

Mechanical properties and wear resistance of medium entropy Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composites

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Abstract

The Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC (volume fraction of TiC, 10%) composites were synthesized in combination of ball milling and spark plasma sintering (SPS) in the present work. Mechanical properties and wear resistance of the Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composites were individually investigated. It was found that TiC particles homogenously distributed in the Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composite after being sintered at 1373 K for 15 min. Meanwhile, grain refinement was observed in the as-sintered composite. Compared with the pure Fe₄₀Mn₄₀Cr₁₀Co₁₀ medium entropy alloy (MEA) matrix grain, addition of 10% TiC particles resulted in an increase in the compressive strength from 1.571 to 2.174 GPa, and the hardness from HV 320 to HV 872. Wear resistance results demonstrated that the friction coefficient, wear depth and width of the composite decreased in comparison with the Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA matrix. Excellent mechanical properties and wear resistance could offer the Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composite a very promising candidate for engineering applications.

Key words

TiC/Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composites; mechanical properties; wear resistance; spark plasma sintering

1 Introduction

High entropy alloys (HEAs)/medium entropy alloys (MEAs) have been accepted great interest in research as a new class of metallic alloys. The unique feature of the MEAs/HEAs is the simple solid solution (FCC/BCC/ HCP) phase comprising of multiple equiatomic or near-equiatomic elements with high configurational entropy. This is significantly different from the conventional alloys consisting of principal elements and minor elements and often crystallized in the formation of solid solution and intermetallic phases [1,2]. Although a number of advantages of HEA/MEAs have been well recognized, the drawbacks of HEA/MEAs including relatively low strength and poor wear resistance, have limited their applications as strong candidates of structural materials. Therefore, the attempts to improve the strength are becoming important in industrial applications.

It has been confirmed that metal matrix composites (MMCs) can offer outstanding mechanical properties and superior wear resistance [3–5]. The alloys reinforced by particulates, such as SiC [6], TiB₂ [7], WS₂ [8], B₄C [9] and Al₂O₃ [10], have been used in various applications. The introduction of particulates into a single-phase matrix of HEA/MEAs can be an effective approach to improve mechanical properties, in particular the wear resistance. Recently, the effect of reinforcement particles on the microstructure and wear resistance of the HEA/MEAs has been the topics of a number of investigations. LIU et al [11] introduced brittle σ and μ phases into a FCC-structured CoCrFeNi MEA through addition of Mo particles. The particle-strengthened alloy exhibits a tensile strength as high as 1.2 GPa and a tensile elongation of ~19%. ROGAL et al [12,13] pointed out that 5% Al₂O₃ (mass fraction) and 5% SiC (mass fraction) particles were uniformly distributed in CoCrFeMnNi HEA, which delivered compressive yield stress and compressive strain as high as 1480 MPa and 1600 MPa, and 31% and 6.5%, respectively. COLOMBINI et al [14] studied the FeCoNiCrAl high entropy alloy reinforced by reactive silicon carbide (SiC) particles through a powder metallurgy synthetic route. The reactive sintering produced a single isomorphic BCC structure and formed stable carbides and silicide particles, which increased the hardness of the alloy. CoCrFeNi self-lubricating composite reinforced by graphite and MoS₂ powder has good mechanical and tribological properties [15].

Fe₄₀Mn₄₀Cr₁₀Co₁₀ is a newly developed MEA with the features of excellent elongation (57.7%) but low yield strength (213 MPa) [16]. It is interesting to study the effect of TiC particles on the reinforcement of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA, in particular the wear resistance. In the present work, Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composites were processed by spark plasma sintering (SPS) to obtain fully dense materials at relatively low temperature and a short time. Mechanical properties and wear resistance of the FCC-type Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composite were assessed in association with the microstructure. The strength and hardness of the composites were compared with those of Fe₄₀Mn₄₀Cr₁₀-Co₁₀ MEA. Discussion focused on the effect of TiC on the strengthening mechanism and the improvement in tribological behavior.

2 Experimental

2.1 Materials and composites manufacturing

The water-atomized Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA powder (with the average particle size of 5–15 μ m) and TiC (99.5 wt.%) particles were used as starting materials. The size of TiC

particles is mainly 1–4 μm . Fe₄₀Mn₄₀Cr₁₀Co₁₀MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite were fabricated by planetary ball milling and SPS. Alcohol was employed as the process control agent (PCA) to avoid welding and oxidation of the powder. The powder was mixed by ball milling for 6 h, followed by consolidating into discs of 40 mm in diameter using SPS at 1373 K for 15 min with a uniaxial pressure of 40 MPa.

2.2 Microstructure characterization

X-ray diffractometer (XRD, Rigaku X–2000) and scanning electron microscope (SEM, FEI nano 230 field emission) equipped with an energy dispersive X-ray (EDAX) spectrometer were used to characterize the microstructure and phase composition. Electron backscattered diffraction (EBSD) was conducted to access the grain structure using a Hitachi S–3400N SEM instrument equipped with a HKL-EABS system. The microstructure and selected area electron diffraction (SADP) were observed using a Tecnai G2 F20 transmission electron microscope (TEM). To prepare thin foil specimens for TEM examination, slices cut from the as-SPSed alloy samples were mechanically ground and cut into discs with 3 mm in diameter. These discs were then ground to a thickness less than 60 μm and finally ion-beam thinned using a Gatan precision ion polishing system (PIPS) at a voltage of 5 kV and an incident angle of 3°–7°.

