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UVC/e-H₂O₂ technology using gas diffusion electrodes with low La loadings on Printex L6 for the removal of emerging contaminants

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This study explored the application of UVC combined with electrogenerated hydrogen peroxide (UVC/e-H₂O₂) for the degradation of benzotriazole (BTA) at pH 6, using Printex L6 (PL6C) carbon-based gas diffusion electrodes (GDEs) modified with trace amounts of lanthanum oxide, La₂O₃ (1.0–10.0% La/PL6C). Oxygen reduction reaction (ORR) studies using the rotating disk electrode technique identified 5% La/PL6C as the optimal condition for H₂O₂ electro-generation. Both modified and unmodified GDEs were tested in a non-divided cell to evaluate H₂O₂ production at different current densities (10, 20, and 30 mA cm⁻²), with 0.05 mol L⁻¹ K₂SO₄ as the supporting electrolyte.

The degradation of 10 mg L⁻¹ of BTA at a current density of 30 mA cm⁻² was analyzed using different processes: anodic oxidation (AO), e-H₂O₂, only UVC, UVC/AO and, UVC/e-H₂O₂. The UVC/e-H₂O₂ process achieved 98.2% removal of BTA in 15 minutes of electrolysis, while mineralization rates reached 62.9% after 1 hour of treatment. The UVC/e-H₂O₂ process was also applied to assess BTA removal in a river water sample, monitoring contaminant mineralization under exhaustive treatment, which reached 83.3% after 5 hours.

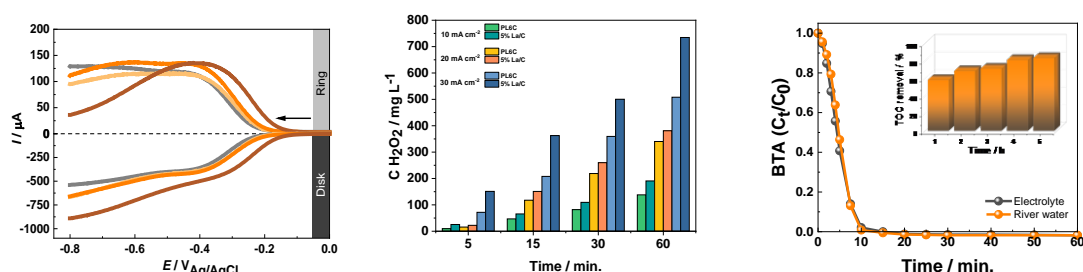


Figure 1. (A) LSV curves for the ORR at 1.0 – 10.0% content of La/PL6C in a 0.05 M K₂SO₄ solution, saturated with O_{2(g)}. Electrode rotation speed: 900 rpm. (B) Concentration of H₂O₂ electrogenerated as a function of the applied current density. (C) BTA degradation by AO, e-H₂O₂, only UVC, UVC/AO and, UVC/e-H₂O₂ process at 30 mA cm⁻².

The low percentages of La₂O₃ used to modify the PL6C-GDEs demonstrated outstanding performance in both H₂O₂ electro-generation and the degradation of emerging contaminants, confirming its effectiveness for water treatment applications ^[1,2].

Acknowledgments

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