- 1 Microstructure and tribological properties of titanium matrix nanocomposites
- 2 through powder metallurgy using graphene oxide nanosheets enhanced copper
- 3 powders and spark plasma sintering

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18 **ABSTRACT**:

- 19 Titanium alloys have been applied for many lightweight structural components in
- 20 the fields of aerospace, automobiles and biomedical implants owing to their
- 21 light-weight, good mechanical properties and biocompatibility. However, poor
- 22 tribological performance often restricts their wide-range applications. In this study, we

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synthesized Cu modified Ti-6Al-4V (TC4) powders with various Cu contents (1, 3, 5, 1 10 wt%), which was further strengthened with 0.3 wt% graphene oxide nanosheets 2 (GONs) using a powder metallurgy technology. These composite powders were then 3 synthesized into titanium matrix composites using spark plasma sintering. Effects of 4 Cu contents on microstructure evolution, phase composition and tribological 5 properties of Ti matrix composites were systematically investigated. The synthesized 6 7 composites were consisted of α -Ti, β -Ti, Ti₂Cu, *in-situ-*formed TiC and remained GONs, and showed better tribological properties than those of TC4 alloy. The average 8 9 coefficient of friction was reduced from 0.168 to a minimum value of 0.120 as the copper content increased from 0 to 3 wt%, meanwhile the wear volume loss was 10 reduced by 49.3%. Whereas further increasing copper contents resulted in the 11 12 increases of both coefficients of friction and wear volume loss. These improvements are mainly attributed to the strengthening effects by nanoscaled Ti-Cu intermetallics 13 and TiC@GONs structure, as well as the self-lubricating effect of GONs. Compared 14 15 with traditional surface modification processes, the new method proposed in this work is cost-effective and promising for improving the tribological performance of titanium 16 alloys in industry applications. 17

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Keywords: Ti matrix composites, Tribological properties, Microstructure, Powder modification, Spark plasma sintering

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1. Introduction

Titanium and its alloys are ideal materials for petrochemical and medical applications due to their attractive properties such as light weight, high specific

strength, and good corrosion resistance and antibacterial properties [1-3]. However, application of new types of high-performance structural materials normally requires not only their strength and toughness, but also their good wear resistance. The poor wear resistance of titanium materials influences not only their surface appearance, but also their sensitivity of crack initiation and shortening of the part life due to early failure under complex loads or environments. To improve the comprehensive mechanical properties especially the tribological properties of titanium materials, considerable research has been focused on the applications of various surface modification and strengthening strategies [4-6]. So far, introduction of various nanoscale ceramic particles (TiC, TiB, Al₂O₃, SiC) into Ti matrix has gained extensive attention [7-10]. To our knowledge, graphene and its derivatives have been demonstrated to be effective for reinforcing metal matrix composites [11-14] owing to their unique 2D structures and overall high performances. However, there are always concerns for the uniform dispersion of graphene and the resulting good enhancement in mechanical properties, especially the strength of the composites. Chu et al. [15] reported that the yield strength of the graphene/copper composites was increased by 37% reaching 300 MPa compared to pure copper when 8 vol.% graphene was added. Compared with Al, Wu et al. [16] found that the tensile strength of the graphene reinforced Al composites was increased by 73.9% meanwhile the ductility maintained more than 10%. Previously, Dong et al [17] showed that using spark plasma sintering (SPS) technique, addition of only 0.3wt.% of GONs into Ti matrix composites (TiMC) led to the yield

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strength and ultimate tensile strength of the composites increased by 7.44% and 9.65%, respectively. Zhang's group fabricated Ni coated graphene nanoplates reinforced Ti composites using mechanical agitation and SPS, and achieved a strength of 793 MPa, which is ~ 40% higher than that of monolithic Ti matrix [18]. However, among all these studies, few of them have been focused on the tribological behavior of graphene/Ti composites. Ti-6Al-4V alloys is the most extensively used Ti alloy among all the Ti materials, however, they have a poor wear resistance, thus limiting their applications where wear resistance is essential. This study aims to explore the relationship between tribological performance and composition as well as microstructure of graphene/Ti composites, and to reveal the fundamental mechanisms on the damage of the composites resulted from wear. To achieve this, copper powders with different contents (1, 3, 5, 10 wt%) and 0.3% GONs were synergistically added into Ti-6Al-4V (TC4) matrix using a novel powder

fundamental mechanisms on the damage of the composites resulted from wear. To achieve this, copper powders with different contents (1, 3, 5, 10 wt%) and 0.3% GONs were synergistically added into Ti-6Al-4V (TC4) matrix using a novel powder modification technology and then the composites were synthesized using SPS in order to improve their tribological properties. The copper was chosen as an alloying element to modify the TC4 powders because it is a β -stabilizing element. According to the equilibrium phase diagram of Ti-Cu binary alloys, Ti-Cu alloys with a low Cu content consists of a matrix of α -Ti and precipitates of Ti₂Cu intermetallic compounds [19]. It was reported that the Ti alloys with appropriate Cu concentrations show good biocompatibility [20], corrosion resistance [21, 22] and mechanical properties [23]. For examples, Kikuchi et al. reported that the mechanical properties of cast Ti- α -Cu (α = 0.5-10 wt%) alloy have been significantly improved compared with those of pure

Ti (e.g., tensile strength and yield strength have been increased up to 30% and 40%,

The SPS was used because of its advantages such as short sintering time, fast

2 respectively [23].

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- heating rate, controllable structure, uniform temperature and simple operation, which
 can achieve uniform microstructure and high density of the sintered materials by
 inhibiting their grain growths [24, 25]. The microstructure and wear behavior of the
 composites were systematically characterized, and the wear mechanisms were
 discussed. The Ti matrix composites showed the highest hardness with a value of 411
- 10 (330 HV). The composites showed the lowest coefficients of friction (0.12) and the

HV with 5 wt.% copper content, which is ~25% higher than that of the TC4 alloy

- wear volume loss (0.155 mm³) among the alloys and composites studied in the present
- work. The improvement of tribological properties is due to the strengthening by the
- 13 nanoscale precipitates (Ti₂Cu and TiC) at high temperatures, self-lubricating effect
- introduced by the pulled-out GONs from the composites, the fragileness of the brittle
- 15 TiC@GONs structure, as well as the enhanced heat conduction resulted from the high
- 16 heat conductivity of the preserved elemental copper and graphene. The
- 17 GONs-xCu@TC4 composites with a good wear resistance developed in this work has
- their promising applications in the field of brake discs and aerospace.

