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Public Health Risks Assessment of Bioaerosols from Different Locations in Bariga Metropolis of Shomolu Local Government Area, Lagos State, Nigeria

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Treatment Consortium

Ogah JO^{1*}; Mba OO²; Okelola CA³ and Kolawole RM³

¹Department of Microbiology, Faculty of Science, University of Lagos, Akoka, Nigeria

²Department of Public Health, Faculty of Health Sciences, National Open University of Nigeria, Lagos

³Department of Cell Biology and Genetics, University of Lagos, Akoka, Nigeria

ORCID ID: 0009-0008-1462-2456

ABSTRACT

The increasing human populations and their daily activities have contributed to bioaerosols generation in Bariga, leading to air pollution with consequent public health threats to exposed individuals. This study aimed to assess public health risks associated with bioaerosols generated from outdoor activities in Bariga community, Shomolu Local Government Area of Lagos State. An analytical observational study design based on the measurement of meteorological parameters in the field and laboratory analysis of bioaerosol particles collected from the study population using a handheld Kestrel 3000 weather meter and standard microbiological procedures, respectively, was adopted. Four hundred (400) samples of bioaerosol particles obtained by the Taro Yamani formula at a 5% level of precision were analyzed. Passive air monitoring using Koch's sedimentation method was employed for total viable counts, while microbial isolates were identified using cultural, morphological, and biochemical characterization. The variation in environmental parameters obtained was largely dependent on time and seasons, with a consequent adverse effect on the presence and movement of bioaerosol particles in the atmosphere. Microbial populations varied in densely and less-densely populated study areas. The study identified Bariga as a high-risk area with CFU above 10² -10⁴ CFU/m³, thus capable of causing bioaerosols-related diseases. A total of twelve (12) bacterial and six (6) fungal genera were isolated, with Gram-positive bacteria having a prevalence rate of 87.02% and Gram-negative bacteria at 12.99%. The predominant fungi, on the other hand, were 87.94% mold and 12.06% yeast. One-way analysis of variance (ANOVA) at 0.05 significance level showed a significant correlation (F-ratio > F-critical value) (p=0.05) between exposure to bioaerosol particles and associated health risks in the exposed individuals. We recommend that Lagos State Government should intensify efforts to reduce the public health effect of bioaerosols through policies. structural planning, development, and education on environmentally friendly activities and personal hygiene.

Keywords: Bioaerosols; Meteorological Parameters; Microorganisms; Public Health; Bariga; COVID-19.

INTRODUCTION

The ubiquitous and diverse nature of microorganisms in an ecosystem could be beneficial and detrimental to the existence of plants and animals within the biosphere or community. The increasing human populations with their daily activities have greatly contributed to air pollution consisting of inorganic (particulate matter) and organic (bioaerosols). The generated bioaerosols could serve as a means of transmitting pathogens and thus pose a potential public health threat to the community. These pollutants have constituted an inevitable environmental issue in the world's developing countries, and it stands out among the array of global environmental hazards facing metropolitan cities such as Lagos [1]. Bioaerosols generated are suspensions of atmospheric particles of biological origin containing living and/or dead microorganisms, pollens, and their derivatives [2]. The concentration of different bioaerosols in the air varies from one location to another. The bioaerosols can be generated during sneezing, talking, laughing, and yawning [3]; pollens from plants or anthropogenic activities can influence the bioaerosol's particle size. composition, and concentration in the atmosphere [4]. Microbial spores suspended in the air could serve as a potential source of allergy and general health problems in different ways [5]. They can easily be transferred from one environment to another because of their small size (0.001-100 µm) and lightweight, which are dependent upon prevailing physicochemical properties of the atmosphere, such as temperature, humidity, solar radiation, wind, precipitation and atmospheric pressure, for transport and survival [6-9] as reported [5]. After the release of biological aerosols into the atmosphere, most of them combine with the ambient particulate matter for diffusion and transportation from one place to another over long distances [8]. The susceptibility to acquiring infections from inhaling bioaerosols is largely determined by the virulence of the biological agent, the pathogenicity of the agent, dosage and the host's immune response. The airs we breathe also contain a mixture of gaseous and particulate matter released from natural and anthropogenic sources [9].

Humans play a significant role as hosts to many microorganisms, some of which are normal flora of the body, while others are pathogens capable of causing infectious diseases in man. In developing countries like Nigeria, air pollution is mainly due to overpopulation and uncontrolled urbanization, coupled with rapid industrialization, disparity among residents and poor environmental

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sustainability management education [10]. The population density of bioaerosols suspended in the atmosphere in a particular environment is influenced by the human population and their interactions with such environment; for instance, human populations have been implicated in the spread of infectious diseases [11]. With the continuous growth of populations in Bariga, human mobility and interactions, production of biological aerosols becomes inevitable in a densely populated area (such as market squares, bus stops and other public gatherings) than in less densely populated area (such as residential and streets within the residential areas) thus increasing the presence of microorganisms putting forward an eco-epidemiological assumption that disease transmission at the population level could be caused by the mixing patterns of individuals within the environment [12]. Respiration and the shedding of microorganisms daily by the human populace as they interact with one another and their surrounding contribute to bioaerosols built-up in such environments [13, 14].

Potential health effects on the community bioaerosols depend on caused bv the pathogenicity or immunogenic potential of specific microorganisms and/or their metabolites, as well as other environmental conditions that can influence their survival in the air, the behaviour of the bio-aerosol particles and immunogenic status of the host [15]. The short-term and long-term exposure to air polluted with bioaerosols by members of the community increases their chances of contracting bioaerosols-related infections or reactions such as acute respiratory diseases, skin reactions, cough, watery eyes, difficulty in breathing and predisposing factors to pneumonia, influenza, measles, asthma, allergies, tuberculosis. cardiovascular diseases and gastrointestinal illness as long term effects [5, 16, 17, 18]. Many countries of the world, including Nigeria, has not recovered from the economic recession due to airborne bioaerosol disease such as COVID-19. Findings revealed that Nigeria's Gross Domestic Products (GDP) fell by 23% during the lockdown, especially in the food chain, with a GDP of 11%, which was primarily due to restrictions of movement that prevented food from the North to the South leading to hunger and poverty on households at 9% [19]. In Nigeria, there is no official data on economic loss due to bioaerosols-originated infections. However, the current fight against COVID-19, which is a bioaerosols-related disease, was estimated to cost the country over N45b [20]. Thus, the influx of migrating population from rural to urban areas with

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increasing settlement in Bariga can lead to public health threats.

Despite these challenges, there was no previous study on the increasing emerging and reemerging airborne-related infectious diseases and how there are influenced by the increasing human population, environmental factors and their implications in the study area. The aim of this study was, therefore, to assess public health risks associated with bioaerosols generated from outdoor activities in different locations in the Bariga metropolis of Shomolu Local Government Area of Lagos State, Nigeria.

