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Kakovost vode - Radij Ra-226 - 2. del: Preskusna metoda z emanometrijo (ISO 13165-2:2014)

Water quality - Radium-226 - Part 2: Test method using emanometry (ISO 13165-2:2014)

Wasserbeschaffenheit - Radium 226 - Teil 2: Untersuchungsverfahren mittels Emanometrie (ISO 13165-2:2014)

Qualité de l'eau - Radium 226 - Partie 2: Méthode d'essai par émanométrie (ISO 13165-2:2014)

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**Water quality — Radium-226 —
Part 2:
Test method using emanometry**

Qualité de l'eau — Radium 226 —

Partie 2: Méthode d'essai par émanométrie

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ISO 13165-2:2014(E)

Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT) see the following URL: Foreword - Supplementary information

The committee responsible for this document is ISO/TC 147, *Water quality*, Subcommittee SC 3, *Radioactivity measurements*.

ISO 13165 consists of the following parts, under the general title *Water quality — Radium-226*:

- *Part 1: Test method using liquid scintillation counting*
- *Part 2: Test method using emanometry*

The following parts are under preparation:

- *Part 3: Test method using coprecipitation and gamma spectrometry*

Introduction

Radioactivity from several naturally occurring and human-made sources is present throughout the environment. Thus, water bodies (surface waters, ground waters, sea waters) can contain radionuclides of natural and human-made origins:

Natural radionuclides, including potassium-40, and those of the thorium and uranium decay series, in particular radium-226, radium-228, uranium-234, uranium-238, and lead-210, can be found in water for natural reasons (e.g. desorption from the soil and wash-off by rain water) or releases from technological processes involving naturally occurring radioactive materials (e.g. the mining and processing of mineral sands or phosphate fertilizer production and use).

Human-made radionuclides such as transuranium elements (americium, plutonium, neptunium, curium), tritium, carbon-14, strontium-90, and some gamma emitters radionuclides can also be found in natural waters as they can be authorized to be routinely released into the environment in small quantities in the effluent discharge from nuclear fuel cycle facilities and following their use in unsealed form in medicine or industry. They are also found in the water due to the past fallout of the explosion in the atmosphere of nuclear devices and those following the Chernobyl and Fukushima accidents.

Drinking water can thus contain radionuclides at activity concentration which could present a risk to human health. In order to assess the quality of drinking water (including mineral waters and spring waters) with respect to its radionuclide content and to provide guidance on reducing health risks by taking measures to decrease radionuclide activity concentrations, water resources (groundwater, river, lake, sea, etc.) and drinking water are monitored for their radioactivity content as recommended by the World Health Organization (WHO).

The need of a standard on a test method of radium-226 activity concentrations in water samples is justified for test laboratories carrying out these measurements, required sometimes by national authorities, as they may have to obtain a specific accreditation for radionuclide measurement in drinking water samples.

Radium-226 activity concentration can vary widely according to local geological and climatic characteristics and ranges from 0,001 Bq l⁻¹ in surface waters up to 50 Bq l⁻¹ in natural groundwaters. The guidance level for radium-226 in drinking water as recommended by WHO is 1 Bq l⁻¹ (see Reference [11]).

NOTE The guidance level is the activity concentration (rounded to the nearest order of magnitude) with an intake of 2 l·d⁻¹ of drinking water for 1 year that results in an effective dose of 0,1 mSv·y⁻¹ for members of the public, an effective dose that represents a very low level of risk that is not expected to give rise to any detectable adverse health effect.

This International Standard is one of a series on determination of the activity concentration of radionuclides in water samples.

Water quality — Radium-226 —

Part 2: Test method using emanometry

WARNING — Persons using this document should be familiar with normal laboratory practice. This document does not purport to address all of the safety problems, if any, associated with its use. It is the responsibility of the user to establish appropriate safety and health practices and to ensure compliance with any national regulatory conditions.

IMPORTANT — It is absolutely essential that tests conducted in accordance with this document be carried out by suitably qualified staff.

1 Scope

This part of ISO 13165 specifies the determination of radium-226 (^{226}Ra) activity concentration in all types of water by emanometry.

The method specified is suitable for the determination of the soluble, suspended, and total ^{226}Ra activity concentration in all types of water with soluble ^{226}Ra activity concentrations greater than $0,02 \text{ Bq l}^{-1}$.

In water containing high activity concentrations of ^{228}Th , interference from ^{220}Rn decay products can lead to overestimation of measured levels (see [Figure A.2](#)).

2 Normative references

The following documents, in whole or in part, are normatively referenced in this document and are indispensable for its application. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 5667-3, *Water quality — Sampling — Part 3: Preservation and handling of water samples*

ISO/IEC 17025, *General requirements for the competence of testing and calibration laboratories*

ISO 80000-10, *Quantities and units — Part 10: Atomic and nuclear physics*

3 Terms, definitions and symbols

3.1 Terms and definitions

For the purposes of this document, the terms and definitions in ISO 80000-10 and the following apply.

