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INTERNATIONAL JOURNAL  
OF INNOVATIVE AND APPLIED RESEARCH

## RESEARCH ARTICLE

Article DOI: 10.58538/IJAR/2110

DOI URL: <http://dx.doi.org/10.58538/IJAR/2110>

### BACTERIOLOGICAL QUALITY OF WELL WATER USED FOR DOMESTIC PURPOSES IN OWO METROPOLIS, ONDO STATE, NIGERIA

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#### Manuscript Info

#### Manuscript History

Received: 28 August 2024  
Final Accepted: 29 September 2024  
Published: September 2024

#### Keywords:

Well-Water, Pollution, Pathogens,  
Water-Borne Illness, Health Risk

#### Abstract

Water from locally dug wells represent the major source of water supply for domestic purposes in most rural and sub-urban communities in Nigeria. Well water is prone to contamination from human and animal activities and from the environment with microorganisms that may include pathogens associated with water-borne illnesses or possess the potential to initiate diseases of opportunities. This study aimed to determine the bacteriological class of well water available for domestic use in Owo Metropolis, Ondo State, Nigeria. Well water samples collected, analyzed by the double-strength and single-strength McConkey broth for the most probable number, MPN of coliform bacteria, and their varieties from subcultures. Antimicrobial susceptibility tests and extended-spectrum beta-lactamase, ESBL were also determined. The MPN in all the samples were several folds higher (15-1800+ MPN/100 mL) than recommended upper limits. Klebsiella species was the most dominant microorganism 46(50%) of 92 isolates. Other isolates included Enterobacter species (25%), Pseudomonas aeruginosa (12.0%), Proteus mirabilis (8.7%) and Serratiamarcescens (4.4%) was the least recovered microorganism. Production of ESBL was highest amongst Klebsiella species (19.7%), Enterobacter species (16.0%), Proteus mirabilis (12.5%) and Pseudomonas aeruginosa (9.1%) and Serratiamarcescens tested negative. The sampled wells were profoundly polluted with microorganisms that carry resistance enzymes incriminated in most nosocomial infections thus, demanding concrete intervention strategies to circumvent major water-borne disease outbreaks.

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#### Introduction:-

Water supply has been a major determinant in location of human settlements for generations and there is always poor accessibility in most low-income countries [1,2]. The quality of water supply is dependent on the source and has critical influence on the health of the community [3]. The natural and traditional supply of water comes from

rivers, springs or underground water sourced through dug wells [4]. Hand dug well water represents a key water supply source for drinking and domestic use in poor-resource economies particularly in the rural and sub-rural settings [5]. Shallow wells are expected to be 15 m deep and deep wells 50 m at the minimum [6]. The geological constitution of the rock from which the well is dug (or karst) and sediment (or aquifer) in the water are key determinants of the water purity [7]. The physical, chemical and microbiological physiognomies of water reflect the properties of its potability [8]. Pollution of water is the consequence of anthropogenic and geogenic activities [9,10]. The pollution of ground water come with grave imports for drinking and domestic purposes [11]. Understandably, underground water, does not favor the growth of bacteria due to deficiency in sufficient nutrients to support multiplication and initiation of illnesses [12,13]. Pollution of water with pathogenic microorganisms is the main recurrent hazard of well water. The consistent presence of opportunistic pathogens in water and the detection of indicator microorganisms such as *Escherichia coli* and or *Enterococcus faecalis* pointedly indicate fecal pollution [1]. Coliform bacteria other than *Escherichia coli* are more abundant in the gut and survive longer in water thus making their presence in water a valuable indicator of fecal contamination [1,15-18]. The quality of dug well water for drinking and domestic use requires assessment to ensure compliance with drinking water quality index, DWQI [19]. Several diseases, especially those capable of originating diarrheal diseases are associated with unsanitary water sources [20,21]. The study aimed to determine the bacteriological quality of well water used for drinking and domestic purposes in Owo Metropolis, Ondo State, Nigeria.

