NATURAL ENVIRONMENTAL RADIOACTIVITY IN THE SOIL OF TERENGGANU STATE, MALAYSIA

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Abstract. The activity concentrations of naturally occurring radionuclides in environmental medium such as soil play an important role in setting national and international average doses to public for radiation protection purposes. This work measures the activity concentrations of the naturally occurring radionuclides (²²⁶ Ra, ²³²Th and ⁴⁰K) in soil samples of Terengganu state, Malaysia using hyper pure germanium (HPGe) spectrometry. The mean (range) activity concentrations of ²²⁶Ra, ²³²Th, and 40K in the soil samples 79 ± 3 (20 ± 1 - 151 ± 5) Bq kg⁻¹ with the mean value of Bq kg⁻¹ for ²²⁶Ra; 84 ± 3 (8 ± 1 - 182 ± 6) Bq kg⁻¹ for ²³²Th, and 545 ± 55 (47 ± 5 - 1056 ± 107) Bq kg-1 for 40K. Upon comparing these values with the world averages for ²²⁶Ra, ²³²Th and ⁴⁰K, which are 33, 36 and 474 Bq kg⁻¹, respectively. It is therefore, evidently seen that the mean activity concentrations of ²²⁶Ra and ²³²Th in the soil of Terengganu are slightly more than twice their corresponding global averages. The mean activity concentration of ⁴⁰K in the soil of Terengganu is ~ 15% higher than the world average.

Keywords: Environmental radioactivity; activity concentrations; 232Th; 226Ra; 40K; Terengganu

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1.0 INTRODUCTION

Natural environmental radioactivity arises mainly from primordial radionuclides such as ⁴⁰K and also from ²³⁸U and ²³²Th decay series, which occur at trace levels in all ground formation [1]. Several studies performed worldwide have measured the activity concentration of natural radionuclides in soil [2-10]. In Malaysia, studies have been conducted to measure the terrestrial natural radioactivity as well as the human health consequences [11-22].

This study is one of the few pioneer which consider the State of Terengganu for the measurement of terrestrial gamma radiation dose rates and the top soil activity concentrations of 226 Ra, 232 Th and 40 K.

2.0 EXPERIMENTAL

2.1 METHODS

2.1.1 Study Area

The study area is located in the North-eastern part of Peninsula Malaysia, covering latitude 4 °00 ′00 to 5 °15 ′00 N and longitude 102 °15 ′00 to 103 ° 45 ′00. It is bordered in the Northwest by Kelantan state, the southwest by Pahang, and the east by the South China Sea. Several outlying islands, including Pulau Perhentian, Pulau Kapas and Pulau Redang, are also a part of the state. The state consists of seven districts (**Figure 2.1**) and has a total area of 13,035 km² and a population of 1,015,776 as of 2010 [23]. The study area is underlain by eight geological formations described by Director of Geological Survey, 1985 shown in (**Figure 2.2**).

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Figure 2.1: Districts map of the study area



Figure 2.2 Geological formations map of Terengganu

2.2 Data Collection

2.2.1 Sample collection and preparation

2 kg soil samples were collected from the state of Terengganu, which widely covered each of the eight districts. Twenty four soil samples were taken from the depth of 10 - 15 cm within a marked cleared area of about 50 cm in diameter and mixed thoroughly, in order to obtain a representative sample of that area. They are taken away from roads, trees, buildings and any other source of obstruction. The sampling locations were recorded with a global positioning system (GPS). The collected samples were packed in a well labeled polythene bags and transported to the laboratory.

In the laboratory, the samples were oven dried at 110 ^oC for 24 hours, cleared of stones and pebbles, crushed and ground to a fine powder and passed through a 400 µm and 250 µm mesh sieved to be homogenized. The homogenized samples were then weighted and packed into a standard 500 mL Marinelli beakers and stored for about thirty days to attain secular equilibrium between radium and its progeny, this procedure have been adopted in past studies [24-26].

2.2.2 Sample analysis

A coaxial high purity germanium (HPGe) detector (GC2018-7500 SL) manufactured by Canberra with a relative efficiency of 20 % relative to a 7.62 cm \times 7.62 cm NaI (Tl) detector and a resolution of 1.8 keV for the 1332 keV gamma ray emission of ⁶⁰Co, Genie 2000 (VI.3) software from Canberra was used to analyze the spectra.

The energy calibration was done using a point source of 241 Am (59.54keV), 137 Cs (661.62 keV) and 60 Co (1173 and1332 keV), whereas efficiency was calibrated using a 500 mL multi-nuclide standard solution. A Marinelli beaker full with de-ionized water was counted to strip the background from the samples. The value of Minimum Detectable Activity (MDA) was 13 Bq kg⁻¹ for 40 K, 1 Bq kg⁻¹ for 226 Ra and 2 Bq kg⁻¹ for 232 Th for a counting time of 21,600 s [27, 28].

