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# INVESTIGATION OF CRYSTALLOGRAPHIC SLIP IN POLYCRYSTALLINE Fe<sub>3</sub>Al USING SLIP TRACE MEASUREMENT AND MICROTEXTURE DETERMINATION

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Abstract—An intermetallic Fe<sub>72</sub>Al<sub>28</sub> alloy (doped with Cr, Zr, Mo, and C) with an imperfectly ordered B2 crystal structure was rolled at 830–860 K to  $\epsilon = 20\%$ . To investigate crystallographic slip an etching technique was developed which allowed slip traces to be determined in grain interiors rather than at the sample surface. To derive the prevalent glide systems both the slip traces and the corresponding orientations were determined in grain scale. Three types of slip systems were identified, namely  $\{110\}\langle 111\rangle$ ,  $\{112\}\langle 111\rangle$ , and  $\{123\}\langle 111\rangle$ . However, the slip traces produced by  $\{123\}\langle 111\rangle$  systems appeared wavy and were interpreted in terms of macroscopic or effective rather than crystallographic slip. The critical resolved shear stress ratio of the slip systems involved was fitted from experiment using a Relaxed Constraints Taylor model. The best correspondence between predicted and experimentally observed slip systems was attained for a critical resolved shear stress ratio of  $\tau_{\{110\}}/\tau_{\{112\}} = 1.05/1.0$ . © 1997 Acta Metallurgica Inc.

### 1. INTRODUCTION

Intermetallic Kurnakov type iron aluminides with compositions near that of Fe<sub>3</sub>Al which are ordered to the DO<sub>3</sub> or imperfectly ordered to the B2 crystal structure increasingly attract attention owing to their good oxidation and sulphidation resistance, high strength at low and intermediate temperatures, low production costs, and low density [1–12]. These characteristics justified the considerable efforts made to establish Fe<sub>3</sub>Al based alloys as a new structural material class.

However, the advantageous property profile of Fe<sub>3</sub>Al based alloys was impaired by the lack of ductility at ambient temperature and a decrease in strength above  $\approx 870 \, \text{K}$ . Since the beginning of research in this field a rapidly increasing body of literature has thus appeared, concentrating on the improvement of room temperature ductility, strength at elevated temperatures, and corrosion resistance by the control of chemical composition and microstructure.

McKamey et al. [7, 13] showed that an enhanced content of Cr, Si, Ta, Mo, Hf, Ti, and Nb significantly increases the yield strength of Fe<sub>3</sub>Al at temperatures in excess of 870 K. Recent studies [10–14] confirmed that ternary additions of 2–6% Cr to DO<sub>3</sub> ordered iron aluminides were most effective in enhancing the room temperature ductility. Maximum values of 20% for elongation after fracture have been recently reported for Cr doped Fe<sub>3</sub>Al by Sun et al. [12]. The results were interpreted in terms of the anti-phase boundary (APB) energies, change of fracture modes, and slip characteristics.

It was also shown that substitutional and interstitial additions affect the stability of the ordered phases involved. Depending on the chemical composition, the phase transformation  $B2 \rightarrow DO_3$  $(T \approx 813 \text{ K})$  can either take place directly or via decomposition into an ordered and a disordered phase, i.e.  $B2 \rightarrow (B2 + A2) \rightarrow (A2 + DO_3) \rightarrow DO_3$ [7, 15, 16]. The formation of the DO<sub>3</sub> phase is typically very slow (DO<sub>3</sub> ordering of pure Fe<sub>3</sub>Al at 800 K requires about 1 week) [7, 10-12]. In the presence of ternary additions like Cr, Mo, Zr, Hf, Nb, C, and B, it may even be considerably delayed [10-12]. Due to such slow transformation kinetics, recent studies have successfully focused on the partial stabilization of the imperfectly ordered B2 phase at room temperature through the employment of a so-called thermomechanical B2 treatment [10-12].

Owing to the optimization of both constitution and resulting microstructure, Fe<sub>3</sub>Al based alloys now have improved mechanical and corrosion properties which are equivalent or even superior to those of many steels. This applies especially to their strength-to-weight ratio. Consequently, Fe<sub>3</sub>Al based alloys are even considered nowadays as potential candidates for applications where ferritic stainless steels are in use [7].

Despite such progress in improving many technical aspects of Fe<sub>3</sub>Al, the mechanisms that are responsible for both the strength and ductility observed are not yet sufficiently understood. This applies especially to the selection of active slip systems during large strain deformation. In fact, a quantitative analysis of how an externally imposed strain state is accomplished by crystallographic slip represents an important

approach to understanding the mechanical properties of polycrystalline Fe<sub>3</sub>Al. Hence, this paper deals mainly with the investigation of crystallographic slip in an imperfectly B2 ordered intermetallic Fe<sub>72</sub>Al<sub>28</sub> alloy, doped with Cr, Zr, Mo, and C.

# 2. SLIP SYSTEMS IN Fe<sub>3</sub>Al

Most direct and indirect studies of prevalent slip systems agree that dislocation movement in Fe<sub>3</sub>Al based alloys primarily takes place on those glide systems that are typically activated in non-ordered body centered cubic (b.c.c.) alloys. Among these, it appears that in particular {110}<111> and {112}<111> slip systems contribute to the accomplishment of externally imposed strains [7].

