Function of Silver in the Mixed ITE Activators for Lead-acid Batteries

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Abstract
The behaviors of Ag+ and activator were studied on Pb anode for lead-acid batteries. The deposition of Ag and dissolution of Ag+ take place at lower potential than redox couple of Pb/PbSO4. This process may be considered to activate the reaction in the redox couple of Pb/PbSO4. The activator molecules are less adsorbable on bare Pb, and adsorbed easily on PbSO4 formed on Pb. The complexes of Ag+ and activator molecule are adsorbable on bare Pb, and PbSO4 is initially formed on the adsorbed layer of complexes.

Keywords
lead-acid battery, anode, electrolyte, silver ion, organic polymer, activator

1. INTRODUCTION
The ITE mixed polymer activator added to electrolyte of lead-acid batteries have been found to extend battery life. The anode (negative electrode) sulfation is the main cause of deterioration in performance of the lead-acid batteries. It has been explained that the activator molecules are adsorbed on anode to produce the beneficial action for battery life.

On the other hand, the active materials are impregnated on grid alloy such as Pb-Ca, Pb-Sb and so on, and the alloy components may be dissolved as metal ions in the electrolyte during charge and discharge cycles. The additional alloy components are added for giving corrosion resistance property.

It is important to study the behaviors of dissolved metal ions from the grid alloy for battery performance. The silver is being used as an alloy component of Pb-Ca grid. This fact is worth paying particular attention, because the participation of Ag+ in the anode reaction of Pb seems to be important for battery performance in the electrolyte containing activator molecules. This paper describes the effects of Ag+ and activator on anode reaction on Pb by cyclic voltammetry.

2. EXPERIMENTAL
Cyclic voltammograms (CV) were recorded as follows. The test electrode was lead plate (99.99%, 1 x 1 cm²), which was chemically polished in CH3COOH-H2O2 solution at room temperature just before measurements of CV. Counter electrode was cylindrical lead with large surface area. Reference electrode was Hg/Hg2SO4/5 M H2SO4.

Electrolyte was 5M H2SO4 with/without additives, which were Ag2SO4 as Ag+ and the ITE mixed activator composed of polyvinylalcohol. The concentration of Ag2SO4 and activator was 2mg/100ml 5M H2SO4 and 1.0wt.%, respectively. The test electrode was immersed in the electrolyte, and scanned negatively from the spontaneous potential (-1.05V) to hydrogen evolution region. Then, the potential was positively scanned to -0.6V, and the scan was repeated between this potential region at the scan rate of 10mV/s.

3. RESULTS
Figure 1 shows the CV curves of Pb electrode in 5M H2SO4 and 5M H2SO4-Ag+ solutions. The redox couple

![Fig. 1 Cyclic voltammograms of Pb in 5M H2SO4 and 5M H2SO4-Ag+](image-url)
of Pb/PbSO₄ appears around -1.1 V, and corresponds to the charge and discharge reaction on anode of lead-acid battery. The hydrogen evolution current was observed below -1.6 V. When Ag⁺ was added to the electrolyte, the reaction of redox couple at -1.1 V was activated to increased the current two times. On the other hand, the hydrogen evolution deviates to positive potential. The Ag is formed on Pb at lower potential and decreases hydrogen overpotential.

Figure 2 shows the CV in 5M H₂SO₄-activator solution with that in 5M H₂SO₄. The addition of activator decreases the current in the redox couple of Pb/PbSO₄, and splits the anodic current of redox couple into two peaks. This suggests that the two processes take part in the formation of PbSO₄ from Pb. There is no effect of activator on hydrogen evolution region.

4. DISCUSSION

Figure 4 shows the mechanisms, in which the addition of Ag₂SO₄ as Ag⁺ increases the current in the redox couple of Pb/PbSO₄. When Pb electrode is polarized negatively near hydrogen evolution potential, Ag⁺ is reduced to deposit Ag metal on Pb. When this electrode is scanned to positive direction, the deposited Ag dissolves as Ag⁺ even before PbSO₄ formation potential. Such deposition and dissolution during potential scan gives fresh and bear surface of Pb to activate PbSO₄ formation. Accordingly, the redox current Pb/PbSO₄ is significantly increased in Ag⁺ containing solution.

Figure 5 shows the illustration to explain the effect of activator on the oxidation of Pb to PbSO₄. It is considered that the adsorption of activator takes an important role on this reaction. Activator has negative charge on oxygen atoms of the molecules, and hence is easily adsorbed on positive center of the electrode. If the Pb electrode is polarized near hydrogen evolution region, activator molecules are less adsorbable. Therefore, there is no deviation of hydrogen evolution potential compared with that in 5M H₂SO₄.

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**Fig. 2** Cyclic voltammograms of Pb in 5M H₂SO₄ and 5M H₂SO₄-activator

**Fig. 3** Cyclic voltammograms of Pb in activator and activator and Ag⁺ solution.

**Fig. 4** Mechanisms of the formation of PbSO₄ in the redox couple of Pb/PbSO₄

**Fig. 5** Mechanism of the formation of PbSO₄ in the activator containing solution
In the positive scan of Pb electrode from hydrogen evolution region, the oxidation of Pb to PbSO$_4$ starts on bare Pb surface. The electrode is covered with PbSO$_4$ by anodic current. The electrode covered with PbSO$_4$ is positively charged, and then the activator molecules are adsorbed on the film of PbSO$_4$. It is considered that this process gives the first peak current. The growth of PbSO$_4$ proceeds on the electrode with adsorbed activator to give the second peak current.

Figure 6 shows the reaction mechanism in the presence of both Ag$^+$ and activator molecules. In this case, the first peak current is decreased to give the shoulder in CV curve. This result indicates the some species are adsorbed on Pb electrode before the initial formation of PbSO$_4$.

**Fig. 6** Mechanism of the formation of PbSO$_4$ in the activator and Ag$^+$ containing solution

The complex ions of Ag$^+$ and activator molecules must be considered for the explanation of CV curve. Activator itself is less adsorbable species on Pb polarized negatively. The activator molecules with negative charge center react with Ag$^+$ to give complex ions. This is more adsorbable on Pb than free activator molecules. Accordingly, the first peak current is decreased to give the shoulder by the PbSO$_4$ formation on the polymer adsorbed layer. Then, the formation and growth of PbSO$_4$ proceed on the electrode adsorbed by activator to give the second peak current in CV.

5. CONCLUSION

The results obtained are summarized as follows.
(1) The addition of Ag$^+$ activates the reaction of Pb/PbSO$_4$ redox couple, and deviates the hydrogen evolution to positive region.
(2) The activator affects the reaction in redox couple of Pb/PbSO$_4$. This is explained by adsorption of activator on Pb and PbSO$_4$. The activator molecules are less adsorbable on Pb and adsorbed on PbSO$_4$.
(3) When both of activator and Ag$^+$ are present, the complex ion of Ag$^+$ and activator molecules is adsorbed on Pb. The initial formation of PbSO$_4$ proceeds on Pb adsorbed by the complex ion.

References


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