

MoS₂ coated with Al₂O₃ for Ni–MoS₂/Al₂O₃ composite coatings by pulse electrodeposition[☆]

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Received 2 August 2007; accepted in revised form 26 November 2007

Available online 5 December 2007

Abstract

The MoS₂ powders were coated with Al₂O₃ (5 wt.%) through controlling hydrolysis of Al(NO₃)₃·9H₂O. MoS₂ powder coated with Al₂O₃ was written as MoS₂/Al₂O₃ hereinafter. MoS₂/Al₂O₃ powders were put into Ni plating electrolyte bath. Cetyltrimethylammonium bromide (CTAB) — the surfactant was also put into the bath. The experiment proves that MoS₂/Al₂O₃ particles were absorbed onto the Ni plate. The amount of MoS₂/Al₂O₃ deposited on Ni plate rises with the increasing concentration of MoS₂/Al₂O₃ in the bath. The microhardness, micro-surface, phase and the tribological property of the MoS₂/Al₂O₃ multi-plating coating were measured and analyzed. The performances of microhardness and wear resistance of the Ni–MoS₂/Al₂O₃ composite are better than those of Ni–MoS₂ composite.

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PACS: 81.15.P

Keywords: MoS₂ coated with Al₂O₃; Electrodeposition; Dendrite growth; Ni–MoS₂/Al₂O₃ coating; Tribological property

1. Introduction

The combination of different materials could produce the improved mechanical, chemical, electrical, magnetic and optical properties. However, these properties depend upon the distribution of phase and the composition of basic phase in the composite.

One type of composite in which ceramic particles were distributed in the metal matrix has been applied widely, especially in machinery for anti-wear and anti-friction. Various cationic surfactants had been added into the plating bath in order to increase the ceramic particles in the composite since the ceramic particles in a particle-dispersed metal matrix composite can share most of the load.

Electroplating has typically been used in the surface treatment industry [1]. It is reported that the composites with ultra-fine particles have been made successfully over the past decades. These composites are combined with metallic powders,

silicon carbide powders, oxide powders, diamond powders, polymers and the multi-plating of metal or alloy. The structures and properties of the mentioned composites have been studied and confirmed [2–4].

It is well known that pulse electroplating is more effective than the traditional DC electroplating to make composite and alloy due to its independent, controllable parameters and the high instant current [5]. The structure of materials or alloys can be changed through adjusting the pulse current and the properties of materials or alloys can be controlled or improved accordingly. Electroplating is one of the most important surfaces coating technology. When there are particles among the plating bath, the particles will be absorbed onto the plate. Such kind of coating plate is called composite plating. Many researchers have confirmed that the multi coating possess of many specific properties due to the different particles in the coating [6–12].

It has been reported that low-friction composite materials such as Ni-polyethylene [13], graphite-brass [14], Ni-PTFE [15], graphite-bronze [16], Ni-P-carbon-nanotube [17] and good wear resistance composite materials such as Ni–Al₂O₃ [18], Ni–P–SiC [19], Ni–ZrO₂ [20] can be obtained via the electroplating technique.

[☆] Foundation item: Project (K1703060819) supported by the National Defense Basic Research of China.

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Table 1
Bath composition and operating conditions

Composition and operating conditions	
Nickel sulfate	330 g/l
Nickel chloride	10 g/l
Boric acid	40 g/l
Cetyltrimethylammonium bromide (CTAB)	50 ppm/l
MoS ₂ or MoS ₂ /Al ₂ O ₃ respectively	0, 5, 15, 20, 25, 30 g/l
Temperature	60 °C
PH	4.5
Pulse current condition density	15 Adm ⁻²
Duty cycle	2/3
Frequency	50 Hz
Magnetic stirring speed	600 rpm

MoS₂ has the specific property of self-lubricate. Ni–MoS₂ composite plating is obtained by adding MoS₂ particles into the Ni bath. Such kind of composite plating is called functional plating in industry. Ni–Al₂O₃ composite plating can be used as anti-friction material due to the good harness performance of Al₂O₃. The composite plating will have self-lubricate and anti-friction properties if MoS₂/Al₂O₃ was absorbed onto the plate. In this experiment, we will show you the procedure of making Ni–MoS₂/Al₂O₃ composite plating and analyze its friction property.

