SLIDING-WEAR BEHAVIOR OF ALUMINUM-MATRIX COMPOSITES REINFORCED WITH GRAPHENE AND SiC NANOPARTICLES

Kaifeng Zheng, Xiaoming Du*, Haotian Qi, Tong Zhao, Fengguo Liu, Bo Sun
School of Materials Science and Engineering, Shenyang Ligong University, Shenyang 86110159, China

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The microstructure, hardness and tribological properties of hot-pressed Al 7075 matrix composites containing the same SiC nanoparticles and different amounts of graphene were investigated. The results show that the Al-SiC-graphene composites exhibit higher hardness and wear resistance compared with Al-SiC composite and Al 7075 alloy. The friction coefficients decrease significantly with the increase of graphene content in the composites, whereas the limited impact on reducing friction is found on SiC nanoparticles. The dominant wear mechanism of Al-SiC-graphene composites transfers from abrasive wear to delamination wear with the increase of graphene content in the composites. It is believed that the hardness and wear-resistance improvements of the Al-SiC-graphene composites are attributed to the unique strengthening effect as well as the good lubricating efficiency of graphene, which demonstrates that graphene is an ideal filler for aluminum-matrix composites, acting as not only a favorable reinforcement but also an impactful lubricant.

Keywords: graphene, SiC nanoparticle, aluminum matrix composite, wear mechanism

1 INTRODUCTION

Aluminum-matrix composites (AMCs) have attracted extensive attention in recent years because of their excellent mechanical properties, high strength, excellent wear resistance and lightweight properties.1,2 They have a wide range of applications in aerospace, military, automotive and other fields. Among the many reinforcing phases, such as graphite,3 carbon nanotube,4 SiC5 B4C6 silicon carbide and graphene are considered to be excellent reinforcements of AMCs because of their good physical and chemical properties.7

At present, silicon-carbide-reinforced metal-matrix composites have been reported in many publications,8,9 so that their wear resistance has been improved well. It is pointed out that increasing ceramic reinforcement phase is one of the main ways to improve the wear resistance of alloys.10 However, it has also been reported that soft-particle lubrication and hard-particle reinforcement are the inevitable trends to improve friction properties.11

Before this, the lubrication effect of graphene on the wear behavior has been reported in many papers.12,13 M. R. Akbarpour et al.14 studied the friction properties and wear behavior of SiC and CNT hybrid reinforced copper-matrix composites, and pointed out that a uniform lubrication layer was formed on the worn surface. SiC and graphite hybrid reinforced A356 composites were prepared.15 The wear properties and friction properties of A356 composites were also studied. The results show that the friction coefficient of hybrid composites is not decreased by the addition of graphite, but the wear rate of hybrid composites is reduced by the presence of SiC particles. Although some valuable research has been concentrated on the effect of pure SiC and graphene-reinforced aluminum composites, there are rare studies on the binary SiC-graphene-reinforced hybrid composites.

In this study, graphene and SiC nanoparticles hybrid reinforced Al7075 composites were prepared by the hot-pressing method. This study aims to investigate and compare the effect of pure SiC and synergistic binary SiC-graphene addition on the density, hardness, trib-
logical properties, and microstructure of aluminum hybrid composites.

2 EXPERIMENTAL PART

2.1 Materials

Al7075 (Al-Zn-Mg-Cu alloy) powder (10 μm, supplied from Beijing Hongyu Materials Company) was used as the matrix material. Graphene and SiC nanoparticles (800 nm, supplied from Shanghai Yunfu Nanotechnology Co., Ltd.) were used as reinforcement phases. The graphene is the commercial agent from Nanjing XFNANO Materials Tech Co., Ltd. All of the above materials were used without further purification.

