

PHOTOCATALYTIC DEGRADATION OF PHENOL DERIVATIVES OVER SILVER SUPPORTED ON MESOPOROUS TITANIA NANOPARTICLES

(Degradasi Fotomangkin Daripada Terbitan Fenol Ke Atas Perak Disokong Pada Nanopartikel Titania Berliang Meso)

Nur Farhana Jaafar^{1*} and Aishah Abdul Jalil^{2,3}

¹School of Chemical Sciences,
Universiti Sains Malaysia, 11800 USM Penang, Malaysia

²Centre for Hydrogen Energy, Institute of Future Energy

³Department of Chemical Engineering, Faculty of Chemical and Energy Engineering
Universiti Teknologi Malaysia, 81310 UTM Johor Bahru, Johor, Malaysia

*Corresponding author: nurfarhana@usm.my

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Abstract

In this study, the potential of silver supported on mesoporous titania nanoparticles (Ag-MTN) as visible-light-driven photocatalyst was investigated for the degradation of phenol derivatives. Characterisation results illustrated that the presence of Ag reduced the band gap of MTN and increased the number of oxygen vacancies (OV) and Ti^{3+} site defect (TSD). The presence of both properties is among the important factors that need to be considered for catalyst properties in order to achieve effective photocatalytic degradation under visible light. The photoactivity of the catalyst was also significantly influenced by the presence of Ag, which acted as an electron trap that enhanced electron-hole separation. The ability of Ag-MTN to degrade phenol derivatives [phenol (Ph), 2-chlorophenol (2-CP), 2,4-dichlorophenol (2,4-diCP), and 4-acetamidophenol (4-AcePh)] was compared with commercial TiO_2 and MTN. The percentage degradation of phenol derivatives was in the following order: $\text{TiO}_2 < \text{MTN} < \text{Ag-MTN}$. This indicated that the presence of Ag in MTN effectively enhanced the photoactivity of MTN towards the degradation of phenol derivatives.

Keywords: Ag-MTN catalyst, electron trapper, visible light, phenol derivatives

Abstrak

Dalam kajian ini, potensi perak disokong pada nanopartikel titania berliang meso (Ag-MTN) sebagai fotomangkin-pemacu cahaya nampak telah disiasat bagi degradasi terbitan fenol. Keputusan pencirian menunjukkan kehadiran Ag mengurangkan ruang jalur MTN dan meningkatkan bilangan kekosongan oksigen (OV) dan permukaan Ti^{3+} cacat (TSD). Kehadiran kedua-dua sifat adalah antara faktor penting yang perlu dipertimbangkan bagi sifat mangkin bagi mencapai degradasi fotomangkin yang efektif di bawah cahaya nampak. Fotoaktiviti bagi mangkin juga sangat dipengaruhi dengan kehadiran Ag di mana bertindak sebagai perangkap elektron yang meningkatkan pemisahan elektron-lubang. Keupayaan Ag-MTN untuk degradasi terbitan fenol [fenol (Ph), 2-klorofenol (2-CP), 2,4-diklorofenol (2,4-diCP) dan 4-asetamidofenol (4-AcePh)] telah dibandingkan dengan TiO_2 komersial dan MTN. Peratusan degradasi bagi terbitan fenol telah mengikuti susunan: $\text{TiO}_2 < \text{MTN} < \text{Ag-MTN}$. Ini menunjukkan kehadiran Ag dalam MTN berkesan meningkatkan fotoaktiviti bagi MTN terhadap degradasi terbitan fenol.

Kata kunci: mangkin Ag-MTN, perangkap elektron, cahaya nampak, terbitan fenol

Introduction

Phenol derivatives are widely used as intermediates in pulp, paper, biocides, cleaning agents, dyes, preservatives, and pesticides industries [1]. However, the release of phenol derivatives into aqueous environmental systems has raised concerns due to their high toxicity, high stability, and potentially carcinogenic [2]. The carbon-halogen bond causes recalcitrance of phenol derivatives, which is cleaved with great difficulty and the stability of their aromatic structure, resulting in their growth in nature. As a result, the Department of Environment (DOE) Malaysia has gazetted the allowable limits as in Environmental Quality (Sewage and Industrial Effluent) Regulations, 1979, that this pollutant should be treated to be less than 1 mg L^{-1} before being discharged to water bodies.