## Materials and Methods:-

### Study design:

Cross-sectional study of the bacteriological quality of well water used for domestic purposes in Owo Metropolis, Ondo State, Nigeria.

### Collection of samples:

The tap source for each covered well water was washed wearing sterile gloves, allowed to flow for 30 sec and with sterile 250 mL bottle, approximately 200 mL of water was collected and kept in cold chamber for transportation to the laboratory for examination within 2 h.

### Examination of samples:

Physical visual assessment of samples for turbidity was determined and measurement of pH with a meter (Ohaus® ST20) and values noted.

Bottles of 5 x 10 mL double-strength McConkey, MCC broth (Oxoid CM 0005) containing Durham tubes inoculated with 10 mL of water aseptically after discarding a third of the sample, single-strength MCC broth of 5 x 5 mL with 1.0 mL, and 5 x 5 mL with 0.1 mL respectively. Control tests involved a set of broth in each category inoculated with sterile water. Thereafter, the trapped gas in each tube was expelled prior incubation at 37°C for 24 h. All negative bottles re-incubated for further examination at 48 h. The production of acid and gas indicated a presumptive positive test. The most probable number, MPN of coliform bacteria present in each sample was obtained using the McCrady's Probability Table for poor water source. Subsequently, sub-cultured onto MCC agar (CM 0007) and Salmonella-Shigella agar, SSA, (Oxoid CM 0099) incubated at 37°C for 24 h. Each growth was picked up for characterization tests and identified by the scheme outlined by Cowan [22]. The double-disc synergy test, (amoxicillin-clavulante 30 µg plus ceftazidime 30 µg combination) was applied for the detection of extended-spectrum beta-lactamase, ESBL. Susceptibility of isolates to antimicrobial agents was determined by the Kirby-Bauer disc diffusion technique with tetracycline (TE 10 µg), ampicillin (PN 10 µg), amoxicillin (A 10 µg), amoxicillin-clavulanate (AMC 30 µg), gentamycin (CN 10 µg), ciprofloxacin (CIP 10 µg), levofloxacin (LVX 10 µg), ceftazidime (CAZ 10 µg), cefotaxime (CTX 10 µg) and azithromycin (AZM 10 µg). The application of the Clinical Laboratory Standards Institute [23] guidelines assisted in the determination of isolates susceptibility.

## Results:-

Forty-seven representing 78.3% of 60 samples were turbid or had color and the pH of all samples ranged from 6.0-6.9. The least range of MPN in 100 mL sample was 15-100 in 22(36.7%) of 60 well water samples (Table 1). The MPN range 101-200/100 mL occurred in 16(26.7%), 201-300 in 8(18.3%), 301-400, 7(11.7%) and the highest MPN/100 mL, ≥401 occurred in 4(6.4%) of the samples.

The distribution of microorganisms recovered from the well water samples is shown in Table 2, with *Klebsiella* species 46(50%) of 92 isolates as the most frequent isolate made up of *Klebsiella pneumoniae* 31(33.7%) and *Klebsiella aerogenes* 15(16.3%). Other microorganisms recovered were *Enterobacter* species 23(25.0%) consisting of *Enterobacter cloacae* 17(18.5%) and *Enterobacter aerogenes* 6(6.5%), *Pseudomonas aeruginosa* 11(12.0%), *Proteus mirabilis* 8(8.7%) and *Serratiamarcescens* 4(4.4%) was the least isolated microorganism. Extended-spectrum beta lactamase, ESBLs detection was highest within isolates of *Klebsiella* species 9(19.6%). The detection rate of ESBLs in isolates of *Enterobacter* species was 17.4%, *Pseudomonas aeruginosa* 9.1% and *Proteus* species 12.5%. Isolates of *Serratiamarcescens* tested negative.

