

Fig. 1 -- Schematic diagram of the weld groove.

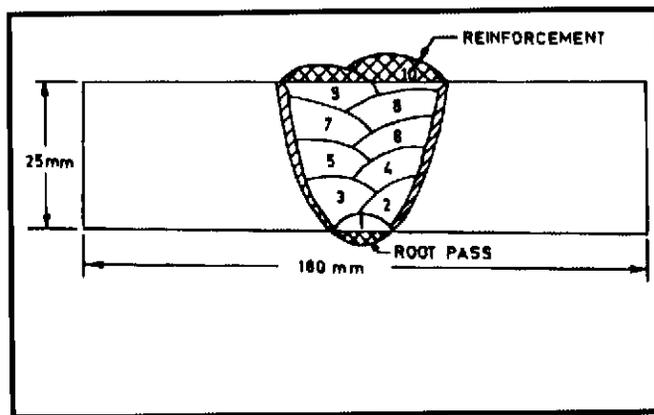


Fig. 2 -- Schematic diagram of the sequence of weld passes.

with the microstructure, zinc pickup, porosity content, and tensile strength of the weld. Finally, the properties of the weldments prepared by multipass pulsed current GMA welding are compared with those of the multipass weldments prepared by conventional continuous current GMA welding.

Experimental Procedure

Pulsed current GMA welding of an extruded section (40 x 25 mm) of Al-Zn-Mg alloy (7005) was carried out using 1.6-mm-diameter filler metal (5183) and commercial argon (99.98%) as shielding gas at a flow rate of 20 L/min. The welding was carried out by filling the weld groove (Fig. 1) in ten passes — Fig. 2. Prior to welding, the plates were thoroughly cleaned to remove the oxide layer and any dirt or grease adhering to the groove surface. The pulsed current welding was carried out at a given mean current of 220 A where the pulse frequency and pulse duration were varied in the range of 25–100 Hz at 4.5–8.5 ms, respectively — Table 1. The mean current was selected on the basis of earlier observations (Refs. 6–13) that showed its effectiveness in improving weld properties. Before welding, a number of pieces were

placed side by side, so their grooves were properly aligned for each weld pass and rigidly held in a fixture to minimize distortion during welding. The welding was carried out on a stainless steel backing plate to obtain uniform root penetration. Both the preheat and interpass temperatures were maintained as 393 K. During the deposition of the fill passes adjacent to the groove edge, a clearance of about 1.5 mm was maintained between the electrode and the groove edge to ensure proper fusion of the base metal. Between subsequent fill passes, the surface of the earlier deposit was thoroughly cleaned using a stainless steel wire brush. For a comparative study, the welds were also produced with conventional continuous current (without pulse) GMA welding process using the same welding current as that of the mean current used in the pulsed current welding. The arc voltage and welding speed were kept constant as 23 V and 300 mm/min, respectively. During welding the pulse characteristics were recorded with an oscilloscope. The peak current (I_p), pulse duration (t_p) and pulse off time (t_{off}) were estimated (Table 1) with the help of a digital oscilloscope; whereas, the mean (I_m) and base (I_b) currents were recorded from an ammeter fixed in the welding power source.

Before collecting specimens for testing, the weldments were stored at room temperature (295 K) for more than 30 days so they would naturally age sufficiently. The weld reinforcement of both the top and bottom surfaces of the weldments was machined flat. The presence of zinc in the weld deposit and the detailed chemical composition of the base and filler metals were analyzed under an atomic absorption spectrometer.

Specimens for metallographic examination were cut from the central part of the weldment to ensure a true representation of the weld characteristics. Transverse sections of the specimens were prepared by standard metallographic technique and suitably etched in Keller's reagent to reveal the microstructure. The microstructure of the weld was studied both under optical and scanning electron microscopes. The quantification of the coarse and fine dendritic portions of the matrix was carried out by the standard lineal intercept method (Ref. 18). An x-ray diffraction study of the powder collected from the weld metal was carried out to determine the presence of any intermetallic precipitate. Porosity content of the weld (excluding reinforcement), revealed as dark spots on the metallographically polished and unetched transverse section of the weld, was estimated under the optical microscope following the standard point counting method (Ref. 18).

A tensile test of the weld joint was performed with a Servo Hydraulic Universal Testing Machine using round tensile specimens (DIN 50215), which had weld metal at their centers as shown in Fig. 3. Tensile properties of the base metal were also determined using the similar round tensile specimen. The tensile test was carried out at a cross head speed of 1 mm/min. Yield strength was determined at 0.2% offset strain on the

Table 1 — Welding Parameters

Specimen Designation	Mean Welding Current (A)	Pulse Frequency (Hz)	Pulse Duration (ms)	Pulse off time Fraction	Base Current (A)	Peak Current (A)
A	220	0	0	—	—	—
B	220	25	4.5	0.8875	190	373
C	220	50	4.5	0.775	160	366
D	220	100	4.5	0.55	110	352
E	220	25	6.5	0.8375	160	438
F	220	50	6.5	0.675	120	421
G	220	100	6.5	0.35	60	361
H	220	25	8.5	0.7875	150	455
I	220	50	8.5	0.575	110	424
J	220	100	8.5	0.15	50	376

load vs. strain diagram obtained with an extensometer fixed on the gauge length portion (50 mm) of the specimen. The elongation of the joint was also estimated over the same gauge length.

