
Ethylene glycol electrooxidation reaction on silver surfaces modified with low noble metal content

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The electrooxidation of ethylene glycol (EG) in alkaline media represents a promising route for sustainable energy generation using direct alcohol fuel cells (DAFCs), owing to EG's high energy density, low toxicity, and renewable origin [1,2]. This study investigates the synthesis and electrochemical performance of silver (Ag) surfaces modified with low contents of Pd, Pt, Au, and Ta, aiming to enhance catalytic activity and long-term durability [3]. The Ag/C electrode exhibited low electrocatalytic activity, whereas its combinations with Pd and Pt significantly improved performance. Among the synthesized materials, the PdAuAg/C catalyst displayed the highest activity, reaching a peak current density of $120 \text{ mA}\cdot\text{cm}^{-2}$ with good cycling stability, suggesting strong synergistic and bifunctional effects. In contrast, Pd/C (Etek) and PdTa/C showed inferior performance and clear signs of deactivation. Although Pt/C (Etek) achieved the highest initial current ($>150 \text{ mA}\cdot\text{cm}^{-2}$), it also experienced current decay, in line with the known stability limitations of platinum-based systems [2]. Chronoamperometric tests supported these findings, with PdAuAg/C and AgAu/C maintaining currents above $30 \text{ mA}\cdot\text{cm}^{-2}$, outperforming both monometallic and commercial benchmarks. These results highlight the potential of multimetallic catalysts supported on modified carbon materials as durable and efficient platforms for EG oxidation in alkaline media [3].

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