

## Antimicrobial Susceptibility Profile Of Bacteria Isolated From Raw Sewage At A Wastewater Treatment Plant In Rio De Janeiro

### ABSTRACT

**Background and Aim:** Given the concentration of microorganisms and emerging micropollutants, wastewater is a favorable environment for the exchange of resistance genes and the selection of antimicrobial-resistant strains, one of the most significant global threats of the 21st century, since inadequate environmental sanitation causes around 88% of deaths worldwide. The present study aimed to identify bacteria isolated from a sewage treatment plant located in Rio de Janeiro and determine their susceptibility profile to antimicrobials.

**Methods:** A 10 µL aliquot of raw sewage was inoculated onto chromogenic agar and presumptive identification of microorganisms were based on their biochemical profiles. Isolates that could not be characterized using traditional biochemical tests were analyzed using polymerase chain reaction (PCR). Following the Brazilian Committee on Antimicrobial Susceptibility Testing, the resistance profile was determined by measuring the diameter of the inhibition zone and the isolates were classified as susceptible (S), resistant (R), multidrug resistant (MDR), extensively resistant (XDR) or pan-resistant (PDR).

**Results:** In this study, 38 strains were isolated, of which 24% were identified as Gram positive (n = 09) and 76% as Gram negative (n = 29). Among the isolated strains, *Enterobacter* sp. (n = 7, 18%) was the most frequent genera and *Escherichia coli* (n = 6, 15%) and *Klebsiella pneumoniae* (n = 5, 13%) were the most frequent species. No pan-resistant strains were identified, however, isolates of *Klebsiella pneumoniae* (60%, n = 3/5), *Enterobacter* sp. (57%, n = 4/7) and *Escherichia coli* (50%, n = 3/6) presented the profile MDR and only one *Enterobacter* isolate was considered XDR (14%). Isolates of *Morganella morgani*, *Proteus mirabilis*, and *Acinetobacter baumannii* were fully sensitive to the antimicrobials tested. Regarding the antimicrobial susceptibility of Gram-positive strains, the *Bacillus* sp., *Enterococcus* sp. and *Staphylococcus aureus* isolates showed different resistance profiles, in which only *Bacillus* sp. and *Microbacterium paraoxydans* were considered high risk to health.

**Conclusion:** The occurrence of potentially pathogenic and antimicrobial-resistant microorganisms in sewage treatment plants is a reality that has been explored over the years. Resistance to beta-lactam drugs is increasingly present in clinical practice, and these were the antimicrobials with the highest number of resistant isolates in the present study, including *Escherichia coli*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa* and *Enterobacter* sp., which are listed as WHO priority. Contact of these microorganisms with the community may cause not only public health problems, but also environmental impacts, increasing the need to implement and improve environmental surveillance of resistant pathogens, establishing stricter control over the use of antimicrobials, to try to mitigate the advance of bacterial resistance.

**Keywords:** Antimicrobial resistance; Enterobacterales; Residual waters; Sewage.

Commented [n1]: 9

Commented [n2]: 7 (18%)

Commented [n3]: 6(15%)

## 1. INTRODUCTION

The integrity of human, animal and environmental health is directly associated with basic sanitation conditions and water quality [1]. Diseases related to inadequate environmental sanitation are responsible for approximately 88% of global deaths, most of which are transmitted by different aquatic matrices [2]. Wastewater, also called sewage, resulting from anthropogenic activity, poses a risk of transmission of pathogenic microorganisms and, mainly, those resistant to antimicrobials [1,3]. In essence, Antimicrobial Resistance Bacteria (ARB), as well as Antimicrobial Resistance Genes (ARG), are considered a global public health problem that tends to increase, especially after the COVID-19 pandemic [4-6].

**Commented [n4]:** A short indent at the beginning of each topic

Approximately 68 million units of antibiotics were sold in Brazil between 2020 and 2023, which may lead to an increase in cases of infections caused by multidrug-resistant or pan-resistant bacteria [6]. The use of these drugs implies the elimination of residues in aquatic matrices, concentrating such substances in different compartments, including sewage system. Considering that effluent treatments currently used are not capable of eliminating these environmental contaminants, the sewage system becomes capable of amplifying the spread of ARBs and ARGs [3,7]. Based on this, the present study aimed to identify bacteria isolated from a sewage treatment plant located in Rio de Janeiro and determine their susceptibility to antimicrobials.

