

RESEARCH TITLE

Natural radioactivity measurements and excessive lifetime cancer risk for some surface Soil samples in (Al-Jabal Al-Gharbi) Mountain, (Awiniya) Region, Libya

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Abstract

Fifteen samples of soil were chosen to achieve this work, the samples were obtained from– Al jabel Algharbi - Awiniya region, Libya. Distributions of natural radionuclides of ²³⁸U, ²³²Th and ⁴⁰K in these samples were determined using a high-purity germanium detector (HPGe) with a specially designed shield. The content of uranium, thorium and potassium is lower than permissible levels in all soil samples. To evaluate the radiological hazard from soil in this area the radium equivalent activity (Raeq), absorbed gamma dose rate (D), annual effective dose equivalent (AEDE), external hazard index (Hex) and excess lifetime cancer risk (ELCR) were calculated in this work. The levels of natural radioactivity and radiation hazard parameters were compared with the international recommended values. All obtained results were found to be below the international recommended limit.

Key Words: Natural radionuclides, Soil samples, Health effects, Gamma spectroscopy, Aljabel - Al.Gharbi (Awiniya region) , Libya

قياسات النشاط الإشعاعي الطبيعي وزيادة خطر الإصابة بالسرطان مدى الحياة لبعض عينات التربة السطحية في (الجبل الغربي) ومنطقة (العوينية) ليبيا

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المستخلص

تم إختيار خمسة عشر عينة من التربة لإنجاز هذا العمل ، وتم الحصول على هذه العينات من الجبل الغربي - منطقة العوينية - ليبيا . حيث تم قياس تركيز النشاط الإشعاعي للنويدات المشعة الطبيعية لكل من ^{238}U ، ^{232}Th و ^{40}K في هذه العينات بإستخدام كاشف الجرمانيوم عالي النقاوة ، أظهرت نتائج هذه الدراسة أن تركيزات اليورانيوم والثوريوم والبوتاسيوم أقل من المستوى المسموح به في جميع عينات التربة في هذه المنطقة. ولتقييم الأثار الإشعاعية التي تسببها النويدات المشعة الأرضية على السكان في هذه المنطقة تمت مقارنة مستويات النشاط الإشعاعي الطبيعي ، ومؤشرات المخاطر الأشعاعية بالقيم الموصى بها دولياً مثل : نشاط مكافئ الراديوم ، ومعدل جرعة جاما الممتصة وما يقابلها من الجرعة السنوية الفعالة ، و زيادة خطر الإصابة بالسرطان مدى الحياة ، و مؤشر الخطر الخارجي وقد تبين أن جميع النتائج التي تم الحصول عليها كانت أقل من الحد الموصى به دولياً.

1. INTRODUCTION

The natural radioactivity in the environment is the main source of radiation exposure for the human body. Natural radionuclide in soil contributes a significant amount of background radiation exposure to the population through inhalation and ingestion. It can be also transferred to plants, foods and drinking water. (Algattawi.A et al., 2019). Natural radiation arises from two main sources, cosmic rays and terrestrial radionuclides. The terrestrial radionuclides are the long-lived primordial radionuclides and their daughter nuclides, which have been present on Earth's crust since the creation of the Earth (M. Degerlier et al., 2008). The soil is the most important source of the terrestrial, The presence of terrestrial radionuclides in soil originates from the disintegrating of rocks through the processes of erosion and deposition, and they are carried to the soil by rain and flows and become an integral component of the soil. The terrestrial component of the natural radiations comes mainly from long-lived primordial radionuclides and their daughter nuclides, which have been present on Earth's crust since the creation of the Earth. The most important primordial radionuclides present in the soil are the radionuclides with half-lives comparable to the age of Earth from the uranium series, thorium series, and ^{40}K , Table (1), (Alajeeli .A , 2019).

Table (1): Radionuclide Half-life (10^9 years).

