

Effect of Cr and Ni on diffusion bonding of Fe₃Al with steel

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Abstract. Microstructure at the diffusion bonding interface between Fe₃Al and steel including Q235 low carbon steel and Cr18–Ni8 stainless steel was analysed and compared by means of scanning electron microscopy and transmission electron microscopy. The effect of Cr and Ni on microstructure at the Fe₃Al/steel diffusion bonding interface was discussed. The experimental results indicate that it is favourable for the diffusion of Cr and Ni at the interface to accelerate combination of Fe₃Al and steel during bonding. Therefore, the width of Fe₃Al/Cr18–Ni8 interface transition zone is more than that of Fe₃Al/Q235. And Fe₃Al dislocation couples with different distances, even dislocation net occurs at the Fe₃Al/Cr18–Ni8 interface because of the dispersive distribution of Cr and Ni in Fe₃Al phase.

Keywords. Cr; Ni; Fe₃Al/steel interface; diffusion bonding.

1. Introduction

Fe₃Al intermetallic is a candidate material for structural application in many industrial fields such as aviation, automobile and energy transition system because of its excellent strength at elevated temperature, resistance to oxidation and low cost (Mckamey *et al* 1991; Sun *et al* 2001). The weldability of Fe₃Al intermetallic has also drawn wide attention of the research community. The welding processes used are gas tungsten arc welding, electron beam welding and vacuum diffusion bonding (David and Zacharia 1993; Gao *et al* 2000; Li and Wang 2003). The results indicate that only by choosing weld wire with the same chemical composition as the base metal and strictly controlling welding heat input, can weld crack be avoided when using gas tungsten arc welding. While using electron beam welding and vacuum diffusion bonding processes, weld crack can be effectively avoided because of the vacuum atmosphere in welding. In addition, the vacuum diffusion bonding of Fe₃Al intermetallic with dissimilar materials has been studied (Wang *et al* 2001, 2003). But reports on the effect of alloying elements on diffusion bonding of Fe₃Al with steel are few.

In this paper, microstructure at the diffusion bonding interface between Fe₃Al and steel including Q235 low carbon steel and Cr18–Ni8 stainless steel was analysed and compared by means of SEM and TEM. The effect of Cr and Ni on microstructure at the Fe₃Al/steel interface was discussed. The results indicate that it is possible to pro-

note favourable microstructure at the Fe₃Al/steel diffusion bonding interface.

2. Experimental

The base metals in the test are Fe₃Al intermetallic, Q235 low carbon steel and Cr18–Ni8 stainless steel. The chemical composition of Q235 steel is (wt%) C 0.14%, Si 0.10%, Mn 0.5%, S 0.035%, P 0.035% and Fe balance. The chemical composition and thermo-physical properties of Fe₃Al and Cr18–Ni8 steel are listed in table 1.

Samples for diffusion bonding were prepared by grinding on sand paper, acetone cleaning, alcohol cleaning, water rinsing and drying. Diffusion bonding was conducted in vacuum chamber of Workhorse II equipment immediately after the above treatments. The diffusion bonding was carried out at 1333 K for 60 min with a pressure of 15 MPa in a vacuum of 4.5×10^{-5} Pa. In the heating process, several holding temperature stages were included to promote uniformity of heat in the samples. The temperature of chamber is brought down to 373 K by water circulation and then it was cooled to room temperature within the furnace.

The samples of Fe₃Al/steel joint were cut and etched with a solution of 75% hydrochloric acid and 25% nitric acid for metallographic observation. Microstructural characteristics and fine structures at the Fe₃Al/Q235 and Fe₃Al/Cr18–Ni8 interface were analysed by means of JXA-840 scanning electron microscope (SEM) and transmission electron microscope (TEM) with PV9900-H-8010 energy spectrum analysis system.