Two direct observation techniques were employed to obtain a more quantitative experimental picture of the predominant slip systems, the mechanisms of dislocation generation and movement, and of corresponding reactions in single- and polycrystalline Fe<sub>3</sub>Al. First, slip trace investigations were conducted on the sample surface using optical microscopy. Second, direct dislocation observations were carried out employing transmission electron microscopy (TEM) [7, 14, 15, 17–21].

Thorough slip trace investigations on polished sample surfaces were conducted by Leamy et al. [22], Marcinkowski and co-workers [23-25], and Taoka and co-workers [26, 27] on polycrystals. These authors concentrated on the investigation of slip markings mainly with respect to their geometrical appearance (wavy or straight). However, they did not carry out complementary microtexture measurements for the determination of the respective glide systems. They reported that samples containing either 25 or 30.6 at.% Al and compressed at 300 K revealed diffuse slip markings which gradually sharpened up to a plastic strain of  $\epsilon = 20\%$  (samples with 25 at.% Al) and  $\epsilon = 30\%$  (samples with 30.6 at.% Al), respectively. Based on results achieved from samples with different chemical compositions and long range ordering, the two types of slip trace were interpreted in terms of deformation via ordinary (wavy) or superlattice (straight) dislocations. Although these slip trace investigations were helpful in assessing the influence of order and disorder on slip, they did not provide information about glide crystallography during polycrystal deformation in grain interiors.

Complementary slip line investigations by Schröer et al. [15] on Fe<sub>70</sub>Al<sub>30</sub> single crystals, where the orientation of the surface plane inspected was known, revealed that both  $\{110\}$  and  $\{112\}$  glide planes had been activated depending on the temperature and orientation of the samples. At low temperatures up to 689 K only straight  $\{110\}\langle 111\rangle$  slip traces were observed. The activation of  $\{112\}\langle 111\rangle$  slip was especially noticed at temperatures above 740 K. At higher temperatures ( $T \ge 870$  K) Schröer et al. [15] reported the occurrence of a slip plane that was

inclined by a 'few degrees away from {110} towards {112}'. This observation hints at the activation of {123} slip planes. Such single crystal experiments were important for extracting information about the critical resolved shear stresses (CRSS) of  $\{110\}\langle 111\rangle$ and {112}(111), the slip asymmetry especially of {112}<111>, and the resulting orientation dependence of the activation of slip. In this context especially the anomalous course of the temperature dependence of the CRSS of Fe<sub>3</sub>Al is of relevance [4, 28]. Single crystal experiments with various starting orientations suggested that between 270 and 550 K a negative, between 550-750 K a positive, and above 750 K again a negative temperature dependence of the CRSS occurs. Depending on the chemical composition of their alloys other authors reported temperatures. transition somewhat different Although these results provide essential insight into the deformation process of Fe<sub>3</sub>Al, they cannot unambiguously be used for the interpretation of polycrystal deformation, where severe strain constraints already enforce multiple slip at low strains.

TEM investigations, which were carried out on deformed single- and polycrystalline material with a broad range of chemical composition and deformation temperature, showed that most Fe<sub>3</sub>Al dislocations contained Burgers vectors parallel to (111) [7, 14, 15, 17–21]. Furthermore, it was observed that dislocations moved on {110} and {112} slip planes. Similar results were reported from in situ TEM experiments, where dislocations were observed during rather than after straining of Fe<sub>70</sub>Al<sub>30</sub> [20]. Most dislocations observed in these experiments had screw rather than edge character. This was attributed to the fact that the mobility of the edge dislocations exceeds that of the screw dislocations, which allows them to leave the TEM foils. Dislocation glide on {110} slip planes was observed over a broad temperature interval, including regimes of both positive and negative temperature dependence of the CRSS. At the temperature of the peak strength,  $T_{\rm p} \approx 750$  K, slip activity was observed on  $\{112\}$  slip planes. At lower temperatures (573 K < T < 693 K) {112} planes were observed to act as cross slip or double cross slip planes.

The strength anomaly observed is commonly discussed in terms of the properties of the dislocations involved. For this purpose the results of the above cited experimental studies on prevalent slip systems has to be complemented with corresponding experimental [7] and theoretical [17–19] investigations about APB energies and dislocation core effects. In long range ordered crystals, superlattice rather than ordinary lattice dislocations are the elementary carriers of plastic flow. While in the B2 phase typically twofold-dissociated, i.e. paired  $a_{\rm B2}/2\langle 111\rangle$  dislocations prevail, in the DO<sub>3</sub> phase even fourfold-dissociated dislocations appear. In the latter case the  $a_{\rm B2}/2\langle 111\rangle$  pairs combine to form quadruplets, or double pairs, respectively, consisting of four partial

