

## Article

# Synthesis of Phosphonated Carbon Nanotubes: New Insight into Carbon Nanotubes Functionalization

Małgorzata Nadolska <sup>1</sup>, Marta Prześniak-Welenc <sup>1</sup> , Marcin Łapiński <sup>1</sup>  and Kamila Sadowska <sup>2,\*</sup> 

<sup>1</sup> Institute of Nanotechnology and Materials Engineering, Gdansk University of Technology, 80-233 Gdansk, Poland; malgorzata.nadolska@pg.edu.pl (M.N.); marwelen@pg.edu.pl (M.P.-W.); marcin.lapinski@pg.edu.pl (M.L.)

<sup>2</sup> Institute of Biocybernetics, Biomedical Engineering of the Polish Academy of Sciences, 02-109 Warsaw, Poland

\* Correspondence: ksadowska@ibib.waw.pl or kamsadow@pg.edu.pl

**Abstract:** Carbon nanotubes were successfully functionalized for the first time in a free radical phosphonylation reaction. Three synthetic protocols were proposed. Carbon nanotubes and diethylphosphite reacted in the presence of known radical initiator, such as azobisisobutyronitrile, single electron oxidant—Mn(OAc)<sub>3</sub>, or under UV radiation. The functionalized material was fully characterized by means of spectroscopic methods, together with microscopic, surface area and thermogravimetric analyses. UV-illumination was found to be the most effective approach for introducing phosphonates onto carbon nanotubes. X-ray photoelectron spectroscopy analysis showed 6% phosphorus in this sample. Moreover, the method was performed at room temperature for only one hour, using diethylphosphite as a reactant and as a solvent. The functionalized carbon nanotubes showed an improved thermal stability, with a decomposition onset temperature increase of more than 130 °C. This makes it very promising material for flame retarding applications.

**Keywords:** carbon nanotubes; chemical functionalization; phosphorus containing groups; phosphonates; free-radical reactions



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## 1. Introduction

Carbon nanotubes (CNTs) receive a great deal of attention due to their impressive properties and variety of applications. Owing to their large specific surface area and high electrical conductivity, CNTs are promising candidates for sensors [1,2], energy storage materials [3,4], and catalyst supports [5,6]. A well-developed surface combined with good chemical stability also makes them effective pollutant adsorbents [7,8] and excellent drug and gene carriers [9,10]. However, in most cases, CNTs cannot be used in their pristine form and their surface properties usually need to be tailored to specific application.

Surface functionalization is a powerful tool that allows to adjust the physical and chemical properties of CNTs, and, as a consequence, increase their processability and applicability. There are many approaches that have been applied for the functionalization of CNTs, including covalent and noncovalent modifications using chemical and physical methods (plasma treating, thermal annealing) [11–14]. One key area of research deals with the introduction of different functional groups onto the surface of CNTs. The most common are definitely oxygen functional groups (e.g., hydroxyl, carboxyl, carbonyl); however, many examples of CNTs with nitrogen (e.g., amine, amide) and sulfur-containing (e.g., thiol, sulfonic) functional groups can also be found in the literature [15–18]. Contrary to the examples mentioned above, phosphorus-containing groups, especially those with a phosphorus atom attached directly to the carbon skeleton, are rarely reported. At the same time, organic compounds containing C–P bonds find a number of applications, such as corrosion inhibitors [19,20], detergents [21], flame retardants [22,23], and others. It is also well known that such compounds possess a high affinity towards various metal ions and can be used in waste water treatment [24,25]. C–P bonds are also widely found in many

biologically active compounds, with applications ranging from enzyme inhibitors to bone-seeking pharmaceuticals [26,27]. The vast majority of literature reports include noncovalent approaches, e.g.,  $\pi$ - $\pi$  stacking interactions between naphthalen-1-ylmethylphosphonic acid and CNTs [28,29] or the attachment of phosphonates via linkage (e.g., amide or carbonyl bonds [30–34]). Such connections are prone to hydrolysis and can suffer from insufficient thermal stability. In the literature, there are only a few examples described in which phosphorus-containing groups are directly connected to the carbon of nanotubes. Dehghani and co-workers used chemical vapor deposition method to obtain multi-walled carbon nanotubes functionalized with phosphonic acid [35,36]. The functionalization degree of such a product was high (11.65%), but the synthesis protocol required a high temperature (1300 °C) and a special system to guarantee a controlled atmosphere and proper gas flow. CNTs functionalized with triphenylphosphine have also been reported [37–39]. The conversion of the carboxyl groups present in CNTs into phosphonic ones was proposed by two groups to produce bis-phosphonic derivatives [40,41].

