Photocatalytic Solar Light H₂ Production by Aqueous Glucose Reforming


Abstract: A series of tungsten and nitrogen doped Pt-TiO₂ samples were prepared with the aim to extend the TiO₂ absorption to the visible light region and to enhance the separation efficiency of the photogenerated electron/hole pairs. The physicochemical features of the powders were characterized by X-ray diffraction (XRD), UV/Vis reflectance spectra, specific surface area (SSA) determinations, and transmission electron microscopy (TEM) analyses. The influence of the presence of different doping agents was evaluated, under anaerobic conditions, in the aqueous photo-reforming of glucose to form H₂ at ambient pressure and temperature under a halogen lamp or natural solar light irradiation. Arabinose, erythrose, and formic acid were the main glucose partial oxidation products observed in the liquid phase, whilst H₂ and CO₂ were measured in the gaseous phase. The highest H₂ production was observed in the presence of the Pt-TiO₂−Wₐ₂Sₕ sample.

1. Introduction

In the last decades, there has been a growing interest in renewable energy sources, due to both the almost exhaustion of the fossil fuels currently used, and the growth of environmental pollution with damaging effects for the ecosystem and health of people. The use of renewable sources can efficiently help to solve these issues and, among them, hydrogen is a clean energy carrier, because its combustion with oxygen produces only water.[1–3]

Nowadays, the dominant industrial processes for hydrogen production are steam reforming from hydrocarbons,[11] biomass conversion processes (such as direct combustion, gasification, pyrolysis)[4–6] and water splitting by means of different technologies: electrolysis, radiolysis, thermoysis, photobiology and photocatalysis.[7–8] Most of these methods are not economically sustainable as they require large equipment, high temperature and pressure, and significant energy consumption.[9]

To ensure that this new energy carrier is effectively “green”, hydrogen production has to be as clean as possible. In this scenario heterogeneous photocatalysis, under ambient conditions and in the presence of green solvents and sunlight irradiation, represents one of the cleanest way to directly split water into hydrogen and oxygen.[10–12] Unfortunately, the direct water splitting is a very hard and disadvantageous process by a thermodynamic and kinetic point of view. Biomass conversion is less energy demanding than water splitting. The use of glucose, glycerol, lignin, etc., as sacrificial agents, can reduce the electron/hole recombination rate and favour selectively the synthesis of high value-added products.[6,13–22] There are still only a relative small number of materials that can efficiently photoreform biomass, and works dedicated to improving the efficiency of photocatalysts, especially under natural solar light irradiation, are useful. TiO₂ is the most employed photocatalyst for this process because of its low cost, chemical and photochemical stabilities, and ease of preparation.[16,23]

TiO₂ photoactivation presents some weaknesses due to its high electron/hole recombination rate and the need of UV light.[24] Many approaches have been explored to improve the TiO₂ efficiency and to better exploit the solar radiation, as doping with metal[25–27] and non-metal species,[28–30] co-doping,[31–33] surface modification by organic species sensitization,[34,35] coupling of different semiconductors.[36–39]

Among the nonmetallic elements, since the work of Asahi et al.[40] a great attention has been addressed to N doped TiO₂ because it is reported that the band gap of TiO₂ becomes smaller, due to formation of a mid-gap energy state above the valence band[41–43] and presence of oxygen vacancies generating color centers.[44] N-TiO₂ samples have been employed also for glucose photo-reforming.[45–46] Ilie et al.[45] compared the photocactivity of a series of Au-TiO₂ samples dispersed on SiO₂ and doped with metal (Mn, Cr, V) or non-metal (N or S) species towards the H₂ production from aqueous sugar solution. The best results were obtained in the presence of non-metals and were attributed to a lower tendency to form recombination centers with respect to samples doped with metals.

As far as the metallic species are concerned, tungsten has attracted the attention of many researchers because the pres-
ence of WO$_3$ in the TiO$_2$ lattice can enhance the separation efficiency of the photogenerated charges, while the presence of cations (W$^{6+}$ and/or W$^{4+}$) substituting Ti$^{4+}$ ions, can modify the TiO$_2$ electronic structure. Due to the different behaviour of metals and non-metals, co-doping was used as a strategy to enhance the photocatalytic activity of TiO$_2$. Various N, W co-doped TiO$_2$ systems have been prepared and tested under solar light irradiation for oxidation reactions. In those cases, isolated N 2$p$ states are located above the valence band edge whilst W 5$d$ states below the conduction band edge.

The obtainment of high value-added chemicals from biomass derived compounds has been the subject of intense research in recent years. In this context the use of glucose appears interesting because it is an abundant and cheap monosaccharide.

