Direct synthesis of mesoporous TiO$_2$ using PVA as surfactant template and assessment of their photocatalytic activities

Ridhawati Thahir$^{1,*}$, Herman Bangngalino$^1$, Abdul Wahid Wahab$^2$, Nursiah La Nafie$^2$, Indah Raya$^2$

$^1$Chemical Engineering Department, State Polytechnic of Ujung Pandang, Jl. Perintis Kemerdekaan Km. 10 Makassar, Indonesia
$^2$Department of Chemistry, Faculty of Mathematics and Natural Science, Hasanuddin University, Jl. Perintis Kemerdekaan Km. 10 Makassar, Indonesia

* Corresponding author email: ridha331@poliupg.ac.id

Abstract. Mesoporous TiO$_2$ with the visible-light active photocatalyst activity in methylene blue adsorption and Cu(II) ions were effectively synthesized using PVA as a surfactant template through hydrothermal treatment. The preparation of mesoporous TiO$_2$ was identified by X-ray diffraction (XRD), nitrogen adsorption, and scanning electron microscopy (SEM) with EDX. The results indicate the presence of mesoporous materials type IV with H2-type hysteresis loop with specific surface area 106 m$^2$/g, pore volume 0.18 cc/g, and pore diameter 11 nm. XRD analysis was obtained anatase phase with tetragonal structure. It was observed that the SEM morphology of mesoporous TiO$_2$ were successful. In addition, the performance of a photocatalytic activity in methylene blue and Cu(II) ions physisorption as a model organic and inorganic pollutants was 96 mg/g and 1620 mg/g. The findings of this study suggest that it is surprising to clear the environment by degradation treatment of wastewater.

Keywords: mesoporous TiO$_2$, PVA, photocatalytic activity, adsorption

1. Introduction

The porous materials are the most potent simply accessible hollow space and used in many applications in the word of the synthesized scientific researcher. The International Union Pure Applied Chemistry (IUPAC) has classified porous material based on the pore diameter, i.e. macroporous (d $>$50 nm), mesoporous (50<d$<$2 nm) and microporous (<2 nm) materials [1]. Investigating the mesoporous materials are a continuing concern within the synthesis of TiO$_2$ mesoporous and assessment of their potential application. The varied application includes photocatalyst, dye-sensitized solar cell, remediation of Pb(II), sensor, antimicrobial activity and photodegradation of organic and inorganic pollutants in water and air [2-9].

Recent evidence suggests that synthesis and photocatalytic activity of mesoporous TiO$_2$ have attracted much practical in recent years. To acquire a right mesoporous material for photocatalyst, many structural parameters are crucial such as large pore diameter, high specific surface area, particle size, and phase structure [10]. Therefore, there are many factors affect the properties of mesoporous TiO$_2$ materials to improve their photocatalytic activity. In addition, some methods such as sol-gel route [5, 7, 11], ultrasonic irradiation [12], hydrothermal route [13], and solvothermal route [14] have been reported for synthesis mesoporous TiO$_2$. Furthermore, another strategy had been investigated for improving characteristic mesoporous TiO$_2$ such as a various surfactants template as a structure directing agent, which consist of P123 copolymer [15], polyacrylamide [11], soluble starch [16],
hexadecyltrimethylammonium bromide [17], combination surfactant CTAB/PEG/sodium dodecyl sulphate [4], polyethylene [18], and bulky organic acid [19].

More recent attention has focused on the provision of design and fabrication of mesoporous TiO$_2$. It was explored that the high surface area and large pore volume can be synthesized with a green template-free method. The result of mesoporous anatase TiO$_2$ with high surface area 119 m$^2$/g, pore volume 0.3 cc/g and photocatalytic decomposition of methyl orange 47.8% in aqueous solution 10 mg/L [6]. The other study that the various surfactant template via the sol-gel method has found the specific surface area 23.27-51.06 m$^2$/g, pore volume 0.0894-0.2108 cc/g, and higher adsorption capacity of Pb(II) are 420.5 mg/g at 1000 mg/L of lead soil solution [4]. One observer has already drawn attention to the paradox in mesoporous TiO$_2$ powders was prepared via a template-free method with large surface area 353 m$^2$/g, pore volume 0.3 cc/g, and adsorption capacity by degradation of rhodamine B of 35 mg/g [20].