METHODOLOGY The Study Area

The study area was Bariga, a district and suburb in Lagos State, South-West, Nigeria, formerly under Shomolu Local Government Development Area (LCDA) (Figure 1). The coordinates of Bariga LCDA are 6°32′6.0″N 3°23′41.0″E with a population of 391,200 people, according to [21]. Due to the high cost of rent in Lagos Island and some parts of the Mainland, with the increasing population of rural-urban migration, low and middle-income earners patronize places like Bariga, where rent is relatively affordable, thus contributing to a high level of aerosols generation with subsequent health hazards on the community.



Figure 1: The Map of Lagos State showing red shaded study area (Bariga-Shomolu)

Study design, sampling size and study population

Multi-stage sampling technique was adopted in the study, namely: observational study design and cluster sampling methods. The amount of airborne biological aerosol particles was observed based on the frequency of anthropogenic activities that contributed to the abundance of bioaerosols in the atmosphere and which emanated from human population indices: densely populated and lessdensely populated parts of the study area. The sampling units, therefore, were aerosol particles collected from bus stops, markets, residential areas and streets in the study area.

In the second stage sampling technique, four hundred (400) samples of bioaerosols from the atmosphere were randomly collected from twenty (20) clusters using random cluster sampling technique from different locations (Table 1). The sample size of approximately four hundred (400) was determined using the Yamani formula [22]. The observation was made from selected clusters to represent the sample populations; this was based on the variables, which were the frequency of anthropogenic activities resulting in the production of biological aerosols in the study area. Thus samples were taken from densely populated clusters (markets and bus stops) and less-densely populated clusters (residential areas and streets) for microbial population density enumeration (bacteria and fungi) and the likely effects it will have on the populace. This is because the more densely the populations of people engaging in different activities, the higher the probability of biological aerosol generation compared to the less densely populations. Table 1 indicates the sampling frame where samples were taken from the clusters.

| S/No | Densely populated clusters | Less-densely populated clusters |
|------|-------------------------------------|---------------------------------|
| 1. | Bariga Market – BM | Olanrewaju Street - OL |
| 2. | Ilaje Bus Stop - IL | Ifelodun Street - IF |
| 3. | New Garage - NG | Community Road - CR |
| 4. | Odo-Eran Market - OE | Temple Residential Area - TEM |
| 5. | Igbo-Igunnu Spare-Parts Market - IG | Adeboye Residential Street - AD |
| 6. | Odunsi Bus Stop - OD | Olalere Street - OLA |
| 7. | Evening Market - EM | Lawal Street - LW |
| 8. | Murtala Bus Stop - MB | Abeokuta Residential Area - AB |
| 9. | Bariga Bus Station - BS | Evans Adelaja Area - EA |
| 10. | Kajola Bus Stop – KB | Oshinfolarin Street – OS |

Table 1: The sampling frame where bioaerosols samples were collected in the study area

Methods of Data Collection

Total viable counts for bacteria and fungi (CFU/m³). temperature (°C), humiditv (%). precipitation (%), wind (km/hr), atmospheric pressure (mBar), and sunlight (Ultraviolet radiation) were primary data obtained from direct observations using quantitative measurement: microbial enumeration and their probable identity, environmental factors and secondary data obtained from review of risks assessment or health effects of biological aerosols from different articles and journals.

Microbiological analyses of bioaerosol particles

The isolation, enumeration and probable identification of microbial isolates followed microbiological standard procedures, including isolation of pure culture, cultural characterization, cellular morphology (Gram staining and fungi and biochemical characterization: staining) catalase, oxidase, starch hydrolysis, coagulase, production, carbohydrate fermentation, gas hydrogen sulphide production, indole, citrate utilization, motility, hemolysis and bile esculin tests according to [23, 24, 25]. Thus, passive air known monitoring, otherwise as Koch's sedimentation method or settling plate technique [26, 28], was adopted for this study. The exposure of nutrient plates to air allowed viable biological particles to sediments out of the air onto the nutrient plate's surface over the period of exposure. The bioaerosols were collected in duplicate plates using a random cluster sampling

technique from designated study locations (Table 1). Ninety millimetres (90 mm) diameter Petri dishes containing nutrient agar (CM0003 Oxoid, UK) and potato dextrose-chloramphenicol agar (M096 Himedia Lab, India) for the isolation of bacteria and fungi, respectively, were exposed to the air above the shoulder level for one h; at 10 mins interval, the exposed nutrient plates were removed and covered with lids until one h was completed. The inoculated plates were transported to the Laboratory of the Department of Microbiology, University of Lagos, Nigeria, for incubation at $35 \pm 2^{\circ}C$ (bacteria) for 24 h and 28 ± 2°C (fungi) for four days along with control plates. The developed colonies were counted in duplicates, and the mean values were taken to determine the colony forming units per cubic meter (CFU/m³) of the isolates and were further estimated according to Polish standard PN89/2-04088/08 [26, 27].

Measurement of environmental factors affecting the survival or dispersion of microbes in bioaerosols

Since environmental factors or parameters play an important role in the survival, growth and dispersal of released bioaerosols [29], they were measured at every sampling location with handheld Kestrel (3000) weather meter and Galaxy A01 Samsung Global Positioning System (GPS); the values for Humidity, Atmospheric temperature, Pressure, Sunlight or Ultraviolet-radiation, Wind and Precipitation were recorded (Table 2A, 2B).

| Environmental Parameters at 95% confidence | | Temperature (°C) | Humidity (%) | UV-radiation | Atmospheric pressure | Precipitation (%) | Wind (mph) |
|--|------------------------------------|---------------------|-----------------|--------------|-------------------------|-------------------|---------------|
| S/N | Locations | | | | (mBar) | | |
| 1. | Bariga Bus-station | 29.2±1.46 | 41.0±2.05 | 06.0±0.3 | 1012.0±50.6 | 00.0±0.0 | 0.8±0.04 |
| 2. | Bariga Market | 29.9±1.50 | 36.1±1.80 | 08.0±0.4 | 1009.0±50.5 | 00.0±0.0 | 1.0±0.05 |
| 3. | Evening Market | 30.8±1.54 | 69.4±4.82 | 11.0±0.6 | 1010.0±50.5 | 20.0±1.0 | 0.6±0.03 |
| 4. | lgbo-lgunnu Spares- Part Market | 33.0±1.65 | 29.4±1.47 | 11.0±0.6 | 1010.0±50.5 | 10.0±0.5 | 1.2±0.06 |
| 5. | Ilaje Bus stop | 31.9±1.60 | 74.4±3.72 | 11.0±0.6 | 1012.0±50.6 | 40.0±2.0 | 0.8±0.04 |
| 6. | Kajola Bus stop | 29.3±1.47 | 68.8±3.44 | 04.0±0.2 | 1010.0±50.5 | 25.0±1.25 | 0.5±0.03 |
| 7. | Murtala Bus stop | 30.5±1.53 | 70.3±3.52 | 09.0±0.5 | 1011.0±50.6 | 10.0±0.5 | 1.5±0.08 |
| 8. | New Garage | 32.1±1.61 | 71.7±3.59 | 11.0±0.6 | 1012.0±50.6 | 70.0±3.5 | 1.3±0.07 |
| 9. | Odo-Eran market | 32.1±1.61 | 71.5±3.58 | 04.0±0.2 | 1009.0±50.5 | 30.0±1.5 | 2.6±0.13 |
| 10. | Odunsi Bus stop | 28.2±1.41 | 36.6±1.83 | 05.0±0.3 | 1012.0±50.6 | 10.0±0.5 | 0.7±0.04 |

