

International Standard

ISO 17294-2

Water quality — Application of inductively coupled plasma mass spectrometry (ICP-MS) —

Part 2:

Determination of selected elements including uranium isotopes

Qualité de l'eau — Application de la spectrométrie de masse avec plasma à couplage inductif (ICP-MS) —

Partie 2: Dosage des éléments sélectionnés y compris les isotopes d'uranium

Third edition 2023-10

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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 document 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).

ISO draws attention to the possibility that the implementation of this document may involve the use of (a) patent(s). ISO takes no position concerning the evidence, validity or applicability of any claimed patent rights in respect thereof. As of the date of publication of this document, ISO had not received notice of (a) patent(s) which may be required to implement this document. However, implementers are cautioned that this may not represent the latest information, which may be obtained from the patent database available at www.iso.org/patents. ISO shall not be held responsible for identifying any or all such patent rights.

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For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 147, *Water quality*, Subcommittee SC 2, *Physical, chemical and biochemical methods*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 230, *Water analysis*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This third edition cancels and replaces the second edition (ISO 17294-2:2016), which has been technically revised.

The main changes are as follows:

- with the incorporation of mercury in the previous edition, mercury has now been excluded as a hydrolysable and has now become a non-hydrolysable element because it was not in line with the other existing standards for the determination of mercury;
- the addition of a modifier has been clarified;
- titanium has been added to the scope.

A list of all parts in the ISO 17294 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at www.iso.org/members.html.

This corrected version of ISO 17294-2:2023 incorporates the following corrections:

- Note 1 to entry has been moved from the terminological entry <u>3.1.26</u> to <u>3.1.16</u>;
- the symbols "k", " α " and " β " have been revised in 3.2 and the symbols " η " and " σ " have been revised in Tables C.3 and C.5, respectively;
- Clause 12 b) has been editorially revised;
- "204TI" has been changed to "205Tl or ¹⁹³Ir" in Clause A.2;

- aula (A.3.

 c descriptions.)

 Addumantis and analysis and Formula (A.3) has been revised from " $R = r \left(\frac{m_A}{m_B}\right) \beta$ " to " $R = r \left(\frac{m_A}{m_B}\right)^{\beta}$ ";

This document is a previous general ded by tills

Water quality — Application of inductively coupled plasma mass spectrometry (ICP-MS) —

Part 2:

Determination of selected elements including uranium isotopes

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.

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

1 Scope

This document specifies a method for the determination of the elements aluminium, antimony, arsenic, barium, beryllium, bismuth, boron, cadmium, caesium, calcium, cerium, chromium, cobalt, copper, dysprosium, erbium, gadolinium, gallium, germanium, gold, hafnium, holmium, indium, iridium, iron, lanthanum, lead, lithium, lutetium, magnesium, manganese, mercury, molybdenum, neodymium, nickel, palladium, phosphorus, platinum, potassium, praseodymium, rubidium, rhenium, rhodium, ruthenium, samarium, scandium, selenium, silver, sodium, strontium, terbium, tellurium, thorium, thallium, tin, titanium, tungsten, uranium and its isotopes, vanadium, yttrium, ytterbium, zinc and zirconium in water (e.g. drinking water, surface water, ground water, waste water and eluates).

Taking into account the specific and additionally occurring interferences, these elements can be determined in water and digests of water and sludge (e.g. digests of water as described in ISO 15587-1 or ISO 15587-2).

The working range depends on the matrix and the interferences encountered. In drinking water and relatively unpolluted waters, the limit of quantification ($L_{\rm OQ}$) lies between 0,002 µg/l and 1,0 µg/l for most elements (see <u>Table 1</u>). The working range typically covers concentrations between several ng/l and mg/l depending on the element and specified requirements.

The quantification limits of most elements are affected by blank contamination and depend predominantly on the laboratory air-handling facilities available on the purity of reagents and the cleanliness of glassware.

The lower limit of quantification is higher in cases where the determination suffers from interferences (see Clause 5) or memory effects (see ISO 17294-1).

Elements other than those mentioned in the scope can also be determined according to this document provided that the user of the document is able to validate the method appropriately (e.g. interferences, sensitivity, repeatability, recovery).

