

Adsorption and Electrochemical Behaviour of Cyt-c on Transparent Mesoporous TiO₂ Films

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The immobilization of biomolecules upon electrodes and other surfaces is important not only for studying their function and structure but also for the development of bioanalytical devices such as biosensors. We investigate in this study the use of mesoporous optically transparent TiO₂ films as substrates for protein immobilization.

TiO₂ films with developed mesoporous architecture are among the best candidates as host matrices for the incorporation of various optoelectroactive guest molecules. The films have a high surface area, are optically transparent, exhibit good electrical conductivity (wide bandgap ~ 3.2 eV) and can be prepared by low cost screen printing technologies for the development of dye sensitized solar cells, electrochromic displays, electrode devices and colorimetric chemical sensors.

Therefore in this study, their ability to provide a novel and versatile immobilization surface for biomolecules and the development of an amperometric biosensor for H₂O₂ will be examined. The resulting TiO₂ films were structurally characterized by a variety of techniques such as SEM (see figure 1), TEM, XRD and BET analysis. In addition a thorough study of the binding of Cytochrome-c upon them [1] and the parameters that influence its adsorption are presented. [2] Finally, with the use of a 3-electrode electrochemical cell, electrochemical techniques such as cyclic voltammetry and spectroelectrochemistry (see figure 2) were used in order to investigate the electrochemical behavior of the immobilized biomolecules and their potential use for the development of an electrochemical biosensor for H₂O₂.

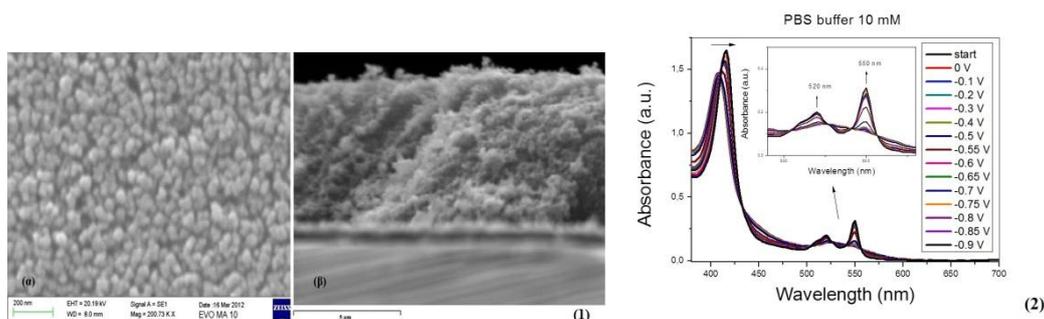


Figure 1: SEM images of TiO₂ thin film (top view and cross section).

Figure 2: Spectroelectrochemical study of Cyt-c/TiO₂ in PBS buffer 10 mM, pH 7.

References

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