Electrospun Nanofibers Embedding ZnO/Ag₂CO₃/Ag₂O Heterojunction Photocatalyst with Enhanced Photocatalytic Activity

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Abstract: The immobilization of photocatalyst onto substrate has a great potential for energy-intensive separation to avoid the costly separation process and unwanted release of photocatalyst into the treated water. In this study, electrospun nanofiber composed of polyvinylidene fluoride (PVDF) with the immobilized ZnO, ZnO/Ag₂CO₃, ZnO/Ag₂CO₃/Ag₂O, and ZnO/Ag₂O photocatalysts were prepared via the electrospinning process. The immobilized ZnO and heterojunctioned ZnO in the PVDF electrospun nanofiber were proven via X-ray diffraction (XRD) and Fourier transform infrared spectroscopy (FTIR). The electrospinning allowed high chemical binding of the nanofiber composite with good physical interaction between the photocatalyst and the electrospun nanofiber. AFM images obtained for the nanofibers were found to be rougher than that of the pristine PVDF electrospun nanofiber. Among the photocatalyst embedded, the immobilized ZnO/Ag₂CO₃/Ag₂O had endowed the nanofiber with an excellent photocatalytic activity and recyclability for the degradation of the RR120 under UV light irradiation. Based on the results, effective immobilization of ZnO/Ag₂CO₃/Ag₂O in PVDF nanofiber with 99.62% photodegradation in 300 min compared to PVDF-ZnO, PVDF-ZnO/Ag₂CO₃, and PVDF-ZnO/Ag₂O of 28.14%, 90.49%, and 96.34%, respectively. The effective ZnO/Ag₂CO₃/Ag₂O immobilization into polymers with affinity toward organic dye pollutants could both increase the efficiency and reduce the energy requirements for water treatment via the photocatalytic application.

Keywords: nanofiber; photocatalytic; nanocomposite; PVDF; ZnO

1. Introduction

In the last decades, the expanding use of synthetic dyes in many industries such as textile, food, leather, paper production, hair coloring, and agriculture has been particularly noticeable, leading to increasing concern because of their widespread presence. As such, these activities compel the continual number of wastewater discharge [1]. Despite the concentration of the dye, oftentimes, the wastewaters are colored and turbid. Dyes, as well as their breakdown products and myriad others, are mutagen and harmful to human health. Consequently, approaches to curbing issues related to these lethal organic materials are instigated, as the toxic materials are being released excessively.
The photocatalytic approach is one of the approaches that are progressively addressed due to it being environmentally friendly technique, which suggests a great potential for environmental protection and energy conversion so organic pollutants can be efficiently degraded. Throughout the process, light irradiation with an increased photon energy or equal to the band gaps of the photocatalysts can be used for generation of reactive oxidation species (ROS) by semiconductor-based photocatalysts [2,3]. In the photo-mineralization of organic pollutants, zinc oxide (ZnO) titanium dioxide (TiO$_2$), ferric oxide (Fe$_2$O$_3$), bismuth oxide (Bi$_2$O$_3$), and many others have been widely used in photocatalytic candidates, owing to its high activity due to their effective degradation efficiency of many pollutants [4–7]. Among these, ZnO and TiO$_2$ are widely used for photocatalytic degradation of many organic pollutants due to their photochemical stability, non-toxic features and high oxidation power [8,9]. Nevertheless, the use of ZnO nanomaterials as photocatalysts is impeded due to its major disadvantages, such as limited visible light application and low quantum yields. Studies have discovered that the photocatalytic activity of the photocatalysts can definitely be improved by modifying the photocatalyst materials, such as doping with transition metals [10–13] and non-metals [13–17], while constituting composite photocatalysts from different semiconductors [18,19] via photo response range extension and increased efficiency in the electron-hole pair separation [20,21]. It is believed that nearly all composite semiconductors had functioned with increasing photocatalytic activity, as compared to a single semiconductor [22,23]. In previous study on the modification of ZnO via heterojunction with mixed-phase semiconductors, Ag$_2$CO$_3$/AgO exhibited better photocatalytic properties than the pristine ZnO. The Ag$_2$CO$_3$/AgO heterostructure has found a relatively low band gap energy (about 2.3 eV), which offered the ability of absorbing a broad solar spectrum, making this novel ZnO/Ag$_2$CO$_3$/AgO photocatalyst effectively functional for photocatalytic application [24].

However, in most photocatalysis studies, the post-treatment for the removal of photocatalyst in slurry form presents several drawbacks: (a) The complicated separation or filtration phases, (b) the challenging use in continuous flow systems, and (c) the particles accumulation or loss, especially in high concentrations [20]. Thus, the process becomes complicated, incompetent and costly [25]. Henceforth, the introduction regarding to photocatalytic membrane in order to support the photocatalyst has been one of outstanding technique to overcome the drawbacks. The photocatalytic membrane basically consists of separation and degradation processes. This integration concept has allowed the photocatalytic technology to take a front role for degrading the target pollutants from complex molecules to simple and less hazardous molecules. Later, the membrane separation takes place as the second role, where it allows simple and less hazardous molecules to pass through the membrane. Undeniably, the photocatalyst can be regarded as the front role, owing to benefits of minimizing the “membrane fouling” and retaining the membrane performances throughout the process [26,27]. It was reported by Lee et al. [28] that the photocatalytic membrane is compromised of the N-TiO$_2$ membranes, creating multifunctional inorganic membranes which are capable of destroying the pollutants and self-cleaning under solar irradiation. However, most of immobilized photocatalyst suffer ineffectively within the membrane layer due to the immersion of photocatalyst in the opaque membrane layer. As a result, the photocatalyst is unable to degrade effectively, owing to the limitation of contact with light irradiation [29]. The ineffectiveness of the photocatalyst impregnated in membrane matrix is reported to be four times lower compared to photocatalyst dispersion, thus, a lowered photocatalytic performance of the membrane would be obtained [20,30].

