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# ABSTRACT

Aim: To determine and compare the effect of used phone batteries on *Nitrosomonas* spp. in tri aquatic bodies

**Comparative Ecotoxicological Assay of E-**

Waste (Phone Batteries) On Some Aquatic

**Micro flora** 

Study design: The study employs experimental design and statistical analysis of the data and interpretation.

**Place and Duration of Study:** Fresh water and Marine samples were collected from Gokana L.G.A, Rivers state, brackish water was collected from Eagle Island Rivers state Nigeria. These samples were transported with ice pack to the microbiology laboratory of Rivers state university, Port Harcourt within 24 hours for microbiological and toxicity testing. The used phone batteries were purchase from Garrison Junction, Port Harcourt. The toxicity testing was done for duration of 4 hours interval for 24 hours respectively at room temperature.

**Methodology:** Standard microbiological techniques were used; Toxicity testing procedures were carried out by preparing mobile phone batteries at concentrations of 0%, 5%, 25%, 50% and 75%, tested for duration of 0h, 4h, 8h, 12h and 24h respectively. The cultures were incubated at 35 C for 18 to 24 hours.  $LC_{50}$  was determine using SPSS version 20

**Results:** The results indicate that percentage logarithm mortality of *Nitrosomonas* species increases with increased toxicants concentration and exposure time. The median lethal concentration (LC<sub>50</sub>) of the mobile phone batteries increases in the following order: (Note: the higher the LC<sub>50</sub> the Lower toxic the toxicant); Nokia phone battery in marine water (65.97%)<Tecno phone battery in Brackish water (65.84%)<Tecno phone battery in brackish water (65.47%)<Nokia phone battery in fresh water (64.17%) Tecno phone battery in fresh water (64.13%). **Conclusion:** Tecno phone battery in fresh water (LC<sub>50</sub> = 64.13%) is the most toxic; having the lowest LC<sub>50</sub> while Nokia phone battery in marine water (LC<sub>50</sub> = 65.97%) has the lowest toxicity effect. These results show that spent phone batteries can inhibit the nitrification process in aquatic ecosystem.

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Keywords: used phone batteries, Toxicity, *Nitrosomonas*, fresh, Brackish, Marine, Nitrification,
 ecosystem.

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# 12 **1. INTRODUCTION**

13 Ecotoxicology is the branch of toxicology concerned with the study of toxic effects caused by natural 14 or synthetic pollutants, to the constituents of an ecosystems; animals (including human), vegetations 15 and microorganisms, (Aquastel, 2007). Normal micro flora of an aquatic ecosystem controls the 16 habitability of the earth through their functions in biogeochemical cycles and food webs. The oceans 17 aquatic environments are also sensitive to durable environmental changes including those of 18 anthropogenic origin such as; E wastes disposal, etc. Microbial activities are essential to how 19 ecosystems transform pollutants which are reflected in biogeochemical cycles and food webs, and 20 how microorganisms respond to toxicant in an ecosystem will partially, if not primarily, determine the 21 fate of that ecosystem when the assimilative capacity have not been exceeded. (Aquastel, 2007)

22 According to Douglas and Nwachukwu (2016) the current general direction in technological 23 advancement and latest discoveries in information and communication technology, Electronic Devices 24 such as Laptop and Phones have become a part of day to day activities. The constant request and 25 Use of Laptops and Phones results in the constant production of large amount of Electronic wastes 26 yearly, these wastes are referred to as E- waste (Beata, 2014). The major components of the phone 27 that makes it harmful to the Environment and human health when disposed is the Heavy metals 28 contains in the battery once released into the environment, they continuously circulate therein, and 29 can cause acute or chronic poisoning (Armstrong et al., 2005). "When they are released into the 30 aquatic environment, they pollute our water bodies and when thrown into 'dump' areas their toxic ingredients are left to seep into the soil, finally to groundwater, causing massive and devastating damage to our natural ecosystem" (Armstrong *et al.*, 2005). Some metals such as mercury, lithium, cadmium, chromium, and lead are especially toxic to aquatic organisms and humans (Douglas, and Nwachukwu, 2016). But most times microorganisms are not considered when discharging wastes into water bodies and these microorganisms play vital role in an ecosystem especially *Nitrosomonas* species.

Therefore the aim of this study is to analysis and compares the toxic effect of different product of
 spent batteries on *Nitrosomonas* species in Fresh Brackish and Marine environments.