There are ongoing studies on the appropriate methods for eliminating these toxic organic compounds from aqueous systems due to the adverse effect caused by phenol derivatives on the environment and humans. Several removal techniques including ion exchange, adsorption, and solvent extraction have been proposed for removing or treating phenol derivatives [3]. However, these techniques incur high cost, is time consuming, and can be responsible for generation of secondary pollutants [4]. Heterogeneous photocatalysts such as TiO_2 , FeOOH , ZnO , CuO , and ZrO_2 are commonly used in advanced oxidation process (AOP) for removal of this recalcitrant pollutant. Among them, TiO_2 shows superior performance due to its strong oxidising power and chemical inertness [5]. However, photocatalytic application of TiO_2 has been limited by its high photogenerated electron-hole recombination rate caused by the rapid transfer of electrons generated from the semiconductors to the metal particles and can only be excited by UV-light irradiation [6]. Hence, many approaches have been reported to overcome this problem such as varying the synthesis method in order to obtain a mesoporous structure for enhancing its efficiency and modification with other semiconductors or noble metals.

Nowadays, self-modified mesoporous titania nanoparticles (MTN) containing oxygen vacancies (OV) and Ti^{3+} site defects (TSD) are one of the approaches in light-absorption modification for TiO_2 to increase its photocatalytic performance [7]. Microwave (MW)-assisted method is one of the efficient methods that have been studied to synthesise MTN since it could provide a uniform and fast reaction environment to produce mesoporous materials [8]. Besides, Ag^0 is one of the potential photocatalysts to be used in the doping with TiO_2 due to its properties that delay the recombination of photogenerated electron-hole pairs by acting as an electron trap [9]. Various methods have been used to activate Ag^+ to Ag^0 for synthesising Ag-MTN including electrochemical method, which is a simple activation of Ag^0 onto MTN without the need of high temperature or longer time. Therefore, we reported the synergistic effect between Ag^0 and MTN towards the degradation of phenol derivatives under visible light.

Materials and Methods

Reagents, material and apparatus

TiO_2 powder catalyst, JRC- TiO_2 -2, was supplied by the Catalysis Society of Japan and titanium (IV) isopropoxide (TTIP) was bought from Sigma Aldrich. Ag and Pt plates of greater than 99% purity were used as electrodes and were obtained from Nilaco, Japan. *N,N*-dimethylformamide (DMF), cetyltrimethylammonium bromide (CTAB), propanol, phenol (Ph), 2-chlorophenol (2-CP), 2,4-dichlorophenol (2,4-diCP), 4-acetamidophenol (4-AcePh), and perchloric acid were purchased from MERCK, Malaysia. Naphthalene and tetraethylammonium bromide solution were obtained from Fluka Chemical and acetone was purchased from HmbG Chemical. Sodium hydroxide (NaOH), ammonium hydroxide (NH_4OH), and hydrochloric acid (HCl) were purchased from QRECTM and methanol was purchased from RPE Reagent Pure Erba. Tetraethylammonium perchlorate (TEAP), which was used as a supporting electrolyte in the electrolysis, was prepared in accordance with the procedure reported in the literature [10].

Preparation of catalyst: Synthesis of mesoporous titania nanoparticles

Mesoporous titania nanoparticles (MTN) was synthesised by dissolving 4.68 g of CTAB in 720 ml of distilled water, 120 ml of propanol, and 29 ml of 28% ammonia solution. The mixture was stirred continuously for 30 minutes at 323 K in water bath. Then, the temperature of water bath was increased to 353 K, followed by the addition of 5.7 mL TTIP, and the mixture was stirred for another 2 hours. The white solution was transferred into a beaker after 2 hours of stirring and placed in a microwave. The heating continued under 0.56 Wg^{-1} as the power density of microwave until a sol-gel of TiO_2 was formed. The obtained product was collected and dried overnight in an oven before calcination at 873 K for 3 hours.