To immobilize the photocatalyst on a suitable solid inert support with the desired morphological structure, the electrospun nanofiber has been considered as the most promising candidate due to its three-dimensional open structure, highly effective surface area, and flexibility in operation [31–33]. These electrospun nanofiber-based photocatalysts have the advantage of optional design nanostructures (such as solid, porous, hollow, core-sheath, hierarchical fibers, etc.) due to flexible nanofibrous structure with tunable diameters, which are easy to control and environmentally compatible [34]. Their applications in photocatalytic are expected in high photocatalytic property, attributed to its advantages of higher surface area/volume ratio [35], compared by incorporating the photocatalyst
nanoparticles into the membrane. In addition, the photocatalytic membrane ensures the photocatalyst recyclability, as their fibrous structure facilitates the difficulty of powder photocatalysts’ separation from reaction mixture as previously mentioned. Therefore, fabrication of a nanofiber structure would be an effective solution to the problem of catalyst recovery [36]. Kareem et al. [37] recently reported that a visible light driven CNTs/ZnO-NH₂ nanocomposite incorporated in polyacrylonitrile (PAN) nanofiber managed to degrade methylene blue and indigo carmine in a short duration of 29 and 115 min, respectively. The immobilization of GO-ZnO nanocomposite into PAN electrospun by Abdel-Mottaleb et al. [38] also exhibited an outstanding photocatalytic property toward methylene blue dye degradation under visible light irradiation. Similarly, the incorporation of Ag-AgBr@Bi₂O₂TiO₃ as core shell over PAN nanofiber electrospun for herbicide isoproturon degradation offered great photocatalytic activity and recyclability under visible light irradiation [39].

Furthermore, an ideal substrate material for photocatalyst immobilization must be stable to avoid catalyst leaching, remain steady against ROS, and act selectively toward desired contaminants [40]. Numerous materials have been studied as photocatalyst supports, such as ceramic, polymers, silica, activated carbon, and alumina [41–45]. Fluoro-polymers, for instance, polyvinylidene fluoride (PVDF), is one of promising substrates, offering high chemical and mechanical permanency attributed to durable C–F bonds [46]. Titanium dioxide embedded PVDF nanofibers are prepared for the effective photo-oxidation of methylene blue pollution [47]. Homo assembled Ag nanoparticles on PVDF nanofibers reported a boosted photocatalytic activity of methyl orange pollutant [48]. Also, high dichlorination and photodegradation of pesticide has been studied on Ag/PAN-g-PAA-Fe-Pd nanofiber hetero-architecture, which was prepared using PVDF/PAN nanofibers as supports and calcined for doping and grafting of C, N, and Fe–Pd elements [49].

In this study, ZnO heterojunction with Ag-based photocatalyst nanoparticles was embedded in hydrophobic PVDF nanofiber by electrospinning method. The hydrophobic nature of the composite nanofiber has allowed the capability of the photocatalyst to degrade the pollutant with the irradiation of light exposure to the nanoparticle. To the best of our knowledge, little or no documentation on the fabrication of heterojunctioned ZnO/Agₓ (Agₓ refers to Ag₂CO₃, Ag₂CO₃/Ag₂O, and Ag₂O)-embedded PVDF composite nanofiber for pollutant degradation is published. The pollutant degradation ability of the composite nanofiber was investigated by determining the color reduction of a model pollutant (Reactive Red 120-RR120). The phase composition, morphological structures, optical absorption, and specific surface areas were among the physicochemical properties of the prepared electrospun nanofiber that was studied. Based on the physicochemical properties and photocatalytic evaluation, the composite nanofiber in this research can be used as an effective candidate for pollutant degradation under UV light and can permit the recyclability of the photocatalytic composite.

