

1 **EFFECTS OF HEAVY METALS ON MICROBIAL FUNCTION**
2 **IN WATER AND SEDIMENT AROUND ABONNEMA ISLAND AT LOWER**
3 **SOMBRIERO RIVER, NIGERIA**
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ABSTRACT

Aims: The interrelationships between heavy metals lead (Pb), Copper (Cu), Cadmium (Cd), Chromium (Cr) and Zinc (Zn) and microorganisms in sediment and overlying water along the shoreline of Abonnema Island were investigated by determining the levels of carbon dioxide (CO₂), Ammonia (NH₃), and Sulphate (SO₄²⁻).

Study design: Water and sediment samples were collected from seven stations approximately 500 metres apart using plastic containers and Erkman grab sampler respectively. Glucose, palm oil, and cowbell powdered milk were separately added to each set of water and sediment samples and each inoculated with 0.05mg/l, 0.15mg/l and 0.5mg/l of copper, zinc, lead, chromium and cadmium and incubated for seven days at about 30°C.

Place and Duration of Study: The study was undertaken in the Institute of Geosciences and Space Technology, University of Science and Technology, Port Harcourt, Nigeria between January 2007 and November 2008.

Methodology: Heavy metals were determined using Atomic absorption spectrophotometer. Carbon dioxide, Ammonia, Sulphate in water and sediment were determined by titration, phenate, turbidimetric and KH₂ PO₄ extraction methods using spectrophotometer. Total heterotrophic counts were carried out on Nutrient agar plates. Total coliform and Total faecal coliform counts were determined using the most probable number (MPN) technique using Mac – Conkey broth. Petroleum degrading bacteria was isolated in petroleum agar.

Results: The results showed that heavy metals had stimulatory effect on microbial functions in some cases and inhibitory effect in others. The stimulatory effect on CO₂ respiration was more in the water than in the sediment while the stimulatory effect on NH₃ respiration was more in the sediment than in the water. The stimulatory effect on SO₄²⁻ respiration was more in the sediment than in the water. The inhibitory effects on NH₃ and SO₄²⁻ respiration were more in the water than in the sediment. The concentrations of CO₂ and SO₄²⁻ were within while NH₃ exceeded permissible limit.

Conclusion: The inhibitory effects of the heavy metals on the levels of CO₂, NH₃ and SO₄²⁻ resulting from the biodegradation of carbohydrate (glucose) and protein (cowbell powdered milk) show that the rate of cycling of carbon, Nitrogen and Sulphur in the coastal waters is being affected by the heavy metals. Thus effluents should be treated before discharge into the river while direct discharge of other heavy metal containing wastes into the river should be discouraged.

Keywords: Heavy metals, Microbial function, Water, Sediment, Sombriero River, inhibition, stimulation, Abonnema Island

1. INTRODUCTION

Coastal areas are among the most biologically productive and economically important areas of the seas (Woodwell *et al*, 1973). Coastal areas are also sites of discharge and accumulation of a range of environmental contaminants (Shaw, 1984).

Microbiological processes in coastal areas contribute in ecosystem changes at larger scales. Where human induced changes have been adjudged to be negative, the natural micro biota or introduced surrogates may be a useful means for some level of restoration through bioremediation, (Capone and Bauer, 1992).

Changes in microbial functions (i.e. modification of communities functions- shift in the liberation of CO₂ and NH₃) in respect to the toxic heavy metals interactions become an indicator of the effect of toxic heavy metals on the microbial functions. Microbial function can then be potentially used as a biomarker of contaminant availability and increase in concentration. Microbial function may be a responsive marker of contaminant stress in the coastal water. The purpose of microbial activities from the microorganisms' point of view is to get energy for cellular activities but from the human point of view is to clean up organic wastes of man, domestic, industry and agriculture by processes of oxidation and reduction. (Tortora, 1989).

Contamination of soils with toxic metals is a major problem world wide; of particular interest to the field of bioremediation is the selection of biological or biochemical markers for the end point of remediation (John *et al*, 1998). Biomarkers indicate that sufficient pollutant is available for enough time to elicit a response or effect (Melanon, 1995). Because biochemical changes generally are detectable before adverse effects are seen at higher levels of biological organization, the biochemical marker approach is often an early warning or proactive tool. This is great advantage because responses at higher levels such as the ecosystem are usually measurable only after significant or permanent damage has occurred. The biomarker can be used as monitoring tool for environmental degradation. The biochemical markers are also useful to monitor the shift back to a normal state after clean up of a contaminated environment. It can also serve as evaluating remediation alternatives for contaminated aquatic system. (Michael and Michael, 2002; Timothy *et al*, 2005).

The effects on the microorganisms, the interactions of contaminants and the resident microorganisms are almost overlooked in many environmental toxicological studies (Capone and Bauer, 1992). However, environmental contaminants may affect microbiological activities. Micro

biota effects can range from stimulation to inhibition depending on the form and concentration of the contaminant. If contaminants are either acutely or chronically toxic to part or all of the microbial population, alterations in the normal path ways of carbon energy flow and productivity within the system may be expected.

At present, however, sufficient information of the effects on the microorganisms of interactions between the natural microorganisms of coastal areas and environmental contaminants especially toxic heavy metals is relatively sparse, making it difficult to predict the fate and effects on the microorganisms of contaminants compounds in affected areas (Capone and Bauer, 1992). By the breakdown of organic wastes, microorganisms contribute decisively to the natural self-purification of coastal waters. During this process the concentration of organic nutrients is diminished so much that eventually the bacterial content of the water decreases correspondingly (Rheinheimer, 1991). Natural self-purification of coastal waters is extremely important because this process keeps removing contaminants from the water so that rivers are reasonably clean again some kilometers away from the point of the organic waste outlet. This process keeps contaminants concentration in sea foods low.

The decisive role is played by bacteria and fungi however, they can break down organic compounds both those which are present in solid form and in solution (Rheinheimer, 1991; Balba and Bewley, 1991).

Heavy metals discharged into aquatic ecosystems are likely to be scavenged by particles leading to their accumulation in sediments (Luoma, 1990). A large reservoir of metals in the sediments can also act as a source to the overlying water column after their input to the ecosystem has ceased (Officer and Lynch, 1989), potentially leading to adverse ecologic effects (Luoma, 1989). However, the extent of the risks is difficult to accurately assess because of the complexity of biologic and chemical interactions that alter the bioavailability of metals. Release from sediments may not only result from re-suspension of particulates, but also through the activities of microorganisms within the sediments and at the sediment-water interface, resulting in biotransformation to more volatile/soluble forms (Gilmour *et al*, 1985). These soluble forms of the heavy metals may be incorporated in crustaceans, finfish and shellfish (NNPC and RPI, 1986).