The susceptibility of isolates to antimicrobial agents (Table 3) shows the *Klebsiella* species and *Enterobacter* species were most susceptible to gentamycin, ciprofloxacin, levofloxacin, ceftazidime and cefotaxime at a range of 65.5-86.7%. The susceptibility of *Pseudomonas aeruginosa* was 36.4-54.5% for these same agents. *Proteus mirabilis* was most susceptible to cefotaxime (75%) and least to amoxicillin-clavulanate (12.5%). *Serratiamarcescens* was the only isolate that showed susceptibility to amoxicillin (25%) and susceptibility was highest with gentamycin (100%). No isolate showed susceptibility to either of tetracycline or ampicillin.

**Table 1:-** The most probable number (MPN) of coliform/100 mL of well water.

MPN Range	No. of Cases (%)
15-100	22(36.7)
101-200	16(26.7)
201-300	11(18.3)
301-400	7(11.7)
≥401	4(6.7)

**Table 2:-** Distribution of isolates and detection rate of ESBL within each genus.

Isolate	No. of Cases (%)	ESBL (%)
<i>Klebsiella</i> spp	46(50.0)	9(19.6)
<i>Enterobacter</i> spp	23(25.0)	4(17.4)
<i>Pseudomonas aeruginosa</i>	11(12.0)	1(9.1)
<i>Proteus mirabilis</i>	8(8.7)	1(12.5)
<i>Serratiamarcescens</i>	4(4.4)	0(0.0)

**Table 3:-** Susceptibility of isolates to antimicrobial agents.

Isolate	Antimicrobial agent (%)										
	TE	PN	A	AMC	CN	CIP	LVX	CAZ	CTX	AZM	
<i>Klebsiella</i> spp	0.0	0.0	0.0	0.0	71.7	69.6	71.7	65.2	67.4	47.2	
<i>Enterobacter</i> spp	0.0	0.0	0.0	4.3	73.3	86.9	65.2	73.9	43.5		
<i>Pseudomonas</i> spp	0.0	0.0	0.0	54.5	45.4	54.5	36.4	54.5	18.2		
<i>Proteus mirabilis</i>	0.0	0.0	12.5	75.0	50.0	50.0	65.5	75.0	75.0		
<i>Serratiamarcescens</i>	0.0	0.0	25.0	50.0	100	75.0	75.0	50.0	50.0	25.0	

## Discussion:-

The study revealed that 78.3% of well water samples were turbid, cloudy or colored as opposed to a clear and colorless appearance of potable water amongst other physical attributes recommended by the World Health [24]. Water turbidity is an indication that the water contains dissolved substances and particulate matter as a sign of microbiological pollution and poor quality [25]. The pH of the samples ranged from 6.5-6.9, all within recommended limits of 6.5-8.5 [24,26-28]. However, acidic drinking water can result in metallic taste due to dissolved chemicals and modulate microbiome constitution and alterations in function [29,30]. The least MPN or coliform organisms in the well water samples was 15/100 mL, this is much higher than the maximum for non-chlorinated water, thus inferring poor quality or pollution. These observations are consistent with reports from other studies [31,32]. Quite unpredictably, the isolated microorganisms did not include *Escherichia coli*, the classical marker of water pollution;

