THE DEVELOPMENT OF ROLLING TEXTURES IN LOW-CARBON STEELS

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The crystallite orientation distribution analysis was applied to the study of the development of the rolling textures in low-carbon steels. It was found that the constraining effect of the grain boundary remarkably influences the rolling textures of polycrystalline iron. This enhances the crystal rotations, which would not be expected to occur in single crystals; and grains having the $\langle 111 \rangle$$\langle 112 \rangle$ orientations are forced to rotate about the $\langle 111 \rangle$ axes lying in the sheet normal direction toward the $\langle 111 \rangle$$\langle 110 \rangle$ orientations. This is the origin of the $\langle 111 \rangle$ fiber texture normally found in the rolling textures of low-carbon steels. The presence of the partial fiber texture having the $\langle 110 \rangle$ axes inclined 30 deg from the sheet normal toward the rolling direction could not be confirmed.

INTRODUCTION

The rolling textures of polycrystalline iron have been described in various ways.\textsuperscript{1-9} According to Barrett and Levenson,$^4$ these consist of two continuous sets of end orientations. The first set is a partial fiber texture with the $\langle 110 \rangle$ fiber axes in the rolling direction, while the second set is a fiber texture having the $\langle 111 \rangle$ axes in the sheet normal direction. The presence of the first set has been confirmed by all investigators. As to the second set, there have been controversies in its description.

Based on the $\langle 110 \rangle$ pole figures of cold rolled carbonyl iron, Haessner and Weik$^2$ described this in terms of a $\langle 110 \rangle$ fiber texture having the $\langle 110 \rangle$ axes inclined 35 deg from the sheet normal toward the rolling direction.

Bennewitz$^4$ found, on the other hand, that the second set of orientations could better be described as a partial fiber texture with the $\langle 110 \rangle$ axes inclined 30 deg from the sheet normal toward the rolling direction. This was designated as a "$\langle 110 \rangle$ 60 deg RD fiber texture". On this basis, he divided the development of the rolling texture into three stages.

However, the presence of the "$\langle 110 \rangle$ 60 deg RD fiber axes" is theoretically difficult to explain. Direct experimental evidence, verifying that individual grains really rotate about these fiber axes, have not been obtained so far. The presence of these fiber axes was only indirectly inferred from the pole figures, above all from $\langle 110 \rangle$ pole figures.

Generally, pole figures describe only the distribution of particular crystallographic axes in the sample coordinate system. Two crystallites, whose orientations are related to each other by a rotation about one of these axes cannot be discriminated in the pole figure. In this respect, the precise distribution of the crystallite around the $\langle 110 \rangle$ fiber axes cannot adequately be described by the (110) pole figure alone. This uncertainty is completely eliminated in the method of the crystallite orientation distribution analysis developed by Roe\textsuperscript{10-13} and Bunge\textsuperscript{14-18}.

It is the purpose of the present investigation, using this method of analysis, to study the formation of the rolling textures in low-carbon steels. For the detailed discussions on the formation of the fiber textures, it is necessary to clarify how individual grains reorient during cold rolling. To this end, several polycrystalline specimens with sharp initial textures were cold rolled, and from their orientation changes the behaviour of a grain having a specific initial orientation in a random polycrystal was indirectly deduced. The results were finally compared with the development of the rolling texture observed in a low-carbon steel having a weak initial texture.

EXPERIMENTAL PROCEDURE AND METHOD OF ANALYSIS

Chemical compositions and hot rolling finishing and coiling temperatures of the starting hot bands
are given in Table I. Steels 1 and 4 are rimmed steels, while Steel 13 is an aluminum killed steel. Steel 18 is an experimentally hot rolled electrolytic iron.

Sharp initial textures were obtained from these steels in the following way.

1. \(\{110\}\langle001\rangle\) texture: Hot bands from Steel 1 were held 5 min at 1250°C and cooled in air. After 70 pct cold reduction, specimens were annealed for 1 hr at 750°C.