The shoreline of Abonnema having served as harbour for more than two decades, before it was abandoned at the wake of the Nigerian Civil war in 1967 by the Nigerian ports Authority, by implication has the toxic heavy metal pollutants found around harbours.

These heavy metals get adsorbed onto the sediment surface at different pH. Heavy metals sop onto suspended particulates; this is a concern because filter feeding organisms such as shellfish are most likely to bioconcentrate the metals associated with these particulates (NNPC and RPI, 1986).

The shoreline of Abonnema is biologically productive, as artisanal fishing activities are always carried out in the coastal waters. The inhabitants of Abonnema depend largely on sea foods as the major source of their protein need. Therefore it is important to keep surveillance by the biomarker approach over increase in concentration of the toxic heavy metals and the purity of the coastal waters where these sea foods are obtained; so that bioavailable toxic heavy metals do not disable the community structure and functions of the major self-purification instruments (the microbiota) of the coastal waters by anthropogenic activities. The shoreline of Abonnema is occasionally inundated by spilled oil from oil spillage when it occurs as it leads to other oil and gas bearing communities like Idama and Soku where oil and gas exploitation activities are at the peak. An oil spill can cause not only hydrocarbon contamination but also heavy metal contamination in extremely small concentrations such as inorganic salts, metal soaps and organic metal complex compounds. (Chaerun *et al*, 2004). The building of the Abonnema - Degema bridge contributed its share of toxic heavy metals. The heavy metal contaminants may be released from the sediment into the water column; leading to the possible contamination of benthic organisms living in contact with them and finally of all the benthic food chain (Calmano *et al*, 1996).

This study aims to determine the effects of lead (Pb), Copper (Cu), Cadmium (Cd), Chromium (Cr) and Zinc (Zn) on microbial functions by determining the levels of carbon dioxide (CO₂), Ammonia (NH₃), and Sulphate (SO₄²⁻) liberated in water and sediment around the shorelines of Abonnema island.

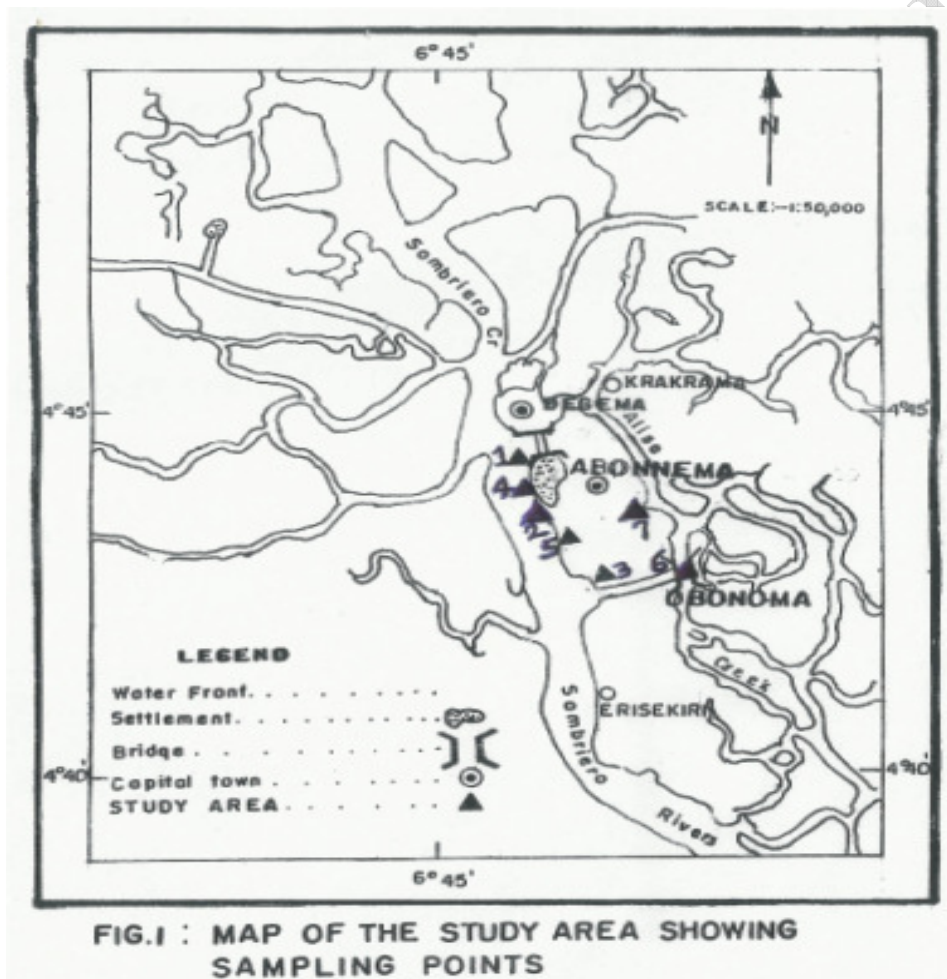
2. MATERIALS AND METHODS

2.1 THE STUDY AREA

Abonnema along whose shoreline the samples for the study were collected is located on the lateritic clay residual deposits in the transition or mangrove (Middle Delta) zone of the Niger Delta along the Sombriero River of Rivers State (Teme, 2002) Fig. 1. This is the mangrove brackish water zone that has numerous intertidal flats and mangrove vegetation subsoil here is characterized by a typical fibrous pervious clayey mud that shows a large value of compressibility and consolidation (Teme, 2002).

The climate of the zone is basically that of tropical monsoon with rainfall occurring almost all through the year except the months of December, January and February, which are not completely free from rainfall in some years (Teme, 2002).

The zone experiences diurnal ebb and flow of the tides with maximum values obtained during the once- a-year spring tide (Teme, 2002). The surface water of the Sombriero River along the shoreline of Abonnema is the sodium chloride (NaCl) subtype of the rock dominated type (Meybeck, 1981), because during the dry season a slight sparking in the river water is noticed when the water is disturbed at night by the observer in a boat. When the river water is collected and evaporated to dryness, sodium chloride (NaCl) crystals form (Ideriah *et. al*, 2012). This salt can be used as table salt after recrystallizations to improve the purity, as practiced by most coastal communities located along the Sombriero River (Ideriah *et. al*, 2012).