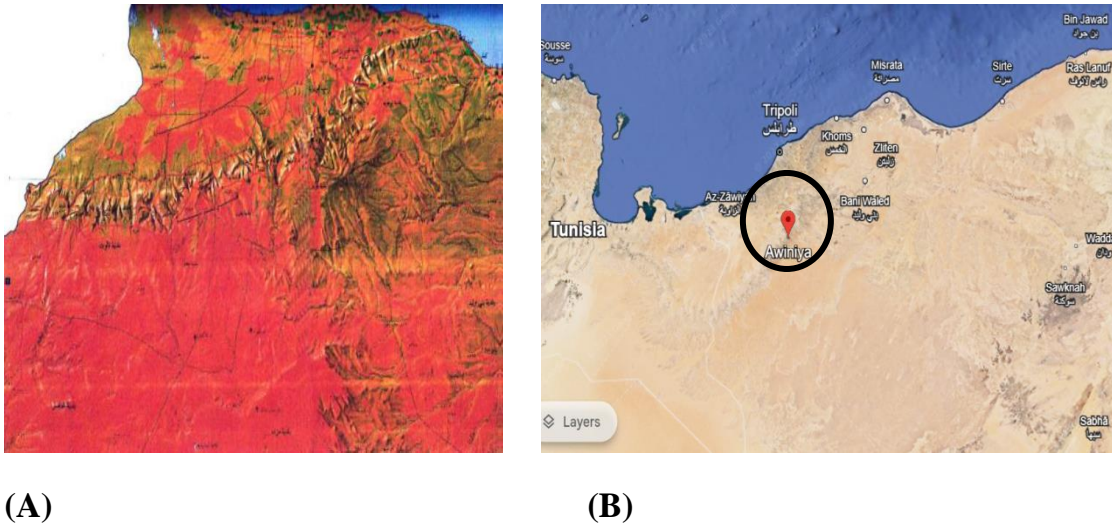
Radionuclide	Half-life (10^9 years)
^{238}U	4.47
^{235}U	0.70
^{232}Th	14.1
^{40}K	1.28

Measurements of natural radioactivity in soil have been performed in many parts of the world, mostly for assessment of the dose and risk resulting from them (Coplestone et al., 2001; Dowdall et al., 2004; El-Bahi et al., 2005). it is necessary to measure and monitor the activity levels of radionuclides in the soil to establish a baseline map and database on environmental radioactivity levels.

2. Materials and methods:

2.1. Site Description of the study area:

Jabal Al-Gharbi Mountain starts from Al-Azeeziah North-East to Wazen West at the Tunisian border, and from Jfarah Valley at the bottom to Nalut at the top. Its height is about 800 m above the sea level (Fig.1 (A)), (Askouri. N et al., 2011).



(A)

(B)

Fig.1: (A) Map of Al-Jabal Al-Gharbi / Libya , (Askouri. N et al., 2011).

(B) Location map of the studied are , Awiniya region .

