









PREPARATION OF S-DOPED TIO₂ VIA SOL-GEL/ HYDROTHERMAL METHOD AND ITS ACTIVITY AS PHOTODEGRADATION OF RB05

Adiansyah^{1,*}, I. Elisabeth Purba², M. Tarigan¹, Lisnadiyanti³, Yusnaidar⁴ and I.P. Mahendra⁵

¹Department of Chemistry, Universitas Sari Mutiara, Indonesia ²Department of Public Health, Universitas Sari Mutiara, Indonesia ³Faculty of Nursing and Midwifery, Universitas Binawan, Indonesia ⁴Department of Chemistry Education, Faculty of Teacher Training and Education,

Universitas Jambi, Indonesia

⁵Department of Chemistry, Faculty of Mathematics and Sciences, Universitas Sumatera Utara *E-mail: adiansyah_skd@yahoo.co.id

ABSTRACT

S-doped TiO₂ photocatalyst has been prepared by the hydrothermal method at 180° C that assisted by the sol-gelel method and its photocatalytic activity was evaluated to reactive black 05. The obtained photocatalyst was characterized by XRD and SEM. The phase composition of S-doped TiO₂ was influenced by the amount of sulfur. **Keywords:** TiO₂, Doping process, Sulfur, Photocatalyst, Reactive Black 05.

© RASĀYAN. All rights reserved

INTRODUCTION

 TiO_2 is classified as semiconductor materials that have excellent property as a catalyst, especially in the photodegradation of organic pollutants, i.e. dyes, phenolic compounds, etc. The previous researches that focused on the photoactivity of TiO_2 , mostly work under UV light due to it shows high activity and stable under UV light.¹⁻³

The issues that were found during its applications, photoactivity of TiO_2 is not fully effective when it utilized under solar light. Several methods have been developed to improve its performance under solar light.⁴⁻⁶ One of these efforts is through the doping technique. The doping technique also influences the bandgap of the doped materials with a range in the visible light area. As the dopant, either metal or non-metal elements can be incorporated with the TiO_2 , it depends to the atomic radius between dopant and oxygen atom of TiO_2 .⁷⁻¹⁰ The previous work about the doping process of TiO_2 has reported results on the substitution of several elements, i.e. nitrogen, oxygen, and sulfur.^{10,11} The incorporating of those elements into TiO_2 showed a significant change in the photoactivity of materials, especially sulfur.

S-doped TiO₂ in the previous work has been prepared using calcination technique, sol-gel, hydrothermal and in the supercritical condition in CS₂/ethanol.^{12,13} From those several methods, the combination method between sol-gel and hydrothermal provides a simple and best method to obtain nanosized of doped TiO₂ photocatalyst. Also, the photocatalyst of TiO₂ that prepared using the hydrothermal method can have a different ratio between retail and amateurs depend on the preparation of raw material and temperature during hydrothermal. Based on the literature review, there have been few reports on the preparation of S-doped TiO₂ using a hydrothermal method that assisted by the sol-gel method. In the present work, the S-doped TiO₂ was prepared using a hydrothermal method that assisted by the sol-gel method. The photoactivity of the obtained S-doped TiO₂ was evaluated for degradation of reactive black 05.

Rasayan J. Chem., 13(1), xxxx-xxxx(2020) http://dx.doi.org/10.31788/RJC.2020.1315461



EXPERIMENTAL

Materials

Sulfur and TTIP (97%) were obtained from Merck and Sigma Aldrich, respectively. Absolute ethanol was provided by Guangdong Guanghua Sci-Tech Co. Ltd.

Preparation of S-doped TiO₂

The doping process was performed using the method that has been described previously.¹⁴ The asprepared S-doped TiO2 then was characterized using several instruments, i.e Bruker D8 Advance X-ray diffractometer for crystallography analysis, scanning electron microscopy TM4000 Hitachi for morphological analysis and UV-Vis Diffuse Reflectance Spectrophotometry Shimadzu UV-2600 Series for determine the band gap value.

Photoactivity Measurements

The photoactivity of S-doped TiO_2 was performed in a cylindrical glass that equipped with a condenser to control the temperature. The treatment was performed using a method that has been described as previously¹⁵. The obtained aliquots were characterized using UV-Vis spectrophotometer to determine the final concentration of imitated wastewater after treatment.

