



Article A Study on TiO₂ Surface Texturing Effect for the Enhancement of Photocatalytic Reaction in a Total Phosphorous Concentration Measurement System

Jae Keon Kim ^{1,2,†}, Seung Deok Kim ^{3,†}, Jae Yong Lee ³, Chang Hee Kim ¹, Hyeon-Su Lee ², Seong Mo Koo ^{2,3}, YoungJin Lee ⁴, Jong-Hoo Paik ⁴, Da Ye Kim ³ and Seong Ho Kong ^{1,2,3,*}

- ¹ Department of Sensor and Display Engineering, Kyungpook National University, Daegu 41566, Korea; kjg@medisentech.com (J.K.K.); changhee@knu.ac.kr (C.H.K.)
- ² Medisentech, Inc., Techno-Building B206, 80 Daehakro, Bukgu, Daegu 41566, Korea; leehs56@medisentech.com (H.-S.L.); smkoo@medisentech.com (S.M.K.)
- ³ School of Electronic and Electrical Engineering, Kyungpook National University, Daegu 41566, Korea; ksd5683@gmail.com (S.D.K.); cheerssss@naver.com (J.Y.L.); dykim0827@gmail.com (D.Y.K.)
- ⁴ Electronic Convergence Division, Korea Institute of Ceramic Engineering & Technology, Jinju-si 52851, Korea; yjlee@kicet.re.kr (Y.L.); jhpaik@kicet.re.kr (J.-H.P.)
- Correspondence: shkong@knu.ac.kr; Tel.: +82-53-950-8679
- Authors contributed equally to this work.

Abstract: Powerful sunlight, a high water temperature, and stagnation in the water flow induce eutrophication in rivers and lakes, which destroys the aquatic ecosystem and threatens the downstream water supply systems. Accordingly, it is very important to perform real-time measurements of nutrients that induce algal growth, especially total phosphorus, to preserve and manage the aquatic ecosystem. To conduct quantitative analysis of the total phosphorus in the aquatic ecosystem, it is essential to perform a pretreatment process and quickly separate the phosphorus, combined with organic and inorganic materials, into a phosphate. In this study, the sandblasting process was used for the physical etching of the wafer, and photocatalytic materials were deposited on the surface with various roughness in order to improve the photocatalytic reaction surface and efficiency. The photocatalytic reaction was applied to combine the pretreated sample with the coloring agent for color development, and the absorbance of the colored sample was analyzed quantitatively to compare and evaluate the characteristics, followed by the surface increase in the photocatalytic materials. In addition, the pretreatment and measurement parts were materialized in a single chip to produce a small and light total phosphorus analysis sensor.

Keywords: total phosphorus; surface texturing; sandblast; photocatalysis

1. Introduction

Various human activities, such as the discharge of agricultural water and factory wastewater and urbanization, lead to excessive quantities of phosphorus in the aquatic ecosystem and cause eutrophication in the water [1–3]. Eutrophication in water quality leads to the over-breeding of algae, and, as a result, this consumes a large amount of oxygen and rapidly reduces the amount of oxygen available for the survival of fish, crustaceans, and various aquatic organisms [4–7]. Accordingly, early detection and response to eutrophication are priorities for water quality management in rivers and water supply sources. Furthermore, phosphorus, one of the causes of eutrophication, is also an indicator of water pollution [8–10]. Therefore, research is being actively conducted worldwide to develop a small total phosphorus monitoring system with real-time measurement to prevent eutrophication in advance [11].



Citation: Kim, J.K.; Kim, S.D.; Lee, J.Y.; Kim, C.H.; Lee, H.-S.; Koo, S.M.; Lee, Y.; Paik, J.-H.; Kim, D.Y.; Kong, S.H. A Study on TiO₂ Surface Texturing Effect for the Enhancement of Photocatalytic Reaction in a Total Phosphorous Concentration Measurement System. *Micromachines* **2021**, *12*, 1163. https:// doi.org/10.3390/mi12101163

Academic Editor: Moongyu Jang

Received: 13 August 2021 Accepted: 16 September 2021 Published: 28 September 2021

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2021 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 (https:// creativecommons.org/licenses/by/ 4.0/). To implement a high-efficiency total phosphorus monitoring system, it is essential to accurately and promptly conduct a pretreatment process in which the phosphorus in the water is separated into phosphate (PO_4^{3-}).

