



PERFORMANCE OF SOLAR PHOTOCATALYSIS AND PHOTO-FENTON DEGRADATION OF PALM OIL MILL EFFLUENT

(Prestasi Fotopemangkinan dan Degradasi Foto-Fenton Menggunakan Sinar Suria
ke atas Efluen Kilang Minyak Kelapa Sawit)

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Abstract

Palm oil mill effluent (POME) contains significant amounts of organic matter, solids, and grease or oil, which requires appropriate treatment prior to being discharged into the environment. In this study, solar radiation was investigated as a possible source of photon in the solar TiO₂ and ZnO photocatalysis, and solar photo-Fenton treatments to reduce the chemical oxygen demand (COD) in POME. The results indicated that solar photo-Fenton was more efficient in reducing COD levels compared to dark Fenton and indoor photo-Fenton. The highest removal was recorded at 89% in the presence of 1:30 ratio of Fe²⁺:H₂O₂ under acidic pH (~2.8) after 3 hours of solar exposure. Increased concentrations of H₂O₂ have greatly influenced the COD removal. Additionally, solar TiO₂ photocatalysis (pH~3.7; TiO₂ = 0.1 g/L) has outperformed solar photolysis and solar ZnO photocatalysis in reducing COD levels in POME. With successive increase of TiO₂ from 0.02 to 0.1 g/L, the removal of COD had linearly increased from 54.3% to 88.5% after 5 hours of solar exposure. Based on the investigated conditions, the optimum TiO₂ concentration of 0.1 g/L was concluded. In conclusion, solar TiO₂ photocatalysis and solar photo-Fenton can be applied as possible means to reduce the organic loads in POME.

Keywords: advanced oxidation process, organic matter, reduction, solar, titanium dioxide

Abstrak

Efluen kilang minyak kelapa sawit (POME) mengandungi jumlah bahan organik, pepejal dan gris atau minyak yang ketara yang memerlukan rawatan sesuai sebelum ia boleh disalurkan ke persekitaran. Dalam kajian ini, sinar suria dikaji sebagai salah satu sumber foton dalam fotopemangkinan TiO₂ dan ZnO sinar suria dan rawatan foto-Fenton menggunakan sinar suria untuk mengurangkan keperluan oksigen kimia (COD) dalam POME. Keputusan kajian menunjukkan foto-Fenton menggunakan sinar suria adalah lebih berkesan dalam menurunkan COD berbanding dengan Fenton bercahaya dalam dan Fenton tanpa cahaya. Penyingkiran tertinggi yang dicatatkan ialah 89% menggunakan nisbah Fe²⁺:H₂O₂ sebanyak 1:30 dalam pH berasid (~2.8) selepas pendedahan kepada sinar suria selama 3 jam. Peningkatan kepekatan H₂O₂ sangat mempengaruhi penyingkiran COD. Tambahan pula, fotopemangkinan TiO₂ menggunakan sinar suria (pH~3.7; TiO₂ = 0.1 g/L) menunjukkan prestasi yang lebih baik berbanding dengan fotolisis dan fotopemangkinan ZnO menggunakan sinar suria dalam mengurangkan kandungan COD dalam POME. Peningkatan TiO₂ secara berturutan dari 0.02 ke 0.1 g/L meningkatkan penyingkiran COD secara linear daripada 54.3% ke 88.5% selepas pendedahan kepada sinar suria selama 5 jam. Berdasarkan keadaan eksperimen yang dikaji dapat dirumuskan bahawa kepekatan optimum TiO₂ ialah sebanyak 0.1 g/L. Kesimpulannya, teknik fotopemangkinan TiO₂ sinar suria dan Fenton sinar suria boleh digunakan untuk mengurangkan kandungan organik dalam POME.

Kata kunci: proses pengoksidaan termaju, bahan organik, penurunan, sinar suria, titanium dioksida

Introduction

Malaysia is known as the world's largest palm oil exporter. The palm oil industry has made significant contribution to the country's economic revenue. The global demand for palm oil, particularly for crude palm oil and its related products, such as palm oil and palm kernel oil, is rapidly growing. Consequently, the palm oil industry is generating a significant amount of waste, namely palm oil mill effluents (POME). Approximately 44 million tons of POME was generated in 2008 in Malaysia [1]. Typically, raw POME is brownish in colour and contains elevated levels of organic matter, such as total solids (405,000 mg/L) and grease/oil (6000 mg/L), as well as high biochemical oxygen demand (BOD) (25,000 mg/L) and chemical oxygen demand (COD) (50,000 mg/L) [2, 3]. POME is also known to be highly acidic (pH 3.8 – 4.5) and biodegradable in nature. The open ponding system is currently used by 85% of palm oil mill operators, as the potential method to treat POME [1], due to its biodegradable nature. A ponding system comprises of a series of anaerobic, facultative and aerobic treatments. This type of system demands vast area of lands, requires lengthy treatment periods (up to 40 – 60 days) for effective treatments to be achieved, and it releases bad odour, despite its ease of operation [4, 5]. A previous study has reported that this biological treatment process is rather inefficient in treating POME [2]. It contributes to various environmental issues due to the high loadings of BOD and COD, while a low pH of POME renders the conventional treatment technique inefficient.

