

Evolution of Grain Boundary Precipitates in a Directionally Solidified Ni-base Superalloy during High Temperature Creep

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Abstract

The evolution of grain boundary phases during high temperature creep has been systematically investigated in a directionally solidified Ni-base superalloy. At the primary creep stage, precipitation of nanosize M_5B_3 boride from γ matrix and phase transformation from $M_{23}(C,B)_6$ borocarbide to M_5B_3 boride occur at grain boundaries. At the steady-state creep stage, precipitation of blocky $M_{23}(C,B)_6$ borocarbide and coarsening of $M_{23}(C,B)_6$ with the dissolution of M_5B_3 boride occur at grain boundaries. At the tertiary creep stage, precipitation of M_5B_3 boride from γ matrix or phase transformation from $M_{23}(C,B)_6$ borocarbide to M_5B_3 boride occur again at grain boundaries. The precipitation of M_5B_3 is due to diffusion of B atoms promoted by applied tensile stress during high temperature creep, and the phase transformation between $M_{23}(C,B)_6$ and M_5B_3 is thought to correlate strongly with the strain rate of different creep stages.

Introduction

Directional solidification (DS) and single crystal (SC) Ni-base superalloys are widely used as blade materials in advanced aircraft and stationary gas turbines because of their excellent mechanical properties at high temperatures [1]. In operating gas turbine engines, an important part of the damage experienced by rotating turbine blades is due to creep deformation [2]. For this reason, a great number of studies have been conducted investigating the creep behavior of superalloys. The two prominent features of DS and SC Ni-base superalloys during high temperature creep deformation are the rapid development of interfacial dislocation networks and directional coarsening or “rafting” of the γ' precipitates [3, 4].

Compared to SC Ni-base superalloys, DS Ni-base superalloys have a particular creep deformation characteristic due to presence of the longitudinal grain boundaries (GBs). The formation of premature cavity and microcrack always concentrates on the GBs during creep, especially at high temperatures where GB sliding becomes a significant deformation mechanism [5-7]. In order to suppress or delay this mode of intergranular failure, trace amount of C and B are added [8, 9].

The beneficial effects of C are derived primarily from the intergranular carbides, and the effects of carbides upon high temperature properties are dependent upon the type of carbide present and its morphology. No all-inclusive statement can be made concerning the most desirable type and morphology of carbides [10, 11]. In general optimum high temperature properties are obtained when $M_{23}C_6$ is present as discrete particles distributed uniformly along the GBs [12-14]. However, much less effort has been devoted to the detailed study of precipitation and

evolution mechanism of intergranular $M_{23}C_6$ during high temperature creep.

It is well accepted that B segregates to the GBs, and a variety of possible strengthening mechanisms include: (1) increasing GBs cohesive strength, lowering GBs diffusivity and GBs surface energy [15, 16], (2) hindering the diffusion of tramp elements to GBs [17, 18], (3) promoting the cross slip of dislocations and the migration of dislocations across the GBs [19, 20], and (4) decreasing the agglomeration of $M_{23}C_6$ concurrently with depletion of γ' from the adjacent matrix and preventing formation of topologically closed-packed phases [21, 22]. However, it should be emphasized that the precise role played by the B element is uncertain. B additions over the solubility limit are shown to result in the intergranular precipitation of the M_5B_2 compound and caused no further improvement or deterioration in mechanical properties [23, 24]. Moreover, it is not clear whether or not presence of any other borides strengthening the GBs.

It should be mentioned that little is known concerning the synergetic effects of C and B, except that B can substitute for C in $M_{23}C_6$ structure [25]. Considering (1) the precipitation and evolution mechanism of intergranular carbide occurred at high temperature creep are not clear, (2) the direct experimental evidence supporting above actual strengthening mechanisms of B is seldom found, and (3) the creep strength of DS Ni-base superalloys correlates very strongly with the microstructure of GBs, it is of utmost important to gather information about the intergranular regions during creep. Therefore, the primary objective of the present work was to employ transmission electron microscopy (TEM) techniques to study the evolution of GB microstructure during high temperature creep, especially focusing on phase precipitation and transformation, trying to obtain the direct experimental evidence revealing the actual strengthening mechanisms of C and B.

