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1.1 These Conditions alone shall govern and be incorporated in every contract for the sale of Goods made by or on behalf of the Supplier with any Customer. Jun 5, 2013 · 5 reasons why android tablets cant replace laptops... URL: of 2,4-dichlorophenoxyacetic acid and 2,4,5-trichlorophenoxyacetic acid in water using TiO<sub>2</sub>, Fe(IV) + MnO<sub>4</sub><sup>-</sup>, and OH radical generating photocatalysts. Two photosensitizer/catalyst systems were tested for the photolysis of 2,4-dichlorophenoxyacetic acid (2,4-D) and 2,4,5-trichlorophenoxyacetic acid (2,4,5-T) at pH 4.35. An aqueous suspension of anatase/rutile-TiO(2) (7:3), which had a pseudo-first order rate constant ( $k_{app}$ ) of 0.24 day<sup>-1</sup>, was tested in the presence of three different catalysts, viz., MnO(4)(-) ( $k_{app}$ )=0.0358 day<sup>-1</sup>), Fe(IV)-H(2)O-MnO(4)(-) ( $k_{app}$ )=0.0134 day<sup>-1</sup>) and OH ( $k_{app}$ )=0.019 day<sup>-1</sup>) [with 4.0% TiO(2)]. Addition of the three catalysts increased the photolysis rate of 2,4-D (30-fold) and 2,4,5-T (2.5-fold). The rate enhancement was strongly dependent on the catalyst concentration (0.04-0.40mM) and pH value (6.4-4.35). The presence of different TOC in the suspensions did not influence the activity of the three catalysts. The relative order of activity was MnO(4)(-)>Fe(IV)-H(2)O-MnO(4)(-)>OH. The order obtained was similar to that observed previously in the reactivity of dioxins and furans with TiO(2). A three-compartment model was used to determine the kinetics of the photode