system is weak [21]. *Bacillus, Paenibacillus, and Chaetomium* were among the bacterial genera reported by Yan et al. [3]. The *Staphylococcus* genus, which was discovered in this study, is found on human and other animals' skin and mucous membranes.

They are a part of the microbial flora of soil and are present all over the planet. Staphylococcus aureus, Staphylococcus epidermidis, and Staphylococcus saprophyticus are the three most significant species of human pathogens [21], with S. aureus being the most pathogenic of all and responsible for serious infections in healthcare facilities like Pneumonia; which most frequently affects people with underlying lung disease, Osteomyelitis; a bone infection that can be brought on by staphylococcal bacteria entering the bloodstream or being directly introduced, as in the case of intravenous (IV) drug abuse, vaginal infections, etc. endocarditis, and many other conditions [22]. Staphylococcus aureus was one of the dominant isolated organisms, this is a common pathogenic bacterium that is associated with various diseases such as respiratory tract, digestive system, and postoperative infections, also urinary tract, and skin disorders, it is resistant to antibiotics, its presence might be due to post-sterilization, or the environment contamination [15]. Anyone can become infected with Staphylococcus, but some people are more susceptible than others, such as those who have diabetes, cancer, etc. The risk is also higher in hospitals where there are patients with compromised immune systems, those who have had specific surgeries, those in intensive care units (ICUs), and those who have had medical devices implanted in their bodies [22]. Although the organism can be found in many places, including on the surface of the skin, in dust, water, and soil, in this investigation, Micrococcus sp. was not particularly prevalent in the medical center. In this investigation, the genus Bacillus was also isolated. These saprophytic organisms are common in the air and several other ecosystems. Medically significant pathogenic species include Bacillus cereus. traditionally considered foodborne pathogens that establish occasional opportunistic infections, and have naturally evolved to cause fatal anthrax-like disease. This serves as a reminder that the field of medical microbiology is constantly changing, posing new challenges that require ongoing vigilance and research [23] and Bacillus anthracis, which produces anthrax. Bacilli endospores are more resistant to heat, drying, and disinfectants than vegetative cells are. They may survive better in the air and can be found in dirt, dust, and plant products [24]. In this study, the gram-positive, and catalasenegative streptococci were the final bacterial genera discovered. These organisms belong to the natural flora but turn pathogenic when they are exposed to an immune system that is weak. Streptococcus pyogenes (group А Streptococcus) is one of the most important bacterial causes of skin and soft tissue infections (SSTIs) worldwide. There is no other pathogen that causes as many diverse clinical entities as S. pyogenes. [25]. Specifically, this organism causes infections in the superficial keratin layer (impetigo), the superficial epidermis (erysipelas), the subcutaneous tissue (cellulitis), the fascia (necrotizing fasciitis), or muscle (myositis and myonecrosis) and it is also the etiologic agent of scarlet fever and Streptococcal Toxic Shock Syndrome (StrepTSS). Impetigo is a non-lifethreatening infection, but can result in poststreptococcal acute glomerulonephritis (AGN). Cellulitis and erysipelas can be mild or moderately severe, while necrotizing fasciitis, myonecrosis, and StrepTSS are life-threatening [25].

The fungal isolates from the different units of Clifford University Medical Centre: male ward, male toilet, female ward, female toilet, theatre, and the environment, were subcultured for 3-5 days at 25°C. Afterward, 5 pure cultures were isolated and grouped based on their different sources of the sample.

