FABRICATION AND CHARACTERIZATION OF ELECTRODEPOSITED AND MAGNETRON-SPUTTERED THIN FILMS

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ABSTRACT
The MnO–Zn thin films were fabricated by radio frequency (RF) magnetron sputtering and compared with pulse electrodeposition (PED) Zn thin films, doped with MnO and ZrO nanoparticles. Surface morphology, structural properties, chemical composition and corrosion resistance of these coatings were investigated by using scanning electron microscopy, X-ray diffraction (XRD), energy-dispersive X-ray spectroscopy, 3-D scanning interferometry and environmental chamber. Surface morphology and degree of crystallinity have different behaviours for different deposition methods. Pulse-coated films have polycrystalline structure with high surface roughness (\(R_a\)), whereas sputtered films are monocristalline with reduced roughness (\(R_a\)). Corrosion tests of both RF sputter and PED films revealed that the distribution of corrosion products formed on the surface of sputter films were not severe in extent as in case of electrodeposited coatings. Results showed that the doping of ZrO nano-sized particles in Zn matrix and Mn–Zn composite films significantly improved the corrosion resistance of PED thin films.

Keywords: Corrosion resistance, electro-deposition, magnetron sputtering, surface analysis, thin films.

1 INTRODUCTION
Zinc oxide (ZnO) thin films are widely used in variety of applications such as thin-film transistor [1], transparent conducting films, solar cell material [2], UV photodetector [3], piezoelectric [4] and wear-resistant films [5], due to its several favorable properties, including wide bandgap, e.g. –3.37 eV [6], large exciton binding energy (60 meV) [6] and also having good transparency, which are excellent parameters for optoelectronic devices.

High electron mobility [7] and room temperature electrical conductivity of ZnO thin films make it perfect material for electronics equipment. Additional advantages of ZnO thin films and micro-nanostructure are abundance and non-toxicity of the ZnO material, low cast and quantum size effect. Depending on the application, tuning properties and band gap of ZnO is possible by doping another material such as: aluminum (Al), gallium (Ga), magnesium (Mg) and manganese (Mn).

Also, there are advanced techniques of growing ZnO thin films and manufacturing such as metal organic chemical vapor deposition [8], radio frequency (RF) magnetron sputtering [9], sol–gel method and pulsed laser deposition [10]. Properties of ZnO thin films are investigated by using molecular beam epitaxy [11].

Zinc plating is extensively used in corrosion protection of steel in many structural and general engineering applications. Zn coating on steel substrate provides with physical and mechanical properties as well as good corrosion resistance [12]. However, a high dissolution rate and low corrosion resistance limit the use of Zn coatings. In recent research and development activity on zinc-based alloy coatings, there is a growing interest in the use of Zn with various alloying elements like Mn and Zr. These elements (Mn and Zr) have an electrically, more negative potential (\(E_{\text{Mn}}^{2+}/\text{Mn} = −1.185\) V vs. SHE (standard hydrogen electrode) and \(E_{\text{Zr}}^{2+}/\text{Zr} = −1.45\) V vs. SHE) when compared with Zn [13]. The alloys Zn–Mn, Zn–Zr and
Zn–Zr–Mn show an interdependent effect with better corrosion resistance than the individual metals. The need for coatings with improved resistance to highly aggressive environments is high as a result of a growing demand for extended safe service life of industrial objects. Conventional coatings such as spray gun coatings perform well in normal environmental conditions; however, when subjected to aggressive environments result in coating failure due to blistering and delamination. Sputtered and electrodeposited coatings can perform well in aggressive environments and are more resistant to failure due to blistering delamination.

This work is the continuation of current research in materials, structural integrity and coatings [14–21]. This paper presents results, which show that how materials’ qualities have been affected by deploying different methods of thin film fabrication. Pulse electrodeposition process is simple with low cost, and deposition can be applied on comparatively larger areas [22]. The RF-magnetron sputtering is a flexible technique in terms of control of composition and microstructure. Zn–MnO thin films can be fabricated by reactive RF magnetron sputtering deposition with high-purity ZnO powder, which results in the formation of films with excellent purity. On the other hand, a vacuum method is a costly procedure and some of scaling process is difficult. Pulse electrodeposition has been widely applied to fabrication of thin films, and results show that the uniformities such as thickness of layer, grain size, roughness and defect density are difficult to control [23].

