



Article

# PMMA-TiO<sub>2</sub> Fibers for the Photocatalytic Degradation of Water Pollutants

Namrata Kanth <sup>1</sup>, Weiheng Xu <sup>2</sup>, Umesh Prasad <sup>2</sup>, Dharnedar Ravichandran <sup>2</sup>,  
Arunachala Mada Kannan <sup>3</sup> and Kenan Song <sup>3,\*</sup>

<sup>1</sup> Materials Science & Engineering, School for Engineering of Matter, Transport and Energy (SEMTE), Ira A. Fulton Schools of Engineering, Arizona State University, Tempe, AZ 85281, USA; nkanth@asu.edu

<sup>2</sup> System Engineering, The Polytechnic School (TPS), Ira A. Fulton Schools of Engineering, Arizona State University, Mesa, AZ 85212, USA; weihengx@asu.edu (W.X.); uprasad1@asu.edu (U.P.); dravich2@asu.edu (D.R.)

<sup>3</sup> The Polytechnic School, School for Engineering of Matter, Transport, and Energy (SEMTE), Ira A. Fulton Schools of Engineering, Arizona State University, Mesa, AZ 85212, USA; amk@asu.edu

\* Correspondence: kenan.song@asu.edu; Tel.: +1-(480)-727-2720

Received: 9 June 2020; Accepted: 21 June 2020; Published: 30 June 2020



**Abstract:** Titanium dioxide (TiO<sub>2</sub>) is a promising photocatalyst that possesses a redox potential suitable for environmental remediation applications. A low photocatalytic yield and high cost have thus far limited the commercial adoption of TiO<sub>2</sub>-based fixed-bed reactors. One solution is to engineer the physical geometry or chemical composition of the substrate to overcome these limitations. In this work, porous polymethyl methacrylate (PMMA) substrates with immobilized TiO<sub>2</sub> nanoparticles in fiber forms were fabricated and analyzed to demonstrate the influence of contaminant transport and light accessibility on the overall photocatalytic performance. The influences of (i) fiber porosity and (ii) fiber architecture on the overall photocatalytic performance were investigated. The porous structure was fabricated using wet phase inversion. The core-shell-structured fibers exhibited much higher mechanical properties than the porous fibers (7.52 GPa vs. non-testability) and maintained the same degradation rates as porous structures (0.059 vs. 0.053/min) in removing methylene blue with comparable specific surface areas. The highest methylene blue (MB) degradation rate ( $k_{MB}$ ) of 0.116 min<sup>-1</sup> was observed due to increases of the exposed surface area, pointing to more efficient photocatalysis by optimizing core-shell dimensions. This research provides an easy-to-manufacture and cost-efficient method for producing PMMA/TiO<sub>2</sub> core-shell fibers with a broad application in water treatment, air purification, and volatile sensors.

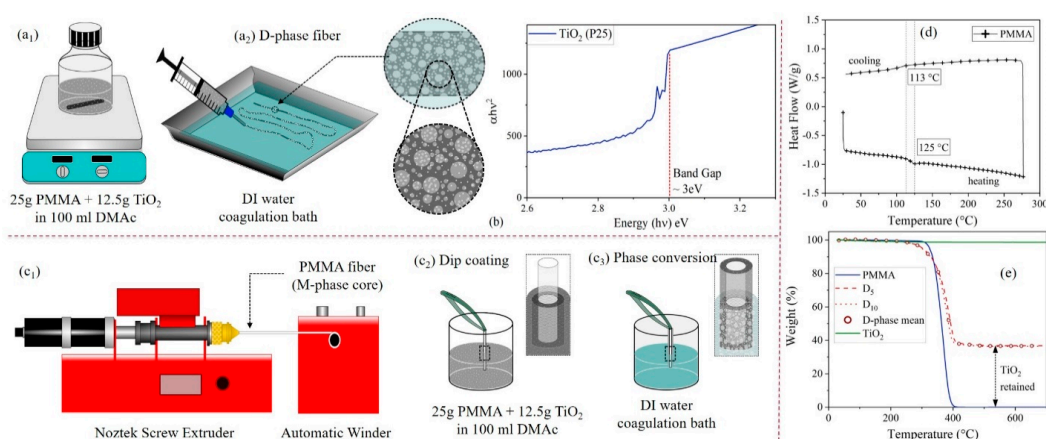
**Keywords:** nanocomposite; fibers; TiO<sub>2</sub>; PMMA; photocatalytic