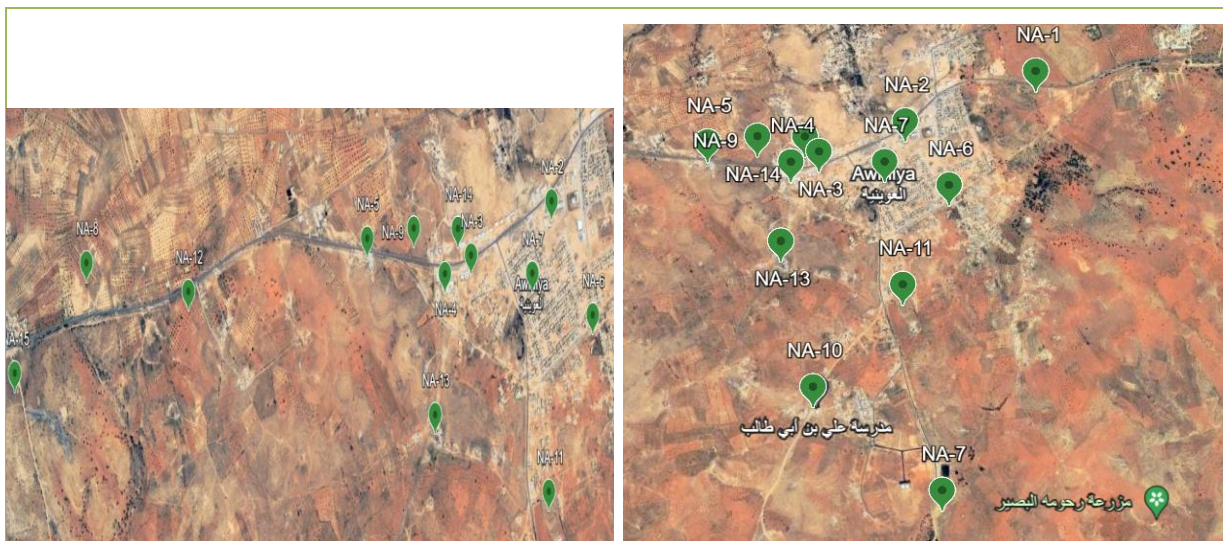


Fig.2: Location map of the studied area samples, Awiniya region .

2.2. Radioactivity measurements :

Gamma-ray spectrometry was done using a high-purity germanium p-type coaxial detector with 30% relative efficiency. Which is connected to a multi-channel (4096) analyzer. The software used is called APTEC 2000 Energy resolution FWHM of the detector is 1.98 Kev at 1332 kev of Co60 Operating voltage is 3000 Manufacturing company of the detector is Eurysis. 226Ra activity was determined indirectly from the gamma rays emitted by its daughter products (214Pb and 214Bi) where; the concentrations of 214Pb and 214Bi were measured from (295.1(18.7%), 351.9 (35.8%)) KeV and (609.3 (45%), 1120.3(14.9%) KeV photopeaks, respectively. The gamma-ray transitions of 228Ac (911 (29%)) KeV, 212Bi (727.3 KeV (7%)) and 208Tl (583.1 KeV (30%)) were used to evaluate the specific activity of 232Th. 40K was determined directly from the 1460 KeV (10.7%) peak energy (**Technical Reports Series, 1989**).

2.3. Calculation of activity concentration :

The radioactivity concentration in the environmental samples was obtained using the equation (1) (Usif. M and Taher. A. , 2008):

$$A = \frac{\text{CPS}}{\text{eff} \times I \times W} \quad (1)$$

Where, A = Activity of the sample in Bq·kg⁻¹; cps = the net counts per second; eff = the counting efficiency of the gamma energy; I = absolute intensity of the gamma-ray; and m is the mass of the sample in kilograms.

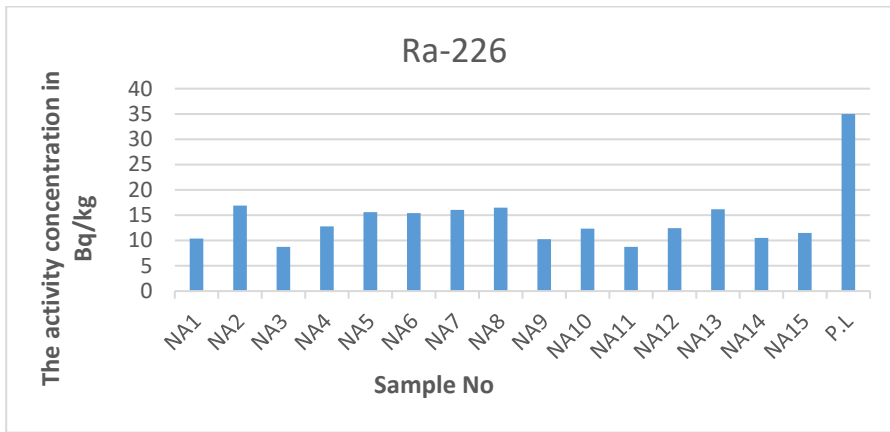
3. Result and Discussion :

