

Graphene / TiO₂ Composite Electrode for Oxygen Reduction

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Abstract

A commercial graphite nanocrystals was oxidized to graphene oxide (GO) using Hummers' method. Later, GO powder was reduced either by placing it under a H₂ gas flow in a controlled gas reactor at 450 °C or using a hydrazine hydrate solution (HH) in a 1000-Watt microwave oven. X-ray photoelectron spectroscopy (XPS) and X-ray diffraction patterns (XRD) confirmed that both of the oxidation and reduction processes of graphite and GO powders, respectively, were incomplete. Also, the surface area of the H₂ gas - reduced GO powder was found to be higher than the case where HH was used to reduce the same powder. Fourier transform infrared spectroscopy (FT-IR) and XPS (Figure 1) have revealed that GO surface consists mainly of hydroxyl, epoxy, carbonyl and carboxylic groups. The electrocatalytic properties of (i) glassy carbon (GC), (ii) commercial TiO₂ (P25-TiO₂/GC), (iii) TiO₂-supported GO (GO/TiO₂/GC), (iv) TiO₂ – supported HH – reduced GO (HHRGO/TiO₂/GC) and (v) TiO₂ – supported H₂ gas – reduced GO (H₂RGO/TiO₂/GC) electrodes towards the oxygen reduction reaction (ORR) in acidic solution in presence and absence of UV radiation were examined. The results have shown that the H₂RGO/TiO₂/GC electrode has the best electrocatalytic activity in terms of current at a certain potential but glassy carbon electrode (GC) was found to be the best in terms of the onset potential of the ORR.

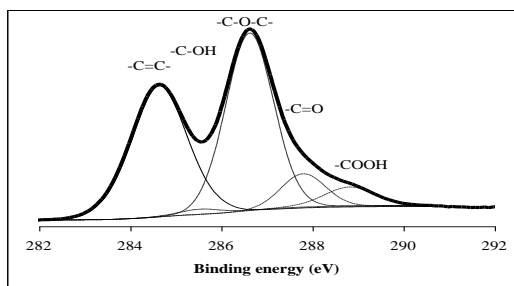


Figure 1: XPS of GO