## 2. METHODS

The methodology proposed in this pilot study consisted of collecting 500 mL of raw sewage, called influent, in sterile graduated bottles. Following all biosafety protocols, the collection was performed by the sewage treatment plant operator, allowing photographic recording of the collection process. The bottle was then properly labeled according to origin, time, type of collection and type of sample. To avoid bacterial competition, transportation took place in an isothermal box, as recommended in the Standard Methods for Examination of Water and Wastewater.

All experiments were carried out at the Microbiology Laboratory (LabMicro), Department of Biological Sciences, Sérgio Arouca National School of Public Health (ENSP), located at the Center for Research, Innovation and Surveillance in Covid-19 and Health Emergencies of the Oswaldo Cruz Foundation (FIOCRUZ).

### 2.1 Bacterial identification

To obtain isolated bacterial colonies, the initial sample was diluted in sterile saline (0.9% NaCl) with different concentrations, ranging from  $10^{-3}$  to  $10^{-7}$  CFU. 10 $\mu$ L of the sample was seeded on chromogenic agar, allowing presumptive differentiation of the microorganisms. After seeding, the plates were incubated in an oven at  $35\pm 1^{\circ}\text{C}$  for 24 hours for bacterial growth. The colonies of interest were replated and identified by Gram staining to confirm the morphotintorial characteristics of the bacteria. Biochemical screening tests, such as Triple Sugar Iron Medium (TSI), Sulfide Indole Motility Medium (SIM), Simmons Citrate and Costa and Vernin Medium (CV) were used for presumptive identification.

**Commented [n5]:** Why used the oven?  
Bacterial culture must incubated in incubator

**Commented [n6]:** Morphological characteristics

In order to identify isolates that could not be characterized using traditional biochemical tests, polymerase chain reaction (PCR) analysis was performed, in which specific oligonucleotide primers for the 16S rRNA gene were amplified using the 16S rRNA full gene PCR Kit (Life Technologies) and the sequences obtained were compared based on similarity with the GenBank database. The DNA extraction protocol used was performed as described by Campos *et al.*<sup>8</sup> at the National Reference Laboratory for Tuberculosis, Professor Hélio Fraga Reference Center, Sérgio Arouca National School of Public Health, Fiocruz.

**Commented [n7]:** italic

## 2.2 Antimicrobial Sensitivity Test (AST)

To perform the sensitivity test, agar diffusion technique was chosen, in which a bacterial suspension in 0.9% saline, corresponding to the 0.5 MacFarland scale, is seeded on Mueller-Hinton agar with the aid of a sterile swab. The antimicrobial discs were placed on agar and the plates were incubated in a bacteriological incubator at 37°C for 24 hours. Following the recommendations of the Brazilian Committee on Antimicrobial Susceptibility Testing (BRCAST), the isolates were tested for the following antimicrobials: amikacin, amoxicillin-clavulanate, ampicillin, cefepime, ceftioxin, ceftriaxone, ciprofloxacin, ertapenem, gentamicin, imipenem, meropenem, penicillin, and piperacycline-tazobactam. The sensitivity or resistance profile was determined by measuring the diameter of the inhibition zone and the isolates were classified as susceptible (S) when they did not present resistance, resistant (R) when they presented resistance to only one class of antimicrobials tested, multidrug resistant (MDR) when they were resistant to up to 3 classes, extensively resistant (XDR) when they were resistant to more than 3 classes, or pan-resistant (PDR) when they did not present susceptibility to any antimicrobial tested (BRCAST, 2024) [9].

Commented [n8]: not technique it is method

Commented [n9]: antimicrobial agents

Commented [n10]: insert the dose each of antibiotics

## 2.3 MAR Index

To determine the health risk, the Multiple Antibiotic Resistance Index (MAR Index) was calculated [10,11], using the following formula:

