



## ASSESSMENT OF SOLID MINERALS AND SOME ELEMENTAL OXIDES IN SOIL OF KWAYA KUSAR LOCAL GOVERNMENT AREA OF BORNO STATE, NORTHEASTERN NIGERIA.

N. Jonathan<sup>1</sup>, H. M. Maina<sup>2</sup>, P. Handawa<sup>3</sup>

<sup>1</sup>Department of Chemistry, Adamawa State College of Education Hong, Nigeria

<sup>2</sup>Department of Chemistry, Modibbo Adama University, Yola, Nigeria

<sup>3</sup>Department of Chemistry, Adamawa State College of Education Hong, Nigeria

Correspondence email: [nelsonjonathanbaba@gmail.com](mailto:nelsonjonathanbaba@gmail.com).

### ABSTRACT

Soil samples were collected from (5) locations, wandali, kwaya kusar, gashina, kubagaya and guwal in kwaya kusar Local Government Areas, Borno State Northeastern Nigeria using Complete Randomized Block Design (CRBD). The coordinate of each location were taken, the soil samples were homogenized and dry at 500°C for 12Hrs, then crushed using iron pestle and mortar and sieved using 150micro meshes, the samples were analyzed analytically using X-Ray Diffraction (XRD) for minerals, the results reveals the presence of Kaolinite, Microcline, Quartz, Albite, Biotite, Hematite, Chlorite and Montmorillonite with Quartz been dominant across all the area studied. Major elemental oxides such as SiO<sub>2</sub>, Fe<sub>2</sub>O<sub>3</sub>, Al<sub>2</sub>O<sub>3</sub>, CaO, TiO<sub>2</sub>, MgO, K<sub>2</sub>O, Na<sub>2</sub>O, MnO and SO<sub>3</sub>. Were also analyzed using X-Ray Fluorescence (XRF) Spectroscopy. SiO<sub>2</sub> was observed to be dominant followed by Al<sub>2</sub>O<sub>3</sub> and Fe<sub>2</sub>O<sub>3</sub> across all the areas studied.

**Keywords:** Assessment, Soil, Solid Mineral, XRD, XRF

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### INTRODUCTION

Geological composition of Biu plateau varies greatly, which provides great varieties of topographical forms, featuring rugged hills of granite, sandstone and volcanic plugs. The plateau developed on sedimentary and volcanic rocks [1]. The breaking down of rocks into smaller pieces under some condition is known as weathering, which leads to the breaking down of the minerals into their various elemental forms and that lead to an increase in the concentration of these elements in the soil [2]. According to Nelson [3], each rock or mineral have a particular chemical makeup, which is combination of two or more elements? Eg Calcite ( $\text{CaCO}_3$ ), Bauxite ( $\text{Al}(\text{OH})_3$ ), etc, while some minerals are made up of single elements, eg gold (Au), Diamond (C) and silver (Ag) [4]. Some elements that are found in combined state in the earth crust and mantle are listed below in their predominantly order, Oxygen 46.6%, Silicon 27.7%, Aluminium 8.1%, Iron 5.0%, Calcium 3.6%, Sodium 2.8%, Potassium 2.6 % Magnesium 2.1% and all others put together 1.5%. [3]. The elements are combined in various proportions to form the rocks of the earth which are the parent materials with differences in their structure, composition and rate of decomposition [5]. It was also observed by Badmus. [6] that deep chemical weathering of rocks in hot humid and tropical climate promote mineral enrichment. It was also observed that dissolution of minerals by weathering process lead to the apparent enrichment of these elements in soil [7]. Some of these oxides have strong oxygen to other elements bond over a wide range of conditions e.g SiO which make their compound very strong and stable [8].

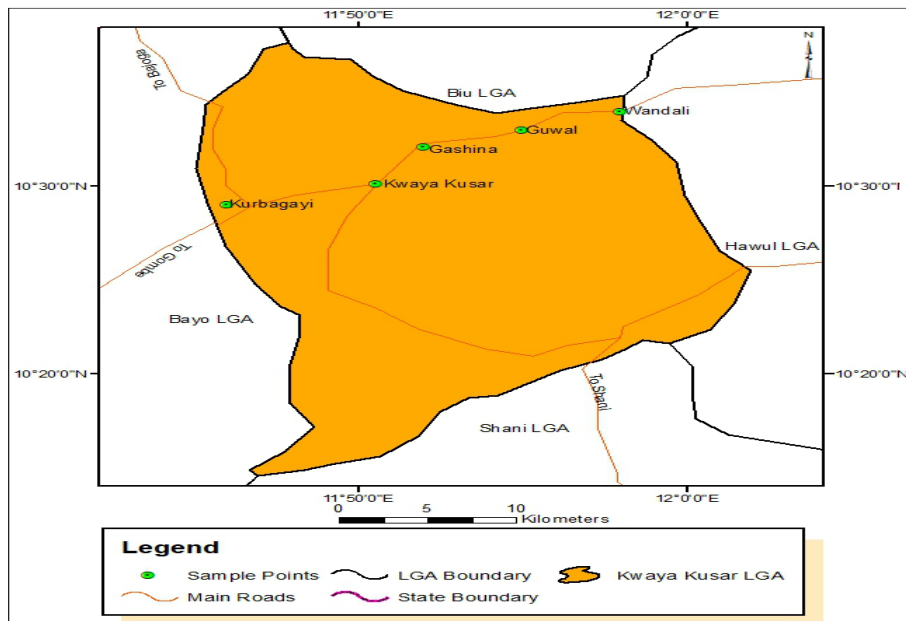
The mineral oxides typically present in soil comprises oxide, hydroxide, oxyhydroxide and hydrated oxides of Si, Fe, Mn, Al and Ti which are predominantly inherited from primary rocks minerals. Most oxide form in soil are metal cations  $\text{Fe}^{2+}$ ,  $\text{Mn}^{2+}$ ,  $\text{Ti}^{4+}$ ,  $\text{Al}^{3+}$  and  $\text{Si}^{4+}$  are release from silicate by weathering. The divalent cation Mn and Fe oxide hydrolyze and precipitate almost exclusively and oxides mineral  $\text{Al}^{3+}$  and  $\text{Si}^{4+}$  have strong tendency to form secondary alumino-silicate clay minerals. The formation of Al and Si oxide requires special condition [9]. All oxides have very low solubility at common soil PH, and are therefore enriched during pedogenesis. Highly weathered soil which has lost a substantial part of the alkali such as Si may contain as much as 50% metal oxide. Mn, Fe and Al may exhibit a high surface area with reactive surface site which strongly bind oxyanion and metal cation, thereby impacting the availability and mobility of plant and toxic metals [10]. The basic structural unit of Fe, Mn, Al and Ti oxides are cationic metal centers bound to six oxygen in relation to the metal cation.

