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Microwave- and Formaldehyde-Assisted Synthesis of Ag–Ag₃PO₄ with Enhanced Photocatalytic Activity for the Degradation of Rhodamine B Dye and Crude Oil Fractions

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1. INTRODUCTION

Environmental pollution and its effects on human civilization are a major global problem. For example, oil spills and the discharge of dyes into water bodies used in painting, printing, plastic, textile, pharmaceutical, food processing, and so forth have a devastating effect on the environment and affect the survival of humans and aquatic species. For oil spills, remediation technologies such as in situ burning, the application of chemical dispersants, and mechanical containment and collection are employed to reduce their environmental impact. Although chemical dispersants are widely accepted as a response strategy for large-scale oil spills, they are reportedly ineffective in remediating the water-soluble components of crude oil. Therefore, there is a strong need for a remediation strategy that is effective at treating the water-soluble components of crude oil (aromatics). Additionally, environmental pollution by water-soluble dyes can be tackled through promoting adsorption, followed by photocatalytic treatments.

Interest in photocatalytic processes presents an attractive solution to numerous wastewater and oil spill remediation challenges. Photocatalysis has been reported to be an effective strategy using materials that are stable and environmentally friendly for environmental pollution control. Photocatalysis involves the following steps: (a) a photocatalyst is irradiated using photons with energy greater than or equal to the band gap energy of the photocatalyst, (b) electron–hole pair is generated as a result of the photoexcitation of electrons from the valence band (VB) to the conduction band (CB), (c) the generated holes and electrons are transported to the surface of the photocatalyst where they are utilized in the creation of reactive oxygen species. The high oxygen potential observed in photocatalytic processes allows for complete breakdown of organic contaminants. In addition, the use of sunlight makes it an attractive solution environmentally. The most widely used semiconductor photocatalyst material is based on TiO₂. TiO₂ is relatively cheap, very stable, and readily available and has high activation. The band gap of TiO₂ is ca. 3.2 eV. This large value has restricted its application to the UV region of the electromagnetic spectrum. Because ultraviolet light represents approximately 4–5% of solar radiation that arrives on the earth’s surface, many researchers have focused on developing

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visible light active photocatalysts. However, TiO₂ and many photocatalysts have limitations because of wide band gap, poor quantum efficiency under visible light, and faster decay of photoexcited electron–hole pairs. These challenges can be addressed by developing a new visible light active photocatalyst or by coupling different photocatalysts to create heterostructures. Ag₃PO₄ which is a visible light active semiconductor in the environmental remediation of water-soluble dyes has been investigated, and direct and indirect band gaps of Ag₃PO₄ are reported to be 2.34 and 2.36 eV, respectively. Ag₃PO₄ has a CB and a deep VB located at +0.45 V versus NHE (normal hydrogen electrode) and +2.9 V versus NHE, respectively, implying that reduction of Ag₃PO₄ to Ag⁰ is thermodynamically more favorable than the reduction of H⁺ to H₂ under light irradiation in water. Hence, the stability of Ag₃PO₄ is an issue. The limited photostability of Ag₃PO₄ has negatively affected its practical application in photocatalysis.

Hence, many researchers are studying Ag₃PO₄ with the main objective of reducing photocorrosion and enhancing the photocatalytic efficiency for practical applications. In order to enhance the photostability and efficiency of Ag₃PO₄, coupling of Ag₃PO₄ with either a metal or a semiconductor is proposed. Ag₃PO₄ modified through the introduction of a heterojunction can significantly outperform Ag₃PO₄ single phases. Electron–hole recombination is reduced when the electron and hole migrate from a semiconductor to the conductor. The reduction of the electron–hole recombination will increase the life span of the photogenerated electrons and holes. In addition, the localized surface plasmon resonance of the silver nanoparticles on Ag₃PO₄ can enhance the photocatalytic efficiency and stability. Coupling Ag₃PO₄ with AgS resulted in an enhanced photocatalytic degradation of methyl orange when compared to pure Ag₃PO₄. Liu et al. developed a one-pot pyridine-assisted hydrothermal synthesis method for Ag/Ag₃PO₄. Another synthesis route for Ag/Ag₃PO₄ involving exchange of ions between Na₂HPO₄ and AgNO₃ followed by a photoreduction route has been reported. Ag/Ag₃PO₄/rGO microspheres with enhanced removal efficiency of Cr(IV) have also been recently reported. In all these, the photocatalytic efficiency and photostability of Ag₃PO₄ were improved by combining it with Ag.

In this work, we show for the first time that a new silver phosphate nanocomposite is synthesized using formaldehyde, and the nanocomposite efficiently degrades crude oil fractions. We have synthesized Ag–Ag₃PO₄ with enhanced photocatalytic efficiency and photostability using formaldehyde and microwave (MW) radiation. Rhodamine B dye and crude oil fractions (benzene, n-hexane, and 1:1 v/v methanol/benzene-soluble fractions) were chosen as the model pollutants to examine the photocatalytic efficiency of Ag–Ag₃PO₄ in aqueous media under visible light irradiation. The results from this study showed that the Ag–Ag₃PO₄ photocatalyst degrades rhodamine B dye and crude oil fractions with high photocatalytic efficiency. It is worth noting that the model pollutants themselves are encountered as real pollutants because of human activities, and thus, the data obtained are significant for direct applications.