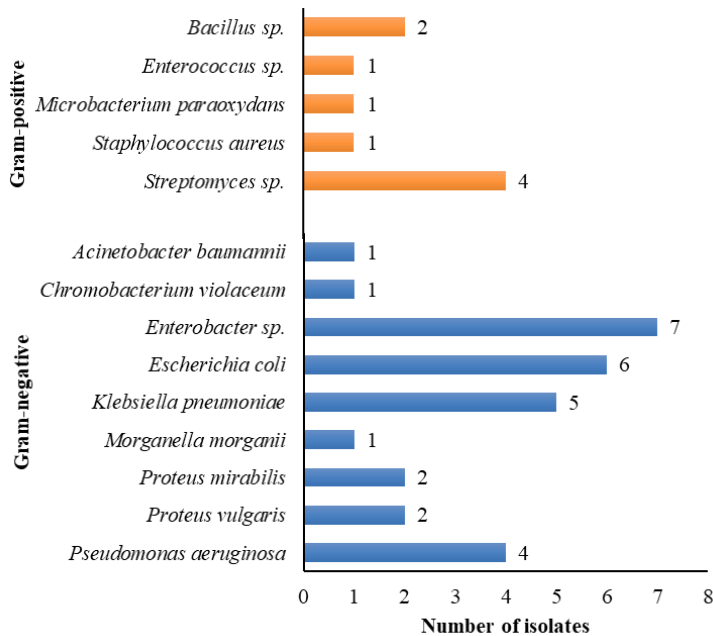
$$\text{MAR Index} = n^{\circ} \text{ ineffective antibiotics} / n^{\circ} \text{ antibiotics tested}$$

Following the interpretation proposed by Zagui (2022), intermediate resistance was considered sensitive and results equal to or greater than 0.2 were considered high risk [10].

Commented [n11]: What is the unit of measurement used?

## 3. RESULTS AND DISCUSSION

Given the concentration of microorganisms and emerging micropollutants, wastewater is a favorable environment for the exchange of resistance genes and the selection of antimicrobial-resistant strains, one of the most significant global threats of the 21st century [12,13]. In the present study, 38 bacterial strains were isolated, of which 24% were identified as Gram positive (n = 09), being 02 cocci and 07 bacilli, and 76% as Gram negative bacilli (n = 29). Only 5 isolates were considered non-fermenters. Among the isolated strains, the most frequent genera were *Enterobacter* sp. (n = 7, 18%), *Escherichia* sp. (n = 6, 15%), *Klebsiella* sp. (n = 5, 12.5%) and *Streptomyces* sp. (n = 4, 10%). The most frequent species were *Escherichia coli* (n = 6, 15%) and *Klebsiella pneumoniae* (n = 5, 13%), as shown in Fig. 1.



**Fig. 1. Total number of isolates identified in raw sewage from a wastewater treatment plant in Rio de Janeiro.**

Source: Created by the authors (2024).

The most frequent isolated species, *Escherichia coli* and *Klebsiella pneumoniae*, can be explained by their presence in the intestinal microbiota of humans<sup>14</sup> while other bacterial isolates, such as *Proteus mirabilis*, *Proteus vulgaris* and *Pseudomonas aeruginosa*, are frequently found in wastewater due to their ability to adapt and tolerate the environment [15,16].

First described in 1872, *Chromobacterium violaceum* is a tropical pathogen that can be commonly found in soil and water bodies in tropical and subtropical regions [17] and usually transmitted through exposure of skin wounds and traumatic injuries to contaminated soil and water [18-21]. Its zoonotic potential makes harmful to humans and animals, given its ability to cause abscesses in the skin and organs, such as the liver, lungs, kidneys, spleen, and lymph nodes, as well as septicemia with necrotizing lesions in multiple organs, resulting in the death of the patient [17,22]. However, the isolate was not subjected to susceptibility testing because it was not viable for growth.

Regarding the antimicrobial susceptibility of Gram-positive strains, the *Bacillus sp.*, *Enterococcus sp.* and *Staphylococcus aureus* isolates tested showed different resistance profiles, in which only *Bacillus sp.* and *Microbacterium paraoxydans* were considered high risk (Table 1). Resistant strains of *Bacillus sp.* have also been reported in other studies [13,23], representing a relevant environmental risk, given the possibility that horizontal gene transfer favors the spread of antibiotic resistance not only among different *Bacillus* species, but mainly among other clinical pathogens, such as *Enterococcus sp.* and *Staphylococcus aureus* [24]. Although they present intermediate resistance to ampicillin and ciprofloxacin,

