

# PRISTINE AND Pt-MODIFIED TiO<sub>2</sub> DRIVEN ORGANIC COMPOUNDS PHOTODEGRADATION

Alexandra ILIE<sup>1</sup>, Crina ANASTASESCU<sup>1</sup>, Luminita PREDOANA<sup>1</sup>, Adriana RUSU<sup>1</sup>, Silviu PREDA<sup>1</sup>,

Daniela C. CULITA<sup>1</sup>, Veronica BRATAN<sup>1</sup>, Valentin Adrian MARALOIU<sup>2</sup>, Valentin S. TEODORESCU<sup>2,3</sup>, Ioan BALINT<sup>1</sup> and Maria ZAHARESCU<sup>1</sup>

<sup>1</sup> „Ilie Murgulescu” Institute of Physical Chemistry of the Romanian Academy, 202 Splaiul Independentei, 060021 Bucharest, 6<sup>th</sup> District, Romania

<sup>2</sup> National Institute of Materials Physics, 405 bis Atomistilor Street, 077125 Magurele-Ilfov, Romania

<sup>3</sup> Academy of Romanian Scientists, 3 Ilfov, 050044 Bucharest, Romania

## INTRODUCTION & AIM

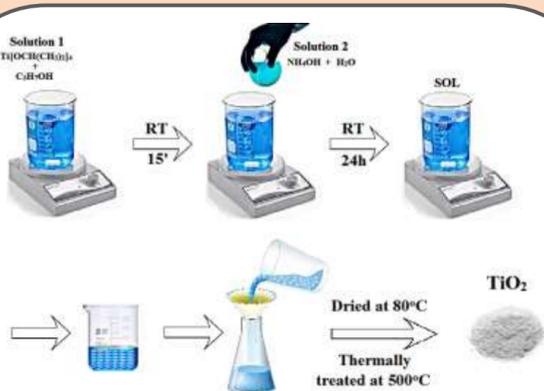
Photocatalysts powders with different compositions, sizes and dimensional distributions were prepared by sol-gel method.

The purpose of this work was to prepare, by the sol-gel method, TiO<sub>2</sub> and Pt-modified TiO<sub>2</sub> with photocatalytic activity for ethanol degradation in gaseous phase under simulated solar light irradiation.

## METHOD

The powders were obtained by sol-gel method and noted: TiO<sub>2</sub> and TiO<sub>2</sub>-Pt (doped by post-synthesis impregnation). The methods used for characterization were infrared spectroscopy (FT-IR), transmission electron microscopy (TEM), X-ray diffraction (XRD), X-ray fluorescence(XRF), UV-Vis and BET-specific surface area determination. The products of oxidative degradation process were analyzed by gas-phase chromatography (GC-TCD and GC-FID).

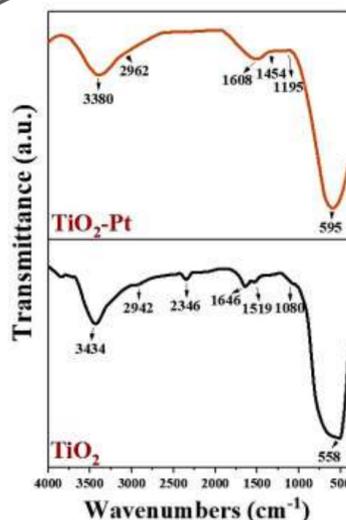
## SYNTHESIS



**Fig. 1.** TiO<sub>2</sub> preparation by the sol-gel method.

- Reagents: Titanium isopropoxide for TiO<sub>2</sub> source and PtCl<sub>4</sub> for dopant (1 mol%).
- The samples obtained were white (TiO<sub>2</sub>) or light grey (TiO<sub>2</sub>-Pt) powders.

## RESULTS & DISCUSSION



**Fig. 2.** IR spectra of thermally treated powders.

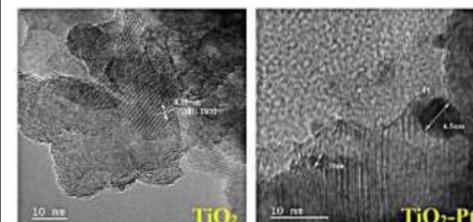
- The small displacement to higher wavelength values in case of TiO<sub>2</sub>-Pt is assigned to the presence of noble metal in the composition of the investigated sample.

## FT-IR

**Tab. 1.** Assignment of vibration bands of thermally treated powders.

Wavenumbers (cm <sup>-1</sup> )	Assignments and vibration bands	
TiO <sub>2</sub>	TiO <sub>2</sub> -Pt	
558	595	Ti-O stretching vibration of TiO <sub>2</sub>
1080	-	C-O
-	1195	-
-	1454	$\nu_s(\text{CO}_2)^2$
1519	-	$\nu_{as}(\text{CO}_2)^2$
1646	1608	$\delta\text{H}_2\text{O}$ (adsorbed water)
2346	-	CO <sub>2</sub> (absorption)
2942	2962	C-H, $\nu_{as}(\text{CH}_2)$
3434	3380	$\nu\text{OH}$ (structural OH group)

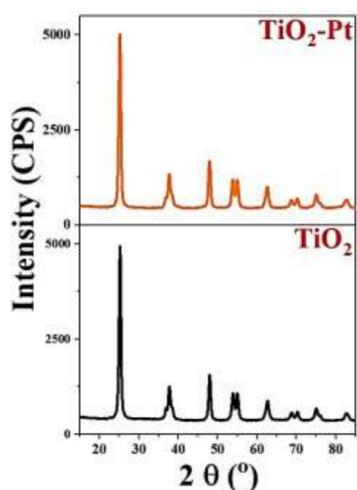
## TEM



**Fig. 3.** TEM micrographs of the samples.

- The TiO<sub>2</sub> sample is formed by aggregates crystallites with dimensions of 20-60 nm.
- TiO<sub>2</sub>-Pt present Pt nanoparticles are spherical and have dimensions from 2.5 nm to 6 or 7 nm.