2. Results and Discussion

2.1. Characterization of Electrospun Nanofiber

2.1.1. XRD Analysis

Figure 1 displays the X-ray diffraction (XRD) patterns, showing the crystallographic structure and phase purity of the as-fabricated samples. From Figure 1f, it is apparent that the prepared ZnO/Ag₂CO₃/Ag₂O photocatalysts displayed the coexistence of the ZnO, Ag₂CO₃, and Ag₂O phases. The major peaks at 2θ = (31.75°, 34.44°, 36.25°, 47.54°, 56.56°, 62.87°, 67.92°, and 69.06°) were assigned to the diffractions of the (100), (002), (101), (102), (110), (103), (112), and (201) characteristic peaks with hexagonal structure of ZnO, as indexed in JCPDS no. 05 0664 [50]. The other series of diffraction peaks at 18.54° and 20.54° corresponded to the (020) and (110) planes of monoclinic phase Ag₂CO₃ (JCPDS 26-0339) [51]. In addition, the diffraction peaks at 32.79° and 38.07° could be indexed to the (111) and (011) planes of cubical Ag₂O phase (JCPDS41-1104) [52]. As shown in Figure 1a, the pristine PVDF nanofiber demonstrated only one broad diffraction peak between 15° and 25°, representing its amorphous form. For the embedded photocatalyst electrospun in Figure 1b–e, the prepared
sample showed the diffraction peaks in accordance with the impregnated photocatalyst. As shown in Figure 1b, the incorporation of ZnO in PVDF nanofibers is indicated by the diffraction peaks that are perfectly indexed to ZnO. As observed in Figure 1c, the diffraction peak of ZnO/Ag$_2$CO$_3$ incorporated in the PVDF polymer nanofiber are well-indexed, indicating the presence of ZnO and Ag$_2$CO$_3$. However, there was an additional peak present at 33.67°, ascribing to the reflections of (−130) planes of the Ag$_2$CO$_3$. These inexistence of (−130) plane was due to the synthesis route of ZnO/Ag$_2$CO$_3$/Ag$_2$O (Figure 1f) that undergoes a low temperature calcination process, causing the Ag$_2$CO$_3$ to be thermally unstable. The Ag$_2$CO$_3$ phase had partially decomposed into Ag$_2$O during the calcination process. Therefore, the phase crystallinity of Ag$_2$CO$_3$ in Figure 1d,e was reduced in accordance to the phase transformation of Ag$_2$CO$_3$ into Ag$_2$O, forming heterojunction with Ag$_2$CO$_3$/Ag$_2$O mixed phase over the ZnO photocatalyst [53]. This would result in an improved electron-hole separation via the smooth charge transfer process, compared to the ZnO/Ag$_2$CO$_3$ and ZnO/Ag$_2$O. Moreover, Figure 1d demonstrated that the diffraction peaks of Ag$_2$CO$_3$, Ag$_2$O, and ZnO are well-incorporated into the PVDF electrospun nanofiber matrix. As observed in Figure 1e, ZnO and Ag$_2$O phases were clearly observed. The invisible Ag$_2$CO$_3$ phase in Figure 1e showed that the Ag$_2$CO$_3$ has completed transformed into the Ag$_2$O phase.

![Figure 1. X-ray diffraction (XRD) spectra of (a) P0: PVDF; (b) P1: PVDF-ZnO; (c) P2: PVDF-ZnO/Ag$_2$CO$_3$; (d) P3: PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O; (e) P4: PVDF-ZnO/Ag$_2$O; and (f) ZnO/Ag$_2$CO$_3$/Ag$_2$O photocatalyst.](image)

2.1.2. Morphological Structure Analysis

The morphological structure and corresponding histograms of the electrospun nanofiber diameter distribution exhibited tremendous difference, compared to the pristine PVDF electrospun nanofiber (refer to Figure 2). It can be clearly seen that the diameter of the prepared electrospun nanofibers were in the range of 500—800 nm. As shown in Figure 2a, the surface of the pristine PVDF electrospun nanofibers is relatively smooth and uniform in diameter, indicating that the parameters used in the electrospinning process were appropriate. Figure 2b–e revealed that the immobilized photocatalyst resulted in the rough surface of the nanofibers. It displayed the existence of ZnO nanorods structure interposed inside the produced nanofiber structures. The results also indicated that the diameter of the nanofibers increased, as the ZnO and ZnO heterojunctioned photocatalyst was embedded. This is due to the effect of the dope solution viscosity. The dope viscosity could be altered by increasing the polymer concentration or by inclusion of insoluble filler (ZnO or ZnO heterojunctioned) [40,54]. Overall, all of the as-spun nanofibers generated the construction of an open three-dimensional (3D)
network, which offers a high surface area and flexible surface functionalities that can promote good contact with the organic pollutants for photocatalytic activity.

The distribution of the photocatalyst on the electrospun fibers in PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O nanofiber was further studied by TEM images (Figure 3). The granule and dark shade on the TEM images indicated the presence of ZnO/Ag$_2$CO$_3$/Ag$_2$O particles, while the light shade bound the photocatalyst particles is the PVDF fiber. From Figure 3, the ZnO/Ag$_2$CO$_3$/Ag$_2$O particles were found homogenously distributed in PVDF nanofiber. The TEM image displayed a curved layer at the edges of ZnO/Ag$_2$CO$_3$/Ag$_2$O particles, revealing that the photocatalyst was enclosed with a layer of PVDF polymer. The selected area electron diffraction (SAED) pattern (the inset in the lower left) of nanofiber composite displayed multiple bright spots rings, indicating the crystalline structure of ZnO/Ag$_2$CO$_3$/Ag$_2$O nanocomposite was persevered after incorporated in PVDF fiber.

The surface morphologies of the electrospun nanofibers were further evaluated by AFM, as shown in Figure 4a-2–e-2. It is revealed that the surface roughness of the PVDF electrospun nanofiber impregnated with photocatalysts was considerably higher than that of the pristine PVDF electrospun nanofiber. Based on the contact angle result, the PVDF-ZnO nanofiber displayed the highest value of 118.75° in Figure 4b-1. The incorporation of ZnO nanorod particles interposed inside PVDF nanofibers led to the hydrophobilization of the nanofibers, which induced the rougher surfaces [40], as presented in Figure 4b-2. In fact, the AFM results of PVDF-ZnO displayed the roughest average surfaces toward others. Thus, this situation led to a higher physical repulsion toward water droplets that generate hydrophobic properties [55–57]. However, the hydrophobicity of the PVDF nanofiber decreased as the ZnO heterojunction with Ag-based photocatalyst was introduced. The AFM result depicts the value of surface roughness that decreased from 0.70 to 0.35 μm. A similar trend found that the wettability behavior of the fabricated electrospun nanofiber showed a reduced contact angle, as shown in Figure 4c-1–e-1. This is due to the presence of beads-on-string morphology, reducing the surface roughness of the nanofiber, hence developing electrospun nanofiber with lower physical repulsion toward the water droplets, thus having lower hydrophobilization.