A considerable amount of the previous studies have been observed on preparation route and surfactant template. The major objectives of this study have evaluated a novel synthesis method for arrangement properties of mesoporous TiO$_2$ to enhance their photocatalytic activity using polyvinyl alcohol as direct surfactant agent and modified technique via sol-gel, ultrasonic and hydrothermal treatment. In this paper, we propose that this project can advance the high performance of mesoporous TiO$_2$ materials as a photocatalyst to degrade of methylene blue and Cu(II) ions pollutants in water. The physisorption analysis, crystallinity, and the morphology of the samples were characterized by X-ray diffraction (XRD), nitrogen adsorption-desorption, and scanning electron microscopy (SEM). To analyse the amount of methylene blue or Cu(II) ions with UV-vis diffuse reflectance spectra.

2. Materials and Method

2.1. Materials

The chemicals and reagents utilized during this preparation are analytical graded and used as received, they are titanium (IV) isopropoxide as a precursor of titania (TTIP, analytical reagent; Sigma Aldrich), polyvinyl alcohol as a surfactant agent (PVA, M$_{\text{w}}$=89,000-98,000 g/mol, analytical reagent; Sigma Aldrich), acetic acid (analytical reagent; Merck), ethanol (analytical reagent; Merck), CuSO$_4$.5H$_2$O (analytical reagent; Merck), methylene blue (analytical reagent; Merck), and distilled deionized water.

![Figure 1. Diagram of preparation pathway of mesoporous TiO$_2$.](image)

2.2. Experimental method

The mesoporous TiO$_2$ were prepared via sol-gel, ultrasonic, hydrothermal treatment, and calcination method. 2.4 g PVA dissolved in deionized water at boiling point and stirred for 30 min. Afterward, this solution was added to another solution containing 15 mL of TTIP hydrolysed in 60 mL of ethanol and 6 mL of acetic acid. The sonication of the solution was performed by an Ultrasonic Disruptor UD-21 until the complete solution for 60 min at ambient condition. The intermediate products were then loaded into a teflon close container for hydrothermal treatment at 105°C for 17 hours. Finally, the products were calcined at 500°C for 5 hours. The corresponding diagram of preparation pathway of mesoporous TiO$_2$ in Fig. 1.
2.3. Characterization

X-ray diffraction (XRD) was carried out using a Bruker D2 Phaser Diffractometer System with Cu Kα radiation source 1.5406 Å run at 40 kV, 30 mA. For XRD analysis in the range of 2θ from 10° to 80°, with scan step size of 0.02°. The measurements of nitrogen sorption isotherms at -196°C were carried out using a Quantachroma NovaWin instrument version 11. The specific surface area (S\textsubscript{BET}) of the sample was calculated with the Brunauer, Emmett and Teller method \[21\], using the adsorption data in the range of relative pressure \[22\]. The total pore volume was also estimated experimentally as the volume adsorbed at P/P\textsubscript{0}=0.95. The sample was previously degassed at 300°C at approximately under vacuum for 3 hours. The morphology and defined areas of the samples were characterized using a scanning electron microscope (SEM with EDX) SU3500 with a working distance of 4940 µm and an electron voltage of 10 kV. EDX is based on the detector of characteristic x-ray emitted of an element as a result of the de-excitation of core electron holes created by a high energy electron beam. Instrument SU3500 EDX spectrum Ti and spectrum O were measured with beam 15 kV for 30 seconds.

