Phytoremediation Of Crude Oil Impacted Soil Using Purple Nutsedge

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ABSTRACT: The present study investigated the viability of purple nutsedge in the phytoremediation of a crude oil-contaminated land in the Kom-Kom community, Oyigbo, Rivers state, Nigeria. 150g of soil samples were randomly collected from two (2) different points on the polluted site and a control site and analyzed for Petroleum Aromatic Hydrocarbons (PAHs), Total Petroleum Hydrocarbons (TPH) and Heavy metals (Pb, Cd, Cr & Ni) in soils and plants before and after phytoremediation. Plants were transplanted into the contaminated and contaminated soil after soil sample collection. After planting, the progress of plant growth was observed and recorded biweekly for 3 months before harvesting. From the results obtained, over 80% and 66% PAHs and TPHs phytodegradation efficiencies were achieved using the plant while Cd, Pb and Cr were removed by 90%, 67% and 39.2% respectively. The Bioaccumulation Factor (B.F) of the heavy metals in study plant were found to be greater than 1 which makes it suitable for phytoextraction of heavy metals. Therefore, the study suggests that purple nutsedge can be useful in the phytoremediation of a crude oil polluted soil.

DOI: https://dx.doi.org/10.4314/jasem.v25i3.25

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Dates: Received: 12 December 2020; Revised: 26 January 2021; Accepted: 12 February 2021

Keywords: Hydrocarbons, Heavy metals, Phytoremediation, Crude oil, Purple nutsedge

The discovery of crude oil in Nigeria has significantly improved her economy due to its importance as a significant source of foreign exchange and in boosting the Gross domestic product (GDP) of the country. However, it’s exploration and exploitation has posed immense danger and harm to the oil-producing communities due to the resultant effect it has on the aquatic life and soil topography of the oil-rich region. The Niger Delta region of Nigeria is well-known for the production of oil that commenced as far back as 1956 and the discovery of oil in the Niger Delta region improved the financial state of the economy but was immediately accompanied by a series of crude oil spillage on land and water giving rise to soil and water pollution (Adekunle et al., 2015). The major cause of oil spills has been identified to be from vandalism and leakages of pipelines. The effects of environmental pollution resulting from the production of oil became a topical issue at both national and international levels (Adekunle et al., 2015). Oil spillage results in contamination of the environment with aliphatic and aromatic hydrocarbons(such as Total Petroleum Hydrocarbon (TPH) & Polycyclic Aromatic hydrocarbon (PAH) as well as heavy metals(such as Pb, Ni, Cr & Cd). Hence, it is essential to develop efficient remediation strategies to reduce the disastrous effect of hydrocarbon and heavy metals pollution in soils in the Niger Delta Region as, high concentrations in soils pose a great risk to humans and the entire ecosystem at large(Nwaichi, 2014; Nwaich et al., 2016). Also, it is pertinent to investigate methods and strategies to clean up oil spillage and prevent its disastrous effect on soil, animals, and humans. One of the most promising biological approaches, to deal with the problem of oil spillage is phytoremediation- which is an In-situ type of bioremediation, and the advantage of this approach over others is that it is cost-effective, eco-friendly, and suitable for the treatment of large volumes of soils in contaminated sites (Sabo et al., 2018). The four distinct mechanisms of phytoremediation include phytostabilization, phytoextraction, phytodegradation, phytovolatilization, and rhizodegradation (Germida et al., 2002). Uptake of organic pollutants by plants is dependent on two major factors namely: abiotic (such as physiochemical properties of the molecule in terms of log octanol /water partition coefficient \(K_{ow}\), its molecular weight and the composition of clays, iron oxides and organic oxides present in the soil) and biotic factors (such as transpiration rates, types of amount of lipids in root cells, enzyme complement, root exudates, and growth dilution) (Collins et al., 2006; Osuoha and Nwaichi, 2020). From recent studies, it has been observed that uptake of PAHs by plants occurs primarily through the roots and secondarily through the leaves while the uptake of metals by plants occurs majorly through the process of phytoextraction. Phytoextraction is the process of
removal of heavy metals from contaminated sites through their uptake into different parts of the plant (Suman et al., 2018). *Cyperus rotundus* (also known as purple nutsedge) is one of the dominant plant species found growing along the contaminated site and is tolerant to the soil and weather conditions of the contaminated area. In previous field researches and investigations, it has been found that *Cyperus rotundus* can be beneficial to the phytoremediation of oily soils due to its great root surface area (Wang et al., 2010). However, the study about the viability of purple nutsedge to phytoremediate a crude oil impacted soil within oil-producing communities in Niger Delta rarely exist. This research aims to evaluate the viability of purple nutsedge to phytoremediate a crude oil impacted soil in Kom-Kom community, Oyigbo Rivers state Nigeria.