2.2 SAMPLING SITE SELECTION AND LOCATION

The sampling site was selected based on representativeness of the site of wastes and metal components discharge and easy accessibility. Water and sediment samples were collected from seven stations approximately 500 metres apart. Station 1 (Owusara) is situated between

longitude 006° 46' 23" E and latitude 04° 44' 24" N, station 2 (Bulk oil produce, BOP, company area) is situated between longitude 006° 46' 11" E and latitude 04° 43' 48" N, station 3 (Alice-Okolo creek junction) is situated between longitude 006° 46' 11" E and latitude 04° 43' 48" N, station 4 (Quick-Penny water side in Georgewill compound) is situated between longitude 006° 46' 9" E and latitude 04° 43' 59" N, station 5 (Timber Poku i.e. Timber water side) is situated between longitude 006° 46' 19" E and latitude 04° 45' 32" N, station 6 (Alice Okolo, Bridge Area) is situated between longitude 006° 46' 36" E and latitude 04° 43' 31" N, station 7 (Okolobio, between Jack and Georgewill compounds) is situated between longitude 006° 46' 30" E and latitude 04° 43' 59" N. The water and sediment samples were labeled (a) and (b) respectively at each station.

2.3 SAMPLE COLLECTION AND PREPARATION

The water samples were collected after recording the pH in-situ using portable digital pH meter and temperature using thermometer, plastic containers were used to collect the water sample. The containers were rinsed thoroughly with the water twice before collection. The sediments samples were collected with an 'Erkman grab' sampler.

Dissolved oxygen (DO) contents of the water samples were fixed in the field by the addition of manganous sulphate solution (Winkler I) and alkaline iodide azide reagent (Winkler II). The sediment was dried, ground, sieved, weighed and stored in polyethylene bags.

The parameters determined are Carbon dioxide, Nitrogen (ammonia), Sulphate, pH, Temperature, and Dissolved Oxygen.

2.4 MATERIALS

Water and sediment samples from Abonnema shoreline for the sources of the bacteria, glucose for carbohydrate, palm oil for lipid, cowbell powdered milk for protein, lead nitrate ($\text{Pb}(\text{NO}_3)_2$), copper nitrate ($\text{Cu}(\text{NO}_3)_2$), and zinc nitrate ($\text{Zn}(\text{NO}_3)_2$), cadmium nitrate ($\text{Cd}(\text{NO}_3)_2$), chromium chloride (CrCl_3) for the sources of the toxic heavy metals. Solar series Atomic Absorption spectrophotometer (AAS), Spectronic 21D, Erlenmeyer flasks, centrifuge tubes, 2mm sieve, Analytical balance, Whatman No 42 filter paper, conical flasks, funnels, mortar and pestle, photometer, Burettes, pipettes, mechanical shaker, Erkman grab sampler, polythene bags, plastic bottles, DO bottles and thermometer.

2.5 EXPERIMENTAL METHODS

2.5.1 Decomposable Organic Matter

Water samples, 200ml were measured into each of six (6) sets of seven 250ml plastic containers, representing the sampling stations, with lids. Into two (2) sets were inoculated 0.9375g of carbohydrate (glucose), and into the second two (2) sets were inoculated 2cm³ of lipid (palm oil) and into the third two (2) sets were inoculated 0.7930g of protein (cowbell powdered milk).

Ten grams (10g) of sediment samples was measured into six (6) sets of seven (7) 250ml plastic containers with lids. Into two (2) sets were inoculated 0.9375g of carbohydrate (glucose) and into the second two (2) sets were inoculated 2cm³ of lipid (palm oil) and into the third two (2) sets were inoculated 0.7930g of protein (cowbell powdered milk).

Ten grams (10g) of sediment samples was measured into six (6) sets of seven (7) 250ml plastic containers with lids. Into two (2) sets were inoculated 0.9375g of carbohydrate (glucose) and into the second two (2) sets were inoculated 2cm³ of lipid (palm oil) and into the third two (2) sets were inoculated 0.7930g of protein (cowbell powdered milk).

One set of the water and sediment samples inoculated with the decomposable organic matter were incubated simultaneously for seven (7) days at about 30°C. At the end of the incubation period all the samples containing carbohydrate and lipid were tested for carbon dioxide. And all the samples containing protein were tested for carbon dioxide (CO₂), Ammonia (NH₃), and Sulphate (SO₄²⁻).

2.5.2 Metal Inoculation

One set of water samples containing carbohydrate, lipid, and cowbell powdered milk, were inoculated with 0.0277g of (Cu (NO₃)₂) containing 0.05mg/l of copper. One (1) set of sediment samples containing carbohydrate, lipid and cowbell powdered milk were inoculated with 0.0277g of (Cu (NO₃)₂). Both the water and sediment samples containing the heavy metal were incubated for seven (7) days at about 30°C. At the end of the incubation period all the samples containing carbohydrate and lipid were tested for carbon dioxide and all the samples containing proteins were tested for carbon dioxide (CO₂), Ammonia (NH₃), and Sulphate (SO₄²⁻).

The differences between the chemical parameters tested for in the samples with heavy metal and those without heavy metals were noted. The sets of containers containing metals were similarly and separately treated (incubated) with the following concentrations of heavy metals: (0.05mg, 0.15mg and 0.5mg)/l of copper; (0.05mg, 0.15mg, 0.5mg)/l of zinc; (0.05mg, 0.15mg, 0.5mg)/l of lead; (0.05mg, 0.15mg, 0.5mg)/l of chromium; (0.05mg, 0.15mg, 0.5mg)/l of cadmium.

2.6 ANALYTICAL METHODS

2.6.1 Heavy metals were determined using solar S. series Atomic absorption spectrophotometer (AAS).

2.6.2 Carbon dioxide was determined by titration method. The sample was titrated with sodium hydroxide solution to the pink colour and point using phenolphthalein indicator.

2.6.3 Ammonia was determined by phenate method. 50ml of the sample solution was treated with 2ml each of phenol and sodium nitroprusside, 5ml each of hypochlorite, trisodium citrate and sodium hydroxide solutions to produce bluish colour. The mixture was measured in spectrophotometer spectronic 2ID at 630nm.

2.6.4 Sulphate in the water samples was determined by turbidimetric method and sulphate in the sediment by KH_2PO_4 extraction method.

2.6.4.1 Turbidimetric method: Sample aliquot, 20ml, was measured into conical flask and 1ml of conditioning reagent added. BaCl_2 crystals were added to the sample and shaken. The absorbance of the sample solution was measured before and after addition of BaCl_2 with spectrophotometer spectronic 2ID (APHA, 1995).

2.6.4.2 KH_2PO_4 Extraction method: Five grams (5gm) of the sediment sample was treated with 25ml of extraction solution KH_2PO_4 solution and filtered through Whatman No. 42 filter papers. Sulphate content was determined by the turbidity method and using spectrophotometer. pH was determined by pH meter. Temperature was determined by Thermometer.

2.6.5 Dissolved oxygen (DO) was determined by the modified Winkler method (APHA 1995). The sample was collected into 75ml DO bottle; 0.5ml of Winkler I solution was added followed by 0.5ml of Winkler II solution. The sample was corked and shaken, and then 0.5ml by 0.5ml of concentrated sulphuric acid H_2SO_4 , was added and shaken. 25ml of the fixed sample was titrated against 0.025 N $\text{Na}_2\text{S}_2\text{O}_3$ with starch indicator.