3.1. Activity concentrations of ²²⁶Ra , ²³²Th and ⁴⁰K :

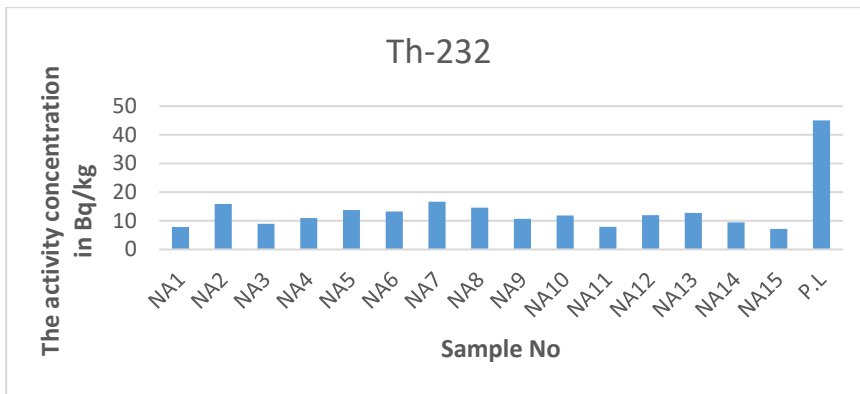
The values of activity concentration of ²²⁶Ra, ²³²Th and ⁴⁰K of all studied soil samples were calculated and illustrated in (Table 2). The activity concentrations for radium ²²⁶Ra range (from 8.72 Bq/kg in sample 3 and sample 11 to 16.88 Bq/kg in sample 2) (Fig.3), ²³²Th from (7.14 Bq/kg in sample 15) to (16.66 Bq/kg in sample 7) (Fig.4) and ⁴⁰K (from 58.05 Bq/kg in sample 3 to 180.18 in sample 8) (Fig.5). The radioelements worldwide average (W.A.) values are 35 Bq/kg for ²²⁶Ra, 45 Bq/kg for ²³²Th, 420 Bq/kg for ⁴⁰K (UNSCEAR, 2010), the activity concentrations of all studied samples for thorium, radium and potassium are lower than the permissible level.

Table (2) : The activity concentrations in (Bq/Kg) of ²²⁶Ra , ²³²Th and ⁴⁰K

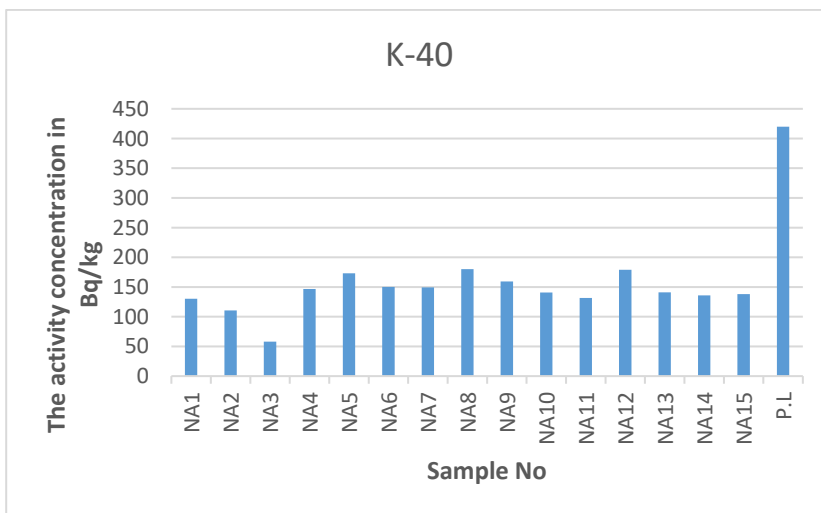
Sample Number	²²⁶ Ra	²³² Th	⁴⁰ K
NA1	10.37	7.84	130.19
NA2	16.88	15.89	110.64
NA3	8.72	8.96	58.05
NA4	12.79	10.95	146.75
NA5	15.61	13.75	173.13
NA6	15.41	13.26	150.19
NA7	16.03	16.66	149.41
NA8	16.47	14.57	180.18
NA9	10.26	10.69	159.40
NA10	12.34	11.87	140.69
NA11	8.72	7.91	131.53
NA12	12.44	11.96	178.92
NA13	16.18	12.77	141.12
NA14	10.50	9.47	136.00
NA15	11.47	7.14	137.97
P.L	35	45	420