Synthesis of silver supported on mesoporous titania nanoparticles

The synthesis of Ag-MTN was carried out using a simple electrochemical method in accordance with procedure in the literature [11]. A closed-system electrolysis cell was fitted with a magnetic stirrer and a platinum plate cathode facing a silver plate anode. Then, MTN and TEAP were added to DMF. The electrolysis was conducted at a constant current of 240 mA cm^{-2} and 273 K under air atmosphere to load 5 wt.% Ag on MTN. The sample was evaporated and dried at 383 K for 12 hours before calcination at 873 K for 3 hours to yield a grey powder catalyst donated Ag-MTN that is ready for characterisation and photocatalytic testing.

Characterisation

The crystalline structures of the catalysts were examined using a Bruker Advance D8 X-ray powder diffractometer (XRD) with Cu K α radiation of $\lambda=1.5418 \text{ \AA}$ at 2θ angle ranging from 22° to 66° . The phases were identified with the aid of the Joint Committee on Powder Diffraction Standards (JCPDS) files. The chemical functional groups present in the catalysts were identified by Fourier transform infrared (FTIR) spectroscopy (Perkin Elmer Spectrum GX FTIR spectrometer). IR absorbance data were obtained over a range of wavenumbers from 390 to 3700 cm^{-1} . The morphological properties of the catalysts were examined by transmission electron microscopy (TEM, JEOL JEM-2100F). Nitrogen adsorption-desorption isotherms were used to determine the textural properties at liquid nitrogen temperatures using a SA 3100 Surface Analyzer (Beckman Coulter). The Brunauer-Emmett-Teller (BET) and non-local density functional theory (NLDFT) methods were used to calculate surface area and pore distribution, respectively. Prior to measurement, all samples were degassed at 300°C and 0.1 Pa . The X-ray photoelectron spectroscopy (XPS) was conducted in the range $0\text{--}800 \text{ eV}$ using a Kratos Ultra spectrometer equipped with Mg K α radiation source (10 mA , 15 kV). A small Inox cylinder was used to press the powdered sample prior to analysis inside an analysis chamber pressure at about $1 \times 10^{-10} \text{ Pa}$ to obtain the data. The binding energy of the C (1s) at peak $284.5 \pm 0.1 \text{ eV}$ was taken as the internal standard or reference as the solution to correct the energy shift. A JEOL JES-FA100 ESR spectrometer was used for the determination of surface defect Ti^{3+} and oxygen vacancy, where the sample was placed at about 2 cm height inside the glass vessel followed by the electron spin resonance (ESR) measurement to identify the g-value at room temperature. UV-Vis diffuse reflectance spectroscopy (UV-Vis DRS) was recorded in air at room temperature using a Perkin Elmer Lambda 900 UV/Vis/NIR spectrometer with an integrating sphere.

Photodegradation of phenol derivatives

The photoactivity of the catalysts was tested for the degradation of phenol derivatives. Four types of phenol derivatives were used: Ph, 2-CP, 2,4-diCP, and 4-AcePh. The stock solutions of the phenol derivatives were prepared with the initial concentration of each derivative of 50 mgL^{-1} . The stock solutions of the phenol derivatives were mixed in the ratio of 1:1:1:1. The photocatalytic experiments were performed in a batch reactor fixed with a cooling system. A 39 W metal halide lamp (400 nm) was used as a visible light source.

