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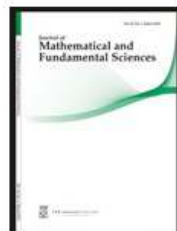
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ISSN: 2337-5760  
E-ISSN: 2338-5510

### Publication History

#### Formerly known as:

- ITB Journal of Science (2008 - 2012)
- Proceedings ITB Science and Technology (2003 - 2007)
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# Effect of V Dopant on Physicochemical Properties of Vanadium-Doped Anatase Synthesized via Simple Reflux Technique

Hari Sutrisno<sup>1</sup>, Ariswan<sup>2</sup> & Dyah Purwaningsih<sup>1</sup>

<sup>1</sup> Department of Chemistry Education, Faculty of Mathematics and Natural Sciences, Yogyakarta State University, Kampus Karangmalang, Jl. Colombo 1, Yogyakarta, 55281, Indonesia

<sup>2</sup> Department of Physics Education, Faculty of Mathematics and Natural Sciences, Yogyakarta State University, Kampus Karangmalang, Jl. Colombo 1, Yogyakarta, 55281, Indonesia  
Email: sutrisnohari@uny.ac.id

**Abstract.** Mesoporous pure TiO<sub>2</sub> (M-TiO<sub>2</sub>) and mesoporous-vanadium-doped TiO<sub>2</sub> (M-V-doped TiO<sub>2</sub>) were successfully synthesized via a facile and simple reflux technique. The purpose of this research was to study the effect of vanadium dopant on the physicochemical properties of all materials obtained. Characterization of the prepared materials was carried out using X-ray diffraction (XRD), scanning electron microscopy-energy dispersive X-ray spectroscopy (SEM-EDS) and N<sub>2</sub>-adsorption-desorption analysis. The presence of Ti and O elements in M-TiO<sub>2</sub> and of Ti, V and O elements in M-V doped TiO<sub>2</sub> could be detected by SEM-EDS, while the patterns of X-ray diffraction of all the prepared samples had a well-crystalline surface of anatase type. All mesoporous vanadium-doped TiO<sub>2</sub> (M-V-doped TiO<sub>2</sub>) materials performed in a highly transparent mode in the visible region at 554 nm ( $E_g = 2.24$  eV) and 588 nm ( $E_g = 2.12$  eV) for 3.3 and 4.9 wt% V doped TiO<sub>2</sub>, respectively. The Rietveld refinement method was applied to extract the structural parameters of the M-TiO<sub>2</sub> and M-V-doped TiO<sub>2</sub> using the Fullprof program in the WinPlotr package. The prepared materials were refined in the crystal system and space group of anatase (tetragonal,  $I4_1/amd$  (141)). The vanadium ion was successfully doped into TiO<sub>2</sub>. The isotherm type of M-TiO<sub>2</sub> and 2.3 wt% V doped TiO<sub>2</sub> were of type IV, with a profile of type H2 hysteresis loops, while the 3.3 and 4.9 wt% vanadium-doped TiO<sub>2</sub> reflected isotherm type III. The Brunauer-Emmett-Teller (BET) results showed a significant reduction in surface area due to increased concentrations of vanadium. The highest values of BET-specific surface area, pore volume and average pore size of M-TiO<sub>2</sub> were 46 m<sup>2</sup>/g, 18.45 nm and 0.2572 cm<sup>3</sup>/g respectively.

**Keywords:** anatase; hysteresis; mesoporous materials; reflux technique; Rietveld analysis.

## 1 Introduction

Titanium dioxide or titania (TiO<sub>2</sub>) is an n-type semiconductor that has a wide band-gap with specific chemical properties and stability. TiO<sub>2</sub> has attracted a great deal of interest over the past decades due to its special physiochemical properties. A

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Received August 4<sup>th</sup>, 2015, 1<sup>st</sup> Revision December 5<sup>th</sup>, 2015, 2<sup>nd</sup> Revision March 4<sup>th</sup>, 2016, 3<sup>rd</sup> Revision March 13<sup>th</sup>, 2016, Accepted for publication March 14<sup>th</sup>, 2016.

