Article

Activation Treatments and SiO₂/Pd Modification of Sol-gel TiO₂ Photocatalysts for Enhanced Photoactivity under UV Radiation

Julien G. Mahy ^{1,2,*} , Valérie Sotrez ², Ludivine Tasseroul ², Sophie Hermans ¹ and Stéphanie D. Lambert ²

¹ Institute of Condensed Matter and Nanosciences (IMCN), Université catholique de Louvain, Place Louis Pasteur 1, 1348 Louvain-la-Neuve, Belgium; sophie.hermans@uclouvain.be

² Department of Chemical Engineering—Nanomaterials, Catalysis & Electrochemistry, University of Liège, B6a, Quartier Agora, Allée du six Août 11, 4000 Liège, Belgium; valerie.sotrez@gmail.com (V.S.); ludivine.tasseroul@gmail.com (L.T.); stephanie.lambert@uliege.be (S.D.L.)

* Correspondence: julien.mahy@uclouvain.be; Tel.: +32-4-3664771

Received: 3 September 2020; Accepted: 13 October 2020; Published: 14 October 2020

Abstract: The objective of this work is to improve the efficiency of TiO₂ photocatalysts by activation treatments and by modification with palladium nanoparticles and doping with SiO₂. The influence of the additive loading was explored, and two activation treatments were performed: UV exposition and H₂ reduction. TiO₂/SiO₂/Pd photocatalysts were synthesized by an original cogelation method: a modified silicon alkoxide, i.e., [3-(2-aminoethyl)aminopropyl]trimethoxysilane (EDAS), was used to complex the palladium ions, thanks to the ethylenediamine group, while the alkoxide groups reacted with TiO₂ precursors. Pure TiO₂ was also synthesized by the sol-gel process for comparison. X-ray diffraction evidenced that the crystallographic structure of TiO₂ was anatase and that Pd was present, either in its oxidized form after calcination, or in its reduced form after reduction. The specific surface area of the samples varied from 5 to 145 m² g^{−1}. Transmission electron microscopy allowed us to observe the homogeneous dispersion and nanometric size of Pd particles in the reduced samples. The width of the band gap for pure TiO₂ sample, measured by UV/Visible diffuse reflectance spectroscopy at approximately 3.2 eV, corresponded to that of anatase. The band gap for the TiO₂/SiO₂/Pd composite samples could not be calculated, due to their high absorption in visible range. The photocatalytic activity of the various catalysts was evaluated by the degradation of a methylene blue solution under UV radiation. The results showed that the photocatalytic activity of the catalysts was inversely proportional to the content of silica present in the matrix. A small amount of silica improved the photocatalytic activity, as compared to the pure TiO₂ sample. By contrast, a high amount of silica delayed the crystallization of TiO₂ in its anatase form. The activation treatment under UV had little influence on photocatalytic efficiency. The introduction of Pd species increased the photocatalytic activity of the samples because it allowed for a decrease in the rate of electron–hole recombinations in TiO₂. The reduction treatment improved the activity of photocatalysts, whatever the palladium content, thanks to the reduction of Ti⁴⁺ into Ti³⁺, and the formation of defects in the crystallographic structure of anatase.

Keywords: TiO₂; sol-gel process; SiO₂; Pd-modification; activation treatments; photocatalysis

1. Introduction

For decades, different processes have been developed to reduce the current pollution in water, air and soil [1,2]. Among these various methods, photocatalysis had emerged as a potential technique to degrade organic pollutants [3,4]. Its principle lies on the use of a semiconductor as photocatalyst,

which is activated by light illumination. Its activation produces a series of redox reactions leading to radicals' formation. These radicals can attack organic pollutants and degrade them in CO_2 and H_2O if the degradation is complete [3,4].

The most used photocatalyst is TiO_2 [5–7]. This material is activated under UV light due to its band gap around 3.2 eV for the anatase phase [4]. TiO_2 presents the advantage of being easily available, inexpensive and non-toxic. The use of TiO_2 as photocatalyst has two main limitations [4]: (i) the fast charge recombination and (ii) the large band gap value. Indeed, if the recombination of the photo-generated species (electrons, e^- , and holes, h^+) is fast, the production of radicals is low and the degradation is less effective. Furthermore, if the band gap is large, the energy required for the electron transfer is high and only UV radiation can be used. This study focused on the improvement of the charge recombination phenomenon.

Regarding the improvement of the recombination time, the major modification of TiO_2 materials is the addition of metallic nanoparticles such as Ag [8–11], Au [12], Pt [12,13] or Pd [8,14]. In this case, the metallic nanoparticle plays the role of electron trap, allowing us to increase the recombination time [9]. Combination with other semiconductors has also been investigated to increase the recombination time [4]. In this case, synergetic effects can be observed leading to a better charge separation or an increased photostability. Different combinations of semiconductors were tested, such as TiO_2/ZnO [15], TiO_2/CdS [4] or $\text{TiO}_2/\text{ZrO}_2$ [16]. The introduction of SiO_2 in the TiO_2 matrix can also enhance the photoactivity due to surface acidity modification, the introduction of defects or increased surface area [9,17]. In order to optimize the modification efficiency, the additive needs to be highly dispersed in the TiO_2 matrix. Previous works have reported that a high dispersion of metallic nanoparticles into a SiO_2 and TiO_2 matrix can be achieved through the use of a complexing agent, e.g., ethylenediamine [9,18,19]. Indeed, by using modified silicon alkoxide, small amounts of Ag, Cu or Ni nanoparticles are dispersed in either TiO_2 or SiO_2 . The use of this silicon alkoxide introduces a SiO_2 doping in the TiO_2 matrix, and this doping is referenced as effective to increase the photoactivity [9,17,20].

In this study, TiO_2 photocatalyst was modified with Pd species, in order to increase the photocatalytic property under UV light for the degradation of organic dye. To finely disperse these Pd species, a modified silicon alkoxide (3-(2-aminoethyl) aminopropyl]trimethoxysilane, EDAS) was cogelled with Ti alkoxide. Indeed, this silicon alkoxide possesses an ethylenediamine function able to complex metallic ions; SiO_2 was also introduced into the materials. Different amounts were added to study the influence of SiO_2 addition on TiO_2 properties. This original preparation method allows the modification in one step. Four different samples were prepared with four Pd loadings; one pure TiO_2 and one $\text{TiO}_2/\text{SiO}_2$ were also prepared for comparison. To further increase the photoefficiency, two activation treatments were applied to the sample: UV activation and H_2 reduction. The physicochemical properties of the resulting materials were determined thanks to nitrogen adsorption–desorption, X-ray diffraction, inductively coupled plasma optical emission spectroscopy, diffuse reflectance and transmission electron microscopy. The photocatalytic activity was followed on the degradation of methylene blue (MB), under UV-A illumination, during 6 h of reaction.

