

## TiO<sub>2</sub>-Assisted photocatalysis of lead-EDTA

Vohra, M.S. , Davis, A.P.

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**Abstract:** Several industrial and remedial processes create aqueous wastes containing complexed heavy metal species. TiO<sub>2</sub>-assisted photocatalysis was evaluated to address contamination problems from Pb-EDTA at both stoichiometric and non-stoichiometric Pb(II)/EDTA ratios. Pb-EDTA destruction occurred rapidly. At higher pH, lead became adsorbed onto the TiO<sub>2</sub> as complexed intermediates. Continued PCO saw reappearance of the lead in solution, primarily as Pb<sup>2+</sup>. Stoichiometric Pb-EDTA studies showed no noticeable pH effects even though Pb-EDTA adsorption depends strongly on pH. This suggests that initial complex adsorption is not required for its destruction and that Pb-EDTA photocatalysis occurs both at the TiO<sub>2</sub> surface and in the aqueous phase. Production of CO<sub>2</sub> and formaldehyde during photocatalysis suggests initial attack at an acetate group on the complex. Acetate and formate were found as minor intermediates. Nitrogen-containing products were ammonium and nitrate. For EDTA > Pb(II) studies, smaller amounts of CO<sub>2</sub> compared with stoichiometric Pb-EDTA were found, suggesting that attack on the acetate group is promoted by lead complexation. Pb(II) > EDTA studies showed decreasing CO<sub>2</sub> production with increasing pH, resulting from a shift in adsorbed Pb-EDTA speciation, or increased adsorption of free lead at higher pH, decreasing the available surface sites for OH<sup>·</sup> production.