2.3 Mechanical properties test

An Instron® 3369 universal testing machine was used for testing the compressive mechanical properties of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite, with a cross-head speed of 1 mm/min. The size of the cylindrical sample was $d5\text{ mm} \times 7\text{ mm}$. The hardness tests were carried out using an HVS–5 hardness tester with the load being 49 N for 10 s. All the reported data were the average of at least 5 specimens.

2.4 Wear resistance test

Wear tests were carried out using an HRS–2M roller friction wear tester at room temperature with a rotation speed of 600 r/min, a test load of 10 N, a duration time of 5 min and a total sliding distance of 60 m. The counter body of roller was a Si₃N₄ ball with a diameter of 5 mm.

3 Results

3.1 Microstructure characterization

SEM–BSE images in Fig. 1 show the micro-structure of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA without and with 10% TiC particles. The microstructure of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA exhibited a homogenous single phase. Pores were observed in the as-SPSed alloy, as marked by the arrows in Fig. 1(a). With the addition of TiC particles, obvious changes in the microstructure were observed in comparison with Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA. The Fe₄₀Mn₄₀Cr₁₀Co₁₀ matrix powder was present in a softening state at sintering temperature during SPS process. Under an external pressure during SPS, TiC particles filled the pores/gaps between their neighboring matrix powder particles to form more compact mixed powder particles. Thus, the grains were refined considerably and much less pores were found in the composite (Fig. 1(b)).

Table 1 gives the chemical compositions of various phases measured by SEM–EDS analysis. The analyzed results from the matrix at Point 1 in Fig. 1(a) are consistent with the initial proportion of alloy powder ($n(\text{Fe}):n(\text{Mn}):n(\text{Cr}):n(\text{Co})=4:4:1:1$). Analysis from the dark grey phase at Point 2 in Fig. 1(b) shows that the phase consists mainly of carbon and titanium. Similarly, the composition at Point 3 in Fig. 1(b) is roughly satisfied with the alloy composition of $n(\text{Fe}):n(\text{Mn}):n(\text{Cr}):n(\text{Co})=4:4:1:1$. It was also seen that the two phases in Fig. 1(b) contained other elements. For example, there were a small quantity of Fe, Mn, Cr and Co in the TiC phase. There were also Ti and C in the matrix phase. This could be attributed to two reasons: one is the measure error as the phase is too fine for SEM beam; the other is that atomic diffusion is resulted from the added particles and the matrix alloy during the sintering process at high temperature, indicating that the interaction between the added particles and the matrix alloy occurred. Elemental mapping on the selected area of the composite was further carried out by SEM–EDS. As shown in Fig. 2, Fe, Cr and Co elements distributed uniformly in the matrix grains; however, Mn and O mainly segregated to the matrix grain boundaries. Due to the high chemical affinity between Mn and O atoms, some MnO may be expected to develop at the matrix grain boundaries. The dark gray phase was composed of Ti and C, confirming that the dark grey phase in Fig. 1(b) was TiC.

The XRD patterns of Fe₄₀Mn₄₀Cr₁₀Co₁₀ and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite are shown in Fig. 3. Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA only showed the peaks of a FCC phase, implying that no other phases were formed except for the matrix phase. This was in good agreement with medium entropy alloy and also agreed with the previous study [16]. On the other hand, Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite consisted of FCC phase and TiC phase with no other phases.

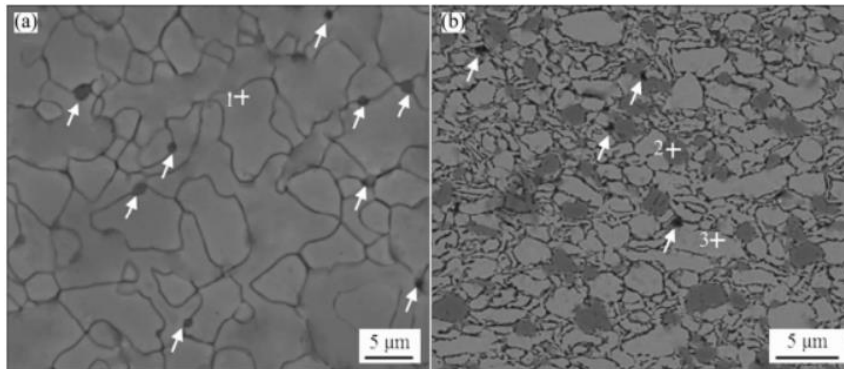


Fig. 1 Backscattered SEM micrographs showing microstructure of Fe₄₀Mn₄₀Co₁₀Cr₁₀ MEA (a) and Fe₄₀Mn₄₀Co₁₀Cr₁₀/10% TiC composite (b) (Arrows mark porosities. “+” marks are locations for composition analysis)

Table 1 EDS results of Fe₄₀Mn₄₀Cr₁₀Co₁₀/10%TiC composite corresponding to Fig. 1 (at.%) Point in Fig. 1

	Fe	Mn	Cr	Co	Ti	C
1	40.33	38.26	11.12	10.29	—	—
2	01.94	01.82	01.09	00.55	49.09	45.51
3	39.16	32.75	09.37	05.94	05.02	07.76