(Fig.3) : The Activity Concentration of ²²⁶Ra



(Fig.4) : The Activity Concentration of ²³²Th



(Fig.5) : The Activity Concentration of ⁴⁰K

3.2. Radiological hazard indices :

Radium Equivalent (Raeq) :

The radium equivalent (R_{aeq}) index in Bq/kg is a widely used radiological hazard index. It is a convenient index to compare the specific activities of samples containing different concentrations of ²²⁶Ra, ²³²Th and ⁴⁰K. It was defined on the assumption that 10 Bq/kg of ²²⁶Ra, 7 Bq/kg of ²³²Th and 130 Bq/kg of ⁴⁰K produce the same gamma dose rate. It was calculated by equation (2) as follows:

$$R_{aeq} = C_{Ra} + 1.43 C_{Th} + 0.077C_k \quad (2)$$

Where C_{Ra} , C_{Th} and C_k are the activity concentrations of ^{226}Ra , ^{232}Th and ^{40}K in Bq/kg, respectively. R_{aeq} was estimated for the collected samples and is given in Table (3), Fig (6). The values of R_{aeq} for all soil samples varied (from 26.01 to 51.36) Bq/kg, and were estimated lower than the recommended maximum value of 370 Bq kg⁻¹ (Ababneh et al., 2010).

External hazard index (H_{ex}) :

The external hazard index (H_{ex}) represents the external radiation exposure associated with gamma irradiation from radionuclides of concern. The value of H_{ex} should not exceed the maximum acceptable value of one to keep the hazard insignificant. The external hazard (H_{ex}) is defined by the equation (3): (Jankovic et al., 2008).

$$H_{ex} = (C_{Ra}/370 + C_{Th}/259 + C_k/4810) \leq 1 \quad (3)$$

Where C_{Ra} , C_{Th} and C_k are the concentrations in (Bq/ Kg) of radium, thorium and potassium respectively. The values of H_{ex} for all soil samples varied from (0.07 - 0.13), and were found to be less than 1 as given in Table (3), Fig (7).

Radiation Level Index (I_γ) :

This index can be used to estimate the level of γ -radiation hazard associated with the natural radionuclides in the samples; it is given by the equation (4) :

$$I_\gamma = \frac{C_{Ra}}{150} + \frac{C_{Th}}{100} + \frac{C_K}{1500} \quad (4)$$

Where C_{Ra} , C_{Th} and C_k are the activity concentration of ^{226}Ra , ^{232}Th , ^{40}K in Bq/kg, respectively. The value of these indexes must be less than unity in order to keep the radiation hazard insignificant, the values of Representative Level Index (I_γ) from (0.18 - 0.37) for all samples are found to be less than 1 given in Table (3), Fig (7).

Absorbed dose rate (D) :

The absorbed dose rate D in outdoor air at 1m above the ground level was assessed from the natural activities of ^{226}Ra , ^{232}Th and ^{40}K (Bq.kg⁻¹) dry Weight was calculated using equation (5) (Ajayi and Kuforiji, 2001; El-Arabi et al., 2006) :

$$D_{out} = 0.436 C_{Ra} + 0.599 C_{Th} + 0.0417 C_K \text{ (nGyh}^{-1}\text{)} \quad (5)$$

The values of D_{out} for all soil samples varied from (11.59 - 23.42) nGyh⁻¹, and were found to be less than 59 nGyh⁻¹ as given in Table (3), Fig (8). This factor is very important to evaluate the annual effective dose in an outdoor environment, AED can be given by equation (6):

$$AED = D_{out} \text{ (nGyh}^{-1}\text{)} \times 20\% \text{ of } 8760 \text{ h} \times 0.7 \text{ (Sv Gy}^{-1}\text{)}$$

$$AED = D_{out} \times 1.226 \mu\text{Sv} \quad (6)$$

The values of AED for all soil samples varied from (0.014 – 0.029) mSvy⁻¹, and were found to be less than 0.07 mSvy⁻¹ as given in Table (3), Fig (9).