2. \(\{110\}\langle110\rangle\) texture: The transverse direction of the \(\{110\}\langle001\rangle\) specimen was chosen as the rolling direction.

3. \(\{001\}\langle110\rangle\) texture: Hot bands from Steel 4 were used in the as-received condition.

4. \(\{111\}\langle110\rangle\) texture: Hot bands from Steel 13 were cold rolled 70 pct. Specimens were then heated to 700°C with a heating rate of 20 deg/hr and held 3 hrs. at 700°C.

5. \(\{111\}\langle112\rangle\) texture: The transverse direction of the \(\{111\}\langle110\rangle\) specimen was chosen as the rolling direction.

These specimens were finally cold rolled under good lubrications up to 95 pct reduction in thickness.

The crystal rotation produced by cold rolling was observed by revealing etch pits on the rolling planes of Steel 18, cold rolled 70 pct. Specimens were ground to the center of their thickness and (001) facet pits were produced. A JEOL–JSM–U3 scanning electron microscope was used for the observation.

Samples for texture determinations were chemically thinned to about 0.04 mm thickness. Using both the transmission and reflection technique, complete (200), (110) and (211) pole figures were determined. After various corrections were made normalized intensities were obtained.

The crystallite orientation distribution analysis was made with Roe's method. The crystallite orientation distribution is usually expanded in a series of generalized spherical harmonics. In the present study, the series expansion was truncated at 22nd order.

The crystallite orientation distribution takes often, in practice, negative value. As this is physically meaningless, the largest negative value on the \(\psi = 45\) deg section was regarded, in the present study, as the total possible error. This was designated by N and shown above each set of figures, which will be shown later. All the calculations were...
carried out using an IBM 360-65 computer. The program was so constructed that the $\phi = 45$ deg section and constant $\psi$ sections of the crystallite orientation distribution appeared directly on a line printer.

The locations of ideal orientations on the $\phi = 45$ deg section are shown in Figure 1. Orientations having the $\langle 110 \rangle$ axes in the rolling direction are located on the $\psi = 90$ deg line (indicated by $RD \parallel [\overline{1}0])$, while those having the $\langle 111 \rangle$ axes in the sheet normal direction are situated on the $\theta = 55$ deg line (indicated by $ND \parallel [\overline{1}1])$. Orientations having the $\langle 110 \rangle$ directions in the transverse direction appear on the $\psi = 0$ deg line (indicated by $TD \parallel [\overline{1}0])$. Locations of the members of the $\langle 110 \rangle$ 60 deg RD fiber textures are, in the range $\phi = 34$ to 45 deg, projected on the $\phi = 45$ deg section and are shown in dashed line (indicated by $[101]$ 60° from RD). For orthotropic physical symmetry and cubic crystal symmetry, the crystallite orientation distribution takes the same value for each member of the family $\{hkl\} <uvw>$. In the present study, the latter notation is adopted.

RESULTS AND DISCUSSIONS

The $\{111\} <110>$ specimen Figure 2(a) is the $\phi = 45$ deg section of the crystallite orientation distribution, showing the $\{111\} <110>$ initial texture, while Figures 2(b) and (c) are $\phi = 45$ deg sections, showing the rolling textures of this specimen at 70 and 85 pct reductions, respectively.

After 70 pct reduction, a high intensity region developed uniformly along the $\theta = 55$ deg line, Figure 2(b). This indicates the formation of the $\{111\}$ fiber texture having the $\{111\}$ fiber axes in the sheet normal direction. The $\{111\} <110>$ initial texture merged into this $\{111\}$ fiber texture. (The origin of the $\{111\}$ fiber texture seems to be found in the $\{554\} <225>$ and $\{111\} <112>$ orientations, which were present in a sufficient amount in the orientation spread of the $\{111\} <110>$ initial texture. Evidence in support of this possibility will be given later.)