nevertheless, the very high number of coliform bacteria (MPN) recorded in each of the water samples make them of equal relevance. *Klebsiella* species was the most prevalent microorganism recovered, representing 50% of all isolates. This observation simulates reports from earlier studies [33,34], and involving markedly higher proportion of *Klebsiella pneumoniae* (33.7%) than other *Klebsiella* species (16.3%), indicating that *Klebsiella pneumoniae* was the most dominant species in these water sources [35]. *Klebsiella* species natural habitat includes fresh water sources, vegetation, and the gut of humans and animals, this interestingly, enhances the capacity of the microorganism to contaminate water easily in poor sanitary settings. This microorganism is associated with complicated urinary tract infections, UTIs and sepsis chiefly in individuals with pre-existing health challenges, posing major fears due to accumulative number of hyper-virulent and carbapenem-resistant phenotypes being reported globally [36,38]. *Enterobacter* species (25.0%) was the next common isolate - a fecal microorganism previously used in combination with *Escherichia coli* as marker organisms for fecal pollution of water, this denotes pollution of the water as other studies have indicated [27,39]. *Pseudomonas aeruginosa* occurred at a rate of 12.0%. The recovery of the microorganism from domestic water source points to a source of peril, mimicking reports from earlier studies [40,41], Water source burdened with *Pseudomonas aeruginosa* is consequentially an indication of water insecurity. This in part due to the microorganism's intrinsic resistance to many antimicrobial agents and a major initiator of nosocomial infections in clinical sceneries [42-45]. The isolation of *Proteus mirabilis* (8.7%) from the water samples infers that the microorganism as a regular contaminant in bodies of fresh water and signifies organic matter pollution of the wells [46-48]. The association of *Proteus* species with domestic water source inevitably portends danger for public health. The microorganism is one of the commonest in catheter-associated urinary tract infections, cUTI owing to the pathogenicity potential via the production of urease that breaks down urea in urine releasing ammonia leading to alkalization and precipitation of nephrolithiasis (or kidney stones) [49]. *Serratia marcescens* (4.4%) was the least frequent isolate. The habitat of this microorganism includes fresh water bodies, soil, insects and vegetation, and a regular environmental pollutant of fresh water, thus showing its presence in water as an index of poor water quality. Many *Serratia marcescens* isolates are multidrug resistant, MDR and linked to infections of opportunities, and as a consequence represents a dangerous water source or channel of health hazard [50,51]. This microorganism typically causes infections in persons with depressed immunity and in individuals with other clinical conditions [52]. In spite of higher susceptibility rates of the isolates to ciprofloxacin, levofloxacin, ceftazidime, cefotaxime and gentamycin (45.4-100%). All isolates showed resistance to tetracycline and ampicillin, this is in consonance with prior observations on susceptibility of the isolates to first-line antimicrobial agents [53]. The production of extended-spectrum beta-lactamase, ESBL was highest amongst *Klebsiella* species (19.7%) and *Serratia marcescens* isolates tested negative. *Klebsiella* species, *Enterobacter* species and *Pseudomonas aeruginosa* classified into the ESKAPE group of bacteria and included in the World Health Organization priority list of microorganisms resistant to a wide range of antimicrobial agents requiring extra precautionary measures in their infections [54]. This in addition, creates dire predicaments in making treatment decisions, hampering the ability to provide effective clinical care [35,52,55-58], and also the leading bacteria involved in hospital-acquired infections, HAIs [59]. The presence of ESKAPE and MDR bacteria in water negates the requirement for freedom of domestic water source from harmful microorganisms as a basic prerequisite for healthy living [60]. This study infers that the isolated microorganisms were regularly present in the environment of the wells as underground water does not support multiplication of bacteria [13]. Antimicrobial resistance, AMR has been a key challenge for humans, animals and the environment in the last half century requiring concerted control efforts and adoption of novel strategies towards the attainment of one health goal [61,62]. Insufficiency or failure to provide potable water in this community is the main driver for well water for home use [63] and unfortunately, these wells are dug by artesianians who possessed no professional competence, further exacerbating the risk of pollution [64]. The responsibility for the safety and or purity of private well water lies with the individual owner unlike public water systems with testing schedules and set standards [65].

### **Conclusion:-**

The study highlights insecure water source exemplified by poor physical and bacteriological properties of well water used for domestic purposes in Owo metropolis laden with ESKAPE member bacteria amongst others including MDR strains that can be contributory to the growing burden of disease and mortality rates in infections related to these microorganisms internationally. The provision of potable water as a right of all citizens in any nation requires the action of appropriate government agency in providing potable water in Owo and education on domestic wells specifications, safety testing and maintenance where potable or pipe-borne chlorinated water is impracticable.