Experimental Procedures

The nominal composition in weight percent (wt. %) of the alloy used in this work was 8Cr, 1Mo, 9Co, 6W, 2Ta, 1Nb, 5Al, 2Ti, 0.1C, 0.01B, balance Ni. The experimental alloy was initially vacuum induction melted and cast into polycrystalline ingots. The ingot was remelted and directionally solidified by the high gradient liquid metal cooling (LMC) process. The ceramic mold used in LMC was designed to cast a cluster of four cylindrical bars with 18 mm in diameter and 220 mm in length.

The solution treatment was performed at 1265 °C for 4 hours after heating at 1160 °C for 2 hours as a pre-homogenization to avoid any chance of incipient melting. A two-step aging treatment was performed, first at 1110 °C for 4 hours and second at 850 °C for 24 hours, both followed by air-cooling.

The heat-treated bars with [001] orientation were machined into creep specimens with gauge length of 50 mm and a gauge diameter of 8 mm. Constant load tensile creep tests were performed in ambient atmosphere at 975 °C under a stress of 255 MPa. Most of the creep tests were interrupted for microstructural examinations, whereas some tests were run to failure.

Heat treated sample was cut, ground and polished. The TEM slices were mechanically thinned down to a thickness of approximately 50 µm. Discs of 3 mm in diameter were punched out of the thin sheets and electro-polished with 90 pct alcohol and 10 pct perchloric acid electrolyte at -25 °C and 20 V, using a double-jet electropolisher.

Electron probe microanalyzer (EPMA) was used to characterize the distribution of B after heat treatment. The TEM was used to identify and analyse the phases using selected area diffraction (SAD) and energy dispersive spectrometer (EDS).

Results

GB Phase before Creep

Figure 1(a) is a typical EPMA mapping of B, from which the enrichment of B along the GB was observed after heat treatment. TEM micrograph of heat treated specimen (Figure 1(b)) shows that the GB was extensively decorated with blocky precipitates. According to the SAD of precipitate and EPMA mapping of B analysis, it can be confirmed that the blocky precipitate was $M_{23}(C,B)_6$ borocarbide. $M_{23}(C,B)_6$ borocarbide had an FCC structure with a cube-cube orientation relationship with the γ matrix:

$[001]_{M_{23}(C,B)_6} // [001]_{\gamma}, (100)_{M_{23}(C,B)_6} // (100)_{\gamma}$
No other borides were observed.

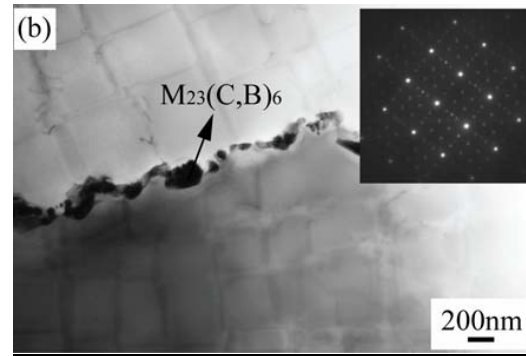
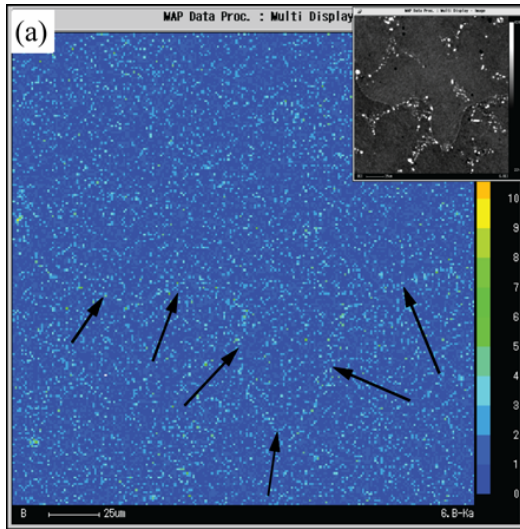


Figure 1. (a) EPMA mapping of B and (b) TEM micrograph of GB after heat treatment are shown.

