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In Situ Photo-Fenton-like Reaction and Oxygen Evolution through Polaron-Mediated WO_3

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ABSTRACT. The effective utilization of both electrons and holes in semiconductor-based photocatalysis is crucial for enhancing light-driven chemical reactions. However, the photocatalytic performance of WO_3 is limited by its low conduction band potential, which restricts its ability to generate reactive species such as superoxide radicals and to facilitate hydrogen production. Consequently, WO_3 primarily relies on holes for redox processes, reducing its overall photocatalytic efficiency. This paper proposes a novel strategy to simultaneously exploit both

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charge carriers by activating a photo-Fenton-like reaction driven by polaronic phenomena of electrons (W^{5+} formation). Electrons are responsible for generating hydroxyl radicals (•OH), enabling the degradation of Acid Orange 7 dye with an activity ten times higher than that of the conventional Fenton reaction. Meanwhile, holes effectively drive the oxygen evolution reaction (OER), achieving an oxygen yield of 1.62 mmol g^{-1} h^{-1}. The polaronic behaviour of WO_3 was further optimized by accelerated supersaturation process through protonation which enhances the surface properties and increase W^{5+} formation during light irradiation. This strategy led to approximately threefold enhancement in Acid Orange 7 degradation and a 2.3-fold increase in OER performance (3.66 mmol g^{-1} h^{-1}, one of the highest reported values). Overall, this study presents a new approach to optimizing charge carrier utilization in photocatalysis, offering a promising pathway toward more efficient and versatile semiconductor-based systems.

1. INTRODUCTION

The photo-Fenton reaction has emerged as a promising solution for wastewater treatment due to its strong oxidative capability in decomposing a wide range of pollutants[1], including synthetic dyes[2,3], pharmaceuticals,[4] and radioactive waste[5]. This process involves the reaction of iron salts with hydrogen peroxide, generating highly reactive hydroxyl radicals (•OH) that act as powerful oxidizing agents. However, the practical application of the Fenton reaction faces a major limitation: the residual iron salts remain in the solution after the reaction and are difficult to recover, potentially causing secondary pollution in water sources.[6] To address this issue, researchers have explored immobilizing iron sources in the form of iron oxides/hydroxides or anchoring them onto various supports such as metal oxide semiconductors[7,8], metal–organic

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frameworks (MOFs)[9], and carbon-based materials[10]. Despite these advances, iron leaching—particularly under acidic conditions—remains a significant challenge,[11,12] limiting the long-term stability and sustainability of the process.

Instead of focusing solely on stabilizing iron ions, a more promising strategy is to develop stable materials that can generate hydroxyl radicals (•OH) through interactions with H_2O_2. For instance, previous studies have reported that TiO_2 with a high concentration of Ti^{3+} defects can serve as effective candidates. However, these Ti^{3+} defect sites are not reversible, leading to gradual performance degradation over time.[13] Other materials also such as CuO with Cu^{2+}[14] content and Mn_3O_4[15] facing low stability for long term usage. In contrast, tungsten trioxide (WO_3) offers a more structurally stable[16] alternative with defect reversibility due to polaronic behavior where the photoexcited electron localize on tungsten sites with the local structure distortion W^{6+} to W^{5+}.[17,18] which is similar with Fe^{2+}/Fe^{3+} making it a promising material for sustained hydroxyl radical generation.

Moreover, the utilization of electrons to generate hydroxyl radicals (•OH) allows the holes in WO_3 to be more freely available for driving chemical reactions which enable WO_3 as photo-Fenton like reaction as well as photocatalytic reaction. Several studies have reported that the holes in WO_3 can promote the Oxygen Evolution Reaction (OER) in the presence of electron scavengers such as Ag^+ and Fe^{3+}.[19] However, the activation of OER in these systems is limited due to the non-recyclable nature of the scavenging process. In contrast, in the Photo-Fenton-like reaction using WO_3, the electron scavenging process is expected to be efficiently recyclable, which could lead to enhanced OER efficiency.

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Therefore, in this study, the novel utilization of both charge carriers in the direct photo-Fenton-like reaction of WO_3 by acknowledging the polaronic behavior of WO_3 was investigated. Recently, interesting polaron behavior in small nanoparticles driven by lattice distortion are reported for metal oxides including particle size effects and surface heterogeneity of polaron,[20–23] and thus nanoparticle synthesis of WO_3 is also expected to exploit polaronic behavior. The utilization of both charge carriers, electrons and holes, by the photo-Fenton reaction was monitored by wastewater removal for electrons and oxygen evolution reaction for holes at the same time. Currently, the usage of photo-Fenton-like reaction of WO_3 always requires iron support to activate the reactions[24,25] which are probably attributed to the inefficient polaron formation of WO_3. Therefore, in this study, the efficiency of the polaron formation was further optimized by understanding the growth of WO_3, which is controlled by the control of protonation process, enabling us to understand the crystallization difference that affects the photo-Fenton like efficiency. This study will provide new insights into the utilization of WO_3 as a catalytic material for promoting chemical reactions.