2. RESULTS AND DISCUSSION

An ion-exchange method was used to synthesize Ag₃PO₄ and decorated with Ag by chemical reduction of the excess AgNO₃ used in the synthesis of Ag₃PO₄ with an organic compound (formaldehyde). Ag₃PO₄ suspended in the excess AgNO₃ solution with formaldehyde was then microwaved. The solution was then centrifuged, and the precipitate was collected and dried at 70 °C. Ag₃PO₄ turned dark yellow/brownish after microwave (Figure 1). This change in color with respect to pure Ag₃PO₄ suggests that the synthesized Ag₃PO₄ is decorated with Ag, and the resulting composite materials have acquired the ability to absorb light. This process is schematically represented in Figure 1.

The diffuse reflectance spectroscopy (DRS) analysis of the synthesized Ag₃PO₄ decorated with Ag using different volumetric amounts of formaldehyde and MW time of 3 min is presented in Figure 2. The DRS of the Ag₃PO₄ composite material revealed an enhanced absorption from 20 to 63% in the visible light region. The absorption increases with the increasing volume of formaldehyde used in the synthesis but leveled off after the addition of 8 mL of formaldehyde. The spectra showed that the addition of formaldehyde did not have...
a dramatic effect on the optical band gap of Ag₃PO₄. The optical band gap of Ag₃PO₄ and Ag₃PO₄ modified using 4 mL of formaldehyde [Ag−Ag₃PO₄(F,MW)] was estimated with the Kubelka–Munk method, and the plot is represented in Figure 3. In [Ag−Ag₃PO₄(F,MW)], F and MW represent formaldehyde and MW, respectively. From Figure 3, 2.47 and 2.41 eV were the estimated optical band gaps for Ag₃PO₄ and Ag−Ag₃PO₄, respectively.

Dye degradation experiments were conducted using rhodamine B dye. A 300 W tungsten halogen lamp was used to illuminate the catalyst. The degradation efficiency as well as the pseudo-first-order rate constants is presented in Figure 4. The rate constants were estimated from eq 1 according to the Langmuir–Hinshelwood kinetics

$$\ln \frac{[C]}{[C_0]} = -kt$$

where C and C₀ are the concentrations of the dye at time t min and 0 min, respectively, and k (min⁻¹) is the pseudo-first-order rate constant. The percent degradation as well as the rate constant increased significantly upon modification with Ag (Figure 4).

The first two bars show the rate constant and the degradation efficiency of bare Ag₃PO₄ giving a % degradation and rate constant of 79.9% and 0.1696 min⁻¹, respectively, in 10 min. The results show a dramatic enhancement in the photocatalytic activity upon the addition of formaldehyde, followed by microwaving for 3 min. Upon the addition of 1 mL of formaldehyde, the % degradation and rate constant increased to 92.3% and 0.5143 min⁻¹, respectively. Further increase in the volumetric amount of formaldehyde to 10 mL resulted in a % degradation and rate constant of 98.8% and 0.854 min⁻¹, respectively. There was no significant difference in the photocatalytic activity of the composite material synthesized with 8 and 10 mL of formaldehyde when the values reached a plateau. The photocatalytic activity (rate constant) of Ag₃PO₄ therefore increased over 400% upon modification with surface Ag.

The above experiments were conducted using 3 min of microwaving time. The effect of microwaving time on optical properties and photocatalytic activity was examined. By increasing the microwaving time from 3 to 5 min, the absorption of light increased slightly in the visible region (Figure 5). The nature of the DRS shows that the band gap remained unchanged upon increasing the microwaving time. A scanning electron microscopy (SEM image) (Figure 8—presented later in this article) revealed an extreme etching of Ag−Ag₃PO₄. Increasing the microwaving time therefore affected the surface morphology.

Figure 2. DRS of Ag₃PO₄ and Ag₃PO₄ modified with Ag at different volumes of formaldehyde used for the modification.

Figure 3. Generated plots from the Kubelka–Munk theory for (A) Ag₃PO₄ and (B) Ag−Ag₃PO₄(F,MW).

Figure 4. Effect of formaldehyde on the rate constant and degradation efficiency of rhodamine B.
from four major peaks at 367.26 eV (Ag^+ 3d_{5/2}), 373.26 eV (Ag^+ 3d_{3/2}), 367.87 eV (Ag^0 3d_{5/2}), and 373.87 eV (Ag^0 3d_{3/2}), respectively. In the spectra, both Ag^+ and Ag^0 show spin orbit splitting with an energy separation (∆E) of 6.0 eV. The result shows the formation of reduced metallic Ag, and from the peak area percentage value, it is approximated that the ratio of Ag^0 to Ag^+ is 1:2. In Figure 7C, the peak at 532.54 eV is attributed to adsorbed oxygen, while the major peak at 530.72 eV is due to lattice oxygen. The K alpha prime (Kα') peak at 234.52 eV (Figure 7B) and 527.74 eV (Figure 7C) is due to a nonmonochromatic X-ray source. The binding energy peak position at 132.52 eV can be ascribed to P 2p, corresponding to P=O in Ag₃PO₄ (Figure 7D). The XPS results confirmed the creation of silver nanoparticles on the surface of Ag₃PO₄. The samples prepared with formaldehyde and MW radiation were therefore designated as Ag–Ag₃PO₄.

