

Table 1—Chemical Composition (wt-%)

Material	C	S	P	Si	Mn	Ni	Cr	Mo	Cu	V	Nb	Ti	Al	N
Steel A	0.14	0.004	0.012	0.24	1.43	0.24	0.04	ND	0.01	0.06	0.019	0.005	0.018	0.0078
Steel B	0.10	0.011	0.015	0.46	1.42	0.28	0.05	ND	0.20	0.08	ND ^(a)	ND ^(a)	0.060	0.0057

(a) ND = not detected (< 0.005).

was -10°C (14°F). After the completion of each test, the specimens were sectioned, polished and etched to ensure that the proper microstructure (*i.e.*, the ICHAZ) was sampled by the fatigue crack.

Microstructural Analysis

Although some optical work was conducted, the base materials and the IC-HAZs were primarily examined using SEM and TEM. The optical and SEM techniques were relatively routine; therefore, only the TEM procedures will be briefly outlined. In order to ensure that the narrow ICHAZ region was in the vicinity of the small hole produced by electropolishing, careful preparation of the HAZ specimen was necessary before punching the 3-mm disk. One side of the sample was polished/etched and the location of interest marked. Then the sample was mechanically thinned to a thickness of 0.005–0.006 in. (0.127–0.152 mm) by grinding from the unmarked side. At this stage the 3-mm disk was punched out making sure that the marked region remained at the center. This procedure maximized the chances of making the ICHAZ region electron transparent. The disks were twin-jet electropolished using an electrolytic solution consisting of 300 mL of methanol, 175 mL of butynol, and 30 mL of per-

Table 2—Mechanical Properties

Material	Transverse Tensile Test Data				Transverse Charpy Data at 1/2 thickness	
	Yield Strength (MPa)	Tensile Strength (MPa)	Elongation (%)	Reduction of Area (%)	35 J ^(a) (°C)	50 J ^(a) (°C)
Steel A	362	518	29	71	-80	-75
Steel B	306	506	28	70	-90	-85

(a) These values refer to the 35 J or 50 J transition temperature.

chloric acid. The polishing was done at -40°C. All of the thin foils were examined in a Philips EM 430 microscope at an accelerating voltage of 300 kV. The microscope was equipped with an EDAX detector (20-deg take-off angle) and a 5500 EDAX analyzer. The TEM procedures included dark-field, bright-field, microdiffraction and EDS analysis.

Results

Charpy and CTOD Data

The Charpy data indicate that the ICHAZ for both steels incurred a reduction in toughness from that of the base metal; the 35-J and 50-J transition tem-

Table 3—ICHAZ Toughness Data

Material	Charpy Tests		CTOD Tests (mm)
	35 J ^(a) (°C)	50 J ^(a) (°C)	
Steel A	-42	-36	2.02, 1.99, 1.75
Steel B	-40	-30	0.01, 0.08, 0.03

(a) These values refer to the 35 J or 50 J transition temperature.

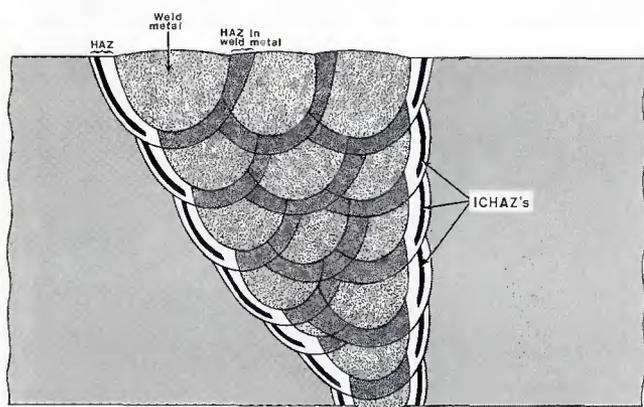


Fig. 3 — Schematic of weld cross-section showing the location of the intercritical HAZs (ICHAZs).