Three samples were prepared by hot pressing in a vacuum at 610 °C for 1.5 h under a pressure of 50 MPa, in which these samples have the same SiC content of 0.25 w/(% and different graphene content, which were 0 w/(% (Al-SiC), 0.25 w/(% (Al-SiC-Gr1) and 0.5 w/(% (Al-SiC-Gr2), respectively. The preparation methods of the composites are reported in our previous work.16

All the composite samples were subjected to solutionizing at a temperature of 470 °C for 2 h followed by quenching using cold water. The quenched samples are then subjected to artificial aging for 16 h at a temperature of 140 °C.

2.2 Characterization and dry sliding wear test

The X-ray diffraction patterns of the samples were recorded on Rigaku Ultima IV X-ray diffractometer using Cu-Kα (λ = 0.15406 nm) radiation. Slow scans of the selected diffraction peaks were carried out in the step mode. The step size is 0.02°. The diffraction angle (2θ) was maintained between 20° and 100°. The average grain size was calculated by Williamson-Hall method.17 The density of the composite materials was measured by Archimedes’ method. The Vickers hardness of the sample was measured by a Vickers hardness instrument (HVS-50, Shanghai, China). The load force was set to 9.8 N and the dwell time was 10 s. The average hardness of the material is measured at least ten times in different areas of each sample.

A reciprocating friction and wear tester (MDW-02, Jinan, China) was used for the dry-sliding wear tests. All the samples were polished before wear testing to ensure that the surface of the sample has the same roughness. The dry wear tests were carried out at room temperature using a load of 10 N. The average sliding speed in the wear tests is 0.02 m/s and the total sliding distance is 24 m. Balls with a diameter 6.35 mm and made of GCr15 bearing steel (HRC63±3) were used for the wear tests. The average friction coefficient is calculated under steady-state conditions in which the wear test reaches the steady state for more than 90 s. The worn surface and balls surface were analyzed using a scanning electron microscope (TESCAN) equipped with EDS. The wear volume of the samples was analyzed based on the depth and width of the wear trajectory for pure and composite samples using a laser scanning confocal microscopy (LSCM) (LEXT OLS4100, Japan). The morphology of the debris produced by different samples in wear test was analyzed by scanning electron microscope (HITACHIS-3400N, Japan). The structures of the graphene during sintering and sliding were investigated by Raman spectroscopy using a Jobin-Yvon microspectrometer (LabRam HR, Jobin-Yvon Co. Ltd., France). The spectra were collected under ambient condition using the 514.5-nm line of an argon-ion laser.

3 RESULTS AND DISCUSSION

3.1. Microstructure

The X-ray diffraction patterns of three composite samples with different graphene content and the same SiC content are shown in Figure 1. The diffraction peaks of α-Al, SiC, MgZn2 and graphene were retrieved. As can be seen in Figure 1, all the samples have major α-Al peaks at 38.4° (111), 44.7° (200), 65.1° (220), 78.2° (311), 82.4° (311) and 99.1° (400). In addition to the diffraction peaks of α-Al matrix, some very weak diffraction peaks were also found. The intensity of these peaks is relatively weak. It is possibly due to the small amount of graphene and SiC nanoparticles in the composites. The weak diffraction peaks located at 26.5° (002) for Al-SiC-Gr2 and Al-SiC-Gr1 samples are of character peak in carbon X-ray diffraction pattern, which indicates that graphene has been integrated into the matrix materials. The result is consistent with that reported in the literature.18 The intensity of the diffraction peak for graphene decreases with the decrease of the content of graphene, such as Al-SiC-Gr1 sample. The weak diffraction peaks located at 35.8° (103) and 60.1° (110) in the three composites are of character peak in SiC with a trigonal crystal structure X-ray diffraction

Figure 1: XRD patterns of Al-SiC, Al-SiC-Gr1 and Al-SiC-Gr2
Moreover, aluminum carbide (Al₄C₃) peaks were not found in this analysis, indicating the better potential of graphene in strengthening Al-based composites.

