Printed in the Republic of Korea ANALYTICAL SCIENCE
& TECHNOLOGY
Vol. 36 No. 4, 191-197, 2023 https://doi.org/10.5806/AST.2023.36.4.191

Radioactivity concentrations of natural radionuclides in fine dust of Jeju, Korea

Chung-Hun Han^{1,2}, Sohyeon Lim¹, and Hee-Jung Im^{1, *}

¹Department of Chemistry, Jeju National University, 102 Jejudaehak-ro, Jeju-si, Jeju Province 63243, Korea ²Institute for Nuclear Science and Technology, Jeju National University, 102 Jejudaehak-ro, Jeju-si, Jeju Province 63243, Korea (Received May 27, 2023; Revised July 5, 2023; Accepted July 17, 2023)

Abstract: Radioactivity concentrations for natural radionuclides were determined from fine dust samples collected in Jeju, Korea according to atmospheric events (Asian dust, haze, fog-mist, and non-event), and radium equivalent activity was calculated. The mean atmospheric radioactivity concentrations for ²³⁸U, ²³²Th, and ⁴⁰K in 127 fine dust samples were 0.49, 0.24, and 7.23 μ Bq m⁻³, respectively, and the radium equivalent activity was 33.25 Bq kg⁻¹. The mean concentrations of ²³⁸U and ²³²Th in the fine dust during the Asian dust period were 1.31 and 1.60 μ Bq m⁻³, respectively, above the global average, while the values for the other three atmospheric events were lower. The ratio of ²³²Th/²³⁸U radioactivity during the Asian dust period was 1.22, higher than the ratio for the other three atmospheric events.

Key words: natural radionuclides, fine dust, atmospheric phenomenon

1. Introduction

Humans are always exposed to natural radiation associated with non-decaying radionuclides such as ⁴⁰K and the daughter radionuclides of the ²³²Th and ²³⁸U decay series. We receive over 80 % of our radiation from these nuclides.^{1,2} Radiation is commonly found in rocks, soils, oceans, houses, and more,³ so, radiation levels vary by region due to different concentrations of nuclides derived from soil, fertilizers, or their equivalents.^{1,4}

Many studies have been performed to measure these radioisotope activities released from rocks and soils, and the dose gamma radiation on human was estimated by calculation.^{1,5-11} Besides the rocks and soils, internal radiation doses from tobacco, ingestion, building materials, fine dust, etc. have also been studied.¹²⁻¹⁶

In 2015, as in 2014, fine dust (PM₁₀ aerosols) was collected from Gosan site on Jeju's Halla mountain in Korea, which is largely unaffected by urban air pollutants and serves as a background for estimating air pollution on the Korean Peninsula.¹⁷ The area is centrally located between China, Taiwan, Japan, and the Korean Peninsula, making it a good place to monitor the characteristics of pollutants traveling

 \star Corresponding author

Phone : +82-(0)64-754-3544 Fax : +82-(0)64-756-3561 E-mail : imhj@jejunu.ac.kr

This is an open access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/ by-nc/3.0) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

long distances from neighboring countries.

In general, natural radionuclides are easily measured by mass spectrometry, because they have too low a gamma-ray emission rate or a very low branching ratio from the parent nuclide to be measured by conventional gamma spectrometry. Therefore, instead of directly measuring gamma radiation, we used inductively coupled plasma - mass spectrometry (ICP-MS) to measure the radioactivity concentrations of ⁴⁰K, ²³²Th and ²³⁸U in the fine dust and compare the results with those from 2014.

2. Experimental

2.1. Sample preparation and analysis

Jeju Island faces the Asian continent, and the Gosan site where the fine dust was collected is located at $33^{\circ}1'N$ and $126^{\circ}10'E$ on the western side of Jeju Island.

Sampling was performed using an automated system, PMS-103 (APM Engineering, Korea), equipped with a Teflon filter (47 mm/2.0 μ m) for 24 hours every three days during whole year of 2015, the same as the previous collection method.¹⁶ It maintained a flow rate of 16.7 L/min with mass flow controller (MFC) to collect fine dust.