After 85 pct reduction, this $\{111\}$ fiber texture did not develop further, Figure 2(c). At this stage the $\{111\} <110>$ components increased remarkably and their presence became evident again on the $\theta = 55$ deg line.

These intensity changes of the $\{111\} <110>$ orientations suggest that grains having the $\{111\} <110>$ initial orientations were stable up to 85 pct reduction in thickness.

To explain the increase of the $\{111\} <110>$ orientations observed above 70 pct reduction, it would be necessary to assume that some part of the $\{111\} <110>$ orientations were formed from the $\{111\}$ fiber texture.

It is to be noted in Figure 2(c) that the exact position of the maximum on the $\psi = 90$ deg line is not at $\theta = 55$ deg, but at $\theta = 50$ deg. This indicates that, although the $\{111\} <110>$ orientations are stable, a slight rotation takes place about the $\langle 110 \rangle$ axes lying in the rolling direction.

Origins of the minor peaks at the $\{110\} <110>$, $\{110\} <001>$ and $\{001\} <310>$ ($\theta = 0$ deg, $\psi = 26.6$ and 63.4 deg) orientations in Figures 2(b) and (c) are not known.
The \{111\}<112> specimen Figures 3(a) and (b) are \(45^\circ\) deg sections, showing the rolling textures of this specimen at 70 and 85 pct reductions, respectively.

At 70 pct reduction, the rolling texture may be described as consisting of the \{111\}<112> initial orientations superposed on the sharp \{111\} fiber texture having the \{111\} axes in the sheet normal direction. It is to be noted that the \{111\} fiber texture derived from the \{111\}<112> initial texture, Figure 3(a), is much stronger than that derived from the \{111\}<110> initial texture, Figure 2(b). As both the specimens were taken from the same original sheet and only the rolling directions were different, the sharpness of the initial textures must be the same in these two cases. In comparison with the \{111\}<110> initial orientations, the \{111\}<112> initial orientations seem to be, therefore, more probable sources of the \{111\} fiber texture. This fiber texture can be formed from the \{111\}<112> initial orientations by the rotation about the sheet plane normal.

At 85 pct reduction, the \{111\}<112> component lost its prominence completely, while the \{111\} fiber texture showed further development and the presence of the \{111\}<110> component became distinct on the \(\theta = 55^\circ\) deg line, Figure 3(b). These orientation changes can be explained again in terms of rotation about the \{111\} axes lying in the sheet normal direction; and the grains having the \{111\}<112> initial orientations appear to become more stable as the \{111\}<110> orientations are approached as a result of this rotation.

Among the members of the \{111\} fiber texture, the \{111\}<110> orientations are most distant from the \{111\}<112> orientations. If a further rotation were to take place about the \{111\} fiber axes, then grains which have reached the \{111\}<110> orientations would be brought back nearer to the \{111\}<112> initial orientations, thus becoming unstable again. From this reasoning, the rotation about the \{111\} fiber axes must terminate at the \{111\}<110> orientations.

As the \{111\}<110> orientations are, at the same time, members of the \{110\} fiber texture having the \{110\} fiber axes in the rolling direction, they constitute the transition points of these fiber textures; and further rotations might be possible about the \{110\} fiber axes lying in the rolling direction.

Taking these rotation paths into account, it would be expected that, at high reductions, the rolling texture of the \{111\}<112> specimen approaches to that of the \{111\}<110> specimen. In fact, it can be seen that a close similarity exists between Figure 2(c) and Figure 3(b), with the exception of the \{001\}<110> component, which showed a remarkable development in the \{111\}<112> specimen.