Table 2A: Environmental factors affecting the survival or dispersal of microbial cells suspended in the bioaerosol particles in the densely populated study area

The meteorological parameters obtained in the tables both in densely and less-densely populated study areas showed a relatively significant rise in temperature as the ultraviolet radiation (sunlight) rises. The rising and falling of environmental factors witnessed in this study were majorly due to variations in time and season when the parameters were measured.

Table 2B: Environmental factors affecting the survival or dispersal of microbial cells suspended in the bioaerosol particles in the less-densely populated study area

| | onmental Parameters % confidence | Temperature (°C) | Humidity (%) | UV- radiation | Atmospheric pressure (mBar) | Precipitation (%) | Wind (mph) |
|-----|-------------------------------------|---------------------|-----------------|------------------|--------------------------------|-------------------|---------------|
| SN | Locations | | | | | | |
| 1. | Abeokuta Street | 29.5±1.48 | 79.0±3.95 | 05.0±0.25 | 1011.0±50.6 | 50.0±2.5 | 0.5±0.03 |
| 2. | Adeboye Street | 30.2±1.51 | 62.8±3.14 | 11.0±0.55 | 1013.0±50.7 | 00.0±0.0 | 1.0±0.05 |
| 3. | Community road | 28.8±1.44 | 73.2±3.66 | 02.0±0.10 | 1010.0±50.5 | 06.0±0.3 | 4.0±0.20 |
| 4. | Evans-Adelaja | 32.2±1.61 | 80.0±4.00 | 11.0±0.55 | 1008.0±50.4 | 10.0±0.5 | 0.6±0.03 |
| 5. | Ifelodun Street | 32.6±1.63 | 69.5±3.48 | 02.0±0.10 | 1010.0±50.5 | 50.0±2.5 | 2.3±0.13 |
| 6. | Lawal Street | 29.1±1.46 | 75.6±3.75 | 07.0±0.35 | 1010.0±50.5 | 06.0±0.3 | 1.5±0.08 |
| 7. | Olalere Street | 34.8±1.74 | 29.5±1.48 | 11.0±0.55 | 1009.0±50.5 | 10.0±0.5 | 1.2±0.06 |
| 8. | Olanrewaju Street | 29.7±1.49 | 80.2±4.01 | 06.0±0.33 | 1012.0±50.6 | 60.0±3.0 | 0.6±0.03 |
| 9. | Oshinfolarin Street | 32.7±1.64 | 59.0±2.95 | 11.0±0.55 | 1010.0±50.5 | 00.0±0.0 | 0.9±0.05 |
| 10. | Temple Area | 32.4±1.62 | 36.5±1.83 | 08.0±0.40 | 1011.0±50.6 | 00.0±0.0 | 0.8±0.04 |

Public health risks assessment of bioaerosols and its economic implications

Public health risks assessment of bioaerosols was evaluated based on Occupational Safety and Health Administration recommendation [6, 30] and empirical knowledge on the pathogenicity of microbial isolates, their potential adverse health effects on the exposed population and their economic implications on the community. The economic implication involved the direct and indirect costs of the treatment of diseases caused by these identified infectious agents.

Exposure duration to pollutants of anthropogenic origin as a risk factor and ways to mitigate them

Pollutants of anthropogenic origin are pollutants that emanate from human activities that affect the ecosystem; some of these activities cause environmental pollution; for example, combustion of fossil fuel, waste disposal, urbanization, industrialization, agricultural activities, deforestation, construction, etc. [31, 32] Short-term and long-term exposure to these pollutants can cause acute and chronic respiratory diseases; it is important that careful evaluation of the findings to establish the relationship between duration of exposure to bioaerosols and health hazards as hypothesized was done using relevant statistical tools and empirical methods. Thus the findings were then reviewed based on the findings of other researchers, and necessary solutions were proffered to reduce their effects.

Data Analysis

The primary data obtained in this study from direct observation or measurement of samples in duplicates were sorted, processed, analyzed, presented and interpreted using mean, tables, charts, One-way analysis of variance (ANOVA) and F-distribution calculator [33].

RESULTS

This study showed variation in environmental parameters (Table 2A, 2B) at a p=0.05 significance level, which was largely dependent on time and seasons with subsequent adverse effects on the presence and movement of biological aerosol droplets suspended in the study area. The data obtained in this study showed that there was variation in the microbial concentration, with densely populated areas having the highest

microbial populations than less densely populated areas. The highest bacterial populations within the exposed period of 1h were recorded in Bariga Market (BM), which ranged from 4.40x10⁵ to ND and lowest in New Garage (NG) from 2.83x10⁴ to 2.46×10^4 (Table S1, Figure 2A), while fungal populations in the same category showed Bariga Market (BM) with highest populations ranging from 2.04×10^4 to 3.93×10^2 and lowest in Odo-Eran Market (OE) with 0.00×10^1 to 1.57×10^3 (Table S2, Figure 2B). The less-densely populated study areas, on the other hand, showed the highest bacterial populations in the Temple area (TEM), ranging from 8.33×10^4 to 4.87×10^4 cfu/m³ and lowest in Olanrewaju Street (OL) from 1.18x10⁴ to 6.94x10³ (Table S3, Figure 3A) while the highest fungal population was observed in Temple area (TEM) 3.14×10^4 to 8.64×10^3 and lowest in Abeokuta Street (AB) from 0.0x10¹ to 2.75x10³ (Table S1, S2, S4 - https://getjournal.org/wpcontent/uploads/2023/05/Supplementary-Table-GJOBOH-2022-031.pdf and Figure 3B). Tables 3A and 3B show the microbial prevalence rate, while Table 4 shows public health risk assessment values compared to a recommended threshold.

