

# **Spectrophotometric analysis of phosphate concentration in agricultural soil samples and water samples using molybdenum blue method**

## **Introduction**

The chemistry of soil indicates that it contains several chemical components ranging from organic to inorganic compounds. Scientifically, it is defined as the outermost or weathered superficial layer of the earth's crust. They are consists mainly of decomposed parent rock material (organic matter), mineral (inorganic) matter, water, air and organisms (Weier, 1973). Soil is a complex system which includes Soil and water is one of the major constituents of the earth. Geographically, the earth crust contains 70% of water (hydrosphere) (Oladeji et al., 2016). From recent researches, scientists discovered that soil and water contains many

metals, chemical constituents or compounds. Some of these include phosphates, nitrates and sulphates amongst others. From soil plant absorbs water and solutes necessary for its continued well-being.

Phosphorus is an important nutrient that occurs widely in the environment. It is the key elements necessary for the growth of plants and animals and also, assists in metabolism. In the earth crust, it is the eleventh most abundant element on the surface of the earth and is most commonly found as phosphate (Chaube and Gupta, 1983).

Eutrophication of surface water is caused by increase level of phosphate in water and this increases the growth of algae thereby reduce the light intensity and level of dissolved oxygen in the water. Researches and recent report indicates the

high utilization of phosphates in detergents, fertilizers, sugar industries amongst others, thus the determination of phosphorus is important to water analyst and limnologist (Kharat and Pagar, 2009).

Due to these reports, various phosphate determination procedures titrimetry, complexogravimetry, colorimetry, atomic absorption spectroscopy, flow injection analysis, and spectrophotometry methods (Verma, 1988; Bøtker et al., 1994). In the attempt to quantitate the phosphate level, spectrophotometric methods have been preferred because it involves no extraction, sophisticated instruments and easily accessible. The commonest spectrophotometric includes molybdovanadate and ammonium molybdate methods (Vogel, 1999). In ammonium molybdate

spectrophotometric method, reducing agents such as tin(II) chloride, ascorbic acid and 1-amino-2-naphthol-4-sulfonic acid are often preferred. It also involves the formation of molybdophosphoric acid from ortho phosphate and an excess of molybdate in acidic solution followed by reduction to give molybdenum blue. The concentrations or phosphate levels in the sample is proportional to the intensity of colour produced and the absorbance is measured spectrophotometrically at a certain wavelength. In the attempt to reduce the health and environmental hazards by phosphate contamination, spectrophotometrically analysis of phosphate in soil and water sample is inevitable.

## **Materials and methods**

### **Chemicals and reagents**

All the chemicals and reagents used were analytical reagent grade unless otherwise stated and the solutions were prepared in double distilled water.

## **Instruments**

The absorbance were determined using ultraviolet visible spectrophotometer, model WPAS 104 UK with 1 cm matched quartz cells.

## **Preparation of standard solutions (stock solution)**

About 0.717 g of potassium dihydrogenphosphate ( $\text{KH}_2\text{PO}_4$ ) was accurately weighed and dissolved in double distilled water and the volume was made 500 mL in a volumetric flask. The working phosphate solutions of different concentrations were prepared by serial dilution of the stock solution (Oladeji et al., 2016).

## **Preparation of working solution of ammonium**

## **molybdate**

About 1.7081 g of ammonium molybdate was accurately weighed and dissolved in 100 mL of warm water; a milky colouration was observed, cooled and transferred into 250 mL volumetric flask.

### **Preparation of dilute hydrochloric acid**

A measured amount of concentrated hydrochloric acid solution was diluted and made up to mark in a 100 mL of volumetric flask.

### **Preparation of dilute trioxonitrate (V) acid**

A measured amount of 50 mL of concentrated trioxonitrate (V) acid solution was diluted and made up to mark in a 100 mL of volumetric flask.

### **Preparation of hydrazine hydrate**

About 0.125 g of hydrazine sulphate was accurately weighed and dissolved with double distilled water in

a clean beaker in a 100 mL standard flask and made up to calibrated mark.

### **Soil sample collection and preparation.**

Soil sample was taken from Goddex and Linkway Farms in Ogbomoso. The sample was dried for 10 days, sieved and stored for further analysis.

### **Water sample collection and preparation**

About 50 mL of water samples were collected in March, 2016 from three major rivers (Arowomole, Odoje and Alapata Rivers) located in the Southern part of Ogbomoso, Nigeria. The water samples were filtered using filter paper No. 42. The filtered samples was then stored in clean sample bottle.

