

UNITED STATES DEPARTMENT OF COMMERCE
WASHINGTON

National Bureau of Standards
Certificate of Analyses
Standard Sample 68B
Ferromanganese

ANALYST	Mn	C	P		S		Si	
	Bismuthate (FeSO ₄ -KMnO ₄)	Combustion	Gravimetric (weighed as Mg ₂ P ₂ O ₇ after removal of arsenic)	Alkali-molybdate	Gravimetric (direct ox- idation and final precipi- tation in reduced solu- tion)	Evolution with HCl ZnS-Iodine (theoretical sulfur titer) ^a	Combustion	Sulfuric acid dehydration
1	79.96	^b 6.75	0.285			0.006	^e 0.005	^d 0.44
2	79.91	^e 6.73		^f 0.293			^e .008	.44
3	^b 80.00	ⁱ 6.77		^j .279	0.005		.004	.43
4	80.06	^k 6.77		^j .297	.005	.007		1.44
	80.00	6.77		^m .303		ⁿ .008	^o .006	.45
6	79.91	^p 6.83	.302	^f .301			^o .007	.42
7	{ 79.93 79.96}	6.76	.285	.292	.006	.006	^o .006	1.44
Average	79.97	6.77	0.291	0.294	0.005	0.007	0.006	0.44
General average	79.97	6.77	0.293		0.006			0.44

^a Value obtained by standardizing the titrating solution by means of sodium oxalate through KMnO₄ and Na₂S₂O₈, and use of the ratio 2I:1S.

^b Sample mixed with ingot iron. Determination made by Charles C. Marshall.

^c 1-g sample covered with alundum and burned in oxygen at 1475° C. SO₂ absorbed in starch-iodine solution and titrated with standard KIO₃ solution based on 93 percent of the theoretical factor. Determination made by John L. Hage.

^d Double dehydration with intervening filtration.

^e Sample mixed with CuO and electrolytic iron.

^f Titrating solution standardized by use of a standard steel.

^g Burned with tin and theoretical factor used for iodate solution.

^h Manganese dioxide precipitated in nitric acid solution. Solution filtered, and the dioxide titrated with FeSO₄ and KMnO₄ standardized by use of NBS ferromanganese, 68a.

ⁱ Sample mixed with red lead.

^j Titrating solution standardized by use of NBS ferromanganese, 68a.

^k Sample mixed with tin and ingot iron.

^l Same value obtained by HClO₄ dehydration.

^m Titrating solution standardized by use of NBS acid potassium phthalate, 84d.

ⁿ KIO₃ standardized by use of NBS Arsenic Trioxide, 83a.

^o Iodate method with copper used as an accelerator.

^p Sample mixed with tin.

^q Burned with tin and SO₂ titrated with iodate standardized by use of a standard steel.

^r Potentiometric titration with KMnO₄ in neutral pyrophosphate solution. See Ind. and Eng. Chem. Anal. Ed. 18, 191 (1946).

^s SO₂ absorbed in neutral H₂O₂ solution titrated with Na₂CO₃, using methyl red indicator.

Values for constituents not as accurately determined as the above are: copper, 0.12; chromium, 0.055; vanadium, 0.043; cobalt, 0.04; and arsenic, 0.09 percent.

List of Analysts

- R. K. Bell, National Bureau of Standards.
- J. J. Furey, Electro Metallurgical Company, Niagara Falls, N. Y.
- W. E. Steiner, Bethlehem Steel Company, Johnstown Plant, Johnstown, Pa.
- Armco Research Chemical Laboratory, Middletown, Ohio, A. H. Thomas in Charge. Analysis by M. Dannis, L. C. Ikenberry, L. Chenault, and I. Shroyer.

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