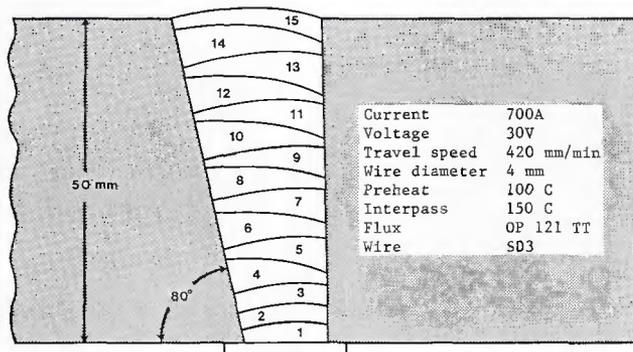


Fig. 4 — Weld procedure geometry and details.

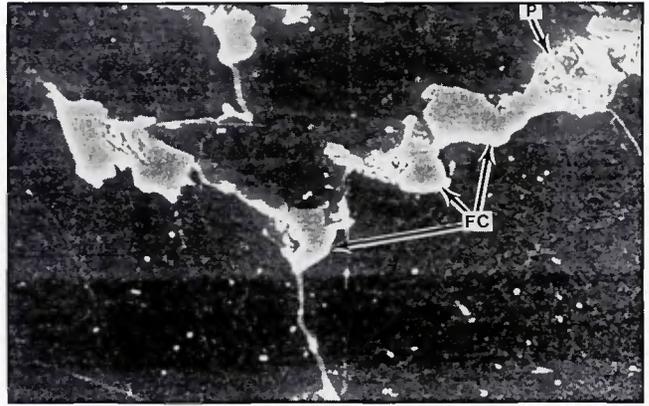
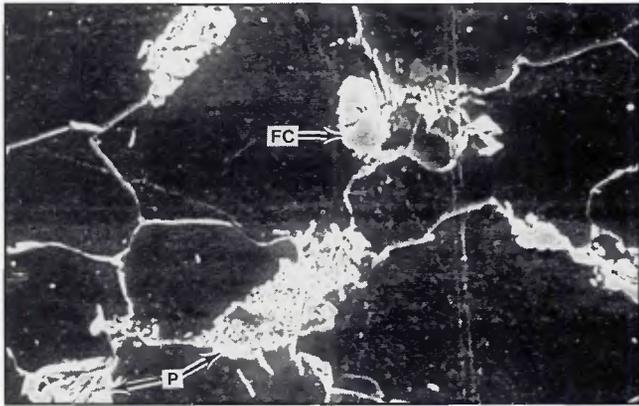


Fig. 7 — SEM micrographs of the ICHAZ for Steels A (left) and B (right). Examples of pearlite are denoted by "P" and the featureless constituent is denoted "FC."

tradition to the Charpy data. Typical required HAZ CTOD values for these steels (Ref. 8) are in the range of 0.15–0.35 mm (0.006–0.014 in.). While Steel A easily meets these required values, Steel B does not.

Microstructure — SEM Examined

The base metal microstructures for both steels, as photographed in the SEM at two different magnifications, are shown in Fig. 6. The steels exhibit a ferrite-pearlite microstructure. From observation of several areas of the samples, it was concluded that no detectable difference in the morphology of the pearlite exists in the two steels, although the amount of pearlite in Steel A is somewhat higher due to a higher carbon content. As seen in Fig. 6, Steel A has a slightly smaller grain size than Steel B; 15 microns compared to 20 microns.