**MATERIALS AND METHODS**

*Description of the study area*: The study area is located in Kom-Kom community, Oyigbo, Rivers state, Nigeria. The Trans Delta Bonny Light Line of an oil company passes through the area. A one-week crude oil spillage was reported in this community in March 2014. Reports showed that the cause of the release was from the vandalism of old above ground pipelines. The type of soil found in the study area is loamy soil and the most common type of weed found growing on the soil is *Cyperus rotundus*. It is important to mention that Kom-Kom Community is a fast-growing urban settlement and therefore needs environmental cleanup. Control soil consisted in a soil from same geographical location but with no history of crude oil pollution for comparison.

*Sample collection*: The samples used in the analysis of the project were collected from Kom-Kom community in Oyigbo, Rivers state. A hand soil auger was used to collect soil samples in replicates of three at a 30cm depth from the contaminated study area and these were bulked. The samples were immediately placed in a sterilized, air tight cellophane bags, labelled and stored at 4°C prior to laboratory analysis. Soil samples were also obtained in replicate of three from the control site. Taxonomic classification of the experimental soil was loamy sand. Three replicate samples of the plant were collected from the study area. The samples were immediately placed in a sterilized, air tight cellophane bags, labelled and stored at 4°C prior to laboratory analysis. The plant was identified by a taxonomist in the department of Plant Biology and Biotechnology, University of Port Har court to be *Cyperus rotundus* (Purple nutsedge).

*Methods*: The study was an experiment carried out at the the contaminated site. Three 12L buckets were filled to the brim with uncontaminated soil for the control samples while three different points in the contaminated sites were mapped out for polluted samples. 150g of soil samples were gotten from the contaminated and uncontaminated soil for laboratory analysis before planting. Plants were transplanted into the contaminated and contaminated soil after soil sample collection. After planting, the progress of plant height and width was observed and recorded biweekly for 3 months before harvesting. The position and terrain of the mapped out sites were the only notable differences between the polluted samples. All samples were prepared in triplicate determination.

*Preparation of samples*: Soil samples were air-dried under room temperature to ensure constant weight. After drying, they were homogenised to obtain finer texture and remove unwanted particles. The air-dried soil samples were then sieved through a 2 mm polythene sieve to obtain only particles less than 2mm mesh size. Plant samples were gently washed under running water, cut into smaller sizes using a knife and oven dried at 60°C for 24 hours to ensure constant weight. The dried samples were then homogenized to a suitable size for digestion and analysis.

*Samples digestion and analysis*

**PAH AND TPH**: Ten grams (10g) of the sample was measured into a solvent rinsed beaker. Thirty (30) ml of dichloromethane was then added to the sample. Sample was then spiked with ortho-terphenyl. Sample was shaken in a vortex mixture for five minutes. Sample was placed in an orbital shaker for thirty minutes. The extract was filtered through a glass funnel with glass wool and anhydrous sodium sulphate. The extract was then transferred to a Teflon lined screw cap vial ready for PAH and TPH subsid ized using a gas chromatography. The extract was then analyzed using gas chromatography.

*Heavy metals*: 0.2g of sample was dissolved in 6ml of concentrated nitric acid (HNO3), 2 ml of concentrated hydrochloric acid (HCl) and 2 ml of hydrofluoric acid (HF) and set on a water bath to dissolve completely. The concentrations of the heavy metals Cd, Cr, Ni, and Pb in the solution were analyzed with Atomic Absorption Spectrometer (AAS).

*Nitrate*: About 5g of soil sample was weighed and 20ml of sodium acetate extracting solution added and shaken for 1 minute and filtered with Whatmann filter paper. About 5ml of filtrate was added to Nitravera 5 powder pillow, shaken for 1 minute and allowed to stand for 5 minutes. The absorbance was read using UV visible spectrophotometer.

*Phosphate*: An aliquot of 2g of soil sample was weighed. 40ml extracting solution was added and filtered. About 5ml of soil extract was added and 25ml
of distilled water. It was left for 10 mins and the absorbance was read.

**Electrical Conductivity:** The conductivity meter was calibrated using the 1000µS/cm conductivity standard. The electron was rinsed with de-ionized water. 10g of dried sample was weighed and added into 10ml of deionized water. It was shaken for 10 minutes and then the conductivity value was taken when the reading was stable.

