



Original Article

Study of effect of tin on the electrochemical properties of cycled PbSn alloys of lead-acid battery

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ABSTRACT

Tin has a beneficial effect on the electrochemical properties of the corrosion film formed on PbSn grids by reducing its resistivity mainly on cycling of the battery. Binary alloys with different tin contents were subjected to several charge-discharge cycles between and vs. Electrode in solution at. The electrochemical properties of alloys in sulfuric acid solution were studied by Cyclic Voltammetry, Linear Voltammetry and Chronpotentiometry. The compositions of the corrosion layer obtained at different numbers of cycles were determined by DRX. We were found that reduces the thickness and favorites a more conductive and dense layer between the grid and the positive active mass.

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1. Introduction

The addition of tin in the grid of valve regulated lead acid battery was as an attempt to illuminate the semiconductive layer constituted mainly of and between the collector and the positive active mass [1,2].

The introduction of tin as an alloy with lead dates back to the invention of the lead-acid battery by Planté in order to support the improvement of the mechanical characteristics of the plates. During the oxidation of PbSn alloys, tin is detached from the anode and deposited on the negative plate. It then decreases the hydrogen release overvoltage which increases the decomposition of water. The accumulator must always be under maintenance. In order to free this accumulator from this maintenance (addition of water), it was necessary to replace the tin by calcium which has good mechanical characteristics similar to those of tin. This replacement of tin does not have the desired results regarding corrosion resistance. The PbCa sulphur alloys of the so-called "PCL = premature capacity lost" or the (early capacity loss) due to the formation of a barrier layer of Sulphate and α -PbO which increase the resistance of the

layer between the collector and the active mass. This layer changes during the service of the accumulator isolating a very large quantity of the active mass and thus the lead accumulator is taken out of service. To remedy this phenomenon, different elements have been tested by which tin, which has given good results as a depassivation element, has been used.

It is in this contest that we have tried to contribute to understand first of all the phenomenon itself on tin-free electrodes and after addition of tin to lead. According to [3-11], PbSn and PbSn(Ca) alloys have a beneficial effect on reducing water decomposition.

Many new grid materials have been investigated in recent years to eliminate the disadvantages of PbSb and PbCa alloys on the performance of lead batteries containing different elements such as Ag, Al, Bi, La, Ce and Li [12-17].

This work aims to provide an explanation of the effect of tin on the electrochemical properties of cycled PbSn alloys of lead acid battery. Using different techniques.

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2. Materials and Methods

2.1. Material

The elaboration of the materials was carried out by fusion of pure metals (Pb and Sn) in a porcelain crucible. The melting was carried out in an electric resistance furnace. At

2.2. The electrochemical cell

The electrochemical experiments are performed with an AUTOLAB PGSTAT302N potentiostat/galvanostat controlled by a computer that records the data with NOVA 2.1 software, allowing the exploitation of the results of this study. The cell and the electrodes are connected directly to the device. The electrochemical cell is made of glass with

a temperature of 328°C, the bath formed was stirred slightly with a stainless steel spoon in order to achieve maximum homogeneity of the material.

The alloy in the liquid state is cast in a parallelepipedal metal mould with dimensions (75 x 25 x 190) mm. The ingot obtained is sawn into several pieces to be used as samples for different analyses.

a hole on one side (for the working electrode) and a glass cover with 02 holes allowing the passage of the electrodes (auxiliary and reference). This cell has a capacity of 200 ml containing the electrolyte in which the three electrodes used are immersed: working electrode, auxiliary electrode and reference electrode as shown in Fig. 1.