2.6.7 Microbiological analysis

The samples were investigated for Total heterotrophic count (THC), Total coliform count (TCC) and Total faecal count (TFC). Petroleum degrading Bacteria count (PDBC).

The total heterotrophic counts were carried out on Nutrient agar plates and incubated at 37°C for 48 hours.

Total coliform and Total faecal coliform count were determined using the most probable number (MPN) technique using Mac – Conkey broth in tubes and incubating at 37°C for 24 hours.

Petroleum degrading bacteria was isolated in petroleum agar. The sediment was added to diluents i.e. 1g of the solid to 10ml of H_2O and 0.1ml was plated directly on the petroleum agar.

The solid nutrient is petroleum and agar for solidification.

Bacteria characterization was by Gram staining method.

3. RESULTS AND DISCUSSION

3.1 RESULTS

The Results of the study are presented in Table 1 below.

Table 1: Levels of Physico-Chemical Parameters Measured in Water and Sediment along the Shoreline of Abonnema before Incubation.

S/No.	Sample No.	pH	DO (mg/l)	SO ₄ ²⁻ (mg/l)	NH ₃ (mg/l)	CO ₂ (mg/l)
1	1a	7.0	15.71	196.10	0.173	12.49
2	1b	6.2	9.32	52.83	0.710	12.49
3	2a	7.1	14.61	163.12	0.034	19.02
4	2b	5.9	5.67	29.11	0.820	17.48
5	3a	7.2	12.42	132.68	0.174	15.68
6	3b	6.1	7.88	88.98	0.560	14.98
7	4a	6.81	5.7	269.7	0.07	16.48
8	4b	6.5	5.62	551.2	1.02	19.76
9	5a	6.9	7.3	373.7	<0.02	16.28
10	5b	7.1	6.82	424.4	0.91	14.78
11	6a	7.06	8.1	287.4	<0.02	15.98
12	6b	6.3	6.78	728.8	0.49	16.0
13	7a	6.99	8.1	218.9	<0.02	10.88
14	7b	6.1	5.93	459.9	1.76	13.98

a = water b = sediment

Table 1 shows the levels of physico-chemical parameters determined in water and sediment before incubation in the decomposable organic matter (glucose, palm oil and cowbell powdered milk) and heavy metals.

3.1.1 pH

pH range of 6.81 – 7.2 was determined in the water and 5.9 – 7.1 in the sediment. The lower limit of the pH in water was determined at station 4 and the upper limit at station 3. The lower limit of the pH in the sediment was determined at station 2 and the upper limit at station 5. The pH was higher in the water than in the sediment.

3.1.2 Dissolved Oxygen (DO)

Concentrations of dissolved oxygen ranged of 5.7 – 15.71mg/l were determined in the water and 5.62 – 9.32mg/l in the sediment. The lower limit of the dissolved oxygen in the water was determined at station 4 in the upper limit at station 1. The lower limit of the dissolved oxygen in

the sediment was determined at station 4 and the upper limit at station 1. Concentrations of dissolved oxygen was higher in the water than in the sediment

3.1.3 NH₃

Concentrations of NH₃ range of < 0.02-0.174mg/l were determined in the water and 0.49 – 1.76mg/l in the sediment. The lower limit of NH₃ in the water was determined at station 5, 6 and 7 and the upper limit at station 3 the lower limit of NH₃ in the sediment was determined at station 6 and the upper limit at station 7. Concentration of NH₃ was higher in the sediment than in the water.

3.1.4 CO₂

Concentrations of CO₂ range of 10.88-19mg/l were determined in the water and 12.49-19.76mg/l in the sediment. The lower limit of CO₂ in the water was determined at station 7 and the upper limit at station 2. The lower limit of CO₂ in the sediment was determined at station 1 and the upper limit at station 4.

3.1.5 SO₄²⁻

Concentrations of SO₄²⁻ range of 132.68-373.7mg/l was determined in the water and 29.11 – 728.8mg/l in the sediment. The lower limit of SO₄²⁻ in the water was determined at station 3 and upper limit at station 5. The lower limit of SO₄²⁻ in the sediment was determined at station 2 and the upper limit at station 4. Concentration of SO₄²⁻ was higher in the sediment than in the water.

3.2 DISCUSSIONS

3.2.1 pH

In Table 1, the lower limits of the pH range observed at stations 7 and 5 were slightly acidic; this may be as a result of high organic carbon load oxidation in those stations which produced large volumes of carbon dioxide that dissolved in the water to produce carbonic acid which lowered the pH. The pH of the water was higher than that of the sediment. Statistical analysis showed significant difference (p<0.05) between the pH in the water and sediment. This was due to organic carbon breakdown in the sediment. On the average the pH range was within the permissible limit of 6.5 – 8.5 (RSMENR, 2002).

The relative changes in pH will be affected by the buffer capacity of the water and amount of substrate utilized by the microorganisms. (Verschuere, 1983; Balba and Bewley, 1991).

3.2.2 Dissolved Oxygen (DO)

The level of dissolved oxygen in the water was higher than that in the sediment. This may be due to photosynthesis by phytoplankton taking place in the water and limited dissolved oxygen penetration into the sediment. Statistical analysis showed significant difference ($p < 0.05$) between the levels of dissolved oxygen in the water and sediment. Dissolved oxygen concentrations in all the stations were within the permissible limit of 5ppm (RSMENR, 2002).

3.2.3 Ammonia (NH₃)

The concentrations of NH₃ was higher in the sediment than in the water but statistical analysis showed that there was no significant difference ($p < 0.05$) between the concentrations of NH₃ in the water and sediment. The concentrations NH₃ in the water and the sediment were higher than the permissible limit of 0.0125ppm (RSMENR, 2002).

3.2.4 Carbon Dioxide (CO₂)

The concentrations of CO₂ in the water and sediment did not vary much. Statistical analysis showed that there was no significant difference ($p > 0.05$) between the concentrations of CO₂ in the water and in the sediment. The concentrations of CO₂ were within the permissible limit of 0 – 15ppm (RSMENR, 2002).

3.2.5 Sulphate (SO₄²⁻)

The concentrations of SO₄²⁻ were higher in the sediment than in the water but statistical analysis showed that there was no significant difference ($p > 0.05$) between them. The mean concentrations of SO₄²⁻ in the water and the sediment were within the permissible limit of 500mg/l (RSMENR, 2002).

3.3 EFFECT OF INCUBATION

The concentrations of CO₂ determined in the water after incubation with glucose were higher than those in the water before incubation. This was as a result of higher population of bacteria produced by the incubation. (Fig.1a).