2. Experiment

Commercial MoS₂ powder was used in this study. The MoS₂ powder was suspended in a mixed mineral acid (H₂SO₄+HCl),

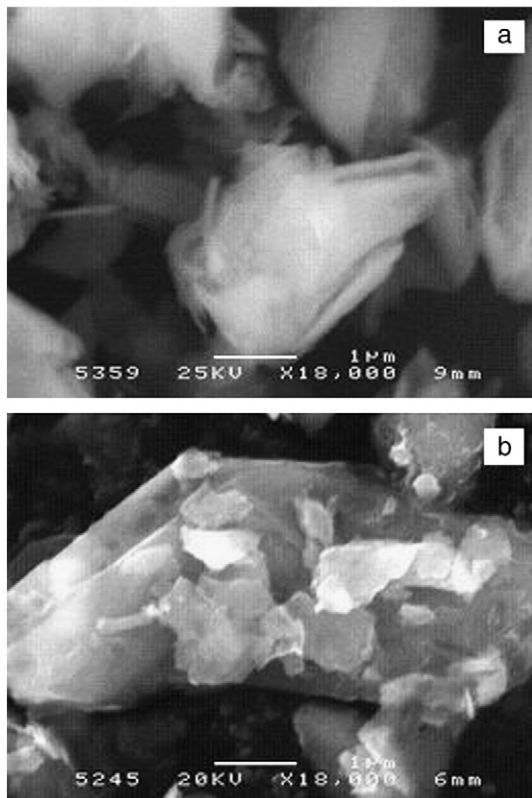


Fig. 1. SEM of powder: (a) pristine MoS₂ powder (b) 5 wt.% Al₂O₃ coated MoS₂.

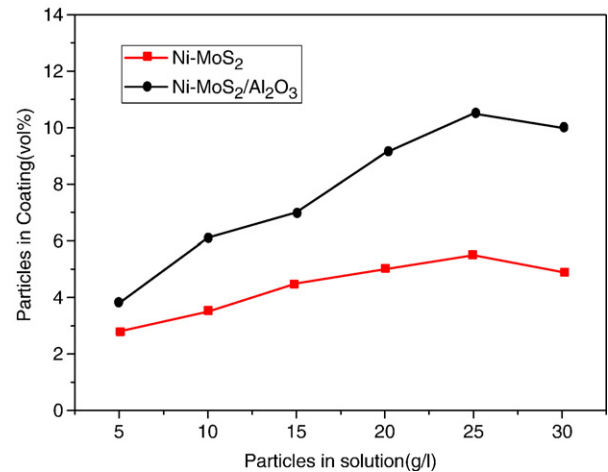


Fig. 2. Comparison of Ni–MoS₂ and Ni–MoS₂/Al₂O₃ coating.

and the acid solution was refluxed for 4 h. The MoS₂ powder treated with the mineral acid was repeatedly washed with water to remove the acid solution, and then the washed MoS₂ powder was dried in a drying oven. The MoS₂ powder 40 g, was suspended in 500 ml distilled water for experiments. Sodium acetate which was used to amortize pH of solution was added to the solution. And pH of the solution was maintained at 6. The beaker was set in a heating mantle on a magnetic stirrer to avoid settling of the MoS₂ powders during the process and subsequently by ultrasonic agitation for 20 min just prior to coating. The solution containing MoS₂ powder was heated at 80± °C for experiments. 32 g of Al (NO₃)₃ · 9H₂O was dissolved in 5 ml distilled water and added to the MoS₂ solution slowly. After completion of the experiment, the powder was collected by filtering and washed with hot distilled water repeatedly until the supernatant showed neutral pH. And then the powder was washed with ethanol and was collected by filtering. The MoS₂ powder treated with the Al(NO₃)₃ · 9H₂O reagent was heated at 300 °C for 3 h.

The MoS₂ and MoS₂/Al₂O₃ powders respectively, were suspended in the plating electrolyte bath, and CTAB, which is a surfactant. Surfactant CTAB was added to reduce the particles particle agglomeration [21]. Additionally, the adsorption of surfactant CTAB on the particles has been proven to enhance the adhesion force to the cathode, enabling larger particles to be embedded [21]. The plating electrolyte was a nickel sulfate bath. The compositions and the range of experimental operating conditions are shown in Table 1. A nickel plate of 120 mm×60 mm was used as the anode, and a low carbon steel disk of about half

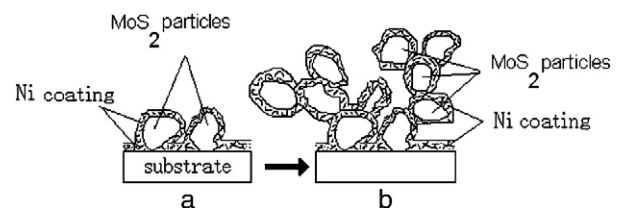


Fig. 3. Schematic of the diacritic growth of the Ni–MoS₂ coating. (a) MoS₂ was incorporated into the deposited Ni. (b) MoS₂ was adsorbed onto the protruding of the MoS₂ resulting in dendrite growth.

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