It is obvious that the properties of ZnO thin films are extremely sensitive to the sample preparation process, and the focus of this research is to compare chemical and physical deposition techniques. Investigate surface morphology, elemental composition and durability of two different methods of ZnMnO thin films fabrication. Additionally, the effect of incorporated MnO and ZrO nanoparticles on corrosion resistance is also investigated for coatings prepared by two different methods.

2 EXPERIMENTAL

2.1 Preparation of sputtered and electrodeposited coatings

In this work, MnO–Zn thin film was deposited on glass substrate by RF magnetron sputtering method. The required thin films have been achieved by using a mixture of high-purity ZnO with 36 wt. % MnO₂. Substrate was conditioned carefully by using ethanol and dried by subjecting the samples to flowing N₂ gas. The pressure of the chamber was kept as 2.9 × 10⁻⁸ mbar. High-purity (99.999 %) Ar (Argon) gas was introduced into the chamber with a total flow rate fixed at 18 sccm. The ZnMnO target was sputter cleaned for 20 min in Ar before allowing any deposition to take place on the substrate. The substrate temperature, sputter pressure, RF power and the distance between target and substrate was set at 500°C, 5.4 × 10⁻³ mbar, 250 W and 35 mm, respectively.

In a second technique, MnO–Zn, ZrO–Zn and ZrO–MnO–Zn coatings were deposited over mild steel (50 mm × 25 mm × 1.4 mm) substrate using pulse electrodeposition method. Zinc sheets (90 mm × 75 mm × 0.8 mm) were used as the cathode.

Substrates were polished with successive finer grade of emery papers of grades: 320, 600 and 1200 followed by burnishing with a very soft cloth. The samples were then washed with tap water, cleaned in acetone using an ultrasonic bath and finally dried with air blow. Electrodeposition was then performed at 30°C temperature using an acidic chloride (pH 4.3).

For the composite coatings, 10 g/L manganese and zirconium oxide particles with diameter about 40–60 nm was added into the bath. To better suspension of nanoparticles, the bath
was stirred about 12 h by using a magnetic stirrer pre electrodeposition process. The concentration of various bath constituents in the electrolytic bath is presented in Table 1.

2.2 Sample pretreatment and corrosion tests

The experiments were performed through accelerated corrosion tests by deploying a temperature-climate simulation chamber. The chamber was heated to 100–105°C. The coated samples were placed in a beaker, with a capacity of 200 ml, of 3.5 wt. % filtered sodium chloride (NaCl) solution of which 3.5g of NaCl was mixed in 100 ml of deionized water.

Corrosion tests were performed in 3.5 wt. % NaCl solution, which is appropriate for simulating corrosion due to the presence of corrosion activators (chloride ions). The beaker contained coated samples, and NaCl solution was placed in a heated chamber for 150 h. The coated samples were continuously monitored, and progress was recorded after every 24 h.

2.3 Characterization of sputtered and electrodeposited coatings

Structural properties of all coatings were investigated through HRXRD. The grain size of the films were estimated with Scherer’s formula [24]

\[ D = \frac{k\lambda}{B\cos \theta} \]

where D is the grain size, k is the constant (=0.9), \( \lambda \) is the wavelength of X-ray (=1.540598 Å), B is the line width (FWHM) and \( \theta \) is the Bragg angle.

Coating surface roughness values were analysed and averaged through five data points per coated sample using 3-D surface scanning interferometry. The microstructure analyses and measuring of incorporated nanoparticles weight percentage of coatings were conducted through scanning electron microscopy and energy dispersive X-ray spectroscopy (EDS), respectively.

3 RESULTS AND DISCUSSION

3.1 Coating surface morphology and structure properties

Figure 1 compares the surface morphologies of thin films manufactured by two different methods. Figure 1a shows that the film produced by sputtering method has high density and
small value in average surface roughness ($R_a \sim 0.017 \, \mu m$) determined by 3D Interferometer. The surface morphology of electrodeposited films can be seen in Fig. 1b–d, which shows that the coating morphology consists of hexagonal platelet morphology varied from mainly granular shape to least pyramidal shapes. Similar morphology of electrodeposited Mn–Zn alloys was also reported by other researchers [13, 25, 26].