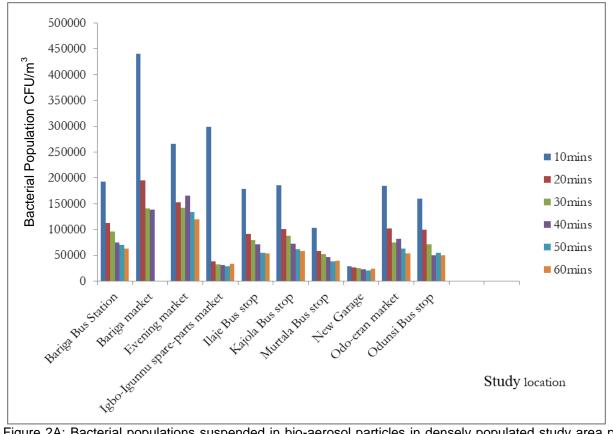


Figure 2A: Bacterial populations suspended in bio-aerosol particles in densely populated study area per hour

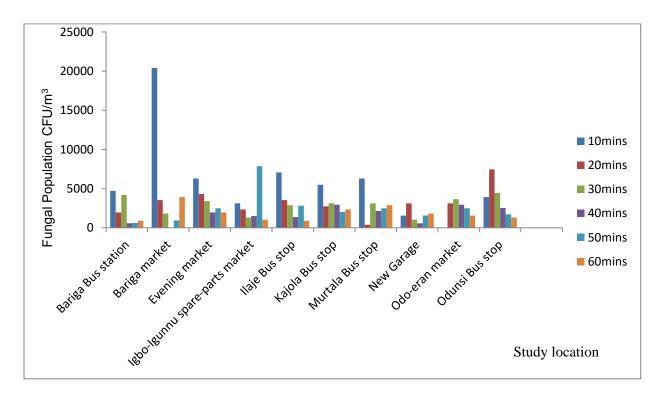


Figure 2B: Fungal populations suspended in bio-aerosol particles in densely populated study area per hour

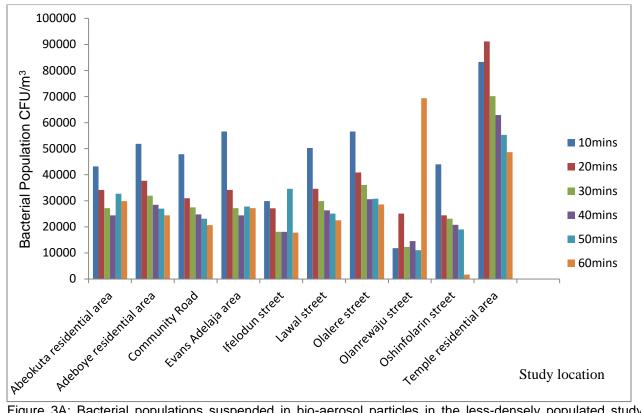


Figure 3A: Bacterial populations suspended in bio-aerosol particles in the less-densely populated study area

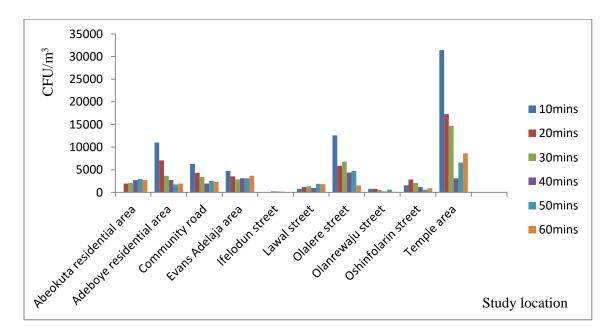


Figure 3B: Fungal populations suspended in bio-aerosol particles in less-densely populated study area per hour

| S/No | Bacteria Genera | Frequency | Percentage (%) | Prevalence rate (%) |
|------------|------------------------|-----------|----------------|---------------------|
| 1. | Staphylococcus | 32 | 41.56 | 8.18 |
| 2. | Bacillus | 26 | 33.77 | 6.65 |
| 3. | Micrococcus | 06 | 7.79 | 1.53 |
| 4. | Pseudomonas | 04 | 5.19 | 1.02 |
| 5. | Actinomycetes | 02 | 2.60 | 0.51 |
| 6. | Streptococcus | 01 | 1.30 | 0.26 |
| 7. | Shigella | 01 | 1.30 | 0.26 |
| 8. | Serratia | 01 | 1.30 | 0.26 |
| 9. | Enterobacter | 01 | 1.30 | 0.26 |
| 10. | Escherichia | 01 | 1.30 | 0.26 |
| 11. 12. | Salmonella Yersinia | 01 01 | 1.30 1.30 | 0.26 0.26 |
| | Total | 77 | 100 | |

Table 3A: Prevalence rate of Bacterial Genera in the study population

Table 3B: Prevalence rate of fungal genera in the study population

| S/No | Fungi Genera | Frequency | Percentage (%) | Prevalence rate (%) |
|------|---------------|-----------|----------------|---------------------|
| 1. | Aspergillus | 21 | 36.21 | 5.37 |
| 2. | Fusarium | 16 | 27.59 | 4.09 |
| 3. | Saccharomyces | 07 | 12.07 | 1.79 |
| ŀ. | Neurospora | 05 | 8.62 | 1.28 |
| 5. | Rhizopus | 05 | 8.62 | 1.28 |
| б. | Mucor | 04 | 6.90 | 1.02 |
| | Total | 58 | 100 | |

These tables state the prevalence rate of bacteria and fungi in the studied population.