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3. Results and discussion

3.1 Effect on microstructure

Microstructure at the Fe₃Al/steel diffusion bonding interface obtained under 1333 K for 60 min with a pressure of 15 MPa is shown in figure 1. It can be seen that there is obvious mutual diffusion and an interface transition zone is formed between Fe₃Al and steel. Microstructure in the interface transition zone and on two sides of Fe₃Al and steel is interlaced. The width of Fe₃Al/Cr18–Ni8 interface transition zone is 35 μm, which is more than that of Fe₃Al/Q235 interface transition zone (29 μm) under the same bonding parameters (1333 K × 60 min, 15 MPa). This is because the diffusion of Cr and Ni from Cr18–Ni8 to Fe₃Al can affect the dynamic balance at the Fe₃Al/Cr18–Ni8 interface and accelerate the diffusion of Al towards the side of Cr18–Ni8 to improve the bonding between Fe₃Al and Cr18–Ni8.

In addition, there are second phases with irregular shape distributed dispersively in the interface transition zone between Fe₃Al and steel. The chemical composition of second phases was analysed by means of electron probe microanalysis (EPMA) and the results are listed in table 2.

C and Cr contents in the second phase of the Fe₃Al/Q235 interface transition zone are more than that in the matrix and Fe and Al contents are less, while in the Fe₃Al/Cr18–Ni8 interface transition zone, Cr and Ni contents are more than that in the matrix and Fe and Al contents are also less. This is because of the insufficient time for C and Cr to diffuse fully in the Fe₃Al/Q235 interface transition zone during diffusion bonding. The second phase rich in Cr and Ni in the Fe₃Al/Cr18–Ni8 interface transition zone can restrain effectively C precipitation to avoid the formation of carbide.

3.2 Effect on fine structure

TEM morphology and electron diffraction pattern of Fe₃Al in the Fe₃Al/Q235 interface transition zone are shown in figures 2a–c. The results from electron diffraction pattern and index schematic diagram indicate that Fe₃Al formed in Fe₃Al/Q235 interface transition zone exhibits a superlattice. It can be seen from [110] orientation that fine structure of Fe₃Al consists of dislocation walls which include several small regions. The production of Fe₃Al dislocation walls is closely related to the

Table 1. Chemical composition and thermo-physical properties of Fe₃Al and Cr18–Ni8 steel.

Chemical compositions (wt%)										
Material	Al	Cr	Nb	Ni	Zr	B	Ce	S	P	Fe
Fe ₃ Al	16.0 ~ 17.0	2.40 ~ 2.55	0.95 ~ 0.98	–	0.05 ~ 0.15	0.01 ~ 0.05	0.05 ~ 0.15	–	–	Bal.
Cr18–Ni8	–	18.21	–	9.43	–	–	–	0.03	0.03	Bal.
Thermo-physical properties										
Material	Structure	Order critical temperature (K)	Young's modulus (GPa)	Melting point (K)	Coefficient of thermal expansion (10 ⁻⁶ .K ⁻¹)	Density (g.cm ⁻³)	Tensile strength (MPa)	Elongation (%)	Hardness	
Fe ₃ Al	DO ₃	813	140	1813	11.5	6.72	455	3	≥ 29 HRC	
Cr18–Ni8	–	–	–	–	16.7	8.03	520	40	70 HRB	

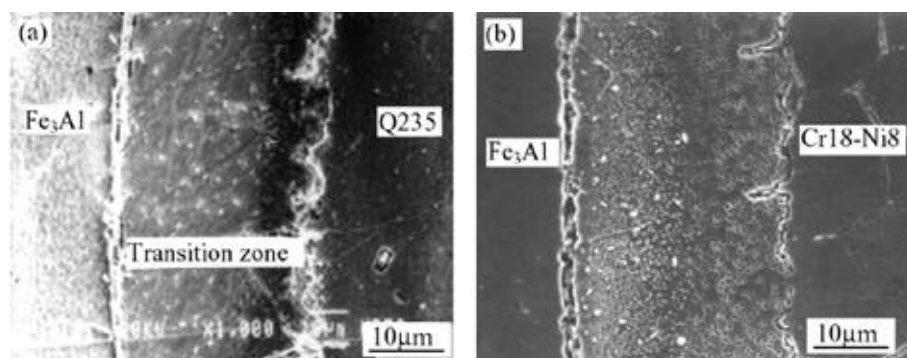


Figure 1. Microstructure at the Fe₃Al/steel diffusion bonding interface: (a) Fe₃Al/Q235 and (b) Fe₃Al/Cr18–Ni8.

motion of dislocations. The dislocation walls can affect dislocation glide and even microstructure at the Fe₃Al/Q235 diffusion bonding interface.

