

Non-enzymatic Amperometric Sensor for H₂O₂ Based on MnCO₃ Thin Film Electrodes

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Abstract: The present study describes development of a non-enzymatic amperometric sensor for detection of H₂O₂ based on MnCO₃ thin film electrodes. The film was deposited on electroconductive FTO coated glass substrates using simple chemical bath deposition method. The phase composition of the thin film was confirmed by X-ray diffraction analysis. The electrochemical properties and the sensor sensitivity towards H₂O₂ were examined using cyclic voltammetry and chronoamperometry in 0.1 M phosphate buffer solution with pH = 7.5. It was revealed that the sensing mechanism is based on electrocatalytic oxidation of H₂O₂, involving Mn species as redox mediators. According to the results, the best sensor response towards H₂O₂ was found at $E = +0.25$ V, with detection limit and sensor sensitivity of 10.0 μ M and 2.64 μ A cm⁻² mM⁻¹ (for the range of 0.09–1.8 mM), respectively, associated with $R^2 = 0.999$.

Keywords: amperometric sensors, hydrogen peroxide, manganese(II) carbonate thin films, electrocatalysis.

INTRODUCTION

THIS research contributes to the application of manganese(II) carbonate (MnCO₃), which already demonstrated interesting behaviour when used for preparing electrochromic materials,^[1] as a precursor for synthesizing perovskites applied in high temperature solid oxide fuel cells,^[2] and especially as an electrode material in supercapacitors.^[3–7] The possibility of controlled deposition of uniform MnCO₃ thin films on FTO-coated substrates,^[1] makes this material eligible to be studied as a working electrode in electrochemical systems for hydrogen peroxide (H₂O₂) sensor applications. H₂O₂ is an important substance that finds a wide use in various fields. Its oxidizing properties enable application in chemical and petro-chemical industry as a strong oxidizer, bleaching agent, disinfectant and propellant.^[8–14] H₂O₂ is also used in medicine, pharmacy, cosmetics, food and beverage industry.^[10,13–16] Apart from industrial applications, H₂O₂ is also important for the living cells.^[17] It is well established that H₂O₂ is formed as a product in the mitochondria due to

enzymatic reactions that involve free radicals.^[17,18] Moreover, the increased mitochondrial production of H₂O₂ causes cytotoxic effects^[11,13,18,19] through activation of several classes essential signalling proteins that compromise the cell reproduction, causing diseases such as cancer, diabetes, cardiovascular and neurodegenerative disorders.^[17,20,21] The presence of H₂O₂ in the cells is significantly detrimental and commonly responsible for proliferation, apoptosis and/or necrosis of the cells, which depends on the cytosolic steady state concentration.^[18,20] From this point of view, an accurate and precise quantification of H₂O₂ is substantially important. Hence, an enormous research strive is in progress in order to develop simple, efficient and reliable methods for detection and quantification of H₂O₂ at relatively low concentrations in biological fluids.^[17] There are numerous methods for detection and quantification of H₂O₂. These include redox titrations,^[22] chemiluminescence,^[23–26] fluorescence and fluorimetry,^[27–29] spectrophotometry,^[30–32] chromatography^[33] and electrochemistry.^[13,34] The electrochemical sensors are based on sensing either reduction or oxidation