



Certificate of Analysis

Standard Reference Material[®] 918b

Potassium Chloride

(Clinical Standard)

This Standard Reference Material (SRM) is intended for use as an analytical standard of known purity. It is intended primarily for use in the calibration and standardization of procedures for potassium (K) and chloride (Cl) determinations employed in clinical analysis and for routine critical evaluation of the daily working standards used in these procedures. This lot of potassium chloride (KCl) was prepared to ensure a material of high purity and homogeneity and has been assayed after heating at 110 °C to 120 °C. A unit of SRM 918b consists of a single glass bottle containing 30 g of the material.

Certified Values: The certified values for this SRM are listed in Table 1. The certified values for this SRM are expressed as mass fractions, w , of KCl, K, and Cl. A NIST certified value is a value for which NIST has the highest confidence in its accuracy in that all known or suspected sources of bias have been investigated or accounted for by NIST [1].

Table 1. Certified Values^(a) for SRM 918b Potassium Chloride

Quantity	Value (%)		
w_{KCl}	99.927	±	0.014
w_{K}	52.4121	±	0.0086
w_{Cl}	47.5284	±	0.0049

^(a) Each result is expressed as the certified value \pm the expanded uncertainty, U , calculated as $U = k u_c$, where u_c is the combined standard uncertainty calculated according to the ISO and NIST Guides [2]. The value of u_c is intended to represent, at the level of one standard deviation, the combined effect of inherent sources of uncertainty of the assay techniques and applicable corrections for interfering trace elements. The value of k is 1.96, which is the coverage factor corresponding to approximately 95 % confidence based on > 1000 overall effective degrees of freedom.

Expiration of Certification: The certification of this SRM is valid until **01 August 2016**, within the measurement uncertainties specified, provided the SRM is handled and stored in accordance with the instructions given in this certificate (see “Instructions for Use”). The certification is nullified if the SRM is damaged, contaminated, or modified.

The coordination of the technical measurements leading to the certification of SRM 918b was provided by K.W. Pratt of the NIST Analytical Chemistry Division.

The coulometric and gravimetric analyses were performed by K.W. Pratt and T.W. Vetter, respectively, of the NIST Analytical Chemistry Division. Trace bromine (Br) determination by X-ray fluorescence was performed in the NIST Analytical Chemistry Division by J.R. Sieber. Additional trace element analyses by glow-discharge mass spectrometry were performed by a commercial laboratory.

Statistical consultation was provided by W.F. Guthrie of the NIST Statistical Engineering Division.

The support aspects involved in the preparation of this SRM were coordinated through the NIST Measurement Services Division.

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Analytical Chemistry Division

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Reference Values: A reference value for the mass fraction of bromine is listed in Table 2. The expanded uncertainty is calculated using $k = 2$. A reference value is a best estimate of the true value provided by NIST where all known or suspected sources of bias have not been fully investigated by NIST [1].

Table 2. Reference Value for SRM 918b

Element	Mass Fraction ($\mu\text{g/g}$)	Expanded Uncertainty ($\mu\text{g/g}$)
Br	130	45

Information Values: Table 3 lists information values for the mass fractions of trace elements in SRM 918b. Information values are non-certified values that may be of interest and use to the SRM user, but insufficient information is available to provide an uncertainty associated with the value [1]. No other elements were detected at a mass fraction greater than $1 \mu\text{g/g}$.

Table 3. Information Values for SRM 918b

Element	Mass Fraction ($\mu\text{g/g}$)
Na	35
Rb	2.6
Si	1.8

Maintenance of Certification: NIST will monitor representative samples from this SRM lot over the period of its certification. If substantive changes occur that affect the certification before the expiration of certification, NIST will notify the purchaser. Registration (see attached sheet) will facilitate notification.

NOTICE AND WARNINGS TO USERS

This SRM is intended for “in vitro” diagnostic use only.

Stability and Storage: This SRM should be stored in its original bottle at room temperature. It must be tightly re-capped after use and protected from moisture.

Homogeneity: This SRM is homogeneous within the uncertainty limits for the nominal sample mass, 250 mg, used for the coulometric assays. Samples less than 250 mg are not recommended in order to avoid possible heterogeneity with smaller sample sizes.