TEM morphology, electron diffraction pattern and index schematic diagram of Fe₃Al in the Fe₃Al/Cr18–Ni8 interface transition zone are shown in figures 3a–c. Dislocations with [001] orientation are distributed in couples and the spacing is small between the dislocation couples. This is because Cr and Ni diffused at the Fe₃Al/Cr18–Ni8 interface can hamper the superlattice dislocation motion in Fe₃Al. The external applied stress can make dislocations decompose into unit dislocations and they slide alternatively. In addition, Cr and Ni can decrease the antiphase boundary energy during the deformation of Fe₃Al (Mckamey *et al* 1990; Yin *et al* 1996). This will cause an increase in the spacing between dislocation couples.

Fe₃Al dislocations with $[1\bar{2}2]$ orientation are mostly unpaired and they are tangled in some regions (figures 4a–c). This has resulted from the diffusion of Cr and Ni at the Fe₃Al/Cr18–Ni8 diffusion bonding interface. The diffused Cr and Ni can be precipitated in the second phase and distributed dispersively in Fe₃Al. When the individual dislocation moves to the second phase, continued glide is hampered and they form a dislocation net.

When the dislocation pairs with small spacing and dislocations in a network at the Fe₃Al/Cr18–Ni8 diffusion bonding interface move during the diffusion bonding, they restrict the smooth glide of each other. Fe₃Al has these various dislocation features at the Fe₃Al/Cr18–Ni8 diffusion bonding interface because of Cr and Ni diffusion which are favourable for improving the strength of the diffusion bonding interface.

Table 2. EPMA analysis on second phase in the transition zone between Fe₃Al and steel (%).

Position	No.	Fe	Al	C	Cr	Nb	Zr	Ni
Fe ₃ Al matrix	1	82.60	16.62	0.14	0.52	0.05	0.07	–
	2	82.80	16.40	0.13	0.49	0.03	0.05	–
	3	81.91	17.10	0.13	0.51	0.03	0.02	–
	4	82.21	16.90	0.13	0.54	0.01	0.01	–
Fe ₃ Al/Q235	1	81.69	16.35	0.55	1.18	0.01	0.02	–
	2	81.94	15.90	0.51	1.28	0.03	0.04	–
	3	82.54	15.47	0.40	1.32	0.04	0.03	–
	4	81.89	16.23	0.22	1.26	0.04	0.06	–
Fe ₃ Al/Cr18–Ni8	1	71.8	12.5	–	8.9	–	–	4.4
	2	72.3	9.5	–	13.2	–	–	4.5
	3	73.9	2.5	–	17.2	–	–	8.0
	4	71.8	8.3	–	14.3	–	–	5.6