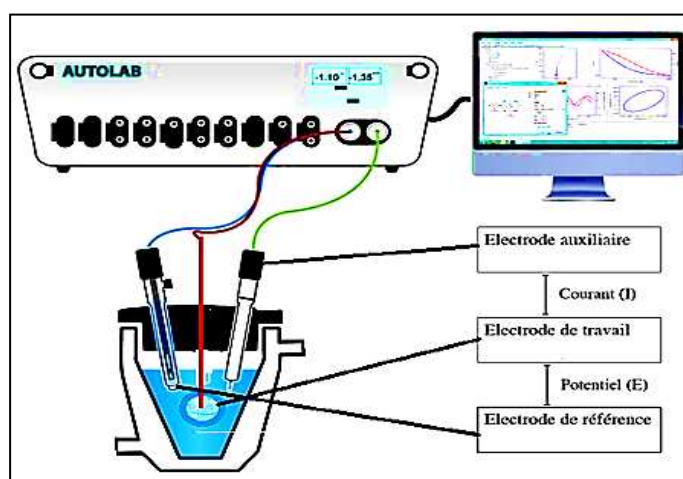


Fig. 1. Schematic representation of the experimental setup

Lead alloys require special care for metallographic analysis because they are very soft and very ductile. Forced polishing could cause deep structural transformations. This is why the entire polishing protocol is done manually. The specimens are prepared by mechanical polishing using an abrasive paper with a magnification of: (1000, 1200 and 2000).

After polishing, the specimens are thoroughly degreased and cleaned using acetone and then washed with distilled water. The electrochemical tests that have been adopted, aim to know the behavior of the alloys in a sulfuric medium (H_2SO_4) at 0.5M, which is the electrolyte used in the manufacture of "batteries" called lead-acid batteries and acid (Lead Acid Batteries) intended for starting vehicles.

The reference electrode used is the mercury sulphate electrode. The electrochemical tests were carried out using an AUTOLAB PGSTAT 302N Potentiostat connected to a microcomputer equipped with computer software used to transform the electrical signals from the electrochemical cell into numerical and graphical data interpreting the electrochemical behaviour of the studied alloys in a sulphuric medium.

2.3. The electrodes used are:

-The reference electrode is a mercury sulphate electrode filled with a saturated potassium sulphate solution, $Hg/Hg_2SO_4/K_2SO_4$.

-The auxiliary electrode is a pure lead electrode with a larger surface area than the working electrode, which ensures the passage of current through the electrical circuit and its measurement.

-The working electrode: we have formed pellets with a diameter of 1 cm and a thickness of 0.5 cm.

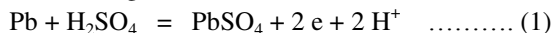
3. Results and Discussion

3.1. Potentiodynamic measurements.

3.1.1. Cyclic Voltammetry

Before any potentiodynamic experiment, the potential of the working electrode is maintained at the hydrogen release potential for 20 minutes in order to reduce any impurities that may exist on the surface. The measurements were carried out using an AUTOLAB PGSTAT302N potentiostat/galvanostat controlled by NOVA 2.1 software, allowing the acquisition and the exploitation of the

obtained results. The ramp is made to scan the potential of the working electrode between -1.2V and 0.7 V with respect to the reference electrode Hg/Hg₂SO₄/K₂SO₄ saturated with a scanning speed of 10 mV/s. A total of 40 relative to the oxidation of Pb to PbSO₄ around -860 mV for the following reaction:



At this moment the electrode is covered with a layer of PbSO₄ and the oxidation current is rapidly decreased (diffusion-controlled kinetics). The cutting of the PbSO₄ crystals is done in such a way as to block any exit when the SO₄²⁻ ions arrive at the surface of the substrate and the electrode is passivated. By increasing the potential a small peak towards -442 mV is noted (A2) corresponding to the oxidation of Pb into α-PbO. This is explained by different

charge-discharge cycles were performed for each alloy. A cycle starting from the hydrogen evolution potential (-1.2V) is eluted on Fig. 2. We notice that when the potential increases anodically, we obtain a noted peak (A1) researchers as being the result of the reaction of Pb²⁺ and the O²⁻ ion which is able to cross the PbSO₄ dense layer because of its small volume compared to that of SO₄²⁻ which has a larger atomic radius than that of the pores left by the PbSO₄ crystals.

The passivation domain will continue after this last peak. In the cathode scan after the arrival at 700 mV, we notice a small peak of reduction of PbO to Pb around -900 mV noted (C1). More cathodically, we obtain another cathodic peak of reduction of PbSO₄ to Pb around -981 noted (C2) Fig. 2.

