13th Joint Conference on Chemistry (13th JCC)

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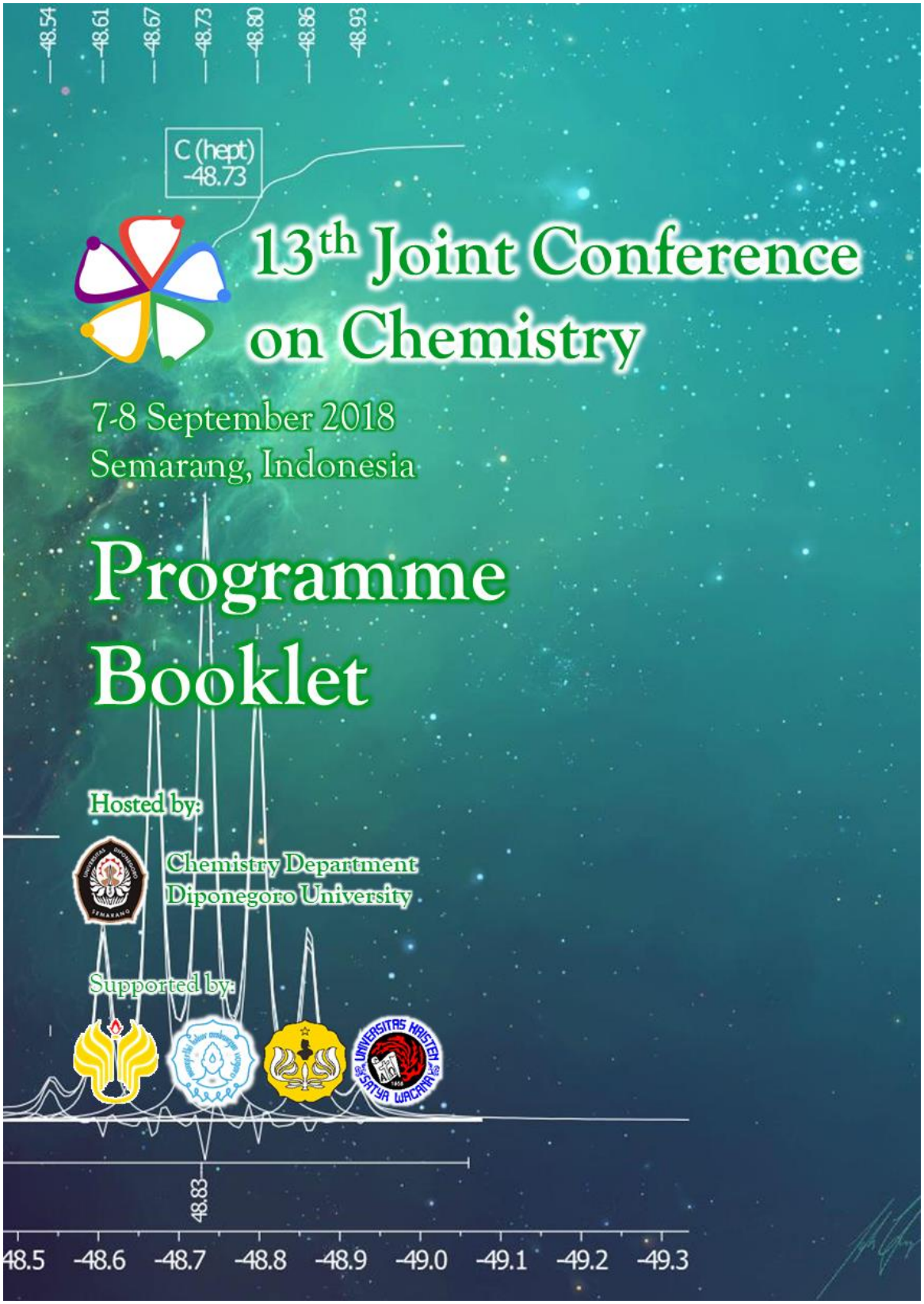

13th Joint Conference on Chemistry

7-8 September 2018

Semarang, Indonesia

Preface

On behalf of the Consortium of Chemistry Department in Central Java, Indonesia and the JCC Committee, I would like to thank you for your participation in the 13th Joint Conference on Chemistry which to be held from 7-8th September 2018 in Semarang, Indonesia. The Joint Conference on Chemistry is an annual conference organized by the consortium of Chemistry Department of five universities in Central Java: Diponegoro University (UNDIP), State University of Semarang (UNNES), Sebelas Maret University (UNS), Jenderal Soedirman University (UNSOED) and Satya Wacana Christian University (UKSW). The JCC has been held since 2006.