Figure 2. SEM images and the corresponding histograms of the electrospun nanofiber diameter distribution (a) P0: PVDF; (b) P1: PVDF-ZnO; (c) P2: PVDF-ZnO/Ag$_2$CO$_3$; (d) P3: PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O; (e) P4: PVDF-ZnO/Ag$_2$O.
The IUPAC classification can be used to analyze the types of nitrogen adsorption-desorption isotherms [58]. As shown in Figure 5, the shapes of nitrogen adsorption-desorption isotherms of samples P0 were depicted as Type III. Meanwhile, P1, P2, P3, and P4 demonstrated a Type IV curve with a H3-type hysteresis loop and with mesoporous materials characterization [59]. Other than that, the abrupt increment in the N₂ uptake at high relative pressures (P/P₀ > 0.9) described the presence of macropores [60]. The outcome established that the embedded heterojunction photocatalyst displayed the hierarchical macro- and mesoporous structure. Figure 5a–e shows the corresponding BJH pore size distribution curves. It was found that the pore size of the pristine PVDF nanofiber fell less than 200 nm, while the pore size of the PVDF nanofiber incorporated with photocatalyst increased up to 500 nm. This indicated that the mesopores and macropores were observed in the PVDF-ZnO composite nanofibers. The enhanced BET surface area and hierarchical pore architecture of the PVDF-ZnO composite nanofibers have facilitated the surface absorption of reactants and mass transfer of degradation products.
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ZnO composite nanofibers have facilitated the surface absorption of reactants and mass transfer of products. This might be due to the strong interaction between the ZnO, Ag2O, or ZnO/Ag2O and PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O. This might be due to the strong interaction between the ZnO, Ag2O, or ZnO/Ag2O and PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O. This might be due to the strong interaction between the ZnO, Ag2O, or ZnO/Ag2O and PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O. This might be due to the strong interaction between the ZnO, Ag2O, or ZnO/Ag2O and PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O. This might be due to the strong interaction between the ZnO, Ag2O, or ZnO/Ag2O and PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O.

Figure 6 illustrates the FTIR spectra of PVDF, PVDF-ZnO, PVDF-ZnO/Ag2CO3, PVDF-ZnO/Ag2CO3/Ag2O, and PVDF-ZnO/Ag2O nanofibers. The results indicated that the PVDF spectra follows similar trends, as reported in a previous study [61]. The main peaks of ZnO, Ag2CO3, and Ag2O appeared, suggesting that the structural integrity of ZnO/Ag2CO3, ZnO/Ag2O, or ZnO/Ag2CO3/Ag2O persist, abounding even when incorporated in PVDF nanofiber. It could be distinguished that a broad band, located in the range of 3000 to 3700, was ascribed to the stretching vibration of –OH group in ZnO [62]. Furthermore, as compared to PVDF-ZnO, it was observed that O-H stretching absorption peaks shifted from the high to low absorption peaks in PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O. This might be due to the strong interaction between the hydroxyl group of ZnO particles either with hydrogen bond interaction of CO3^2- or Ag-O bonds in

Table 1. The surface area and pore properties of the electrospun nanofibers.

<table>
<thead>
<tr>
<th>Samples</th>
<th>BET Surface Area (m^2g^-1)</th>
<th>Average Pore Size (nm)</th>
<th>Pore Volume (x10^-2 cm^3g^-1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>P0</td>
<td>3.75</td>
<td>23.97</td>
<td>2.94</td>
</tr>
<tr>
<td>P1</td>
<td>4.12</td>
<td>22.47</td>
<td>2.31</td>
</tr>
<tr>
<td>P2</td>
<td>6.61</td>
<td>13.86</td>
<td>2.29</td>
</tr>
<tr>
<td>P3</td>
<td>6.50</td>
<td>11.26</td>
<td>1.83</td>
</tr>
<tr>
<td>P4</td>
<td>6.22</td>
<td>18.91</td>
<td>2.80</td>
</tr>
</tbody>
</table>

2.1.4. FTIR Analysis

The incorporation of ZnO in PVDF nanofiber can be further proven by Fourier transform infrared (FTIR) spectroscopy analysis. Figure 6 illustrates the FTIR spectra of PVDF, PVDF-ZnO, PVDF-ZnO/Ag2CO3, PVDF-ZnO/Ag2CO3/Ag2O, and PVDF-ZnO/Ag2O nanofibers. The results indicated that the PVDF spectra follows similar trends, as reported in a previous study [61]. The main peaks of ZnO, Ag2CO3, and Ag2O appeared, suggesting that the structural integrity of ZnO/Ag2CO3, ZnO/Ag2O, or ZnO/Ag2CO3/Ag2O persist, abounding even when incorporated in PVDF nanofiber. It could be distinguished that a broad band, located in the range of 3000 to 3700, was ascribed to the stretching vibration of –OH group in ZnO [62]. Furthermore, as compared to PVDF-ZnO, it was observed that O-H stretching absorption peaks shifted from the high to low absorption peaks in PVDF-ZnO/Ag2CO3 and PVDF-ZnO/Ag2O. This might be due to the strong interaction between the hydroxyl group of ZnO particles either with hydrogen bond interaction of CO3^2- or Ag-O bonds in
Ag$_2$CO$_3$ and Ag$_2$O, respectively [63]. Besides, all of the embedded heterojunctioned photocatalysts in nanofiber show characteristic features of CO$_3^{2-}$– and Ag-O stretching vibration modes with absorption bands around 1630–1650 cm$^{-1}$ [64,65]. Meanwhile, the broad peaks from 1500 to 700 cm$^{-1}$ prove that the mixture of Ag$_2$CO$_3$ and Ag$_2$O was formed in the PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O nanofiber [66]. Overall, the main absorption peaks of Ag$_2$CO$_3$, Ag$_2$O and ZnO were all observed, indicating a strong interaction of photocatalyst impregnated in PVDF nanofiber.

