

Determination of ²¹⁰Po and ²¹⁰Pb depositions in lichen and soil samples collected from Köprübaşı-Manisa, Turkey

Sermin Çam Kaynar¹ · Umit H. Kaynar² · Umran Hiçsönmez³ · Omer S. Sevinç⁴

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Abstract In this study, we aimed to determine the accumulations of ²¹⁰Po and ²¹⁰Pb in soil and lichen samples in Köprübaşı. The Köprübaşı district is home to the largest uranium deposits in Turkey. To date, there has been no study recorded in the literature related to ²¹⁰Po and ²¹⁰Pb depositions in lichens in Köprübaşı. Six different lichen species (Cladonia convoluta, Parmelina tiliacea, Physcia stellaris, Pleurosticta acetabulum, Xanthoparmelia conspersa, and Xanthoria parietina) as well as soil samples were collected from seven sampling locations around Köprübaşı. Lichens were used as biomonitors for ²¹⁰Po and ²¹⁰Pb deposition. The ²¹⁰Po and ²¹⁰Pb activity concentrations were measured in all the samples by alpha spectrometry. The activity concentrations in the lichen samples ranged from 64 to 577 Bq kg^{-1} with an average of 266 Bq kg⁻¹ for ²¹⁰Po and from 78 to 565 Bq kg⁻¹ with an average of 333 Bq kg^{-1} for ²¹⁰Pb. The activity ratios of 210 Po/ 210 Pb ranged from 0.80 to 1.99. In the lichen species, the mean ²¹⁰Po activity values varied from 154 Bq kg⁻¹ in

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Sermin Çam Kaynar Sermin.cam@cbu.edu.tr

- ¹ Department of Physics, Faculty of Sciences and Arts, Manisa Celal Bayar University, 45140 Manisa, Turkey
- ² Science Education, Faculty of Education, Manisa Celal Bayar University, 45900 Manisa, Turkey
- ³ Department of Chemistry, Faculty of Sciences and Arts, Manisa Celal Bayar University, 45140 Manisa, Turkey
- ⁴ Primary School Science Teaching, Duzce University, 81620 Manisa, Turkey

Pleurosticta acetabulum to 390 Bq kg⁻¹ in *Xanthoparmelia conspersa*. The range of the mean ²¹⁰Pb activity was between 153 Bq kg⁻¹ in *Cladonia convoluta* and 378 Bq kg⁻¹ in *Parmelina tiliacea*. In the soil samples, ²¹⁰Po and ²¹⁰Pb activity concentrations were ranged from 14 to 1268 Bq kg⁻¹ and from 19 to 1113 Bq kg⁻¹, respectively. While the values of ²¹⁰Po and ²¹⁰Pb measured in the lichen samples are comparable with those of the literature, the results of ²¹⁰Po and ²¹⁰Pb in the soil taken from the uranium mine are higher than the results of the literature studies.

Keywords ²¹⁰Po · ²¹⁰Pb · Manisa · Lichen

1 Introduction

Lichens are symbiotic units formed from fungi and algae [1]. They have no roots or specialized structures for water and gas exchange. They need only a few nutrients and grow relatively slowly. They are perennial and live in various growth places (on soil, rocks, mosses, and trees) [2]. They can only live in places with clean air. They show a susceptibility to dirty air and are, therefore, a good indicator to imply whether the air of an area is clean or not [3]. Lichens have the ability to effectively accumulate a high level of different pollutants (SO2, NO2, ozone, heavy metals, and radionuclides) from the environment. They can be used as biomonitors of air pollution [4], because they highly depend on the atmospheric deposition for nutrients and easily collect pollutants in their thallus in line with atmospheric concentrations [4, 5]. Since the lichens absorb ²¹⁰Po from the air, they act as bioindicators for determining the amount of ²¹⁰Po in the atmosphere of that region.

