

Enhanced photocatalytic degradation of oxytetracycline using LaFeO₃ supported on Mullite ceramic foam

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INTRODUCTION & AIM

The immobilization of a photocatalyst on a solid support is essential for practical environmental applications, as the photocatalyst is then not dispersed in the solution with contaminants and can be recycled several times. LaFeO₃ powder and LaFeO₃ supported on a mullite-based ceramic foam are promising materials for the photocatalytic degradation of oxytetracycline (OTC) under visible light irradiation and in the presence of H₂O₂.

METHOD



Figure 1. (A) LaFeO₃ powder, (B) pristine mullite foam; (C) LaFeO₃ supported on the mullite foam.

Table 1. experimental conditions

Photocatalyst	Light Source	[OTC] (mol L ⁻¹)	[H ₂ O ₂] (mol L ⁻¹)
LaFeO ₃ supported on a mullite-based ceramic foam	2 Daylight, 8W each	5.0 × 10 ⁻⁶	3.0 × 10 ⁻³

Parallel tests were carried out, one with the LaFeO₃ supported on the mullite foam (Fig. 1C), the other with the mullite foam without catalyst (Fig 1B). The mixtures were stirred in the dark for 20 minutes to allow adsorption/desorption equilibrium on the mullite foam and catalyst surface. Then the two fluorescent lamps (daylight, 8 W each, irradiance on the sample 89 Wm⁻²) emitting in the 380–780 nm region were turned on for 240 min.

RESULTS & DISCUSSION

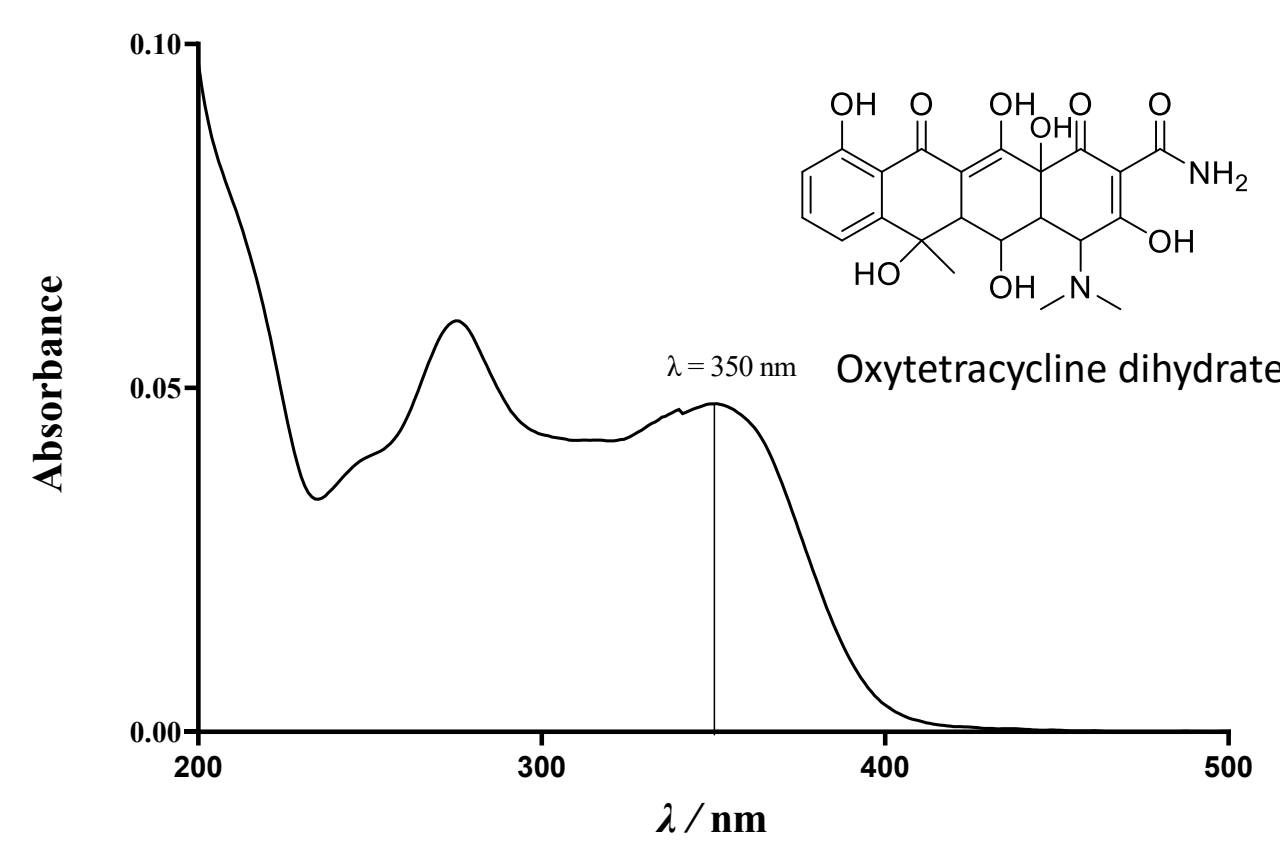


Figure 2. UV–Vis absorption spectra of Oxytetracycline and its chemical structure.