RESULTS AND DISCUSSION

The crystallographic of doped TiO_2 is shown in Fig-1. The obtained S-doped TiO_2 with the incorporation of 0.5% S only consisted of anatase phase and rutile, no other phase can be found. However, when the amount of incorporated S increases the phase of TiO_2 was dominated by anatase. This result indicated the ratio of anatase, or rutile phase depends on the amount of S that doped to TiO_2 . The phase composition of TiO_2 was obtained using Match! software version 3.8.0.137 and the 0.5% S-doped TiO_2 showed a combination signal of JCPDS 96-900-1681 and JCPDS 96-720-6076, while 1.0% S-doped TiO_2 has similar signal to JCPDS 96-720-6076.



Figure-2 shows the SEM image of doped TiO_2 obtained by hydrothermal assisted by sol-gel method. The effect of the amount of S that doped to TiO_2 can be seen in the particle size of the photocatalyst. The particle size of TiO_2 was reduced by the increase of the amount of S, the particle size of the TiO2. The nanoparticle of TiO_2 with particle size in the range $> 1 \ \mu m$ was changed to 40-60 nm when the concentration of S increases.

The photodegradation of RB05 was evaluated by determining the concentration of dyes in the system at a specific time. In Fig-3, the photodegradation of RB05 by amateurs, 0.5% S-doped TiO₂ and 1.0% S-doped

PREPARATION OF S-DOPED TIO2

TiO₂ were 83.10; 20.73 and 98.31%, respectively. The as-prepared 1.0% S-doped TiO₂ showed to be more photoactive than undoped TiO₂ with a rate constant of 4.03×10^{-2} . The 0.5% S-doped exhibited the lowest photoactivity with a rate constant of 2.24×10^{-3} .



Fig.-2 The SEM Images of (a) 0.5% S-doped TiO₂ and (b) 1.0% S-doped TiO₂



Fig.-3 (a) Photocatalytic Activity of doped TiO₂ and (b) Rate of Photocatalytic Activity of doped TiO₂

The process of dyes photodegradation could be said as a complicated process that cannot be only viewed from one perspective, but it should be observed from all perspectives, i.e. particle size, morphology, and phase/ chemical composition of photocatalysts^{16–20}. In this study, the photoactivity of doped TiO₂ was influenced by two factors, i.e. the particle size of doped TiO₂ and phase composition of doped TiO₂. Based on kinetic theory, particle size has a big influence on the reaction rate. Smaller particle size will have a larger surface area and it can facilitate the dyes photodegradation be more effective as the impact of the increase of photogenerated electrons. The other perspectives, the phase composition of doped TiO₂has an important role on the dye photodegradation process. The ratio of rutile phase in the 0.5% S-doped TiO₂ was 88%, the high rutile phase content in S-doped TiO₂ phase was dominated by anatase and improved the photocatalytic activity. This result also supported by the previous work, the ratio of rutile and anatase phase has an important role to the photocatalytic activity, when the rutile ratio is high the photocatalyst will show lower activity and vice versa.^{4,16,21,22}

CONCLUSION

S-doped TiO_2 was successfully synthesized using hydrothermal assisted sol-gel method. The morphological and photocatalytic activity of doped TiO_2 was totally dependent on the S amount that incorporated into TiO_2 .

ACKNOWLEDGEMENT

This research was supported by Direktorat Riset dan Pengabdian Masyarakat Direktorat Jenderal Riset dan Pengembangan Kementerian Riset, Teknologi, dan Pendidikan Tinggi Penelitian through Hibah Dosen Pemula.