Again, as in reference 16, the aerosol filter sample was placed in a perfluoroalkoxy Teflon container in a microwave digestion system, and 10 mL of a 5.55 % HNO₃/16.75 % HCl acid solution was added.¹⁸ This solution was heated in a 180 °C and 1000 W microwave for 15 min to digest the particulate matter, and the digested solution was filtered through a 45 µm PVDF syringe filter. To the solution, 5 mL of 3 % HNO₃/ 8 % HCl acid solution and ultrapure water were added until the volume was 25 mL.¹⁹ Three species, ³⁹K, ²³²Th and ²³⁸U, were measured by ICP-MS (ICP-DRC-MS, Perkin Elmer, ELAN DRC-II, USA) under the conditions of 1500 W RF power, 40 MHz RF frequency, 15.0 L min⁻¹ cooling water, 0.9-1.05 L min⁻¹ Ar flow rate, and assist 1.2 L min⁻¹.

2.2. Concentrations of isotopic activity Based on ICP-MS measurements of potassium, thorium, and uranium in fine dust, the radioactivity concentrations of the isotopes ⁴⁰K, ²³²Th, and ²³⁸U could be calculated as follows:²⁰

$$A_i = \frac{\ln 2}{T_{1/2}} \times \frac{R_i \times m_e}{M_i} \times N \tag{1}$$

Here, A_i : radioactivity concentration (Bq m⁻³)

 $T_{1/2}$: half-life time (s) of isotope *i* (⁴⁰K; 1.28×10⁹ y, ²³²Th; 1.405×10¹⁰ y and ²³⁸U; 4.468×10⁹ y)

 R_i : isotopic ratio (natural abundance) of isotope *i* (⁴⁰K; 0.000117, ²³²Th; 1.0 and ²³⁸U 0.99275, based on the IUPAC report 2009²¹)

 m_e : mass concentration of element e corresponding to isotope *i* (g m⁻³)

 M_i : atomic mass (g mol⁻¹)

N: Avogadro's number $(6.022 \times 10^{23} \text{ mol}^{-1})$

2.3. Radium equivalent activity

The radioactivity concentrations per unit mass for the dust above were used to calculate the radium equivalent activity (Ra_{eq}). The Ra_{eq} is an indicator of radiation hazard from dose of daughter radionuclides contained in a material, which is often expressed as a biohazard index.²² This Ra_{eq} was set under the assumption that the γ dose rate is the same for nuclides ²³⁸U = 370, ²³²Th = 259, and ⁴⁰K = 4810 Bq kg⁻¹ per unit mass. In this study, the concentrations of the nuclides (⁴⁰K, ²³²Th, and ²³⁸U) in PM₁₀ samples collected in 2015 were calculated as dust masses, and the Ra_{eq} was calculated by utilizing them. The results were then checked to see if they were below the limit of 370 Bq kg^{-1.23}

$$Ra_{eq} = \left(\frac{A_U}{370} + \frac{A_{Th}}{259} + \frac{A_K}{4810}\right) \times 370$$
(2)

Here, Ra_{cq} : radium equivalent activity A_U : activity concentration (Bq kg⁻¹) of ²³⁸U A_{Th} : activity concentration (Bq kg⁻¹) of ²³²Th A_k : activity concentration (Bq kg⁻¹) of ⁴⁰K

3. Results and Discussion

3.1. Radioactivity concentration per dust mass Total 127 fine dust samples were collected at

192

Table 1.	Comparison	of ⁴⁰ K,	232 Th, and	²³⁸ U	mass	and	radioactivity	concentrations	per	dust	mass	of	Gosan	fine	dust
----------	------------	---------------------	------------------	------------------	------	-----	---------------	----------------	-----	------	------	----	-------	------	------