Copyright © 2016 Published by ITB Journal Publisher, ISSN: 2337-5760, DOI: 10.5614/j.math.fund.sci.2016.48.1.8

number of potential applications of TiO<sub>2</sub> in both fundamental research and practical development work, have been developed, including in photocatalysis [1,2], photovoltaic cells [3,4], photoinduced super-hydrophilicity [5,6] and anti-bacterial applications [7,8].

The properties of TiO<sub>2</sub> depend on the phase, surface area, dopant, degree of crystallization, particle size, pore size and morphology. For TiO<sub>2</sub> photocatalysis and photoinduced super-hydrophilicity, efforts have been made in two aspects: the first is to narrow the wide band gap semiconductor of TiO<sub>2</sub> to extend the spectral response to the visible region or ultra violet radiation, among others by doping with nonmetallic dopants such as N [9], C [10] and S [11]. The other is to minimize the recombination rate of the photogenerated electron-hole pairs by doping with transition metals such as Ag [12,13], V [14] and Cr [15] into TiO<sub>2</sub> to develop visible-photocatalysts. Among transition metal ions, vanadium ion is attractive as dopant because it will generate an intermediate band so that absorption will happen in the visible region. Various methods have been selected to prepare V doped TiO<sub>2</sub> photocatalysts, such as a hydrothermal method [16], electro spinning technique [17], wet chemical method [14,18], sol-gel method [19,20], flame spray pyrolysis technique [21], and anodic oxidation [22].

In this study, pure and 2.3, 3.3, 4.9 wt% V doped mesoporous-TiO<sub>2</sub> were prepared by reflux technique at 150 °C for 6 hours. Mesoporous material can be synthesized by using templates, for example surfactant, lipid, polymer etc. and free-templates. This research synthesized pure and V-doped TiO<sub>2</sub> mesoporous materials without using a template. In previous studies, mesoporous materials were produced with free templates at low temperatures, for example by hydrothermal and solvothermal methods [23,24]. At low temperatures, the polycondensation process takes place very slowly, especially propagation, which consists of oligation and oxolation that allow to control the formation of pores, the crystal structure and morphology [25]. The morphology and the influence of V doped-TiO<sub>2</sub> were investigated using scanning electron microscopy (SEM) and energy dispersive X-ray spectroscopy (EDS), respectively. The purpose of the research was to observe the role of the V dopants with respect to their physicochemical properties.

## 2 Experiment

### 2.1 Materials

Ammonium hydroxide (NH<sub>4</sub>OH, 28-30% NH<sub>3</sub> basis) solution, hydrogen peroxide solution (H<sub>2</sub>O<sub>2</sub>, 10 wt. % in H<sub>2</sub>O), ammonium metavanadate (NH<sub>4</sub>VO<sub>3</sub>, 99%), titanium (IV) chloride (TiCl<sub>4</sub>, 99%) manufactured by Sigma-Aldrich were purchased. All the reagents were used without purification. Ti(O<sub>2</sub>)O.2H<sub>2</sub>O was obtained from the reaction of TiCl<sub>4</sub> and H<sub>2</sub>O<sub>2</sub> [26]. In a typical procedure, 50 ml of

TiCl<sub>4</sub> was added into a 500 ml glass flask loaded in an ice water bath and then 25 ml of H<sub>2</sub>O<sub>2</sub> was added slowly into the reaction vessel under magnetic stirring.