Creep Testing

A typical creep curve for the present alloy at 975 °C / 255 MPa is shown in Figure 2. Microstructure at four stages during the creep was examined: after 2 h, 14 h, 40 h and 100 h of creep deformation (points A, B, C and D in Figure 2), respectively.

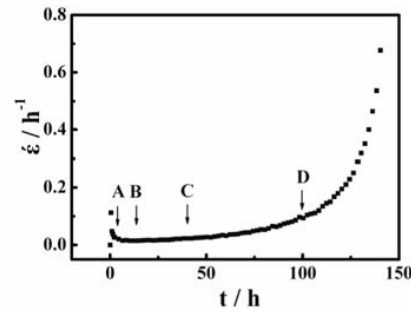


Figure 2. Creep curve of the experimental alloy at 975 °C / 255 MPa.

GB Phase at Primary Creep Stage

Although the creep time was very short at point A (2 h), a low magnification TEM micrograph (Figure 3(a)) shows that some dislocations were activated and have propagated in the γ matrix, and a continuous film-like precipitate was observed along GB. Closer examination at higher magnification (Figure 3(b)) indicated that the film-like precipitate was composed of many closely contacted fine nanosize particles. The SAD analysis demonstrated that these precipitates possessed the same crystal structure and they were Cr_5B_3 -type M_5B_3 boride with a body centered tetragonal structure. Based on the analyses of SAD, the orientation relationship between the boride and γ matrix can be determined as follows:

$[001]_{M_5B_3} // [001]_{\gamma}, (\bar{3}10)_{M_5B_3} // (0\bar{2}0)_{\gamma}$

Based on Figure 3(b), it can be also found that besides precipitation of M_5B_3 boride from γ matrix, phase transformation from $M_{23}(C,B)_6$ to M_5B_3 also occurred at the primary creep stage.

Microchemical analysis of the particles by TEM-EDS showed that

the composition of M_5B_3 boride contained a very high level of W, Mo and Cr (Figure 3(c)).

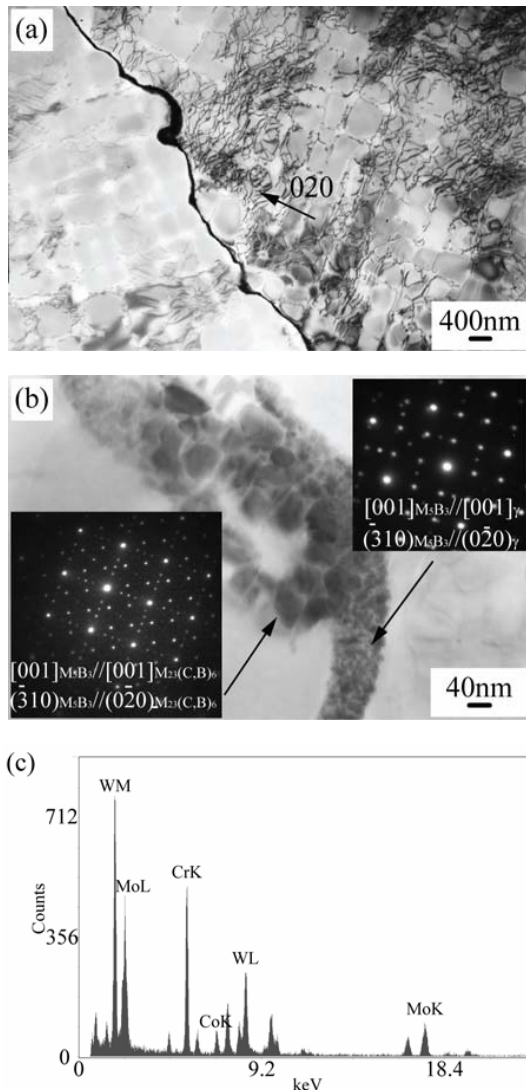


Figure 3. TEM micrographs of sample after 2 h of creep deformation at 975 °C / 255 MPa are shown. (a) Low-magnification GB micrograph. (b) High-magnification GB micrograph and the corresponding diffraction diagram of precipitate. (c) Typical EDS obtained from the precipitate in Figure 3 (b).