This conference provides an interactive international forum to provide for sharing and exchange information on the latest research on Chemistry and related sciences, to enhance the capacities for creating innovation system, to contribute in the formulation of global strategies in advancing science role as well as developing policy initiatives in community, to stimulate future collaborations among industries, researchers, governments and other stakeholders who apply science and technology for better live. The speakers and participants of the 13th JCC are up to 250 coming from various countries extending from Indonesia, Malaysia, Philippine, Australia, South Korea, Japan, Iran, Nigeria, UK and India.

We received nearly 200 papers submitted to be included in the proceedings of this conference and after the review and revision process we finally got 158 papers to be published

I would like to thank for the endeavour of committee from Chemistry Department - UNDIP and the consortium member. In addition, the conference committee acknowledges the technical and financial support from Diponegoro University.

Adi Darmawan, Ph.D
The Chair of 13th Joint Conference of Chemistry
Chemistry Department, Faculty of Science and Mathematics, Diponegoro University

Committee

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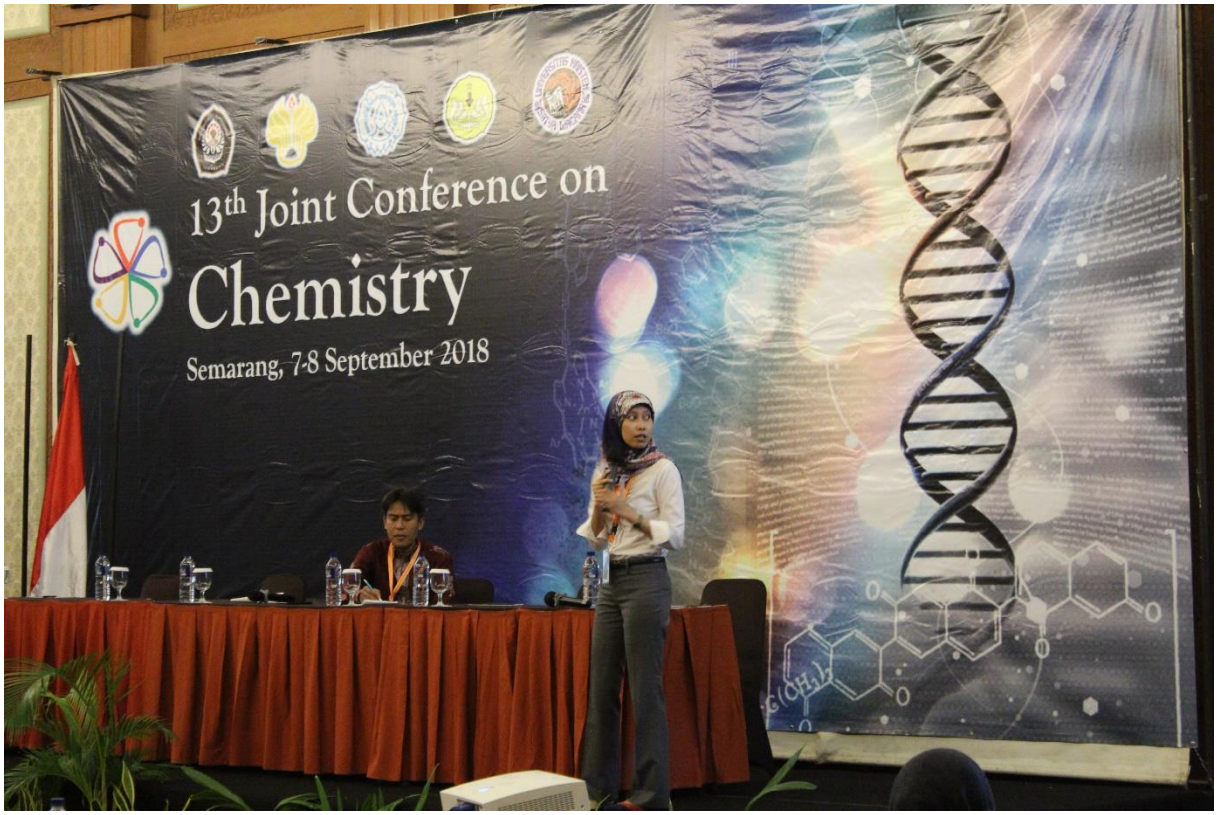
Documentation


13th Joint Conference on
Chemistry
Semarang, 7-8 September 2018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Muktiningsih Nurjayadi, Dwi Ariastuti, Kurnia Agustini, Asri Sulfianti and Wibowo Mangunwardoyo