## 40 2. MATERIAL AND METHODS

## 41 **2.1 Sample Collection/Study Area**

Fresh water sample was collected in sterile (4) litre plastic container from Biara stream, while marine water was collected from Bodo city both in Gokana L.G.A, Rivers state, also, brackish water was collected from Eagle Island River in Port Harcourt L.G.A Rivers state Nigeria with a four (4) litre sterile plastic container. These samples were transported to the microbiology laboratory with ice pack within 24 hours.

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# 2.2 Microbiological Analysis

## 50 2.2.1 Total Heterotrophic Bacteria (THB)

51 Total heterotrophic bacteria for each water samples were enumerated using spread plate method. An aliquot (0.1ml) of the dilution of 10<sup>-6</sup> were aseptically transferred unto properly dried nutrient agar 52 53 plates in duplicate, spread evenly using bent glass rod and incubate at 37 C for 24 to 48 hours, after 54 incubation, the bacterial colonies that grew on the plates were counted and sub-cultured unto fresh 55 nutrient agar plates using the streak plate technique. Discrete colonies on the plates were aseptically 56 transferred into 10% (v/v) glycerol suspension, well label and stored as stock cultures for preservation 57 and identification (Amadi et al., 2014). Total Heterotrophic Bacteria (THB) Counts for each sample 58 were then calculated using the below formula:

- 59
- 60 THC (cfu/g) = <u>Number of Colonies</u>
- 61

## Dilution $(10^{-6})$ x Volume plated (0.1ml) (Nrior and Odokuma 2015).

## 62 2.2.2 Total Heterotrophic Fungi

The total fungi in each of water samples were enumerated using spread plate method. An aliquot (0.1ml) of the dilution of 10<sup>-4</sup> dilution was aseptically transferred unto properly dried Sabouround Dextrose Agar plates containing antibiotic (Tetracycline and Penicillin) to inhibit bacterial growth, in duplicate, spread evenly using bent rod and incubate at 37 C for 3days, pure culture of fungal isolates were counted and sub-cultured unto Sabouround Dextrose Agar slant in bijou bottles for preservation and identification (Odokuma and Okpokwasili 1992).

Total Heterotrophic Fungi (THB) Counts for each sample were then calculated using the below formula:

- 71 THFC (cfu/g) = <u>Number of Colonies</u>
- 72 Dilution  $(10^{-4})$  x Volume plated (0.1 ml)

## 73 2.2.3 Isolation Of *Nitrosomonas0* Species

Winogradsky Agar medium composition as modified by Odokuma and Nrior, 2015 was used: Agar 74 agar 15.0g/l, FeSO<sub>4</sub>.7H<sub>2</sub>O 0.4g/l, NaCl 2.0g/l, K<sub>2</sub>HPO<sub>4 1.0g</sub>/l, MgSO<sub>4</sub> .7H<sub>2</sub>O 0.5g/l, and (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> 75 76 2.0g/l were dissolved in 1000ml of Distilled water and autoclaved at 121°C for 15minutes (psi) after 77 which it was allowed to reduce to about 40<sup>o</sup>C and the medium was poured on the Petri-dishes. Then, 78 the medium was allowed to solidify before progress to the hot air oven to dry the moisture. An aliquot 79 from fresh, brackish and marine water respectively were inoculated unto the Winogradsky agar 80 and incubate aerobically for 2 - 3days at room temperature (30± 2°c), greyish, mucoid, flat colonies 81 revealed pear-shaped, and Gram negative of Nitrosomonas (Odokuma and Nrior, 2015). 82

83 **2.2.4** Confirmation of *Nitrosomonas* species

84 Suspected Nitrosomonas species were subcultured on a fresh Winogradsky agar medium and 85 transfered into a broth containing Ammonium sulphate and sodium nitrate and incubated at about (30± 2°c) for 2 - 3 days. 1ml of sulfanilic acid, dimethylnapthalamine and zinc dust was added to 86 87 medium after (2) days of incubation. Red colouration indicated by nitrate production from ammonium 88 sulphate was a confirmation of Nitrosomonas species. 89

### 90 Preparation of Stock Toxicant. 2.3

91 The phone Batteries (Nokia and Techno) were aseptically forced open and 4grams of each product 92 was weighed on an electric weighing balance into 100ml of autoclaved fresh, brackish and marine 93 water respectively as stock toxicant. 94