2. EXPERIMENTAL DETAIL

2.1 WO_3 synthesis

Initially, 3.3 g of disodium tungstate dihydrate ($Na_2WO_4 \cdot 2H_2O$) (Fujifilm Wako Pure Chemical co.) was dissolved in 200 mL of deionized water under continuous stirring with a magnetic stirrer. Once fully dissolved, the solution was protonated by adding varying amounts of 37% HCl (Fujifilm Wako Pure Chemical co.). The protonation process was controlled by adjusting the volume of HCl to 15 mL, 30 mL, 60 mL, and 180 mL, corresponding to proton concentration ($[H^+]$) were 0.8, 1.6, 2.8 and 5.7 M **(Supporting Information, (S1))**, hereafter referred to as $[H^+]$

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= 0.8, $[H^+]$ = 1.6, $[H^+]$ = 2.8, and $[H^+]$ = 5.7, respectively. The solution was stirred continuously for 12 hours at room temperature. After stirring, the precipitate was collected by centrifugation and washed with deionized water until the pH of the supernatant reached 4–5. The washed sample was then dried under vacuum at 60 °C for 12 hours. Finally, the dried products underwent annealing at 400 °C for 2 hours under ambient air atmosphere, yielding the final WO_3 product.

2.1 Materials characterization

The morphology of the particles was investigated by transmission electron microscope (JEM-2100Plus, JEOL, Japan). Crystal structure and crystallite size were determined using X-ray powder diffraction (XRD, Rigaku SmartLab 9MTP, Japan) with a Cu Kα (λ =1.5418 Å) radiation sources at 3˚/min. The 2θ scanning range is 20-80˚. The chemical states of the samples were investigated by X-ray photoelectron spectroscopy (XPS; PHI5600, ULVAC-PHI, Japan.) which was performed using monochromatized X-rays of Al Kα with 300 W power source, the fitting of tungsten elements were conducted from 33 to 40 eV with Shirley baseline methods. The optical absorption of was monitored by UV-Vis-NIR V-770TWK, JASCO, Japan and the luminescence analysis was investigated by Fluorescence Spectrophotometer (FL), F-2700 Hitachi High-Technologies, Japan. The local structure of WO_3 was further characterized by a microscopic confocal Raman spectroscopy (HORIBA LabRAM HR-800, Japan) with an excitation wavelength of 532 nm. The defect analysis then investigated by electron paramagnetic resonance (EPR) (B Bruker BioSpin, Bruker, Germany).

2.2. Photo-Fenton like reaction

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The photo-Fenton-like reaction was evaluated by monitoring the degradation of Acid Orange 7 in the presence of WO_3, H_2O_2, and UV light (detailed experimental set-up see **Supporting Information, (S2)**). Typically, a 20 mg L^{-1} Acid Orange 7 solution was prepared in 50 mL of water. The pH was then adjusted to 2 by adding a 10% HCl solution. Afterwards, 40 mg L^{-1} of the WO_3 catalyst was added, and the suspension was stirred for 30 minutes under a 200 mL min^{-1} N_2 atmosphere to ensure sufficient adsorption–desorption equilibrium. Subsequently, 2 mL of 30% H_2O_2 was introduced and stirred for 5 minutes. The mixture was then irradiated using a 22 W UV lamp (approx.: 70 mW cm^{-2}, 365 nm). A 3 mL aliquot of the solution was collected, centrifuged to remove residual particles, and analyzed by UV–Vis spectroscopy (JASCO V-570, Japan). Meanwhile, 0.5 mL of the evolved gas was sampled and injected into a gas chromatograph (GC–Shimadzu 2014, Japan) equipped with a thermal conductivity detector to identify gaseous products (detailed calibration procedures are provided in the **Supporting Information, (S2)**. For •OH detection via the terephthalic acid fluorescence method, 0.04 g of WO_3 catalyst was dispersed in a ~4.8×10^{-4} M terephthalic acid solution, followed by the addition of 2 mL of H_2O_2. The suspension was then irradiated under UV light. After irradiation, fluorescence intensity was measured using a Fluorescence Spectrophotometer (F-2700, Hitachi High-Technologies, Japan) at an excitation wavelength of 325 nm. For the scavenger experiments, two scavengers were employed: 0.059 mmol silver nitrate ($AgNO_3$) as an electron scavenger and 21 mmol tert-butanol as an •OH radical scavenger. Each scavenger was added separately to the photo-Fenton-like reaction system, and the corresponding degradation performance was subsequently monitored.

3. RESULTS AND DISCUSSION

Figure 1(a) illustrates the degradation of Acid Orange 7 under different reaction conditions. Under WO_3 with UV light and WO_3 with H_2O_2 there are no degradations observed, indicating that neither

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process alone facilitates a redox reaction. However, when H_2O_2 is exposed to UV light (H_2O_2 + UV), approximately 10% degradation occurs, likely due to the generation of hydroxyl radicals from H_2O_2 under UV irradiation.[26] In contrast, a significant increase in degradation efficiency is observed when combining UV light, WO_3, and H_2O_2, reaching approximately 50% degradation. This strong degradation suggests a synergistic effect among UV light, WO_3, and H_2O_2.