The aim of the work reported in this paper was to synthesize Pt-TiO$_2$-rutile based materials doped with N or W species which were used under ultraviolet (UV), halogen lamp (HL) or natural solar light (SOL) irradiations to obtain H$_2$ by photo-reforming of a glucose aqueous solution. Rutile TiO$_2$ polymorph was chosen because it has been reported to show a photocatalytic H$_2$ production efficiency higher than that of the anatase phase, under similar experimental conditions.

2. Results and Discussion

2.1 Characterization of the Photocatalysts

The XRD patterns of the pristine and doped samples are shown in Figure 1. The bare TiO$_2$ consisted of pure rutile; the presence of nitrogen did not alter the phase composition (Figure 1A), whilst the appearance of anatase was observed in the presence of tungsten, and it increased with the amount of the dopant (Figure 1B), according to what previously observed. Broad peaks were obtained, suggesting a poor crystallinity of the samples due to the low preparation temperature. The absence of peaks related to N and W species can be ascribed to the low amount of dopants and to a high degree of dispersion of the loaded species on the TiO$_2$ surface. The crystallites size ranged between 5.6 and 12.5 nm (Table 1) and the highest values were measured for the samples containing the highest percentages of dopant species. The slight broadening of the peaks after doping may be due to the formation of structural defects in the presence of foreign ions.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Phase</th>
<th>Φ [nm]</th>
<th>$E_g$ [eV]</th>
<th>SSA [m$^2$ g$^{-1}$]</th>
</tr>
</thead>
<tbody>
<tr>
<td>TiO$_2$</td>
<td>R</td>
<td>5.6</td>
<td>2.98</td>
<td>21</td>
</tr>
<tr>
<td>Pt-TiO$_2$</td>
<td>R</td>
<td>7.7</td>
<td>2.90 (2.45)</td>
<td>7</td>
</tr>
<tr>
<td>Pt-TiO$_2$-N0.5</td>
<td>R</td>
<td>7.9</td>
<td>2.86 (2.47)</td>
<td>1</td>
</tr>
<tr>
<td>Pt-TiO$_2$-N1</td>
<td>R</td>
<td>12.5</td>
<td>2.74 (2.41)</td>
<td>1</td>
</tr>
<tr>
<td>Pt-TiO$_2$-N2</td>
<td>R</td>
<td>12.1</td>
<td>2.72 (2.42)</td>
<td>1</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W0.25</td>
<td>R</td>
<td>8.8</td>
<td>3.26 (2.95)</td>
<td>66</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W0.5</td>
<td>R+A</td>
<td>7.7</td>
<td>3.16 (2.96)</td>
<td>64</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W1</td>
<td>R+A</td>
<td>12.4</td>
<td>2.92 (2.57)</td>
<td>18</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W2</td>
<td>A+R</td>
<td>10.3</td>
<td>2.86 (2.49)</td>
<td>28</td>
</tr>
<tr>
<td>Pt-TiO$_2$-N0.5W0.5</td>
<td>R</td>
<td>7.5</td>
<td>2.88 (2.48)</td>
<td>3.4</td>
</tr>
</tbody>
</table>

The bare TiO$_2$ exhibited a specific surface area (SSA) of 21 m$^2$ g$^{-1}$, that decreased in the presence of Pt and nitrogen, whilst an increase was observed when W was loaded. The samples containing N showed very low values of SSA (about 1 m$^2$ g$^{-1}$), and when tungsten was present, the increase was

Figure 1. XRD patterns of N-doped (A) and W-doped (B) TiO$_2$ samples. A = Anatase, R = Rutile.
the highest in the presence of the lowest amounts of metal (0.25 and 0.5 weight %) whilst only small variations occurred with the highest quantities. The pore volume was 0.018 cm$^3$ g$^{-1}$ for pristine TiO$_2$, decreased by adding Pt (0.005 cm$^3$ g$^{-1}$) and further with nitrogen down to 0.003 cm$^3$ g$^{-1}$ whilst an increase occurred in the presence of tungsten, reaching the value of 0.03 cm$^3$ g$^{-1}$. The increase of the pore volume for the W-doped samples with respect to the N-doped ones (0.03 vs. 0.003 cm$^3$ g$^{-1}$) may partly justify the higher surface areas of the first samples, although the crystallites size for all of the doped samples is not dramatically different. The surface area and pore volume decrease observed after addition of Pt can be due to the pore blockage by the metal. Their variation in the presence of foreign species, instead, can be ascribed to surface modifications, as confirmed by TEM analyses.

