

Indonesian Journal of Science & Technology



Journal homepage: <u>http://ejournal.upi.edu/index.php/ijost/</u>

Photodecomposition Profile of Organic Material during the Partial Solar Eclipse of 9 March 2016 and Its Correlation with Organic Material Concentration and Photocatalyst Amount

Asep Bayu Dani Nandiyanto *, Desri Sofiani, Novie Permatasari, Tranmissia Noviska Sucahya, Ani Siti Wiryani, Annisa Purnamasari, Arvina Rusli, and Eka Cahya Prima

Departemen Kimia, Universitas Pendidikan Indonesia, Jl. Dr. Setiabudi no 229, Bandung 40154, Indonesia

* Correspondence: Nandiyanto@upi.edu (A.B.D. Nandiyanto); Tel.: +62-22-5000-279

ABSTRACT

The present study investigated the photodecomposition of an organic material during the partial (88.76%) solar eclipse phenomenon of 9 March 2016 in Bandung, Indonesia. Curcumin and anatase titanium dioxide (TiO₂) particles were used as models of organic material and photocatalyst, respectively. The influence of the process parameters (i.e., curcumin concentration and the amount of TiO₂) on the photodecomposition process was also investigated. The results showed that the curcumin was decomposed along with the solar light irradiation time. During the solar eclipse, the photodecomposition rate is suddenly down. The partial solar eclipse provided a slower photodecomposition process than the process under sunny days (conducted on 8 and 10 March 2016). The concentration of curcumin and the amount of TiO₂ also have played an important role in the photodecomposition, in which the lower concentration of curcumin and the higher amount of TiO₂ have a correlation to the obtainment of higher photodecomposition rate. © 2016 Tim Pengembang Journal UPI

ARTICLE INFO

Article History: Submitted/Received 03 May 2016 First Revised 31 May 2016 Accepted 16 June 2016 First Available online 09 Aug 2016 Publication Date 01 Sep 2016

Keyword:

Curcumin, Organic material, Titania, Light intensity.

1. INTRODUCTION

Studies on photodecomposition of organic materials using metal oxide catalysts (such as TiO₂, ZnO, WO₃, and others) have attracted tremendous attention.(Kormann *et al.*, 1991) Photodecomposition process has shown considerable potentials for research and applications, particularly for solar-related applications, such as waste water treatment, dye-sensitized solar cell, and many others.

A number of studies have reported the photodecomposition development of process. Their reports have been completed with information about parameters involved in the photodecomposition, such as photocatalyst dosage, (Eskizeybek et al., 2012) initial concentration of organic materials,(Matthews 1991) properties of organic materials and catalyst, (Iskandar et al., 2007; Nandiyanto *et al.,* 2009; Nandiyanto et al., 2013; Arutanti et al., 2014) temperature, pH,(Sun et al., 2008) reaction volume, light intensity, type of light sources(Matthews 1991; Eskizeybek et al., 2012) (i.e., artificial UV/visible light source and natural sunlight), turbidity, oxygen gas flow rate , additional chemical (i.e., oxidant,(Sun et al., 2008), reducing agent, co-catalyst, (Arutanti et al., 2014; Arutanti et al., 2015) and dopant(Arutanti et al., 2015)), and consideration of inhibitor. However, there is no report on the photodecomposition of organic materials during the solar eclipse phenomenon. In fact, when the solar eclipse occurs, the Sun is masked by the Moon and this causes the darkness for a shorter duration.(Fabian et al., 2001; Foken et al., 2001; Nishanth et al., 2011) The solar intensity, light spectrum, and UV irradiation decrease, (Fabian et al., 2001; Foken et al., 2001; Koepke et al., 2001; Zerefos et al., 2001; Nishanth et al.,

2011) and this disturbs the light-dependent chemical reaction in the Earth, such as photosynthesis process.(Abram *et al.*, 2000)

Solar eclipse phenomenon has attracted researchers from various fields. Many investigated the researchers incidents related to the meteorological parameters (e.g., temperature, (Foken et al., 2001; Nishanth et al., 2011) wind speed, (Nishanth et al., 2011) gravity,(Altadill et al., 2001; Šauli et al., 2006) humidity,(Nishanth et al., 2011) geomagnetic field, (Özcan et al., 2004) ionospheric electron content,(Liu et al., 1999) and light irradiation(Koepke et al., 2001; Zerefos et al., 2001; Nishanth et al., 2011; Libra *et al.*, 2016)) and the photochemical properties in the atmosphere(Hunt 1965; Bojkov 1968; Eastman et al., 1980; Abram et al., 2000; Chudzyński et al., 2001; Tzanis et al., 2008; Sharma et al., 2010; Nishanth et al., 2011). However, no research investigates on the impact of solar eclipse the to photodecomposition of organic material in the Earth. Although some reports examined the photochemical properties, (Hunt 1965; Bojkov 1968; Eastman et al., 1980; Abram et al., 2000; Chudzyński et al., 2001; Tzanis et al., 2008; Sharma et al., 2010; Nishanth et al., 2011) they were limited to the chemical components in the atmosphere and did not consider the incidents in the ground, such as photodecomposition process.

Taking a lesson from our previous approaches in the photocatalytic processes, (Iskandar et al., 2007; Nandiyanto et al., 2009; Nandiyanto et al., 2013; Arutanti et al., 2014; Arutanti et al., 2014; Arutanti et al., 2015; Nandiyanto et al., investigated 2016) we the photodecomposition of organic materials during the partial solar eclipse phenomenon in Bandung, Indonesia. As models of organic material and photocatalyst, curcumin and anatase titanium dioxide (TiO2) powder were used, respectively.

Curcumin was selected because it can be easily obtained from turmeric, which is commonly used and largely available in Asia.(Sogi et al., 2010) Therefore, understanding the photodecomposition of curcumin will give benefits due to high usage of this material (e.g., food, drugs, antioxidant, solar-related materials, and many others). Furthermore, the amount of curcumin in the aqueous solution relates to the color and the turbidity of the mixed solution. Therefore, the decrease in its concentration can be distinguished easily even using direct visual observation. (Bonsnes et al., 1945) Although curcumin can be decomposed by exposing light, its decomposition rate is relatively low.(Zebib et al., 2010) Thus, curcuminoid compounds are used as a photoprotective molecules for lignocellulosic materials. Curcumin is also highly stable to heat(Sogi et al., 2010). Thus, the change in the temperature will have no impact to the decomposition of curcumin. These reasons make photodecomposition of curcumin being studied. (Kumavat et al., 2013; Buddee et al., 2014)

Another material used in the present study was TiO2 as the photocatalyst. This material was selected due to its affinity for light, and its ability to degrade organic compounds and transfer energy (known as solar cell).(Nandiyanto *et al.*, 2009) In addition, this material is relatively cheap, easily accessible, and non-toxic (Buddee *et al.*, 2014). Anatase-TiO₂ was used because it has been proven to be the best for photocatalytic activity.