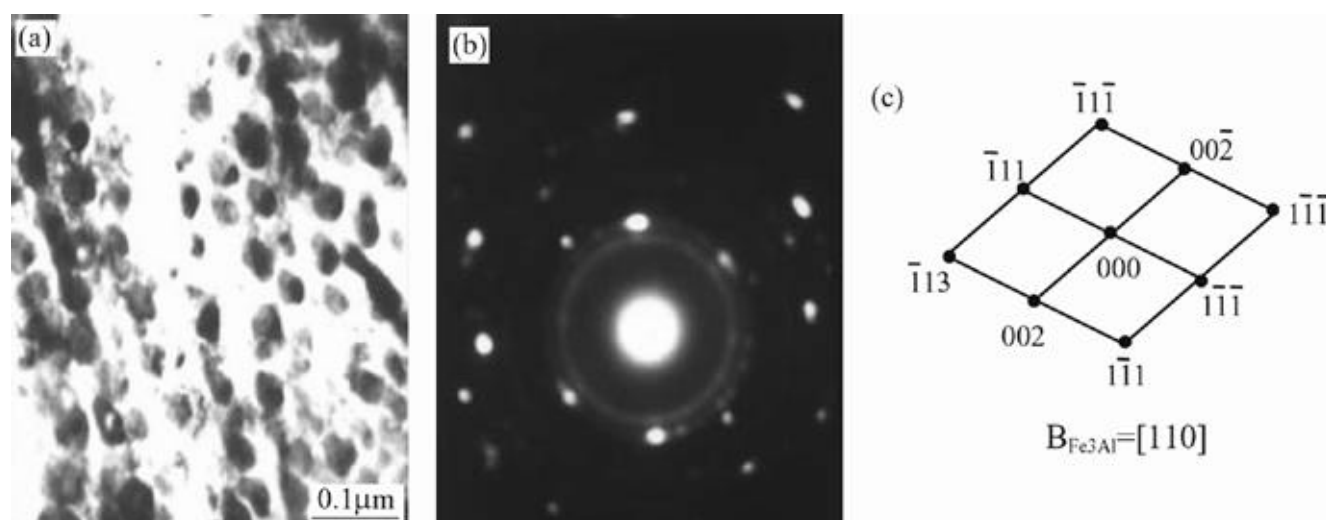


Figure 2. Fine structure of Fe₃Al in the Fe₃Al/Q235 diffusion bonding interface: (a) TEM morphology, (b) electron diffraction pattern and (c) index schematic diagram.

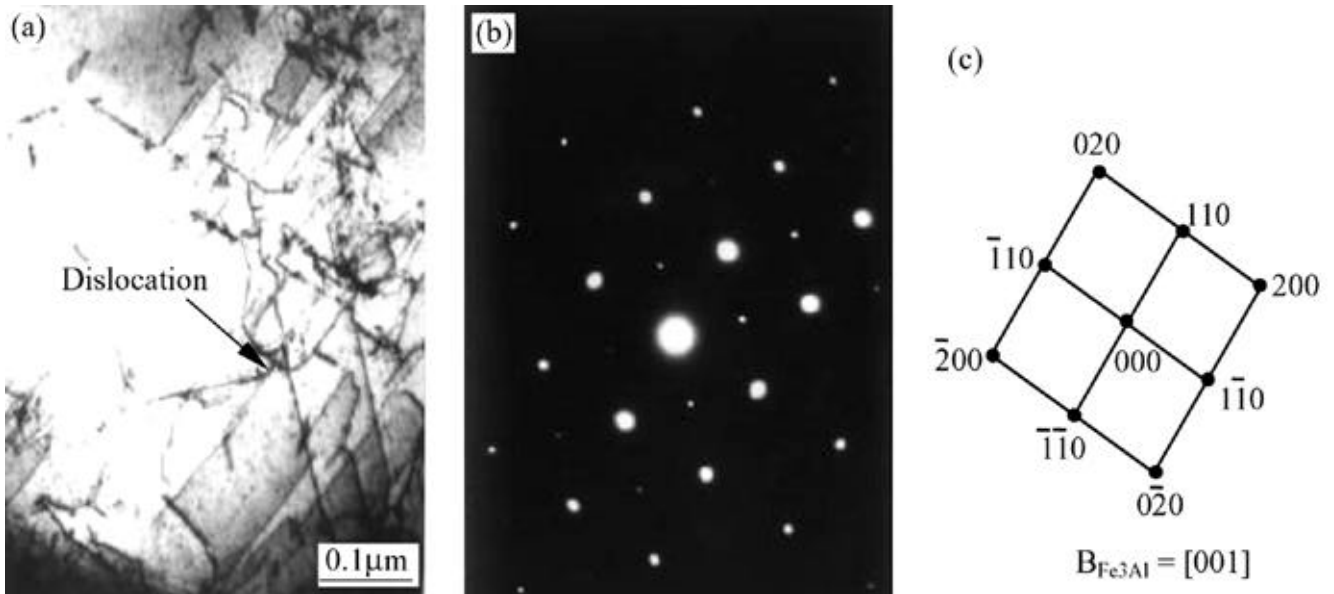


Figure 3. Fine structure of Fe_3Al from $[001]$ orientation in the $\text{Fe}_3\text{Al}/\text{Cr18-Ni8}$ diffusion bonding interface: (a) TEM morphology, (b) electron diffraction pattern and (c) index schematic diagram.