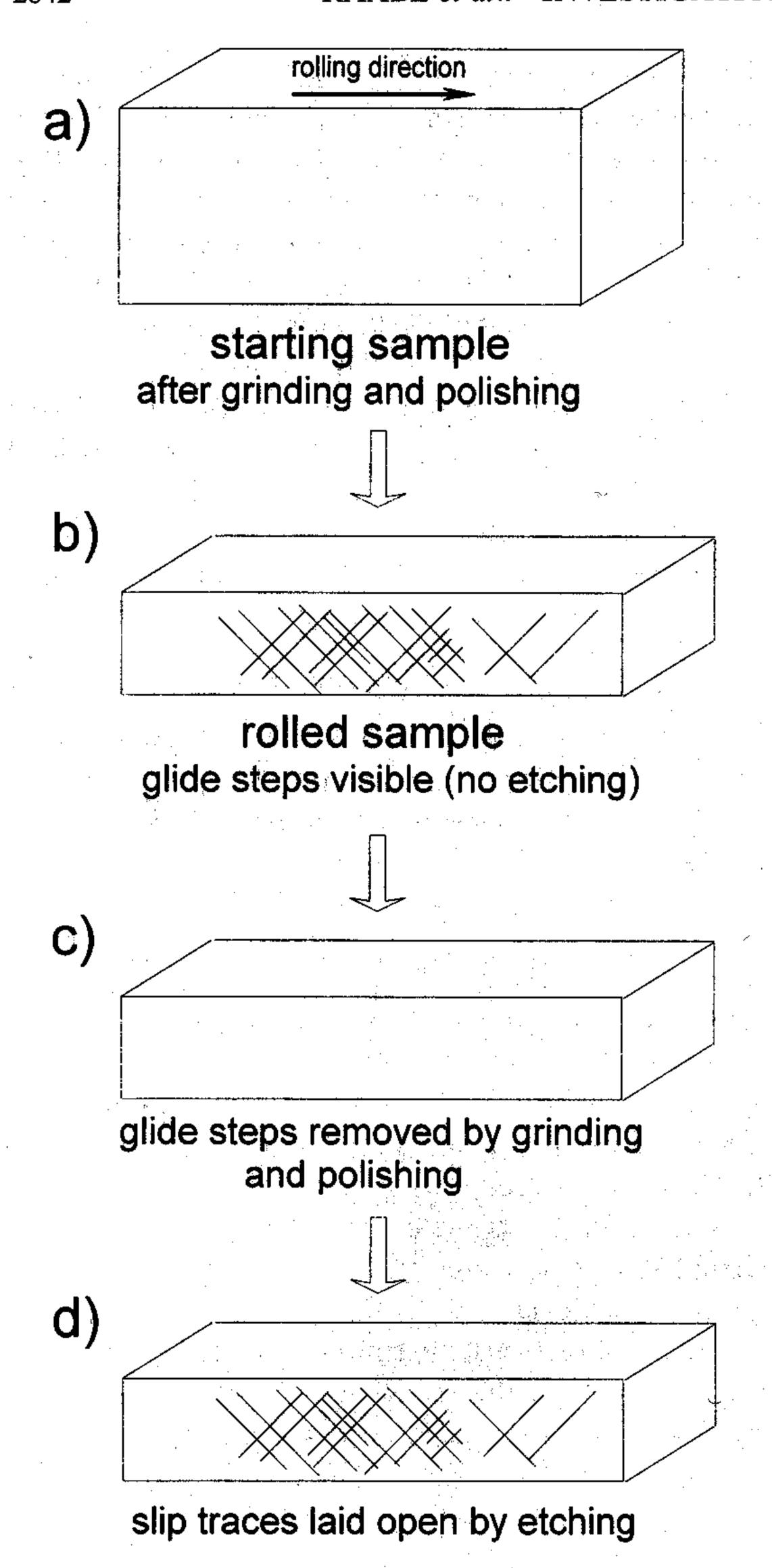


Fig. 1. Schematical explanation of the experimental procedure which was employed to prove identity of conventional surface glide steps and slip traces obtained via etching. (a) Specimen prior to rolling, (b) rolled sample with visible glide steps on its surface, (c) glide steps entirely removed by grinding and polishing, (d) etching the sample in pure HCl produced slip traces shortly before they were removed.

preparation of specimens by grinding or cutting after rolling necessarily leads to the removal of the initial glide steps on the sample surface. Nevertheless, in order to investigate slip markings in the sample interior we used a simple etching technique which allowed slip traces to be made visible after cutting and polishing, i.e. after removing the slip traces on the specimen surface. The first type of marking is hereafter referred to as (surface) glide steps and the second one as bulk slip traces.

At first it was demonstrated that the markings obtained after etching were really identical to bulk slip traces. For this purpose some specimens were cut, ground, and polished prior to deformation [Fig. 1(a)].

After rolling the surface glide steps (longitudinal sections) were investigated without any etching treatment [Fig. 1(b)]. Figures 2(a) and 3(a) clearly show that up to three different groups of such slip markings could be identified within most grains. Often some groups of glide step appeared much more distinct than others. While some slip lines were straight or only slightly curved, especially in the vicinity of grain boundaries, others appeared wavy, indicating frequent cross slip. After a thorough analysis of these surface glide steps, the samples were again ground and polished until all slip markings were entirely removed [Fig. 1(c)]. In order to demonstrate the identity of the original glide steps exposed via polishing and rolling and the bulk slip traces revealed via etching, the polished samples were then etched in pure HCl for 1-3 min. To study deformations in excess of 20%, a solution of 10 ml H<sub>2</sub>O<sub>2</sub> and 1 ml HF (30 s) was used. This etching procedure indeed led again to the appearance of the slip lines which were removed shortly before and thus demonstrated their identity with the initial glide steps generated after rolling on the polished sample surface [Fig. 1(d)]. Although Figs 2(b) and 3(b) show that after etching some slip markings appeared weaker and others clearer than before, a maximum information was nevertheless extracted by using sufficiently high resolution in an SEM to allow a thorough analysis of the faint slip traces also.