Statistical analysis showed that there was significant difference ($p < 0.05$) between them with very high positive correlation (0.7164). The concentrations of CO₂ measured in the sediment after incubation with glucose were higher than those in the water before incubation (Fig 2a). This was as a result increased microbial activities resulting from incubation which increased the population. Statistical analysis showed no significant difference ($p > 0.05$) between them with negative correlation (- 0.1973). The concentrations of CO₂ measured in the water after incubation with palm oil were similar with those before incubation (Fig.1b). Statistical analysis showed that there was no significant difference ($p > 0.05$) between them with negative correlation between them (- 0.1000). There was also no significant difference ($p > 0.05$) between the concentrations of CO₂

measured in the sediment after incubation with palm oil and those before incubation with negative correlation between them (-0.2429) (Fig. 2b).

The concentrations of CO_2 measured in the water after incubation with protein were higher than those in the water before incubation (Fig. 1c), but statistical analysis showed no significant difference ($p < 0.05$) with positive correlation (0.0527) between them. The concentrations of CO_2 measured in the sediment after incubation with protein were higher than those before incubation (Fig. 2c) but statistical analysis showed no significant difference ($p < 0.05$) with very high positive correlation (0.7061) between them.

The concentrations of NH_3 determined in the water after incubation with cowbell powdered milk were higher than those in the water before incubation (Fig. 3). This was as a result of bacteria population increase due to incubation. Statistical analysis showed no significant difference ($p < 0.05$) between them and the correlation between them was positive (0.2499). The concentrations of NH_3 measured in the sediment after incubation with cowbell powdered milk was higher than those in the sediment before incubation (Fig. 4) but statistical analysis showed that there was no significant difference ($p < 0.05$) between them with negative correlation between them (-0.2747).

The concentrations of SO_4^{2-} measured in the water after incubation with cowbell powdered milk was lower than those in the water before incubation (Fig. 5). This was as a result of low concentrations of the protein for the incubation compared to the high protein load in the shoreline. Statistical analysis showed that there was significant difference ($p < 0.05$) between them with negative correlation (-0.1053). The concentrations of SO_4^{2-} measured in sediment before incubation was higher than those in the sediment after incubation with cowbell powdered milk (Fig. 6). Statistical analysis showed that there was significant difference ($p < 0.05$) between them with high positive correlation between them (0.6841). This was as a result of low concentration of SO_4^{2-} used for the incubation compared with the high protein load along the shoreline.

3.4 EFFECTS OF HEAVY METALS

The concentrations of CO_2 measured in the water containing glucose before incubation was higher than those in the water after incubation with a mixture of glucose and Cu (Figs. 7a – 7b). Statistical analysis showed that there was no significant difference ($p < 0.05$) between them with negative correlation between them (-0.2377). The concentrations of CO_2 in the sediment treated with glucose before incubation was higher than those in the sediment after incubation with a mixture of glucose and Cu (Fig 10a – 10b) but statistical analysis showed that there was no significant difference ($p > 0.05$) between them with negative correlation (-0.3823).

The concentrations of CO_2 in the water after incubation with palm oil and Cu was higher than those in the water treated with palm oil (Figs. 7a – 7b) but statistical analysis showed no significant difference ($p < 0.05$) with positively high correlation (0.6336). The concentrations of CO_2

in the sediment containing palm oil and Cu were higher than those treated with palm oil after incubation (Fig. 10a – 10b) but statistical analysis showed that there was no significant difference ($p>0.05$) between them with positive correlation (0.6888) between them. The concentrations of CO_2 in water containing cowbell powdered milk and Cu after incubation was higher than those in the water treated with cowbell powdered milk before incubation (Fig. 7a – 7b) but statistical analysis showed significant difference ($p>0.05$) with positive correlation (0.3846) between them. The concentrations of CO_2 in the sediment containing cowbell powdered milk and Cu after incubation were higher than those in the sediment treated with cowbell powdered milk before incubation (Figs. 10a – 10b). Statistical analysis showed that there was no significant difference ($p>0.05$) between them with positive correlation (0.0025).

The concentrations of NH_3 in the water containing cowbell powdered milk and Cu after incubation was higher than those in the water treated with cowbell powdered milk before incubation (Fig. 8a – 8b) but statistical analysis showed that there was significant difference ($p>0.05$) between them with positive correlation (0.2471). The concentrations of NH_3 in the sediment treated with cowbell powdered milk and Cu after incubation was higher than those in the sediment treated with cowbell powdered milk after incubation (Figs. 11a – 11b) but statistical analysis showed that there was significant difference ($p<0.05$) between them with positive correlation (0.2707) between them.

The concentrations of SO_4^{2-} in the water treated with cowbell powdered milk and Cu after incubation was higher than those in the water treated with cowbell powdered milk before incubation (Figs. 9a – 9b) but statistical analysis showed there was no significant difference ($p>0.05$) between them with positive correlation (0.1626) between them. The concentrations of SO_4^{2-} in the sediment treated with cowbell powdered milk and Cu before incubation was higher than those in the sediment treated with cowbell powdered milk before incubation (Figs. 12a – 12b) but statistical analysis showed that there was no significant difference ($p>0.05$) between them but the correlation between them was positively very high (0.8950).

Cu inhibited the oxidation of glucose to produce CO_2 in the mixture of glucose and Cu in the water and sediment but stimulated CO_2 production in the mixture of palm oil and Cu in the water and sediment; Cu also stimulated CO_2 production NH_3 production in the mixture of cowbell powdered milk and Cu in the water and sediment but inhibited SO_4^{2-} in the sediment and stimulated SO_4^{2-} oxidation in the water (Capone and Bauer, 1992).

Similar observations were made with Zn, Pb, Cd and Cr. (Figs. 13 – 22).

From the results of this research work it was observed that the interfaces of toxicity did not depend on heavy metals concentration gradients but on bioavailability of the metals.

In some cases lower concentrations of the metals had higher toxic effects on the microorganisms which were either stimulatory or inhibitory than the higher concentrations of the metals (Capone and Bauer, 1992). This may be due to the fact that toxicity is related to bioavailability of a particular metal for uptake (NNPC and RPI, 1986). However, bioavailability or toxicity is correlated with the free metal concentrations (Borgmann, 1983).

4. CONCLUSION

The inhibitory effects of the heavy metals on the levels of CO₂, NH₃ and SO₄²⁻ resulting from the biodegradation of carbohydrate (glucose) and protein (cowbell powdered milk) show that the rate of cycling of carbon, Nitrogen and Sulphur in the coastal waters is being affected by the heavy metals.

The Government should introduce domestic waste treatment methods so that only treated wastes will be discharged into the coastal waters.