The high degradation efficiency observed only in the presence of WO_3, UV light, and H_2O_2 has similarity with the Photo-Fenton reaction, where Fe^{2+} reacts with H_2O_2 and UV light to accelerate hydroxyl radical generation.[26,27] Given this similarity, it is highly likely that a similar mechanism occurs in WO_3. To clarify this assumption, XPS analysis of the W 4f state before and after UV irradiation was performed, as shown in **Figures 1(b)–(c).** The results reveal a gradual increase in the W^{5+} content from 2.5% to 3.6% after UV irradiation. This transition is also visually confirmed by the photographic evidence in **Figure 1(d)**, which shows a distinct color change from bright yellow to a greenish hue, attributed to the formation of W^{5+}, a phenomenon known as photochromism.[28,29]

The transition from W^{6+} to W^{5+} plays a crucial role in promoting the Photo-Fenton-like reaction, as described by the following reaction pathway:

$$UV + WO_3 \rightarrow e^- + h^+ \quad (1)$$

$$e^- + W^{6+} \rightarrow W^{5+} \quad (2)$$

$$W^{5+} + H_2O_2 \rightarrow \cdot OH + W^{6+} + OH^- \quad (3)$$

$$4h^+ + 2H_2O \rightarrow O_2 + 4H^+ \quad (4)$$

$$h^+ + OH^- \rightarrow \cdot OH + H^+ \quad (5)$$

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Under UV irradiation, WO_3 generates electron-hole pairs, where photogenerated electrons reduce W^{6+} to W^{5+} on the surface while holes on O 2p states trapped in oxygen sites ready for promoting possible chemical reaction.[30] The W^{5+} species then interact with H_2O_2, further promoting the formation of hydroxyl radicals which decompose Acid Orange 7 into carbon dioxide (Direct evidence of CO_2 evolution has been shown in the **Supporting Information, (S3**) and water, as illustrated in **Figure 1(e).** Interestingly, the Photo-Fenton-like activity of WO_3 shows better catalytic reaction than traditional Fenton reaction either by using $FeCl_2$ ion and Fe_2O_3 nanoparticles as shown in **Table 1** which highlight the strong potential of WO_3 as an effective material for waste-water degradation by photo-Fenton reactions.

While the electron continuously generates •OH through photo-Fenton like reaction process, holes in the valence band are ready to promote chemical reactions. There are two possible reactions for holes in these reaction conditions. The first one is the reaction with OH^- for generating •OH. This reaction is one of the common reactions for WO_3, where WO_3 has been acknowledged as good oxidative photocatalyst.[31] However, based on the catalytic reaction process, if •OH is formed, the degradation of Acid Orange 7 should occur even without H_2O_2. It indicates that the •OH radicals are not formed in this reaction condition. The second possibility is the water oxidation process where holes directly oxidize water into O_2 and H^+.[19]

To confirm the second reaction possibility, oxygen evolution during the Photo-Fenton-like reaction was investigated. To eliminate the possibility of O_2 release from direct H_2O_2 decomposition, a control experiment was conducted using H_2O_2 under UV irradiation alone. The calibration details for the O_2 evolution reaction are provided in **Supporting Information, (S2)**. As shown in **Figure 1(f)**, UV irradiation of H_2O_2 alone generates approximately 0.62 mmol g^{-1} of O_2,

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with an O_2 evolution rate of about 0.21 mmol·g^{-1}·h^{-1}. In the presence of WO_3 under UV irradiation, the O_2 production increases significantly to 3.1 mmol·g^{-1}, confirming the enhanced photocatalytic activity. To exclude the contribution from H_2O_2 photodecomposition, the O_2 evolution rates were corrected by subtracting 0.21 mmol g^{-1} h^{-1}. After correction, the O_2 evolution rate for WO_3 under UV irradiation is approximately 0.80 mmol g^{-1} h^{-1} and further increases to about 1.62 mmol g^{-1} h^{-1} when WO_3 is combined with H_2O_2 (see **Supporting Information, (S4)**). These results demonstrate that both photogenerated electrons and holes in WO_3 are effectively utilized, overcoming the previously reported limitations associated with WO_3 photocatalysis. Furthermore, the O_2 evolution performance achieved in this work is higher than values reported in the literature (**Table 1**), reinforcing the enhanced photo-Fenton-like activity of WO_3.