## 1. Introduction

Ever-increasing pollution calls for the development of a robust and cheap solution for environmental remediation [1]. Among the numerous methods that can be employed to tackle this problem [2], harnessing the photocatalytic activity of titanium dioxide (TiO<sub>2</sub>) can offer a promising solution. TiO<sub>2</sub> is a wideband semiconductor material that possesses a redox potential capable of decomposing organic contaminants into carbon dioxide and water [3]. This advanced oxidation capability not only makes it suitable for purification applications, such as pollutant capture [4], the treatment of industrial waste [5], and wastewater disinfection [6], but means that it can also be used for the reformation of hydrocarbon fuels [7]. The commercial adaptation of TiO<sub>2</sub> has thus far been limited by a low throughput and the high cost of photoreactors [8,9]. These limitations can be addressed by a fixed-bed reactor where nano-scale TiO<sub>2</sub> is immobilized on inert substrates.

Nanoparticles of  $\text{TiO}_2$  provide a high surface to volume ratio, improving the transport of contaminants. Additionally, immobilization eliminates the need for expensive post-treatment separation of  $\text{TiO}_2$  [10]. However, this system suffers from drawbacks, such as a poor light distribution and hindered charge transfer, due to the encapsulation of  $\text{TiO}_2$  in substrates [11]. A cost-effective solution is to use polymers as a substrate, since they can be easily molded to the desired shape, size, and microstructure [12]. Past research has reported anchoring  $\text{TiO}_2$  on polymeric beads [11], membranes [13], and electrospun fibers [14]. Incorporating porosity in these substrates enhances mass transport. Another approach is to coat radially emitting optical fibers with  $\text{TiO}_2$  [15,16]. This unique architecture serves two purposes: It creates a greater surface area by utilizing the fiber's longitudinal morphology and enables remote light accessibility through optical fibers [17]. However, to the best of our knowledge, previous studies have not compared the photocatalytic merits of improving mass transfer and enhancing light accessibility, which is discussed here.

In this paper, we report the fabrication and characterization of polymethyl methacrylate (PMMA)- $\text{TiO}_2$  fibers, manufactured by low-cost and scalable methods (Figure 1a–c). In this investigation, two fiber architectures were studied [18]. The first was a porous PMMA fiber containing dispersed  $\text{TiO}_2$  (D-phase), where the porous morphology entailed enhanced mass transport (Figure 1c<sub>1,2</sub>). The second was a multiphase fiber (M-phase) with a solid PMMA core and a porous PMMA- $\text{TiO}_2$  sheath (Figure 1c<sub>3</sub>). The solid PMMA core in the M-phase serves as an optical fiber for remote light transmission, and the porous sheath provides enhanced mass transport similar to the D-phase. PMMA was chosen as a substrate material because of its excellent ultraviolet (UV) transparency [19] and stability against strong oxidative radicals [20]. Supporting materials, such as glass fibers, have been traditionally used for immobilizing  $\text{TiO}_2$ ; however, the rigidity and labor-intensive manufacturing of glass fibers mean that they are not optimal materials in water treatment applications [9,12,21–23]. The solid PMMA core was fabricated using melt spinning and the porous matrix (D-phase and M-phase sheath) was fabricated by wet phase inversion. Their photocatalytic performance was assessed by comparing the methylene blue (MB) concentrations in the dark and in the presence of UV (~365 nm) radiation. Inferences were drawn related to the effect of architecture/morphology on light transport and mass transport, which are critical parameters in reactor scale-up [9].



**Figure 1.** Schematic of the fabrication procedures for (a<sub>1</sub>,a<sub>2</sub>) preparing D-phase containing the (b)  $\text{TiO}_2$  with a bandgap of 3 eV (Tauc plot) and (c<sub>1</sub>) PMMA fiber extrusion, (c<sub>2</sub>) dip coating of extruded PMMA fiber and (c<sub>3</sub>) phase conversion for M-phase fibers. (d) Differential scanning calorimetry (DSC) of polymethyl methacrylate (PMMA) raw material. (e) Back-to-back thermogravimetric analysis (TGA) conducted to quantify  $\text{TiO}_2$  retention after wet phase inversion.

## 2. Materials and Methods

### 2.1. Chemicals

PMMA ( $M_w \sim 120,000$ ), *N,N*-dimethylacetamide (DMAc, ACS Reagent 99%), tetrahydrofuran (THF) and MB ( $\geq 82\%$  purity) were purchased from Sigma Aldrich (St. Louis, MO, USA). TiO<sub>2</sub> (AEROXIDE® TiO<sub>2</sub> P 25,  $\sim 53 \text{ m}^2/\text{g}$ ) was obtained from Evonik Industries (Essen, Germany). All polymers and solvents were used as received. Deionized (DI) water from a Millipore water purification system (Milli-Q Academic) (Sigma Aldrich, St. Louis, MO, USA) was used.

