

## National Bureau of Standards

# Certificate of Analysis

## Standard Reference Material 1093

## Oxygen in Valve Steel

This Standard Reference Material (SRM) is intended primarily for the determination of oxygen in ferrous materials. The high manganese content of the valve steel material (9 to 10%) maker the determination of oxygen by vacuum and inert gas fusion methods difficult because of the gettering action of the manganese. However, it is suitable for use with 14 MeV neutron activation analysis. The material for this standard was furnished to NBS by the Allegheny Ludlum Steel Corp., Brackenridge, Pennsylvania.

SRM No.	Description	exygen, ppin	11.	Uncertainty, ppm
		X V	(by weight)	
1093	Valve Strel	00		5 <sup>b</sup>

<sup>&</sup>lt;sup>a</sup>Results determined by 1 MeV neutron activation analysis on 12 g sample weights [1-3].

SRM 1093 is supplied as and 1/4 in (0.5 mm) in diameter and 4 in (8.2 cm) long.

Caution: Oxygen determinations around be made only on thoroughly and freshly cleaned samples that represent the full cross-section of the rods.

Analyses were performed at NJS by S.S. Nargolwalla, G. W. Smith, and J. E. Suddueth.

The overall direction and coordination of technical measurements leading to the certification were under the chairmanship of P. D. LaFleur.

The technical and support aspects concerning the preparation, certification, and issuance of this Standard Reference Material were coordinated through the Office of Standard Reference Materials by R. E. Michaelis.

Gaithersburg, MD 20899 November 5, 1984 (Revision of certificate dated 3-4-69) Stanley D. Rasberry, Chief
Office of Standard Reference Materials

(over)

bUncertainty at the 95 recent confidence level is expressed as  $ts/\sqrt{n}$ , where n= number of samples. Eight determinations per sample of seven samples were made and the uncertainty shown is given as a measure of precision only and not accuracy sin e only one analytical technique was used.

### SUPPLEMENTARY INFORMATION

Although not certified, nitrogen was measured both by a pressure-bomb distillation-titrimetric method ( $4807 \pm 10 \text{ ppm}$ ) and by vacuum fusion (3640  $\pm$  76 ppm).

### PREPARATION FOR THE DETERMINATION OF OXYGEN:

- (1) Samples should be cut from the original rod in such a manner as to minimize heating of the sample; i.e., by a hand hacksaw.
- (2) All surfaces of the cut sample should be thoroughly cleaned with a fine file.
- (3) Samples should be washed with ether, acetone, or other suitable solvent, dried in a stream of warm clean air, and thereafter handled only with clean forceps.
- (4) Analyses should be made as soon after cleaning the sample as possible

Conditions for Analysis at NBS: The rods were cut to 2 in lengths with a hard hacksay and channel in 1,1,1-trichloroethane in an ultrasonic bath for 30 minutes. The samples were dried it a dry nitrogen a mosphere and analyzed with 14 MeV neutrons using the <sup>16</sup>O(n,p) <sup>16</sup>N reaction.

### **REFERENCES:**

- 666 (1968). [1] Nargolwalla, S.S., Crambes, M.R., and DeVoe, J. Anal. Chem
- Chen, shidueth, J.1 [2] Lundgren, F.A. and Nargolwalla, S.S., Angh Chem. 40, 672 (1968).
- [3] Nargolwalla, S.S., Przybylowicz, E.P., Suldueth, J.E., and Pirkhead, S.L., Anal. Chem. 41, 168 (1969).