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One-step synthesis of sandwich-type Cu/graphene/Cu ultrathin foil with enhanced property via electrochemical route

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HIGHLIGHTS

- The Cu/Gr/Cu ultrathin foil is synthesized by one-step electrochemical route.
- Cu is deposited successively on substrate and GO under different overpotentials.
- The foil is of thickness as thin as 4–5 \( \mu \)m and twice tension strength enhancement.
- The high performances are attributed to strong interface bonding and graphene.
- This work can also be used to other metal/Gr/metal layered composites.

GRAPHICAL ABSTRACT

A layered structure has a better effect on improving performance of the graphene-reinforced composites due to its unique two-dimensional structure and excellent properties. In this paper, a novel "one-step" electrochemical route was proposed for synthesizing the graphene-reinforced ultrathin copper (Cu) foil with high performance. The process includes: 1) A loose graphene oxide (GO) membrane, was prepared by electrophoresis deposition (EPD), that allows Cu ions passing through; 2) According to the difference of Cu deposition potential on different substrates, a potential step was designed for electrodepositing Cu successively on both sides of the GO membrane, i.e., the bottom Cu layer forms under low over-potential, while the top Cu layer forms under high over-potential. The experimental results show that the foil thickness reaches to as thin as 4–5 \( \mu \)m, and the tensile strength is almost twice as large as that of pure Cu foil. The process is simple, controllable and possible mass production, and expected to further practical applications in fields of Cu clad plate, printed circuit board and lithium-ion battery cathode collector system for saving raw material and also the space. In addition, this work proposes a new idea for preparing the layered composites via electrochemical route.

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1. Introduction

As an important raw material, electrolytic copper (Cu) foil has been widely used in areas, such as Cu clad plate (CCL), printed circuit board (PCB), and lithium ion battery cathode collector system [1]. For instance, in lithium ion battery, Cu foil is not only the carrier of negative electrode, but also acts as the electron collection and conduction [2]. In commercial applications, to produce ultrathin Cu foil with high-strength and high-conductivity is still a challenge [3]. In general, ultrathin copper foil is defined as thickness of <12 μm [4]. In order to improve its performance, many strategies have been proposed including optimizing the electrodeposition process [5] and adding nano-reinforcements, such as Si₃N₄ [6], graphite, [7] carbon nanotubes, [8,9] graphene [10,11], and so on.

As a two-dimensional planar structure of a single atomic layer of carbon, Graphene (Gr) is of a theoretical thickness 0.35 nm and its carbon atoms are all tightly connected with each other in the form of sp² hybridization. Therefore, Gr each lattice consists of a stable hexagonal structure formed by three sigma bonds, which results in unique properties, such as carrier mobility reach up to 200,000 cm²/(V·s) at room temperature [12,13], and tensile strength up to 130 GPa and elastic modulus to 1.1 TPa. In addition to broad application prospects in the fields of microelectronics, energy, information and so on, Gr is especially considered to be an ideal reinforcement in composites [14–16].

In recent years, high performance Cu-Gr composites prepared by electrochemical deposition have attracted much attentions [17–19]. The basic approach is to add Gr or GO or its derivatives into the electrolyte, and treats it with ultrasound or surfacetant for uniform dispersion. And then the co-deposition of copper and graphene are occurred on the cathode under electrical current to form a film or coating. Table 1 lists the summary of researches on Cu-based composites containing Gr and carbon nanotubes (CNTs). Obviously, the Gr co-deposition greatly increases the mechanical properties of Cu foil, and also makes its electrical conductivity slightly improve or remains unchanged. In comparison, some work on layered Cu-carbon nanocomposite films exhibit better enhancing effect on mechanical properties, such as up to 1.5 GPa compressive strength. The ultra-high strengths of the Cu-Gr nanolayered structures indicate the effectiveness of graphene in blocking dislocation propagation across the Cu-Gr interface [22]. Besides, Yi et al. [23] prepared a sandwich structural CNTs/Cu/CNTs composite, where a template method was used for preparing the freestanding CNTs network film. Then, using the film as the cathode, two anodic electrodes were applied during electrodeposition, so that the electrodeposition could occur simultaneously on both sides of the CNTs film.