The SEM images of Ag₃PO₄ and Ag–Ag₃PO₄ photocatalysts are presented in Figure 8. The Ag₃PO₄ particles exhibited an irregular spherical morphology with a nonuniform diameter. The particle sizes were estimated to be between 200 and 500 nm. The composite Ag–Ag₃PO₄ exhibited a similar morphology but with the deposits of Ag nanoparticles attached to the surface of Ag₃PO₄ (Figure 8B). The energy-dispersive X-ray spectroscopy (EDX) of Ag₃PO₄ and Ag–Ag₃PO₄ is presented in Figure 8E–F. The EDX mapping of Ag₃PO₄ (Figure 8C,D) revealed even distribution of the elements present in Ag₃PO₄. The silver composition was estimated to be 74 wt %, and it is close to the theoretical value of ca. 77 wt %. For Ag–Ag₃PO₄ modified using formaldehyde and MW irradiation (Figure 8E,F), the EDX mapping revealed a higher concentration of Ag (blue color) with an estimated Ag wt % of 83. This confirms the formation of Ag on the surface of Ag₃PO₄ as deduced from the XPS analysis. EDX analysis of the smaller nanoparticle deposits on the surface of Ag₃PO₄ revealed that it has higher concentrations of Ag and a relatively lower concentration of phosphorus and oxygen.

The SEM images of the composite Ag–Ag₃PO₄ synthesized by increasing the MW irradiation time from 3 to 5 min revealed extreme etching (patches) of the Ag₃PO₄ surface. SEM–EDX analysis of the patches that appeared on the surface of Ag₃PO₄ revealed a higher concentration of silver (Figure 9) and lower concentrations of phosphorus and oxygen. The concentration of silver for two different regions in Figure 9A,B was estimated to be 95.2 and 84.9 wt %, respectively. This depicts inhomogeneous silver distribution on the surface of Ag₃PO₄.

The X-ray diffraction (XRD) pattern (Figure 10) for Ag₃PO₄ has a well-defined crystalline Ag₃PO₄ and exhibits a body-centered cubic crystal structure of Ag₃PO₄ (JCPDS no. 06-0505). Despite the increase in the silver concentration on the surface of Ag₃PO₄, XRD spectra for Ag₃PO₄ and that of Ag–Ag₃PO₄(F,MW) were similar without the appearance of Ag peaks. Because the crystal structure of Ag₃PO₄ was preserved for Ag–Ag₃PO₄ despite the relatively lower concentrations of phosphorus and oxygen detected by the EDX analysis, it can be concluded that the modification of Ag₃PO₄ in the presence of formaldehyde and MW radiation was restricted to the surface of Ag₃PO₄. Coto et al. reported a similar observation when they decorated the surface of a 25 nm TiO₂ nanoparticle with Ag nanoparticles using formaldehyde. Reduction of Ag⁺ into Ag⁰ using formaldehyde and conventional heating resulted in the formation of a thin layer of Ag nanoparticles coating on the surface of TiO₂. Unlike XPS, XRD could not detect the presence of Ag nanoparticles. This further confirms that the reduction of Ag⁺ to Ag happened only on the surface of the Ag₃PO₄ particles. That is, the synthesis method adopted resulted in the surface of Ag₃PO₄ particles being coated with a thin layer of Ag nanoparticles. Because XPS is a surface-sensitive characterization technique (∼10 nm), the

![Figure 5](http://pubs.acs.org/journal/acsodf)  Effect of microwaving time on optical properties.

![Figure 6](http://pubs.acs.org/journal/acsodf)  Effect of microwaving time on photocatalytic activity.

95%. The rate constant also decreased from 0.7656 to 0.5703 min⁻¹. A similar trend was observed for all the samples synthesized with different volumetric amounts of formaldehyde.

X-ray photoelectron spectroscopy (XPS) analysis was conducted to confirm that the sample prepared using formaldehyde and MW (Ag₃PO₄–10 mL F–3 min MW) resulted in the formation of metallic Ag nanoparticles deposited on Ag₃PO₄. The survey scan of the sample shows the characteristic peaks of P 2p (132.5 eV), Mo 3d (225.0 and 227.9 eV), C 1s (284.8 eV), Ag 3d (367.5 and 373.5 eV), O 1s (531.1 eV), Ag 3p (574.1 eV), and O KLL (743 eV) (Figure 7A). The Mo 3d peak is contributed by the sample holder. The deconvoluted spectra for Ag 3d⁵/₂ and Ag 3d₃/₂ (Figure 7B) show four major peaks at 367.26 eV (Ag⁺ 3d₃/₂), 373.26 eV (Ag⁺ 3d₅/₂), 373.87 eV (Ag⁰ 3d₅/₂), and 373.87 eV (Ag⁰ 3d₃/₂), in which Ag⁺ and Ag⁰ correspond to Ag₃PO₄ and metallic Ag, respectively. In the spectra, both Ag⁺ and Ag⁰ show spin orbit splitting with an energy separation (∆E) of 6.0 eV. The result shows the formation of reduced metallic Ag, and from the peak area percentage value, it is approximated that the ratio of Ag⁰ to Ag⁺ is 1:2. In Figure 7C, the peak at 532.54 eV is attributed to adsorbed oxygen, while the major peak at 530.72 eV is due to lattice oxygen. The K alpha prime (Kα’) peak at 234.52 eV (Figure 7B) and 527.74 eV (Figure 7C) is due to a nonmonochromatic X-ray source. The binding energy peak position at 132.52 eV can be ascribed to P 2p, corresponding to P=O in Ag₃PO₄ (Figure 7D). The XPS results confirmed the creation of silver nanoparticles on the surface of Ag₃PO₄.

