

UNITED STATES DEPARTMENT OF COMMERCE
WASHINGTON

National Bureau of Standards
Certificate of Analyses
Standard Sample 63c
Phosphor Bronze Bearing Metal

ANALYST	COPPER Electrolytic	LEAD Weighed as PbSO ₄	TIN SnCl ₂ -KIO ₃	ANTIMONY	NICKEL Weighed as nickel dimethylglyoxime	PHOSPHORUS Weighed as Mg ₂ P ₂ O ₇	ZINC ZnS-ZnO	SULFUR	ARSENIC	IRON
1.....	^a 80.46	^b 9.35	^c 9.03	^d 0.50	0.32	0.145	0.090	^e 0.059	^f 0.025	^g 0.0010
2.....	80.43	9.33	^h 9.04	ⁱ .54	.32	{ .148 i.15 }	.088	^k .056	^l .017	^m .002
3.....	80.49	9.34	ⁿ 9.06	^o .53	.31	i.150	.084	^p .065	^q .029	.0015
4.....	^p 80.52	{ 9.38 ^q 9.36 }	^r 8.96	^s .52	^t .31	.144	.096	^k .056	^v .022	^w .0015
5.....	80.48	9.32	^x 9.05	^y .52	.32	i.138	.098	^u .062	^z .022	^{aa} .0012
6.....	80.47	9.35	^{ab} 9.04	.50	^{ac} .31	i.143	^{ad} .102	^k .063	^l .021	^{ae} .0009
Range.....	80.48	9.35	9.03	0.52	0.32	0.145	0.093	0.060	0.023	0.0013

Five-gram sample dissolved in 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 overnight, using a current density of 0.5 amp/dm². Metastannic-acid precipitate treated with HNO₃-HClO₄-HBr and the residual solution combined with the first electrolyte. Residual copper in the filtrate from PbSO₄ (footnote b) precipitated with H₂S and determined by electrolysis.

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

^c Two-gram sample dissolved in HCl-HNO₃, 15 mg of ingot iron as FeCl₃ added, and tin precipitated twice with NH₄OH. Precipitate dissolved in HCl, tin reduced with nickel and titrated with KIO₃ standardized with pure tin.

^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 with NBS standard sodium oxalate.

^e Combustion-iodate method. Determination made by R. E. McIntyre.

^f Distillation-As₂S₃ method.

^g Orthophenanthroline-photometric method.

^h Tin reduced with nickel in the presence of antimony and titrated with KIO₃.

ⁱ Metastannic-acid precipitate separated and treated with Na₂SO₄ and H₂SO₄. Antimony reduced with Na₂SO₃ and titrated with KBrO₃. Correction made for arsenic.

^j Molybdivanadophosphoric acid-photometric method. See ASTM method E62-50T.

^k Combustion-iodate method.

^l Molybdenum-blue photometric method.

^m Tin reduced with iron in the presence of antimony and titrated with iodine standardized with tin and NBS standard 127a.

ⁿ Antimony titrated with KBrO₃ after separation of arsenic by distillation.

^o HBr evolution, weighed as BaSO₄.

^p Copper deposited in the presence of tin in an HNO₃-HF solution of a 5-g sample.

^q Metastannic-acid precipitate from a 1-gram sample separated and lead determined electrolytically as PbO₂.

^r Tin reduced with iron in the presence of antimony and titrated with KIO₃.

^s Metastannic-acid precipitate separated and digested in H₂SO₄. Arsenic separated as AsCl₃ and antimony in residual solution titrated with KBrO₃.

^t Dimethylglyoxime precipitate titrated with cyanide.

^u Same value obtained by the dimethylglyoxime-photometric method.

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

^w Thioglycolic acid-photometric method.

^x Tin reduced with aluminum and titrated with KIO₃.

^y Iodide-photometric method. See Anal. Chem. 19, 353 (1947).

^z Solution in nitric acid. Sulfur weighed as BaSO₄.

^{aa} Distillation-iodimetric method.

^{ab} NH₄CNS-photometric method.

^{ac} Tin reduced with aluminum and titrated with iodine.

^{ad} Dimethylglyoxime-photometric method.

^{ae} Zinc separated from the copper electrolyte with H₂S, and determined by electrolysis.

List of Analysts

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| 1. Nonferrous Laboratory, National Bureau of Standards.
R. K. Bell, in charge. Analysis by E. E. Maczkowski, B. B. Bendigo, and L. A. Machlan. | 4. A. B. Shapiro, H. Kramer and Co., Chicago, Ill. |
| 2. W. A. Eddie and J. T. Krantz, National Bearing Division, American Brake Shoe Co., St. Louis, Mo. | 5. J. D. Kopp, Scovill Manufacturing Co., Waterbury, Conn. |
| 3. B. A. Stoltz, Ajax Metal Division, H. Kramer and Co., Philadelphia, Pa. | 6. C. E. Potts, F. deF. Camp, Kathleen M. O'Brien, O. P. Case, and E. M. Horton, The American Brass Co., Waterbury, Conn. |

The metal for the preparation of this standard was furnished by National Bearing Division of American Brake Shoe Co.

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A. V. ASTIN, Director.