For photoactivity evaluation, 0.375 gL^{-1} of MTN was added to the phenol derivatives solution (200 ml) and stirred for 1 hour in the dark to achieve adsorption-desorption equilibrium. The initial pH of the solution was pH 5 and the reaction was carried out at 30°C . Then, the reaction was carried out for another 6 hours under light irradiation under continuous stirring. The concentration of the sample was measured using UV-Vis spectrometer at the absorption bands of 270, 274, 285, and 248 nm for Ph, 2-CP, 2,4-diCP, and 4-AcePh, respectively. Similar procedure was used for TiO_2 and Ag-MTN.

Results and Discussion

Figure 1 shows the XRD patterns of TiO_2 , MTN, and Ag-MTN in the range of $2\theta = 22\text{--}66^\circ$. A series of characteristic peaks was observed for TiO_2 (Fig. 1a) with peaks at 25° (101), 36.8° (103), 37.5° (004), 39° (112), 47.5° (200), 54° (105), 56° (211), and 63° (204) that was consistent with the TiO_2 anatase phase (JCPDS file no. 00-004-0477). The XRD pattern of MTN was similar to TiO_2 . However, the peak intensity of MTN (Fig. 1b) seemed to be more intense compared with TiO_2 , which indicated that the crystallinity of MTN is better than TiO_2 . The XRD pattern of Ag-MTN catalyst (Fig. 1c) was similar to MTN (Fig.1b), which illustrated that the loading of Ag did not intrude the structure of MTN [12]. However, the peak intensity of MTN seemed to decrease with the addition of Ag, which indicated the degree of orderliness in MTN and mesoporous uniformity decreased but did not remarkably

intrude the structure of catalyst [13]. Meanwhile, new diffraction peaks were observed at 32.35° (200), 33.15° ($\bar{1}11$), and 57.2° ($\bar{1}13$) after the Ag loading due to the plane of Ag for AgO moieties in Ag-MTN [14]. In addition, another two diffraction peaks at 27.5° and 46.3° showed the presence of traces of rutile and brookite. The crystallite size of the catalysts was calculated by the Debye-Scherrer equation based on the major peak $2\theta = 25^\circ$ as follows:

$$\tau = \frac{k\lambda}{\beta \cos \theta} \quad (1)$$

where τ is the particle size, λ is the wavelength of X-ray radiation ($\text{Cu K}\alpha = 0.1542 \text{ nm}$), k is the shape factor ($k = 0.94$), β is the line width at half-maximum height, and θ is the angular position of the peak maximum. It was found that the size of TiO_2 and MTN were 30.0 and 11.3 nm, respectively. However, the addition of Ag onto MTN demonstrated the decrease in the crystallite size from 11.3 to 7.21 nm, possibly due to the interaction of Ag and MTN. A similar observation was reported by Das et al., which revealed that the interaction of titania species with ZrP and TiP was the reason for the decrease in crystallite size of catalysts [15].