To simplify the investigation on the photodecomposition process during the solar eclipse, we conducted an experimental procedure in a 500-mL batch-process photoreactor system. The reactor was equipped with a mixer, bubbler (connected to the air compressor), and several sensors (i.e., a light sensor, a voltage measurement analysis, a thermocouple, conductivity meter, and pH meter). The sensors were connected to the computer system; thus, a real-time decomposition of curcumin in the solution can be investigated precisely.

Different from other reports on the photodecomposition process that utilize spectrophotometer for measuring the concentration, this study employed a turbidity analysis based on the light penetration into the curcumin solution to real-time measure the analysis of concentration. In addition, the proposal of the photodecomposition mechanism during the solar eclipse phenomenon was added to explain qualitatively the experimental observations.

As the present study focused on the effect of solar eclipse on the photodecomposition of curcumin, commercial anatase titanium dioxide particles were used to simplify the experimental procedure. To confirm the photodecomposition behavior of curcumin during the solar eclipse, the photodecomposition process in the solar eclipse day (9 March 2016) and two sunny days (8 and 10 March 2016) were compared. The influence of the process parameters (i.e., curcumin concentration (10 and 25 ppm) and the amount of TiO2 (3 and 6 ppm) on the photodecomposition process was examined. In addition, also higher concentration of curcumin and TiO2 in this study was not used because of limitation of our sensor (higher concentrations of these materials have correlation to lowering preciosity of the measurement analysis).

In addition, although this study has no direct correlation to the industrial applications, understanding the phenomena during the solar eclipse is important in the scientific point of view. In general, this study can be used for assisting further scientific studies and technological developments concerning the solar-related applications.

2. HYPHOTETICAL PHOTODECOMPOSITION OF ORGANIC COMPONENT USING PHOTOCATALYST AND SOLAR ECLIPSE PHENOMENON

2.1. Photodecomposition of organic component using TiO₂ photocatalyst

The illustration of the photodecomposition of organic material using a photocatalyst in the common process is shown in **Figure 1**. In the common process, the photocatalytic activity is mainly influenced by the existence of photon, which is provided from the solar light. (Sucahya *et al.*, 2016)

The catalyst is activated by appropriate amount of energy from photons (**Route R1**). Photon leads to the formation of holes (h^{+}) in the valence band (VB) and electrons (e⁻) in

the conduction band (CB) (**Route R2**) . TiO₂ (as a model of catalyst) with a band gap of 3.35 eV is activated by absorbing UV light.(Arutanti, *et al.*, 2014; Kormann, *et al.*, 1991;Devipriya & Yesodharan, 2005)

When there are e^{-} and h^{+} in the catalyst, unique phenomenon (known as charge separation) occurs. In the presence of water, e^{-} can attract and convert oxygen (O₂) (**Route R3**), whereas h^+ can convert water molecule (H₂O) (Route R4).(Arutanti, 2014; Arutanti, 2015) Both reactions enhance the production of OH radicals, in which the OH radical is effective for degrading organic compound $(C_xH_vO_z)$ (Route R5).(Arutanti, 2014; Arutanti, 2015; Arutanti, 2014; Nandiyanto, 2013; Kormann, 1991; Devipriya & Yesodharan, 2005) Therefore, when there is a curcumin in the photocatalytic system, the concentration of presumably curcumin would decrease during the solar light illumination.(Buddee, 2014)



Figure 1. The proposal of photodecomposition of organic material (C_xH_yO_z) using TiO₂ catalyst under solar light irradiation

2.2. Solar eclipse phenomenon

Solar eclipse is an astronomical phenomenon when the Sun's disc is masked by the Moon. During the solar eclipse, several incidents can be observed, such as the changing color of the sky, the moving shadow, shadows bands, the Sun's chromospheres, the corona and its prominences, the diamond ring as well as planets and stars that suddenly become visible against the darkening sky.(Fabian, et al. 2001) The solar eclipse also causes almost complete darkness for a shorter duration.(Nishanth, et al., 2011; Fabian, et *al.*, 2001; Foken, *et al.*, 2001)

An illustration of phases occurring during the solar eclipse phenomenon is depicted in **Figure 2**. There are three types of solar eclipses depending on the distance of the Moon from the Earth:(Tiwari, 2015)

- (i) Total solar eclipse occurs when the Moon obscures the Sun (see Figure 2a). In this type of eclipse, the apparent size of the Moon is bigger than that of the Sun, allowing the solar corona to be visible. This eclipse happens only in umbra region (the darkest part of the Moon's shadow).
- (ii) Annular solar eclipse occurs when the Sun and the Moon are exactly in line but the Moon does not cover the Sun completely (see Figure 2b). In this type of eclipse, the Moon is far from the Earth; thus, the apparent size of the Moon is smaller than that of the Sun, allowing the bright ring (annulus) to be visible. This eclipse happens only in antumbra region (the part of the Moon's shadow that extends beyond the umbra).
- (iii) Partial solar eclipse occurs when the Sun and the Moon are not exactly in line and the Moon partially obscures the Sun (see Figure 2c). This eclipse happens only in the penumbra region (the weak or the pale part of the Moon's shadow).

During the solar eclipse, the following phases happen:

(i) The first contact. The edge of the Moon starts to overlap the edge of the Sun.

(ii) The second contact. Almost the entire disc of the Sun is covered by the Moon.

(iii) The maximum eclipse phase. For the total solar eclipse, the Moon completely covers the Sun's disc. For the annular and the partial solar eclipse, the distance between the center of the Moon's disc and the Sun's disc is minimum. In addition, in the total solar eclipse and the annular solar eclipse, this phase is known as totality and annularity, respectively.