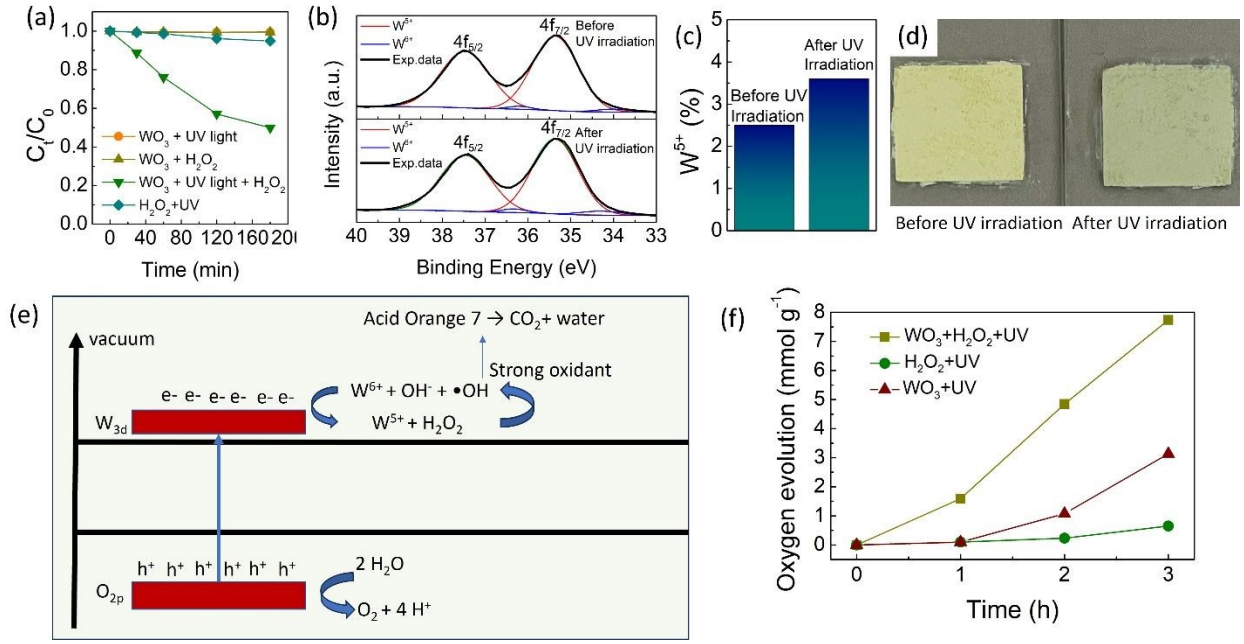

Figure 1. (a) Acid Orange 7 degradation under various systems. (b) XPS analysis of WO_3 before and after UV irradiation. (c) W^{5+} percentage obtained from XPS deconvolution. (d) Photographic images of WO_3 before and after reaction. (e) Proposed photo-Fenton reaction by using WO_3. (f) Oxygen evolution reaction by WO_3+H_2O_2+UV compared with WO_3+UV and H_2O_2+UV.

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Table 1. Photocatalytic performance for Acid Orange 7 degradation and oxygen evolution reaction of Polaron $WO_3 + H_2O_2$ compared with other reports and data.

Sample	degradation rates (h^{-1})	References	Sample	O_2 Evolution rate (mmol g^{-1} h^{-1})	References
$Fe_2O_3 + H_2O_2$	0.06	[32]	$Fe + TiO_2$	0.54	[33]
$Fe_2O_3 + H_2O_2$	0.13	[34]	$Ag + TiO_2$	0.06	[35]
$Fe + H_2O_2$	0.03	This works	$FeOOH + WO_3 + H_2O_2$	0.03	[25]
Polaron $WO_3 + H_2O_2$	0.18	This works	Polaron $WO_3 + H_2O_2$	1.62	This works

The WO_3-based photo-Fenton-like reaction has been demonstrated; however, the degradation efficiency remains relatively low (50% in 180 minutes). Therefore, further optimization strategies are necessary. The key to enhancing the photo-Fenton-like activity in WO_3 lies in the generation of W^{5+} species. Increasing the formation of W^{5+} under UV irradiation is essential. Previous studies have proposed that defect engineering can play a crucial role in promoting W^{5+} generation,[36] such as hydrogen annealing[37] or doping[38,39], can improve photochromism. However, these approaches introduce additional processing steps and materials, reducing overall efficiency.

Different from the above-mentioned techniques, the control of the nucleation and growth process during synthesis also provides a controllable of lattice behavior including strain and defect, which is insightful for improving W^{5+} formation in WO_3. In this study, W^{5+} formation was improved by regulating crystal growth through modulating the protonation level during synthesis. The proton concentration varied from $[H^+] = 0.8$, 1.6, 2.8 and 5.7. Initially the precursors are still in the form of (H_4WO_5) (see XRD and TEM results in **Supporting Information, (S5)**) and reduces

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the particle size of the precursor as increase of the protonation indicating the increase of the supersaturation process due to high protonation. After annealing treatment, the stable WO_3 is formed (detail structure analysis at **Figure 4(a)-(b)**). The change in the acid concentration significantly affects W^{5+} formation in the WO_3 structure. **Figure 2(a)** shows images of WO_3 at different protonation levels after UV irradiation. Clear evidence demonstrates that after UV irradiation, the color gradually changed to dark green as protonation increased, indicating an enhancement in W^{5+} generation. XPS analysis of W4f after and before UV irradiation has confirmed that the W^{5+} gradually increase with increasing protonation level from $[H^+]$ 0.8 to 5.7 (**Figure 2(b)**). **Table 2** also shows the detail of W^{5+} percentages after UV irradiation which indicates a gradually increase with increase of protonation. Besides W^{5+} formation, the N_2 adsorption desorption analysis results (**Figure 2(c)**) have revealed that the high protonation also increases the surface area, pore size and pore volumes of WO_3 as tabulated in **Table 2**.

Table 2. W^{5+} concentration, and physical surface properties of WO_3 at different protonation.