The fact that slip traces can be conveniently made visible in Fe<sub>3</sub>Al by etching offers two main experimental advantages. First, it allows crystallographic slip to be investigated not only at the sample surface but also in the sample interior. This point is of importance for a proper analysis of the results considering surface artifacts. although etching produces slip traces that are somewhat less sharp than those exposed without etching, it is more sensitive and sometimes lays open a larger number of slip markings than the conventional method. For this reason the slip traces were exclusively studied using the etching rather than the glide step technique. In order to make sure that the investigated slip traces were generated by crystallographic slip or microbands rather than by non-crystallographic shear bands, the investigations were carried out only in weakly deformed specimens  $(\epsilon = 20\%)$ .

The textures of the specimens were examined in grain scale exploiting the anisotropy of electron back scatter diffraction in the SEM according to the method of Venables and Harland [33]. Details of the latter technique are explained elsewhere [34]. The active slip systems were derived on the basis of the experimentally observed slip traces and the corresponding local orientations. In order to obtain unambiguous solutions we skipped the analysis of special grain orientations where {110}, {112}, and {123} planes generated similarly oriented slip traces with a mutual inclination of less than 5°.

#### 3.3. Fitting procedure

For fitting the CRSS ratio of the slip systems involved, the experimentally observed prevalent glide systems were compared to those predicted by a Taylor Relaxed Constraints model (relaxation of 613 and  $\epsilon_{23}$  where 1 = rolling direction, 2 = transverse direction, and 3 = normal direction) [29-32]. The micrographs with the clearest slip markings were used for calibration. The ratio of the CRSS was successively changed in the Taylor simulation until a maximum agreement between the experimentally determined and the predicted slip system was achieved. In the present case the CRSS represents a somewhat artificial value. However, comprising various drags like the nucleation of superlattice dislocations, core effects, order effects, and the Peierls stress, it represents a rough measure of whether particular glide systems substantially contribute to slip or not. The increase in the CRSS with increasing

strain was assumed to be identical for all types of operative slip system. This assumption implies identical latent hardening for all slip systems and thus an isotropic growth of the yield surface. For the fitting procedure the externally imposed strain state during warm rolling was idealized as an ideal plane strain state. Such simplification seems justifiable since all investigated grains were located close to the center layer. The predications were carried out considering  $\{110\}\langle 111 \rangle$  and  $\{112\}\langle 111 \rangle$  slip systems. As will be discussed later the consideration of {123}(111) slip seemed to be less meaningful due to the non-crystallographic character of the corresponding slip traces. As was shown in previous texture simulations for non-ordered b.c.c. alloys [35, 36] the asymmetry of {112}(111) slip seems to be of minor importance and was hence neglected. In the Taylor model there is no need to explicitly distinguish deformation via ordinary or superlattice dislocations. This is due to the fact that in Taylor theory the glide systems are

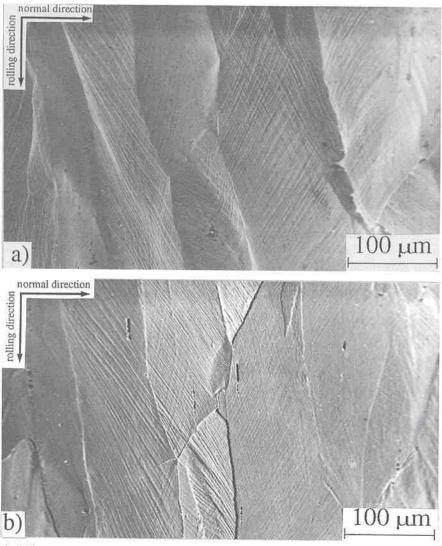


Fig. 2. Different groups of slip trace at surface of a rolled sample: (a) after rolling without etching, (b) after rolling, grinding, polishing, and etching.

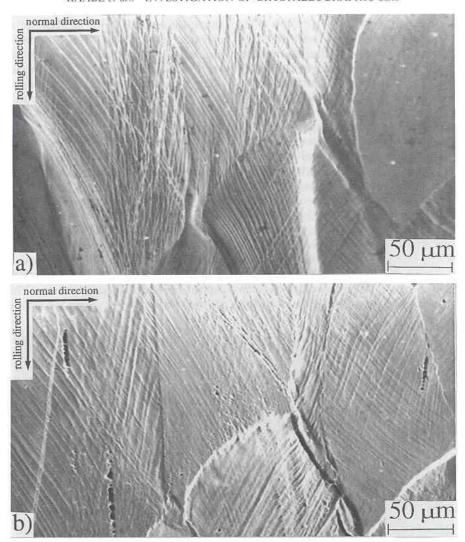


Fig. 3. Different groups of slip trace at surface of a rolled sample: (a) after rolling without etching, (b) after rolling, grinding, polishing, and etching.

reduced to their geometrical function, i.e. the entire framework of dislocation dynamics is completely neglected.

#### 4. RESULTS

Warm rolling ( $\epsilon = 20\%$ ) was carried out within the temperature range 830–860 K. The rolled specimens were sectioned parallel to the transverse plane, polished, and etched. Both the slip traces and the microtexture were determined in the interior of 56 different grains. In each crystal up to three groups of differently oriented slip traces were analysed. In 93 cases the observed slip markings could be unambiguously associated to  $\{110\}$ ,  $\{112\}$ , or net  $\{123\}$  planes.