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The activity of ²²⁶Ra was determined based on gamma ray emissions of ²¹⁴Pb (352 keV) and ²¹⁴Bi (609), ²³²Th was determined based on the emissions of ²⁰⁸Tl (583.1 keV) and ²²⁸Ac (911.2 keV) and that of ⁴⁰K was determined directly from it emission energy of 1461.8 keV. The concentrations of ²²⁶Ra and ²³²Th were calculated from the weighted mean activity values determined for various emissions. IAEA Soil 6, IAEA S-14 and IAEA S-16 were used for quality control and assurance.

3.0 RESULTS AND DISCUSSION

Activity concentrations of naturally occurring radionuclides were measured for 24 soil samples collected using (HPGe) detector. The measured activity concentration of ²²⁶Ra varied from 20 ± 1 to 151 ± 5 Bq kg⁻¹ with a mean value of 79 ± 3 Bq kg⁻¹, ²³²Th activity concentrations ranged from 8 ± 1 to 182 ± 6 Bq kg⁻¹ with a mean value of 84 ± 3 Bq kg⁻¹ and the activity concentrations of ⁴⁰K ranged from 47 ± 5 to 1056 ± 107 Bq kg⁻¹ with a mean value of 545 ± 55 Bq kg⁻¹as shown in Table 1.

District -	K (Bq kg ⁻¹)		Ra (Bq kg ⁻¹)		Th (Bq kg ⁻¹)	
	Mean	Range	Mean	Range	Mean	range
Besut	424 ± 45	83 - 1056	83 ± 4	20 - 115	82 ± 4	8 - 160
Dungun	589 ± 58	266 - 929	74 ± 2	39 - 114	106 ± 4	51 - 182
Hulu Terengganu	431 ± 44	262 - 828	91 ± 3	60 - 148	93 ± 4	49 - 166
Kemaman	588 ± 58	47 - 986	81 ± 3	29 - 151	84 ± 3	34 - 131
Kuala Terengganu	504 ± 50	459 - 549	33 ± 1	22 - 43	51 ± 2	45 - 57
Marang	475 ± 48	475 - 475	55 ± 2	55 - 55	79 ± 3	79 - 79
Setiu	762 ± 75	655 - 832	89 ± 3	51 - 121	79 ± 3	47 - 132
Terangganu State	545 ± 55	47 - 1056	79 ± 3	20 - 151	84 ± 3	8 - 182

 Table 1.0
 Activity concentrations of natural radionuclides in the districts within the study area

The measured activity concentration of of 226 Ra and 232 Th in the soil of Terengganu were slightly more than twice their corresponding global averages. The mean activity concentration of 40 K in the soil of Terengganu was ~ 15% higher than the world average. It can be observed from (**Figure 3.1**) that the highest 226 Ra activity concentrations were found in the Hulu Terengganu with mean activity concentrations of 91 ± 3 Bq kg⁻¹.

Dungun has the highest activity concentrations of 232 Th with a mean value of 106 ± 4 Bq kg⁻¹ whereas the highest concentration of 40 K was found in Setiu district with a mean value of 762 ± 75 Bq kg⁻¹. The areas which are covered by acid intrusive geological formations and overlain by Haplic Acrisol-Haplic Acrisol - Gleyic Luvisol soil type has the highest mean activity concentration of 226 Ra, while the area covered by mostly acid intrusive geological formation and overlain by Dystric Fluvisols - Dystric Gleysol soil type has the highest activity concentration of 232 Th.

The lower concentrations was observed in areas which are covered by carboniferous geological formation and overlain by Haplic Acrisol- Haplic Acrisol - Gleyic Luvisol soil type for both ²²⁶Ra and ²³²Th respectively. ⁴⁰K activity concentration was found to be higher in carboniferous geological formation and overlain by Haplic Acrisol- Haplic Acrisol - Gleyic Luvisol soil types.



Figure 3.1: A plot of district's Ra, Th and K mean activity conc. in Terengganu

4.0 CONCLUSION

The mean activity concentration of 226 Ra, 232 Th and 40 K was found to be 79 Bq kg⁻¹, 84 Bq kg⁻¹ and 545 Bq kg⁻¹ respectively. The mean activity concentration of 226 Ra and 232 Th are about two times higher than the world average whereas the mean activity concentration of 40 K is slightly higher than the world average value of 420 Bq kg⁻¹.

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