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"Production of Formaldehyde through Photocatalytic Reduction of CO₂ over rGO Grafted NiO-CeO₂ Heterostructure Nanocomposite"

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Resumen

CO₂ emission led by excessive fossil fuel usage has increased its concentration in the earth's atmosphere to unprecedented level. Therefore, finding ways to prevent the climatic change driven by CO₂ surge is one of the most urgent tasks for the scientific community. While a variety of solutions such as capturing CO₂ in rocks, making CO₂-based copolymers such as polypropylene carbonate (PPC), and electrochemical CO₂ reduction have been suggested for the capture and utilization of CO₂, all these methods require external energy to activate, reduce, and convert CO₂ into fuel products due to high thermodynamic stability of CO₂. As the search for cost-effective efficient methods for CO₂ reduction is continued, utilization of cheap renewable energy such as solar light might be a viable alternative for producing high energy chemicals through CO₂ photoreduction in cheaper way. During the past two decades, various semiconductors such as TiO₂, CdS, CaFe₂O₄, ZnO, and TaON have been tested as photocatalysts for CO₂ reduction. However, low efficiency and poor product selectivity of CO₂ reduction remained the principal obstacles for practical application of these photocatalysts. Solar light is one of the sustainable, abundant, and cheaper energy sources, which has been utilized for the reduction of CO₂ and its conversion to fuel. Photoelectrochemical and photocatalytic CO₂ reductions have been carried out using solar light. Taking the advantage of solar light as cheap energy source, PEC and PC CO₂ reductions have been performed over several semiconductor photocatalysts. Use of different semiconductor photocatalysts yielded different solar fuels such as methanol, ethanol and formaldehyde. During the past two decades, various semiconductors such as TiO₂, CdS, CaFe₂O₄, ZnO, and TaON have been tested as photocatalysts for CO₂ reduction. However, low efficiency and poor product selectivity of CO₂ reduction remained the principal obstacles for practical application of these photocatalysts. We combined p-type NiO particles with n-type CeO₂ particles to form p-n junction interfaces, which can suppress the charge carrier recombination process at the surface of CeO₂ nanoparticles. Further, we introduced reduced graphene oxide (rGO) in the composite to exploit its high electrical conductivity and charge trapping behavior, which allowed an easy accumulation of electrons at the hybrid composite photocatalyst surface and subsequent sequential multi-electron transfer at each of the intermediate steps of CO₂ reduction process. The sequential multielectron transfer process induced photocatalytic reduction of CO₂ molecules adsorbed and activated at CeO₂ surface, producing formaldehyde as liquid fuel at 421.09 μ mol g⁻¹ h⁻¹ rate.

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