## XRD



**Fig. 4.** XRD patterns of the samples.

- Both samples exhibit a single-phase anatase TiO<sub>2</sub> structure.
- Addition of Pt does not significantly alter the anatase phase composition.

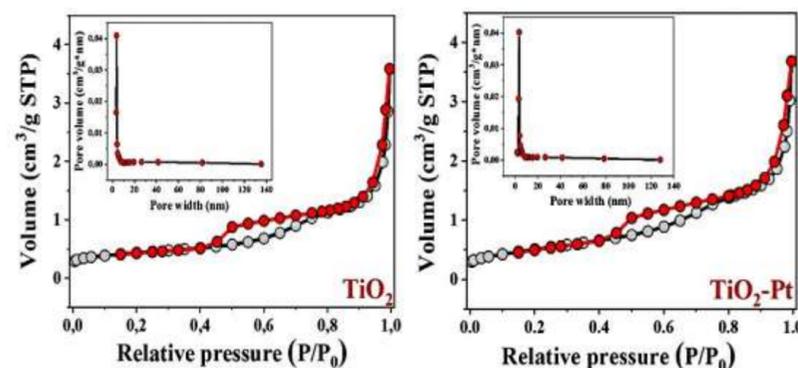
## XRF

**Tab. 2.** Elemental composition of the powders.

Sample	Composition	Values (Mass %)	Line
TiO <sub>2</sub>	Ti	59.25	Ti-KA
	O	39.57	O-KA
	Traces	1.18	
TiO <sub>2</sub> -Pt	Ti	59.32	Ti-KA
	O	38.25	O-KA
	Pt	1.05	Pt-LA
	Traces	1.38	

- The analysis confirmed the presence of platinum (Pt) in the Pt-doped sample in quantities closely matching the initially calculated composition, indicating successful incorporation of Pt into the material.

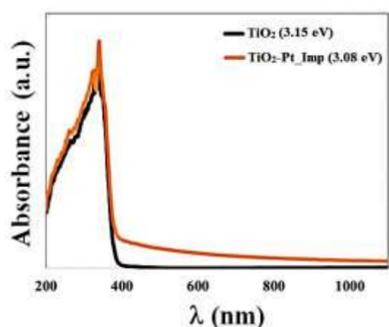
## BET



**Fig. 5.** N<sub>2</sub> adsorption-desorption isotherms and pore size distributions (insert in the figures) of powders.

- Both samples exhibit type IV(a) isotherms which is characteristic of mesoporous materials.

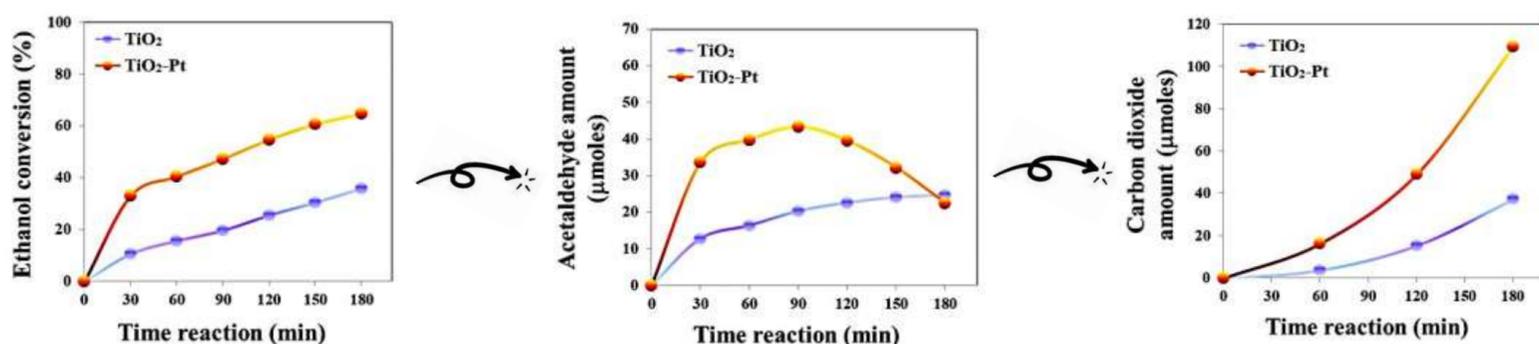
## UV-Vis



**Fig. 6.** UV-Vis absorption spectra.

- The samples present a strong absorption in the UV region, with a sharp peak at ~340 nm, characteristic to TiO<sub>2</sub>.

## Oxidative photodegradation of ethanol in gaseous phase under solar simulated light



- Oxidative degradation of ethanol leads to acetaldehyde (CH<sub>3</sub>CHO) and then, through successive reactions, to the intermediate formation of formic acid (HCOOH - in trace amounts) and finally to CO<sub>2</sub>.

## CONCLUSIONS

- TiO<sub>2</sub> powder were obtained using the sol-gel method, and doping with Pt by post-synthesis impregnation.
- The structural and morphological characterizations of the obtained photoactive materials were correlated with their photocatalytic activity.
- After 3 hours of solar simulated light irradiation, TiO<sub>2</sub>-Pt powder achieved the highest conversion of ethanol photodegradation (64.82%).
- Addition of Pt improves the catalytic activity by: co-catalytic effect, decrease of bandgap, better light absorption and separation of charges.