Fatigue life (N cycle) of the weldments was studied using three specimens from each welding parameter. The specimens, having a cross-sectional area of 12.5 x 7 mm and gauge length of 50 mm with the weld at its center (Fig. 4), were tested according to the ASTM (E466-82) standard (Ref. 19). Tests were carried out under uniaxial loading on a hydraulically operated pulsator operated at a mean stress of 80 N/mm², a stress ratio (minimum stress/maximum stress) R of 0.5, and a stress cycle of 5 Hz. The number of cycles to fracture was recorded.

Results and Discussion

Chemical analysis

The chemical composition of the base and filler metals is shown in Table 2. The table shows that the filler metal does not contain zinc but the base material has a considerable amount of zinc in it. The selection of (Al-Mg) filler metal containing no zinc was made as per recommended practice (Ref. 20), where the weld deposit provides good mechanical properties and corrosion resistance. However, the weld deposit has been found to pick up zinc from the base metal due to dilution (Table 3). The influence of pulse frequency duration and off time fraction ($t_o/(t_p + t_o)$) on the zinc pickup of the weld deposit are given in Table 3. At a given pulse frequency, the increase in pulse duration was found (Table 3) to enhance the zinc pickup in weld deposit; whereas, at a given pulse frequency, the increase in pulse off time fraction was found (Table 3) to reduce the zinc pickup in the weld. The existence of zinc in the weld deposit caused precipitation of $Mg_3Zn_3Al_2$ in the matrix, as identified (Table 4) by the presence of (Al,Zn)₄₉Mg₃₂ under the x-ray diffraction studies. The presence of (Al,Zn)₄₉Mg₃₂, which is in close approximation by nature (Ref. 21) to the $Mg_3Zn_3Al_2$, has been identified and confirmed by the interplanar spacing (d-value) where, no other interfering d value is observed. The stoichiometric amount of precipitate has been found to increase with the enhancement of zinc pickup as shown in Table 4.

Microstructure

The microstructure of the multipass weld was found to consist of a mixture of coarse columnar dendritic and relatively finer dendritic regions. These two regions are largely defined by their different dendrite arm spacing and the aspect ratio of dendrite arms measured under a scanning electron microscope (SEM). It was observed that the dendrite arm spacing of the coarse columnar dendrite was

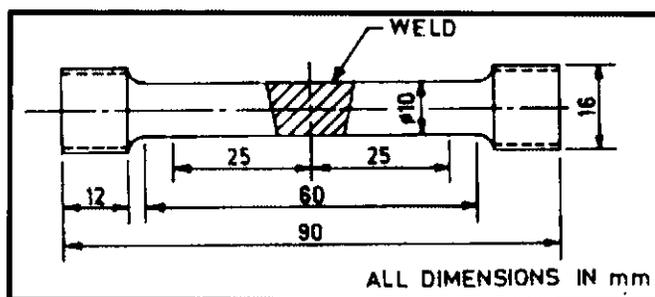


Fig. 3 — Schematic diagram of the tensile specimen.

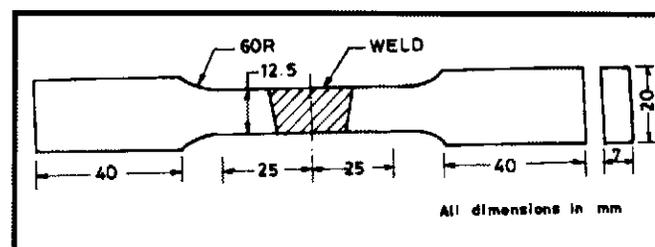


Fig. 4 — Schematic diagram of the fatigue test specimen.

Table 2 — Chemical Composition of Base Metal and Filler Metal

Material	Zn	Mg	Mn	Fe	Si	Cu	Cr	Al
Base metal (Wt-%)	4.5	1.20	0.47	0.32	0.30	0.02	0.15	Rest
Filler Metal (Wt-%)	—	4.5	0.65	—	—	—	—	Rest

Table 3 — Dilution, Zinc Content, Amount of Precipitate, Fine Dendrite Fraction and Porosity Content of the Weld Deposit at Different Welding Parameters

Specimen Designation	Dilution (%)	Zinc Pickup (Wt-%)	Amount of (Al, Zn) ₄₉ Mg ₃₂ Precipitate (Wt-%)	Finer Dendrite Fraction (Area-%)	Porosity Content (Vol-%)
A	17.5	0.64	1.0539	39	2.4
B	15.5	0.60	0.9880	51	3.4
C	13	0.53	0.8563	59	2.9
D	14.5	0.56	0.9180	49	2.3
E	17.5	0.67	1.1033	60	3.0
F	15.5	0.58	0.9551	65	2.7
G	18	0.69	1.1362	56	2.1
H	19.5	0.72	1.1856	55	2.25
I	17.5	0.68	1.1198	61	1.80
J	21	0.78	1.2845	52	1.40

Table 4 — X-ray Diffraction Analysis of the Weld Bead

Sin θ	d A°	(h k l)
0.761	1.0124	(400) Al
0.659	1.169	(222) Al
0.631	1.221	(311) Al
0.538	1.431	(220) Al
0.536	1.437	(941, 853) (Al, Zn) ₄₉ Mg ₃₂
0.383	2.012	(710, 550) (Al, Zn) ₄₉ Mg ₃₂
0.380	2.024	(200) Al
0.333	2.310	(611, 532) (Al, Zn) ₄₉ Mg ₃₂
0.330	2.338	(111) Al