**References:-**

1. Lusic DV, Maestro N, Cenov A, Lusic D, Smolcic K, Tolic S, et al. Occurrence of *Pseudomonas aeruginosa* in water intended for human consumption and in swimming pool water. *Environ*, 2021; 8(12): doi.org/10.3390/enviro/s8120312.
2. Some S, Mondal R, Mitra D, Jan D Verma D, Das S. Microbial pollution of water with special reference to coliform bacteria and their nexus with environment. *Energy Nexus*, 2021; 1(2020): 100008, doi.org/10.1016/j.nexus.2021.100008.
3. Rahman MM, Kunwar SB, Bahara AK. The interconnection between water quality level and studies: analysis of *Escherichia coli* contamination and drinking water from Nepal. *Water Res Economics*, 2021; 14: 100179.
4. Kurniawan V. Spring water as the water source for Cirebon Kanigan and Malalengka region. *IOP ConfSer: Mater SciEng*, 2019; 508: 012012, doi.10.1088/1757-899x/508/1/012012.
5. Djaouda M, Njime T, Liang S, Meraje DE, Gake B, Hubert S. Bacteriological quality of well water in Garoua North Cameroon. *Water Quality Exposure Health*, 2014; 6: 161-76.
6. Ojo OI, Otiero FA, Ochieng GM. Characteristics, qualities, pollutions and treatments: an overview. *Int J Water Res Environ Eng*, 2012; 4(6): 162-70.
7. Mahmoud N, Zayad O, Petrusevski B. Ground water quality of drinking water wells in the West Bank Palestine. *Water*, 2022; 14(3): 37, doi.org/10.3390/w14030377.
8. Hassan ON. Water quality parameters [Internet], Water quality, science assessments and policy. *Intech Open*, 2020; doi.org/10.5772/intechopen.89657.
9. Musa JJ, Akpoebidimiyen OE, Musa MT, Dala PO, Musa ET. Evaluation of temporal changes in deep well water quality in Igabi Local Government Area of Kaduna State. *J Environ Protection*, 2020; 11: 22-33.
10. Keerthen L, Ramya-Priya R, Elango L. Geogenic and anthropogenic contamination in river water and groundwater of the Lower Cauver Basin, India. *Front Environ Sci*, 2023; 11: doi.org/10.3390/fenvs.1001052.
11. Ramirez E, Robles E, Gonzalez ME, Martinez ME. Microbiological and physiochemical quality of well water used as a source of public supply. *Air Soil Water Res*, 2020; 3(1): doi.org/10.1177/ASWR.s4823.
12. Mile II, Jande JA, Dagba BI. Bacteriological contamination of well water in Makurdi town, Benue State, Nigeria. *Pat J BiolSci*, 2012; 15(2): 1048-51.
13. Ganpurev D, Gombosuren O, Andarai T, Battulga D, Amaarjargal I, Zboot B, et al. Microbiological and hydrological parameters of deep wells used for drinking water in Ulaatar, Mongolia. *Proceedings of the 5<sup>th</sup> International Conference on Chemical Investigation and Utilization of Natural Water Source (ICCUNR-2020)*. Atlantis Highlights Chem Pharm Sci, 2021: 2.
14. Nabeela F, Azzullah A, Bibi R, Uzma S Murad W, Shakir SK, et al. Microbiological contamination of drinking water in Pakistan: a review. *Environ Sci Res Int*, 2014 21(24): 13929-42.
15. Pathak SP, Gopal K. Prevalence of bacterial contamination with enterotoxigenic fecal coliforms in treated drinking water. *J Toxicol Environ Health A*, 2008; 71(7): 423-33.
16. Garba MB, Aliyu MS, Wada-Kaura A, Olonitola OS. Prevalence of *Escherichia coli* in some public water sources in Gusau Municipal, North Western Nigeria. *Bayero J Pure ApplSci*, 2009; 134-137.
17. Navab-Daneshmand T, Friedman MA, Gachter M, Nhiwatiwa T, Mosler HJ, Julian TR. *Escherichia coli* contamination across multiple environmental compartments: soil, hands, drinking water: correlation and risk factors. *Am J Trop Med Hyg*, 2018; 98(3): 803-13.
18. Obikpo L, Onyia FC, Offe IM, Ezeilo CM, Ezebilu C, Afunwa RA. Bacteriological quality of community well water and public health concerns in Enugu Urban, Nigeria. *Afri J ClinExperMicrobiol*, 2022; 23(2): 190-200.
19. Oluwagbayide SD, Abulude FO. Assessment of the quality of dug well water samples in Nigeria and their suitability for drinking and irrigation purposes. *Sustain Water Res Manag*, 2022; 8: 149, doi.org/10.1007/s40899.
20. Nazemi K, Saari S, Eskandani MA. Assessment of the *Escherichia coli* pollution in drinking water and water sources in Sistan. *Iran J Water Reuse Desalination*, 2018; 8(3): 386-92.
21. Odonkor ST, Mahami T. *Escherichia coli* as a tool for disease risk assessment of drinking water sources. *Int J Microbiol*, 2020; (2020): 2534130, doi.10.1155/2020/25344130.
22. Cowan ST. *Gowan and Steel's Manual for the identification of medical bacteria (2<sup>nd</sup> edn.)* Cambridge University press London, 1974: pp.1-180.
23. Clinical Laboratory Standards Institute. (30th ed), M100. CLSI, 2020; Wayne, PA, USA. pp 1-332.
24. World Health Organization. *Water quality and health - review of turbidity: Information for regulators and water suppliers*. WHO, 2017: WHO/FWC/WSH?17.01.

