

ORIGINAL PAPER

Non-enzymatic hydrogen peroxide sensor based on a nanoporous gold electrode modified with platinum nanoparticles

^{a,b}Guang Yin, ^bLing Xing, ^bXiu-Ju Ma, ^aJun Wan*

^aKey Laboratory of Eco-chemical Engineering, Ministry of Education, College of Environment and Safety Engineering, Qingdao, ^bCollege of Chemistry and Molecular Engineering, Qingdao University of Science and Technology, Qingdao 266042, China

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A novel non-enzymatic electrochemical sensor based on a nanoporous gold electrode modified with platinum nanoparticles was constructed for the determination of hydrogen peroxide (H₂O₂). Platinum nanoparticles exhibit good electrocatalytic activity towards hydrogen peroxide. The nanoporous gold (NPG) increases the effective surface area and has the capacity to promote electron-transfer reactions. With electrodeposition of Pt nanoparticles (NPs) on the surface of the nanoporous gold, the modified Au electrode afforded a fast, sensitive and selective electrochemical method for the determination of H₂O₂. The linear range for the detection of H₂O₂ was from 1.0×10^{-7} M to 2.0×10^{-5} M while the calculated limit of detection was 7.2×10^{-8} M on the basis of the 3σ /slope (σ represents the standard deviation of the blank samples). These findings could lead to the widespread use of electrochemical sensors to detect H₂O₂.

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Introduction

H₂O₂ is cytotoxic to cellular life and acts as an indicator of oxidative stress (Lases et al., 2000; Yorek, 2003); it is the product of several biological, enzyme-catalysed reactions (Clark, 1970; Ruiz et al., 1993; Yang et al., 1995, 2006; Thomé-Duret et al., 1996; Niwa et al., 1998; Armstrong & Wilson, 2000; You et al., 2003). This has motivated the development of stable hydrogen peroxide sensors with high sensitivity, low limit of detection and a rapid and accurate response for utilisation in the food industry, in environmental waste management and in medical diagnostics. Many H₂O₂ analytical methods have been developed, including titrimetry, spectrophotometry, chemiluminescence and electrochemistry (Wang et al., 2005; Thenmozhi & Narayanan, 2007; Zhang et al., 2008a; Chen et al., 2009; Ma et al., 2009).

Many enzyme-based biosensors have been constructed to achieve H₂O₂ oxidation (Vianello et al.,

2007; Kafi et al., 2008; Mala Ekanayake et al., 2008; Zhou et al., 2010), but the application of these H₂O₂ enzymatic biosensors is limited, in part due to the disadvantages represented by their rigorous environment, high cost, the instability of enzymes and the complicated process to achieve enzyme immobilisation (Jia et al., 2002; Zhou & Ju, 2004; Lin et al., 2007). In comparison with enzyme-based sensors, non-enzymatic sensors exhibit a high stability, free from the influence of temperature.

Metal nanoparticles have been studied extensively because of their unique functions (Lewis, 1993; Heath, 1998; Bönnemann & Richards, 2001; Roucoux et al., 2002; Link & El-Sayed, 2003). The catalytic behaviours of the metal NPs have been investigated because of their high catalytic activity in many chemical reactions. In particular, Pt NPs have been demonstrated to reduce the H₂O₂ oxidation–reduction over-voltage efficiently and investigated for their catalytic activity in H₂O₂ reduction. Pt NPs have found many

*Corresponding author, e-mail: wanjundz@sohu.com