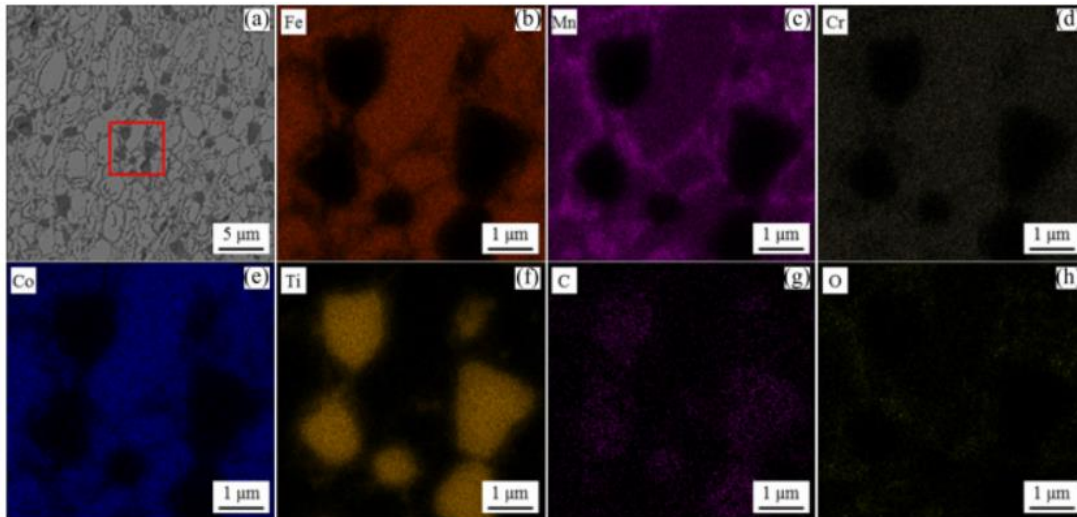


Fig. 2 SEM–EDS mapping images showing element distribution of Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite: (a) SEM image; (b–h) EDS mapping images

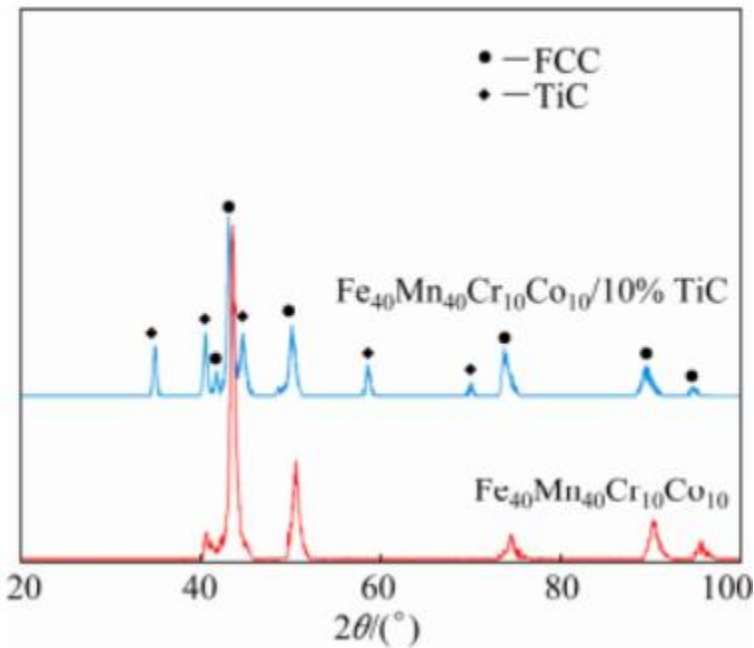


Fig. 3 XRD patterns of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite

EBSDF results in Fig. 4 show the grain structure of Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite in comparison with Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA. The addition of the TiC particles resulted in an obvious grain refinement and much less pores for the composite. The average grain sizes of the FCC phase were 4.85 μm for Fe₄₀Mn₄₀-Cr₁₀Co₁₀ MEA and 2.64 μm for the composite. The inverse pole figures (IPFs) of the FCC phase shown as inserts in Fig. 4 confirmed that the FCC grains have no obvious preferred orientation in Fe₄₀Mn₄₀Cr₁₀Co₁₀ with and without TiC addition.

TEM bright field image in Fig. 5(a) shows the microstructure of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA, and the SADP (insert in Fig. 5(a)), taken from area L1, verified the FCC structure. TEM bright field image taken from the composite is shown in Fig. 5(b), where two phases can be

seen in the microstructure. The SADP from area L2 showed that the matrix was also FCC structure. The dark particles marked as L3 were identified as TiC, with the SADPs shown in the insert. High resolution TEM (HRTEM) images in Figs. 5(c) and (d) show the interface between TiC particles and the FCC matrix. It is seen that the interface is sharp and clean (without secondary phase). Fast Fourier transformation (FFT) patterns from the lattice image confirm the orientation relationship of the FCC solid solution and TiC particle. The FCC phase is along the [011] zone axis (see L4) and TiC is along its [122] zone axis (see L5).