The existing total phosphorus analysis procedure involves performing the pretreatment of the sample under high-temperature (>120 °C) and high-pressure (>1.1 kg/cm⁻²) conditions in order to achieve the oxidative decomposition of the phosphorus. In addition, there are disadvantages, such as the bulkiness of the equipment, since it consists of a thermal oxidation device, mixing device, and detection device, as well as the high cost and long time required for analysis.

Meanwhile, the pretreatment method of decomposing the phosphorus into phosphate (PO_4^{3-}) using a photocatalyst is a technology that utilizes the chemical reaction that results from the light under high-temperature and high-pressure conditions, so there are advantages such as being economical and also easy to handle and safe. In particular, among the photocatalytic materials, TiO₂ does not become decomposed by light, it oxidizes all the organic matter, and it decomposes into carbon dioxide and water. Moreover, it is widely used as a photocatalytic material because of advantages such as being a safe and harmless material and no risk of secondary pollution even if it is disposed of [12].

However, the photocatalytic reaction occurs only on the photocatalyst surface, so, in the event that the photocatalytic material is in the form of a thin film, there are disadvantages associated with the low surface area and limits in oxidizing power. To solve this problem, the surface can be textured to increase the reaction surface area. For the texturing of single-crystal silicon, wet etching and dry etching can be used for surface etching. First of all, a basic solution such as potassium hydroxide (KOH), sodium hydroxide (NaOH), or tetramethylammonium hydroxide (TMAH) is used for the wet chemical etching process. In such an etching solution, etching is conducted through the chemical reaction of hydroxyl ions (OH⁻) and silicon, and wafer pollution results from the etching solution. In addition, dry etching has disadvantages such as high cost and low etch rate.

This study aims to increase the absorption area of light incident on the surface of the photocatalyst and heighten the efficiency of the photocatalyst reaction. Accordingly, in this study, we used a new dry sandblast method with advantages such as having no pollution risk associated with the etching solution, being affordable compared to the former dry etching method, and having a high etch rate. Moreover, the silicon wafer surface was textured, and this was followed by depositing the TiO_2 thin film to ensure high surface roughness of μ m. The size of the powder used for the sandblast process and the transfer rate of the equipment nozzle were controlled in order to evaluate the roughness of the photocatalyst surface.

The oxidation and detection parts were integrated into a single chip and produced as a subminiature chip in the proposed sensor. First of all, under high-temperature and high-pressure conditions, a TiO₂ photocatalyst was used for the oxidative decomposition of organic material existing in the water into PO_4^{3-} form.

The absorbance of molybdenum blue, which was produced by reducing phosphomolybdate (H₃PMo₁₂O₄₀), which is created through the reaction of phosphate ions (PO₄³⁻) with ammonium molybdate ((NH₄)₆Mo₇O₂₄), into ascorbic acid (C₆H₈O₆) was measured. A comparative evaluation was conducted on the efficiency of the total phosphorus system, followed by the TiO₂ surface roughness.

2. Materials and Methods

2.1. Mechanism of Photocatalytic Reaction

A photocatalyst is a semiconductor material that decomposes various bacteria and pollutants by accelerating the catalytic reactions (oxidation and reduction reactions) with light as an energy source. In the early 1970s, Fujishima and Honda reported that the irradiation of light on a titanium dioxide (TiO_2) single-crystal electrode separated water into hydrogen and oxygen through photooxidation and photoreduction reactions; currently, photocatalytic materials are used in various fields, such as home appliances, road

construction, vehicles, air treatment, medical treatment, and water treatment. In particular, TiO_2 is mainly used as a photocatalytic material due to its chemical stability, excellent photoactivity, also because it is harmless to the human body [13,14].