Single crystals having the (111)[112] orientation are reported to be stable up to 70 pct reduction in thickness.\(^1\) Above 90 pct reduction, crystal rotation occurred about the [110] axis lying in the transverse direction, producing a weak \{001\}<110> texture.\(^2\)\(^6\)

The crystal rotation about the sheet plane normal was observed only under restricted conditions. Barrett and Levenson cold rolled (111)[112] single crystals under constraints up to 97 pct reduction in thickness and observed a large orientation spread about the sheet plane normal.\(^1\) Hu and Cline observed such orientation spread at low reductions in the surface layers of a 2 pct Al-Fe single crystal having the (111)[112] initial orientation.\(^2\)\(^6\) They
ascribed this to the higher stress on the specimen surface due to friction between the rolls and the specimen.

Taking these results into account, it might be deduced that, in the present \{111\}{\langle}112\rangle poly-crystalline specimen, the crystal rotation about the sheet plane normal would be caused by the constraining effect of the grain boundary. Such possibility appears to be indicated by Figure 4, which shows (001) facet pits on the rolling plane of Steel 18 cold rolled 70 pct. It can readily be seen that, in a (111) grain, the pyramids rotate continuously about the sheet plane normal, as the grain boundary is approached.

Thus, most of the orientation changes up to 70 pct reduction occurred along the $\psi = 0$ deg line and these would be described as a crystal rotation about the $\langle 110 \rangle$ axes lying in the transverse direction.

On further cold rolling to a reduction of 85 pct, Figure 5(e), broad peaks at the $\{111\}\{112\}$ orientations developed into the $\{111\}$ fiber texture having the $\langle 111 \rangle$ axes in the sheet normal direction. A new peak was formed now at about $\theta = 47$ deg on the $\psi = 90$ deg line, indicating the increase of the members of the $\langle 110 \rangle$ fiber texture having the $\langle 110 \rangle$ fiber axes in the rolling direction.

After 91 pct cold reduction, both this and the $\langle 111 \rangle$ fiber texture increased their intensities, whereas a preference for the $\{111\}\{112\}$ orientation became less evident on the $\theta = 55$ deg line, Figure 5(f).

Thus, the orientation changes above 70 pct reduction occurred predominantly on the $\theta = 55$ deg line and on the $\psi = 90$ deg line, and were essentially the same as those observed on the $\{111\}\{112\}$ and $\{111\}\{110\}$ specimens. Note that, aside from the presence of the (001) component and aside from the differences in the sharpness, Figures 2(c), 3(b) and 5(f) represent qualitatively the same texture. It might be that, although starting points are different, the $\{110\}$, $\{111\}$ and $\{111\}$, $\{110\}$ specimens follow the same rotation path to reach the same stable end orientation. The rotation path seems to consist of three stages. At low reductions, rotations occur about the $\langle 110 \rangle$ axes lying in the transverse direction; and grains having the initial orientations in the range $\{110\}$ to $\{111\}$ are brought into the $\{111\}$ orientations. Thus, at medium reductions, a $\{111\}$ texture develops. Then rotations occur about the $\langle 111 \rangle$ axes lying in the sheet normal direction. This leads to the formation of a $\langle 111 \rangle$ fiber texture having the $\langle 111 \rangle$ fiber axes in the sheet normal direction. Those grains which have arrived at the $\{111\}$ orientations undergo additional rotation about the $\langle 110 \rangle$ axes in the rolling direction.

Bennewitz also divided the development of the rolling textures into three stages, but in his case the rotation path is different. In his model, the most basic assumption consists of the presence of the $\{110\}$ 60 deg RD fiber texture. Members of this fiber texture were, in the range $\phi = 34$ to 45 deg, projected on the $\phi = 45$ deg section and were shown in dashed lines in Figures 5(a) to (d). (Members lying in the range $\langle 545\rangle$ to $\{12\}$ $\{110\}$ are located quite close to the $\phi = 45$ deg

\noindent FIGURE 4 (100) facet pits on the rolling plane of Steel 18 rolled 70 pct. (The sample was polished to the center of the specimen thickness).