**Table 1** summarizes the relative density and mean crystal grain size of the three composites. The relative densities of all the composite samples are more than 99 %, which shows that the prepared composites are dense, and the porosity of the composite is low. The relative density of the composites shows a decrease with increasing graphene contents. This may be attributed to the low density of graphene.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Relative density (%)</th>
<th>Grain size (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Al7075</td>
<td>–</td>
<td>3225.00±3.67</td>
</tr>
<tr>
<td>Al-SiC</td>
<td>99.4</td>
<td>1119.80±0.45</td>
</tr>
<tr>
<td>Al-SiC-Gr1</td>
<td>99.4</td>
<td>790.20±3.63</td>
</tr>
<tr>
<td>Al-SiC-Gr2</td>
<td>99.2</td>
<td>427.20±9.39</td>
</tr>
</tbody>
</table>

From the mean grain size of the three composites obtained from XRD in **Table 1**, it can be seen that the content of graphene has a significant effect on the grain size of aluminum matrix composites. A fine dispersion of the SiC nanoparticles and graphene in the matrix may prevent the grain growth of the matrix alloy during the hot-pressing process. Grain size decreases with increasing the graphene content. With an increase in the content of the graphene, the area of interface boundaries between the graphene and Al matrix increases and the dislocations are piled up at the interface boundaries similar to the grain boundaries. Apart from the dispersion strengthening effect, the addition of the graphene refines the grain size and introduces a high dislocation density in the matrix.

### 3.2 Hardness

Results from the Vickers hardness test of Al-SiC, Al-SiC-Gr1 and Al-SiC-Gr2 are shown in **Figure 2**. In previous work, the Vickers hardness of Al7075 alloy prepared by the same method and parameters was 92.0±4.6 HV.²⁶ It can be seen from **Figure 2** that the hardness of the three composites samples is significantly higher than that of the Al7075 alloy, which represents an increase of 26.6 %, 21.9 % and 38.9 % for Al-SiC, Al-SiC-Gr1 and Al-SiC-Gr2, respectively. The increase of the hardness results from dispersion strengthening of SiC nanoparticles and graphene, refining grain strengthening of the aluminum matrix and thermal mismatches between α-Al and SiC, graphene. The mismatch strain due to the difference of the coefficient of thermal expansions between the SiC nanoparticles, graphene and Al matrix develop high-density dislocations, which strengthens the matrix. SiC nanoparticles, graphene obstruct the movement of high density dislocations, resulting in an improvement of the hardness.

### 3.3 Tribological properties

The variation in the measured average coefficient of friction (COFaverage) with the weight percent of graphene in the hybrid-reinforced composites is shown in **Table 2**. The results show that within the set of test conditions used, the average coefficient of friction in the Al7075 alloy is larger than that of the composites. And the coefficient of friction in composites significantly decreased with an increase of the graphene content in the composites. The coefficient of friction decreased by about 22 % and 43 % with the reinforcement of 0.25 % and 0.5 % graphene, respectively. This shows that graphene plays a role in the lubrication in the wear test of the composites. The existence of graphene makes the matrix form a uniform and continuous self-lubricating layer on the worn surface during the wear test. The extremely effective lubrication of graphene was previously reported.¹²,¹³

<table>
<thead>
<tr>
<th>Sample</th>
<th>Average coefficient of friction</th>
<th>Depth (μm)</th>
<th>Width (μm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exp. Cal.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Al7075</td>
<td>0.39</td>
<td>65</td>
<td>270</td>
</tr>
<tr>
<td>Al-SiC</td>
<td>0.37</td>
<td>54</td>
<td>225</td>
</tr>
<tr>
<td>Al-SiC-Gr1</td>
<td>0.29</td>
<td>33</td>
<td>144</td>
</tr>
<tr>
<td>Al-SiC-Gr2</td>
<td>0.21</td>
<td>21</td>
<td>93</td>
</tr>
</tbody>
</table>