The nitrogen adsorption-desorption isotherms (Figure 2) corresponded to Type IV isotherm$^{[61]}$ and showed a hysteresis loop of H4 type, with the two branches almost horizontal and parallel over a wide range of p/p° for the samples TiO$_2$, Pt-TiO$_2$, and Pt-TiO$_2$-N0.5, while for the sample Pt-TiO$_2$-W0.5 the hysteresis loop is intermediate between the types H3 and H4. These results suggest that the appearance of the hysteresis loop is more probably due to agglomeration of the particles, although also pores smaller than 100 Å have been identified by the size distribution analysis for the samples containing W.

A situation even more different was observed for the powders containing N or W. The sample Pt-TiO$_2$-N0.5, for instance, shows the presence of strange long needle-like particles (Figure 4A–B) whilst big and thick particles with irregular shape, whose dimension range from few tens up to hundreds of nanometers, can be noticed for the Pt-TiO$_2$-W0.5 sample (Figure 4C–D).

These results indicate that the different dopants influenced also the morphology of the samples.

In Figure 4(E–F) the sample Pt-TiO$_2$-W0.25 before (E) and after (F) its use in photocatalytic runs is shown. After the photocatalytic test no significant differences of shape and dimension of the particles can be noticed, but the clusters appear slightly bigger (Figure 4F), due to some aggregation which took place during the run.

High Resolution TEM (HR-TEM) and selected area electron diffraction (SAED) analysis of sample Pt-TiO$_2$-W0.25 (Figure 5) show that the obtained rutile is well crystallized with clear parallel lattice fringes. The fringe spacing is approximately 0.32 nm, which correspond to the (1 1 0) lattice spacing of rutile TiO$_2$.$^{[62]}$

In the inset of Figure 5 a sharper image (processed through an Inverse Fast Fourier Transform) of the highlighted area and the histogram showing the exact calculation of d-space are reported. Moreover, HR-TEM micrographs confirm that the crystalline structure did not change after the catalytic test.

In Figure 6 the reflectance spectra of bare TiO$_2$, Pt-TiO$_2$, Pt-TiO$_2$-N and Pt-TiO$_2$-W samples are reported. In the presence of Pt an increase of the absorption in the visible light region, with respect to the bare TiO$_2$, occurred, whilst a slight decrease was observed in the presence of nitrogen. The light absorption above 500 nm of the Pt-TiO$_2$-N and Pt-TiO$_2$-W samples can be attributed to oxygen vacancies, in accordance with the litera-
Figure 4. TEM micrographs of the samples Pt-TiO$_2$-N0.5 (A and B), Pt-TiO$_2$-W0.5 (C and D) and Pt-TiO$_2$-W0.25 before (E) and after (F) the photocatalytic runs.

Figure 5. High resolution TEM images of the sample Pt-TiO$_2$-W0.25 before (A) and after (B) the photocatalytic runs. Inset: processed image of highlighted area and plot profile of atomic fringe showing the calculations of d-spacing.

In Figure 6B it can be noticed that the spectrum of Pt-TiO$_2$ is red shifted with respect to those of the samples containing W because the sample mainly consists of rutile phase. Tungsten stabilized the anatase phase, its presence blue shifted the absorbance of the samples and increased the percentage of reflectance. The band gap values (Table 1) were obtained by extrapolating a linear fitting in the Tauc plot diagram (Figure 7). Bare TiO$_2$ exhibited a band gap of 2.98 eV which is typical of the rutile polymorph, reduced to 2.90 eV after Pt adding. A progressive decrease of the band gap values was observed (Table 1) at increasing nitrogen amount. This is probably due to a broadening of the valence band following the hybridization of the O 2p states with the N 2p states, with a consequent upward shift of the valence band maximum. In the presence of tungsten there was a band-gap increase, with the exception of the samples with the highest tungsten percentages and the Pt-TiO$_2$-N0.5W0.5 sample for which a decrease of the band-gap occurred. The band gap increase in the samples containing W can be attributed to the formation of the anatase phase (the band gap of anatase is higher than that of rutile) whose quantity rises with the W amount. For the highest W percentage two phenomena can take place which contribute in opposite way to the band gap value: the increase of the anatase fraction and the shift towards higher potential values of the CB edge with the W amount.$^{[66]}$ As reported in Table 1,
Indeed, there is a progressive band gap reduction with increasing tungsten percentage despite the increased quantity of anatase. The presence of surface oxygen vacancies in the doped samples could also play a role in determining the band gap value.

The spectra of the TiO₂ and Pt-TiO₂ samples exhibited a single sharp edge while those of the doped samples showed two linear sections, being the second attributable to the existence of a mid-gap band located above the valence band [42,64] or below the conduction band, due to the additional energy levels created by the doping agents.