3.2 Mechanical properties

Figure 6 shows the compressive stress–strain curves of Fe40Mn10Cr10Co10 with addition of TiC in comparison with some other HEA/MEAs. Fe40Mn40Cr10Co10 MEA had a compressive strength of 1.571 GPa with corresponding compressive strain of 28.5%. The addition

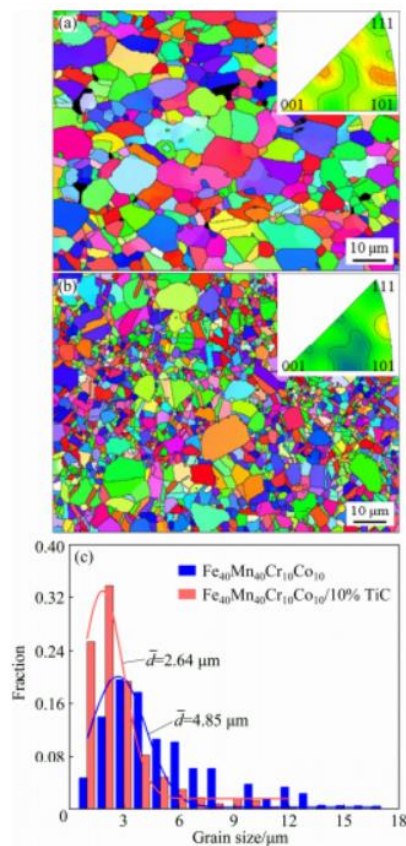


Fig. 4 EBSD mapping-based microstructures showing variation of grain sizes of Fe40Mn40Cr10Co10 MEA (a), Fe40Mn40Cr10Co10 /10% TiC composite (b) and distribution of average grain size (c) (Inserts in (a) and (b) are inverse polefigures (IPFs) of FCC phase in alloy)

of 10% TiC led to an obvious increase in compressive strength to 2.174 GPa and a slight increase in compressive strain to 32.2% for the composite. Considering the fact that compressive strain normally decreases with the addition of the ceramic particles, a slight increase in compressive strain could be ascribed to two factors. One is the effect of grain refinement, the other is the reduction of pores. As shown in Table 2, the hardness is also

significantly increased from HV 320 to HV 872. The difference in fracture morphology is shown in Figs. 6(c) and (d). Fe40Mn40Cr10Co10 MEA showed a single slip fracture surface, and the fracture consisted of a large number of slip bands. While the fracture surface of the composite was of a typical ductile-brittle mode. Moreover, there were many voids left on the surface of the composite after TiC particles were pulled out.

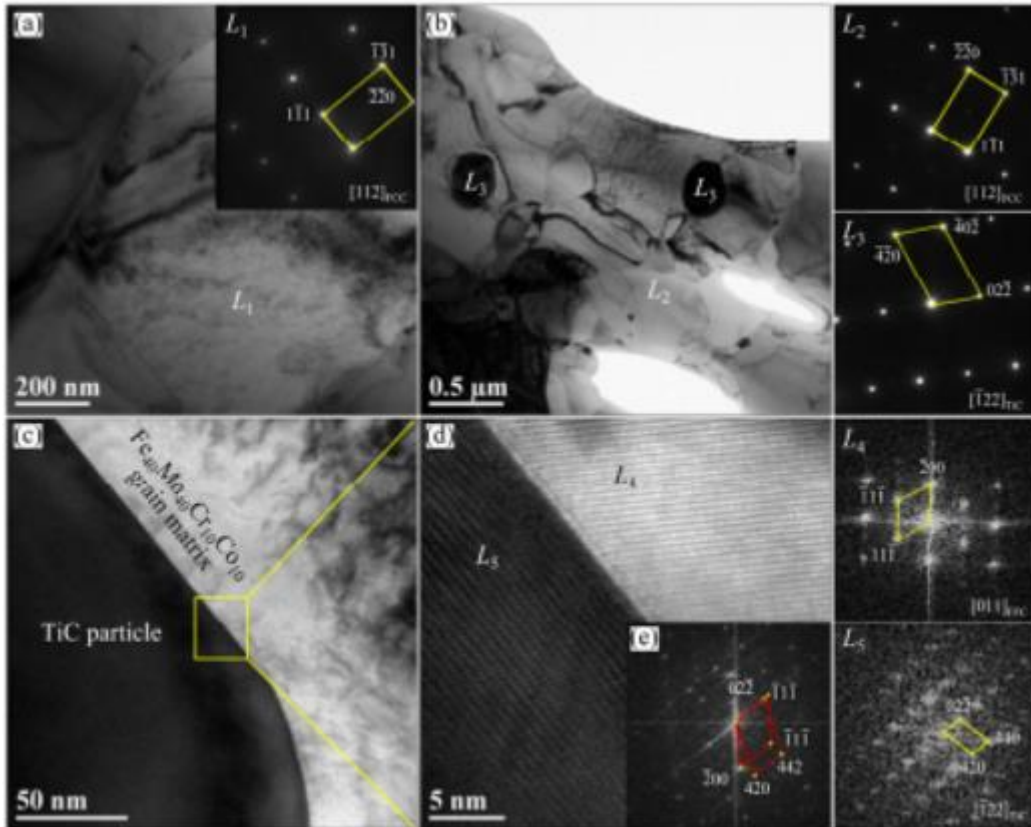


Fig. 5 TEM bright field micrographs of Fe40Mn40Cr10Co10 MEA with SADP revealing single FCC solid solution (a), Fe40Mn40Cr10Co10/10% TiC composite with corresponding SADP patterns (b) and image showing interfacial relationship between matrix phase and TiC (c), TEM high resolution image of rectangular area (d) in (c) and indexed schematic of corresponding interface relationship (e)