In the studies of P. V. Trinh et al.¹⁹ and M. R. Akbarpour et al.,¹⁴ a theoretical calculation method for the coefficient of friction based on hybrid criterion is proposed, and it is pointed out that the content of the reinforcing phase on the friction surface is independent of the load force applied in the wear test. At the same time, the calculation formulas of theoretical coefficient of friction are also proposed. The formulas for calculating Al-SiC and Al-SiC-Gr composites are as follows in Equations (1) and (2):

\[
\mu_{\text{Al-SiC}} = \frac{\mu_{\text{Al}}(A - A_{\text{SiC}})E_{\text{Al}} + \mu_{\text{SiC}}A_{\text{SiC}}E_{\text{SiC}}}{A_{\text{SiC}}E_{\text{SiC}} + (A - A_{\text{SiC}})E_{\text{Al}}} \tag{1}
\]
\[ \mu_{\text{Al-SiC-Gr}} = \mu_{\text{Al}}(A - (A_{\text{Gr}} + A_{\text{SiC}}))E_{\text{Al}} + \mu_{\text{Gr}}A_{\text{Gr}} + \mu_{\text{SiC}}A_{\text{SiC}}E_{\text{SiC}} - \mu_{\text{Al}}A_{\text{Al}}E_{\text{Al}} + A_{\text{Gr}}E_{\text{Gr}} + A_{\text{SiC}}E_{\text{SiC}} + (A - (A_{\text{Gr}} + A_{\text{SiC}}))E_{\text{Al}} \]  

(2)

where \( \mu_{\text{Al-SiC-Gr}} \), \( \mu_{\text{Al-SiC}} \), \( \mu_{\text{Al}} \), \( \mu_{\text{Gr}} \) and \( \mu_{\text{SiC}} \) are the coefficients of friction of Al-SiC composites, Al-SiC-Gr composites, Al7075, graphene and SiC, respectively. \( A \), \( A_{\text{Gr}} \) and \( A_{\text{SiC}} \) represent the mass fraction of Al7075, graphene and SiC, respectively. \( E_{\text{Al}} \), \( E_{\text{Gr}} \) and \( E_{\text{SiC}} \) represent the elastic modulus of Al7075, graphene and SiC, respectively. The data required to calculate the friction coefficient are shown in Table 3.

**Table 3:** The data for calculating theoretical friction coefficients

<table>
<thead>
<tr>
<th>Symbol</th>
<th>( E_{\text{Al}} ) (GPa)</th>
<th>( E_{\text{Gr}} ) (GPa)</th>
<th>( E_{\text{SiC}} ) (GPa)</th>
<th>( \mu_{\text{Graphite}} )</th>
<th>( \mu_{\text{SiC}} )</th>
<th>( \mu_{\text{Al,average}} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>Value</td>
<td>70</td>
<td>1000(^{[20]})</td>
<td>420(^{[14]})</td>
<td>0.1</td>
<td>0.7(^{[14]})</td>
<td>0.39</td>
</tr>
</tbody>
</table>

Because the coefficient of friction of graphene has not been reported at present, it is replaced by the friction coefficient of graphite. However, there are also many reports that the coefficient of friction of graphene will be much lower than that of graphite.\(^{21-22}\) The theoretical and experimental values of the coefficient of friction of the three composite samples are shown in Table 2. It can be seen that there is a certain difference between the theoretical values and the experimental values. The difference increases with the increase of graphene content. There are possibly two reasons for this phenomenon. Firstly, the dispersion of the reinforcing phase in the Al7075 matrix alloy is assumed to be completely dispersed and uniform in the calculation of the theoretical coefficient of friction. However, the dispersion of the reinforcing phase in the Al7075 matrix alloy is not completely uniform. Moreover, the coefficient of friction of graphene is replaced by the coefficient of friction of graphite that should be greater than that of graphene. This results in the theoretical value of friction coefficient of composites being higher than the experimental value. With the increase of graphene content, this deviation becomes more serious.