Table 1 — Lower limits of quantification for unpolluted water

Element	Isotope often used	L_{OQ}^{a}	Element	Isotope often used	L_{0Q}^{a}	Element	Isotope often used	$L_{\mathrm{OQ}}^{\mathbf{a}}$
		μg/l			μg/l			μg/l
	¹⁰⁷ Ag	0,5	Hf	¹⁷⁸ Hf	0,1	Ru	¹⁰² Ru	0,1
Ag	¹⁰⁹ Ag	0,5	Нg	²⁰² Hg	0,05	Sb	¹²¹ Sb	0,2
				²⁰¹ Hg	0,1			
Al	²⁷ Al	1	Но	¹⁶⁵ Ho	0,1		¹²³ Sb	0,2
As	⁷⁵ As ^c	0,1	In	¹¹⁵ In	0,1	Sc	⁴⁵ Sc	5
Au	¹⁹⁷ Au	0,5	Ir	¹⁹³ Ir	0,1		⁷⁷ Se ^c	1
	10B	1	К	³⁹ KC	5	Se	⁷⁸ Se ^c	0,1
В	11B	1	La	¹³⁹ La	0,1		⁸² Se	1
	¹³⁷ Ba	3		⁶ Li	10	Sm	¹⁴⁷ Sm	0,1
Ва	¹³⁸ Ba	0,5	Li	⁷ Li	1		¹¹⁸ Sn	1
Ве	⁹ Be	0,1	Lu	¹⁷⁵ Lu	0,1	Sn	¹²⁰ Sn	1
Bi	²⁰⁹ Bi	0,5		²⁴ Mg	1	Sr	⁸⁶ Sr	0,5
	⁴³ Ca	100	Mg	²⁵ Mg	10		⁸⁸ Sr	0,3
Ca	⁴⁴ Ca	50	Mn	⁵⁵ Mn	0,1	Tb	¹⁵⁹ Tb	0,1
	⁴⁰ Ca	10	Ω	⁹⁵ Mo	0,5	Te	¹²⁶ Te	2
	¹¹¹ Cd	0,1	Mo	⁹⁸ Mo	0,3	Th	²³² Th	0,1
Cd	¹¹⁴ Cd	0,5	Na	²³ Na	10	m1	²⁰³ Tl	0,2
	¹⁴⁰ Ce	0,1		5	0,1	Tl	²⁰⁵ Tl	0,1
_			Nd	116		Ti	⁴⁷ Ti	10
Ce				¹⁴⁶ Nd			⁴⁸ Ti	1
							⁴⁹ Ti	10
Со	⁵⁹ Co	0,2		⁵⁸ Nic	0,1	Tm	¹⁶⁹ Tm	0,1
-	⁵² Cr ^c	0,1	Ni	⁶⁰ Nic	0,1		238U	0,1
Cr	⁵³ Cr	5	Р	31p	5	U	235U	1,10-4
Cs	¹³³ Cs	0,1		²⁰⁶ Pb ^b	0,2		²³⁴ U	1,10-5
-	⁶³ Cu	0,1	Pb	²⁰⁷ Pb ^b	0,2	V	51 V c	0,1
Cu	⁶⁵ Cu	0,1		²⁰⁸ Pb ^b	0,1		¹⁸² W	0,3
Dy	¹⁶³ Dy	0,1	Pd	¹⁰⁸ Pd	0,5	W	¹⁸⁴ W	0,3
Er	¹⁶⁶ Er	0,1	Pr	¹⁴¹ Pr	0,1	Y	89Υ	0,1
Fe	⁵⁶ Fe ^c	5	Pt	¹⁹⁵ Pt	0,5		¹⁷² Yb	0,2
	⁶⁹ Ga	0,3	Rb	⁸⁵ Rb	0,1	Yb	¹⁷⁴ Yb	0,2
Ga	⁷¹ Ga	0,3	ъ	¹⁸⁵ Re	0,1	7/6	⁶⁴ Zn	1
C 1	¹⁵⁷ Gd	0,1	Re	¹⁸⁷ Re	0,1	Zn	⁶⁶ Zn	1
Gd	¹⁵⁸ Gd	0,1	Rh	¹⁰³ Rh	0,1		⁶⁸ Zn	1
Ge	⁷⁴ Ge	0,3	Ru	¹⁰¹ Ru	0,2	Zr	⁹⁰ Zr	0,2

 $^{^{\}rm a}$ $\,$ Depending on the instrumentation, significantly lower limits can be achieved.

2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

Lead (Pb) is reported as the sum of the signal intensities of 206 Pb, 207 Pb and 208 Pb.

These limits are achieved by the use of a collision/reaction cell.

ISO 5667-1, Water quality — Sampling — Part 1: Guidance on the design of sampling programmes and sampling techniques

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

ISO 8466-1, Water quality — Calibration and evaluation of analytical methods — Part 1: Linear calibration function

ISO 15587-1, Water quality — Digestion for the determination of selected elements in water — Part 1: Aqua regia digestion

ISO 15587-2, Water quality — Digestion for the determination of selected elements in water — Part 2: Nitric acid digestion

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

ISO 17294-1:2004, Water quality — Application of inductively coupled plasma mass spectrometry (ICP-MS) — Part 1: General guidelines

3 Terms, definitions and symbols

3.1 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 17294-1 and the following apply.

ISO and IEC maintain terminology databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at https://www.iso.org/obp
- IEC Electropedia: available at https://www.electropedia.org/

3.1.1

analyte

element to be determined

3.1.2

background

 N_0

counts for a given mass in the blank solution

Note 1 to entry: Background is expressed in Counts.

3.1.3

blank calibration solution

solution prepared in the same way as the *calibration solution* (3.1.4) but leaving out the *analyte* (3.1.1)

3.1.4

calibration solution

solution used to calibrate the instrument, prepared from a $stock\ solution(s)\ (3.1.24)$ or from a certified standard

3.1.5

determination

entire process from preparing the *test sample solution* (3.1.26) up to and including the measurement and calculation of the final *result* (3.1.22)

3.1.6

expanded uncertainty

U

product of the standard uncertainty, u(C), and the coverage factor, k, with k = 1, 2, ..., as follows: $U = k \cdot u(C)$

Note 1 to entry: Expanded uncertainty is expressed in the unit of the quantity ${\sf C}.$