Many oxide minerals such as geothnrite. Silicon is bounded to four oxygen forming a tetrahedron; these tetrahedrons are connecting via corner only [11].

Geologists defined mineral as crystalline solid which occurs naturally with specific chemical composition, structural arrangement of component atoms and physical properties [12]. According to Pidwirny [13], there are specific elements or combinations of elements which are known to be associated with specific types of minerals, therefore it is possible to evaluate the elements which are found in a given area [14]. Considering the diversity of mineral groups in nature, the initial mineralogical and chemical examination of mineral ores can be used to indicate the suitability of minerals for different applications [15]. According to Pipkin and Trend [14], minerals are originated through the process of rock formation occurring in nature. A common phenomenon occurring for the origin is the cooling of molten magma (rock materials) from the earth's crust. The magma that reaches the surface is called lava, as this material crystallized and grew into rock [16].

According to Alexander *et al.* [2], the mineral resource potential of an area is a measurement of the likelihood of occurrence of valuable minerals that are valuable in feature. It is against this background that the research is motivated to investigate quantity and quality of minerals and elemental composition in soil of Kwaya Kusar local government area. Biu Plateau has not been extensively studied but some geologists have made an appreciable attempt in identifying the types of rocks and soil in the study area. This work investigates the types of minerals and elemental composition in soil of Kwaya Kusar local government area which is part of Biu Plateau using some analytical techniques.

### **Study area**



### Map of kwaya kusar local government area

#### Soil sampling

Surface soil was taken from 0 cm to 50 cm depths. A soil probe was used for collecting soil samples as described by Alexander *et al.* [17]. A spade was used to dig down the soil profiles. Composite sample of five (5) samples across the study area was obtained. The soils collected was mixed thoroughly and put into well label polythene bags, the soil samples were homogenized and dry at 500<sup>0</sup> C for 12 hours. The soil samples were crushed using iron pestle and mortar then sieve by using 150 micro meshes and ready for analytical assessment of the in the laboratory.

#### Determination of Major Elemental oxides from Soils Samples using X-Ray Fluorescence (XRF) Spectroscopy

Major elemental oxides in soil were determined using X-ray fluorescence spectroscopic method Axion cement Pan Analytical model. 20g of each powder soil was fuse with 0.40g of stearic acid in 20ml platinum crucibles and pass with Herzog D4500 model hydraulic press to form pellet. These fused buttons was X- rayed and counted, and the major elemental oxides was determined (Swanson and Huffman, 1976)[18]. An annular 25 mCi 109Cd was used as an excitation source, which emit Ag-K X-ray (22.1KeV) in which case, the element with lower characteristics excitation energies was accessible for detection in the samples.

Quantitative analysis of samples was carried out using the emission transmission (E – T) method, for which a number of quantifications was developed and apply; these quantification methods provide different approaches to correct the matrix absorption as well as enhancement effect. In this work, quantification was carry out using a modified version of Emission-Transmission (E-T) method and the use of pure target material (Mo) was employ to measure the absorption factor in the samples. The (Mo) target served as monochromatic X-ray which was excited through the sample by primary radiation and penetrate the sample on the way to detector. The absorption factor was experimentally determined which was used by the program in the quantification of concentration of the elements. In addition, the contribution to the Mo-K peak intensity by the Zr-K was subtracted from each sample. The spectra for the sample were collected for 3000s with the 109 Cd source and the spectra were evaluated using the AXIL-QXAS program. 109Cd was used for analysis of the metal oxides.