Sample	W^{5+} % after irradiation	Surface Area ($m^2\ g^{-1}$)	Pore Size (nm)	Pore Volume ($cc\ g^{-1}$)
$[H^+] = 0.8$	2.2	15	3.061	0.066
$[H^+] = 1.6$	3.6	24	2.178	0.244
$[H^+] = 2.8$	5.5	29	29.47	0.331
$[H^+] = 5.7$	6.7	37	32.50	0.247

The influence of increasing protonation was further evaluated through the photo-Fenton reaction. **Figure 2(d)–(e)** presents the Acid Orange 7 degradation profiles at different protonation levels. The degradation efficiency of WO_3 increases progressively with higher protonation, with

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the reaction rate rising from 0.18 to 0.68 h^{-1}. This trend clearly indicates that protonation enhances the photo-Fenton activity. At this stage, two major factors appear to affect the photo-Fenton performance: (i) the formation of W^{5+} species and (ii) the changes in surface properties. To clarify their respective contributions, several additional analyses were performed. First, the adsorption of Acid Orange 7 on WO_3 was examined to evaluate the influence of surface characteristics. As shown in the **Supporting Information, (S6)**, Acid Orange 7 adsorption increases slightly with increasing surface area, with approximately 5% removal observed. This indicates that surface properties also contribute to the degradation process. This conclusion is further supported by the surface charge analysis and photo-Fenton-like reactions performed at different pH values (**Supporting Information, (S7)**) where the surface charge and types of pollutant affected the photo-Fenton reactions, both of which confirm that surface characteristics play a role in accelerating the photo-Fenton reaction. On the other hands, to assess the contribution of W^{5+} species, photo-Fenton reactions were conducted using samples with comparable surface areas but different internal defect (W^{5+}) concentrations conducting by heated the $[H^+] = 5.7$ precursor under air and nitrogen atmosphere (see **Supporting Information, (S8)**). The results clearly show that the higher W^{5+} significantly enhances the degradation rate although the surface area nearly similar. Taken together, these data demonstrate that the improved photo-Fenton performance arises from the synergistic effect between increased formation of W^{5+} and the enhancement of surface properties induced by protonation. While surface properties mainly influence the attachment of Acid Orange 7 onto WO_3, the formation of W^{5+} is primarily responsible for promoting the conversion of H_2O_2 into •OH radicals.

Figure 2(f) presents the terephthalic acid fluorescence analysis, confirming •OH generation under irradiation. The samples with higher protonation exhibit the highest fluorescence

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intensity, indicating enhanced •OH production which correlated with improvement of Acid Orange 7 degradation. However, although •OH plays a key role in Acid Orange 7 degradation, the initial process was starting from the photoexcitation of electrons that generate W^{5+}, which then reacts with H_2O_2 to produce •OH radicals. This conclusion is further supported by the scavenger experiments shown in **Figure 2(g)**, where the presence of electron scavengers results in the greatest suppression of catalytic activity, clearly demonstrating that electron-driven polaron formation is the major contributor to the overall reaction.

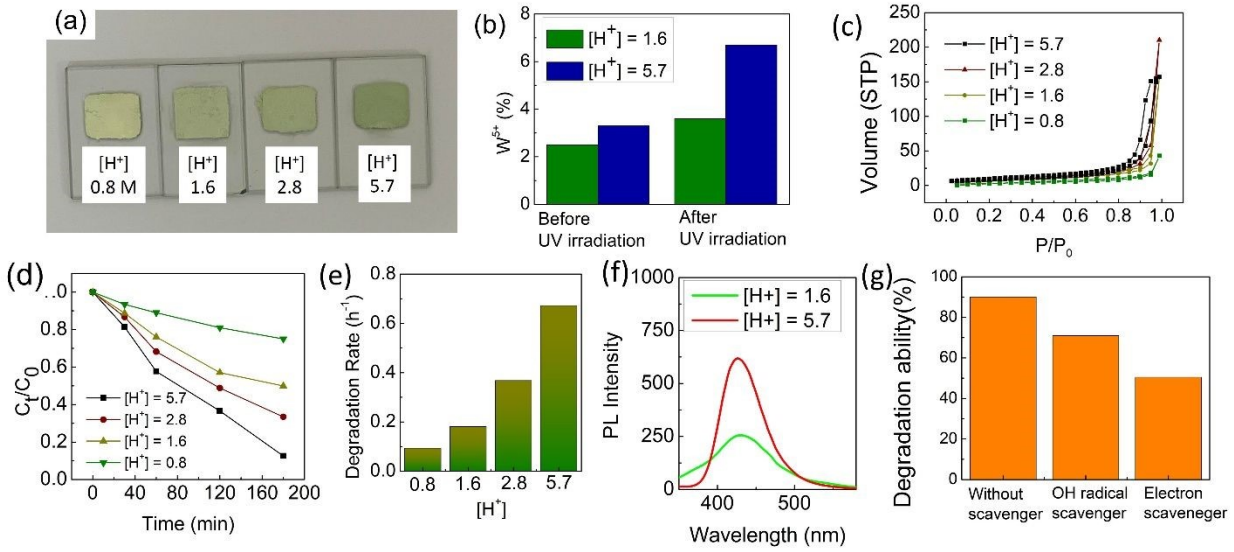

Figure 2. (a) Photographic images of WO_3 samples with different protonation levels after UV irradiation. (b) W^{5+} percentages before and after UV irradiation, obtained from XPS analysis of the W 4f region. (c) N_2 adsorption–desorption isotherms of WO_3 samples with different protonation levels. (d) Acid Orange 7 degradation profiles for WO_3 samples with varying protonation. (e) Degradation rate constants for Acid Orange 7 using WO_3 with different protonation levels. (f) Fluorescence analysis using terephthalic acid to detect •OH radical generation for $[H^+]$ = 1.6 and 5.7, and (g)scavenger tests using •OH scavengers and electron scavengers.

