

Anodic Production of Hydrogen Peroxide on Carbon-based Materials

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The production of hydrogen peroxide (H₂O₂) using renewable electricity, oxygen (O₂) and water (H₂O) is an alternate “green” route to the current anthraquinone process^[1]. There are two approaches for the electrochemical synthesis of H₂O₂. One pathway is the partial reduction of O₂ at cathodes employing either noble metal alloys or doped carbon. Another possible route is the two-electron oxidation of water to H₂O₂, which competes with the four-electron oxidation of H₂O to O₂^[2]. This anodic production of H₂O₂ is desirable since it can be coupled with other cathodic reactions, such as CO₂ reduction reaction to valuable products. However, it remains a great challenge to develop efficient electrocatalysts for H₂O₂ anodic synthesis, which can suppress the thermodynamically favored oxygen evolution reaction (OER). The objective of our research is to identify the parameters that will allow maximizing the efficiency of the catalyst and the electrochemical process for the anodic production of H₂O₂. Carbon has been used as an electrocatalyst due to its large surface area, conductivity, chemical stability in alkaline electrolytes, and low cost. Selective oxidation of water to hydrogen peroxide on different carbon materials was conducted using a two compartment H-cell, where copper was used as a counter electrode. The impact of electrolyte concentration, current density, pH values of the electrolyte, temperature, and ion exchange membranes, have been investigated in order to optimize reaction conditions for the anodic production of H₂O₂. A concentration of 1.73 mM for H₂O₂ was achieved at the applied current density of 100 mA cm⁻² in 2.7 M KHCO₃ electrolyte using a cation exchange membrane in a flow cell. (**Figure 1**).

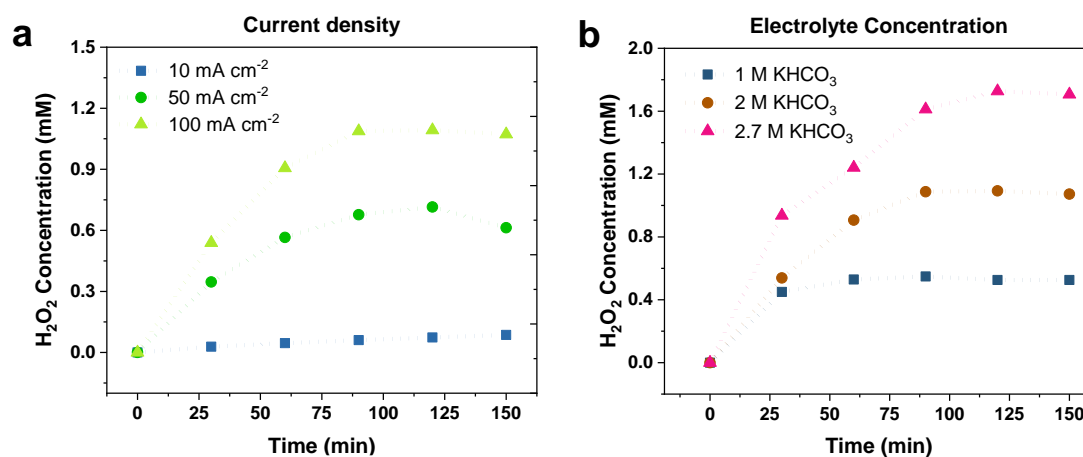


Figure 1: Hydrogen peroxide concentration in the anolyte during electrolysis at (a) different current densities using 2 M KHCO₃ and (b) in different electrolyte concentrations of KHCO₃ at 100 mA cm⁻²

References:

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