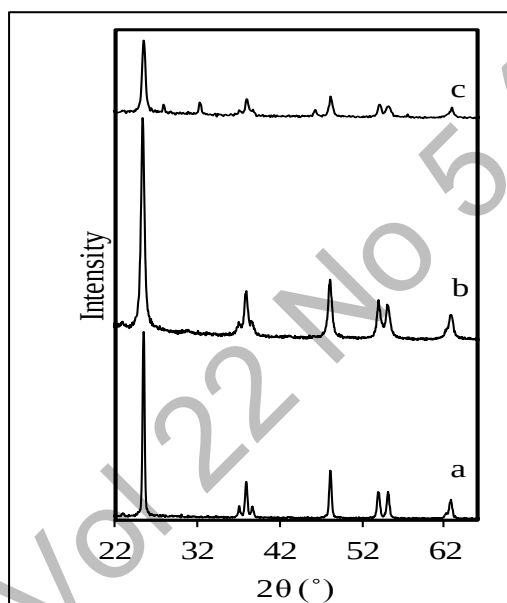


Figure 1. XRD diffractograms of (a) TiO_2 (b) MTN and (c) Ag-MTN

The textural properties for all catalysts are summarised in Table 1. The results showed that MTN has higher surface area and pore volume compared to TiO_2 . This phenomenon indicates that the balance of heat distribution by microwave during the synthesis for hydrolysis and condensation may contribute to the framework cross-linking for better growth of smaller pores [16]. In contrast, the introduction of Ag onto MTN shows a decrease in surface area and pore volume probably due to the aggregation of Ag nanoparticles that blocks the pore surface of MTN. In addition, the introduction of Ag onto MTN also decreased the band gap from 3.11 to 3.05 eV. This result demonstrates that the anchoring of Ag to MTN surface helps to reduce the band gap of Ag-MTN catalyst. A similar trend was also observed by Khan et al. for the synthesis of Ag@TiO_2 composite, in which the band gap of TiO_2 decreased after the introduction of Ag from 3.33 to 2.86 eV [17].

The morphology of the catalysts was then examined by TEM analyses and the images are shown in Figure 2. The TEM images of TiO_2 (Fig. 2a) and MTN (Fig. 2b) show translucent property with irregular shaped particles. Then, the TEM image of MTN (Fig. 2b) was compared with Ag-MTN (Fig. 2c) to study their morphology properties. The original morphology of MTN was interrupted by the introduction of Ag but by comparing with the translucent image of MTN, the distribution of Ag was clearly observed in Ag-MTN. This result confirmed that Ag nanoparticles dispersed well on the surface of MTN.

Table 1. Textural properties of catalysts

Catalyst	Surface Area (m^2g^{-1})	Pore Volume (cm^3g^{-1})	^a Band Gap (eV)
TiO ₂	12.5	0.22	3.31
MTN	181	0.35	3.11
Ag-MTN	100	0.31	3.05

^a Band gap calculated using $E_g = 1240/\lambda$

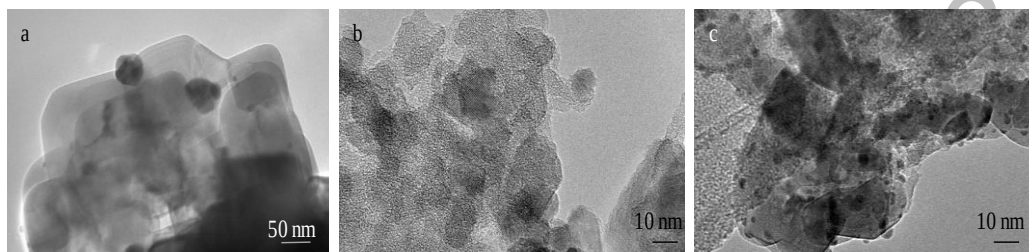


Figure 2. TEM images of (a) TiO₂ (b) MTN and (c) Ag-MTN

Figure 3 shows the FTIR spectroscopy in the range of 2000–390 cm^{-1} for all the catalysts. All catalysts showed a band at 1625 cm^{-1} that corresponded to OH vibration of the surface-adsorbed water [18]. The bands observed at 1100 and 455 cm^{-1} for MTN (Fig. 3b) and Ag-MTN (Fig. 3c) were assigned to Ti-O-Ti asymmetric stretching and bending vibration modes, respectively [19]. However, TiO₂ (Fig. 3a) illustrated that both of the bands were absent but a sharp band could be detected at 530 cm^{-1} , which was attributed to the Ti-O-Ti bond of TiO₂ anatase phase [20]. Besides, another new peak was also observed for Ag-MTN at 650 cm^{-1} attributed to the Ag-O bond, which illustrated that Ag nanoparticles were successfully introduced on MTN.

