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Formation and crystallography of nano/ultrafine-trimorphic structure in metastable β titanium alloy Ti-5Al-5Mo-5V-3Cr-0.5Fe processed by dynamic deformation at low temperature

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ABSTRACT

The dynamic $\beta \rightarrow \alpha$ phase transformation of metastable β titanium alloy Ti-5553 during the hot deformation process was investigated, including the crystallographic orientation relationship between α phase and β matrix and the variant selection mechanism of nanoscale α precipitates. It is revealed that nano/ultrafine-trimorphic structure can be successfully produced dynamically Ti-5553 alloy through warm compression without severe plastic deformation or subsequent aging treatment. Ultrafine equiaxed α/β grains do not respect Burgers orientation relationship were observed in the slip bands together with the remaining regions consisting of the lamellar α structure maintaining the Burgers orientation relationship with β matrix. Moreover, the irregular α grains were mainly attributed to the fragmentation of α lamellae in some areas. There was variation in the proportion and distribution of the precipitates in the different areas depending on deformation. Furthermore, the strain induced variant selection of α happened drastically.

1. Introduction

Over the last decades, structural applications of metastable β titanium alloys have always been popular in the aerospace industry because of their higher yield strength, better hardenability, better fatigue and crack propagation properties than $\alpha + \beta$ titanium alloys, in which strengthening by grain refinement and precipitation of fine α phase yields similar strength levels as low alloyed and even some stainless steels [1,2]. Microstructural development in these alloys is sensitively linked to a range of processing parameters, such as route of thermo-mechanical process (TMP), temperature and strain. In general, fine or even nanostructured materials are expected due to the superior mechanical properties could be obtained. Hence, extensive efforts have been devoted to the ultrafine microstructure and its evolution of titanium alloys related to various thermomechanical processes in the past decades owing to their considerable scientific and practical interest [3–6].

The distribution and morphology of α precipitates in the microstructure largely depend on the nucleation mechanism. In the β phase matrix, α precipitates can nucleate either homogeneously or

heterogeneously. The homogeneous nucleation of α phase involves a pseudo-spinodal decomposition of the metastable β phase, which leads to small compositional fluctuations in the matrix [7]. On the other hand, nucleation of α phase can occur heterogeneously at grain boundaries, subgrain boundaries, dislocations or precursors such as ω phase, β' phase or martensite [8–10]. The α phase formed during the heat treatment process generally possess platelet morphology in metastable β titanium alloys. To obtain the fine multi-morphology structure, deformation must be considered necessarily. A commonly adopted approach is the introduction of severe plastic deformation prior to aging, thus ultrafine-duplex ($\alpha + \beta$) structure attributed to the α nucleation at abundant crystal defects would be obtained. S.L. Semiatin et al. [11] have also shown that the final grain size and the required strain can be significantly reduced through lowering the deformation temperature and/or utilizing a finer initial microstructure (i.e., reducing the α platelet thickness in particular). Nevertheless, the nano/ultrafine-trimorphic structure has never been reported in previous researches of metastable β titanium alloys. Furthermore, none of studies have identified the exact locations of the different morphologies respectively, nor have they clarified its dependence on deformation

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Fig. 1. (a) EBSD band contrast micrograph of the single β phase initial microstructure; (b) EBSD CD axis Inverse Pole Figure (IPF) micrograph; (c) Micro- and macro- $\{110\}$ pole figures; (d) Correlated misorientation angle distribution.

parameters. In addition, R. Shi et al. [12] found that, for a given undercooling, the development of a transformation texture (also called microtexture) of the α phase due to variant selection during precipitation is determined by the interplay between externally applied stress or strain and internal stress generated by the precipitation reaction itself. M. Klimova et al. [13] have found that the average deviation from the Burgers orientation relationship (BOR) [14] reaches a value of 34.5° at a strain of 1.2 in Ti-5Al-5Mo-5V-1Cr-1Fe. Therefore, hot deformation process could give a significant effect on the α variant selection mechanism and α/β crystallographic orientation.

In our present work, the low temperature compressive deformation was performed on the typical metastable β titanium alloy Ti-5Al-5Mo-5V-3Cr-0.5Fe (Ti-5553) with single β initial microstructure. The dynamic $\beta \rightarrow \alpha$ phase transformation behavior during the hot deformation was investigated, including microstructure evolution and variant selection mechanism of nanoscale α precipitates. The results could provide a valuable understanding of the interrelation between phase transformation and thermomechanical processing.

