Fabrication and characterization of electroless Ni–P–ZrO₂ nano-composite coatings

Yongjian Yang · Weiwei Chen · Chungen Zhou · Huibin Xu · Wei Gao

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Abstract A novel technique has been developed to produce nano-particle oxide reinforced metal coatings. This method is based on electroless deposition process by adding ZrO₂ sol into conventional electroless Ni–P plating bath. Ni–P–ZrO₂ nano-composite coatings have been produced with highly dispersive ZrO₂ nano-particles inside the alloy coating matrix. The as plated nano-composite coating exhibits much increased microhardness up to 1045 HV₂₀₀ and remarkably improved wear resistance. X-ray and electron diffraction patterns show a phase transformation in the Ni matrix of the coating from amorphous to nanocrystalline when ZrO₂ sol is introduced into the coating. By comparison with the plain Ni–P coating and conventional Ni–P–ZrO₂ composite coating incorporated with solid ZrO₂ powders, two mechanisms for the increased mechanical properties are proposed based on nano-particle dispersion strengthening and phase transformation strengthening. The formation mechanism of ZrO₂ nano-particle is also discussed.

Keywords Electroless plating · Ni–P–ZrO₂ nano-composite coatings · ZrO₂ sol · Oxide nano-particle dispersion · Microhardness · Wear resistance

Introduction

Electroless plated nickel coatings (EP Ni–P coatings) have found broad applications due to their good properties and convenient process. The property improvement includes high hardness, good wear and corrosion resistance (Balaraju et al. 2003; Hari Krishnan et al. 2006; Peelers et al. 2001). For enhancing their performance, further development has been achieved mainly by two ways (Balaraju et al. 2003; Agarwala et al. 2006; Apachitei et al. 1998; Mallory and Hajdu 2004; Riedel 1991): incorporating hard or soft particles into the Ni–P matrix and heat treatment. In the past few years, various EP nickel-based composite coatings such as three-component Ni–P–Al₂O₃ (Alirezai et al. 2007), Ni–P–ZrO₂ (Song et al. 2007; Szczygiel et al. 2008), Ni–P–SiC (Apachitei and Duszczyk 2000; Huang et al. 2004), Ni–P–TiO₂ (Novakovic et al. 2006), Ni–P–diamond (Xu et al. 2005), and Ni–P–B₄C (Araghi and Paydar 2010) and four-component Ni–P–ZrO₂–Al₂O₃ (Sharma et al. 2002) and Ni–W–P–ZrO₂ (Szczygiel et al. 2008) have been explored by suspension of solid ceramic particles in the plating solutions. However, incorporating solid particles can only improve the properties (hardness and wear resistance) of the coatings to a limited level due to the agglomeration of nano- or micro-particles with high surface energy. This severely reduces the particle dispersion strengthening effect in the metal matrix according to the Orowan mechanism (Balaraju et al. 2006; Feng et al. 2007; Hou et al. 2006; Zhang and Chen 2006). On the other hand, although heat treatment can improve the hardness and wear resistance of EP Ni-P deposits to certain extent, it has limited applications to certain types coating systems. For instance, on substrates such as aluminum and magnesium alloys, it can cause undesirable deterioration of their mechanical properties (Apachitei and Duszczyk 2000).
Recently, a novel technique combining sol-gel and electroless plating processing has been developed by (Chen and Gao 2009; Chen et al. 2010) to synthesize EP Ni–P–TiO$_2$ nano-composite coatings with highly dispersed TiO$_2$ particles. Without any heat treatment, the novel Ni–P–TiO$_2$ nano-composite coatings obtained much improved micro-hardness and wear resistance. This technique utilizes sol solution containing the desirable components to form oxide nano-particles in the electroless plating solution, which can be co-deposited with Ni–P alloy to form nanostructured oxide dispersive composite coatings.