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presence of metallic Ag nanoparticles could be detected. However, XRD is a bulk-sensitive characterization technique, and the presence of metallic Ag nanoparticles in very low concentrations (<∼5%) when compared to the bulk Ag₃PO₄ could not be detected. The thin Ag nanoparticle coating and the rough surface morphology produced from the MW radiation are responsible for the relatively higher photocatalytic activity of Ag−Ag₃PO₄ when compared with that of Ag−Ag₃PO₄ synthesized using other methods. Surface etching of Ag−Ag₃PO₄ by the MW radiation resulted in surface morphology that aids in the higher separation and transfer efficiency of the photogenerated holes and electrons. Different parameters were investigated for their effect on the photocatalytic activity of the modified and bare Ag₃PO₄. Can the deposition of Ag on the surface of Ag₃PO₄ occur by heating the precipitated Ag₃PO₄ in formaldehyde and excess AgNO₃ solution instead of microwaving? The results show that the photocatalytic efficiency of Ag₃PO₄ can be enhanced by conventional heating (heating in a water bath at 80 °C) of Ag₃PO₄ in formaldehyde and excess AgNO₃. However, the samples prepared by heating [Ag−Ag₃PO₄(F, Heating)] had a relatively lower photocatalytic activity (Figure 11) and rate constant (Figure 12) when compared to [Ag−Ag₃PO₄(F, MW)]. It has been reported that MW radiation create Ag nanoparticles from AgNO₃ solution even in the absence of a reducing agent. The calculated rate constant (0.5581 min⁻¹) was higher than that of the unmodified Ag₃PO₄ (0.1696 min⁻¹). It is therefore probable that the MW radiation aided in the generation of more Ag nanoparticles. This observation implies that a heat source (either through microwaving or conventional heating) is needed in the modification process. The extreme etching (patches) observed in the [Ag−Ag₃PO₄(F,MW)] samples (Figure 9) was not observed in [Ag−Ag₃PO₄(F, Heating)], which indicates that the etching resulted from the MW radiation. The MW heating increases the rate of heating versus the water bath, thereby promoting rapid reduction of silver salts and nucleation over growth. This reduced the particle size of Ag while increasing the etching. As already mentioned, the etching of Ag₃PO₄ resulted in a surface morphology that has been reported to promote the transfer efficiency and separation of photogenerated holes and electrons. The EDX analysis of the few patches (Figure 13A,C) observed with the heated samples has a relatively lower silver concentration (ca. 67.4 wt %) when compared with that of [Ag−Ag₃PO₄(F, MW)]. It should however be noted that the smooth surface of [Ag−Ag₃PO₄(F, Heating)] has a relatively higher Ag concentration (80.8 wt %) (Figure 13B,C). The photocatalyst obtained without microwaving or heating but with the addition of formaldehyde [Ag−Ag₃PO₄(F)] showed a photocatalytic activity (k = 0.432 min⁻¹) that was higher than that of bare Ag₃PO₄ but lower than that of [Ag−Ag₃PO₄(F, Heating)] and [Ag−Ag₃PO₄(F, MW)]. This implies that MW radiation and conventional heating influenced the photocatalytic activity of the modified Ag₃PO₄.

After precipitating Ag₃PO₄, formaldehyde was added, and the entire mixture was stirred for 30 min. The liquid phase was decanted, and the precipitated photocatalyst was dispersed in DI water and microwaved. This photocatalyst was designated as [Ag−Ag₃PO₄(F-Decanting, MW)]. As can be seen in Figures A-C, the XPS analysis of Ag−Ag₃PO₄ (F, MW = 3 min) showed that the sample contained Ag0, Ag+ 3d₃/₂ and 3d₅/₂ peaks confirming the presence of Ag⁰. (C) XPS spectrum for O 1s, and (D) XPS spectrum for P 2p.
11 and 12, the photocatalytic activity \( k = 0.3914 \text{ min}^{-1} \)
obtained from this sample was higher than that of \( \text{Ag}_3\text{PO}_4 \) but
lower than that of \( [\text{Ag}−\text{Ag}_3\text{PO}_4(\text{F, Heating})], [\text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW})], \) and \( [\text{Ag}−\text{Ag}_3\text{PO}_4(\text{F})] \). The concentration
of Ag on the surface of \( [\text{Ag}−\text{Ag}_3\text{PO}_4(\text{F- Decanting,MW})] \) was
estimated through EDX (Figure 14A,C) as 55.2 wt %. This
further proves that for optimum photocatalytic activity, both
formaldehyde addition as the reducing agent and MW
radiation as heating are needed together.