²¹⁰Po is the decay product of ²¹⁰Pb in the ²³⁸U decay series. It widely spreads in nature, the atmosphere, and oceans. ²¹⁰Po has short half-life ($t_{1/2}$, 138.4 days), but it stays in the atmosphere for a long time because of its parent radionuclides [²¹⁰Pb ($t_{1/2}$ =22.3 years), ²²⁶Ra ($t_{1/2}$ =1602 years], which have long half-lives. Therefore, the amount of ²¹⁰Po in nature depends largely on the amount of ²¹⁰Pb [6].

²²²Rn emanation from the earth surface into the atmosphere is the main source of ²¹⁰Po. It travels back to the earth surface as attached to airborne particles [7]. Uranium mines are also sources [8]. Extra atmospheric ²¹⁰Po can originate from the external sources, such as volcanic emissions, inflow of the air, and anthropogenic emissions, e.g., emission from coal combustion, waste discharge from the gas, phosphate, and oil industries [7, 9].

²¹⁰Po decays directly to ²⁰⁶Pb by emitting an alpha particle with 5.30 MeV energy [6, 7, 9]. Alpha particles are the most dangerous radiation type and are 400 times more radioactive than uranium. They have a low energy; therefore, they do not pass through a human skin, but easily permeate into the body from the respiratory tract, mouth or open wounds in the skin and pass through living cells [9]. If ²¹⁰Po is taken inside the body, large part of ingested ²¹⁰Po passes through the gastrointestinal tract within a few days. It is excreted with the feces. The retained ²¹⁰Po passes into the blood, it is stored by the soft tissue including the bone marrow. The biologic half-life of ²¹⁰Po is almost 50 days [10]. ²¹⁰Po and other radon products inhaled from the air can lead to lung cancer. Moreover, when ²¹⁰Po is ingested, it is hazardous to human health [9]. Radiation doses occurring in humans increase the risk of cancer. Very high radiation doses cause damage to the tissue and organs and overdoses can be fatal.

Studies on ²¹⁰Po in the environment are important as tracing the atmospheric emissions in populated areas is related to human health [9]. There have been several studies related to ²¹⁰Po and ²¹⁰Pb depositions using lichens [1, 6, 11–14]. However, no study has been performed on the lichens radioactivity in Köprübaşı-Manisa. In the present study, we determined ²¹⁰Po and ²¹⁰Pb activity concentrations in lichen and soil samples collected from the Köprübaşı district of Manisa. The determination of ²¹⁰Po and ²¹⁰Pb accumulations in the studied area is important in determining the environmental pollution and the radiation exposure of the inhabitants.

2 Materials and methods

2.1 Study area

Köprübaşı district is located 120 km northeast of Manisa in the Aegean region of Turkey. It is bordered by Gördes in the northwest, Demirci in the northeast, Gölmarmara in the west, Demirköprü Dam in the southeast, Salihli in the southwest, Kula and Alaşehir in the south, and Selendi in the east. Köprübaşı is located at 38°44'N latitude and 28° 24'E longitude. It has a surface area of 447 km². The Dibek Mountains are in the southwest of the district, the Çanak Mountains in the northeast, and Kayran Mountains in the north [15]. The elevation of Köprübaşı is 250 m. It has a Mediterranean climate with hot and dry summers, and rainy and mild winters.

According to the General Directorate of Mineral Research and Exploration of Turkey (MTA) inventory of mining in the Aegean region, the Köprübaşı area contains marble, feldspar, phosphate, sulfur, titanium, and zeolite deposits [16]. There are also uranium deposits in fluvial sedimentary rocks. The Köprübaşı uranium deposit was discovered by MTA in 1961 [17]. It is the largest known uranium deposit in Turkey [17]. Apatite, biotite, feldspar, ilmenite-magnetite, muscovite, quartz, rutile, tourmaline, and zircon minerals are found in Köprübaşı uranium deposits [18].

In this work, lichen and soil samples were collected from seven sampling locations in Köprübaşı in July 2013. The sampling locations are shown in Fig. 1.

The lichen samples were collected from the surfaces of trees and then placed into the paper bags after wrapping in paper towels. The sampling locations, location number, date, type of substrate, altitude, and coordinate information determined by Garmin brand GPS devices were noted. Soil samples were also obtained at each of the locations.

