

## U. S. DEPARTMENT OF COMMERCE

**National Bureau of Standards**  
**Certificate of Analyses**  
**OF**  
**STANDARD SAMPLE 20D**  
**ACID OPEN-HEARTH STEEL, 0.4% CARBON**

ANALYST*	C	Mn	P	S	Si								
	<i>Direct combustion</i>	<i>Bismuthate (FeSO<sub>4</sub>-KMnO<sub>4</sub>)</i>	<i>Persulfate Arsenite</i>	<i>Granimetric (weighed as Mg<sub>2</sub>P<sub>2</sub>O<sub>7</sub> after removal of arsenic)</i>	<i>Alkali-molybdate<sup>a</sup></i>	<i>Granimetric (direct oxidation and final precipitation in reduced solution)</i>	<i>Evolution with HCl (1-1) ZnS-Iodine (theoretical sulfur titre)<sup>b</sup></i>	<i>Sulfuric acid dehydration</i>	<i>COPPER H<sub>2</sub>S-CuS-CuO</i>	<i>NICKEL Weighed as nickel dimethylglyoxime</i>	<i>CHROMIUM FeSO<sub>4</sub>-KMnO<sub>4</sub> titration</i>	<i>VANADIUM</i>	<i>MOLYBDENUM Colorimetric</i>
1	0.408	0.916	0.912	0.047	0.047	0.100	0.096	d. 0.256	e. 0.166	0.230	f. 0.281	g. 0.045	h. 0.061
2	.413	g. 918	.048	g. 049	.097	.095	.250	e. 169	.218	.285		b. 057	
3	.408	g. 918		g. 051		g. 094	j. 263	e. 160	.217	.291	.049	.065	
4	.408	g. 900	k. 049	.047	.095	.096	d. 255	e. 161	.223	.270	l. 055	m. 065	
5	.413	.924		.052	.095	.090	.245	e. 161	.219	.291	.048	.064	
6	.414	g. 910	.045	g. 046	.097	g. 099	n. 255	o. 166	.230	.297	.049	.065	
7	.410	f. 913		.047	.101	.093	.252	.158	.239	.278			
8	.412	.92		.050	.101	e. 096	d. 261	.165	.228	f. 272			
9	.414	g. 916		.050	.100	.088	d. 252	o. 166	.235	.287			
10	.409	.914	.915		.050	.097	p. 096	.255	.171	.234	.28	a. 046	.06
Averages	<b>0.411</b>	<b>0.917</b>	<b>0.915</b>	<b>0.047</b>	<b>0.049</b>	<b>0.098</b>	<b>0.093</b>	<b>0.254</b>	<b>0.164</b>	<b>0.227</b>	<b>0.283</b>	<b>0.049</b>	<b>0.062</b>
Recommended values	<b>0.411</b>	<b>0.916</b>		<b>0.048</b>		<b>0.098</b>		<b>0.254</b>	<b>0.164</b>	<b>0.227</b>	<b>0.283</b>	<b>0.049</b>	<b>0.062</b>

\* Precipitated at 40° C, washed with a 1-percent solution of  $KNO_3$  and titrated with alkali standardized by the use of National Bureau of Standards acid potassium phthalate and the ratio 23 NaOH:1 P.

<sup>b</sup> Value obtained by standardizing the titrating solution by means of sodium oxalate through  $KMnO_4$  and  $Na_2S_2O_3$ .

<sup>c</sup> Solution in concentrated HCl.

<sup>d</sup> Double dehydration.

<sup>e</sup> Finished by electrolysis.

<sup>f</sup> Potentiometric titration.

<sup>a</sup> Titrating solution standardized by the use of a standard steel.

<sup>b</sup> Initial  $H_2S$  precipitation. Copper separated with  $NaOH$ , molybdenum again precipitated with  $H_2S$ , and ignited to  $MoO_3$ .

<sup>i</sup> Bismuthate-arsenite method.

<sup>j</sup> Nitric-sulfuric acid dehydration.

<sup>k</sup> Weighed as  $(NH_4)_3PO_4 \cdot 12 MoO_3$ .

<sup>l</sup> Vanadium separated from the bulk of the iron (ferrous) by precipitation with cupferron. Deter-

mination finished by the ferrous sulfate-persulfate method.

<sup>m</sup> a-Benzoinoxime method. See BS J. Research 9, 1 (1932) RP453.

<sup>n</sup> Perchloric acid dehydration.

<sup>o</sup>  $KI-Na_2S_2O_3$  titration.

<sup>p</sup> Sample ignited in oxygen, gases passed into  $H_2O$ , and  $H_2SO_4$  titrated with 0.01 N  $NaOH$ .

<sup>q</sup> Mercury cathode separation, titrated with  $KMnO_4$ .

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