In the absence of formaldehyde, the precipitated \( \text{Ag}_3\text{PO}_4 \)
was microwaved, and the photocatalytic activity was examined.
Again, the photocatalytic activity of the microwaved \( \text{Ag}_3\text{PO}_4 \)
\( [\text{Ag}_3\text{PO}_4(\text{MW})] \) with a rate constant of 0.3474 min\(^{-1}\) was
higher than that of \( \text{Ag}_3\text{PO}_4 \). This implies that microwaving also
affected the photocatalytic activity. The SEM–EDX spectra of
\( \text{Ag}_3\text{PO}_4(\text{MW}) \) are shown in Figure 14B,D. The concentration
of Ag on the surface was estimated to be ca. 63.08 wt %.

From the results presented above, it can be concluded that
modification of \( \text{Ag}_3\text{PO}_4 \) using formaldehyde and MW radiation
resulted in an increase in Ag concentration on the surface of
\( \text{Ag}_3\text{PO}_4 \), which enhanced the photocatalytic activity. The authors investigated the effect of \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW})/\text{dye} \)
weight ratio and catalyst concentration on the photocatalytic
activity. The results are presented in Figure 15. Figure 15 is
generated taking into consideration the time required to attain
almost 100% dye removal by \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW}) \).

Generally, as the wt ratio of the photocatalyst/dye and
photocatalyst concentration increased, the time required to
attain complete removal of the dye decreased. With a
minimum photocatalyst concentration of 0.125 g/L, the entire
concentration of dye was removed within 20 min. Increasing
the photocatalyst concentration decreased significantly the
time required to attain 100% dye removal. The time for
complete removal of dye leveled off (at 4 min) when the
catalyst concentration was increased beyond 0.375 g/L. A
similar trend was observed for the variation of wt ratio of
photocatalyst/dye with time required for complete dye
removal. The time required for complete dye removal leveled
off at 4 min when the wt ratio of the photocatalyst/dye was
increased beyond 75. These results are impressive because it
implies that a smaller quantity of \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW}) \) is
required to achieve complete dye removal. The excellent
photocatalytic degradation efficiency of \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW}) \)
could be attributed to the VB position and inductive effect of
\( \text{PO}_4^{3−} \). These promote efficient separation of photogenerated
electrons and holes as well as the localized surface plasmon
resonance of Ag nanoparticles.25 The photostability of the
composite photocatalyst was examined. The cycling degrada-
tion of rhodamine B dye by \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW}) \) is
presented in Figure 16. It can be deduced from Figure 16 that
the composite photocatalyst remained stable even after five cycles
of repeated photoreaction. There was approximately 10% reduction in the photodegradation efficiency after five cycles.

The potential application of the synthesized \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW}) \) in degrading crude oil was examined. Crude
oil was fractionated into benzene, \( n\)-hexane, and 1:1 v/v
methanol/benzene-soluble fractions. Each of these compo-
nents including the crude oil was subjected to photocatalysis
using \( \text{Ag}−\text{Ag}_3\text{PO}_4(\text{F,MW}) \). In a typical experiment, 100 mg of
the photocatalyst was added to crude oil or crude oil fraction

Figure 8. SEM images of (A) \( \text{Ag}_3\text{PO}_4 \) and (B) \( \text{Ag}−\text{Ag}_3\text{PO}_4 \). EDX mapping of (C,D) \( \text{Ag}_3\text{PO}_4 \) and (E,F) \( \text{Ag}−\text{Ag}_3\text{PO}_4 \) (F, MW = 3 min).
(0.2 mL) and synthetic seawater (120 mL) and irradiated using solar energy (sunlight) for 6 h. The degraded crude oil or crude oil fraction was extracted with dichloromethane (DCM) and analyzed with UV−vis and GC/MS. The UV−vis spectra for the degraded and undegraded crude oil and crude oil fractions are presented in Figure 17. It can be inferred from Figure 17 that the concentration of the crude oil and the various crude oil fractions decreased significantly after exposure to sunlight and Ag−Ag₃PO₄(F,MW) for 6 h. Ag−Ag₃PO₄(F,MW) was therefore effective at degrading crude oil. GC/MS analysis was conducted on the various crude oil fractions to identify the compounds that remained and
disappeared and intermediate compounds formed after photocatalysis. The chromatogram of the undegraded and degraded benzene-soluble fraction of the crude oil is presented in Figure 18. GC/MS analysis revealed the disappearance and the formation of new compounds. Nonadecane was the only compound found in both the degraded and undegraded benzene-soluble fractions. The undegraded benzene-soluble fractions contain several aromatics and paraffins. However, after the photocatalytic degradation, the aromatics: naphthalene-1-methyl (RT = 6.400 s), decahydro-4,8,9,10 penta methyl naphthalene (RT = 7.130 s), and naphthalene 1,4 dimethyl (RT = 7.745 s) disappeared, while a new aromatic compound diisooctyl phthalate (RT = 20.721 s) was formed. One noticeable observation was that though all the paraffins apart from nonadecane present in the benzene-soluble fraction disappeared after the photocatalysis, several new paraffinic compounds were formed. This implies that aromatics and cyclic alkanes are very sensitive to photocatalytic degradation and may be completely removed through photocatalysis, while the paraffins are less susceptible. Similar observation has been reported by D’Auria et al.26 Because aromatic compounds present in the crude oil are sensitive to photocatalytic degradation while paraffins are easily remediated through biodegradation,1h a more efficient oil spill remediation strategy can be developed by combining photocatalysis and chemical dispersant application.