### Statistical analysis

All analysis was performed in triplicates. Result was presented as mean  $\pm$  SD. Statistical significant Figure were establish using one-way analysis of variance. (ANOVA). Mean was separated according to Duncans Multiple Range Analysis.  $P \leq 0.05$  was considered statistical significant

### Results

The results of some major elemental oxides in soil samples of kwaya kusar local government area is shown in table 1 bellow. The quantitative analysis reveals that  $\text{SiO}_2$  is present in the samples of all locations with relatively high concentration than any other metal oxide, with concentration ranges from  $35.72 \pm 26.6$  % in guwal to  $60.62 \pm 1.70$  % in kwaya kusar with the total mean of 46.32 % across the study area,  $\text{Fe}_2\text{O}_3$  and  $\text{Al}_2\text{O}_3$  was observed across all the areas studied with concentration ranges from  $8.78 \pm 1.52$  % in kwaya kusar and  $16.64 \pm 0.42$  % in Gashina to  $27.50 \pm 1.09$  % in kubagaya and  $20.46 \pm 0.24$  % in Kubagaya respectively. CaO was observed across all the areas studied with concentration ranges from  $1.96 \pm 1.05$  % in wandali to  $7.46 \pm 1.55$  % in Guwal with the mean concentration of 3.99 %.  $\text{TiO}_2$ , MgO and  $\text{K}_2\text{O}$  occurs

across all the areas studied with concentration ranging from  $2.31 \pm 0.71$  % in guwal,  $1.60 \pm 0.19$  % in Gashina and  $1.45 \pm 0.36$  % in Kubagaya respectively to  $4.41 \pm 0.36$  % in Gashina  $2.53 \pm 0.25$  % in kubagaya and  $7.13 \pm 1.89$  % in kwaya kusar respectively.  $\text{Na}_2\text{O}$  was observed only in wandali with the concentration of  $0.23 \pm 0.21$  %,  $\text{MnO}$  and  $\text{SO}_3$  was detected across all the area studied with relatively low concentration. the concentration ranging from  $0.23 \pm 0.02$  % in kwaya kusar and  $0.43 \pm 0.27$  % in wandali to  $0.54 \pm 0.12$  % in kubagaya and  $0.82 \pm 0.27$  % respectively.  $\text{LOI}$  was observed to vary from one location to another with relatively low concentration of less than 2% across the areas studied.



**Table 1: Present the mean Percentage Concentration of Some Major Elemental Oxides in Soils of Kwaya Kusar Local Government Area**

| Location                  | SiO <sub>2</sub>           | Fe <sub>2</sub> O <sub>3</sub> | Al <sub>2</sub> O <sub>3</sub> | CaO                      | TiO <sub>2</sub>          | MgO                      | K <sub>2</sub> O          | Na <sub>2</sub> O        | MnO                      | SO <sub>3</sub>          | LOI   |
|---------------------------|----------------------------|--------------------------------|--------------------------------|--------------------------|---------------------------|--------------------------|---------------------------|--------------------------|--------------------------|--------------------------|-------|
| Wandali                   | 41.75 ± 0.72 <sup>ab</sup> | 19.56 ± 1.30 <sup>b</sup>      | 19.26 ± 2.11 <sup>ab</sup>     | 1.96 ± 1.05 <sup>a</sup> | 3.31 ± 1.04 <sup>ab</sup> | 1.67 ± 0.59 <sup>a</sup> | 5.23 ± 1.02 <sup>bc</sup> | 0.23 ± 0.21 <sup>b</sup> | 0.49 ± 0.40 <sup>a</sup> | 0.43 ± 0.27 <sup>a</sup> | 0.17  |
| KwayaKusar                | 60.62 ± 1.70 <sup>b</sup>  | 8.79 ± 1.52 <sup>a</sup>       | 18.99 ± 2.06 <sup>ab</sup>     | 2.88 ± 1.51 <sup>a</sup> | 2.63 ± 0.74 <sup>a</sup>  | 2.31 ± 1.29 <sup>a</sup> | 7.13 ± 1.89 <sup>c</sup>  | 0.00 ± 0.00 <sup>a</sup> | 0.23 ± 0.07 <sup>a</sup> | 0.47 ± 0.20 <sup>a</sup> | 1.11  |
| Gashina                   | 54.06 ± 0.32 <sup>ab</sup> | 17.73 ± 0.49 <sup>b</sup>      | 16.64 ± 0.42 <sup>a</sup>      | 1.96 ± 0.10 <sup>a</sup> | 4.41 ± 0.36 <sup>b</sup>  | 1.60 ± 0.19 <sup>a</sup> | 2.93 ± 0.15 <sup>ab</sup> | 0.00 ± 0.00 <sup>a</sup> | 0.44 ± 0.07 <sup>a</sup> | 0.60 ± 0.17 <sup>a</sup> | 1.19  |
| Guwal                     | 35.72±26.6 <sup>a</sup>    | 17.55±1.42 <sup>b</sup>        | 19.48±1.41 <sup>ab</sup>       | 7.46±1.55 <sup>b</sup>   | 2.31±0.71 <sup>a</sup>    | 1.67±0.61 <sup>a</sup>   | 7.60±2.16 <sup>c</sup>    | 0.00±0.00 <sup>a</sup>   | 0.39±0.12 <sup>a</sup>   | 0.82±0.27 <sup>a</sup>   | 1.07  |
| kubagaya                  | 38.99 ± 0.22 <sup>ab</sup> | 27.50 ± 1.09 <sup>c</sup>      | 20.46 ± 0.24 <sup>b</sup>      | 5.70 ± 0.60 <sup>b</sup> | 3.45 ± 0.38 <sup>ab</sup> | 2.53 ± 0.25 <sup>a</sup> | 1.45 ± 0.17 <sup>a</sup>  | 0.00 ± 0.00 <sup>a</sup> | 0.54 ± 0.12 <sup>a</sup> | 0.48 ± 0.15 <sup>a</sup> | 10.92 |
| <b>Range</b>              | 4.92 .62.17                | 7.13-28.40                     | 16.16-21.3                     | 1.17-9.13                | 1.74-4.82                 | 1.00-3.80                | 1.30-9.28                 | 0.00-0.41                | 0.17-0.94                | 0.19-1.02                | 0.28  |
| <b>Total Mean</b>         | 46.23                      | 18.22                          | 18.97                          | 3.99                     | 3.22                      | 1.95                     | 4.69                      | 0.13                     | 0.42                     | 0.56                     | 0.12  |
| <b>Standard Deviation</b> | 14.11                      | 6.25                           | 1.81                           | 2.47                     | 0.95                      | 0.72                     | 2.64                      | 0.05                     | 0.20                     | 0.24                     | 0.49  |
| <b>C.V. (%)</b>           | 30.52                      | 34.30                          | 9.54                           | 61.90                    | 29.5                      | 36.92                    | 56.29                     | 38.46                    | 47.62                    | 42.86                    | 2.14  |