3.1.1

reference measurement standard

measurement standard designated for the calibration of other measurement standards for quantities of a given kind in a given organization or at a given location

3.1.2

working measurement standard

measurement standard that is used routinely to calibrate or verify measuring instruments or measuring systems

Note 1 to entry: A working measurement standard can be used as a solution of known activity concentration obtained by precise dilution or dissolution of a reference standard.

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3.2 Symbols

For the purposes of this document, the symbols in ISO 80000-10 and the following apply.

c_A	^{226}Ra activity concentration in water, in becquerel per litre
c_A^*	decision threshold, in becquerel per litre
$c_A^\#$	detection limit, in becquerel per litre
$c_A^\triangleleft, c_A^\triangleright$	lower and upper limits of the confidence interval, in becquerel per litre
f_a	correction factor for ingrowth of ^{222}Rn in the bubbler, dimensionless
f_d	correction factor for the decay of ^{222}Rn in the detection volume, dimensionless
n	number of counting cycle
n_α	number of alpha-emitters present in the cell per becquerel of radon after a waiting time period between the filling time and the counting time of the cell (n_α is approximately 3 at a waiting time of 3 h for 1 Bq of radon)
N_0	number of background counts
N	number of gross counts
t_c	counting time (common to N, N_0), in seconds
t_i	time of the different steps of the measurement procedure, $i = 0, 1$ and 2
U	expanded uncertainty calculated by $U = ku(c_A)$ with $k = 2$
$u(c_A)$	standard uncertainty associated with the measurement result
V	volume of the test sample, in litre
ε	total efficiency including degassing efficiency and counting efficiency of the system for a count carried out with a radioactive equilibrium between ^{222}Rn and its short-lived decay products, in pulses per second per becquerel
λ	decay constant of the ^{222}Rn , per second

4 Principle

This test method is based on the emanation and scintillation counting of ^{222}Rn , a gaseous daughter product of ^{226}Ra , from a solution (see Reference [6]).

The measurement of ^{226}Ra activity concentration in water is carried out following two separate steps, the sample preparation followed by the measurement of the test sample.

Preparation consists of:

- dissolution when total or particulate radium is to be assayed;
- filtration when soluble radium is to be measured.

It is followed by pre-concentration, if necessary, and an accumulation of decay products without an initial separation.

After filtration and acidification, the test sample is placed in a bubbler (see [Figure B.1](#)) and stored for ingrowth of ^{222}Rn .

After a suitable ingrowth period, the radon gas is removed from the solution by purging with the radon-free gas and transferring it to a scintillation cell, whose internal surface is coated with silver-activated zinc sulfide, $\text{ZnS}(\text{Ag})$ (see [Figure C.1](#)) (see Reference [8]).

The alpha-particles produced by the decay of ^{222}Rn and its short-lived decay products (^{218}Po , ^{214}Po) transfer their energy as they pass through the scintillation medium. As they return to their ground state, the excitation electrons in the scintillation medium emit photons from the $\text{ZnS}(\text{Ag})$ coating that can be detected by a photomultiplier (PMT). The photomultiplier converts the photons into electrical pulses that are then counted. The pulse count is directly proportional to the activity concentration of radon and its decay products present in the scintillation cell.

The soluble ^{226}Ra activity concentration is calculated, taking into account the known steady state between ^{226}Ra and ^{222}Rn after transferring ^{222}Rn into a scintillation cell.

Given its high power of emanation, radon can also escape from particles suspended in water. In the case of the analysis of raw water, it is therefore advisable to dissolve the particulate fraction (see Reference [7]).

5 Reagents and equipment

5.1 Reagents

Unless otherwise stated, use only reagents of recognized analytical grade and distilled or demineralized water or water of equivalent purity and no undesirable radioactivity.

5.1.1 Concentrated nitric acid solution, HNO_3 .

5.1.2 Dilute nitric acid solution, less than or equal to 100 g l^{-1} , with no alpha-radioactivity.

5.1.3 Reference solution of ^{226}Ra .

5.1.4 Cellulose ester filter, $0,45 \mu\text{m}$ porosity.

5.1.5 Hydrochloric acid solution, if the particulate fraction of ^{226}Ra requires dissolution.

5.2 Equipment

Usual laboratory equipment and, in particular, the following.

5.2.1 Hotplates.

5.2.2 Needles, length of approximately 30 mm, diameter of approximately 1,5 mm.

5.2.3 Bubblers, minimum volume of 125 ml with two (poly)tetrafluoroethylene needle valves (see [Figure B.1](#)).

5.2.4 Specialized solid-state scintillation detection set, equipped with a photomultiplier.

5.2.5 Manometer, to measure pressure.

5.2.6 Flat-bottomed glass scintillation cell, volume of 250 ml to 500 ml.

The internal surface of the cell, apart from the bottom, is coated with silver-activated zinc sulfide, $\text{ZnS}(\text{Ag})$. The external surface is coated with a light-excluding material except on its flat bottom, which forms the measurement window (see [Figure C.1](#)).

5.2.7 Vacuum pump, to obtain a pressure reduction of about 300 Pa.