Controllable two-scale network architecture and enhanced mechanical properties of (Ti$_5$Si$_3$ + TiBw)/Ti6Al4V composites

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Recent studies reveal that titanium matrix composites (TMCs); especially, discontinuously reinforced titanium matrix composites (DRTMCs) reinforced with ceramic particles are gaining widespread attention, due to their outstanding mechanical properties. DRTMCs have become promising materials for aerospace, automotive and military applications due to their isotropic properties, high specific strength, high modulus, good wear resistance and high temperature durability$^{1,2}$. Among various manufacturing methods of DRTMCs, a combination of reaction hot pressing (RHP) and in-situ technology has been widely investigated and developed, because of its near net-shape processing and low cost, especially its ability to make composites obtain superior mechanical properties, clean interface and strong interface bonding$^6$. In considerable amount of literatures studying DRTMCs, in-situ TiB whisker (TiBw)$^4$-6 and TiC particles (TiCp)$^7,8$ have been unanimously considered as the best reinforcements, due to their desirable properties, such as high strength, modulus, good chemical stability and similar coefficient of thermal expansion (CTE) with Ti matrix. In addition, hybrid reinforcements, such as TiBw and TiCp$^9$, TiBw and La$_2$O$_3$$^{10}$ were extensively utilized to pursue superior mechanical properties. However, Ti$_5$Si$_3$ particles have rarely been used as reinforcement during the fabrication of TMCs via in-situ methods. Ti$_5$Si$_3$ phase possesses many excellent properties, such as high melting point (2130 °C), low density (4.26 g/cm$^3$), retaining strength up to 1200 °C, especially good oxidation and creep resistance at elevated temperatures$^{11}$. Furthermore, the CTE of Ti$_5$Si$_3$ is also close to that of Ti alloys$^{12}$. Ti$_5$Si$_3$ can potentially be regarded as an attractive wear-resistant and corrosion-resistant material due to its inherent high hardness, unique chemical composition and strong covalent bonds$^{13}$. Therefore, Ti alloy matrix composites reinforced with Ti$_5$Si$_3$ particles are potential candidates for high temperature applications.

Inadvertently, previous investigations have usually pursued DRTMCs with a homogeneous microstructure. Unfortunately, the homogeneous composites have always showed inferior ductility, even extreme brittleness for those fabricated by conventional power metallurgy, except for limited improvement in strength. In recent years, on the basis of Hashin-Shtrikman (H-S) theory, Huang et al.$^{1,4,5}$ designed and fabricated a series of DRTMCs with a quasi-continuous network structure of reinforcements using low-energy milling and RHP processes combined with in-situ technology. Sufficient results illustrate that the network architecture is beneficial to inspire
strengthening effect of reinforcement and toughening effect of matrix alloy in DRTMCs. However, the large network structure limits reinforcement fraction, which restricted further improvement in the strength of composites. Microstructural architectures with material specific design have proven to be powerful in nature and engineering applications. Natural materials, such as shells, tooth, and bone, exhibit multi-scale hierarchical structures spanning from microscopic to macroscopic length scales and show significantly improved mechanical properties (e.g. strength and toughness) compared to base materials. For this reason, there are increasing efforts to artificially create or imitate natural materials’ structures in composites.

In this study, not only the quasi-continuous network architecture and the best TiBw reinforcement were adopted, but also low-cost micro Si particles were used to achieve Ti5Si3 reinforcement in the novel lower secondary-scale structure by precipitation, in order to further improve the mechanical properties of the composites. Ti6Al4V alloy as a typical one of dual-phase titanium alloys was selected as the matrix due to its superior mechanical properties and extensive usage. Therefore, in the present work, the aim is to tailor the distribution of Ti5Si3 and TiBw reinforcements, so as to create a controllable two-scale network structure and enhance the mechanical properties of titanium alloy matrix composites.