### 95 2.4 Toxicity Test Procedure

96 The toxicity tests were done by setting up fifteen test tubes aseptically covered with cotton wool. The 97 test was carried out in five (5) separate test tubes containing appropriately autoclaved water samples 98 from fresh, marine and brackish water from the habitat of the organism separately. In each of the test 99 tubes, the four toxicant concentrations (5%, 25%, 50%, and 75%) were added separately. while the 100 control consists of fresh, marine and brackish water respectively (Nrior and Gboto, 2017). One 101 millilitre (1ml) of the test organism was added to each toxicant concentration in the test tubes 102 containing (5%, 25%, 50%, 75% and control respectively). Then an aliquot (0.1ml) from each of the 103 concentrations of the effluent were then plated out using spread plate technique on Winogradsky agar 104 immediately after inoculation as zero (0) hour, inoculation and spreading continues after 4, 8, 12 and 105 24hours respectively and was incubated for 24 to 48 hours at room temperature (37± 2°C). After 106 which the colonies on the plates were counted (Odukuma and Nrior 2015).

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### 108 2.4.1 Toxicity test of bacteria Nitrosomonas species in mobile phone batteries.

109 The percentage log survival of the *Nitrosomonas* species isolates in the mobile phone batteries 110 effluent were calculated according to formula used by Nrior and Obire (2015). The percentage log 111 survival of the *Nitrosomonas* isolates in the effluent was calculated by obtaining the log of the count in 112 toxicant concentration, divided by the log of the count in the zero toxicant concentration and 113 multiplying by 100. Thus:

114 Percentage (%) log survival =  $Log C \times 100$ 

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Log c Where: Log C=log of the count in each toxicant concentration Log c = log of count in the control (zero 116 117 toxicant concentration).

- 118 Percentage (%) log mortality = 100 - % log survival
- 119

### 120 3. RESULTS AND DISCUSSION

121 The Total Heterotrophic bacterial and fungal counts of the triaquatic bodies is presented in figure 122 4.1 below, the result revealed that brackish water have the highest microbial load followed by marine

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### 124 Figure 4.1: Total Heterotrophic bacterial and total heterotrophic fungal counts 125 expressed in Log<sub>10</sub>.

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129 The logarithm counts of *Nitrosomonas* species exposed to Nokia and Techno spent phone

- 130 batteries toxicants in Fresh, Brackish, and Marine water are revealed in table 1 and 2
- 131 respectively. Percentage logarithm mortality of the counts is presented in the figures below.

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### Table 1: Log Counts Of Nitrosomonas Species With Nokia Phone Battery In Fresh, Brackish, And Marine Water

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136		Fresh	Water	+ Nc	okia		Brackish	Water	+ No	okia	М	arine W	ater -	+ Noki	a	
137	Conc./time	0H	4H	8H	12H	24H	0H	4H	8H	12H	24H	0Н	4H	8H	12H	24H
	Control	2.40	2.42	2.45	2.51	2.54	2.41	2.43	2.50	2.50	2.54	2.41	2.42	2.45	2.48	2.50
	5%	2.39	2.38	2.14	2.38	2.40	2.39	2.38	2.41	2.42	2.40	2.38	2.34	2.33	2.34	2.31
	25%	2.32	2.30	2.28	2.27	2.38	2.34	2.35	2.36	2.33	2.25	2.26	2.26	2.22	2.20	2.20
	50%	2.21	2.19	2.16	2.14	2.15	2.27	2.26	2.24	2.22	2.02	2.13	2.15	2.12	2.04	2.04
	75%	2.00	1.95	1.87	1.85	2.05	2.08	2.10	2.06	2.00	1.85	1.99	2.00	2.01	1.94	1.90

141	TABLE 2: Log Counts of Nitrosomonas Species with Techno Phone Battery In Fresh, Brackish,
142	And Marine Water

3		Fre	sh Wa	ter +	Nokia		Bracki	sh Wa	ter +]	Nokia		Marine V	Vater	+ Nok	ia		
	CONC./TIM E	0Н	4H	8H	12H	24H	0H	4H	8H	12H	24H	OH	4H	8H	12H	24H	
	CONTROL	2.40 2.39	2.42	2.45 2.38	2.51 2.38	2.58 2.40	2.41 2.39	2.43 2.38	2.50 2.41	2.50 2.41	2.54 2.40	2.41 2.38	2.42 2.34	2.45 2.33	2.48 2.34	2.50 2.31	
	25%	2.32	2.30	2.28	2.27	2.32	2.34	2.35	2.36	2.33	2.25	2.26	2.26	2.22	2.20	2.20	
	50%	2.21	2.19	2.16	2.14	2.05	2.27	2.26	2.24	2.22	2.02	2.13	2.15	2.12	2.04	2.04	
	75%	2.00	1.95	1.87	1.85	1.85	2.08	2.10	2.06	2.00	1.85	1.99	2.00	2.01	1.94	1.90	