Samplas		⁴⁰ K	232	Th	23	Dof		
Samples	mg kg ⁻¹ -dust	Bq kg ⁻¹ -dust	mg kg ⁻¹ -dust	Bq kg ⁻¹ -dust	mg kg ⁻¹ -dust	Bq kg ⁻¹ -dust	Kel.	
2014 (n=115)	0.56±0.38	146.90±101.84	1.03 ± 0.73	4.16±2.98	0.53 ± 0.60	6.48±7.43	16	
2015 (n=127)	0.62±0.41	163.31±107.69	1.15±1.16	4.65±4.71	1.14 ± 0.91	14.05±11.16		

Gosan site during 2015. Among them, 7 samples on Asian dust events, 62 samples on normal weather days (non-event days), 10 samples on haze days, and remaining 48 samples on fog-mist days were collected. All of these samples were analyzed by ICP-MS to determine the radioactivity concentrations of ⁴⁰K, ²³²Th, and ²³⁸U in the fine dust.

Table 1 shows the concentrations of the masses of the isotopes ⁴⁰K, ²³²Th, and ²³⁸U of whole fine dust samples in 2014 and 2015, as well as the radioactivity concentration per mass of dust. The mean mass concentration of whole fine dust samples in 2015 was 41.54 μ Bq m⁻³, which was similar to the 2014 concentration.¹⁶ The mean mass concentrations of ⁴⁰K, ²³²Th, and ²³⁸U were 0.62±0.4, 1.15±1.2, and 1.14±0.9 mg kg⁻¹-dust respectively. In addition, the mean radioactivity concentrations per dust mass of ⁴⁰K, ²³²Th, and ²³⁸U were 163.31±107.7 Bq kg⁻¹-dust (range: 3.16~468.23 Bq kg⁻¹-dust), 4.65±4.7 Bq kg⁻¹-dust (range: 0 ~ 42.69 Bq kg⁻¹-dust), and 14.05 ± 11.2 Bq kg⁻¹-dust (range: 2.58~78.82 Bq kg⁻¹-dust) respectively. The global average radiation concentrations of

 ^{232}Th and ^{238}U are 30 and 35 Bq kg^-1-dust, so the concentrations in the measured samples are much lower.^1

3.2. Activity concentrations per air volume and radium equivalent activity

Table 2 and Fig. 1 show the activity concentrations per air volume and radium equivalent activity by month. The overall high levels in February and March are due to the effects of the 2015 Asian dust, which was concentrated on February 22-23 and March 21-22. However, the high radium equivalent activity in September is unusual, and the reasons for it are not yet fully understood. Nevertheless, all ²³²Th and ²³⁸U activity concentrations and radium equivalent activity values are below than the global average of mean atmospheric activity concentrations and radium equivalent activity which are 0.5, 1.0 μ Bq m⁻³, and 310 Bq kg⁻¹, respectively.¹ This is expected due to the nature of the Gosan region, where relatively low dust levels are present, whereas the global average is obtained based on total suspended particles.

Table 2. The activity concentrations of ⁴⁰K, ²³²Th, and ²³⁸U per air volume and radium equivalent activity per month at Gosan

Year-month (n)	⁴⁰ K (µBq m ⁻³)	²³² Th (µBq m ⁻³)	²³⁸ U (µBq m ⁻³)	$Ra_{eq} (Bq \ kg^{-1})$
2015-Jan (8)	7.19±7.02	0.14±0.12	0.31±0.20	22.95±10.24
2015-Feb (13)	15.59±13.83	0.77 ± 1.10	0.92 ± 0.75	40.68±17.04
2015-Mar (11)	14.52±13.77	0.49±0.51	0.63 ± 0.45	30.54±14.53
2015-Apr (6)	5.73±7.11	$0.20{\pm}0.21$	$0.40{\pm}0.38$	21.78±10.61
2015-May (11)	2.56±1.78	0.08 ± 0.07	0.30 ± 0.27	21.88±15.53
2015-Jun (11)	6.71±7.47	0.16±0.31	0.28±0.37	22.21±14.35
2015-Jul (11)	3.04±1.54	$0.10{\pm}0.04$	0.45 ± 0.14	31.43±16.39
2015-Aug (11)	4.52±4.21	$0.12{\pm}0.07$	0.59 ± 0.43	31.76±23.62
2015-Sep (14)	4.95±2.25	0.23 ± 0.42	0.50 ± 0.22	54.28±25.52
2015-Oct (12)	10.98 ± 5.84	0.25±0.12	0.52±0.21	44.27±15.21
2015-Nov (9)	5.21±4.29	0.09 ± 0.05	0.33±0.10	34.08 ± 8.46
2015-Dec (10)	3.14±1.83	0.09 ± 0.04	0.44 ± 0.28	26.63±13.89
Total mean (127)	7.23±8.29	0.24±0.46	0.49±0.40	33.25±19.17