2.2 Determination of ²¹⁰Po and ²¹⁰Pb activity concentrations

The inessential materials were cleaned from each sample in the laboratory. Samples were dried at room temperature and then grinded and sieved. One gram of sample was weighed and concentrated acids (HNO₃ and H₂O₂) were added, before leaving the samples overnight. The following day, the solution was heated on a hot plate and evaporated to dryness. Mixed acids (HNO₃ and HCl) were added to the dried residue. The solution was then evaporated to dryness again by heating on a hot plate. This process was repeated a total of four times. After evaporation, 0.5 M HCl acid was added to the dry residue. The solution was filtered with filter paper (particle retention 10–



Fig. 1 (Color online) Map showing the sampling locations (1, 2, 3, 4, 5, 6 and 7) at Köprübaşı-Manisa: (1) Çarıklar village, (2) Kozakli village, (3) Kasar uranium deposit, (4) West of Borlu, (5) Northeast of Köprübaşı, (6) the Köprübaşı-Gordes road, and (7) the Salihli-Simav road

15 μ m, 125 mm diameter, and 84 g m⁻² weight). Ascorbic acid was added to the solution to reduce Fe³⁺. A copper disc of diameter 2.5 cm was prepared. ²¹⁰Po was spontaneously accumulated onto the copper disc at 70 °C for 5 h.

The ²¹⁰Po accumulated onto the copper disc was counted for 5 h with an alpha spectrometer equipped with a PIPS detector placed in a vacuum chamber connected to a 1024 multichannel analyzer (Canberra). The energy resolution was ≤ 20 keV with a detector source spacing equal to the detector diameter. The detector efficiency was $\geq 25\%$ of the detector source spacing of ≤ 10 mm. The background was ≤ 1 count/hour above 3 MeV. The system calibration was performed with an ²⁴¹Am point source.

The measurement of ²¹⁰Po was performed using the alpha particle emission peak with 5.30 MeV energy, using ²⁰⁹Po as the internal tracer (National Institute of Standards & Technology, SRM 4326 consists of radioactive polonium-209 chloride and hydrochloric acid dissolved in 5 ml of distilled water. The solution mass was 5.160 ± 0.003 g. It is enclosed in a flame sealed NIST borosilicate glass ampoule. The ²⁰⁹Po massic activity of the solution was 85.42 Bq g⁻¹).

In this study, the chemical efficiency was calculated as 37.7% using the ²⁰⁹Po standard. The total efficiency was found to be 102%. The ²¹⁰Po activity concentration for each sample was corrected for recovery using the total efficiency.

The ²¹⁰Pb activity measurement was performed indirectly from its measured product ²¹⁰Po activity after reaching radioactive equilibrium. The ²¹⁰Pb and ²¹⁰Po isotopes reached secular equilibrium at least 6 months after the first ²¹⁰Po electro-deposition. At this time, ²¹⁰Po grew from the ²¹⁰Pb of the sample. The ²¹⁰Po deposition in the lichen samples for the ²¹⁰Pb activity was collected on the cupper discs at 70 °C for 5 h and counted for 5 h by the alpha spectrometer. The ²¹⁰Pb activity concentration from the measured ²¹⁰Po activity in the analyzed samples was calculated by the Bateman equation [19]:

$$A_0(^{210}\text{Pb}) = \frac{A_2(^{210}\text{Po})}{1 - e^{-\lambda(t_2 - t_1)}},$$

where A_0 (²¹⁰Pb) is the activity of a sample during the collection time, A_2 (²¹⁰Po) is the activity of ²¹⁰Po ingrown from ²¹⁰Pb after the second electro-deposition, t_1 is the time between the sample collection time and the first ²¹⁰Po

activity measurement, t_2 is the time between the sample collection date and the second ²¹⁰Po activity measurement, and λ is the decay constant of ²¹⁰Po [19].

