Measurement of Natural Occurring Radio nuclides

(NORM_s) in Soil using Gamma- ray Spectrometry

R. Obid Hussain, E. Kadum. Abbas

Physics Dept. College of science, kufa-university

Abstract :

The aim of this work is focused on preparation and using gamma-ray spectrometry to investigate and demonstrate a method to determine and evaluate natural radio nuclides which are uranium, thorium and potassium and their activity concentrations, radium equivalent activity, absorbed dose rates in air at one meter above the ground surface and the annual external and internal effective dose rates. The spectrometer consists of NaI (TI) detector coupled with preamplifier and ADC of 4096 Chanel memory, CASSY type. The suitable gain was determined. The efficiency, energy calibration and resolution had been measured by using some standard source .. In order to evaluate the radiological impact for the population, four soil samples have been Collected from different localities in middle area of Najaf province, and measured in which ²³⁸U 232 Th and 40 K were noted. The value of activity concentration ranged from 4.84±0.52 to 17.52±0.94, 5.92 ± 0.31 to 7.07 ± 0.344 and 103.39 ± 2.52 to 185.48 ± 2.09 (Bq kg⁻¹) for ²³⁸U, ²³²Th and 40 K respectively. The calculated values of radium equivalent activity (Raea) ranged from 25.25 to 33.94 $(Bq kg^{-1})$ whereas absorbed doses varied from 12.35 to 16.09 $(Bq kg^{-1})$. Annual effective Doses were found to be in the range (0.061) to (0.079) and (0.015) to (0.020) (mSv) for indoor and Outdoor respectively. The comparison between our results and standard levels recommended by the IAEA and UNSCEAR show low radiation level.

الخلاصة:

ان هدف هذا العمل يتركز على استخدام مطياف اشعة كاما لغرض در اسة وتقييم النويدات المشعة وهي اليور انيوم والثوريوم والبوتاسيوم ووليداتها في التربة وقياس مستوى خطورة الأشعاع عن طريق قياس تركيز نشاطها الأشعاعي النوعي وجرعة اليور انيوم المكافئة وكذلك معدلات الجرعة الممتصة على ارتفاع واحد متر من سطح الارض اضافة الى الجرعة السنوية المؤثرة الداخلية والخارجية يتكون المطياف من كاشف ايوديد الصوديوم المرتبط مباشرة بمضخم ابتدائي والذي يتصل بدوره بوحدة المحول الخطي الى رقمي الذي يحوي على وكذلك معدلات الجرعة الموتسة. على ارتفاع واحد متر من سطح الارض اضافة الى الجرعة المحول الخطي الى رقمي الذي يحوي على المطياف من كاشف ايوديد الصوديوم المرتبط مباشرة بمضخم ابتدائي والذي يتصل بدوره بوحدة المحول الخطي الى رقمي الذي يحوي على 4006 قناة من نوع كاسي . تم قياس كفاءة الكاشف وقدرته التحليلية للطاقة بأستخدام عدد من المصادر المشعة القياسية . ولغرض تقدير خطورة الإشعاع جمعت اربع نماذج من مواقع مختلفة من وسط محافظة النجف وسجلت الاطياف الكامية لها ولوحظت قمم الطاقة العائدة الى كل من الشعاع جمعت اربع نماذج من مواقع مختلفة من وسط محافظة النجف وسجلت الاطياف الكامية لها ولوحظت قمم الطاقة العائدة الى لكل من اليور انيوم والثوريوم و22 وكذلك البوتاسيوم 40 . كما اشارت النتائج بأن قيم النشاط الاشعاعي النوعي اليور انيم والثوريوم والبوتاسيوم 10 . كما الشارت النتائج بأن قيم النشاط الاشعاعي النوعي اليور انيم والثوريوم والبوتاسيوم اليور انيوم والبوتاسيوم 10. كما تراوحت من 25.2±20 لله الى 25.9±20 المالي 2000 المرى تراوحت من 25.2±20 الي 2004 المرى تراوحت من 25.2±20 للك كل كل كيلي غرام لكل من اليور انيوم واللوتاسيوم على التوالي . وكانت القيم المحسوبة لجر عة الراديوم المحاف عرام في حين كما كل كيلي غرام لكل من اليور انيوم والبوتاسيوم على التوالي . وكانت القيم المتصعة على ارتفاع واحد متر من سطح الارص تراوحت من 25.25 الي 20.4 لكل من اليور انيوم والبوتسيوم على التوالي . وكانت القيم المحسوبة لجر عة الراديوم مالمدى من 25.25 للمتصعة على ارتفاع واحد متر من سطح الارض تراوحت من 25.25 لي 20.5 لي عام كل كل كيلي غرام لكل من اليوريوم والبوتاسيوم على التوالي . وكانت القيم المحسوبة لعر ام في حين قيم الحرعة الممتصة على ارتفاع واحد متر من سطح الارض تراوحت من 25.25 لي 20.5 المكان