Mean ± Standard Deviation within a column with different superscript letters are significantly different at  $P \leq 0.05$  according to Duncan Multiple Range Test. CV = Co-efficient of Variation

### **Mineral Identified in Soil of Kwaya Kusar Using X – Ray Diffraction Ppectroscopy (XRD)**

The mineral identified in soil of kwaya kusar Local Government Area are presented in Table 2. The result from Kubagaya revealed the presence of Chlinochlore, Hematite and montmorillonite with the percentage composition of 3 %, 86 % and 11 % respectively as shown in Figure 1. The sample of wandali showed the presence of 30 % Augite, 36 % quartz, 12 % Hematite and 22 % kaolinite as shown in the Figure 2 bellow. The mineral recorded in the soil of Guwal were 33 % Ablite, 41 % microcline, 10 % Biotite and 16 % Quarts as shown in the Figure 3: The soils of Gashina contain microcline and Quarts with percentage composition of 21 % and 79% respectively as shown in the Figure 4 below. The mineral identified in the soil of kwaya kusar were kaolinite, microcline and Quarts with the percentage composition of 13 %, 44 % and 43 % respectively as show on Figure 5.



**Table 16: Mineral Identified in Soil of Kwaya Kusar Local Government Area Using X – Ray Diffraction Spectroscopy (XRD)**

| Location          | Minerals        | Compound name                                         | Chemical formula                                                                      | Crystal system |
|-------------------|-----------------|-------------------------------------------------------|---------------------------------------------------------------------------------------|----------------|
| <b>Kwayakusar</b> | Kaolinite       | Aluminium Silicate Hydroxide                          | $\text{Al}_2\text{Si}_2\text{O}_5(\text{OH})_4$                                       | Anorthic       |
|                   | Microchine      | Potassium Aluminum silicate                           | $\text{KAlSi}_3\text{O}_8$                                                            | Anorthic       |
|                   | Quartz          | Silicon Oxide                                         | $\text{SiO}_2$                                                                        | Hexagonal      |
| <b>Gashina</b>    | Microchine      | Aluminium Silicate Hydroxide                          | $\text{KAlSi}_3\text{O}_8$                                                            | Anorthic       |
|                   | Quartz          | Silicon Oxide                                         | $\text{SiO}_2$                                                                        | Hexagonal      |
| <b>Guwal</b>      | Albite          | Sodium Aluminium Silicate                             | $\text{NaAlSi}_3\text{O}_8$                                                           | Anorthic       |
|                   | Biotite         | Potassium ironAluminiumMagnesium Silicate             | $\text{K}(\text{Mg,Fe})_3(\text{AlSiO}_{810})$                                        | Anorthic       |
|                   | Microchine      | Potassium Aluminium Silicate                          | $\text{KAlSi}_3\text{O}_8$                                                            | Anorthic       |
|                   | Quartz          | Silicon Oxide                                         | $\text{SiO}_2$                                                                        | Hexagonal      |
| <b>Wandali</b>    | Augite          | Calcium iron magnesium silicate                       | $\text{Ca}(\text{Mg,Fe})\text{Si}_2\text{O}_6$                                        | Monochinic     |
|                   | Hematite        | Iron oxide                                            | $\text{Fe}_2\text{O}_3$                                                               | Hexagonal      |
|                   | Kaolinite       | Aluminium Silicate Hydroxide                          | $\text{Al}_2\text{Si}_2\text{O}_5(\text{OH})_4$                                       | Hexagonal      |
|                   | Quartz          | Silicon Oxide                                         | $\text{SiO}_2$                                                                        | Hexagonal      |
| <b>Kubagaya</b>   | Chinochore      |                                                       |                                                                                       | Anorthic       |
|                   | Hematite        | Iron oxide                                            | $\text{Fe}_2\text{O}_3$                                                               | Hexagonal      |
|                   | Montmorillonite | Sodium magnesium Aluminium Silicate Hydroxide Hydrate | $\text{Na}_2(\text{AlMg})_2\text{Si}_4\text{O}_{10}\text{OH}_{2.6}\text{H}_2\text{O}$ | Monochinic     |