2. Results and Discussion

Six samples were prepared: one pure titania (named TiO_2), one titania doped with silica (named $\text{TiO}_2/\text{SiO}_2$) and four titania samples doped with silica and modified with Pd named $\text{TiO}_2/\text{SiO}_2/\text{PdX}$, where X corresponds to the theoretical wt.% of Pd and can vary between 1, 5, 12 and 22 wt.% of Pd. Table 1 summarizes the amounts of reactants for each sample. The weight amount of SiO_2 is in the same range as Pd. The UV-treated samples are denoted with “-UV”, while those reduced under H_2 are denoted with “- H_2 ”.

Table 1. Synthesis operating variables and additive loadings determined by ICP.

Sample	n_{Solvent} (mmol)	n_{Pd} (mmol)	n_{EDAS} (mmol)	n_{TIPT} (mmol)	n_{Water} (mmol)	Gelification Time (min)	Theoretical Loading SiO ₂ /Pd (wt.%)	Actual Loading SiO ₂ /Pd (wt.%)
Pure TiO ₂	1830	-	-	91.5	183	9	-	-
TiO ₂ /SiO ₂	1830	-	6.88	84.6	180	8	5.8/0	4.9/0
TiO ₂ /SiO ₂ /Pd1	1830	0.69	1.38	90.1	182	>60	0.98/1	1.2/1.1
TiO ₂ /SiO ₂ /Pd5	1830	3.44	6.88	84.6	180	40	4.9/5	4.7/4.2
TiO ₂ /SiO ₂ /Pd12	1830	8.60	17.20	74.3	174	>60	11.76/12	12.9/12.2
TiO ₂ /SiO ₂ /Pd22	1830	17.4	34.90	56.6	166	>60	21.56/22	24.6/21.8

n_{Solvent} = amount of 2-methoxyethanol; n_{Pd} = amount of palladium acetylacacetate; n_{EDAS} = amount of EDAS;
 n_{TIPT} = amount of TIPT; n_{Water} = amount of water.

2.1. Colors and Compositions of Photocatalytic Samples

The pure and TiO₂/SiO₂ samples had a white color, characteristic of TiO₂ and SiO₂ materials. The Pd-modified TiO₂/SiO₂ samples were brown after calcination and darker when the amount of Pd increased. The brown color is characteristic of PdO material. The samples, after UV activation, kept their initial coloration, while after H₂ reduction, the Pd-modified ones became black, which is characteristic of metallic palladium. The pictures of the samples are shown in Supplementary Materials Figure S1.

The actual loading (measured by ICP) is given in Table 1 and was close to the theoretical one, calculated on the basis of the engaged amounts. In each composite sample, the amount of SiO₂ is similar to the amount of Pd, as expected.

2.2. Crystallinity

The XRD patterns of all calcined samples are presented in Figure 1a. For pure TiO₂ sample, the characteristic peaks of TiO₂ anatase were detected (JCPDS 21-1272: 25.3°, 37.9°, 48.0°, 54.6°, 62.8° and 68.8°). When additive was added, the intensity of anatase peaks decreased and even disappeared for TiO₂/SiO₂/Pd22 sample. For the samples modified with Pd, additional peaks appeared at 34°, 42°, 54.9° and 60.3° (JCPDS 75-584), which corresponded to palladium oxide. The intensity of these peaks increased with the amount of Pd (Figure 1a).

Figure 1. Cont.

Figure 1. (a) XRD patterns of calcined samples: (x) pure TiO₂, (•) TiO₂/SiO₂, (■) TiO₂/SiO₂/Pd1, (◆) TiO₂/SiO₂/Pd5, (▲) TiO₂/SiO₂/Pd12, and (♣) TiO₂/SiO₂/Pd22; and (b) XRD patterns of reduced samples: (x) pure TiO₂—H₂, (•) TiO₂/SiO₂—H₂, (■) TiO₂/SiO₂/Pd1—H₂, (◆) TiO₂/SiO₂/Pd5—H₂, (▲) TiO₂/SiO₂/Pd12—H₂ and (♣) TiO₂/SiO₂/Pd22—H₂. (A) Reference pattern of anatase, (PdO) reference pattern of palladium oxide and (Pd) reference pattern of palladium.

After UV activation, no modification of the crystallinity was observed (XRD patterns identical to Figure 1a, not shown). Contrarily, after H₂ reduction (Figure 1b), the peaks of metallic palladium (JCPDS 87-0638: 39.8°, 46.2° and 67.6°) were observed replacing those of palladium oxide, as expected.

Concerning the crystallite size evaluated by using the Scherrer equation, the TiO₂ crystallites ($d_{\text{XRD Ti}}$) decreased with the additive content: from 34 nm for pure TiO₂ to 7 nm for TiO₂/SiO₂/Pd12 calcined sample (Table 2). For the PdO crystallites ($d_{\text{XRD PdO}}$, for the calcined samples), they increased with the amount of Pd: from 8 nm for TiO₂/SiO₂/Pd5 to 25 nm for TiO₂/SiO₂/Pd22 sample. The metallic Pd nanoparticles ($d_{\text{XRD Pd}}$ for the reduced samples) had a slightly smaller size than PdO crystallites (Table 2).

Table 2. Samples' crystalline, textural and optical properties.

Sample	$d_{\text{XRD Ti}}$ (nm) ±1	$d_{\text{XRD PdO}}$ (nm) ±1	$d_{\text{XRD Pd}}$ (nm) ±1	S_{BET} (m ² g ^{−1}) ±5	V_{DR} (cm ³ g ^{−1}) ±0.01	V_{P} (cm ³ g ^{−1}) ±0.1	$d_{\text{TEM Ti}}$ (nm) ±5	$d_{\text{TEM PdO}}$ (nm) ±4	$d_{\text{TEM Pd}}$ (nm) ±4	E_{g} (eV) ±0.01
Pure TiO ₂	34	— ^b	— ^b	<5	—	0.04	25	— ^b	— ^b	3.17
Pure TiO ₂ -H ₂	34	— ^b	— ^b	<5	—	0.04	— ^a	— ^b	— ^b	3.16
Pure TiO ₂ -UV	34	— ^b	— ^b	<5	—	0.04	25	— ^b	— ^b	3.25
TiO ₂ /SiO ₂	8	— ^b	— ^b	100	0.04	0.2	10	— ^b	— ^b	3.25
TiO ₂ /SiO ₂ -H ₂	10	— ^b	— ^b	145	0.06	0.4	— ^a	— ^b	— ^b	3.23
TiO ₂ /SiO ₂ -UV	8	— ^b	— ^b	100	0.04	0.2	10	— ^b	— ^b	3.27
TiO ₂ /SiO ₂ /Pd1	19	— ^c	— ^b	40	0.02	0.1	18	— ^c	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd1-H ₂	25	— ^b	— ^c	60	0.02	0.2	— ^a	— ^b	— ^c	— ^d
TiO ₂ /SiO ₂ /Pd1-UV	19	— ^c	— ^b	40	0.02	0.1	18	— ^c	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd5	11	8	— ^b	45	0.02	0.2	12	8	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd5-H ₂	11	— ^b	17	60	0.02	0.2	— ^a	— ^b	10	— ^d
TiO ₂ /SiO ₂ /Pd5-UV	10	8	— ^b	45	0.02	0.2	12	9	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd12	7	17	— ^b	20	0.01	0.6	10	12	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd12-H ₂	6	— ^b	9	25	0.01	0.4	— ^a	— ^b	11	— ^d
TiO ₂ /SiO ₂ /Pd12-UV	7	17	— ^b	20	0.01	0.6	10	12	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd22	— ^b	25	— ^b	45	0.02	0.4	— ^b	14	— ^b	— ^d
TiO ₂ /SiO ₂ /Pd22-H ₂	— ^b	— ^b	12	100	0.05	0.1	— ^b	— ^b	11	— ^d
TiO ₂ /SiO ₂ /Pd22-UV	— ^b	25	— ^b	45	0.02	0.4	— ^b	14	— ^b	— ^d