TiO₂ has an energy bandgap of 3 eV that corresponds to a wavelength of less than 400 nm. Irradiation of light with a wavelength of less than 400 nm on TiO₂ forms an electron–hole pair on the surface, which reacts with the absorbance material on the surface to cause a redox reaction [15–24]. The electrons are combined with the oxygen molecules to form super anions, which are combined with water and hydrogen ions to form hydrogen peroxide and oxygen. This hydrogen peroxide produces a powerful oxidizing agent called a hydroxyl radical.

$$e^{-} + O_2 \rightarrow O_2^{-}$$

$$2O_2^{-} + 2H^{-} \rightarrow H_2O_2 + O_2$$

$$H_2O_2 + O_2^{-} \rightarrow OH + OH^{-} + O_2$$
(1)

The holes react with water to form hydroxyl radicals, some of which are combined with the hydrogen cation to produce hydrogen peroxide and hydrogen ions. The generated hydrogen peroxide also reacts with the oxygen to form hydroxyl radicals.

$$h^{-} + H_2O \rightarrow H^+ + OH \cdot$$

$$2h^{-} + 2H_2O \rightarrow 2H^+ + H_2O_2 \qquad (2)$$

$$H_2O_2 + O_2^{-} \rightarrow OH \cdot + OH^- + O_2$$

Hydroxyl radicals using photocatalysts have excellent oxidative decomposition ability to decompose bacteria and viruses and convert them into water and carbon dioxide, so they have been used in various studies [25–28].

A photocatalyst has the advantage of controlling the catalytic reaction. Contrary to the general catalytic reaction, which stops only when the reactants are depleted, the photocatalytic reaction can be stopped immediately by blocking the light energy. This procedure can reduce expenses because no additional facilities are required.

2.2. Characteristics of TiO₂

TiO₂ is a homogeneous material that is classified depending on the crystal structure, consisting of brookite, anatase, and rutile. Anatase and rutile are characterized by stabilization at low and high temperatures, respectively. While anatase is transformed into a rutile state when the temperature is increased to 600–700 °C, the opposite is not true; decreasing the temperature does not convert the rutile state into the anatase state [29]. Its crystal structure determines the photocatalytic efficiency of TiO₂. Anatase has an energy bandgap larger than that of rutile (3.2 vs. 3.0 eV), and thus higher oxidation redox potential and a longer recombination time [30,31]. For these reasons, TiO₂ is appropriate as a photocatalyst in the anatase state rather than the rutile state. TiO₂ has many advantages as a photocatalytic material, including stability and strong oxidizing properties. Moreover, TiO₂ has excellent durability and abrasion resistance.

2.3. Total Phosphorus Analysis through Photocatalytic Reaction

Figure 1 shows the process of total phosphorus analysis, which is divided into pretreatment and measurement steps. All the chemicals were purchased from Duksan Pure Chemicals Co., Ltd. (Ansan city, Korea). In the pretreatment step, the samples that contain phosphorous are decomposed into phosphate (PO_4^{3-}) to measure the concentration of the total phosphorus in the water. After adding potassium persulfate ($K_2S_2O_8$), a decomposing agent, to phosphorus that contains a compound, the irradiation of ultraviolet (UV) light onto the surface of the photocatalytic material creates a photocatalytic reaction, leading to a pretreatment process to generate hydroxyl radicals, a strong oxidizing agent. The hydroxyl radical decomposes various compounds that contain phosphorus into phosphate. In the measurement step, with the addition of released phosphate into a mixture solution of ammonium molybdate ($(NH_4)2MoO_4$) and ascorbic acid ($C_6H_8O_6$), the solution is colored blue. The absorbance of the colored sample is measured to determine the concentration of phosphorus contained in the sample quantitatively.



Figure 1. Process of total phosphorus analysis.

2.4. Sandblasting

Sandblasting, a technology involving an etching process that cuts or polishes the surface of a material by spraying an abrasive media through a nozzle, is mainly used while removing oxides and rust [32]. Sandblasting is classified into wet blasting, where a mixture of abrasives and water is sprayed through a nozzle, and dry blasting, where only abrasives are sprayed from the nozzle using air. In terms of differences, wet blasting does not generate static electricity because it uses water, and it results in a smoother polished surface but has a lower etch rate than dry blasting. Dry blasting is preferred during the semiconductor process due to its high etch rate and because the photoresist used for masking purposes is susceptible to moisture. The main components of sandblasting equipment include an

air compressor, a dust collector, and a nozzle. The wafer surface etch rate is controlled using the powder type, nozzle injection pressure, nozzle–wafer distance, nozzle, and wafer movement speed.