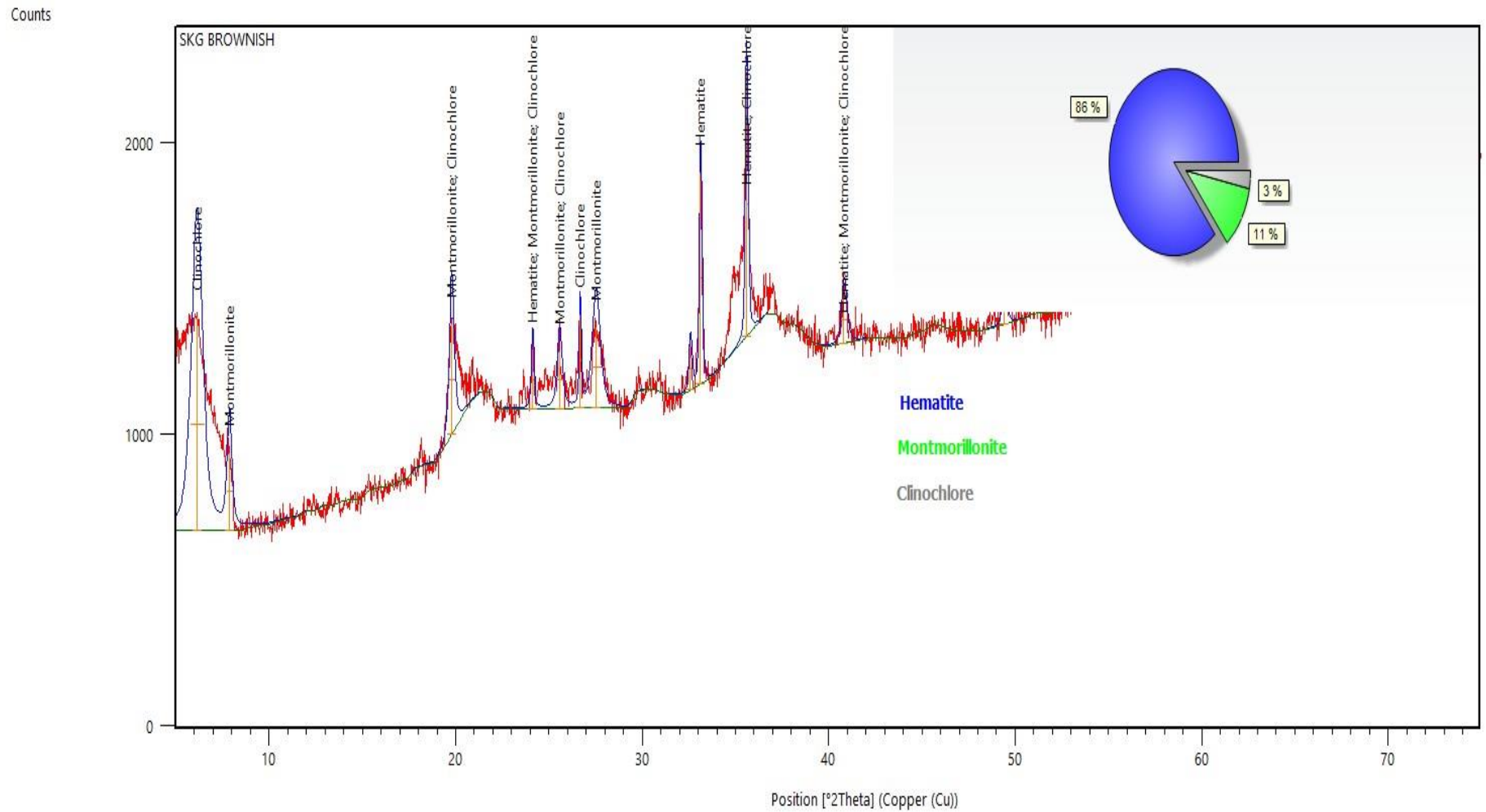


Figure. 1 Percentage composition of minerals in Soil of Kubagaya

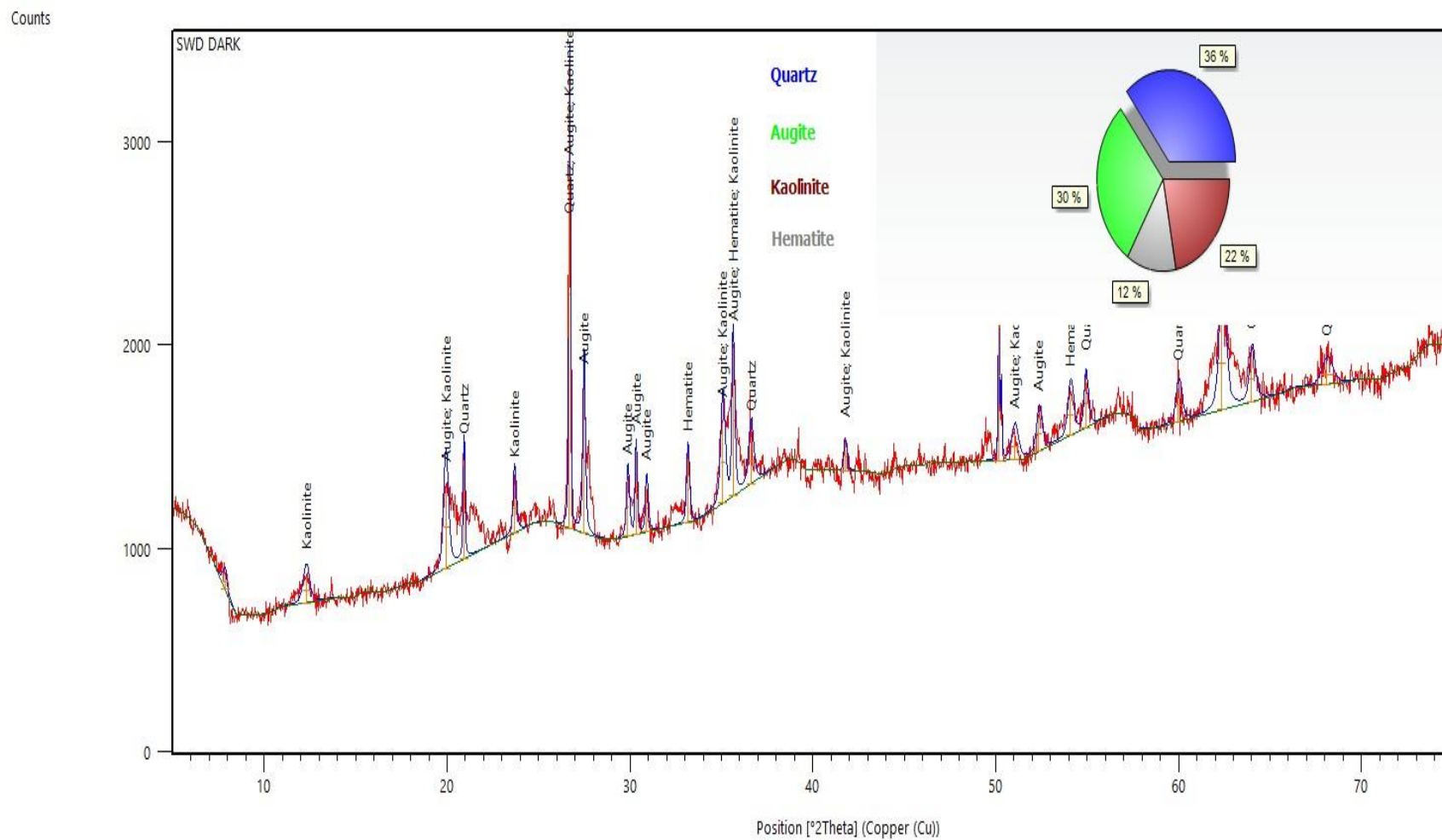