Excess lifetime cancer risk (ELCR) :

For outdoor exposure, ELCR can be calculated from outdoor annual effective dose AED by using the following equation (7) :

$$ELCR = AED \times 66 \times 0.05 \quad (7)$$

Where 66 is the average life expectancy (approximately 66 years) and 0.05 is the fatal risk factor per Sievert, (**Qureshi et al, 2014**). The values of *AED* for all soil samples varied from $(0.047 - 0.095) \times 10^{-3}$, and were found to be less than (0.29×10^{-3}) as given in Table (3), Fig (10).

Table (3) : The values of Radium equivalent activity, Radioactivity level index, External hazard index, absorbed doses rate D, the annual effective doses AED and excess lifetime cancer risk ELCR for the area under investigation.

Sample Numbe	Ra _{eq} Bq/Kg	I gamma	H _{ex}	D(out) nGyh ⁻¹	E(out) mSvy ⁻¹	ELCR(out) × 10 ⁻³
NA1	31.61	0.23	0.08	14.65	0.018	0.059
NA2	48.13	0.34	0.13	21.49	0.026	0.087
NA3	26.01	0.18	0.07	11.59	0.014	0.047
NA4	39.75	0.29	0.10	18.25	0.022	0.074
NA5	48.60	0.35	0.13	22.26	0.027	0.090
NA6	45.94	0.33	0.12	20.92	0.026	0.085
NA7	51.36	0.37	0.13	23.20	0.028	0.094
NA8	51.19	0.37	0.13	23.42	0.029	0.095
NA9	37.82	0.28	0.10	17.52	0.022	0.071
NA10	40.15	0.29	0.10	18.35	0.023	0.074
NA11	30.16	0.22	0.08	14.02	0.017	0.057
NA12	43.33	0.32	0.11	20.05	0.025	0.081
NA13	45.31	0.32	0.12	20.59	0.025	0.083
NA14	34.53	0.25	0.09	15.92	0.020	0.064
NA15	32.32	0.23	0.08	15.03	0.018	0.061
P.L	370	1	1	59	0.07	0.29

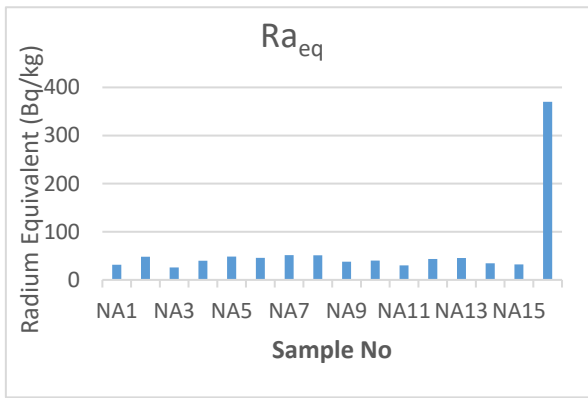


Fig (6) : Radium equivalent (Ra_{eq}) in Bq/kg.

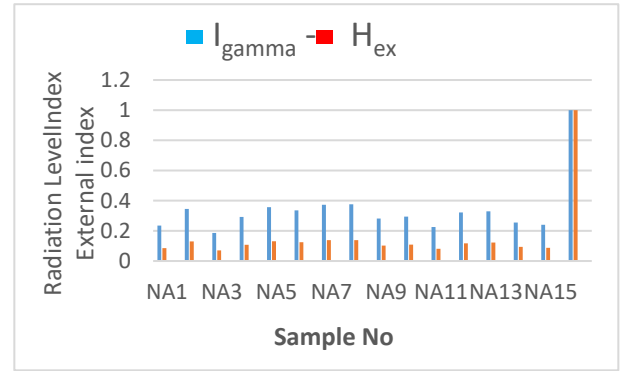


Fig (7) : External hazard index (H_{ex}) and Radiation Level Index (I_γ).