In 45% of all investigated grains {110}<111> slip was identified. Figure 4 shows two grains in a longitudinal section. Each grain reveals two groups of straight slip trace. In both grains all traces could be attributed to {110}<111> glide systems. In many grains the occurrences of straight slip traces were due to {110} glide. The slip trace spacing varied between

1 and  $15\,\mu\text{m}$ . In 46% of all examined grains  $\{112\}\langle111\rangle$  slip was observed. In the grains depicted in Fig. 5 both  $\{112\}\langle111\rangle$  (grain A) and  $\{110\}\langle111\rangle$  (grain B) slip traces were identified. Both groups revealed straight slip markings and a homogeneous slip line distribution throughout the grains. The average slip trace spacing was again between 1 and  $15\,\mu\text{m}$ .

Of all inspected grains 77% contained net \$\{123\\<111\}\$ traces. Figure 6 shows such a grain, which is characterized by an inhomogeneous distribution of the slip traces. From this micrograph three main observations can be extracted. First, two different groups of slip markings can be identified. Position A reveals traces from intersecting \$\{112\}\$ and \$\{123\}\$ planes. At positions B and C only \$\{123\}\$ planes are apparent. Second, the slip markings reveal both straight and wavy shapes. Third, not only the active slip systems but also the average slip trace spacings vary. The inhomogeneity of the slip traces was also apparent in the microtexture which revealed con-

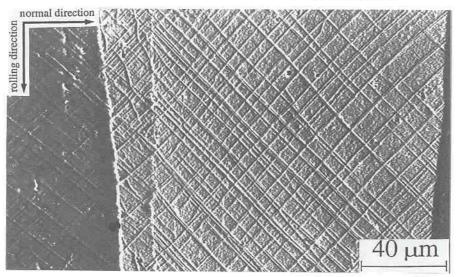


Fig. 4. Two grains in longitudinal section. Each grain contains two groups of straight slip trace.

siderable local misorientations in the grain interior. In the longitudinal section shown in Fig. 7 a large contribution of very wavy slip planes can be observed. For deriving net planes from the wavy traces their net direction rather than their local tangent direction was used. For this reason the derived planes should not be referred to as slip planes in a crystallographic sense. In most cases these wavy traces corresponded to net {123} planes.

The experimental data are summarized in Table 2 considering {110}<111> (crystallographic), {112}<111> (crystallographic), and {123}<111> (noncrystallographic) slip systems. In the columns a '+' indicates the occurrence and a '-' the absence of the corresponding slip system. The sharpness of an etched slip trace generated by a particular slip system does not necessarily reflect its quantitative contribution to the total shear. Consequently, the analysis

is confined to the qualitative statements 'slip system is active (+)' or 'slip system is not active (-)'.

#### 5. DISCUSSION

#### 5.1. Orientation dependence of the experimental data

Figure 8(a) shows the symmetrized {100} pole figure of all investigated grain orientations as determined by EBSD. The pole figure shows a weak texture with a slight pronunciation of an orientation close to the cube component having a maximum pole density of only 2.3 times random. Consequently, there is no need to take texture into account to avoid an over- or underestimation of certain slip systems.

A summary of the observed slip activity reveals that 45% of the grains show slip on {110}, 46% on {112}, and even 77% on net {123} planes. In order to investigate whether {110} or non-{110} slip

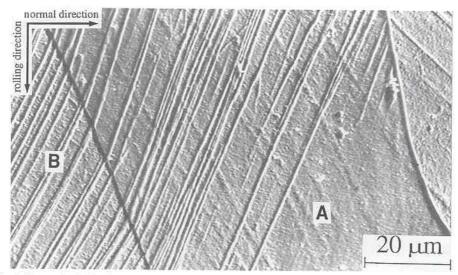


Fig. 5. Two grains in longitudinal section. Grain A reveals straight {112}<111> slip traces. Grain B shows straight {110}<111> slip traces.

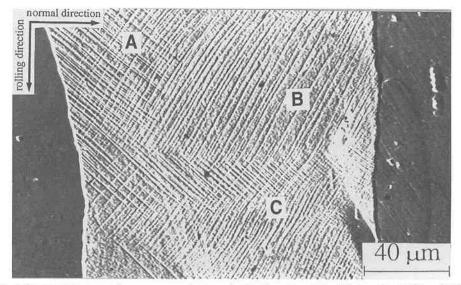


Fig. 6. Grain with a more inhomogeneous slip trace distribution. At position A interesting {112} and {123} slip planes were determined. At positions B and C {123} slip planes prevail. The slip markings reveal both straight and wavy shapes.

depends on grain orientation one has to examine the orientations in which the respective type prevails. Figure 8(b) shows the symmetrized {100} pole figure of grains in which exclusively {110} slip planes were identified. Figures 8(a) and (b) are practically identical. That means {110} slip was not confined to particular grain orientations. Figure 8(c) shows the symmetrized {100} pole figure of grains in which exclusively non-{110} slip planes were identified. Again, it does not reveal any substantial difference to the other pole figures [Fig. 8(a), (b)]. This similarity between the various pole figures shows that within the statistical limit of the present study there are no special grain orientations which promote the activation of {110} or non-{110} slip.

