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*A. Hussain*  
PRINCIPAL  
Sree Narayana Gurukulam  
College of Engineering  
Kadayiruppu, Kolenchery-682 311

DEPARTMENT OF CIVIL ENGINEERING

**Sree Narayana Gurukulam College of Engineering**  
Kadayiruppu, Kolenchery, Kerala, India



Proceedings of International Conference on Sustainable Construction Materials (ICSCM-23)



*A. Hussain*

PRINCIPAL  
Sree Narayana Gurukulam  
College of Engineering  
Kadayiruppu, Kolenchery-682 311



Sree Narayana Gurukulam College of Engineering Kadayiruppu, Kolenchery, Kerala, India

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# Mineralization of a Model Pollutant over Platinum Loaded Nanotitania Photocatalyst under Sunlight

Dr. Suprabha T<sup>1</sup>, Dr. Preetha Varghese<sup>2</sup>

<sup>1</sup>Science and Humanities Department, Sree Narayana Gurukulam College of Engineering, Kadayiruppu, Ernakulam 682311

<sup>2</sup>Science and Humanities Department, Sree Narayana Gurukulam College of Engineering, Kadayiruppu, Ernakulam 682311

<sup>1</sup>[suprabha@sngce.ac.in](mailto:suprabha@sngce.ac.in)

<sup>2</sup>[preethavarghese@sngce.ac.in](mailto:preethavarghese@sngce.ac.in)

**Abstract**—Mineralization of a model pollutant, methylene blue over bare and platinum loaded titania nanostructures was conducted. Nanotitania with three different morphologies were synthesized by microwave irradiation technique. Platinum particles were loaded on nanotitania. Synthesized nanostructures were characterized. The dye degradation capabilities of Pt/TiO<sub>2</sub> were found superior to bare titania nanostructures.

**Keywords**— Photocatalysis, methylene blue, nanotitania, Pt/TiO<sub>2</sub>, microwave technique

## 1. INTRODUCTION

Microwave synthesis and photocatalysis have played pivotal roles in recent years toward the goal of switching to increasingly efficient and benign processes (Balachandran, Dar, Yang and Zheng). As a most widely used photocatalyst, TiO<sub>2</sub> has a relatively large band gap of 3.2 eV and TiO<sub>2</sub> has to be activated by a UV light source. The general approach is to modify TiO<sub>2</sub> by creating intra-band gap states that are close to the conduction or valence band edges, and absorb visible light at sub-band gap energies of less than 3.2 eV (Kisch).

We have already reported a simple microwave method to synthesize phase pure anatase and rutile nanotitania with different morphologies viz., cubes, spheres and rods (Suprabha). In an attempt to modify the optical properties of titania synthesized by microwave irradiation we were successful in loading Au and Ag into titania structure which shifted the absorption into visible region (Suprabha). In the present work, we could enhance the visible light photocatalysis of titania nanostructures by loading platinum by metal sol method (Grunwaldt). To the best of our knowledge no

reports are available with regard to the preparation of Pt loaded TiO<sub>2</sub> using TiO<sub>2</sub> synthesized by microwave irradiation. The catalytic performances of three morphologically different nanotitania and their platinum loaded nanostructures (Pt/TiO<sub>2</sub>) were studied for dye degradation of methylene blue under sunlight.

## 2. MATERIALS AND METHOD

All reagents were purchased from Merck, Germany. Titanium trichloride (15 wt. % TiCl<sub>3</sub>, 10 wt. % HCl) was used as the titanium precursor. NH<sub>4</sub>OH (1.5 M), NaCl (5 M), NH<sub>4</sub>Cl (5 M), hexachloroplatinic acid (H<sub>2</sub>PtCl<sub>6</sub>), NaBH<sub>4</sub>, and polyvinyl alcohol (PVA) were employed for the synthesis. A typical microwave oven (Whirlpool, 1200 W) operating at a frequency of 2450 MHz was used for the synthesis.

The three TiO<sub>2</sub> nanostructures with different morphologies viz., cube, sphere and rod (S1, S2, S3) synthesized as per our early report (Suprabha), were loaded with 1 wt% Pt employing metal sol method. For this a suitable amount of PVA solution (1wt %) was added to an aqueous hexachloroplatinic acid solution (0.1 mg Pt/ml) under vigorous stirring. Then, a freshly prepared solution of NaBH<sub>4</sub> (molar ratio NaBH<sub>4</sub>/ Pt = 4) was slowly added dropwise into the mixture. The TiO<sub>2</sub> samples were added after the pH of Pt sol was adjusted to 6.0 by NH<sub>4</sub>OH or HCl solution. The mixture was stirred at 60°C for 3 hrs, then filtered and washed until no Cl<sup>-</sup> was detected by AgNO<sub>3</sub> solution. After being dried at 100°C overnight, the samples were calcined at 550°C for 4 hrs. The platinum loaded on samples S1, S2 and S3 are represented as PtS1, PtS2 and PtS3 respectively.

Photocatalytic activity of TiO<sub>2</sub> was evaluated by the degradation of the dye, methylene blue (MB) in



*A. Anwar*

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Sree Narayana Gurukulam  
College of Engineering  
Kadayiruppu, Kolenchery - 682 311

ISBN 978-81-956872-5-1