GB Phase at Initial Stage of Steady-State Creep Stage

Figure 4(a) is a representative TEM micrograph of sample after 14 h of creep deformation at 975 °C / 255 MPa. Although the dislocations grew continuously in the γ matrix passages with the progress of creep, the SAD analysis demonstrated that these precipitates were still M_5B_3 boride. No significant change for M_5B_3 boride in morphology was noticed by comparison of Figure 3 and Figure 4.

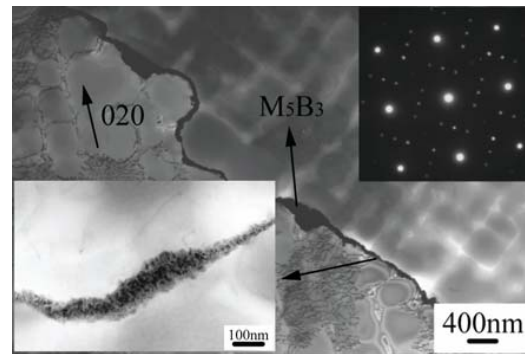
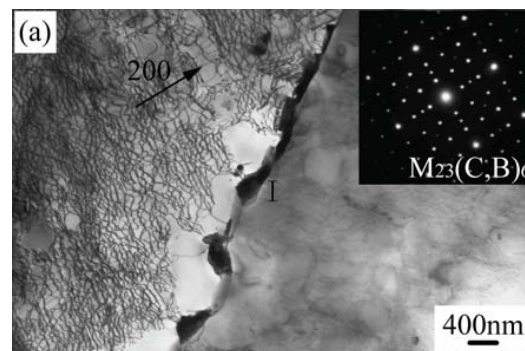


Figure 4. TEM micrographs of sample after 14 h of creep deformation at 975 °C / 255 MPa is shown.

GB Phase at Middle Stage of Steady-State Creep Stage

As the creep time reached 40 h (point C in Figure 2), the interfacial dislocation networks were formed. Meanwhile, the precipitate morphology at GB changed from continuous film-like to discrete block-like (Figure 5(a)). The analysis of SAD from the particles demonstrated that these precipitates were mainly $M_{23}(C,B)_6$ borocarbide. However, detailed SAD analysis (Figure 5(b)) indicated that small fraction of M_5B_3 boride may also exist at the GB, although M_5B_3 can not be found from Figure 5(a). This indicated that M_5B_3 boride was dissolving gradually during the steady-state creep stage.

According to the orientation relationship between M_5B_3 and γ matrix (Figure 3(b)), it can be inferred that the $[020]$ orientation of γ matrix from which M_5B_3 precipitated was parallel (or vertical) to direction II or III, as shown in Figure 5(b). And the $[020]$ orientation of γ matrix from which $M_{23}(C,B)_6$ precipitated was parallel (or vertical) to direction IV, as shown in Figure 5(b). Therefore, it was interesting to note that $M_{23}(C,B)_6$ particles were precipitated from the adjoining grain of the grain from which M_5B_3 precipitated and coarsening with the dissolution of the M_5B_3 during high temperature creep. By qualitative analysis the typical TEM-EDS results shown in Figure 3(c) and Figure 5(c), it can be found that the composition of $M_{23}(C,B)_6$ and M_5B_3 was similar, both containing a very high level of W, Mo and Cr. The major difference was the content of Cr in the $M_{23}(C,B)_6$ was higher compared to that in M_5B_3 .



