Morphology of alpha-lead dioxide electrodeposited on aluminum substrate electrode

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Abstract: Alpha-lead dioxide was deposited by anodization of alkaline solution containing HPbO\textsuperscript{2−} anions. Scanning electron microscopy (SEM) results show that the morphology is remarkably affected by the current density, concentration of HPbO\textsuperscript{2−} anions, bath temperature and electroplating time. Compact and well adherent layers are possibly obtained under conditions of current densities $\leq 3$ mA/cm\textsuperscript{2}, electrolyte containing 4 mol/L NaOH and 0.12$-0.14$ mol/L lead (II), bath temperature of 40 ℃, and electroplating time of 2 h. EDS analyses show that the PbO\textsubscript{2} deposited in alkaline condition is highly non-stoichiometric at high current density.

Key words: alpha-lead dioxide; morphology; electrodeposition; composition; aluminum substrate

1 Introduction

Lead dioxides have been used frequently in industry because of their excellent properties such as good conductivity, low cost, high stability and relatively high service life. A great number of applications have been reported for lead dioxide as positive active material in lead acid batteries[1], oxidation of organic compounds in waste water[2–4], oxidation of glucose[5], ozone evolution[6–7], oxidation of phenol[8–9] and Cr\textsuperscript{3+}[10].

The lead dioxide from electrodepositing is known to exist in two polymorphs: orthorhombic $\alpha$-lead dioxide and tetragonal $\beta$-lead dioxide[11]. It is well known that the crystal structure of PbO\textsubscript{2} deposits depends on the pH of the electroplating solution. Alpha-PbO\textsubscript{2} is obtained from bases and $\beta$-PbO\textsubscript{2} from acids[11]. Alpha-PbO\textsubscript{2}, which results in better contact between particles, has a more compact structure than $\beta$-PbO\textsubscript{2}. The more compact structure makes $\alpha$-PbO\textsubscript{2} more difficultly discharged compared with $\beta$-PbO\textsubscript{2}[12]. However, results from the work by RÜETSCHI[13] and FENG and JOHNSON[14] showed that $\alpha$-PbO\textsubscript{2} had a higher catalytic activity than $\beta$-PbO\textsubscript{2} in dilute H\textsubscript{2}SO\textsubscript{4} solution.

A new type of PbO\textsubscript{2}-coated metal anode has been widely used in electrolysis[10, 15]. This electrode is made up of four layers: the base is made of titanium plate, and covered with a conductive undercoating (being necessary for protecting the substrate from passivation) as bottom; the intermediate coating is composed of $\alpha$-PbO\textsubscript{2} and plated with $\beta$-PbO\textsubscript{2} as surface layer. Titanium is not a viable substrate for practical electrodes in electrodepositing of nonferrous metals. Aluminum is relatively cheap and has good conductivity. The electrode material by electrodepositing lead dioxide on Al substrate has huge market prospects. A stress-free intermediate $\alpha$-PbO\textsubscript{2} coating is produced by electrodeposition from an alkaline lead bath[16], which plays the role of binder on the top $\beta$-PbO\textsubscript{2} coating and can improve the servo life of electrode. However, the $\alpha$-PbO\textsubscript{2} films deposited from basic solution were highly porous[17–18] and the research work was confined to the internal stress measurement[16].

Most of the previous studies on the morphology has concerned $\beta$-polymorph[2, 19–20]. However, there is little research about the morphology of $\alpha$-PbO\textsubscript{2} coating, and the structure of the intermediate coating is likely to be important for the lead-acid battery and as an electro-

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catalytic material for oxygen transfer reactions. In this work, $\alpha$-PbO$_2$ films with different morphologies electrodeposited on aluminum substrate in alkaline bath were discussed. An analysis of cyclic voltammetry experiments with $\alpha$-PbO$_2$ electrodes in alkaline and acid bath was also given, respectively.

