

Electrochemical Sensing of Vitamin D₃: A Comparative Use of Glassy Carbon and Unmodified Screen-Printed Carbon Electrodes

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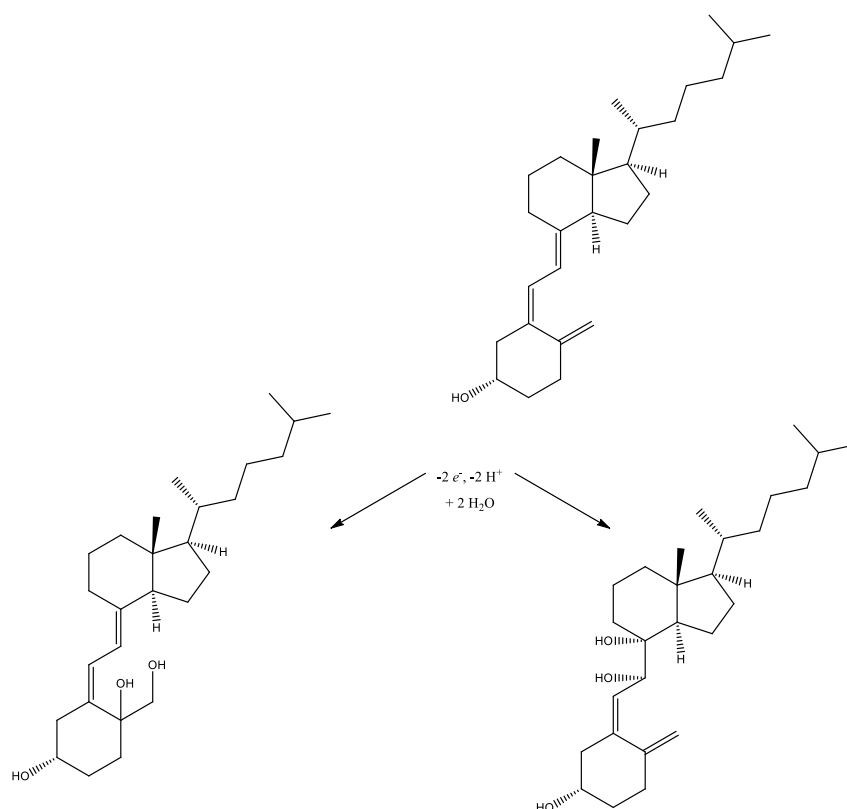


Figure S1 – Two possible oxidation mechanisms of Vitamin D₃.

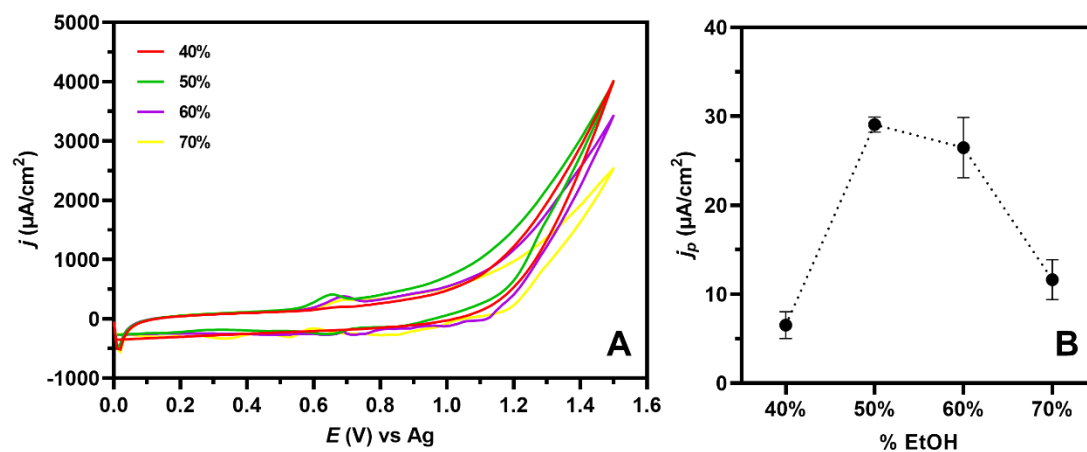


Figure S2 – Cyclic voltammograms of 0.90 mmol L⁻¹ vitamin D₃ in 0.1 M LiClO₄ prepared in different proportions of ethanol/water using a SPCE. Voltammograms were measured between 0 V and + 1.5 V; scan rate of 50 mV s⁻¹; step potential of 10 mV. Three replicates were performed for each ethanol content.