## 2.2 Preparation of M-TiO<sub>2</sub> and M-V doped TiO<sub>2</sub>

A series of vanadium doped TiO<sub>2</sub> at various weight percentages of V doped TiO<sub>2</sub> was prepared by reflux technique. Mesoporous-TiO<sub>2</sub> (M-TiO<sub>2</sub>) and mesoporous-vanadium-doped TiO<sub>2</sub> (M-V doped TiO<sub>2</sub>) were prepared. In a typical synthesis, 10 g Ti(O<sub>2</sub>)O.2H<sub>2</sub>O was dissolved in 50 ml of distilled water under vigorous stirring. The solution was stirred for 4 hours to obtain a colloidal A. To determine the effect of the NH<sub>4</sub>VO<sub>3</sub> concentration, 0, 3, 6 and 9 wt% V doped TiO<sub>2</sub> respectively were adopted in a separated beaker. The samples were thoroughly dissolved in 20 mL of distilled water under vigorous stirring to obtain solution B1, B2, B3, and B4 respectively. Each B1, B2, B3 and B4 colloid was then slowly added to each colloid A. The four types of final solution mixtures were sealed and further stirred for 2 h, after which NH<sub>4</sub>OH was added dropwise until the pH value reached about 8-10. Finally, the solution mixtures were heated with a magnetic stirrer in reflux equipment at 150°C for 6 hours. The obtained precipitate was filtered, washed with distilled water and dried at 70°C for 3 hours. Finally, the precipitate materials were calcined at 600°C for 2 hours.

## 2.3 Characterization

The morphologies were characterized using a Phenom ProX Desktop scanning electron microscope (SEM). The influence of the vanadium load on the structural characteristics of the vanadium-doped TiO<sub>2</sub> and the presence of Ti, V and O elements in the prepared materials were analyzed using energy dispersive X-ray spectroscopy (EDS).

All samples were suspended in ethanol solution. The colloidal samples were evaluated using a Shimadzu 2450 UV-vis diode array spectrophotometer.

The prepared materials were examined using powder X-ray diffraction (XRD). The XRD patterns were obtained on a Rigaku Miniflex 600-Benchtop XRD instrument, operated in the Bragg configuration using Cu K $\alpha$  radiation ( $\lambda = 1.5406 \text{ \AA}$ ). The XRD instrument was run at 40 kV and 15 mA. The XRD data were collected in steps of 0.02° with a count time of 15 s/step. The intensities were determined in the 2 $\theta$  interval ranging from 20° to 90°. To refine the crystal structures from the powder diffraction data, the Rietveld analysis was carried out using the Fullprof program by Roisnel and Rodriguez Carbajal in the WinPlotr package [27]. The following parameters were refined: unit cell, scale factor, and full width at half-maximum (FWHM).

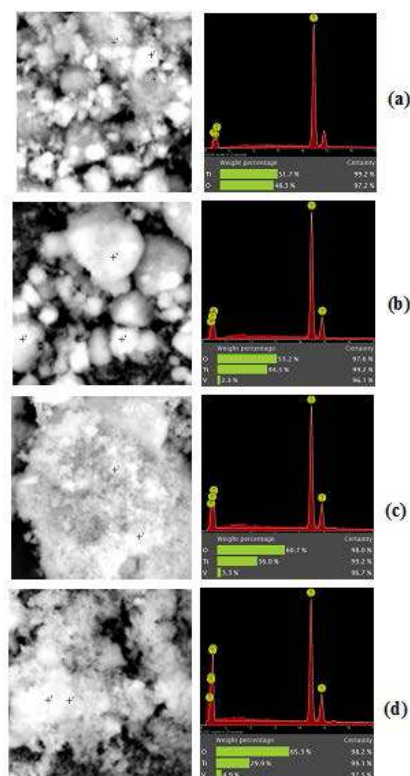
The conventional analysis of N<sub>2</sub> adsorption-desorption isotherms was carried out at 77 K with a Micromeritics ASAP 2020 instrument based on adsorption-desorption

data (55 points). The samples were degassed at 150 °C under vacuum for 4 h prior to analysis with a vacuum set point of 10  $\mu\text{mHg}$ . The Brunauer-Emmett-Teller (BET) specific surface area ( $S_{\text{BET}}$ ) data were collected based on adsorption data (10 point) in the multi-point BET measurement from (P/P0) of  $\sim 0.06$  to  $\sim 0.30$  [28]. The pore volume and average pore size were evaluated from the adsorption-desorption isotherms by the procedure developed by Barrett, Joyner and Halenda (BJH) [29]. The BET and BJH calculations were done with the ASAP 2020 V4.01 software from Micromeritics.