Structural properties of both types of coatings are shown in Fig. 2. It can be seen that the sputter-deposited samples have a preferred (002) orientation, whereas pulse-coated films including MnO–Zn, ZrO–Zn and ZrO–MnO–Zn have a polycrystalline structure. The varied crystallographic orientations in electrodeposited coatings cause enhanced porosity and consequently increased average surface roughness of $R_a \sim 0.39 \, \mu m$.

The reinforcement content of nanoparticles in coatings calculated by using EDS analysis is presented in Table 2. Grain sizes in electrodeposited films are significantly small compared with those sputtered coatings. However, the porous structure of pulsed coatings is a disadvantage in terms of corrosion resistance property compared with dense sputtered films.

3.2 Experimental set-up and corrosion study

These experiments are the part of research being carried out in the area of corrosion within our research group [21, 27–34]. The details for the corrosion tests and sample preparation have been discussed in Section 2.2.
The corrosion product (rust) forms on the surface of metals as a result of electrochemical reactions. The atoms on metal surface undergo changes by losing electrons and become positively charged ions. This allows them to bond to other groups of atoms that are negatively charged resulting in oxidation at anodic sites.

\[ \text{Metal} \rightarrow \text{Metal}^{n+} + n \text{electrons} \]

The electrons flow from the anodic sites to cathodic sites, i.e. the electrons migrate through a metallic path. This migration occurs due to a voltage difference between the anodic and cathodic reactions. The excess of dissolved oxygen in water results in cathodic reaction and forms cathodic sites on the metal surface

\[ \text{O}_2 + 2\text{H}_2\text{O} + 4\text{e}^- \rightarrow 4\text{OH}^- \]

The anodic and cathodic reactions result in the formation of corrosion products (rust) on the metal surface. The metal ions (Metal\(^{n+}\)) coming from the reaction of metal at anodic site will combine with hydroxyl ions (OH\(^-\)) producing metal hydroxide.

\[ \text{Metal}^{n+} + 2\text{OH}^- \rightarrow \text{metal hydroxide} \]
Metal hydroxide is further oxidized to form hydrated metal oxide, which is dark brown due to iron.

3.3 Pre- and post-corrosion observations

The possibility of occurring sputter damage is known but insufficiently investigated for corrosion products of Zn-based steel coatings such as Zn–Mn. Post-corrosion test observations show that a compound that is formed when Zn and Zn-based protection layers are exposed to a corrosive environment is ZnO as shown in Fig. 3. Microscopic surface analysis and comparison with pulse electrodeposited coating revealed that the distribution of corrosion products formed on the surface of sputter coating was not severe in extent as in case of electrodeposited coatings.

Table 2 shows the front face and 3-D surface images of Zn–Mn, Zn–Zr and Zn–Zr–Mn coatings deposited on steel substrate. Post-corrosion test observations show that the surface morphology of the electrodeposits varied from a densely red-brown flaky rust (Zn–Mn) to a fine white/grey (Zn–Zr–Mn) unstable rust, depending upon the type of alloy composition. These micrographs indicate that incorporation of a small quantity of Mn and Zr into Zn matrices significantly alters the corrosion morphology of the Zn–Mn and Zn–Zr–Mn alloys.

3D scanning interferometry technique was deployed to evaluate the extent of corrosion of Zn–Mn and Zn–Zr–Mn alloys using average surface roughness ($R_a$) and peak surface height (PSH) as two measurement parameters. The results were analysed and compared in order to identify the coating with best corrosion resistivity.

Comparison of pre- and post-corrosion observations of Zn–Mn sample revealed that the PSH, after the development of rust changed from 2.94 to 109.73 µm. Similarly, the average surface roughness ($R_a$) changed from 0.57 to 2.79 µm. Similarly, for Zn–Zr sample, the PSH, after the development of rust changed from 3.26 to 17.54 µm and the average surface roughness ($R_a$) changed from 0.4 to 1.75 µm. Moreover, for Zn–Zr–Mn sample, the PSH changed from 4.31 to 9.29 µm and the average surface roughness ($R_a$) changed from 0.64 to 1.45 µm.