### **Soil sample analysis for phosphates**

About 2 g of the soil sample was accurately weighed and placed in a 250 mL conical flask, 60

mL of HCl and HNO<sub>3</sub> (ratio of 3 to 1) was added and shaken in a mechanical shaker for 30 min. The digestion was done for 6 h:30 min. This was then filtered using filter paper Whatmann No. 41. About 15 mL of the filtered digested sample was taken, 3 mL of ammonium molybdate, 2 mL of hydrazine sulphate was added and kept in a water bath for 30 min. The blue colour observed was measured spectrophotometrically (Amponsah et al., 2014).

### **Water sample analysis for phosphates**

About 15 mL of the filtered water sample was accurately measured, 3 mL of ammonium molybdate, 2 mL of hydrazine sulphate was added and kept in a water bath for 30 min. The blue colour observed was measured spectrophotometrically (Amponsah et al., 2014).

## **Results and discussion**

### **Determination of maximum absorbance**

The absorption spectral data of the potassium dihydrogenphosphate used as standard against the reagent blank (water) exhibit the wave length of maximum absorbance around 740 nm.

### **Calibration curve for the standard (KH<sub>2</sub>PO<sub>4</sub>)**

The molybdenum blue method is based on the formation of phosphomolybdate. At effective analytical conditions, the amount of phosphates is proportional to the intensity of colour (blue) in the stock solution (KH<sub>2</sub>PO<sub>4</sub>). The calibration curve of the absorbance from the UV-visible spectrophotometer of concentrations 0.5-3.0 mg/L, respectively, is given below in Figure 1 from Table 1.

### **Phosphate analysis in soil and water samples**

The analytical results of phosphate analysis in soil and water samples are shown in Table 2. The phosphates concentration in soil and water was obtained by interpolating the calibration curve in Fig 1. The highest concentration of phosphate was obtained from Goddex Farms (SF1) with a concentration of 2.80 mg/L, the lowest being Odoje River with concentration (W2) of 1.480 mg/L. The uses of fertilizers, detergents, fungicides and even paints have been discovered to be the major pollutants in Ogbomoso, Nigeria. There is no health-based guideline proposed for phosphate consumption, though, medically high intake of phosphate can cause health problems like cancer, ulcers, kidney problems, brain damage and gastrointestinal effects. The use of fertilizers,

herbicides, fungicides in farm help improve the quality of farm produce. The environmental protection agency has a recommended limit for phosphates levels in soil and water bodies. For streams and rivers the Environmental Protection Agency recommended limit is

1.1 mg/L. The following recommendations are also given for phosphate consumptions: total phosphorus; no more than 0.1 mg/L for streams which do not empty into reservoir, 0.05 mg/L for streams discharging into reservoirs and 0.025 mg/L for reservoirs (Oladeji et al., 2016). It is recommended that health authorities be notified of sources of drinking. Man made sources of phosphate include human sewage, agricultural run-off from crops, sewage from animal feedlots, pulp and paper industry, vegetable and fruit processing, chemical and fertilizer manufacturing, and detergents (McCoy, 2011).

1.2 The highest concentration of phosphates at Goddex Farms (SF1) could be attributed to the mixed farming activities in the farm. There was

