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Investigation of Carbon Corrosion Resistance of CNT Containing Electrode

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Carbon support corrosion is one of the major degradation mechanisms of polymer electrolyte membrane (PEM) fuel cell. Carbon oxidation occurs in PEM electrode and is accelerated at high potential created by adverse operating conditions and improper distribution of reactants and products [1, 2, 3]. Carbon corrosion can lead to the thinning of the electrode layer and severe performance degradation. The detailed mechanisms of carbon support corrosion induced performance loss are still not fully understood; it is believed that the following events contribute to the decay: (1) structural collapse of the porous electrode due to the loss of carbon; (2) carbon surface modification due to the formation of hydrophilic surface groups which can induce water accumulation and flooding of the electrode; (3) detachment and dissolution of platinum, which results in the reduction of platinum surface area. Together, these processes contribute to the loss of electrode performance.

Carbon nanotubes (CNT) have been used as alternative Pt catalyst support, it has been reported that CNT support offers better resistance to carbon corrosion [4]. This can be partly attributed to the high intrinsic stability of CNT in comparison with carbon black (CB). In addition, CNT-containing electrode is more structurally robust, because CNT can reinforce the electrode layer to avoid and delay the collapse of electrode structure. To help elucidate the mechanisms for the improved carbon corrosion resistance of CNT-containing electrode, three types of electrodes were prepared: (1) Pt catalyst supported on carbon black from Tanaka Kikinzoku Kogyo (TKK), (2) Pt catalyst supported on the CNT, (3) A mixture of Pt catalysts supported CB and supported on CNT. The cathodes were subjected to an accelerated catalyst support corrosion test protocol. Periodic performance and electrochemical diagnostic tests were conducted to track the decay behavior. The initial test results are shown in Figure 1-4, the decay rate of the electrode with mixed Pt/CB and Pt/CNT is decreasing compared with that of Pt/CB electrode. These results will be compared with electrode with Pt/CNT catalysts. Polarization was collected at 80°C, 85% RH, ambient pressure with H₂/Air on anode/cathode. The cyclic voltammetry was plotted at the same cell temperature and with a scan rate of 100 mV/sec. The cell area was 25 cm², and the anode was used as a reference.

Figure 1. Performance decay with Pt/CB catalysts.

Figure 2. Performance decay with mixed Pt/CB and Pt/CNT catalysts.

Figure 3. Change of ECA with Pt/CB catalysts.

Figure 4. Change of ECA with mixed Pt/CB and Pt/CNT catalyst.

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