Possible Interfering Species: It is the responsibility of the user to evaluate which species may interfere with the application of this SRM and to apply any necessary corrections that affect the given application. The following information and the values in Tables 2 and 3 may be useful in this evaluation.

The certified value for w_{KCl} is obtained from a weighted combination of the results of independent coulometric analyses, corrected for the potassium bromide (KBr), rubidium chloride (RbCl), and sodium chloride (NaCl) impurities; and gravimetric analyses, corrected for the KBr impurity.

The certified value for w_{K} is obtained from an equally-weighted combination of w_{K} obtained directly from the gravimetric analyses and the indirect w_{K} , which is calculated from the coulometric w_{KCl} and the additional K in the KBr impurity.

The certified value for w_{Cl} is obtained from the coulometric analyses, corrected for interfering bromide.

The corrections for bromide, sodium, and rubidium were obtained from the trace element determinations and the appropriate gravimetric factors [3]. A portion of the K is present in SRM 918b as KBr, and a portion of the chloride is present as NaCl and RbCl. Therefore, the sum of the certified values for w_{K} and w_{Cl} does not equal the certified value for w_{KCl} .

INSTRUCTIONS FOR USE

Drying Instructions: Dry the material at 110 °C to 120 °C for 4 h. After the SRM has been dried, store it in a desiccator over anhydrous magnesium perchlorate.

Source of Material: The KCl used for this SRM was obtained from a commercial supplier. The material was examined for compliance with the specification for reagent grade KCl as specified by the American Chemical Society [4]. The material was found to meet or exceed the minimum requirements in every respect.

Assay Techniques: The coulometric assay value was obtained by automated titration [5] with coulometrically generated Ag^+ using potentiometric detection of the endpoint. The gravimetric assay value was obtained by ion-exchange separation of the K fraction and conversion to potassium sulfate (K_2SO_4), including corrections for instrumentally-determined K not collected with the K fraction and for trace contaminants in the K_2SO_4 (procedure based on reference 6).

REFERENCES

- [1] May, W.; Parris, R.; Beck, C.; Fassett, J.; Greenberg, R.; Guenther, F.; Kramer, G.; Wise, S.; Gills, T.; Colbert, J.; Gettings, R.; MacDonald, B.; *Definitions of Terms and Modes Used at NIST for Value-Assignment of Reference Materials for Chemical Measurements* NIST Special Publication 260-136, U.S. Government Printing Office: Washington, DC (2000); available at http://www.cstl.nist.gov/nist839/special_pubs/SP260136.pdf.
- [2] ISO; *Guide to the Expression of Uncertainty in Measurement*; ISBN 92-67-10188-9, 1st ed.; International Organization for Standardization: Geneva, Switzerland (1993); see also Taylor, B.N.; Kuyatt, C.E.; *Guidelines for Evaluating and Expressing the Uncertainty of NIST Measurement Results*; NIST Technical Note 1297, U.S. Government Printing Office: Washington, DC (1994); available at <http://physics.nist.gov/Pubs>.
- [3] *IUPAC Commission of Atomic Weights and Isotopic Abundances*; Pure & Appl. Chem., Vol. 75 (8), pp. 1107–1122 (2003).
- [4] *Reagent Chemicals*; 9th ed., American Chemical Society: Washington, DC (1999).
- [5] Pratt, K.W.; *Automated, High-Precision Coulometric Titrimetry Part II. Strong and Weak Acids and Bases*; Anal. Chim. Acta, Vol. 289 (2), 135–142 (1994).
- [6] Moody, J.R.; Vetter, T.W.; *Development of the Ion Exchange-Gravimetric Method for Sodium in Serum as a Definitive Method*; J. Res. Natl. Inst. Stand. Technol., Vol. 101, pp. 155–164 (1996); available at <http://nvl.nist.gov/pub/nistpubs/jres/101/2/j2mood.pdf>.

Users of this SRM should ensure that the certificate in their possession is current. This can be accomplished by contacting the SRM Program at: telephone (301) 975-6776; fax (301) 926-4751; e-mail srminfo@nist.gov; or via the Internet at <http://www.nist.gov/srm>