The depth and width of the wear track for Al7075 alloy and the hybrid composites are listed in Table 2. Both the wear-track depth and width, which are criteria of the volume of worn material, were reduced due to adding SiC and increase of the amount of graphene to 0.5 w/\%. The reason for the improved wear resistance of the composite samples is their hardness (Figure 2), which is the result of the presence of SiC nanoparticles and graphene. In this way, the reinforcing phases act as load-bearing elements and reduce the actual contact area between the counterface and Al matrix. This improves the wear resistance.

In order to investigate the wear mechanism, the surfaces of the worn samples were examined under the SEM. Figure 3 shows the SEM morphology of the worn surfaces for three composites under dry-sliding conditions. In Figure 3a and Figure 3b it can be found that there are very obvious abrasive grooves in the sliding direction. This indicates that the wear process mainly took place by abrasive wear for the Al-SiC composite. This is consistent with the wear mechanism of Al-SiC.
composites reported by J. Zhang et al.\textsuperscript{23} The deep abrasive grooves observed in Figure 3a most probably produced by pushing the SiC nanoparticles on the surface of the Al matrix. This process is equivalent to micro-cutting of aluminum matrix surface with SiC particles. A three-body wear state is formed between the worn sample surface and the friction pair.\textsuperscript{6} In addition, the surface patch state and wear pits\textsuperscript{11} caused by adhesive wear can be observed on the worn surface. It is also found that there are many wear debris adhered to the worn surface. EDS results show that these wear debris mainly originate from the aluminum matrix.

In Figure 3c and Figure 3d there are still some abrasive grooves. However, those abrasive grooves are shallow and which of surfaces are nearly covered by adhesive layers. It is indicated that the wear mechanism of Al-SiC-Gr1 composite is the combination of abrasive wear and delamination wear. In Figure 3e and Figure 3f the abrasive grooves on the worn surface of the Al-SiC-Gr2 composite almost disappear. On the worn surfaces, areas with a fractured appearance can be observed. It can be seen that the layer of material has been removed as debris from the surface and that the debris is in the form of thin sheets. In reference to SEM studies of worn surfaces, it could be observed that wear damage was caused by the plastic flow of the matrix, indicating that the delamination wear is the dominant wear mechanism of the Al-SiC-Gr2 composite. It is concluded that the wear mechanism of Al-SiC-graphene composites transfers from abrasive wear to the delamination wear with the increase of graphene content in composites.

Figure 4 shows SEM micrographs of the worn surfaces of the balls corresponding to the three composite samples. The large number of abrasive grooves can be found in Figure 4a. This confirms that the three-body abrasive wear exists in the dry-sliding wear test for Al-SiC composite. For Al-SiC-Gr1 composite, there are only a few abrasive grooves and adhesive layers on the worn surfaces of the ball (Figure 4b). However, there are only few abrasive grooves and a lot of adhesive layers on the worn surfaces of the ball for Al-SiC-Gr2 composite, as shown in Figure 4c. Moreover, a large amount of wear debris is also coated on the surface of the balls. Al adhesion can be found on the surface of the ball from the EDS result (Figure 4d). These results are consistent with the wear-surface analysis results of the composites mentioned above.

Studying the shapes and sizes of the wear debris provides clues to the wear mechanisms extant and brings information about the wear state. Figure 5 shows SEM micrographs of the wear debris of three composite samples. There are three types of wear debris according to their morphology and sizes:

1) flake wear debris. This wear debris is the most common form, with sizes ranging from 100 \( \mu \text{m} \) to 300 \( \mu \text{m} \). Flakes can be observed in Figure 5a and Figure 5b, ranging in thickness from 10 \( \mu \text{m} \) to 30 \( \mu \text{m} \). This debris is caused by adhesion in the sliding direction and delamination of the balls.