To assess the potential of this photo-Fenton-like reaction by applications, the degradation of other types of pollutants was also investigated (**Figure 3(a)**). Cationic dyes, represented by methylene blue, and pharmaceutical waste, represented by 4-Nitrophenol, were studied. The

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results show that the photo-Fenton reaction using WO_3 with high W^{5+} effective in degrading various types of pollutants, demonstrating its potential for future wastewater management applications.

The improvement of electron site efficiency is directly correlated with the improvement in the hole efficiency. As shown in **Figures 3(b)**, the OER rate increases steadily from 1.62 mmol g^{-1} h^{-1} to 3.66 mmol g^{-1} h^{-1}. This data demonstrates a clear correlation between the enhanced photo-Fenton-like reaction, driven by increase in W^{5+} attributed to the increase of protonation. However, as shown in the mechanism section **Figure 1(e)**, the recycling process should occur during reaction which means that the WO_3 should have good recyclability to be used several times. For clarifying these phenomena, the repeatability test was also conducted to assess the recyclability of the photo-Fenton reaction. The data plotted in **Figure 3(c)** shows the good stability of oxygen evolution reaction and dye degradation after 4 recycle experiments which further elevated the potential of this concept for practical applications. This stability test confirmed the advantages of photo-Fenton like reaction by using WO_3 which shows better catalytic stability compared to traditional iron ion, TiO_2,[13] CuO,[14] and Mn_3O_4.[15]

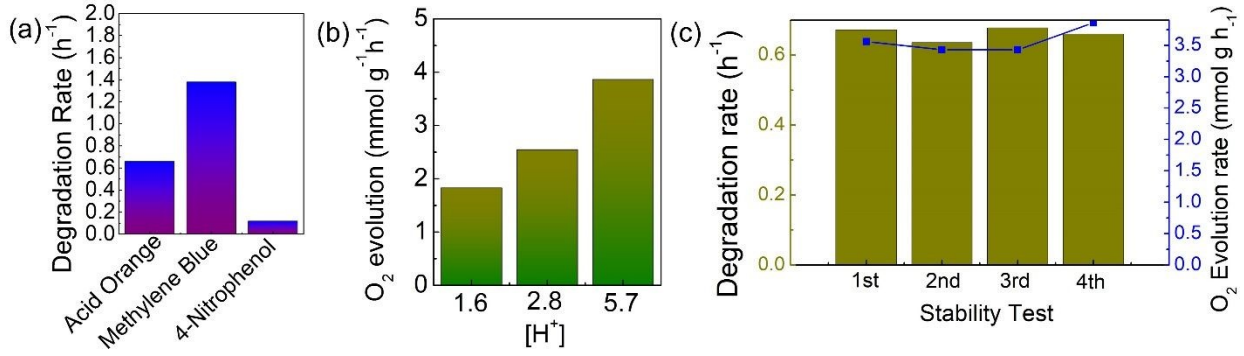

Figure 3. (a) Degradation ability of $[H^+] = 5.7$ sample for degrading other types of wastewaters. (b) Oxygen evolution rate for different protonation samples. (c) Stability of the $[H^+] = 5.7$ catalysts in terms of the reaction rate for Acid Orange 7 removal and oxygen evolution reaction.

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The enhancement of the photo-Fenton-like reaction has been clarified with relatively satisfactory results through increasing protonation. However, the origin of the increase in surface area and W^{5+} formation during UV irradiation, along with the gradual increase in protonation, needs to be clarified in detail. One major question is clarifying the crystal growth behavior under different protonation levels, which affects the ease of W^{5+} formation and surface properties. Initially, XRD patterns were analyzed to investigate the crystal structure of WO_3 at different protonation levels. As shown in **Figure 4(a)**, the diffraction patterns exhibit the characteristic peaks corresponding to monoclinic WO_3 (JCPDS No. 83-0950) with a space group of *P21/n*. Rietveld refinement performed using the Profex (open-source) software confirms that all patterns can be well-fitted with a single WO_3 phase, indicating the absence of any secondary phases. No diffraction peaks associated with H_4WO_5 were detected in any of the samples. **Supporting information (S9)** further shows that at 400 ˚C, H_4WO_5 is fully transformed into the WO_3 structure.