25. De-Roos AJ, Gurian PL, Robinson LF, Rai A, Zakari I, Kondo M. Review of epidemiological studies of drinking water turbidity in relation to acute gastrointestinal illness. *Environ Health Perspect*, 2019; 125(8): doi.10.1189/EPH1090
26. Dirisu CG, Mafiana MO, Dirisu GB, Amodu R. Level of pH drinking water of oil and gas producing community and perceived biological and health implications. *Eur J Basic Appl Sci*, 2016; 3(3): ISSN 2059-3058.
27. Adesakin TA, Oyewale AT, Barje IB, Assessment of bacteriological quality and physicochemical parameters of domestic water sources in Samaru Community, Zaria North-West, Nigeria. *Cell Press*, 2020; 6(8): e04773.
28. Ali AI, Sandi S, Sahara E, Rofiq MN, Dahlanuddin. Effects of acid drinking water on nutrient utilization, water balance and growth of goats under hot-humid tropical environment. *Small Ruminant Research*, 2022; 210: 106689, doi.org/10.1016/j.smallrumres.2022.106689.
29. Hansen TH, Thomasssen MT, Madsen ML, Kern T, Bak EG, kashani A. The effects of drinking water pH on the human gut microbiota and glucose regulation: results of a randomized controlled cross over intervention. *Sci Rep*, 2018; 8: 16626, doi.10.1038/s41598-018-34761-5.
30. Whipple B, Agar J, Zhao J, Pearce DA, Kovacs AD. The acidified drinking water-induced changes in the behavior and gut microbiota of wild-type mice depend on the acidification mode. *Sci Rep*, 2021; 2877, doi.org/10.1038/s41598-021-82570-0.
31. Niba RN, Nchang C. Bacteriological analysis of well water sources in the Banbui student residential area. *J Water Res Protect*, 2013; 5(11): doi.10.4236/jwrp.2013.511108
32. Alabi OS, Akintayo I, Odeyemi JS, Oloche JJ, Baballa CM, Nwimo, et al. Suboptimal bacteriological quality of household water in Municipal Ibadan, Nigeria. *Am J Trop Med Hyg*, 2024; 110(2): 346-55.
33. Barati A, Ghaderpour A, Chew LL, Bong CW, honk KL, Chai LC. Isolation and characterization of aquatic-borne *Klebsiella pneumoniae* from Tropical Estuaries. *Int J Environ Res Public Health*, 2016; 13(4): 426, doi.10.3390/ijerph13040426.
34. Araujo S Silva V, Dapkevicius ML, Martins A, Igrejas G, Poeta P. Comprehensive profiling of *Klebsiella* in surface water from Northern Portugal: understanding patterns in prevalence, antibiotic resistance and biofilm formation. *Water*, 2024; 16(9): doi.10.3390/w.16091257.