In generally, commercial Cu foil is divided into several thickness ranges, i.e., thick foil (>70 μm), conventional foil (18 μm–70 μm), thin foil (12 μm–18 μm) and ultrathin foil (<12 μm), respectively [25]. And, the tensile strength is around 350 MPa. Therefore, the value higher than 400 MPa is called the high-performance Cu foil.

Obviously, because of low strength, the ultrathin foil is prone to wrinkle and tearing, which make its production, transportation and application very difficult. At present, the carrier method is adopted to prepare the ultrathin Cu foil in industry, which refers to a process, i.e., deposit the ultrathin Cu foil firstly on a thick Cu foil, and then peels it off [26]. Obviously, the operation of this process is tedious and the industrial production cost is high, meanwhile the performance of the ultrathin Cu foil has not been improved.

Electrophoretic deposition (EPD) is a surface treatment technology that has been widely used in processing advanced ceramics and coatings. In brief, EPD is a process for depositing charged colloidal particles on electrode under electric field. By using EPD technology many researchers prepared the dense GO films as corrosion resistant coatings [27–30].

This paper proposed a novel process to prepare the layered Cu/Gr/Cu composite foil, i.e., according to the difference of Cu deposition potential on different substrates, a potential step was designed for electrochemical deposition, which achieved the one-step preparation of the Cu/Gr/Cu ultrathin foil, as shown in Fig. 1. The foil's thickness reached to as thin as 4–5 μm and had a twice tension strength enhancement, which was firstly reported. The process is of advantages, such as simple and controllable, and expects to further mass production in industry. Further practical applications are expected in the areas, such as the Cu clad plate (CCL), printed circuit board (PCB), and lithium ion battery cathode collector system for saving raw material and also the space. In addition, this work provides a new idea for preparing the layered composites via electrochemical route.

2. Experimental section

2.1. Preparation of the EPD-GO film

All experimental reagents were in analytical grades, and deionized water was utilized to prepare the electrolyte. Fig. 1 illustrates the preparation process of the Sandwich-type Cu/Gr/Cu ultrathin foil. The detailed process for preparing the EPD-GO film was as follows:

1) Graphene oxide (GO) was synthesized from natural graphite powders by a modified Hummers’ method and the detailed process operations was described in supporting information.

2) GO was first dispersed in deionized water and sonicated for 10 min at room temperature. A uniform and stable aqueous solution containing 0.5 mg·ml⁻¹ of GO was obtained.

3) A direct current (DC) electrophoresis method was used by using a DC power supply (KXX-3010D, Zhaoxin, China). Both the cathode and anode were Type 304 stainless steel with a size of 30 mm × 50 mm × 0.5 mm and the applied voltage was fixed at 5 V, while the electrophoresis time varied from 10 s to 60 s.

4) A hydrous EPD-GO film was obtained on the anode and dried in the air for later use. This loose film allowed Cu ions to permeate through and contacted with the stainless steel substrate.

2.2. Preparation of the Sandwich-type Cu/Gr/Cu layered composite foil

In view of the difference of the Cu deposition potential on different substrates, a potential step method in a three-electrode system was
designed for preparing the sandwich-type Cu/Gr/Cu foil by using an electrochemical workstation (CHI660E, Chenhua, China). The system consisted of a platinum foil with a large area, a mercury/mercuric sulfate (Hg/Hg₂SO₄) as the counter electrode, and a reference electrode. As shown in Fig. 2, the process was performed in an electrolyte consisting of 100 g·L⁻¹ CuSO₄, 30 g·L⁻¹ H₂SO₄, and the specific steps were: 1) had a thin EPD-GO film on the SS substrate; 2) carried out electrochemical deposition at a low overpotential (−0.55 V vs. Hg/Hg₂SO₄) to allow Cu ions passing through the GO film and get to SS substrate to form the bottom Cu layer; 3) Shift to a high overpotential (−0.65 V vs. Hg/Hg₂SO₄) to obtain the top Cu layer. In general, the Cu/Gr/Cu ultrathin foil could be stripped directly from the stainless steel substrate due to the inferior adhesive property, which was similar to the continuous production in industry. It also described in our previous work [20]. The foil samples were blown dry by hair drier for the subsequent characterizations and tests. For comparison, pure Cu foil was also prepared by using the same method.