#### 5.2. Wavy and straight slip traces

In the longitudinal sections (Figs 4–7) both straight and wavy slip traces were observed. Traces which were generated by crystallographic slip on {110} or {112} planes were mostly straight. In contrast, slip markings which were attributed in the analysis to net {123} planes often had a wavy rather than a straight shape (Fig. 7). The latter result means that the observed net traces did not necessarily follow crystallographic {123} slip planes. In a first approach, the occurrence of such non-crystallographic slip can be understood in terms of combined {110} and {112} slip portions in atomistic scale [37–41].

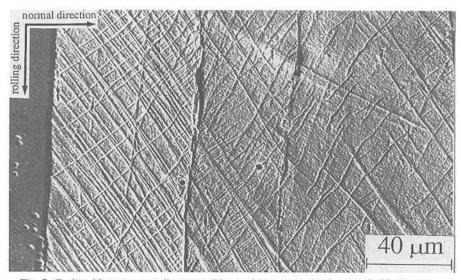


Fig. 7. Grain with very wavy slip traces. The markings correspond to net {123} slip planes.

Table 2. Results regarding local slip activity obtained on hot rolled Fe<sub>3</sub>Al specimens, deformed at 830-860 K to  $\epsilon = 20\%$  thickness reduction

			Experimental		
	~ Crystal orientation		{110}	Experiment {112}	nai Net {123}
	$\{20-98-2\}\langle -30-5-95\rangle$		1	()	- (00 (120)
. <b>.</b> .	$\{-98-7\ 19\}\langle 1-95-31\rangle$		<b>+</b> 		<del>-</del>
	$\{-4\ 41-91\}-96-27-7$			<u> </u>	<u> </u>
	${94-14-31}\langle 21\ 96\ 20\rangle$		<u>!</u> .	+	
	$\{-38 \ 32 - 87\}\langle 92 \ 3 - 40\rangle$				<del>_</del>
	$\{-94\ 35\ 6\}\langle -10-10-99\rangle$		·		· · · · · · · · · · · · · · · · · · ·
	${29 - 95 \ 10}(79 \ 18-59)$		· _ ·	+	<u> </u>
	{100-0-3}<0 100-0>		· · ·	· •	+
	$\{80-36, 48\}\langle -24-93-29\rangle$		—	+	+
•	$\{-91\ 39\ 14\}\langle -30-85\ 43\rangle$		, j <del></del> ,	+	<del> </del>
	$\{-25\ 5\ 97\}\langle 91\ 35\ 22\rangle$	:`	+	· · · —	+
	$\{-95-28,15\}\langle -9-21-97\rangle$		_	<u> </u>	+
	$\{-94, 15, 30\}\langle -10, -98, 18\rangle$		: <u> </u>	· · · · · · · · · · · · · · · · · · ·	+
•	$\{-96\ 24-10\}\langle 4-25-97\rangle$		+		+
	$\{-94-24-25\}\langle 17-94\ 29 \rangle$		+	<del></del>	
	$\{1-90-43\}\langle 95, 14-27 \rangle$		. ·	<del>+</del>	+ .
•	$\{-15 \ 32 - 94\} \langle 97 \ 25 - 7 \rangle$		_	+	+
	$\{-36-64-67\}\langle 28-77.58 \rangle$		+	. — ·	+
	$\{19-98-4\}\langle -64-10-76\rangle$		+	+	+
	$\{-93\ 19\ 33\}\langle 1-85\ 52\rangle$		+	· -	<del>.</del> —
	$\{-85-12-51\}\langle -4-95\ 30 \rangle$		+	+	+
	${39 - 86 \ 32} < -47 \ 12 \ 88$		+ .	: <del>  </del>	+
	$\{22\ 33-92\}\langle 97-5\ 22\rangle$		_	<del></del> .	+
	$\{-95-32\ 6\}\langle 31-94-17\rangle$			<del>-</del>	+
	$\{-31\ 19-93\}\langle -44-90-3\rangle$	· .	. <del></del>	+	+
	$\{-98 \ 6 \ 19\}\langle 2 - 92 \ 39\rangle$		+	<del></del>	· <del>- · ·</del>
	$\{-18-3-98\}\langle 96\ 22-18\rangle$		+	<del>-</del>	<del></del>
, y	$\{-92\ 24\ 29\}\langle -26\ 16-95\rangle$	· · · · · · · · · · · · · · · · · · ·	<del>-</del>	<del>.</del> +	+
	$\{14\ 13 - 98\}\langle -31\ 95\ 9\rangle$		+	<del></del>	· <del></del> · .
	$\{98 - 197\}\langle 1897 - 17\rangle$			+	<u></u>
	$\{-87 - 39 - 31\}\langle 13 \ 44 - 89\rangle$	· · :	+	+	+
	$\{55, 56 - 61\} \langle 70 - 71 - 2 \rangle$	· :	· · · · · · · · · · · · · · · · · · ·	+	· <del>· ·</del>
	$\{86\ 35 - 39\}\langle 12\ 60\ 79\rangle$			+	+
	$\{-39, 2-92\}\langle -0, 100, 2\rangle$		+	en e	<del>-</del>
	$\{-13, 28 - 95\} \langle 97 - 16 - 17 \rangle$		. <del></del> .		+
	$\{13.91 - 39\} \langle -95.23.21 \rangle$	· ·		+	+
	$\{46 - 89 \ 2\} \langle -33 - 15 \ 93 \rangle$		+	+	+
	$\{-27 - 96\ 1\}\langle 31 - 10 - 95\rangle$ $\{28 - 87\ 41\}\langle 91\ 9 - 41\rangle$	•		+	+
	$\{96 - 10 \ 25\} \langle 22 - 28 - 94 \rangle$		<del>-</del> .	+	+
	$\{-7.24 - 97\}\langle 16 - 96 - 25\rangle$		_		+
	$\{-8.83 - 55\}\langle -52.43.73\rangle$				+
	$\{92 - 18 - 35\}\$		 I	_	+
	$\{-38 \ 22 - 90\} \langle 84 \ 49 - 23 \rangle$	• •	· +		+
	$\{-32\ 22 - 92\}$ \(\lambda 49 - 23\rangle		<del>" -</del> 1		
	$\{-67, 41 - 62\}\langle -8, 79, 60\rangle$	· · .	, <del>†</del>	+	+
•	$\{-37\ 30 - 88\} \langle 91\ 28 - 30 \rangle$	•	<del>_</del> .		· · · · · · · · · · · · · · · · · · ·
	$\{-6.97.23\}\langle -95.2-32\rangle$	·	. <b>T</b>	<b>+</b>	+
	$\{-6.7 - 100\}\langle -96.26.8 \rangle$			. †	<del>- -</del> •
	$\{28\ 44 - 85\}\langle -42\ 85\ 31\rangle$		<del>-</del>	<del>-</del>	+
	$\{96-28-9\}\$ $\{25.93-27\}$	:	+	<u> </u>	+
	$\{57 - 47 - 67\}\$			+	<u> </u>
	$\{-46.86 - 23\}\langle -53 - 6.85 \rangle$		<del>†</del>		+
	$\{-77-52-37\}\langle 36-83,42 \rangle$				+
	$\{97-8-23\}\langle -9 77-63\rangle$	· ·	<b>-fr</b> **	+	+
	$\{9.99.15\}\langle -91.2.42 \rangle$			; <b>+</b>	+
· · · · · · · · · · · · · · · · · · ·	(~ ~ ~ ~ ~ ) \	,	<b>十</b> .		+