2. Materials and Research Methodology

The as-received material of the present study is ($\alpha + \beta$) forged Ti-5553 bar with chemical composition (wt%) of Al: 5.26; Mo: 4.99; V: 4.80; Cr: 2.86; Fe: 0.42; Ti: balance. The β transus temperature of Ti-5553 alloy determined by metallographic analyses is approximately 870°C . Cylindrical specimens, 10 mm in diameter and 15 mm in height, were machined from the forged bar. They were then solution treated at 900°C for 30 min followed by water quenching in order to obtain single β phase state with homogeneous microstructure. The macro-texture of the initial microstructure was measured by using Brüker D8 Advance X-Ray Diffraction (XRD). Cu K α ($\lambda = 1.5418 \text{ \AA}$) was used, under 40 kV and 40 mA.

Uniaxial hot compression test was carried out with a Gleeble-3500 thermo-mechanical simulator along the cylindrical axis of the specimen. Prior to compression, each specimen was heated to the processing temperatures at a rate of 25°C/s and then held for 10 s before the commencement of compression to ensure an establishment of homogeneous temperature field in the specimen. The temperature was controlled by a thermocouple welded to the central part of the specimen

surface. Graphite lubrication was applied to the anvil and the specimen in order to reduce the relative friction between the two parts to enhance deformation homogeneity and also to prevent adhesion. The temperature (600°C), strain rate (0.001 s^{-1}) and deformation strain (0.7) were automatically controlled and recorded by the Gleeble-3500 thermal simulator system.

The deformed specimens were sectioned parallel to the compression axis and all microstructural and crystallographic characterizations were performed on this specimen section. For SEM (JSM-6500F) observations and EBSD measurements (OXFORD EBSD acquisition camera and the OXFORD-AZTEC online acquisition software), the specimens were first mechanically polished and then electro-polished with a solution of 20% perchloric acid in methanol under a voltage of 35 V at 10°C for 6 s. And then they were etched in a modified Kroll's reagent of 10 vol% HF, 10 vol% HNO $_3$ and 80 vol% H $_2\text{O}$ for 10 s. For TEM (Philips CM200 operated at 200 kV) analyses, foil specimens were prepared first by mechanical thinning to about 80 μm in thickness. Then discs with 3 mm in diameter were punched off and further thinned to be electron transparent by twin-jet electro-polishing using a Struers Tenupol-5 twin-jet electropolisher in a solution of 20% perchloric acid in methanol under 10 V at -35°C . The orientations of the α/β grains of the two phases were determined from the Kikuchi patterns produced under the convergent electron beam with the home made orientation determination software Euclid Phantasies (EP) [15,16] and represented with a triplet of Euler angles in Bunge notation [17].

3. Results and Discussion

3.1. Initial Microstructure of Ti-5553 Alloy

The initial single β microstructure obtained by solution treatment at 900°C for 30 min followed by water quenching, as shown in Fig. 1(a), is composed of single β phase with an average grain size of $\sim 300 \mu\text{m}$ in equiaxed shape. Except for few low angle grain boundaries (LAGBs, 2° – 10°), almost all of the β grains are surrounded by high angle grain boundaries (HAGBs, $> 10^\circ$) (Fig. 1(b)). The correlated misorientation angle distribution histogram in Fig. 1(d) displays that the initial microstructure possesses a disorientation angle distribution close to that of randomly oriented polycrystals (the line profile), suggesting that the

Fig. 2. SEM micrographs of the nano/ultrafine-trimorphic structure in the central area of the deformed Ti-5553 alloy: (a) equiaxed α ; (b) lamellar α ; (c) fragmented α .

alloy is probably not sharply textured. That is consistent to the implication of the micro/macro $\{110\}$ pole figures in Fig. 1(c).

3.2. Nano/Ultrafine-Trimorphic Structure and Orientation Relationship

The microstructural observation of the center of the deformed specimen using SEM/TEM shows in Fig. 2. Obviously, three different morphologies of α phase could be detected simultaneously in the center, i.e., equiaxed α grains (50–200 nm) (Fig. 2(a)), lamellar α (~ 20 nm in width) (Fig. 2(b)) and fragmented α grains (~ 20 nm) (Fig. 2(c)). What needs to be stressed is that all of the various α phase precipitated dynamically during the compression test. The equiaxed microstructure seems form irregular net which filled by the lamellar and fragmented α (Fig. 2(a)).