2 Experimental

20 2.1 Raw Materials

- 21 Ti-6Al-4V spherical powders with an average particle size of 15-53 μm were
- used as the matrix material, and they were purchased from Xi'an Sino-Euro Materials

Technologies Co., Ltd., China. Figure 1a shows a scanning electron microscope 1 (SEM, JEOL JSM-6700F) image, and their chemical composition is listed in **Table 1**. 2 3 The particle size of the TC4 powders conforms to a normal distribution with an average value of 37 um (inset in Figure 1a). The wide distribution of the particle size 4 is beneficial for the full densification of the powders as the fine particles can fill the spaces in the triangle intersections of coarse particles during the following SPS 6 process [26]. The average size of the Cu powders is 48 µm, and they were supplied by 7 Beijing Xin Rong Yuan Technology Co., Ltd., China. An SEM image of the pure Cu 8 9 powders shown in Figure 1b reveals their flower-like and flocculent patterns which are composed of fine copper particles. GONs with diameters of 0.5~5 um and 10 thicknesses of 1~3 nm were purchased from XFNANO Technology Co., Ltd., China. 11 12 A transmission electron microscope (TEM) was used to characterize the GONs. For the TEM observation, the GON nanopowders were firstly dispersed into ethanol and 13 then drop casted onto an amorphous carbon coated copper grid [27]. An TEM image 14 15 in Figure 1c shows a crumpled and thin-layered sheet structure, which is the typical morphology of graphene oxides. A high-resolution TEM (HRTEM) image shown in 16 Figure 1d presents that the GONs have a few layer structure. The selected area 17 electron diffraction (SAED) pattern of the GONs displays two diffraction rings, 18 corresponding to (1011) and (1120) planes of the graphene (Figure 1e). Table 1 lists 19 chemical compositions of both the TC4 and Cu powders. 20

Table 1. Chemical composition of Ti-6Al-4V and copper powders (wt %).

| Materials Al V Fe C N O H Cu II | Materials | Al | V | Fe | C | N | O | Н | Cu | Ti |
|---------------------------------|-----------|----|---|----|---|---|---|---|----|----|
|---------------------------------|-----------|----|---|----|---|---|---|---|----|----|

| Ti6Al4V | 6.17 | 4.23 | 0.17 | 0.008 | < 0.003 | 0.10 | 0.0048 | - | Bal. |
|---------|------|------|------|-------|---------|-------|--------|------|------|
| Cu | - | - | - | - | - | 0.012 | 0.002 | Bal. | - |

2.2 Fabrication of the GONs-xCu@TC4 composites

2.2.1 Preparation of mixture powders

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In order to disperse the copper powders and GONs uniformly in the TC4 3 powders, a two-step shift-speed ball milling process was adopted to prepare mixture 4 powders. (1) The mixed powders of copper and TC4 were ball-milled for 2 hours in 5 an Ar atmosphere at a speed of 300 r/min with a ball ratio of 3:1 in the same direction. 6 (2) Based on the previous study [17], we have chosen 0.3 wt% GONs to be added into 7 8 the Cu modified TC4 powders using a low-energy ball milling, with a speed of 150 r/min for another 3 hrs. The aim is to coat the GONs uniformly onto the composite 9 powder surfaces. Five types of mixing powders were prepared, i.e., mixtures with 10 11 copper contents of 1 wt%, 3 wt%, 5 wt%, 10 wt% and a mixture without copper. **Figure 1f** shows an SEM image of the composite powders after two-step ball milling. 12 Clearly the TC4 powders are slightly deformed. Whereas copper powders are severely 13 14 deformed, i.e. their shapes are transformed into a smaller flake-shape, and they are tightly attached to the surfaces of TC4 powders. The GONs are uniformly dispersed 15 onto the surface of TC4, which is clearly indicated by the energy dispersion X-ray 16 spectrum (EDS) (Figure 1g). 17

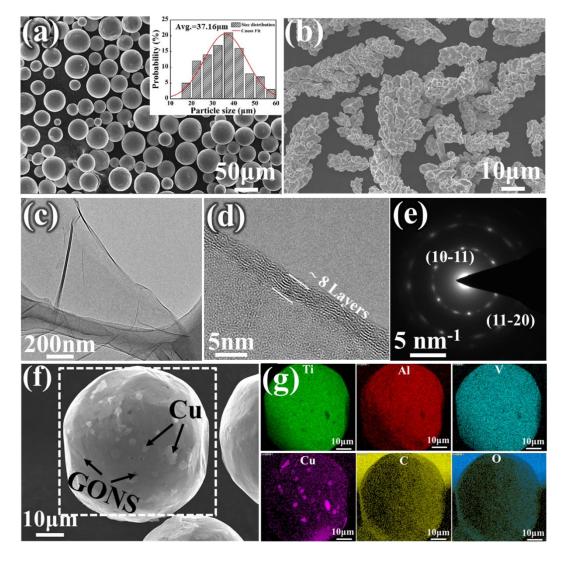


Figure 1. (a) SEM of raw TC4 powders and (b) copper powders; (c) TEM, (d)

- 3 HRTEM images of GONs and (e) the corresponding SAED pattern; (f)
- 4 GONs-1Cu/TC4 composites powders; (g) EDS mapping images of Ti, Al, V, Cu, C, O
- 5 elements, respectively.