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\textit{The \{110\}\{001\} specimen} Figures 5(a) to (f) are $\phi = 45$ deg sections of the crystallite orientation, showing the textures of this specimen at each stage of cold rolling. Figure 5(a) represents a sharp initial texture. After 30 pct cold rolling, the maximum at the $\{110\}$ orientation disappeared, while two broad peaks emerged at the positions corresponding to the $\{554\}$ orientations, Figure 5(b). At 50 pct reduction, these peaks started to leave the $\{554\}$ orientation, aiming at the $\{111\}$ orientation, Figure 5(c). At 70 pct reduction, these were reached exactly. Note that, in Figure 5(d), one of the main peaks is located at $\theta = 55$ deg on the $\psi = 0$ deg line. The height and sharpness of these peaks were remarkably increased.
section.) If grains gather around the \( \{554\}<225> \) orientations at low reductions and rotate further about the \( <110> \) 60 deg RD fiber axes toward the \( \{112\}<110> \) orientations, as Bennewitz predicted, then it should be observed in the present \( \{110\}<001> \) specimen, that one of the sharp maxima which were formed at the \( \{554\}<225> \) orientations at 30 pct reduction would move with increasing reduction along this dashed line, aiming at the \( \{112\}<110> \) orientations.

However, the observed orientation changes did not occur along the dashed line, but along the \( \theta = 55 \) deg line, as described above. In addition, it was noted earlier, in Figure 5(d), that the centers of the maxima arrived exactly at the \( \{111\}<112> \) orientations, passing through the \( \{554\}<225> \)
orientations. Thus, the significance of specifying the \{554\}\{225\} orientations seems to be doubtful in the case of the rolling texture.

The process occurring in the present \{110\}\{001\} specimen below 70 pct reduction is qualitatively the same as those observed on the single crystals having this orientation.\textsuperscript{27,30} In the present polycrystalline specimen, however, the \{001\}\{110\} component showed a remarkable development above 85 pct reduction. The same phenomenon was observed also in the \{111\}\{112\} specimen cold rolled 85 pct, Figure 3(b). This suggests again that the rotation paths are partly common in both cases.

The \{110\}\{110\} specimen Figures 6(a) to (d) are \(\phi = 45\) deg sections of the crystallite orientation distribution, showing the development of the rolling textures in this specimen. Figure 6(a) shows the \{110\}\{110\} initial texture.

After 30 pct cold reduction, the sharpness of the initial texture decreased and minor peaks appeared at the \{001\}\{110\}, \{001\}\{100\}, \{111\}\{110\} and \{111\}\{112\} orientations. However, in a marked contrast to the \{110\}\{001\} specimen, large amounts of the crystals retained their initial orientations, showing their high stabilities. A continuous orientation spread was produced on the \(\psi = 90\) deg line between the \{111\}\{110\} and \{110\}\{110\} orientations, Figure 6(b).

At 50 pct reduction, the original peak at the \{110\}\{110\} orientations disappeared completely and, instead, weak \{111\} and \{100\} fiber textures, having their fiber axes in the sheet normal direction, developed, Figure 6(c).

After 70 pct cold rolling, a broad peak was formed on the \(\psi = 90\) deg line. This peak extended preferentially along the \(\theta = 50\) deg line, Figure 6(d).

Of these orientation changes, those along the \(\psi = 90\) deg line are described as a rotation about the \{110\} axes in the rolling direction. As a result of this rotation, grains having the \{110\}\{110\} initial orientations could thus be brought as far as \(\theta = 50\) deg on the \(\psi = 90\) deg line after 70 pct reduction. Although single crystals having the \{110\}\{110\} orientations tend also to rotate slightly about the rolling direction, they retain their initial orientations up to 70 pct reduction.\textsuperscript{27,29} Taking these results into account, it might be concluded that this

![Figure 6](image-url)
rotation occurs more readily in polycrystals due to the constraining effect of the grain boundary.