### 3 Results and Discussions

#### 3.1 SEM images and EDS analysis

Typical SEM images and EDS analysis of M-TiO<sub>2</sub> and M-V doped TiO<sub>2</sub> nanocrystalline are shown in Figure 1. Figures 1(a)-1(d) show that the particles were highly agglomerated.



**Figure 1** SEM images, EDS analysis and weight percentage (%) of Ti, O and V in M-TiO<sub>2</sub> (a), 2.3 wt% V doped TiO<sub>2</sub> (b), 3.3 wt% V doped TiO<sub>2</sub> (c), and 4.9 wt% V doped TiO<sub>2</sub>(d).



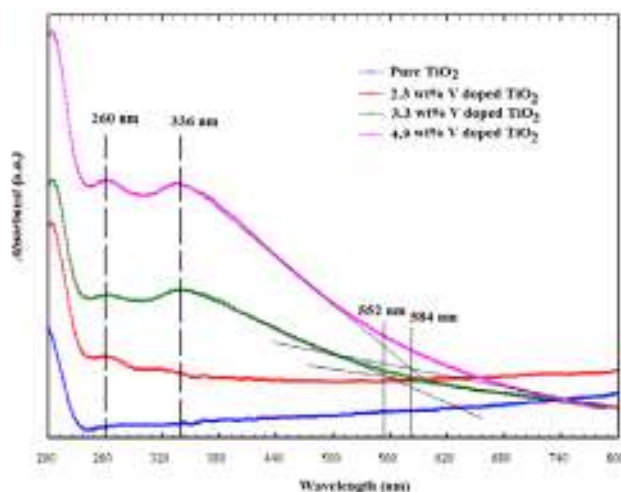
The EDS analysis identified the presence of Ti and O elements in the M-TiO<sub>2</sub> and the presence of Ti, V and O elements in the M-V doped TiO<sub>2</sub>. As shown in Figure 1, the increasing concentration of NH<sub>4</sub>VO<sub>3</sub> also increased the quantity of V in the M-V doped TiO<sub>2</sub>. The crystal of V<sub>2</sub>O<sub>5</sub> was found in the 4.9 wt% V doped TiO<sub>2</sub>, which was confirmed by the XRD pattern (Figure 2(d)). Table 1 shows that the addition of 3, 6 and 9 wt% V doped TiO<sub>2</sub> produced only 2.3, 3.3 and 4.9 wt% V doped TiO<sub>2</sub> respectively.

**Table 1** Weight Percentage of Ti, O and V of M-TiO<sub>2</sub> and M-V Doped TiO<sub>2</sub>.

Samples	Weight Percentage (wt%)			Mole ratio = Ti:V
	Ti	V	O	
Pure TiO <sub>2</sub>	51.7	0	48.3	1.079 : 0.000
2.3 wt% V doped TiO <sub>2</sub>	44.5	2.3	53.2	0.958 : 0.040
3.3 wt% V doped TiO <sub>2</sub>	36.0	3.3	60.7	0.922 : 0.074
4.9 wt% V doped TiO <sub>2</sub>	29.9	4.9	65.3	0.868 : 0.126