In recent times, advanced oxidation processes (AOPs) have been considered as suitable methods to treat effluents in various wastewaters. AOPs are based on short-lived hydroxyl radicals ($\text{HO}\cdot$), which are powerful oxidizing agents that can decompose organic compounds in water [6]. AOPs, such as photocatalysis, ozonation, wet oxidation, and photo-Fenton have been documented to efficiently eliminate or mineralize organic pollutants in different types of wastewaters such as textile wastewater [7], pharmaceutical wastewater [8, 9], and pulp mill wastewater [10]. In this study, titanium dioxide (TiO_2) and zinc oxide (ZnO) photocatalysis, and photo-Fenton oxidation have been narrowed down as suitable methods to reduce the organic loads in POME. The photo-Fenton process uses Fenton reagents (H_2O_2 and Fe^{2+}) in the presence of light to produce $\text{HO}\cdot$ radicals. This process involves the oxidation of ferrous ions (Fe^{2+}) to ferric ions (Fe^{3+}) in an acidic aqueous solution to produce $\text{HO}\cdot$ radicals, which will trigger the oxidation of organic compounds. It is important to keep the solution at pH 3 because the Fenton reaction works best in an acidic condition [11].

In the case of TiO_2 and ZnO photocatalysis, photon illumination ($\lambda < 400 \text{ nm}$) onto the TiO_2 and ZnO surfaces, of greater than or equal to the band gap energy, will result in the formation of electron-hole pairs. These pairs will be involved in the oxidative and reductive reactions with molecules present at/or near the surface of the semiconductor [12]. Both TiO_2 and ZnO have similar band gap energy of 3.2 eV [13]. Nonetheless, TiO_2 is the most applied photocatalyst in various environmental applications, primarily due to its non-toxicity, low cost, and photostability. ZnO , despite being highly photosensitive and absorbs a larger portion of the solar spectrum compared to TiO_2 , suffers from photocorrosion in acidic aqueous solution and the formation of $\text{Zn}(\text{OH})_2$ on its surface due to dissolution [13]. Thus, photocatalysis is of special interest since sunlight can be used as a photon source. Solar ZnO and TiO_2 photocatalysis can be activated by sunlight of lower than 390 nm, while photo-Fenton oxidation requires sunlight of up to 500 nm. The degradation of organic pollutants using AOPs with sunlight as the energy source is advantageous in lowering costs. Previous studies on solar photo-Fenton [14], and solar ZnO and TiO_2 photocatalysis [15, 16] have recognized the roles of these mechanisms in reducing or degrading organic compounds in wastewater effluents. Nonetheless, there remains a paucity of knowledge regarding the feasibility of these treatments on POME. Aris et al. [17] reported that solar photo-Fenton has resulted in a better COD and colour from biologically treated POME compared to ambient photo-Fenton. On the other hand, TiO_2 photocatalysis was able to reduce the COD level of POME to 78% within 20 hours of irradiation [18].

This study aims to assess the efficiency of solar ZnO photocatalysis, solar TiO_2 photocatalysis, and solar photo-Fenton in reducing the COD level of POME. The dependence of solar photodegradation rate on intrinsic parameters, such as ZnO and TiO_2 concentrations, and the ratio of $\text{Fe}^{2+}:\text{H}_2\text{O}_2$ was investigated.

Materials and Methods

Sample collection

The POME samples used in this study were collected during the month of November 2014, from SALCRA Palm Oil Mill Plant located in Bau, Sarawak, Malaysia. The POME samples were collected from the cooling ponds of the palm oil mill, and stored in polyethylene bottles. The air-tight bottles were transported to the laboratory and kept refrigerated at 4 °C until further analysis. Characterization and solar photocatalytic oxidation treatments were performed using diluted POME.

Characterization of palm oil mill effluents

Diluted POME was used for water quality analysis. Distilled water was used to dilute 1 mL of POME effluent to 1000 mL. The levels of biological oxygen demand (BOD₅), chemical oxygen demand (COD), total suspended solids (TSS), and dissolve oxygen (DO) were determined. All analyses were performed according to the Standard Methods of Water and Wastewater Treatment [19].

