
Synthesis of various Ag-based electrocatalysts for fuel cell applications via plasma discharge in solution

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Ag is considered to be a promising electrocatalyst since it is roughly 100 times cheaper than Pt and is more stable than Pt in base. However, their synthetic route has been still issued because the synthesis of Ag nanocrystals is mostly performed by a chemical method. Recently, a method for the synthesis of the Ag nanocrystals by means of plasma discharge in liquid phase is introduced and plasma discharge in liquid phase is an attractive method for the synthesis of nanoparticles because it is faster (microseconds), simpler (one-step method) and more cost-effective (only requiring an ion source) than traditional colloid methods. In this paper, reported was the preparation of the Ag-based electrocatalysts (*e.g.* Ag nanowires, nanodendrites, and Ag@Co₃O₄ core-shell nanocrystals) using bi-polar pulsed plasma discharge in water for catalytic oxygen reduction and ethanol oxidation in alkaline solutions. The Ag nanowires and nanodendrites were achieved by tailoring the reduction kinetics with adjusted electron densities in plasma. From the linear sweep voltammetry, the specific activity of Ag nanowires exhibited the 2.5 times higher than Ag nanodendrites, and the electron transfer number was calculated to be 3.83, which is indicative of a four-electron ORR pathway. For the Ag@Co₃O₄ core-shell hybrid nanostructures, it was found that the reactive oxygen species generated from plasma-assisted decomposition of water molecules played an important role in controlling the phase structure of the nanoparticles. The electrochemical measurements for oxygen reduction reaction revealed that the electrocatalytic activity of the Ag@Co₃O₄ core-shell nanocrystals were higher than that of pure Ag and Co₃O₄ nanoparticles.

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