

U. S. DEPARTMENT OF COMMERCE

# National Bureau of Standards

## Certificate of Analyses

OF

STANDARD SAMPLE 55 B

OPEN-HEARTH IRON

ANALYST*	C	Mn	P		S		Si	COPPER H <sub>2</sub> S-CuS-CuO	NICKEL Weighed as nickel dimethylglyoxime	CHROMIUM FeSO <sub>4</sub> -KMnO <sub>4</sub> titration	VANADIUM	MOLYBDENUM Colorimetric	COBALT	TIN	ALUMINUM (total)	ALUMINUM OXIDE (Al <sub>2</sub> O <sub>3</sub> )	ARSENIC	NITROGEN
	Gravimetric (weighed as Mg <sub>2</sub> P <sub>2</sub> O <sub>7</sub> after removal of arsenic)	Alkali-Molybdate <sup>a</sup>	Gravimetric (direct oxidation and final precipitation in reduced solution)	Evolution with HCl (1-1) ZnS-Iodine (theoretical sulfur titre) <sup>b</sup>	Sulfuric acid dehydration													
1	0.010 <sub>7</sub>	0.015	0.002	0.003	0.019	0.020	0.001	0.042	0.016	0.002	<0.001	0.003	0.006	0.005	0.002	0.001	0.009	0.003
2	0.009 <sub>4</sub>	0.017	0.003	0.003	0.018	0.018	0.001	0.039	0.016	0.003	0.004	0.004	0.005	0.007	0.002	0.002		
3	0.010 <sub>2</sub>	0.018	0.004	0.004	0.019	0.019	0.001	0.042	0.016	0.004	0.004	0.004	0.006	0.006	0.003			
4	0.009 <sub>7</sub>	0.014	0.003	0.003	0.018	0.019	0.001	0.042										
5	0.011 <sub>5</sub>	0.017	0.003	0.003	0.016	0.018	0.001	0.039	0.015			0.003	0.008					
6	0.011 <sub>6</sub>	0.015	0.002	0.002	0.021	0.021	0.001	0.046	0.019	0.001		0.005	0.005	0.007	0.003			
7	0.010 <sub>9</sub>																	
ges	0.010 <sub>5</sub>	0.016	0.003	0.003	0.018	0.019	0.001	0.042	0.016	0.003	<0.001	0.004	0.006	0.006	0.003	0.002	0.009	0.003
General average	0.010 <sub>5</sub>	0.016	0.003	0.003	0.018	0.019	0.001	0.042	0.016	0.003	<0.001	0.004	0.006	0.006	0.003	0.002	0.009	0.003

<sup>a</sup> Precipitated at 40° C, washed with a 1-percent solution of KNO<sub>3</sub>, and titrated with alkali standardized by the use of acid potassium phthalate and the ratio 23NaOH:1P.  
<sup>b</sup> Value obtained by standardizing the titrating solution by means of sodium oxalate through KMnO<sub>4</sub> and Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>, and the use of the ratio 2I:1S.  
<sup>c</sup> Direct combustion.  
<sup>d</sup> 10-g samples extracted with ether. Bismuthate (FeSO<sub>4</sub>-KMnO<sub>4</sub>) method.  
<sup>e</sup> Molybdenum-blue photometric method. See J. Research NBS 26, 405 (1941) RP1386.  
<sup>f</sup> 10-g sample, bicarbonate hydrolysis, persulfate oxidation and potentiometric titration with ferrous ammonium sulfate.  
<sup>g</sup> Ether separation on a 10-g sample. Double ZnO precipitation and cobalt precipitated with α-nitroso-β-naphthol. Ignited precipitate dissolved and acid solution treated with H<sub>2</sub>S, filtered, cobalt precipitated with α-nitroso-β-naphthol, ignited and weighed as Co<sub>2</sub>O<sub>3</sub>. See BS J. Research 7, 883 (1931) RP380.  
<sup>h</sup> 50-g sample, H<sub>2</sub>S-distillation-cupferron method. Determination made by R. K. Bell. See BS J. Research 8,