2.3 Determination of gamma radioactivity in soil samples

The soil samples collected from the seven sampling locations in Köprübaşı were milled and dried in an oven in the laboratory, before being sieved and weighed at 100 g in a polyethylene beaker. They were tightly closed and stored for as a minimum of 4 weeks to permit the ²³⁸U and ²³²Th to arrive at equilibrium with their decay products. The gamma measurements were performed by a gamma ray spectrometer using a 3×3 inch NaI(Tl) (ORTEC-905-4) detector. The system calibration was made with the standard samples (52% K, 625 ppm eU and 150 ppm eTh) under appropriate conditions. The best available resolution was < 7.5% for the gamma peak of ¹³⁷Cs (662 keV). The activity concentrations for ⁴⁰K, ²³⁸U, and ²³²Th were evaluated from the radioactive potassium peak (1.46 MeV), the 214 Bi peak (1.76 MeV), and the 208 Tl peak (2.61 MeV), respectively. All the samples were counted for 7200 s. The ⁴⁰K, ²³⁸U and ²³²Th activities were then calculated.

3 Results and discussion

3.1 Results of ²¹⁰Po and ²¹⁰Pb activity concentrations in samples

The 17 lichen samples of six different species (*Cladonia convoluta, Parmelina tiliacea, Physcia stellaris, Pleurosticta acetabulum, Xanthoparmelia conspersa,* and *Xanthoria parietina*) and seven soil samples were collected from seven sampling locations in Köprübaşı district. The lichen species are presented in Fig. 2. The results of the ²¹⁰Po and ²¹⁰Pb activity measured in the samples, according to sampling location, are presented in Table 1.

At location 1, according to Table 1, there were two different lichen species and the highest ²¹⁰Po and ²¹⁰Pb activities of 577 and 493 Bq kg⁻¹, respectively, were obtained in *Xanthoria parietina*. These values were the highest activities among all lichen samples. In location 2, while the highest value for ²¹⁰Po was obtained as 290 Bq kg⁻¹ in *Xanthoparmelia conspersa*, the highest value for ²¹⁰Pb was calculated as 270 Bq kg⁻¹ in *Physcia stellaris*. There were six different lichen species in location 3, where the Kasar uranium deposit is located. The highest activities for ²¹⁰Po and ²¹⁰Pb of 504 and 565 Bq kg⁻¹, respectively, were obtained for *Parmelina tiliacea*. In location 4, there was only one lichen species, namely *Xanthoria parietina*. In location 5, there were three

different lichen types, with the highest obtained activities for 210 Po (171 Bq kg⁻¹) and 210 Pb (144 Bq kg⁻¹) in Xanthoria parietina. In location 6, the highest activities for 210 Po (176 Bq kg⁻¹) and 210 Pb (190 Bq kg⁻¹) were detected in Parmelina tiliacea. In location 7, there was only one lichen species. The activities of ²¹⁰Po and ²¹⁰Pb in Xanthoria parietina were 450 and 227 Bq kg⁻¹, respectively. The polonium accumulation is also affected by such factors as the age, health, and type of the lichen species, including their positions on trees, surface structure [20], altitudes where the lichens grow, and meteorological conditions (wind and rain). The mean annual rainfall of Köprübası is 575.8 mm and average annual temperatures range from 4.1 to 27.3 °C in 2013. Xanthoria parietina is the most common lichen species and was found at all locations except for location 2. It was assumed to be the most suitable bioindicator for pollution in the study area due to its high capture efficiency, geographical and climatic suitability, and common occurrence.

While the ²¹⁰Po activity levels in soil samples collected from six locations, except for location 3, varied between 14 and 53 Bq kg⁻¹, the ²¹⁰Pb activity levels ranged from 19 to 55 Bq kg⁻¹. It can be seen that the activity concentration in the samples in location 3 (1268 Bq kg⁻¹ for ²¹⁰Po and 1113 Bq kg⁻¹ for ²¹⁰Pb) were higher than in the samples from the other locations. Uranium mining had previous occurred in this area (location 3). In the other locations, ²¹⁰Po and ²¹⁰Pb concentrations in the soil samples were lower than those for lichen. This implies that lichens do not feed from the soil as they do not have roots and are affected by atmospheric fallouts.