(iv) The third contact. The Moon starts to move away from the Sun's disc. The maximum eclipse phase end.

(v) The fourth contact. The Moon's disc stops to overlap the Sun's disc. The solar eclipse ends.

The interval between the second and the third contacts is known as the length of maximum eclipse phase. For the partial solar eclipse, the second and the third contacts cannot be observed easily; hence, these phases sometimes are known as the **partial phase I** and **II**, respectively.

addition, In when solar eclipse phenomenon occurs, solar intensity, light spectrum, and UV irradiation are suddenly down,(Nishanth et al., 2011; Zerefos, et al., 2001) specifically just after the first contact.(Founda, et al., 2007; Zerefos, et al., 2001) Therefore, this condition disturbs some photochemical reactions both in atmosphere and in the ground.(Sharma, et al., 2010; Tzanis, et al., 2008; Bojkov, 1968; Nishanth, et al., 2011; Eastman & Stedmand, 1980;Hunt, 1965; Chudzyński, et al., 2001; Abram, et al., 2000).



Indeed, when the photodecomposition process of organic material was conducted during the solar eclipse, the concentration of organic material was presumably constant. Then, when the solar eclipse ended, the photodecomposition would occur normally.

3. EXPERIMENTAL METHOD

3.1. Materials

The study was conducted using the following raw materials: curcumin and TiO_2 particles (anatase titanium dioxide; Bratachem Co., Indonesia). Curcumin was extracted from Turmeric (Curcuma longa L) collected from Bandung, Indonesia. The purchased TiO_2 was used without further purification.

To get the extracted curcumin, a piece of turmeric was washed and cut into small pieces (sizes of about 1×1 cm). The small pieces of turmeric were then dried at 70°C to remove the water content in the turmeric. Next, the dried sample was dissolved into the ethanol solution (95%) and heated at 50°C for 1 hour in the water bath. The heated solution was then filtered, and the filtrate was subsequently put into the rotary evaporator (at 90°C) to get curcumin gel.

3.2. Photodecomposition process

The photodecomposition of curcumin investigations was carried out at the experimental eld of the Universitas Pendidikan Indonesia (6.86340 SL,107.59430 EL) in Bandung, Indonesia (See **Figure 3**). The site was selected because Bandung was thorough the path of the solar eclipse (about 88.76%) on 9 March 2016.

In the experimental procedure for the photodecomposition process, curcumin and TiO_2 particles were dissolved in the aqueous solution and placed in the batch photoreactor. The mixed suspension was then illuminated by a solar light naturally. The reactor itself consisted of a batch-glass reactor (500 mL), a magnetic stirrer (800 rpm), and a bubbler. The reactor was also

equipped with several sensors (i.e., a light sensor (lux meter; BH1750FVI, Rohm Co. Ltd., Japan), a voltage measurement analysis (DIY Mini Solar Cell; 3V, 0.42 Watt; sizes of 54 x 54 x 3 mm; Guangzhou Future Solar Technology Co. Ltd., China), and a thermocouple. During the photodecomposition process, 600 mL/min of air was bubbled into the reactor to keep the concentration of dissolved oxygen in the solution constant.



Figure 3. The path of the total solar eclipse over Indonesia on 9 March 2016. The field site was located at Universitas Pendidikan Indonesia, Bandung, about 550 km from the central axis of the eclipse totality, and experienced a 88,76 % eclipse at 07:21 local time (LT) (00:21 Universal Time (UT)). The insert figures in the right side are the illustration of solar eclipse at 100%, 90%, and 80% of coverage of Moon's disc to the Sun's disc. The map was adopted from http://asiapacific.anu.edu.au/mapsonline/base-maps/indonesia-relief , whereas the solar eclipse path was from http://eclipse.gsfc.nasa.gov/SEgoogle/SEgoogle2001/SE2016Mar09Tgoogle.html and Badan Meteorologi, Klimatologi, dan Geofisika (BMKG) Indonesia.

The concentrations of curcumin were 10 and 25 ppm, whereas the amounts of TiO_2 were 3 and 6 ppm. As a comparison, the experiment using curcumin solution only (without additional TiO_2) was also conducted. In this study, the photodecomposition process in the solar eclipse day (9 March 2016) and two sunny days (8 and 10 March 2016) was compared.

3.3. Characterizations

To confirm the type of raw materials used in this study, several analyses were conducted, such as a Scanning Electron Microscope (SEM; JEOL JSM-6360 LA, JEOL Ltd., Japan), an X-Ray Diffraction (XRD; PANalytical X'Pert PRO PW3040/x0, PANalytical, the Netherlands), and a Fourier Transform Infra Red (FTIR; Prestige 21, Shimadzu Corp., Japan)).

To measure the concentration of curcumin over the photodecomposition process time, we approximated the concentration using a turbidity analysis based on the light penetration into the mixed suspension. The analysis was conducted in the batch photoreactor based on the light sensor and voltage measurement analysis that were integrated with the computer system to get information in the real-time measurement of concentration. A spectrophotometer (UV-Vis mini 1240, Shimadzu Corp., Japan) was also used to confirm the concentration of curcumin gained from the turbidity analysis in the real-time measurement.

To confirm the analysis of the photodecomposition process, measurement of the solar intensity was conducted using a lux meter (BH1750FVI, Rohm Co. Ltd., Japan) and an ultraviolet sensor (UV sensor; ML8511). To measure the solar irradiation precisely, all sensors to the computer were connected. Since the solar eclipse occurred in the morning, the sensors were oriented to be perpendicular to the ground to get the maximum absorption of solar light.

In addition, the Langmuir–Hinshelwood kinetic model was used to calculate the photodecomposition rate that involves

solid–solute photochemical reactions. Based on this model, the photodecomposition rate can be written as

$$-\frac{dC}{dt} = \frac{k_1 \cdot k_2 \cdot C}{1 + k_2 \cdot C}$$
(1)

where *C* and *t* are the concentration and the photoreaction time, respectively. k_1 and k_2 are the apparent reaction rate constant and the apparent equilibrium constant for adsorption of the chemical on the catalyst surface, respectively.(Nandiyanto, *et al.*, 2009) Since the concentration of curcumin used in this study was very low (less than 100 ppm), the **Equation (1)** can be re-expressed as a first-order rate constant, as follows:

$$-\frac{dC}{dt} = k.C$$
 (2)

where k is the reaction rate constant. Then, the photodecomposition rate (as k value) was obtained by solving the differential problem in the **Equation (2)**.

