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## Article

# Photoelectrochemical Enhancement of Graphene@WS<sub>2</sub> Nanosheets for Water Splitting Reaction

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**Abstract:** Tungsten disulfide nanosheets were successfully prepared by one-step chemical vapor deposition using tungsten oxide and thiourea in an inert gas environment. The size of the obtained nanosheets was subsequently reduced down to below 20 nm in width and 150 nm in length using high-energy ball milling, followed by 0.5 and 1 wt% graphene loading. The corresponding vibrational and structural characterizations are consistent with the fabrication of a pure WS<sub>2</sub> structure for neat sampling and the presence of the graphene characteristic vibration modes in graphene@WS<sub>2</sub> compounds. Additional morphological and crystal structures were examined and confirmed by high-resolution electron microscopy. Subsequently, the investigations of the optical properties evidenced the high optical absorption (98%) and lower band gap (1.75 eV) for the graphene@WS<sub>2</sub> compared to the other samples, with good band-edge alignment to water-splitting reaction. In addition, the photoelectrochemical measurements revealed that the graphene@WS<sub>2</sub> (1 wt%) exhibits an excellent photocurrent density (95 μA/cm<sup>2</sup> at 1.23 V bias) compared with RHE and higher applied bias potential efficiency under standard simulated solar illumination AM1.5G. Precisely, graphene@WS<sub>2</sub> (1 wt%) exhibits 3.3 times higher performance compared to pristine WS<sub>2</sub> and higher charge transfer ability, as measured by electrical impedance spectroscopy, suggesting its potential use as an efficient photoanode for hydrogen evolution reaction.

**Keywords:** WS<sub>2</sub> nanosheets; graphene@WS<sub>2</sub>; photochemical measurements; water splitting; reversible hydrogen electrode (RHE); photoanode

## 1. Introduction

Hydrogen fuel is considered a promising source of clean energy that could partly replace fossil fuels at the origin of greenhouse gas emissions. Today, there is a growing universal tendency to focus research and development on green hydrogen production by considering the general ecosystem to lower its costs and make it a more affordable clean-energy source, especially through water splitting (WS) [1–5]. In this context, several advanced materials were successfully examined and tested for enhancing the WS reaction, such as noble metals combined with highly catalytic semiconductors, which have shown an improved hydrogen evolution reaction (HER). However, their complex implementation and high cost prohibited their adoption in realistic WS plants [6–9]. Therefore, cost-efficient

and easy-to-produce alternative materials with improved WS performances have attracted a great interest. To this end, WS<sub>2</sub>, a transition-metal dichalcogenide (TMD) semiconductor, constitutes a highly attractive option owing to its excellent band gap tunability, its stability at high temperatures in corrosive media, and its easy and low-cost fabrication. Recently, both computational and experimental studies carried out on WS<sub>2</sub> have shown its effective role in catalyzing HER with a relatively good yield [10–16]. WS<sub>2</sub>-like materials, such as vertically aligned MoS<sub>2</sub>, have shown increasingly exposed edges as active sites for HER electrocatalysis, while the basal plane is catalytically inert [17]. The increasing interest in using TMDs as supporting electrodes is justified by their easy and low-cost process [15,18–20]. To further improve WS<sub>2</sub> performances, it is crucial to select an appropriate catalyst-supporting material that can prevent high-surface-area WS<sub>2</sub> nanostructures from agglomerating, enhance electron mobility, and overcome the surface overpotential to improve the overall HER performance. Graphene is the other two-dimensional carbon material that would meet the aforementioned criteria. Due to its high ability in electronic excitation and mobility, its mechanical properties, and its high stability, even at the nanoscale, graphene has received rigorous scientific and industrial attention for a long time [21–26]. Indeed, graphene is reported to possess high specific surface area (~2600 m<sup>2</sup>/g), high electron mobility  $\mu_e$  at room temperature (250,000 cm<sup>2</sup>/Vs), outstanding thermal conductivity (5000 W m<sup>-1</sup> K<sup>-1</sup>), and high electrical conductivity (5:6.4 × 10<sup>6</sup> S/m), as well as a very low weight. These outstanding properties enable graphene to be considered an excellent electrocatalyst support for WS<sub>2</sub> nanostructures [21,22].

In this work, we first produced low-cost graphene@WS<sub>2</sub> nanocomposite as electrodes for improved WS reactions. Next, we conducted a systematic structural characterization and assessment of the optical properties to evaluate the performance of the fabricated samples in photocatalysis-driven WS.

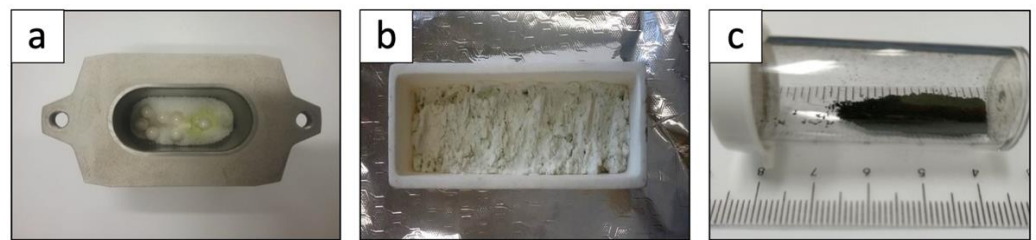
## 2. Materials and Methods

The raw materials, namely tungsten oxide (WO<sub>3</sub>) and thiourea (CH<sub>4</sub>N<sub>2</sub>S) powders, were acquired from Alfa Aesar™ (Thermo Fisher GmbH, Kandel, Germany). Graphene nanoplatelets were purchased from Sigma Aldrich™ St. Louis, MO, USA. The indium tin oxide (ITO) substrates used in this study are from Lumtec™ (Tapei, Taiwan), exhibiting a resistance < 10 Ω/cm<sup>2</sup>. All used substrates were cleaned by immersion in detergent, rinsed successively with acetone, ethanol, and deionized water, and dried under nitrogen flow. WS<sub>2</sub> nanosheets were prepared using four processing conditions consisting of various WO<sub>3</sub> and thiourea contents until achieving the complete suppression of the oxygen residue and obtaining pure WS<sub>2</sub> nanosheets. The initial materials' contents and the corresponding samples are summarized in the Table 1.

**Table 1.** The precursor masses used to prepare each WS<sub>2</sub> sample.

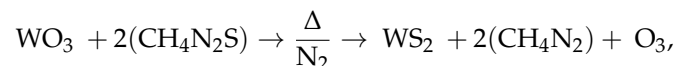
Sample	S1	S2	S3	S4
Thiourea (g)	4.5	5	5	7
WO <sub>3</sub> (g)	0.23	0.23	0.22	0.2

In a typical processing route, WS<sub>2</sub> nanosheets are prepared by mixing WO<sub>3</sub> with thiourea using a high-energy ball milling process E-Max Retsh™ (GmbH, Düsseldorf, Germany) machine with tungsten carbide container and Zirconia oxide balls (Figure 1a). After milling at 400 rpm for 30 min, the obtained mixture was collected into 50 mL alumina crucible (Figure 1b). A tube furnace was used for the chemical vapor deposition (CVD) reaction. The processing temperature was set to 850 °C under nitrogen gas flow of 500 sccm at 20 °C/s heating rate. The mixture was introduced into the tube furnace at T = 400 °C, and the CVD reaction took place for 1 h of dwell time at 850 °C, followed by air cooling. The obtained WS<sub>2</sub> exhibited its typical black color (Figure 1c).