2.3. Characterizations

The morphologies and chemical compositions of the GO and the foil samples were characterized using several analysis instruments, including a scanning electron microscope (SEM) (S-4800, Hitachi, Japan, and SIRON, FEI, The Netherlands) equipped with an energy-dispersive X-ray spectroscopy (EDS); a transmission electron microscope (TEM) (JEM-2100, JEOL, Japan), with the acceleration voltage of 200 kV; a Fourier transform infrared spectrometer (FT-IR, Nicolet iS10, Thermo Fisher, USA) with a scanning range of 525–4000 cm⁻¹ and a laser scanning confocal micro-Raman spectrometer (Raman) (LabRAM HR, HORIBA, France) with a laser excitation wavelength of 488 nm and scans with an extended range of 1000–3000 cm⁻¹.

Absorbance of GO was measured by using a diffuse reflectance spectra of a UV–vis spectrophotometer (UV-2550; Shimadzu, Kyoto, Japan) in absorption mode, in which deionized water was used as a background between 190 nm and 800 nm scopes.

2.4. Property measurements

The tensile strength of the samples was measured by an electronic universal testing machine (model CMT6203, MTS systems Co., Ltd.) in accordance with the testing method IPC-TM-650, where the foil width was 12.7 mm and tensile rate was 2 mm·min⁻¹. After tensile test, the cross-section fracture surfaces were observed by using SEM. The Gr strengthening mechanism was expressed by the following equation [31,32]:

$$\sigma_c = \sigma_m + \Delta\sigma_{LT} + \Delta\sigma_D + \Delta\sigma_{GR}$$  \hspace{1cm} (1)

where $\sigma_c$ and $\sigma_m$ were tensile strength of the composite and matrix, respectively. $\sigma_{LT}$, $\sigma_D$ and $\sigma_{GR}$ were the strengthening contributions from load transfer, dislocation strengthening and grain refinement, respectively. The square resistance of the foil samples was measured using a four-point probe instrument (RTS-9, 4 Probes Tech, China). The resistivity of all samples was calculated using the following equation:

$$\rho = d \cdot R_s$$  \hspace{1cm} (2)

where $\rho$, $d$ and $R_s$ were resistivity, thickness and square resistance of the foils, respectively.
3. Experimental results

Fig. 3 shows the microstructural characterizations of GO. SEM and TEM observations revealed that GO was of a nearly transparent lamellar structure and crimped at the edges. The strong diffraction spots of selected-area electron diffraction (SAED) pattern demonstrated it a well-defined hexagonal structure, and high crystallinity [33]. The FT-IR spectrum showed the GO surface contained oxygen-containing functional groups [34], involving O–H (hydroxy) stretching vibration near 3190 cm\(^{-1}\), C=O (carboxyl/carbonyl) stretching near 1719 cm\(^{-1}\), C=C (aromaticring) near 1621 cm\(^{-1}\), and C–O (alkoxy) stretching near 1043 cm\(^{-1}\). These groups would improve the dispersion of GO in aqueous. The UV–vis spectra had a peak at 227 nm, which was due to \(\pi \rightarrow \pi^*\) of C=C bond, and a shoulder at ~290–300 nm, corresponding to n \(\rightarrow\) \(\pi^*\) transition of the C=O bond [35]. The Raman spectrum indicated it was a typical GO, i.e., D peak near 1360 cm\(^{-1}\) revealing the defect density due to the lattice vibration leaving the Brillouin zone center; G peak near 1586 cm\(^{-1}\) was caused by sp\(^2\) hybridization of carbon atoms of in-plane vibration; 2D peak near 2700 cm\(^{-1}\) was the phonon resonance second-order Raman peak indicating the way of carbon atoms stack in graphene; and the scattering of D peak and 2D peak in the valley generates the D+ G peak near 2935 cm\(^{-1}\) [36–38].

Fig. 4 shows the SEM morphologies of the EPD-GO films with different durations. When the EPD time was short, the GO film exhibited a flat surface with fewer folds. The EDS element mapping, as shown in Fig. S1, also indicates that GO was successfully electrodeposited on the SS substrate. However, for the longer deposition time, the GO film became higher stack and fold significantly, due to the increased thickness.

