Study of Radioactivity and its Impact Activity in the District of al-Qasim / Babil Province

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Abstract: In this research study the level of background radiation for four selected areas of soil the district of al-Qasim/Babil province by using the counting system and electronic analysis to detect radiation gamma and the detector NaI(Tl) radionuclide resulting from the dissolution chains of 238U and 232Th (214Bi, 214Pb 234Th 226 Ra, 212Bi) are produced in addition to radionuclide potassium individual (40K), the paper also counts the radioactivity of each radionuclide and its concentration within the Sample unit (mg/Kg), and calculate the ratios of natural uranium, depleted uranium, the creation of ratio between 238U/235U, Where results proved the presence of varying proportions of depleted uranium contamination of the sites (1,3,4), ranging between (0.005466_0.006456), which is located within the limits allowed by the International Atomic Energy Agency, (0.0046).

Keywords: Radioactivity, Radiation Gamma, Natural Uranium

1. Introduction

The typical radioactivity researches for many places in Iraq were done by many workers for a soil samples Al_Saji in 1999 [1],studied the effect of the radioactive wopens on the water south of Iraq and the found the ²²⁶Ra isotope. In 2000, Nashwan Shewket [2] the radioactive pollution in Neinow province and found ²²⁶Ra and ¹³⁷Cs concentration more than the natural level. In 2001 [3], Al

Kinany studied the radioactivity pollution with the depleted uranium in a Basra province and he calculated the ²³⁵U / ²³⁸ U, the results shows a pollution with different ratios . Al_Gurabi 2002 [4] , show that on increase in the radioactive pollution in the south of Iraq . In 2003 [5], Murtada Shaker studied the radioactivity for the water in Babylon province and the results shows on existence some isotopes belong to ²³⁸U and ²³²Th series . In 2004 [6] Al_Baiaty studied the concentration of the depleted uranium and the radioactive pollution in the soil samples in Al Tamim province. In 2005 [7], Al_Azawi calculated the ²³⁴Th and ²³⁵U concentrations and the ²³⁵U / ²³⁸U ratio in Al Basra south of Iraq.

The aim of the present work is study radioactive background in different places in district of al-Qasim / Babil province, middle of Iraq and determine the type and the concentration for the radioactive nuclides.

2. The Nuclear Detector System

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In the present work , the nuclear detection system , figure (1) , type (UCS 30) $\,$ (Spectrum Techniques LLC) with NaI(Tl) size of crystal ($3.8\times2.5\,$ cm)

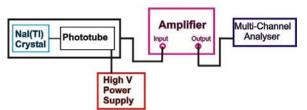


Figure 1: Detection system used

The NaI(Tl) detector consist of two pants, the NaI(Tl) crystal and the photo multiplier, the detection system consist of two amplifiers, pave and main, the job of this amplifiers is shaping the electronic signal, amplification and release the electronic noise. The electronic signals go to the multichannel analyzer and show the result as an energy spectrum as shown in figure (2) [8].



Figure 2: Parts of the system Detect

3. The Detection System Calibration

Two calibration were done for the gamma ray nuclear detection system, the first for the detection efficiency and the second is the energy, by using a standrad radioactive sources putted in a container of 0.25 L volume. The calculate efficiency (ξ) is given by [9].

$$\xi = \frac{N/T}{A \cdot I \gamma} \qquad \dots (1)$$

N: count rate under photo peak position.

T: time measurement.

A: activity of radioactive sources using of calibration.

 I_{γ} : relative intensity of each energy source of the energies of the radioactive

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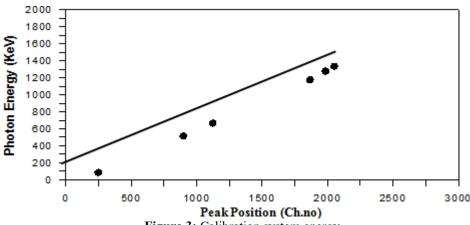


Figure 3: Calibration system energy

And the energy resolution (ER) is given by, fig. (3), by using 60 Co source [9].

$$E.R = \frac{F.H.W.M}{Ch.no.} \times 100\% \dots (2)$$

F.W.H.M: Full width high maximum. Ch.no: photo peak position.

4. The soil samples preparation

Four selected places from different region in Babylon province, middle of Iraq, figure (4) show the map of Babylon province. The soil samples prepared for the detection.

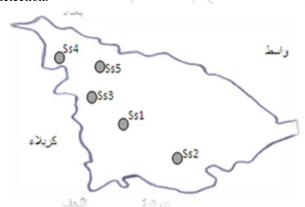


Figure 4: The region selected in the studied

5. The radioactive samples test

The radiation background (B.G) were calculated to sub it from the energy spectrum for the studied sample as in following equation [10].

$$B.G(Bq) = \frac{Area}{I_{\nu}\%\xi\%T} \times 100\% \dots (3)$$

The specials radioactivity was calculated for each sample with (18000) s time and it calculated from [11]:

$$Specific\ Activity = \frac{^{Area}/_{T-B.G}}{_{I_{\gamma}\% \xi \%\ m}}\ x100\% \ \dots \dots (4)$$

Where:

Area: net area under the photo peak position. m: mass of sample unit (Kg).