Besides TiO$_2$ sol, there are also other metallic oxide sols such as ZrO$_2$ and Al$_2$O$_3$ sols (Connor et al. 1995; Liang et al. 2009; Shukla et al. 2002) that can be used to make sol-enhanced composite coatings. This work explores the possibility of using ZrO$_2$ sol to synthesize EP Ni–P–ZrO$_2$ nano-composite coatings with improved mechanical properties. The formation mechanism of ZrO$_2$ nano particles is also discussed.

**Experimental**

The substrate material was the commercial AZ31 Mg alloy sheet with a composition (wt.%) of 2.33%Al, 1.27%Zn, 0.68%Mn, 0.68%Fe and balanced with Mg. Specimen (15 × 10 × 3 mm$^3$) pre-treatment is identical to that described in reference (Chen et al. 2010).

Transparent ZrO$_2$ sol was prepared by two steps detailed elsewhere (Liang et al. 2009). In the first step, 11.3 mL zirconium(IV) n-propoxide (70 wt.% solution in 1-propa-nol) was dissolved in a mixture containing 30.9 mL anhydrous ethanol and 2.8 mL diethanolamine (DEA). The solution was then kept stirring to achieve a complete chelation between Zr(OPr)$_3$ and DEA. In the second step, a mixture of 0.46 mL deionized water and 4.5 mL ethanol solution was mixed with the solution under vigorous stirring to form a fresh sol.

The bath composition and plating parameters in the present study is presented in Table 1. From the beginning of electroless deposition, ZrO$_2$ sol was simultaneously dripped into the plating bath with a rate of 0.3–0.4 mL/min. At the same time the solution was kept homogeneous by magnetic stirring at a speed of 200 rpm.

| Bath constituents and parameters | Quantity | Bath constituents and parameters | Quantity |
|---------------------------------|----------|---------------------------------|----------|
| NiSO$_4$·6H$_2$O                 | 15 g/L   | Thiourea                        | 1 mg/L   |
| Na$_2$H$_2$PO$_4$·2H$_2$O        | 14 g/L   | ZrO$_2$ sol                     | 120 mL/L |
| NaCH$_3$COO                     | 13 g/L   | Temperature                     | 80 ± 2°C |
| HF (40%)                        | 12 mL/L  | pH value                        | 6.4 ± 0.2|
| NH$_4$HF$_2$                     | 8 g/L    | Time                            | 90 min   |

For comparison purposes, plain EP Ni–P coatings and solid powder mixing enhanced “conventional” EP Ni–P–ZrO$_2$ composite coatings were also prepared with the identical bath composition and plating parameters. In the case of conventional electroless Ni–P–ZrO$_2$ composite deposition, commercial ZrO$_2$ powder (Aldrich Chemical Company, Inc.) with a grain size smaller than 5 μm was well dispersed (5 g/L) into the plating bath by stirring before deposition. The content of ZrO$_2$ particles in the composite coatings was obtained in the same way detailed elsewhere (Chen et al. 2010) by the following equation:

\[ C = \frac{[\text{Weight(ZrO}_2)/\text{Weight(Coating)}] \times 100\%}{1} \quad (1) \]

Since ZrO$_2$ particles do not react with HNO$_3$, they were extracted by dissolving the metal matrix into HNO$_3$ solution. An electronic balance with an accuracy of 0.01 mg was used to measure the quantity of oxide.

The surface and cross-section of the coating were investigated with Philips XL30S FSEM. To determine the coating composition, EDX analysis was performed with every specimen. X-ray diffractometer (Bruker D8) with Cu-K$_\alpha$ was used to study the phase structure of the coatings. Crystal size was determined using the Debye–Scherrer relation. The morphologies and distribution of the ZrO$_2$ particles in the composite coatings were also characterized by JEOL JEM-2100F and FEI Tecnai 12 transmission electron microscopes (TEM). Microhardness of the coatings was determined on a Leco M400 microhardness tester with a Vickers diamond indenter under a 200 g load for a holding time of 15 s. Eight measurements were conducted for each sample and the average was used as the hardness value. Wear tests were conducted on a tribometer (Nanovea TRB) with a friction counterpart of a ruby ball of 6 mm in diameter. A load of 7 N and a sliding speed of 50 mm/s were used without lubrication at ambient temperature. The length of the wear track was 10 mm long and the total elapsed time was 100 min.