TABLE 3: Median lethal conc. (LC50) from percentage (%) log mortality of nokia battery on nitrosomonas sp. in fresh water

Concentration	% mortality	Mean % mortality	Conc. different	∑ of Conc. diff. × mean % mortality
0	0	-	-	-
5	16	3.2	5	16
25	43	8.6	20	172
50	79	15.8	25	395
	100	00	25	500
75	100	20	20	500
75 LC <sub>50</sub> = LC <sub>100</sub> - <u>∑ C(</u>	DNC. DIFF. × MEAN %	20 MORTALITY	20	1083
75 LC <sub>50</sub> = LC <sub>100</sub> - <u>ΣCC</u>	DNC. DIFF. × MEAN % % CONTROL	20 MORTALITY	25	1083
75 LC₅₀ = LC₁₀₀ - <u>∑ C(</u> LD₅₀ <mark>=</mark> 75 – <u>1083</u>	DNC. DIFF. × MEAN % % CONTROL	20 MORTALITY	25	1083
75 LC <sub>50</sub> = LC <sub>100</sub> - ∑C( LD <sub>50</sub> <del>=</del> 75 - <u>1083</u> 100	DNC. DIFF. × MEAN % % CONTROL	20 MORTALITY	25	1083
75 LC <sub>50</sub> = LC <sub>100</sub> - ∑ <u>C(</u> LD <sub>50</sub> = 75 - <u>1083</u> 100 LD <sub>50</sub> = 75 - 10.83	DNC. DIFF. × MEAN % % CONTROL	20 MORTALITY	25	1083

Concentration	% mortality	Mean % mortality	Conc. different	∑ of Conc. diff. × mean % mortality
0	0	-	-	-
5	27	5.4	5	27
25	34	6.8	20	136
50	64	12.8	25	320
75	94	18.8	25	470
				953
$LC_{50} = LC_{100} - \sum$	Conc. diff. × mean	<u>% mortality</u>		
<mark>LD₅₀</mark> = 75 – <u>953</u>				
100				
$LD_{50} = 75 - 9.53$				

Table 4: Median Lethal Conc. (LC50) from Percentage (%) log mortality of Nokia battery on Nitrosomonas sp. in brackish water

Table 5: Median Lethal Dose (LD50) from Percentage (%) log mortality of Tecno battery on Nitrosomonas sp. in marine water

Concentration	% mortality	Mean % mortality	Conc. different	∑ of Conc. diff. × mean % mortality
0	0	-	-	-
5	19	3.8	5	19
25	37	7.4	20	148
50	66	13.2	25	330
75	118	23.6	25	590
				1087
$LC_{50} = LC_{100} - \sum_{n=1}^{\infty}$	Conc. diff. × mean % control	<u>% mortality</u>		
LD₅₀ = 75 – <u>1087</u> 100				
LD <sub>50</sub> = 75 – 10.87				
LD <sub>50</sub> = 64.13%				

Table 6: Median Lethal conc. (LC50) from Percentage (%) log mortality of Tecno battery on Nitrosomonas sp. in Fresh water 

Concentration	% mortality	Mean % mortality	Conc. different	∑ of Conc. diff. × mean % mortality
0	0	-	-	-
5	18	3.6	5	18
25	32	6.4	20	128
50	60	12	25	300
75	94	18.8	25	470
				916
$LC_{50} = LC_{100} - \sum$	Conc. diff. × mean	<u>% mortality</u>		
	% control			
<mark>_D₅₀</mark> = 75 – <u>916</u>				
100	1			
LD <sub>50</sub> = 75 – 9.16				
LD <sub>50</sub> = 65.84%				

Table 7: Median Lethal Conc. (LC50) from Percentage (%) log mortality of Tecno battery on Nitrosomonas sp.

Concentration	% mortality	Mean % mortality	Conc. different	∑ of Conc. diff. × mean % mortality
0	0	-	-	-
5	17	3.4	5	27
25	34	6.8	20	136
50	64	12.8	25	320
75	94	18.8	25	470
				943
$LC_{50} = LC_{100} - \sum_{i=1}^{n}$	Conc. diff. × mean	% mortality		
	% control			
<mark>LD₅₀</mark> = 75 – <u>943</u> 100				
LD₅₀= 75 – 9.43				
$LD_{50} = 65.57\%$				

176 177 178

171
172 Table 8: Median Lethal Dose (LD50) from Percentage (%) log mortality of Nokia battery on Nitrosomonas sp.
173 in Marine water

