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Certificate of Analysis Standard Reference Material 1633b



National Institute of Standards & Technology

Certificate of Analysis

Standard Reference Material 1633b

Constituent Elements in Coal Fly Ash

This Standard Reference Material (SRM) is intended for use in the evaluation of analytical methods for the determination of constituent elements in coal fly ash or materials with a similar matrix. SRM 1633b is a bituminous coal fly ash that was sieved through a nominal sieve opening of 90 km (170 mesh) and then blended to assure homogeneity. A unit of SRM 1633b consists of 75 g of powdered material

The certified values for the constituent elements are given in Table 1. The values, except for Hg, are based on measurements using one definitive method or two or more independent and reliable analytical techniques. Noncertified values for a number of elements are given in Table 2 as additional information on the composition of the material. The noncertified values should not be used for calibration or quality control. Analytical methods used for the certification of this SRM are given in Table 3 along with analysts and cooperating laboratories. All values are based on measurements using a dry sample weight of at least 250 mg.

NOTICE AND WARNING TO USERS

Expiration of Certification: This certification is valid for 5 years from the date of shipment from NIST. Should any of the certified values change before the expiration of the certification, the purchaser will be notified by NIST.

Stability: This material is considered to be stable; however, its stability has not been rigorously assessed. NIST will monitor this material and will report any substantive changes in certification to the purchaser.

Use: A minimum dry sample weight (see Instructions for Drying) of 250 mg should be used for analytical determinations to be related to the certified values on this Certificate of Analysis.

To obtain the certified values, sample preparation procedures should be designed to affect complete dissolution. If volatile elements (e.g., Hg, As, Se) are to be determined, precautions should be taken in the dissolution of SRM 1633b to avoid volatilization losses.

Statistical consultation was provided by S.B. Schiller of the NIST Statistical Engineering Division.

The overall direction and coordination of the analyses were under the chairmanship of R.R. Greenberg of the NIST Inorganic Analytical Research Division.

The technical and support aspects involved in the preparation, certification, and issuance of this SRM were coordinated through the Standard Reference Materials Program by J.S. Kane.

Gaithersburg, MD 20899 June 22, 1993

Thomas E. Gills, Acting Chief Standard Reference Materials Program

Instructions for Drying: When non-volatile elements are being determined, this material should be dried to constant weight before using. Recommended procedures for drying are: 1) Vacuum drying for 24 h at ambient temperature using a cold trap at or below -50 °C and a pressure not greater than 0.2 mm Hg (30 Pa); 2) drying for 2 h in an oven of 105 °C. Samples of the dried material weighing at least 250 mg should be used for analysis. When not in use, the material should be kept in a tightly sealed bottle. Volatile elements should be determined on an as-received basis, and corrected to dry weight. Correction should be based on a separate determination of moisture, using one of the above drying procedures.

Source and Preparation of the Material: The fly ash was supplied by a coal fired power plant and is the product, of Pennsylvania and West Virginia coals. It was selected as a typical bituminous coal fly ash and is not intended as a fly ash from a specific coal or combustion process. The material was air dried, sieved, and blended for 24 h, before being placed in a series of bulk containers. X-ray fluorescence and industively coupled plasma atomic emission analyses were performed on ten grab samples taken from the bulk for a pretiminary homogeneity assessment before proceeding with bottling the material in 75 g units.

Analysis: The homogeneity of the bottled material was assessed by X-ray fluorescence spectrometry and instrumental neutron activation analysis, using selected elements as indicators. In some cases, statistically significant differences between samples were seen, and the variance due to material inhomogeneity is included in the overall uncertainties of the certified values. The estimated relative standard deviation for material inhomogeneity is less than 1% for those elements for which homogeneity was assessed, except Th, for which material inhomogeneity was estimated to be 2%.

Certified Values and Uncertainties: The certified values are weighted means of results of two or more independent analytical methods, or the means of results from a single definitive method, except for mercury. Mercury certification is based on cold vapor atomic absorption spectrometry measurements performed at NIST. The weights for the weighted means were computed according to the iterative procedure of Paule and Mandel (NBS Journai of Research 87, 1982, pp. 377-385). The stated uncertainty includes allowances for measurement imprecision, material variability, and differences among analytical methods. Each uncertainty is the sum of the half-width of a 95% prediction interval, and includes an allowance for the systematic error among the methods used. In the absence of systematic error, a 95% prediction interval predicts where the true concentrations of 95% of the samples of this SRM lie.