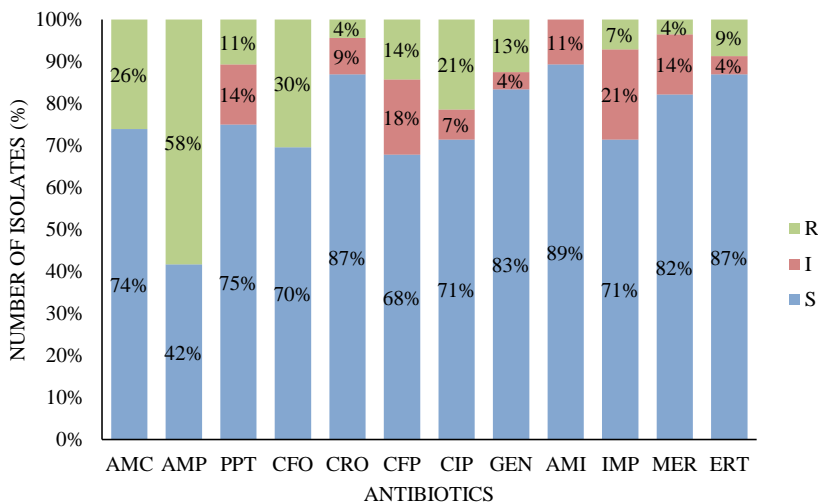


	<i>sp.</i>													
EB3B	<i>Proteus mirabilis</i>	S	S	S	S	S	S	S	S	S	S	S	S	0,00
EB3C	<i>Escherichia coli</i>	S	S	S	S	S	S	S	S	S	S	S	S	0,00
EB3C	<i>Pseudomonas aeruginosa</i>	NA	NA	I	NA	NA	I	R	NA	S	I	S	N	0,20
EB3D	<i>Morganella morganii</i>	S	S	S	S	S	S	S	S	S	S	S	S	0,00
EB3E	<i>Klebsiella pneumoniae</i>	S	R	S	R	S	S	R	S	S	I	S	S	0,25
EB3F	<i>Pseudomonas aeruginosa</i>	NA	NA	I	NA	NA	I	R	NA	I	I	I	N	0,17
EB3G	<i>Proteus mirabilis</i>	S	S	S	S	S	S	S	S	S	S	S	S	0,00
EB4B	<i>Klebsiella pneumoniae</i>	S	R	S	S	S	S	S	S	S	S	S	S	0,08
EB4C	<i>Klebsiella pneumoniae</i>	S	S	S	S	S	S	S	S	S	S	S	S	0,00
EB5A	<i>Proteus vulgaris</i>	S	R	S	S	S	S	S	S	S	S	S	S	0,08
EB5B	<i>Enterobacter sp.</i>	S	S	S	S	S	S	S	S	S	S	R	R	0,17
EB6A	<i>Escherichia coli</i>	S	R	S	S	S	S	S	S	S	I	S	S	0,08
EB6B	<i>Enterobacter sp.</i>	R	R	S	R	S	R	S	S	S	S	S	S	0,33
EB6C	<i>Klebsiella pneumoniae</i>	R	R	R	R	R	R	S	S	S	S	S	S	0,50
EB7A	<i>Proteus vulgaris</i>	S	R	S	S	S	S	S	S	S	S	S	S	0,08
EB7E	<i>Escherichia coli</i>	S	S	S	S	S	S	S	S	S	S	S	S	0,00
EB7F	<i>Enterobacter sp.</i>	S	R	R	S	S	S	R	R	S	R	S	S	0,42
EB7G	<i>Escherichia coli</i>	S	R	S	S	S	I	R	R	S	S	S	S	0,25
EB8A	<i>Enterobacter sp.</i>	S	R	S	S	S	S	S	I	S	S	S	S	0,08
EB8B	<i>Pseudomonas aeruginosa</i>	NA	NA	I	NA	NA	I	I	NA	I	I	S	N	0,00
EB9B	<i>Enterobacter sp.</i>	R	S	S	R	S	S	S	S	S	S	S	S	0,17
EB11	<i>Escherichia coli</i>	R	R	R	S	S	S	S	R	S	S	S	S	0,33
EB11	<i>Pseudomonas aeruginosa</i>	NA	NA	I	NA	NA	I	R	NA	I	I	I	N	0,17
EB11	<i>Enterobacter sp.</i>	S	R	S	R	S	S	S	S	S	S	S	S	0,17
EB11	<i>Klebsiella pneumoniae</i>	R	R	S	R	I	S	S	S	S	R	I	I	0,33

Legend: AMC – Amoxicillin-Clavulanate; AMP – Ampicillin; PPT – Piperacilina-Tazobactam; CFO – Cefoxitin; CTX – Ceftriaxone; CFP – Cefepime; CIP – Ciprofloxacin; GEN – Gentamicin;

AMI - Amikacin; IMP – Imipenem; MER – Meropenem; ERT – Ertapenem; NA – Not applicable; R – Resistant; I – Intermediate; S – Sensitive. The authors (2024).

