

U. S. DEPARTMENT OF COMMERCE

National Bureau of Standards

Certificate of Analyses

OF

STANDARD SAMPLE 63B

PHOSPHOR BRONZE BEARING METAL

ANALYST*	COPPER Electrolytic	TIN	LEAD Weighed as PbSO ₄	ZINC ZnS-ZnO	ANTIMONY	IRON	PHOSPHORUS Weighed as Mg ₂ P ₂ O ₇	NICKEL Weighed as nickel dimethylglyoxime	SULFUR	SILICON	ALUMINUM	SILVER Weighed as AgCl	ARSENIC
1	^a 77.90	^b 9.78	^c 9.35	0.71	^d 0.55	^e 0.47	0.44	^f 0.33	{ ^g 0.17 ^h 0.16}	ⁱ 0.13	^j 0.05	0.04	^k 0.014
2	77.95	^l 9.79	9.35	.71	^m .55	ⁿ .48	^o .44	.33		^b .17	^p .13	^q .05	.04
3	77.92	^s 9.76	^a 9.40	^t .71	^u .54	^v .47	.44	.33	^h .16	^e .11	^w .05	.04	^x .014
4	77.96	^y 9.79	{ ^{9.34} ^{9.36} }	^{z1} .72	^{z2} .52	^{z3} .45	.44	.33	^{z4} .16	.13	-----	.05	^{z5} .016
5	77.96	¹ 9.78	9.33	.70	^m .53	^e .48	⁶ .44	.33	⁴ .16	ⁱ .12	⁷ .06	⁸ .04	⁹ .016
Averages	77.94	9.78	9.36	0.71	0.54	0.47	0.44	0.33	0.16	0.12	0.05	0.04	0.015

* Five-gram sample dissolved with 55 ml of HNO₃ (1:1). Solution digested on a steam bath overnight, filtered, and the precipitate washed with hot HNO₃ (1:99). Filtrate diluted to 350 ml, 2 drops of 0.1 N HCl added, and solution electrolyzed by the use of a current density of 0.5 amp/dm². Metastannic acid precipitate and paper treated with HNO₃-H₂SO₄. Tin, antimony, and arsenic volatilized by HBr-Br₂. Residual solution combined with the first electrolyte and lead separated as PbSO₄. Residual copper in filtrate from PbSO₄ precipitated with H₂S and determined by electrolysis. Correction made for silver in cathode deposit.

^b Tin separated by distillation from a 1-g sample, precipitated with cupferron, and ignited to SnO₂. (See J. Research NBS 33, 307 (1944) RP1810).

^c First anode deposit (footnote a) dissolved in nitric acid and a little alcohol. Solution combined with the first electrolyte and lead determined as PbSO₄.

^d Antimony separated by distillation from a 5-g sample, precipitated with H₂S, and titrated with KMnO₄ as described in J. Research NBS 21, 95 (1938) RP1116. KMnO₄ standardized on high-purity antimony.

^e SnCl₂-K₂Cr₂O₇ method.

^f Dimethylglyoxime-photometric method.

^g Weighed as BaSO₄.

^h Combustion-iodate method.

ⁱ Tin, antimony, and arsenic volatilized by HBr-HClO₄ treatment. Double dehydration with HClO₄ with intervening filtration.

^j Weighed as AlPO₄.

^k Molybdenum-blue photometric method. See J. Research NBS 24, 7 (1940) RP1267.

^l Tin reduced with nickel and titrated with KIO₃.

^m Metastannic acid precipitate separated and antimony titrated with KBrO₃.

ⁿ Alkali-molybdate method.

^o Sulfuric acid dehydration.

^p Mercury cathode-phenylhydrazine method.

^q Distillation-As₂S₃ method.

^r Tin reduced with iron in the presence of added antimony and titrated with KBrO₃.

^s Same value obtained by electrodeposition and weighing as PbO₂.

^t Same value obtained by 8-hydroxyquinoline-KBrO₃ method.

^u Metastannic acid precipitate separated and digested in H₂SO₄, (NH₄)₂S₂O₈. Tartaric acid solution added and solution treated with H₂S. Sulfides separated and dissolved in HNO₃-H₂SO₄. Antimony reduced with

tartaric acid, and titrated with KMnO₄ standardized on Na₂C₂O₄.

^v Iron reduced with granular zinc. Precipitated copper and tin removed, and FeCl₃ titrated with Ce(SO₄)₂.

^w Mercury cathode-8-hydroxyquinoline-KBrO₃ method.

^x Arsenic distilled from a 10-g sample and titrated with iodine.

^y Tin reduced with iron and titrated with iodine.

^z PbCrO₄ method.

¹ K₄Fe(CN)₆ method.

² Metastannic acid precipitate separated, and digested in H₂SO₄. Antimony reduced and titrated with KMnO₄. Same value obtained by pyridine-iodide-colorimetric method.

³ SnCl₂-KMnO₄ method.

⁴ HBr evolution method.

⁵ Spectrographic determination.

⁶ Molybdivanadophosphoric-photometric method. See ASTM method E-82.

⁷ Mercury cathode-aluminum oxyquinolate.

⁸ Internal electrolysis. See ASTM method E-37.

⁹ Arsenic distilled from a 5-g sample and titrated with KBrO₃.

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