2 Experimental

2.1 Preparation of PbO$_2$/Al anode

The PbO$_2$/Al anode was produced by applying $\alpha$-PbO$_2$ to an Al substrate with a conductive undercoating, followed by an intermediate coating consisting of $\alpha$-PbO$_2$ deposit.

2.2 Application of coatings

The substrates were aluminum (or aluminum alloy) rods of $d$ = 9 mm $\times$ 60 mm, which were roughened by sand-blasting, degreased and chemically etched firstly, then coated by a conductive coating. The procedure was described as follows: firstly, the conductive solution was applied to the substrate by brushing; secondly, the substrate was surface dried under ultraviolet lamp, and finally dried in electricity box at 423 K for 2 h. The undercoating produced in this study was about 20−30 µm thick, and the details were presented in Ref.[21]. Thereafter, the composition and process conditions of the PbO$_2$ plating bath were shown as follows: 4 mol/L NaOH with litharge PbO(s) (the soluble Pb(II) species were HPbO$_2^-$ anions), pH > 14, anode current density of 1−5 mA/cm$^2$, mild stirring using a magnetic stirrer, bath temperature of 25−55 $^\circ$C, and electroplating time of 1−3 h.

2.3 Characterization of PbO$_2$/Al electrode

The electrochemical performance was measured by the three-electrode cell: Al/conductive coating/PbO$_2$ anode as the working electrode, Hg/Hg$_2$Cl$_2$ (KC1, saturated) as the reference electrode, and a graphite as the auxiliary electrode. Cyclic voltammetric curves were measured in an alkaline and acid lead bath, respectively. The alkaline bath (called S1) contained 4 mol/L NaOH with litharge PbO(s) (the soluble Pb(II) species were HPbO$_2^-$ anions), pH > 14, anode current density of 1−5 mA/cm$^2$, mild stirring using a magnetic stirrer, bath temperature of 25−55 $^\circ$C, and electroplating time of 1−3 h.

3 Results and discussion

The cyclic voltammograms of Al/conductive coating/$\alpha$-PbO$_2$ electrode in basic solution S1(Fig.1(a)) and acid solution S2(Fig.1(b)) at a scan rate of 50 mV/s are shown in Fig.1. As shown in Fig.1(a), the potential of the beginning deposition of $\alpha$-PbO$_2$ is higher than 0.25 V[10]. Two anodic current density peaks at $\phi_{p1}$=0.29 V and $\phi_{p2}$=0.49 V are observed on the positive branch of the cyclic voltammogram. These peaks probably correspond to the formation of Pb$_3$O$_4$ and PbO$_2$, respectively. The cathodic current density peak is observed at approximately $\phi_{p3}$=0.10 V on the negative potential scan due to cathodic dissolution of PbO$_2$ deposited during the previous positive scan[14]. The exponential growth of the anodic current density at potential over 1.0 V is due to oxygen evolution.

![Fig.1](image-url) Cyclic voltammograms performed in different solutions with working electrode of Al/conductive coating/$\alpha$-PbO$_2$ at 50 mV/s: (a) Solution S1; (b) Solution S2

It is obvious from Fig.1(b) that the potential of the beginning deposition of $\beta$-PbO$_2$ is higher than 1.60 V. The anodic current density peak at $\phi_{p4}$=1.66 V is observed on the positive branch and the cathodic current...
density peak at $\phi_p=0.90\text{V}$ is observed on the negative branch. The anodic peak corresponds to the formation of surface lead dioxide or oxygen evolution and the cathodic peak is probably due to the formation of the oxidation of a soluble Pb(II).