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Antioxidant activity from limonene encapsulated by chitosan

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Ridhawati Thahir, Herman Bangngalino, Abdul Wahid Wahab, Nursiah La Nafie and Indah Raya

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Development of heterogeneous catalyst from chicken bone and catalytic testing for biodiesel with simultaneous processing

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Saprizal Hadisaputra, Agus Abhi Purwoko, Saprini Hamdiani and N Nuryono

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Agus Abhi Purwoko, Veni Rori Setiawati and Saprizal Hadisaputra

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Siti Mariyah Ulfa, Dita Prihartini, Adam Mahfud, Rizqi Munandar and Indah Nur Pramesti

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Electrosynthesis of coordination polymers containing magnesium(II) and benzene 1,3,5-tricarboxylate: the influence of solvents and electrolytes toward the dimensionality

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Synthesis and characterization of composite gels starch-graftacrylic acid/bentonite (St-g-AA/B) using N'Nmethylenebisacrylamide (MBA)

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Synthesis of silver orthophosphate under dimethyl sulfoxide solvent and their photocatalytic properties

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Enhanced hydrogen sorption properties over Mg^{2+} modified solvothermal synthesized HKUST-1 (Mg^{2+}/HKUST-1)

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Synthesis of activated carbon/bismuth oxide composite and its characterization for battery electrode

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Optimization of conventional and ultrasound assisted extraction of inulin from gembili tubers (*Dioscorea esculenta* L.) using response surface methodology (RSM)

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Anti-atherosclerosis potency of *Pandanus tectorius* fruit rich by trangeretin and ethyl trans-caffeate, and their cytotoxicity against HepG2 cell line

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Discovering anticancer compound of ethyl acetate extract from RL1 code endophytic fungi culture derived by *Phyllanthus niruri* Linn leaves through cell cycle modulation in T47d cells

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Direct synthesis of mesoporous TiO_2 using PVA as surfactant template and assessment of their photocatalytic activities

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Direct synthesis of mesoporous TiO_2 using PVA as surfactant template and assessment of their photocatalytic activities

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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 nitrogen adsorption on TiO_2 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 indicated 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) 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_{wt}=89,000-98,000 g/mol, analytical reagent; Sigma Aldrich), acetic acid (analytical reagent; Merck), ethanol (analytical reagent; Merck), $CuSO_4.5H_2O$ (analytical reagent; Merck), methylene blue (analytical reagent; Merck), and distilled deionized water.

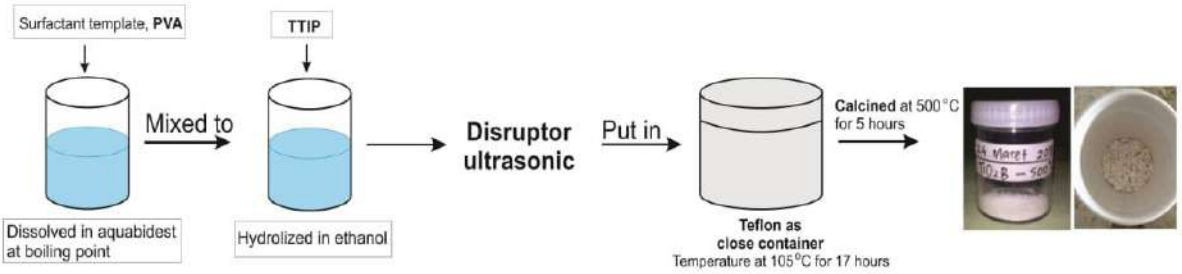

Figure 1. Diagram of preparation pathway of mesoporous TiO_2.

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 Quantachoma NovaWin instrument version 11. The specific surface area (S_{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_o=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_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_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^{-1} and P2; 50, 100, 200, 300, 500 mgL^{-1}). The adsorption amount, q_e (mg/g), was calculated as follows:

$$q_e = \frac{(Co-Ce)V}{m}$$

Where C_o and C_e (mgL^{-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

$$RE(\%) = \frac{(Ci-Ct)}{Ci} x\ 100$$

Where C_i and C_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_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_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_2 is the most active crystalline for photocatalytic activity [6].