The EB6C strain stands out due to its resistance to various beta-lactams, being classified as an extended-spectrum beta-lactamase (ESBL) *Klebsiella pneumoniae*, which confers resistance to a wide range of antibiotics, hindering the treatment of infections and increasing the risk of dissemination [31,32]. Being capable of causing a series of infections in hospitalized patients, with the advancement of antibiotic therapy, resistant strains of *K. pneumoniae* are increasingly prevalent, favoring the dissemination of different resistance genes, especially in the hospital environment, being considered one of the greatest challenges for global public health [31,33]. According to Scarpate and Cossatis (2009), the prevalence of ESBL-positive *K. pneumoniae* in Latin America is higher than the world average (20-30%) [33], which may be directly related to the widespread and indiscriminate use of third-generation cephalosporins, such as ceftriaxone, and carbapenems, in which meropenem is considered a last resort treatment, disseminating this multidrug-resistant microorganism to the community, making clinical management of infections difficult [32-37]. Among the antimicrobials tested, beta-lactams showed the highest resistance rates, including penicillins, such as ampicillin (58%, n = 14/24), ceftioxin (30%, n = 7/23), amoxicillin associated with potassium clavulanate (26%, n = 6/23) (Fig. 2).



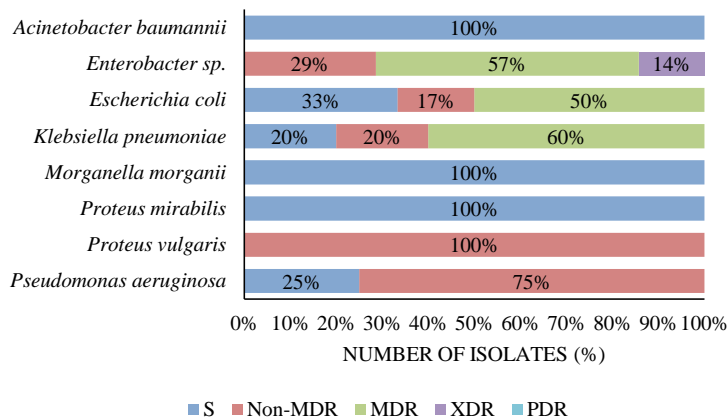
**Fig. 2. Strains resistant to the tested antibiotics isolated from a sewage treatment plant in Rio de Janeiro.**

Legend: AMC – Amoxicillin-Clavulanate; AMP – Ampicillin; PPT – Piperaciline-Tazobactam; CFO – Cefoxitin; CTX – Ceftriaxone; CFP - Cefepime; CIP – Ciprofloxacin; GEN – Gentamicin; AMI - Amikacin; IMP – Imipenem; MER – Meropenem; ERT – Ertapenem; R – Resistant; I – Intermediate; S – Sensitive. Source: Created by the authors (2024).

Among the carbapenems tested, 9% of the strains tested were resistant to ertapenem (n = 2/23), 7% were resistant to imipenem (n = 2/28) and only 4% were resistant to meropenem (n = 1/28), which may be linked to the fact that carbapenems are the drugs of choice for the treatment of ESBL-positive strains, in which meropenem is considered a last resort treatment [36,37]. Of the aminoglycosides tested, only cases of resistance to gentamicin were identified (13%, n = 3/24). Resistance to aminoglycosides may originate from cross-

resistance, given that amikacin is a substrate for only a few aminoglycosidases, bacterial enzymes capable of modifying the structure of aminoglycosides, making strains resistant to other aminoglycosides, such as gentamicin, likely to be susceptible to amikacin [38,39]. This statement corroborates the results found in the present study, in which, only cases of resistance to gentamicin were identified, a drug primarily recommended in cases of severe infections, but which, due to its low cost, began to be used indiscriminately, favoring the selection of resistant strains [39].

Furthermore, no pan-resistant strains were identified. However, isolates of *Klebsiella* sp. (60%, n = 3/5), *Enterobacter* sp. (57%, n = 4/7), and *Escherichia* sp. (50%, n = 3/6) presented the MDR profile, and only one *Enterobacter* isolate was considered XDR (14%). Isolates of *Morganella morganii*, *Proteus mirabilis*, and *Acinetobacter baumannii* were fully sensitive to the antimicrobials tested (Fig. 3).