3.1 Effect of current density

Fig.2 shows SEM images of deposited $\alpha$-PbO$_2$ on A1/conductive coating electrode from solution containing 4 mol/L NaOH saturated with litharge PbO(s) at different current densities. The surface of PbO$_2$ deposited at 1 mA/cm$^2$ consists of crystals with large size and surface uniformity (Fig.2(a)). At current density of 2 mA/cm$^2$ (Fig.2(b)), the PbO$_2$ surface becomes more uniform, consisting of rod-like grains of 50−60 nm. Further increasing the current density, there are a large number of small crystals without clear crystal edges (Fig.2(c)). The film exhibits a fibre texture with good orientation and recrystallizes into the oriented fibre texture standing on surface layer of the formed film[18]. At current densities of 4 mA/cm$^2$ or higher, however, the PbO$_2$ films of the fibre texture are randomly oriented and are highly porous. A comparison between Fig.2(d) and (e) reveals that the clear crystal edges appear at current density of 4 mA/cm$^2$. It is interesting that increasing current density from 1 to 5 mA/cm$^2$ causes the decrease of the diameter of fibres by approximately 20 nm.

This phenomenon may be explained as follows. The structure of the $\alpha$-PbO$_2$ deposit maybe result from complexation of lead cation in the electrodeposition solution. It has been reported[18, 22] that PbO dissolves in alkaline solutions with the formation of HPbO$_2^-$ and, to a less extent, polynuclear complexes. It seems probably that oxidation of such complexes can lead to polynuclear complexes containing PbO$_3^{2-}$. The hydrolysis of such species, following oxidation at the electrode to precipitate $\alpha$-PbO$_2$, is slow. At higher current densities, the solid oxide may precipitate at some distance from the electrode surface due to the increased concentration of the complexed Pb($IV$) cation. The agglomeration of such precipitates on the surface coatings can lead to a randomly oriented deposit. Also, a higher current causes the domination of oxygen evolution and, consequently, the formation of a series of

Fig.2 SEM images of $\alpha$-PbO$_2$ prepared from 4 mol/L NaOH saturated with litharge PbO(s) on A1/conductive coating electrode at 40 $^\circ\text{C}$ for 2 h with different current densities: (a) 1 mA/cm$^2$; (b) 2 mA/cm$^2$; (c) 3 mA/cm$^2$; (d) 4 mA/cm$^2$; (e) 5 mA/cm$^2$
pores on the surface of anodic coating. At lower current densities, precipitation will take place near the electrode at a slower rate. There is enough time for the fibre texture to find the most suitable position and to form equilibrium state with the $\alpha$-PbO$_2$ particles. The adsorption of anions on the crystallites may lead to the stability of ordered fibre structure. In addition, at lower current densities, most of current is used for the formation of $\alpha$-PbO$_2$[23].

3.2 Effect of variables on electrodeposition of $\alpha$-PbO$_2$

Fig.3 shows the effect of HPbO$_2^-$ concentration on the morphology of $\alpha$-PbO$_2$ coatings. The coatings were prepared from solutions containing 4 mol/L NaOH and 0.10–0.14 mol/L HPbO$_2^-$. As shown in Fig.3(a), a structure in leaves shape has been obtained at lower concentration of HPbO$_2^-$ (0.10 mol/L). With increasing concentration up to 0.12 mol/L, a well-structured morphology has been obtained and the pore has not been found (Fig.3(b)). A further increase in HPbO$_2^-$ concentration up to 0.14 mol/L results in the formation of larger PbO$_2$ particles with clear crystal edges(Fig.3(c)). At lower HPbO$_2^-$ concentrations, it seems that the mass transfer is a limiting step and the concentration polarization occurs on the electrode surface. At higher HPbO$_2^-$ concentrations, mass transfer process is quick enough compared with electron transfer and a well-formed structure of electrodeposited surface has been obtained[24]. Also, with increasing HPbO$_2^-$ concentration, the nucleation rate is correspondingly increased.