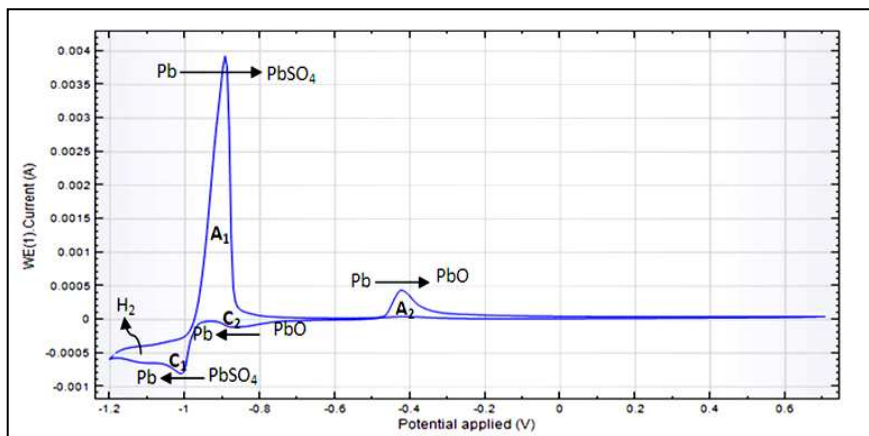


Fig. 2. Different pairs in the potential range from -1.2 to 0.7 V

3.1.2. Linear Voltammetry

Linear Voltammetry is also carried out by scanning the potential of the working electrode between -1.2V and 0.7 V with respect to the reference electrode Hg/Hg₂SO₄/K₂SO₄ saturated with a scanning rate of 10 mV/s in sulphuric acid at 0.5 M at 25°C. The results obtained for the current and corrosion potential at the 10th

and 40th cycle were recorded. A typical voltammogram is shown in Fig. 3. On this voltammogram we observe a peak of oxidation of Pb to PbSO₄. A lead passivation step is then obtained. At this stage the entire surface of the alloy is covered with α -PbO under a layer of lead sulphate. In the area of PbO formation, the appearance of a small peak around 440 mV corresponds to the oxidation of Pb to PbO.

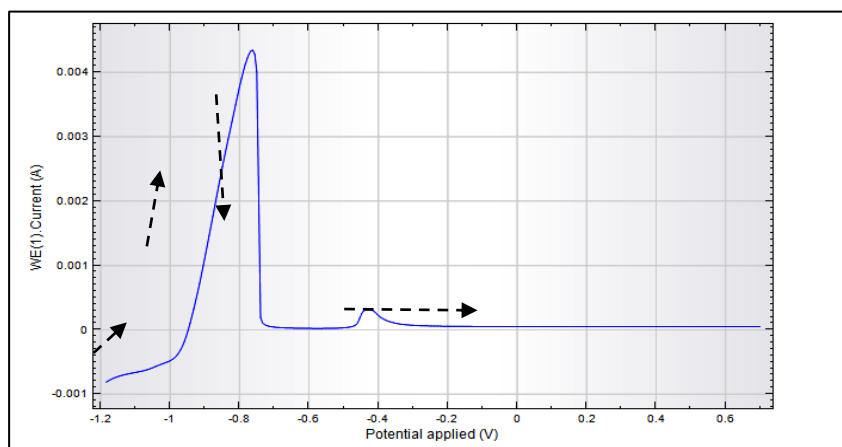


Fig. 3. Example of a linear voltammogram

3.1.3. Tafel curves

The superposition of the Tafel curves of the Pb, Pb -0,2 wt.%Sn and Pb-2 wt.%Sn electrodes are given in Fig. 4.