**Results and discussion**

Surface and cross-sectional morfologies of coatings

Figure 1 shows the surface morphologies of the plain Ni–P, conventional Ni–P–ZrO$_2$ composite and sol enhanced...
novel Ni–P–ZrO\textsubscript{2} composite coatings. Typical spherical nodular structures can be seen on all the three kinds of electroless nickel coatings whereas the two composite coatings have rougher surfaces in comparison with the plain Ni–P. In the inset of Fig. 1b, microsized ZrO\textsubscript{2} particles can be seen clearly on the conventional Ni–P–ZrO\textsubscript{2} composite coating. For the sol processed Ni–P–ZrO\textsubscript{2} composite coating, the inset of Fig. 1c shows a granular morphology throughout the surface, which is also distinct from the smooth surface morphology of Ni–P–TiO\textsubscript{2} composite coatings by others (Chen et al. 2010).

Figure 2 shows the cross-sectional SEM images and qualitative elemental distribution across the plain Ni–P, conventional Ni–P–ZrO\textsubscript{2} composite and novel Ni–P–ZrO\textsubscript{2} composite coatings. All three coatings are compact with uniform thickness of \( \sim 20 \mu m \) regardless of any surface irregularity, giving a plating rate in the range of 12–14 \( \mu m/h \). For the conventional Ni–P–ZrO\textsubscript{2} composite coatings, clusters of ZrO\textsubscript{2} particles sized from 200 to 600 nm can be seen in the inset of Fig. 2b1, with a high content of 24.4 \( \pm 0.8 \) wt%. By contrast, the sol enhanced novel Ni–P–ZrO\textsubscript{2} composite coating exhibits a homogeneous structure without any visible ZrO\textsubscript{2} particles (inset of Fig. 2c1), probably owing to their extremely small size and relatively low content of 4.2 \( \pm 0.5 \) wt%. The elemental distributions of the three coatings in Fig. 2 show a less quantity but more uniform distribution of Zr in novel Ni–P–ZrO\textsubscript{2} composite coating than in the conventional Ni–P–ZrO\textsubscript{2} composite coating, corresponding to the more dispersive ZrO\textsubscript{2} small particles in the novel Ni–P–ZrO\textsubscript{2} composite coating compared to the micron sized ZrO\textsubscript{2} particles in the conventional Ni–P–ZrO\textsubscript{2} composite coating. At the same time, the lower content of P (\( \sim 1.34 \) wt%) in the novel Ni–P–ZrO\textsubscript{2} composite coating in Fig. 2c2 compared to the other two coatings shown in Fig. 2a2, b2 can be one of the reasons for phase structure transformation in the novel Ni–P–ZrO\textsubscript{2} composite coating which will be detailed in the ensuing section.

TEM observations of coating microstructure and particle morphology

TEM bright field micrographs of the three electroless coatings are shown in Fig. 3. The plain Ni–P coating (Fig. 3a) shows a homogeneous contrast with mixed amorphous and crystalline structures, evidenced by the diffuse rings in the corresponding electron diffraction pattern. For the conventional Ni–P–ZrO\textsubscript{2} composite coating, because the ZrO\textsubscript{2} micron particles are large and have already been confirmed in the above cross-sectional SEM observation, the micrograph (Fig. 3b) only presents the Ni–P matrix structure. As illustrated in the diffraction pattern (inset of Fig. 3b), the microstructure of the conventional Ni–P–ZrO\textsubscript{2} composite coating contains more crystal portion than amorphous phase with sharper rings compared to the plain Ni–P coating. In contrast to both Ni–P and conventional Ni–P–ZrO\textsubscript{2} composite coatings, the novel Ni–P–ZrO\textsubscript{2} composite coating shows a crystal structure containing highly dispersive ZrO\textsubscript{2} nano-particles with the size less than 20 nm as pointed by the arrows (Fig. 3c).