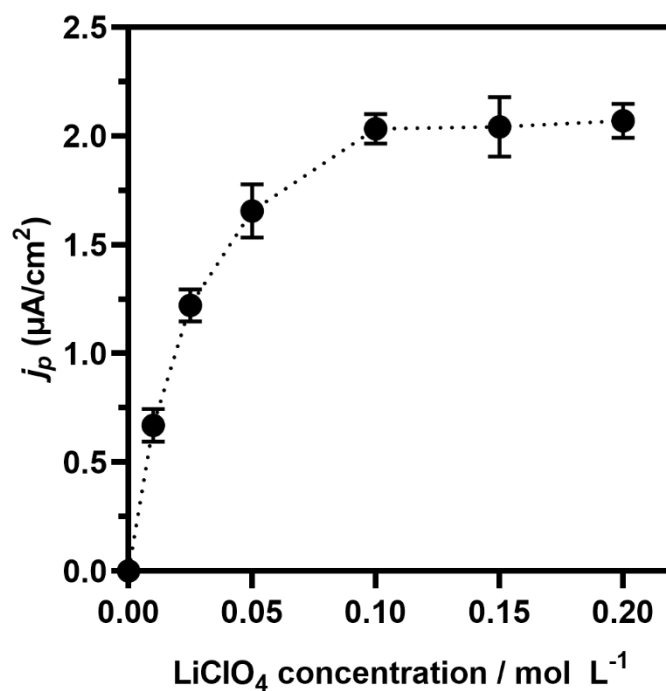


Figure S3 – Dependence of peak current on the concentration of lithium perchlorate in the supporting electrolyte solution, prepared in 50/50 ethanol/water. Measurements, in triplicate, were performed on the GCE system by SWV on a 0.1 mmol L⁻¹ Vitamin D₃ solution. Voltammograms were measured between 0 V and + 1.5 V; frequency of 50 Hz; amplitude of 50 mV and step potential of 5 mV.

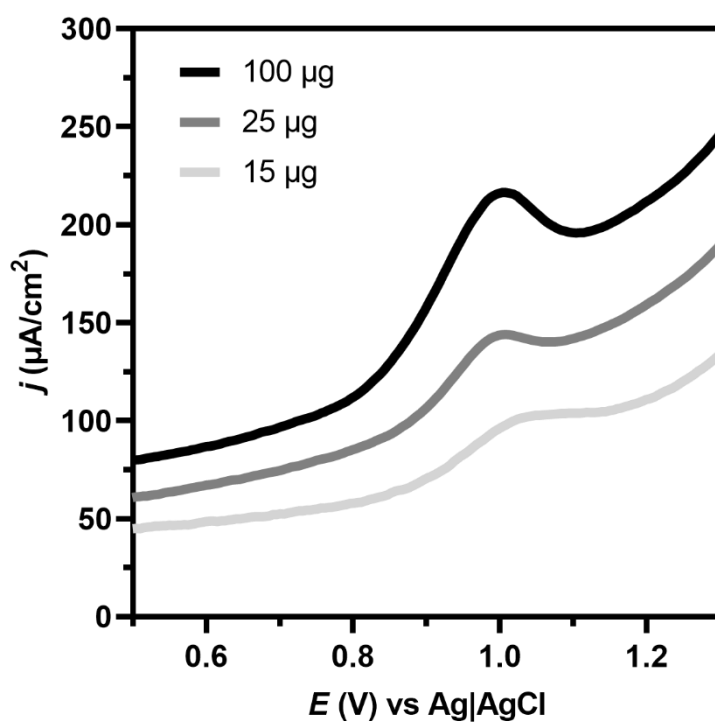


Figure S4 - Representative square-wave voltammograms on the GCE of the analysed Vitamin D₃ samples, obtained in 50/50 ethanol/water solution. SWV was performed using the optimized electrochemical conditions.