

# CERTIFICATE OF ANALYSIS

## CERTIFIED REFERENCE MATERIAL

**Magnesium (Mg 2+) 1000 mg/l in H2O for IC**

Lot N: 1116948  
Barcode: 83171987

Ref N: 12922145

Certification Date: 14.05.2025

Component	Certified Value and uncertainty [mg/l]	Metrological traceability
Mg <sup>2+</sup>	1001.3 ± 2.8 <sup>(y)</sup>	NIST SRM No 3131a Lot 140110; NIST SRM No 3168a Lot 120629

### Notes:

(y) WQP 5.15.1.24 The certified value was obtained by a weighted mean of the results of two independent testing methods among: Classical Volumetric, Primary Gravimetric, Instrumental (ICP, ICP/MS or IC)

Density\* 1.001 g/cm<sup>3</sup> at 20°C

Starting Material, Purity*	Batch
MgCl <sub>2</sub> 99.995%	82089887

\* These values are not certified

Storage Conditions: Store under normal laboratory conditions, at temperatures between 15°C to 25°C

Expiry date: 14.06.2028

### Concept of Certification and traceability statement:

This certified reference material (CRM) is produced using a high purity starting material, acid from sub-boiling and 18 MOhm deionised water and filtered through a 0.2 micron filter.

The reported expanded uncertainty of measurement is stated as the standard uncertainty of measurement multiplied by the coverage factor k = 2, which for a normal distribution corresponds to a coverage probability of approximately 95%. The standard uncertainty of measurement has been determined in accordance with EA 4/02

Property of the result of a measurement whereby it can be related to stated references, usually national or international standards, through an unbroken chain of comparisons all having stated uncertainties (ISO VIM)

The metrological traceability is using certified reference material traceable to SI of NIST (SRM) or BAM (CRM). All contributions in relation to the certification of standard solutions are considered when evaluating the uncertainty.

The measurement results are traceable to SI. All analytical balances used for the preparation of the solution are calibrated yearly under an in-house procedure with analytical weights, traceable to DKD, and are checked daily.

Class A laboratory glassware is used.

The results from temperature measurement are traceable to SI. The thermometers used for solution's calibration are calibrated from an ISO 17025 accredited laboratory. The ambient conditions are controlled with a hygrometer calibrated from an ISO 17025 accredited laboratory.

### Intended use: For Laboratory Use Only

Calibration of Ion Chromatographs  
Preparation of "working reference samples"

This statement is not intended to restrict the use for other purposes.

Validation of analytical methods  
Detection limit and linearity studies

### Instructions for the correct use of this reference material:

This certified reference material can be used directly or can be diluted in an appropriate high purity matrix. Only a clean class A glassware should be used. Do not pipet from container. Obtained concentration (in mg/l) after dilution is a result from the multiplication of certified value of CRM concentration and the CRM's volume used for dilution and divided into the flask's volume used for dilution.

### Stability and storage:

This CRM is with a guaranteed stability until ±0.5% of the certified concentration within its shelf life. Stability is guaranteed, provided that the solution is kept in its original packaging, tightly closed stored, as written in the section: Storage Conditions. If storage of a partially used bottle is necessary, the cap should be tightly sealed and the bottle should be stored in refrigerator to minimize transpiration rate. The laboratory performs stability tests according to MQP 5.14.1 therefore solutions with one and the same bar-code number might have different expiration dates.

#### Hazardous situation:

The normal laboratory safety precautions should be observed when working with this CRM. Further details for the handling of this CRM are available as safety data sheet.

#### Level of homogeneity:

The material was tested for homogeneity by analyzing randomly selected samples according to an in-house procedure. The level of homogeneity proved satisfactory for a sample volume of 20 ml. The uncertainty incorporates the sample standard deviation combined with the uncertainty calculated from homogeneity and stability studies.

To ensure sufficient homogeneity of the sample prior to use thoroughly mix by inversion.

#### Names of certifying officers:

Laboratory:

*Stoyanov*

Tihomir Stoyanov

Manager:

*Taralova*

Krassimira Taralova

*This document QF 5.17.1/1 version 1 is designed and the certified value(s) and uncertainty(ies) are determined in accordance with ISO Guide 31, ISO Guide 35, and Eurachem / CITAC Guides*

*This certificate relates solely to the lot number given above.*

*All processes (including generating of this certificate) are completely controlled by the specialized Computer-Aided-Manufacturing (CAM) software.*

*This Certified Reference Material was produced under a quality management system that is:*

- Registered to ISO 9001 Quality Management System (Lloyd's Register Quality Assurance Ltd Cert No 0039638)
- Accredited according to ISO/IEC 17025
- Accredited according to ISO 17034

#### Trace impurities in the actual solution reported in ppm:

(all values below are nominal and not certified)

Trace impurities in the actual solution reported in ppm:	
Ca <sup>2+</sup>	0.062
K <sup>+</sup>	0.069
Li <sup>+</sup>	<0.012
Mg <sup>2+</sup>	*
Na <sup>+</sup>	0.061
NH <sub>4</sub> <sup>+</sup>	0.017

Operating Conditions Ion Chromatography:	
Column:	IonPack CS12A 4 mm
Cation Self regenerating	DIONEXCSRS 300 4 mm
Suppressor:	
Eluent Flow Rate:	1.0 ml/min
Eluent:	15 mM CH <sub>3</sub> SO <sub>3</sub> H
Sample Concentration:	10 mg/l
Sample Volume:	25 µl loop