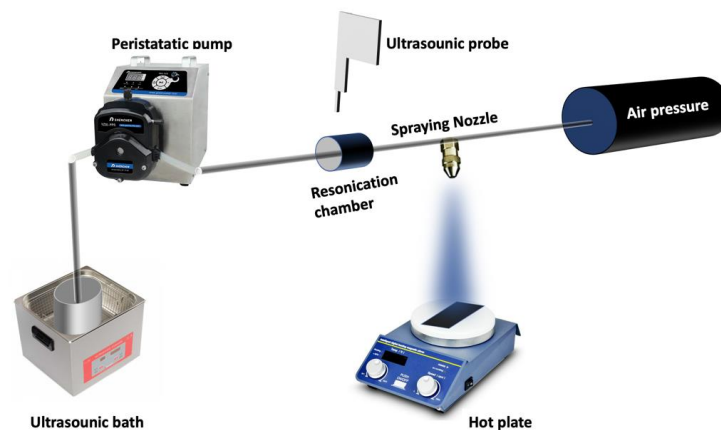


**Figure 1.** (a) Tungsten carbide container including sample mixture and ZrO<sub>2</sub> balls; (b) ball-milled mixture collected in alumina boat; (c) WS<sub>2</sub> nanosheets obtained after CVD reaction.

The chemical reaction occurring during the CVD process is expressed as follows:



For the graphene@WS<sub>2</sub> nanocomposites, two graphene contents (0.5 wt% and 1 wt%) were added to the optimized fabricated WS<sub>2</sub> nanosheets (S4), followed by high-energy ball milling at 400 rpm for 1 h. To investigate the different physical properties of the resulting compounds, thin films were prepared on top of various substrates, such as glass, quartz, and ITO. Figure 2 shows an illustration of the enhanced spray technique, where two ultrasonic-solution dispersive devices were utilized to prevent WS<sub>2</sub> nanosheets from aggregating in the transferring tube. Ethanol was used as the host solution.



**Figure 2.** Schematic diagram of the spray pyrolysis system used to prepare thin films out of the graphene@WS<sub>2</sub>.

For all samples, the spraying procedure consisted of 0.1 g of a mixture made of WS<sub>2</sub> with or without graphene, which was poured into a 40-megaliter ethanol container followed by water-bath sonication, as shown in Figure 2. During the solution distribution through a peristaltic pump, a second sonication (ultrasound probe) was used to resuspend the mixture towards the nozzle. Finally, an air pressure of 1.2 bar allowed spraying the dispersed suspension through a nozzle over the desired substrate, which was kept at 45 °C with a hot plate. The nozzle–substrate distance was maintained at 20 cm and the deposition rate was set to 1 mL/min for a total deposition time of 30 min. The obtained film thickness was in the range of 150 to 200 nm.

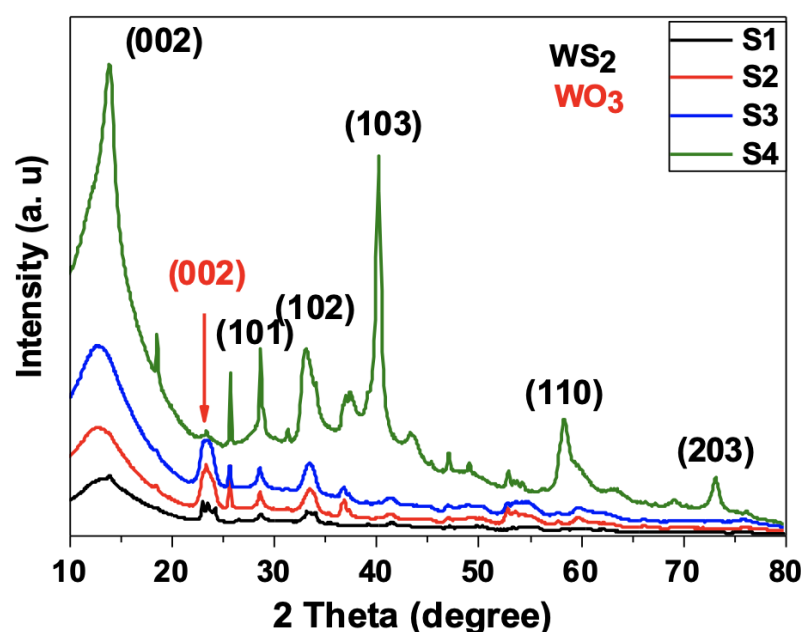
Subsequently, the structural characterization was carried out using a Bruker™ D4 Endeavor X-ray diffractometer with a 1.54 Å CuKα source, and the vibrational analyses were performed with Raman spectroscopy Renishaw™ (Wotton-under-Edge, UK) using a green laser excitation source (532 nm). The microstructure analysis was performed with a Zeiss™ (Oberkochen, Germany) Gemini 500 ultra-high resolution field emission electron microscope (FESEM) operating at low voltage (1 kV), using an In-lens detector. The crystal structure and the morphology of the obtained nanocomposite were investigated

by high-resolution transmission electron microscopy (HRTEM) Cs-corrected Titan from Thermo Fisher Scientific™ (Waltham, MA, USA) operating at 300 kV. TEM samples were prepared on holey carbon Cu grids using drop-casting method. The analysis of optical properties was conducted on a UV-vis-near IR spectrometer V-700 JASCO™ (Easton, MD, USA) and Fourier transform infrared spectroscopy (FTIR) from Thermo Fisher Scientific™ (Waltham, MA, USA). The electrical impedance spectroscopy (EIS) and the photoelectrochemical (PEC) measurements were performed using PalmSens4™ (Houten, Netherlands) EIS electrochemical interface.

### 3. Results and Discussion

#### 3.1. XRD Analysis

The XRD diagrams obtained for the four prepared WS<sub>2</sub> samples are shown in Figure 3. The XRD patterns were recorded in the range of 2-theta 10–80° and compared to the observed reflection planes (002), (101), (102), (103), (110) and (203) with the standard diffraction data file (JCPDF card no. 01-084-1398). These diffraction peaks were indexed with the hexagonal WS<sub>2</sub> phase of cell constants  $a = b = 3.15 \text{ \AA}$ ,  $c = 12.32 \text{ \AA}$ .

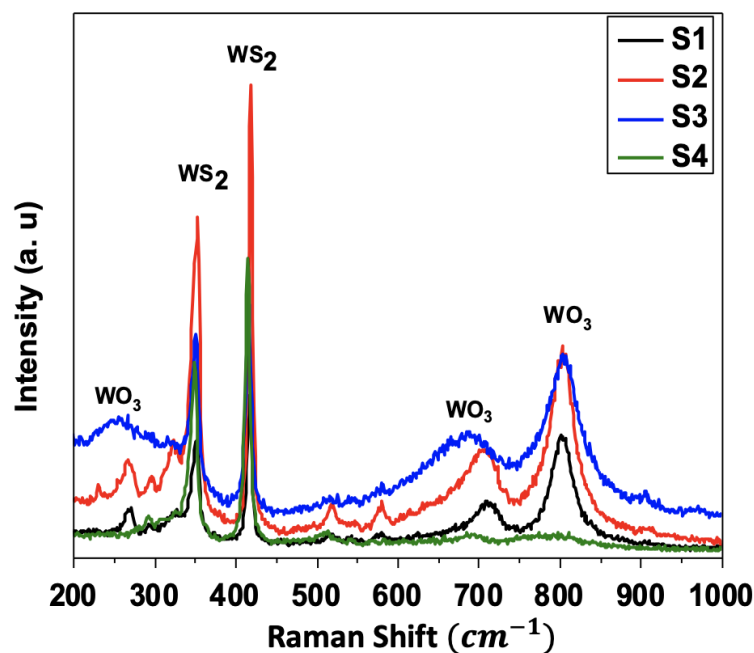


**Figure 3.** XRD diagrams for the four processed samples. Note the absence of WO<sub>3</sub> peak in sample S4 indicating the high purity WS<sub>2</sub> nanosheets.