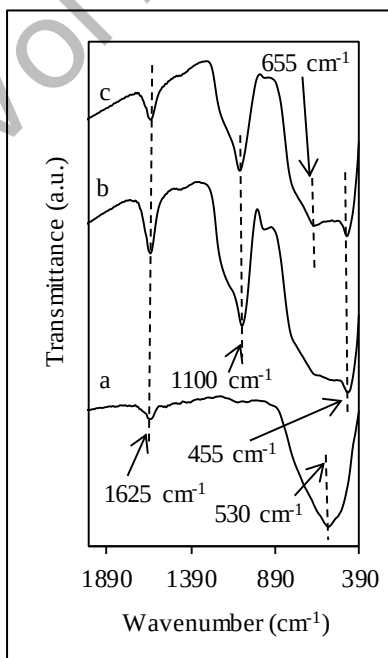


Figure 3. FTIR spectra in region 2000–390 cm^{-1} of (a) TiO₂ (b) MTN and (c) Ag-MTN

Chemical states of MTN and Ag-MTN were determined using XPS analysis as shown in Figure 4. Fig. 4A shows the wide scan for MTN and Ag-MTN. Titanium (Ti) and oxygen (O) were detected for both catalysts, while Ag could only be detected in Ag-MTN, as indexed in both catalysts. Fig. 4B(a) displays the region of Ti 2p_{3/2} with Ti³⁺ were observed at 457.7 and 458.9 eV while the binding energies at 458.3 and 459.3 eV were assigned to Ti⁴⁺ for MTN [21]. The region O 1s of MTN (Fig. 4C(a)) illustrated the existence of Ti³⁺-O at 530.8 and 532.5 eV, whereas the binding energies at 529.5 and 534.5 eV were attributed to Ti⁴⁺-O and OH or OH⁻, respectively [22]. These results confirmed the presence of TSD in MTN. The XPS analysis of Ag-MTN showed that new regions were observed other than Ti 2p_{3/2} (Fig. 4B(b)) and O 1s (Fig. 4C(b)), which were Ti 2p_{1/2} (Fig. 4B(b)) and Ag 3d (Fig. 4D) prior to the introduction of Ag onto MTN. Fig. 4B(b) illustrated four binding energies in the region of Ti 2p_{3/2} at 458 (Ti³⁺), 459 (Ti³⁺), 458.4 (Ti⁴⁺), and 459.5 eV (Ti⁴⁺), with two binding energies at 462.8 (Ti⁴⁺) and 464.7 eV (Ti⁴⁺) for the region of Ti 2p_{1/2} [17]. Meanwhile, O 1s (Fig. 4C(b)) region showed the binding energies at 530.4 and 532.7 eV for peak Ti³⁺-O, whereas the peaks at Ti⁴⁺-O and OH or OH⁻ were observed at 529.5 and 534.5 eV, respectively. The presence of Ag nanoparticles was confirmed in Fig. 4D by the presence of binding energies at 368 and 374 eV, which were attributed to Ag 3d_{5/2} and Ag 3d_{3/2}, respectively [23]. This result agrees with XRD (Fig. 1c) and FTIR (Fig. 3) analysis where Ag and AgO were detected.