It is suggested that in some grains slip follows non-crystallographic planes with the maximum resolved shear stress rather than crystallographic {110} and {112} slip planes with a lower shear stress. This assumption is underlined by the fact that in grains where net {123} traces dominated, a maximum orientation factor close to 0.5 occurred on the corresponding pseudo {123}(111) system. This observation shows that in some grains the slip geometry reflects quasi-continuum rather than strictly crystallographic character.

Instead of this atomistic interpretation of non-crystallographic slip one could also think of a simple

cross slip mechanism which is not necessarily confined to the atomistic scale. In recently published in situ TEM studies [20] it was observed that most dislocations in Fe<sub>3</sub>Al had screw rather than edge character. This result was in accord with previous studies [42, 43]. The lack of edge dislocations was interpreted in terms of their high mobility, which allows them to leave the TEM foils [20]. However, in the bulk sample the contribution of edge type dislocations might be larger than suggested by the TEM studies. The low mobility of the screw dislocations was attributed to the non-planar spreading of their cores [20, 37–41]. This observation

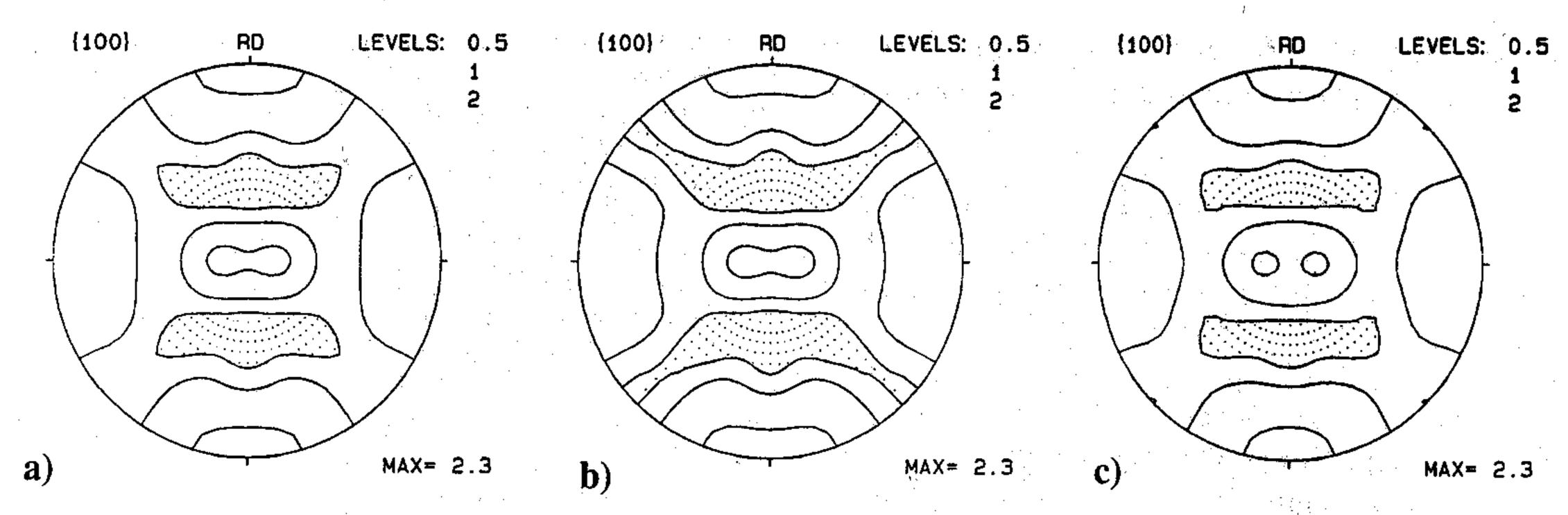


Fig. 8. Symmetrized {100} pole figures as determined from EBSD data. (a) Texture of all investigated grains, (b) texture of grains in which {110} slip planes were exclusively identified, (c) texture of grains in which non-{110} slip planes were exclusively identified.

would support the assumption of frequent cross slip. In the in situ TEM study [20] indeed {112} planes were frequently observed to act as cross slip or double cross slip planes for the {110} slip systems. However, the direct observation of slip on crystal {123} planes was not confirmed. These findings are in good accord with previous investigations on pure b.c.c. metals [37–39, 41]. On the basis of the present SEM experiments (resolution up to 20.000 × ) it cannot be decided whether non-crystallographic slip has to be interpreted in terms of alternating {110} and {112} slips in atomistic scale [e.g. 37-40] or in terms of simple cross slip [41]. However, one may conclude that in accord with previous TEM results on Fe<sub>3</sub>Al [20, 42, 43], slip on crystallographic {123} slip planes is not observed.

# 5.3. Activation of {110} and {112} slip planes

The generation and movement of superlattice dislocations in B2 and DO<sub>3</sub> ordered alloys is mainly determined by two properties. First by the dislocation core structure [17-19], which determines the mobility of superlattice dislocations, and second by the energy of the APB, which affects both their generation and interaction [5, 6]. The dislocation core of superpartials in the B2 lattice is similar to the core of screw dislocations in the b.c.c. lattice. Vitek [38], Wütherich [39], Duesbery [37] investigated the properties of screw and non-screw type dislocations in the b.c.c. lattice in atomistic simulations both with and without an externally imposed stress. They showed that under external stresses the non-planar core of the screw dislocations led to a net path consisting of atomistic steps on  $(\overline{1}01)$ ,  $(0\overline{1}1)$ , and  $(\overline{1}10)$ . The properties of screw dislocations with (111) Burgers vector in ordered B2 alloys were correspondingly simulated by Takeuchi [40]. He observed that the cores of the partial superlattice screw dislocations were very similar to those of the ordinary b.c.c. lattice, i.e. they extended on three intersecting {110} planes where they create stacking faults [17–19, 40]. The lowest energy arrangement of the superpartial-APB complex was a configuration where the fractional dislocations in the cores were oriented in such a way that they did