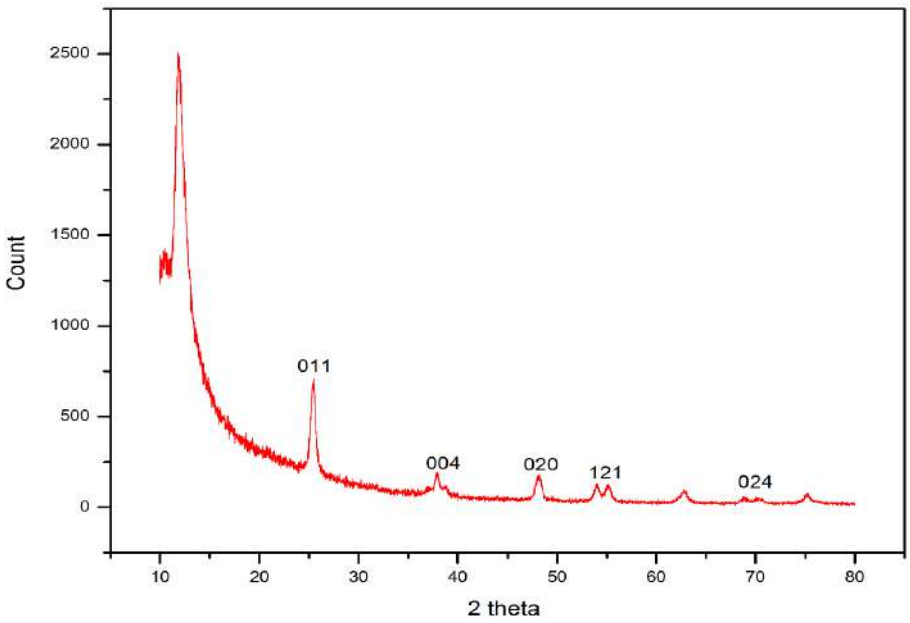

Figure 2. X-ray diffraction pattern of mesoporous TiO_2.

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].

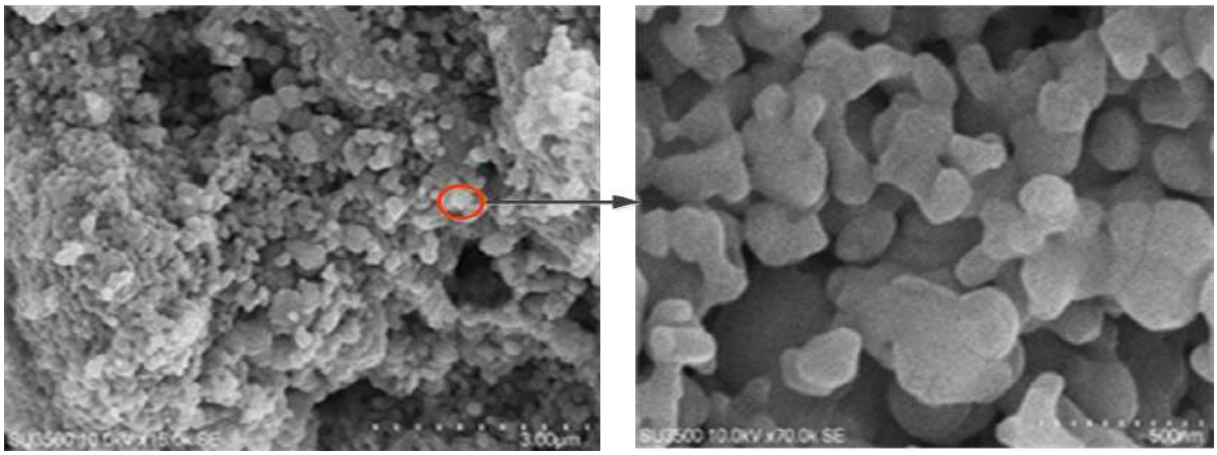

Figure 3. SEM image of mesoporous TiO_2 for macro (3 µm) and micro-morphology (500 nm).

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%.

