

Photocatalytic Degradation of Carbon Monoxide (CO) Using Bi₂O₃ Modified TiO₂ Photocatalyst

Nusaiba Islam¹, Sharmin Jahan Prama¹, Ashiqur Rahman^{1,2*}
and Ashok Kumar Chakraborty¹

¹Nanomaterials Laboratory, Department of Applied Chemistry and Chemical Engineering, Islamic University, Kushtia-7003, Bangladesh.

²Department of Chemical Engineering, Faculty of Engineering and Built Environment, School of Engineering, The University of Newcastle, Callaghan, New South Wales 2308, Australia.

Authors' contributions

This work was carried out in collaboration between all authors. Authors NI and AR designed the study, performed the statistical analysis, wrote the protocol and wrote the first draft of the manuscript. Authors AR and SJP managed the analyses of the study. Authors AR and AKC managed the literature searches. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/IRJPAC/2017/34828

Editor(s):

(1) Francisco Marquez-Linares, Full Professor of Chemistry, Nanomaterials Research Group, School of Science and Technology, University of Turabo, USA.

Reviewers:

(1) Ashish Kumar, Agra College, India.

(2) Anukorn Phuruangrat, Prince of Songkla University, Thailand.

(3) Tariq Mahmood, Quaid-i-Azam University, Pakistan.

Complete Peer review History: <http://www.sciencedomain.org/review-history/20011>

Original Research Article

Received 14th June 2017

Accepted 10th July 2017

Published 12th July 2017

ABSTRACT

TiO₂ modified Bi₂O₃ nanoheterojunction was prepared using maleic acid as an organic linker. TiO₂ photocatalyst cannot perform photocatalytic reaction in the visible region. It only can work in the UV light region because of its band gap 3.2 eV. On the other hand Bi₂O₃ modified TiO₂ can perform photo activity in UV as well as visible region because Bi₂O₃ can absorb photon in the visible light. The Bi₂O₃/TiO₂ nanocomposite was characterized Transmission Electron Microscope (TEM), X-ray diffraction (XRD), UV-Vis spectroscopy and Gas Chromatography (GC). The as modified composite exhibited high photo oxidation activity under visible region of λ up to 430 nm, whereof the modified photocatalyst can oxidize CO in UV as well as visible light. Because of the development of anti-air pollution technology, our modified nanoheterojunction could be a significant photocatalyst to oxidize pollutants in air to keep the environment clean.

*Corresponding author: E-mail: ashiq.chem@yahoo.com

Keywords: Photo oxidation; $\text{Bi}_2\text{O}_3/\text{TiO}_2$ nanocomposite; carbon monoxide (CO); air pollution.

1. INTRODUCTION

In recent years photocatalytic oxidation (PCO) is widely studied in the field of water and air pollution control to clean-up the environmental hazardous pollutants [1-5]. The research on oxidation of carbon monoxide (CO) has become an important topic during the last few years. CO is a harmful air pollutant for the living environment and generally it produce in partial combustion of hydrocarbons. Photocatalytic oxidation process is one of the most promising resolutions to decompose CO. Especially TiO_2 photocatalyst can successfully degrade the volatile organic compounds present in air under normal temperature [6-9]. However, the limitation is TiO_2 only can work under UV light because of its large band gap of 3.2 eV [10]. Therefore, modification of TiO_2 is mandatory to absorb visible light in the photocatalytic process to enhance the reaction rate.

There are different ways to modify TiO_2 photocatalyst. Among them doping by metal and non-metal, photosensitizing with dye on TiO_2 surface, modification by semiconductors having narrow bandgap and noble metal depositions are widely used. Of the many processes modification of TiO_2 using noble metal deposition on its surface have recently been placed as one of the most feasible modification process for enhancing the photocatalytic oxidation reactions [11-13].

In this research, TiO_2 was modified by bismuth (III) oxide (Bi_2O_3) to enhance the photocatalytic activity of TiO_2 . Maleic acid was used as an organic linker to develop the $\text{Bi}_2\text{O}_3/\text{TiO}_2$ nanoheterojunction. The presence of Bi_2O_3 on TiO_2 surface significantly enhances the photocatalytic oxidation rate of CO because the PCO reaction can possible in presence of UV as well as visible light [14]. On the other hand, research literatures have confirmed that Bi_2O_3 is comparatively cheap and less toxic than other metals like lead, iron and antimony [10,14].