Results and Discussion

Microstructure. Figure 1 shows the X-ray diffraction patterns of the as-sintered composites with different Ti5Si3 volume fractions fabricated at 1300 °C for 1.5 h. It presents that the as-sintered composites mainly consist of Ti, Ti5Si3 and TiB phases. Moreover, no Si and TiB2 phases were detected. This result demonstrates that the Ti5Si3 volume fractions fabricated at 1300 °C for 1.5 h.

Figure 2a shows SEM micrograph of the 4vol.%Ti5Si3/Ti6Al4V composite. It can be seen from Fig. 2a that the microstructure of the 4vol.%Ti5Si3/Ti6Al4V composite is similar to that of the Ti6Al4V alloy, due to the absence of the first-scaled TiBw/Ti6Al4V network structure. The observed microstructure is a typical α + β lamellar structure, which belongs to the Widmanstätten microstructure. When α + β two-phase Ti alloys are cooled slowly (furnace cooling) from a temperature above the β transus, the Widmanstätten microstructure is usually obtained. In addition, β grains with large size of about 1 mm replace Ti6Al4V particle with 150 μm, which indicates that the Ti6Al4V particles entirely merged, while Si particles entirely solid soluted into the β phase. During sintering process, the Si element totally diffused into the β phase. Figure 2b shows needle-like Ti5Si3 precipitated and distributed in the β phase, which was around the α phase. This phenomenon can be interpreted as: during furnace cooling, the solubility of Si element in the β phase decreases with decreasing temperature; the fraction of β phase also decreases due to the phase transition (β → α).

Figure 2c–h show SEM micrographs of the (Ti5Si3 + TiBw)/Ti6Al4V composites with different fractions of Ti5Si3 reinforcement. Only Ti, Ti5Si3, and TiB phases were observed, which is consistent with the XRD result (Fig. 1). It can be clearly seen from Fig. 2c that the TiBw reinforcement distributed around the Ti6Al4V particles, and formed an analogous “grain boundary” structure. The “grain” size is about 150 μm equal to that of the Ti6Al4V raw material. On the whole, the TiBw reinforcement formed the regular first-scale network structure (FSNS, Fig. 2c), as reported in the TiBw/Ti6Al4V composite. On the lower scale, few Ti5Si3 particles precipitated in the β-Ti phase, which distributed around the α-Ti phase, and formed the secondary-scale network structure (SSNS, Fig. 2d), due to the first-scale network structure. Since the two reinforcements distributed in different scales, these two-scale networked composites are different from the conventional hybrid-reinforced composites with homogeneous dispersion of two reinforcements. The lower SSNS of Ti5Si3 reinforcement can strengthen the interior matrix particles effectively, due to the precipitation of fine Ti5Si3 needles. Moreover, the SSNS can increase deformation compatibility, which is beneficial to the strength and ductility of composites. Previous results have indicated that the precipitation of Si (silicide) leads to an increase in the strength...
of (TiB + La₂O₃)/Ti composites. When the fraction of Ti₅Si₃ reinforcement is increased to 4 vol.%, as shown in Fig. 2f, more needle-like Ti₅Si₃ particles precipitated in the β-Ti phase compared with that in Fig. 2d. Figure 2e,f show some fine and needle-like Ti₅Si₃ particles formed in the β-Ti phase. The possible reasons for this phenomenon are as follows. According to the Ti-Si phase diagram, the solubility of Si in β-Ti phase is around 3 wt.% at 1300 °C, while it nearly does not solid solute in α-Ti at room temperature. Therefore, the entire Si can dissolve into β-Ti phase at 1300 °C. However, when cooling from 1300 °C, the volume fraction of β-Ti phase decreases due to phase transformation from β-Ti to α-Ti. The saturation of Si element leads to the precipitation of Ti₅Si₃ phase in the β-Ti phase. It is worth pointing out that Si element is considered to work as a β-stabilizer in Ti alloys. Therefore, the volume fraction of β-Ti can be slightly increased, which may prove to be beneficial to the ductility of composites. The needle-like Ti₅Si₃ reinforcement distributed in the β-Ti phase around the near equiaxed α-Ti phase, and formed the secondary-scale network structure (SSNS). Gu et al. studied the mechanical alloying of Ti-Si powder mixture using high-energy ball milling at ambient temperature. A significant increase in solid solubility of Si in Ti was achieved by mechanical alloying. The shrinkage of Ti lattice was caused by diffusion of Si atoms into Ti. In a previous literature, La atoms would solid solute into β-Ti firstly, followed by their precipitation from β-Ti during the phase transformation from β-Ti to α-Ti. Following which the La atoms reacted with oxygen inside the matrix alloy, forming fine and homogeneously distributed La₂O₃ particles on the grain boundary of the titaniuim alloy matrix.

