

using different travel speeds. For the GMA welds, welding parameters were 180–260 A, 26–31 V, direct current electrode positive (DCEP), 38 cm/min welding speed and 15 L/min gas flow. All three levels of moisture in the shielding gas (as supplied, 500 ppm and 1000 ppm) were used. The system was purged with dry nitrogen during intervening periods.

During welding, atmospheric conditions were monitored and were within the 16–19°C temperature and 40–60% relative humidity range. All welding was manual.

For GTA welding, a water-cooled torch was used with a 11.5-mm-diameter gas cup, with provisions for a laminar gas flow. The GMA gun had a 16-mm-diameter conic gas cup.

Test Program

In order to be able to study the relationship between process variables and hydrogen content, specimens were welded according to ISO 3690/IIW II-1155-91 standards with the exception that all welds were made on 22%Cr duplex base metal. The AWS or ISO standards for ferritic mild steel are not appropriate for measuring the hydrogen content in (super) duplex stainless steel weldments, as can be explained from phase transformations of duplex stainless steels (Ref. 10).

Based on previous experiments by the authors (Refs. 1, 2), Fekken (Ref. 9) and Lundin (Ref. 3), an extraction in a quartz capsule for 72 h at 400°C was applied initially for determination of hydrogen content in (super) duplex stainless steel. A recent group-sponsored project at TWI (Ref. 11) indicated that significant levels of hydrogen might remain in the sample after encapsulation; therefore, all sam-

ples were analyzed at 900°C after hydrogen extraction for 72 h at 400°C.

Hydrogen content in GTA and GMA welds was determined using both duplex and super duplex consumables. Gas tungsten arc filler metal of a 2.4-mm diameter was used with argon 4.0 and argon + 30% helium shielding gas. Two different welding speeds resulted in heat inputs varying from 0.8 to 2.6 kJ/mm. For 1.2-mm-diameter GMA welding wires, argon + 2% oxygen and argon + 2% carbon dioxide were used as shielding gases, and welding was performed with heat inputs ranging from 0.8 through 1.3 kJ/mm.

The weld metal ferrite content was determined for both hydrogen test samples and the restrained welding specimen using Magne Gage according to ANSI/AWS 4.2-91.

The eight most severe conditions (as indicated in Table 5) were tested for cracking susceptibility using a three-point bend test as shown in Fig. 3. Here a longitudinal bead-on-plate weld was made on a 16-mm duplex stainless steel base plate, which, immediately after cooling in air, was placed in the bend test unit for 24 h, where it was bent to a 20-deg angle using a 50-mm former diameter. This configuration results in approximately 12% strain on the top surface of the weld (calculated), which was maintained for 24 h.

After this load, the specimens were examined using the liquid penetrant technique. When no cracks were identified, a further bend of the specimen to 90 deg, which is equivalent to approximately 27% (calculated) strain on the top surface of the weld, was applied. The test welds were inspected microscopically for the presence of cracks. Figure 4 illus-

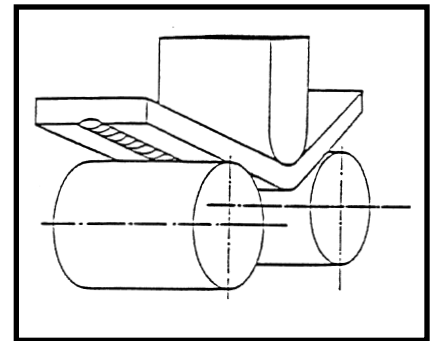


Fig. 3 — Schematic bend test setup with position former, support and weld bead.

trates a rutile SMA welded specimen bent to 90 deg.

Results

In an earlier program (Ref. 2), where test conditions were identical to this investigation, cracking data were presented for shielded metal arc, flux cored arc, gas tungsten arc and submerged arc welds as shown in Fig. 5. In that investi-

Table 5 shows a summary of these variables. None of the investigated welds showed any (micro) cracks, even after bending the specimen to a 90-deg angle, which is equivalent to approximately 27% plastic strain at the top surface in the weld. An overview of the results is illustrated in Figs. 14 and 15.

Discussion

Comparison tests, performed in several laboratories over the last years, demonstrated a good relationship between vacuum hot extraction (VHE) and the encapsulation technique in quartz, followed by an extraction for 72 h at 400°C.

A recent publication out of a group-sponsored project at TWI (Ref. 11), however, indicated that not all hydrogen in the weld metal might evolve using the encapsulation technique. In that publication, an additional hot extraction method at 900°C, following the encapsulation test method, gave an extra volume of hydrogen. The hydrogen release from a sample in a capsule was concluded to be hindered by the development of a positive hydrogen partial pressure within the capsule.

Since the encapsulation technique has been used in a number of investigations and is the basis for several publications worldwide, this project used the encapsulation technique, followed by hot extraction at 900°C.

Hydrogen present in the base metal and the welding consumables previously was identified as the first potential source for weld metal hydrogen. This hydrogen level in consumables may vary from one heat to another. However, the initial hydrogen level of the consumable appeared to be of less importance. Both GTA and GMA weldments, regardless of

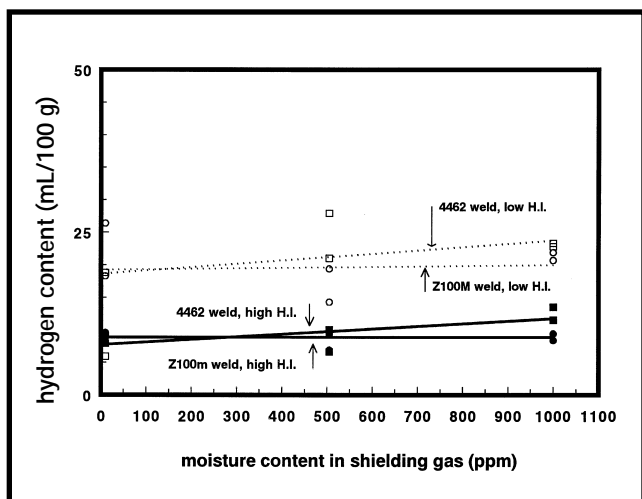


Fig. 7 — Effect of moisture content in Ar + 30%He shielding gas on hydrogen content for GTA welds.

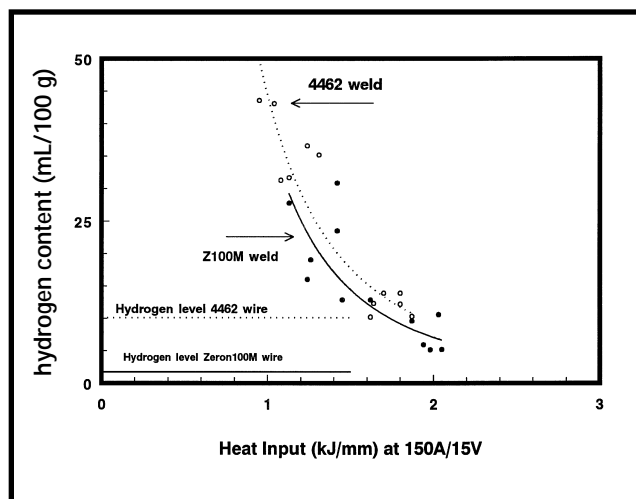


Fig. 8 — Effect of heat input (welding speed) on hydrogen content for GTAW with Ar 4.0.