**Formulas for hydrocarbon and heavy metal remediation**

Amount Remediated \((AR)_{\text{Parameter}}\) = \(\text{Initial Conc. (Ic)} - \text{Final Conc. (Fc)}\)

\[%\;\text{Bioremediation} = \frac{AR}{Ic} \times 100\]

Bioaccumulation Factor \((B.F) = \frac{[C_p]}{[C_s]}\)

Where \([C_p]\) means metal Conc. in Plant Tissue (whole plant) and \([C_s]\) means initial Conc. of Metal in Substrate i.e soil

**Data analysis:** Statistical analysis was performed using statistical packages for social sciences (version 20). Result were expressed as mean ± standard deviation of triplicate determination. The data were by one way ANOVA followed by Duncan’s multiple range test, to determine the level of significance expressed at 95% confidence level \((p< 0.05).\)

**RESULTS AND DISCUSSION**

Petroleum hydrocarbons and heavy metals are accumulated in soils as a result of oil spillage. Results of Polycyclic Aromatic hydrocarbons (PAH) and Total Petroleum Hydrocarbons (TPH) in soil samples and Heavy metals (Pb, Ni, Cr, Cd) found in soil and plant samples are presented in Tables 1, 2 & 3. There is an increasing problem of crude oil pollution which has affected several oil producing countries especially Nigeria. Crude oil contamination has negatively impacted plants and the ecosystem at large (Odukayo et al., 2019). Though crude oil contains majorly aliphatic and aromatic compounds, it also contains trace element like nickel, iron, aluminium and some heavy metals like lead, cadmium and chromium (Wilberforce, 2016). From Table 1, the PAHs and TPHs decreased significantly \((p<0.05)\) after phytoremediation, giving a 80% and 66% reduction in concentrations of PAHs and TPHs respectively; this is in agreement with the study reported by Hovet et al. (2016). The decrease observed in the soil samples indicated that *C. rotundus* was effective in taking up the PAH into its system. Also, there was a significant difference between the contaminated and uncontaminated soil. High levels of PAHs and TPHs has a negative impact on the availability of nutrients needed for plant growth and can lead to death of aquatic and terrestrial organisms.

Table 1: Hydrocarbons content (PAH & TPH) (mg/kg) of remedied and control soils

| Hydrocarbons | Initial sample | Final sample | Initial control | Final control |
|--------------|----------------|--------------|----------------|--------------|
| PAH          | 0.6±0.03       | 0.12±0.01    | 0.04±0.01      | 0.01±0.00    |
| TPH          | 5726.3±90.00   | 1963.0±10.00 | 222.5±10.00    | 7.8±0.10     |

Data are expressed as mean ± standard deviations of triplicate determinations at a significant difference \((p<0.05).\)

Table 2: Heavy metals of (Pb, Ni, Cr & Cd) of remedied and control soils

| Heavy Metals | Initial sample | Final sample | Initial control | Final control |
|--------------|----------------|--------------|----------------|--------------|
| Lead (Pb)   | 0.045±0.001    | 0.019±0.0170 | 0.120±0.0200   | 0.001±0.0000 |
| Nickel (Ni) | 0.0001±0.0000  | 0.0001±0.0000 | 0.0001±0.0000  | 0.001±0.0000 |
| Chromium (Cr)| 0.3750±1.0560 | 0.2820±0.1370 | 0.2820±0.0020  | 0.001±0.0000 |
| Cadmium (Cd)| 0.010±0.0000   | 0.0001±0.0000 | 0.001±0.0000   | 0.001±0.0000 |

Data are expressed as mean ± standard deviations of triplicate determinations. Values with the symbol * in the same row are not significantly different while others are significantly different at \(p<0.05).\)

Table 3: Heavy metals (Pb, Ni, Cr & Cd) of plants from remedied and control soils

| Plant     | Lead (Pb) | Chromium (Cr) | Cadmium (Cd) | Nickel (Ni) |
|-----------|-----------|---------------|--------------|-------------|
| Sample    | 0.16±0.001* | 0.55±0.01    | 0.006±0.001 | 0.0009±0.001* |
| Control   | 0.18±0.001* | 0.06±0.1    | 0.005±0.001 | 0.0008±0.001* |

Data are expressed as mean ± standard deviations of triplicate determinations. Values with the symbol * in the same column are significantly different while others are not significantly different at \(p<0.05).\)

Table 2 shows the rate at which the level of the heavy metals was reduced in the contaminated soils. Pb, Cd and Cr was significantly \((p<0.05)\) reduced after phytoremediation while there was no significant \((p<0.005)\) difference in Ni after phytoremediation. The Total Removal Percentage was calculated and it showed that Cd, Pb and Cr were removed by 90%, 67% and 39.2% respectively while there was no removal of Ni from the polluted soil. This means that the plant is a good hyperaccumulator of Pb, Cd, Ni and Cr which is in line with the study of Messou et al., 2012. The bioaccumulation factor \((B.F)\) of the heavy metals in the analyzed plants gave a ratio greater than 1 which suggests that *C. rotundus* is suitable for