Figure. 2 Percentage compositions of minerals in Soil of Wandali.

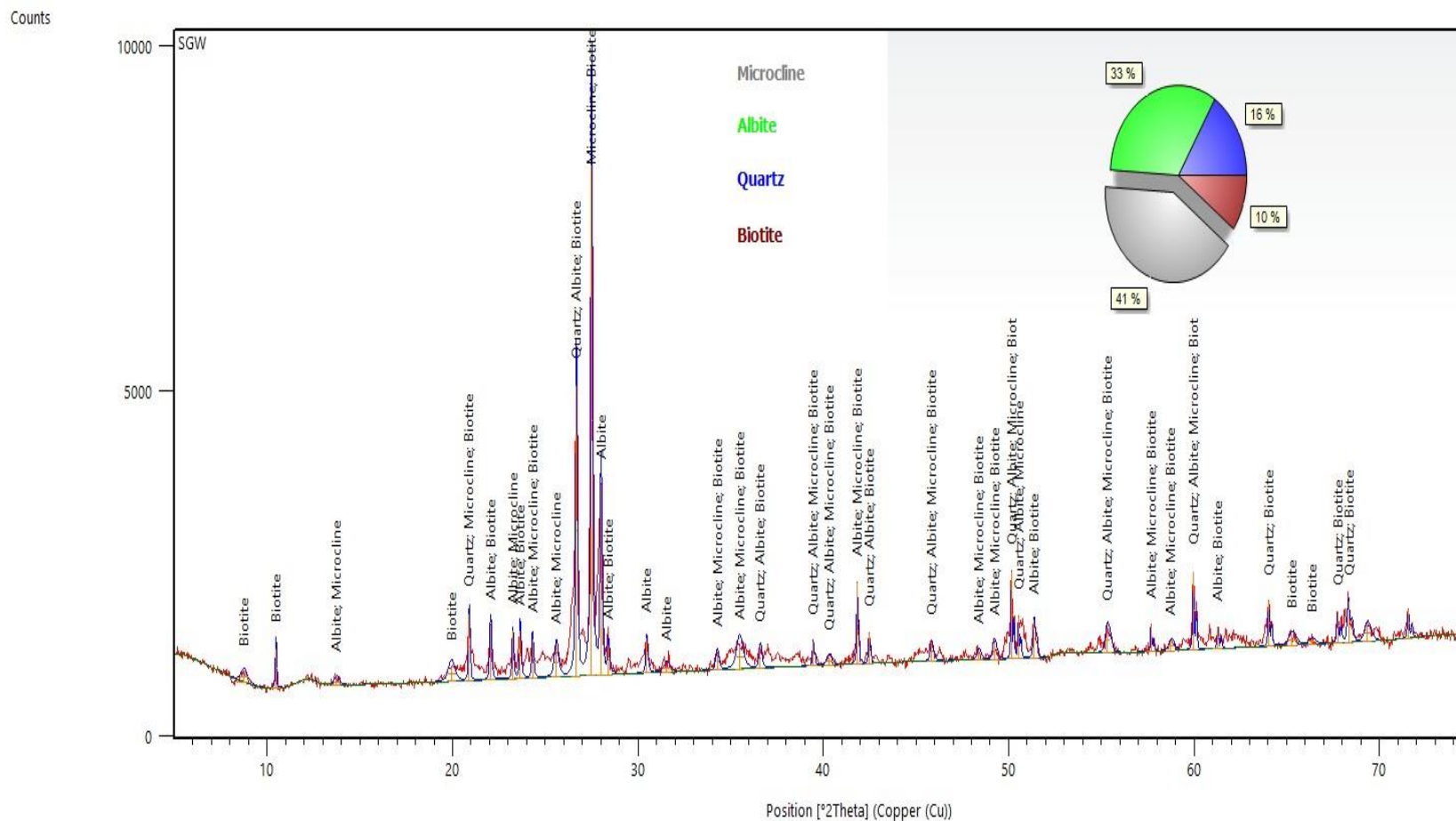


Figure. 3: Percentage composition of minerals in Soil of Guwal.