Solar photodegradation of palm oil mill effluents

POME solar photodegradation study was conducted on the ground floor of the Faculty of Resource Science and Technology, Universiti Malaysia Sarawak, Malaysia. All solar treatments were performed for 2 – 5 hours on sunny days. Beakers filled with diluted POME and the required quantities of reagents were exposed to sunlight under continuous magnetic stirring. The beaker used for solar study was covered with a polyethylene wrap to avoid evaporation. A portable digital Lux meter (TEX 1335) was used to measure the intensity of sunlight. Temperature, pH, and solar intensity were periodically measured during periods of solar exposure. The recorded sunlight intensity and temperature during all solar experiments had ranged from 30 to 180 Klux and 30 to 38 °C, respectively. The temperatures recorded during the solar photo-Fenton and solar ZnO and TiO₂ photocatalysis had varied from 28 to 38 °C. In the case of indoor photo-Fenton experiments, the recorded temperatures had varied from 23 to 24.5 °C. All experiments were performed in duplicate. COD reduction was calculated using Equation 1:

$$\text{COD reduction (\%)} = \frac{\text{COD}_{\text{initial}} - \text{COD}_{\text{final}}}{\text{COD}_{\text{initial}}} \times 100 \quad (1)$$

Palm oil mill effluent concentration for treatment

As the concentration of POME in the photocatalytic treatments would affect the overall COD removal efficiency, three dilution ratios (POME: water) were considered: 1:10, 1:100 and 1:1000. Diluted POME sample of 1:10 ratio (Figure 1a) appeared to be more brownish, thicker, and murkier compared to the other diluted samples (Figure 1b and 1c). To ensure the maximum performance of the photocatalysts or reagents in these treatments, and to ensure adequate light penetration, the dilution ratio of 1:1000 was used. At high pollutant initial concentrations, all catalytic sites are occupied [20].

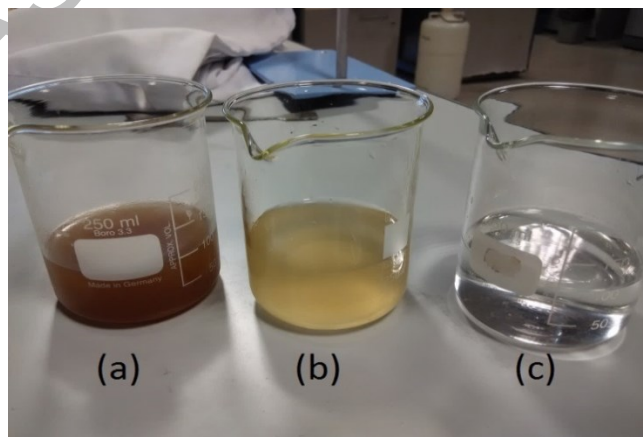


Figure 1. Photograph of diluted POME solutions (POME: water) (a) 1:10, (b) 1:100, and (c) 1:1000

Treatment of palm oil mill effluent using solar photocatalysis

Solar ZnO and TiO₂ photocatalysis were conducted by varying the concentrations of TiO₂ and ZnO from 0.02 to 0.2 g/L and 0.5 to 4 g/L, respectively in 150 mL of diluted POME (1:1000). The mixture of POME and photocatalyst (either ZnO or TiO₂) was magnetically stirred for 30 minutes in a dark environment (covered with polyethylene wrap) to establish equilibrium between the two substances. Then, the reaction slurry in the beaker was uncovered before being exposed to direct sunlight. The beaker was magnetically stirred during solar exposure and sample collection was done at pre-determined intervals using a syringe. A parallel control experiment (without photocatalyst) was also conducted. Supernatant obtained *via* filtration was used for COD analysis.

Treatment of palm oil mill effluent using solar photo-Fenton

Indoor photo-Fenton (laboratory) treatments with ambient light and solar photo-Fenton treatments were performed to compare their efficiency in reducing COD levels in POME. The concentration of Fe²⁺ was kept constant, while the concentration of H₂O₂ was varied in the photo-Fenton experiments. Similar ratios of Fenton reagents, Fe²⁺:H₂O₂, which include 1:10, 1:20, 1:30, and 1:40, were investigated for both treatments. The pH of the solution mixture was kept in the range of pH 2 to 3 by adding H₂SO₄ accordingly. Then, the mixture was continuously stirred in the laboratory to ensure homogeneity. It was subsequently exposed to sunlight. Samples were collected at fixed intervals. At the end of the solar exposure period, the pH of the sample was adjusted to range between pH 11 and 12 by adding NaOH pellets. The sample was stirred again for 15 minutes before being left overnight to ensure the decomposition of H₂O₂. After the sample has stood overnight, filtration was performed, which was followed by COD analysis. A blank sample (POME only) was also exposed to sunlight alongside the photo-Fenton treatment. Similar procedures were applied for photo-Fenton treatments in the laboratory. In this case, the source of photon was from the visible light in the laboratory. The beaker used for dark Fenton treatment was wrapped with aluminium foil to prevent any form of light from penetrating the reaction mixture.

Characterization

Analytical grade TiO₂ photocatalysts (99.8% trace metals basis, anatase) and ZnO were supplied by Sigma-Aldrich and Bendosen, respectively. The crystal phase of the TiO₂ and ZnO samples were determined using the X-ray Diffraction (XRD, PaNalytical X'pert Pro) method with Cu K α radiation ($\lambda = 0.154$ nm) in the scanning range of 2θ between 10° and 80°, at a rate of 0.04° per second. The accelerating voltage and applied current were 45 kV and 40 mA, respectively. Field emission scanning electron microscopy (FESEM) was performed using the Carl Zeiss, SUPRA 40VP Scanning Microscope to analyse the surface morphology of the samples. A low electron beam voltage of 5 kV was used. Therefore, no coating was required prior to imaging.