Figure 7 shows SEM micrographs of the ICHAZ for the two steels. The difference in grain size is again apparent. Both steels show a primary microstructure of ferrite with second-phase networks (is-

lands) distributed around the ferrite grain boundaries. Considering the relative amounts of the two phases, it is believed that during welding the peak temperature in this region of the ICHAZ was about 780°C (1436°F). It was apparent that the structural makeup of the second phases was different between Steels A and B. In Steel A, the second phase is predominantly pearlite with the remainder being a featureless constituent (featureless at the magnification level afforded by SEM). In Steel B, pearlite comprises only a minor fraction of the second phase, while the major portion is a featureless constituent. Figure 7 shows that the featureless constituent in Steel B is relatively continuous and well connected around the ferrite grain boundaries. TEM analyses were required to determine the nature of the featureless constituent and to observe any microalloy precipitates present in the ferrite.

Microstructure — TEM Examined

Figure 8 shows TEM micrographs of the base metal ferrite in both steels. A

large number of microalloy precipitates in the ferrite of Steel A are apparent. These precipitates were identified as primarily Nb-V carbonitrides (Nb,V(C,N)). In contrast, the ferrite of Steel B contains few precipitates. The few observed are primarily V-bearing carbonitrides (V(C,N)). Apparently, a significant amount of the V in Steel B is in solid solution in the ferrite matrix.

TEM micrographs of the ferrite phase from the ICHAZ of both steels are shown in Fig. 9. It is clear from this figure that the microalloy precipitates seen in the base metal ferrite of Steel A are also present in the ICHAZ. The ICHAZ ferrite of Steel B is largely void of microalloy precipitates, as was the base metal ferrite. These observations indicate that the microalloy precipitation pattern present in the base metal ferrite survives the ICHAZ thermal cycle.

TEM micrographs of the second phase from the ICHAZ of Steel A are shown in Fig. 10. The predominant second-phase constituent is pearlite (including degenerate pearlite) and is shown in Fig. 10A. The minor con-

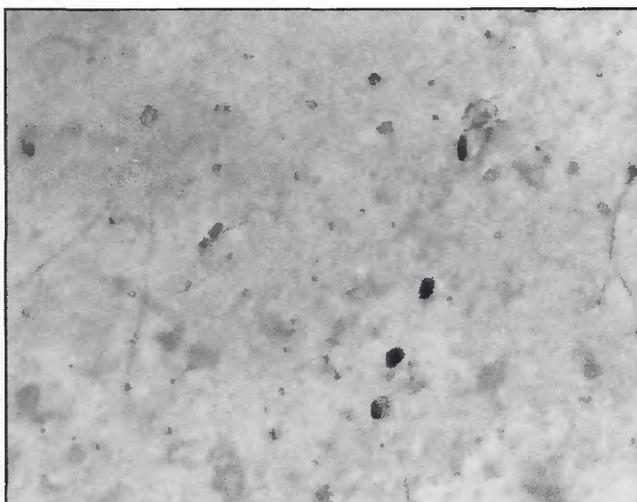


Fig. 8 — TEM micrographs of the base metal ferrite in Steels A (left) and B (right).

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Interpretive Report on Dynamic Analysis of Pressure Components—Fourth Edition

This fourth edition represents a major revision of WRC Bulletin 303 issued in 1985. It retains the three sections on pressure transients, fluid structure interaction and seismic analysis. Significant revisions were made to make them current. A new section has been included on Dynamic Stress Criteria which emphasizes the importance of this technology. A new section has also been included on Dynamic Restraints that primarily addresses snubbers, but also discusses alternatives to snubbers, such as limit stop devices and flexible steel plate energy absorbers.

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Calculation of Electrical and Thermal Conductivities of Metallurgical Plasmas

By G. J. Dunn and T. W. Eagar

There has been increasing interest in modeling arc welding processes and other metallurgical processes involving plasmas. In many cases, the published properties of pure argon or helium gases are used in calculations of transport phenomena in the arc. Since a welding arc contains significant quantities of metal vapor, and this vapor has a considerably lower ionization potential than the inert gases, the assumption of pure inert gas properties may lead to considerable error. A simple method for calculating the electrical and thermal conductivities of multicomponent plasmas is presented in this Bulletin.

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