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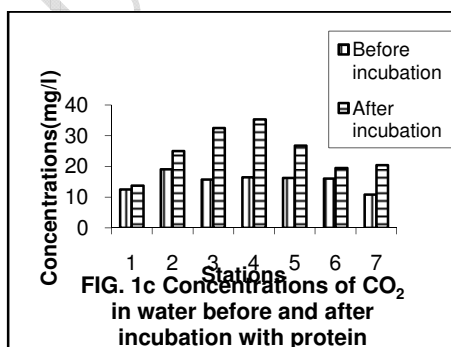
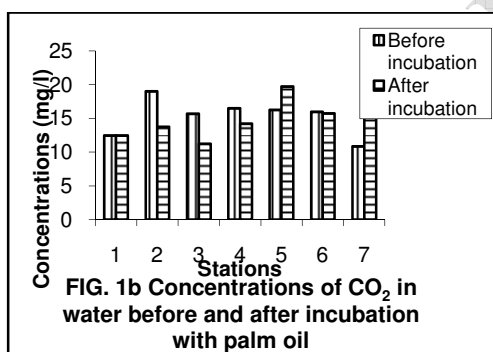
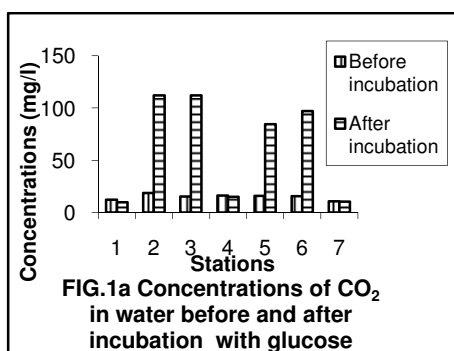
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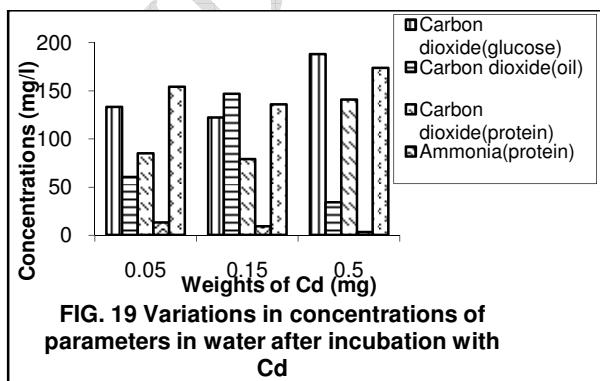
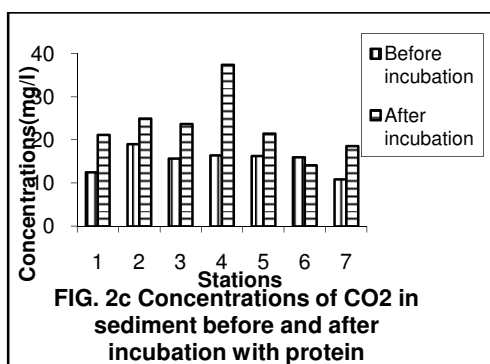
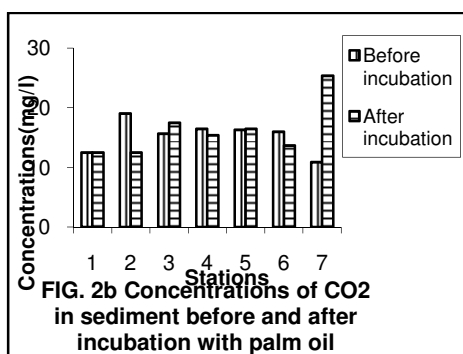
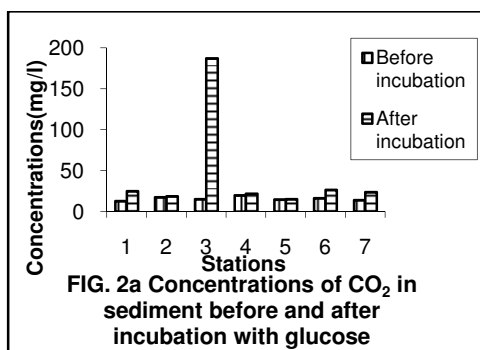
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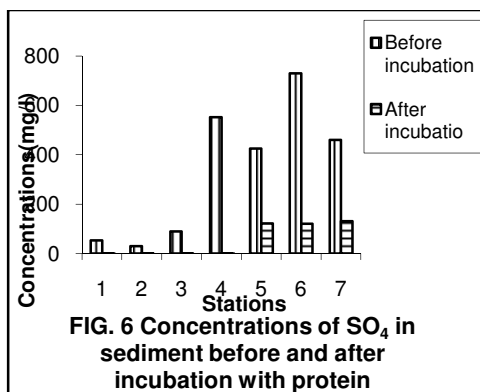
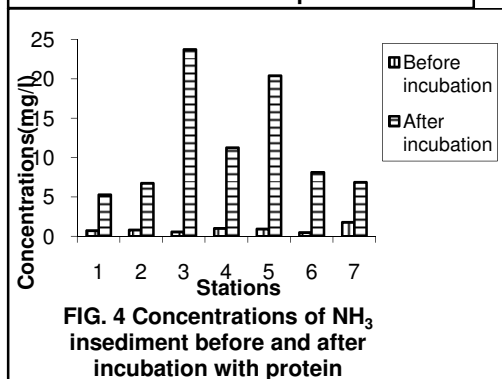
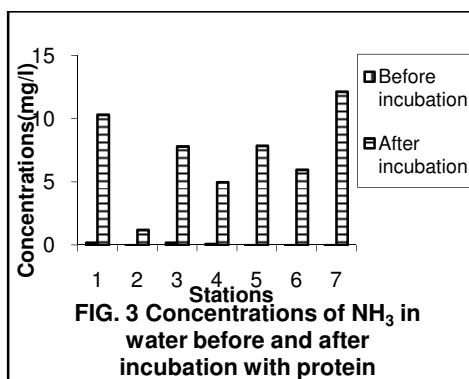
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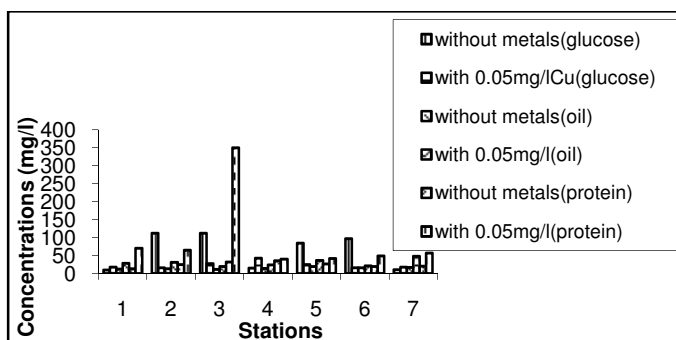