Vol. 36, No. 4, 2023



Fig. 1. Histogram of *Table* 2 (The activity concentrations of ⁴⁰K, ²³²Th, and ²³⁸U per air volume and radium equivalent activity per month at Gosan).

Table 3. The activity ratio of 40 K, 232 Th, and 238 U activity concentrations (μ Bq m⁻³) by atmospheric phenomenon at Gosan site in Jeju

Atmospheric	⁴⁰ K	²³² Th	²³⁸ U	²³² Th/ ²³⁸ U	
Phenomenon (<i>n</i>)	(µBq m ⁻³)	(µBq m ⁻³)	(µBq m ⁻³)		
Asian Dust (7)	27.98±14.6	1.60±1.2	1.31 ± 0.7	1.22	
Haze (10)	18.39±8.4	0.41±0.3	0.77±0.5	0.53	
Fog-Mist (48)	4.65±4.7	0.11±0.1	0.39±0.3	0.28	
Non-Event (62)	5.09±3.7	0.17±0.2	0.46±0.3	0.37	
Whole (127)	7.23±8.3	0.24±0.5	0.49±0.4	0.49	

The radioactivity concentrations by atmospheric phenomenon are summarized in *Table* 3. The mean radioactivity concentrations of 40 K, 232 Th, and 238 U were 7.23±8.3 μ Bq m⁻³, 0.24±0.5 μ Bq m⁻³, and 0.49±0.3 μ Bq m⁻³, respectively. The radium equivalent activity was 33.25 Bq kg⁻¹.

The mean atmospheric radioactivity concentrations of 232 Th and 238 U per air volume obtained from 7 fine dust samples during the Asian dust event were 1.60 ± 1.2 μ Bq m⁻³ (range: 0.23 to 3.63 μ Bq m⁻³) and 1.31 ± 0.7 μ Bq m⁻³ (range: 0.37 to 2.18 μ Bq m⁻³), respectively,

which exceeded the global reference values. These were 5.5, 9.4, and 2.9 times higher than the mean atmospheric radioactivity concentrations of 40 K, 232 Th and 238 U per air volume, respectively, from 62 samples during the non-event period. The 232 Th/ 238 U ratio was also high for Asian dust, at 1.22, compared to other atmospheric phenomena (0.28-0.53), including haze, fog-mist, and non-event cases.

Fig. 2 shows the correlations of the investigated natural isotopes. The correlation between 232 Th and 238 U activity concentrations in total fine dust was



Fig. 2. Correlation between 232 Th and 238 U concentrations in (a) whole fine dust and (b) Asian dust.

positive ($R^2 = 0.5092$, *p*-value < 0.0001). This confirms that the two isotopes come from the same source, i.e., the Earth's crust (*Fig.* 2(a)). The correlation between ²³²Th and ²³⁸U activity concentrations in Asian dust is positive ($R^2 = 0.8472$, *p*-value = 0.125), suggesting that dust from China has a higher ²³²Th content than dust from other regions (*Fig.* 2(b)).