![Figure 6. Fourier transform infrared (FTIR) spectra of: (a) P0: PVDF; (b) P1: PVDF-ZnO, (c) P2: PVDF-ZnO/Ag$_2$CO$_3$; (d) P3: PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O; (e) P4: PVDF-ZnO/Ag$_2$O.](image)

2.1.5. Optical Properties

Optical properties were determined by characterizing the response of electrospun nanofibers towards electromagnetic radiant energy, including visible and ultraviolet (UV) light. The optical properties of all of the prepared nanofibers were analyzed using UV–Vis spectroscopy, as shown in Figure 7a. It was observed that the pristine PVDF electrospun fiber mats showed very weak absorption across the wavelength, hence the band gap energy of pristine PVDF could not be presented. Meanwhile, all the embedded heterojunctioned photocatalysts in the PVDF nanofibers demonstrated an excellent optical response toward UV and visible light irradiation.

In comparison to the PVDF-ZnO nanofiber, the absorption edge shifted toward visible light range in the UV–Vis light spectrum, thus revealing the capability of the ZnO heterojunctioned photocatalyst in the PVDF electrospun nanofiber performed in visible light irradiation. A similar observation by Kočí et al., [67] found the heterojunctioned photocatalyst shifted the absorption edge toward higher wavelength, that is, toward the visible region. These shifted absorption edges signified that the heterojunctioned photocatalysts could efficiently extend absorption of nanocomposites to the visible region, hence increasing the photocatalytic capability in the visible light region.

The influence of electronic properties of all of the prepared samples toward the photocatalytic activity was determined by measuring the energy gaps between the valence and conduction band.
The band gap energy of the prepared electrospun nanofiber was estimated using Tauc plot relation by the following equation:

\[
(ah\nu) = A(h\nu - E_g)^{n/2}
\]

where \(\alpha\), \(A\), \(E_g\), \(h\nu\), and \(n\) represent the absorption coefficient, constant, band gap, photon energy, and an integer, respectively, while the value of integer \(n\) depends on the characteristic of optical transition (\(n = 1\) or \(4\) for direct or indirect band transition, respectively). The band gap energies of the electrospun nanofibers are illustrated in Figure 7b. The \((ah\nu)^2\) versus the energy absorbed light was plotted, where both zinc and silver are indirect transition semiconductors [68]. According to the result, the obtained band gap values of PVDF-ZnO, PVDF-ZnO/Ag_2CO_3, PVDF-ZnO/Ag_2O, and PVDF-ZnO/Ag_2CO_3/Ag_2O were found to be about 3.2, 3.13, 3.08, and 2.95 eV, respectively. Hence, these results proved that PVDF electrospun nanofibers embedded with photocatalysts were capable of undergoing the photocatalytic reaction.

![Figure 7. (a) Absorbance spectra, (b) band gap determination by drawing the line at \((ah\nu)^2\) of the electrospun nanofibers.](image)

2.2. Evaluation of Photocatalytic Activity

Photocatalytic activities of the prepared electrospun nanofibers were tested out for the degradation of organic pollutant under UV light irradiation by using the reactive red 12 (RR120) as a model pollutant. The change in optical absorption spectra of RR120 by the electrospun nanofiber under UV light irradiation as a function of time is shown in Figure 8a. It was found that the adsorption of RR120 had increased, as the photocatalyst was incorporated into the PVDF nanofibers. After 90 min of adsorption, the PVDF-ZnO/Ag_2CO_3 showed the highest RR120 adsorption at 6.76%, compared to PVDF-ZnO, PVDF-ZnO/Ag_2CO_3/Ag_2O, and PVDF-ZnO/Ag_2O at 4.62%, 4.33%, and 5.19%, respectively. The difference in RR120 adsorption is due to the difference in their surface properties (e.g., surface area and surface roughness) [69,70]. Based on the analysis, BET surface area and surface roughness of PVDF embedded with ZnO heterojunctioned photocatalyst are higher compared to the pristine PVDF and PVDF-ZnO nanofiber. The higher BET surface area and surface roughness would contribute to higher percentage of adsorption, hence increasing the chances for UV to reach and come into contact with the photocatalyst through its highly porous interconnected nanofiber media. The photolysis of the RR120 solution without the addition of electrospun nanofiber was carried out as a controlled experiment. As shown in Figure 8a, there was a slight prominent change for the RR120 solution after UV light irradiation for 300 min, in which the RR120 can be categorized as having good photostability and potential resistance under the UV light irradiation [71]. The photocatalytic activity using pristine PVDF nanofiber without the presence of photocatalyst was also performed for comparison purposes.
As portrayed in Figure 8a, the pristine PVDF nanofiber exhibited an approximately 10.43% adsorption of RR120 after 300 min of UV light irradiation.