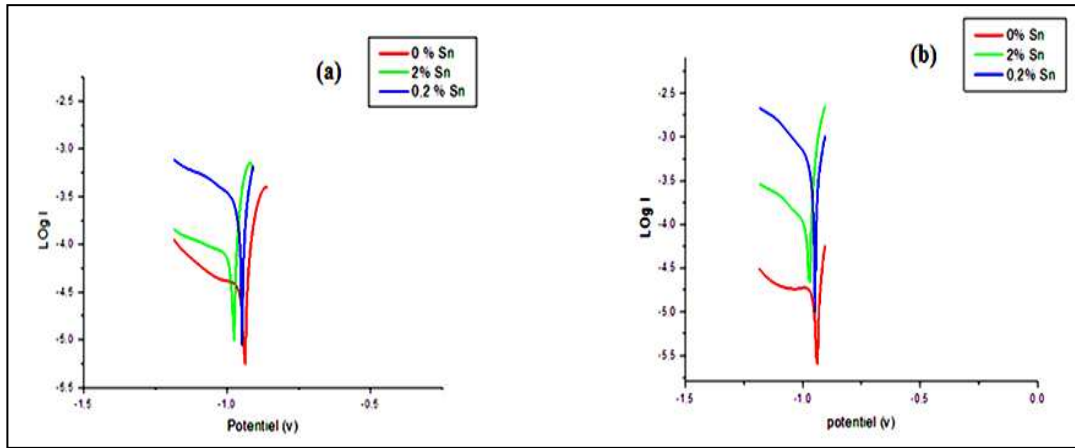


Fig. 4 .Tafel plots: (a) 10 cycles;(b) 40 cycles

The data collected from these Tafel curves are shown in Table 1.

Table 1. Results of I_{corr} and R_p of the different alloys at different cycles according to the Tafel plots.

N° of cycles	Pb			Pb -0,2 wt.%Sn			Pb -2 wt.%Sn		
	I_{corr} (μA)	E_{corr} (mv)	R_p (Ω)	I_{corr} (μA)	E_{corr} (mv)	R_p (Ω)	I_{corr} (μA)	E_{corr} (mv)	R_p (Ω)
10	19	-937.59	498.97	34.27	-969.82	221.81	107.1	-943.28	103.4
40	7.063	-940.65	1419.8	15.35	-971.87	160.28	106.1	-943.82	69

To better examine the influence of tin on the corrosion layer resistance of PbSn alloys, we have reported the results obtained from the corrosion current density as a function of number of cycles for the three samples studied on the histogram Fig. 5.

We clearly see that the corrosion current decreases with cycling. This is due to the growth of the corrosion layer consisting mainly of the mixture of PbSO_4 , $\alpha\text{-PbO}$ for Pb

and PbSO_4 electrodes, $\alpha\text{-PbO}$ and some tin oxides for Pb-0.2 wt.%Sn and Pb-2 wt.%Sn alloys. On the other hand, for a well-defined number of cycles, the corrosion current increases by increasing the tin content within the alloy from 0 to 2%. Tin increases corrosion resistance as the number of cycles increases. In other words, tin decreases the polarization resistance of PbSn alloys.

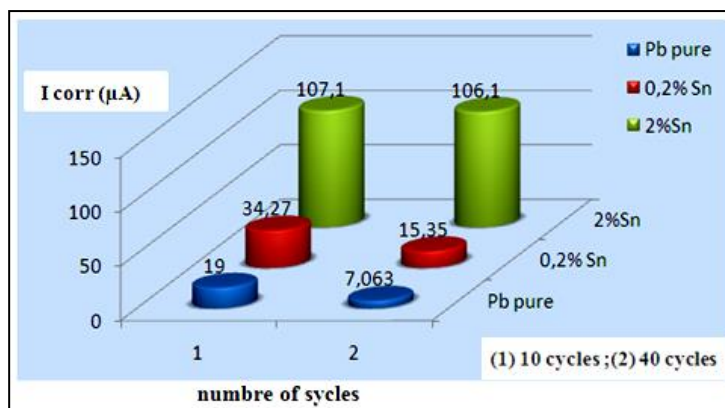
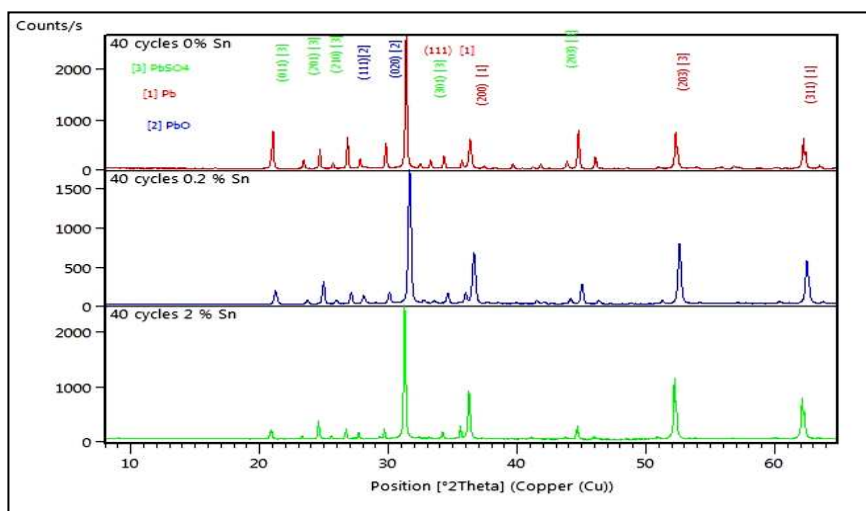