2. EXPERIMENTAL

Carbon Monoxide (CO), titanium dioxide (TiO_2) nanoparticle, bismuth oxide (Bi_2O_3), maleic acid ($\text{C}_4\text{H}_4\text{O}_4$) and absolute ethanol ($\text{CH}_3\text{CH}_2\text{OH}$) were purchased from Sigma Aldrich (Germany) and were used without further purification.

2.1 Modification of TiO_2 with Bi_2O_3

Different molar ratios of Bi_2O_3 modified TiO_2 were prepared using maleic as an organic linker. 5/95, $\text{Bi}_2\text{O}_3/\text{TiO}_2$ was synthesized using 5 mol% of Bi_2O_3 and 95 mole% of TiO_2 . During synthesizing process 0.24 g of Bi_2O_3 was dissolved in 50 mL of ethanol solution. Then 0.2 g of maleic acid was added in to the solution. Afterward, 0.76 g of TiO_2 was added in to the mixture and stirred for 5 h at room temperature until a homogenous solution. Then the suspension was stand in an oven for overnight at 60°C for drying. To increase the bonding between TiO_2 and Bi_2O_3 the prepared composite was annealed at 120°C in a muffle furnace. The same methodology was used to synthesizing at ratios of 20/80, 35/65, 50/50, 65/35, $\text{Bi}_2\text{O}_3/\text{TiO}_2$ nanoheterojunction [14].

2.2 Photocatalytic Oxidation Test

Photocatalytic oxidation of CO was carried out in a gas reactor prepared in the laboratory. Gas chromatography (GC) was used to measure the remnant of CO after photocatalytic oxidation reaction. Our modified photocatalyst was placed inside the gas reactor and the light source was placed in front of the photocatalyst. The CO gas was inserted in to the gas reactor and after oxidation the remnant of CO was collected and was analyzed using GC.

2.3 Characterization Techniques

The surface structure of the pure and modified TiO_2 were observed using fei-TECNAI G2 Transmission Electron Microscopy (TEM). A D8 Bruker Advance X-ray diffractometer (Bruker, Germany) was used to analyze the X-ray diffraction measurements. A Shimadzu UV-1601 spectrophotometer was used to record UV-Vis reflectance spectra. To observe the photocatalytic oxidation rate of CO a Gas Chromatograph was used with the gas reactor.

3. RESULTS AND DISCUSSION

3.1 Crystallite Shape and Compositional Analysis

In Fig. 1 the surface morphology of 65/35, $\text{TiO}_2/\text{Bi}_2\text{O}_3$ nanocomposite was analyzed using transmission electron microscope (TEM) and high resolution electron microscopy (HR-TEM).

The crystallite structure of TiO_2 nanoparticles can be clearly noticed from Fig. 1(a-b). In Fig. 1(c-d) the modification of TiO_2 using Bi_2O_3 was observed. Thus, the TEM analysis asserts that the Bi_2O_3 was finely distributed on the TiO_2 surface and no other unwanted particles observed.

The Energy-dispersive X-ray spectroscopy (EDX) elemental analysis was performed on Bi_2O_3 modified TiO_2 nanocomposite in a weight ratio of $\text{TiO}_2:\text{Bi}_2\text{O}_3$; 65:35 which shown in Table 1. From the analysis, it is clear that at the end of the modification process the rest of the maleic acid was evaporated from the nanocomposite. Only Ti, Bi and O elements were found from the analysis that represent the presence of TiO_2 and Bi_2O_3 in the nanocomposite.

Table 1. EDX data of Bi_2O_3 modified TiO_2 nanocomposite

| Elements | Bi_2O_3 modified TiO_2 nanocomposite Wt.% |
|----------|--|
| Ti | 41.56 |
| Bi | 4.08 |
| O | 54.36 |

(Weight ratio of $\text{TiO}_2:\text{Bi}_2\text{O}_3$; 65:35)

3.2 XRD Analysis

The XRD patterns of TiO_2 , Bi_2O_3 and Bi_2O_3 modified TiO_2 nanocomposite is presented in Fig. 2. The TiO_2 nanoparticle curve showed

peaks at 25.3° , 37.8° , 54° , and 62.10° , which indicate the present of the anatase phase and at 43.90° , 56.90° , and 65.10° , exhibited the rutile phase. The Bi_2O_3 curve also exhibited some diffraction peaks at 21.90° , 26.90° , 32.80° , 47.19° , and 55.20° . The XRD patterns of Bi_2O_3 modified TiO_2 nanocomposite equalized the diffraction peaks of TiO_2 and Bi_2O_3 without added of any other compounds phases. This observation declares that in the composite formation process no other chemical reactions occur between Bi_2O_3 and TiO_2 nanoparticle.