| Table 4: Public health | | | | |
|--------------------------------------|--------------------------------|--------------------------|--|---|
| Study location | Mean total | Mean total | Guideline | Remarks |
| | Bacteria CFU/m ³ | Fungi CFU/m ³ | CFU/m ³ | |
| L1-Bariga Bus Station | 2.88 x 10 ⁵ | 5.50 x10 ³ | <1.0 x 10 ² | Low |
| | | | 10 ² –1.0 x 10 ³ | Intermediate |
| | F | 2 | >1.0 x 10 ³ | High |
| L2-Bariga Market | 4.97 x 10 ⁵ | 7.86 x 10 ³ | For houses | |
| | | | $<2.0 \times 10^{2}$ | Low |
| | | | $<1.0 \times 10^{3}$ | Intermediate |
| | 4 57 405 | 0.50 403 | $<1.0 \times 10^{4}$ | High |
| L3-Evening Market | 4.57 x 10 ⁵ | 9.59 x 10 ³ | >1.0 x 10 ³ | Indicates contamination |
| | | | >1.06 x | Indicates contamination |
| | | | 10 ² fungi/g of dust | |
| L4-Igbo-Igunnu | 1.12 x 10 ⁵ | 4.72 x 10 ³ | $a = 4a^2$ | NI I |
| L C Illeia Dua Otan | 2.47 x 10 ⁵ | 8.02 x 10 ³ | <8.0 x 10 ² <3.0 x 10 ² | Normal |
| L5-Ilaje Bus Stop | 2.47 x 10° | 8.02 x 10° | <3.0 x 10 ⁻ | Common fungi are accepted |
| | | | <1.5 x 10 ² | Mixedfungi other than pathogenic |
| L6-Kajola Bus Stop | 2.67 x 10 ⁵ | 9.43 x 10 ³ | <7.5 x 10 ² | is accepted Total airborne bacteria and fungi are accepted if species |
| 17 Murtolo Duo Ston | 1.65 x 10 ⁵ | 1.04 x 10 ⁴ | | are not infective or allergenic |
| L7-Murtala Bus Stop L8-New Garage | 8.33×10^4 | 6.83×10^{3} | | |
| L9-Odo-Eran Market | 2.63×10^{5} | 8.80×10^{3} | | |
| L10-Odunsi Bus Stop | 2.33×10^{5} | 9.82×10^{3} | >1.04 x 10 ² | Total fungi is a threat to health |
| | | | >5.0 x 10 ² | One species of potentially pathogenic nature is a threat to |
| | 4.05 4.05 | 0.00 403 | | health |
| L11-Abeokuta Street | 1.05 x 10 ⁵ | 8.80 x 10 ³ | | |
| L12-Adeboye Street | 1.03 x 10 ⁵ | 1.12 x 10 ⁴ | | |
| L13-Community Road | 8.86 x 10 ⁴ | 9.98 x 10 ³ | | |
| L14-Evan Adelaja | 1.01 x 10 ⁵ | 1.18 x 10⁴ | | |
| L15-Ifelodun Street | 8.17 x 10 ⁴ | 7.86 x 10 ² | | |
| L16-Lawal Street | 9.57 x 10 ⁴ | 5.27 x 10^{3} | | |
| L17-Olalere Street | 1.16 x 10 ⁵ | 1.66 x 10 ^₄ | | |
| L18-Olanrewaju Street | 4.24 x 10 ⁴ | 1.57 x 10 ³ | | |
| L19-Oshinfolarin Street | 7.39 x 10 ⁴ | 4.56 x 10 ³ | | |
| L20-Temple Area | 2.16 x 10 ⁵ | 3.46 x 10 ⁴ | | agerosals and their impact on human |

Table 4: Public health risks or hazard assessment of bioaerosols on the study population

This remark was made based on guidelines adopted from Kim, K.-H.et al. (2017). Airborne bioaerosols and their impact on human health, J. Environ. Sci. (2017), https://doi.org/10.1016/j.jes.2017.08.027

DISCUSSION

As a result of daily anthropogenic activities in the Bariga metropolis of Shomolu, Lagos, Nigeria, microorganisms are released in the form of bioaerosol droplet nuclei, which float in the atmosphere. People are exposed to these bioaerosols being carried about by wind and other environmental factors through the exchange of gases (respiration), contaminated air containing pathogens is inhaled and equally released via talking, coughing, sneezing, yawning and laughing. The exposed individuals, especially the vulnerable ones, may come down with one form of infection or another, thus causing public health threats. Some of these environmental factors have been observed to be relatively less densely populated than less densely populated study areas, thus affecting the microbial distribution and leading to a high population of microorganisms in densely populated areas than in less densely populated study areas. This assertion was corroborated by Samaranayakea et al. [34], who stated that aerosols could be transported or suspended in the air for considerable periods of time depending on the humidity, airflow, and temperature of the environment into which they are expelled. Górny [35] also observed that microclimate parameters such as temperature and relative humidity of the air, together with ultraviolet radiation, oxygen and other physical factors, contribute to the influx of airborne microorganisms. These environmental factors also influence the survival of airborne microbes and affect their ability to colonize surfaces after deposition. As the ultraviolet radiation rises over time, there is a corresponding gradual rise in temperature. This rise in temperature between 28.2-34.8°C, as observed in this study, favoured the survival and viability of mesophiles which are generally known to be human pathogens and capable of surviving at body temperature (37°C). Humidity and precipitation support the growth of microorganisms, as bacterial cells are composed mainly of water. Though the high the precipitation, the less the microbial population is due to the fact that rainfall washes the microorganisms from the atmosphere unto open surfaces. The total microbial population densities varied from location to location depending on the human population densities. It was observed that mean microbial colonies increase with the increase in exposure period with a decline in CFU/m³ when estimated with Polish standard PN89/2-04088/08. The high microbial population recorded in densely populated areas is attributable to the influx of people and their daily activities that resulted in the generation of bioaerosols leading high microbial to contamination above the standard limits of 0 - 1.0x 10[°] CFU/m[°] [15]. Bariga market recorded high microbial populations. Market squares and bus terminals are heterogeneous settings where several activities are taking place daily. Human, animal, vehicular movement and unhygienic environmental practices become the order of the day with the resultant generation of one form of aerosols or another, thus polluting the air and exposing the market and bus patrons to the danger of respiratory diseases, skin diseases, systemic diseases and allergic reactions. This study is in line with Adams et al. [14], who affirmed that the population density of microorganisms in bioaerosol suspended in the atmosphere in a particular environment is influenced by human populations and their interactions with such environment. On the other hand, the high microbial populations witnessed at Temple residential area compared to others in the same categories could be a result of the poor environmental sanitation and lack of personal hygiene practised by the residents. The

place was observed to be more of a 'ghetto' settlement inhabited by low-income earners that will and capacity to lacked the practice environmental hygiene compared to the Olanrewaju residential area, where the residents were middle-income earners and educated. Thus the level of environmental and personal hygiene was relatively acceptable compared to other study locations within the populations. This study shows that the predominant bacteria isolated were Grampositive bacteria (87.02%), with the genus, population Staphylococcus having (41.56%) frequency and the least prevalence rate (1.79%). The high occurrence of Gram-positive bacteria and moulds in this study may be due to their ubiquitous nature, ability to survive longer in the air, colonization of the skin and other body orifices, presence of thick layers of peptidoglycan in their cell wall (20 - 80 nm), possession of spores by some genus Bacillus and mold, presence of small protective molecules such as sugars, amino acids, alcohols and betaine, expression of heat shock proteins [36]; whereas least occurrence of Gramnegative bacteria which are majorly enteric bacteria recorded in this study could be due to their inability to survive outside their natural habitat (human body) and the presence of a thin layer of peptidoglycan in their cell wall (2 - 3 nm) surrounded by an outer membrane containing lipopolysaccharide [37]. This observation is in conformity with earlier findings [1, 6, 26, 38] which isolated some predominant bacterial genera of Bacillus. Staphylococcus, Streptococcus. Escherichia, Micrococcus, Pseudomonas, Serratia, Salmonella. Shigella and fungal genera of Aspergillus, Fusarium, Rhizopus. and Saccharomyces. Some of the bioaerosols in this study are capable of causing infections in exposed individuals to contaminated air; for instance, Staphylococcus sp is a potential cause of sepsis, infective endocarditis, osteomyelitis, skin and soft tissue infections [39]; Bacillus sp has also been implicated in previous studies as a potential cause of abscesses, bacteremia, wound and burn infections, ear infections, endocarditis, meningitis, ophthalmitis, osteomyelitis, peritonitis, respiratory, urinary tract infections and food poisoning [40]. Micrococcus spp. are capable of causing opportunistic infections in immuno-compromised HIV patients [41], Actinomyces which causes Actinomycosis [42, 43], Streptococcus has been identified in previous studies to be responsible for Strep throat infection, scarlet fever, impetigo, cellulitis, streptococcal toxic shock syndrome and rheumatic fever [44]. Pseudomonas, Shigella, Serratia, Enterobacter, Escherichia, Salmonella and Yersinia are Gram-negative, enteric bacteria