### 2.2. Characterization

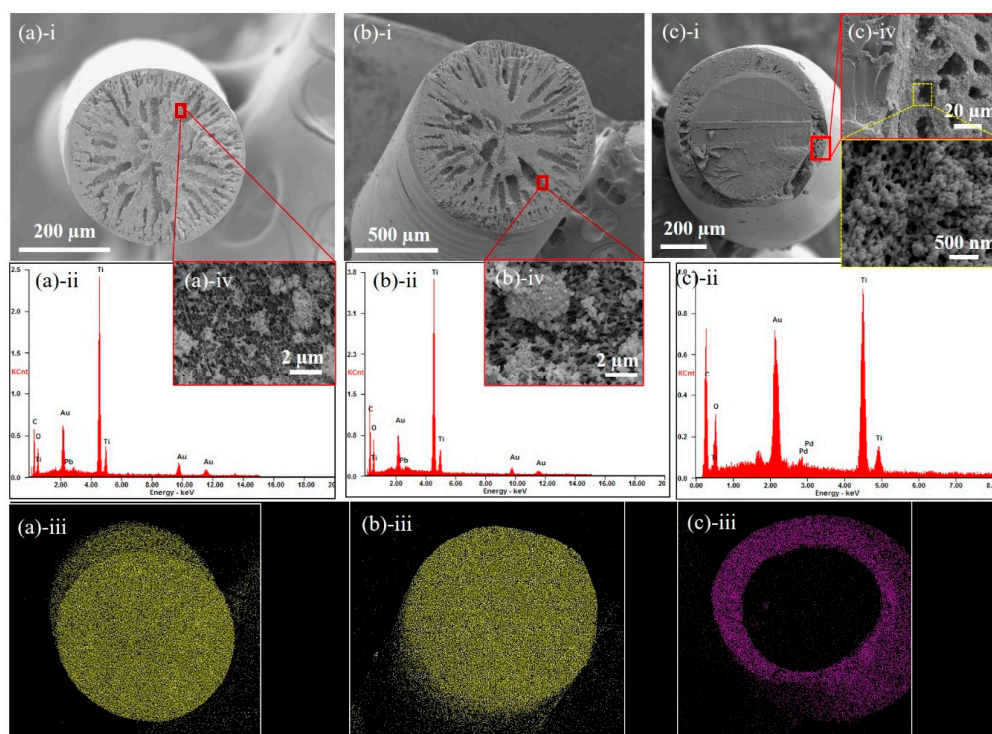
The morphology of the fibers was observed using a Scanning Electron Microscope (SEM) (XL30 SEM-FEG) (Philips, Amsterdam, The Netherlands). For SEM, all samples were sputter-coated with 15 nm thick gold. The dispersion of TiO<sub>2</sub> across the cross-section of the fibers was observed using Electron Dispersive Spectroscopy (EDS) (XL30 SEM-FEG) (Philips, Amsterdam, The Netherlands). The specific surface area (SSA) values were measured and obtained using Brunauer, Emmett, and Teller (BET, Tristar II 3020) (Micromeritics, Norcross, GA, USA) Surface Area and Porosity Analysis. The optical properties were measured using UV-vis spectrophotometry (UV-vis, GENESYS 150) (Thermo Fisher Scientific, Walham, MA, USA). Thermogravimetric (TGA) studies were conducted using TGA550 (TA instruments, New Castle, DE, USA) in an air atmosphere at a ramping rate of 20 °C/min from room temperature (RT) to 700 °C. Differential Scanning Calorimetry (DSC) was conducted in a nitrogen environment with a purging rate of 50 mL/min using DSC250 (TA instruments, New Castle, DE, USA). The mechanical property was characterized by conducting a tensile test using Discovery Hybrid Rheometer (HR-2) (TA instruments, New Castle, DE, USA) for ten fiber samples. For assessing the photocatalytic performance, MB concentration was monitored in the dark and in the presence of UV radiation. MB concentration was measured as per Beer-Lambert Law, by monitoring the absorption peak at 665 nm (UV-vis). The UV radiation source was 365 nm cross-linking oven (DYMEX ECE 5000) (Dymax, Torrington, CT, USA).