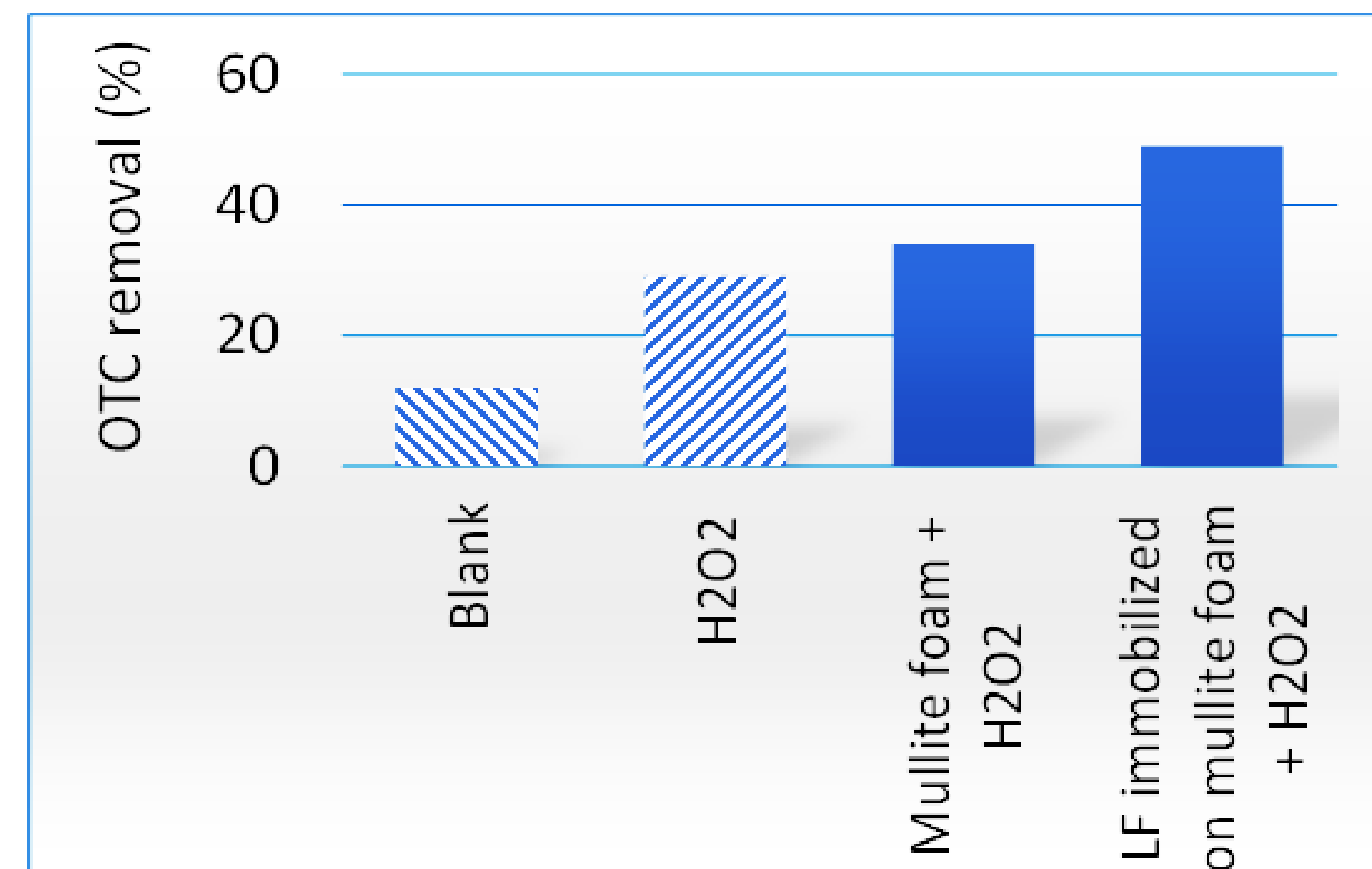


Figure 4. photocatalytic degradation of OTC after 240 min of visible light irradiation. LF = LaFeO₃. C₀=5·10⁻⁶ M, the OTC solution is phosphate buffered at pH = 5.0 since at pH=5.0 OTC is poorly photolyzed.

The point of zero charge (PZC) for LaFeO₃ is at pH= 8.9 suggesting that the surface charge of catalyst is positive in our tests. OTC exists predominantly as a zwitterion between pH 3.6 and 7.5 resulting from the loss of proton from the phenolic diketone moiety.

CONCLUSION

- Plain mullite foam and LaFeO₃ supported on the mullite foam degraded 34% and 50% of OTC respectively, after 240 min of irradiation, in presence of H₂O₂.
- A small addition of H₂O₂ increases the photodegradation rate of organic pollutants by removing the surface-trapped electrons, thereby lowering the electron-hole recombination rate and increasing the efficiency of hole utilization for reactions, such as OH⁻ + h⁺ → •OH.
- In this study the OTC solution is buffered at pH = 5.0, so OTC has zwitterionic form. Physisorption of OTC molecules on the catalyst promotes photodegradation of the contaminant by their surface diffusion to the photocatalytic sites.