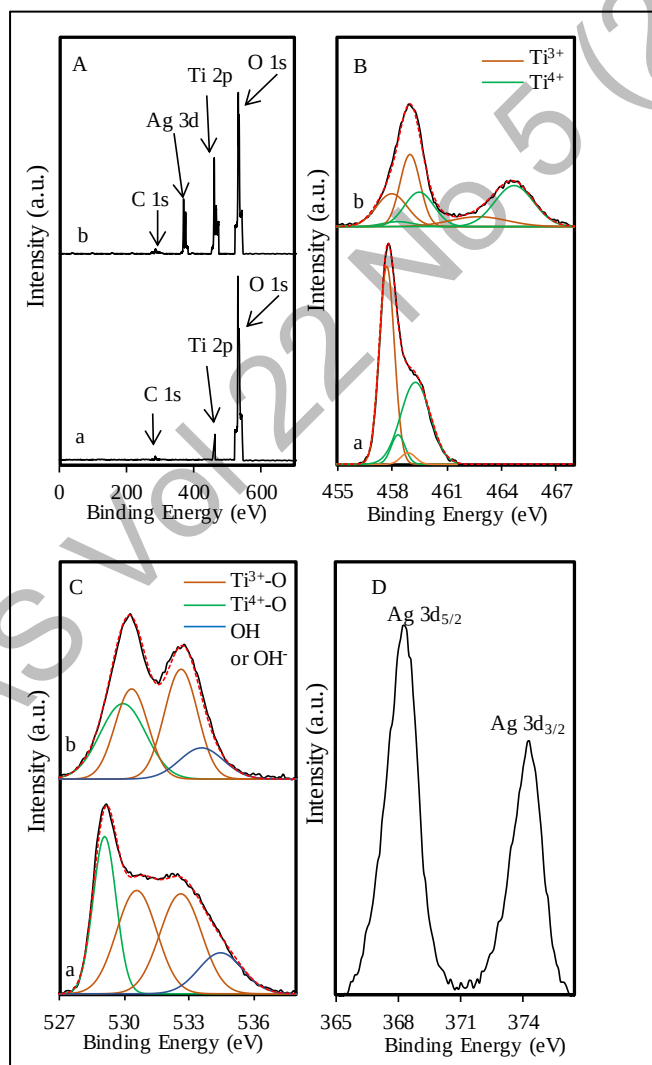


Figure 4. XPS spectra of (A) wide scan; (B) Ti 2p; (C) O 1s of (a) MTN and (b) Ag-MTN; and (D) Ag 3d of Ag-MTN

In addition, ESR also further confirmed the presence of TSD and oxygen vacancies (OV) for the catalysts and the results are shown in Figure 5. Two signals corresponded to OV and TSD sites were observed at $g=1.99$ and $g=1.96$, respectively. The signal intensity of Ag-MTN (Fig. 5c) was the highest for both OV and TSD prior to the introduction of Ag onto MTN compared to MTN. However, the signal intensities of MTN (Fig. 5b) were higher compared to TiO_2 (Fig. 5a). These results signify the presence of OV and TSD in all catalysts where these species are important elements in the photocatalytic process since this process mainly depends on OV and TSD [24].

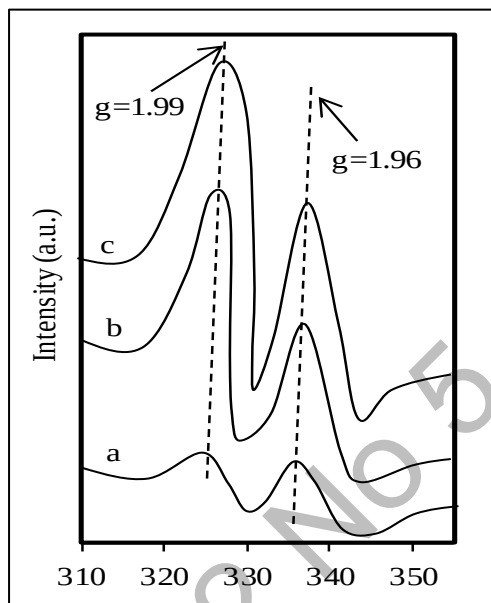


Figure 5. ESR of (a) TiO_2 (b) MTN and (c) Ag-MTN

Figure 6 demonstrates the photocatalytic activities of the catalysts on the degradation of phenol derivatives. It could be observed that MTN showed higher performance on the degradation of phenol derivatives compared to TiO_2 . Photocatalytic degradation of phenol derivatives using MTN successfully degraded 80, 92, 70, and 75% of Ph, 2-CP, 2,4-diCP, and 4-AcePh, respectively. In contrast, TiO_2 only managed to degrade 15, 20, 25, and 13% of Ph, 2-CP, 2,4-diCP, and 4-AcePh, respectively. This phenomenon occurs may be due to numbers of OV and TSD for TiO_2 that are lower than MTN, where the larger values of both species, the higher the photoactivity. These two species could act as the electron acceptor in order to decrease the electron-hole recombination, as well as the promoter in the oxidation of phenol derivatives since it could also adsorb O_2 [25].