The XRD diagrams of the samples S1, S2, and S3 exhibit an extra peak at (002) position which corresponds to the WO<sub>3</sub> structure. This indicates that the reaction of the WO<sub>3</sub> with the thiourea was not complete; hence, the resulting WS<sub>2</sub> of these samples contained WO<sub>3</sub> residue. It is worth noting that the strength of the WO<sub>3</sub> residue peak was observed to decrease with increasing thiourea content (from sample S1 to S3). By contrast, the WO<sub>3</sub> peak was no longer visible at higher thiourea content as it disappeared for the sample S4. This result suggests that the excess thiourea added to the mixture made it possible to obtain the pure WS<sub>2</sub> nanosheets.

#### 3.2. Raman Spectroscopy

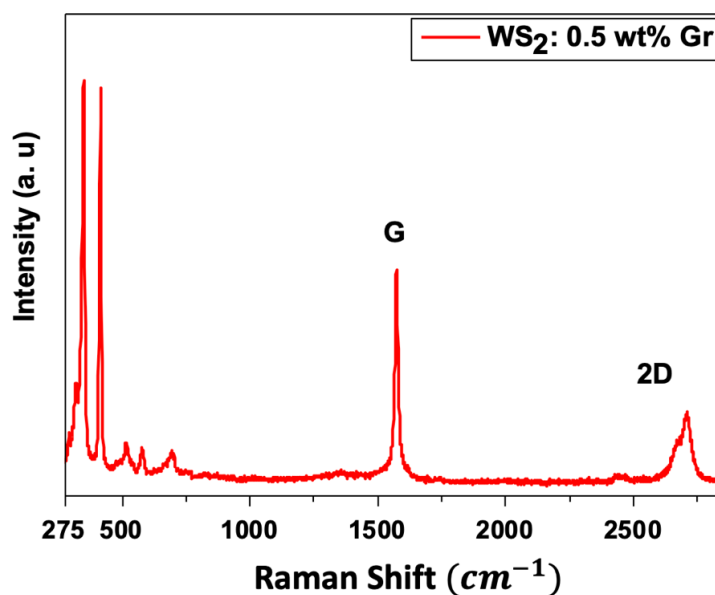
Figure 4 illustrates the vibrational modes of the four processed WS<sub>2</sub> samples. All the samples show common WS<sub>2</sub> characteristic peaks, i.e., two strong peaks attributed to E<sub>2g1</sub> in-plane and A<sub>1g</sub> out-of-plane vibrational modes appearing at the 350 and 420 cm<sup>-1</sup> positions, respectively. The E<sub>2g1</sub> mode involves a displacement of W and S atoms, whereas the A<sub>1g</sub> mode concerns only the S atoms.



**Figure 4.** Raman spectra for the four prepared WS<sub>2</sub> samples. Note the absence of WO<sub>3</sub> vibration modes at 286 cm<sup>-1</sup>, 666 cm<sup>-1</sup>, and 820 cm<sup>-1</sup> in sample S4, which highlights the high purity of WS<sub>2</sub>.

Similar to the XRD results, the Raman spectra of the S1, S2, and S3 showed the presence of WO<sub>3</sub> vibration modes at the 293 cm<sup>-1</sup>, 675 cm<sup>-1</sup>, and 836 cm<sup>-1</sup> positions, respectively, while the S4 only exhibited the vibration modes E<sub>1g</sub><sup>1</sup> and A<sub>1g</sub> for WS<sub>2</sub> occurring at 351 cm<sup>-1</sup> and 415 cm<sup>-1</sup>, respectively. As mentioned above, these results confirm that the CVD reaction was fully completed for the S4 and the oxygen was entirely consumed.

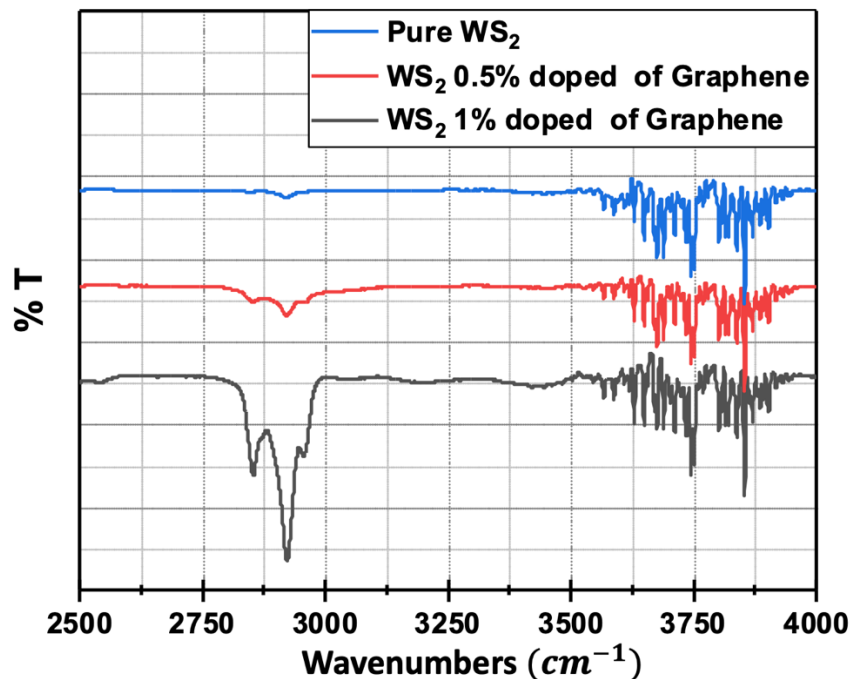
Furthermore, the Raman spectroscopy was carried out on graphene@WS<sub>2</sub> samples, and the typical Raman spectra are depicted in Figure 5. In addition to the presence of the common WS<sub>2</sub> peaks, graphene vibrational modes were also obtained for both samples at 1560–1575 cm<sup>-1</sup> and at 2647–2700 cm<sup>-1</sup> for the G-band and 2D-band, respectively. This confirms the presence of the graphene in the processed graphene@WS<sub>2</sub> samples.



**Figure 5.** Raman spectra for graphene@WS<sub>2</sub> samples. Typical G-band and 2D-band vibrational modes of graphene are recorded.

### 3.3. FTIR Spectroscopy

To further screen the presence of the graphene in the processed graphene@WS<sub>2</sub> nanocomposite samples, FTIR absorption spectroscopy was conducted on pure WS<sub>2</sub>, WS<sub>2</sub>: 0.5 wt% Gr, and WS<sub>2</sub>: 1 wt% Gr, respectively, as shown in Figure 6.



**Figure 6.** FTIR absorption spectra of WS<sub>2</sub>, WS<sub>2</sub>: 0.5 wt% Gr, and WS<sub>2</sub>: 1 wt% Gr. Note the presence of the typical graphene absorption peak at 2780–3000 cm<sup>−1</sup> for both graphene@WS<sub>2</sub> samples.

