

Facile synthesis of CaFe_2O_4 for visible light driven treatment of polluting palm oil mill effluent: Photokinetic and scavenging study

Ashwin Charles^a Maksudur R.Khan^a; Kim HoongNg^c; Ta Yeong Wu^d; Jun Wei Lim^e; Suwimol Wongsakulphasatch^f; ThongthaiWitoon^g; Chin KuiCheng^{ab}

^a Faculty of Chemical & Natural Resources Engineering, Universiti Malaysia Pahang, Lebuhraya Tun Razak, 26300 Gambang Kuantan, Pahang, Malaysia

^b Centre of Excellence for Advanced Research in Fluid Flow (CARIFF), Universiti Malaysia Pahang, Lebuhraya Tun Razak, 26300 Gambang Kuantan, Pahang, Malaysia

^c Chemistry & Chemical Engineering, Xiamen University Malaysia, Jalan Sunsuria, Bandar Sunsuria, 43900 Sepang, Selangor, Malaysia

^d Chemical Engineering Discipline, School of Engineering, Monash University, Jalan Lagoon Selatan, 47500 Bandar Sunway, Selangor, Malaysia

^e Department of Fundamental and Applied Sciences, Universiti Teknologi PETRONAS, 32610 Seri Iskandar, Perak, Malaysia

^f Center of Ecomaterials and Cleaner Technology, Department of Chemical Engineering, Faculty of Engineering, King Mongkut's University of Technology North Bangkok, Bangkok 10800, Thailand

^g Center of Excellence on Petrochemical and Materials Technology, Department of Chemical Engineering, Faculty of Engineering, Kasetsart University, Bangkok 10900, Thailand

ABSTRACT

In this paper, a facile synthesis method for CaFe_2O_4 is introduced that produces a catalyst capable of significant photocatalytic degradation of POME under visible light irradiation. The co-precipitation method was used to produce two catalysts at calcination temperatures of 550 °C and 700 °C dubbed CP550 and CP700. CP550 demonstrated the maximum COD removal of 69.0% at 0.75 g/L catalyst loading after 8 h of visible light irradiation which dropped to 61.0% after three consecutive cycles. SEM images indicated that the higher calcination temperature of CP700 led to annealing which reduced the pore volume ($0.025 \text{ cm}^3/\text{g}$) and pore diameter (10.3 nm) while simultaneously creating a smoother and more spherical surface with lower S_{BET} ($9.73 \text{ m}^2/\text{g}$). In comparison, CP550 had a rough hair-like surface with higher S_{BET} ($27.28 \text{ m}^2/\text{g}$) and pore volume ($0.077 \text{ cm}^3/\text{g}$) as evidenced by BET analysis. XRD data indicated the presence of CaFe_5O_7 in the CP550 composition which was not present in CP700. The presence of Wustite-like FeO structures in CaFe_5O_7 are likely the cause for lower photoluminescence intensity profile and hence better charge separation of CP550 as these structures in CaFe_2O_4 have been known to increase resistivity and electron localization. The COD removal of CP550 dropped from 69.0% to just 7.0% upon adding a small quantity of isopropanol into the reaction mixture indicating hydroxyl radicals as the primary reactive oxidative species.

KEYWORDS:

CaFe_2O_4 ; Photocatalysis; POME; Visible light; Co-precipitation