

U. S. DEPARTMENT OF COMMERCE

# National Bureau of Standards

## Certificate of Analyses

OF  
STANDARD SAMPLE 133  
CHROMIUM-MOLYBDENUM STEEL

ANALYST	C	Mn	P		S		Si			Cr	Mo				
	Direct combustion	Bismuthate (FeSO <sub>4</sub> -KMnO <sub>4</sub> )	Persulfate-Arsenite	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 precipitation after reduction of iron)	Combustion	Perochloric acid dehydration	COPPER H <sub>2</sub> S-CuS-CuO	NICKEL Weighed as nickel dimethylglyoxime	FeSO <sub>4</sub> -KMnO <sub>4</sub> titration	VANADIUM	Gravimetric	Colorimetric	NITROGEN
1	0.116	<sup>b</sup> 0.802	<sup>c</sup> 0.800	0.020	<sup>d</sup> 0.020	0.362		<sup>e</sup> 0.434	0.060	0.292	<sup>f</sup> 13.60	<sup>g</sup> 0.018	<sup>h</sup> 0.554	0.549	<sup>i</sup> 0.046
2	.118		<sup>c</sup> 0.785		<sup>i</sup> 0.021	<sup>k</sup> 0.352	<sup>k</sup> 0.373	.440	<sup>l</sup> 0.058	.28	<sup>m</sup> 13.59	.024		<sup>n</sup> 0.565	<sup>n</sup> 0.043
3	.116		<sup>b</sup> 0.782		.023	<sup>k</sup> 0.357		.431		.283	13.59	.025	<sup>c</sup> 0.554		
4	.122		<sup>b</sup> 0.814	<sup>p</sup> 0.023	.024	<sup>q</sup> 0.354		.431	<sup>r</sup> 0.061	<sup>r</sup> 0.276	<sup>m</sup> 13.54	<sup>s</sup> 0.016	<sup>b</sup> 0.568		<sup>s</sup> 0.047
5	.115	.814	<sup>b</sup> 0.792		.022		<sup>k</sup> 0.369	.433	<sup>t</sup> 0.058	<sup>r</sup> 0.30	13.57	<sup>u</sup> 0.020	<sup>v</sup> 0.569	<sup>v</sup> 0.567	<sup>s</sup> 0.047
6	.116		<sup>c</sup> 0.793	.019	<sup>i</sup> 0.019	<sup>k</sup> 0.348	<sup>w</sup> 0.349	<sup>e</sup> 0.436	<sup>x</sup> 0.061	<sup>y</sup> 0.291	13.60	<sup>z</sup> 0.020	<sup>v</sup> 0.571	<sup>v</sup> 0.573	<sup>z</sup> 0.047
7	.120	<sup>z</sup> 0.784			.022			<sup>e</sup> 0.438	<sup>y</sup> 0.069	<sup>y</sup> 0.281	<sup>m</sup> 13.57	<sup>z</sup> 0.021		<sup>z</sup> 0.57	<sup>z</sup> 0.044
	.110	<sup>b</sup> 0.807	<sup>c</sup> 0.808		.022	<sup>q</sup> 0.353	<sup>k</sup> 0.362	<sup>e</sup> 0.436	<sup>1</sup> 0.064	.287	13.55	<sup>u</sup> 0.021	<sup>b</sup> 0.551		<sup>z</sup> 0.046
<sup>g</sup>	.122		<sup>c</sup> 0.798	.021	.022	<sup>k</sup> 0.354	<sup>w</sup> 0.353	.433	<sup>x</sup> 0.060	.304	13.61	.014	<sup>v</sup> 0.535		.048
10	.123		<sup>b</sup> 0.80		<sup>i</sup> 0.023	<sup>k</sup> 0.363	<sup>w</sup> 0.358	.428	<sup>l</sup> 0.057	<sup>y</sup> 0.285	<sup>m</sup> 13.62			<sup>v</sup> 0.562	<sup>s</sup> 0.043
11	.121		<sup>c</sup> 0.798		.022	<sup>q</sup> 0.348	<sup>z</sup> 0.347	.420	<sup>t</sup> 0.065	<sup>y</sup> 0.27	<sup>f</sup> 13.60	<sup>z</sup> 0.024	<sup>b</sup> 0.540		.046
Averages	0.118	0.802	0.797	0.021	0.022	0.355	0.359	0.433	0.061	0.286	13.59	0.020	0.555	0.564	0.046
General averages	0.118	0.799		0.022		0.356		0.433	0.061	0.286	13.59	0.020	0.559		0.046