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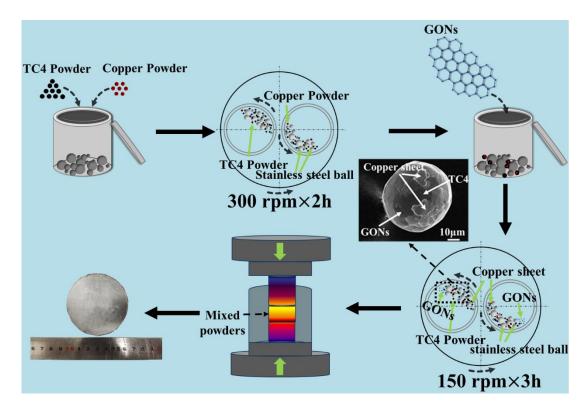
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2.2.2 Fabrication of Ti matrix composites

- 8 SPS is one of the best sintering processes for preparing metal matrix composites
- 9 (MMCs) owing to its high efficiency. Figure 2 shows the powder metallurgy and SPS
- 10 processes for fabrication of TiMCs in this work. The mixed powders were loaded into

- a graphite grinder with an internal diameter of 60 mm and an external diameter of 100
- 2 mm. The graphite paper with a thickness of 1~2 mm were placed around the powders
- 3 to prevent the reaction and bonding of TC4 powder with the graphite mold.
- The vacuum of SPS furnace was maintained at $\sim 10^{-2} \sim 10^{-3}$ Pa during sintering.
- The holding pressure, heating rate, and sintering temperature were 45 MPa, 100
- 6 °C/min, 1000 °C, respectively. There was a temperature monitoring hole with a
- 7 diameter of 10 mm in the middle of the graphite mold. During the sintering, the
- 8 temperature was measured using an infrared camera, which was positioned at about 5
- 9 mm from the sample center. After sintering for about 5 minutes, the samples were
- cooled down with the furnace. Pure TC4 alloy and GONs/TC4 composites without Cu
- were also sintered under the same conditions for comparisons. The selection of
- 12 detailed sintering parameters was described in a previous study [17]. For
- simplification, hereafter the as-sintered TiMCs with different Cu contents were named
- as GONs-xCu/TC4 composites (x=0, 1, 3, 5 and 10 wt%). The relative densities of
- these sintered Ti composites were determined using the Archimedes' method [11], and
- the obtained results are listed in Table 1. Clearly all the composites show a higher
- 17 relative density than that of pure TC4.



2 Figure 2. Schematic illustrations of fabrication procedures for the GONs-Cu/TC4

3 composites using powder modification and SPS.

5 Table 1. The theoretical density, measured density and relative density of fabricated

6 Ti matrix composites.

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| Materials | Theoretical density | Measured density | Relative density | |
|---------------|---------------------|-------------------|------------------|--|
| | (g/cm^3) | (g/cm^3) | (%) | |
| TC4 | 4.510 | 4.407 ± 0.009 | 97.72 | |
| GONs/TC4 | 4.494 | 4.405 ± 0.005 | 98.02 | |
| GONs-1Cu/TC4 | 4.516 | 4.439 ± 0.008 | 98.29 | |
| GONs-3Cu/TC4 | 4.560 | 4.486 ± 0.010 | 98.38 | |
| GONs-5Cu/TC4 | 4.603 | 4.513±0.013 | 98.04 | |
| GONs-10Cu/TC4 | 4.707 | 4.616 ± 0.015 | 98.07 | |

2.2.3 Wear and hardness tests

9 Before testing hardness and wear behavior of the composites, all the sintered

samples were polished to 2000 grits, and then ultrasonically cleaned with acetone and

2 ethanol. Vickers hardness was tested using an HVS-1000 machine with a load of 3 N

and a dwell time of 15 s at 10 different locations. Then the average value was

4 obtained to represent the hardness of the sample.

The wear behavior of the composites was tested using a MS-T3001 rotary friction testing machine (Lanzhou Huahui Instrument Technology Co.,) by dry sliding at room temperature in air. A Si₃N₄ ball with a diameter of 4 mm was used as the counterpart. The test was conducted with the normal load of 50 N, the rotary speed of 400 r/min and the rotation radius of 3 mm for 120 min. The friction coefficients of the samples were recorded automatically. Profiles of the worn surfaces were measured using a MicroXAM 3D surface profilometer system in order to determine the wear volume loss. The morphologies of the worn surface with different wear conditions were characterized using the SEM.

2.2.4 Microstructures of composite materials

Microstructures of the composites and the prepared powders were characterized using the SEM. Before observation, the Ti matrix composites were polished with 2000 grits of silicon carbide sandpaper, cleaned ultrasonically in an alcohol solution and then polished with alumina polishing solution. After that, the surface of the composites was etched using a solution of hydrofluoric acid, nitric acid and water, with ratios of 1:3:5 (vol.%). The phases in the composites were identified using X-ray diffraction (XRD, Bruker D8 ADVANCE) with Cu- K_{α} radiation and a scanning rate

- 1 of 10°/min.
- 2 Raman spectroscopy was used to investigate defects in the GONs and presence of
- 3 TiC, and the scan range of laser beam was from 300 to 3000 cm⁻¹. The tests were
- 4 performed at room temperature using a Laser Raman Spectrometer (LabRAM HR
- 5 Ecolution) with an excitation wavelength of 532 nm.
- An X-ray photoelectron spectroscope (XPS, Thermo SCIENTIFIC ESCALAB
- 7 250Xi, Al- K_{α} source) with an analysis chamber pressure of 4×10^{-8} Pa, was applied to
- 8 investigate the chemical valences of surface elements of sintered Ti matrix
- 9 composites.

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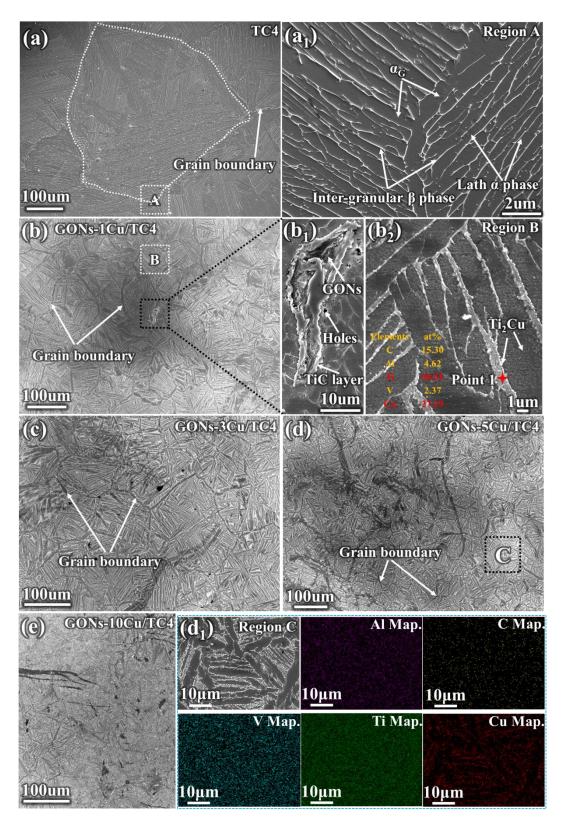
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3 Results and discussions