This study found that the air at Clifford University Medical Centre contained a great diversity of funai. White Mold, Trichophyton terrestre, Aspergillus flavus, Aspergillus niger, and Penicillium were among the fungi that were isolated; Aspergillus species predominated in the medical center, with the female ward having the highest fungal count (4 CFU). In a related study, Saccharomyces cerevisiae, Aspergillus sp., Rhizopus stolonifera, and Alternaria sp. were isolated in the study of Indoor Air Microflora of some Daycare Centres in Illorin south local government area [26]. Cladosporium, Asperaillus. Penicillium. Paecilomvces. Staphylococcus, and Micrococcus spp were the most prevalent genera isolated from an academic pediatric hospital [27]. Due to their sporulate and reputation ability to for pathogenicity, the Aspergillus spp. isolated in this investigation dominated the genera of fungus isolated. These are Aspergillus niger, which can cause aspergillosis and generates oxalic acid, malformin, and other chemicals, and Aspergillus flavus, which is known to create mycotoxins. Aspergillus flavus, one of the two isolated species, is known to produce aflatoxin [28]. In several offices at the University of Ilorin in Ilorin, Nigeria, previous research also noted the presence of Aspergillus niger, Aspergillus flavus, Aspergillus glaucus, Aspergillus versicolor. Alternaria alternata, Geotrichum candidum, and Rhizopus [29].

Penicillium spp. are among the most prevalent fungi in the environment and are typically thought to not be harmful to people. This result is in agreement with Jalili [13] who reported the isolation of species of yeast, Aspergillus, and Penicillium in Shahrekord Hospitals in Iran, these organisms are highly pathogenic or opportunistic pathogens/medically important microorganisms. Most fungi are known to be associated with asthma in both children and adults [13] These fungi lead to pulmonary aspergillosis when inhaled [30]. Cases of contamination occur mostly in people with underlying illnesses and low immunity levels [31]. Approximately 49.1% of Aspergillus sp. outbreaks within hospitals can be attributed to the construction work in or around hospitals [32]. In a study conducted in Benin City. Nigeria, Penicillium, Mucor, Aspergillus, and Fusarium were among the fungi species isolated from the hospital indoor air [33]. Airborne fungi and their spores have the potential to be blown into buildings with natural ventilation. They can pose a threat to the life of immunocompromised patients when are blown in through the windows of the relevant wards [34]. A study conducted in hospitals in Sari, Iran isolated predominantly pathogens which are Mucorales sp., Candida sp., Fusarium sp., and Aspergillus sp. [35]. Francisco et al. [36] reported the isolation of Trichosporon asahii and T. inkin from urine and blood samples collected from 24 Medical Centres.

5. CONCLUSION AND RECOMMENDA-TION

5.1 Conclusion

The results of this study showed that the Clifford University Medical Center's indoor and outdoor air harbour a variety of genera of Gram-positive cocci and rod bacteria, Staphylococcus sp predominated in the hospital environment and the male ward having the highest bacterial count (115 CFU). Of the fungi isolated, Aspergillus sp has the greatest fungal count (4 CFU). A few pathogenic species were isolated among bacteria and fungi such as Staphylococcus, Aspergillus, Bacillus. Streptococcus, and Trichophyton spp respectively. Therefore, the air microflora in this medical centre may have possibly contributed to the infection rate in the Medical Centre.

5.2 Recommendation

By refraining from any actions that could cause dust to be produced, such as sweeping, the amount of bioburden in the medical facility can be reduced. It is preferable to mop the floor rather than sweep it. The vaccination of children against respiratory illnesses like diphtheria, whooping cough, etc. is advised to parents. In order to stop the spread of infectious droplets to other patients, people with respiratory illnesses like rhinitis, frequent sneezing, etc., as well as other transmissible diseases, should be assigned to different wards. The hospital wards should be fumigated with chemical disinfectant monthly in order to decontaminate the room. To keep the restrooms sanitary and germ-free, various disinfection techniques should be offered. Water should always be available at the hospital so that toilets may be flushed after each usage.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- 1. Pepper IL, Gerba CP. Aeromicrobiology environmental microbiology. 2015;89–110 DOI: 10.1016/B978-0-12-394626-3.00005-3
- [Aydogdu H, Asan A, Tatman OM. Indoor and outdoor airborne bacteria in child daycare centres in Edirne City (Turkey). Seasonal Distribution and Influence of Meteorological Factors. Environmental Monitoring and Assessment. 2010;164:53– 66.
- Yan D, Zhang T, Bai JL, Su J, Zhao LL, Wang H, Fang XM, Zhang YQ, Liu HY, and Yu LY. Isolation, Characterization, and antimicrobial activity of bacterial and fungal representatives associated with particulate matter during Haze and Non-haze Days. Frontiers Microbiol. 2022; 12:793037. DOI:10.3389/fmicb.2021.793037
- 4. Pavan R, Manjunath K. Qualitative analysis of indoor and outdoor air. Airborne fungi in Cowshed. Journal of Mycology. 2014;2:1–8.
- 5. Jain A, Jain R, Jain S. Isolation of microorganisms from air. In the book: Basic techniques in biochemistry, microbiology, and molecular biology. 2020; 119–120.

