



**Table 1—Chemical Composition of Weld Deposits<sup>(a)</sup>**

Code	Aluminum in coating, %	C	Mn	Si %	S	P	Al <sup>(b)</sup>	Ti <sup>(b)</sup>	O <sup>(c)</sup> ppm	N <sup>(d)</sup> Total	N <sup>(e)</sup> Residual
P	0	0.069	1.36	0.30	0.007	0.009	<5	37	432	67	33
—	1	0.073	1.39	0.33	0.007	0.008	20	39	427	66	n.d.
—	2	0.078	1.39	0.32	0.007	0.009	44	42	436	66	n.d.
Q	3	0.080	1.41	0.33	0.006	0.009	78	43	432	61	34
—	4	0.080	1.42	0.37	0.006	0.009	120	43	439	54	n.d.
—	5	0.076	1.36	0.37	0.005	0.009	190	36	422	54	24
R	6	0.078	1.31	0.44	0.005	0.008	340	36	431	52	18
—	6.5	0.076	1.30	0.51	0.005	0.008	490	43	423	50	17
S	7	0.079	1.32	0.57	0.004	0.008	610	38	422	48	18

(a) Nb, V, B: <5 ppm.  
 (b) ICP-AES.  
 (c) BALZERS.  
 (d) LECO.  
 (e) TWI, ref. 7.

notch location (Ref. 1). However, the average columnar grain width in the top beads was observed to differ, firstly increasing with increasing aluminum, then decreasing and finally increasing again, as indicated in Fig. 3.

**As-Deposited Weld Metal**—Top beads of six chosen deposits were optically examined and metallographic measurements were made, following the current guidelines (Ref. 8) of IIW Subcommittee IX J, to quantify the major microstructural components, namely: primary ferrite (PF), ferrite with second phase (FS) and acicular ferrite (AF).

The point count results obtained are plotted in Fig. 4 and reveal a complex situation. Initially, aluminum induced a decrease in the volume fraction of acicular ferrite before increasing it again. The changes were compensated for by a reverse trend in the volume fraction of ferrite with second phase, the amount of primary ferrite remaining relatively constant.

Photomicrographs of as-deposited columnar regions of welds containing 5, 78, 340 and 610 ppm Al—designated P, Q, R

and S, respectively—are shown at X100 magnification in Fig. 5. Of note, even at this low magnification, is the coarse structure exhibited at the highest level of aluminum investigated. Further detail, illustrating columnar regions, is shown in Fig. 6. Ferrite with second phase is particularly evident in S, and coarsening of the acicular ferrite laths and the associated microphases is seen to have occurred.

**Reheated Weld Metal**—Examination of the high-temperature regions directly below the top beads revealed similar changes to those encountered in the as-deposited metal, as shown in Fig. 7. In the extreme case (S), ferrite envelopes still delineated the prior austenite grain boundaries, but ferrite with aligned second phase had replaced approximately 50% of the fine acicular structure within the grains.

Modifications were also observed in the low-temperature reheated regions, as seen in Fig. 8, on comparison of the two extreme aluminum levels, P and S. Grain size measurements (Ref. 9) are plotted in Fig. 9 and indicate grain refinement up to approximately 300 ppm Al and thereafter a slight grain coarsening. At high aluminum contents, however, the so-called

fine-grained reheated regions became more duplex in character, and increasing difficulty in measurement, at the recommended (Ref. 9) magnification of 630X, was encountered. The same problem has been reported on adding major alloying elements, such as Mo (Ref. 10) and Cr (Ref. 11), to the C-Mn system.

More detailed examination, using the scanning electron microscope (SEM), revealed that the microphases associated with the equiaxed ferrite grains were martensite/austenite (M/A) and cementite films. Little change in microphase morphology could be detected with increasing aluminum, and the volume fraction increased only marginally from 8 up to approximately 10%. The grain boundary cementite films, however, were perceptibly thicker and possibly more continuous at high aluminum contents, as seen on comparison of the two micrographs shown in Fig. 10.

**Nonmetallic Inclusions**

Six deposits were examined (Ref. 6) at SINTEF (Norway) to determine the composition of the inclusions and the mean

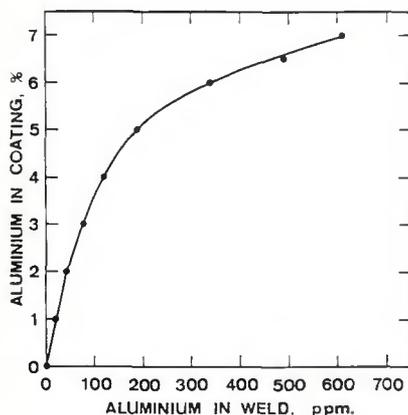


Fig. 1—Plot of aluminum in electrode coating against aluminum in weld metal.

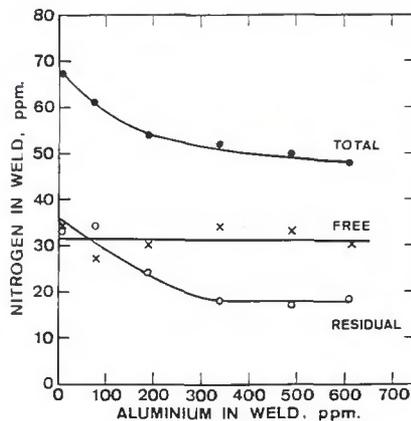


Fig. 2—Plot of residual, free and total nitrogen content against aluminum in weld metal.

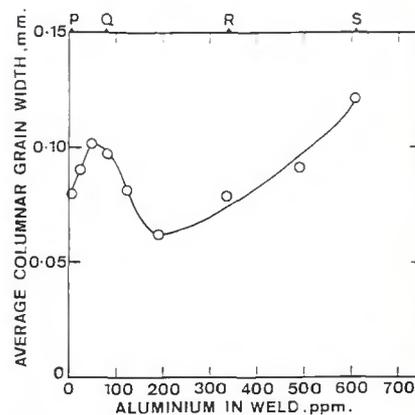


Fig. 3—Effect of aluminum on the average columnar grain width (top bead).











a compromise is required in practice.

The present study is limited to the extent that the results are specific to a weld metal oxygen content of 430 ppm and, consequently, other degrees of deoxidation are currently being explored, so as to define the Al-O system. In addition, since the titanium level in the present case was standardized, further work is envisaged to characterize the Al-Ti system. By this means, it is intended to separate the individual roles of aluminum and titanium and clarify the influence of nonmetallic inclusion interfaces on solidification substructure, prior austenite grain size and acicular ferrite formation. The basis for the work has already been set, the Mn-O system having been studied (Ref.17) in the absence of both aluminum and titanium.

## Conclusions

For a balanced basic low-hydrogen electrode, of a specific slag base type, the following occurred on increasing from zero the amount of aluminum in the coating:

- 1) The microstructure of as-deposited weld metal was modified, with the volume fraction of acicular ferrite initially decreasing, then increasing and finally decreasing again.
- 2) The mean composition of the non-metallic inclusions changed, MnO and SiO<sub>2</sub> being progressively replaced by Al<sub>2</sub>O<sub>3</sub>.
- 3) The hardness of as-deposited metal tended, in the main, to increase.
- 4) The tensile properties of the deposits marginally increased.
- 5) Optimum notch toughness characteristics were exhibited at the zero aluminum level.
- 6) Following degradation from zero aluminum, a small recovery in toughness was encountered at an aluminum content of 350 ppm.

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