3.1 Microstructure Evolution and Phase Analysis

SEM images of the sintered GONs/TC4 composites with various copper contents 13 are shown in Figure 3. The monolithic TC4 alloy without the addition of copper and 14 15 GONs has a typical Widmanstätten lamellar microstructure (Figure 3a), consisting of the lath-shaped α phase, the α phase at grain boundaries (α_G) and the intergranular β 16 17 phase (Figure 3a₁) The size of the primary β grains is much larger (~450 um) than that of the raw TC4 powder (~38 um, Figure 1a), which is due to the slow cooling 18 process from the β transformation temperature [28]. However, owing to the 19 introduction of Cu and GONs, the grain size of primary β phases in Figures 3b~3e is 20 21 much smaller than that of the TC4 monolithic alloy (Figure 3a), which is beneficial to the enhancement of their mechanical properties. When the content of copper reaches 22

- 1 10 wt.% in the composites, the primary β grain boundary disappears as shown in
 2 Figure 3e.
- 3 Besides the α phase and the β phase, the unreacted GONs, the *in-situ* formed TiC particles or layers (Figure 3b₁) and the nanoscale Ti₂Cu precipitates co-exist in the 4 5 composites as shown in Figures 3b~3e. The EDS mapping analysis in Figure 3d₁ shows that the Cu is preferred to distribute around the β phase because the Cu is a β 6 stabilizing element. C element is relatively homogeneously distributed inside the Ti 7 matrix without obvious agglomerations. The formation of TiC is attributed to the 8 initiative reactions between C and Ti matrix [29], which is beneficial to the interfacial 9 10 strength and tribological properties. The presence of Ti₂Cu is evidenced by the energy dispersion spectrum (EDS) results (Figure 3b2, inset). The microstructure near point 11 1 in Figure 3b₂ consists of Ti (50.51 at.%) and Cu (27.19 at%), indicating that the 12 Ti₂Cu phase was nucleated and grown during the SPS process. 13



2 **Figure 3.** SEM images of GONs-xCu/TC4 composites with different Cu contents. (a)

- $3 \sim (a_1)$ pure TC4; (b)~(b₂) GONs-1Cu/TC4; (c) GONs-3Cu/TC4; (d) GONs-5Cu/TC4;
- 4 (e) GONs-10Cu/TC4; (a₁) and (b₁) \sim (b₂) are enlarged images of corresponding marked

region in Figure (a) and Figure (b), respectively. (d₁) EDS mapping analysis of the marked region C in Figure (d).

To further verify the phases of the sintered composites, XRD analysis was 3 carried out and the results are shown in Figures 4a and 4b. A peak at ~ 39.49° (PDF # 4 15-717) corresponding to the (103) of Ti₂Cu is identified in the XRD results and its intensity increases significantly with the addition of copper. Meanwhile, Cu, β-Ti 6 phase, TiC and GONs could not be detected by XRD for all the composites and TC4 7 alloy. The XRD results indicate that the main phase is α -Ti and the secondary phase in 8 9 the composites is mainly Ti₂Cu. No elemental copper is found in the composites because the reactions between copper and titanium during sintering result in the 10 formation of Ti₂Cu. Although other phases, i.e., GONs, TiC and β-Ti phase are 11 12 observed by SEM, their contents are out of the detecting limit of the XRD instrument used in the present work [30]. 13

According to the phase diagram of titanium-graphite, TiC is the only type of intermetallics that could form in the Ti-C system. Although the composite is a multi-elements system containing Ti, Al, V, Cu, C and O, the mixing of C, Cu with the matrix TC4 is not at an atomic level but at a microscale level. Taking into account the short sintering time of SPS, the local environment of Ti-C or Ti-Cu could be considered as binary systems. At the sintering temperature of 1273 K, the change of Gibbs energy is estimated to be -84.4 kJ/mol for the following reaction [31, 32]:

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$$Ti+C=TiC$$
 $\Delta G_1=-92400+6.275T \text{ J/mol}$ (1)

Hence, TiC is resulted from the spontaneous reaction between the matrix Ti and

- the carbon from the GONs [29]. Similarly, Ti₂Cu is the preferable phase when the
- 2 content of Cu is less than 33.3 at.%, according to the binary phase diagram of Ti-Cu.
- 3 The change of Gibbs energy can be estimated as -6.2 kJ/mol at 1273 K for the
- 4 following reaction [31, 32]:

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$$2T_1+C_0=T_12C_0$$
 $\Delta G_2=-12131+4.688T \text{ J/mol}$ (2)

- 6 which also indicates that the formation of Ti₂Cu is spontaneous.
- 7 To further verify the existence of TiC intermetallics and GONs in the composites,
- 8 Raman and XPS spectroscopies were used. The Raman spectra of the raw GONs and
- 9 the TiMCs are shown in Figure 4c and the main peaks are listed in Table 2. The
- 10 characteristic Raman peaks for graphene are similar to those of the composites, i.e.,
- 11 the G band at ~ 1580 cm⁻¹ is originated from the in-plane vibration of sp² carbon
- atoms, the D band at $\sim 1350 \text{ cm}^{-1}$ corresponds to the doubly resonant disorder-induced
- mode related to the defects, and the 2D band at ~ 2700 cm⁻¹ represents the feature of
- graphene [27, 33]. Similar characteristic bands between the composites and the raw
- GONs confirm that the GONs is retained in the TiMCs. It is also noticed that the I_D/I_G
- values for all the composites are significantly higher than the corresponding value for
- the GONs. Since I_D increases with the density of defects [27], the increase of the I_D/I_G
- value for the composites suggests the introduction of defects into the GONs during
- 19 the ball milling or sintering process.
- 20 Besides the characteristic peaks of graphene in the composites, two additional
- Raman peaks located at $\sim 418 \text{ cm}^{-1}$ and $\sim 605 \text{ cm}^{-1}$ suggest the presence of TiC [34].
- 22 Further investigation using the XPS was carried out, and Figure 4d and Figure 4e

show that the peaks at 454.6 eV and 460.2 eV correspond to the binding energy of the Ti-C bond, which verifies the existence of the TiC intermetallics. In addition to the main peaks of Ti located at 453.8 eV for Ti_{2p}^{3/2} and 459.95 eV for Ti_{2p}^{1/2} in the XPS spectrum, peaks at 932.5 eV for Cu_{2p}^{1/2} and 952.4 eV for Cu_{2p}^{3/2} as shown in **Figure**4f correspond to the binding energy of the Cu-O bond. XPS analysis proves the formation of the Cu-O bonds in the composites, which is helpful for improving the interfacial bonding strength between the Cu and GONs [35].