Results and Discussion

Characteristics of palm oil mill effluent

Table 1 lists the results of pH, BOD₅, COD, and TSS. POME is a thick brownish colloidal mixture of water, oil, and fine suspended solids. The pH of POME was found to be acidic, resulting from the organic acids produced during the fermentation process [5]. This indicates that raw POME is unsuitable to be directly discharged into water bodies as the permissible level set by DOE falls between 5 and 9 (Table 1). The BOD and COD values obtained in this study implied the elevated levels of organic matters in POME. A comparison between the obtained values and those outlined by the effluent discharge standard of the Environmental Quality Act 1974 [5], showed that these values had exceeded the permissible levels. The ratio of BOD₅/COD, which is normally used to express the biodegradability of wastewater, was calculated to determine whether POME could be biodegraded using biological treatments [21]. Clearly, the obtained BOD₅/COD ratio of 0.10 (Table 1) implied that POME was not suitable to be biodegraded using biological treatments. The characterization of POME suggested that the open ponding system, as practiced by palm oil mill operators, has failed to reduce these parameters to comply with the discharge limits set by the Environmental Quality Act 1974. Based on the obtained BOD₅/COD ratio, this method is inefficient to improve the water qualities of POME to an acceptable level. Therefore, other treatments must be sought.

Table 1. Characteristics of POME

Parameter	Mean $\pm$ SD	Effluent discharged standard for crude palm oil (Environmental Quality Act 1974)*
pH	4.85 $\pm$ 0.05	5-9
BOD ₃ , 30°C (mg/L)	7600 $\pm$ 15.2	100
COD (mg/L)	73 150 $\pm$ 124.5	1000
BOD ₅ /COD	0.10	NA
TSS (mg/L)	18 000 $\pm$ 5000	400

*Parameters Limit of Environmental Quality (Prescribed Premises) (Crude Palm Oil) (Amendment) Regulation 1997, and NA: Not applicable

Reducing chemical oxygen demand using solar heterogeneous photocatalysis

Two types of photocatalyst, TiO₂ and ZnO were applied during the solar heterogeneous photocatalytic treatment of POME. The effect of photolysis (without photocatalyst) was studied for two reasons: (i) to act as a control, and (ii) to confirm the contribution of solar heterogeneous photocatalysis towards reducing COD level. COD reduction, by means of photolysis, has ranged between 7 to 29%, thus indicating that the presence of TiO₂ or ZnO is crucial to reduce the COD level in POME.

An optimum concentration of photocatalyst under studied experimental conditions is an important factor towards achieving efficient removal of COD. Thus, the effect of TiO₂ (0.02 to 0.2 g/L) and ZnO (0.5 to 2.0 g/L) concentrations on the reduction of COD was studied. The same experimental conditions were applied for both photocatalytic treatments, where 150 mL of diluted 1:1000 POME was exposed under sunlight for 5 hours. The optimum concentration of photocatalyst was then fixed for other parameters studied. Figure 2 shows the percentage of COD reduction after 5 hours of solar TiO₂ photocatalysis. As expected, the percentage of COD reduction was increased from 54.3 to 88.5% concomitantly with the increased amount of TiO₂ from 0.02 to 0.1 g/L. However, further increase of TiO₂ to 0.2 g/L has yielded only 40% of COD reduction. Under the studied experimental conditions, 0.1 g/L of TiO₂ has led to a maximum reduction of 88.5% of COD (Figure 2). TiO₂ concentrations of higher than the optimum value (0.1 g/L) could increase the opacity of the suspension due to the excess amount of photocatalyst. Furthermore, penetration of solar radiation could have been impeded by the excess catalysts [22], contributing to the retardation of COD reduction. With 0.1 g/L of TiO₂, COD was reduced from 73,150 mg/L to 8,000 mg/L after 5 hours of solar exposure (Figure 3).

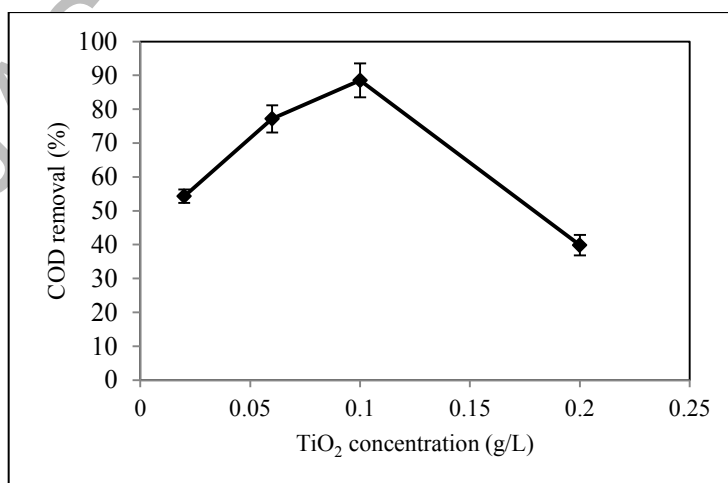


Figure 2. Effect of TiO₂ concentration on COD reduction efficiency using solar TiO₂ photocatalysis (error bars indicate standard deviation)