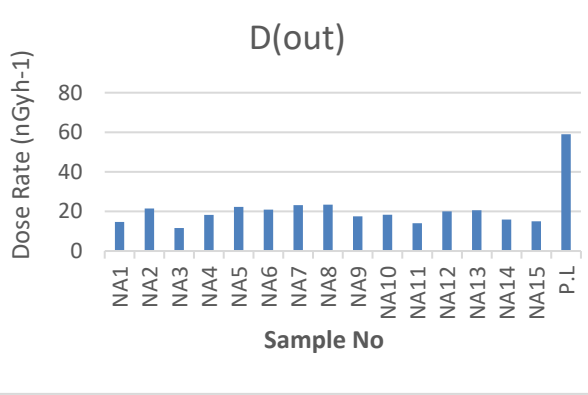


Fig (8) : absorbed dose rate D in outdoor in (nGyh⁻¹).

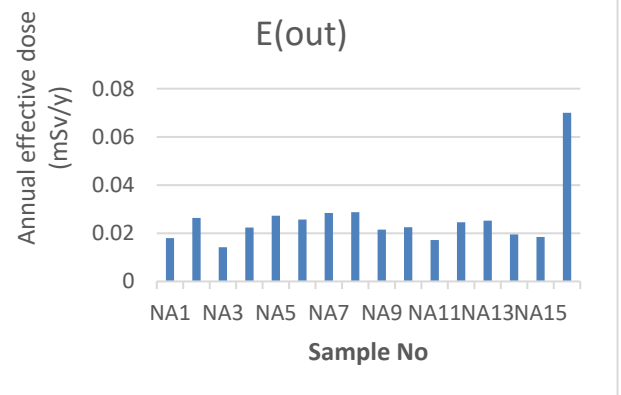


Fig (9) : the annual effective dose in an outdoor mSvy⁻¹.

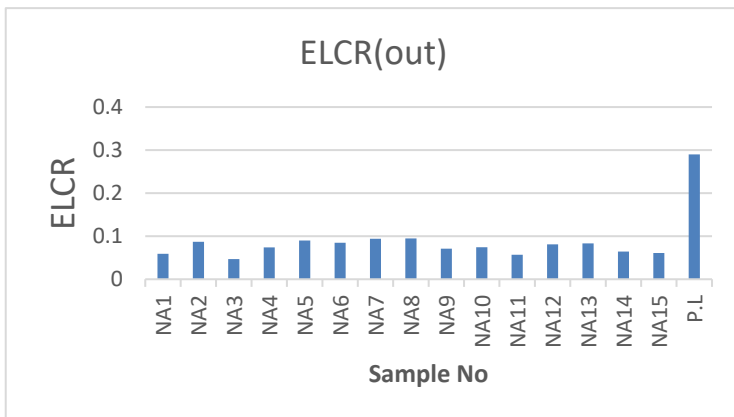


Fig (10) :The excess lifetime cancer risk ELCR.

5 . Conclusions :

In the work presented here, The obtained results illustrate the following important observation: The activity concentrations of ^{226}Ra , ^{232}Th and ^{40}K were measured by a high-purity germanium (HPGe) detector for the 15 soil samples. The measured activity concentrations of ^{226}Ra , ^{232}Th and ^{40}K in all soil samples are lower than the permissible limit reported in **UNSCEAR (2010)**. The radium equivalent, external hazard index, and dose rate have low values. The ELCR factor assessed during this work based on outdoor effective dose (Eout) was found to be lower than the permissible level. This indicates that the soils considered were safe and can be used in the building of dwellings and sources of construction material without giving any radiological hazard to the local population. The results of measurements will serve as baseline data and background reference level for the Awiniya region, Libya, for drawing a radiological map of this region.

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