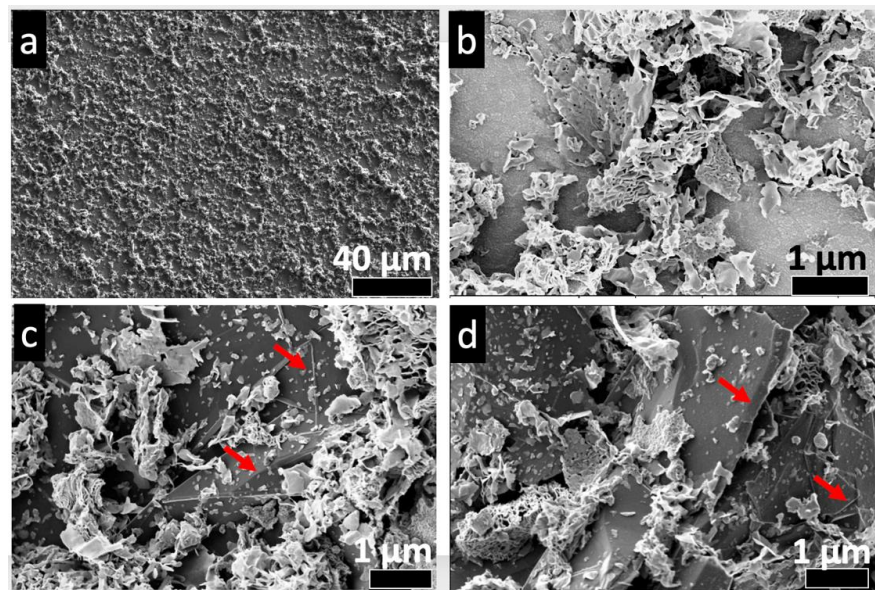
As can be seen, the typical graphene absorption peak is present in the graphene@WS<sub>2</sub> nanocomposites and the strength of this peak is more pronounced for the sample with the higher graphene content.

### 3.4. Microstructure Analysis

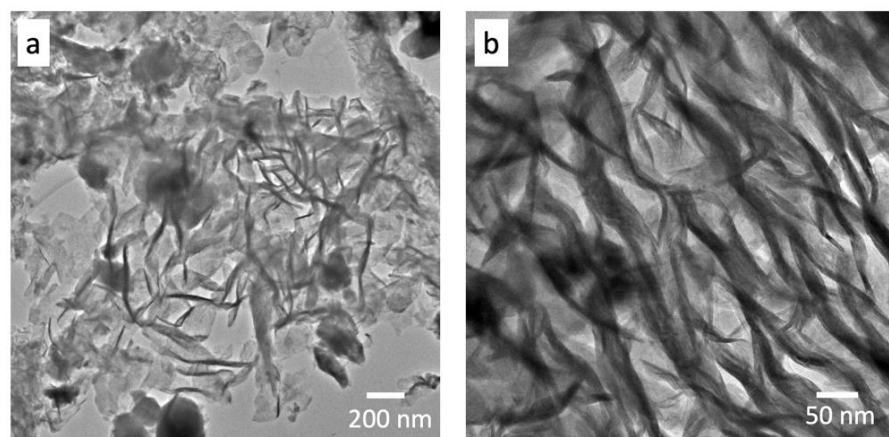
A general view of the sprayed samples on the ITO substrate is given in Figure 7a. As can be seen, the nanosheets are evenly distributed with relatively uniform thickness. The insets shown in Figure 7b–d highlight the inner structures of the neat WS<sub>2</sub>, WS<sub>2</sub>: 0.5 wt% Gr, and WS<sub>2</sub>: 1 wt% Gr, respectively. Both the WS<sub>2</sub> nanosheets and the graphene platelets are visible. These samples were further investigated by HRTEM and used for the photoelectrochemical measurements.

Figure 8 depicts the bright-field TEM image of the typical microstructure encountered in WS<sub>2</sub>: 1 wt% Gr. The WS<sub>2</sub> nanosheets (dark contrast) appeared to be encapsulated in the graphene nanoplatelets (light contrast), as shown in Figure 8a. A higher-magnification TEM image is given in Figure 8b; it shows a focus on entangled graphene@WS<sub>2</sub> nanosheets. The quality of the nanocomposite was further verified using HRTEM.

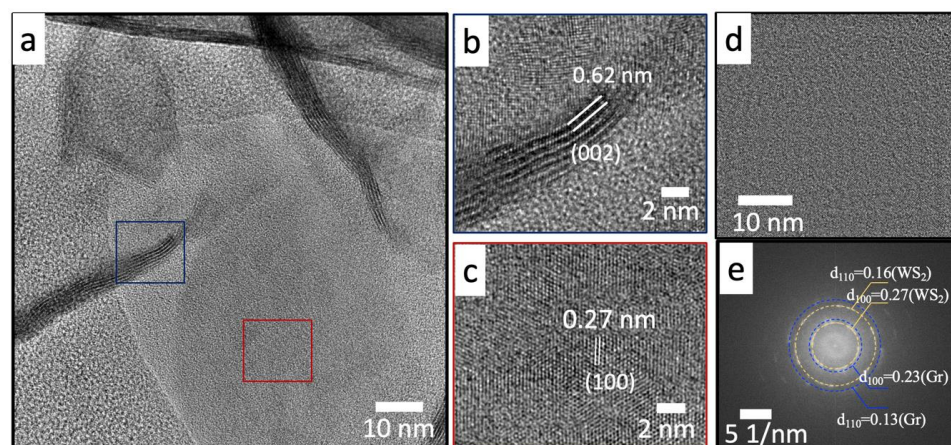
Figure 9a clearly shows the interconnections between the WS<sub>2</sub> nanosheets and the graphene nanoplatelets, delimited by the blue box highlighted in Figure 9b. Their corresponding crystal structure conformed with the 2H-WS<sub>2</sub> materials, as demonstrated by the interplanar distances  $d_{(002)} = 0.62$  nm and  $d_{(100)} = 0.27$  nm, illustrated in the red box in Figure 9c. The analysis of the interconnected WS<sub>2</sub> and graphene regions given in Figure 9d revealed the presence of both graphene and WS<sub>2</sub>, as demonstrated by the corresponding fast Fourier transform (FFT) intensity image depicted in Figure 9e.



**Figure 7.** Low kV SEM micrographs: (a) typical general view for all samples; (b) zoom-in on neat WS<sub>2</sub> nanosheets; (c) zoom-in on WS<sub>2</sub>: 0.5 wt% sample; and (d) zoom-in on WS<sub>2</sub>: 1 wt% sample. Red arrows identify the graphene platelets on graphene@WS<sub>2</sub> samples.



**Figure 8.** Bright-field TEM images of graphene@WS<sub>2</sub> nanocomposites: (a) low and (b) high magnification.



**Figure 9.** HRTEM images of graphene@WS<sub>2</sub> nanocomposites: (a) general view; (b) blue-box zoom-in, showing the crystal lattice of WS<sub>2</sub> d<sub>002</sub>; (c) red-box zoom-in showing the crystal lattice of WS<sub>2</sub> d<sub>100</sub>; (d) HRTEM image of interconnected graphene ws2 region and corresponding (e) FFT image.



### 3.5. Optical Properties

The investigation of the optical properties was carried out using an optical spectrometer in the 350–800 nm range. Figure 10 shows the optical absorption of all the considered samples.

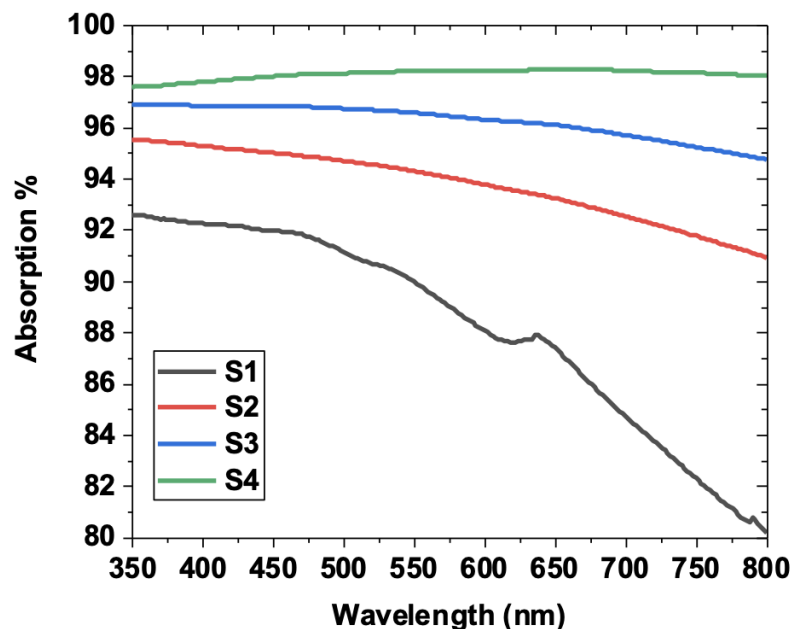


Figure 10. Optical absorption for the four considered samples. Note excitons position at 620 nm.