![Figure 4: SEM micrographs of the surfaces of balls used by: a) Al-SiC, b) Al-SiC-Gr1, c) Al-SiC-Gr2, d) the EDS of the selected area in c)](image-url)
2) fine debris. Contrary to large wear debris, a small number of fine particles as well as sheet-like wear debris with a range of 1 μm to 20 μm in planar are also collected (Figure 5b and Figure 5c). The generation of fine wear debris is attributed to an abrasive micro-cutting effect. Generally, the appearance of a smaller size of wear debris is related to better anti-wear performance of the material. 3) corrugated debris. The amount of the debris is very small. It can be observed that there are many obvious layers on the surface of the debris. It is mainly due to the joint action of shear force and constant friction force in the sliding direction. The results of EDS analysis indicate that most of the wear particles origin from aluminum matrix alloy.

In order to investigate the evolution of graphene in the process of hot-pressing and sliding, Raman spectroscopy analysis was performed on graphene, as-prepared Al-SiC-Gr2 composite and the wear track of Al-SiC-Gr2 composite, respectively, as shown in Figure 6. It is clear from Figure 6 that the typical D-band (from defect), G-band (from graphite), and 2D band (shape of the second-order Raman bands) of graphitic carbons are detected at 1335 cm⁻¹, 1590 cm⁻¹, and 2773 cm⁻¹ respectively, for the initial material. During hot pressing, the band shift and broadening can occur by structural defects and domain size in graphitic materials, and damage of crystalline graphitic structure. However, the band broadening can only be observed during sliding.

This shift and broadening can be attributed to the residual strain in the graphitic materials, which is induced by the compressive stress during hot-pressing and the shear stress at the contact surface during sliding, resulting in the structure disruption of the graphene. The ratio of the intensity of the D band to the intensity of the G band \((I_D/I_G)\) is normally used to characterize the degree of defects of the graphite-based material. The larger \(I_D/I_G\) ratio represents the higher defect density in the graphitic structure. The results show that the \(I_D/I_G = 1.30\) of worn surface is higher than that of the initial graphene \((I_D/I_G = 1.01)\) and non-worn surface \((I_D/I_G = 1.25)\). This indicates more exfoliated and fragmented graphene flakes contribute to the formation of a lubricating tribolayer on the contact interface, resulting in an improved wear resistance. A similar result has been observed by other authors for copper-matrix composites containing graphene.

4 CONCLUSIONS

In the present work, graphene and SiC nanoparticles hybrid reinforced Al7075 composites were prepared by hot pressing. The microstructure, hardness and wear behavior were studied. X-ray diffraction analysis showed the presence of the carbon peaks and no proof of Al₄C₃ formation. The graphene content in the base Al 7075 gives rise to a significant improvement in the hardness, density and wear resistance. The extensive increase in hardness of the Al 7075 matrix composites is due to the dispersion strengthening of the SiC nanoparticles and graphene, refining grain strengthening of \(\alpha\)-Al matrix and thermal mismatches between \(\alpha\)-Al and SiC, graphene. Consistent with the hardness improvements, were marked improvements in wear resistance. On increasing the graphene content, the coefficient of friction values were reduced and were consistent with the theoretically calculated friction coefficient of composite materials. The coefficient of friction decreased by about 22 % and 43 % with the reinforcement of 0.25 w% and 0.5 w% graphene, respectively. SEM observation along with EDS for the worn surfaces, counter-face and debris shows that the wear mechanism
of Al-SiC-graphene composites transfers from abrasive wear to delamination wear with the increase of graphene content in composites and confirms the formation of graphene lubricating tribolayer between the wear surfaces, which influences the effect of wear behaviour and coefficient of friction values in the composites. The Raman spectroscopy confirms the increase of \( I_D/I_G \) ratio of the graphene after hot pressing, indicating the graphene is to some extent disrupted in the structure during the hot-pressing process. The \( I_D/I_G \) ratio reaches a maximum after sliding, which demonstrates the formation of a graphene-rich lubricating tribolayer at the contact surface, which hinders the friction and wear.

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