Noticeably, comparing Fig. 2a with Fig. 2e, a remarkable difference in the Ti₅Si₃ reinforcement was observed between the monolithically reinforced composite and the composite with a two-scale microstructure. In the (Ti₅Si₃ + TiBw)/Ti₆Al₄V composite, a small quantity of needle-like Ti₅Si₃ reinforcements distributed near the TiBw reinforcements. Most of the Ti₅Si₃ particles were observed in the β-Ti phase of the SSNS. Li et al. found that few silicides are distributed in the α platelet. Previous results have shown that Nb₅Si₃ particles precipitated at the γ/γ interface and the γ/α₂ interface. However, entire Ti₅Si₃ particles precipitated in the β-Ti phase in the Ti₅Si₃/Ti₆Al₄V composite. Owing to the absence of distribution of the TiBw reinforcement in the first-scale network region (FSNR), the formation of the FSNS did not take place.

Besides the central β phase of the Ti₆Al₄V particles, few Ti₅Si₃ particles were observed around the TiBw reinforcements (Fig. 2e). This phenomenon may be due to the addition of TiBw, which increased the amount of defects. Sun et al. prepared TiAl alloys using non-consumable electrode arc melting in an argon atmosphere. Ti₅Si₃ particles preferred to precipitate at stacking faults. The growth of needle-like Ti₅Si₃ particles at the γ/γ boundaries was controlled by interfacial diffusion of Si, with an incoherent γ/γ interface. In addition, the Ti₅Si₃ phase distributed around the TiBw reinforcement with a size of around 1 μm; whereas, the size of the Ti₅Si₃ precipitation in the β-Ti phase is about 400 nm. Si element prefers to diffuse to areas with defects and precipitate during cooling process.
However, when the fractions of Ti5Si3 reinforcement is as high as 8vol.%, as shown in Fig. 2h, lesser Ti5Si3 particles precipitated in the β-Ti phase of the central Ti6Al4V particles as compared to that in Fig. 2f. In contrast, coarse Ti5Si3 particles formed in the vicinity of TiBw reinforcements. This may be attributed to the limited solubility of Si in the β-Ti phase. Redundant Si reacted with Ti to in-situ synthesize Ti5Si3 particles. This increased the local volume fraction of ceramic phases in the FSNR, whose energy was increased. Therefore, Si element in the β-Ti phase prefers to nucleate and grow in the FSNR during cooling process. The connected and coarse Ti5Si3 particles are certainly harmful to the properties of the composites23.

According to the Ti-Si phase diagram19, the silicide in the equilibrium microstructure of Ti-Si should be Ti5Si3. However, there are many contradictions in previous literatures, about the most thermodynamically favorable structure of precipitation at low concentrations of Si26–28. Poletaev et al.26 applied a recently developed thermodynamic model to predict the structure of Ti-Si precipitations in the α-Ti matrix. They discovered that formation of the Ti5Si3 phase is more favorable than that of the Ti3Si phase in contradiction. The theoretical framework was confirmed with experimental investigations of microstructure.

To further determine the crystal structure of the precipitated particles, TEM analysis was conducted. The corresponding morphologies and selected area diffraction (SAD) patterns are shown in Fig. 3. Figure 3a further confirmed that the precipitations are Ti5Si3 phase rather than Ti3Si phase. Ti5Si3 phase is a common precipitation in Si-containing Ti alloys. SAD analysis in Fig. 3b found that the Ti5Si3 particle has certain crystal relationships corresponding morphologies and selected area diffraction (SAD) patterns. This is also in agreement with the previous results showing that very fine silicides were formed at α/β interfaces23.