As can be seen, samples S1, S2, and S3 exhibited similar optical behavior, consisting of a steady decrease in the optical absorption, which started at 93–97% and hit 80% for S1. This suggests that the presence of  $\text{WO}_3$  impurities induces a decrease in the optical absorption in the visible region. By contrast, the optimized sample S4 showed a relatively unchanged optical absorption. In particular, it remained stable until it reached the excitons position (around 620 nm), at which point it slightly increased. It is clear that the optimized  $\text{WS}_2$  sample S4 exhibited high broadband light absorption in the visible region, reaching more than 98% compared to the other samples. Using the Tauc formula  $(\alpha h\nu)^n = A(h\nu - E_g)$ , the band-gap energy was obtained, as depicted in Figure 11.

The obtained band-gap energies were 1.91 eV, 1.87 eV, 1.82 eV, and 1.8 eV for S1, S2, S3, and S4, respectively. The S4 showed the lowest energy, which was in agreement with the broadband optical absorption performances achieved.

To evaluate the effect of the graphene's incorporation on the  $\text{WS}_2$  nanosheets, the optical absorption was also measured for both graphene@ $\text{WS}_2$  samples, namely 0.5 wt% Gr and 1 wt% Gr, and the results are shown in Figure 12.

The addition of  $\text{WS}_2$ : 0.5 wt% Gr does not seem to enhance the optical absorption (Figure 10 vs. Figure 12a). Indeed, the optical absorption of this sample reaches 98% absorption but only in the 550–750 nm region in contrast to pristine  $\text{WS}_2$ , which exhibits a broadband absorption. Nevertheless, the  $\text{WS}_2$ : 0.5 wt% Gr band gap energy appears to slight decrease to 1.78 eV (Figure 12c). In contrast, the  $\text{WS}_2$ : 1 wt% Gr sample does exhibit an optical absorption exceeding 99% across the entire visible light spectrum. Moreover, its band gap energy is further decreased to reach 1.75 eV. Considering these performances,  $\text{WS}_2$  and  $\text{WS}_2$ : 1 wt% Gr samples were selected to perform photochemical measurements.

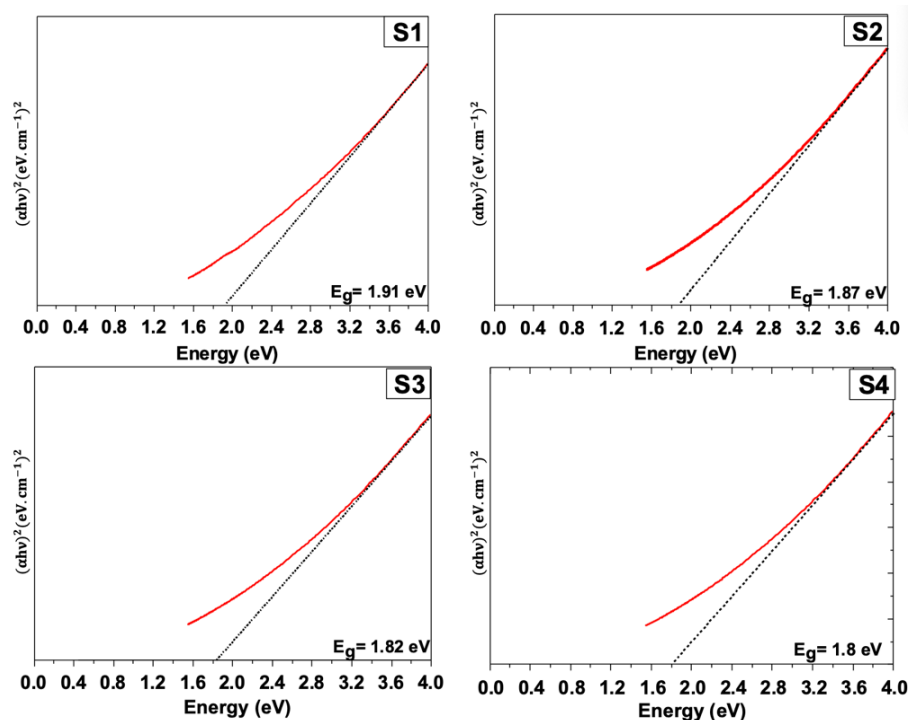


Figure 11. Tauc plots for all samples showing the lowest band-gap energy value for the sample S4.

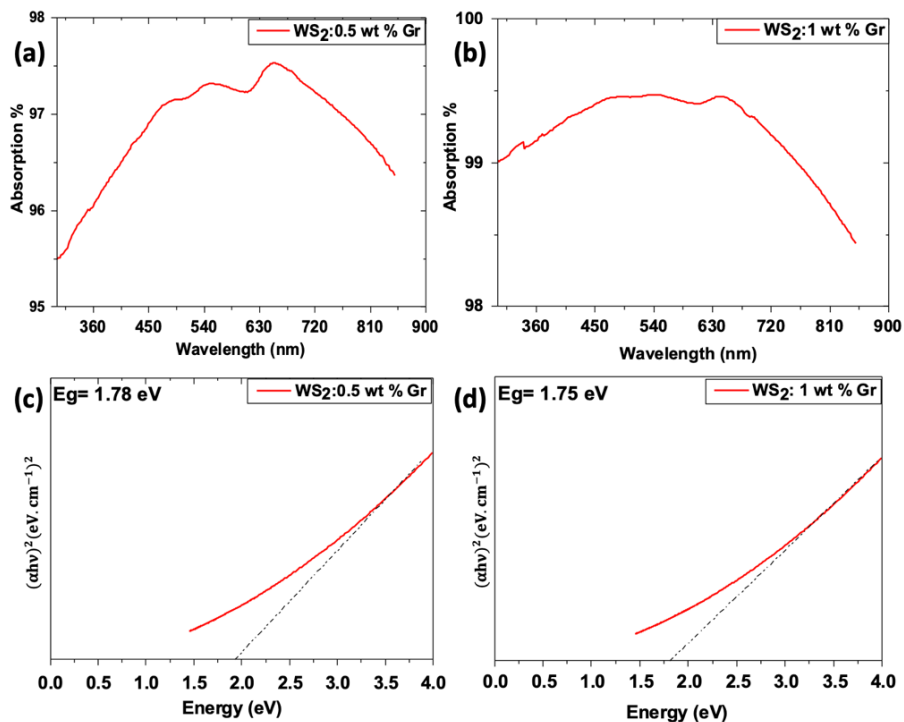


Figure 12. Optical properties of graphene@WS<sub>2</sub> samples: (a,b) optical absorption of WS<sub>2</sub> 0.5 wt% and 1 wt%, respectively; (c,d) corresponding Tauc plots.