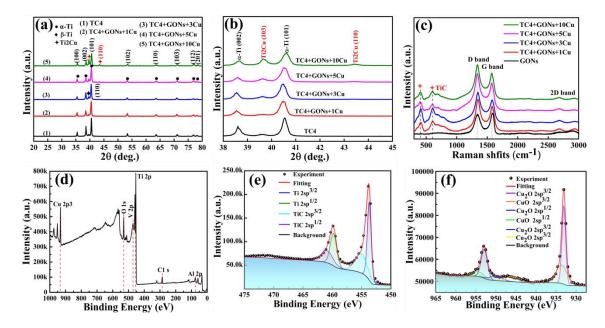


Figure 4. XRD, Raman and XPS analysis results for GONs-xCu/TC4 composites: (a) XRD patterns; (b) An enlarged XRD patterns at 2θ=38~45° in Figure (a); (c) Raman spectra of composites; (d) XPS survey spectra of GONs-5Cu/TC4 composites; High resolution XPS spectrum of Ti_{2p}; High resolution XPS spectrum of (e) Ti_{2p} and (f) Cu_{2p}, respectively.

Table 2. The detailed Raman spectra results of the sintered GONs-xCu/TC4 composites.

| Materials | D band | G band | 2D band | I_{D}/I_{G} |
|---------------|---------|---------|---------|---------------|
| GONs | 1351.28 | 1587.31 | 2695.91 | 0.95 |
| GONs-1Cu/TC4 | 1346.64 | 1581.16 | 2679.29 | 1.23 |
| GONs-3Cu/TC4 | 1349.62 | 1581.15 | 2685.78 | 1.28 |
| GONs-5Cu/TC4 | 1348.48 | 1579.44 | 2685.44 | 1.30 |
| GONs-10Cu/TC4 | 1346.13 | 1577.16 | 2679.94 | 1.31 |

3.2 Tribological properties

Figure 5a presents the effect of Cu contents on the hardness of sintered Ti matrix composites. The Vickers hardness of the composites increases from 330 HV to 411 HV when the content of Cu is increased from 0 to 5 wt.%. According to the characterization of the microstructure aforementioned, the hardness enhancement of the composites is attributed to the strengthening effect due to the smaller grain sizes, the fine second phases of Ti₂Cu, TiC and the preserved GONs, especially a large number of Ti₂Cu precipitates. Further addition of Cu results in a slight decrease of the hardness, due to the softening effect induced by the elemental copper which is not reacted with the titanium matrix during the short sintering time of SPS.

The coefficients of friction (COFs) vary with the sliding time during the wear tests as shown in **Figure 5b**. The friction experiences an unstable initial stage for all the samples. The duration of this unstable stage is much longer for the TC4 alloy (~65 min) and the GONs-1Cu/TC4 composite (~50 min) than that for the composite with the content of copper above 1 wt.% (~10 min). After the running-in stage, the COF

curves of the samples stabilize gradually. The initial fluctuation of the COFs is due to 1 the "break-in" or "running-in" period and an unstable surface roughness of the 2 specimens during the initial sliding [36, 37]. However, the COFs of the GONs/TC4 3 composites are reduced after ~ 40 min, which is attributed to the presence of GONs, 4 whose delamination can induce a lubrication effect during the sliding process (inset, Figure 8b₁). 6 The average COFs versus the copper content of the composites are plotted in 7 Figure 5c. The COF of the composite with 3 wt.% of Cu is the minimum among the 8 9 five types of samples. The increase of deviation for the Cu content from 3 wt.% accompanies the increase of the COFs. The wear depth and width were determined 10 from the cross sectional profiles of the wear scars, and the results are shown in Figure 11 12 **6.** They show similar trends to the COFs, i.e., the minimum values are around 3wt.% of Cu as shown in Figure 5d. The variation trends of the wear volume losses with the 13 copper content (Figure 5e) are similar to those of the COFs. Compared to that of the 14 15 pure TC4 alloy, the wear volume losses of the composites with GONs and Cu are 16 significantly lower. In particular, the minimum wear volume loss is found in the composite with a copper content of 3 wt.%, and the wear volume loss of this kind of 17 composite is about half the value of the monolithic TC4 alloy. These reveal that the 18 addition of Cu and GONs can remarkably improve the wear resistance properties 19 compared with the unreinforced matrix alloy. The wear volume loss was also 20

evaluated based on the 3D morphology of the wear tracks (Figure 6). The wear

volume loss as well as the average COFs are listed in **Table 3**.

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Table 3. The wear properties of the sintered GONs-xCu/TC4 composites.

| Wear properties | Cu contents | | | | | | |
|-------------------------------------|--------------|-------------------|-------------|-------------|-------------|-------------|--|
| | TC4 | 0 wt.% | 1 wt.% | 3 wt.% | 5 wt.% | 10 wt.% | |
| Average COFs | 0.1680±0.011 | 0.1389±0.017 | 0.150±0.012 | 0.120±0.015 | 0.130±0.01 | 0.140±0.015 | |
| Wear width (µm) | 3533±45 | 3190±28 | 2895±30 | 2953±35 | 3341±38 | 3353±45 | |
| Wear height (µm) | 472±5 | 439±6 | 352±7 | 335±8 | 371±9 | 468±5 | |
| Wear volume loss (mm ³) | 0.306±0.015 | 0.168 ± 0.008 | 0.176±0.009 | 0.155±0.01 | 0.202±0.012 | 0.230±0.008 | |