110} layer faults extended on the {112} planes at their tips. In his simulations Takeuchi [40] used two different potentials to investigate the influence of both a high and a low energy APB. For the low energy case, which corresponds to the present alloy, he observed that the dislocations moved in a similar manner as in a non-ordered b.c.c. lattice.

According to Umakoshi and Yamaguchi [17–19] the energy associated with the APB on {110} is lower than on {112}. However, they mentioned that the dislocation core is probably more extended on {110} glide planes. Prior to its displacement the complexly dissociated dislocation core must first be converted into a planar configuration which contributes to the glide resistance. In contrast superpartials on {112} planes create complicated multiplayer faults, which suggests that their dislocation cores are very narrow. Although they have a higher APB energy than dislocations on {110}, their core can easily transform into a glissile configuration prior to its movement. Altogether this is assumed to cause a somewhat lower glide resistance on {112} compared to {110} glide planes.

These structural arguments for the dominance of {112} slip planes are supported by the fitting procedure. The best accord between the Taylor prediction and experiment is obtained when the CRSS of {110} \langle 111 \rangle exceeds that of {112} \langle 111 \rangle by 5%. This CRSS ratio is smaller than suggested by previous values obtained by comparing macroscopic deformation textures with Taylor type simulations [44, 45]. It suggests that the tendency for {112} slip is not to be interpreted in terms of the APB energy but in terms of dislocation core effects. The contribution of the APB energy to the increase in energy that is required to generate new {112} \langle 111 \rangle dislocations seems to be of minor importance.

### 6. CONCLUSIONS

An intermetallic Fe<sub>72</sub>Al<sub>28</sub> alloy was warm rolled at 830–860 K to  $\epsilon = 20\%$  thickness reduction. The active crystallographic slip systems were derived by

determination of slip traces in grain interiors and the corresponding grain orientations. The main conclusions are:

- In the experiments both {110}(111) and {112}(111) slip systems were identified with a similar slip activity. Corresponding slip traces typically had a straight shape.
- Slip on {123}<111> slip systems appeared as macroscopic or effective rather than crystallographic slip. Corresponding slip traces typically had a wavy shape. It was suggested that wavy {123}<111> slip can be explained by combined {110}<111> and {112}<111> slip.
- The critical resolved shear stresses of the involved slip systems were fitted from experiment using a Taylor Relaxed Constraints model. The best correspondence between predicted and experimentally observed slip systems was attained for a critical resolved shear stress ratio of  $\tau_{\{110\}}/\tau_{\{112\}} = 1.05/1.0$ .

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