4. RESULTS AND DISCUSSION 4.1. Physicochemical properties of

extracted curcumin and commercial TiO₂

Figure 4 shows the FTIR results of TiO_2 particles, dried turmeric, and curcumin (obtained from the isolation of turmeric). The result showed that the peaks of dried turmeric and curcumin are almost similar. A broad peak at about 3404 cm⁻¹ indicates the presence of OH. Characteristics peaks of curcumin were identified, located at 3404, 1625, and 1512 cm⁻¹.(Kumavat, 2013)



Regarding the TiO₂ sample, several peaks at about 3448, 1382, 680, and 543 cm⁻¹ were detected in the FTIR analysis. Peaks at 3448 and 1382 cm⁻¹ related to stretching hydroxyl (O-H) and O-H bending. The other peaks at 680 and 543 cm⁻¹ was assigned to the Ti-O stretching bands. Although those peaks are a typical pattern of anatase, the detection of peaks can not be used directly to clarify the structure of material. Therefore, additional analyses (e.g., XRD) are required.

Figure 5a depicts the SEM images of TiO₂ particles used in this study. The Ferret analysis indicated that the mean size of TiO₂ particles was about 28 micrometers. **Figure 5b** shows the XRD analysis of TiO₂ particles. The detection of diffraction peaks at 20 of 25.30; 37.78; 47.88; 54.50; and 63.32° replied to the crystallographic structure of anatase phase (based on JCPDS No. 84-1286).(Nandiyanto, *et al.*, 2009) No other phase was detected in the pattern, confirming that the sample was pure anatase.



Figure 5. SEM image (a) and XRD analysis result (b) of TiO₂ particles

4.2. SOLAR RADIATION DURING THE SOLAR ECLIPSE

The photograph images of the Sun during the solar eclipse phenomenon are depicted in **Figure 6**. The day of the eclipse was characterized by mostly clear skies, with some light cloud in the beginning of the solar eclipse, slight westerly winds, and comparatively low pollution. The photograph images of Sun were taken at 06:20; 06:46; 07:08; 07:18; 07:21; 07:37; 08:00; and 08:32 LT, corresponding to **Figures 6a-h**, respectively. An illustration image attached in each photograph image was added to make clearer discussion about the partial solar eclipse phenomenon.

The first contact occurred at 06:21 LT as depicted in Figure 6a. However, since the sky was cloudy, the first contact could not be observed clearly. The first contact occurred when the edge of the Moon started to overlap the edge of the Sun. This contact indicated the beginning of solar eclipse. Then, the significant reduction of the solar radiation was observed just after the first contact(Founda, et al., 2007). The Moon obscured a few part of the Sun's disc, indicating the occurrence of partial phase (Figures 6b-d). The Sun could be clearly observed as the clouds moved away from it. This partial phase continuously occurred until the distance between the center of the

Moon's disc and the Sun's disc was minimum (Figure 6e). As the Moon blocked the Sun, the sky turned gray. Since Bandung experienced the maximum Sun's obscuration about 88.76% at 07:21 LT (00:21 UT) for about two minutes, the Sun was observed to be red and crescent shape by the camera. The ending of the maximum phase was indicated when the Moon's disc moved away from the Sun, and the Sun reappeared (Figures 6f-g). At 08:32 LT (Figure 6h), the Moon stopped overlapping the Sun's disc and the daylight returned. This step was known as the fourth contact, indicating the ending of solar eclipse.



Figure 6. Photograph images of the Sun during the partial solar eclipse with illustration model. The pictures were taken in the specific local time: (a) 06:20; (b) 06:46; (c) 07:08; (d) 07:18; (e) 07:21; (f) 07:37; (g) 08:00; and (h) 08:32 LT. **Figure 7** shows a plot of the solar radiation intensity during three sunny days (from 8 to 10 March 2016) between near to sunrise (06:00 LT) and morning (09:00 LT). The results showed that the intensity of the solar radiation increased along with the

increasing time from sunrise to morning. The results were in a good agreement with the Kimball result that the midday has richer luminous radiation than the morning. (Kimball, 1924)



The intensity of solar radiation during the solar eclipse (9 March) was identified to be lower than that during two sunny days (8 and 10 March). The intensities in both two sunny days were from about 300 to 45,000 lux with the rate of increasing solar intensity was about 1,200 lux/min. The rate of increasing solar intensity during the solar eclipse depends on the phases. Before the partial phase I (06:00-06:20 LT), the rate was similar to that in the sunny days. The rate during the partial phase I (06:20-07:21 LT) was about 75 lux/min. Then, in the maximum eclipse phase (07:21-07:23 LT), the rate was almost zero. Finally, in the partial phase II (07:23-08:32 LT), the rate was about 1,430 lux/min.

The decreases in the radiation intensity in the partial phase I were due to the increases in the blockage area of the Sun by the Moon's disc. Then, in the maximum phase, the rate was almost zero because the blockage of Moon disc reached the maximum. The intensity in the maximum phase was about 10,000 lux, whereas the intensity of sunny day was more than 60,000 lux with regard to the minor differences of measurement conditions in two consecutive sunny days in 8 and 10 March 2016. Approximating the ratio of intensities, we found that the intensity in the maximum eclipse phase was at least 16% of the intensity of solar radiation in the sunny time. During the eclipse, part of the Sun's disc was covered by the Moon, leading the reduction of solar radiation intensity(Koepke, et al., 2001). The result was in a good correlation to the shinning area of the Sun's disc that was about 12% (the coverage area of the Sun's disc by the Moon in Bandung was about 88.76%; see Figure 6e). The decrease in solar intensity

during the eclipse has been mentioned in the current literatures (Libra, 2016;Koepke, 2001) When the partial phase II started, the rate increased rapidly (rate = 1,430 lux/min), confirming the appearance of the Sun to reach normal. In addition, the intensities after 08:20 in all samples fluctuated between 30,000 and 60,000 mV due to the hysteresis and the limitation of our lux sensors.

A plot of the UV radiation during the three sunny days between near to sunrise and morning is shown in **Figure 8**. The results showed that the intensity of UV radiation increased along with the increasing time from sunrise to morning.



Figure 8. The profile of UV radiation over time during the solar eclipse day and two sunny days.