Further TEM micrographs and SAED patterns are shown in Fig. 3. The equiaxed constituents consist of mixed fine α and β grains. The SAED pattern of the fine equiaxed microstructure shows that the α/β reflections tend to form arcs, suggesting the random crystallographic orientations between α/β phases (Fig. 3(a)). For the lamellar constituents, they are composed of alternatively distributed α and β platelets, maintaining BOR ($\{0001\}_{\alpha} \parallel \{110\}_{\beta}$, $\langle 11\bar{2}0 \rangle_{\alpha} \parallel \langle 111 \rangle_{\beta}$ [14]), as shown in Fig. 3(b). The SAED pattern indicates that the α lamellae in one colony possesses the unique crystallographic orientation.

To investigate the orientation relationship (OR) between the fine α and β phases in the equiaxed constituents zones, the deformed specimens was further studied by TEM electron diffraction under the convergent beam. Fig. 4 is a TEM bright field image of the equiaxed α and β grains. The orientations of the α/β grains of the two phases were determined from the Kikuchi patterns produced under the convergent electron beam with the home made orientation determination software package Euclid Phantasies (EP) [15,16] and represented with a triplet of Euler angles in Bunge notation [17]. The table in Fig. 4 shows the Euler angles of the α and β grains indicated in the TEM bright field micrograph. The pole figures of the adjacent three β grains and the four α grains are displayed in the Fig. 5. Unfortunately, no ORs including the typical BOR could be found between the adjacent β and α grains.

Whereas, compared with the case of heat treatment process [3], all the α phase (grain boundary α and intragranular α) generally maintains the BOR with the β matrix. Obviously, the external hot deformation has a significant influence on the obeying of BOR during $\beta \rightarrow \alpha$ phase transformation.

3.3. Variation of Microstructure in the Compression Disc

Fig. 6 shows the variation of precipitation morphologies along the compression disc radius and thickness. In the area along the thickness near the edge of the specimen (denoted by A), except for fine equiaxed α grains (~ 150 nm) nuclear at the initial β grain boundaries, the dense cluster of ultrafine α lamellae (~ 20 nm in width) forming as colonies distributed in the β grains (Fig. 6(a₁)). This proves that area A has experienced relatively limited deformation during deformation. Meanwhile, only two or three α variants with different crystallographic orientations distributed in one β grain (Fig. 6(a₂)). In the area along the thickness near the central of the specimen (denoted by B), the straight multi-slip bands are formed significantly and the microstructure consisted of both fine equiaxed α/β grains and the α lamellae structure (Fig. 6(b₁)). The equiaxed and parallel lamellar α precipitates are formed inside and outside the slip bands, respectively (Fig. 6(b₂)). Straight multi-slip bands and equiaxed grains as well as fragmented α platelets are also easily fund in the area of C (Fig. 6(c₁₋₂)) (similar to B). For area D, the microstructure is kinked but no more significant slip bands (Fig. 6(d₁₋₂)). The characteristics of B and C are closer to the central microstructure, which contributed to the relatively larger strain applied to B/C than A/D. In general, the nearer to the central of specimen, the easier to obtain a nano/ultrafine-trimorphic structure. Actually, the irregular equiaxed microstructure nets in central area are the result of drastic kinking and accumulation of slip bands during deformation. All α platelets distributed as certain directions, i.e., the α variants were selected severely, which is totally different from the α platelet precipitates usually nucleate and eventually form a basket-weave structure in the metastable β titanium alloys [18,19].

Fig. 3. TEM micrographs and the corresponding SAED patterns: (a) bright field TEM micrograph and its SAED pattern of the fine equiaxed microstructure; (b) dark field TEM micrographs and its SAED pattern of the α/β lamellae microstructure.

Grain	Euler angle (ϕ_1, Φ, ϕ_2)
α_1	280.17, 50.44, 260.50
α_2	162.47, 39.50, 179.71
α_3	140.81, 88.80, 210.79
α_4	220.49, 66.62, 270.55
β_1	132.80, 19.47, 130.90
β_2	160.64, 4.50, 55.26
β_3	199.68, 77.24, 320.93

Fig. 4. TEM bright field micrograph of the equiaxed α and β grains and the corresponding Euler angles of the marked α and β grains with dotted line in the micrograph.