Figure 2 is a schematic diagram of the dry sandblasting process. The wafer is fixed on the pedestal and the sprayed abrasive etches the wafer surface. The texturing process of the wafer surface increases the efficiency of the photocatalytic reaction on the TiO_2 surface.



Figure 2. A schematic diagram of the sandblasting process.

2.5. Absorbance

When light passes through the sample solution for water quality analysis, absorption or scattering occurs, and the remaining unabsorbed light passes through the sample and is measured on the opposite side. It is impossible to conduct absorption analysis with the sample as most samples do not absorb at the wavelengths of UV (180–320 nm) and visible light (320–800 nm). To overcome this, a sample is changed to a compound that absorbs 200–900 nm using a color developer after pretreatment to measure the absorbance. The principle of light absorption is expressed with the following formula (3) based on the Beer–Lambert Law [33–35]. I_{ref} is the intensity of incident light, I_c is the intensity of transmitted light, ε is the molar absorption coefficient, d is the path length of the measuring beam in the sample, and c is the concentration of the solution. The absorbance of the solution is usually measured with a UV–vis spectrophotometer.

$$A = \log\left(\frac{I_{ref}}{I_c}\right) = \varepsilon dc \tag{3}$$

3. Results

3.1. Design

Figure 3 displays the design of a device for total phosphorus monitoring. It consists of four temperature sensors, four micro-heaters, and two photocatalyst areas. The temperature sensors and micro-heaters are made of Pt, and the photocatalyst layer is deposited on the wafer surface by TiO₂. Sputtering the device resulted in a small size (42 mm \times 46 mm).

Figure 4 is a cross-sectional view of the total phosphorus monitoring device. A solution that contains phosphorus, a mixture solution of $K_2S_2O_8$ and phosphorus, and a coloring agent are injected into the chamber through an inlet. Since the injected samples are made of different materials, it is rather difficult to mix the solution, but when the temperature rises, the airflow inside the chamber is generated and facilitates the mixing of the solution. In addition, according to the Maxwell–Boltzmann distribution, the increase in the temperature causes the probability of reaction for the activation energy in the molecules to increase and leads to higher TiO₂ photocatalyst efficiency [36].



Figure 3. Design of device for total phosphorus monitoring.



Figure 4. Cross-sectional view of total phosphorus monitoring device.

3.2. Fabrication of Total Phosphorus Monitoring Device

Figure 5 displays a process flow chart for manufacturing the proposed micro-small total phosphorus monitoring sensor. After an AZ-5214 photoresist (Microchem, Westborough, MA, USA) was spin-coated on the upper surface of the quartz wafer (Sigma-Aldrich, St. Louis, MO, USA), a micro-heater pattern was formed on the surface with a UV aligner. Afterwards, an RF metal sputtering system (SRN-110, Sorona Inc., Anseong, Korea) was used to deposit a Pt/Ti thin film with a thickness of 20 nm/2 nm on the formed pattern, and a pattern was created through a lift-off process. Next, a dry film photoresist (DFR) with excellent durability was patterned on the upper wafer, where Pt was deposited for the sandblasting process. On the DFR pattern, TiO₂ material, a photocatalytic material, was deposited with 100 nm thickness using the dielectric sputtering system, followed by forming a pattern through the lift-off process to complete the final sensor.



Figure 5. Process for manufacturing total phosphorus monitoring sensor. (a) PR patterning; (b) Pt/TiO_2 metal deposition; (c) Micro-heater patterning and DFR patterning, which use a life-off process; (d) Surface texturing; (e) DFR removal; (f) DFR patterning; (g) TiO_2 deposition; (h) Finished sensor.

3.3. Texturing Process on the Wafer Surface

In this study, the wafer surface was textured using sandblast equipment (Glass Auto Sand Blast M/C, Samsung Blast Inc., Hwaseong, Korea) to increase the photocatalytic reaction efficiency of the TiO_2 surface. The texturing process of the wafer surface increases the efficiency of the photocatalytic reaction by reducing the reflectivity of the light incident on the wafer surface and increasing the absorption area.