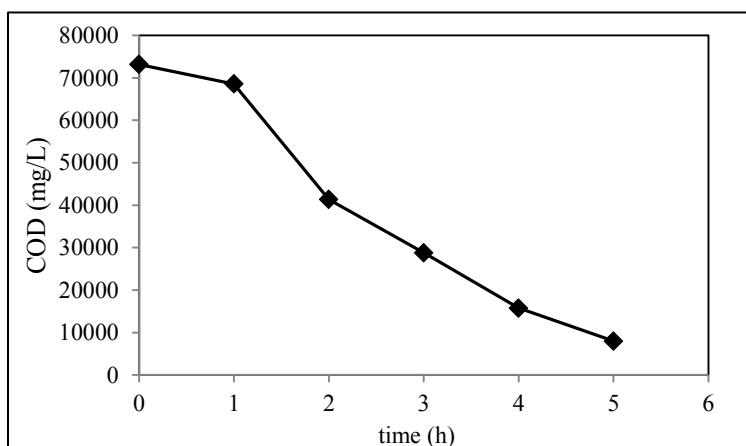


Figure 3. Reduction of COD with the optimum concentration of TiO_2 (0.1 g/L) using solar TiO_2 photocatalysis

In the case of ZnO , the percentage of COD reduction has increased from 26.3 to 60.5% with increased ZnO concentration, from 0.5 to 2 g/L (Figure 4). Further increase of ZnO concentration to 4 g/L has resulted in the declining percentage of COD reduction to 36.6%. The reasons elucidated for TiO_2 photocatalysis could also be applied to the phenomenon observed during the ZnO photocatalysis. When the performances of ZnO and TiO_2 photocatalysis were compared based on the effect of photocatalyst concentration on COD reduction, the latter has demonstrated better reduction efficiency under the investigated experimental conditions. The higher percentage of COD reduction of 88.5% was achieved with a low TiO_2 concentration (0.1 g/L). In contrast, 2 g/L of ZnO was required to obtain the maximum percentage of COD reduction of 60.5%. Several studies have reported that ZnO photocatalysis had performed better than TiO_2 photocatalysis for pollutant degradation [16, 23] because ZnO can absorb a larger fraction of the solar spectrum compared to TiO_2 . However, in this study, the performance of TiO_2 was superior to ZnO .

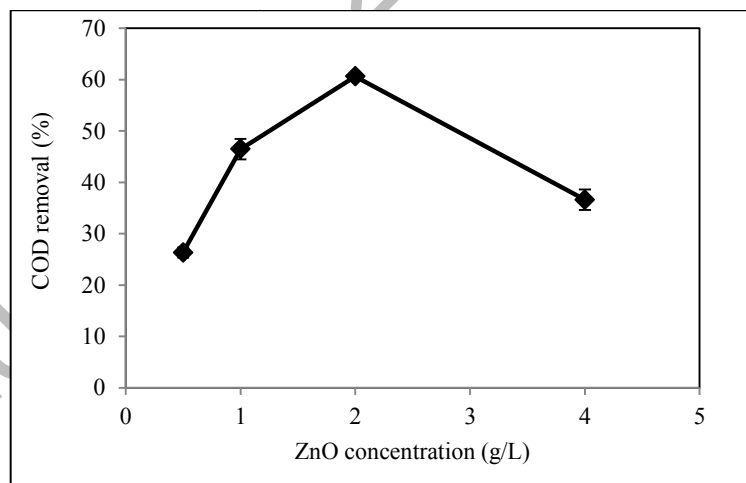


Figure 4. Effect of ZnO concentration on COD reduction efficiency using solar ZnO photocatalysis (error bars indicate standard deviation)

The performance of photocatalysts in degrading organic matter is known to be closely linked to their synthetic nature. The XRD patterns in Figure 5 clearly corresponded to the typical pure anatase TiO_2 . The existence of strong crystalline peaks at 2θ values of 25.3° , 37.6° , 48.0° , 53.9° , and 54.9° , which corresponded to the crystal planes of

(101), (004), (200), (105), and (211), respectively, were indicative of pure anatase TiO₂ (Figure 5a). As for ZnO, the crystalline peaks at 2 θ values of 31.8°, 34.4°, 36.2°, and 47.5° corresponded to the crystal planes of (100), (002), (101), and (102), respectively. These peaks were indicative of the predominant crystalline phase of ZnO (Figure 5b). FESEM images were also acquired for the commercial TiO₂ and ZnO, as shown in Figure 6. Figure 6a shows the surface of TiO₂ to be smooth and spherical in shape, with slightly varied particle sizes. Meanwhile, ZnO particles were predominantly rod-shaped (Figure 6b). Irregular sizes of ZnO particles in spherical and rectangular shapes can also be observed.