3.3. Air mass inflow paths

Using the National Oceanic and Atmospheric Administration's (NOAA) Hybrid Single Particle Lagrangian Integrated Trajectory (HYSPLIT4) model,²⁴ the 5-day backward trajectories of air entering the Gosan site was analyzed. The percentage of air entering the Gosan site on the day of the measurement was



Fig. 3. Results of Gosan site backtracking for 5 days during the study period.

33.9%, 38.6%, 15.0%, and 12.6% for Sector I (Mainland China), Sector II (Korean Peninsula), Sector III (Japan), and Sector IV (North Pacific), respectively (*Fig.* 3).

The ⁴⁰K, ²³²Th and ²³⁸U concentrations in sector I were 11.23, 0.43, and 0.65 μ Bq m⁻³, respectively, which were greater than those in other sectors (sector II: 6.28±6.9, 0.19±0.3, 0.40±0.3 μ Bq m⁻³; Sector III: 4.13±3.2, 0.08±0.1, 0.46±0.4 μ Bq m⁻³; Sector IV: 3.08±3.1, 0.10±0.1, 0.38±0.3 μ Bq m⁻³). They are, however, still lower than the global averages of the mean atmospheric activity concentrations of ²³²Th and ²³⁸U. The ratio of ²³²Th/²³⁸U activity in Sector I was 0.66, also higher than the other sector ratios (Sector II: 0.48, Sector III: 0.17, Sector IV: 0.26). Since Sector I is the dominant source of atmospheric inputs on Asian dust days, it can be said that this sector has a higher ²³²Th content than dust sources in other regions.

4. Conclusions

Radioactivity concentrations of natural radionuclides by atmospheric event (Asian dust, haze, fog-mist, and non-event) were measured in 127 particulate matter samples from Jeju Island collected throughout 2015.

The mean atmospheric radioactivity concentrations of 238 U, 232 Th, and 40 K in all samples were 0.49, 0.24, and 7.23 μ Bq m⁻³, respectively, which is lower than the global average concentration in fine dust. The

 232 Th/ 238 U radioactivity ratio for the entire sample was 0.49. The average concentrations of 238 U and 232 Th in fine dust during the Asian dust period were 1.31 and 1.60 µBq m⁻³, respectively, which were higher than the global average concentrations and the values for the other three atmospheric events. The 232 Th/ 238 U radioactivity ratio for the Asian dust period was 1.22, which was higher than the ratios for the other three atmospheric events.

After analyzing the concentrations by dividing the airborne pathways by 4, it was found that the mainland China pathway had higher concentrations than the other three pathways. The ²³²Th/²³⁸U radioactivity ratio of the mainland China pathway was 0.66, which was higher than the other pathway ratios.

Acknowledgements

This work was studied with the support of Jeju Green Environment Center in 2023 and was funded by the Nuclear Research and Development program of the National Research Foundation of Korea (NRF) grant funded by the Korean government (Ministry of Science and ICT; RS-2023-00248338). The authors thank Prof. C.-H. Kang and Dr. J.-M. Song at Jeju National University for their help with sample preparation.

References

- United Nations Scientific Committee on Effects of Atomic Radiation, *Sources and Effects of Ionizing Radiation*, UNSCEAR 2000 Report Vol. I. United Nations, New York, 2000.
- D. Shahbazi-Gahrouei, M. Gholami, and S. Setayandeh, Adv. Biomed. Res., 2(3), 65 (2013). https://doi.org/10.4103/ 2277-9175.115821
- International Atomic Energy Agency, 'Control of radioactive waste disposal into the marine environment, Safety Series No.61.', International Atomic Energy Agency, Vienna, 1978.
- T. V. Ramachandran, *Iran J. Radiat. Res.*, 9, 63-76 (2011).
- 5. S. A. Al-Kahtani, M. A. Farouk, and A. A. Al-Zahrani,