For the two sunny days, the intensity increased with the increasing rate of 0.02 mW/cm².min. However, the intensity of UV radiation during the solar eclipse was identified to be lower than that during two sunny days. Prior to the partial phase I, the increasing rate was similar to that during two sunny days. In the partial phase I from 06:20 to 07:00 LT, the intensity of UV radiation increased (increasing rate of about 0.006 mW/cm^2 .min). Then, the intensity decreased when reaching the maximum phase. The UV intensity did not reach zero because Bandung experienced the partial solar eclipse. Thus, Sun still provides light during the maximum phase. Finally, when the maximum phase passed, the intensity drastically increased (increasing rate in the partial phase II of about 0.05 mW/cm².min). The present result in the profile of UV radiation was in a good agreement with the current literatures.(Zerefos, et al., 2001; Koepke, *et al.*, 2001)

4.3. Photodecomposition of curcumin under TiO₂ powder during the solar eclipse and sunny days

The UV-vis spectra of 25-ppm curcumin solutions photodecomposed by TiO₂ particles during the partial solar eclipse is depicted in **Figure 9**. The insert figures are the photograph image of curcumin solution during the solar eclipse, respectively. **Figures 9a** and **b** are the curcumin solutions with 6 and 3 ppm of TiO₂ particles, respectively.

Curcumin concentrations during the experiments were determined based on the Beer Law(Arutanti, et al., 2014; Arutanti, et 2015; Nandiyanto, et al., 2009; al., Nandiyanto, 2013) at the absorbance of 425 nm wavelength light. At the initial condition (at 06:00 LT), curcumin solution spectra had a maximum absorption at a wavelength of around 425 nm, confirming the yellow color. at all The absorbance wavelengths decreased over the time, which was due to the degradation of curcumin during the partial solar eclipse.(dos Santos, et al., 2014) The decreases in absorbance of curcumin solution over time during the solar eclipse were in a good agreement with the change of color by a visual observation (displayed in the insert figures in Figures 7a and b), in which the color of solution became brighter.

In the case of sample containing 6 ppm of TiO_2 (Figure 7a), the absorbance decreased down to about 60% at 06:20 LT. Further decreases in the absorbance were found at 07:21 LT (in the beginning of maximum solar eclipse) and 09:00 LT (in the end experiment), in which the of absorbances were about 40 and 20%, respectively. Regarding the sample containing 3 ppm of TiO_2 (Figure 7b), the absorbance decreased down to about 75% at 06:20 LT and 55% at 09:00 LT. Based on the results, the final absorbance of sample containing 6 ppm of TiO₂ was lower than that containing 3 ppm of TiO₂. The result confirms that the TiO₂ plays a role in the photodecomposition process.



Figure 9. UV-vis spectra analysis results of samples containing 25-ppm curcumin with TiO_2 catalyst during the partial solar eclipse. Figures (a) and (b) are the samples with 6 and 3 ppm of TiO_2 particles, respectively.

4.3.1. Effect of solar eclipse

When suspension containing curcumin and TiO_2 was irradiated by natural sunlight, changes in physical appearances (i.e., degradation of color) were observed in all variations. The experiments conducted in both two sunny days (8 and 10 March 2016) resulted in the degradation of color from clear amber to transparent color. Then, in the solar eclipse day, the color changed from clear amber to light yellow. The different changes in color confirmed that the photodecomposition rates of solar eclipse and two sunny days were different. **Figure 10** presents the effect of solar eclipse on the photodecomposition of curcumin with various curcumin concentrations and TiO₂ amounts. **Figures 10a** and **b** are the profile of photodecomposition of 25-ppm curcumin,

whereas **Figures 10c** and **d** are the profile when using 10 ppm of curcumin. **Figures 10a** and **c** are the profile of the process using 6 ppm of TiO_2 , whereas **Figures 10b** and **d** are the process using 3 ppm of TiO_2 .



The experimental results showed that the concentration of curcumin decreased along with the photodecomposition time from sunrise to morning for all variations. Then, the changes in the experimental condition allowed to the obtainment of different photodecomposition profiles.

In all variations in **Figure 10**, the experiments performed in the solar eclipse day (showed as dashed line) had a lower photodecomposition rate than that in the sunny days (showed as solid line), confirmed

by the higher final concentration (C/Co). The fundamental reason for the lower photodecomposition rate in the solar eclipse because the solar eclipse day is phenomenon provides the decreases in the solar radiation and UV intensity (verified in Figures 7 and 8). Light intensity is a major factor in the photocatalytic reactions because electron-hole pairs are produced only by the light. As the consequence, the decrease in light intensity would disturb the photodecomposition process.(Abram, et al., 2000)

To confirm the effect of the solar eclipse condition on the photodecomposition of the rates for the photocurcumin, decomposition process of 25 ppm of curcumin were calculated based on five phases in the solar eclipse (see **Table 1**): (i) before the solar eclipse (at 06:00-06:20 LT), (ii) the partial phase I (at 06:20-07:21 LT), (iii) the maximum phase (at 07:21-07:23 LT); (iv) the partial phase II (at 07:23-08:32 LT), and (v) after the solar eclipse (at 08:32-09:00 LT). As shown in this table, the rate of photodecomposition during the solar eclipse was slower compared with that in the sunny day.

Before solar eclipse, the rate of photocatalysis in the solar eclipse day was identical to that in the sunny day, indicated by the identical *k* values for samples conducted in solar eclipse and sunny day. The identical light intensities (as shown in **Figures 7** and **8**) in this phase were the main reason for the same photochemical reaction rates in all samples.

In the partial phase I, the rate constant of photodecomposition in the solar eclipse day obtained was slower than that in sunny days. For the case of samples using 25 ppm of curcumin and 6 ppm of TiO₂, conducting the photodecomposition process in the solar eclipse and the sunny days resulted in the k values of 1.47 10^{-2} and 1.74 10^{-2} min⁻¹, respectively. The fundamental reason is due to the decreases in solar intensity that affects the intensity of UV radiation.

In the maximum eclipse phase, the rate constant of photodecomposition of 25 ppm of curcumin with 6 ppm of TiO₂ in the solar eclipse reached was 0.97 10⁻² min⁻¹. The different concentrations of samples before and after the maximum phase were about 1%. In this phase, the rate did not reach to because experimental zero the site (Bandung) experienced the partial solar eclipse event. Therefore, the Sun still provides solar light for the photodecomposition process.