Combing with the above analysis, the formation and distribution mechanisms of secondary-scale Ti5Si3 phase can be illustrated in Fig. 4. During sintering process, Si completely diffused into the Ti6Al4V matrix from the TiBw whisker, a large proportion of Si element distributed in the residual α-Ti phase. Ti5Si3 needles precipitated with the decrease in temperature.

In order to adjust the precipitation sites of the Ti5Si3 phase, (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composites were heat treated at 990°C, 1100°C and 1200°C for 40min. SEM micrographs of the heat treated composites are shown in Fig. 5. After heat treatment, there is no change in the microstructure and distribution of the TiBw reinforcement, whereas there is a clear change in the microstructure of the Ti6Al4V matrix and Ti5Si3 reinforcement. The bright β phase in the Ti6Al4V matrix is converted into the transformed β microstructure (βT), which consists of the residual β phase, martensite α' phase formed in the WQ process. Furthermore, the contrast between the primary α phase and the transformed β microstructure decreases with increasing quenching temperatures, which demonstrates that the increasing fraction of the martensite and the decreasing fraction of β phase in βT. The increased fraction of the martensite is beneficial to the hardness and strength of composites29. Comparing Fig. 5 with Fig. 2, the volume fraction of βT in the heat treated composite is much higher than that of the stable β phase in the as-sintered composite. Figure 5 also indicates that the volume fraction of βT increases with increasing WQ temperatures. While, the fraction of Ti5Si3 reinforcement distributed in the vicinity of TiBw reinforcement decreased with increasing quenching temperatures. When the heat treatment temperature reached 1200°C, Ti5Si3 phase disappeared near TiBw reinforcement and in the SSNS. The size of the Ti5Si3 precipitation in the as-sintered composites is around 400 nm, while some Ti5Si3 precipitation in the heat treated composites even increased to 2μm or decreased to 0. Therefore, heat treatment can control the size of Ti5Si3 precipitation. Li et al.18 reported that both particle size and volume fraction of silicides increase with increasing thermal exposure temperature.
To detect the distribution of Si in the composites, EDX (element surface scanning) analysis was carried out, as shown in Fig. 5. According to the results of element surface scanning, the fraction of Si element in the vicinity of TiBw reinforcement decreased with increasing WQ temperatures. A small quantity of the Ti5Si3 phase grew and much of the Si element dissolved into the $\beta_T$ when the heat treatment temperature was 990 °C. When the temperature was increased to 1100 °C, more Ti5Si3 phase became fine and acicular. When the temperature reached 1200 °C, Si was not detected in the vicinity of TiBw reinforcement. This indicates that the Si element entirely dissolved into the $\beta_T$.

**Mechanical properties.** Figure 6a shows the typical tensile stress-strain curves of the as-sintered Ti6Al4V alloy and composites at room temperature. It is obvious that the strength of the (Ti5Si3 + TiBw)/Ti6Al4V
composites significantly increased when compared to those of the monolithically reinforced Ti₅Si₃/Ti₆Al₄V composite and the as-sintered Ti₆Al₄V alloy. For example, the ultimate tensile strength of the (2vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composite increased to 1130 MPa from 1030 MPa, when compared to that of the monolithically reinforced Ti₅Si₃/Ti₆Al₄V composite. The yield strength of the (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composite increased to 1004 MPa from 769 MPa, compared with the Ti₆Al₄V alloy. Moreover, the elongation of the (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composite was retained at 4%, which is much higher than that of the Ti₅Si₃/Ti₆Al₄V composite. This phenomenon can be attributed to the two-scale network microstructure with high shear resistance. When compared to the STA Ti₆Al₄V alloy, the strength of the composites with a two-scale network decreased significantly. However, the elongations show a slight difference. In addition, it is clear that the strength and elongation of the composites increased with increasing volume fraction of Ti₅Si₃ from 2vol.% to 4vol.%. The increased elongation is attributed to more Ti₅Si₃ precipitations with nano-size in the β-Ti phase, which increase the deformation compatibility of the composites. The deformation compatibility of network unit was improved, and the moving dislocations can bypass the particles instead of shearing them, and then the elongation to fracture can be improved. However, the strength and elongation of the (8vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composite were inferior over the other composites. The composite did not undergo any plastic deformation, resulting in brittle fracture. This is consistent with the high local volume fraction of reinforcement in the FSNS and the coarse Ti₅Si₃ particles in the FSNR. In the fabrication of metal matrix composites, reproducibility is a significant concern. Figure 6b shows reproducibility of the (Ti₅Si₃ + 3.4vol.%TiB₆) composites with different Ti₅Si₃ fractions. In the curves, 3-1, 3-2, 3-3 represents three similar tensile curves of the (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆) composites, respectively. From the tensile curves, it is obvious that the tensile properties of the composites with a two-scale network structure are reproducible.