Figure 3d is the TEM micrograph and diffraction pattern of the ZrO\textsubscript{2} particles extracted from the novel Ni–P–ZrO\textsubscript{2} coating.
Fig. 2 Cross sections and line scanning analyses results of electroless nickel coatings. a1, a2 Plain Ni–P coating, b1, b2 conventional Ni–P–ZrO$_2$ composite coating, c1, c2 novel Ni–P–ZrO$_2$ nano-composite coating
composite coatings. The apparent particle agglomeration shown in Fig. 3d is likely to form after extraction as no such big particles were found in the coatings under SEM and TEM examinations. However, the small particles as arrowed in Fig. 3d represent their original size in the alloy matrix before agglomeration. On the other hand, indication from the diffraction pattern in Fig. 3d does not show a clear crystal structure, which is consistent with the results by electron diffraction pattern of the novel Ni–P–ZrO₂ composite coating, showing no characteristic rings from crystal ZrO₂ particles.

Phase analysis of coatings

Figure 4 is the XRD spectra showing significant differences in phase structure between Ni–P, conventional Ni–P–ZrO₂ composite and the novel Ni–P–ZrO₂ nano-composite coatings. As indicated in the electron diffraction patterns in the former section, both Ni–P and conventional composite coatings are composed of amorphous and crystal phases while the novel Ni–P–ZrO₂ coating is seen fully crystallized with strong diffraction peaks. Since the well-known amorphous structure of the electroless Ni–P coating is attributed to the distortion of crystal lattice of nickel by the phosphorus atoms (Agarwala et al. 2006; Mallory and Hajdu 2004; Sharma et al. 2002), and previous results (“Surface and cross-sectional morphologies of coatings”) indicated the lower P content for the novel Ni–P–ZrO₂ coating in contrast with the others, it can be concluded that...
the addition of ZrO$_2$ sol into the electroless Ni–P plating solution decreases the incorporation of P into the coating.

For the conventional Ni–P–ZrO$_2$ coating, the diffraction peaks from ZrO$_2$ powders are clearly shown in Fig. 3, consistent with the results from cross-sectional observation which elucidate the existence of crystal ZrO$_2$ particles in the coating. By contrast, the XRD pattern for novel Ni–P–ZrO$_2$ nano-composite coating shows no peaks from ZrO$_2$ particles, again possibly due to the low quantity of the particles (~4.2 wt%) and their amorphous state. Further calculation of grain size by Scherrer method revealed that the crystal sizes for the novel Ni–P–ZrO$_2$ composite coatings are in the level of ~64 nm, confirming the nano-crystalline structure of the matrix of novel Ni–P–ZrO$_2$ composite coatings.

**Mechanical properties of coatings**

Microhardness tests were performed on the plain Ni–P, conventional Ni–P–ZrO$_2$ and novel Ni–P–ZrO$_2$ coatings as listed in Table 2. By adding ZrO$_2$ sol into the electroless Ni–P plating solution to form an optimal nano-composite structure, the microhardness of the novel Ni–P–ZrO$_2$ composite coating has been increased to a much higher level of 1,045 ± 30 HV$_{200}$ compared to the Ni–P coating of 619 ± 20 HV$_{200}$ and conventional Ni–P–ZrO$_2$ composite coating of 759 ± 25 HV$_{200}$.