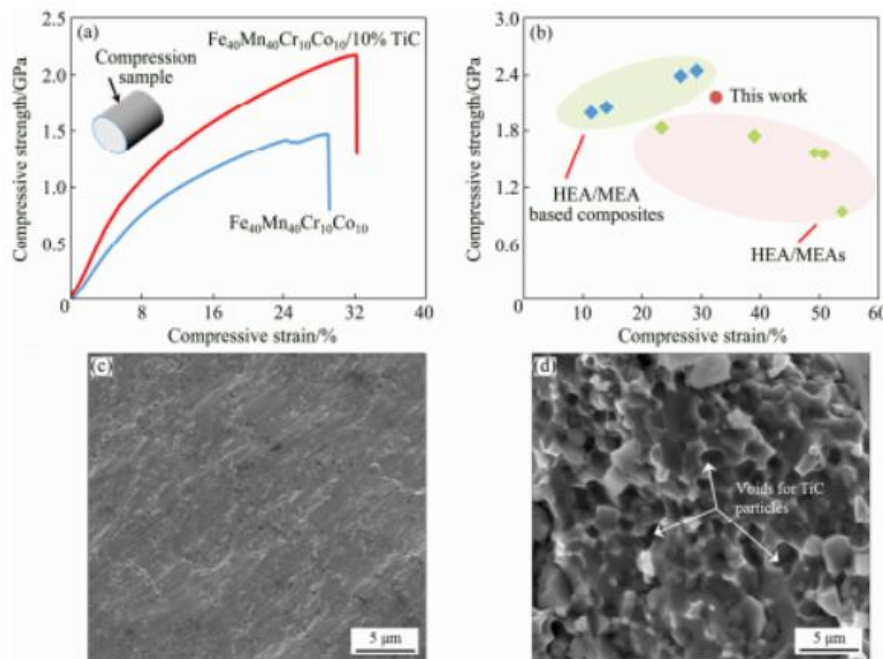


Fig. 6 Compressive stress–strain curves of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite (a), comparison of compressive strength vs compressive strain of typical HEA/MEAs [17–19] and HEA/MEA based composites [12–13, 20–21] (b), fracture surfaces of as-sintered Fe₄₀Mn₄₀Cr₁₀Co₁₀ (c) and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC (d) (Voids of post-deformed samples are highlighted by white arrows in (d))

3.3 Wear resistance

Figure 7 shows the 3D optical wear tracks and the corresponding 2D cross-section depth profiles of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₁₀Cr₁₀Co₁₀/10% TiC composite under a load of 10 N. As expected, the wear track of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA was deeper than that of the composite. The depth of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA was 1.9 about times that of Fe₄₀Mn₁₀Cr₁₀Co₁₀/10% TiC composite. The friction coefficients were 0.6 and 0.26 for Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and the composite, respectively (see Table 3). The reduced depth and friction coefficient confirmed that the wear resistance of the composite was improved.

SEM images in Fig. 8 show the morphologies of the wear tracks. For Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA, numerous wear particles and micro-ploughing were found inside the wear track (Fig. 8(a)). However, there were no visible micro-ploughs and wear particles on the worn surface of the composite (Fig. 8(d)). SEM images of the wear track at higher magnification in Figs. 8(b) and (e) showed that wear particles were clearly visible inside the wear track of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA. These particles were identified as metallic oxides based on the SEM–EDS results shown in Fig. 8(c). The observation of the worn surface indicates that the wear modes of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA included abrasive wear and oxidation wear, while the worn surface of the composite (Fig. 8(e)) was rough, which may be caused by the debonding of TiC and matrix. And the shedding TiC particles acted as abrasive particles and the primary wear mode of the composite was abrasion wear.

The XPS analysis results of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite are shown in Figs. 9 and 10, respectively. From Fig. 9, the metallic oxides (mainly Fe₂O₃, Cr₂O₃ and Mn₂O₃) were formed on the worn surface, which was in good agreement

with the SEM–EDS results in Fig. 8(c). Similarly, the metallic oxides were also found on the worn surface of Fe40Mn40Cr10Co10/10% TiC composites (Fig. 10), although no wear particles were visible in Fig. 8(e). The results of XPS showed that both Fe40Mn40Cr10Co10 and Fe40Mn40Cr10Co10/10% TiC composite were obviously oxidized during wear test. Therefore, oxidation wear is the major wear mechanism for Fe40Mn40Cr10Co10/ 10% TiC composite.

Table 2 Mechanical properties of Fe40Mn10Cr10Co10 MEA and Fe40Mn10Cr10Co10/10% TiC composite

Alloy Yield strength/GPa Compressive strength/GPa Compressive strain/% Hardness (HV)