FIG. 7a Concentrations of CO₂ in water after incubation with and without metal

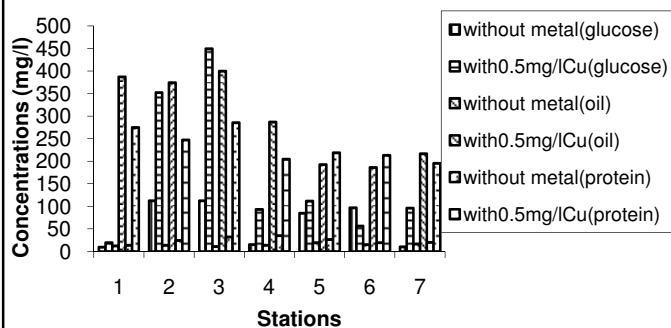


FIG. 7b Concentrations of CO₂ in water after incubation with and without metals

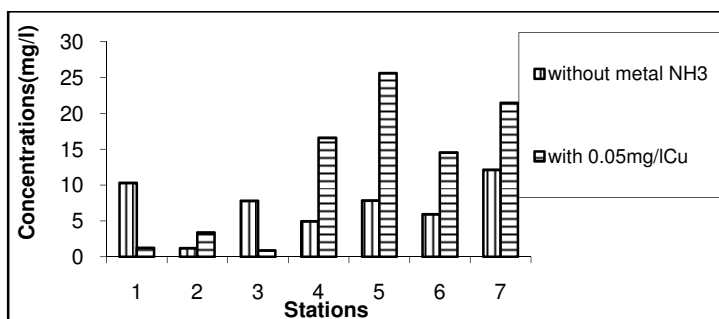


FIG. 8a Concentrations of NH_3 in sediment after incubation with and without metal

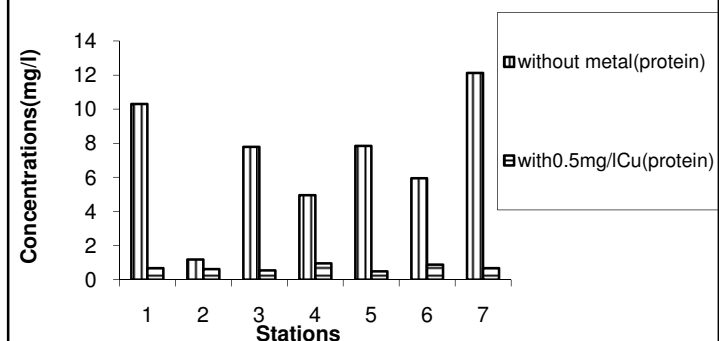


FIG. 8b Concentrations of NH_3 in sediment after incubation with and without metal

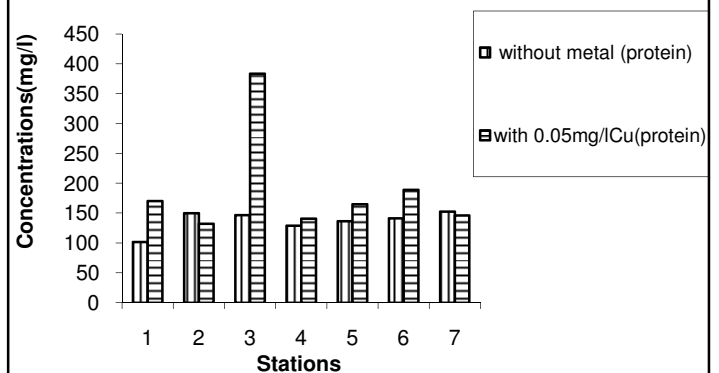


FIG. 9a Concentrations of SO_4^{2-} in water after incubation with and without metal

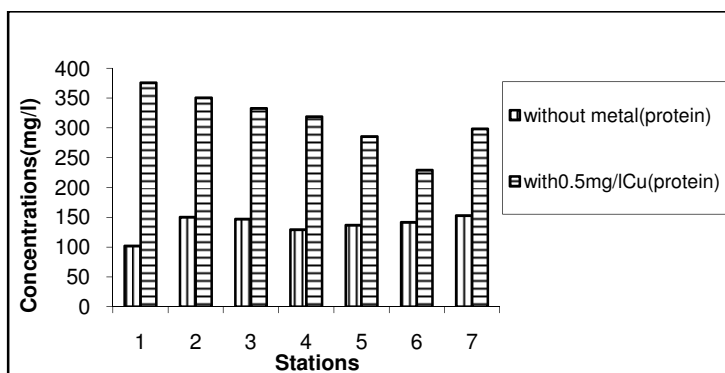


FIG. 9b Concentrations of SO_4^{2-} in water after incubation with and without metal