After the addition of ZnO, the photodegradation of RR120 was found to have increased up to 28.14%. Meanwhile, in the impregnation of ZnO/Ag₂CO₃ heterojunctioned into the PVDF nanofiber, the photocatalytic activity of the samples increased, as compared to the PVDF-ZnO nanofiber. The highest photocatalytic activity was attained by PVDF-ZnO/Ag₂CO₃/Ag₂O with 99.62% degradation after 300 min of reaction, followed by PVDF-ZnO/Ag₂O and PVDF-ZnO/Ag₂CO₃ of 96.34% and 90.49%, respectively. The lowest photocatalytic activity was attained for PVDF-ZnO/Ag₂O. This is possibly due to the smaller band gap possessed by the Ag₂O. Hence, the potential recombination of electron-hole pair could exist, which would indirectly lower the production of hydroxyl radical, as well as photocatalytic reaction. The change in the absorption spectrum of RR120 with PVDF-ZnO/Ag₂CO₃/Ag₂O electrospun at different time intervals is presented in Figure 8b. As the reaction time increased, all the peaks decreased gradually, and the spectrum scanning model distorted apparently after 300 min. The magnitude of the absorption peak at 533 nm slowly declined, indicating the reduction of RR120 concentration. According to Panakoulias et al. [72], the visible band at 533 nm owed to the RR120 nitro groups breakdown (discoloration of the solution) (Figure 8b inset), while it progressed with the cleavage of the aromatic ring at absorption bands 265 and 330 nm. The elimination of absorbance at 330 and 265 nm caused it to continue producing aliphatic intermediates that would be mineralized to carbon dioxide. A similar observation has also been made by Balachandran et al. [73], where a similar pattern of the absorption peak reduction without the appearance of new absorption peaks during the degradation process revealed the intermediates do not absorb at the analytical wavelengths of 285 and 533 nm.

The Langmuir-Hinshelwood model was extensively applied for the analysis of photocatalytic degradation kinetics of pollutant in aqueous phase, as follows [74]:

\[
\ln \left( \frac{C_i}{C_0} \right) = -kt
\]

where, \(C_0\) and \(C_t\) refer to the concentrations of pollutant (mgL\(^{-1}\)) at the respective irradiation time \(0\) and \(t\) min, while \(k\) is the apparent rate (min\(^{-1}\)). The pseudo-first-order kinetic model refers to the photocatalytic activity of RR120 in aqueous solution by the prepared electrospun nanofiber, in which Figure 8c shows the equal value of rate constant \(k\) to the corresponding slope of the fitting line. Figure 8d exhibits the reaction rate constant of all of the prepared nanofibers.

The pristine PVDF electrospun nanofiber showed that the relatively low reaction rate was obtained for nanofiber without photocatalyst. The main reason for the reduction of RR120 concentration is mainly due to the adsorption process. However, when ZnO, ZnO/Ag₂CO₃, ZnO/Ag₂CO₃/Ag₂O, and ZnO/Ag₂O were embedded in the PVDF electrospun nanofiber, a higher reduction of RR120 concentration was observed. This is because two processes were involved: The adsorption and photocatalysis processes. The photocatalyst provided the surface that was responsible for the photocatalysis process. Overall, PVDF nanofiber incorporated with the heterojunctioned ZnO composite photocatalyst significantly increased the photocatalytic activity. The PVDF-ZnO/Ag₂CO₃/Ag₂O exhibited the highest reaction rate constant of \(118 \times 10^{-4}\) min\(^{-1}\), which was about 1.26, 1.42, and 5.57 times higher than those of PVDF-ZnO/Ag₂O, PVDF-ZnO/Ag₂CO₃, and PVDF-ZnO, respectively.

The competency of the as-spun photocatalytic nanofibers is essentially accomplished through the synergetic combination of a highly porous structure and a high surface area generated by the nanofiber microstructure, which accurately aids in the adsorption migration-photodegradation process [75]. The RR120 is likely to be adsorbed by the PVDF nanofibers because of the lower hydrophobization took place via the PVDF-ZnO nanofiber while shifting to the photocatalyst nanoparticles, and finally degraded under the UV irradiation [47].

Figure 8e shows the recyclability test, demonstrating that the PVDF-ZnO/Ag₂CO₃/Ag₂O nanofiber indicated a comparable degradation performance for the five consecutive cycles. It was suggested that the PVDF-ZnO/Ag₂CO₃/Ag₂O was relatively stable after multiple cycles, even though there was a slight decline in the photodegradation rate for RR120. This proposes a robust physical interlocking of
photocatalyst in the PVDF nanofiber matrix [76]. In fact, no leaching of photocatalyst was observed during the experiment. The remarkable recyclability of PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O electrospun nanofiber proved their potential for practical applications in a large scale.

Based on the above results, a proposed mechanism was discussed toward the development of the photocatalytic properties of the PVDF-ZnO/Ag$_2$CO$_3$/Ag$_2$O composite nanofiber. In the present work, fibers with an average diameter of 645 ± 0.48 nm loaded with ZnO/Ag$_2$CO$_3$/Ag$_2$O photocatalyst nanoparticles create groups of higher fiber diameter (Figure 2), hence offering a much higher specific surface area with a larger interaction of the dye and the inorganic filler, suggesting a faster degradation mechanism. However, due to the hydrophobic nature of PVDF polymer (Figure 4), the degradation of pink dye was only possible due to the interaction of the ZnO/Ag$_2$CO$_3$/Ag$_2$O photocatalyst with the dye. In this sense, the enhanced photocatalytic degradation is primarily due to the synergetic combination of porous structure and higher surface area formed by the fiber microstructure, which competently allowed the adsorption migration-photodegradation process as aforementioned during degradation.
of RR120. In this way, the RR120 is possibly adsorbed by the polymer fibers, then migrated to the ZnO/Ag_2CO_3/Ag_2O nanoparticles, and finally degraded by ZnO/Ag_2CO_3/Ag_2O under UV radiation.