**Fig. 3. Resistance profile of Gram-negative rods isolated from sewage treatment plant in Rio de Janeiro.**

Legend: S – Sensitive; MDR – Multidrug resistant; XDR – Extensively resistant; PDR – Pan-resistant. Source: Created by the authors (2024).

Even though pan-resistant strains were not identified in this research, it is important to highlight that *A. baumannii* has intrinsic resistance to some antimicrobials and a greater tendency to acquire resistance genes from the environment [40]. Further analyses are necessary to obtain more robust data. However, the results obtained corroborate the study by Abrantes (2022), in which the author states that bacteria of the order Enterobacteriales have a high profile of non-sensitivity to antimicrobials, including strains whose resistance occurs through exposure (intermediate), mainly to beta-lactams, including cephalosporins and carbapenems, and quinolones [41]. Based on these findings, it is crucial to highlight the role of effluents in the dissemination of antimicrobial-resistant pathogens, the importance of creating epidemiological surveillance networks in these matrices, implement effective methodologies for the removal and inactivation of ARBs and/or ARGs, helping to reduce risks to human, animal and environmental health and mitigate the advance of bacterial resistance.

#### 4. CONCLUSION

Although these are preliminary results, it allows us to identify the role of effluents in bacterial resistance, with emphasis on *Bacillus* sp. and, mainly, Gram-negatives, including species

such as *Escherichia coli*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa* and *Enterobacter* sp., which are listed as WHO priority. Of the isolates identified, only 8 may have originated from an environment with no risk to health, in which there was no direct contact with drugs, given their sensitivity to the antimicrobials tested. However, resistance to beta-lactam drugs, such as ampicillin, amoxicillin and some cephalosporins, is increasingly present in clinical practice, and these were the antimicrobials with the highest number of resistant isolates in the present study.

The occurrence of potentially pathogenic and antimicrobial-resistant microorganisms in sewage treatment plant is a reality that has been explored over the years. Bacteria carrying antimicrobial resistance genes can be easily disseminated through waterways. Contact of these microorganisms with community may cause not only public health problems, but also environmental impacts, increasing the need to implement and improve environmental surveillance of resistant pathogens, establishing stricter control over the use of antimicrobials, in order to try to mitigate the advance of bacterial resistance.

## REFERENCES

1. Monteiro, M. et al. Occurrence of Antimicrobials in River Water Samples from Rural Region of the State of Rio de Janeiro, Brazil. *Journal of Environmental Protection*, 2016; 7(2): 230–241.
2. Monteiro, G. et al. Impactos na saúde pública pós-implantação de sistema integrado de saneamento rural: resultados no SISAR Moxotó em Pernambuco. *Hygeia - Revista Brasileira de Geografia Médica e da Saúde*, 2024; 20:e2059–e2059.
3. Joseph, S. M. et al. Longitudinal Comparison of Bacterial Diversity and Antibiotic Resistance Genes in New York City Sewage. *mSystems*, 2019; 4(4).
4. Amarasiri, M.; Sano, D.; Suzuki, S. Understanding human health risks caused by antibiotic resistant bacteria (ARB) and antibiotic resistance genes (ARG) in water environments: Current knowledge and questions to be answered. *Critical Reviews in Environmental Science and Technology*, 2020; 50(19):2016–2059.
5. Silva, L. O. P.; Abrantes, J. A.; Nogueira, J. M. da R. O uso indiscriminado de antimicrobianos durante a pandemia da COVID-19 como possível fator influenciador da resistência bacteriana em efluentes hospitalares no Brasil. *Contribuciones A Las Ciencias Sociales*, 2024; 17(3): e4360–e4360.
6. Wang, J. et al. Risk control of antibiotics, antibiotic resistance genes (ARGs) and antibiotic resistant bacteria (ARB) during sewage sludge treatment and disposal: A review. *Science of The Total Environment*, 2023; 877:162772.
7. Manaia, C. M. et al. Antibiotic resistance in wastewater treatment plants: Tackling the black box. *Environment International*, 2018; 115:312–324.
8. Campos, C.E.D. et al. First isolation of *Mycobacterium kyorinense* from clinical specimens in Brazil. *Journal of clinical microbiology* 2012; 50(7):2477-2478.
9. BRCAS. *Tabelas de pontos de corte para interpretação de CIMs e diâmetros de halos*, 2024.
10. Zagui, G. S. *Antibióticos e bactérias multirresistentes em esgoto e águas superficiais receptoras: riscos da propagação de genes codificadores de ESBL, de carbapenemases e de tolerância aos metais no contexto da saúde única*. 2022. text – Universidade de São Paulo, 2022.
11. Afunwa, R. A. et al. Multiple Antibiotic Resistant Index of Gram-Negative Bacteria from Bird Droppings in Two Commercial Poultry in Enugu, Nigeria. *Open Journal of Medical Microbiology* 2020; 10(4):171-181.