### 3.6. Photochemical Measurements

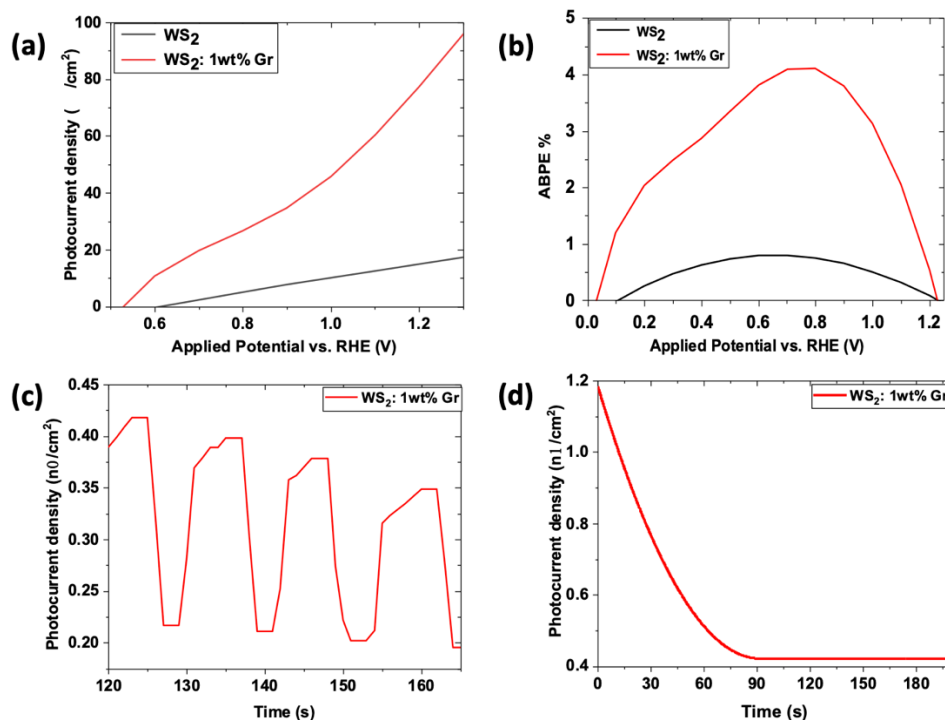
In this section, the photocatalytic performances of neat WS<sub>2</sub> and WS<sub>2</sub>: 1 wt% Gr samples are screened. Both samples were deposited on ITO substrate and immersed in deionized water medium (pH = 6). Then, a linear sweep voltammetry (LSV) and chronoamperometry experiments were carried under standard solar simulator (energy~AM1.5G).

To extract the potential of RHE, the following Nernst equation was used:

$$E_{\text{RHE}} = E_{(\text{Ag}/\text{AgCl})} + (0.059 \times \text{pH}) + E_0$$

where  $E_0 \approx 0.197$  V at 25 °C and  $E_{\text{Ag}/\text{AgCl}}$  is the applied potential.

For both samples, the active surface is about 1 cm<sup>2</sup> and the distance between the three electrodes was kept at 5 mm. A Pt fishnet (0.5 cm outer diameter) was used as counter electrode, while standard Ag/AgCl was used as the reference electrode. The LSV scan rate was set to 0.1 V/s along 0–1.5 V range. Prior to LSV experiments, all measurements were stabilized for 200 s under zero applied potential. The results are depicted in Figure 13.



**Figure 13.** Photoelectrochemical measurements carried out on optimized WS<sub>2</sub> S4 and graphene@WS<sub>2</sub>: 1 wt%: (a) generated current density as function of applied potential vs. RHE; (b) applied bias potential efficiency as function of applied bias with respect to RHE. Chronoamperometry experiment performed on WS<sub>2</sub>: 1 wt% Gr using solar simulator of AM1.5G; (c) cyclic and (d) steady state tests.

As can be seen in Figure 13a, a high-density current was obtained for the graphene@WS<sub>2</sub>, which was observed to quickly increase with increasing RHE with respect to the applied potential. Note that the highest current density obtained for the S4 WS<sub>2</sub> at high applied voltage was reached by the graphene@WS<sub>2</sub> sample at very low applied voltage. Hence, the addition of graphene dramatically enhanced the generated density current, six-folds, which was beneficial to HER. To produce WS reactions, a theoretical potential value of 1.23 V versus RHE is required without considering the surface overpotential and the voltage loss due to electron transport. As can be seen, the S4 WS<sub>2</sub> nanosheets showed the lowest photocurrent density, of 17 μA/cm<sup>2</sup>, at 1.23 V versus RHE over the entire potential range. When the graphene was added, the photocurrent density increased vigorously to 95 μA/cm<sup>2</sup>. The onset potentials of the S4 WS<sub>2</sub> and WS<sub>2</sub>: 1 wt% Gr nanosheets were 0.61 and 0.52 V, respectively. The cathodic shift in the onset potentials indicated an enhanced charge transport; hence, a higher separation efficiency was obtained, even at low ranges of applied potential.

Moreover, the applied bias potential efficiency (ABPE) was evaluated using the following equation:

$$ABPE = \frac{J(1.23 - V_{\text{bias}})}{P_{\text{light}}}$$

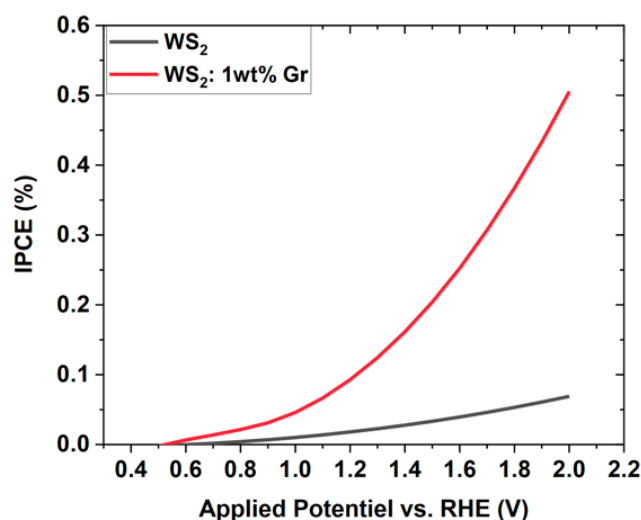
where  $J$  is the current density,  $V_{\text{bias}}$  is the bias potential, and  $P_{\text{light}}$  is the light power.

The ABPE equation translates how much the cell device using the processed samples as photoanodes is able to produce ionization current under an external applied voltage at constant solar irradiation. Here, the light power was AM1.5G. The applied potentials were converted into the corresponding potential versus RHE using the Nernst equation. The ABPE findings given in Figure 13b indicate that the WS<sub>2</sub>/ITO photoanode exhibited an ABPE of 0.79% at around 0.75 V versus RHE, while the WS<sub>2</sub>: 1 wt% Gr/ITO photoanode reached 4.11% at the same voltage versus RHE. This increase in ABPE in the graphene@WS<sub>2</sub> sample represented a fourfold-higher performance than that of the pristine WS<sub>2</sub> nanosheets. This underlines the beneficial effect of WS<sub>2</sub> loaded with graphene on photocatalytic WS reactions.