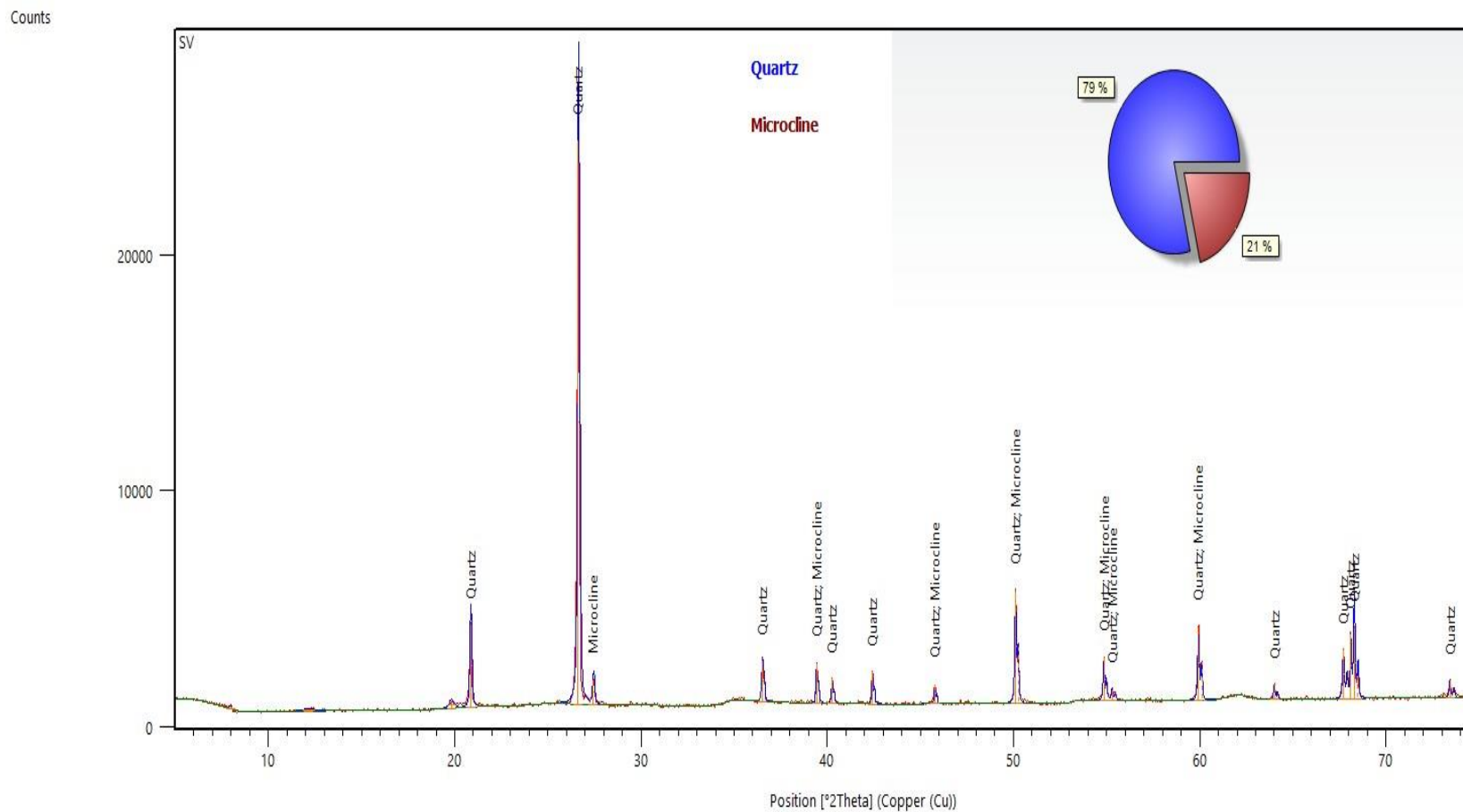


Figure. 4: Percentage composition of minerals in Soil of Gashina.