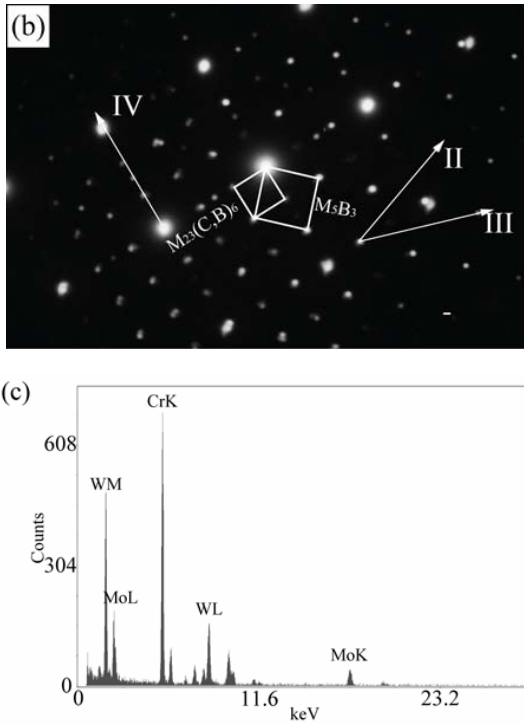


Figure 5. TEM micrographs of sample after 40 h of creep deformation at 975 °C / 255 MPa are shown. (a) GB micrograph and the corresponding diffraction diagram of precipitate. (b) Corresponding diffraction diagram of I position in Figure 5(a). (c) Typical EDS obtained from the precipitate in Figure 5(a).

GB Phase at Initial Stage of Tertiary Creep Stage

As the creep time reached 100 h (point D in Figure 2), the interfacial dislocation networks at the γ/γ' interface were locally damaged, some dislocations cut through the γ' particles, leaving dislocation couples in the γ' . A continuous film-like M_5B_3 boride was again observed along GB, besides block-like $M_{23}(C,B)_6$ borocarbide (Figure 6). From Figure 6, it can be found that the size of M_5B_3 boride was also in nanometer scale. However, it was unknown that if M_5B_3 boride at the tertiary creep stage was precipitated from γ matrix or transformed from $M_{23}(C,B)_6$.

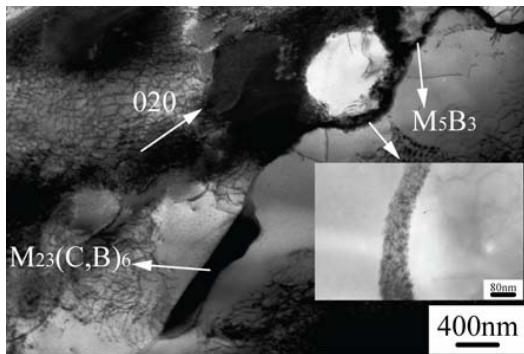


Figure 6. TEM micrographs of sample after 100 h of creep deformation at 975 °C / 255 MPa is shown.

Discussion

The most striking characteristics of GBs during 975 °C / 255 MPa creep deformation were GB phase evolutions. The precipitation of M_5B_3 from γ matrix and phase transformation from $M_{23}(C,B)_6$ to M_5B_3 occurred at the primary creep stage; precipitation of $M_{23}(C,B)_6$ from γ matrix and coarsening of $M_{23}(C,B)_6$ with the dissolution of M_5B_3 occurred at the steady-state creep stage; precipitation of M_5B_3 from γ matrix or phase transformation from $M_{23}(C,B)_6$ to M_5B_3 occurred at the tertiary creep stage.