DOI:10.1007/978-1-4939-9861-1 33.

- Acosta-Martinez V, Van Pelt S, Moore-Kucera J, Baddock M.C. and Zobeck TM. Microbiology of Wind–Eroded Sediments: Current Knowledge and Future Research Directions. Aeolian Research. 2015;18:99– 113.
- Moreno D, Alcami A. Monitoring of airborne biological particles in outdoor atmosphere. International Microbiology. 2016;19(1):1–13.
- 8. Chuaybamroong Ρ, Choomseer Ρ. Ρ. Sribenialux Comparison between hospital single Air unit and central Air unit for Ventilation Performances and Airborne Microbes. Aerosol and Air Quality Research. 2008;8(1):28-36.
- 9. Obbard JP, Fang LS. Airborne concentration of bacteria in a hospital

environment in Singapore. Water, Air, and Soil pollution. 2013;144:333–341.

- Hambraeus A. Aerobiology in the operating room-a review. J Hosp Infect. 1988 Feb;11 Suppl A:68-76. DOI:10.1016/0195-6701(88)90169-7. PMID: 2896749.
- Emmerson AM. The impact of surveys on hospital infection. J Hosp Infect. 1995 Jun;30 Suppl:421-40. DOI:10.1016/0195-6701(95)90047-0. PMID: 7560981.
- 12. Augustowska M, Dutkiewicz. Variability of airborne microflora in a hospital ward within a period of one year. Annals of Agricultural and Environmental Medicine. 2006;13(1):99–106.
- Jalili D, Dehghani MH, Fadaei A, Alimohammadi M. Assessment of airborne bacterial and fungal communities in Shahrekord Hospitals. Journal of Environmental and Public Health. 2021; 7 pages. Available:https://doi.org/10.1155/2021/886
- 4051
 14. Li X, Li L, Zhuo Z, Zhang G, Du X, Li X, Huang Z, Zhou Z, Cheng Z. Bioaerosol identification by wide particle size range single particle mass spectrometry. Atmosphere. 2023;14(6):1017 Available:https://doi.org/10.3390/atmos140 61017
- 15. Sanaa O, Yagoub, Amani El Agbash. Isolation of potential pathogenic bacteria from the air of hospital-delivery and nursing rooms. Journal of Applied Sciences. 2010;10:1011-1014.
- 16. Cheesbrough M. district laboratory practice in tropical countries. Cambridge University Press. 2005;62–70.
- Clinical, and Laboratory Standard Institute (CLSI). Performance Standard for Antimicrobial Susceptibility Testing. CLSI approved standard M100S. Clinical and Laboratory Standard Institute, Wayne, PA.
- Shin SK, Kim J, Ha SM, Oh HS, Chun J, Sohn J, Yi H. Metagenomic insights into the bioaerosols in the Indoor and Outdoor Environments of Childcare Facilities. PLoS ONE. Journal.Plos.Org. 2015;10(5) 27–37.
- 19. Shiaka GP, Yakubu SE. Comparative analysis of airborne microbial concentrations in the indoor environment of two selected clinical laboratories. IOSR Journal of Pharmacy and Biological Sciences. 2013;8(4):13–19.