2.2 Photocatalytic Runs

No formation of H₂ occurred in the presence of bare TiO₂, confirming the essential role of Pt. The results of the photocatalytic runs carried out under UV light irradiation in the presence of Pt-TiO₂, Pt-TiO₂-N0.5, and Pt-TiO₂-W0.5 samples are reported in Figure 8. Concentration of glucose and its partial oxidation products in liquid phase, along with H₂ in gas phase, vs. irradiation time are reported in Figure 8A–C whilst the amounts of H₂ and CO₂ per gram of catalyst after 6 h of irradiation are reported in Figure 8D. Notably, no CO and/or CH₄ were detected under the experimental conditions used. Arabinose, erythrose and formic acid were the main observed species deriving from the partial glucose oxidation, according to previous results [22].

In Table 2 the glucose conversion and the selectivity towards the different by-products are reported. After 6 h of UV irradiation the substrate conversion was 45.8 % by using the Pt-TiO₂ sample, a small increase was obtained by adding 0.5 % of nitrogen, and a significant reduction was observed in the presence of 0.5 % of tungsten. With regard to the selectivity towards arabinose, no great variation was observed in the presence of the three samples; the selectivity to formic acid was reduced by the presence of foreign elements, and a difference was obtained in the distribution of the oxidation products. In particular, no erythrose was detected in the presence of the Pt-TiO₂-W0.5 sample. The results confirm that glucose oxidation occurred with the C₁–C₂ break, following the mechanism reported in Figure 9, in accordance with literature [19,22]. The highest H₂ production (715 μmol g⁻¹ after 6 h of irradiation, Figure 8D) was measured with the non-doped sample and both N and W caused a decrease, obtaining the lowest amount with the addition of W. Similar CO₂ amounts were detected in the presence of the three samples.

In Figure 10 the trend of the concentration of all of the species measured during the photocatalytic runs carried out under a halogen lamp irradiation in the presence of the different samples is reported. The amount of both CO₂ and H₂ increased.
as the glucose concentration decreased. Glucose oxidation can occur via direct hole transfer to glucose bound to TiO₂ surface, due to the strong adsorption of the substrate, as reported in literature.[67–70] Photooxidation of carbohydrates in water was reported to involve sequential cleavage of formic acid to form the shorter aldose and H₂. The most active samples for H₂ production were those containing the lowest W percentages.

In Table 3 the glucose conversion and the selectivity to arabinose, erythrose and formic acid after 6 h of HL irradiation are reported. The behaviour of powders for glucose degradation was similar to that observed under UV light irradiation and the same oxidation products were obtained. The results showed that the photocatalytic glucose degradation mechanism is the same, regardless of the irradiation source used. In the presence of N-doped catalysts, the glucose conversion was very similar or slightly lower than that measured by using Pt-TiO₂, exhibiting the different N-doped samples a bell shaped trend of the photocatalytic efficiency. A noticeable negative effect, instead, was observed in the presence of W-doped catalyst, being the lowest value obtained by using the Pt-TiO₂-W0.5 catalyst. By adding N, the selectivity to arabinose and formic acid was higher than that found in the presence of the Pt-TiO₂ samples for all of the dopant percentages, whilst a bell shaped behaviour was observed by increasing the W dopant amount. The selectivity to erythrose was low and negligible in the presence of N and W, respectively, with the exception of the sample Pt-TiO₂-W2.
Table 3. Glucose conversion, selectivity to arabinose, erythrose and formic acid and amounts of CO\textsubscript{2} and H\textsubscript{2} per grams of catalyst after 6 h of HL irradiation.