—^a, not measured; —^b, not present; —^c, not detectable; —^d, unmeasurable; $d_{\text{XRD Ti}}$, mean diameter of TiO₂ crystallites measured by the Scherrer method; $d_{\text{XRD PdO}}$, mean diameter of PdO crystallites measured by the Scherrer method; $d_{\text{XRD Pd}}$, mean diameter of Pd crystallites measured by the Scherrer method; S_{BET} , specific surface area determined by the BET method; V_{DR} , specific micropore volume determined by Dubinin–Raduskevitch theory; V_{P} , specific liquid volume adsorbed at the saturation pressure of nitrogen; $d_{\text{TEM Ti}}$, mean diameter of TiO₂ nanoparticles measured by TEM; $d_{\text{TEM PdO}}$, mean diameter of PdO nanoparticles measured by TEM; $d_{\text{TEM Pd}}$, mean diameter of Pd nanoparticles measured by TEM; E_{g} , optical band-gap values calculated by using the transformed Kubelka–Munk function.

The decrease of the TiO_2 crystallinity can be explained by the introduction of SiO_2 in the TiO_2 matrix. Indeed, silica was introduced by the EDAS, which was used to disperse Pd in the TiO_2 matrix. The introduction of silica delayed the crystallization of TiO_2 [9,21,22], leading to amorphous TiO_2 for the sample with the highest content of SiO_2 ($\text{TiO}_2/\text{SiO}_2/\text{Pd}22$ sample).

There was no modification of the crystallite sizes, both for TiO_2 and PdO, for all UV-activated samples (Table 2). For the H_2 reduced samples, as explained above, the PdO species are converted to Pd species and the evolution of TiO_2 crystallite follows the evolution of the non-reduced samples.

2.3. Morphology

The nitrogen adsorption–desorption isotherms are presented in Figure 2a for all calcined samples. Pure TiO_2 and all Pd-modified TiO_2 samples were non-porous solids with a very low surface area ($S_{\text{BET}} < 50 \text{ m}^2 \text{ g}^{-1}$, Table 2). Only $\text{TiO}_2/\text{SiO}_2$ sample was a mesoporous solid (type IV isotherm [23]) with a hysteresis between 0.4 and 0.6 in relative pressure. Indeed, the introduction of sol-gel silica increases in the TiO_2 leads to more porous solid [22]. The Pd modification seemed to have low influence on the texture of the samples.

After H_2 reduction (Figure 2b), the S_{BET} increased for all samples (Table 2). Indeed, some remaining organic contaminants could be removed by this thermal treatment, leading to an increase of the microporosity (V_{DR} increased in the majority of the reduced samples, Table 2).

The UV activation had no influence on the textural properties (nitrogen adsorption–desorption isotherms identical to Figure 2a, not shown).

The samples were observed by TEM, as shown in Figure 3, for four representative samples. The estimation of the different sizes of particles (TiO_2 , PdO and Pd) are given in Table 2. The TiO_2 nanoparticles ($d_{\text{TEM Ti}}$) were observed in the size range of 10–18 nm for all samples doped with SiO_2 (Figure 3a–d). For $\text{TiO}_2/\text{SiO}_2/\text{Pd}22$ sample, no TiO_2 crystallite was observed in the matrix, confirming the amorphous $\text{TiO}_2/\text{SiO}_2$ materials. For pure TiO_2 , the size was around 25 nm. This result was coherent with the XRD patterns (Figure 1) where crystallization delays were obtained with SiO_2 doping [9,21]. For the Pd-modified ones, darker nanoparticles highly dispersed in the $\text{TiO}_2/\text{SiO}_2$ matrix were observed which corresponded to the PdO (calcined samples) or Pd (reduced samples) nanoparticles (Figure 3b–d). The Pd species were highly dispersed in the oxide matrix, thanks to the use of EDAS [24,25]. When the amount of Pd became too high, the size of the nanoparticles increased (d_{TEMPdO} and d_{TEMPd} , Table 2). Indeed, the palladium species were closer to each other, leading to sintering when the samples were heated. The size of PdO and metallic Pd was quite similar for all samples, before and after H_2 reduction (Table 2). The UV activation treatment had no influence on the size of the materials.

The XRD crystallite sizes and the nanoparticle sizes estimated with TEM, for the Pd and PdO species, were in the same range. The small differences result from the fact that the crystallites sizes determined by XRD correspond to the average diameter in volume of crystallites. Therefore, XRD gives more statistical significance to large particles, compared to TEM.

Figure 2. Nitrogen adsorption–desorption isotherms of (a) calcined samples and (b) reduced samples.

Figure 3. TEM micrographs of calcined and reduced samples: (a) $\text{TiO}_2/\text{SiO}_2$, (b) $\text{TiO}_2/\text{SiO}_2/\text{Pd5-H}_2$, (c) $\text{TiO}_2/\text{SiO}_2/\text{Pd12-H}_2$ and (d) $\text{TiO}_2/\text{SiO}_2/\text{Pd22}$.

2.4. Optical Properties

The diffuse reflectance UV/Visible spectra of the calcined samples are presented in Figure 4. The pure TiO_2 and $\text{TiO}_2/\text{SiO}_2$ samples had the characteristic spectrum of titania materials with absorption in UV range from around 400 nm. The Pd-modified samples had a larger band of absorption in the UV/Visible region due to the presence of the PdO nanoparticles. As the reduced samples were black, they were not analyzed with this technique. The band gap can be calculated only for the pure TiO_2 and $\text{TiO}_2/\text{SiO}_2$ samples, and is given in Table 2. For both samples, values around 3.2 eV were found in agreement with titania band gap [26].