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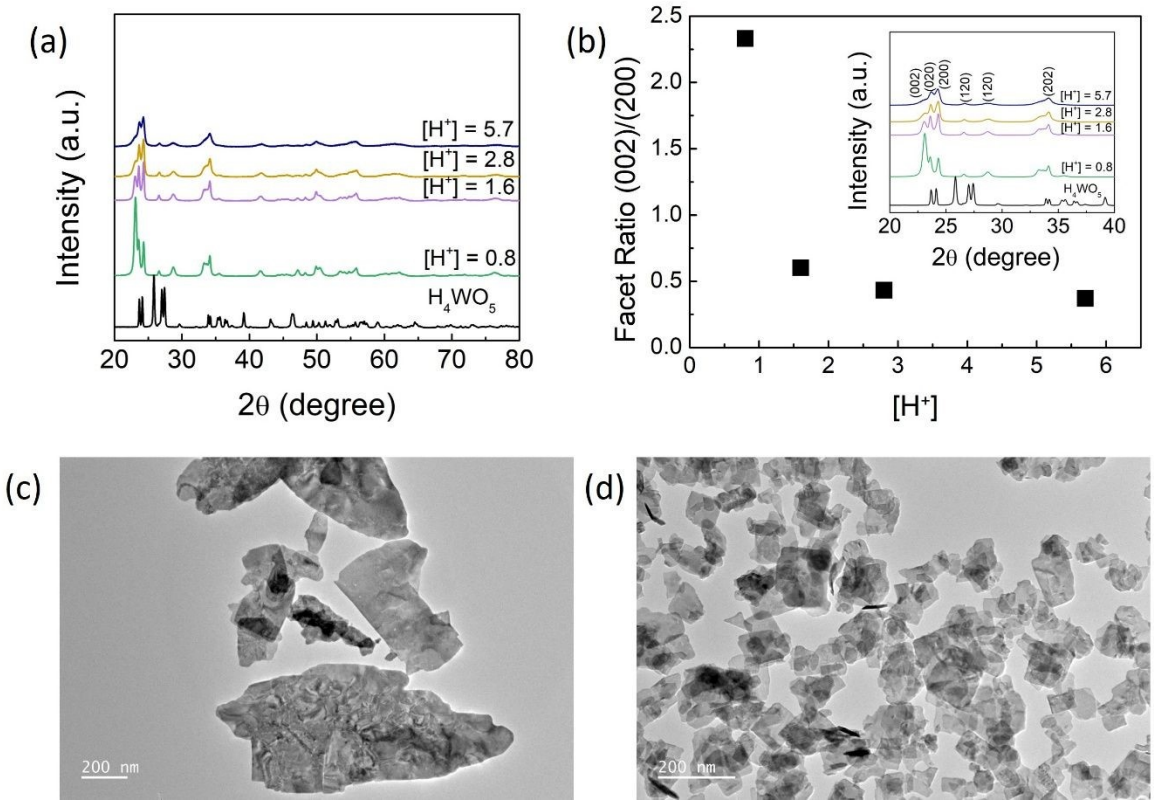

Figure 4. (a) XRD spectra of WO_3 prepared at different protonation. (b) Intensity ratio between (002) facet with (200) facet (inset shows XRD spectra of WO_3 prepared at different protonation with narrow angle ranges). TEM images of WO_3 for (c) $[H^+] = 0.8$ and (d) $[H^+] = 5.7$ samples.

Although all samples crystallize in the monoclinic WO_3 phase, the intensity of the (002) reflection decreases progressively with increasing protonation (inset of **Figure 4(b)**). The intensity ratio of the (002) to (200) planes decrease sharply from 2.33 for $[H^+] = 0.8$ to 0.33 for $[H^+] = 5.7$, indicating a significant change in the relative growth orientation associated with protonation. The (002) peak is usually correlated with the lateral dimension of the material. With the gradual decrease in the (002) peak, there is a possibility that the lateral dimension also decreases, indicating a reduction in particle size as confirmed by TEM analysis (**Figure 4(c)** and **(d))** attributed to the increase in supersaturation degree which increase the number of nuclei formations and decrease the particle growth. It is easy to understand that this smaller size correlated with surface area

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enhancement. On the other hand, to find the origin of high W^{5+} formation during UV irradiation, additional analysis is required.

The detail crystal structure analysis at different protonation levels then analyzed by examining the changes in lattice volume and lattice parameters, as shown in **Figure 5(a)**. An expansion in lattice volume was observed with increasing supersaturation process attributed to the different protonation levels during synthesis. The lattice strain then estimated by comparing the lattice structure of high crystalline WO_3 with the current samples (see **Supporting Information, (S10)**). The results show that the higher protonation, the increase in the tensile strain from 0.21 % to 0.79 %. The expansion of the lattice might be correlated with the distortion of the lattice crystals by protonation. For confirming the distortion in the crystal, Raman spectroscopy was performed to analyze the chemical bonding between tungsten and oxygen atoms (**Figure 5(b)**). Two distinct peaks were observed in the range of 500–1000 cm^{-1} at 806 cm^{-1} and 712 cm^{-1}, corresponding to the stretching vibrations of W–O–W bonds[41,42]. Notably, peak broadening at 712 cm^{-1} became more pronounced with increasing protonation, indicating increased lattice distortion. Additionally, a shoulder peak at 637 cm^{-1} emerged and intensified with higher protonation levels. This feature, absent in highly crystalline WO_3, further confirms the progressive distortion of the crystal structure with increasing protonation.

