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Synthesis of a Mesoporous Silica/Titanium Dioxide Nanocomposite and Evaluation of Photoactivity William Adams¹, Martin G. Bakker¹*, and I. Atly Jefcoat² ¹Department of Chemistry, The University of Alabama, Tuscaloosa, AL 35487-0336 ²Dave C. Swalm School of Chemical Engineering, Mississippi State University, Starkville, MS 39762

There is considerable interest in the development of Advanced Oxidation Processes, especially those that use wide band gap semiconductors for photo-oxidation of pollutants in waste water streams. The photo-oxidation is a surface process and so the greatest photo-activity is displayed by colloidal particles. Unfortunately the particles do not settle from aqueous dispersions and so, to avoid a costly separations step, it is desirable to develop a supported form of the photo-catalyst.

We have encapsulated Degussa P-25 titanium dioxide into a thin film of mesoporous silica as a means of confining the photo-catalyst while retaining maximum access to the photo-catalyst for the aqueous solutes. Physical characterization of the medium was carried out by powder X-Ray Diffraction supplemented by Variable Angle Spectroscopic Ellipsometry. Chemical characterization focused on the photodecomposition of 2,4-dichlorophenol using 350 nm light. The photodegradation was monitored by both UV-vis and HPLC.

Tests showed that in the absence of a polymer template the silica/titanium dioxide film formed was not mesoporous. Such films were found to have very low photoactivity. On a per photon absorbed basis the mesoporous silica/titanium dioxide films were approximately 2/3rds as efficient as an aqueous dispersion of the same amount of P-25. The mesoporous silica has an hexagonal arrangement of 7 nm diameter pores. Experiments with other mesoporous structures; cubic and lamellar phases, showed much lower photoactivity, comparable to that of non-templated silica/P-25 composites.

Duty cycle experiments show that the photoactivity decreases for the first 50 hours of irradiation time, but then becomes steady at approximately 80% of the initial efficiency.