Fig. 5. Histogram of the evolution of corrosion current as a function of number of cycles at different alloys

Fig. 7. Diffractogram of sample at 40th cycles of different alloys

According to the table regrouping the relative intensities of the characteristic sulphate and lead monoxide lines of the three electrodes in the 10th cycle

Table 3 we can see that tin reduces the rate of the two phases of the corrosion layer

Table 3. Comparison of peak characteristics at different alloys and different cycles

10 th	Picks	hkl	2θ (°)	d [Å°]	Reference data(*)		Intensity I [%]		
					2θ (°)	d [Å°]	Pb	Pb -0,2 wt.%Sn	Pb -2 wt.%Sn
	PbSO4	0 1 1	20,828	4,26499	20,886	4,2497	12,24	14,4	7,72
	PbSO4	2 0 1	24,6027	3,61852	24,659	3,6074	39,4	28,54	8,46
	PbSO4	2 1 0	26,7955	3,32891	26,808	3,3228	14,93	19,25	45,03
	PbSO4	3 0 1	34,3447	2,61855	34,34	2,6094	13,12	11,49	16,16
	PbSO4	2 0 3	44,6823	2,02646	44,674	2,0268	20,31	26,32	22,63
	PbO	1 1 1	28,9992	3,08449	29,071	3,0692	29,48	57,03	48,86
	PbO	0 2 0	30,0001	2,8033	32,576	2,7465	70,52	42,97	51,14
40 th	PbSO4	0 1 1	20,8777	4,2485	20,886	4,24968	27	13,47	12,56
	PbSO4	2 0 1	24,6434	3,60264	24,659	3,60743	14	30,97	33,33
	PbSO4	2 1 0	26,7851	3,32844	26,808	3,32284	22	14,4	18,72
	PbSO4	3 0 1	34,3451	2,6163	34,34	2,60937	8,6	13,55	11,47
	PbSO4	2 0 3	44,6851	2,02533	44,674	2,02684	28	27,6	23,91
	PbO	1 1 1	29,01631	3,21337	29,071	3,06922	27	39,91	39,91
	PbO	0 2 0	30,29556	3,00258	32,576	2,7465	73	60,09	60,09

(*): ICDD PDF4+

4. CONCLUSION

This work aimed to study the effect of tin on the electrochemical properties of cycled PbSn alloys in 0.5 M H₂SO₄ compared to Hg/Hg₂SO₄/K₂SO₄ saturated reference electrode. Using different techniques (potentiometric, X-ray diffraction characterization) .we have reached the following conclusions:

- The corrosion layer consists of a thin layer mainly composed of PbSO₄ and α-PbO confirmed by DRX in the 3rd cycle.

- The quantities of these phases increase with the increase in the number of cycles (growth of the corrosion layer). Their quantities are lower in PbSn alloys.

- Tin is beneficial for PbO removal. Its action lies in the reduction of thickness by the formation of conductive oxides such as SnO_x.

Conflict of Interest

The authors declare that they have no conflict of interest

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