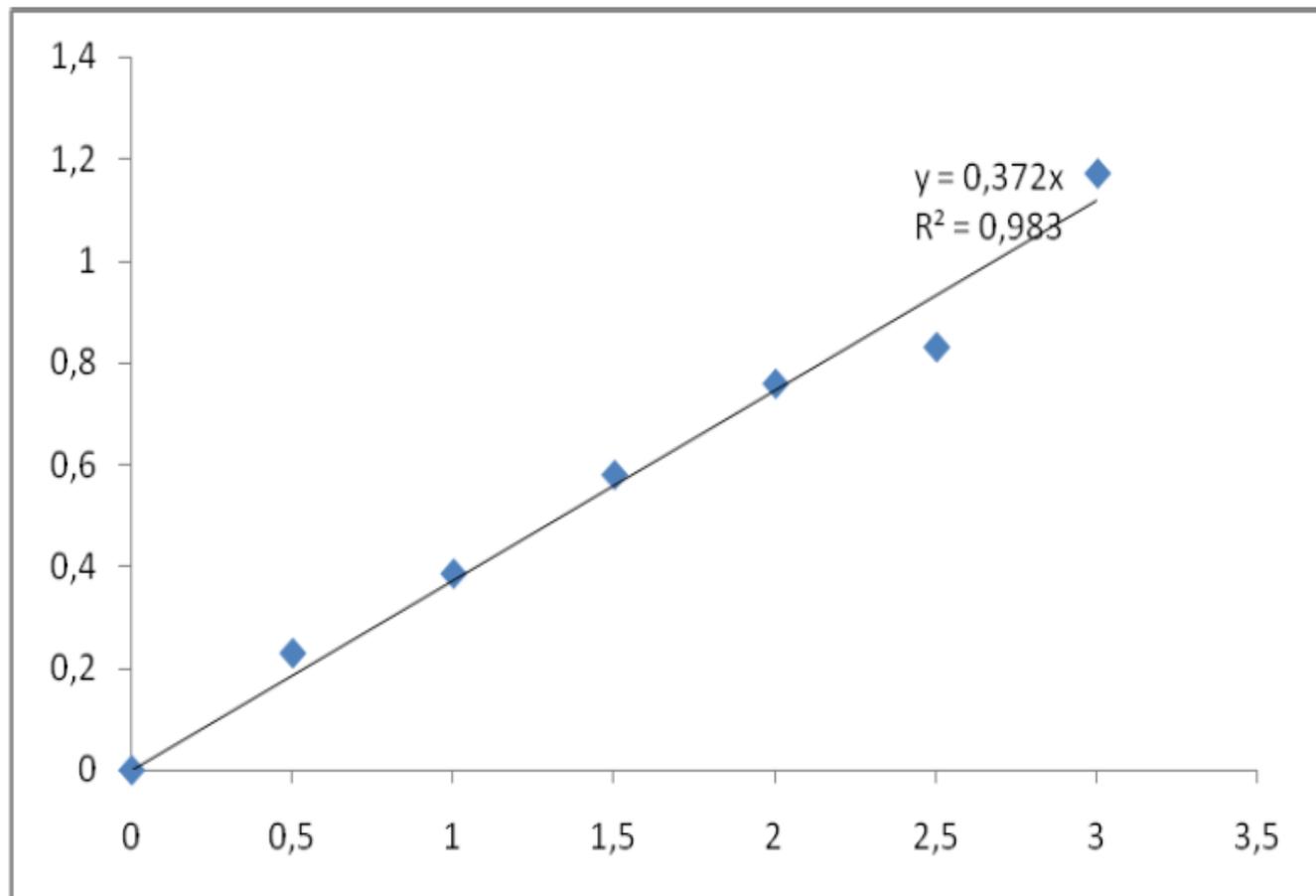
an indication that the use of manure (cow dung) and inorganic fertilizers (phosphate fertilizers) was the usual practice in this farm. The use of fertilizers in agriculture is paramount when compare with herbicides, fungicides and nematocides uses. The high level of phosphates in soil samples taken from maize fields (SF1 and SF2) could be due to the use of phosphates fertilizers and have higher level when compared to soil samples taken from lands sprayed with phosphates fungicides. In water samples, the high level of phosphates was recorded in Arowomole River (W1). This could be due to the industrial, human and farming activities in the community. The dumping of refuse, sewage, human waste could also be a major factor. The

laundry activities, car-washing, block industry and also farming activities are also major contributing factors (McCoy, 2011). Also, in Alapata River, W3, residents close to this water source uses phosphate containing detergents.

**Table 1.** The absorbance,  $A_T$  obtained from different concentration (mg/L) of potassium dihydrogenphosphate.

	$C_T$	$A_T$
1.	0.00	0.000
2.	0.50	0.230
3.	1.00	0.386
4.	1.50	0.580
5.	2.00	0.759
6.	2.50	0.831
7.	3.00	1.172

Where  $C_T$  = concentration (mg/L);  $A_T$  = absorbance.



**Figure 1.** The calibration curve for the standard solution ( $KH_2PO_4$ )

## Conclusion

The spectrophotometric analysis using ammonium molybdate as reductants proved effective analytical for the determination of phosphate. It involves the reduction of phosphomolybdate by potassium dihydrogenphosphate. The phosphate concentration in the samples was measured at 740 nm. The phosphates levels are high in all the samples when compared to the Environmental Protection Agency. This could be due to natural such as erosion and weathering, agricultural and also human activities. In the attempt to prevent the effects of high phosphates, the use of phosphate containing materials such be reduce and the people in these area should be enlighten on the health effects on these phosphate levels can cause.