<table>
<thead>
<tr>
<th>Sample</th>
<th>X\textsubscript{glucose}</th>
<th>S\textsubscript{arabinose}</th>
<th>S\textsubscript{erythrose}</th>
<th>S\textsubscript{formic acid}</th>
<th>CO\textsubscript{2} [μmol g\textsuperscript{-1}]</th>
<th>H\textsubscript{2} [μmol g\textsuperscript{-1}]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt-TiO\textsubscript{2}</td>
<td>62.1</td>
<td>35.9</td>
<td>20.1</td>
<td>49.9</td>
<td>33</td>
<td>578</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-N0.5</td>
<td>43.7</td>
<td>44.0</td>
<td>16.4</td>
<td>61.4</td>
<td>18</td>
<td>491</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-N1</td>
<td>61.3</td>
<td>45.9</td>
<td>16.0</td>
<td>61.8</td>
<td>30</td>
<td>278</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-N2</td>
<td>51.9</td>
<td>54.3</td>
<td>4.7</td>
<td>58.4</td>
<td>31</td>
<td>385</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-W0.25</td>
<td>37.4</td>
<td>33.5</td>
<td>–</td>
<td>32.3</td>
<td>85</td>
<td>708</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-W0.5</td>
<td>32.5</td>
<td>36.7</td>
<td>–</td>
<td>28.6</td>
<td>66</td>
<td>671</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-W1</td>
<td>34.5</td>
<td>41.5</td>
<td>–</td>
<td>41.5</td>
<td>49</td>
<td>45</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-W2</td>
<td>48.1</td>
<td>28.9</td>
<td>15.8</td>
<td>26.8</td>
<td>65</td>
<td>–</td>
</tr>
<tr>
<td>Pt-TiO\textsubscript{2}-N0.5W0.5</td>
<td>36.1</td>
<td>44.5</td>
<td>–</td>
<td>68.7</td>
<td>23</td>
<td>79</td>
</tr>
</tbody>
</table>

This behaviour indicates the occurrence of a different glucose reaction mechanism in the presence of the various photocatalysts,\textsuperscript{22} due to different effects of the foreign species (which is not easy to fully explain) on the surface properties. The glucose conversion in the presence of the co-doped sample was similar to that observed in the presence of the Pt-TiO\textsubscript{2}-W samples. Notably, no improvement was noticed with respect to the Pt-TiO\textsubscript{2} sample.

In Figure 11 the amounts of H\textsubscript{2} and CO\textsubscript{2} after 6 h of HL irradiation are reported. A negative effect for H\textsubscript{2} and CO\textsubscript{2} formation with respect to the Pt-TiO\textsubscript{2} sample was noticed in the presence of nitrogen whilst an increase of the formation of both products was observed by using low amounts of tungsten, despite the lower glucose conversion (Table 3) compared to that obtained with the N-doped samples. The highest W amounts had a detrimental effect on the photocatalytic H\textsubscript{2} production, being the latter negligible and equal to 708 μmol g\textsuperscript{-1} by using the Pt-TiO\textsubscript{2}-W2 and Pt-TiO\textsubscript{2}-W0.25 samples, respectively. H\textsubscript{2} evolution was low also when the co-doped Pt-TiO\textsubscript{2}-N0.5W0.5 catalyst was employed. By comparing the amounts of CO\textsubscript{2} produced under UV irradiation with those obtained under HL irradiation, in the latter a general more significant mineralization was obtained with a maximum in the presence of Pt-TiO\textsubscript{2}-W0.25 sample. The finding that the activity of the co-doped sample was not higher than that of the corresponding single doped samples, indicates that there was not a synergistic effect between the two dopant species, and probably the high defects number favoured the e\textsuperscript{-}/h\textsuperscript{+} recombination.

In order the reduction of H\textsuperscript{+} to H\textsubscript{2} take place, it is necessary that the conduction band level of the photocatalyst is more negative than the potential of the H\textsuperscript{+}/H\textsubscript{2} couple (E\textsubscript{H+/H2} = 0 V vs. NHE at pH = 0) (Figure 12). Generally, the TiO\textsubscript{2} rutile CB edge has a slight negative value,\textsuperscript{21} and it is reported that the presence of N in TiO\textsubscript{2} causes a VB shift towards less positive values whilst the incorporation of W induces a CB shift towards less negative values.\textsuperscript{66} These considerations could justify the drop in H\textsubscript{2} production by using the W-doped samples for the highest metal amounts. It is worth noticing that the value of

![Figure 11. CO\textsubscript{2} and H\textsubscript{2} amount per gram of catalyst after 6 h of HL irradiation. The error of analysis is less than ±5.](image1)

![Figure 12. Photocatalytic hypothesized mechanism of glucose oxidation and H\textsubscript{2} production.](image2)
the edge of the conduction band progressively moves to less negative values as the amount of tungsten increases, reaching values very close to the hydrogen reduction potential. In Table 4 the energy values of the conduction band ($E_{CB}$) and valence band ($E_{VB}$) edges, calculated in accordance with Xu and Schoonen,[72] are reported. The values were calculated by the empirical Equation (1) and Equation (2):[73]

$$E_{CB} = \chi - E^0 - 0.5 E_g$$  \hspace{1cm} (1)

$$E_{VB} = E_{CB} + E_g$$  \hspace{1cm} (2)

Table 4. Energy values of the conduction band ($E_{CB}$) and valence band ($E_{VB}$) edges.