3.3 UV-Vis Analysis

The UV-vis absorption spectra of TiO_2 , Bi_2O_3 and the different ratios of $\text{Bi}_2\text{O}_3/\text{TiO}_2$ nanocomposite are illustrated in Fig. 3. The bandgap of TiO_2 is wide than Bi_2O_3 and were investigated to be 3.2 eV and 2.8 eV, respectively [15,16]. The optical absorptions of Bi_2O_3 is greater than of 400 nm and for that reason Bi_2O_3 can absorb the photon from the visible light of the solar spectrum. TiO_2 optical absorption is less than 385 nm therefore, TiO_2 is photo inactive in the visible light. On the other hand, the optical absorption of Bi_2O_3 modified TiO_2 nanocomposites are greater than 400 nm because of the presence of Bi_2O_3 . The photon absorption rate in the visible light increase with the increasing of the molar percentage of Bi_2O_3 in the composite. This analysis demonstrate that $\text{Bi}_2\text{O}_3/\text{TiO}_2$ nanoheterojunction is effective for the photon absorption in the visible light.

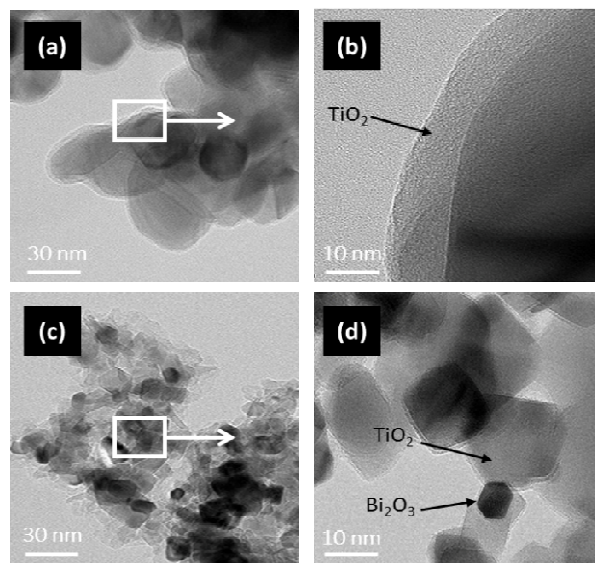


Fig. 1. TEM images of (a, b) pure TiO_2 nanoparticle (c, d) Bi_2O_3 modified TiO_2 nanocomposite