Precipitation Mechanism of M_5B_3

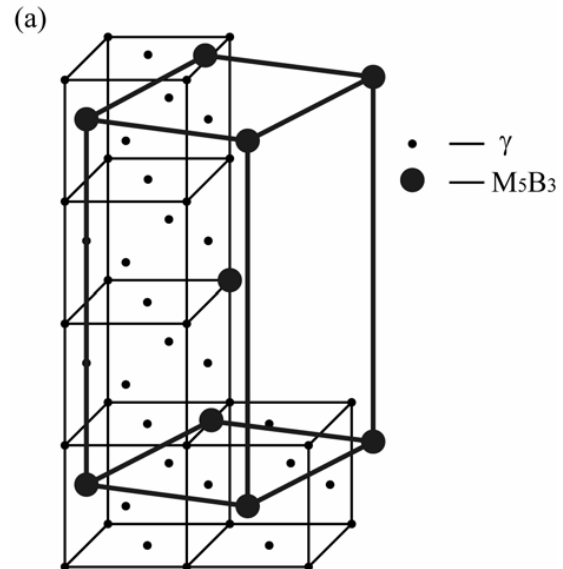
In general, for DS Ni-base superalloys containing high content of Cr, $M_{23}C_6$ or $M_{23}(C,B)_6$ usually formed at GBs during heat treatment or high temperature creep. However, precipitation of M_5B_3 borides was seldom observed at GBs [26], especially during high temperature creep.

In view of crystal structures of M_5B_3 and γ matrix, the lattice correspondence between M_5B_3 and γ matrix can be inferred based on Figure 3(b), as shown in Figure 7(a). The corresponding relationship between (001) atomic plane of M_5B_3 and (002) atomic plane of γ matrix is depicted in Figure 7 (b). According to Figure 7, the lattice parameters of M_5B_3 can be calculated (the lattice parameter of γ matrix a_γ approximately equals 3.6 nm):

$$a_{M_5B_3} = \sqrt{\left(\frac{3}{2}\right)^2 + \left(\frac{1}{2}\right)^2} a_\gamma = 1.58a_\gamma \approx 0.569nm \quad (1)$$

$$c_{M_5B_3} = 3a_\gamma = 1.08nm \quad (2)$$

The calculated lattice parameters of M_5B_3 coincided well with the data reported in handbook ($a_0=0.5699nm$, $c_0=1.088nm$) [27]. Therefore, Figure 7 probably exhibited correctly the lattice relationship between M_5B_3 and γ matrix.



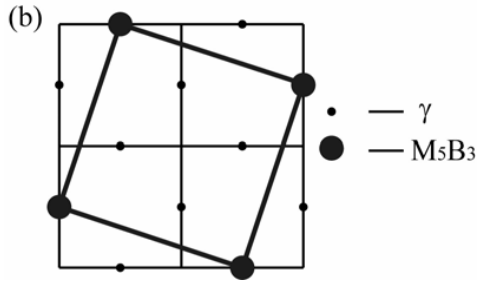


Figure 7. (a) Depiction of lattice correspondence between M_5B_3 and γ matrix. (b) Depiction of relationship between (001) atomic plane of M_5B_3 and (002) atomic plane of γ matrix.

According to Figure 7, eight corner atoms of M_5B_3 were all located at γ matrix face centered sites, and the center atom of M_5B_3 boride was located at γ matrix cornered site. It is well accepted that B atoms stay at octahedral interstitial sites of γ lattice as schematically illustrated in Figure 8. Therefore, the formation of M_5B_3 required B atoms diffusing from the octahedral interstitial sites to the face centered sites. In general, this diffusion was difficult to occur, thus M_5B_3 was seldom found after heat treatment. However, it has been well established that DS Ni-base superalloys deformed by operation of $a/2\langle 101 \rangle$ dislocations on $\{111\}$ planes in γ matrix during high temperature creep when the loading direction was parallel to $\langle 001 \rangle$. This activated the movement of matrix atoms, from site 2 to site 1 in Figure 8. During this process, B atoms may diffuse easily under an applied uniaxial tensile stress, from site 3 to site 2. It seemed that the nucleation kinetics of M_5B_3 was rather rapid during the primary creep stage, since extensive nanosize particles were observed in Figure 3. High strain rate promoted this process.