The 210 Po and 210 Pb activities in the lichen and soil samples in this study were compared with those in the literature in Table 2. In the study, the mean 210 Po and 210 Pb activity levels in the lichen samples were similar to those of similar studies in the literature. The 210 Po/ 210 Pb ratio in the present study was found to be unity, implying the equilibrium between the 210 Po and 210 Pb radioisotopes. The 210 Po measured in lichens originates from 210 Pb, which is its parent in the uranium chain.

The mean 210 Po and 210 Pb activities according to the lichen species are shown in Fig. 3. The mean activity ratios of 210 Po/ 210 Pb are presented in Table 3.

The mean ²¹⁰Po and ²¹⁰Pb activities in lichen species ranged from 154 to 390 Bq kg⁻¹ and from 153 to 378 Bq kg⁻¹, respectively. While the highest mean activity for ²¹⁰Po was detected in *Xanthoparmelia conspersa*, the lowest mean activity was seen in *Pleurosticta acetabulum*. For ²¹⁰Pb, while the highest mean activity was seen in *Parmelina tiliacea*, the lowest activity was found in *Cladonia convolute* (Table 3).

The mean activity ratios of 210 Po/ 210 Pb in the lichen species ranged from 0.85 to 1.52. The activity ratio of









Fig. 2 (Color online)Lichen species collected from the Köprübaşı district: a Cladonia convoluta, b Parmelina tiliacea, c Physica stellaris, d Pleurosticta acetabulum, e Xanthoparmelia conspersa, and f Xanthoria parietina

Table 1 ²¹⁰ Po and ²¹⁰ Pb activity concentrations of the samples according to sampling location (Bo	l kg	(-1)	
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Samples (n: sample number)	Location 1		Location 2		Location 3		Location 4		Location 5		Location 6		Location 7	
	²¹⁰ Po	²¹⁰ Pb												
Lichen species														
Cladonia convoluta (n: 2)	138	105	-	-	328	201	-	-	-	-	-	-	-	-
Parmelina tiliacea (n: 2)	_	_	_	_	504	565	_	_	_	_	176	190	_	_
Physcia stellaris (n: 3)	_	_	222	270	331	378	_	_	66	79	_	_	_	_
Pleurosticta acetabulum (n: 2)	-	-	-	-	194	215	-	-	114	143	-	-	-	-
Xanthoparmelia conspersa (n: 2)	-	-	290	256	489	457	-	-	-	-	-	-	-	-
Xanthoria parietina (n: 6)	577	493	_	_	245	267	64	78	171	144	148	154	450	227
Soil samples	45	51	53	55	1268	1113	24	40	14	22	19	24	21	19
(<i>n</i> : 7)														

²¹⁰Po/²¹⁰Pb was unity in Xanthoparmelia conspersa. This demonstrates the equilibrium between the ²¹⁰Po and ²¹⁰Pb radioisotopes. In three lichen species (Parmelina tiliacea, Physcia stellaris, and Pleurosticta acetabulum), the ²¹⁰Pb deposition was greater than the ²¹⁰Po deposition. This disequilibrium indicated the presence of ²¹⁰Pb due to the atmospheric deposition in the lichen species. Since the ground surface is the main source of airborne ²¹⁰Pb, the air concentrations of ²¹⁰Pb show the local, geological, and global climatological background of the areas observed