## 3. Results and Discussion

### 3.1. Fiber Morphology

The fiber morphology influences both the mechanical durability and photocatalytic behaviors. During our fiber-making process, the choices of solvent (i.e., DMAc) and nonsolvent (i.e., DI water) were critical in forming the desirable microstructures (Figure 1a). PMMA is a suitable substrate material, since it is  $\sim 80\%$  transparent to radiation with a wavelength of 250 nm or above. However, the introduction of pores in the substrate hampers its optical properties. A porous substrate demonstrates poor light transmittance through its thickness since it scatters most of the light falling on the surface (Figure S1). Our TiO<sub>2</sub> sample had a bandgap of 3 eV (or  $\lambda = 413 \text{ nm}$ , Figure 1b), which means that the substrate should have been transparent to radiation with a wavelength of less than or equal to 413 nm. TiO<sub>2</sub> only anchored to the surface of the porous fibers that had access to light essential for photocatalysis. This corroborates the need for lean fiber-like architectures whose three-dimensional (3D) structure provides a greater avenue for light impingement. The overall structure of the D-phase was porous, with uniformly dispersed TiO<sub>2</sub> (Figure 2a,b). PMMA filaments extruded at a temperature of 275 °C (i.e., higher than the glass transition from Differential Scanning Calorimetry in Figure 1d) had 0.5 and 1.0 mm diameters, respectively (i.e., D<sub>5</sub> and D<sub>10</sub>), based on the extruder sizes. The D<sub>5</sub> fiber had a  $\sim$ two times higher specific surface area (SSA) compared to D<sub>10</sub> (Table 1). This was because of the different sizes of the fiber samples, which influenced the coagulation dynamics [24]. The solvent leaching process was more effective for a smaller fiber diameter (D<sub>5</sub>) due to the shorter diffusion scale for the coagulant, resulting in a more porous structure. A consequence of porosity was mechanical fragility, as can be seen in the D-phase fibers (Table 1). As a comparison, the M-phase fibers with a

densely-packed core region surrounded by a porous sheath (Figure 2c) and uniform TiO<sub>2</sub> dispersions exhibited a high mechanical robustness (Table 1). Note that, as shown in Figure 2a-iii to Figure 2c-iii, the TiO<sub>2</sub> was homogeneously dispersed in both the dispersed and core-shell fibers, which is critical for studying the microstructural effects on their water treatment efficiency. The TiO<sub>2</sub> concentration in both the D-phase and M-phase surface was 36 wt.% (Figure 1e). This concentration of TiO<sub>2</sub> was selected due to the fact that higher TiO<sub>2</sub> concentrations in DMAc will cause particle aggregation and fast sedimentation that will influence the coating quality on PMMA fibers. A densely packed core was desirable, since it implies that the core has no visible pores or voids, which might hamper light transmission.



**Figure 2.** Morphological and compositional characterization of (a) D5, (b) D10, and (c) M-phase fibers with (a-i to c-i) the cross-sectional surface topology, (a-ii to c-ii) EDS of the surface showing the element identification, (a-iii to c-iii) distribution of TiO<sub>2</sub>, and (a-iv to c-iv) zoom-in regions in (a-ii to c-i) showing the dispersion of TiO<sub>2</sub>.

**Table 1.** Physical properties of the as-fabricated fibers.

Fiber Morphologies	Sample	Dimensions ( <i>d</i> , Core Diameter; <i>t</i> , Sheath Thickness; mm)	Mechanical Properties §		BET Specific Surface Area (m <sup>2</sup> /g)	Langmuir–Hinshelwood Pseudo-First-Order Kinetics Parameters	
			Young's Modulus (GPa)	Tensile Strength (MPa)		Initial Concentration C <sub>0</sub> , mg/L	Degradation Rate <i>k</i> <sub>MB</sub> , min <sup>-1</sup>
D-phase	D <sub>5</sub>	<i>d</i> = 0.5	Fragile		33.2	0.706	0.116
	D <sub>10</sub>	<i>d</i> = 1.0			17.4	2.466	0.053
M-phase core-shell	M <sub>C</sub> S	<i>d</i> = 0.5, <i>t</i> = 0.1	7.52 ± 2.73	117.7 ± 43.2	15.7	2.545	0.059
	M <sub>C</sub> #1					1.364	0.0077
	M <sub>C</sub> #2					0.581	0.0069