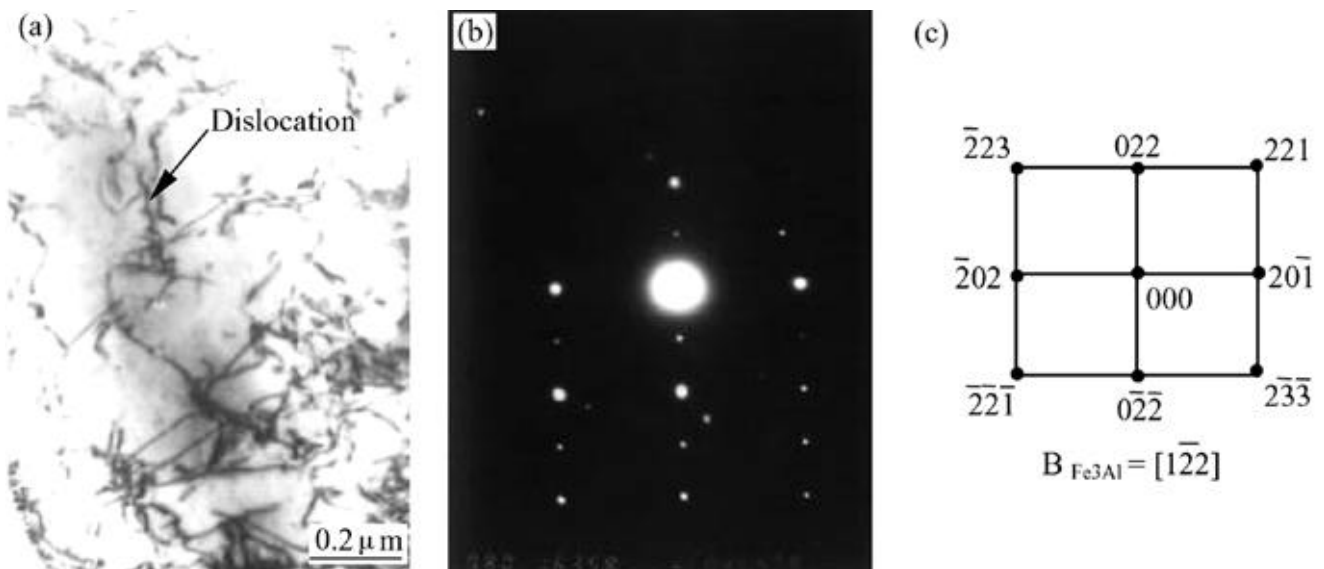


Figure 4. Dislocations in Fe_3Al from $[1\bar{2}\bar{2}]$ orientation in the $\text{Fe}_3\text{Al}/\text{Cr18-Ni8}$ diffusion bonding interface: (a) TEM morphology, (b) electron diffraction pattern and (c) index schematic diagram.

4. Conclusions

The diffusion of Cr and Ni at the interface is favourable to improve the strength of the interface between Fe_3Al and steel. The width of $\text{Fe}_3\text{Al}/\text{Cr18-Ni8}$ interface transition zone is $35\ \mu\text{m}$. It is larger than that of $\text{Fe}_3\text{Al}/\text{Q235}$ interface transition zone ($29\ \mu\text{m}$) under the condition of $1333\ \text{K} \times 60\ \text{min}$ and a pressure of $15\ \text{MPa}$. The second phase rich in Cr and Ni distributes dispersively in Fe_3Al formed at the $\text{Fe}_3\text{Al}/\text{Cr18-Ni8}$ interface to cause dislocation couples with different spacing and even dislocation

network. This is favourable to improve the joint strength at the $\text{Fe}_3\text{Al}/\text{Cr18-Ni8}$ interface.

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