### 3.2 Optical Properties

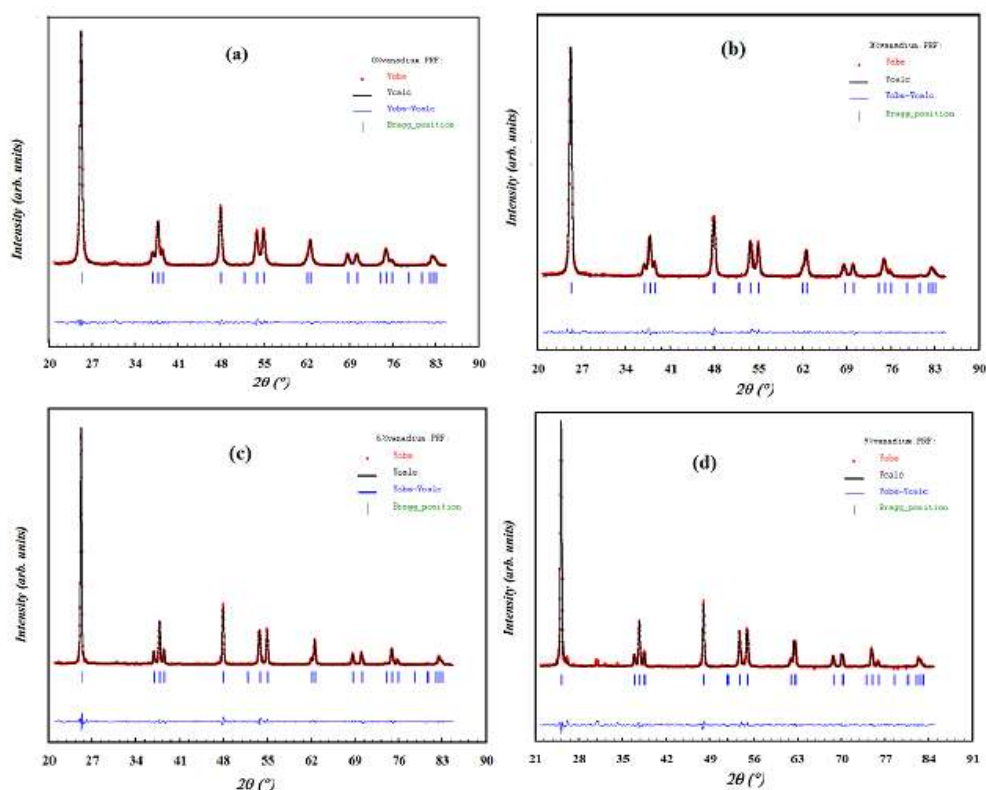
In order to investigate the optical properties of the M-TiO<sub>2</sub> and M-V doped TiO<sub>2</sub> samples at 2.3, 3.3 and 4.9 wt% V doped TiO<sub>2</sub>, the absorbance was measured as a function of wavelength in the range of 200-800 nm. The UV-visible spectra for the M-V doped TiO<sub>2</sub> samples at 2.3, 3.3 and 4.9 wt% V doped TiO<sub>2</sub> are shown in Figure 2. For comparison purposes, the spectrum of undoped TiO<sub>2</sub> (M-TiO<sub>2</sub>) is also displayed. All mesoporous V-doped TiO<sub>2</sub> (M-V doped TiO<sub>2</sub>) powders performed in a highly transparent mode in the visible region at 554 nm ( $E_g = 2.24$  eV) and 588 nm ( $E_g = 2.12$  eV) for the 3.3 and 4.9 wt% V doped TiO<sub>2</sub>, respectively.



**Figure 2** UV-Vis spectra of undoped TiO<sub>2</sub> and M-V doped TiO<sub>2</sub> at 2.3, 3.3 and 4.9 wt% V doped TiO<sub>2</sub> suspended in ethanol solution.

### 3.3 XRD Analysis

The wide-angle XRD patterns of the M-TiO<sub>2</sub> and the M-V doped TiO<sub>2</sub> are presented in Figure 3. The peak positions at  $2\theta$ : 25.36, 37.04, 37.88, 38.61, 48.11, 53.96, 55.10, 62.11, 62.69, 68.75, 70.26, 75.01, 75.96, and 82.65° are in accordance with the TiO<sub>2</sub> anatase phase. The main diffraction peaks can be indexed as the (101), (103), (004), (112), (200), (105), (211), (213), (204), (116), (220), (215), (301), and (224) reflections of anatase crystalline phase matching as shown in JCPDS card No. 21-1272. The reflections of TiO<sub>2</sub>(B) phase can be observed in the prepared M-TiO<sub>2</sub>. The presence of TiO<sub>2</sub>(B) phase is characterized by the presence of diffraction peaks at  $2\theta = 30.7^\circ$  [30].