Meanwhile, Ag-MTN showed the highest degradation of phenol derivatives compared to TiO_2 and MTN, in which Ag-MTN successfully degraded 85, 95, 80, and 79% of Ph, 2-CP, 2,4-diCP, and 4-AcePh, respectively. This result is probably because Ag-MTN has the highest number of OV and TSD according to the ESR result (Fig. 5), which benefits the photocatalytic process by reducing the band gap energy of the catalyst. The decrease in the band gap of Ag-MTN could also improve photoactivity because it is expected that Ag nanoparticles will act as the electron acceptor after the visible light irradiation on MTN that leads to electron excitation from the valence band (V_B) to the existing OV and TSD before reaching the conduction band (C_B). Consequently, this increases the concentration of photogenerated holes that play an important role at the catalyst surface. Besides, this result also illustrated the role of Ag as the electron trapper, which is attributed to the Schottky energy barrier formed between the interaction of Ag and MTN and plasmonic sensitizer to suppress the electron-hole recombination, where the synergistic effect between Ag and MTN plays an important role in the system.

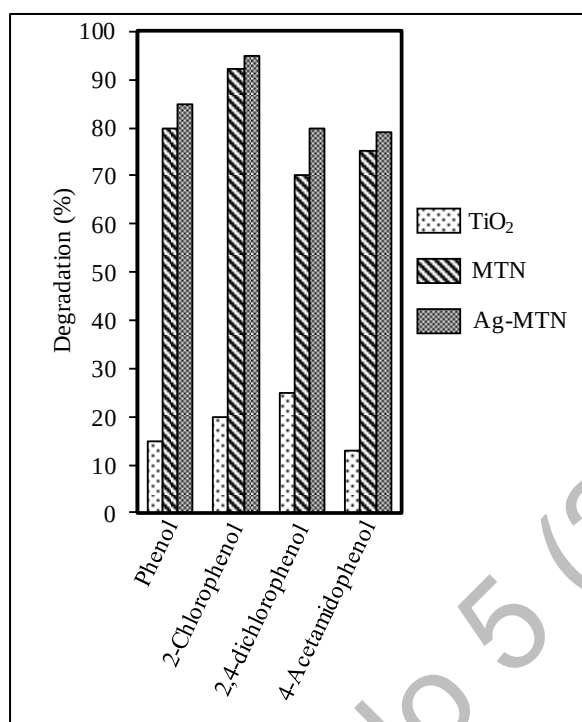


Figure 6. Photodegradation activity of TiO₂, MTN and Ag-MTN for degradation of simulated phenol derivatives [C_{Phenol derivatives}=50 mg L⁻¹, pH=5, W= 0.375 g L⁻¹, t=6 hours, 303 K]

Conclusion

The introduction of Ag onto MTN demonstrated that it can be used as an effective photocatalyst for the degradation of phenol derivatives under visible light. The physicochemical properties of the catalysts were studied using XRD, TEM, FTIR, ESR, N₂ adsorption desorption, UV-Vis DRS, and XPS. The analyses of UV-Vis DRS, ESR, and XPS verified that the introduction of Ag reduced the band gap of MTN and increased the number of OV and TSD, which seemed to play important roles for application under visible light. Ag-MTN gave the highest percentage degradation of phenol derivatives followed by MTN, while TiO₂ showed the lowest degradation. The presence of Ag not only improved the properties of MTN to be applied under visible light but also acted as electron trapper and plasmonic sensitizer to enhance the photoactivity of catalyst. Thus, this modified catalyst has a great potential to be extended for degradation of various pollutants such as heavy metal, organic pollutants, and pharmaceutical wastes. Besides, by using the reported techniques (microwave-assisted method and electrochemical method), it is believed that further modification of other semiconductors can be also carried out to create various properties for various applications.

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