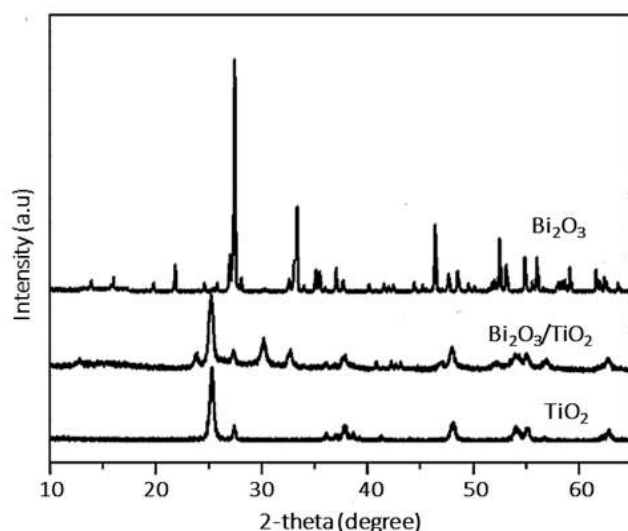


Fig. 2. XRD analysis of TiO_2 , Bi_2O_3 and Bi_2O_3 modified TiO_2 nanocomposite

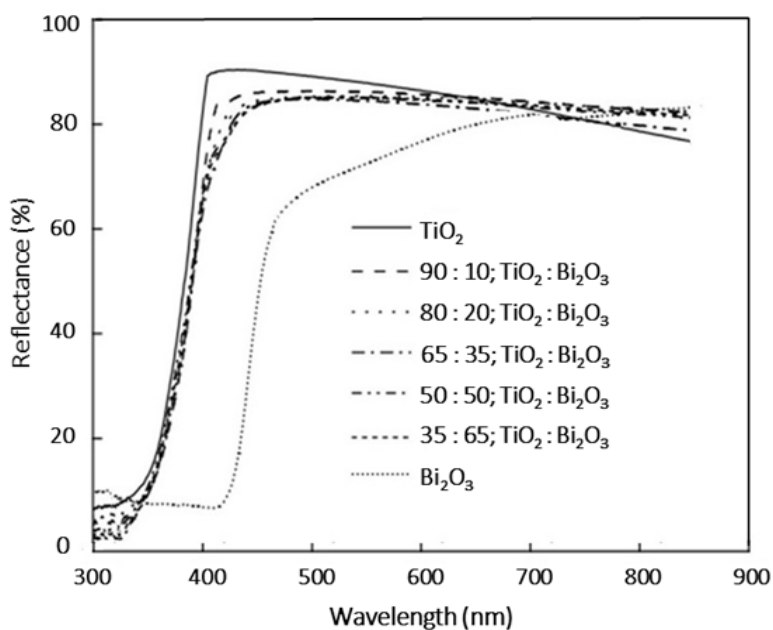


Fig. 3. UV-vis absorption spectra of TiO_2 , Bi_2O_3 and different ratios of Bi_2O_3 modified TiO_2 nanocomposite

3.4 Photocatalytic Oxidation of CO

Fig. 4 illustrate the photocatalytic oxidation of CO on pure TiO_2 and Bi_2O_3 modified TiO_2 under visible light. In case of pure TiO_2 no photo oxidation occurs because TiO_2 cannot absorb

photon in the visible light. The different ratios of $\text{Bi}_2\text{O}_3/\text{TiO}_2$ composites perform different photocatalytic activity. With increasing the percentage of Bi_2O_3 in the composite the rate of oxidation also increase but 65/35, $\text{TiO}_2/\text{Bi}_2\text{O}_3$ exhibited highest photo oxidation of CO.

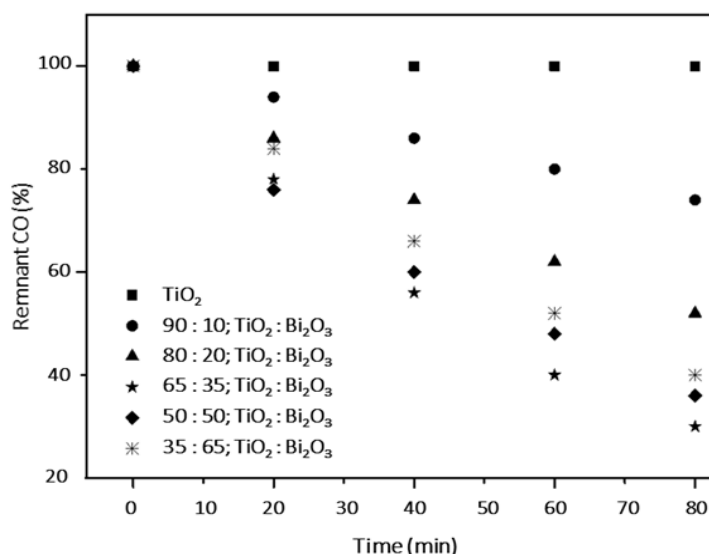


Fig. 4. Remnant CO after photocatalytic oxidation of CO on TiO₂, and different ratios of Bi₂O₃ modified TiO₂ nanocomposites