1-Introduction

The knowledge of radionuclide distribution levels in the environment is important for assessing the effect of radiation exposure due to both terrestrial and cosmogenic sources .Terrestrial radiation is due to radioactive nuclei present in varying amounts in soil, building materials ,water, rocks [1]. The transference of some radio nuclides from these sources to man was happened through food chain or inhalations. The internal exposure is due to irradiation and external exposures occur because of inhalation and ingestion. The particular importance of evaluation of gamma radiation dose from natural sources comes from largest contributor of natural radiation to the external dose of word population [2]. The wide ²³⁸U, ²³²Th, ⁴⁰K and their daughter product dangers effect to human health is the main motivation behind several studies of measuring radiation in environment including the soil and building materials [3,4,5]. For this reason and from the natural risk point of view, it is necessary to obtain a system by which we can observe and measure the natural radiation levels [5,6,7]. Gamma ray spectrometry is a good technique used to measured and analysis of natural and artificial radionuclide [8,9]. The result of this study will b used to establish a baseline map for Najaf area which will be use a reference in formation to record any change in the radioactivity background level due to the change in the topography of the location , other developments and settlement around it and any artificial influences on the environment. The aim of this work is to estimate and demonstrate a method based on the use of gamma- ray spectroscopy technique to determine the natural occurring radionuclide (NOR) in soil in middle of Najaf city .

2-Theoretical part

2-1 Specific activity

The calculation of specific activity was based on equation (1):

Where : A : specific activity,

- C : area under the photo peak ,
- **ε** : counting efficiency ,

 I_{γ} : probability of gamma transition , t : time of measurement , m : mass of the sample .

2-2 Ra_{eq} (Radium equivalent activity) :

T o assess the radiological hazard of soil , it is useful to calculate an index called the radium equivalent activity (Ra_{eq}) [10] defined according to estimation that 1Bq kg⁻¹ of 226 Ra , 0.7Bq kg⁻¹ of 232 Th and13 Bq kg⁻¹ of 40 K produce that the same γ -ray dose [10,11] . Ra_{eq} is given as .

$$Ra_{eq} = A_U + B * A_{Th} + C * A_K \dots (2)$$

Where : $A_{U,}\;A_{Th}$ and $A_{K\;:}$ the specific activity of ^{238}U , ^{232}Th and ^{40}K respectively . B , C : are conversion factors .

2-3 Absorbed doses

The gamma dose rate in the outdoor air at one meter above the ground is calculated by using equation (3)[13].

$$D(nGy/h) = C_1 A_U + C_2 A_{Th} + C_3 A_K \dots (3)$$

Where : C1, C2 and C3 : are the absorbed dose rate conversion factors (DRCF) ; AU ,ATh and AK are the activity concentrations of ^{248}U , ^{232}Th and ^{40}K in (Bq kg^-1) respectively .

2-4 Effective dose

To convert the absorbed dose rate to human effective dose equivalent with an indoor occupancy of 80 % and 20% for outdoor , a conversion factors (A) and (B) in $(SvGy^{-1})$ were used for indoor and outdoor respectively [1,13].

The annual effective doses were determined as following [14].

Indoor (nSv) = (Absorbed Dose) nGyh⁻¹ × 8760h × (A) × 0.7 SvGy⁻¹(4)

Outdoor (nSv) = (Absorbed Dose) nGyh⁻¹ × 8760h × (B) × 0.7 SvGy⁻¹(5)

Where : A and B are conversion factors .

3- Experimental Part :

3-1 Sample preparation

Four surface soil samples were collected from different locations in the middle area of Najaf province . from each location, one sample was collected by digging a hole at a depth of 10 cm before the ground surface . the samples were, crushed, sieved through 0.2 mm mesh and dried in an oven at a temperature of 100 C^0 for 24 h . One kg of each sample was packed in polyethylene marginally beaker of 1.4 L volume . The samples were counted for a period of 10000 second.

3-2 Shielding

The block diagram of gamma ray spectrometry was shown in fig (1). The detector is shielded by a chamber of 2 layer starting from inside with copper (2 mm thick), followed by lead 10 cm thick from out side .



Figure (1) : Block diagram of spectrometer



Figure (2): Different processes in the NaI(T ℓ) crystal

3-3 The gain

To obtain a suitable value of gain , we recorded several spectra of 60 Co standard source for different values of gain as shown in figures (3,4,5).



Figure(3) : Measured spectrum of 60 Co , gain = (-8) .