responsible for gastrointestinal diseases in infants, adults and the elderly with characteristics of abdominal cramps and diarrhoea, nausea and vomiting, gastroenteritis, fever, and other clinical infections such as neonatal meningitis and pneumonia [45-47]. Moulds and Bacillus are generally spore-producing organisms which can withstand unfavourable environmental conditions; their spores can easily be carried about from one place to another by the wind until they come in contact with a suitable medium that supports their growth. The presence of these spores in the study area is a public health threat, thus exposing individuals, especially those with underlying diseases such as asthma and respiratory diseases, to inhaling contaminated air leading to allergic reactions and severe respiratory infections. Some Aspergillus species have been implicated in causing mycoses in humans, allergic sinusitis, and systemic and respiratory diseases in exposed hosts [1, 45, 48]. Fusarium spp. is known to cause superficial, locally invasive, diffuse infections and keratomycosis in humans [49]. Saccharomyces sp can cause opportunistic infections on exposed hosts [52]; Neurospora sp is a non-human pathogen useful as an experimental model in genetics [48]. Rhizopus sp and Mucor are known for causing Mucormycosis [51]. The predominance of bioaerosols in the study area and their ability to survive harsh conditions of the atmosphere other than their physiochemical properties might be due to the presence of dust in the air, which supports their existence in spore form [16]. This is in conformity with the World Health Organization, which postulated that exposure to microbial agents of aerosol origin increases the risks of rare health conditions. The variation and prevalence of bacterial and fungal genera in the air, as observed here, are due to differences in stability and viability in the air. Gorny [35] asserted that air as a biotope does not support the survival of biological agents, but he further stated that several studies showed that numerous fine particles of biological origin were able to maintain their viability and immunological reactivity in the air much longer than bigger organisms. This also explains why the bacterial population is high in this study, considering their size, which is between 0.5 - 2.0 µm, than fungi, whose individual cell size can vary widely from $2 - 3 \mu m$ to $20 - 50 \mu m$ in length.

The potential health hazards associated with frequent exposure to bioaerosols in the study population were shown to be very high in all locations. According to Occupational Safety and Health Administration [30] and other regulatory bodies provided a guideline for microbial load threshold between $10^2 - 10^4$ CFU/m³ (low – high

risk); though this value varies from country to country. In Nigeria, for instance, there is no official data from environmental and health regulatory bodies on biological aerosols threshold; the data obtained in this study were interpreted based on standards/guidelines for bioaerosols proposed by different governmental and private organizations; based on European literature databases on residential indoor air quality as reported by Kumar et al. [5]. This study showed mean microbial load of 7.86 x $10^2 - 4.97$ x 10^5 CFU/m³. Hence it can be deduced that there is a high public health risk of frequent exposure to biological aerosols in the Bariga community. Therefore, adequate hazard mitigation should be employed to reduce the risk, thus promoting healthy living.

A significant correlation (p=0.05) between bioaerosols exposure duration and the likelihood of developing the onset of diseases in the susceptible, exposed host was observed. This is supported by Franchitti et al. [52], who reported a significant correlation statistically between bioaerosols exposure and the onset of health effects on humans. The economic implication of pathogens of bioaerosols origin has been estimated: the cost of treatment of infections or diseases (\$10 - 720,167.22) per episode or annum, low productivity and deaths [53]. It can be challenging to treat some of these pathogens because of their multi drugs resistance, thus putting their hosts in grave danger and other lifethreatening situations. Preventive measures to curb the pandemic emanating from bioaerosols include guarantine, preparation of health facilities, isolating infectious cases, and tracing contacts involving public health resources. human resources and implementation costs. It also involves health system expenditures to provide health facilities for infectious cases and the arrangement of consumables such as antibiotics, and personal protective medical supplies, equipment [56]. In Nigeria, there is no official data on economic loss from bioaerosol-originated infections. However, the current fight against COVID-19 (bio-aerosol borne) disease was reported to cost the country over N45b [25]. Bioaerosols-related infectious diseases have been reported to be responsible for economic losses, including medical expenditures of about US\$21 billion globally, economic productivity losses, environmental degradation and material losses [20]. This cost is significantly high in this era of economic recessions, which can create global implications of bioaerosols-related pandemic impact on the health system and simultaneously the growth prospects of the whole countries [8]. The health risk of outbreaks of infectious diseases.

the fear and worry that accompany them, low productivity due to illness and absenteeism from work and school lead to various economic risks, thus increasing poverty within the community and the nation at large.

CONCLUSION

This study has shown a significant relationship between the frequencies of exposure to bioaerosols and associated public health risks on the exposed individuals. The statistical analysis one-way analysis of variance (ANOVA) at 0.05 significance level between exposure to bioaerosols and subsequent health hazards on the general public. The variation in environmental parameters is largely dependent on time and seasons, with subsequent adverse effects on the presence and movement of biological aerosol droplets suspended in the atmosphere. The study also revealed that the study area is a high-risk area (10^2) 10⁴CFU/m³) for biological aerosol-related infections or diseases. Hence, the prevalence of genera, Staphylococcus, Bacillus, Micrococcus, Streptococcus. Pseudomonas, Actinomycetes, Shigella, Serratia. Enterobacter, Escherichia, Salmonella. Yersinia, Aspergillus, Fusarium, Saccharomyces, Neurospora, Rhizopus and Mucor is an indication of microbial contamination. The knowledge of the presence of biological agents and adaptation of lifestyles by the residents in the study area can help in environmental and public health sustainability, thus reducing bioaerosolsrelated pollution. It is therefore recommended that adequate proactive, preventive and control measures such as structural and aesthetic modification in line with urban plans should be put in place to reduce the survival, distribution, colonization and establishment of bioaerosols in the Bariga community, thus promoting the health of the residents. Lagos State Government should also intensify efforts to reduce the public health effect of bioaerosols through policies, structural planning, development, and education on environmentally friendly activities and personal hygiene.