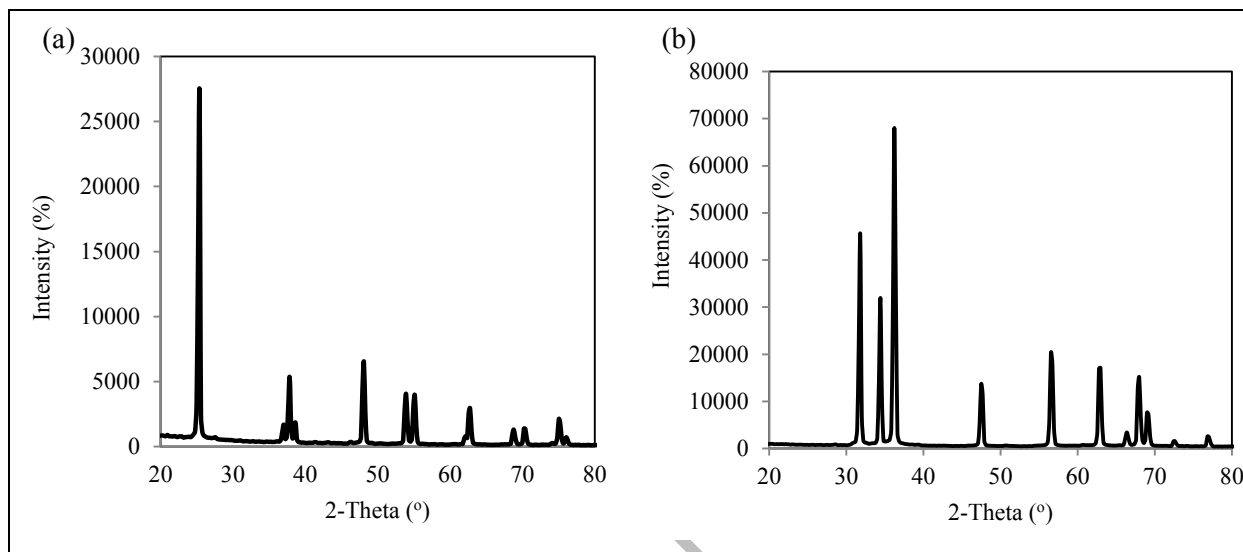


Figure 5. Comparison of XRD patterns between commercial: (a) TiO₂, and (b) ZnO

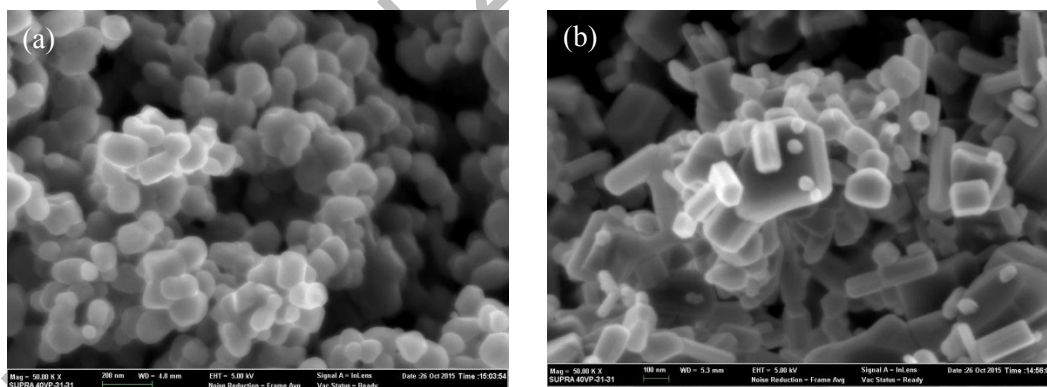


Figure 6. FESEM images of commercial: (a) TiO₂, and (b) ZnO

High crystallinity and non-porosity of both ZnO and TiO₂ could have contributed to the oxidation or mineralization of the organic matter in POME. The Brunauer-Emmett-Teller specific surface areas (S_{BET}) for TiO₂ and ZnO were 55 m²/g and 25 m²/g, respectively. The average particle size of TiO₂ could also be estimated from the S_{BET} by assuming that all particles have the same spherical size and shape. The particle size, D is given by Equation 2 [24]:

$$D = \frac{6000}{S_{BET} \times \rho} \quad (2)$$

where ρ is the true density for TiO_2 , which in this study, was 4.2 g/mL and the estimated particle size was 26 nm. However, estimating the particle size of ZnO from Equation 2 would be inaccurate as ZnO has distinct irregular shapes, ranging from rod to rectangular. By using the Image J software, the particle size of ZnO was estimated to be 300 – 400 nm, with some 46 – 50 nm spherical particles. In this study, COD removal efficiencies could largely be attributed to fine spherical nanoparticles that have higher surface areas compared to the larger particles. With its smaller particle size, TiO_2 could absorb more photon and subsequently, more active sites were created, where more hydroxyl radicals were induced to enhance COD removal. In contrast, ZnO has lower absorption potential due to its relatively larger particle size. A similar observation was also made by Rojviroon et al. [25] and Vineetha et al. [26].