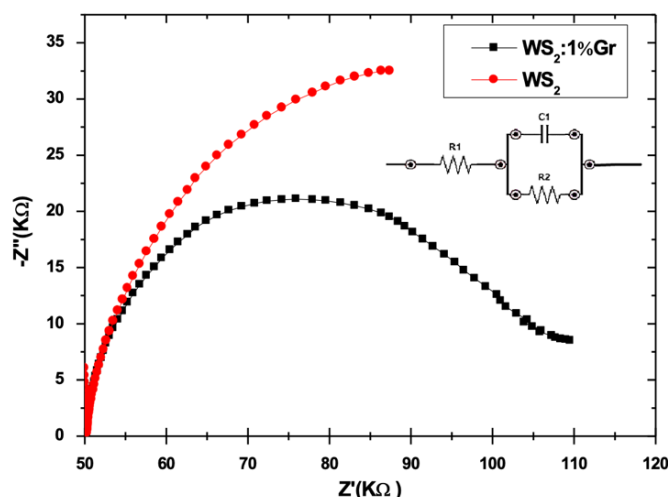
To examine the photoresponse of the photoanode WS<sub>2</sub>: 1 wt% Gr/ITO over time, the transient photocurrent was recorded at 0 V of bias with the light on/off cycles at AM1.5G/cm<sup>2</sup>, using a monitored mechanical shutter. The results depicted in Figure 13c demonstrate a fair stability profile over more than 160 s and the fast response of the excited photoanode over a duration of less than 10 s and a dwell time of 20 s. The stability of the current density was screened by the steady-state measurements of the current density generated (Figure 13d). The photocurrent stability of the WS<sub>2</sub>: 1 wt% Gr photoanode was examined at a bias potential of 0 V under AM1.5G illumination. A fast decay occurred for the first 100 s. Subsequently, a plateau was recorded, indicating that the photocurrent had reached a value of 0.4 nA/cm<sup>2</sup> over the following 100 s. During this time, the photogenerated electrons were transferred to the Pt counter electrode to boost the HER. Consequently, the generated holes could actively participate in the oxidation process at the photoanode site. In our case, the initiation of the WS experiment induced the accumulation of the photogenerated holes at the photoanode site, since no bias potential was applied. These holes could barely be scavenged by water molecules, and recombination occurred, resulting in a decrease in the photocurrent with time. After about 100 s, the rate of generation and consumption of the holes became constant and the photocurrent is stable [15]. Furthermore, we accounted for the efficiency of our photoelectrochemical process for the neat WS<sub>2</sub> NSs and WS<sub>2</sub>: 1 wt% Gr by evaluating the incident photon-to-current efficiency (IPCE), which gives a good estimation of the number of produced electrons with respect of the number of incident photons. The IPCE was determined by the following expression: IPCE (%) =  $J \times V$  / incident Power. The obtained IPCE was plotted against the applied potential (V vs. RHE) and depicted in Figure 14.

Figure 14 shows an increase in IPCE for both samples as a function of the applied bias. The IPCE reached 0.1% for the neat sample, whereas it approached more than 0.5% for the WS<sub>2</sub>: 1 wt% Gr. Therefore, the incorporation of the graphene dramatically enhanced the IPCE, which exhibited an exponential function profile. This indicates that the graphene provided additional electrons circulating in the photochemical cell, which is further proof of the beneficial effect of graphene in enhancing photochemical reactions.

To further examine the performance of the WS<sub>2</sub>: 1 wt% Gr photoanode, an EIS experiment was carried out to evaluate its charge transfer capabilities. The EIS was conducted on a three-electrode cell and deionized water electrolyte. A Pt fishnet and Ag/AgCl were utilized as the counter and reference electrodes, respectively. The applied voltage was set to 1 V, with a frequency sweep in the range of 0.01 Hz–100 KHz under visible light irradiation at 100 mW/cm<sup>2</sup> of power density. The exposed surface area of all the samples was set to 1 cm<sup>2</sup>. Nyquist plots of the optimized WS<sub>2</sub> and WS<sub>2</sub>:1 wt% Gr and the corresponding equivalent electrical circuit are shown in Figure 15.



**Figure 14.** Variation in the incident photon-to-current efficiency (IPCE) as function of applied potential (V vs. RHE) calculated for neat WS<sub>2</sub> NSs and WS<sub>2</sub>: 1 wt% Gr.



**Figure 15.** EIS measurements of WS<sub>2</sub> and WS<sub>2</sub>: 1 wt% Gr samples. The inset is the equivalent circuit used to extract the resistance of both samples.

The comparison of the EIS measurements with the equivalent electrical circuit indicated the higher resistance of the S4 WS<sub>2</sub> nanosheets (45.8 KΩ) compared to the very low resistance obtained for WS<sub>2</sub>: 1 wt% Gr (10 KΩ). This result clearly shows that the WS<sub>2</sub> loaded with graphene promoted a fourfold better charge transfer compared to the neat WS<sub>2</sub> sample. This performance undoubtedly suggests that graphene@WS<sub>2</sub> is more suitable for use as a photoanode for HER.

In summary, in contrast to previously reported work [2,27], the materials used in the present study were produced using a one-pot fabrication process yielding a nanocomposite material made of WS<sub>2</sub> NSs and graphene. Our findings demonstrate that processing composite materials based on TMD materials combined with a semimetal achieves four-to-five-times better current density at 1.23 V (V vs. RHE), which is necessary to drive WS reactions and hydrogen-evolution reactions. This contrasts with the findings of a previous study [23], in which the r-GO semiconductor was mixed with TMDs to produce heterostructures exhibiting good performances at lower applied potentials (i.e., −0.3 V for WS<sub>2</sub>/rGO and −0.4 V for neat WS<sub>2</sub>). These values remain low compared to the theoretical potential of 1.23 V required to achieve WS and HER.

#### 4. Conclusions

Nanocomposite materials consisting of graphene and WS<sub>2</sub> were produced in order to be used as highly efficient photoanodes for hydrogen production via a water-splitting reaction. The successful fabrication of the optimized WS<sub>2</sub> nanosheets as well as the graphene@WS<sub>2</sub> with two graphene contents was confirmed by several combined techniques, including XRD, Raman and FTIR spectroscopies, SEM, and HRTEM. Our main findings indicate that the sample WS<sub>2</sub> with 1 wt% graphene exhibited a broadband visible light absorption reaching 98% and an appropriate band gap of 1.75 eV for water-splitting reactions. These outstanding optical properties led to an enhancement of the photogenerated electrons and to a higher charge transfer, as recorded by the photochemical and electron impedance spectroscopy measurements at ambient conditions. Furthermore, associating graphene with WS<sub>2</sub> at only 1 wt% content led to an increase in the current density from 17 to 95  $\mu\text{A}/\text{cm}^2$  at 1.23 V versus RHE under AM1.5G illumination with ABPE, which was fourfold higher than the pristine WS<sub>2</sub> nanosheets. Hence, the graphene@WS<sub>2</sub> could be considered a desirable high-efficient photoanode for hydrogen-evolution reactions.