Fe40Mn10Cr10Co10 MEA 0.48 1.571 28.5 320

Fe40Mn10Cr10Co10 MEA/10% TiC 0.73 2.174 32.2 872

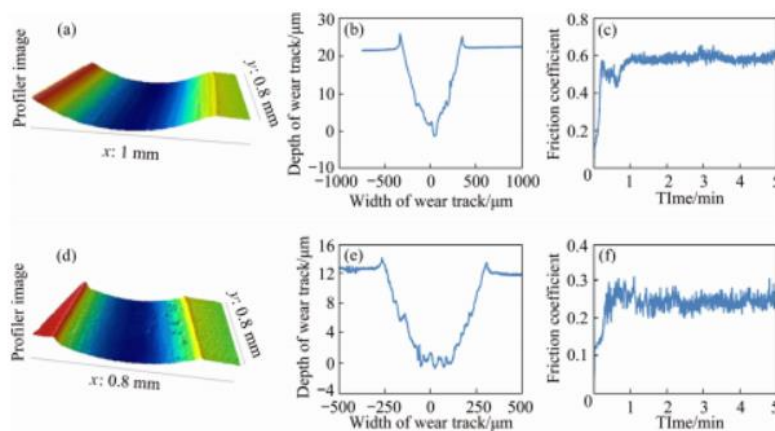


Fig. 7 3D optical wear tracks of Fe40Mn40Cr10Co10 MEA (a) and Fe40Mn40Cr10Co10/10% TiC composite (d) under 10 N load, cross-section depth profiles of Fe40Mn40Cr10Co10 MEA (b) and Fe40Mn40Cr10Co10/10% TiC composite (e), and friction coefficients of Fe40Mn40Cr10Co10 MEA (c) and Fe40Mn40Cr10Co10/10% TiC composite (f)

Table 3 Wear resistance properties of Fe40Mn10Cr10Co10 MEA and Fe40Mn10Cr10Co10/10% TiC composite

Alloy Depth of wear track/ μm Width of wear track/ μm Friction coefficient
 MEA 23.64 674.87 0.6
 MEA/10% TiC 12.44 567.18 0.26

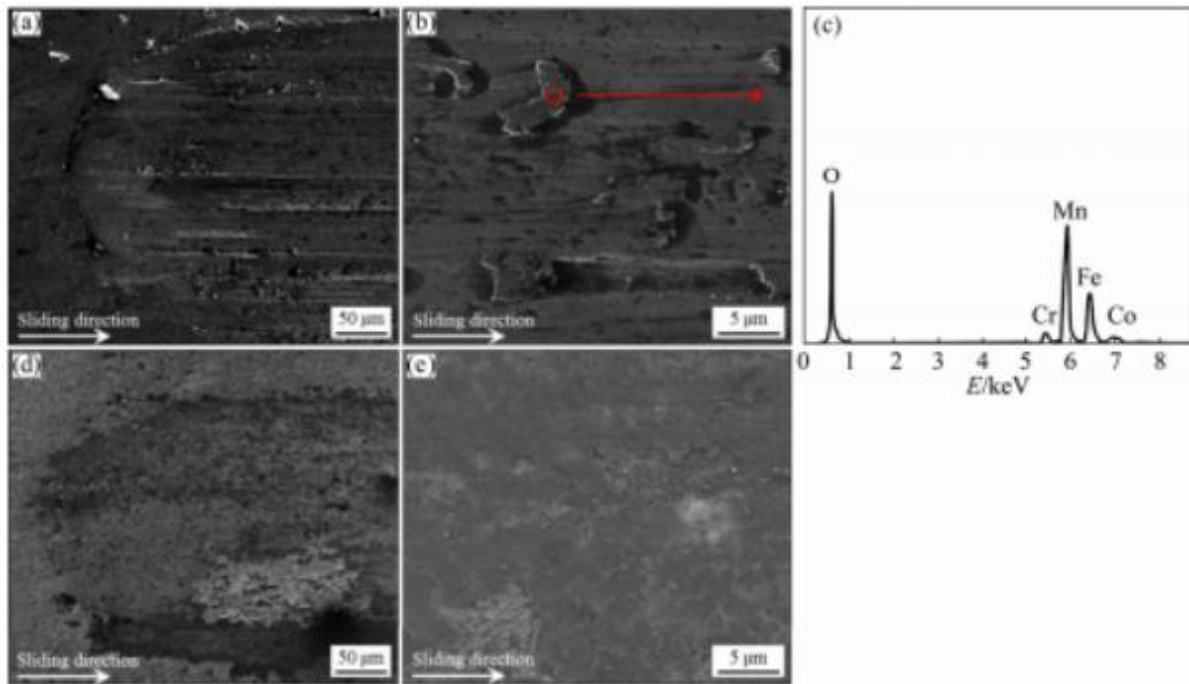


Fig. 8 SEM images showing wear tracks on Fe40Mn40Cr10Co10 MEA (a) and Fe40Mn40Cr10Co10/10% TiC composite (d), high magnification SEM images of wear tracks on Fe40Mn40Cr10Co10 MEA (b) and Fe40Mn40Cr10Co10/10% TiC composite (e), and corresponding EDS pattern of denoted wear particle in (b) (c)