Next, the partial phase II had similar trends to the partial phase I and the maximum phase, where the photodecomposition rates of samples conducted in the solar eclipse day were lower than that in the sunny days. Finally, after the solar eclipse, the rates were still lower. This was due to the limitation of photodecomposition process time (conducting from 08:32 to 09:00 LC).

Sample*	Final concentration at 9:00 LT (%)	<i>k</i> value (min ⁻¹)				
		Before SE* (06:00 – 06:21 LT)	Partial phase I (06:20 – 07:21 LT)	Maximum phase (07:21 – 07:23 LT)	Partial phase II (07:23 – 08:32 LT)	After SE* (08:32 – 09:00 LT)
Curcumin only	95	0.21 x 10 ⁻²	0.06 x 10 ⁻²	0.04 x 10 ⁻²	0.04 x 10 ⁻²	0.03 x 10 ⁻²
$TiO_2 = 6 ppm (SD)$	6	4.60 x 10 ⁻²	1.74 x 10 ⁻²	1.45 x 10 ⁻²	1.39 x 10 ⁻²	1.57 x 10 ⁻²
$TiO_2 = 6 ppm (SE)$	30	4.61 x 10 ⁻²	1.47 x 10 ⁻²	0.97 x 10 ⁻²	0.87 x 10 ⁻²	0.69 x 10 ⁻²
$TiO_2 = 3 ppm (SD)$	23	2.15 x 10 ⁻²	0.84 x 10 ⁻²	0.72 x 10 ⁻²	0.80 x 10 ⁻²	0.81 x 10 ⁻²
$TiO_2 = 3 ppm (SE)$	60	2.19 x 10 ⁻²	0.58 x 10 ⁻²	0.40 x 10 ⁻²	0.35 x 10 ⁻²	0.29 x 10 ⁻²

Tabel 1. The rate of photodecomposition of 25-ppm curcumin under various conditions

Note: *SE is the experiment provided in solar eclipse, and SD is experiment conducted in sunny day

4.3.2. Effect of photocatalyst amount

The profile of photodecomposition of curcumin using various catalyst amounts (i.e., 0, 3, and 6 ppm of TiO_2) is shown in Figure 11. The reaction was performed from near sunrise (06:00 LT) to morning (09:00 LT), which was an ample time for comparing the effect of catalyst amount on the photodecomposition rate. To confirm the effect of catalyst amount, the figure based on the constant curcumin concentration was classified. The profile of photodecomposition process using 25 ppm of curcumin is depicted in Figures 11a and c, and that using 10 ppm of curcumin is in Figures 11b and d. In addition, to confirm that catalyst has an impact on the photodecomposition in all process conditions, the catalyst amounts were varied in the experiments conducted in the sunny day (Figures 11(a-b)) and the solar eclipse day (Figures 11 (c-d)).

The results showed that the concentration of curcumin decreased along with the photodecomposition time from sunrise to morning, and the photodecomposition rate increased as the increases in the amount of TiO₂. In contrast,

the photodecomposition rate was the lowest for all cases in the absence of the TiO_2 .

The photodecomposition rate is strongly influenced by the number of active sites and the photoabsorption ability of the catalyst (Sakthivel, et al., 2003). The increase in number of active sites available would have a positive impact to the increasing photocatalytic activity and efficiency. This was confirmed by the fact that the absence of the TiO₂ resulted in the obtainment of the lowest photodecomposition rate in both solar eclipse and sunny day cases. In general, the higher amount of TiO₂ catalyst resulted in more effective photocatalytic process than that using lower concentration of TiO₂. This corroborates the hypothesis of other studies.(Zhu, et al., 2014; Sobana & Swaminathan 2007; Mills, 1993) In addition, although the concentration of curcumin decreases, the required processing time for curcumin decomposing takes longer compared to previous study. It occurs due to the use of less concentration of catalyst.



Figure 11.(a-b) Effect of TiO₂ amount on the photodecomposition of curcumin. Figures (a) and (c) are the samples using 25 ppm of curcumin, whereas Figures (b) and (d) are using 10 ppm of curcumin. Figures (a) and (b) are the process conducted in the sunny day.



respectively.

4.3.3. Effect of initial organic material concentration

Figure 12 presents the effect of initial concentration of curcumin (i.e., 10 and 25 ppm) on the photodecomposition rate during the sunny day (from near to sunrise (06:00 LT) to morning (09:00 LT)). The profile of photodecomposition process using 6 ppm of TiO₂ is shown in Figures 12a and c, whereas that of the process using 3 ppm of TiO₂ is shown in Figures 12b and d. The comparison of the process in the sunny day and the solar eclipse day is shown in Figures 12 (a-b) and (c-d), respectively. The result showed that the concentration of curcumin decreased along with the photodecomposition time from sunrise to morning.

As shown in **Figures 12a** and **b**, at the constant TiO_2 amount, the photodecomposition rate depended on the initial curcumin concentration. The lower concentration of curcumin led to the obtainment of faster photodecomposition process.

The photodecomposition of curcumin during the partial solar eclipse (shown in **Figures 12c** and **d)** yielded some unique

phenomena. Although the curcumin concentration varied, the decomposition profile during the solar eclipse (specifically from 06:20 to 07:23 LT) was almost similar. Also, the concentration of curcumin was almost unchanging during the eclipse. It started to decrease drastically after the passing the solar eclipse (after 08:32). The results confirmed that the solar eclipse phenomenon had higher impact to the photodecomposition process than the initial concentration of curcumin. In addition to factor, the effect of curcumin this concentration on the photodecomposition rate was obtained. The use of lower concentration of curcumin had faster photodecomposition process than that of higher concentration.

When high concentration of organic molecules (i.e., curcumin) is used, more organic molecules absorb on the surface of catalyst. The amount of absorbed organic molecule is thought to have an inhibitive effect. Further, high concentration of organic molecules may absorb a significant amount of solar light (photon).(Mills, *et al.*, 1993) Thus, the less number of photons are available to reach the catalyst surface, making fewer catalyst areas be activated by solar light and less OH radicals be formed.(Byrappa, *et al.*, 2006) Therefore, increases in the concentration have a negative impact to the photodegradation rate.(dos Santos, *et al.*, 2014; Zhu, *et al.*, 2014; Sobana, *et al.*, 2007; Mills, *et al.*, 1993; Byrappa, *et al.*, 2006).



