

# Commercial TiO<sub>2</sub> mixtures for photocatalytic hydrogen production

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**1. Introduction** – The work on water-photo-splitting of Fujishima and Honda in 1972 [1] inspired subsequent researches to obtain hydrogen in eco-friendly way. The heterogeneous photocatalysis using TiO<sub>2</sub> can be an alternative [2]. If Pt nanoparticles are added to the TiO<sub>2</sub> surface, hydrogen production is highly favored [3] with respect to that from unmodified TiO<sub>2</sub>. In this work, we used mixtures of the most productive commercial photocatalysts (Aeroxide P25 and Kronos) with 1% wt. photo-deposited Pt with in order to study if there is a synergistic effect between both catalysts.

**2. Experimental** - Mixtures of Kronos and P25 catalysts were prepared with different weight percentages (80-20, 60-40, 40-60 and 20-80). A borosilicate reactor was then filled with [methanol] 6.17 M, [photocatalyst] 1 g·L<sup>-1</sup> and the calculated amount of hexachloroplatinic acid to obtain 1% wt Pt loading. The generated hydrogen was carried by a He continuous flow at 20 mL·min<sup>-1</sup> to a gas chromatographer to be quantified. The radiation was supplied by a fluorescent emitting between 200 and 400 nm with a maximum at 365 nm.

**3. Results and Discussion** - The mixtures were characterized by XRD, SEM-EDX and DRS. The XRD showed that the Pt deposition process did not modify the crystalline structure. The SEM-EDX studies confirmed that the amount of deposited Pt corresponded to that theoretically calculated. The DRS spectra showed a shift to a larger wavelength compared to that of bare catalysts. The most productive photocatalyst was pure P25 (3767 μmol·h<sup>-1</sup>). The mixtures exhibited slightly lower maximum productions and these were between those of the bare catalysts. However, maximum production was reached faster for 40-60 and 60-40 mixtures. The intermediates detected were, in gas phase, CH<sub>4</sub> and CO<sub>2</sub> and in the liquid phase HCHO and HCOOH.

**4. Conclusions** – The presence of Kronos in a 40 and 60% wt. decreased significantly the time necessary to reach the *plateau* of maximum production.

## 5. References

- [1] A. Fujishima, K. Honda, Electrochemical photolysis of water at a semiconductor electrode, *Nature* 238 (1972) 37-38.
- [2] E. Pulido Melián et al., Efficient and affordable hydrogen production by water photo-splitting using TiO<sub>2</sub>-based photocatalysts, *International Journal of Hydrogen Energy* 38 (2013) 2144-2155.
- [3] E. Pulido Melián et al., Hydrogen production using Pt-loaded TiO<sub>2</sub> photocatalysts, *International Journal of Hydrogen Energy* 38 (2013) 11737-11748.