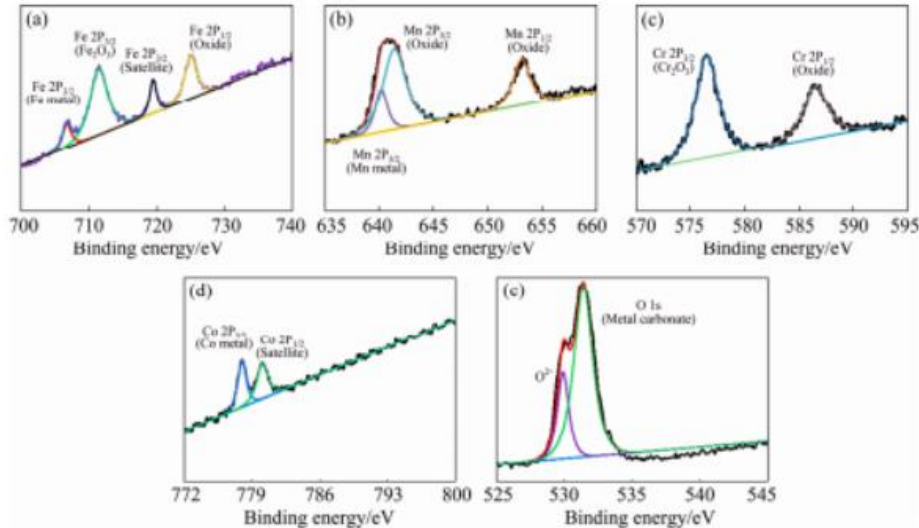


Fig. 9 XPS analysis results of Fe40Mn40Cr10Co10 MEA in air: (a) Fe; (b) Mn; (c) Cr; (d) Co; (e) O

4 Discussion

4.1 Improvement in mechanical properties by adding TiC particles

The present study has confirmed that Fe40Mn40- Cr10Co10/10% TiC composite fabricated by powder metallurgy with SPS exhibited improved mechanical properties in comparison with Fe40Mn40Cr10Co10MEA, as shown in Fig. 6. Fe40Mn40Cr10Co10/10% TiC composite also presented superior comprehensive mechanical properties in counterpart with

HEA/ MEAs [17–19] and their based composites [12–13,20,21]. An orientation relationship (OR), i.e., [011]//[122], (111)//(022), was determined, as shown in Fig. 5(d). This OR indicates that a coherent interface ((111)//(022)) exists between the matrix and the nanoscale particle. Thus, Orowan strengthening would be introduced through the TiC nanoscale particles. The addition of the TiC particles can inhibit the grain growth during sintering process and hinder dislocation motion during the deformation period. Grain refinement, effective load transfer effect and Orowan strengthening mechanism are responsible for these improved mechanical properties. In the following sections, the possible strengthening mechanisms will be discussed separately.

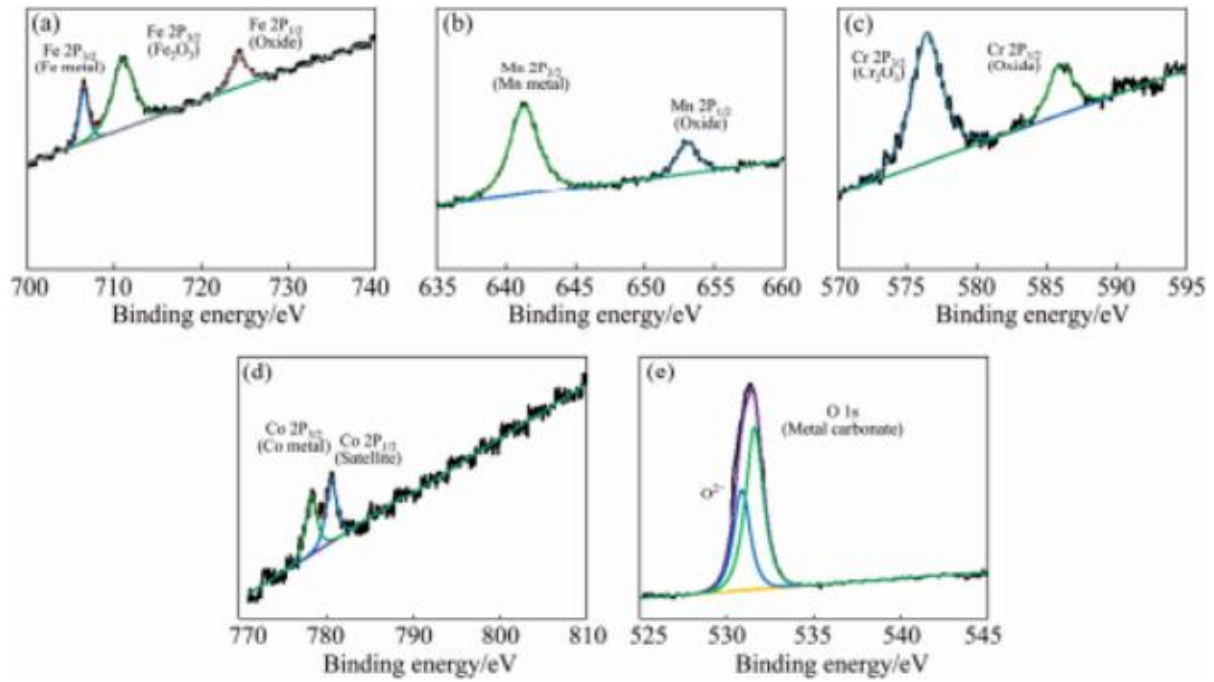


Fig. 10 XPS analysis results of Fe40Mn40Cr10Co10/10% TiC composite in air: (a) Fe; (b) Mn; (c) Cr; (d) Co; (e) O

(1) The load transfer of interface between Fe40Mn40Cr10Co10 and TiC particle

The load can transfer from matrix to the uniformly distributed reinforcement particles efficiently during deformation [22,23]. As depicted in Fig. 6(d), there are many voids left on the surface of the composite after TiC particles were pulled out. The interface can transfer load from Fe40Mn40Cr10Co10 matrix to TiC effectively, which contributes to the improvement of the strength of Fe40Mn40Cr10Co10/10% TiC composite. According to the modified shear lag model reported in Ref. [21], the strength contribution of load transfer effect in PRMMCs can be expressed by the following expression:

$$\Delta\sigma_{\text{Load}} = 0.5VP\sigma_m \quad (1)$$

where σ_m is the yield strength of Fe40Mn40Cr10Co10 MEA, as presented in Table 2. VP is the volume fraction of the TiC particles. Equation (1) exhibits that $\Delta\sigma_{\text{Load}}$ mainly depends on the volume fraction of TiC particles. The increase in VP brings a higher strength contribution.