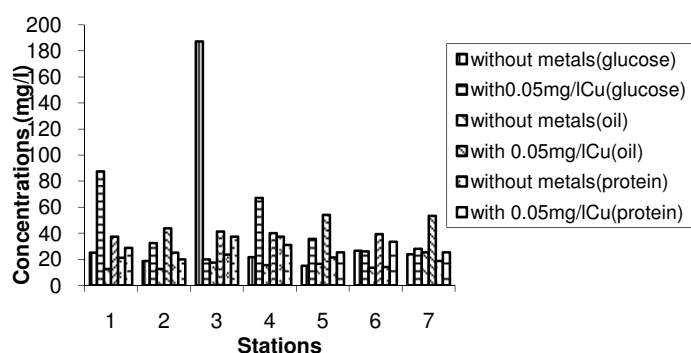


FIG. 10a Concentrations of CO_2 in water after incubation with and without metal

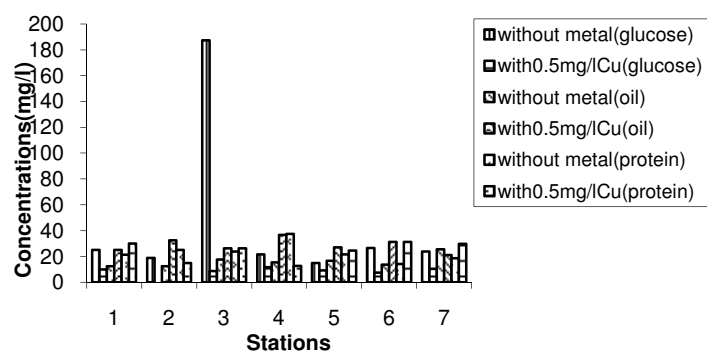
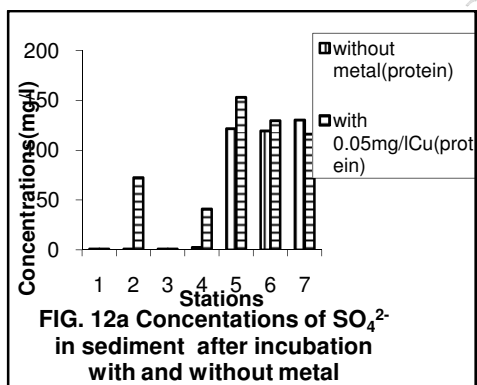
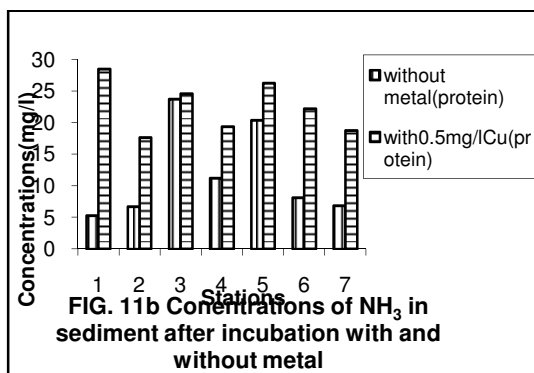
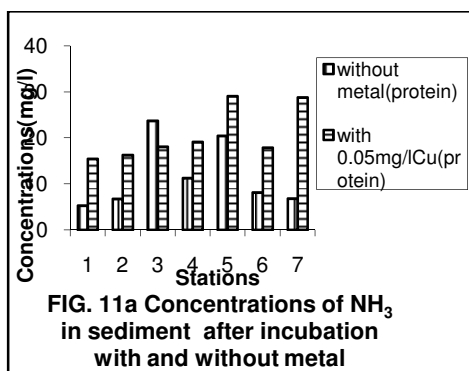
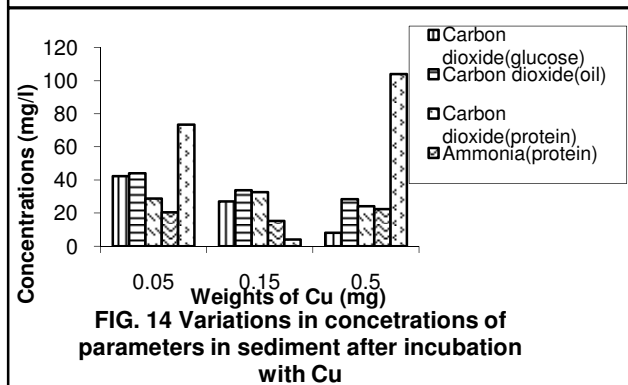
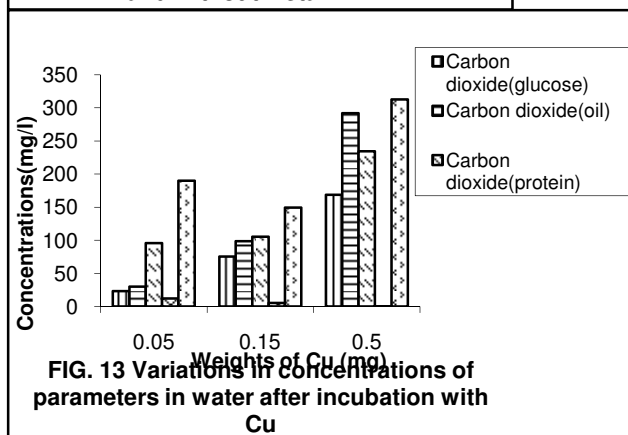
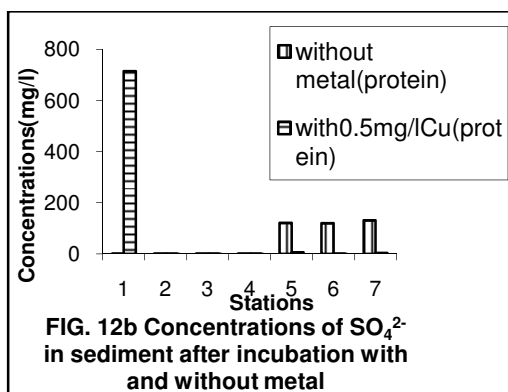


FIG. 10b Concentrations of CO_2 in water after incubation with and without metal





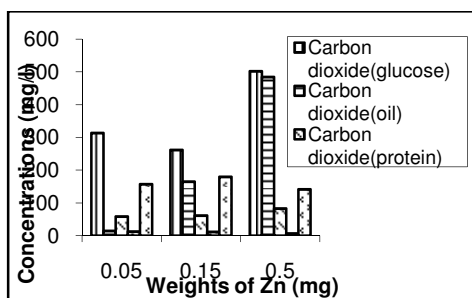


FIG. 15 Variations in concentrations of parameters in water after incubation with Zn

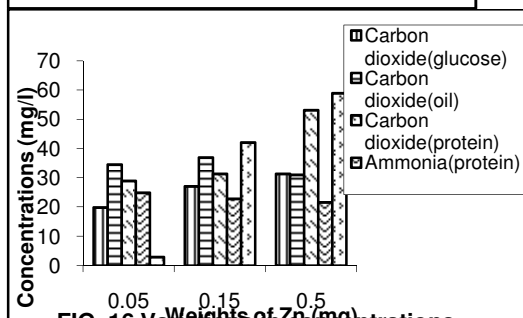


FIG. 16 Variations in concentrations of parameters in sediment after incubation with Zn

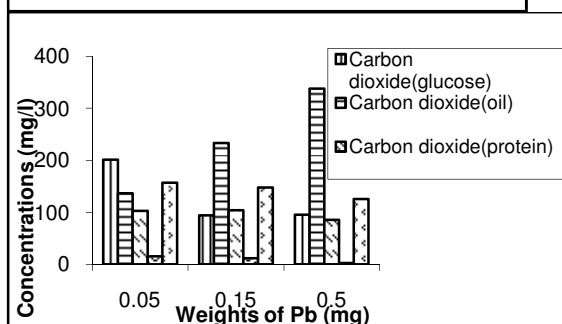


FIG. 17 Variations in concentrations of parameters in water after incubation with Pb

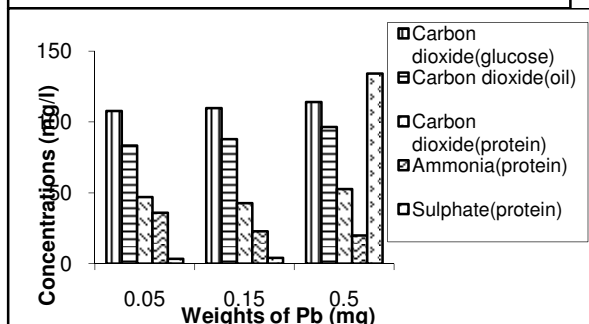


FIG. 18 Variations in concentrations of parameters in sediment after incubation with Pb

