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Certificate of Analysis

CERTIFIED REFERENCE MATERIAL

This document is designed and the certified value(s) and uncertainty(ies) are determined in accordance with ISO Guide 31 [1], ISO Guide 35[2], EA 4/02 and Eurachem / CITAC Guides[3]

Potassium Dichromate Solution for Absorbance Control at 430 nm

Lot N: XXXXXX **Barcode:** XXXXXXXX **Certification Date:** XXXXXXXXXX

Ref N: UV2225PDB.L08

Ph Eur N: This product conforms to the specifications given in 2.2.25 of the old European Pharmacopoeia

Description: 570 - 630 mg/l Potassium dichromate R in 0.005 M sulphuric acid and blank of 0.005 M sulphuric acid for the control of absorbance at 430 nm

Certified values/Uncertainties:

Absorbance	Specific Absorbance*/ Tolerance	Wavelenght* (nm)
0.960 +/- 0.006	16.0 (15.7 to 16.1)	430.0 +/- 0.1 nm

Concentration
600.0 +/- 1.1 mg/l Potassium dichromate

Measured values are at 20 +/- 1°C
Path length: 10 +/- 0.016 nm
* *These values are not certified*

Traceability and Instrument Qualification: The metrological traceability is assured through gravimetric measurement and dissolving the certified reference material from BAM ref N 42019 and traceable to SI. Certified absorbance values are traceable to SI (cert N 72172)
The spectrophotometer is checked prior to use for:
Control of wavelengths: VWR Ref N 86130 (Holmium Oxide Solution for Wavelenghts Control acc. 2.2.25 of Ph Eur)
Control of absorbance: VWR Ref N 86124 (Set of 2x10 ml Potassium Dichromate Solution for Absorbance Control 235 - 350 nm and 6x10 ml blank acc. 2.2.25 of Ph Eur)
Limit of stray light: VWR Ref N 86128 (Potassium Chloride Solution for Stray Light Limit acc. 2.2.25 of Ph Eur)

Calibration method: CRM's calibration procedure (WQP 5.15.1/31)

Concept of Certification and traceability statement: 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, trough an unbroken chain of comparisons all having stated uncertainties (ISO VIM [5]).
All analytical balances used for the preparation of the solution are calibrated yearly under an in-house procedure with class E1 and class E2 analytical weights, traceable to SI (DKD) and are checked daily.
Class A laboratory glassware is used.
The vernier caliper used for the cuvette check is calibrated from an ISO 17025 accredited laboratory.
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.

Ph Eur N: This product conforms to the specifications given in 2.2.25 of the European Pharmacopoeia (old edition)

Expiry date: XXXXXX

Intended use: For Laboratory Use Only
Calibration/ Accuracy testing of UV/VIS spectrophotometer
Validation of analytical methods
Preparation of ?working reference samples?

LRQA CERTIFIED ISO 9001 · UKAS MANAGEMENT SYSTEMS 0001 · ilac-MRA · ANAB ACCREDITED ISO 17034 REFERENCE MATERIAL PRODUCER

CPAchem Ltd. is ISO 17034 (Cert No AR-1835)
and ISO/IEC 17025 (Cert No AT-1836) accredited by ANAB

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SAMPLE DOMINIQUE DUTSCHER SAS

Detection limit and linearity studies
These statements are not intended to restrict the use for other purposes.

Instructions for the correct use of this reference material: for direct use only
Set the spectrophotometer to measure absorbance at 430 nm. Fill a cell with Blank (refer to ampoule labelling), and autozero the spectrophotometer. Fill a second cell with this CRM 600mg/L, and record the absorbance.

Stability and storage:
This CRM is with a guaranteed stability until ?0.5% of the certified value within its shelf-life. Stability is guaranteed provided that the solution is kept in its original packaging, tightly closed under normal laboratory conditions. According to an in-house procedure the producer will monitor this CRM at appropriate intervals and the purchasers will be notified of any significant changes resulting in recertification or with withdrawal of the CRM during the state period of the validity of the certificate.

Hazardous situation:

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

Level of homogeneity:
This CRM was mixed according to an in-house procedure and is guaranteed to be homogeneous.
To ensure sufficient homogeneity of the sample prior to use thoroughly mix by inversion.

Names of certifying officers:
Laboratory: Dinko Gospodinov

Manager: Krassimira Taralova

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

[1] ISO Guide 31: Reference materials - Contents of certificates and labels
[2] ISO Guide 35: Reference materials - General and statistical principles for certification
[3] EURACHEM/CITAC Guide: Quantifying Uncertainty in Analytical Measurement
[4] EA 4/02: Expression of the Uncertainty of Measurement in Calibration
[5] ISO/IEC Guide 99: International Vocabulary of Metrology-Basic and general concepts and associated terms (VIM)
[6] ISO/IEC 17025: General requirements for the competence of testing and calibration laboratories
[7] ISO 17034: General Requirements for the Competence of Reference Material Producers

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.

Signed by: Taralova, Chemical Production Manager

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