In order to further investigate the contribution of the two-scale network distribution to the tensile properties, the (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composites were compared with the (TiB₆ + TiCp)/Ti₆Al₄V and TiB₆/Ti₆Al₄V composites with a one-scale network structure. Firstly, compared with the monolithically reinforced 5vol.%TiB₆/Ti₆Al₄V composite, the ultimate tensile strength and yield strength of the (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composite increased by 70 MPa and 64 MPa. Moreover, the elongation increased from 3.6% to 4%. Secondly, in the (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composite, the tensile strength and elongation increased from 1129 MPa and 2.4% to 1160 MPa and 4%, as compared to the hybrid-reinforced 3vol.%TiB₆ + TiCp)/Ti₆Al₄V composite. All the results show that the tensile properties of the (Ti₅Si₃ + TiB₆)/Ti₆Al₄V composite with a two-scale network architecture are superior over those of the Ti₅Si₃/Ti₆Al₄V, TiB₆/Ti₆Al₄V and hybrid-reinforced (TiB₆ + TiCp)/Ti₆Al₄V composites with a one-scale structure. When compared to the composites fabricated by casting, the present strengthening effect can be viewed as a superior strengthening effect, because not only the strength but also the elongation was improved. This may be attributed to the combination of the FSNS and SSNS. The fine Ti₅Si₃ needles in the β-Ti phase within the Ti₆Al₄V matrix can strengthen the softer β-Ti phase and provide dispersion hardening acting as a barrier for the dislocation movement. Moreover, TiB₆ distributed in the FSNS strengthened the grain boundary of the composites. Additionally, the formed fine SSNS can effectively increase deformation compatibility and prohibit necking during tensile deformation, resulting in elongation improvement.

Table 1 shows the tensile properties of the as-sintered and heat treated (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composites tested at room temperature, in order to further demonstrate the effect of heat treatment to adjust the distribution of Ti₅Si₃ precipitation. With increasing quenching temperatures, the tensile strength of the heat treated composites increased. However, compared with that of the as-sintered composites, the tensile strength after water quenching decreased significantly. Moreover, in those composites, brittle fracture took place immediately. The decreased strength and brittleness may be due to the excessive martensite or hardness, which is generated by the high quenching temperatures.

In order to evaluate the evolution of properties, compressive tests were performed on the heat treated composites which showed brittle fracture under tensile test conditions. Figure 7 shows the variations in the compressive strength of the composites after water quenching. Furthermore, the compressive properties including yield compressive strength (YCS), ultimate compressive strength (UCS) and fracture strain (ε) of the as-sintered and heat treated (4vol.%Ti₅Si₃ + 3.4vol.%TiB₆)/Ti₆Al₄V composites tested at room temperature are shown in Table 2. It