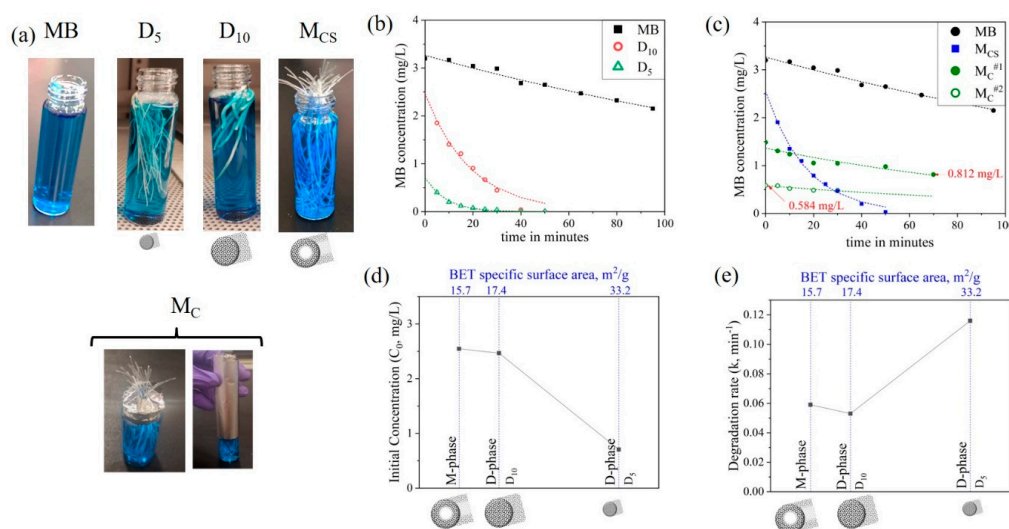
§ M-phase core-shell and M-phase core showed comparable mechanical properties. Note that due to the porous feature of the M-phase core-shell, it is challenging to precisely estimate the cross-section area. See BET data in Figure S2. #1,2 are core-shell fibers used for the second-cycle measurement of photocatalysis.



Methylene blue is frequently used to dye office supplies and color papers, as well as to tone up silk colors. In veterinary and human medicine, MB has been largely used in several therapeutic and diagnostic procedures. Natural degradation through conventional water treatment procedures is challenging because of MB's resistance to light, temperature, water, and chemicals. The long-term existence of MB in nature can not only pollute the water system, but also pose dangers to human health. Photocatalytic oxidation is considered to be one of the most effective ways of degrading MB.

Four sets of fiber samples ( $D_5$ ,  $D_{10}$ ,  $M_C$ , and  $M_{CS}$ ), each containing  $\sim 50$  mg of  $TiO_2$ , were placed in individual glass vials containing 8 mL MB solution (3.2 mg/L) (Figure 3a). The glass vials were kept in the dark (i.e., inside a closed opaque box to protect them from light exposure) for five days to achieve the adsorption-desorption equilibrium of MB. Then, the samples were exposed to UV radiation and the MB dye concentrations were monitored over time (Figure 3b,c). The photocatalytic reaction kinetics was analyzed using the Langmuir–Hinshelwood pseudo-first-order kinetics model [25].

$$\ln\left(\frac{C}{C_0}\right) = -k_{MB}t \quad (1)$$



**Figure 3.** (a) Samples used for photocatalytic testing. Photocatalytic removal of methylene blue (MB) in the presence of UV light in the (b) D-phase, and (c) M-phase fibers. Degradation parameters (d)  $C_0$  and (e)  $k$  estimated using regression fit as a function of the BET specific surface area (SSA).

Here,  $C_0$  and  $C$  represent the dye concentration after the five-day soaking period and during UV exposure, respectively.  $k_{MB}$  represents the degradation rate and time,  $t$ , was recorded from the instant when the sample was exposed to UV radiation. The parameters  $C_0$  and  $k$  were estimated by regression fit (Table 1). The experiments were categorized into two sets. First, to analyze the overall photocatalytic performance of D-phase ( $D_5$  and  $D_{10}$ ) and M-phase ( $M_{CS}$ ). Second, to quantify the effectiveness of the fiber core in M-phase fiber ( $M_C$ ). The sample was covered in aluminum foil to only allow light transmission through the core (Figure 3). An MB solution without any fibers was used as a control test.