4. CONCLUSION

Bi₂O₃ modified TiO₂ photocatalyst was prepared using maleic acid as an organic linker. The Bi₂O₃ has finely distributed on the TiO₂ nanoheterojunction. The Bi₂O₃/TiO₂ composite successfully performed photocatalytic oxidation of CO in the visible light region. The role of Bi₂O₃ in the composite was to absorb the photon from the visible region and to be active of the Bi₂O₃/TiO₂ nanoheterojunction for the photo oxidation of CO in the UV as well as visible light.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- Hoffmann MR, Martin ST, Choi W, Bahnemann DW. Environmental applications of semiconductor photocatalysis. *Chemical Reviews*. 1995;95(1):69-96.
- Choi W, Hong SJ, Chang YS, Cho Y. Photocatalytic degradation of polychlorinated dibenzo-p-dioxins on TiO₂ film under UV or solar light irradiation. *Environmental Science & Technology*. 2000;34(22):4810-5.
- Cho Y, Choi W, Lee CH, Hyeon T, Lee HI. Visible light-induced degradation of carbon tetrachloride on dye-sensitized TiO₂. *Environmental Science & Technology*. 2001;35(5):966-70.
- Kim S, Choi W. Kinetics and mechanisms of photocatalytic degradation of (CH₃)_nNH₄ⁿ⁺ (0 ≤ n ≤ 4) in TiO₂ suspension: The role of OH radicals. *Environmental Science & Technology*. 2002;36(9):2019-25.
- Lee H, Choi W. Photocatalytic oxidation of arsenite in TiO₂ suspension: Kinetics and mechanisms. *Environmental Science & Technology*. 2002;36(17):3872-8.
- Driessen MD, Goodman AL, Miller TM, Zaharias GA, Grassian VH. Gas-phase photooxidation of trichloroethylene on TiO₂ and ZnO: Influence of trichloroethylene pressure, oxygen pressure, and the photocatalyst surface on the product distribution. *The Journal of Physical Chemistry B*. 1998;102(3):549-56.
- Cho Y, Yu Y, Lim J, Park J. Effect of Cu and Co addition on CO gas-sensing properties of TiO₂ prepared by oxidation of mechanically-synthesized TiN composites. *Journal of Ceramic Processing Research*. 2016;17(8):825-30.

8. Peral J, Domenech X, Ollis DF. Heterogeneous photocatalysis for purification, decontamination and deodorization of air. *Journal of Chemical Technology and Biotechnology*. 1997; 70(2):117-40.
9. Choi W, Ko JY, Park H, Chung JS. Investigation on TiO₂-coated optical fibers for gas-phase photocatalytic oxidation of acetone. *Applied Catalysis B: Environmental*. 2001;31(3):209-20.
10. Sobana N, Muruganadham M, Swaminathan M. Nano-Ag particles doped TiO₂ for efficient photodegradation of direct azo dyes. *Journal of Molecular Catalysis A: Chemical*. 2006;258(1):124-32.
11. Hufschmidt D, Bahnemann D, Testa JJ, Emilio CA, Litter MI. Enhancement of the photocatalytic activity of various TiO₂ materials by platinisation. *Journal of Photochemistry and Photobiology A: Chemistry*. 2002;148(1):223-31.
12. Dobosz A, Sobczyński A. The influence of silver additives on titania photoactivity in the photooxidation of phenol. *Water Research*. 2003;37(7):1489-96.
13. Arana J, Dona-Rodríguez JM, González-Díaz O, Rendon ET, Melián JH, Colon G, Navio JA, Peña JP. Gas-phase ethanol photocatalytic degradation study with TiO₂ doped with Fe, Pd and Cu. *Journal of Molecular Catalysis A: Chemical*. 2004;215(1):153-60.
14. Rahman A, Ching YC, Ching KY, Awanis N, Chakraborty AK, Chuah CH, Liou NS. Surface modification of natural fiber using Bi₂O₃/TiO₂ composite for photocatalytic self-cleaning. *BioResources*. 2015;10(4): 7405-18.
15. Hameed A, Montini T, Gombac V, Fornasiero P. Surface phases and photocatalytic activity correlation of Bi₂O₃/Bi₂O_{4-x} nanocomposite. *Journal of the American Chemical Society*. 2008;130(30):9658-9.
16. Xu Y, Schoonen MA. The absolute energy positions of conduction and valence bands of selected semiconducting minerals. *American Mineralogist*. 2000;85(3-4):543-56.

© 2017 Islam et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:

*The peer review history for this paper can be accessed here:
<http://sciencedomain.org/review-history/20011>*