Figure 4. Normalized diffuse reflectance UV/Visible spectra of all calcined samples: (x) pure TiO_2 , (•) $\text{TiO}_2/\text{SiO}_2$, (■) $\text{TiO}_2/\text{SiO}_2/\text{Pd1}$, (♦) $\text{TiO}_2/\text{SiO}_2/\text{Pd5}$, (▲) $\text{TiO}_2/\text{SiO}_2/\text{Pd12}$ and (♣) $\text{TiO}_2/\text{SiO}_2/\text{Pd22}$.

2.5. Photocatalytic Activity

The evolution of the MB concentration during the photocatalytic experiments is given in Figure 5 for all samples. For each sample, curves obtained with samples treated by both UV and H_2 are also represented. No photolysis of MB was observed under illumination without catalyst (blank experiment). Thanks to the dark experiments, the adsorption of MB on the samples without illumination was deduced from the observed decrease of MB concentration, under illumination, to measure only the MB degradation.

Figure 5. Cont.

Figure 5. Methylene blue (MB) concentration evolution ($[MB]/[MB]_0$ ratio) over time (6 h) for all sample series under UV-A illumination: (a) pure TiO_2 , (b) TiO_2/SiO_2 , (c) $TiO_2/SiO_2/Pd1$, (d) $TiO_2/SiO_2/Pd5$, (e) $TiO_2/SiO_2/Pd12$ and (f) $TiO_2/SiO_2/Pd22$. The contribution of MB adsorption was subtracted for all samples.

For samples without palladium (pure TiO_2 and TiO_2/SiO_2), the H_2 reduction treatment did not seem to have much influence on the percentage of MB degradation, while the activation treatment under UV seemed to improve it slightly for the pure TiO_2 sample, and even more for TiO_2/SiO_2 sample. For samples with low palladium content ($TiO_2/SiO_2/Pd1$ and $TiO_2/SiO_2/Pd5$ samples), the opposite phenomenon was observed: Activation under UV seemed to decrease the photoactivity, while H_2 reduction greatly improved it for the $TiO_2/SiO_2/Pd1$ sample and slightly more for the $TiO_2/SiO_2/Pd5$ and $TiO_2/SiO_2/Pd12$ samples. The $TiO_2/SiO_2/Pd22$ sample had a low photocatalytic activity, regardless of the treatment, due to its amorphous nature (Figure 1). The MB conversion for all the samples is given in Table 3, together with the percentage of MB adsorption.

Table 3. Photocatalytic experiments for MB degradation after 6 h, under UV-A illumination.

Sample	MB Adsorption after 6 h (Blank Test—%) ±5	MB Conversion after 6 h (Blank Deduced—%) ±5	Kinetic Constant <i>k</i> (h ^{−1})
Pure TiO ₂	0	35	0.069
Pure TiO ₂ -H ₂	5	50	0.116
Pure TiO ₂ -UV	0	45	0.092
TiO ₂ /SiO ₂	50	50	0.119
TiO ₂ /SiO ₂ -H ₂	60	40	0.082
TiO ₂ /SiO ₂ -UV	15	85	0.333
TiO ₂ /SiO ₂ /Pd1	0	60	0.147
TiO ₂ /SiO ₂ /Pd1-H ₂	15	75	0.233
TiO ₂ /SiO ₂ /Pd1-UV	0	35	0.070
TiO ₂ /SiO ₂ /Pd5	0	55	0.131
TiO ₂ /SiO ₂ /Pd5-H ₂	5	60	0.147
TiO ₂ /SiO ₂ /Pd5-UV	0	35	0.069
TiO ₂ /SiO ₂ /Pd12	20	25	0.039
TiO ₂ /SiO ₂ /Pd12-H ₂	15	30	0.059
TiO ₂ /SiO ₂ /Pd12-UV	15	25	0.050
TiO ₂ /SiO ₂ /Pd22	10	15	0.030
TiO ₂ /SiO ₂ /Pd22-H ₂	10	0	0.005
TiO ₂ /SiO ₂ /Pd22-UV	10	0	0.000

The pure TiO₂ samples had very low MB adsorption by opposition to TiO₂/SiO₂ samples. This difference could come from the textural properties of the TiO₂/SiO₂ samples, which presented higher S_{BET} values, as compared to pure TiO₂ samples (< 5 vs. 100–145 m² g^{−1}, Table 2). The Pd-modified samples presented S_{BET} values between pure TiO₂ and TiO₂/SiO₂ samples (~ 40 m² g^{−1}, Table 2), so their MB adsorption behavior was intermediate with a small adsorption (~10%), except for the TiO₂/SiO₂/Pd12 series, which adsorbed around 20%. The MB adsorption was therefore directly linked to the surface properties of the samples [27].

On the one hand, the photoactivity of TiO₂ was increased by the introduction of a small amount of silica into its network [9,28]. The TiO₂/SiO₂/Pd1 sample degraded 60% of MB against 35% for the pure TiO₂ sample (Table 3). On the other hand, when the quantity of silica introduced became too large, the crystallization of TiO₂ in its anatase form was delayed [21]. Thus, in the samples containing a large amount of palladium (TiO₂/SiO₂/Pd5, TiO₂/SiO₂/Pd12 and TiO₂/SiO₂/Pd22), and therefore a large amount of silica, photocatalytic efficiency decreased: We obtained 60% MB degradation for the TiO₂/SiO₂/Pd1 sample against 15% MB degradation for the TiO₂/SiO₂/Pd22 sample (Table 3). In all cases, except for the TiO₂/SiO₂/Pd22 sample, which contains only amorphous TiO₂, the H₂ reduction treatment had the effect of increasing the photocatalytic activity. Indeed, H₂ reduction allowed us to decrease the rate of recombination of the “electron–hole” pairs present in TiO₂ via two distinct phenomena: (i) The first was the possible creation of oxygen vacancies on the surface of TiO₂ by reduction of Ti⁴⁺ ions to Ti³⁺ [29,30], and (ii) the second was the reduction of PdO particles to metallic palladium acting as an electron trap and thus limiting the recombination’s of the “electron–hole” pairs [31,32]. Activation processing under UV, on the other hand, has little influence on the photocatalytic activity of calcined samples.

The optimal sample is TiO₂/SiO₂/Pd1, which combines the advantages of both additives (SiO₂ and Pd) with a weight loading percentage of 1% for each. Indeed, the introduction of SiO₂ allows us to increase the specific surface area, as compared to pure TiO₂ (Table 2), while the crystallinity of anatase is maintained (Figure 1). Moreover, Pd introduction allows an absorption in the visible range (Figure 4).

