

National Bureau of Standards

Certificate of Analysis

Standard Reference Material 1568a

Rice Flour

This Standard Reference Material (SRM) is intended primarily for calibrating instruments and evaluating the reliability of analytical methods for the determination of minor and trace elements in rice flour and similar agricultural food products.

Certified Concentrations of Constituent Elements: The certified concentrations of the constituent elements are shown in Table 1. Except for sulfur, the concentrations are based on results obtained by two or more independent, reliable analytical methods. Sulfur is certified based on its determination by a definitive method, isotope dilution thermal ionization mass spectrometry. Non-certified values, which are given for information only, appear in Table 2. All values are based on a minimum sample size of 500 mg and are reported on a "dry weight" basis. (See "Instructions for Drying").

Notice and Warnings to Users:

Expiration of Certification: This certification will be invalid after 5 years from the date of shipping. Should it be invalidated before then, purchasers will be notified by NBS.

Storage: The material should be kept in its original bottle and stored at temperatures between 10-30 °C. It should not be exposed to intense sources of radiation, including ultraviolet lamps or sunlight. Ideally, the bottle should be kept in a desiccator, in the dark at the temperature indicated.

Use: The following procedures should be followed to relate the analytical determinations to the values reported in this certificate. The bottle should be shaken well before each use, and a minimum sample of 500 mg of the material should be used. Mercury should be determined without drying and the concentration values adjusted for the moisture content of the material using separate samples. Other elements may be determined either on samples without drying as indicated above or on samples vacuum-dried for 24 hours as indicated under "Instructions for Drying."

Coordination of some technical measurements leading to this certificate was performed by M.S. Epstein of the Inorganic Analytical Research Division.

Statistical analysis of the experimental data was performed by K.R. Eberhardt of the NBS Statistical Engineering Division.

The technical and support aspects involved in the preparation, certification, and issuance of this Standard Reference Material were coordinated through the Office of Standard Reference Materials by R. Alvarez.

Gaithersburg, MD 20899
January 20, 1988

Stanley D. Rasberry, Chief
Office of Standard Reference Materials

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Table 1. Certified Concentrations of Constituent Elements^a

Minor Elements

<u>Element^b</u>	<u>Concentration, Percent by Weight^c</u>
Calcium ^{1a,2b}	0.0118 ± 0.0006
Magnesium ^{1a,2a,4a}	0.056 ± 0.002
Phosphorus ^{2a,2c,5}	0.153 ± 0.008
Potassium ^{2b,4a}	0.1280 ± 0.0008
Sulfur ^{3b}	0.120 ± 0.002

Trace Elements

<u>Element^b</u>	<u>Concentration, µg/g^c</u>
Aluminum ^{2a,4a}	4.4 ± 1.0
Arsenic ^{1c,4a}	0.29 ± 0.03
Cadmium ^{1b,4b}	0.022 ± 0.002
Copper ^{1a,4a,4b}	2.4 ± 0.3
Iron ^{1a,3b,4a}	7.4 ± 0.9
Manganese ^{1a,4a}	20.0 ± 1.6
Mercury ^{1d,4b}	0.0058 ± 0.0005
Molybdenum ^{2c,3a,4a}	1.46 ± 0.08
Rubidium ^{2b,4a}	6.14 ± 0.09
Selenium ^{1c,4a}	0.38 ± 0.04
Sodium ^{2b,4a}	6.6 ± 0.8
Zinc ^{1a,4a}	19.4 ± 0.5

^a Analytical values are based on the "dry-weight" of material (see "Instructions for Drying"). Mercury should be determined on samples without drying and the results adjusted to a "dry-weight" basis by determining moisture on separate samples.

^b Number and letter code, as superscripts, indicate methods used for certification. (See "Analytical Methods").

^c The certified concentration is the weighted mean computed according to the procedure described by R.C. Paule and J. Mandel (NBS Journal of Research, 87, 1982, pp. 377-385). The uncertainty is stated as a 95% confidence interval plus an additional allowance for systematic error among the methods used. The allowance for systematic error is the greatest difference between the weighted mean and the component means for the analytical methods used. For manganese, an additional allowance for material inhomogeneity is included, so that the uncertainty represents a 95% expected coverage statistical tolerance interval.

Analytical Methods

1. Atomic Absorption Spectrometry
 - a. Flame
 - b. Graphite Furnace
 - c. Hydride Generation
 - d. Cold Vapor
2. Atomic Emission Spectrometry
 - a. DC Plasma
 - b. Flame
 - c. Inductively Coupled Plasma
3. Mass Spectrometry
 - a. Isotope Dilution Inductively Coupled Plasma
 - b. Isotope Dilution Thermal Ionization
4. Neutron Activation Analysis
 - a. Instrumental
 - b. Radiochemical
5. Spectrophotometry

Table 2. Non-Certified Concentrations of Constituent Elements^a

NOTE: The values shown in this table are not certified because they are not based on the results of either two or more independent reliable methods or a definitive method of known high accuracy. These values are included for information only and therefore no uncertainty limits are provided.

Trace Elements			
Element	Concentration, μg/g	Element	Concentration, μg/g
Antimony	(0.0005)	Lead	(<0.010)
Bromine	(8)	Tin	(0.0047)
Chlorine	(300)	Tungsten	(0.0012)
Cobalt	(0.018)	Uranium	(0.0003)
Iodine	(0.009)	Vanadium	(0.007)

^a Analytical values are based on the "dry-weight" of material (see "Instructions for Drying").

Preparation of Material: The rice flour for this Standard Reference Material was described by the supplier as 100% long grain from Arkansas. At NBS, the material was passed through a sieve with openings of 425 μm (No. 40) and blended. The bottled material was then subjected to 2.5 megarads of 60Co radiation for microbiological control at Neutron Products, Inc., Dickerson, MD.

Homogeneity Assessment: A preliminary evaluation of the homogeneity was made by instrumental neutron activation using samples of approximately 500 mg. The uncertainties for the concentrations in Table 1 incorporate these results.

Instructions for Drying: Except for mercury, elements should be determined on samples that have been dried as follows:

Vacuum-dry the material at approximately 25 °C for 24 hours at a pressure not greater than 70 Pa (0.5 mm Hg) with a cold trap at a temperature of about -30 °C or below.

Mercury should be determined on undried samples. However, because the certificate values are reported on a "dry-weight" basis, the elemental concentration determined on undried samples should be adjusted for the moisture content of separately measured samples. The moisture content, which was approximately 8% when bottled, should be determined on separate samples by the vacuum-drying procedure described above. Samples for analysis should not be oven-dried lest elements be lost by volatilization.

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