### 3.2. Overall Photocatalytic Performance

The MB concentrations ( $c$ ) as a function of time ( $t$ ) for  $D_5$ ,  $D_{10}$ , and  $M_{CS}$  fibers exposed in MB solutions are shown in Figure 3b,c. The y-intercept depicts the initial MB concentration  $C_0$ , which is characteristic of the adsorption capability of the fiber.  $D_5$  fibers demonstrated the lowest  $C_0$ , corresponding to the highest porosity (Table 1) for contaminant adsorption. On the contrary,  $D_{10}$  and  $M_{CS}$  had a comparable BET SSA, resulting in comparable  $C_0$  (Figure 3d). The decrease in MB concentrations over time ( $y$ -axis in Figure 3b,c) was a result of the photo-inactivation of adsorbed

contaminants. UV radiation decomposed the adsorbed dye, which facilitated the adsorption of fresh MB from the solution into the replenished sites in the fiber. Therefore, the degradation rate (column  $k_{MB}$ , Table 1) was a measure of the substrate's morphological ability to deliver UV light to the immobilized  $TiO_2$ . As discussed earlier, the good dispersion in both the dispersed and core-shell fibers (Figure 2a-iii to Figure 2c-iii) excluded the influence of  $TiO_2$  agglomerations, and thus, the differences in the degradation rate and initial concentrations were the results of the fiber microstructures. Amongst all of the samples, D<sub>5</sub> demonstrated the highest degradation rate of  $0.116 \text{ min}^{-1}$ . This was attributed to the lean architecture of D<sub>5</sub>, which ensured better light delivery. M<sub>C5</sub> and D<sub>10</sub> exhibited comparable degradation rates, in spite of M<sub>C5</sub> being comprised of a fiber core, which entailed better light accessibility. It should be noted that our test of the degradation of water pollutants was based on a simple setup with UV light radiating toward a bundle of randomly oriented fibers with both cross-section and surfaces exposed. This setup eases the practical application; however, precise UV transition toward exposed fiber ends may reveal the role of the core in pollutant degradation, especially for M<sub>C5</sub> fibers when the core is expected to transmit UV-light.

### 3.3. Effectiveness of the M-Phase Core

M-phase fibers with only fiber cross-section areas exposed in MB solutions were used to check the UV-transmittance efficiency of the core (M<sub>C</sub><sup>#</sup>). For M<sub>C</sub><sup>#1</sup>, the MB concentration reduced during UV exposure, indicating that the fiber core was successful in delivering light in the M-phase architecture (Figure 3c); however, this degradation rate, compared to that facilitated by the surface pores, was one order of magnitude smaller (Table 1). Then, the same sample was left in a dark room for another day and retested in the presence of UV (represented by M<sub>C</sub><sup>#2</sup>). MB concentrations after the first cycle of exposure (M<sub>C</sub><sup>#1</sup>) and prior to the second cycle of exposure (M<sub>C</sub><sup>#2</sup>) differed by 0.226 mg/L, implying continued dye adsorption. During M<sub>C</sub><sup>#2</sup>, the MB concentration reduced again and eventually saturated in ~50 min. Dye decomposition during M<sub>C</sub><sup>#1</sup> and M<sub>C</sub><sup>#2</sup> indicated that UV exposure facilitated dye adsorption by purging the  $TiO_2$  sites from the previously adsorbed dye, producing similar degradation rates (i.e., 0.0077 vs. 0.0069, shown in Table 1). The saturation of M<sub>C</sub><sup>#1</sup> and M<sub>C</sub><sup>#2</sup> curves indicates the asymptotic nature of adsorption, which slows down due to a reduction of concentration gradients. Therefore, the pace of adsorption limits the photocatalytic performance of the fibers.

## 4. Conclusions

In this study, we developed novel porous PMMA- $TiO_2$  fibers using wet phase inversion and analyzed their photocatalytic performance. Two fixed-bed architectures were explored, namely, a dispersed phase PMMA- $TiO_2$  fiber (D-phase) and multiphase fiber (M-phase) with a solid PMMA core and dispersed PMMA- $TiO_2$  sheath. D-phase exhibited the highest degradation rate of  $0.116 \text{ min}^{-1}$  due to its high porosity of  $33.2 \text{ m}^2/\text{g}$ . Size-controlled D-phase and M-phase samples with a comparable porosity displayed similar performance characteristics, despite operating on different mechanisms of light exposure. This means that the overall rate was limited by adsorption of the support structure, even though the overall photocatalytic process required concurrent adsorption and photoactivation. From the perspective of product development, internally illuminated M-phase fibers will result in a compact reactor design, since the fibers can be bundled together, whereas the light distribution in externally illuminated D-phase becomes a limiting factor. The influences of a porous morphology and solid core on light transport characteristics were also explored in this study. To further improve the performance of porous fixed-bed reactors, the dimensions and adsorption sites of the fiber must be optimized to balance both mass transport and light accessibility.