ABBREVIATIONS

ASHRAE - The American Society of Heating, Refrigerating, and Air-Conditioning Engineers. CFU - Colony-forming Unit COVID-19 - Coronavirus disease GDP - Gross Domestic Product GPS - Global Positioning System LCDA - Local Council Development Area ND – Not Determined OECD - The Organization for Economic Cooperation and Development

OSHA - Occupational Safety and Health Administration TNTC - Too Numerous to Count UK - United Kingdom WHO - World Health Organization

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REFERENCES

[1] Fashola MO, Grillo JA, Obayori OS, Opere BO, Eguakun EA. Microbial Assessment and Antibiogram of Bacteria Isolated from Air Samples around Dumpsites in Igando, Lagos, Nigeria. NJM. 2019; 34(1) 4829 – 4841.

[2] Fernandez MO, Thomas RJ, Garton NJ, Hudson A, Haddrell A, Reid JP. Assessing the Airborne Survival Of Bacteria in Populations of Aerosol Droplets With a Novel Technology. J R Soc Interface. 2019; 16:20180779. http://dx.doi.org/10.1098/rsif.2018.0779.

[3] ASHRAE, American Society of Heating, Refrigerating and Air-Conditioning Engineers. ASHRAE Position Document on Airborne Infectious Aerosols. 1791 Tullie Circle, NE Atlanta, Georgia 30329-2305 404-636-8400. 2020. (www.ashrae.org).

[4] Fröhlich-Nowoisky J, Kampf CJ, Weber B, Huffman JA, Pöhlker C, Andreae MO, et al. Bioaerosols in the Earth System: Climate, Health, and Ecosystem Interactions. Atmos Res. 2016; 182:346-376.

https://doi.org/10.1016/j.atmosres.2016.07.018.

[5] Kumar P, Singh AB, Singh R. Spatial Variation and Comprehensive Health Risk Assessment of Microbial Indoor Air Quality in Microenvironments of North Delhi. Res Sq. 2021. https://doi.org/10.21203/rs.3.rs-445730/v1.

[6] Kim KH, Kabir E, Jahan SA. Airborne Bioaerosols and Their Impact on Human Health. J Environ Res. 2017.

https://doi.org/10.1016/j.jes.2017.08.027.

[7] Van-Leuken JPG, Swart AN, Havelaar AH, Van Pul A, Van der HW, Heederik D. Atmospheric Dispersion Modelling of Bioaerosols that are Pathogenic to Humans and Livestock—A Review to Inform Risk Assessment Studies. Microb. Risk Anal. 2016; 1:19–39. In: Kim et al. Airborne bioaerosols and their impact on human health. J Environ Sci. 2017.

https://doi.org/10.1016/j.jes.2017.08.027.

[8] Wenwen X, Yanpeng L, Wenyan B, Junli H, Tianfeng M, Xuelin Z, et al. The Source and Transport of Bioaerosols in the Air" A Review. Front. Environ. Sci. Eng. 2021; 15: (3)44. <u>https://doi.org/10.1007/s11783-020-1336-8</u>.

[9] Manisalidis I, Stavropoulou E, Stavropoulos A, Bezirtzoglou E. Environmental and Health Impacts of Air Pollution: A Review. Front Public Health. 2020;8:4.

https://doi.org/10.3389/fpubh.2020.00014.

[10] Adams RI, Bhangar S, Pasut W, Arens EA, Taylor JW, Lindow SE et al. Chamber Bio-

Aerosol Study: Outdoor Air and Human Occupants as Sources of Indoor Airborne Microbes. <u>PLoS</u> <u>One.</u> 2015; 10(7): e0133221. https://doi.org/10.1371%2Fjournal.pone.0133221.

[11] Prussin II AJ, Marr LC. Sources of Airborne Microorganisms in the Built Environment.

Microbiome. 2015; 3(78). https://doi.org/10.1186/s40168-015-0144-z.

[12] Banerjee S. Environmental Factors Affecting Microbial Growth. The Biology Notes. 2022. https://thebiologynotes.com/environmental-factorsaffecting-microbial-growth/ (Accessed 10th October 2022).

[13] ASM, American Society for Microbiology. Why Studying Microorganisms in The Air is Vital. 2020. https://asm.org/Articles/2020/December/Why-

Studying-Microorganisms-in-the-Air-Is-Vital.

(Accessed 15th November 2021).

[14] Adams RI, Bhangar S, Pasut W, Arens EA, Taylor JW, Lindow SE et al. Chamber Bio-Aerosol Study: Outdoor Air and Human Occupants as Sources of Indoor Airborne Microbes. PLoS One. 2017.

https://doi.org/10.1371/journal.pone.0128022

[15] Mbareche H, Morawska L, Duchaine C. On The Interpretation of Bioaerosol Exposure

Measurements and Impacts on Health. J Air Waste Manag Assoc. 2019; 69:(7):789-804.

[16] Chretien JP, Anyamba A, Small J, Britch S, Sanchez JL, Halbach AC. et al. Global Climate Anomalies and Potential Infectious Disease Risks 2014–2015. PLoS Curr. 2015. 26:17. https://doi.org/10.1371/currents.outbreaks.95fbc4a 8fb4695e049baabfc2fc8289f.

[17] Cleveland Clinic. Infectious Diseases. 2021. https://my.clevelandclinic.org/health/diseases/1772 4-infectious-diseases. (Accessed 6th October 2021).

[18] Éze IC, Schaffner E, Fischer E, Schikowski T, Adam M, Imboden M et al. Long-Term Air Pollution Exposure and Diabetes in a Population-Based Swiss Cohort. Environ Int. 2014; 70:95–105.

[19] Adam K, Edeh H, Oboh V, Pauw K, Thurlow J. Impacts of COVID-19 on Food Systems and Poverty in Nigeria. Advances in Food Security and Sustainability. 2020; 5:145–73.

[20] Muanya C. How COVID-19 Treatments Cost Government Over N44.9b. Guardian Newspaper. 2020 7th August. (Accessed 9th November 2021).

[21] NPC, National Population Commission. Bariga History. 2006. https://wikipedia.com. (Accessed 24th August 2021).

[22] Yamani T, Statistics: An Introductory Analysis. 2nd Edition. New York: Harper and Row. 1967.

[23] Willey JM, Sherwood LM, Woolverton CJ, Prescott LM, Harley JP, Klein DA.

Prescott, Harley, and Klein's Microbiology 10th Edition. McGraw-Hill, New York. 2017.

[24] Aryal S. Biochemical Test for Bacteria. Microbiology info.com.

https://microbiologyinfo.com/category/biochemicaltest-of-bacteria/. (Accessed 4th October 2022).

[25] Cheesbrough M. District Laboratory Practice in Tropical Countries Part 2. Second Edition. Cambridge, University Press, New York. 2018.