The performance of solar photocatalytic reaction is greatly influenced by pH and the concentration of catalyst [27]. In this study, solar ZnO and TiO_2 photocatalysis were performed without any pH adjustment. POME is known to be very acidic in nature. In this study, pH adjustment was not considered as it will increase the cost of treatment. Nonetheless, pH readings were recorded during sample collection to investigate whether there was any direct relationship between the pH of the suspension and COD reduction. The pH values of the collected samples during this treatment ranged between 1.2 and 4.9. The recorded mean pH for the highest percentage of COD reduction, which was attained using 0.1 g/L TiO_2 , was at 3.68. Higher TiO_2 loading of 0.2 g/L had not only resulted in a very acidic pH (~1.2), it had also produced the lowest percentage of COD reduction. The aggregation of TiO_2 particles at 0.2 g/L could have reduced the ability of the suspension to absorb light [24], thus affecting the percentage reduction of COD. In addition, TiO_2 photocatalytic oxidation of POME may also produce other intermediary or degradation products, which could also have been acidic in nature, and directly affected COD reduction. Hence, this study has shown that COD reduction in POME is possible by means of solar TiO_2 photocatalysis without any prior pH adjustments and a small amount of TiO_2 .

Reduction of chemical oxygen demand by solar photo-Fenton

Solar photo-Fenton treatment was conducted by varying the $\text{Fe}^{2+}:\text{H}_2\text{O}_2$ ratio to investigate its effect on COD reduction efficiency. Experiments were conducted by varying the concentrations of H_2O_2 in difference ratios of $\text{Fe}^{2+}:\text{H}_2\text{O}_2$ (1:10, 1:20, 1:30, and 1:40). A predetermined volume of NaOH was added to the samples at the end of the photo-Fenton treatments. The NaOH will lead to the formation of iron precipitate, $(\text{Fe}(\text{OH})_3)$, which could accelerate the decomposition of H_2O_2 . The precipitation of $\text{Fe}(\text{OH})_3$ was observed to be slightly reduced in samples treated *via* photo-Fenton. To avoid the interference of excess H_2O_2 in the COD analysis, titration with potassium permanganate was conducted to ensure the complete decomposition of excess H_2O_2 . In the presence of H_2O_2 , manganese(VII) was reduced to colourless manganese(II), as shown in Equation 3.



Blank diluted POME (1:1000) was added with $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ and H_2O_2 separately (without both reacting simultaneously) to determine the effect of the addition of these reagents on COD reduction. POME samples that contained H_2O_2 had turned bright red immediately after the addition of Ferriin indicator, thus no titration was performed on these samples. POME samples with $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ had required similar amount of ferrous ammonium sulphate as the blank sample during titration. This observation implies that the addition of these reagents did not affect the experimental results of COD reduction in the photo-Fenton treatment.

The amount of $\text{Fe}(\text{II})$ catalyst was fixed, while the amount of H_2O_2 was varied to determine its optimum ratio. Ferrous ions react rapidly with H_2O_2 to produce a large amount of $\text{HO}\cdot$ radicals (Equation 4). Based on the properties of this catalyst, only a small amount was required to speed up the reaction. Higher loadings of ferrous ions may only contribute to the waste of chemical.



Figure 7 shows the reduction of COD using dark Fenton, indoor photo-Fenton and solar photo-Fenton after 3 h of reaction time. Dark Fenton oxidation has contributed to COD reduction percentage that ranged between 20 to 45%. The COD reduction efficiency was increased when the ratio of H_2O_2 concentration was increased from 1:10 to 1:30 for both indoor (50 – 78%) and solar photo-Fenton (62 – 89%) treatments. When the ratio was increased to 1:40,

COD reduction efficiency had declined to 40% for indoor photo-Fenton and 56% for solar photo-Fenton. The excessive H_2O_2 loading could have resulted in the production of $\text{HO}_2^\bullet$ radicals, which are less productive, compared to $\text{HO}^\bullet$ radicals (Equation 5). $\text{HO}_2^\bullet$ radicals could also have formed with the incorporation of $\text{HO}^\bullet$ radicals in H_2O_2 (Equation 6). In addition, the $\text{HO}^\bullet$ radicals could have recombined to form H_2O_2 , without favouring any reactions (Equation 7). Based on Figure 7, the reaction was concluded to be optimized when the $\text{Fe}^{2+}:\text{H}_2\text{O}_2$ ratio was 1:30. With this ratio, the maximum percentages of COD reduction of 78% (from 73,000 to 16,000 mg/L) and 89% (from 73,400 to 7,855 mg/L) were obtained with indoor photo-Fenton and solar photo-Fenton treatments, respectively after 3 h of reaction time (Figure 8). A comparison of COD reductions between dark Fenton oxidation, solar photo-Fenton, and indoor photo-Fenton has shown that photon is necessary to activate the Fenton reagents. COD reduction using dark Fenton oxidation was rather low, although the trend was similar to those observed in the photo-Fenton treatments (indoor and solar).

