INTERNATIONAL STANDARD



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Nickel alloys — Determination of tantalum — Inductively coupled plasma atomic emission spectrometric method

Alliages de nickel — Dosage du tantale — Méthode par spectrométrie d'émission atomique avec source à plasma induit par haute fréquence



Reference number ISO 22725:2007(E)

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Contents

Forewo	ordi	iv
1	Scope	1
2	Normative references	1
3	Principle	1
4	Reagents .	2
5	Apparatus	3
6	Sampling and sample preparation	4
7 7.1 7.2 7.3 7.4 7.5 7.6	Procedure	4 4 4 4 5 5 6
8 8.1 8.2 8.3	Expression of results	6 6 6 7
9	Test report	7
Annex	A (normative) Checking the performance of an ICP instrument	8
Annex	B (informative) Notes on the test programme	0

Foreword

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Nickel alloys — Determination of tantalum — Inductively coupled plasma atomic emission spectrometric method

1 Scope 🥒

This International Standard specifies an inductively coupled plasma atomic emission spectrometric method for the determination of the mass fraction of tantalum between 0,1 % and 5 % in nickel alloys.

2 Normative reference

The following referenced documents are indispensable for the application 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.

ISO 648:—¹⁾, Laboratory glassware — Single volume pipettes

ISO 1042:1998, Laboratory glassware — Se-mark volumetric flasks

ISO 3696:1987, Water for analytical laboratory de — Specification and test methods

ISO 5725-1:1994, Accuracy (trueness and precision) of measurement methods and results — Part 1: General principles and definitions

ISO 5725-2:1994, Accuracy (trueness and precision) of measurement methods and results — Part 2: Basic method for the determination of repeatability and reproduciblity of a standard measurement method

ISO 5725-3:1994, Accuracy (trueness and precision) of measurement methods and results — Part 3: Intermediate measures of the precision of a standard measurement method

3 Principle

Dissolution of a test portion in a mixture of hydrofluoric, hydrochloric, nitro and phosphoric acid and fuming after addition of perchloric acid. Addition of hydrofluoric acid and, if desired of an internal reference element and dilution of the solution to known volume. Nebulization of the solution into an inductively coupled plasma atomic emission spectrometer and measurement of the intensity of the emitter light from tantalum, and eventually from the internal reference element, simultaneously.

An example of the analytical line for tantalum is given in Table 1.

The method uses a calibration based on a very close matrix-matching of the calibration solutions to the sample and bracketing of the mass fractions between 0,75 and 1,25 of the approximate concentration of tantalum in the sample to be analysed. The concentration of all elements in the sample has, therefore, to be approximately known. If the concentrations are not known, the sample has to be analysed by some semiquantitative method. The advantage of this procedure is that all possible interferences from the matrix will be automatically compensated, which will result in high accuracy. This is most important for spectral interferences,

¹⁾ To be published. (Revision of ISO 648:1977)