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#### References

1. Cortright, R.D.; Davda, R.R.; Dumesic, J.A. Hydrogen from Catalytic Reforming of Biomass-Derived Hydrocarbons in Liquid Water. *Nature* **2002**, *418*, 964–967. [[CrossRef](#)] [[PubMed](#)]
2. Kudo, A.; Miseki, Y. Heterogeneous Photocatalyst Materials for Water Splitting. *Chem. Soc. Rev.* **2009**, *38*, 253–278. [[CrossRef](#)] [[PubMed](#)]
3. Yilanci, A.; Dincer, I.; Ozturk, H.K. A Review on Solar-Hydrogen/Fuel Cell Hybrid Energy Systems for Stationary Applications. *Prog. Energy Combust. Sci.* **2009**, *35*, 231–244. [[CrossRef](#)]
4. Rajeshwar, K.; McConnell, R.; Licht, S. *Solar Hydrogen Generation*; Springer: New York, NY, USA, 2008. [[CrossRef](#)]
5. Khaselev, O.; Turner, J.A. A Monolithic Photovoltaic-Photoelectrochemical Device for Hydrogen Production via Water Splitting. *Science* **1998**, *280*, 425–427. [[CrossRef](#)]
6. Sakthivel, S.; Shankar, M.V.; Palanichamy, M.; Arabindoo, B.; Bahnemann, D.W.; Murugesan, V. Enhancement of Photocatalytic Activity by Metal Deposition: Characterisation and Photonic Efficiency of Pt, Au and Pd Deposited on TiO<sub>2</sub> Catalyst. *Water Res.* **2004**, *38*, 3001–3008. [[CrossRef](#)] [[PubMed](#)]
7. Linic, S.; Christopher, P.; Ingram, D.B. Plasmonic-Metal Nanostructures for Efficient Conversion of Solar to Chemical Energy. *Nat. Mater.* **2011**, *10*, 911–921. [[CrossRef](#)]
8. Abed, J.; Rajput, N.S.; El Moutaouakil, A.; Jouiad, M. Recent Advances in the Design of Plasmonic Au/TiO<sub>2</sub> Nanostructures for Enhanced Photocatalytic Water Splitting. *Nanomaterials* **2020**, *10*, 2260. [[CrossRef](#)]
9. Wu, B.-H.; Liu, W.-T.; Chen, T.-Y.; Perng, T.-P.; Huang, J.-H.; Chen, L.-J. Plasmon-Enhanced Photocatalytic Hydrogen Production on Au/TiO<sub>2</sub> Hybrid Nanocrystal Arrays. *Nano Energy* **2016**, *27*, 412–419. [[CrossRef](#)]
10. Xu, S.; Li, D.; Wu, P. One-Pot, Facile, and Versatile Synthesis of Monolayer MoS<sub>2</sub>/WS<sub>2</sub> Quantum Dots as Bioimaging Probes and Efficient Electrocatalysts for Hydrogen Evolution Reaction. *Adv. Funct. Mater.* **2015**, *25*, 1127–1136. [[CrossRef](#)]
11. Ahmadi, A.; Shoushtari, M.Z.; Farbod, M. Photoelectrochemical Application of WS<sub>2</sub> Nanosheets Prepared via a Low-Temperature CVD Method. *J. Mater. Sci. Mater. Electron.* **2019**, *30*, 6342–6349. [[CrossRef](#)]
12. Chen, Y.; Sun, M. Two-Dimensional WS<sub>2</sub>/MoS<sub>2</sub> Heterostructures: Properties and Applications. *Nanoscale* **2021**, *13*, 5594–5619. [[CrossRef](#)] [[PubMed](#)]
13. Thiehmmed, Z.; Shakoor, A.; Altahtamouni, T. Recent Advances in WS<sub>2</sub> and Its Based Heterostructures for Water-Splitting Applications. *Catalysts* **2021**, *11*, 1283. [[CrossRef](#)]
14. Pan, Y.; Zheng, F.; Wang, X.; Qin, H.; Liu, E.; Sha, J.; Zhao, N.; Zhang, P.; Ma, L. Enhanced Electrochemical Hydrogen Evolution Performance of WS<sub>2</sub> Nanosheets by Te Doping. *J. Catal.* **2020**, *382*, 204–211. [[CrossRef](#)]

15. Han, A.; Zhou, X.; Wang, X.; Liu, S.; Xiong, Q.; Zhang, Q.; Gu, L.; Zhuang, Z.; Zhang, W.; Li, F.; et al. One-Step Synthesis of Single-Site Vanadium Substitution in 1T-WS<sub>2</sub> Monolayers for Enhanced Hydrogen Evolution Catalysis. *Nat. Commun.* **2021**, *12*, 709. [[CrossRef](#)]
16. Ahmadi, A.; Zargar Shoushtari, M. Enhancing the Photoelectrochemical Water Splitting Performance of WS<sub>2</sub> Nanosheets by Doping Titanium and Molybdenum via a Low Temperature CVD Method. *J. Electroanal. Chem.* **2019**, *849*, 113361. [[CrossRef](#)]
17. Kibsgaard, J.; Chen, Z.; Reinecke, B.N.; Jaramillo, T.F. Engineering the Surface Structure of MoS<sub>2</sub> To Preferentially Expose Active Edge Sites For Electro catalysis. *Nat. Mater.* **2012**, *11*, 963–969. [[CrossRef](#)]
18. Deokar, G.; Rajput, N.S.; Vancsó, P.; Ravoux, F.; Jouiad, M.; Vignaud, D.; Cecchet, F.; Colomer, J.F. Large Area Growth of Vertically Aligned Luminescent MoS<sub>2</sub> nanosheets. *Nanoscale* **2017**, *9*, 277–287. [[CrossRef](#)]
19. Mishra, A.K.; Lakshmi, K.V.; Huang, L. Eco-Friendly Synthesis of Metal Dichalcogenides Nanosheets and Their Environmental Remediation Potential Driven by Visible Light. *Sci. Rep.* **2015**, *5*, 15718. [[CrossRef](#)]
20. Hai, X.; Chang, K.; Pang, H.; Li, M.; Li, P.; Liu, H.; Shi, L.; Ye, J. Engineering the Edges of MoS<sub>2</sub> (WS<sub>2</sub>) Crystals for Direct Exfoliation into Monolayers in Polar Micromolecular Solvents. *J. Am. Chem. Soc.* **2016**, *138*, 14962–14969. [[CrossRef](#)]
21. Chen, Y.; Ren, R.; Wen, Z.; Ci, S.; Chang, J.; Mao, S.; Chen, J. Superior Electrocatalysis for Hydrogen Evolution with Crumpled Graphene/Tungsten Disulfide/Tungsten Trioxide Ternary Nanohybrids. *Nano Energy* **2018**, *47*, 66–73. [[CrossRef](#)]
22. Li, C.; Cao, Q.; Wang, F.; Xiao, Y.; Li, Y.; Delaunay, J.J.; Zhu, H. Engineering Graphene and TMDs Based van Der Waals Heterostructures for Photovoltaic and Photoelectrochemical Solar Energy Conversion. *Chem. Soc. Rev.* **2018**, *47*, 4981–5037. [[CrossRef](#)]
23. Shifa, T.A.; Wang, F.; Cheng, Z.; Zhan, X.; Wang, Z.; Liu, K.; Safdar, M.; Sun, L.; He, J. A Vertical-Oriented WS<sub>2</sub> Nanosheet Sensitized by Graphene: An Advanced Electrocatalyst for Hydrogen Evolution Reaction. *Nanoscale* **2015**, *7*, 14760–14765. [[CrossRef](#)] [[PubMed](#)]
24. Balandin, A.A.; Ghosh, S.; Bao, W.; Calizo, I.; Teweldebrhan, D.; Miao, F.; Lau, C.N. Superior Thermal Conductivity of Single-Layer Graphene. *Nano Lett.* **2008**, *8*, 902–907. [[CrossRef](#)] [[PubMed](#)]
25. Castro Neto, A.H.; Guinea, F.; Peres, N.M.R.; Novoselov, K.S.; Geim, A.K. The Electronic Properties of Graphene. *Rev. Mod. Phys.* **2009**, *81*, 109–162. [[CrossRef](#)]
26. Das Sarma, S.; Adam, S.; Hwang, E.H.; Rossi, E. Electronic Transport in Two-Dimensional Graphene. *Rev. Mod. Phys.* **2011**, *83*, 407–470. [[CrossRef](#)]
27. Zhou, H.; Yu, F.; Sun, J.; He, R.; Wang, Y.; Guo, C.F.; Wang, F.; Lan, Y.; Ren, Z.; Chen, S. Highly active and durable self-standing WS<sub>2</sub>/graphene hybrid catalysts for hydrogen evolution reaction. *J. Mater. Chem. A* **2016**, *4*, 9472–9476. [[CrossRef](#)]