Oxidation of ethanol under potential scanning conditions

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The electrochemical oxidation of ethanol in fuel cells is currently considered as an important alternative energy source. For high energy efficiency, complete ethanol oxidation is required, and so CO_2 should be the major product.

In this work a non-dispersive infrared carbon dioxide monitor has been used to measure the amount of CO_2 produced from the oxidation of ethanol in a direct ethanol fuel cell (DEFC). During the oxidation of ethanol, absorbed CO accumulates on the surface of the Pt anode. Linear sweep and cyclic voltammetry in combination with the CO_2 monitor can be used to investigate its oxidation to CO_2 , and how this influences the yield of CO_2 . The periodic oxidation of absorbed CO increases the CO_2 yield and so can increase DEFC efficiency.

Linear sweep experiments

As seen in Fig. 1, curve a, stripping of CO from the anode by linear sweep voltammetry while monitoring the CO₂ concentration in the fuel cell exhaust resulted in only a transient production of CO₂, even when the potential was held at the upper limit of +0.9V for an extended period. This procedure leads to oxidation of absorbed CO to CO₂, but negligible sustained ethanol oxidation.

Cyclic voltammetry experiments

In contrast, it has been found that with cyclic voltammetry, the CO_2 produced increased significantly and was sustained for the duration of the potential cycling period (Fig. 1, curve b). Thus CO that is stripped from the anode at high potentials is replaced by ethanol adsorption and partial oxidation at lower potentials.

Additional results obtained over a range of temperatures and with a variety of anode catalysts will be presented.



Figure 1. CO_2 *vs.* time curves for (a) linear sweep voltammetry (+0.1 V to +0.9 V at 10 mV/s, followed by 28 min at +0.9 V) (b) cyclic voltammetry (+0.1 V to +0.9 V at 10 mV/s) with 0.1M ethanol at the anode and N₂ at the cathode of a DEFC.

References:

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