2.4. Photocatalytic activity experiments

Methylene blue (MB) was chosen as a model organic pollutant and Cu(II) ions as an inorganic pollutant to investigate the photocatalytic activity of the mesoporous TiO\textsubscript{2} with a 150W mercury lamp. In the initial 100 mL pollutant (P1=MB=100 ppm; P2=Cu(II) ions=500 ppm). Afterward, 0.1 g TiO\textsubscript{2} powders were added, then the solutions were stirred in the dark for 60 min until adsorption/desorption stability. A fixed quantity of each P1 and P2 solution was taken at a regular interval 10 min. The solutions were filtered and analysed the amount of MB or Cu(II) ions with UV-Vis absorption spectra. The filtrates were calibrated by 5 control solutions (for P1; 20, 40, 60, 80, 100 mgL\textsuperscript{-1} and P2; 50, 100, 200, 300, 500 mgL\textsuperscript{-1}). The adsorption amount, q\textsubscript{e} (mg/g), was calculated as follows:

\[
q_e = \frac{(c_0-c_e)V}{m}
\]

Where C\textsubscript{0} and C\textsubscript{e} (mgL\textsuperscript{-1}) in this case were estimated to be the amount of MB and Cu(II) ions at initial and equilibrium state, m is the mass adsorbent (g), and V is the solution volume. The adsorption removal efficiency was calculated

\[
RE(\%) = \frac{(c_i-c_t)}{c_t} \times 100
\]

Where C\textsubscript{i} and C\textsubscript{t} set as the initial and final of P1 or P2 concentration (at a certain time t), respectively.

3. Result and Discussion

3.1. Crystal structure and morphology

The crystalline structure was determined using XRD and morphology structure by SEM with EDX. XRD patterns were indicated to evaluate the phase of the sample. Fig. 2 presents that the XRD patterns of mesoporous TiO\textsubscript{2} were indicated anatase phase in all sharp peaks observed from the XRD pattern. Noticeable diffraction peaks positioned at 2θ=25.29°, 37.94°, 48.76° and 53.95° which observed on the spectra attribute to (011), (004), (020), and (121) which match with the JCPDS card No 21-1276. The orientation plane of TiO\textsubscript{2} samples is tetragonal structure and only anatase phase. The results of this study will now be compared to the finding of previous work \[16, 19\]. The anatase phase of TiO\textsubscript{2} is the most active crystalline for photocatalytic activity \[6\].
In most recent studies, it is generally believed that characteristic of mesoporous materials is dependent on the specific surface area, pore volume and pore diameter of the particle. Scanning electron microscopy (SEM) of the TiO$_2$ materials were implemented for the sample which indicated better porous materials to evaluate the morphology. Fig. 3 shows the inter correlation among the porous of TiO$_2$ sample. The more surprising image from the using PVA as surfactant template can enhance the pore diameter until 11 nm. These results are consistent with those of other studies and suggest that the surfactant agents were created of mesoporous materials [15]. There are similarities between the attitudes expressed by PVA in this study and those described by Abdolahi Sadatlu and Mozaffari [15].

The results obtained from the preliminary analysis of mesoporous TiO$_2$ are presented in Fig. 4. In addition, the SEM images show that the obtained surfaces are porous. On the basis of EDX peaks, it can be concluded that the amount of Ti=39.91% and O=60.91%.

Figure 2. X-ray diffraction pattern of mesoporous TiO$_2$.

Figure 3. SEM image of mesoporous TiO$_2$ for macro (3 µm) and micro-morphology (500 nm).
Surface area and pore analysis of mesoporous TiO$_2$ were obtained with nitrogen adsorption-desorption isotherm which is shown by Fig. 5. Using the BET method, the surface areas were calculated 106 m$^2$/g. The average pore size distributions were evaluated with the BJH method. The pore sizes were 11 nm and pore volume 0.18 cc/g.