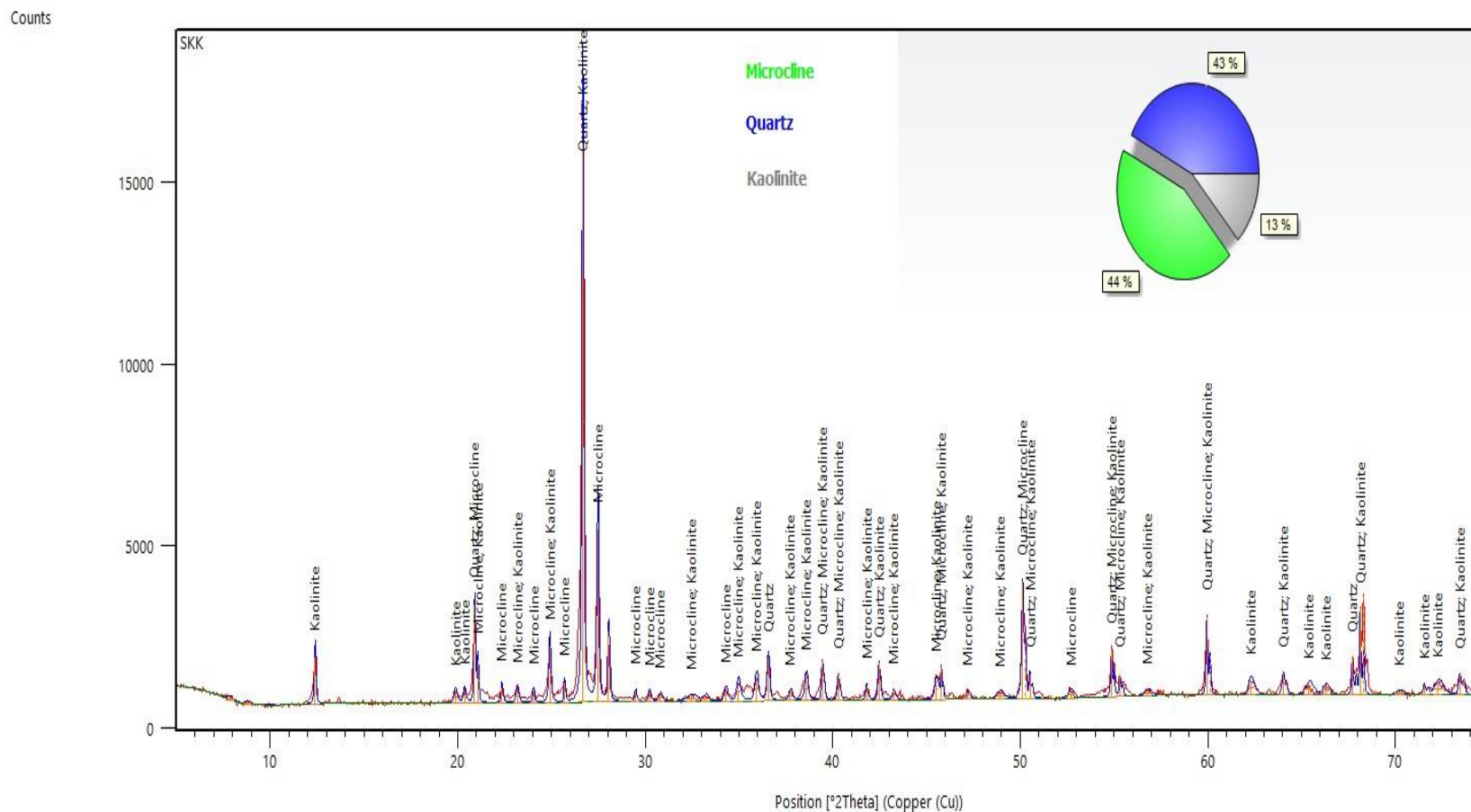


Figure. 5: Percentage composition of minerals in Soil of Kwaya kusar

## DISCUSSION

Base on the information given in the table above,  $\text{SiO}_2$  was observed to be relatively higher than Any other elemental oxides, similar results was obtained by Osemeahon *et al.* [19], in a research titled survey of solid minerals in rocks of Ditera and Waltadi in song local government area of Adamawa state. The high concentration of  $\text{SiO}_2$  could be due to the fact that silicon and oxygen are the most abundant element in the earth crust and mantle, they are strongly bonded over a wide range of condition which make their compound to be stable, as recorded by Alexander *et al.* [2]. This study confirm that the soil of the study area have high percentage of silicate minerals such as quartz, biotite and microcline as reported by Tadzabia *et al.* [20].  $\text{Al}_2\text{O}_3$  was also observed to be relatively high in concentration; this study is in line with the research conducted by Ako *et al.* [21]. The high concentration could be due to the fact that  $\text{Al}_2\text{O}_3$  are less mobile elements in soil in which they are lock up especially in alimino-silicate minerals as reported by Penuel *et al.* [22]. The relatively high concentration of  $\text{Al}_2\text{O}_3$  signals the reason why bytownite, phlogopite, microcline, Kaolinite and chlinichlore was detected in the soil as reported by Magili and Maina, [4].  $\text{Fe}_2\text{O}_3$  was observed across all the area studied with relatively high concentration, similar results were also recorded by Alexander *et al.* [2] and Osemeahon *et al.* [19]. The high concentration of  $\text{Fe}_2\text{O}_3$  across all location could be due to the fact that, Fe is relatively abundant in earth crust, the presence of  $\text{Fe}_2\text{O}_3$  justify the existence of minerals hematite and albite in the study area as reported by Angus [23].  $\text{TiO}_2$  was observed across all the study area with relatively low concentration which could be due to leaching or due to ionic exchange or reaction that took place in the soil. The low concentration of  $\text{TiO}_2$  justifies the absence of minerals such as Barite in the soil of the area studied.  $\text{K}_2\text{O}$ ,  $\text{MgO}$  and  $\text{MnO}$  were observed in all the samples studied with relatively low concentration but the concentration varies from one location to another, the variation could be due to complex distribution of elements in the soil [24]. In Rubio *et al* [25] reported that the variation from one location to another could be due to instability of the carbonate and organic materials content of the soil samples.  $\text{Na}_2\text{O}$  and  $\text{SO}_3$  were observed to have low concentration and does not occurs in all sample , it could be due to leaching in the soil and instability of their chemical bond.

Lol was observed to vary from one location to another with very low concentration of less than 1%, this indicate that the minerals does not contain decomposable organic matter such as carbonate etc.

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