![Figure 6](image-url)
can be concluded that the in-situ synthesis of TiBw reinforcement and precipitation of Ti5Si3 phase significantly increased the strength of the as-sintered composite, while decreasing the plasticity correspondingly. The YCS and UCS of the heat treated composites increased with increasing quenching temperatures. The YCS and UCS can be increased to 1687 MPa and 1753 MPa from 1225 MPa and 1402 MPa, respectively. The fraction of the βT in the matrix and the saturation of Si element increased with increasing quenching temperatures, which led to a significant improvement in strength. The other reason is the presence of fine Ti5Si3 precipitation in the βT. Li et al.31 reported that the influence of heat treatment played an important role in enhancing the strength of the (TiB + La2O3)/Ti composites. After thermal exposure, the strength of the heat treated specimens increased due to the precipitation of Ti3Al and silicides. When compared to the monolithically reinforced 5vol.%TiBw/Ti6Al4V and 5vol.%TiCp/Ti6Al4V composites, the yield strength of the (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composite increased by 28.9%32 and 44.1%33, respectively. The first-scale TiBw reinforcement and the secondary-scale Ti5Si3 particles distributed in different regions. The increased reinforcement fractions enhanced the composites strength. Additionally, fine Ti5Si3 particles can bear higher stress than TiB whiskers. Consequently, the strengthening mechanisms of the composites may be mainly attributed to the “grain boundary” strengthening from first-scale networked TiBw reinforcement and dispersion strengthening from the secondary-scale networked Ti5Si3 reinforcement. Furthermore, according to the literatures34,35 and our experience, the composites with a two-scale network structure display superior mechanical properties and oxidation resistance at high temperatures.

Ti alloys are unlikely to have adequate wear resistance for using as structural materials without surface treatment. To overcome the poor wear resistance of Ti alloys, particulate-reinforced TMCs are a suitable solution. A previous study showed that the (TiB + TiC)/Ti composite displays improvement in wear resistance46. The coefficients of friction (COFs) of the Ti6Al4V alloy, the as-sintered and heat treated (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composites under contact load of 10N are shown in Fig. 8. The average COFs and mass loss of the samples are listed in Table 3. It can be seen from Fig. 8 that all the samples showed a shorter and
unstable friction period (about 300s) in the initial stage of friction tests. The COF underwent an initial fluctu-
ation followed by a steady tendency. When compared to the as-sintered Ti6Al4V alloy, the average COF of the
as-sintered (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composite decreased by 7.0%. This suggests the addition of
reinforcements resulted in a change in wear mechanism and a decrease in COF compared with the unreinforced
matrix alloy. On one hand, the strength and hardness of two-scale network structured composites are higher
than the Ti6Al4V alloy. Especially, the finer Ti5Si3 particles improved the hardness and strength of the softer
β-Ti matrix. The adhesive effect of the composites against their counterparts was limited and the proportion of
asperities was reduced during friction process. On the other hand, the increased flash temperature resulted in the
softening of the Ti6Al4V matrix. However, the introduction of the reinforcements improved the high temper-
ature strength of the composites. Therefore, the scuffing effect of the composites against their counterparts was
alleviated. The mass loss of the as-sintered composite is lower than that of the Ti6Al4V alloy. It means that the
composite has superior wear resistance over the matrix alloy, due to the reinforcement with high hardness.

It is worth pointing out that the mass loss of the heat treated composites is slightly higher than that of the
as-sintered composites. The mass loss of the heat treated composites decreased with increasing quenching tem-
peratures. When the heat treatment temperature was 990 °C, some growing Ti5Si3 phases resulted in a decrease
in wear resistance. When the temperature was increased to 1100 °C, more Ti5Si3 phases became fine and acicular,
which is beneficial to the wear resistance of the composites. At 1200 °C, the excessive martensite resulted in high
hardness and Si element entirely dissolved into the β-Ti strengthened the matrix alloy. Therefore, the wear resist-
ance of the composite after water quenching at 1200 °C is superior over the as-sintered composite.

