

# Bureau of Standards

## Certificate of Analysis

OF

STANDARD SAMPLE No. 30b

# CHROME-VANADIUM STEEL

MATERIAL FURNISHED BY BETHLEHEM STEEL COMPANY, SOUTH BETHLEHEM, PA.

ANALYST*	C	Mn	P	S	Si	COOPER H <sub>2</sub> S-CuS-CuO	NICKEL Weighed as nickel dimethylglyoxime	Cr		V		MOLYBDENUM	COBALT	ARSENIC			
	CARBON (Direct combustion)	MANGANESE 1. Bismuthate (FeSO <sub>4</sub> -KMnO <sub>4</sub> titration after removing Cr and V). 2. Other methods	PHOSPHORUS 1. Alkali-molybdate (vanadium reduced before molybdate precipitation). 2. Gravimetric (weighed as Mg <sub>3</sub> P <sub>2</sub> O <sub>8</sub> after elimination of arsenic)	SULPHUR 1. Sulphur (direct oxidation and final precipitation in reduced solution) 2. Sulphur Evolution ZnS-Iodine (theoretical sulphur titre)	SILICON Sulphuric acid dehydration			CHROMIUM 1. Persulphate oxidation (FeSO <sub>4</sub> -KMnO <sub>4</sub> titration) 2. Other methods	VANADIUM 1. FeSO <sub>4</sub> -(NH <sub>4</sub> ) <sub>2</sub> S <sub>2</sub> O <sub>8</sub> -KMnO <sub>4</sub> after chromium determination 2. Other methods								
1.....	0.293	0.500	0.502 <sup>c</sup>	0.024	0.024	0.032	0.030	0.215	{0.134 .131 <sup>d</sup> }	0.286	1.03	1.03 <sup>e</sup>	0.207	0.212 <sup>f</sup>	0.018	0.024	{0.021 <sup>g</sup> .018 <sup>h</sup> }
2.....	.298	.504	.505 <sup>i</sup>	.024	.024	{.032 <sup>j</sup> .033 <sup>k</sup> }	.031	.208	.141	.299	1.04	1.02 <sup>l</sup>	.205	.218 <sup>i</sup>	.013	.....	.....
3.....	.291	.....	.488 <sup>c</sup>	.024	.....	.032	.027	.215	.144	.27 <sup>l</sup>	1.02	.....	.....	.22 <sup>m</sup>	.....	.....	.....
4.....	.299	.501	.511 <sup>n</sup>	.023	.....	.033	.028	.211	{.139 .129 <sup>d</sup> }	.291	1.03	.....	.206	{.199 <sup>f</sup> .202 <sup>o</sup> }	.012	.....	.....
5.....	.288	.505	.51 <sup>p</sup>	.024	.....	.033	.032	{.204 <sup>q</sup> .207	.125	{.291 .293 <sup>r</sup> }	1.01	.....	.20	.....	.012	.....	.....
6.....	.30	.50	{.49 <sup>p</sup> .49 <sup>s</sup> }	.....	.....	.032	.....	{.214 <sup>t</sup> .213	.126 <sup>d</sup>	{.29 <sup>u</sup> .29 <sup>r</sup> }	1.01	1.02 <sup>v</sup>	.....	.....	.015	.....	.....
7.....	.301	.496	{.496 <sup>n</sup> .404 <sup>p</sup> }	.026	.....	.032	.030	{.205 <sup>q</sup> .206	.12	.....	1.03	.....	.20 <sup>w</sup>	.....	.....	.....	.....
.....	.285	.51	.....	.026	.026	.032	.028	.215	.124	.29	1.04	.....	.....	{.206 <sup>a1</sup> .210 <sup>f</sup> .210 <sup>e</sup> .201 <sup>o</sup> }	.....	.....	.....
9.....	.286	.501	.....	.022	.....	{.032 .033 <sup>j</sup> }	.....	{.219 <sup>q</sup> .215	.133 <sup>d</sup>	.256	1.05	1.05 <sup>x</sup>	.....	.....	.....	.....	.....
10.....	.288	.496	.....	.024	.....	.030	.029	.220	{.111 <sup>d</sup> .122	.28 <sup>u</sup>	1.01 <sup>y</sup>	1.98	.200 <sup>z</sup>	.....	.....	.....	.....
11.....	.294	.494	.....	.023	.....	{.030 <sup>j</sup> .031	.....	.218	.146	.....	1.01 <sup>i</sup>	.....	.218 <sup>i</sup>	.....	.....	.....	.....
12.....	.285	.....	.50 <sup>p</sup>	.022	.....	.032	.030	.21	.....	.25	1.02	.....	.217	.218 <sup>a1</sup>	.....	.....	.026
13.....	.290	.....	.492 <sup>z</sup>	.023	.....	.033 <sup>j</sup>	.....	.217	{.133 <sup>a2</sup> .127 <sup>z</sup> }	.288 <sup>r</sup>	1.01 <sup>z</sup>	.....	.218 <sup>a3</sup>	.....	.....	.....	.....
AVERAGES.....	.292	.501	.498	.024	.025	.032	.029	.212	.130	.282	1.03	1.02	.206	.209	.014	.024	.022
General Averages..	.292	.499	.024	.032†	.029	.212	.130	.282	1.03	.208†	.014	.024	.022				