The GC/MS chromatogram of the undegraded crude oil fraction soluble in a 1:1 v/v methanol/benzene mixture is presented in Figure 19A. The undegraded fraction was made up of two paraffins: octadecane 2-methyl (RT = 17.974 s) and...
octadecane 3-ethyl-5-(2-ethylbutyl) (RT = 23.287 s) and several unidentified complex mixtures (UCMs). After photodegradation using Ag−Ag₃PO₄(F,MW), almost all the UCMs observed in Figure 19A were resolved, and several new compounds were formed (Figure 19B). The two paraffins identified in the undegraded fraction disappeared after the photocatalysis. However, several paraffins were identified in the degraded photoproducts formed.

The GC/MS chromatogram for the degraded and undegraded n-hexane-soluble fraction is presented in Figure 20. Most of the compounds identified in this fraction are paraffins. Three of the compounds were aromatics: benzocycloheptatriene (RT = 6.190 s), decahydro-4,4,8,9,10 pentam-
ethylnaphthalene (RT = 7.345 s), and naphthalene 1,8 dimethyl (RT = 7.80). After degradation (Figure 20B), all the parafins in the undegraded fraction disappeared except for nonadecane. The area under the nonadecane peak for the degraded and undegraded fractions was almost similar. Several new paraffins were identified in the degraded fraction. In addition, the three identified aromatics in the undegraded fraction also disappeared after photocatalysis. However, diisoodyl phthalate (RT = 20.776), a new aromatic compound, was formed after photocatalysis.

From the GC/MS results, it can be concluded that photocatalysis is effective at degrading completely aromatic components in crude oil while forming fewer aromatic compounds as photoproducts. On the other hand, though photocatalysis can degrade paraffins, it leads to the formation of several new paraffin compounds.

From the results presented above, Ag−Ag3PO4(F,MW) has excellent photocatalytic activity for the degradation of dyes and crude oil fractions. High degradation efficiency can be attained within shorter visible light exposure time even at relatively lower wt ratio of photocatalyst/dye and lower photocatalyst concentration. To understand the degradation mechanism, the radical and hole trapping experiment was conducted. The reactive specie(s) responsible for the efficient photocatalytic activity of Ag−Ag3PO4(F,MW) was examined using t-BuOH, benzoquinone, and EDTA−Na2 as scavengers. Benzoquinone, t-BuOH, and EDTA−Na2 serve as scavengers for superoxide radicals (O2•−), hydroxyl radicals (OH•), and photogenerated holes (h+), respectively. Addition of benzoquinone, t-BuOH,
and EDTA−Na₂ reduced the photocatalytic efficiency from 98.1 to 48.2, 71.3, and 19.7%, respectively. This indicates that photogenerated holes (h⁺) are the main reactive species responsible for the photodegradation of rhodamine B by Ag−Ag₃PO₄(F,MW). This observation is consistent with literature reports.27 A possible photocatalytic degradation mechanism is presented below.

From the SEM−EDX images and analysis, a schematic diagram of the synthesized Ag−Ag₃PO₄ is presented in Figure 21A. Upon visible light irradiation, photons can be absorbed by both Ag₃PO₄ and Ag. This will result in the generation of electrons (e⁻) and holes (h⁺). The electrons in the VB of Ag₃PO₄ are then excited to the CB (Figure 20B). The localized surface plasmon effect produced by the collective oscillation of surface electrons on the Ag nanoparticles could generate a local inner magnetic field, which can enhance the separation of electrons and holes (Figure 21C,D). Because of the localized magnetic field and the good electron conductivity of Ag, the photogenerated electrons are quickly transferred from Ag₃PO₄. Because the electrons are transferred far from the Ag⁺ ions in Ag₃PO₄, the stability of Ag−Ag₃PO₄ is enhanced. That is, the electrons generated are transferred to the Ag

13651

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nanoparticles with Ag acting as the electron acceptor. The photogenerated holes on the other hand remain on the surface of Ag$_3$PO$_4$ reducing the electron–hole recombination rate. Another possible explanation for the enhanced photocatalytic activity is presented as follows. Because Ag$_3$PO$_4$ is an n-type semiconductor, its Fermi level is close to the CB edge (4.95 eV). When the Ag–Ag$_3$PO$_4$ heterojunction is created, electrons flow from Ag$_3$PO$_4$ to Ag. The Ag–Ag$_3$PO$_4$ heterojunction results in an equilibration of the Fermi level. The new Fermi level attained from this equilibration is close to the CB of Ag$_3$PO$_4$. Hence upon visible light irradiation, Ag acts as an electron acceptor, allowing the available photoexcited holes to be available for photocatalysis because the electron–hole recombination is reduced. Tastuma and Takada have reported that increase in electron concentration on the Ag nanoparticles resulting from the equilibration of the Fermi level may initiate a series of reduction reactions, while the high concentration of photoexcited holes on Ag$_3$PO$_4$ will initiate a series of photooxidation reactions that may result in the degradation of organic pollutants such as dyes and crude oil fractions. Furthermore, the phosphate ions in Ag$_3$PO$_4$ have strong affinity for water. Hence, H$_2$O easily binds to the surface of Ag$_3$PO$_4$. With the presence of holes on Ag$_3$PO$_4$, hydroxyl radicals can be easily formed by the oxidation of H$_2$O molecules by holes. These hydroxyl radicals oxidize dyes and crude oil fractions into photogenerated byproducts and ultimately carbon dioxide and water. However, the radical and hole trapping experiment revealed that photoexcited holes are the main reactive species responsible for the degradation of dyes; hence, the contribution from hydroxyl radicals may not be significant. The relatively lower photocatalytic activity of bare Ag$_3$PO$_4$ can be attributed to fast electron–hole recombination and the relatively lower photostability.