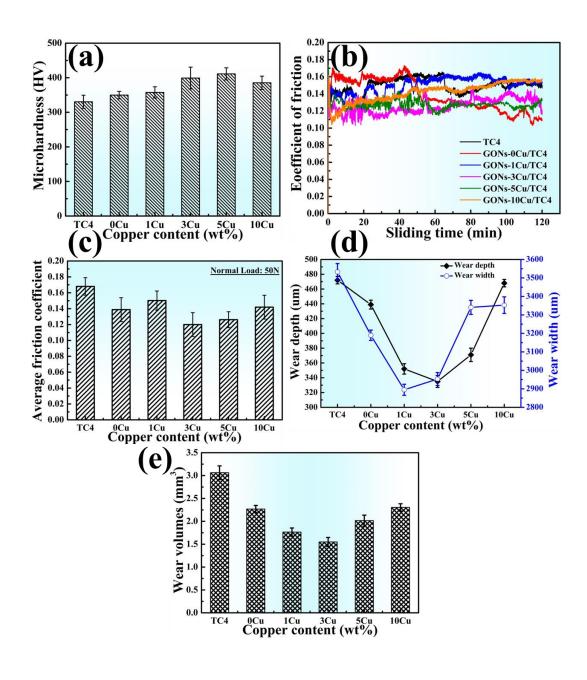


Figure 5. (a) Hardness of sintered TiMCs samples, (b) Variations of friction

1 coefficient of GONs-xCu/TC4 composites under a normal load of 50 N, (c) The

average friction coefficients of all samples, (d) Wear depth and the maximum wear

depth of scars for sintered composites, and (e) Wear volumes loss of different samples,

4 respectively.

The wear debris at the surface of the wear scars were examined using SEM and EDS, and the results are shown in Figure 7. The shape of the wear debris changes with the increase of copper content in the composites. On the specimen surface of the TC4 alloy, there exists two types of wear debris, i.e., the plate-like debris with an average size of ~80 μm and the fragments of fine particles with an average size of ~10 μm (Figure 7a). With the addition of copper and GONs into the TC4 matrix, the plate-like debris are still the main feature, but the number of the fine particle fragments is reduced drastically and almost becomes zero when the copper content is 10 wt.%. The debris surfaces of the composites with copper contents of 5 wt.% (Figure 7d1) and 10 wt.% (Figure 7e2) show deep grooves, indicating significant ploughing features during sliding wear. It was also noticed that the oxygen content in the wear debris of the composites is lower than the corresponding value of the TC4 alloy as shown in the EDS results in Figure 7.

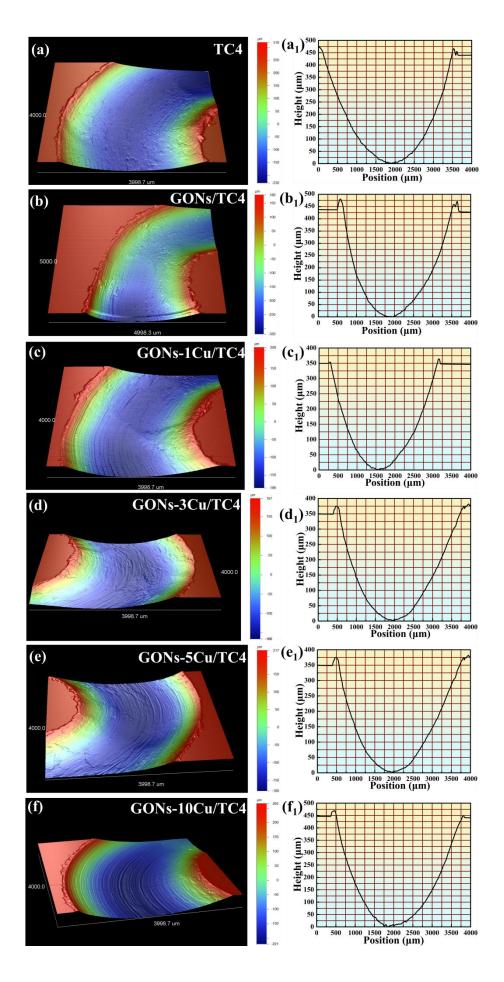


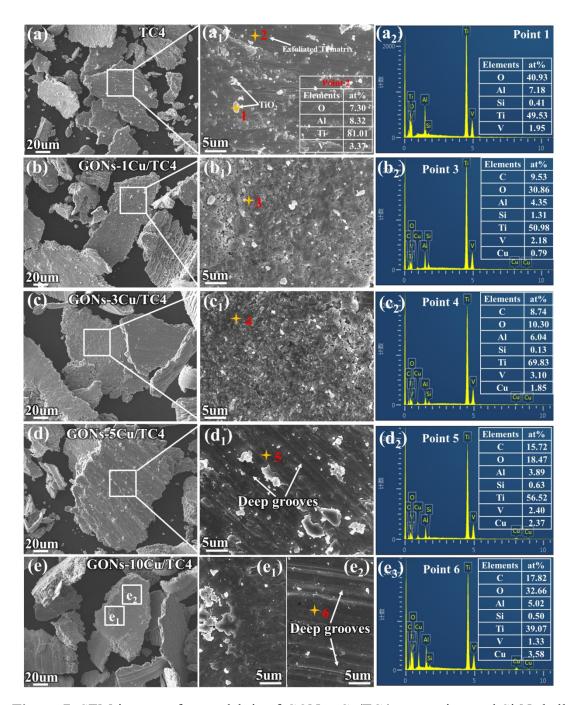
Figure 6. (a \sim f) Surface 3D morphologies of the wear scars for sintered composites

2 with various Cu contents and (a₁, b₁, c₁, d₁, e₁ and f₁) corresponding 2D cross-section

profiles of wear tracks, respectively.

The corresponding worn surfaces and the wear debris were also examined using the SEM and EDS. The results are shown in **Figure 8** and the obtained compositions are listed in **Table 4**. The amount of wear debris on the worn surface decreases with increasing copper content in the composites. The worn surface on the samples of the TC4 alloy is consisted of deep grooves and parallel furrows with some detached edges, indicating significant ploughing features during sliding wear. Ploughing or plastic deformation is a typical characteristic of the abrasion and adhesion wear [38]. Elements detected in the grooves are mainly Ti, Al, V and O (Point 1 in **Figure 8a**1 and spectrum in **Figure 8f**), and the content of oxygen is as high as 30.21 at.%, indicating significant oxidation of the alloy in the surface layer during the reciprocal wear process.