Figure(4) : measured spectrum of $^{\circ\circ}$ Co , gain = (+3) .







Figure. (6) Spectrum of ¹³⁷Cs

From the three figures (3,4,5), the best value of the gain was of -0.98. For energy calibration we requiring a spectrum from radioactive standard sources of known energies like (¹³⁷Cs, $E_{\gamma} = 662 \text{ KeV}$) and (⁶⁰Co , $E_{\gamma 1} = 1173 \text{ KeV}$ and, $E_{\gamma 2} = 1332 \text{ KeV}$) as shown in figures (5,6).

3-4 Counting efficiency

To calculate the counting efficiency of the detector we use equation (6) [1,4].

$$\varepsilon = A_{re} / A_t \cdot t \times 100 \% \qquad \dots (6)$$

Where :

 A_{re} : recorded area under the photo peak , A_t : Activity of standard source ,

We used equation (7) to correct equation (1)[1].

$$\dots \dots (7) A_{t} = A_0 \ e^{-\lambda t_d}$$

Where:

A_o: The primary activity at manufactured date.

t_d: The period of time between the date of manufacturing and the date of measuring in second.

3-5 Energy resolution:

The resolution of spectrometer is defined as the ability to separate two peaks that closed together in energy

The resolution of photo peach is found by solving the followin

$$\mathbf{R} = \frac{\Delta \mathbf{ch}}{\mathbf{P} \cdot \mathbf{P}} \times \mathbf{100\%} \dots \dots (\mathbf{8})$$

Where :**R**: The resolution ; Δch = The Full width of photo peak at half of the maximum count level (FWHM) measured in number of channels; **P.P** : The channel number at the centre of the photo peak.

4- Result and Discussion:

The counting efficiency of the system was calculated by using 137 Cs, 60 Co and 22 Na ,standard sources. The efficiency curve was plotted in figure (7)



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E(KeV) Figure (7) : Counting efficiency of the detector

respectively.



Figure (8) Spectrum of soil sample s_1



Figure (9) Spectrum of soil sample s_2



Figure (10) Spectrum of soil sample s_3

Table (1) represents the activity concentrations of soil sample obtained by direct gamma-ray From these spectra , three gamma-ray lines spectrometry \cdot the activity concentrations of ${}^{40}K$ of 214 Bi (1760 KeV), 208 T ℓ (2615 KeV) and was found to be in the range 103.39±2.52 to 40 K (1460 KeV) were used to determine the 185.48±2.09 (Bq kg⁻¹) while for 238 U it was in the activity concentration of 238 U, 232 Th and 40 K range 4.84±0.52 Bq kg⁻¹ to 17.52±0.94 Bq kg⁻¹. On the other hand

Table (1): Activity concentrations of the sample

Sample	Activity concentration Bq kg ⁻¹		
Code	²³⁸ U	²³² Th	40K
\mathbf{S}_1	9.18±0.67	6.38±0.32	163.2±2.86
S_2	17.52±0.94	5.92±0.31	103.39±2.32
S_3	4.84±0.52	6.21±0.33	149.69±2.9
S_4	9.85±0.70	7.07±0.34	185.48±2.09

the activity concentrations of ²³²Th was found to be in the range5.92±0.31 to kg^{-1} . The results 7.07±0.344 Bq are comparable to the world average activity concentration . which are 40.35 and 420 Ba kg⁻¹. For 238 U , 232 Th and 40 K . respectivity [13]. Table 1 indicates also that the activity concentrations of 238 U , 232 Th and 40 K are under the worldwide average.



Figure (11) Spectrum of soil sample s_4

The air-absorbed dose rate due to the gamma-ray emission from the samples ranged from 12.35 to 16.09 $nGyh^{-1}$ which is also in the permissible limit [13]. the obtained results showed that the effective doses were lower than world average which is 1.0 Svy⁻¹.

Ra _{eq} (Bq kg ⁻¹)	AD (nGyh ⁻¹)	Indoor (AED [*])(mSv)	Outdoor (AED [*])(mSv)
30.87	15.02	0.074	0.018
33.94	16.09	0.079	0.020
25.25	12.35	0.061	0.015
26.54	12.52	0.061	0.015

5- Conclusions

gamma ray spectrometry has some advantages which are , it is a good technique extensively used in determination of radio nuclides , the wide range of natural and artificial radio nuclides that can be determined and its applicability in a variety of sample matrices . the value of activity concertinos , radium equivalent activity and absorbed dose rate of 238 U, 232 Th and 40 K were found to be lower than the world average . the activity concentration of 40 K was higher than that of 238 U and 232 Th . in general the obtained results showed that indoor and outdoor effective dose due to natural radioactivity of soil samples were less than permissible limit which is 1.0 Svy ⁻¹

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