6. Calculated the Ratio between $^{235}\mathrm{U}$ / $^{238}\mathrm{U}$

The radioactivity per 1 mg of ²³⁸U is given by [6, 12].

$$Ac = \lambda N \dots (5)$$

 A_c : Radioactivity unit (Bq)

 λ : Decay constant = $\ln 2 / t_{1/2}$

 $t_{1/2}$: half live of ²³⁸U isotope

N: number of atoms and is given by of the equation :.

$$N=No \ (W)/A.....(6)$$

 N_0 : Avocado number.

W: mass of isotope unit (Kg).

A: mass number of isotope radioactivity.

And one can calculated the radioactivity for ²³⁸U series (Ac) from the equation following:-

$$A_c = Ln2/t_{1/2} \times No \times W/A \qquad (7)$$

That Ac from equation (7) is (12,35 Bq).

To calculate the percentage for depleted uranium (X%) in the soil samples, one can use the following equation [9]:

$$X\% = DU(mg/kg)/CsU - 238(mg/kg) \times 100$$
(8)

Where:

DU: Quantity of depleted uranium is found that in a soil sample .

CsU: Concentration 238U in a soil sample of unit (mg / Kg).

And to gut the 235 U / 238 U ratio (Ru) for the soil contain the natural uranium of 238 U with 99.2745 % ratio and 235 U with 0.72 % ratio and 234 U 0.0054 % ratio and depleted uranium contain an increase of 238 U with 0,35 % , and decrease 235 U ratio with the same quantity we can find it by the following equation .

$$Ru = \frac{0.72 - 0.37X}{99.2745 + 0.3755X} \quad \dots \tag{9}$$

Where

X : Ratio of depleted uranium in a soil sample .

7. The Results

In the present work , the determination of the radioactive nuclides resulting from decay of the $^{238}\mathrm{U}$, $^{232}\mathrm{Th}$ series and the single $^{40}\mathrm{K}$ nuclide and $^{137}\mathrm{Cs}$, and the specific radioactivity for

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the (1,2,3,4) samples were done and the results shown in

		Table (1) radioactivity	of the soil sample (1,2,3,4)		
4	3	2	1	Number of Sample	Energy	Nuclide
S4	S3	S2	S1	Region	K(eV)	radioactive
				Ratio between ²²⁶ Ra/ ²³⁴ Th		
0.958	0.897	0.812	0.894		40	²¹² Bi
4.230	11.223	10.112	11.778	Radioactivity of unit (Bq /Kg)	63	²³⁴ Th
19.128	36.286	33.796	15.108		186.2	²²⁶ Ra
21.476	38.204	39.013	19.568		351.9	²¹⁴ Pb
2.234	4.796	3.727	1.512		609.3	²¹⁴ Bi
3.511	2.908	3.794	2.547		1461.5	⁴⁰ K
479.984	518.791	502.859	372.549		40	²¹² Bi

Also, the natural to depleted uranium and $^{235}U/^{238}U$ ratio were calculated and the results shown in table (2).

Table 2: Ratio between depleted uranium and natural and found that the ratio between ²³⁵U/²³⁸U

	Percent ratio of natural	Percent ratio of depleted	Concentration depleted	Concentration	Number of
Ru	uranium (100- X %)	uranium (X %)	uranium (mg/Kg)	²³⁸ U (mg/Kg)	sample
0.005466	52.31601	47.68399	0.919664	1.928664	1
0.007036	94.21707	5.782928	0.061931	1.070931	2
0.006456	78.7534	21.2466	0.272215	1.281215	3
0.005466	52.33359	47.66641	0.919016	1.928016	4

8. The Discussion result

In the present work , the study of the specific radioactivity for a selected soil samples in Babylon province middle of Iraq were done for the position (1,2,3,4) It is found that the ratio between the daughters in ^{238}U series in the start of the series decay of values_between (0,958_0,812) and it represent to a radiation equilibrium approach to the ideal case (1) . But in the end the decay series it is found radiation equilibrium because of the production of ^{226}Ra gas which result from the decay of the ^{226}Ra isotope .

Also is found that the radioactivity of the ($^{212}\mathrm{Bi}$, $^{234}\mathrm{Th}$, $^{226}\mathrm{Ra}$, $^{214}\mathrm{Bi}$, $^{214}\mathrm{Pb}$) nuclides are given of the different ratio depend on concentration isotope of the sample and depended on the geological nature of the soil sample .

The radioactivity of potassium nuclide ^{40}K the highest concentration found in S3(518.791) Bq/Kg for the presence of agriculture land containing phosphate fertilizer , which increased the concentration of isotope ^{40}K .

Also been I identified radioactive region contaminated areas by calculating the pollution of depilated uranium to natural uranium and to find that the ratio between the ^{235}U ,as the ratio between then is the nature (0,0072) , if the ratio between (0.00562 - 0.0072), this means that concentration ranging from depleted uranium (0% - 40 %) [12 , 6]. This is what has been observed in model of the soil, were he found that the sites (1, 3, 4) contain the quantities and different properties of depleted uranium .

9. Conclusions

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1) Fond that the ratio ²³⁵U/²³⁸U in our measurement of soil sample that the site (1.3,4) rang between (0.006456– 0.005466) which indicate the presence varying ratio depleted uranium contamination, which

- is located within limits allowed by the international atomic energy agency, (0,0046) [14, 15] .
- 2) Found that the ratio of ²³⁴Th/²²⁶Ra be in the rang (0.958–0.812), which indicates the radioactive equilibrium which is approaching the ideal case (1), but did not notice this equilibrium after the radium due to the consumption of radon ²²²Rn from the container used the measurement of radioactivity specific model.

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