- 20. Dong UP, Jeong-Kwan Y, Won JK, Kyeong- Min L. Assessment of the levels of airborne gram-positive Bacteria and Fungi in Hospital Lobbies. International Journal of Environmental Resources and Public Health. 2013;10:541–555.
- 21. Arora DR, Arora BB. Textbook of microbiology, 4th ed., CBS publishers and distributions, PVT Ltd., New Delhi, India. 2012;674–677.
- 22. Center for disease control (CDC): Staphylococcus aureus in healthcare settings. Centers for disease control and prevention, national center for emerging and zoonotic infectious diseases (NCEZID), division of healthcare quality promotion (DHQP) Last reviewed: January 17, 2011. Retrieved Online on 24/08/2023.
- Baldwin VM. You Can't B. cereus A Review of Bacillus cereus Strains That Cause Anthrax-Like Disease. Front. Microbiol. 2020;11. Available:https://doi.org/10.3389/fmicb.202 0.01731
- Seino K, Takano T, Nakamura K, Watanabe M. An evidential example of airborne bacteria in a Crowded, Underground Public Course in Tokyo. Atmospheric Environment. 2005;39:337– 341.
- Stevens DL, Bryant AE. Impetigo, erysipelas and cellulitis in: Streptococcus pyogenes: Basic Biology to clinical manifestations [Internet]. By Ferretti J. J., and Stevens D. L, and Fischetti V. A, editors. Oklahoma City (OK): University of Oklahoma Health Sciences Center; 2016.
- Sule IO, Agbabiaka TO, Saliu BK, Odebisi-Omokanye, Zakariyah RF, Ali AA. Analysis of indoor air microflora of some daycare centres in Ilorin South Local Government Area, Nigeria. Al-Hikman Journal of Pure and Applied Sciences. 2017;4:31–35.
- 27. Mirhoseini SH, Didehdar M, Akbari M, Moradzadeh R, Jamshidi R, Torabi S. Indoor exposure to airborne bacteria and fungi in sensitive wards of an academic pediatric hospital. Aerobiologia. 2020 Jun; 36:225-32.
- 28. Khan AAH, Karuppayil SM. Fungal pollution of indoor environments and its management. Saudi Journal of Biological Sciences. 2012;19(4):405–426.
- 29. Adetitun DO, Oladele IL. Airborne microbial load and diversity in some offices in University of Ilorin. Nigerian Journal of

Pure and Applied Science. 2016;29:2715–2723.

- 30. Jaakkola JJK, Hwang BF, Jaakkola MS. Home dampness and molds as determinants of allergic rhinitis in childhood: A 6-year, population-based cohort study," American Journal of Epidemiology. 2010;172(4):451-459.
- Shiaka G, Yakubu S. Comparative analysis of airborne microbial concentrations in the indoor environment of two selected clinical laboratories," IOSR Journal of Pharmacy and Biological Sciences (IOSR-JPBS). 2013;8:13–19.
- Vonberg RP, Gastmeier P. Nosocomial aspergillosis in outbreak settings. Journal of Hospital Infection. 2006;63:246– 254.
- Ekhaise FO, Ogboghodo B. Microbiological indoor and outdoor air quality of two major hospitals in Benin City, Nigeria. Sierra Leone Journal of Biomedical Research. 2011;3:169–174.

- Tang JW, Nicolle A, Pantelic J, et al. Different types of door-opening motions as contributing factors to containment failures in hospital isolation rooms. PLoS One. 2013;8: Article ID e66663.
- Moazeni Nabili 35. Μ, Asgari S. Μ. Nosocomial fungal infections: epidemiology, diagnosis, treatment and Journal Mazandaran prevention. of University of Medical Sciences. 2018;28: 182 - 212.
- 36. Francisco EC, de Almeida Junior JN, de Queiroz Telles F, Aquino VR, Mendes AV, de Andrade Barberino MG, Castro PD, Guimarães T, Hahn RC, Padovan AC, Chaves GM. Species distribution and antifungal susceptibility of 358 Trichosporon clinical isolates collected in 24 medical centres. Clinical Microbiology and Infection. 2019 Jul:25(7):909.e1-909.e5. DOI:10.1016/i.cmi.2019.03.026. Epub 2019 Apr 13. PMID: 30991116.

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