(2) Orowan strengthening of TiC particles σ_{Orowan} can be calculated as follows [24]:

$\sigma_{\text{Orowan}} = 0.13 \ln 2 G / b r b \sigma \lambda = (2)$ where G is the shear modulus of matrix, b is the Burgers vector value, λ is the inter-particle spacing, and r is the radius of the particle. In present work, TiC

particles are hard (the hardness is as high as HV 2000–3000), which incorporate with the soft ductile Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA matrix and result in an increase in the hardness of the composite from HV 320 to HV 872 (see Table 2). These hard and dispersive particles are distributed to the grain boundaries and create resistance to dislocation propagation and movement. This suggests that higher shear stresses are needed to bypass dislocations through impeding the TiC particles, which eventually result in strengthening of the Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite.

(3) Dislocation strengthening caused by the difference in coefficient of thermal expansion (CTE)

Large thermal mismatch stress can be generated owing to the difference in coefficient of thermal expansion between the matrix and the reinforcement during deformation [25,26]. Hence, a lot of thermal mismatch dislocations are generated in the composite to relax these thermal stresses. In present work, the CTEs of TiC particle and Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA are 7.4×10^{-6} [27] and $23 \times 10^{-6} \text{ K}^{-1}$ [28], respectively, which lead to high dislocation density in Fe₄₀Mn₄₀Cr₁₀Co₁₀/TiC composite. These increased dislocations interact with each other, then impede their own motions and further increase the resistance to bypass the TiC particles.

(4) The grain refinement effect of TiC particle

The particles segregated at grain boundaries can provide a strong resistance between two neighboring grains. Grain boundary motion can be inhibited during sintering and achieve improved mechanical properties. For example, the grain size of CoCrFeNiMn MEA is decreased from 0.8 to 0.4 μm by in-situ yttrium nano-oxides particles [29]. The addition of TiC particles, locating at the grain boundaries, can decrease the average grain size of the FCC phase, as indicated by the results of SEM and EBSD and theoretically described by the Zener pinning [30]. The grain size and the second phase particle size satisfy the following relationship:

$$d_m = 4\alpha d_p / 3v_{SP} \quad (3)$$

where d_m is the grain diameter, d_p is the second particle diameter, v_{SP} is the volume fraction of the second particle, and α is the proportionality constant. This equation ties the grain radius of the matrix, the second particle diameter and volume fraction of the second particle so tightly together. Obviously, the decrease in d_p and the increase in v_{SP} bring a finer grain. This phenomenon has also previously been verified in Refs. [31–33].

4.2 Improvement in friction performance by adding TiC particles

TiC-based ceramic metals have been reported to show a high abrasive wear resistance [34]. Figure 7 shows that the wear depth and friction coefficient of alloy decreased by adding TiC particles. The improvement in wear resistance of Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC is attributed to the improved strength and hardness (see Table 2). TiC particles bonded with the FCC matrix and acted as plowing stopper to confine the soft FCC phases pulling out from the composite. Therefore, the hard TiC particles could protect the soft FCC matrix against the Si₃N₄ balls during the sliding and the composite exhibited improved resistance and lower friction coefficient. ZHANG et al [35] found that the wear resistance of Fe–28Al–5Cr alloy was significantly improved by the addition of TiC and confirmed that the high hardness of

TiC was the main reason. The wear resistance of Al2219 alloy was also reinforced by TiC particles at different loads and sliding distances [36]. The uniform distribution of TiC particulates in matrix enhanced the wear resistance of the composite.

5 Conclusions

(1) Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composites can be produced by powder metallurgy route through SPS at 1373 K for 15 min under 40 MPa. Fe₄₀Mn₄₀Cr₁₀Co₁₀MEA exhibits a single FCC phase and Fe₄₀Mn₄₀Cr₁₀Co₁₀/ 10% TiC composite consists of FCC phase and TiC phase.

(2) Grain refinement is obtained by the addition of TiC in Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA. The average grain size was 2.64 μm for Fe₄₀Mn₄₀Cr₁₀Co₁₀ /10% TiC composite, which is finer than that for Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA with an average grain size of 4.85 μm.

(3) The significant increase in strength is achieved for Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite. The compressive strength increases from 1.571 to 2.174 GPa and the hardness improves from HV 320 to HV 872 for Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite.

(4) Wear resistance of Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite is superior to that of Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA under the same condition. The wear modes in Fe₄₀Mn₄₀Cr₁₀Co₁₀ MEA and Fe₄₀Mn₄₀Cr₁₀Co₁₀/10% TiC composite include the micro-abrasive wear and oxidation wear.

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