309 (1932) RP415 and J. Research NBS 33, 307 (1944) RP1610.  
<sup>i</sup> 50-g sample dissolved in diluted HNO<sub>3</sub> (1:4).  
<sup>j</sup> Distillation-molybdenum-blue photometric method. See J. Research NBS 24, 7 (1940) RP1267.  
<sup>k</sup> Determination made by Marie C. Wallace by the vacuum-fusion method. See BS J. Research 7, 375 (1931) RP346.  
<sup>l</sup> Same value obtained by evolution and combustion methods.  
<sup>m</sup> Sulfuric-nitric acid dehydration.  
<sup>n</sup> KI-Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub> titration.  
<sup>o</sup> 10-g sample, phenylhydrazine concentration, diphenylcarbazide colorimetric method.  
<sup>p</sup> Solution in diluted HNO<sub>3</sub>, tin precipitated with H<sub>2</sub>S, dissolved by fuming in H<sub>2</sub>SO<sub>4</sub>-HNO<sub>3</sub>, and precipitated with NH<sub>4</sub>OH. Precipitate dissolved in HCl, and tin reduced with antimony and titrated with KIO<sub>4</sub> solution standardized on high-purity tin.  
<sup>q</sup> 50-g sample dissolved in diluted HCl (1:1).  
<sup>r</sup> Low pressure-combustion method.  
<sup>s</sup> Persulfate-arsenite method.

<sup>t</sup> Titrating solution standardized by use of a standard steel.  
<sup>u</sup> Combustion method, SO<sub>2</sub> bubbled through H<sub>2</sub>O<sub>2</sub>-NaOH solution, and excess alkali titrated with H<sub>2</sub>SO<sub>4</sub>.  
<sup>v</sup> Perchloric acid dehydration.  
<sup>w</sup> ZnO-α-nitroso-β-naphthol method.  
<sup>x</sup> Solution in HCl, titration with KIO<sub>4</sub> solution.  
<sup>y</sup> Periodate photometric method.  
<sup>z</sup> Weighed as P<sub>2</sub>O<sub>5</sub>·24MoO<sub>3</sub>.  
<sup>aa</sup> Iron extracted with isopropyl ether.  
<sup>ab</sup> Copper-ammonia-complex colorimetric method.  
<sup>ac</sup> Finished by electrolysis.  
<sup>ad</sup> MoS<sub>3</sub>-MoO<sub>3</sub> method.  
<sup>ae</sup> Iron removed by ether and basic acetate separations, copper by H<sub>2</sub>S. Cobalt separated with KNO<sub>3</sub>, and weighed as CoSO<sub>4</sub>.  
<sup>af</sup> Ether extraction, spectrochemical determination.  
<sup>ag</sup> Bicarbonate hydrolysis, peroxide fusion, K<sub>2</sub>CrO<sub>4</sub> colorimetric method.  
<sup>ah</sup> Solution in HCl, iodimetric titration.

### \* LIST OF ANALYSTS

1. Ferrous Laboratory, National Bureau of Standards, John L. Hague in charge. Analysis by J. I. Shultz, J. P. Hewlett, Jr., and C. Litsey.
2. ARMCO Research Chemical Laboratory, Middletown, Ohio, Arba Thomas in charge. Analysis by L. C. Ikenberry, C. Perrine, O. H. Fritzsche, and LoRene Chenault.
3. D. P. Bartell, Allegheny Ludlum Steel Corporation, Brackenridge, Pa.
4. S. E. Q. Ashley and W. M. Murray, General Electric Co., Pittsfield, Mass.
5. R. H. Wynne and E. W. Beiter, Research Laboratories, Westinghouse Electric & Manufacturing Co., Pittsburgh, Pa.
6. B. L. Clarke, Bell Telephone Laboratories, New York, N. Y.
7. L. A. Wooten, Bell Telephone Laboratories, Murray Hill, N. J.
8. J. B. Austin, Research Laboratory, U. S. Steel Corporation, Kearny, N. J.

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E. U. CONDON, *Director*.