Figure 12. Effect of initial curcumin concentration on the photodecomposition of curcumin.
 Figures (a) and (c) are the samples using 6 ppm of TiO₂, whereas Figures (b) and (d) are using 3 ppm of TiO₂. Figures (a) and (b) are the process conducted in the sunny day, whereas Figures (c) and (d) are that performed in the solar eclipse day. SE and SD are the samples for solar eclipse and sunny day, respectively.

5. CONCLUSION

This study investigated the behavior of photodecomposition of organic material during the partial (88.76%) solar eclipse phenomenon of 9 March 2016 in Bandung, Indonesia. In the experiments, curcumin and anatase titanium dioxide_particles were used as models of organic material and photocatalyst, respectively. These materials were dissolved in water, mixed, and placed into the reactor that was integrated to the computer system to get a real-time concentration analysis. The results showed that the solar eclipse provides a change in the solar light intensity. Indeed, the intensity disturbed the photodecomposition process. To confirm the influence of solar eclipse phenomenon on the photodecomposition process, curcumin concentration and the amount of catalyst were also varied.

6. ACKNOWLEDGEMENTS

A.B.D.N. and N.P. acknowledged RISTEK

DIKTI (Grant: Program Unggulan Perguruan Tinggi (PUPT) and Program Kreativitas Mahasiswa (PKM-P)). The authors also thank to Dian Herdiana and Togema (Tim Observasi Gerhana Matahari) UPI for their assistance in this experiment.

7. AUTHORS' NOTE

The author(s) declare(s) that there is no conflict of interest regarding the publication of this article. Authors confirmed that the data and the paper are free of plagiarism.

8. REFERENCES

- Abram, J. P., D. J. Creasey, D. E. Heard, J. D. Lee and M. J. Pilling (2000). Hydroxyl radical and ozone measurements in England during the solar eclipse of 11 August 1999. *Geophysical research letters*, 27(21), 3437-3440.
- Altadill, D., J. Sole and E. Apostolov (2001). Vertical structure of a gravity wave like oscillation in the ionosphere generated by the solar eclipse of August 11, 1999. *Journal of geophysical research: Space physics, 106*(A10), 21419-21428.
- Arutanti, O., A. B. D. Nandiyanto, T. Ogi, F. Iskandar, T. O. Kim and K. Okuyama (2014). Synthesis of composite WO₃/TiO₂ nanoparticles by flame-assisted spray pyrolysis and their photocatalytic activity. *Journal of alloys and compounds*, *591*, 121-126.
- Arutanti, O., A. B. D. Nandiyanto, T. Ogi, T. O. Kim and K. Okuyama (2015). Influences of Porous Structurization and Pt Addition on the Improvement of Photocatalytic Performance of WO₃ Particles. *ACS applied materials and interfaces*, 7(5), 3009-3017.
- Bojkov, R. D. (1968). The ozone variations during the solar eclipse of 20 May 1966. *Tellus*, 20(3), 417-421.
- Bonsnes, R. W. and H. H. Taussky (1945). On the colorimetric determination of creatinine by the Jaffe reaction. *Journal of biological chemistry*, *158*(3), 581-591.
- Buddee, S., S. Wongnawa, P. Sriprang and C. Sriwong (2014). Curcumin-sensitized TiO₂ for enhanced photodegradation of dyes under visible light. *Journal of nanoparticle research*, 16(4), 1-21.
- Byrappa, K., A. Subramani, S. Ananda, K. L. Rai, R. Dinesh and M. Yoshimura (2006). Photocatalytic degradation of rhodamine B dye using hydrothermally synthesized ZnO. *Bulletin of materials science*, *29*(5), 433-438.
- Chudzyński, S., A. Czyżewski, K. Ernst, A. Pietruczuk, W. Skubiszak, T. Stacewicz, K. Stelmaszczyk, A. Szymański, I. Sowka and A. Zwoździak (2001). Observation of ozone concentration during the solar eclipse. *Atmospheric research*, *57*(1), 43-49.
- Devipriya, S. and S. Yesodharan (2005). Photocatalytic degradation of pesticide contaminants in water. *Solar energy materials and solar cells*, *86*(3), 309-348.
- dos Santos, T. C., G. J. Zocolo, D. A. Morales, G. de Aragão Umbuzeiro and M. V. B. Zanoni (2014). Assessment of the breakdown products of solar/UV induced photolytic degradation of food dye tartrazine. *Food and chemical toxicology*, *68*, 307-315.

- Eastman, J. A. and D. H. Stedman (1980). Variations in the ambient ozone concentration during the 26 February 1979 solar eclipse. *Atmospheric environment*, *14*(6), 731-732.
- Eskizeybek, V., F. Sarı, H. Gülce, A. Gülce and A. Avcı (2012). Preparation of the new polyaniline/ZnO nanocomposite and its photocatalytic activity for degradation of methylene blue and malachite green dyes under UV and natural Sun lights irradiations. *Applied catalysis B: Environmental*, *119*, 197-206.
- Fabian, P., M. Winterhalter, B. Rappenglück, H. Reitmayer, A. Stohl, P. Koepke, H. Schlager,
 H. Berresheim, T. Foken and B. Wichura (2001). The BAYSOFI Campaign–
 Measurements carried out during the total solar eclipse of August 11, 1999.
 Meteorologische zeitschrift, 10(3), 165-170.
- Foken, T., B. Wichura, O. Klemm, J. Gerchau, M. Winterhalter and T. Weidinger (2001). Micrometeorological measurements during the total solar eclipse of August 11, 1999. *Meteorologische zeitschrift*, 10(3), 171-178.
- Founda, D., D. Melas, S. Lykoudis, I. Lisaridis, E. Gerasopoulos, G. Kouvarakis, M. Petrakis and C. Zerefos (2007). The effect of the total solar eclipse of 29 March 2006 on meteorological variables in Greece. *Atmospheric chemistry and physics*, 7(21), 5543-5553.
- Hunt, B. (1965). A theoretical study of the changes occurring in the ozonosphere during a total eclipse of the Sun. *Tellus*, *17*(4), 516-523.
- Iskandar, F., A. B. D. Nandiyanto, K. M. Yun, C. J. Hogan, K. Okuyama and P. Biswas (2007). Enhanced photocatalytic performance of brookite TiO₂ macroporous particles prepared by spray drying with colloidal templating. *Advanced materials*, 19(10), 1408-1412.
- Kimball, H. H. (1924). Records of Total Solar Radiation Intensity and Their Relation to Daylight Intensity. *Monthly weather review*, *52*(10), 473-478.
- Koepke, P., J. Reuder and J. Schween (2001). Spectral variation of the solar radiation during an eclipse. *Meteorologische Zeitschrift*, *10*(3), 179-186.
- Kormann, C., D. Bahnemann and M. R. Hoffmann (1991). Photolysis of chloroform and other organic molecules in aqueous titanium dioxide suspensions. *Environmental science and technology*, *25*(3), 494-500.
- Kumavat, S. D., Y. S. Chaudhari, P. Borole, P. Mishra, K. Shenghani and P. Duvvuri (2013). Degradation studies of curcumin. *International journal of pharmacy review research*, 3(2), 50-55.
- Libra, M., P. Kouřím and V. Poulek (2016). Behavior of Photovoltaic System during Solar Eclipse in Prague. *International journal of photoenergy*, 2016, 1-6.
- Liu, J., H. Tsai, L.-C. Tsai and M. Chen (1999). Ionospheric total electron content observed during the 24 October 1995 solar eclipse. *Advances in space research*, *24*(11), 1495-1498.