Commented [n13]: Mention the full names of the researchers

12. Fröhler, H. S.; Guimarães, R. E.; Frazzon, A. P. G. Avaliação da existência de resistência bacteriana em estação de tratamento de esgoto. In: *Congresso Brasileiro de Engenharia Sanitária e Ambiental*, 2023.
13. Patra, M.; Pandey, B.; Dubey, S. K. Prevalence of diverse antimicrobial resistance genes and bacteria in sewage treatment plant-derived sludge environment. *FEMS Microbes*, 2024; 5:xtae004.
14. Mills, M. C.; Lee, J. The threat of carbapenem-resistant bacteria in the environment: Evidence of widespread contamination of reservoirs at a global scale. *Environmental Pollution*, 2019; 255(1): 113143.
15. Araújo, S. et al. Carbapenem-resistant bacteria over a wastewater treatment process: Carbapenem-resistant Enterobacteriaceae in untreated wastewater and intrinsically-resistant bacteria in final effluent. *Science of The Total Environment*, 2021; 782:146892.
16. Zhang, L. et al. The Prevalence and Characterization of Extended-Spectrum  $\beta$ -Lactamase- and Carbapenemase-Producing Bacteria from Hospital Sewage, Treated Effluents and Receiving Rivers. *International Journal of Environmental Research and Public Health*, 2020; 17(4):1183.
17. Hota, S. R. et al. Characterization and Whole Genome Sequencing of Chromobacterium violaceum Ouat\_2017: A Zoonotic Pathogen Found Fatal to a Wild Asiatic Elephant. *Indian Journal of Microbiology*, 2022; 62(4):627-633.
18. Ansari, S. et al. Chromobacterium violaceum Isolated from a Wound Sepsis: A Case Study from Nepal. *Case Reports in Infectious Diseases*, 2015; 2015:1-4.
19. Baker, S. et al. Fatal Wound Infection Caused by Chromobacterium violaceum in Ho Chi Minh City, Vietnam. *Journal of Clinical Microbiology*, 2008; 46(11):3853-3855.
20. Chowdhury, M.; Lee, N.; Wey, E. Q. Chromobacterium violaceum causing disseminated soft tissue and pulmonary abscesses in a traveller returning from the Azores. *Access Microbiology*, 2021; 3(8).
21. Menezes, C. B. A. et al. Chromobacterium amazonense sp. nov. isolated from water samples from the Rio Negro, Amazon, Brazil. *Antonie van Leeuwenhoek*, 2015; 107(4):1057-1063.
22. Soares, R. L. et al. Chromobacteriosis (Chromobacterium violaceum) in a calf from Brazil - case report. *Arquivo Brasileiro de Medicina Veterinária e Zootecnia* 2019; 71(6):1929-1933.
23. Kalra, S.; Bhatnagar, A.; Thakral, N. Bacillus aryabhatai: A Multi Metal Resistant Sewage Water Bacteria and Bioremediatory Tool for Sewage Water Pollutants. *Egyptian Journal of Aquatic Biology and Fisheries*, 2024; 28(3):329-340.
24. Zhai, Z. et al. Prevalence, antimicrobial susceptibility, and antibiotic resistance gene transfer of Bacillus strains isolated from pasteurized milk. *Journal of Dairy Science*, 2023; 106 (1):75-83.
25. CLSI. *Normas de Desempenho para Testes de Sensibilidade Antimicrobiana: 15o Suplemento Informativo*. [S. l.]: Clinical and Laboratory Standards Institute, 2005. Disponível em: [https://bvsm.s.saude.gov.br/bvs/publicacoes/metodo\\_ref\\_testes\\_diluicao\\_modulo4.pdf](https://bvsm.s.saude.gov.br/bvs/publicacoes/metodo_ref_testes_diluicao_modulo4.pdf).
26. Amano, J. et al. Catheter-related bloodstream infection by Microbacterium paraoxydans in a pediatric patient with B-cell precursor acute lymphocytic leukemia: A case report and review of literature on Microbacterium bacteremia. *Journal of Infection and Chemotherapy*, 2019; 25(10):806-810.
27. Bernard, K.; Pacheco, A. L. In Vitro Activity of 22 Antimicrobial Agents against Corynebacterium and Microbacterium Species Referred to the Canadian National Microbiology Laboratory. *Clinical Microbiology Newsletter*, 2015; 37(23):187-198.