NOTE.—By the use of methods employing empirical titres for evolution sulphur, an average of 0.032% was obtained by six analysts.

† Recommended values: 0.215% vanadium; .032% sulphur.  
<sup>a</sup> Value obtained by standardization of titrating solution against sodium oxalate through KMnO<sub>4</sub> and Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub>.  
<sup>b</sup> Described in "Methods of the U. S. Steel Corporation for the Sampling and Analysis of Alloy Steels," 1921, page 47, and in A. S. T. M., "Proposed Tentative Methods of Chemical Analysis of Alloy Steels," serial designation A55-22T.  
<sup>c</sup> Bismuthate-arsenite.  
<sup>d</sup> Finished by electrolysis.  
<sup>e</sup> Peroxide fusi on method.  
<sup>f</sup> Modification of J. R. Cain's method. Reprint 161 from Bulletin of the Bureau of Standards, vol. 7, No. 3, and J. Ind. & Eng. Chem. 3, 476 (1911).  
<sup>g</sup> Weighed as As<sub>2</sub>S<sub>3</sub>.  
<sup>h</sup> Converted the sulphide to arsenate, precipitated as Ag<sub>3</sub>AsO<sub>4</sub>, dissolved in HNO<sub>3</sub> and titrated with KCNS.  
<sup>i</sup> Electrometric titration.  
<sup>j</sup> Precipitated in FeCl<sub>3</sub> solution.  
<sup>k</sup> Same result obtained by Meineke's method.  
<sup>l</sup> Weighed as NiO.

<sup>m</sup> Garratt's method. J. Ind. & Eng. Chem. 4 (1912), 256. Same result obtained by Blair's method. Blair, "The Chemical Analysis of Iron," 7th edition, p. 209.  
<sup>n</sup> Ford Williams' method.  
<sup>o</sup> Cain & Hostetter's method.  
<sup>p</sup> Persulphate-arsenite.  
<sup>q</sup> Solution in HNO<sub>3</sub> and H<sub>2</sub>SO<sub>4</sub>.  
<sup>r</sup> Dissolved the glyoxime precipitate and titrated with KCN.  
<sup>s</sup> ZnO separation-persulphate arsenite.  
<sup>t</sup> Solution in HCl and HClO<sub>4</sub>.  
<sup>u</sup> Direct KCN titration.  
<sup>v</sup> HNO<sub>3</sub>-NaClO<sub>2</sub>.  
<sup>w</sup> HCl reduction.  
<sup>x</sup> Barba's method.  
<sup>y</sup> Bismuthate oxidation—KMnO<sub>4</sub> titration.  
<sup>z</sup> Johnson's method. Johnson, "Chemical Analysis of Special Steels, etc.," 3d edition.  
<sup>a1</sup> Color method.  
<sup>a2</sup> KCN titration.  
<sup>a3</sup> Johnson's method after removal of copper.

### \*INDEX TO ANALYSTS

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