Further, the compactness of the EPD-GO film was tested by measuring the open circuit potential (E\(_{ocp}\)) - time (T) curve and linear sweep voltammetry (LSV) curve in the copper sulphate electrolyte, respectively, as shown in Fig. 5. In general, the E\(_{ocp}\)-T curve is used to study...
the sensitivity of the interaction between working electrode and electrolyte as well as the probability of corrosion reaction. The more negative the $E_{\text{ocp}}$, the stronger the action, and the stronger corrosion. In the present case, as shown in Fig. 5(a), $E_{\text{ocp}}$ of the EPD-GO film varied from the positive potential close to the freestanding GO film to the negative potential of the SS as time increasing, which revealed that the electrolyte could pass through the EPD-GO film and interacted with the SS substrate. Similarly, Cu ions exhibited different deposition potentials on SS, EPD-GO and freestanding GO film, as shown in Fig. 5(b). Therefore, these results provided a possibility to prepare the Cu/Gr/Cu layered foil by using such an uncompact EPD-GO film.

Fig. 6 shows the SEM surface morphologies of the bottom Cu layer. As the deposition time increased, the Cu deposition on the SS surface continuously took place and became more and more compact, and eventually formed a thin Cu layer. The EDS elements mapping also confirmed the Cu deposition growth on the SS substrate, as shown in Figs. S2 and S3. In addition, the Cu surface showed a good appearance, as shown in Fig. 6(g), and the portion within the black dotted frame will be used for cross-section observation after tensile test, As shown in Fig. 6(h).

Numerous studies have demonstrated that GO can be reduced to reduced GO (or Gr) during electrochemical deposition. In the present work, from Raman spectra (Fig. S4), it could be seen that the peak strength ratio ($I_D/I_G$) of the Gr in the Cu/Gr/Cu foil decreased significantly, when compared with the EPD-GO film, which indicated that GO was reduced to Gr during the foil preparation. In addition, in order to observe the Sandwich structure more clearly under SEM, two kinds of composite foils with different Gr interlayers thickness were prepared, i.e., the Cu/Gr1/Cu foil with thinner Gr interlayer, and the Cu/Gr2/Cu foil with relatively thicker Gr interlayer. Fig. 7 shows the EDS elemental mappings of the cross-sectional Cu/Gr2/Cu foil. The Sandwich structure was clearly observed and the Gr
thickness was about 0.7 μm. The presence of O element was attributed to the cuprous oxide formation during electrodeposition.

Fig. 8 shows the SEM morphologies of the foils’ tensile fracture surfaces. It could be seen that the total foil thickness was about 4.2–4.5 μm including the Gr interlayer < 0.7 μm, which was much less than the thickness 10 μm of commercial ultrathin foil, while in Fig. 8 (b), the Gr interlayer was too thin to be observed. In addition, from the top view, the edges of the pure Cu foil fracture surface were relatively flat, while the two composite foils showed jagged and tearing edges, which indicated a high ductility and strong interface bonding strength between Gr and Cu [39–41]. Fig. 8 shows the SEM morphologies of the bottom Cu layer obtained with different deposition time at different magnifications; (a) and (b) 1 min; (c) and (d) 5 min; (e) and (f) 10 min; (g) general view of the foil; (h) foil after tensile test.

Fig. 9 illustrates the test results of tensile strength and electrical properties of the different foils. Obviously, compared with pure Cu foil (270 MPa), the Cu/Gr1/Cu foil with thinner Gr interlayer exhibited the highest tensile strength up to 535 MPa, i.e., almost double the value. In regarding to the electrical conductivity of the foils, the resistivity was calculated by using the following equation:

$$ \rho = d \cdot R_s $$

where $\rho$ is the resistivity, $d$ is the thickness and $R_s$ is the square resistance. As shown in Fig. 9 (b), the resistivities of the Cu/Gr1/Cu foil with thinner Gr interlayer and pure Cu foil were comparable, but the
Cu/Gr2/Cu foil with relatively thicker Gr interlayer increased significantly.

**4. Discussion**

In general, there are three possible strengthening mechanisms accounting for the enhanced strength of the metal - Gr composites, i.e., load transfer strengthening [44], dislocation strengthening [22] and grain refinement [21], and it can be expressed by the following equation [31,33]:

\[ \sigma_c = \sigma_m + \Delta \sigma_{LT} + \Delta \sigma_D + \Delta \sigma_{GR} \]  

where \( \sigma_c \) and \( \sigma_m \) represent tensile strength of the composite and matrix, respectively; \( \Delta \sigma_{LT} \), \( \Delta \sigma_D \) and \( \Delta \sigma_{GR} \) are the strengthening contributions from load transfer, dislocation strengthening and grain refinement, respectively.