The \{001\}<110\> specimen Figures 7(a) and (b) are \(\phi = 45^\circ\) deg sections of the crystallite orientation distribution, showing the textures of this specimen before and after 70 pct cold rolling. The initial texture contained a large orientation spread along the \(\psi = 90^\circ\) deg line, Figure 7(a). After 70 pct reduction, the \{001\}<110\> orientations decreased remarkably and the broad peak on the \(\psi = 90^\circ\) deg line was shifted along this line to the \{114\}<110\> orientations, Figure 7(b). Further cold rolling up to 90 pct produced no substantial change in the rolling texture.

Thus, in this specimen, grains rotate at lower reductions about the <110> axes in the rolling direction; and above 70 pct reduction, they are stable.

The instability of the \{001\}<110\> initial texture was also confirmed by Richards and Ormay\(^9\) in a 1.5 pct Si steel. As single crystals having the (001) [110] orientation are generally stable,\(^{25,27}\) this instability might be attributed to the constraining effect of the grain boundary.

If this explanation is correct, and if the results of the \{111\}<112\> and \{110\}<110\> specimens are taken into account, it would be concluded that this constraining effect enhances crystal rotations, which would not be expected to take place in single crystals. This might be possible in \(\alpha\)-Fe, in which slip can occur on any of the \{110\}, \{112\} and \{123\} planes. In this case, it would be expected that the constraining effect of the grain boundary could be relaxed relatively easily, accompanying large orientation changes in the neighbourhood of the grain boundary.

Aspden\(^{31}\) studied the effect of constraints in 3 pct Si-Fe crystals having the (001)[100] initial orientation. He found that the amount of rotation toward the \{001\}<110\> stable end orientations was decreased by using constraints during rolling.

Thus, this effect seems to depend on the orientation of the grain in question. Furthermore, it may be influenced by the shape, grain size and orientation of neighbouring grains. At present, this problem is too difficult to be discussed theoretically.

Low-carbon steel Figures 8(a) to (f) are \(\phi = 45^\circ\) deg sections, showing the development of the rolling textures in Steel 13.

The texture of the starting hot-band contained weak components such as \{001\}<110\>, \{221\> <114\>(\(\theta = 70.5^\circ\) deg, \(\psi = 0^\circ\) deg) and \{221\> <110\>(\(\theta = 70.5^\circ\) deg, \(\psi = 90^\circ\) deg), Figure 8(a).

After 15 pct reduction, a distinct peak was formed at the \{332\> <113\>(\(\theta = 64.8^\circ\) deg, \(\psi = 0^\circ\) deg) orientations. The \{001\> <110\> orientations also sharpened somewhat, while other orientations on the \(\theta = 0^\circ\) deg line decreased, Figure 8(b). These orientation changes may be explained in terms of the observations of Koh and Dunn.\(^{25}\) Their results on single crystals having a [110] axis in the transverse direction indicate that grains having an initial orientation in the range \{110\}<001\> to \{112\> <111\> tended to rotate to the \{111\> <112\> orientations, while those in the range \{112\> <111\> to \{001\> <110\> rotated to the \{001\> <110\> orientations.

On the \(\psi = 90^\circ\) deg line, a broad peak was formed at about \(\theta = 10^\circ\) deg. As was already shown in Figure 7(b), this peak might be derived from the \{001\> <110\> initial orientations by the rotation about the <110> axes in the rolling direction. This
FIGURE 8  $\phi = 45$ deg sections, showing the development of the rolling textures in an aluminum killed steel.

may explain the fact that the increase of the $\{001\langle110\rangle$ orientations after 15 pct cold rolling was relatively small. From this broad peak, contour lines were bulging to the center of the $\phi = 45$ deg section, forming a ridge.

At 30 pct reduction, the $\{554\langle225\rangle$ orientations became one of the main components of the rolling texture, Figure 8(c). Comparing Figure 8(c) with Figure 1, it would suggest that the ridge, which extended to the center of $\phi = 45$ deg section, was associated also with the $\{554\langle225\rangle$ orientations. At the beginning of this stage, grains having the initial orientations in the range $\{110\langle110\rangle$ to $\{221\langle110\rangle$ began to rotate about the $\langle110\rangle$ axes in the rolling direction, producing a small peak at the $\{332\langle110\rangle$ orientations ($\theta = 64.8$ deg, $\psi = 90$ deg).