3. CONCLUSIONS

In summary, a highly efficient Ag–Ag$_3$PO$_4$ visible light photocatalyst was synthesized by an ion-exchange method and modified by using excess AgNO$_3$, formaldehyde, and MW radiation. The formaldehyde and MW radiation helped in the deposition of Ag on the surface of Ag$_3$PO$_4$ and the formation of high Ag-concentrated patches on the surface of Ag$_3$PO$_4$. Instead of MW radiation, a conventional heating method can be used but with a relatively reduced photocatalytic activity when compared with the composite photocatalyst synthesized with MW radiation. Ag–Ag$_3$PO$_4$ was stable and efficient for the degradation of rhodamine B dyes and crude oil fractions. GC/MS analysis of the photoproducts generated from the degradation of the crude oil fraction revealed that photocatalysis is effective in completely degrading aromatic components of crude oil with little generation of aromatic photoproducts. On the other hand, photodegradation of paraffins results in the generation of several paraffin photoproducts. The enhanced photocatalytic activity was attributed to the localized surface plasmon effect of Ag and the reduction of the electron–hole recombination.

4. EXPERIMENTAL SECTION

4.1. Materials. AgNO$_3$ (ACS reagent, ≥99%), Na$_3$HPO$_4$ (99.95%), rhodamine B dye (97%), benzene (analytical standard), hexane (laboratory reagent, ≥95%), methanol (HPLC 99.9%), and formaldehyde (37 wt % in H$_2$O) were purchased from Sigma-Aldrich, UK. Crude oil was obtained from Tullow Oil, Ghana.

4.2. Experimental Procedure. 4.2.1. Synthesis of Ag–Ag$_3$PO$_4$. In a typical synthesis procedure for making the photocatalytic system, 50 mL of 0.05 M Na$_3$HPO$_4$ was added dropwise to 50 mL of 0.2 M AgNO$_3$ while stirring vigorously at ambient temperature. The resulting solution was stirred for 30 min while forming the Ag$_3$PO$_4$ precipitate. A known volume of formaldehyde solution was added and continuously stirred for another 30 min. The resulting solution with the precipitate was then subjected to MW radiation of 700 W at 2450 MHz for 3 min. The resulting (shown to be Ag–Ag$_3$PO$_4$) particles were collected by centrifugation and dried at 70 °C for 5 h. In another synthesis variant, after adding formaldehyde and stirring for 30 min, the resulting solution was heated in a water bath set at 80 °C for 3 min. The precipitate was collected by centrifugation and dried at 70 °C for 5 h. The effect of
formaldehyde (F) and MW or thermal heating on the resulting materials was investigated. The conditions for the experiments are summarized in Table 1.

4.2.2. Fractionating of Crude Oil. The crude oil was separated into three fractions. This was achieved by adding 50 mL of crude oil to 181 g of silica gel in a chromatographic column. n-Hexane (500 mL) was first added to the crude oil in the column to elute the n-hexane soluble fraction. Second, 500 mL of benzene was also added to the residual crude in the column for the elution of the benzene-soluble fraction. Lastly, a solution of 250 mL of methanol and 250 mL of benzene was again added to the residual crude after the second elution to elute the 1:1 v/v methanol/benzene-soluble fraction. The eluate samples of the various soluble fraction concentrates were formed by evaporating the solvent–crude oil fraction mixture using the rotary evaporator at 40 °C.

4.3. Product Characterization. 4.3.1. XRD. The powder XRD patterns were obtained with a Bruker D8 ADVANCED focus diffractometer fitted with a position-sensitive detector (LynxEye) and a standard detector. Cu Kα radiation (λ = 0.15405 nm) and a 20 angular range of 10–80° were used.

4.3.2. X-ray Photoelectron Spectroscopy. XPS of the Ag3PO4 sample was analyzed using a Kratos XSAM 800 photoelectron spectrometer connected with dual-anode X-ray source. Nonmonochromatic Mg Kα (1253.6 eV) X-ray source was used. The pressure in the analyzing chamber was less than 10⁻⁹ mbar. The sample spectra were analyzed using the CasaXPS software, and the peak shift due to any apparent charging was calibrated with respect to the C 1s peak set to 284.8 eV. A standard Shirley background with the product Gaussian (50%) and Lorentzian (50%) functions was used to fit each component spectrum.