**Supplementary Materials:** The following are available online at <http://www.mdpi.com/2079-4991/10/7/1279/s1>, Figure S1. Schematic sketch of light interaction in the (a) S-polymethyl methacrylate (PMMA)<sub>film</sub>, (b) P-PMMA<sub>film</sub>, and (c) Ti-PMMA<sub>film</sub>. Optical properties characterized using UV-vis (d) absorbance (e) transmittance and (f) reflectance. Figure S2. (a) N<sub>2</sub> adsorption-desorption curve and (b) Barrett-Joyner-Halenda (BJH) pore size distribution for D<sub>5</sub>. Figure S3. (a) N<sub>2</sub> adsorption-desorption curve and (b) BJH pore size distribution for D10. Figure S4. (a) N<sub>2</sub> adsorption-desorption curve and (b) BJH pore size distribution for M-phase.

**Author Contributions:** Fabrication, photocatalytic study experiments, and manuscript writing, N.K.; BET, EDS, and preliminary fiber fabrication, W.X.; UV-vis experiments, U.P.; SEM imaging, D.R.; photocatalytic studies, A.M.K.; design of experiments and advisor, K.S. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was funded by Master's Opportunity for Research in Engineering, provided by Arizona State University (ASU).

**Acknowledgments:** The authors would like to acknowledge Akshay Phadnis for stimulating discussions on and suggestions for the manuscript.

**Conflicts of Interest:** The authors declare no conflict of interest.

## References

1. Zhang, Y.; Jiang, Z.; Huang, J.; Lim, L.Y.; Li, W.; Deng, J.; Gong, D.; Tang, Y.; Lai, Y.; Chen, Z. Titanate and titania nanostructured materials for environmental and energy applications: A review. *RSC Adv.* **2015**, *5*, 79479–79510. [[CrossRef](#)]
2. Khin, M.M.; Nair, A.S.; Babu, V.J.; Murugan, R.; Ramakrishna, S. A review on nanomaterials for environmental remediation. *Energy Environ. Sci.* **2012**, *5*, 8075–8109. [[CrossRef](#)]
3. Fujishima, A.; Rao, T.N.; Tryk, D.A. Titanium dioxide photocatalysis. *J. Photochem. Photobiol. A* **2000**, *1*, 1–21. [[CrossRef](#)]
4. Nonami, T.; Hase, H.; Funakoshi, K. Apatite-coated titanium dioxide photocatalyst for air purification. *Catal. Today* **2004**, *96*, 113–118. [[CrossRef](#)]
5. Wong, C.L.; Tan, Y.N.; Mohamed, A.R. A review on the formation of titania nanotube photocatalysts by hydrothermal treatment. *J. Environ. Manage.* **2011**, *92*, 1669–1680. [[CrossRef](#)]
6. Shannon, M.A.; Bohn, P.W.; Elimelech, M.; Georgiadis, J.G.; Marias, B.J.; Mayes, A.M. Science and technology for water purification in the coming decades. *Nature* **2008**, *452*, 301–310. [[CrossRef](#)]
7. Tu, W.; Zhou, Y.; Liu, Q.; Tian, Z.; Gao, J.; Chen, X.; Zhang, H.; Liu, J.; Zou, Z. Robust hollow spheres consisting of alternating titania nanosheets and graphene nanosheets with high photocatalytic activity for CO<sub>2</sub> conversion into renewable fuels. *Adv. Funct. Mater.* **2012**, *22*, 1215–1221. [[CrossRef](#)]
8. Linsebigler, A.L.; Lu, G.; Yates, J.T. Photocatalysis on TiO<sub>2</sub> surfaces: Principles, mechanisms, and selected results. *Chem. Rev.* **1995**, *95*, 735–758. [[CrossRef](#)]
9. Ola, O.; Maroto-Valer, M.M. Review of material design and reactor engineering on TiO<sub>2</sub> photocatalysis for CO<sub>2</sub> reduction. *J. Photochem. Photobiol. C* **2015**, *24*, 16–42. [[CrossRef](#)]
10. Singh, S.; Mahalingam, H.; Singh, P.K. Polymer-supported titanium dioxide photocatalysts for environmental remediation: A review. *Appl. Catal. A* **2013**, *462*, 178–195. [[CrossRef](#)]
11. Chong, M.N.; Jin, B.; Chow, C.W.K.; Saint, C. Recent developments in photocatalytic water treatment technology: A review. *Water Res.* **2010**, *44*, 2997–3027. [[CrossRef](#)] [[PubMed](#)]
12. Shan, A.Y.; Ghazi, T.I.M.; Rashid, S.A. Immobilisation of titanium dioxide onto supporting materials in heterogeneous photocatalysis: A review. *Appl. Catal. A* **2010**, *389*, 1–8. [[CrossRef](#)]
13. Ng, L.Y.; Mohammad, A.W.; Leo, C.P.; Hilal, N. Polymeric membranes incorporated with metal/metal oxide nanoparticles: A comprehensive review. *Desalination* **2013**, *308*, 15–33. [[CrossRef](#)]
14. Lee, C.G.; Javed, H.; Zhang, D.; Kim, J.H.; Westerhoff, P.; Li, Q.; Alvarez, P.J.J. Porous electrospun fibers embedding TiO<sub>2</sub> for adsorption and photocatalytic degradation of water pollutants. *Environ. Sci. Technol.* **2018**, *52*, 4285–4293. [[CrossRef](#)] [[PubMed](#)]
15. Peill, N.J.; Hoffmann, M.R. Development and optimization of a TiO<sub>2</sub>-coated fiber-optic cable reactor: Photocatalytic degradation of 4-chlorophenol. *Environ. Sci. Technol.* **1995**, *29*, 2974–2981. [[CrossRef](#)] [[PubMed](#)]
16. Kim, S.; Kim, M.; Lim, S.K.; Park, Y. Titania-coated plastic optical fiber fabrics for remote photocatalytic degradation of aqueous pollutants. *J. Environ. Chem. Eng.* **2017**, *5*, 1899–1905. [[CrossRef](#)]
17. Neal, H.O.; Garcia-segura, S.; Hristovski, K.; Westerhoff, P. Compact light-emitting diode optical fiber immobilized TiO<sub>2</sub> reactor for photocatalytic water treatment. *Sci. Total Environ.* **2018**, *613*, 1331–1338. [[CrossRef](#)]