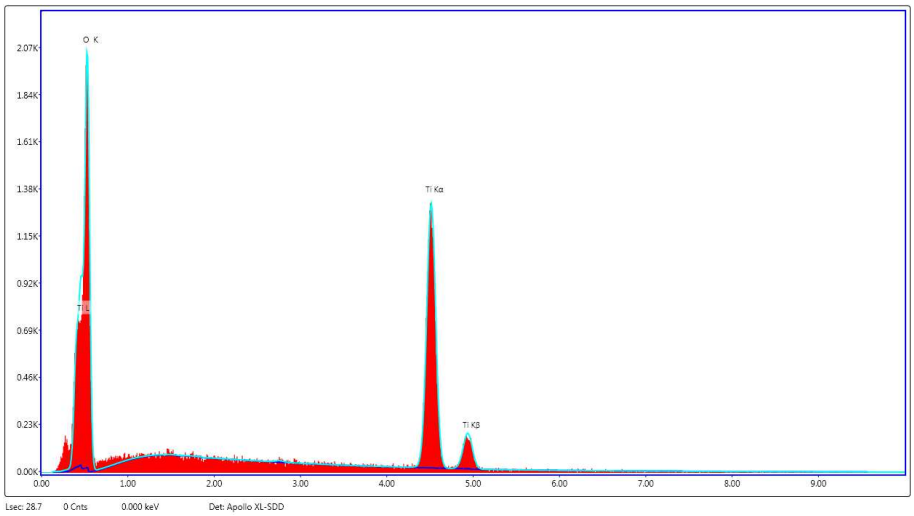

Figure 4. SEM-EDX micrograph of mesoporous TiO_2

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.

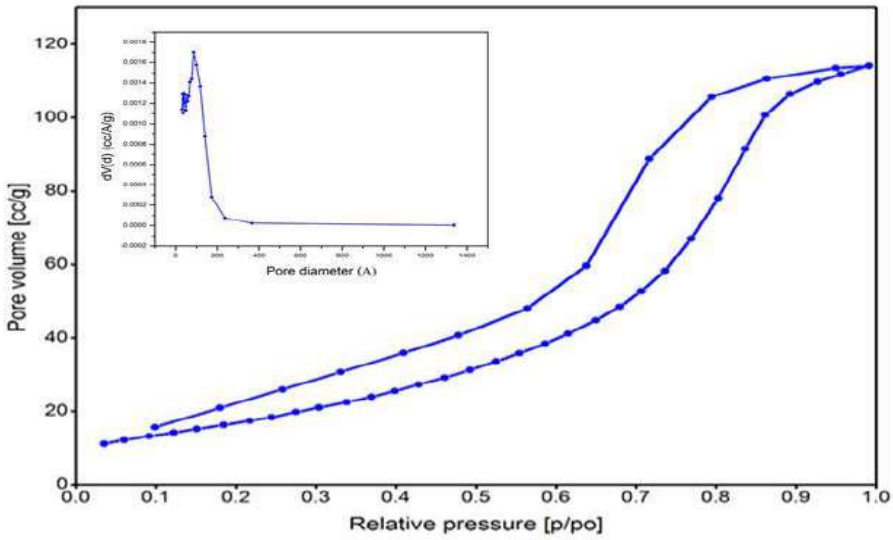

Figure 5. Nitrogen adsorption-desorption isotherm of mesoporous TiO_2.

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.

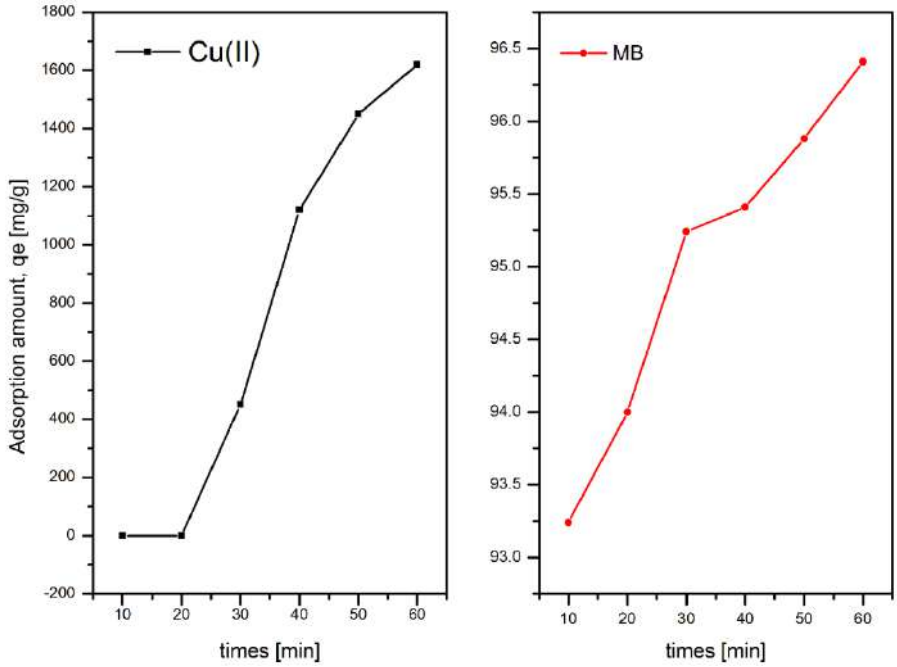

Figure 6. The amount of adsorption capacity of mesoporous TiO_2.

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.

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