Study area	Lichen			Soil	Reference		
	²¹⁰ Po	²¹⁰ Pb	²¹⁰ Po/ ²¹⁰ Pb	²¹⁰ Po	²¹⁰ Pb	²¹⁰ Po/ ²¹⁰ Pb	
Norway	70–212	_	_	_	_	_	[13]
	(140.5)						
India	_	_	_	6.6-35.3	11.7-85.7	0.1-1.5	[21]
				(19.3)	(38.5)	(0.45)	
Norway	39-137.5	150-188	0.3-0.7	36.9-42.87	39.4-46.05	0.86-1	[12]
	(88.3)	(169)	(0.5)	(39.8)	(42.7)	(0.9)	
Hungary	_	_	_	37–184	_	_	[7]
				(85.3)			
Turkey-Emendere	185	_	_	66	_	_	[14]
Turkey-Çan	98.4-206.1	153.6-326.4	0.5-0.78	42.8-135.7	30.8-177.6	0.62-1.86	[11]
	(161.6)	(259.1)	(0.6)	(79.9)	(91.5)	(0.87)	
Western Turkey	151–593	97-360	_	_	-	_	[22]
	(378)	(233)					
Western Turkey	117-569	84-291	1.39-2.33	_	-	_	[1]
	(365)	(206)	(1.7)				
Köprübaşı-Turkey	154–390	153-378	0.79-1.98	14-1268	19–1113	0.74-1.13	This study
	(267)	(256)	(1.04)	(29.3)	(35.2)	(0.87)	

Table 2 ²¹⁰Po and ²¹⁰Pb activity results (Bq kg⁻¹) and ²¹⁰Po/²¹⁰Pb ratios in the lichen and soil samples in the literature



Fig. 3 Mean ²¹⁰Po and ²¹⁰Pb activity in the lichen species at Köprübaşı

[23]. In the *Cladonia convoluta* and *Xanthoria parietina* lichen species, the ²¹⁰Po deposition was higher than the ²¹⁰Pb deposition (Table 3). The activity ratio of ²¹⁰Po/²¹⁰Pb in the *Cladonia convolute* lichen species is higher than the other results. This lichen species was collected at two locations (locations 1 and 3). When the activity ratios for this species were calculated according to the sampling locations and were 1.3 and 1.63 in locations 1 and 3 (Kasar uranium deposit), respectively. According to the results, the ²¹⁰Po activity concentrations were higher than the ²¹⁰Pb

activity concentrations. This result shows that the polonium was a contribution not only from the predominant ²¹⁰Pb but also from the environment. The Kasar uranium deposit is located in location 3 and it can be concluded that the uranium mine affected the polonium activity. The polonium accumulation is affected by the deposits of marble, feldspar, phosphate, sulfur, titanium, and zeolite. This is because the level of ²¹⁰Po in the atmosphere of tungsten, molybdenum, iron, and phosphate rocks is higher [24, 25]. The Köprübaşı uranium deposits are found in apatite,

Table 3 Mean ²¹⁰Po and ²¹⁰Pb activities and the activity ratios of ²¹⁰Po/²¹⁰Pb of the lichen species

Lichen species	²¹⁰ Po (Bq kg ⁻¹)	210 Pb (Bq kg ⁻¹)	The activity ratio of ²¹⁰ Po/ ²¹⁰ Pb
Cladonia convoluta (n=2)	233	153	1.52
Parmelina tiliacea $(n=2)$	340	378	0.90
Physcia stellaris (n=3)	206	242	0.85
Pleurosticta acetabulum $(n=2)$	154	179	0.86
<i>Xanthoparmelia conspersa (n=2)</i>	390	356	1.09
Xanthoria parietina (n=6)	276	227	1.21
Min.	154	153	0.85
Max.	390	378	1.52
Mean	267	256	1.04

biotite, feldspar, ilmenite-magnetite, muscovite, quartz, rutile, tourmaline, and zircon minerals.

A correlation graphic of ²¹⁰Po activity versus ²¹⁰Pb activity in the lichen samples is given in Fig. 4. It can be seen that there was a positive correlation of 0.91 between the two radionuclides, with an R^2 value of 0.78.

3.2 Results of gamma radioactivity in soil samples

The 40 K, 238 U, and 232 Th activities were measured by gamma spectrometry and the obtained results are given in Table 4.

Standard error values for the 40 K, 238 U, and 232 Th activities were calculated separately and were 129, 161, and 8, respectively. These are shown in Fig. 5.