The grooves on the samples of the composites become gradually smoothened with increasing the copper content. In addition to the elements existed in the grooves on the surface of the TC4 alloy, Si, C and Cu elements are also found on the surface of the composite samples. The element Si is transferred from the Si_3N_4 balls, indicating that the adhesion wear occurs between the composite and Si_3N_4 ceramic ball.



2 **Figure 7.** SEM images of wear debris of GONs-xCu/TC4 composites and Si₃N₄ ball

- 3 frication pair after sliding wear test under 50 N load. (a) TC4, (b) GONs-1Cu/TC4, (c)
- 4 GONs-3Cu/TC4, (d) GONs-5Cu/TC4, (e) GONs-10Cu/TC4, (a₁)~(e₁) enlarged
- 5 images of corresponding marked region in Figures (a)~(e) and (a₂)~(e₃) EDS result of
- 6 points 1, 2, 3, 4, 5 and 6, respectively.

Besides the grooves and the wear debris, delamination phenomena are also 1 observed on the sample surface of the composites (Figures 8c1, d1 and e1). At some 2 delaminated regions (Figure 8d₁), high content of carbon is detected by EDS (Figure 3 8f), revealing the pulling out of the GONs from the interfaces in the composites. 4 Furthermore, microscale voids and cracks are found on the sample surface of composites with the content of copper higher than 3wt.% (Figures 8d₁ and e₁). 6 According to the aforementioned phenomenon, the wear on the TC4 alloy or the 7 TC4 based composites is a comprehensive mechanical-chemical coupling process. 8 9 The chemical reactions mainly involve the oxidation of the metals in the alloys or composites. The sliding process of the Si₃N₄ ball on the surface applies a complicated 10 dynamically localized loading onto the sample, and this added load fluctuates 11 12 continuously with changes of worn morphology, local temperature of sample, and formation of wear debris. Under wear process, the mechanical behavior of the TC4 13 alloy or the TC4 based composites involves ploughing or plastic deformation, work 14 15 hardening, local crack initiation and propagation, local welding and pulling out of the GONs. 16 17 Generally, the harder or stronger the metallic materials is, the lower the coefficients of friction and the better the tribological properties are [39]. However Ti 18 and its alloys are differnt, i.e., the higher of their strength is, the higher is the 19 coefficients of friction and the poorer of their tribological properties [40]. In this study, 20 we provide a solution for improving the wear resistance of the TC4 based metallic 21

materials by forming the composites via SPS using TC4 powders modified with Cu

and GONs, and we have achieved both high strength and improved tribological properties of Ti matrix composites (**Figure 5**).

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A schematic illustration on the tribological behavior and the wear mechanism of the Ti matrix composites is shown in Figure 9. After sintering, the Widmanstätten microstructure is formed in the TC4 alloys (Figure 3a and Figure 9a₁). The microstructures of the composites include Ti₂Cu, TiC and GONs in addition to the Widmanstätten microstructure (Figure 9b₁). Since the heat conductivity of the Ti alloys is low, the heat generated during sliding is accumulated, thus leading to the increase of the temperature. On one hand, the high temperature accelerates the oxidation of the alloy thus resulting in the exfoliation of the oxides (Figure 9a₂). On the other hand, the high temperature causes the softening of the alloy, thus leading to the enlargement of the contact area under the same load. Besides, under the locally and instantaneously high temperature at the contact asperities, the hard protrusions at the surface of the Si₃N₄ ball and its severe penetration into the soft matrix leave significant furrows on the sample surface and remove some materials in the grooves (Figure 9a₂). All these factors cause the increase of the wear volume loss and reduce the wear resistance of the Ti alloy. Introducing GONs and copper to the interfaces of the TC4 powders leads to the formation of the nanoscale precipitates of Ti₂Cu and TiC during the SPS. These fine

and reduce the contact area during sliding. The increase of strength also increases the

and stable precipitates act as hard obstacles to prevent dislocations' movement under

the high temperature, thus resulting in the increase of the high temperature strength

1 resistance of the composites to the formation of furrows, thus resulting in the

2 reduction of both the groove depth and the roughness of the surface (Figure 9b₂).

3 Furthermore, the remaining elemental copper and GONs existed at the interfaces

between the matrix grains enhance the heat conduction due to their high heat

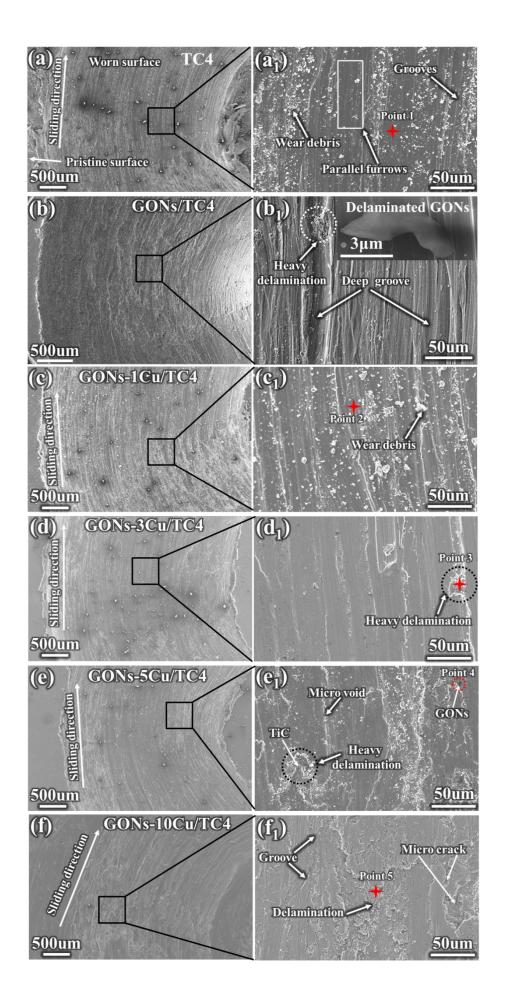
conductivities. This can lower the increase rate of the temperature caused by sliding,