Concentration	% mortality	Mean % mortality	Conc. different	∑ of Conc. diff. × mean % mortality
0	0	-	-	-
5	18	3.6	5	18
25	32	6.4	20	130
50	61	12.2	25	305
75	90	18	25	450
				903
$LC_{50} = LC_{100} - \sum_{n=1}^{\infty}$	<u>Conc. diff. × mean %</u> control	<u>% mortality</u>		
LD <sub>50</sub> = 75 - <u>903</u> 100	1			
LD <sub>50</sub> = 75 - 9.03				
LD <sub>50</sub> = 65.97%				

FIGURE 1 : Summary of median lethal conc. (LC50) of mobile phone batteries (Nokia
 and Tecno) on *Nitrosomonas sp.* in freshwater, brackish water and marine.







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Figure A and B show the Percentage Logarithm Mortality of Nitrosomonas species with Nokia and Techno phone battery in fresh water respectively.



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Figure C and D show the Percentage Logarithm Mortality of Nitrosomonas species with Nokia and 186 Techno phone battery in brackish water respectively. 187



188 189

Figure E and F show the Percentage Logarithm Mortality of Nitrosomonas species with Nokia and 190 Techno phone battery in Marine water respectively

192 The results obtained in this study revealed that the toxicants can inhibit the nitrification process by 193 Nitrosomonas species. Similar observations have been reported by (Wang, 1985, Obire and Nrior, 194 2014, Nrior and Gboto 2017). An increase in the percentage logarithmic of mortality of Nitrosomonas 195 species in Fresh, Marine and Brackish water after 24 hours of exposure to the toxicant concentrations were observed (figures A to F ) respectively. This study also revealed that the 196

<sup>191</sup> 

toxicant (Techno product) toxicant is more toxic to the organism than the Nokia product. This may be
as a result of types and level of heavy metals, according to Sander *et al*,.(1985) and the site of action
of any toxicant depends on the nature of the toxicant.

200 The percent log survival of Nitrosomonas species during 0hr, 4hr, 8hr, 12hr, and 24hr exposure 201 periods to these phone battery products carried out in fresh, brackish water and Marine environments 202 (Table 4.1 and 4.2) respectively shows that both Nokia and Techno batteries exhibited little effect on 203 the test organism in fresh water than brackish water followed by Marine. This may be due to saline 204 nature of the marine and brackish water. The percent log mortality of Nitrosomonas species during 205 Ohr, 4hr, 8hr, 12hr, and 24hr exposure periods to the different concentrations of the toxicants shows 206 that the mortality rate on Techno is higher than that of Nokia battery (figures A to F). Hence, the 207 results of this study suggest that both toxicants caused cell death which resulted reduction in the 208 viable counts. This may be due to inhibition of the nitrification process within the 24hour exposure 209 period. Similar observation was reported by Nrior and Odokuma (2015) who worked on the Toxicity of 210 domestic washing bleach (Calcium hypochloride) and detergents on Escherichia coli.

211 Nitrosomonas sp. mortality expressed as Median Lethal concentration (LC<sub>50</sub>) was used as indices to 212 monitor toxicity (Nrior and Gboto, 2017). The sensitivity of the bacterium to the toxicity of the different 213 concentration of used mobile phone batteries (Nokia and Tecno) with the different water (freshwater, 214 brackish water and marine water) The median lethal Concentration (LC<sub>50</sub>) of the mobile phone 215 batteries used increases in the following order: (Note: the higher the LC<sub>50</sub> the Lower toxic the toxicant 216 and vice-vesa); Tecno phone battery in marine water (65.97%) < Tecno phone battery in Brackish 217 water (65.84%) < Nokia phone battery in marine water (65.57%) < Nokia phone battery in brackish 218 water (65.47%) < Nokia phone battery in fresh water (64.17%) Tecno phone battery in fresh water (64.13%).Conclusively, Tecno phone battery in fresh water (LC<sub>50</sub> = 64.13%) is the most toxic; having 219 220 the lowest  $LC_{50}$  while Nokia phone battery in marine water ( $LC_{50}$ = 65.97%) has the lowest toxicity 221 effect. (Table 4.3-4.8, Fig. 1) 222

### 223 Conclusion and Recommendation

The result revealed that, different concentrations of the toxicants have negative effect s on the survival rate of test organism which show that the content of these batteries can cause environmental pollution affecting *Nitrosomonas* species and other microorganism that play vital functions in an ecosystem not only that but also, batteries can also cause divers kind of acute and chronic health problems in humans and plants if released into the environment.

Therefore it is recommended that phone batteries should not be disposed directly into aquatic environment especially fresh water but rather it should be recycled.

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