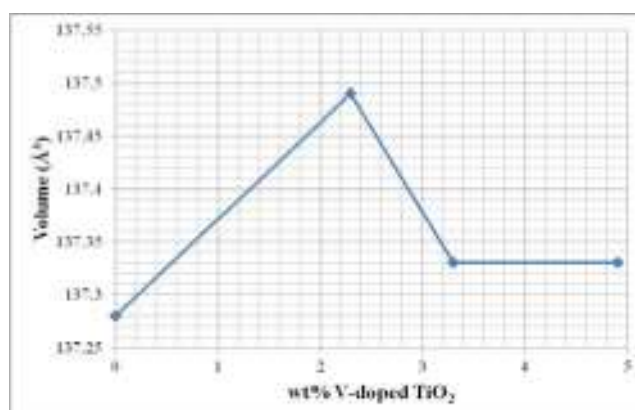
**Figure 3** Rietveld refinement patterns of M-TiO<sub>2</sub> (a), 2.3 wt% V doped TiO<sub>2</sub> (b), 3.3 wt% V doped TiO<sub>2</sub> (c), and 4.9 wt% V doped TiO<sub>2</sub> (d).

The characteristic peaks from V<sub>2</sub>O<sub>5</sub> were found in the XRD patterns of the 4.9 wt% V doped TiO<sub>2</sub>. Rietveld refinement was carried out using the Fullprof program on the M-TiO<sub>2</sub> and the M-V doped TiO<sub>2</sub> samples. All the prepared materials consisted of anatase phase (major) and the Rietveld refinements are shown in Figure 2(a), 2(b), 2(c) and 2(d) respectively. The theoretical data are represented by the solid

line and the experimental points are expressed as dots (.). The difference between the experimental and the theoretical data is represented by the line at the bottom. The vertical lines (blue lines) represent the Bragg's allowed peaks. The results of the crystal system, cell parameters (a, b, c) and atomic position (x, y, z) are presented in Table 2.

**Table 2** Crystal System, Cell Parameters (a, b, c), Volume and Atomic Position (x,y,z) of M-TiO<sub>2</sub> and M-V Doped TiO<sub>2</sub>

Prepared material	TiO <sub>2</sub> phase	Unit cell		Vol. (Å <sup>3</sup> )	Atom	x	y	Z	Rp	Rwp	χ <sup>2</sup>
		a = b (Å)	C (Å)								
Pure TiO <sub>2</sub>	Anatase (tetragonal, I4 <sub>1</sub> /amd)	3.7951	9.5316	137,28	Ti <sup>+4</sup> O <sup>2-</sup>	0.00 0.00	0.00 -0.50	0.50 0.5453	7.44	10.5	0.188
2.3 wt% V doped TiO <sub>2</sub>	Anatase (tetragonal, I4 <sub>1</sub> /amd)	3.7959	9.5384	137.44	Ti <sup>+4</sup> O <sup>2-</sup>	0.00 0.00	0.00 -0.50	0.50 0.5437	7.26	10.9	0.208
3.3 wt% V doped TiO <sub>2</sub>	Anatase (tetragonal, I4 <sub>1</sub> /amd)	3.7949	9.5361	137.33	Ti <sup>+4</sup> O <sup>2-</sup>	0.00 0.00	0.00 -0.50	0.50 0.5424	11.1	14.7	0.390
4.9 wt% V doped TiO <sub>2</sub>	Anatase (tetragonal, I4 <sub>1</sub> /amd)	3.7947	9.5368	137.33	Ti <sup>+4</sup> O <sup>2-</sup>	0.00 0.00	0.00 -0.50	0.50 0,5419	12.7	15.9	0.508



**Figure 4** The influence of V doped TiO<sub>2</sub> on tetragonal lattice parameters (cell volume).

Figure 4 shows the influence of V doped TiO<sub>2</sub> on cell volume. In this case, the ionic radius of V<sup>5+</sup> (0.054 nm) is compatible with that of Ti<sup>4+</sup> (0.0605 nm) [31]. When the Ti<sup>4+</sup> ion is replaced by the V<sup>5+</sup> ion, the cell volume should become smaller, but in fact the cell volume becomes larger. This is because the insertion of oxygen (O<sup>2-</sup>) to balance the cationic charge. Furthermore, the role of the ionic radius of V<sup>5+</sup> (0.54 Å) is obvious on increasing the V-doped TiO<sub>2</sub> resulted in a decreased cell volume.