Assuming that the photocatalyst is homogeneously dispersed in a perfectly agitated medium, the derivative of the material balance in a semi-continuous reactor can be expressed as follows [33]:

$$\frac{dC}{dt} = -r_V \frac{m_{cata,0}}{V_0} \quad (1)$$

where C is the concentration of methylene blue in the solution (kmol/m^3), t is the time (h), r_V is the specific rate of the degradation of methylene blue ($\text{kmol}/\text{kg}_{\text{catalyst}} \cdot \text{h}$), $m_{cata,0}$ is the initial catalyst mass homogeneously dispersed in the solution (kg of catalyst) and V_0 is the initial solution volume (m^3).

In general, the degradation curves of methylene blue in photocatalysis had an exponential appearance. The degradation kinetics of methylene blue could then be modeled by assuming that the reaction was of order 1 under the operating conditions used. The kinetic equation could thus be expressed as follows:

$$\frac{dC}{dt} = -k \times C \times \frac{m_{cata,0}}{V_0} \quad (2)$$

where k is the kinetic constant ($\text{m}^3/\text{kg}_{\text{catalyst}} \cdot \text{h}$).

As part of this work, the photocatalytic activity of the samples was evaluated in media containing 50 mg of catalysts dispersed in 50 mL of MB solution at $20 \mu\text{mol}/\text{L}$ concentration. The $m_{cata,0}/V_0$ ratio was thus equal to 1. The kinetic equation could then be developed as follows:

$$\frac{dC}{dt} = -k \times C \quad (3)$$

$$\frac{dC}{C} = -k \times dt \quad (4)$$

$$\ln\left(\frac{C}{C_0}\right) = -k \times t \quad (5)$$

$$C = C_0 \times e^{-k \times t} \quad (6)$$

where C_0 is the initial MB concentration in the solution (kmol/m^3).

Figure 6 compares the theoretical kinetic model to the experimental values for the $\text{TiO}_2/\text{SiO}_2/\text{Pd1}$ sample. The two curves superimposed perfectly, justifying the hypothesis of an order 1 kinetics of degradation.

Figure 6. Kinetic analysis of the MB degradation for $\text{TiO}_2/\text{SiO}_2/\text{Pd1}$ sample: (—) theoretical evolution and (■) experimental points.

Equation (6) is commonly found in the literature [34,35] for photocatalytic process. Knowing the evolution over time of the concentration of methylene blue in the solutions, the kinetic constant, k , could be calculated for all samples. The values obtained are listed in Table 3.

The evolution of the kinetic constants values corresponds to the observations made previously. Indeed, the kinetic constant, k , was increased by the introduction of a small quantity of silica into the TiO_2 network: The $\text{TiO}_2/\text{SiO}_2/\text{Pd1}$ sample had a kinetic constant equal to 0.147 h^{-1} versus 0.069 h^{-1} for the pure TiO_2 sample (Table 3). On the other hand, when the quantity of silica introduced became too large, the crystallization of TiO_2 in its anatase form was delayed [9,21]. Thus, in samples containing a large amount of palladium ($\text{TiO}_2/\text{SiO}_2/\text{Pd5}$, $\text{TiO}_2/\text{SiO}_2/\text{Pd12}$ and $\text{TiO}_2/\text{SiO}_2/\text{Pd22}$ samples), and therefore a large quantity of silica, the kinetic constant decreased, i.e., 0.147 h^{-1} for the $\text{TiO}_2/\text{SiO}_2/\text{Pd1}$ sample, as compared to 0.030 h^{-1} for the $\text{TiO}_2/\text{SiO}_2/\text{Pd22}$ sample. In all cases (except for the sample $\text{TiO}_2/\text{SiO}_2/\text{Pd22}$, which contained only amorphous TiO_2), the H_2 reduction treatment had the consequence of increasing the kinetic constant of the samples. Activation processing under UV had little influence on the kinetic constant of the calcined samples.

With the combination of Pd/SiO_2 modification and the reduction of the photocatalyst, the photoactivity for the degradation of MB can be greatly increase compared to the pure material. Moreover, the use of modified silicon alkoxide allowed a good distribution of the metal in the titania matrix. This cogelation synthesis method appears as an easy process for homogeneous doping in titania photocatalysts.

3. Materials and Methods

3.1. Photocatalyst Synthesis

The pure and modified TiO_2 samples were prepared by a sol-gel cogelation method inspired from Reference [21]. The steps were the following: Palladium acetylacetonate ($\text{Pd}(\text{acac})_2$, Sigma Aldrich, 99%) was dissolved in 2-methoxyethanol (Sigma Aldrich, 99.8%); then 3-(2-aminoethyl) aminopropyl]trimethoxysilane (EDAS, Merck, 97%) was added to the solution and stirred for 1 h, for complexation. The resulting solution became yellow. Then titanium isopropoxide (TIPT, Sigma Aldrich, 98%) was added, under stirring. In a second vial, distilled water was mixed with 2-methoxyethanol. This second solution was added, dropwise, to the first one, in 5 min, under stirring. The resulting suspension was placed in an oven, for gelation, at 80°C , for 72 h. Then, the gel was dried under vacuum at 80°C for 48 h, until the pressure fell at 1000 Pa. At this pressure, the temperature was increased to 150°C for 24 h. The dried powder was crushed with a mortar and calcined at 550°C for 5 h.

Six samples were prepared with the following protocol: one pure TiO_2 (without $\text{Pd}(\text{acac})_2$ and EDAS), one $\text{TiO}_2/\text{SiO}_2$ (without $\text{Pd}(\text{acac})_2$) and four $\text{TiO}_2/\text{SiO}_2/\text{Pd}$ samples (1, 5, 12 and 22 theoretical wt.% of Pd). Table 1 summarizes the amounts of reactants for each sample: the weight amount of SiO_2 is in the same range as Pd in all cases.

3.2. Activation Treatments

In order to increase the photoactivity of the samples, they underwent two different activation treatments after calcination: UV activation and H_2 reduction.

Concerning the UV activation, the calcined powder was mixed in 50 mL of distilled water, in a Petri dish that was illuminated for 24 h, under stirring, at 25°C . The lamp (Osram Sylvania, Blacklight-Bleu Lamp, F18W/BLB-T8) emitted UV-A radiation considered to be monochromatic, with a wavelength of 365 nm and an intensity of $1.2 \text{ mW}/\text{cm}^2$. After the UV treatment, the photocatalyst was filtered and dried for 24 h, at 120°C . The UV-treated samples are denoted with “- UV”.

Concerning the H_2 reduction, the calcined powder was heated under H_2 , for 5 h, at 450°C . The reduced samples are denoted with “- H_2 ”.

3.3. Characterizations

The nitrogen adsorption–desorption isotherms were obtained thanks to a Fisons Sorptomatic 1900 device. The specific surface area, S_{BET} , the microporous volume, V_{DR} , and the porous volume, V_{P} , were obtained from the isotherms, thanks to BET and Dubinin–Raduskevich theories [23].