On the mechanism of ZnO/Ag_2CO_3/Ag_2O surfaces, the heterojunctioned of Ag_2CO_3/Ag_2O mixed phase were believed to exhibit cooperative effects between Ag_2CO_3, Ag_2O, and ZnO. Under UV irradiation, the ZnO, Ag_2CO_3, and Ag_2O absorbed the photon energy produced the excite electron in from VB to CB, leaving the positive holes at VB. Since the position of the Ag_2CO_3 CB is less negative than ZnO and Ag_2O, the photogenerated electron produced by ZnO and Ag_2O should be transferred to Ag_2CO_3, while the photogenerated positive holes were left in VB migrated to Ag_2O. The positive holes accumulated in the surface of Ag_2O were eventually trapped by surface hydroxyl groups (or H_2O) to harvest OH∙ radicals. Dissolved oxygen molecules react to the electrons (e⁻) at Ag_2CO_3 surfaces to produce superoxide anion radicals, while O_2^− yields the hydroperoxyl, HO_2−radicals, and directly generated hydroxyl radical OH. These super oxide radicals, O_2^− and OH, were very energetic radicals to phenyl or naphthyl rings of the aromatic compounds containing in RR120 [77,78]. Thus, the mechanism for the photocatalytic degradation of RR120 in the experiment was proposed as follows:

\[
\text{ZnO/Ag}_2\text{CO}_3/\text{Ag}_2\text{O} + h\nu \rightarrow e^- (\text{Ag}_2\text{CO}_3) + h^+ (\text{Ag}_2\text{O})
\]

\[
e^- + \text{O}_2 \rightarrow \text{O}_2^-
\]

\[
h^+ + \text{OH}^- \rightarrow \text{OH}
\]

\[
\text{O}_2^- + \text{H}_2\text{O} \rightarrow \text{HO}_2 + \text{OH}^-
\]

\[
\text{HO}_2 + \text{H}_2\text{O} \rightarrow \text{H}_2\text{O}_2 + \text{OH}
\]

\[
\text{H}_2\text{O}_2 \rightarrow 2\text{OH}
\]

\[
\text{OH} + \text{RR120} \rightarrow \text{CO}_2 + \text{H}_2\text{O}
\]

The overall separation and transfer of photogenerated electron-holes pairs in ZnO/Ag_2CO_3/Ag_2O across the PVDF nanofiber and photodegradation of RR120 under UV light irradiation is schematically presented in Figure 9.

![Figure 9](image-url)  

**Figure 9.** Schematic diagram of the band edge alignment of the ZnO/Ag_2CO_3/Ag_2O system along with the proposed mechanism of charge transfer.

3. Materials and Methods

3.1. Material

The zinc acetate dihydrate (C_4H_8O_4Zn·2H_2O) supplied from GmbH company (Haan, Germany) was used as a zinc precursor. Silver nitrate (AgNO_3) and sodium bicarbonate (NaHCO_3) were purchased from Sigma Aldrich (St. Louis, MI, USA) and Bendosen (Bendosen, Norway), respectively.
Polyvinylidene fluoride (PVDF, MW = 7400) and N, N-dimethylacetamide (DMAc) was produced from Merck (Kenilworth, NJ, USA), while acetone (>99.6% purity, HPLC grade) was obtained from QReC (Chonburi, Thailand), and the Reactive Red 120 (RR120) was purchased from Sigma Aldrich (St. Louis, MI, USA). All of the chemicals and reagents were analytical reagent grade and no purification step was performed.

### 3.2. Preparation of the Electrospun Nanofiber

The photocatalyst was synthesized according to our previous study [24]. First, 1.95 g of PVDF pellets was added into 13 mL of DMAc solution prepared with acetone at the mass ratio of 2.3 and stirred at 60 °C until the polymer was dissolved. After that, 0.65 g of ZnO was added into the dope solution and stirred continuously at room temperature until a homogenous solution was obtained. The dope solution was then used in the fabrication of electrospun nanofiber via electrospinning, as shown in Figure 10. Seven milliliters of the dope solution was placed in a 10 mL syringe, equipped with a blunt metal needle. The electrospun was at a spinning rate of 1 mL/h, with a voltage of 8 kV applied between the spinneret and rotating drum placed 150 mm apart. Meanwhile, the collector of the rotating drum was set to 170 rpm. The flat sheet of the fabricated nanofibers was then allowed to dry at room temperature for the hydrolysis step before it was further characterized and tested. The same procedures were repeated for different photocatalysts of ZnO/Ag₂CO₃, ZnO/Ag₂CO₃/Ag₂O, and ZnO/Ag₂O. The resultant samples were denoted, as depicted in Table 2.

![Figure 10. Schematic diagram of the electrospinning process.](image)

#### Table 2. Fabricated electrospun nanofiber.

<table>
<thead>
<tr>
<th>Electrospun Nanofiber</th>
<th>Sample</th>
</tr>
</thead>
<tbody>
<tr>
<td>PVDF</td>
<td>P0</td>
</tr>
<tr>
<td>PVDF-ZnO</td>
<td>P1</td>
</tr>
<tr>
<td>PVDF-ZnO/Ag₂CO₃</td>
<td>P2</td>
</tr>
<tr>
<td>PVDF-ZnO/Ag₂CO₃/Ag₂O</td>
<td>P3</td>
</tr>
<tr>
<td>PVDF-ZnO/Ag₂O</td>
<td>P4</td>
</tr>
</tbody>
</table>