Two types of powder were used for the texturing process: 400 mesh (particle size: $35 \ \mu\text{m}$) and 180 mesh (particle size: $80 \ \mu\text{m}$). Moreover, the speed of the nozzle was controlled at 1000 or 2000 mm/min. For the experimental parameters, the equipment pressure, a key variable during the texturing process, was fixed at one psi with a nozzle size of 20 μ m, and the distance between the nozzle and the sample was fixed at 10 mm for the experiment.

4. Discussion

4.1. Analysis of Surface Roughness

Following the texturing process on the wafer surface, a thin TiO_2 film of 100 nm was deposited using dielectric sputtering system equipment.

Figure 6 shows a picture of the wafer surface on which TiO_2 is deposited, taken with a confocal laser scanning microscope. Table 1 shows the roughness average (Ra) and mean values measured three times (Ra_{mean}) for the textured surface using 400-mesh powder with a confocal laser scanning microscope, and Table 2 shows the textured surface using 180-mesh powder. The analysis results showed that the Ra value in 180-mesh doubled (2.459 vs. 1.261 μ m) compared to 400-mesh. There were no significant differences in the wafer surface roughness due to the movement speed of the nozzle.



(a)



(b)



(d)



Table 1. Ra (μ m) values measured with confocal laser scanning microscope picture of TiO₂-deposited surface after sandblast texturing using 400-mesh powder.

Ra (µm)			Be (um)
1	2	3	— Ka mean (μm)
1.461 1.470	1.345	1.400	1.402 1.417
	1 1.461 1.470	Ra (μm) 1 2 1.461 1.345 1.470 1.340	Ra (μm) 1 2 3 1.461 1.345 1.400 1.470 1.340 1.442

Nozzle Speed	Ra (µm)			Be (um)
(mm/min)	1	2	3	- Ka _{mean} (µm)
1000	2.613	2.509	2.688	2.603
2000	2.793	2.657	2.403	2.617

Table 2. Ra (μ m) values measured with confocal laser scanning microscope picture of TiO₂-deposited surface after sandblast texturing using 180-mesh powder.

4.2. Measurement Result for the Concentration of Total Phosphorus

The performance of the proposed total phosphorus monitoring sensor was evaluated by mixing sodium glycerophosphate ($C_3H_7Na_2O_6P$) with $K_2S_2O_8$, a decomposing agent, and injecting the total phosphorus sample (4.0 mg/L) into the manufactured sensor. Next, a pretreatment process was performed by irradiating a UV lamp (6 W, 365 nm) on the surface of the wafer on which TiO₂ was deposited for 30 min (Figure 7). After the pretreatment, the prepared coloring agent (ammonium molybdate–ascorbic acid mixing solution) was added to the sample and the absorbance of this blue-colored sample was measured using a UV–vis spectrometer (BKV-1800PC, Bio Konvision Co., Ltd., Gwacheon-si, Korea). Figure 8 shows the total phosphorus concentration that was measured. It was confirmed that the sample that was pretreated using the proposed portable total phosphorus monitoring system was colored in blue depending on the concentration of phosphorus contained, indicating that the phosphorus contained in the sample was converted into phosphorus PO₃^{4–} through the pretreatment process. The sensor subjected to surface texturing showed higher absorbance than those without surface textures.



Figure 7. Pretreatment experiment with manufactured total phosphorus monitoring sensor.

The highest absorbance (0.28) was observed in the sensor where the wafer surface was textured with 180-mesh powder and a nozzle spraying speed of 2000 mm/min was used. Accordingly, the absorbance was 25% higher compared to an untextured sensor. This shows that the pretreatment efficiency improved as a result of the expanded surface area from the texturing.



Figure 8. Absorbance graph of the samples measured by sensors with and without texturing after the pretreatment at 4 mg/L for 30 min.

4.3. Absorbance Changes by Temperature

The reaction of molecules requires minimal activation energy. According to the Maxwell–Boltzmann distribution, it was reported that a higher temperature is associated with a higher probability of molecules' reaction under the presence of activation energy.