- Matthews, R. W. (1991). Photooxidative degradation of coloured organics in water using supported catalysts. TiO₂ on sand. *Water research*, *25*(10), 1169-1176.
- Mills, A., R. H. Davies and D. Worsley (1993). Water purification by semiconductor photocatalysis. *Chemical society review*, 22(6), 417-425.
- Nandiyanto, A. B. D., F. Iskandar and K. Okuyama (2009). Macroporous anatase titania particle: Aerosol self-assembly fabrication with photocatalytic performance. *Chemical engineering journal*, *152*(1), 293-296.
- Nandiyanto, A. B. D., O. Arutanti, T. Ogi, F. Iskandar, T. O. Kim and K. Okuyama (2013). Synthesis of spherical macroporous WO₃ particles and their high photocatalytic performance. *Chemical engineering science*, *101*, 523-532.
- Nandiyanto, A. B. D., Munawaroh, H. S. H., Kurniawan, T., & Mudzakir, A. (2016). Influences of Temperature on the Conversion of Ammonium Tungstate Pentahydrate to Tungsten Oxide Particles with Controllable Sizes, Crystallinities, and Physical Properties. *Indonesian Journal of Chemistry*, *16*(2), 124-129.
- Nishanth, T., N. Ojha, M. S. Kumar and M. Naja (2011). Influence of solar eclipse of 15 January 2010 on surface ozone. *Atmospheric environment*, *45*(9), 1752-1758.
- Özcan, O. and M. Aydoğdu (2004). Possible effects of the total solar eclipse of August 11, 1999 on the geomagnetic field variations over Elaziğ-Turkey. *Journal of atmospheric and solar-terrestrial physics*, *66*(11), 997-1000.
- Sakthivel, S., B. Neppolian, M. Shankar, B. Arabindoo, M. Palanichamy and V. Murugesan (2003). Solar photocatalytic degradation of azo dye: comparison of photocatalytic efficiency of ZnO and TiO₂. *Solar energy materials and solar cells*, 77(1), 65-82.
- Šauli, P., P. Abry, J. Boška and L. Duchayne (2006). Wavelet characterisation of ionospheric acoustic and gravity waves occurring during the solar eclipse of August 11, 1999. *Journal of atmospheric and solar-terrestrial physics*, *68*(3), 586-598.
- Sharma, S., T. Mandal, B. Arya, M. Saxena, D. Shukla, A. Mukherjee, R. Bhatnagar, S. Nath, S. Yadav and R. Gautam (2010). Effects of the solar eclipse on 15 January 2010 on the surface O₃, NO, NO₂, NH₃, CO mixing ratio and the meteorological parameters at Thiruvanathapuram, India. *Annales geophysicae*, 28, 1199-1205.
- Sobana, N. and M. Swaminathan (2007). The effect of operational parameters on the photocatalytic degradation of acid red 18 by ZnO. *Separation and purification technology*, *56*(1), 101-107.
- Sogi, D. S., S. Sharma, D. P. S. Oberoi and I. Wani (2010). Effect of extraction parameters on curcumin yield from turmeric. *Journal of food science and technology*, *47*(3), 300-304.
- Sucahya, T.N., Permatasari, N. and Nandiyanto, A.B.D., 2016. Review: Fotokatalis untuk pengolahan limbah cair. *Jurnal integrasi proses*, *6*(2), 1-15.

- Sun, J., L. Qiao, S. Sun and G. Wang (2008). Photocatalytic degradation of Orange G on nitrogen-doped TiO₂ catalysts under visible light and sunlight irradiation. *Journal of hazardous materials*, 155(1), 312-319.
- Tiwari, N. K. (2015). Eclipse. International journal of sanskrit research, 1(3), 38-40.
- Tzanis, C., C. Varotsos and L. Viras (2008). Impacts of the solar eclipse of 29 March 2006 on the surface ozone concentration, the solar ultraviolet radiation and the meteorological parameters at Athens, Greece. Atmospheric chemistry and physics, 8(2), 425-430.
- Zebib, B., Z. Mouloungui and V. Noirot (2010). Stabilization of curcumin by complexation with divalent cations in glycerol/water system. *Bioinorganic chemistry and applications*, 2010, 1-8.
- Zerefos, C., D. Balis, P. Zanis, C. Meleti, A. Bais, K. Tourpali, D. Melas, I. Ziomas, E. Galani and K. Kourtidis (2001). Changes in surface UV solar irradiance and ozone over the Balkans during the eclipse of August 11, 1999. Advances in space research, 27(12), 1955-1963.
- Zhu, H.-Y., J. Yao, R. Jiang, Y.-Q. Fu, Y.-H. Wu and G.-M. Zeng (2014). Enhanced decolorization of azo dye solution by cadmium sulfide/multi-walled carbon nanotubes/polymer composite in combination with hydrogen peroxide under simulated solar light irradiation. *Ceramics international*, 40(2), 3769-3777.