In the soil samples, the 40 K activity ranged between 250.25 and 1064.60 Bq kg⁻¹ with an average value of 506.19 Bq kg⁻¹. The highest concentration of 40 K was measured in the sample collected from location 5 (Table 4 and Fig. 5). The world average for 40 K is accepted as

Table 4 40 K, 238 U, and 232 Th activity concentrations of soil samples in the Köprübaşı district (Bq kg⁻¹)

Soil samples	⁴⁰ K	²³⁸ U	²³² Th	
1	601.91	53.86	16.41	
2	250.25	52.84	34.61	
3 ^a	_	1267.8	80.57	
4	703.81	53.77	32.14	
5	1064.60	42.60	27.19	
6	569.92	52.21	33.26	
7	352.80	41.73	14.61	
Min.	250.25	41.73	14.61	
Max.	1064.60	1267.8	80.57	
Mean	506.19	49.5	39.92	

^a Kasar uranium deposit

400 Bq kg⁻¹ [26], and therefore, 71.5% of the results in this study exceeded the world average.







The ²³⁸U activity in the soil samples ranged from 41.73 to 1267.8 Bq kg⁻¹, with an average value of 49.5 Bq kg⁻¹, except at location 3. The ²³⁸U activity concentration in all soil samples was higher than the world average for ²³⁸U which is accepted as 35 Bq kg⁻¹ [26]. In particular, the ²³⁸U activity concentration in the soil sample taken from the Kasar uranium mine (location 3) is very high (approximately 36 times the world average).

The ²³²Th activity varied from 14.61 to 111.92 Bq kg⁻¹ with a mean value of 39.92 Bq kg⁻¹. The results of two samples (location 2 and location 3) were lower than the world average of 30 Bq kg⁻¹ [26]. The mean of the ⁴⁰K, ²³⁸U, and ²³²Th activities in this study was exceeded the accepted values (400, 35, and 30 Bq kg⁻¹, respectively).

In the soil samples, the correlation between the 238 U activity concentration measured by gamma spectrometry and 210 Po activity concentration measured by alpha spectrometry is demonstrated in Fig. 6. It can be seen that the correlation between the 238 U and 210 Po activity is a positive (0.75) with an R^2 value of 0.998.

4 Conclusion

In this study, the ²¹⁰Po and ²¹⁰Pb activities in six different lichen species were measured by alpha spectrometry. The most common of the lichen species was *Xanthoria parietina* in the Köprübaşı district. The highest activities for ²¹⁰Po (577 Bq kg⁻¹) and ²¹⁰Pb (565 Bq kg⁻¹) in the lichen samples were detected in *Xanthoria parietina* and *Parmelina tiliacea*, respectively.

The highest mean activities of ²¹⁰Po and ²¹⁰Pb were detected in *Xanthoparmelia conspersa* and in *Parmelina tiliacea*, respectively. The lowest mean activities were seen in *Pleurosticta acetabulum* and *Cladonia convolute* for ²¹⁰Po and ²¹⁰Pb, respectively. The age, health, and locations of the lichens differed, and therefore, these factors can cause the accumulation of radionuclides in different proportions in the same species.

According to the lichen species, the activity ratio of ²¹⁰Po/²¹⁰Pb was unity in *Xanthoparmelia conspersa*, demonstrating the equilibrium between the ²¹⁰Po and ²¹⁰Pb



radioisotopes. However, this ratio in the other lichen species is varied from 0.85 to 1.52. In the lichen samples, there was a positive correlation of 0.91 between the ²¹⁰Po and ²¹⁰Pb activity with an R^2 value of 0.78. In addition, in the soil samples, there is a positive relation between the ²³⁸U activity measured by gamma spectrometry and ²¹⁰Po activity measured by alpha spectrometry with an R^2 value of 0.998.

In this study, the ²¹⁰Po activity concentrations in the air using lichens were studied to investigate the dose limits affecting the human health of those living in this area. The ²¹⁰Po and ²¹⁰Pb levels of this area were determined and the obtained data important, because no such study has been performed in this region before. The results of this study can be used as the basic data in future studies, e.g., production of distribution maps.

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