The pre- and post-corrosion results for all the samples showed that PSH and $R_a$ for each

![Figure 3: Samples prepared using RF sputtering technique with Zn–Mn sputtered coating: (a) pre-test surface and (b) post-test surface.](image-url)
Table 3: Corrosion test results (pre and post) and their comparison.

<table>
<thead>
<tr>
<th>Sample type</th>
<th>Analysis type</th>
<th>Pre-corrosion test</th>
<th>Post-corrosion test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zn–Mn</td>
<td>Front face image</td>
<td><img src="Mn-Zn.png" alt="Image" /></td>
<td><img src="Mn-Zn.png" alt="Image" /></td>
</tr>
<tr>
<td></td>
<td>3D surface scanning</td>
<td>$R_s = 0.57 \mu m$</td>
<td>$R_s = 2.79 \mu m$</td>
</tr>
<tr>
<td></td>
<td></td>
<td>$+2.42 \mu m$</td>
<td>$R_s = 109.73 \mu m$</td>
</tr>
<tr>
<td></td>
<td></td>
<td>$-8.10 \mu m$</td>
<td>$-5.58 \mu m$</td>
</tr>
<tr>
<td>Zn–Zr</td>
<td>Front face image</td>
<td><img src="Zr-Zn.png" alt="Image" /></td>
<td><img src="Zr-Zn.png" alt="Image" /></td>
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<tr>
<td></td>
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<td>$R_s = 1.75 \mu m$</td>
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<tr>
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<td></td>
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<td>$+17.54 \mu m$</td>
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<td></td>
<td></td>
<td>$-4.88 \mu m$</td>
<td>$-7.39 \mu m$</td>
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<tr>
<td>Zn–Zr–Mn</td>
<td>Front face image</td>
<td><img src="Mn-Zr-Zn.png" alt="Image" /></td>
<td><img src="Mn-Zr-Zn.png" alt="Image" /></td>
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<tr>
<td></td>
<td>3D surface scanning</td>
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<td>$R_s = 1.45 \mu m$</td>
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<td>$+9.29 \mu m$</td>
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<td>$-9.87 \mu m$</td>
<td>$-8.63 \mu m$</td>
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</table>
sample increased due to the development of rust, after they have been treated with NaCl solution. However, the highest % increase in PSH and $R_a$ was found for Zn–Mn while lowest was found for Zn–Mn–Zr.

These results indicate that Mn and Zr, both combined (Zn–Zr–Mn), are the most beneficial composition, compared with individual additions of Mn and Zr into Zn matrices. Both Mn and Zn combined can significantly improve the resistance of the coatings to atmospheric corrosion.

4 CONCLUSIONS

In conclusion, ZnMnO thin films can be manufactured successfully by chemical and physical methods. Structural and morphological properties of the two methods (sputtering and electrodeposited) are investigated. The coating produced by sputtering method has high density and small value in average surface roughness. However, the surface morphology of electrodeposited coating consists of hexagonal platelet morphology varying from mainly granular shape to least pyramidal shapes. The varied crystallographic orientations in electrodeposited coatings cause enhanced porosity and consequently increase the average surface roughness. The corrosion tests were performed by exposing the samples (sputtered and electrodeposited) to salt solution. The purpose of the exposure was to analyse the corrosion resistance of all the samples while keeping the coating thickness constant for all three samples. The microscopic surface analysis and comparison of sputtered and pulse electrodeposited coatings revealed that the distribution of corrosion products formed on the surface of sputtered coating were not severe in extent as in case of electrodeposited coatings. For the case of electrodeposited coatings, the corrosion test results indicate that Mn and Zr, both combined (Zn–Zr–Mn), are the most beneficial composition, compared with individual additions of Mn and Zr into Zn matrices. Both Mn and Zn combined can significantly improve the resistance of the coatings to atmospheric corrosion. Also, the sputter coatings are more resistant to corrosion failures compared with electrodeposited coatings.

REFERENCES