Element	wt %		Element		ing kg
Aluminum Calcium Iron Magnesium Potassium Silicon Sodium Sulfur Titanium	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	0.06 0.23 0.008 0.03 0.08 0.003 0.0011 0.014	Arsenic Barium Cadmium Chromium Copper Lead Manganese Mercury Vickel Gelenium Crontium Chorium Vanadium	136.2 709 0.784 198.2 112.8 68.2 131.8 0.141 120.6 10.26 1041 25.7	= 26 = 27 = 0006 = 4.7 = 2.6 = 1.1 = 1.7 = 0.019 1.8 1.7 = 1.8 1.7 = 1.8 1.7 = 3.6 = 3.6

Table 2. Noncertified Values

Element	mg/kg	Element	mg/kg
Antimony	6	Phosphorus	2300
Bromine	2.9	Rubidium	140
Cerium	190	Scandium	41
Cobalt	50	Samarium	20
Cesium .	11	Tantalum	1.8
Dysprosium	(V)	. Terbium	2.6
Europium 🔘 📉	4 \i\	Thallium	3.9
Gadolinum	13	Thulium / \ \ \	2.1
Hafnium \	6.8	Tungsten	5.6
Holmium \	3.5	Ytterbium	7.6
Lanthanum	94	Zinc	210
Lutetium	1.2		
Neodymium	85		

Table 3. Analytical Methods Used for Certification Analyses of SRM 1633b

EI	ement	Certification Methods
	Al	INAA, XRF
	.A.ş	FIA-HAAS, INAA
	Ba	ICP-MS, INAA
	Ca	
	Cd	ICP, INAA, XRF
	Cr '	ETAAS, IDTIMS
	Cu	FAAS, INAA
	Cu Fe	FAAS, ICP-MS
		INAA, XRF
	Hg	CVAAS
	K	FAES, INAA. XRF
	Mg	ICP, IDTIMS
	Мn	FAAS, INAA
	Va.	FAES INAA
	٧i	ETAAS IEP
	РЪ	ETAAS, ICP-MS
	₹b	FAES, INAA
S	5	IDTIMS
	Sb	ETAAS, INAA
	Se (FIA-HAAS, INAA
S		GRAV, XRF
> (2)	(r \	FAES, INAA, IDTIMS
/ J	(hf.)	ICP-MS, INAA
\ \ T	ì	INAA, XRF
O f	J	ICP-MS. INAA
\ \	/	ICP, INAA

ID-TIMS - Isotope dilution thermal ionization mass spectrometry, mixed acid digestion.

CP-MS - Inductively coupled plasma mass spectrometry; mixed acid digestion.

INAA - Instrumental neutron activation analysis.

XRF - Wavelength dispersive X-ray fluorescence on fused borate discs.

ICP-AES - Inductively coupled plasma atomic emission spectrometry; mixed acid digestion.

ETAAS - Electrothermal atomic absorption spectrometry; mixed acid digestion.

CVAAS - Cold vapor atomic absorption spectrometry.

FIA-HAAS - Flow injection analyses - Hydride generation atomic absorption spectrometry.

FAAS Flame atomic absorption spectrometry; mixed acid digestion except for Au, leached with HBr-Br₂.

GRAV Gravimetry; sodium carbonate fusion.

Most information values were determined by INAA only; P was determined by ICP-AES and XRF. TI was determined by ICP-MS, and Zn was determined by FAAS and ICP-AES.

Participating NIST Analysts

"Rocio Arvizu Ellyn S. Beary Diane S. Braverman Michael S. Epstein John D. Fassett Karen M. Garrity Robert R. Greenberg W. Robert Kelly Elizabeth A. Mackey John R. Moody Karen E. Murphy Paul J. Paulsen Theresa A. Rush Rajananda Saraswati Johanna M. Smeller Thomas W. Vetter Robert D. Vocke Robert L. Watters, Jr.

Participating Laboratories

JoAnne Delles, Howard Kanare Construction Technology Laboratories, Inc. Skokie, IL 60077

Paul Briggs, David Siems U. S. Geological Survey Branch of Geochemistry Lakewood, CO 80225

ภาคผนวก ข

ปริมาณฝุ่นแขวนลอยขนาดเล็กกว่า 10 ใมครอน ปริมาณฝุ่นโครเมียมแขวนลอยขนาดเล็กกว่า 10 ใมครอน ปริมาณสัมผัสฝุ่นโครเมียมแขวนลอยขนาดเล็กกว่า 10 ใมครอน ความเสี่ยงที่เกิดจากฝุ่นโครเมียมแขวนลอยขนาดเล็กกว่า 10 ใมครอน