<table>
<thead>
<tr>
<th>Sample</th>
<th>$E_{CB}$ [eV]</th>
<th>$E_{VB}$ [eV]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt-TiO$_2$</td>
<td>-0.29</td>
<td>2.97</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W0.5</td>
<td>-0.24</td>
<td>2.92</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W1</td>
<td>-0.12</td>
<td>2.80</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W2</td>
<td>-0.09</td>
<td>2.77</td>
</tr>
</tbody>
</table>

where $E_g$ is the band gap, $E^0$ is the energy of free electrons vs. hydrogen (4.5 eV), and $\chi$ is the electronegativity of the semiconductor, calculated as the geometric mean of Mulliken's electronegativities of the constituent atoms. For TiO$_2$ $\chi$ is 5.84 eV.[72] Even if absolute values cannot be obtained by means of this method because the structural factors are not taken into account, it can provide a rough estimation and a trend of the relative energy of CB and VB vs. normal hydrogen electrode (NHE). By increasing the W amount, a progressive decrease of the CB edge was estimated (Table 4), and for the sample Pt-TiO$_2$-W2 the value is very close to 0 V. This can justify the inactivity of this sample for H$_2$ production.

Some of the synthesized samples were also tested under natural solar light irradiation for 3.5 h. Figure 13 reports the reactivity results obtained by using Pt-TiO$_2$ and Pt-TiO$_2$-W0.25 samples, chosen as the most representative ones, vs. the cumulative irradiation energy. Both samples were more active under natural solar light irradiation than under HL irradiation, and the increase of activity was higher in the presence of the Pt-TiO$_2$-W0.25 sample, being the highest H$_2$ production equal to 3770 μmol g$^{-1}$ under natural solar light irradiation (Figure 13B). The above results showed the positive effect of tungsten on the H$_2$ formation by using the solar energy.

By comparing the photoactivity of Pt-TiO$_2$ and Pt-TiO$_2$-W0.25 samples under natural solar light irradiation and at the same cumulative energy value (Table 5), a higher glucose conversion and H$_2$ and CO$_2$ production can be noticed in the presence of tungsten. This finding indicates the beneficial role of W in the activation of TiO$_2$ for H$_2$ production by using the solar irradiation.

Table 5. Glucose conversion (X), selectivity (S) to arabinose, erythrose, formic acid, and amounts of CO$_2$ and H$_2$ per gram of catalyst under natural solar irradiation at the same cumulative energy value (2500 J).

<table>
<thead>
<tr>
<th>Sample</th>
<th>X$_{glucose}$</th>
<th>X$_{arabinose}$</th>
<th>X$_{erythrose}$</th>
<th>X$_{formic acid}$</th>
<th>CO$_2$ [μmol g$^{-1}$]</th>
<th>H$_2$ [μmol g$^{-1}$]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt-TiO$_2$</td>
<td>64.40</td>
<td>49.96</td>
<td>25.54</td>
<td>67.63</td>
<td>74</td>
<td>2218</td>
</tr>
<tr>
<td>Pt-TiO$_2$-W0.25</td>
<td>81.47</td>
<td>17.62</td>
<td>0.00</td>
<td>13.48</td>
<td>135</td>
<td>3734</td>
</tr>
</tbody>
</table>

The results found are in accord with those reported in the literature.[16] Many papers report the H$_2$ production from glucose photoreforming[16] in the presence of different types of TiO$_2$ doped materials and by using Xe, Hg, UVA, UV LED, and solar simulating lamps as the light sources. By using a solar light simulator, Kondarides et al.[74] observed H$_2$ production rates equal to 2.4 mmol gcat$^{-1}$ h$^{-1}$ in the presence of Pt-TiO$_2$ (P25) with the complete mineralization of glucose to CO$_2$ and without the formation of high value-added products. By using a 350 W Xe lamp as the irradiation source, 6.7 mmol gcat$^{-1}$ h$^{-1}$ of H$_2$ were obtained over Pt/TiO$_2$ nanosheets (Pt 2 %) with exposed the {001} facet.[75] Pdshell–Aucore nanoparticles immobilized on TiO$_2$ exhibited a high efficiency for H$_2$ production under UV irradiation (8.8 mmol gcat$^{-1}$ h$^{-1}$).[76] To the best of our knowledge no data about the activity under natural solar light irradiation are reported in literature. In our system the best H$_2$ production rates was ca. 1.0 mmol gcat$^{-1}$ h$^{-1}$ by using Pt-TiO$_2$-W0.25 under natural solar irradiation.