The crystallographic phases were determined with a Siemens D5000 device, by measurements between 20° and 70° , with a step size of 0.05° and a step time of 5 s. The crystallite size was estimated thanks to the Scherrer equation [36].

The atomic composition of the samples was determined by inductively coupled plasma atomic emission spectrometry (ICP-AES), with an ICAP 6500 device from Thermo Scientific. Before introducing the sample in the device, it was mineralized as follows: 75 mg of sample was mixed with 1.6 g of $\text{Li}_2\text{B}_4\text{O}_7$ and 0.4 g of Li_3BO_3 in a graphitic crucible. Then it was heated to 1000°C for 5 min. After cooling, the pearl was dissolved, under stirring, in hot HNO_3 2 M.

The sample suspensions were observed with transmission electron microscopy (TEM), using a Phillips CM 100 device (accelerating voltage 200 kV). The average nanoparticle size was estimated by measuring approximatively 100 nanoparticles with the TEM software.

Optical properties of the sample materials in the range of 200–800 nm region were obtained with diffuse reflectance measurements on a Varian Cary 500 UV–Vis–NIR spectrophotometer equipped with an integrating sphere (Varian External DRA-2500), and using BaSO_4 as reference. The absorbance spectra were transformed by using the Kubelka–Munk function [37–39] to produce a signal, normalized for comparison between samples, allowing us to calculate the band gap (E_g). The details of this treatment method are widely described elsewhere [40–42].

3.4. Photocatalytic Activity on Methylene Blue Degradation

The photoactivity of the samples was determined by following the methylene blue (MB) degradation, under UV-A illumination. The lamp (Osram Sylvania, Blacklight-Bleu Lamp, F18W/BLB-T8) emitted UV-A radiation considered as monochromatic, with a wavelength of 365 nm and an intensity of 1.2 mW/cm^2 (same as for the UV activation, Section 3.2). A scheme of similar installation was presented in Reference [43].

Then, 50 mL of MB solution with a concentration of $20\text{ }\mu\text{mol/L}$ was mixed with the photocatalyst (1 g/L). The suspension was illuminated for 6 h, at a constant temperature of 25°C , under UV-A illumination. The concentration of MB was followed by recording UV/VIS spectrum between 300 and 800 nm with GENESYS 105 UV-VIS spectrophotometer from Thermo Scientific. A calibration curve was made to link the absorbance to the concentration. Aliquots were taken every 30 min during the first 2 h of the experiment, and then each hour, until a total of 6 h. Each sample was tested 3 times to obtain mean degradation values; the variation on the sample concentration was around 5%. Blank experiment without photocatalyst was also performed to ensure that MB did not undergo photobleaching under UV-A illumination. Dark tests without illumination were performed to assess the MB adsorption on the samples.

4. Conclusions

In this study, TiO_2 photocatalysts were cogelled with silicon alkoxide, in order to finely disperse Pd species in TiO_2 matrix and dope them with SiO_2 ; these modifications were made in one step during the synthesis. This Pd modification was made in order to increase the photoefficiency on methylene blue degradation. Two activation treatments were also performed in the same goal: UV activation and H_2 reduction. A pure and a mixed $\text{TiO}_2/\text{SiO}_2$ sample were also synthesized as reference.

Results showed that the crystallographic structure of TiO_2 is the anatase phase and that Pd is present either in its oxidized form after the calcination stage, or in its reduced form after the reduction stage. The introduction of SiO_2 delayed the crystallization of TiO_2 , leading to amorphous TiO_2 material when the highest Pd loading (22 wt.%) was produced. The specific surface area of the samples,

determined by nitrogen adsorption–desorption, varied from 5 to 145 m² g^{−1}, with the highest S_{BET} value obtained after H₂ reduction treatment. Transmission electron microscopy allowed us to observe the homogeneous dispersion and nanometric size of Pd particles in the reduced samples. The size of Pd crystallites increased with the Pd amount. The band gap measured by UV/Visible diffuse reflectance spectroscopy corresponded to that of the TiO₂ anatase, approximately 3.2 eV for the pure and SiO₂ doped TiO₂. The band gaps for Pd-modified samples were not determined due to the dark color of the samples and the high absorption in the UV/Visible range. The UV activation did not modify the physicochemical properties of the samples, and it had little effect on photocatalytic activity of calcined samples.

The photoactivity of TiO₂ was increased by the introduction of a small amount of silica into its network. However, when the quantity of silica introduced became too large, the crystallization of TiO₂ in its anatase form was delayed, and the photocatalytic activity of samples decreased. The introduction of Pd in the samples also increased the photoactivity. Moreover, the H₂ reduction treatment further increased the photocatalytic activity of the samples, because this treatment allowed us to decrease the rate of electron–hole recombination in TiO₂ via two distinct phenomena: (i) the creation of oxygen vacancies on the surface of TiO₂ by reduction of Ti⁴⁺ ions to Ti³⁺; (ii) the reduction of PdO particles to palladium metallic nanoparticles acting as an electron trap and thus limiting the electron–hole recombination. The optimal loading is 1 wt.% for each (SiO₂ and Pd) additive.

Supplementary Materials: The following are available online at <http://www.mdpi.com/2073-4344/10/10/1184/s1>, Figure S1. Pictures of the calcined samples: from left to right: pure TiO₂, TiO₂/SiO₂, TiO₂/SiO₂/Pd1, TiO₂/SiO₂/Pd5, TiO₂/SiO₂/Pd12 and TiO₂/SiO₂/Pd22.

Author Contributions: Conceptualization, methodology, investigation, analysis and writing, V.S., L.T., J.G.M. and S.D.L.; writing—original draft preparation, J.G.M., S.H. and S.D.L.; supervision, funding acquisition and project administration, S.H. and S.D.L. All the authors corrected the paper before submission and during the revision process. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Acknowledgments: S.D.L. thanks the Belgian National Funds for Scientific Research (F.R.S.-FNRS) for her Associate Researcher position. The authors acknowledge the Ministère de la Région Wallonne Direction Générale des Technologies, de la Recherche et de l’Energie (DGO6), the Fonds de Bay and the Fonds de Recherche Fondamentale et Collective for financial support. J.G.M. and S.H. also thank Innoviris Brussels for financial support through the Bridge project—COLORES.