No.	Sample no.	Date/Station	PM-10	Cr	· I	Risk	Month	Station
			$(\mu g/m^3)$	$(\mu g/m^3)$	(mg/Kg/day)			•
1	22c	5 Mar 99/B	51	6.35E-02	2.72E-03	1.11E-01	3	2
2	23c	11 Mar 99/B	58	4.31E-02	1.85E-03	7.57E-02	3	2
3	24c	17 Mar 99/B	56	3.60E-02	1.54E-03	6.33E-02	3	2
4	Ic	1 May 99/K	18	3.88E-02	1.66E-03	6.82E-02	5	1
5	2c	7 May 99/K	33	2.44E-02	1.04E-03	4.28E-02	5	1
6	3c	18 May 99/K	42	3.85E-02	1.65E-03	6.76E-02	5	1
7	4c	22 May 99/K	25	6.19E-02	2.65E-03	1,09E-01	5	1
8	5c	28 May 99/K	46	2.94E-02	1.26E-03	5.17E-02	5	1
9	25c	7 May 99/B	29	3.87E-02	1.66E-03	6.81E-02	5	2
10	26c	18 May 99/B	38	4.45E-02	1.91E-03	7.83E-02	5	2
11	27c	22 May 99/B	23	7.26E-02	3.11E-03	1.28E-01	5	2
12	28e	28 May 99/B	30	6.42E-02	2.75E-03	1.13E-01	5	2
13	6¢/	3 Jun 99/K	22	4.97E-02	2.13E-03	8.73E-02	6	1
	70	9 Jun 99/K	36	4.53E-02	1,94E-03	7.96E-02	6	1
15	8c	15 Jun 99/K	34	8.51E-02	3.65E-03	1.50E-01	6	1
16	9c	21 Jun 99/K	42	3.95E-02	1.69E-03	6.94E-02	6	1
17	10c	27 Jun 99/K	38/	2.43E-02	1.04E-03	4.26E-02	6	1
18	29c	3 Jun-99/B	25	6.10E-02	2.61E-03	1.07E-01	6	2
19	30c	9 Jun 99/B	30	5.17E-02	2.21E-03	9.08E-02	6	2
20	31c	15/Jun 99/B	36	5.55E-02	2.38E-03	9.75E-02	6	2
21	32c	21 Jun 99/B	44	4.69E-02	2.01E-03	8.24E-02	6	2
22	33c	27 Jun 99/B	30	3.42E-02	1.47E-03	6.01E-02	6	2
23	llc	2 Jul 99/K	44	4.34E-02	1.86E-03	7.63E-02	. 7	1
24	12c	10 Jul 99/K	28	4.52E-02	1.94E-03	7.95E-02	7	1
25	·13c	26 Jul 99/K	61	5.27E-02	2.26E-03	9.26E-02	7	1
26	34c	2 Jul 99/B	41	5.75E-02	2.46E-03	1.01E-01	7	2
27	35c	10 Jul 99/B	27	4.85E-02	2.08E-03	8.52E-02	7	2
28	36c	26 Jul 99/B	52	2.88E-02	1.23E-03	5.05E-02	7	2
29	14c	i Aug 99/K	31	6.01E-02	2.57E-03	1.06E-01	8	1
30	15c	14 Aug 99/K	29	1.63E-02	6.98E-04	2.86E-02	- 8	1

No.	Sample no.	Date/Station	PM-10	Cr	I	Risk	Month	Station
			$(\mu g/m^3)$	$(\mu g/m^3)$	(mg/Kg/day)			
31	16c	26 Aug 99/K	38	4.70E-02	2.01E-03	8.26E-02	8	1
32	38c	8 Aug 99/B	44	4.06E-02	1.74E-03	7.14E-02	8	2
33	39c	14 Aug 99/B	40	2.12E-02	9.07E-04	3.72E-02	8	2
34	41c	25 Aug 99/B	38	3.37E-02	1.44E-03	5.92E-02	8	2
35	17c	1 Sep 99/K	25	4.77E-02	2.05E-03	8.39E-02	9	1
36	18c	7 Sep 99/K	36	6.51E-02	2.79E-03	1.14E-01	9	1
37	19c	13 Sep 99/K	52	6.01E-02	2.58E-03	1.06E-01	9	1
38	42c	1 Sep 99/B	27	6.52E-02	2:80E-03	1.15E-01	9	2
39	43c	7 Sep 99/B	29	5.92E-02	2.54E-03	1.04E-01	9	2
40	44c	13 Sep 99/B.	43	6.26E-02	2.68E-03	1,10E-01	9-	2
41	45c	25 Oct 99/B	(215)	5.88E-02	2.52E-03	1.03E-01	lø	2
42	20c	27 Nov 99/K	82	6.34E-02	2.72E-03	1.11E-01	11	$\mathcal{D}_{\mathbf{l}}$
43	46c	6 Nov 99/B	30	5.24E-02	2.24E-03	9.20E-02	11	2
34	47c	12 Nov 99/B	40	6.23E-02	2,67E-03	1.09E-01	11	2
45	48c	25 Nov 99/B	33	4.98E-02	2,13E-03	8.75E-02	11	2
46	49c	27 Nov 99/B	45	3.38E-02	1.45E-03	5.95E-02	11	2
47	50c	30 Nov 99/B	37/	3.69E-02	1.58E-03	6.49E-02	11	2
48	21c	12 Dec/99/K	189	6.41E-02	2.75E-03	1.13E-01	12	1
49	51c	12 Dec 99/B	68	1.16E-01	4.97E-03	2.04E-01	12	2
50	52¢	24 Dec 99/B	73	5.83E-02	2.50E-03	1.02E-01	12	2

i : Khao Din School

2 : Bang Phuang Health Center