This observed pore size around 11 nm indicates a void space between the one-dimensional tetragonal of the TiO$_2$ sample. This finding has important implications for developing properties of the photocatalytic activity of mesoporous TiO$_2$. Typical type IV isotherms are observed for TiO$_2$ sample, showing a hysteresis loop H2-type with capillary condensation. These study produced results which corroborate the findings of a great deal of the previous work in this field [19].

### 3.2. Photocatalytic activity

The morphology and pore size analysis of the mesoporous TiO$_2$ have a great influence on the adsorption capacity. The photocatalytic activity was investigated by degradation of methylene blue as a model organic pollutants and Cu(II) ions as an inorganic pollutant. From the data on Fig. 6 shows, it is observed
that the degradation of MB shows lower adsorption capacity (96 mg/g) than the degradation of inorganic Cu(II) ions (1620 mg/g). There are several possible explanations for this result.

The particle size of Cu(II) ions is bigger than the pore diameter of mesoporous TiO$_2$. The interaction of Cu(II) ions between adsorbent of TiO$_2$ occurred on surfaces of the adsorbent. The high surface area can support interaction Cu(II) ions between adsorbent of TiO$_2$. It is different for the degradation of MB. However, with a small sample size, caution must be applied, as the finding might not be available interaction on the surface of adsorbent but adsorbable into the pore volume. This is an important issue for future research.

Figure 6. The amount of adsorption capacity of mesoporous TiO$_2$.

The adsorption removal efficiency (RE, %) for degradation of Cu(II) ions and MB were maximized at 99% and 96.4% respectively at initial concentration of 500 ppm and 100 ppm. The ability use of the mesoporous TiO$_2$ as potential adsorbents will be assessed for removal organic and inorganic pollutants in water and air. The finding of this study has a number of important implication for future practice.

4. Conclusion
Mesoporous TiO$_2$ materials were synthesized using PVA as a surfactant direct agent via sol-gel, ultrasonic, and hydrothermal treatment. This investigation assessed the photocatalytic activity of mesoporous TiO$_2$ for degradation of methylene blue and Cu(II) ions as a model organic and inorganic pollutants, respectively. The mesoporous TiO$_2$ has been proven to be able to remove inorganic and organic pollutants as affected by the high surface area of the absorbent for the first and large pore volume as well as pore diameter for the later. The TiO$_2$ is valuable for further application in wastewater treatment.

Acknowledgment
This work was supported by the research grant of State Polytechnic of Ujung Pandang, No. 018/PL.10.13/PL/2018 and we wish to thank those individuals involved in this work.

References
[1] Sing K S W, Everett D H, Haul R A W, Moscou L, Pierotti R A, Rouquerol J and Siemieniewska
T 1985 Reporting physisorption data for gas/solid systems with special reference to the
determination of surface area and porosity (Recommendations 1984) pac 57 603-19

[2] Kumar N, Hazarika S N, Limbu S, Boruah R, Deb P, Namsa N D and Das S K 2015 Hydrothermal
synthesis of anatase titanium dioxide mesoporous microspheres and their antimicrobial
activity Microporous Mesoporous Mater. 213 181-7

[3] Mao-Xiang J, Xue-Qin J, Wang-Xing L, Dong-Hong L and Zhou W 2009 Preparation and
Photocatalytic Activity of Mesoporous TiO2 Microspheres Micro and Nanosystems 1 12-6

[4] Khalaf M M, Abdallah N Y and Abd El-Lateef H M 2018 Fine-template synthetic process of
mesoporous TiO2 using ionic/nonionic surfactants as potential remediation of Pb(II) from
contaminated soil Int. J. Environ. Sci. Technol.

[5] Muniz E C, Góes M S, Silva J J, Varela J A, Joanni E, Parra R and Bueno P R 2011 Synthesis
and characterization of mesoporous TiO2 nanostructured films prepared by a modified sol–
gel method for application in dye solar cells Ceram. Int. 37 3 1017-24

[6] Lu T, Wang Y, Wang Y, Zhou L, Yang X and Su Y 2017 Synthesis of Mesoporous Anatase TiO2
Sphere with High Surface Area and Enhanced Photocatalytic Activity J. Mater. Sci. Technol.