Figure 5 shows the wear tracks and Table 2 listed the wear volume losses for the three types of coatings. The narrower wear track and shallower plough lines show much improved wear resistance of the novel Ni–P–ZrO$_2$ composite coatings compared to the plain Ni–P and conventional Ni–P–ZrO$_2$ composite coatings. While the improved wear resistance of the conventional Ni–P–ZrO$_2$ composite coating compared to that of plain Ni–P coating can be attributed to the dispersion strengthening and load support function of ZrO$_2$ particles in the coating (Song et al. 2007), the reasons for the higher microhardness and improved wear resistance of the novel Ni–P–ZrO$_2$ nano-composite coating possibly come from two perspectives. On one hand, the highly dispersed ZrO$_2$ nano-particles (<20 nm) can strengthen the Ni–P matrix more effectively than the coarse ZrO$_2$ particles (200–400 nm) based upon the Orowan mechanism (Hou et al. 2006; Zhang and Chen 2006; Dieter 1976); on the other hand, according to many researchers’ results (Apachitei and Duszczyk 2000; Jeong et al. 2002; Van Swygenhoven and Caro 1997), a nano-crystalline structure of the Ni–P-matrix is possessing a higher hardness than an amorphous structure, for which most of explanations are based on a transition from dislocation-controlled deformation to other deformation mechanisms. As illustrated in the XRD and TEM diffraction results, the novel Ni–P–ZrO$_2$ composite coating show a better crystallized structure than the plain Ni–P and conventional Ni–P–ZrO$_2$ composite coatings, resulting in a stronger...
matrix which contributes to the higher hardness and therefore better wear resistance of the nano-composite coatings.

ZrO₂ nano-particle formation mechanism

The formation of the nano-particles during the deposition process can be complicated. A suggested mechanism is presented below. The sol-synthesis method in this study follows the chelating sol-gel route, which produces sols directly based on the hydrolysis-condensation of alkoxides with water (Liang et al. 2009; Qiu et al. 2007). Details about formation mechanism of the metallic oxides particles from the sols have been described in others’ works (Gopal et al. 1997; Oleshko et al. 2003; Shukla and Seal 2003; Tang et al. 2003). According to these studies, in the preparation of ZrO₂ sol utilizing neutral ethanol as solvent, condensation process of organic metal macromolecule ions started before completion of hydrolysis, and the formation of ordered structure was hindered, hence the as-synthesized ZrO₂ nano-particles were found to be amorphous as shown in TEM results. The reaction during the preparation can be described as follows:

Hydrolysis:

\[
\text{Zr(OC}_3\text{H}_7)_4 + 4\text{H}_2\text{O} \rightarrow \text{Zr(OH)}_4 + 4\text{C}_3\text{H}_4\text{OH}
\]  

Condensation:

\[
\text{Zr(OH)}_4 \rightarrow \text{ZrO}_2 + 2\text{H}_2\text{O}
\]  

Net reaction:

\[
\text{Zr(OC}_3\text{H}_7)_4 + 2\text{H}_2\text{O} \rightarrow \text{ZrO}_2 + 4\text{C}_3\text{H}_4\text{OH}
\]

On the other hand, because the nano-particles in the sol were added dropwise into the plating solution at a controlled speed, it is believed that the particles were well dispersed and some of them were co-deposited into the Ni–P matrix to form the EP nano-composite coatings. This is further supported by TEM observation of the EP Ni–P–ZrO₂ nano-composite coatings, in which particles are seen to uniformly distribute in the Ni–P matrix, with the size smaller than 20 nm.

Conclusion

EP Ni–P–ZrO₂ nano-composite coating has been successfully synthesized via a new process which utilizes metal oxide sol to introduce highly dispersive particles to strengthen the coating. The microhardness for the as-deposited Ni–P–ZrO₂ nano-composite coating was improved to ~1,045 HV₂₀₀ compared to 619 and 759 of the plain Ni–P coating and Ni–P–ZrO₂ coating made by solid particle mixing. Consequently the coating obtains significantly improved wear resistance. The strengthening effects were discussed based on the Orowan dispersion mechanism and properties of different phase structures. The ZrO₂ particle formation mechanism was also discussed in terms of the hydrolysis–condensation reaction. All results from the present study have proved that this new method is an effective way to produce nano-composite coatings with high hardness and wear resistance for promising industrial applications.

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