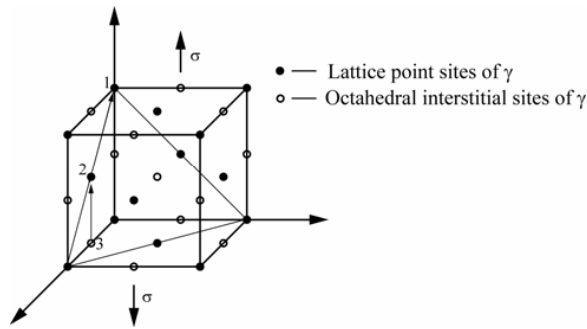


Figure 8. Illustration of B atoms diffusion under uniaxial tensile stress is shown.

Phase Transformation Mechanism from $M_{23}(C,B)_6$ to M_5B_3

The strain rate at the initial stage of primary creep and tertiary creep was very high. Therefore, many dislocations were activated and propagated in γ matrix at the GBs quickly. However, due to presence of precipitate particles, these dislocations piled up at the γ /precipitate interface, which resulted in high strain energy at GBs. In general, the larger the size of the precipitate, the higher strain energy was expected. Thus, in order to decrease strain energy, blocky $M_{23}(C,B)_6$ borocarbide might transform into nanosize M_5B_3 boride, whose composition was similar to $M_{23}(C,B)_6$, but size was apparently smaller than that of $M_{23}(C,B)_6$. This phase

transformation can avoid stress concentration at the $\gamma/M_{23}(C,B)_6$ interface, retard the onset of creep cavitation and rupture. Generally, this phase transformation can not occur in B-free alloys. Therefore, it can be inferred that avoiding stress concentration by phase transformation was perhaps one of the strengthening mechanisms of B.

Precipitation and Coarsening Mechanisms of $M_{23}(C,B)_6$

The strain rate at steady-state creep stage was very low, creep process was governed by thermal diffusion. The formation of M_5B_3 boride caused W, Mo, and Cr depletion at GBs during primary creep stage. Accompanied by high temperature creep, stress may induce the flow of W, Mo, Cr atoms from intragranular regions towards GBs. Considering the large diffusion rate of Cr and its small atom radii, it may be expected that diffusion of Cr was more pronounced. Consequently, $M_{23}(C,B)_6$ borocarbide would form in view of strong segregations of C and B atoms at the GBs.

Based on above results, it can be found that $M_{23}(C,B)_6$ borocarbide not only formed, but also coarsened with the dissolution of M_5B_3 boride driven by the decrease in interface energy at steady-state creep stage. The reason why nanosize M_5B_3 boride did not coarsen was that: The lattice misfit between M_5B_3 and matrix was non-coherent, however, according to Reference [28], it can be inferred that the atomic configuration of $\{111\}$ planes in the $M_{23}(C,B)_6$ and matrix lattices was similar and the lattice misfit between $M_{23}(C,B)_6$ and matrix was very small and be partially coherent. Hence, the elastic energy that coarsening of M_5B_3 per unit volume overcame was higher than that of $M_{23}(C,B)_6$.

Conclusion

The following conclusions can be drawn from analysis of the results obtained in this work:

(1) During high temperature creep, applied tensile stress promotes diffusion of B atoms from octahedral interstitial sites of γ lattice to face centered sites of γ lattice, resulted in precipitation of many nanosize M_5B_3 boride particles at GBs. The orientation relationship between M_5B_3 boride and γ matrix is:

$$[001]_{M_5B_3} // [001]_{\gamma}, (\bar{3}10)_{M_5B_3} // (020)_{\gamma}$$

(2) The strain rate is found to have a significant influence on the evolution of GB phase during high temperature creep. High strain rate induces phase transformation from $M_{23}(C,B)_6$ to M_5B_3 , and low strain rate induces precipitation of $M_{23}(C,B)_6$ from γ matrix and coarsening of $M_{23}(C,B)_6$ with the dissolution of M_5B_3 .

Acknowledgement

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