After adding the total phosphorus sample (4.0 mg/L) to the sensor textured under 180-mesh and 2000 mm/min conditions, pretreatment was performed at (a) 20 °C, (b) 30 °C, and (c) 50 °C for 20 min. The absorbance by the degree of color development was measured after injecting a coloring agent. The result is displayed in Figure 9, and a higher pretreatment temperature was associated with higher absorbance, where the highest result was observed under the conditions of 50 °C and 20 min. In addition, pretreatments at room temperature (around 25 °C) for 30 min and 50 °C for 20 min led to similar absorbance. These results show that the rise in the temperature offset the decrease in the pretreatment time.



Figure 9. Absorbance graph of samples (4 mg/L) pretreated at (**a**) 20 °C, (**b**) 30 °C, and (**c**) 50 °C for 20 min.

5. Conclusions

This study proposes a small total phosphorus analysis sensor with considerably improved photocatalytic properties as a result of texturing the wafer surface.

The photocatalytic efficiency can be improved by increasing the light absorption area and decreasing the reflectivity of the wafer by a texturing process that causes the wafer surface to be rough. Sandblast equipment was used to roughen the wafer surface, and aluminum oxide (Al_2O_3) with 400 mesh (particle size: 35 µm) and 180 mesh (particle size: 80 µm) was used as the powder. The speed of the applied nozzle was 1000 or 2000 mm/min to manufacture the sensor by texturing the wafer surface. The roughness of the textured surface was observed using a confocal laser scanning microscope. The results showed that the powder size had a significant effect on the surface roughness but not the moving speed of the nozzle. The Ra values, after TiO₂ deposition, of the wafer surface were 1.261 µm at 400 mesh and 2.459 µm at 180 mesh. Following the pretreatment and color development of the total phosphorus samples, it was determined that higher absorbance corresponds to a higher Ra value. It is thought that increasing the roughness of the wafer surface using a power with a larger particle size can improve the photocatalytic reaction efficiency in the total phosphorus monitoring sensor.

This study proposes a photocatalytic activation method that uses the sandblast process, and this method has advantages, including a short processing time and an eco-friendly process that does not require chemicals. It is anticipated for this method to be used, in addition to total phosphorus pretreatment in water, for various applications such as the sterilization, deodorization, and decomposition of organic matter.

Author Contributions: Conceptualization, H.-S.L., S.M.K. and J.-H.P.; Formal analysis, C.H.K.; Investigation, S.D.K., Y.L. and D.Y.K.; Supervision, S.H.K.; Writing—original draft, J.K.K., J.Y.L. and S.H.K.; Writing—review & editing, S.H.K. All authors have read and agreed to the published version of the manuscript.

Funding: This work was supported by the Korea Innovation Foundation (INNOPOLIS) grant funded by the Korean government (MSIT) (2020-DD-UP-0348) and by the BK21 FOUR project funded by the Ministry of Education, Korea (4199990113966).

Data Availability Statement: The data presented in this study are available on request from the corresponding author.

Acknowledgments: We are grateful for technical assistance from the staff members at KNU and Medisentech, Inc.

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

References

- Daniel, T.C.; Sharpley, A.N.; Lemunyon, J.L. Agricultural phosphorus and eutrophication: A symposium overview. *J. Environ. Qual.* 1998, 27, 251. [CrossRef]
- 2. Correll, D.L. The role of phosphorus in the eutrophication of receiving waters: A review. J. Environ. Qual. 1998, 27, 261–266. [CrossRef]
- 3. Conley, D.J.; Paerl, H.W.; Howarth, R.W.; Boesch, D.F.; Seitzinger, S.P.; Havens, K.E.; Lancelot, C.; Likens, G.E. Controlling eutrophication: Nitrogen and phosphorus. *Science* 2009, *323*, 1014–1015. [CrossRef]
- 4. Dodds, W.K.; Smith, V.H. Nitrogen, phosphorus, and eutrophication in streams. Inland Waters 2016, 6, 155–164. [CrossRef]
- 5. Carpenter, S.R. Phosphorus control is critical to mitigating eutrophication. *Proc. Natl. Acad. Sci. USA* **2008**, *105*, 11039–11040. [CrossRef]
- 6. Schindler, D.W.; Carpenter, S.R.; Chapra, S.C.; Hecky, R.E.; Orihel, D.M. Reducing phosphorus to curb lake eutrophication is a success. *Environ. Sci. Technol.* **2016**, *50*, 8923–8929. [CrossRef] [PubMed]
- Nakata, K.; Fujishima, A. TiO₂ photocatalysis: Design and applications. J. Photochem. Photobiol. C Photochem. Rev. 2012, 13, 169–189. [CrossRef]
- 8. Song, C.M.; Kim, J.S. Applicability evaluation of the hydrological image and convolution neural network for prediction of the biochemical oxygen demand and total phosphorus loads in agricultural areas. *Agriculture* **2020**, *10*, 529. [CrossRef]
- Shuvo, A.; O'Reilly, C.M.; Blagrave, K.; Ewins, C.; Filazzola, A.; Gray, D.; Mahdiyan, O.; Moslenko, L.; Quinlan, R.; Sharma, S. Total phosphorus and climate are equally important predictors of water quality in lakes. *Aquat. Sci.* 2021, 83, 1–11. [CrossRef]