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

References

1. Pignatello, J.J.; Oliveros, E.; MacKay, A. Advanced oxidation processes for organic contaminant destruction based on the fenton reaction and related chemistry. *Crit. Rev. Environ. Sci. Technol.* **2006**, *36*, 1–84. [\[CrossRef\]](#)
2. Kuyukina, M.S.; Ivshina, I.B. Application of *Rhodococcus* in Bioremediation of Contaminated Environments. In *Biology of Rhodococcus*; Alvarez, H., Ed.; Springer: Berlin/Heidelberg, Germany, 2010; pp. 231–262. ISBN 9783642129377.
3. Linsebigler, A.L.; Lu, G.; Yates, J.T. Photocatalysis on TiO₂ Surfaces: Principles, Mechanisms, and Selected Results. *Chem. Rev.* **1995**, *95*, 735–758. [\[CrossRef\]](#)
4. Pelaez, M.; Nolan, N.T.; Pillai, S.C.; Seery, M.K.; Falaras, P.; Kontos, A.G.; Dunlop, P.S.M.; Hamilton, J.W.J.; Byrne, J.A.; O’Shea, K.; et al. A review on the visible light active titanium dioxide photocatalysts for environmental applications. *Appl. Catal. B Environ.* **2012**, *125*, 331–349. [\[CrossRef\]](#)
5. Oseghe, E.O.; Ofomaja, A.E. Study on light emission diode/carbon modified TiO₂ system for tetracycline hydrochloride degradation. *J. Photochem. Photobiol. A Chem.* **2018**, *360*, 242–248. [\[CrossRef\]](#)
6. Mahy, J.G.; Paez, C.A.; Carcel, C.; Bied, C.; Tatton, A.S.; Damblon, C.; Heinrichs, B.; Wong Chi Man, M.; Lambert, S.D. Porphyrin-based hybrid silica-titania as a visible-light photocatalyst. *J. Photochem. Photobiol. A Chem.* **2019**, *373*, 66–76. [\[CrossRef\]](#)
7. Banerjee, S.; Dionysiou, D.D.; Pillai, S.C. Self-cleaning applications of TiO₂ by photo-induced hydrophilicity and photocatalysis. *Appl. Catal. B Environ.* **2015**, *176*, 396–428. [\[CrossRef\]](#)

8. Espino-Estévez, M.R.; Fernández-Rodríguez, C.; González-Díaz, O.M. Effect of TiO₂—Pd and TiO₂—Ag on the photocatalytic oxidation of diclofenac, isoproturon and phenol. *Chem. Eng. J.* **2016**, *298*, 82–95. [[CrossRef](#)]
9. Léonard, G.L.-M.; Malengreux, C.M.; Mélotte, Q.; Lambert, S.D.; Bruneel, E.; Van Driessche, I.; Heinrichs, B. Doped sol-gel films vs. powders TiO₂: On the positive effect induced by the presence of a substrate. *J. Environ. Chem. Eng.* **2016**, *4*, 449–459. [[CrossRef](#)]
10. Tunc, I. The effect of the presence of Ag nanoparticles on the photocatalytic degradation of oxalic acid adsorbed on TiO₂ nanoparticles monitored by ATR-FTIR. *Mater. Chem. Phys.* **2014**, *144*, 444–450. [[CrossRef](#)]
11. Bodson, C.J.; Heinrichs, B.; Tasseroul, L.; Bied, C.; Mahy, J.G.; Wong Chi Man, M.; Lambert, S.D. Efficient P- and Ag-doped titania for the photocatalytic degradation of waste water organic pollutants. *J. Alloy. Compd.* **2016**, *682*, 144–153. [[CrossRef](#)]
12. Vaiano, V.; Iervolino, G.; Sannino, D.; Murcia, J.J.; Hidalgo, M.C.; Ciambelli, P.; Navío, J.A. Photocatalytic removal of patent blue V dye on Au—TiO₂ and Pt—TiO₂ catalysts. *Appl. Catal. B Environ.* **2016**, *188*, 134–146. [[CrossRef](#)]
13. Borzyszkowska, A.F.; Stepnowski, P.; Ofiarska, A.; Pieczynska, A.; Siedlecka, E.M. Pt—TiO₂-assisted photocatalytic degradation of the cytostatic drugs ifosfamide and cyclophosphamide under artificial sunlight. *Chem. Eng. J.* **2016**, *285*, 417–427.
14. Abdelaal, M.Y.; Mohamed, R.M. Novel Pd/TiO₂ nanocomposite prepared by modified sol-gel method for photocatalytic degradation of methylene blue dye under visible light irradiation. *J. Alloy. Compd.* **2013**, *576*, 201–207. [[CrossRef](#)]
15. Léonard, G.L.-M.; Pàez, C.A.; Ramírez, A.E.; Mahy, J.G.; Heinrichs, B. Interactions between Zn²⁺ or ZnO with TiO₂ to produce an efficient photocatalytic, superhydrophilic and aesthetic glass. *J. Photochem. Photobiol. A Chem.* **2018**, *350*, 32–43. [[CrossRef](#)]
16. Mahy, J.G.; Lambert, S.D.; Tilkin, R.G.; Poelman, D.; Wolfs, C.; Devred, F.; Gaigneaux, E.M.; Douven, S. Ambient temperature ZrO₂-doped TiO₂ crystalline photocatalysts: Highly efficient powders and films for water depollution. *Mater. Today Energy* **2019**, *13*, 312–322. [[CrossRef](#)]
17. Chen, Q.; Shi, W.; Xu, Y.; Wu, D.; Sun, Y.; Si, A. Visible-light-responsive Ag—Si codoped anatase TiO₂ photocatalyst with enhanced thermal stability. *Mater. Chem. Phys.* **2011**, *125*, 825–832. [[CrossRef](#)]
18. Pirard, S.L.; Mahy, J.G.; Pirard, J.-P.; Heinrichs, B.; Raskinet, L.; Lambert, S.D. Development by the sol-gel process of highly dispersed Ni—Cu/SiO₂ xerogel catalysts for selective 1,2-dichloroethane hydrodechlorination into ethylene. *Microporous Mesoporous Mater.* **2015**, *209*, 197–207. [[CrossRef](#)]
19. Belet, A.; Wolfs, C.; Mahy, J.G.; Poelman, D.; Vreuls, C. Sol-Gel Syntheses of Photocatalysts for the Removal of Pharmaceutical Products in Water. *Nanomaterials* **2019**, *9*, 126. [[CrossRef](#)]
20. Yang, J.; Xu, X.; Liu, Y.; Gao, Y.; Chen, H.; Li, H. Preparation of SiO₂@TiO₂ composite nanosheets and their application in photocatalytic degradation of malachite green at emulsion interface. *Colloids Surf. A Physicochem. Eng. Asp.* **2019**, *582*, 123858. [[CrossRef](#)]
21. Braconnier, B.; Páez, C.A.; Lambert, S.; Alié, C.; Henrist, C.; Poelman, D.; Pirard, J.P.; Cloots, R.; Heinrichs, B. Ag- and SiO₂-doped porous TiO₂ with enhanced thermal stability. *Microporous Mesoporous Mater.* **2009**, *122*, 247–254. [[CrossRef](#)]
22. Bodson, C.J.; Lambert, S.D.; Alié, C.; Cattoën, X.; Pirard, J.; Bied, C.; Wong Chi Man, M.; Heinrichs, B. Effects of additives and solvents on the gel formation rate and on the texture of P- and Si-doped TiO₂ materials. *Microporous Mesoporous Mater.* **2010**, *134*, 157–164. [[CrossRef](#)]
23. Lecloux, A.J. Texture of Catalysts. In *Catalysis: Science and Technology*; Anderson, J.R., Boudart, M., Eds.; Springer: Berlin/Heidelberg, Germany, 1981; Volume 2, pp. 171–230.
24. Mahy, J.G.; Claude, V.; Sacco, L.; Lambert, S.D. Ethylene polymerization and hydrodechlorination of 1,2-dichloroethane mediated by nickel(II) covalently anchored to silica xerogels. *J. Sol-Gel Sci. Technol.* **2017**, *81*, 59–68. [[CrossRef](#)]
25. Lambert, S.; Cellier, C.; Grange, P.; Pirard, J.P.; Heinrichs, B. Synthesis of Pd/SiO₂, Ag/SiO₂, and Cu/SiO₂ cogelled xerogel catalysts: Study of metal dispersion and catalytic activity. *J. Catal.* **2004**, *221*, 335–346. [[CrossRef](#)]
26. Khaki, M.R.D.; Shafeeyan, M.S.; Raman, A.A.A.; Daud, W.M.A.W. Application of doped photocatalysts for organic pollutant degradation—A review. *J. Environ. Manag.* **2017**, *198*, 78–94. [[CrossRef](#)]
27. Yukselen, Y.; Kaya, A. Suitability of the methylene blue test for surface area, cation exchange capacity and swell potential determination of clayey soils. *Eng. Geol.* **2008**, *102*, 38–45. [[CrossRef](#)]