Some runs were carried out under HL irradiation by using, instead of glucose, formic acid (initial concentration 1 mM) as the sacrificial reagent, due to the simplicity of this molecule and the direct conversion to CO$_2$ and H$_2$, in accordance with...
literature.\textsuperscript{27,78} The evaluation of the stoichiometric ratio between the initial amount of formic acid and formed hydrogen allowed to indirectly hypothesize the occurrence of water splitting. Indeed, the amount of H\textsubscript{2} produced was higher than that which would have been derived simply from formic acid.

The substrate was completely degraded after 2 h of irradiation; afterwards the CO\textsubscript{2} concentration remained constant whilst the amount of H\textsubscript{2} began to decrease. The H\textsubscript{2} maximum production was ca. 1160 µmol (corresponding to 1449 µmol g\textsuperscript{−1}), almost 1.5 times greater than the formic acid initial molar quantity, whilst the CO\textsubscript{2} amount was a little lower than the stoichiometric value. This finding can be explained by considering that a fraction of formed CO\textsubscript{2} remained probably dissolved in water and/or adsorbed on the photocatalyst surface. By considering the formic acid decomposition (dehydrogenation) reaction\textsuperscript{27,78} according to Equation (3):

\[
\text{HCOOH} \rightarrow \text{CO}_2 + \text{H}_2
\] (3)

The finding that the molar ratio between H\textsubscript{2} and CO\textsubscript{2} was higher than 1 and that the H\textsubscript{2} amount was higher than that corresponding to the total mineralization of formic acid, suggests that the produced H\textsubscript{2} derived from both formic acid and water splitting, with the simultaneous formation of O\textsubscript{2}. Moreover, the decrease of the H\textsubscript{2} amount after formic acid disappearance was probably due to the occurrence of the back reaction between H\textsubscript{2} and O\textsubscript{2} and/or to the Ti\textsuperscript{4+} reduction by H\textsubscript{2}. Probably, the former reaction occurred also during the first hours of irradiation, but its rate increased by increasing the quantity of O\textsubscript{2} formed and present in the solution. When the substrate was still present, the amount of H\textsubscript{2} that recombined with oxygen was balanced by that coming from formic acid. This explains the observed decrease of H\textsubscript{2} production reaction rate by increasing the irradiation time, until an almost constant value or a decrease of its concentration was reached (Figure 14).

Notably, no production of H\textsubscript{2} was observed in the absence of formic acid, confirming the essential role of the sacrificial agents in the electron/hole separation. Some of these considerations based on the experimental results could also be extended to the H\textsubscript{2} production from glucose, because also in this case a decrease of the evolution rate of H\textsubscript{2} over time was noticed. Nevertheless, it was not possible to rationalize these findings or to check a mass balance as glucose was not completely mineralized, and other valuable products were obtained from its transformation.

3. Conclusions

N and/or W doped Pt-TiO\textsubscript{2} samples were used for the anaerobic glucose photo-conversion in aqueous solution under a halogen lamp and natural solar light irradiation with the dual purpose of obtaining high value-added products from the partial oxidation of the substrate and H\textsubscript{2} from the photo-reforming process. Different results were obtained in the presence of the two dopant species evidencing a different influence on the properties of TiO\textsubscript{2}. The presence of N in the Pt-TiO\textsubscript{2} sample generally did not influence the glucose conversion but increased the selectivity to arabinose and formic acid and decreased the H\textsubscript{2} production. Tungsten had a negative effect on the substrate conversion and a beneficial role on the H\textsubscript{2} evolution, with the exception of the highest dopant amounts. Moreover, arabinose, erythrose and formic acid were the main products of the partial oxidation of glucose when using the N-doped samples, whilst erythrose was found in the presence of W only for the highest dopant amount. These findings can be attributed to different TiO\textsubscript{2} surface properties when N and W are present. Moreover, also a difference in the electronic properties was observed in the presence of the two different dopants, in particular, N caused a VB shift towards less positive values whilst W induced a CB shift towards less negative values. The photocatalysts showed a good activity for H\textsubscript{2} production also under direct solar light irradiation, demonstrating the applicability of this type of materials under real-life conditions.