35. De-Boeck H, Miwanda B, Lunguya-Metila D, Muyembe, Tamfum J, Stobbeerngh E, et al. ESBL-positive *Enterobacteria* isolates in drinking water. *Emerg Infect Dis* 2012; 18(6): 1019-20.
36. Effah CY, Sun L, Liu S, Wu Y. *Klebsiella pneumoniae*: an increasing threat to public health. *Ann Clin Microbiol Antimicrob*, 2020; 19(1): doi.org/10.1186/s12941-019-0343-8.
37. Chang D, Sharma L, Dela-Cruz CS, Zhang D. Clinical epidemiology, risk factors and control strategies of *Klebsiella pneumoniae* infection. *Front Microbiol*, 2012; 12: doi.org/10.3389/fmicb.2021.750662.
38. Patel CB, Shanker R, Gupta VK, Upadhyay RS. Q-PCR based culture-independent environmental human pathogen in riverine systems and public water. *Front Microbiol*, 2016; 7: doi.org/10.3389/fmicb.2016.00172.
39. Bhumbla U, Majundar S, Jain S, Dalal AS. A study of isolation and identification of bacteria from lake water and around Udaipur. *J Fam Med Prim Care*, 2020; 9(2): 751-54.
40. Mena KD, Gerba. Risk assessment of *Pseudomonas aeruginosa* in water. *Rev Environ Contam Toxicol*, 2009; 201: 71-115.
41. Balfour-Lynn IM. Environmental risk of *Pseudomonas aeruginosa* - what to advice patients and parents. *J Cyst Fibros*, 2021; 20(1): 1724.
42. Labovska S. *Pseudomonas aeruginosa* as a cause of nosocomial infections. *Intech Open*, 2021; 2021: doi.10.5772/intechopen.95908.
43. Spagnolo AM, Sartini M, Cristina M, Maria L. *Pseudomonas aeruginosa* in the healthcare facility setting. *Rev Med Microbiol*, 2021; 32(3): 169-75.
44. Litwin A, Rojek S, Gozdzik W, Duszynska W. *Pseudomonas aeruginosa* device-associated healthcare infections and its intensive care unit of university hospital: Polish 8.5 years prospective study, single-centre study. *BMC Infect Dis*, 2021; 21(180): doi.org/10.1186/s12879-021-05883-5.
45. Volling C, Mataseje L, Grana-Miraglia L, McGueer A, Guttman DS, Mulvey MR. Epidemiology of healthcare associated *Pseudomonas aeruginosa* in intensive care units: are sinks, drains to blame? *J Hosp Infect*, 2024; 148: 77-86.
46. Drzewiecka D. Significance and roles of *Proteus* spp. Bacteria in natural environment. *Microbiol Ecol*, 2016; 72(2): 74158.
47. Otokpa OJ. Overview of major bacterial contaminants of drinking water in Nigeria: a review. *J Pathogen Res*, 2019; 2(3): 1-9.
48. Erkinovick NJ. Bacteria of the genus *Proteus* as sanitary indicative microorganisms of waterbodies, *J Environ Microbiol*, 2022; 4(1): 1-3.