Fig. 5. $\{110\}_\beta / \langle 111 \rangle_\beta$ and $\{0001\}_\alpha / \langle 11\bar{2}0 \rangle_\alpha$ plane/direction pole figures of the three β grains and the four α grains selected in Fig. 4.

3.4. Variant Selection Mechanism of α Phase

All the 12 α variants would nucleate with equal probability during $\beta \rightarrow \alpha$ phase transformation in an ideal case, as shown in schematic diagram Fig. 7(a). The different colors in Fig. 7 represent corresponding α variants, respectively. One can detect up to 12 α variants in one β grain theoretically [20]. Therefore, in the present results, the only reason of just two or three orientations of α lamellae formation during the deformation must be the strong variant selection. For instance, the α_{v1} (red) and α_{v2} (green) in the one deformed β grain (Fig. 7(b)). The present variant selection of intragranular α (α_i) during the dynamic precipitation is determined by the interplay between externally applied stress (σ^{ex}) and internal stress generated by the phase transformation (σ^{tr}) [12,21]. Thus, the composite stress (σ^{com}) must be considered in the present work. When a coherent or semi-coherent crystallographic orientation is formed during the phase precipitation process, the elastic strain will be built up by lattice misfits between the second phase and

matrix. Definitely, the application of an external directional elastic stress may cause contraction or expansion along the precipitating habit planes, which will lead to either enhancement or release of the misfit strain. The energy change caused by the variations of the misfit strain is the interaction energy between external stress and the stress field of the misfitting precipitate. The variations of the interaction energy of different variants of precipitate have been quantitatively analyzed with respect to external stress direction in the frame of Eshelby's elastic inclusion model [22]. On the other hand, the volume change under external stress during the precipitated platelets replacing the matrix materials contributed to the variation of potential energy of system [23]. Whereupon potential energy change from the volume change under external stress could be expressed as

$$\Delta G_E^{\text{ex}} = \sigma \cos \theta \Delta V \quad (1)$$

where θ is the angle between the external stress axis and the α platelet normal, as shown in Fig. 7(c). ΔV is the volume change during the $\beta \rightarrow \alpha$ phase transformation.

In addition, the interaction energy between external stress and the stress field of the misfitting interface of α/β could be expressed as [23]

$$\Delta G_E^{\text{tr}} = \sigma \cos \theta V_\alpha \epsilon \quad (2)$$

where V_α is the volume of alpha precipitate, ϵ is the coherency strain between α/β . Therefore, the potential energy change of different α platelet variants differs from each other due to different θ values with stress axis, expressed as

$$\Delta G = V_\alpha \Delta G_V + A \Delta G_R + \Delta G_E^{\text{ex}} + \Delta G_E^{\text{tr}} = V_\alpha \Delta G_V + A \Delta G_R + \sigma \cos \theta (\Delta V + V_\alpha \epsilon) \quad (3)$$

where ΔG_V and ΔG_R are the volumic free energy change and interfacial energy and elastic strain energy respectively. A is the α phase surface area. The α variant with the most favorable θ value will get the largest decrease of potential energy and priority to nucleate and grow (Fig. 7(c)), which resulting in only several α variants were selected.

3.5. Microstructure Evolution of Ti-5553 Alloy During the Hot Deformation Process

As mentioned above, all of the slip bands are occupied by fine equiaxed α and β grains, while irregular fragmentized α particles and parallel α lamellae are distributed in the gaps between the bands. Because the initiation of the slip bands produced by dense dislocations is accompanied by the nucleation of α phase during the hot deformation, the slip bands would be the preferential nucleation sites [24].

Fig. 6. Variation of precipitation morphologies along the compression disc radius and thickness.

Abundant cumulate dislocations and the drastic lattice rotation in the bands would result in α nuclei growing up to a fine equiaxed structure, as Fig. 8(a) shows. These precipitates underwent a rotation and increased misorientations to accommodate the continuous strain (Figs. 4–5). Subsequently, both the size of α particles and the width of slip bands would increase gradually with the increased strain. Another important point is that the diffusion rate of alloy elements is faster at GBs and slip bands, which would contribute to the nucleation and growth of α phase [25,26]. That is why the size of α precipitates on slip bands and GBs is larger than the width of α_1 platelets.