Fig.4 shows SEM images of PbO$_2$ prepared from solution containing 4 mol/L NaOH solutions saturated with litharge PbO(s) at 2.5 mA/cm$^2$ on A1/conductive coating electrode for 2 h: (a) 30 °C; (b) 40 °C; (c) 50 °C.
coating electrode for 2 h at different temperatures. At a lower temperature (30 °C), a loose structure consisting of a big fibre texture with obvious boundaries is deposited on the surface of the electrode. At a higher temperature (40 °C), a higher adherence coating is obtained with a pronounced change in the morphology. Further increasing temperature to 50 °C results in the structure of similar pattern with well-distributed fibre textures and more clearly crystal edges. However, either very low temperature or very high temperature gives a lump deposit [25]. Therefore, the effect of temperature is probably due to a decrease of the potential for the onset of electrodeposition [26]. At relatively low potentials, the formation of polycrystalline assemblies is less probable and disorder effects are less common. Considering that the growth rate along different crystallographic directions may depend only slightly on temperature, preferred crystallographic orientations can become more pronounced at higher temperature, which is mainly due to the fact that the number of the polycrystalline assemblies on the electrode surface is small.

Fig. 5 shows the SEM images of lead dioxide deposited on Al/conductive coating electrode from 4 mol/L NaOH solutions saturated with litharge PbO(s) at 3 mA/cm² for different plating time. As shown in Fig. 5(a), rod-like grains of the fibre texture are obtained in a shorter deposition time (1 h). However, with the prolonging of time to 2 h (Fig. 5(b)), a number of crystals with clear crystal edges and a few pores have been found. Further prolonging time to 3 h (Fig. 5(c)) results in a great number of small crystals without clear crystal edges and many small pores. It is also found that the process of electrodeposition of a new phase at Al/conductive coating electrode involves the laying down of a number of nucleation centers followed by the growth of centers, which coalesce to form a complete layer of the new phase [11]. It is interesting to note that H⁺ ions are generated at the anode during PbO₂ deposition. At the imposed current densities and under forced mass-transfer conditions, local pH in the basic solution is unlikely to become markedly lower than the original solution with the prolonging of time. However, the HPbO₂⁻ concentration decreases with increasing the plating time, and the nucleation rate is relatively decreased. A lower HPbO₂⁻ concentration is not high enough to contribute effectively in surface modification. The PbO₂ coating with a few pores is obtained in a very long time of electrodeposition.

3.3 Compositional analysis

EDX was used for identification of each sample by liner sweep scanning of sample. The compositional analysis of α-PbO₂ electrodeposits was performed under different experimental conditions using energy dispersive X-ray analyzer (EDAX). EDAX spectra of PbO₂ films deposited under different conditions are shown in Table 1 and Fig. 6. It is found from Table 1 that Pb contents also increase with the increase of HPbO₂⁻ concentration and the prolonging of the plating time. However, a remarkable change is obtained about the Pb content of α-PbO₂ film at different current densities. The Pb contents increase from 32.17% to 40.42% with current density form 1 mA/cm² to 4 mA/cm², and then decrease to 33.59% at the current density of 5 mA/cm². It is also found the Pb content of α-PbO₂ film is higher at 40 °C.

These results prove that the anodic layer is non-stoichiometric PbO₂. This phenomenon may be explained as follows. H⁺ ions are generated at the anode during PbO₂ deposition, which results in a decrease of

Table 1 Compositional analysis of PbO₂ films deposited under different conditions (molar fraction, %)