At 50 pct reduction, this crystal rotation proceeded further and the $\{332\langle110\rangle$ peak merged into the ridge which ran obliquely across the $\phi = 45$ deg section, Figure 8(d). On the other hand, the broad peak, which had been located at $\theta = 10$ deg on the $\psi = 90$ deg line collapsed and extended into a plateau running parallel to the $\psi = 90$ deg line. In this way, two crystal rotations were occurring on the $\psi = 90$ deg line in the opposite directions.

At 70 pct reduction, a sharp maximum was formed at the $\{335\langle110\rangle$ orientations ($\theta = 40$ deg, $\psi = 90$ deg), Figure 8(e). This might be due to the superposition of these two rotations occurring
along the $\psi = 90$ deg line. On the $\psi = 0$ deg line, the peak at the \{554\}\{225\} orientations tended to leave these orientations behind, and the presence of the \{111\} fiber texture having the \{111\} axes in the sheet normal direction became evident.

At 90 pct reduction, both the \{335\}\{110\} peak and the \{111\} fiber texture developed remarkably, Figure 8(f). In the latter, some preference for the \{111\}\{110\} orientations seems to be present. The \{554\}\{225\} orientations were no more at the center of the peak on the $\psi = 0$ deg line.

Thus, the rolling textures of this specimen having a weak initial texture could be explained in terms of the crystal rotations which were observed in specimens having sharp initial textures. It can be constructed approximately by the superpositions of various components, which follow their predetermined rotation path to reach their stable end orientations.

Dashed lines in Figures 8(a) to (f) have the same meaning as those in Figures 5(a) to (d). Also in this case, the texture development did not proceed along these lines. With increasing cold reduction, the dashed line seemed to be rather traversed by the ridge running diagonally across the $\phi = 45$ deg section. At higher reductions, this ridge became more vertical, and this is just the process through which the \{111\} fiber texture having the \{111\} axes in the sheet normal direction becomes more complete.

Bennewitz's model was examined from another viewpoint. Figures 9(a) to (d) are the constant
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ψ sections of the crystallite orientation distribution of Steel 13 cold rolled 70 pct. For the present purpose, sections in the range ψ = 65 to 90 deg are sufficient. Members of the ⟨110⟩ 60 deg RD fiber texture in the range {554}⟨225⟩ to {112}⟨110⟩ are contained in these sections and they are shown by black dots in the figures. Although a black dot was situated always near the center of the peak at each constant ψ sections, the deviation between them was sometimes larger than the experimental error. The disagreement became largest at the ψ = 90 deg section at which the crystallite orientation distribution exhibited a maximum. At other reductions, the skeleton line deviated from the members of the ⟨110⟩ 60 deg RD fiber texture much more definitely.

From these observations it can be concluded that the whole features of the orientation changes occurring continuously during cold rolling cannot be properly described in terms of the ⟨110⟩ 60 deg RD fiber texture.

The description in terms of the ⟨111⟩ fiber texture having the ⟨111⟩ axes in the sheet normal direction seems to be a better one. The same conclusion was reached by Takechi, Kato and Nagashima, Bunge and Heckler and Granzow. In the latter two investigations, the crystallite orientation distribution analysis was adopted.

CONCLUSION

The development of the rolling textures in low-carbon steels having sharp initial textures was studied by using the crystallite orientation distribution analysis. Based on the result of this analysis, the rotation path of a grain having a specific initial orientation in a random polycrystal was deduced.

