Book of Abstracts

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Influence of nitrogen on boron doped diamond electrodes in the nitrate electrochemical response

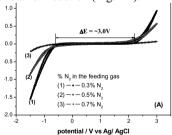
J. T. Matsushima¹, L. L. G. Silva², M. R. Baldan¹ and <u>N. G. Ferreira¹</u>

¹LAS - Instituto Nacional de Pesquisas Espaciais – INPE

²LAP- Instituto Nacional de Pesquisas Espaciais – INPE

Caixa Postal: 515, CEP: 12245-070, São José dos Campos, São Paulo, Brasil neidenei@las.inpe.br

Diamond films grown by chemical vapor deposition has been optimized during recent years because of their excellent physico-chemical properties. Due to its wide working potential window and high corrosion resistance, diamond electrodes have been proposed to be beneficial in a range of electroanalytical applications, which are depending of the material aspects¹. One of these aspects is the film doping, which can be carried out using boron or nitrogen to turn the diamond films as a semiconductor. Boron doped diamond (BDD) electrodes has been already studied as a new material with extraordinary properties that turn them as a chemical sensors². Particularly, high overpotential presented by BDD electrodes for hydrogen evolution has demonstrated to be a decisive attribute for studying the detection and reduction of several chemical species. Besides, the nitrogen addition during the BDD growth process (N and B codoped electrodes) may also enhance this cathodic overpotential³. This work presents the influence of nitrogen on BDD electrodes in response of nitrate ions reduction. Actually, the large use of nitrogen-based compounds in agriculture has increased their ion concentration in the biosphere promoting a significant pollution problem. The working potential window of the N and B co-doped diamond electrode in 1.0 M H₂SO₄ (Fig. 1A) about of 3.0 V, justifies their performance improvement for nitrate reduction and detection. Furthermore, the variation of N2 concentration during the BDD growth process was crucial in their electrochemical behavior. Increasing the N₂ concentration favored the occurrence of the nitrate reduction without the interference of hydrogen evolution reaction (Fig. 1B).



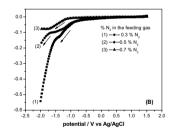


Fig 1: **(A)** cyclic voltammograms and **(B)** linear sweep voltammograms of nitrate reduction for the BDD electrodes grown in different N₂ concentration. Solutions used (a) 1.0 M H₂SO₄ and (B) 0,1 M KNO₃ and scan rate at 0.1 Vs⁻¹

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