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Development of an Electrochemical Tyrosinase Biosensor Using Sulfonic Acid-Modified Magnetic Nanoparticles for Dopamine and Phenolic Compound Detection

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Abstract

An electrochemical biosensor based on tyrosinase (Tyr) was developed for the detection of dopamine and phenolic compounds across medical, environmental, and industrial applications. The biosensor utilized sulfonic acid (SO₃H)-modified magnetic nanoparticles (MNPs) prepared via a hydrothermal synthesis method followed by in-situ surface functionalization. The MNPs enhanced the electrical conductivity and enzyme adsorption on the electrode surface, significantly improving sensor performance. A magnetic carbon paste electrode was fabricated with an optimal graphite-to-paraffin weight ratio. Subsequently, MNP suspension was deposited on the electrode surface, followed by the addition of 15 μ L of 4 mg/mL Tyr solution. The biosensor was allowed to stabilize for 24 hours before undergoing electrochemical characterization. Cyclic voltammetry (CV) revealed an apparent electron transfer rate constant (k_s) of 0.008 s⁻¹ and a formal potential (E°) of 0.21 V. Differential pulse voltammetry (DPV) demonstrated the biosensor's performance for catecholamine detection. For caffeic acid, the limit of detection (LOD) was 45.4 μ M within a linear range of 1–74 μ M, with a sensitivity of 1.68 μ A $\cdot\mu$ M⁻¹ \cdot cm⁻² and a Michaelis-Menten constant (K_m) of 30 μ M. For catechol, the LOD was 68 μ M in a linear range of 1–107 μ M, with a sensitivity of 1.8 μ A $\cdot\mu$ M⁻¹ \cdot cm⁻² and K_m of 55.3 μ M. For L-DOPA, the LOD was 50.6 μ M in a linear range of 1–137 μ M, with a sensitivity of 0.9 μ A $\cdot\mu$ M⁻¹ \cdot cm⁻² and K_m of 68.92 μ M. The results suggest that incorporating cross-linking agents such as glutaraldehyde to covalently stabilize the enzyme on the nanoparticles could further enhance the biosensor's kinetic and electrochemical properties, offering improved reliability and sensitivity.

Key words

Magnetic nanoparticles, sulfonic acid modification, polyphenols, L-DOPA, and carbon paste electrode