### 3.4 N<sub>2</sub> Adsorption/Desorption Measurement

The N<sub>2</sub> adsorption-desorption isotherms of representative prepared samples are displayed in Figure 5. According to the IUPAC classification, for pure TiO<sub>2</sub> (M-TiO<sub>2</sub>) and 2.3 wt% vanadium-doped TiO<sub>2</sub> (Fig. 5a and 5b), the isotherms are of type IV, with a profile of type H2 hysteresis loops, while the 3.3 and 4.9 wt% vanadium doped TiO<sub>2</sub> both reflect isotherm type III. The isotherm type IV represents a capillary condensation phenomenon. The isotherm type is characteristic of a material that contains mesoporosity and high energy adsorption. Based on the type H2 hysteresis loop profile, the prepared materials have pores with narrow, wide sections and possible interconnecting channels. The isotherm type III explains the multilayer formation. This is characteristic of a material, that is not porous, or possibly macroporous, and has low energy adsorption [32,33].

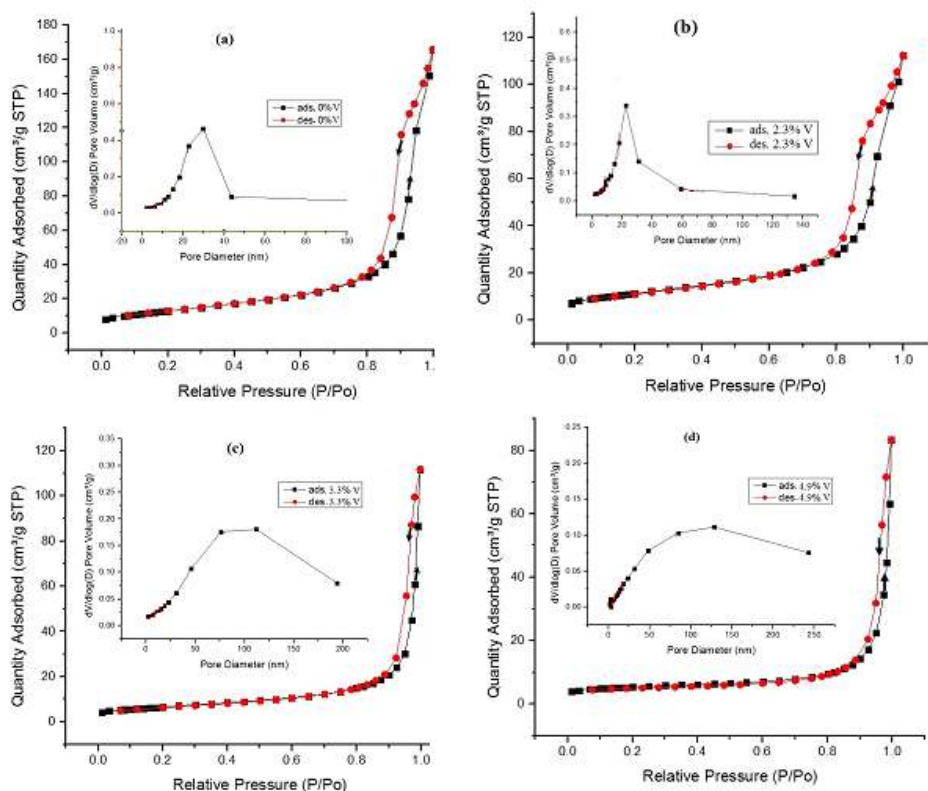
**Table 3** Surface Area, Volume and Pore Size Distribution of M-TiO<sub>2</sub> and M-V Doped TiO<sub>2</sub> from Nitrogen Adsorption-desorption Isotherm Measurements