<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 National Bureau of Standards acid potassium phthalate and the ratio 23NaOH:H<sub>2</sub>P.  
<sup>b</sup> Chromium removed by precipitation with ZnO.  
<sup>c</sup> Chromium volatilized as CrO<sub>2</sub>Cl<sub>2</sub>.  
<sup>d</sup> Colorimetric method. See J. Research NBS 26, 405 (1941) RP1383.  
<sup>e</sup> Double dehydration.  
<sup>f</sup> Persulfate oxidation, potentiometric titration with ferrous ammonium sulfate standardized with recrystallized potassium dichromate.  
<sup>g</sup> Nitric acid oxidation, potentiometric titration with ferrous ammonium sulfate solution standardized with recrystallized potassium dichromate.  
<sup>h</sup> Alpha-benzoinoxime method. See BS J. Research 9, 1 (1932) RP453.  
<sup>i</sup> Determination made by Vernon C. Holm by the vacuum fusion method. See BS J. Research 7, 375 (1931) RP346.

<sup>j</sup> Titrating solution standardized by use of a standard steel.  
<sup>k</sup> Sulfur dioxide absorbed in starch-iodide solution and titrated with KIO<sub>3</sub> solution standardized with standard steels.  
<sup>l</sup> KI-Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> titration.  
<sup>m</sup> Perochloric acid oxidation.  
<sup>n</sup> Solution-distillation method. Sample dissolved in dilute sulfuric acid.  
<sup>o</sup> Weighed as PbMoO<sub>4</sub>.  
<sup>p</sup> Weighed as ammonium phosphomolybdate.  
<sup>q</sup> Meinel's method.  
<sup>r</sup> Glyoxime precipitate ignited and weighed as NiO.  
<sup>s</sup> Solution-distillation method. Sample dissolved in dilute HCl.  
<sup>t</sup> H<sub>2</sub>S-alpha benzoinoxime-CuO.  
<sup>u</sup> Vanadium separated by electrolysis with a mercury cathode, and finally titrated with potassium permanganate solution.

<sup>v</sup> Molybdenum precipitated with H<sub>2</sub>S and weighed as MoO<sub>3</sub>.  
<sup>w</sup> Sulfur gases absorbed in NaOH-H<sub>2</sub>O<sub>2</sub> solution, and excess NaOH titrated with H<sub>2</sub>SO<sub>4</sub>.  
<sup>x</sup> Finished by electrolysis.  
<sup>y</sup> Glyoxime precipitate titrated with NaCN.  
<sup>z</sup> Vanadium precipitated with cupferron, and determined by ammonium persulfate-permanganate method.  
<sup>1</sup> Dissolved in H<sub>2</sub>SO<sub>4</sub>-H<sub>2</sub>PO<sub>4</sub>. Selenium added and solution fumed. Distillation-titration method.  
<sup>2</sup> Chromium removed by precipitation with PbNO<sub>2</sub>.  
<sup>3</sup> Chromium separated as PbCrO<sub>4</sub>. Vanadium determined by differential titration with FeSO<sub>4</sub>-KMnO<sub>4</sub> using o-phenanthroline indicator.  
<sup>4</sup> C. M. Johnson's method. See Iron Age, p. 11, July 26, 1934.  
<sup>5</sup> Sulfur gases absorbed in neutral H<sub>2</sub>O<sub>2</sub> solution, titrated with standard NaOH solution.

\*LIST OF ANALYSTS

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The steel for the preparation of this standard was furnished by the Republic Steel Corporation

WASHINGTON, April 18, 1944.

LYMAN J. BRIGGS, *Director.*