REFERENCES

- E. Carpio, P. Zúñiga, S. Ponce, J. Solis, J. Rodriguez, W. Estrada. J Mol Catal A Chem, 228, (2005), DOI: 10.1016/j.molcata.2004.09.066.
- 2 R.J. Tayade, P.K. Surolia, R.G. Kulkarni, R. V. Jasra. Sci Technol Adv Mater, 8, (2007), DOI: 10.1016/j.stam.2007.05.006.
- B. Liu, X. Wang, G. Cai, L. Wen, Y. Song, X. Zhao. J Hazard Mater, 169, (2009), DOI: 10.1016/j.jhazmat.2009.04.068.
- 4 H. Xu, L. Zhang. *J Phys Chem C*, **113**, (2009), DOI: 10.1021/jp8089903.
- 5 A. Ramchiary, S.K. Samdarshi. *Sol Energy Mater Sol Cells*, **134**, (2015), DOI: 10.1016/j.solmat.2014.12.031.
- 6 N.T. Nolan, D.W. Synnott, M.K. Seery, S.J. Hinder, A. Van Wassenhoven, S.C. Pillai. J Hazard Mater, 211–212, (2012), DOI: 10.1016/j.jhazmat.2011.08.074.
- 7 N.R. Mathews, M.A. Cortes Jacome, C. Angeles-Chavez, J.A. Toledo Antonio. J Mater Sci Mater Electron, 26, (2015), DOI: 10.1007/s10854-014-2294-3.
- 8 M. Bettinelli, V. Dallacasa, D. Falcomer, P. Fornasiero, V. Gombac, T. Montini, *et al. J Hazard Mater*, **146**, (2007), DOI: 10.1016/j.jhazmat.2007.04.053.
- 9 J. Wang, D.N. Tafen, J.P. Lewis, Z. Hong, A. Manivannan, M. Zhi, et al. J Am Chem Soc, 131, (2009), DOI: 10.1021/ja903781h.
- T. Ohno, M. Akiyoshi, T. Umebayashi, K. Asai, T. Mitsui, M. Matsumura. *Appl Catal A Gen*, 265, (2004), DOI: 10.1016/j.apcata.2004.01.007.
- 11 Y. Cong, J. Zhang, F. Chen, M. Anpo. *J Phys Chem C*, **111**, (2007)., DOI: 10.1021/jp0685030
- 12 T. Umebayashi, T. Yamaki, H. Itoh, K. Asai. *Appl Phys Lett*, **81**, (2002), DOI: 10.1063/1.1493647.
- 13 H. Li, X. Zhang, Y. Huo, J. Zhu. *Environ Sci Technol*, **41**, (2007), DOI: 10.1021/ES062680X.
- 14 I.P. Mahendra, A. Huda, H.M. Ngoc, P.T. Nghia, Tamrin, B. Wirjosentono. J Arab J Basic Appl Sci, 26, (2019). doi:10.1080/25765299.2019.1610209.
- 15 A. Huda, I.P. Mahendra, R. Ichwani, C.T. Handoko, H.M. Ngoc, B. Yudono, *et al. Rasayan J Chem*, **12**, (2019), DOI: 10.31788/RJC.2019.1215084.
- 16 H. Tian, J. Ma, K. Li, J. Li. *Ceram Int*, **35**, (2009), DOI: 10.1016/j.ceramint.2008.05.003.
- 17 M. Anpo, T. Shima, S. Kodama, Y. Kubokawa. *J Phys Chem*, **91**, (1987), DOI: 10.1021/j100300a021.
- T. Kawahara, Y. Konishi, H. Tada, N. Tohge, J. Nishii, S. Ito. *Angwandte Chemie*, 41, (2002), DOI: 10.1002/1521-3773(20020802)41:15<2811::AID-ANIE2811>3.0.CO;2-.
- T. Jeyapaul, K. Prakash, S. Harikengaram, A. Chellamani, V. Selvam. *Rasayan J Chem*, 11, (2018), DOI: 10.31788/RJC.2018.1143076.
- H. Sutanto, S. Wibowo, Z. Arifin, K.S. Firdausi, E. Hidayanto, Hadiyanto. *Rasayan J Chem*, 10, (2017), DOI: 10.7324/rjc.2017.1041793.
- 21 S. Mahshid, M. Askari, M. Sasani Ghamsari, N. Afshar, S. Lahuti. J Alloys Compd, 478, (2009), DOI: 10.1016/j.jallcom.2008.11.094.
- 22 R.G. Nair, S. Paul, S.K. Samdarshi. Sol Energy Mater Sol Cells, 95, (2011), DOI: 10.1016/j.solmat.2011.02.017.

[RJC-5461/2019]

Author's Queries (To be sent with Galley Proofs)

1. Check Figure numbers, their citation in text and presentation(as per the journal, refer a sample paper).

- 2. Biological names must be italic trough out the manuscript (if any).
- 3. Check the Title, Names of Author(s), their affiliations and E-mail of Corresponding Author once again.
- 4. Check the complete Manuscript once again for Grammar Check.
- 5. References must be strict as per the STYLE of the journal (Please refer Guidelines and a Published Paper from the current issue), which may otherwise cause unnecessary delay in publication of your paper. Also, Mention DOI with references, wherever possible and please mention complete name of the journal in the reference, not abbreviations.
- 6. Check the terms highlighted (if any) in the Galley Proofs, if it is correct? If not, please correct the same throughout the manuscript in Red/Blue ink, so that it could be visible to us.
- 7. Check/ correct all Figure and Table numbers and their proper citation/position/alignment in the text.