4. Experimental Section

4.1 Photocatalysts Preparation: Bare TiO\textsubscript{2} powders were prepared by adding 10 mL of TiCl\textsubscript{4} (Sigma–Aldrich) to 40 mL of distilled water at room temperature. After ca. 20 min of stirring, the clear solution was placed in a Teflon-lined stainless steel autoclave at room temperature, and afterwards heated at 100 °C for 48 h. The solid was recovered by drying at 60 °C, and it was named TiO\textsubscript{2}. Pt-TiO\textsubscript{2} samples (nominal Pt weight percentage 0.5 %) were obtained by adding the proper amount of PtCl\textsubscript{4} (Sigma–Aldrich) to the TiCl\textsubscript{4}/H\textsubscript{2}O solution prior to the hydrothermal treatment. Pt-Nitrogen, Pt-Tungsten or Pt-Nitrogen-Tungsten loading were performed under mild conditions: appropriate amounts of NH\textsubscript{4}Cl and WCl\textsubscript{6} were added to the clear Pt-TiCl\textsubscript{4}/H\textsubscript{2}O solution to obtain samples containing different weight percentages of N and W during the hydrothermal treatment. The samples were indicated as: Pt-TiO\textsubscript{2}-N0.5, Pt-TiO\textsubscript{2}-N1, Pt-TiO\textsubscript{2}-N2, Pt-TiO\textsubscript{2}-W0.25, Pt-TiO\textsubscript{2}-W0.5, Pt-TiO\textsubscript{2}-W1, Pt-TiO\textsubscript{2}-W2, Pt-TiO\textsubscript{2}-N0.5W0.5, the figures indicating the weight percentages of N and/or W.

4.2 Characterization of the Samples: X-ray diffraction (XRD) patterns of the powders were obtained by a PANalytical X′pertPro X-ray diffractometer with Cu-K\textalpha\textsubscript{1} radiation and 2θ scan rate of 2 °C/ min. The crystalline sizes of the samples were determined by using the Scherrer equation. The diffuse reflectance spectra (DRS) in the 250–800 nm wavelength range were obtained by using a Shimadzu

![Figure 14. Concentration of: (Orange ball) formic acid, (Blue diamond) CO\textsubscript{2} and (Green square) H\textsubscript{2} vs. time under HL irradiation in the presence Pt-TiO\textsubscript{2}-W0.25 sample. The error of analysis is less than ±5.](image-url)
UV-2401 PC spectrophotometer with BaSO₄ as the reference material. Band gap values were determined by plotting the modified Kubelka–Munk function, \([F(R')/hν]^{1/2}\), vs. the energy of the exciting light. The specific surface area (SSA) and the pore size distribution (PSD) of the photocatalysts were determined by nitrogen adsorption-desorption measurements with a Micromeritics ASAP 2020 apparatus. SSA’s were calculated by the BET equation in the p/p₀ interval 0.05–0.33 while the PSD’s were calculated by using the Barrett Joyner Halenda (BJH) methods.

Transmission electron microscopy (TEM) measurements were carried out by a JEOL (Japan) JEM-2100 electron microscope operating at 200 kV, while HR-TEM images SAED (Selected Area Electron Diffraction) patterns were obtained by a Gatan (USA) MSC camera. Samples were prepared by dispersing them in 2 mL of deionized water and by sonicating them for 5 minutes. A drop of each dispersion was spread on a copper holey carbon coated grid and dried overnight prior TEM analysis. Particle size and atomic layers spacing were measured by Gatan (USA) Digital Micrograph software.

4.3 Photocatalytic Activity: Some preliminary runs were carried out under UV light (by using a 125 W medium pressure Hg lamp) to test the efficiency of the catalysts. The lamp radiant power, measured by a radiometer Delta Ohm DO9721, was ca. 120 W m⁻² in the range 315–400 nm. Successively, all the prepared samples were tested in a 100 mL glucose solution. The lamp radiant power was 2100 W m⁻² in the 450–750 nm range, and 17 W m⁻² in the range 315–400 nm. N₂ was bubbled into suspensions for 0.5 h under dark to remove O₂ from the solution, then the reactor was sealed and the lamp switched off for 6 h. The best photocatalysts were tested also under natural solar light irradiation by using a closed batch Pyrex reactor whose volume was 250 mL, filled with 100 mL of glucose solution. In all cases, the amount of catalyst used was 1.0 g L⁻¹ of suspension. The runs under natural solar light irradiation lasted 3.5 h. Glucose was used both as sacrificial agent for H₂ production and as reagent to obtain high value chemicals from its partial oxidation. For the sake of comparison some experiments were performed by using formic acid instead of glucose. The concentrations of glucose, photo-products and formic acid were measured by HPLC equipped with a Diode Array and a refractive index detectors apparatus. SSA’s were calculated by the BET equation in the p/p₀ interval 0.05–0.33 while the PSD’s were calculated by using the Barrett Joyner Halenda (BJH) methods.

Band gap values were determined by plotting the modified Kubelka–Munk function, \([F(R')/hν]^{1/2}\), vs. the energy of the exciting light. Glucose reforming · Hydrogen evolution · Oxidation

Keywords: Photocatalysis · Doping · Titanium dioxide · Glucose reforming · Hydrogen evolution · Oxidation


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