18. Xu, W.; Jambhulkar, S.; Verma, R.; Franklin, R.; Ravichandran, D.; Song, K. In situ alignment of graphene nanoplatelets in poly(vinyl alcohol) nanocomposite fibers with controlled stepwise interfacial exfoliation. *Nanoscale Adv.* **2019**, *1*, 2510–2517. [[CrossRef](#)]
19. Raja, V.; Sarma, A.K.; Rao, V.V.R.N. Optical properties of pure and doped PMMA-CO-P4VPNO polymer films. *Mater. Lett.* **2003**, *57*, 4678–4683. [[CrossRef](#)]
20. Camara, R.M.; Portela, R.; Gutierrez-Martin, M.; Sánchez, B. Evaluation of several commercial polymers as support for TiO<sub>2</sub> in photocatalytic applications. *Glob. NEST J.* **2014**, *16*, 525–535.
21. Hofstadler, K.; Bauer, R.; Novallc, S.; Heisler, G. New reactor design for photocatalytic wastewater treatment with TiO<sub>2</sub> immobilized on fused-silica glass fibers: Photomineralization of 4-chlorophenol. *Environ. Sci. Technol.* **1994**, *28*, 670–674. [[CrossRef](#)] [[PubMed](#)]
22. Leong, S.; Razmjou, A.; Wang, K.; Hapgood, K.; Zhang, X.; Wang, H. TiO<sub>2</sub> based photocatalytic membranes: A review. *J. Memb. Sci.* **2014**, *472*, 167–184. [[CrossRef](#)]
23. Chen, F.; Gong, A.S.; Zhu, M.; Chen, G.; Lacey, S.D.; Jiang, F.; Li, Y.; Wang, Y.; Dai, J.; Yao, Y.; et al. Mesoporous, three-dimensional wood membrane decorated with nanoparticles for highly efficient water treatment. *ACS Nano* **2017**, *11*, 4275–4282. [[CrossRef](#)] [[PubMed](#)]
24. McKelvey, S.; Koros, W.J. Phase separation, vitrification, and the manifestation of macrovoids in polymeric asymmetric membranes. *J. Memb. Sci.* **2003**, *112*, 29–39. [[CrossRef](#)]
25. Vild, A.; Teixeira, S.; Kühn, K.; Cuniberti, G.; Sencadas, V. Orthogonal experimental design of titanium dioxide—Poly(methyl methacrylate) electrospun nanocomposite membranes for photocatalytic applications. *J. Environ. Chem. Eng.* **2016**, *4*, 3151–3158. [[CrossRef](#)]



© 2020 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<http://creativecommons.org/licenses/by/4.0/>).