On the other hand, the α platelets with certain stretching directions are favorable for nucleation and growth due to variant selection under the applied force. According to the above SEM/TEM observations on the deformed lamellar microstructure, the detailed process of the evolution of α/β lamellae under the hot compression can be shown in Fig. 8(b) and described below.

When lower compression strain is applied, severe kinking of α/β lamellae occurred in the α/β lamellar colonies under the complicated shear stress state (upper half of Fig. 8(b)). It should also be noted that the kinking of α/β lamellae should be related to the original orientation of the lamellae relative to the loading direction [27]. Bieler et al. [28] observed that globularization is difficult in colonies having their c -axes

parallel to the compression direction so that the prism and basal slip systems cannot be activated. And then, globularization (fragmentation) of the α/β lamellae may occur under the effect of dense dislocation configurations with the increased strain. Actually, the kinking of α/β lamellae is the preparatory stage of fragmentation.

When higher compression strain is applied, the flat-smooth cutting might form directly and across α/β lamellae gradually (lower half of Fig. 8(b)). Thus, the α platelets are easier to be fragmented under the high shear strain. Actually, the initiation of the flat-smooth cutting might be accompanied by the fragmentation of α/β lamellar around the fracture surface by intense dislocation activities. Refs. [9,29] described that the refinement of grains in deformation bands as a recrystallization behavior. However, the α/β lamellar structures are really thin with ~ 20 nm in width in the present work. Thus, the fragmentation and globularization is speculated to play an important role in the grain refinement process. Moreover, the increased deviation from BOR of α/β due to the deformation would accelerate the kinetics of α lamellae fragmentation [30]. The fragmented α/β lamellae might be further broken and rotated in order to release the stresses, which led to the formation of randomly orientated nanocrystallites. With the continuous increase of the strain, the adjacent α precipitates and β blocks would be swept into the localized strain region. In other words, the fracture

Fig. 7. Illustration of α variant selection mechanism during the hot deformation process: (a) all α variants nucleate theoretically with equal probability during $\beta \rightarrow \alpha$ phase transformation; (b) α variant selection during the hot deformation; (c) schematic of α phase under the external stress. (α phase variants are represented by various colors.) (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Fig. 8. Schematic diagram of the dynamic precipitation and evolution of α phase during the isothermal deformation: (a) α phase in slip bands; (b) α lamellae colony.

surface thickens at the expense of the surrounding α precipitates and β blocks.

It is believed that the flat-smooth cutting may not develop in α colony until a given applied stress is large enough to make the activated dislocations in the α platelet piling up at the α/β interface, which could cut through the interface and glide into the β lamella. Such a applied stress to trigger cutting should be associated with the scale of the lamella and the deformation temperature [31]. On the other hand, only the kinking of α/β lamellar happened in somewhere when the applied stress is not up to the critical value (σ_c) [32]. This evaluation is one of the causes for the flow softening and play an important role in the workability and formability of the alloy during hot working process, which is not only found in the present work but also is often observed in α/β lamellar structured Ti alloys.

4. Conclusions

The microstructure evolution and variant selection mechanism of α phase during the thermo-mechanical treatment at low temperature were studied in the Ti-5553 metastable β titanium alloy with single β phase initial microstructure. The following main conclusions can be drawn:

- (1) The nano/ultrafine-trimorphic structure, consisting of equiaxed α , irregular fragmented α and lamellar α , can be successfully produced dynamically in a Ti-5553 alloy through warm deformation of a single β initial microstructure. All of these varied α precipitates were formed dynamically during the warm deformation process, which is distinctively different with the previous related studies.
- (2) Ultrafine equiaxed α/β grains do not respect BOR were observed in the slip bands together with the remaining regions consisting of the lamellar α structure maintaining BOR with β matrix. A dense array of irregular α grains could be detected in some places, which mainly attributed to the fragmentation of the lamellar α structure. Moreover, there is significant variation in the proportion and distribution of the α precipitates along the compression disc radius and thickness depending on the amount of deformation.
- (3) Only two or three α variants would nucleate in local β matrix under the hot deformation condition due to the drastic strain induced variant selection.

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