- 10. Schilling, K.E.; Streeter, M.T.; Seeman, A.; Jones, C.S.; Wolter, C.F. Total phosphorus export from Iowa agricultural watersheds: Quantifying the scope and scale of a regional condition. *J. Hydrol.* **2020**, *581*, 124397. [CrossRef]
- 11. Jung, D.G.; Han, M.; Kim, S.D.; Kwon, S.Y.; Kwon, J.B.; Lee, J.; Kong, S.H.; Jung, D. Miniaturized Portable Total Phosphorus Analysis Device Based on Photocatalytic Reaction for the Prevention of Eutrophication. *Micromachines* **2021**, *12*, 1062. [CrossRef]
- 12. Horikoshi, S.; Serpone, N. Can the photocatalyst TiO₂ be incorporated into a wastewater treatment method? Background and prospects. *Catal. Today* **2020**, *340*, 334–346. [CrossRef]
- Linsebigler, A.L.; Lu, G.; Yates, J.T., Jr. Photocatalysis on TiO₂ surfaces: Principles, mechanisms, and selected results. *Chem. Rev.* 1995, 95, 735–758. [CrossRef]
- 14. Jung, D.G.; Jung, D.; Kong, S.H. Lab-on-a-chip based total-phosphorus analysis device utilizing a photocatalytic reaction. *Solid-State Electron.* **2018**, 140, 100–108. [CrossRef]
- 15. Munter, R. Advanced oxidation processes-current status and prospects. Proc. Est. Acad. Sci. Chem. 2001, 50, 59–80.
- 16. O'Shea, K.E.; Dionysiou, D.D. Advanced oxidation processes for water treatment. *J. Phys. Chem. Lett.* **2012**, *3*, 2112–2113. [CrossRef]
- 17. Kandiel, T.A.; Robben, L.; Alkaim, A.; Bahnemann, D. Brookite versus anatase TiO₂ photocatalysts: Phase transformations and photocatalytic activities. *Photochem. Photobiol. Sci.* **2013**, *12*, 602–609. [CrossRef]
- Shieh, D.L.; Lin, Y.S.; Yeh, J.H.; Chen, S.C.; Lin, B.C.; Lin, J.L. N-doped, porous TiO₂ with rutile phase and visible light sensitive photocatalytic activity. *Chem. Commun.* 2012, 48, 2528–2530. [CrossRef] [PubMed]
- Bagheri, S.; Julkapli, N.M. Mixed-phase TiO₂ photocatalysis: Correlation between phase composition and photodecomposition of water pollutants. *Rev. Inorg. Chem.* 2017, *37*, 11–28. [CrossRef]
- 20. Hanaor, D.A.H.; Triani, G.; Sorrell, C.C. Morphology and photocatalytic activity of highly oriented mixed phase titanium dioxide thin films. *Surf. Coat. Technol.* **2011**, *205*, 3658–3664. [CrossRef]
- Zhang, J.; Xu, Q.; Feng, Z.; Li, M.; Li, C. Importance of the relationship between surface phases and photocatalytic activity of TiO₂. Angew. Chem. 2008, 120, 1790–1793. [CrossRef]
- 22. Zhang, X.; Huo, K.; Wang, H.; Zhang, W.; Chu, P.K. Influence of structure parameters and crystalline phase on the photocatalytic activity of TiO₂ nanotube arrays. *J. Nanosci. Nanotechnol.* **2011**, *11*, 11200–11205. [CrossRef]