[7] Liu J, Zhou Y, Han F, Chen D and Chen L 2017 Synthesis of mesoporous Au-TiO2 nanocomposites via a one-pot sol-gel process with enhanced photocatalytic activity Mater. Lett. 207 109-12

[8] Islam S, Bidin N, Riaz S, Rahman R A, Naseem S and Marsin F M 2015 Mesoporous SiO2 –
TiO2 nanocomposite for pH sensing Sens. Actuators B, Chem. 221 993-1002

[9] Zheng J, Xiong F-Q, Zou M, Thomas T, Jiang H, Tian Y and Yang M 2016 Enhanced
photocatalytic degradation of rhodamine B under visible light irradiation on mesoporous
anatase TiO2 microspheres by codoping with W and N Solid State Sci. 54 49-53

[10] Li W, Wu Z, Wang J, Elzatahry A A and Zhao D 2013 A Perspective on Mesoporous TiO2
Materials Chem. Mater. 26 1 287-98

[11] Yin Q, Xiang J, Wang X, Guo X and Zhang T 2016 Preparation of highly crystalline mesoporous
TiO2by sol–gel method combined with two-step calcining process J. Exp. Nanosci. 11 14
1127-37

[12] Swapna M V and Haridas K R 2015 An easier method of preparation of mesoporous anatase TiO2
nanoparticles via ultrasonic irradiation J. Exp. Nanosci. 11 7 540-9

[13] Wang J, Chen G, Yin J, Luo C and Zhao X 2017 Enhanced electrorheological performance and
antisedimentation property of mesoporous anatase TiO2 shell prepared by hydrothermal
process Smart Mater. Struct. 26 3 035036

[14] Zhang Y, Zhang S, Wang K, Ding F and Wu J 2013 Surfactant-Free Solvothermal Method for
Synthesis of Mesoporous Nanocrystalline TiO2 Microspheres with Tailored Pore Size J.
Nanomater. 2013 1-7

[15] Abdolahi Sadatlu M A and Mozaffari N 2016 Synthesis of mesoporous TiO2 structures through
P123 copolymer as the structural directing agent and assessment of their performance in dye-
sensitized solar cells Sol. Energy 133 24-34

[16] Muniani S S, Mohd Kaus N H, Jiang Z-T, Altarawneh M and Lee H L 2017 Green synthesis of
mesoporous anatase TiO2 nanoparticles and their photocatalytic activities RSC Adv. 7 76
48083-94

[17] Gharakhlou A R and Sarvi M N 2017 Synthesis of mesoporous nanoparticles of TiO2 from
ilmenite Mater. Res. Express 4 2 025027

[18] Jin X, Yuan K, Xu C, Wang X, Zhu L, Zhang G and Xu D 2018 Water steam modified
crystallization and microstructure of mesoporous TiO2 2 nanofibers Ceram. Int. 44 2 2158-64

[19] Bakre P V and Tilve S G 2018 Direct access to highly crystalline mesoporous nano TiO2 using
sterically bulky organic acid templates J. Phys. Chem. Solids 116 234-40

[20] Ovodok E, Maltanava H, Poznyak S, Ivanovskaya M, Kudlash A, Scharnagl N and Tedim J 2017
Synthesis and characterization of efficient TiO2 2 mesoporous photocatalysts Mater. Today:
Proceedings 4 11 11526-33

[21] Storck S, Bretinger H and Maier W F 1998 Characterization of micro- and mesoporous solids by physisorption methods and pore-size analysis *Appl. Catal. A, Gen.* 174 137-46

[22] Brunauer S, Deming L S, Deming W E and Teller E 1940 On a Theory of the van der Waals Adsorption of Gases *J. Am. Chem. Soc.* 62 7 1723-32