Electrochemical growth of trijunction semiconductors for Renewable Energy applications

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Abstract

In the present work, we report the growth of an electrode based on TiO₂/Fe₂O₃/MnO₂ trijunctionon stainless steel substrate,to develop a new electrode for electrochemical supercapacitor. The synthesized electrodes are characterized by X-ray diffraction (XRD) and Fourier Transform Infrared, Spectroscopy (FTIR). Electrochemical properties are studied using cyclic voltammetry (CV), and electrochemical impedance spectroscopy techniques. The observed results suggest that the trijunction electrode of the three metal oxide materials has a high electrochemical property due to its large surface area, porous structure, and low resistance to charge transfer. In fact, it exhibits capacitive in 1 M KOH electrolyte, indicating a promising electrode material for electrochemical supercapacitors.

Keywords: metal oxides thin films, electrochemical technique, trijunction electrode, supercapacitor application

Fabrication of supercapacitor electrodes:

We have successfully developed an original process to synthesize TiO₂/Fe₂O₃/MnO₂ trijunction metal oxides onto stainless steel substrate as a supercapacitor electrode for energy storage, using three-step strategy methods. To understand the synthetic process of trijunction SS/TiO₂/Fe₂O₃/MnO₂, we have schematized the synthesis conditions in Fig. 1.

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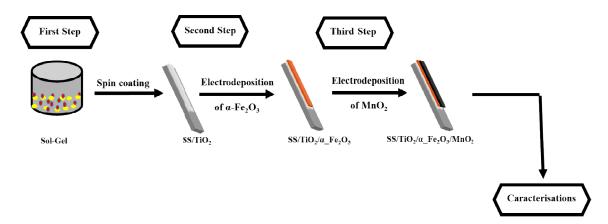


Figure 1: Electrochemical growth of TiO₂/Fe₂O₃/MnO₂ trijunction electrode

Results and discussions

1.Structural analysis

Figure (2a) reveals the XRD pattern recorded for the stainless steel substrate (SS), The TiO_2,α -Fe₂O₃ and the heterojunction TiO_2/α -Fe₂O₃. The dashed lines represent the contribution of the 304L stainless substrate at $2\theta = 43.8^{\circ}$, 44.6° and 50.8° . According to JCPDS (Joint committee on Power Difraction) database with card number 00–023-0298. The (+) represents the hematite peaks, it was observed that the rhombohedral α -Fe₂O₃ structural peaks at 2θ values of 24.21° , 33.23° , 35.70° were indexed to the (012), (104) and (110) planes, respectively (JCPDS card No. 86–0550)[1, 2]. The peak indicated by (*) correspond to TiO₂ anatase phase characterized by the main orientations (101) at $2\theta = 25.3^{\circ}$ [3]. It is recalled that the crystal lattice of anatase is tetragonal (JCPDS-ICDD maps n° 01-084-1286). By comparison with the diffractograms of the materials alone, the peaks of the heterojunction were attributed to the SS substrate, TiO₂ anatase phase and hematite Fe₂O₃. This suggests on the one hand that the rhombohedral structure of α -Fe₂O₃ is well formed on the TiO₂ layer and on the other hand that thin layers of TiO₂ are stable during the second process of hematite synthesis.

For the electrodeposition of MnO₂(Figure 2b), we will not see any peak attributed to manganese oxide except that the characteristic peaks of the stainless-steel substrate. This behavior can be explained by the fact that this compound is amorphous and not detectable by XRD analysis [4]. Hammami and his group [5]showed that even with a heat treatment at 450°C for one hour the MnO₂material remained amorphous and no change in the XRD spectrum was observed. By comparison with TiO₂/Fe₂O₃ heterojunction electrode and MnO₂

spectrum, the spectrum of trijunction electrode contain only the peaks attributed to TiO_2 (101), Fe_2O_3 (012), (104) and (110) planes and any peaks related to MnO_2 are obtained. This confirm another time that le manganese oxide is not detected by XRD analysis.

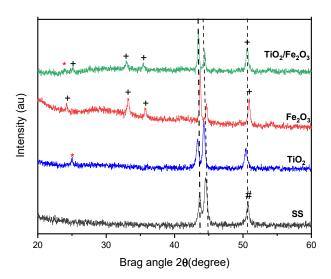


Figure 2. XRD diffraction pattern of thestainless-steel substrate (SS), TiO_2 , α -Fe₂O₃ and the heterojunction TiO_2/α -Fe₂O₃

2. Electrochemical performance

In this work, the electrochemical capacitive performance of the metal oxide materials thin films grown by electrodeposition method onto SS substrate is determined using CV measurements in 1 M KOH solution at a fixed scan rate of 50 mV.s⁻¹ (Figure 3).To neglect the influence of the substrate, the CV measurement of stainless steel substrate was also performed with the same conditions. A slight difference is observed between the CV curve of the Stainless-steel substrate (SS) and TiO₂/Fe₂O₃, indicating that the combination of both TiO₂ and Fe₂O₃ is not suitable enoughto affect the capacitance electrode behavior. However, when TiO₂/Fe₂O₃ is recovered by MnO₂ (pink curve), we can see that the trijunction electrode at a potential scan rate of 50mV.s⁻¹exhibit nearly ideal EDLC behavior with quasi-rectangular form of CV curve along the potential current axis without evident redox peaks, indicating that the trijunction electrode has an ideal electric double-layer capacitive behavior [6]. Besides, the remarkable increase of the area of CV curve is indicating the capacitive nature of this trijunction. Comparing the trijunction electrode to only MnO₂ thin film electrode (black curve), we can clearly see that the prepared TiO₂/Fe₂O₃/MnO₂ electrode has a higher area curve, which means that the obtained trijunction has higher charge storage properties. In fact,

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we can note that the addition of MnO₂ on the TiO₂/a-Fe₂O₃ junction forms a porous structure, allowing an efficient ion exchange between the electrolyte and the active electrode, which promotes the rapid oxidation-reduction reaction to obtain a high specific capacity. All these characteristics ensure full contact between the electrolyte and the electrode in order to accelerate the diffusion of ions.

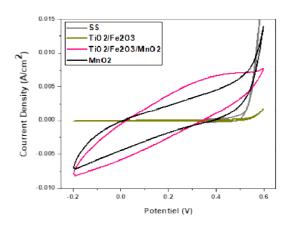


Figure 3: Superposition of CVs forSS, TiO₂/ Fe₂O₃, MnO₂, TiO₂/ Fe₂O₃/MnO₂of various films studied (at a scanning rate of 50 mV/s).

Conclusion

In summary, TiO_2/α -Fe₂O₃/MnO₂ have been successfully elaborated onto SS substrate for the first time by a two-step process of combined Sol-Gel/spin coating technique followed by electrodeposition method. The capacitive effect of MnO₂ on the electrical properties of SS/TiO₂/Fe₂O₃electrode was investigated. The trijunction electrode exhibited high efficiency supercapacitor properties comparing to Fe₂O₃/TiO₂ electrode. From this study, we can conclude that the proposed trijunction electrode can be available for electrochemical performance for supercapacitor application.

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