

Contents lists available at ScienceDirect**Biosensors and Bioelectronics**journal homepage: www.elsevier.com/locate/bios**Tyramine detection using PEDOT:PSS/AuNPs/1-methyl-4-mercaptopyridine modified screen-printed carbon electrode with molecularly imprinted polymer solid phase extraction**Ying Li^a, Cheng-Hung Hsieh^b, Chi-Wei Lai^b, Ying-Feng Chang^a, Hsin-Yi Chan^b, Chang-Feng Tsai^b, Ja-an Annie Ho^a, Li-chen Wu^{b,*}^a *BioAnalytical Chemistry and Nanobiomedicine Laboratory, Department of Biochemical Science and Technology, National Taiwan University, Taipei, 10617 Taiwan*^b *Department of Applied Chemistry, National Chi Nan University, Puli, Nantou, 545 Taiwan***A B S T R A C T**

Tyramine (4-hydroxyphenethylamine), which is a monoamine metabolized by monoamine oxidase (MAO), exists widely in plants, animals, fermented foods, and salted foods. The incidence of hypertension, or “cheese effect”, which is associated with a large dietary intake of tyramine while taking MAO inhibitors has been reported; therefore, the measurement of tyramine is an urgent concern. Herein, an efficient approach that integrates a molecular imprinting polymer for solid phase extraction (MISPE) technique with a sensitive electrochemical sensing platform (SPCE/PEDOT: PSS/AuNP/1-m-4-MP) for the quantification of tyramine is presented. Enhanced electrode conductivity was achieved sequentially by constructing a conductive polymer (PEDOT: PSS) on a screen-printed carbon electrode (SPCE), followed by electrodeposition with gold nanoparticles (AuNPs) and, finally, by modification with positively charged 1-methyl-4-mercaptopyridine (1-m-4-MP) using an Au-S bond. Tyramine was isolated selectively and pre-concentrated by the MISPE technique; electroanalysis that used differential pulse voltammetry (DPV) in NaOH (0.1 M, pH 13) was conducted successively. Experimental parameters (such as modes of electrode modification, ratio of PEDOT: PSS, pH of electrolyte, time required for AuNP deposition, and 1-m-4-MP concentrations) that were associated with optimal detection conditions were evaluated also. We obtained a linear concentration range (5–100 nM, $R^2=0.9939$) with LOD and sensitivity at 2.31 nM, and $3.11 \mu\text{A nM}^{-1} \text{cm}^{-2}$, respectively. The applicability of our technique was demonstrated by analyzing tyramine in spiked serum and milk. The feature of our newly developed analytical methods that coupled sample pre-treatment (sample clean-up and pre-concentration) with sensitive detection makes it a promising tool for quantifying of tyramine.

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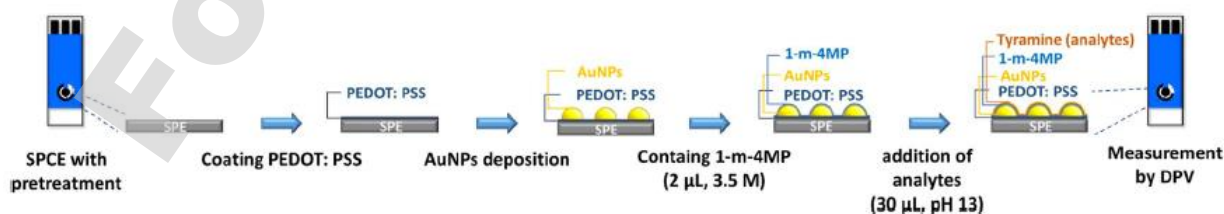


Fig. 1. Schematic diagram of the modification of screen-printed carbon electrodes (SPCE) and detection of tyramine.



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