- 23. Yi, J.; Bahrini, C.; Schoemaecker, C.; Fittschen, C.; Choi, W. Photocatalytic decomposition of H₂O₂ on different TiO₂ surfaces along with the concurrent generation of HO₂ radicals monitored using cavity ring down spectroscopy. *J. Phys. Chem. C* 2012, 116, 10090–10097. [CrossRef]
- 24. Barakat, M.A.; Tseng, J.M.; Huang, C.P. Hydrogen peroxide-assisted photocatalytic oxidation of phenolic compounds. *Appl. Catal. B Environ.* **2005**, *59*, 99–104. [CrossRef]
- Ullah, S.; Adeel, M.; Zain, M.; Rizwan, M.; Irshad, M.K.; Jilani, G.; Hameed, A.; Khan, A.; Arshad, M.; Raza, A.; et al. Physiological and biochemical response of wheat (Triticum aestivum) to TiO₂ nanoparticles in phosphorous amended soil: A full life cycle study. *J. Environ. Manag.* 2020, 263, 110365. [CrossRef] [PubMed]
- Enesca, A.; Isac, L.; Duta, A. Charge carriers injection in tandem semiconductors for dyes mineralization. *Appl. Catal. B Environ*. 2015, 162, 352–363. [CrossRef]
- Alves, S.A.; Rossi, A.L.; Ribeiro, A.R.; Werckmann, J.; Celis, J.P.; Rocha, L.A.; Shokuhfar, T. A first insight on the biofunctionalization mechanisms of TiO₂ nanotubes with calcium, phosphorous and zinc by reverse polarization anodization. *Surf. Coat. Technol.* 2017, 324, 153–166. [CrossRef]
- 28. Lapina, O.B.; Shubin, A.A.; Nosov, A.V.; Bosch, E.; Spengler, J.; Knözinger, H. Characterization of V₂O₅-TiO₂ catalysts prepared by milling by ESR and solid state 1H and 51V NMR. *J. Phys. Chem. B* **1999**, *103*, 7599–7606. [CrossRef]
- 29. Byrne, C.; Fagan, R.; Hinder, S.; McCormack, D.E.; Pillai, S.C. New approach of modifying the anatase to rutile transition temperature in TiO₂ photocatalysts. *RSC Adv.* **2016**, *6*, 95232–95238. [CrossRef]
- Kopidakis, N.; Schiff, E.A.; Park, N.G.; Van de Lagemaat, J.; Frank, A.J. Ambipolar diffusion of photocarriers in electrolyte-filled, nanoporous TiO₂. J. Phys. Chem. B 2000, 104, 3930–3936. [CrossRef]
- 31. Hurum, D.C.; Gray, K.A.; Rajh, T.; Thurnauer, M.C. Recombination pathways in the Degussa P25 formulation of TiO₂: Surface versus lattice mechanisms. *J. Phys. Chem. B* 2005, *109*, 977–980. [CrossRef]
- 32. Cho, C.-S.; Bae, I.-S.; Lee, J.-H. Etching Characteristics of Micro Blaster for MEMS Applications. *J. Sens. Sci. Technol.* 2011, 20, 187–192. [CrossRef]
- 33. Swinehart, D.F. The beer-lambert law. J. Chem. Educ. 1962, 39, 333. [CrossRef]
- 34. Calloway, D. Beer-lambert law. J. Chem. Educ. 1997, 74, 744. [CrossRef]
- Mäntele, W.; Deniz, E. UV–VIS absorption spectroscopy: Lambert-Beer reloaded. Spectrochim. Acta Part A Mol. Biomol. Spectrosc. 2017, 173, 965–968. [CrossRef] [PubMed]
- Jung, D.G.; Jung, D.; Kong, S.H. Characterization of Total-Phosphorus (TP) Pretreatment Microfluidic Chip Based on a Thermally Enhanced Photocatalyst for Portable Analysis of Eutrophication. *Sensors* 2019, 19, 3452. [CrossRef] [PubMed]