J. Radioanal. Nucl. Chem., 250(1), 93-95 (2001). https://doi.org/10.1023/A:1013224516065

- S. Singh, A. Rani, and R. K. Mahajan, *Radiat. Meas.*, 39(4), 431-439 (2005). https://doi.org/10.1016/j.radmeas. 2004.09.003
- G. D. Kim, C. H. Eum, and J. H. Bang, *Anal. Sci. Tech.*, 20(6), 460-467 (2007). https://doi.org/10.5806/AST.2007. 20.6.460
- R. Mehra, K. Badhan, R. G. Sonkawade, S. Kansal, and S. Singh, *Indian J. Pure Appl. Phys.*, 48, 805-808 (2010).
- 9. A. Abdul-Hadi, W. Al-Qadhi, and E. El-Zeen, J. Radioanal. Nucl. Chem., 290, 261-266 (2011). https://doi.org/10.1007/s10967-011-1214-8
- H. Al-Sulaiti, T. Nasir, K. S. Al-Mugren, N. Alkhomashi, N. Al-Dahan, M. Al-Dosari, D. A. Bradley, S. Bukhari, M. Matthews, P. H. Regan, T. Santawamaitre, D. Malain, and A. Habib, *Appl. Radiat. Isotopes*, **70**, 1344-1350 (2012). https://doi.org/10.1016/j.apradiso.2011.11.015
- R. Mehra and M. Singh, J. Phys., 36, 289-297 (2012). https://doi.org/10.3906/fiz-1105-9
- Z. A. Tayyeb, A. R. Kinsara, and S. M. Farid, *J. Environ. Radioact.*, **38**(1), 97-104 (1998). https://doi.org/10.1016/ S0265-931X(97)00014-3
- 13. M. W. Kadi, JKAU: Sci., 17, 197-204 (2005).
- W. R. Alharbi and J. H. Alzahrani, J. Am. Sci., 8(10), 651-656 (2012). https://doi.org/10.7537/marsjas081012.88
- S. M. Farid, Med. J. Islam. World Acad. Sci., 20(3), 84-93 (2012).
- C. H. Han and J. W. Park, J. Radioanal. Nucl. Chem., 316, 1173-1179 (2018). https://doi.org/10.1007/s10967-018-5873-6
- K. J. Kim, S. H. Lee, D. R. Hyeon, H. J. Ko, W. H. Kim, and C. H. Kang, *Anal. Sci. Tech.*, **27**(1), 1-10 (2014). https://doi.org/10.5806/AST.2014.27.1.1
- S. Melaku, R. Dams, and L. Moens, *Anal. Chim. Acta.*, 543(1-2), 117-123 (2005). https://doi.org/10.1016/j.aca. 2005.04.055
- A. Mainey and T. William, 'Compendium of methods for the determination of inorganic compounds in ambient air (Method IO-3.1; Selection, Preparation and Extraction of Filter Material). US EPA/625/R-96/010a', 1-27, 1999.
- 20. J. Magill and J. Galy, 'Radioactivity, Radionuclides,

Analytical Science & Technology

Radiation', Springer, Berlin, 2005.

- M. Berglund and M. E. Wieser, *Pure Appl. Chem.*, 83(2), 397-410 (2011). https://doi.org/10.1351/PAC-REP-10-06-02
- M. A. Zytoon, H. M. Aburas, and M. I. Abdulsalam, J. Environ. Radioact., 129, 148-155 (2014). https://doi.org/ 10.1016/j.jenvrad.2014.01.003
- A. K. Sam and N. Abbas, *Radiation Protection Dosimetry*, 93(3), 275-277 (2001). https://doi.org/10.1093/oxfordjournals. rpd.a006440
- W. H. Kim, C. H. Kang, D. S. Jung, H. J. Go, and W. Lee, *Anal. Sci. Tech.*, **21**(4), 304-315 (2008). https://doi.org/10.5806/AST.2008.21.4.304

Authors' Positions

Chung-Hun Han: Researcher & Part-time lecturerSohyeon Lim: Researcher & Graduate studentHee-Jung Im: Professor

Vol. 36, No. 4, 2023