Grains having the {110}{001} initial orientations rotate at low reductions about their ⟨110⟩ axes in the transverse direction and reach the {111}⟨112⟩ orientations. Subsequently, they rotate about the ⟨111⟩ axes in the sheet normal direction, forming a ⟨111⟩ fiber texture. This rotation terminates, however, at the {111}⟨110⟩ orientations. Since these orientations are also members of the fiber texture having the ⟨110⟩ axes in the rolling direction, grains rotate then about these ⟨110⟩ axes, aiming at the {112}⟨110⟩ orientations.

Although starting points are different, grains having either {111}⟨112⟩ or {111}⟨110⟩ orientations seem to follow the same rotation path. The rotation about the ⟨111⟩ axes in the sheet normal direction, mentioned above, would be caused by the constraining effect of the grain boundary.

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REFERENCES

1. C. S. Barrett and L. H. Levenson, AIME Trans. 145, 281 (1941).
2. F. Haessner and H. Weik, Arch. Eisenhüttenw. 27, 153 (1956).
3. H. Möller and H. Stäblein, Arch. Eisenhüttenw. 29, 377 (1958).
4. J. Bennewitz, Arch. Eisenhüttenw. 33, 393 (1962).
5. I. L. Dillamore and W. T. Roberts, Acta Met. 12, 281 (1964).
6. C. A. Stickles, Trans. Met. Soc. AIME 233, 1550 (1965).
7. R. A. Vandermeer and J. C. Ogle, Trans. Met. Soc. AIME 242, 1317 (1968).
8. H. Takechi, H. Kato and S. Nagashima, Trans. Met. Soc. AIME 242, 56 (1968).
9. P. N. Richards and M. K. Ormay, Trans. Met. Soc. AIME 245, 715 (1969).
10. R. J. Roe and W. R. Kriegbaum, J. Chem. Phys. 40, 2608 (1964).
11. R. J. Roe and W. R. Kriegbaum, J. Chem. Phys., 41, 737 (1964).
12. R. J. Roe, J. Appl. Phys., 36, 2024 (1965).
13. R. J. Roe, J. Appl. Phys. 37, 2069 (1966).
14. H. J. Bunge, Mathematische Methoden der Texturanalyse (Akademie Verlag, Berlin, 1969).
15. H. J. Bunge and F. Haessner, J. Appl. Phys. 39, 5503 (1968).
16. H. J. Bunge, Phys. Stat. Sol. 26, 167 (1968).
17. H. J. Bunge, *Kristall und Technik* 3, 439 (1968).
18. H. J. Bunge and W. T. Roberts, *J. Appl. Cryst.* 2, 116 (1969).
19. T. Taoka, E. Furubayashi and S. Takeuchi, *Japanese J. Appl. Phys.* 4, 120 (1965).
20. B. F. Decker, E. T. Asp and D. Harker, *J. Appl. Phys.* 19, 388 (1948).
21. L. G. Schulz, *J. Appl. Phys.* 20, 1030 (1948).
22. J. Grewen, D. Sauer and H. P. Wahl, *Scripta Met.* 3, 53 (1965).
23. P. R. Morris and A. J. Heckler, in *Advances in X-Ray Analysis*, (Plenum Press, New York, 1968), Vol. II, p. 454.
24. A. J. Heckler and W. G. Grazow, *Met. Trans.* 1, 2089 (1970).
25. P. K. Koh and C. G. Dunn, *AIME Trans.* 203, 401 (1955).
26. H. Hu and R. S. Cline, *Trans. Met. Soc. AIME* 224, 784 (1962).
27. T. Taoka, E. Furubayashi and S. Takeuchi, *Trans. Iron Steel Inst. Japan* 6, 290 (1966).
28. C. G. Dunn, *Acta Met.* 2, 173 (1954).
29. J. L. Walter and W. R. Hibbard, Jr., *Trans. Met. Soc. AIME* 212, 731 (1958).
30. H. Hu, *Trans. Met. Soc. AIME* 221, 130 (1951).
31. R. G. Aspden, *Trans. Met. Soc. AIME* 215, 986 (1959).