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phytoextraction of heavy metals. Since there was no significant decrease in the level of Ni in the soil after phytoremediation, further studies should be done to ascertain what factors could be responsible for these outcome. Heavy metals were found to be below the guideline levels (DPR, 2018). Heavy metals exert toxic effects to soil by reducing the activity of soil microorganisms and inhibiting the physiological metabolism of plant even in low quantities (Singh and Kalamdhad, 2011). Accumulation of heavy metals along the food chain can have a deleterious effect on the health of humans and animals. Absorption and accumulation of heavy metals by plants is dependent on factors such as pH, moisture and availability of nutrient. Toxicity and phytotoxicity of heavy metals to plants leads to chlorosis, weak growth and reduced nutrient uptake.

The physical examination conducted on the soil showed that the taxonomic classification of the soil was loamy sand. Analytical results showed that the soil was slightly acidic with a mean value pH of (6.4, & 4.6) for experimental and control soil respectively which was found to be significantly different (p<0.05). The presence of trees which caused the littering of leaves could be responsible for the lower pH of the control soil which is in line with the study of Vinje (2018). Soil pH plays an important role in the adsorption of heavy metals and controls the hydrolysis and solubility of metal hydroxide, carbonate and phosphates (Tokalioglu et al., 2006). The Electrical conductivity (EC) of the soil had mean values of 264μS/cm and 209μS/cm for experimental and control soils respectively and no significant (p<0.05) difference was observed before and after phytoremediation. EC does not directly affect plant growth but can be used to indicate the amount of nutrients available for uptake by plants and the salinity levels of soils which can impede growth and microbial activity (USDA, 2011). Macro nutrients such as Nitrogen, Potassium and Phosphorus are necessary for plant growth and were found to be below the soil agricultural standards (HSE-ENV, 2004). The nutrients in the soil (Nitrates & Phosphates) were also seen to be affected by the phytoremediation process. The level of nitrates of experimental soil significantly (p<0.05) increased from 5.44mg/kg to 16.01mg/kg mean values after the phytoremediation process which is important for the uptake of other positive ions such as magnesium and calcium. However, the levels of phosphates dropped drastically from 13.31mg/kg to 6.5mg/kg mean values after the phytoremediation. No significant difference (p<0.05) was observed in the control soil before and after phytoremediation. The slopy terrain of the area could have resulted to the loss of phosphates through erosion and run-off as explained by USDA (2011). Potassium of the soil had mean values of 3.982mg/kg and 1.287mg/kg for experimental and control soils respectively and no significant (p<0.05) difference was observed after phytoremediation. Potassium is essential for the regulation of CO₂ intake by plants.

Phytoremediation efficiency of Cyperus rotundus: In the phytoremediation of a crude oil impacted soil, the optimal growth of the plant is important (Anyasi et al., 2018; Nwaichi and Chuku, 2018). However, before the optimal growth of the plant in the contaminated soil, it must be able to withstand the phytotoxic effects of the contamination (Anyasi and Atagana 2014; Tanhan et al. 2011). The toxic nature of the hydrocarbons is due to their volatility and hydrophobicity and this leads to low aeration and water infiltration which is needed for optimal growth of plants (Anyasi et al., 2018). The rate of growth of purple nutsedge at the contaminated site was high. This shows that the plant was able to withstand the toxic nature of the hydrocarbons found in the soil. However, it was observed that there was reduction in the height and root length of plants in contaminated samples after harvesting as compared to the control sample which corresponds to the findings made by Basumatary et al. (2013). Exposure to heavy metals can have an adverse effect on the growth of plants. Cd is the most toxic heavy metals for plants and can inhibit plant growth and reduce crop yield. Decolouration in leaves’ color can be associated with the presence of Cd which decreased the content of chlorophyll and carotenoids and increased non-phytochemical quenching as evidenced in the study of Garba et. al. (2018).

Conclusion: This study involved the evaluation of the viability of purple nutsedge to phytoremediate a hydrocarbon impacted soil. Purple nutsedge showed an ability to thrive in a hydrocarbon impacted soil and demonstrated good potentials for the phytoremediation of polycyclic aromatic hydrocarbons, and total petroleum hydrocarbon, and heavy metals like cadmium and lead. Obtained high bioaccumulation factor for studied heavy metals in purple nutsedge and the high hydrocarbons degradation efficiencies make it a suitable plant for such phytoremediation.

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