For the Sandwich-type Cu/Gr2/Cu layered composite foil, the strengthening effect of was proposed to the following mechanism:

1) Effective load transfer is the most important strengthening mechanism, due to Gr large specific surface area and high aspect ratio, and the load transfer efficiency depends on the interfacial bonding strength. In the present work, such interface was built by the formation of Cu-O-C bonding during electrodeposition, which not only strengthened the load transfer effect, but also fully utilized the intrinsic performance of Gr.

2) Dislocation strengthening effect. Many studies indicated that Gr played a barrier to block dislocation slip [32,45], which caused the high dislocation density at the interface.

3) Grain refinement effect. In our previous work, it was demonstrated that Gr provided more active sites for metal electrodeposition, which hindered the growth of metal crystal nucleus, and resulted in the refined grains [21,46].

However, for the Cu/Gr2/Cu foil with thick Gr interlayer, the tensile strength decreased obviously, as shown in Fig. 9(a), which could be attributed to the impeding of the Cu ions penetration through the thicker Gr layers. Therefore, the top Cu layer would quickly nucleate and grow under high overpotential and further hindered the Cu ions penetration. Alternatively, the GO film failed to form a strong interface bonding with Cu. The detailed mechanism was schematically illustrated in Fig. 10. Additionally, Fig. S5 also confirmed this mechanism.

Regarding electrical property of the Sandwich-type Cu/Gr/Cu layered composite foil, the resistance comes mainly from three factors, i.e., Cu, Gr and the Cu/Gr interface [47–49]. Obviously, the resistance of Cu is fixed, and therefore, the final variation determined by the other two factors, i.e., reduction degree of GO into Gr during electrodeposition and the Cu/Gr interfacial resistance. In general, the dense transition layer is in favor of higher electrical conductivity, while the loose one tends to hinder the electron conduction in composites [50,51]. Consequently, for the present samples, the thinner Gr interlayer was of a strong bonding interface between Cu and Gr, which would result in small resistivity, while the relatively thicker Gr interlayer had a higher resistivity due to the loose Gr structure. This phenomenon could also be interpreted diagrammatically from Fig. 10.

**5. Conclusions**

In order to challenge the mass production of ultrathin Cu foil in industry, a new electrochemical route is proposed to synthesize a kind of Sandwich-type Cu/Gr/Cu layer composite foil, regarding Gr’s strong mechanical and electrical properties.

1) The “one-step” preparation process is based upon two principles, i.e., the loose GO film from EPD method allows Cu ions penetration, and there are different Cu deposition potentials on different
substrates. Therefore, the bottom Cu layer forms during low potential step, while the top Cu layer forms during high potential step.

2) By adjusting the electrodeposition time and using thinner GO film, the thickness of the Cu/Gr/Cu foil can reaches to as thin as 4–5 μm, and comparing to pure Cu foil, it tensile strength increases to almost twice value, while the electrical conductivity is basically unchanged.

3) This process is of advantages, such as simple, controllable and possible large-scale production in industry. It is expected to be broadly

Fig. 8. Lateral view of fracture SEM morphologies of the foil fractural surfaces after tensile test. (a) and (b) pure Cu foil; (c) and (d) Cu/Gr1/Cu foil; (e) and (f) Cu/Gr2/Cu foil.

Fig. 9. Mechanical and electrical properties of the foils. (a) Stress-strain curves; (b) resistivity.
used in the areas, such as the Cu clad plate (CCL), printed circuit board (PCB), and lithium ion battery cathode collector system for saving raw material and also the space.

4) This work introduces a new idea for preparing the layered composites via electrochemical process, and it can be applied in other materials.

CRediT authorship contribution statement

Gongsheng Song: Conceptualization, Formal analysis, Investigation, Methodology, Writing - original draft. Qing Wang: Writing - review & editing. Li Sun: Visualization. Sishi Li: Data curation. Yafei Sun: Funding acquisition, Resources. Qiang Fu: Software. Chunxu Pan: Project administration, Supervision, Validation, Resources, Writing - review & editing.

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

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References