49. Armbruster CE, Mobley HL, Pearson MM. Pathogenesis of *Proteus mirabilis* infection. *Ecosal Plus*, 2018; 8: doi.10.1128/ecosalplus.ESP-0009-2017.
50. Gadhiya GK, Golden J, PanupongH, Balchander D, Cook E, Enriquez K, Smith D. Severe skin infections due to *Serratiamarcescens*: a case with cat scratch in a patient with liver disease and review of the literature. *Infect Dis ClinPract*, 2021; 29(3): e146-50.
51. Khayat MT, Elbaramawi SS, Nazeih SI, Safo MK, KhafagyESAli MA, et al. Diminishing the pathogenesis of foodborne pathogen *Serratiamarcescens* by low doses of sodium citrate. *Biology*, 2023; 12(4): 504, doi.10.3390/biology1240504.
52. Cristinna ML, Sartini M, Spagnolo AM. *Serratiamarcescens* infections in neonatal intensive care units (NICUs). *Int J Environ Res Public Health*, 2019; 16(4): 610, doi.10.3390/ijerph16040610.
53. Denissen J, Reyneke B, Barnard T, Khan S, Khan W. Risk assessment of *Enterococcus faecum*, *Klebsiella pneumoniae* and *Pseudomonas aeruginosa* in environmental water sources, development for antibiotic resistance genes. *Sci Total Environ*, 2023 901: 166217, doi.org/10.1016/j.scitoenv.2023.166217.
54. Nisa TT, Nakatani D, Kaneko F, Takeda T, Nakata K. Antimicrobial resistance patterns of WHO priority pathogens isolated in hospitalized patients in Japan: a tertiary centre observational study. *PLoS ONE*, 2024; 19(1): e0294229.
55. Cabral JP. Water microbiology, bacterial pathogens and water. *Int J Environ Res Public Health*, 2010; 7(10): 3657-703.
56. Asem SS, Lekshmi M, Sreepriya P, Binaya BN, Kumar S. Multiple antibiotic resistant extended-spectrum  $\beta$ -lactamase (ESBL)-producing *Enterobacteria* in fresh seafood. *Microorganisms*, 2017; 5(3): 53, doi.10.3390/microorganisms5030053.
57. De-Oliveira DM, Forde BM, Kidd TJ, Harris PN, Schembri MA, Beatson SA, et al. Antimicrobial resistance in ESKAPE pathogens. *ClinMicrobiol Rev*, 2020; 33(3): e00181-19.
58. Dhengesu D, Lemma H, Asefa L, Tilahun D. Antimicrobial resistance profile of *Enterobacteriaceae* and drinking water quality among households in Blue Thora Town, South Ethiopia. *Risk Manag Healthcare Policy*, 2022; 15: 1569-80
59. Alvarez-Ainza ML, Fong-Coronado PA, Ruiz\_Bustos E, Castillon-Campana LG, Quintero-Reyes IE, Duarte\_Zambrano LA, et al. Antibiotic resistance of ESKAPE group-microorganisms in health institutions from Hermosillo and Ciudad Obregon, Sonora, Mexico. *Front Cell Infect Microbiol*, 2024; 14: doi.org/10.3389/fcimb.2024.1348093.
60. Ahmad M, Jamal A, Tang XW, Al-Sughaiyer MA, Al-Ahmadi HM, Ahmad F. Assessing potable water quality and identifying areas of water-borne diarrheal and fluorosis health risk using spatial interpretation on Peshawar, Pakistan. *Water*, 2020; 12(8): doi.10.3390/w.12082163.
61. Jin L, Pruden A, Boechm AB, Alvarez PJ, Kohn T, Li X. Integrating environmental dimension of one health to resistance: essential research needs. *Environ SciTechnol*, 2022; 56(2): 14871-74.
62. Serna C, Gonzalez-Zorn B. Antimicrobial resistance and one health. *Rev EspQuimioter*, 2024; 35(Suppl 3): 37-40.
63. Auta KI, Mohammed SS, Abubakar MI. Assessment of bacteriological quality of well water around Dogon-Dawa Disrict in Binin-Gwari Local Government, Kaduna. *Sci World J*, 2017; 12(4): scinceworldjournal.org/ISSN1597-6343.
64. Bassey CB, Ogah TA, Magaji JI, Oladeinde OS. The suitability of well water for domestic purpose in Gwagwalada Area Council, Abuja, Nigeria. *Global J Pure ApplSci*, 2021; 27(2): doi.10.4314/gjpas.v27i2.7.
65. Farrel-Poe K, Jones-McLean L, McLean S. Well water testing and understanding the results. The University of Arizona College of Agriculture and Life Science, 2011; AZ1486f