28. Calleja, G.; Serrano, D.P.; Sanz, R.; Pizarro, P. Mesostructured SiO₂-doped TiO₂ with enhanced thermal stability prepared by a soft-templating sol-gel route. *Microporous Mesoporous Mater.* **2008**, *111*, 429–440. [CrossRef]
29. Zhang, Z.; Long, J.; Xie, X.; Zhuang, H.; Zhou, Y.; Lin, H.; Yuan, R.; Dai, W.; Ding, Z.; Wang, X.; et al. Controlling the synergistic effect of oxygen vacancies and N dopants to enhance photocatalytic activity of N-doped TiO₂ by H₂ reduction. *Appl. Catal. A Gen.* **2012**, *425–426*, 117–124. [CrossRef]
30. Páez, C.A.; Lambert, S.D.; Poelman, D.; Pirard, J.P.; Heinrichs, B. Improvement in the methylene blue adsorption capacity and photocatalytic activity of H₂-reduced rutile-TiO₂ caused by Ni(II)porphyrin preadsorption. *Appl. Catal. B Environ.* **2011**, *106*, 220–227.
31. Liu, H.; Ma, H.T.; Li, X.Z.; Li, W.Z.; Wu, M.; Bao, X.H. The enhancement of TiO₂ photocatalytic activity by hydrogen thermal treatment. *Chemosphere* **2003**, *50*, 39–46. [CrossRef]
32. Tong, H.-X.; Chen, Q.-Y.; Yin, Z.-L.; Hu, H.-P.; Wu, D.-X.; Yang, Y.-H. Preparation, characterization and photo-catalytic behavior of WO₃-TiO₂ catalysts with oxygen vacancies. *Trans. Nonferrous Met. Soc. China* **2009**, *19*, 1483–1488. [CrossRef]
33. Pirard, S.L.; Malengreux, C.M.; Toye, D.; Heinrichs, B. How to correctly determine the kinetics of a photocatalytic degradation reaction? *Chem. Eng. J.* **2014**, *249*, 1–5. [CrossRef]
34. Arabatzis, I.M.; Stergiopoulos, T.; Andreeva, D.; Kitova, S.; Neophytides, S.G.; Falaras, P. Characterization and photocatalytic activity of Au/TiO₂ thin films for azo-dye degradation. *J. Catal.* **2003**, *220*, 127–135. [CrossRef]
35. Fisher, M.B.; Keane, D.A.; Fernández-Ibáñez, P.; Colreavy, J.; Hinder, S.J.; McGuigan, K.G.; Pillai, S.C. Nitrogen and copper doped solar light active TiO₂ photocatalysts for water decontamination. *Appl. Catal. B Environ.* **2013**, *130–131*, 8–13. [CrossRef]
36. Patterson, A.L. The Scherrer Formula for X-Ray Particle Size Determination. *Phys. Rev.* **1939**, *56*, 978–982. [CrossRef]
37. Malengreux, C.M.; Douven, S.; Poelman, D.; Heinrichs, B.; Bartlett, J.R. An ambient temperature aqueous sol-gel processing of efficient nanocrystalline doped TiO₂-based photocatalysts for the degradation of organic pollutants. *J. Sol-Gel Sci. Technol.* **2014**, *71*, 557–570. [CrossRef]
38. Kubelka, P. Ein Beitrag zur Optik der Farbanstriche. *Z. Tech. Phys.* **1931**, *12*, 593–601.
39. Kubelka, P. New contributions to the optics of intensely light-scattering materials. *J. Opt. Soc. Am.* **1948**, *38*, 448–457. [CrossRef]
40. Mahy, J.G.; Lambert, S.D.; Léonard, G.L.-M.; Zubiaur, A.; Olu, P.-Y.; Mahmoud, A.; Boschini, F.; Heinrichs, B. Towards a large scale aqueous sol-gel synthesis of doped TiO₂: Study of various metallic dopings for the photocatalytic degradation of p-nitrophenol. *J. Photochem. Photobiol. A Chem.* **2016**, *329*, 189–202. [CrossRef]
41. Mahy, J.G.; Cerfontaine, V.; Poelman, D.; Devred, F.; Gaigneaux, E.M.; Heinrichs, B.; Lambert, S.D. Highly efficient low-temperature N-doped TiO₂ catalysts for visible light photocatalytic applications. *Materials* **2018**, *11*, 584. [CrossRef]
42. Malengreux, C.M.; Pirard, S.L.; Léonard, G.; Mahy, J.G.; Herlitschke, M.; Klobes, B.; Hermann, R.; Heinrichs, B.; Bartlett, J.R. Study of the photocatalytic activity of Fe³⁺, Cr³⁺, La³⁺ and Eu³⁺ single-doped and co-doped TiO₂ catalysts produced by aqueous sol-gel processing. *J. Alloy. Compd.* **2017**, *691*, 726–738. [CrossRef]
43. Tasseroul, L.; Pirard, S.L.; Lambert, S.D.; Páez, C.A.; Poelman, D.; Pirard, J.P.; Heinrichs, B. Kinetic study of p-nitrophenol photodegradation with modified TiO₂ xerogels. *Chem. Eng. J.* **2012**, *191*, 441–450. [CrossRef]

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

© 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/>).