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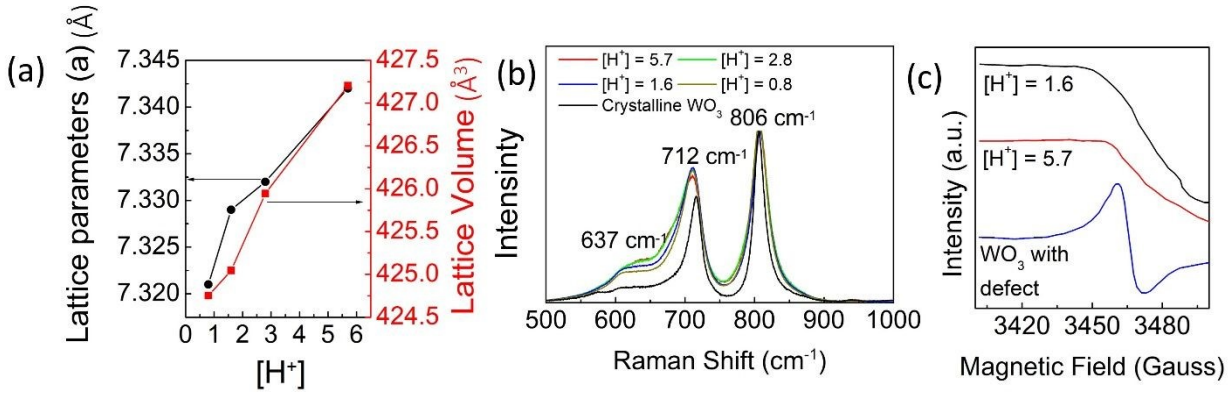

Figure 5. (a) Lattice parameters analysis of WO_3 with different protonation levels. (b) Raman spectra of WO_3 prepared at different protonation levels. (c) EPR analysis of WO_3 for $[H^+]$ = 1.6, $[H^+]$ = 5.7, and defect vacancy existed WO_3.

The origin of lattice distortion can be attributed to several factors, including strain and oxygen vacancy defects. To determine the primary cause, electron paramagnetic resonance (EPR) analysis was performed on WO_3 samples with different protonation levels to confirm the formation of lattice defects (vacancies) (**Figure 5(c)**). Additionally, defective WO_3 was provided as a comparison to confirm the presence of oxygen vacancies by annealing the $[H^+]$ = 5.7 under N_2 atmosphere for 400 ˚C for 2h. A distinct signal was observed at 3450–3480 gauss, characteristic of oxygen vacancy formation. However, no corresponding oxygen vacancy signals were detected for WO_3 samples with $[H^+]$ = 1.6 and $[H^+]$ = 5.7. Instead, distortion is likely induced by strain attributed to the rapid supersaturation process where at high supersaturation rapid nucleation occurs, which limits crystal growth and introduces structural disorder.

The strain-modulated enhancement of W^{5+} formation and its influence on the reaction process has been confirmed. However, the precise origin of the strain effect on W^{5+} formation remains unclear. Several researchers have suggested that strain can induce electron localization,

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allowing electrons to remain longer on the surface.[23,43,44] This phenomenon was further validated in the case of WO_3. As shown in the photoluminescence analysis (**Supporting information, (S11)**), the peak intensity at $[H^+] = 5.7$ is lower than that at $[H^+] = 1.6$, indicating that electrons spend more time in the conduction band, thereby promoting polaron formation. Enhanced electron localization due to strain was also confirmed by spin-polarized density functional theory calculations, which revealed an increase in total magnetization from 2.83 μB in the unstrained structure to 3.14 μB in the strained structure (**Supporting Information, (S12)**). Collectively, these results demonstrate that strained WO_3 more readily localizes electrons and promotes polaronic behavior, thereby accelerating the photo-Fenton reaction.

CONCLUSION

In this study, a novel concept has been successfully introduced for the utilization of charge carriers in WO_3 through a photo-Fenton-like reaction. Light induced on WO_3 promotes the excitation of electrons, reducing W^{6+} to W^{5+}, which interacts with H_2O_2 to produce •OH that was successfully utilized for wastewater remediation. On the other hand, the holes in the valence band efficiently conduct water splitting to produce O_2, attributed to the trapping of electrons that promotes the photo-Fenton-like reaction. The photo-Fenton-like reaction was optimized by controlling protonation during synthesis, which regulates crystal growth. Higher protonation resulted in smaller particle sizes, larger surface areas, and increased W^{5+} content, attributed to the higher crystal strain. These changes in physical and electronic properties enhance the ability of WO_3 to drive the photo-Fenton-like reaction for Acid Orange 7 degradation and the oxygen evolution reaction, owing to the synergistic effect between surface chemistry and W^{5+} formation. Overall, these findings demonstrate the efficient charge utilization of WO_3 in photo-Fenton-like processes.

View Article Online
DOI: 10.1039/D5TA07868G

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Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the Supplementary Information.

Author Contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

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- The data supporting this article have been included as part of the Supplementary Information.