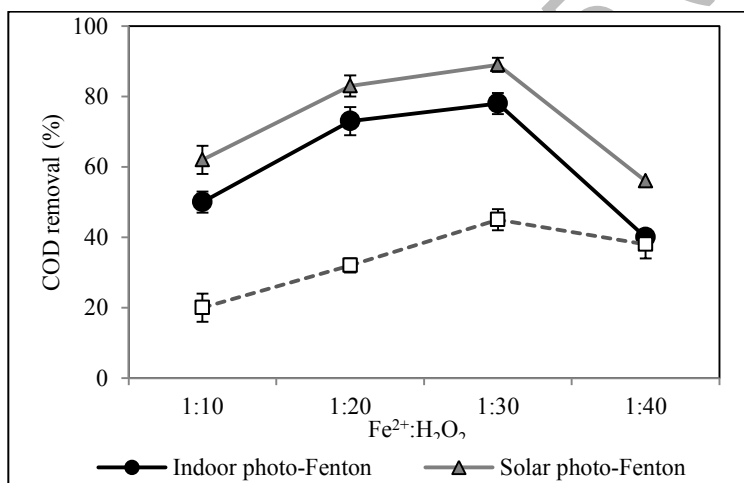


Figure 7. Reduction of COD by dark Fenton, solar photo-Fenton, and indoor photo-Fenton processes

In a Fenton treatment, pH plays an important role. The optimal pH condition in a Fenton reaction system is between 2.8 and 3.0 [28]. Therefore, the pH of POME solutions during each treatment was kept between 2.8 and 2.9. The purpose of the acidic pH is to avoid the precipitation of iron, which might interfere with the absorption of solar irradiation by the exposed medium. Prevention of iron precipitation will lead to the continuous photoreduction of ferric iron complexes at a longer wavelength.

In this study, solar photo-Fenton was found to be more efficient than indoor photo-Fenton. This result was in agreement with the findings reported by Aris et al. [17]. Several other studies have also concluded that solar photo-Fenton had performed better than other AOPs in various conditions, such as for textile effluent [29] and pharmaceutical wastewater [30]. The findings from the present study suggested that solar photo-Fenton should be considered as a valuable option for the reduction of COD in POME.

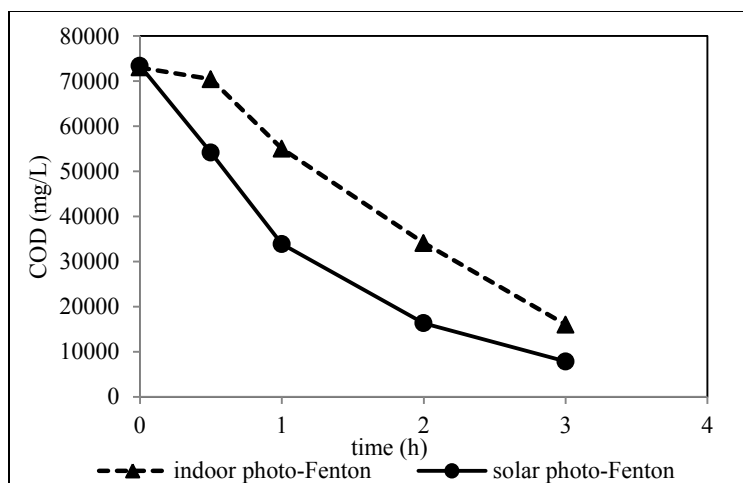


Figure 8. Reduction of COD by solar photo-Fenton and indoor photo-Fenton using the optimum $\text{Fe}^{2+}:\text{H}_2\text{O}_2$ ratio (1:30)

Conclusion

A higher percentage of COD reduction was accomplished with solar photo-Fenton compared to indoor photo-Fenton and dark Fenton under similar solar exposure time. The reduction of COD in POME was found to be governed by the concentration of the photocatalyst in solar ZnO and TiO_2 photocatalysis. On the other hand, H_2O_2 concentration was the determinant for better solar photo-Fenton and indoor photo-Fenton treatments. In addition, the pH value in photo-Fenton systems has also played an important role, where the optimum pH value was found to be between 2.8 and 2.9. Decreasing the pH to 1.2 or increasing the pH to an alkaline level may result in poor COD reduction. Both solar photo-Fenton and solar TiO_2 photocatalysis were able to produce up to 89% of COD reduction. Nevertheless, solar photo-Fenton took only 3h of exposure time to achieve the highest reduction compared to solar TiO_2 photocatalysis that took up to 5h to achieve a similar level. This study has shown that the reduction of COD in POME generated from palm oil milling activities can be performed using solar photo-Fenton and TiO_2 photocatalysis. Thus, solar photo-Fenton and TiO_2 photocatalysis can be considered as pre-treatment steps prior to the biological treatment.

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