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Uranium isotopes as a tracer of groundwater transport studies

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The uranium chain radionuclides are usually used for a wide range of application in the Earth Sciences. The $^{234}\text{U}/^{238}\text{U}$ activity ratio is used as a geochemical tool to investigate transport and flow relationships in major hydrological reservoirs, groundwater pattern and it is highly useful for interpreting timescales of weathering. In rocks older than a few million years, $^{234}\text{U}/^{238}\text{U}$ activity ratios should be in secular equilibrium. Nevertheless, the highly energetic alpha decay of ^{238}U damages a mineral's crystalline lattice and allows ^{234}U to be more mobile during weathering of rock by atmospheric water [1].

In this work, the activity concentration and isotopic ratio of uranium isotopes ($^{234}\text{U}/^{238}\text{U}$) were determined in selected thermal groundwater, ground and surface water samples from central Poland. Additionally the radionuclides concentration in one sample of thermal groundwater from southwest Poland were determined. Uranium isotopes after co-precipitation with hydrated manganese dioxide were separated from other natural radionuclides by extraction chromatography resin (Dowex 1x8). The activity concentrations were determined by using α spectrometry with PIPS detector (Camberra).

Concentration of ^{234}U and ^{238}U in the examined waters varied from 0.75 mBq/dm³ to 16.8 mBq/dm³ and from 0.64 mBq/dm³ to 45.5 mBq/dm³ respectively. The highest uranium concentrations were measured in the thermal groundwater from Mszczonow and Cieplice, while the lowest were observed in ground water from Plock which are got out from Cretaceous geological formation. The $^{234}\text{U}/^{238}\text{U}$ activity ratio varied from 0.370 (Cieplice) to 1.205 (Plock –Tertiary geological formation). It confirms that the uranium isotopic ratio can be useful indicator for determination of surface water infiltration input into the underground water reservoirs.

[1] Camacho A., Devesa R., Valles I., Serrano I., Soler J., Blazquez S., Ortega X., Matia L.: J. Environ. Radioact. 101, 1048 (2010)

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