Figure 8. Friction coefficient of the Ti6Al4V alloy, the as-sintered and heat treated (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composites.

| Materials | Mass loss (mg) | COF  |
|-----------|----------------|------|
| Ti6Al4V   | 7.98 ± 0.01    | 0.298|
| (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V | 6.65 ± 0.04 | 0.277|
| 990 °C WQ | 7.71 ± 0.02    | 0.286|
| 1100 °C WQ | 6.73 ± 0.02  | 0.266|
| 1200 °C WQ | 6.30 ± 0.03  | 0.280|

Table 3. Mass loss and COF for the Ti6Al4V alloy, the as-sintered and heat treated (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composites.
Materials and Methods

To manufacture the composites, spherical Ti6Al4V powders with an average particle size of 150 μm, prismatic TiB2 powders (3 μm) and fine Si powders (3 μm) were used. At first, Ti6Al4V, TiB2 and Si powders were mixed using low-energy ball milling for 5 h at 230 rpm under an argon atmosphere. The ratio of milled media to material is 5:1. Afterwards, the blended powder mixtures were hot pressed in vacuum (10−2) at 1300 °C for 90 min under a pressure of 20 MPa. TiBw reinforcements around the Ti6Al4V matrix were synthesized via in situ reaction between TiB2 and Ti. The reaction can be described as the equation (1)4. Ti5Si3 reinforcements in the Ti phase were achieved according to the equation (2) in the Ti-Si phase diagram10.

\[Ti + TiB_2 = 2TiB\] (1)

\[Ti + Si \rightarrow Ti_5Si_3\] (2)

According the above processes, 4vol.%Ti5Si3/Ti6Al4V, (2vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V, (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V and (8vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composites were prepared. In order to adjust the distribution of Ti5Si3 particles and investigate the contribution of the unique structure to the mechanical properties, the (4vol.%Ti5Si3 + 3.4vol.%TiBw)/Ti6Al4V composites were water quenched (WQ) after solid solution strengthening at 990 °C, 1100 °C, and 1200 °C for 40 min.

A scanning electron microscope (SEM; ZEISS Supra 55 SAPHIRE) instrument and a transmission electron microscope (TEM; TecnaiF2F30) instrument were used to observe the microstructure of the fabricated composites. The tensile and compressive properties of the composites were carried out using an Instron-5569 universal testing device, at a crosshead speed of 0.5 mm/min. The tensile specimens were machined with the gauge size of 15 mm × 5 mm × 1.5 mm, and compressive specimens had dimensions of ø4 mm × 6 mm. At least five tests were performed on each condition and the average values were used.

The wear properties of the Ti6Al4V alloy and the fabricated composites were evaluated using a pin-on-disk tribometer (Teer POD-I). During the tests, the specimens were rotated against a stationary GCr15 steel ball of 5 mm diameter at the speed of 200 r/min for 1800 s under a contact load of 10N. The radius of the wear track was about 5 mm. The friction coefficient was continuously recorded during the tests. All tests were conducted in air (temperature 20 °C, humidity 40% RH). Before and after each wear test, the specimens were cleaned with acetone and then dried. The mass loss of the specimens was measured using an electronic balance with an accuracy of 0.01 mg. Each data point represents the average value of three test results.

Conclusions

To sum up, novel two-scale network structured (Ti5Si3 + TiBw)/Ti6Al4V composites were successfully in situ fabricated using low-energy milling and reaction hot pressing. A summary of the key findings in this paper are:

1. TiBw was synthesized around the Ti6Al4V particles, and formed the first-scale network structure. Ti5Si3 precipitated in the β-Ti phase around the α-Ti phase, and formed the secondary-scale network architecture.
2. Heat treatment can adjust the size and distribution of the Ti5Si3 reinforcement. The ratio of the TiBw and Ti5Si3 reinforcements distributed in different scales can be controlled by adjusting the proportions of TiBw and Si raw materials.
3. Ti5Si3 phase precipitated in the β-Ti phase has certain crystal relationships with the β-Ti phase: [103]Ti5Si3// [110]β-Ti, (321)Ti5Si3//[211]β-Ti.
4. The composites with two-scale network architecture exhibit superior mechanical properties over the composites with one-scale architecture and the Ti6Al4V alloy.

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