### 3.3. Characterization

The phase compositions of the prepared electrospun nanofiber derived from PVDF embedded with heterojunctioned ZnO were analyzed using the Rint 2200 Ultima-III X-ray Diffractometer (Rigaku, Corp., Tokyo, Japan). The morphological structure of the electrospun nanofibers was observed by scanning electron microscope (SEM, TM-1000, HITACHI, Tokyo, Japan) operated at 5 kV. The surface roughness of the electrospun nanofiber was further measured using the atomic force microscopy (AFM, Park XE 100, Schaefer Technologie GmbH, Filderstadt, Germany). Nitrogen adsorption/desorption measurement was carried out at a low temperature (77.38 K) and a relative pressure (0 to 1) to determine the BET surface area and pore size distribution (using BET adsorption analyzer, (Rigaku, Corp., Tokyo, Japan)). The optical properties of the electrospun nanofibers were analyzed by UV -Vis-NIR spectrophotometer (UV-3600 Plus, SHIMADZU, MI, USA). The optical contact angle machine (Filderstadt, Germany) was used to observe the wettabilities of the electrospun nanofibers. 

The surface roughness of the electrospun nanofiber was further measured using the atomic force microscopy (AFM, Park XE 100). The optical properties of the electrospun nanofibers were analyzed by UV –Vis-NIR spectrophotometer (UV-3600 Plus, SHIMADZU, MI, USA). The optical contact angle machine (Filderstadt, Germany) was used to observe the wettabilities of the electrospun nanofibers.
roughness of the electrospun nanofiber was further measured using the atomic force microscopy (AFM, PARK XE-100, Schaefer Technologie GmbH, Langen, Germany). The surface wettabilities of the electrospun nanofiber were determined by contact angle measurements with an optical contact angle machine (OCA 40 Micro, Dataphysics, Filderstadt, Germany). Nitrogen adsorption/desorption measurement was carried out at a low temperature (77.38 K) and a relative pressure range of \( P/P_0 = 0.02 \) to 1 to determine the BET surface area and pore size distribution (using Barrett Joyner Halenda (BJH) equation). Prior to the measurement, the electrospun nanofibers were degassed at 373 K for 3 h in the degas port of the adsorption analyzer. Fourier transform infrared spectroscopy (FTIR: Spectrum 100, PerkinElmer, Waltham, MA, USA) was used to observe the presence of photocatalyst in the PVDF electrospun nanofiber matrix. The optical properties of the electrospun nanofibers were analyzed by UV–Vis-NIR spectrophotometer (UV-3600 Plus Shidmadzu, Japan) in the range of 200 to 800 nm. The collected spectra were converted into Kubelka-Munk function to measure the band gap of the prepared PVDF-ZnO/Ag₂X electrospun nanofiber.

### 3.4. Photocatalytic Activity Measurement

The photocatalytic activity of the prepared electrospun nanofiber containing PVDF embedded with the heterojunctioned ZnO was performed under UV light irradiation using RR120 as a model pollutant. A 30 W lamp (312 nm) was used as a UV light source. The prepared electrospun nanofiber were cut into spherical shape (diameter of 8 cm, weight of 0.1 g) and immersed in 100 mL of RR120 solution. The photocatalytic experiment was first evaluated in the dark to observe the adsorption/desorption equilibrium of RR120 on the prepared nanofibers. During the photocatalytic tests, fixed amounts of the reacted solution were taken out at given time intervals. The final concentration of the RR120 solution was measured by collecting its absorbance at a wavelength of 533 nm with a UV–Vis spectrophotometer (Lambda 25, Perkin Elmer, Waltham, MA, USA). The degradation percentages of RR120 in the aqueous solution was calculated as follows: Degradation (\%) = \((C_0 - C_t)/C_0\) 100%; where \(C_0\) is the initial concentration at time \(t = 0\), and \(C_t\) is the concentration time interval. The used electrospun nanofiber was collected and dried overnight at 60 °C in an oven. For the recyclability study, all of the electrospun nanofibers were reused under similar photocatalytic conditions for five cycles. The experiment was conducted at room temperature and in a neutral pH.

### 4. Conclusions

In summary, a series of PVDF electrospun nanofiber embedded with ZnO heterojunction photocatalysts with different Ag-based compounds were successfully fabricated via a facile electrospinning process. The higher dye removal efficiency of the composite electrospun nanofiber was mainly due to a novel synergistic effect of adsorption and photocatalytic degradation in the presence of heterojunctioned catalyst, which was ZnO/Ag₂CO₃, ZnO/Ag₂CO₃/Ag₂O, and ZnO/Ag₂O. The heterojunctioned catalyst embedded into the PVDF electrospun possessed lower hydrophobization and higher surface roughness, which facilitated the dye molecule adsorption adjacent to their photocatalytic surface sites. The embedding of ZnO/Ag₂CO₃/Ag₂O into PVDF nanofiber led to a fast degradation of dye molecule among the electrospun nanocomposites under UV light irradiation, which was mainly attributed to the heterojunction between the ZnO and Ag₂CO₃/Ag₂O mixed phase, narrowing the band gap. Hence, this effectively facilitated the charge transfer and suppressed the recombination of photogenerated electrons and holes for high photodegradation activity. Furthermore, the PVDF-ZnO/Ag₂CO₃/Ag₂O electrospun nanofiber had maintained a relatively high photocatalytic activity after five cycles, which made them a promising candidate for a wide range of photocatalytic applications. However, to move toward commercialization, drawbacks of lacking nanoscale selectivity, mechanical weakness, and low wettability must be achieved.

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