Samples	Surface Area BET ( $S_{\text{BET}}$ ) (m <sup>2</sup> /g)	t-Plot Method			Pore Volume at $P/P_0 \approx$ 0.99 (cm <sup>3</sup> /g)	Pore Size (nm) BJH ads.
		External Surface Area (m <sup>2</sup> /g)	Micropore Volume (cm <sup>3</sup> /g)	Micropore Area (m <sup>2</sup> /g)		
Pure TiO <sub>2</sub>	46	46	-	-	0.2565	18.45
2.3 wt% V doped TiO <sub>2</sub>	40	39	0.00044	1	0.1738	15.46
3.3 wt% V doped TiO <sub>2</sub>	23	22	0.00063	1	0.1729	28.72
4.9 wt% V doped TiO <sub>2</sub>	17	10	0.00365	7	0.1289	41.20

The surface area, volume and pore size distribution of the prepared samples are summarized in Table 3. From the table, it can be seen that the concentration of V in the M-V doped TiO<sub>2</sub> increased while the BET surface area and mean pore size decreased with smallest surface area 17 m<sup>2</sup>/g, which corresponds to a mean porous size of 41.20 nm in the 4.9 wt% V doped TiO<sub>2</sub>. The BET surface area of the M-TiO<sub>2</sub> exhibited a maximum surface area of 46 m<sup>2</sup>/g, corresponding to a mean porous size of 18.45 nm calculated by the BJH<sub>ads</sub> method.

The pore size distribution curve was calculated from the adsorption-desorption isotherms by the procedure developed by Barrett, Joyner and Halenda (BJH). Figure 3(b)-3(d) (inset) depicts the pore size distribution of the M-TiO<sub>2</sub> and the M-V doped TiO<sub>2</sub> calculated by the BJH<sub>ads</sub> method. The BJH adsorption-desorption analyses show that the prepared materials exhibited mesoporous material. The results of the BJH-adsorption analysis showed that the M-TiO<sub>2</sub> had a pore size of 18.45 nm, while the 2.3, 3.3, 4.9 wt% V doped TiO<sub>2</sub> had pore sizes of 15.46, 28.72,

41.20 nm, respectively. The pore sizes of the M-TiO<sub>2</sub> and the M-V doped TiO<sub>2</sub> are presented in Table 3.



**Figure 5** Nitrogen adsorption-desorption isotherms of M-TiO<sub>2</sub> (a), 2.3 wt% V doped TiO<sub>2</sub> (b), 3.3 wt% V doped TiO<sub>2</sub> (c), and 4.9 wt% V doped TiO<sub>2</sub> (d) (*Inset: pore size distribution of the prepared samples from the adsorption isotherm measurements*).

#### 4 Conclusions

Mesoporous-TiO<sub>2</sub> (M-TiO<sub>2</sub>) and mesoporous vanadium-doped TiO<sub>2</sub> (M-V doped TiO<sub>2</sub>) at 2.3, 3.3 and 4.9 wt% vanadium-doped TiO<sub>2</sub> were successfully synthesized via reflux technique. All mesoporous vanadium-doped TiO<sub>2</sub> (M-V doped TiO<sub>2</sub>) powders performed in a highly transparent mode in the visible region at 554 nm ( $E_g = 2.24$  eV) and 588 nm  $E_g = 2.12$  eV for the 3.3 and 4.9 wt% V doped TiO<sub>2</sub>, respectively. All the prepared mesoporous materials were in anatase phase (tetragonal,  $I4_1/amd$ ). The isotherm type of the pure TiO<sub>2</sub> (M-TiO<sub>2</sub>) and the 2.3 wt% vanadium-doped TiO<sub>2</sub> were of type IV with a profile of type H2 hysteresis loops, while the 3.3 and 4.9 wt% vanadium-doped TiO<sub>2</sub> reflected isotherm type III.

The smallest pore size of 18.45 nm was obtained for pure TiO<sub>2</sub> (M-TiO<sub>2</sub>), while the largest pore size of 41.20 nm was obtained for 4.9 wt% vanadium-doped TiO<sub>2</sub> (M-V doped TiO<sub>2</sub>).

### Acknowledgements

The research was funded by the Directorate General of Higher Education, Ministry of Education and Culture, Republic of Indonesia based on PUPT 2014 Grant, No. 230/UPT-BOPTN/UN34.21/2014

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