[26] Ambrose I, Nweke CO, Umeh SCI, Braide W. Prevalence Of Bio-Aerosols in the Outdoor Air Environment in Uyo Urban, Akwa Ibom State, Nigeria. Res J Microbiol. 2015; 6(2):012-019.

[27] Shiferaw T, Gebre-silasse L, Mulisa G, Zewidu A, Belachew F, Muleta D et al. Bacterial Indoor-Air Load and Its Implications For Healthcare-Acquired Infections in a Teaching Hospital in Ethiopia. Int J Infect Control. 2016; 12(1):1-9.

[28] Merck K. Settle Plates for Microbial Air Monitoring: Darmstadt, Germany and/or its affiliates. 2021. (Accessed 18th August 2021).

[29] Aryal S, O'Neill F. Basic Microbiology: Environmental Factors Affecting Microbial Growth. The Biology Note. 2021. https://thebiologynotes.com. (Accessed 30th August 2021).

[30] OSHA, Occupational Safety and Health Administration, "Indoor air quality-proposed rule" notice of proposed rulemaking. Fed. Regist. 1994; 59 (65): 15968–16039. In: Kim, K.-H., et al. Airborne Bioaerosols and Their Impact on Human Health", J. Environ. Sci. 2017. https:// doi.org/10.1016/j.jes.2017.08.027.

[31] Bali AS, Kumar V. Bioremediation: Physiology, Molecular and Biotechnological Intervention. Handbook of Bioremediation. 2021; 727-737.

[32] Conaway N, Dunn T. Anthropogenic Climate Change Factors and Impacts. 2021. https://study.com/academy/lesson/anthropogenic-

<u>climate-change-definition-factors.html</u>. (Accessed 19th February 2023).

[33] Zach. F Distribution Calculator. 2018. https://www.statology.org/f-distribution-calculator. (Accessed 2nd April 2022). [34] Samaranayakea LP, Fakhruddina KS, Buranawat B, Panduwawala C. The Efficacy of Bio-Aerosol Reducing Procedures Used in Dentistry a Systematic Review. Acta Odontol Scand. 2021;.79(1): 69-80.

[35] Górny RL. Bioaerosols and OSH. 2020. https://test-

oshwiki.osha.europa.eu/wiki/Bioaerosols_and_OS H. (Accessed 1st March 2022).

[36] Bruslind L. Bacterial Cell Walls. General Microbiology.

https://open.oregonstate.education/generalmicrobi ology/chapter/bacteria-cell-walls/. (Accessed 19th February 2023).

[37] Gary K. The Gram-Negative Cell Wall. LibreTexts Biology. 2022. open. Oregon State.education/general

microbiology/chapter/bacteria-cell-walls/.

(Accessed 19th February 2023).

[38] Ohagim PI, Ikon GM, Matthew PC, Ohagim GA. Microbiological Assessment of Indoor Air in Public Toilets across Selected Motor Parks in Owerri Metropolis, Nigeria. J Microbiol Exp. 2017; 5(6):00166. <u>10.15406/men.2017.05.00166</u>

[39] Tong SYC, <u>Davis</u> JS, <u>Eichenberger</u> E, Holland TL, Fowler Jr VG. *Staphylococcus aureus* Infections: Epidemiology, Pathophysiology, Clinical Manifestations and Management. Clin. Microbiol. Rev. 2015; 28: (3)603–61.

[40] Paoli CJ, Reynolds MA, Sinha M, Gitlin M, Crouser E. Epidemiology and Costs of Sepsis in the United States—An Analysis Based on Timing of Diagnosis and Severity Level. Crit Care Med. 42018; 6: (12)1889–189. https://doi.org/10.1097/ccm.00000000003342.

[41] Chukwu VA, Nnadozie TN, Nsemoh HE, Obiekezie SO. Physicochemical and Microbiological Studies of Urine Contaminated Soil in Abia State University, Uturu Campus. Curr Trends Biomed Eng Biosci. 2018; 14(1):009-015.

[42] Brazier Y. What you Should Know About Actinomycosis? 2018.

https://www.medicalnewstoday.com/articles/24514 4. (Accessed 5th April 2022).

[43] J. F. Okulicz. Actinomycosis In: Drug and Infectious Disease. October 2019. https://emedicine.medscape.com/article/211587overview. (Accessed 22nd April 2022). [44] CDC, Centers for Diseases Control and Prevention. Diseases Caused by Group A Strep. 2022. <u>https://www.cdc.gov/groupastrep/diseases-</u> public. (Accessed 12th October 2022).

[45] Madappa T. Escherichia coli (E coli) InfectionsIninfectiousDiseases.2019.https://emedicine.medscape.com/article/217485-

overview. (Accessed 8th March 2022).

[46] Ramirez D, Giron M. *Enterobacter* Infections. 2022.

https://www.ncbi.nlm.nih.gov/books/NBK559296/. (Accessed 8th September 2022).

[47] Cleveland Clinic. Infectious Diseases. 2022. https://my.clevelandclinic.org/health/diseases/1772 4-infectious-diseases. (Accessed 6th October 2022).

[48] Bongomin F, Asio LG, Baluku JB, Kwizera R, Denning DW. Chronic Pulmonary Aspergillosis: Notes for a Clinician in a Resource-Limited Setting Where There Is No Mycologist. J Fungus. <u>2020</u>; <u>6(2):75. https://doi.org/10.3390/jof6020075</u>.

(Accessed 12th October 2022).

[49] Askun T. Introductory Chapter: *Fusarium*: Pathogenicity, Infections, Diseases, Mycotoxins and Management. 2018.

http://dx.doi.org/10.5772/intechopen.76507.

[50] Algazaq JN, Akrami K, Martinez F, McCutchan A, Bhart A. *Saccharomyces cerevisiae* Laryngitis and Oral Lesions in a Patient with Laryngeal Carcinoma. Case Rep Infect Dis. 2017; 2941527. <u>https://doi.org/10.1155/2017/2941527</u>. (Accessed 28th March 2022).

[51] Sandhu A. Mucormycosis (Zygomycosis). July 2021.

https://emedicine.medscape.com/article/222551overview. (Accessed 8th March 2022).

[52] Franchitti E, Pascale E, Fea E, Anedda E, Traversi D. Methods for Bioaerosol

Characterization: Limits and Perspectives for Human Health Risk Assessment in Organic Waste Treatment. Atmosphere. 2020; 11(5):452. <u>https://doi.org/10.3390/atmos11050452</u>. (Accessed 15th November 2021).

[53] Yunfeng S, Haiwei L, Ren Z. Effects of Pandemic Outbreak on Economies: Evidence from Business History Context. Front. Public Health. 2021; 9.

https://doi.org/10.3389/fpubh.2021.632043.