4.3.3. Scanning Electron Microscopy with Energy-Dispersive X-ray Spectroscopy. The morphology of the synthesized nanocomposites was examined with a scanning electron microscope (FEI Nova NanoSem) connected to the EDX acquisition detector. The elemental composition was determined through EDX.

4.3.4. Diffuse Reflectance Spectroscopy (DRS). DRS was conducted using an Ocean Optics USB-4000 UV–vis spectrometer equipped with a dedicated reflectance probe. Glass slides were used to compress the synthesized nanocomposite into a flat film. The DRS measurement was conducted using polytetrafluoroethylene reflectance standard for reflectance calibration.

4.3.5. Photocatalytic Degradation of Rhodamine Blue. The photocatalytic activity of the synthesized Ag3PO4 was examined using rhodamine B dye. In a typical photocatalytic degradation test, 200 mL of rhodamine B solution (5 mg/L) in deionized water was used with an appropriate Ag3PO4 suspension concentration. A jacketed glass reactor with a quartz tube immersion well was used with illumination from a 300 W tungsten halogen lamp. For each of the experiments, the solution was stirred vigorously for 30 min in the dark to attain an adsorption–desorption equilibrium. An aliquot (2 mL) was removed at specific time intervals and centrifuged at 6000 rpm for 5 min to separate the powdered photocatalyst. The dye absorbance was measured using a UV–vis spectrophotometer (Ocean Optics 4000 USB) at an absorbance difference of 554–400 nm. The effect of photocatalyst concentration was examined by using varying amounts of Ag3PO4.

4.3.6. Photocatalytic Degradation of Crude Oil Fractions. The crude oil fraction (0.2 mL) was measured and added to 120 mL of synthetic seawater in a 200 mL beaker. The photocatalyst (100 mg) was added to the mixture of the crude oil fraction and distilled water. The resultant mixture was irradiated under sunlight with continuous stirring at 350 rpm to ensure that the photocatalyst is dispersed well in the water–crude oil mix. Irradiation was carried out continuously for 6 h, after which 10 mL of DCM was added to the resultant mixture to dissolve the degraded crude oil. DCM with the dissolved degraded crude oil fraction was separated from the photocatalyst and synthetic seawater using a separation funnel. The dissolved oil is further centrifuged at 6000 rpm for 15 min to ensure that all residual photocatalyst particles are separated. DCM with the dissolved crude oil fractions was characterized with a UV–vis spectrometer and a GC–MS spectrometer.

4.3.7. Gas Chromatography–Mass Spectrometry. DCM with the dissolved crude oil fraction was analyzed with GC/MS to identify the degraded byproducts from the photocatalysis. The following GC/MS conditions were employed; the oven was heated to an initial temperature of 80 °C, then ramped at 10 °C/min to 240 °C, held for 2 min, then ramped at 5 °C/ min to 280 °C, and held for 10 min; the injection temperature selected was 250 °C, with an injected volume of 0 μL. The split ratio used was 20:1. Helium was selected as the carrier gas. The solvent delay time was 2.50 min. The transfer temperature used was 250 °C, while the source temperature was 150 °C. The GC/MS column dimension was 27.0 m × 250 μm.

Table 1. Summary of Experimental Conditions Used for the Preparation of the Photocatalysts

<table>
<thead>
<tr>
<th>sample</th>
<th>formaldehyde (F)</th>
<th>microwaving for 3 min (MW)</th>
<th>heating at 80 °C for 3 min (Heating)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ag3PO4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ag3PO4 (MW)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ag–Ag3PO4(F, Heating)</td>
<td>√</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ag–Ag3PO4(F, Decanted, MW)</td>
<td>(the resulting solution after the addition of F was decanted, and then the precipitates were dispersed in DI water and MW)</td>
<td>√</td>
<td></td>
</tr>
</tbody>
</table>
Complete contact information is available at: https://pubs.acs.org/10.1021/acsomega.0c00670

Notes
The authors declare no competing financial interest.

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REFERENCES
(12) (a) Nayak, S.; Parida, K. M. Dynamics of Charge-Transfer Behavior in a Plasmon-Induced Quasi-Type-II p–n–n–n Dual Heterojunction in Ag@Ag3PO4/g-C3N4/NaF LDH Nanocomposites for Photocatalytic Cr(VI) Reduction and Phenol Oxidation. ACS Omega 2018, 3, 7324–7343. (b) Shi, W.; Li, M.; Huang, X.; Ren, H.; Yan, C.; Guo, F. Facile synthesis of 2D/2D Co3(PO4)2 heterojunction for highly photocatalytic overall water splitting under visible light. Chem. Eng. J. 2020, 382, 122960.


(20) Teng, W.; Li, X.; Zhao, Q.; Chen, G. Fabrication of Ag/Ag3PO4/TiO2 heterostructure photoelectrodes for efficient decomposition of 2-chlorophenol under visible light irradiation. J. Mater. Chem. A 2013, 1, 9060−9068.


(30) Teng, W.; Li, X.; Zhao, Q.; Zhao, J.; Zhang, D. In situ capture of active species and oxidation mechanism of RhB and MB dyes over sunlight-driven Ag/Ag3PO4 plasmonic nanocatalyst. Appl. Catal., B 2012, 125, 538−545.