<table>
<thead>
<tr>
<th>Solution composition</th>
<th>Electrical parameter</th>
<th>Temperature/℃</th>
<th>x/%</th>
<th>Pb</th>
<th>O</th>
</tr>
</thead>
<tbody>
<tr>
<td>4 mol/L NaOH saturated with PbO(s)</td>
<td>J=1 mA/cm², t=2 h</td>
<td>40</td>
<td>32.17</td>
<td>67.83</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with PbO(s)</td>
<td>J=2 mA/cm², t=2 h</td>
<td>40</td>
<td>34.47</td>
<td>65.53</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with PbO(s)</td>
<td>J=3 mA/cm², t=2 h</td>
<td>40</td>
<td>38.86</td>
<td>61.14</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with PbO(s)</td>
<td>J=4 mA/cm², t=2 h</td>
<td>40</td>
<td>40.42</td>
<td>59.58</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with PbO(s)</td>
<td>J=5 mA/cm², t=2 h</td>
<td>40</td>
<td>33.59</td>
<td>66.41</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH +0.10 mol/LH₂PO₄−</td>
<td>J=1.8 mA/cm², t=2 h</td>
<td>40</td>
<td>34.17</td>
<td>65.83</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH +0.12 mol/LH₂PO₄−</td>
<td>J=1.8 mA/cm², t=2 h</td>
<td>40</td>
<td>37.26</td>
<td>62.74</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH +0.14 mol/LH₂PO₄−</td>
<td>J=1.8 mA/cm², t=2 h</td>
<td>40</td>
<td>37.48</td>
<td>62.52</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with litharge PbO(s)</td>
<td>J=2.5 mA/cm², t=2 h</td>
<td>40</td>
<td>35.22</td>
<td>64.78</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with litharge PbO(s)</td>
<td>J=3 mA/cm², t=1 h</td>
<td>40</td>
<td>35.54</td>
<td>64.46</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with litharge PbO(s)</td>
<td>J=3 mA/cm², t=2 h</td>
<td>40</td>
<td>38.08</td>
<td>61.92</td>
<td></td>
</tr>
<tr>
<td>4 mol/L NaOH saturated with litharge PbO(s)</td>
<td>J=3 mA/cm², t=3 h</td>
<td>40</td>
<td>38.65</td>
<td>61.35</td>
<td></td>
</tr>
</tbody>
</table>

Fig.6 EDAX spectrum of PbO₂ film

Fig.7 Typical XRD pattern of PbO₂ prepared in solution containing 4 mol/L NaOH solutions (40 ℃) saturated with litharge PbO(s) at 2 mA/cm² on A1/conductive coating electrode for 2 h

identify the structure of electrodeposited PbO₂. A typical XRD pattern of PbO₂ deposited on A1/conductive coating electrode at constant current density of 2 mA/cm² for 2 h in solution containing 4 mol/L NaOH solutions (40 ℃) saturated with litharge PbO(s) is shown in Fig.7. In Fig.7, α-PbO₂ can be identified from its intense lines (002, at 38.16°), (200, at 42.28°) and (132, at 71.54°) and PbO from its intense line (320, at 68.2°). The XRD pattern of α-PbO₂ prepared from alkaline solution indicates the presence of PbO, but the characteristic peaks of β-PbO₂, remarked by Miller number, have not been observed. The impurity is attributed to the co-deposition of an insoluble Pb(II) compound from the alkaline plating solution containing

the local pH at the interface. DELAHAY et al[22] showed that the solubility of Pb(II) exhibited a minimum at pH=9.4. Therefore, the local decrease of pH from a highly alkaline solution induces a decrease of the solubility of Pb(II) at the interface, and precipitation of lead hydroxide is possible from the concentrated plumbite solution[27]. On the other hand, a common feature should be noted that a porous morphology is favorable to increasing Pb contents of the PbO₂ films. And highly porous film is normally formed at large current density or potential polarization. Such highly porous structure greatly increases effective area of PbO₂ film.

3.4 XRD identification

Since PbO₂ can exist in two phases, XRD is used to

![Fig.6 EDAX spectrum of PbO₂ film](image)

![Fig.7 Typical XRD pattern of PbO₂](image)
lead hydroxide Pb(OH)₂ or the oxide PbO.

4 Conclusions

1) The current density during the deposition of PbO₂ has a strong influence on the morphology of the prepared film. A compact and uniform layer of lead dioxide is obtained at the current density not higher than 3 mA/cm². A further increase in current density results in a smaller particle with a high porosity. The optimum current density is 2 mA/cm².

2) The morphology and particle size distribution can be improved when the HPbO₂ concentration is 0.12 mol/L and the bath temperature is 40 °C. However, α-PbO₂ coating with a few pores is obtained in a very long plating time.

3) It is found from EDS that the PbO₂ deposited in alkaline conditions is highly non stoichiometric, which can be attributed to a porous behavior of deposited films. XRD pattern reveals that α-PbO₂ is dominated in the deposit.

References


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