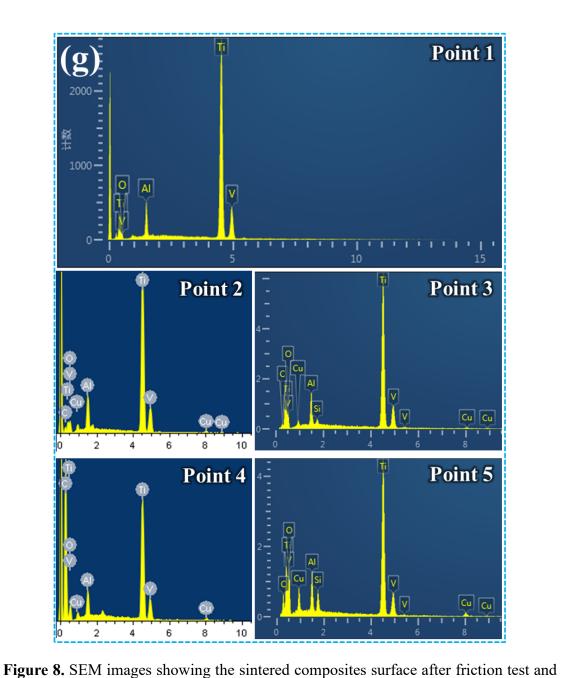
slow down the oxidation rate, and reduce the oxide wear debris.

However, with the further increase of the content of copper, the density of the Ti₂Cu precipitates increases, thus resulting in the increase of the crack initiation probability at the interfaces between the Ti₂Cu particles and the matrix due to stress localization at these interfaces under sliding wear. Therefore, excessive amount of Ti₂Cu precipitates in the Ti matrix composites have caused a deleterious effect on the wear resistance of the composites.

Besides, the pulling-out of the GONs from the interfaces between the matrix grains separates locally the Si₃N₄ ball and the composites, preventing the direct contact and acting as a lubricant (**Figure 9b₂**). Furthermore, since the TiC forms *in situ* at the interface between the TC4 matrix and the GONs, the TiC and the remaining GONs are fused into a brittle structure, which is denoted as TiC@GONs here. During sliding, the brittle TiC@GONs are crushed and the TiC and GONs are separated from each other, thus reducing the local penetration depth of the abrasive asperities when in contact, finally lowering the wear rate [41]. Hence, the improvement of the tribological properties of GONs-xCu/TC4 composites is mainly due to the strengthening by the nanoscale precipitates (i.e. Ti₂Cu and TiC), the fragileness of the

- brittle TiC@GONs structure, and the enhanced heat conduction resulted from the high
- 2 heat conductivity of the preserved elemental copper and GONs.





the EDS analysis results on the worn surfaces. (a) TC4, (b) GONs/TC4, (c) GONs-Cu/TC4, (d) GONs-3Cu/TC4, (e) GONs-5Cu/TC4 and (f) GONs-10Cu/TC4,

(g) corresponding to EDS analysis on point 1, 2, 3, 4 and 5 in Figure (a₁), (c₁), (d₁),

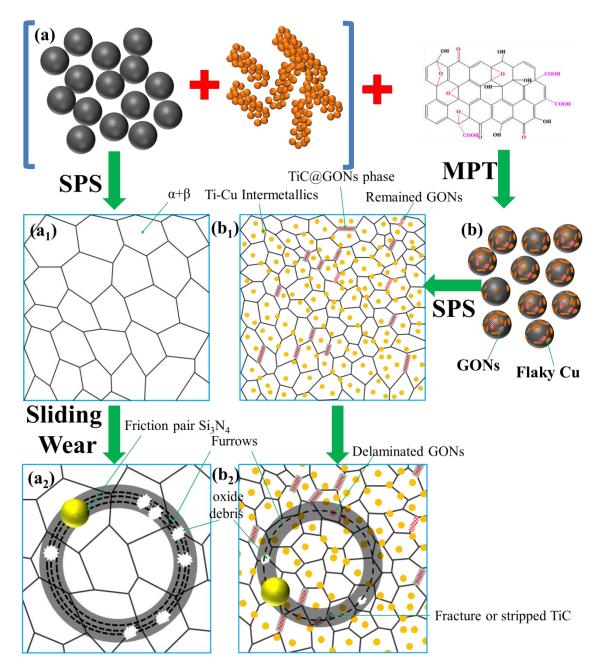
6 (e_1), and (f_1), respectively.

1 Table 4. Chemical compositions (in at. %) of different positions on the worn surface

2 in Figure 8.

| Position | Ti | Al | V | 0 | Si | C | Cu |
|----------|-------|------|------|-------|------|-------|------|
| Point 1 | 61.09 | 5.32 | 3.38 | 30.21 | 0 | 0 | 0 |
| Point 2 | 61.54 | 5.22 | 3.02 | 26.92 | 0.85 | 1.54 | 0.91 |
| Point 3 | 58.13 | 5.68 | 1.78 | 20.33 | 1.07 | 12.77 | 0.24 |
| Point 4 | 4.36 | 1.67 | 0.41 | 6.46 | 0.56 | 85.75 | 0.89 |
| Point 5 | 34.26 | 4.17 | 1.04 | 41.21 | 2.07 | 13.58 | 3.67 |

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- Figure 9. Schematic diagram and mechanism of tribological behavior of Ti matrix
- 2 composites co-strengthen with GONs and Cu in this work. *MPT=Modification

powders technology

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4 Conclusions

In this study, powder metallurgy and SPS technology have been applied to improve the microstructure and tribological properties of TC4 matrix composites with the reinforcement of 0.3 wt% GONs and Cu. The unreacted GONs, in-situ formed TiC particles or layers and Ti₂Cu nanoscale precipitates were generated in the Ti matrix composites, leading to excellent hardness and tribological properties. The peak hardness (411 HV) appears at the composites with 5 wt.% copper, and it is ~25% higher than the hardness of the TC4 alloy (330 HV). The change of the coefficients of friction and the wear volume loss with the content of copper show similar trends, i.e., decreasing when the content of copper is less than 3wt.% but increasing when the concentration is larger 3 wt.%. The GONs-xCu/TC4 composites show the lowest coefficients of friction (0.12) and the wear volume loss (0.155 mm³) among the alloys and composites studied in the present work. The improvement of the tribological properties is due to the strengthening by the nanoscale precipitates (Ti₂Cu and TiC) formed at high temperatures, self-lubricating effect introduced by the pulled out GONs from the composites, the fragileness of the brittle TiC@GONs structure, and the enhanced heat conduction resulted from the high heat conductivity of the preserved elemental copper and graphene oxides.

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