28. Ko, K. S. et al. A new *Microbacterium* species isolated from the blood of a patient with fever: *Microbacterium pyrexiae* sp. nov. *Diagnostic Microbiology and Infectious Disease*, 2007; 57(4):393-397.
29. Pinchman, E. et al. *Acinetobacter radioresistens* and *Microbacterium paraoxydans* endocarditis in patient with indwelling catheter and metastatic carcinoma. *BMJ Case Reports*, 2023; 16(6):e254877.
30. WHO. *WHO bacterial priority pathogens list, 2024: Bacterial pathogens of public health importance to guide research, development and strategies to prevent and control antimicrobial resistance*. 2024. Disponível em: <https://www.who.int/publications/i/item/9789240093461>. Acesso em: 21 jul. 2024.
31. Li, Y. et al. Characteristics of antibiotic resistance mechanisms and genes of *Klebsiella pneumoniae*. *Open Medicine*, 2023; 18(1).
32. Salawudeen, A. et al. Epidemiology of multidrug-resistant *Klebsiella pneumoniae* infection in clinical setting in South-Eastern Asia: a systematic review and meta-analysis. *Antimicrobial Resistance & Infection Control*, 2023; 12(1):142.
33. Scarpate, E. C. B.; Cossatis, J. J. A presença da *Klebsiella pneumoniae* produtora de  $\beta$ -lactamase de espectro estendido no ambiente hospitalar. *Saúde e Ambiente*, 2009; 4(1).
34. Khanfar, H. S. et al. Extended spectrum beta-lactamases (ESBL) in *Escherichia coli* and *Klebsiella pneumoniae*: trends in the hospital and community settings. *The Journal of Infection in Developing Countries*, 2009; 3(4):295-299.
35. Demirdag, K.; Hosoglu, S. Epidemiology and risk factors for ESBL-producing *Klebsiella pneumoniae*: a case control study. *The Journal of Infection in Developing Countries*, 2010; 4(11):717-722.
36. Silva, A. E. F.; Junior, O. M. R. Resistência bacteriana pelo uso indiscriminado dos carbapenêmicos meropenem e imipenem: uma revisão integrativa. *Research, Society and Development*, 2022; 11(7):e44711730195–e44711730195.
37. Silva, K. M. *Caracterização de Enterobacterales multirresistentes em estações de tratamento de esgoto e avaliação de sua remoção por processo foto-Fenton heterogêneo mediado por resíduo de mineração*. 2023. Tese (Doutorado em Ciências), Programa de Pós-graduação em Saúde Pública e Meio Ambiente, Escola Nacional de Saúde Pública Sérgio Arouca, Fundação Oswaldo Cruz, 2023.
38. Bennett, J. E.; Dolin, R.; Blaser, M. J. Mandell, Douglas, and Bennett's Principles and Practice of Infectious Diseases: 2-Volume Set. 9th edition. Philadelphia, PA: Elsevier, 2019.
39. Gazineo, J. L. D. et al. Aminoglicosídeos no século 21: Revisão e atualização, com ênfase na nefrotoxicidade. *Saúde Dinâmica*, 2023; 5(2):57-76.
40. Barbosa, K.W.P. et al. Infecções por *Acinetobacter baumannii* e mecanismos de resistência: revisão de literatura. *Brazilian Journal of Health Review*, 2023; 6(6):31679-31695.
41. Abrantes, J. A. *Avaliação da resistência bacteriana em Estações de Tratamento de Esgoto da Fiocruz com ênfase no perfil fenotípico e molecular para beta-lactamases em enterobactérias*. 2022. Tese (Doutorado em Ciências), Programa de Pós-graduação em Saúde Pública e Meio Ambiente, Escola Nacional de Saúde Pública Sérgio Arouca, Fundação Oswaldo Cruz, 2022.