In the present work, we grafted phosphorus-containing groups directly to the carbon skeleton in a free-radical phosphorylation reaction. It should be noted that the reaction of phosphorus radicals with  $sp^2$  carbon to form C–P bond compounds is known from classical organic chemistry [42]. However, studies on the reaction of phosphorus radicals with carbonaceous nanostructures are rare. To the best of our knowledge, reported works include only fullerenes [43,44]. Herein, we adopt this method for multi-walled carbon nanotubes (MWCNTs). Moreover, we proposed two new protocols, in which the formation of phosphorus-centred free radicals was thermally generated in the presence of an initiator (azobisisobutyronitrile (AIBN)) or by means of UV-irradiation, respectively. The functionalized MWCNTs were thoroughly characterized using spectroscopic (Fourier-Transform Infrared Spectroscopy (FTIR), X-ray Photoelectron Spectroscopy (XPS) and Energy Dispersive X-ray spectroscopy (EDX) and microscopic methods (Scanning Electron Microscopy (SEM)). Furthermore, thermal stability and surface area (adsorption–desorption nitrogen isotherms) analyses have been carried out.

## 2. Materials and Methods

Multi-walled carbon nanotubes (MWCNTs, average outer diameter: 8–15 nm, purity >95 wt%) were ordered from Cheaptubes Inc., Grafton, VT, USA. Diethyl phosphite ( $(C_2H_5O)_2P(O)H$ ), manganese (III) acetate dihydrate ( $Mn(OAc)_3 \cdot 2H_2O$ ), azobisisobutyronitrile (AIBN) and 1,2-dichlorobenzene, tetrahydrofuran (THF) and N,N-dimethylformamide (DMF) were purchased from Acros Organics, Antwerp, Belgium. The chemicals were of analytical grade and were used without further purification.

Multi-walled carbon nanotubes were functionalized through free-radical reactions. In all syntheses, the same amount of carbon material (100 mg) and diethyl phosphite (3 mL), which acted as a phosphate group source, were used. Reactions were carried out in the presence of known radical initiators, such as  $Mn(OAc)_3$  and AIBN, or under UV radiation. The UV-promoted reaction was carried out in a photoreactor prototype designed by Dariusz Wysiecki MSc., Eng. and constructed in cooperation with the Enviklim Company (Gdańsk, Poland). The reactor was equipped with 3 UVA diode arrays: 2xUV-D6565-4LED, 40 W and 1xUV-D6565-15LED, 150 W (Enviklim Company, Gdańsk, Poland) with a 365–370 nm wavelength [45,46]. Detailed conditions are summarized in Table 1. Afterwards, products were washed with DMF and THF, and dried under reduced pressure. The obtained functionalized MWCNTs were labelled as MWCNT-Phos-Mn and MWCNT-Phos-AIBN, in the case of using  $Mn(OAc)_3$  and AIBN, respectively, and MWCNT-Phos-UV when UV-irradiation was used.

**Table 1.** Detailed conditions of the functionalization reactions.

Sample ID	MWCNTs	Radical Initiator	Phosphate Groups Source	Reaction Medium	Atm	Temp. (°C)	Time (h)
MWCNT-Phos-Mn	100 mg	90 mg Mn(OAc) <sub>3</sub> ·2H <sub>2</sub> O	3 mL of (C <sub>2</sub> H <sub>5</sub> O) <sub>2</sub> P(O)H	3 mL of 1,2-dichlorobenzene	Ar	135	5
MWCNT-Phos-AIBN	100 mg	(150 mg + 150 mg) AIBN	3 mL of (C <sub>2</sub> H <sub>5</sub> O) <sub>2</sub> P(O)H	(C <sub>2</sub> H <sub>5</sub> O) <sub>2</sub> P(O)H	Ar	65	5
MWCNT-Phos-UV	100 mg	UV radiation (365–370 nm)	3 mL of (C <sub>2</sub> H <sub>5</sub> O) <sub>2</sub> P(O)H	(C <sub>2</sub> H <sub>5</sub> O) <sub>2</sub> P(O)H	Air	25	1

### Characterization

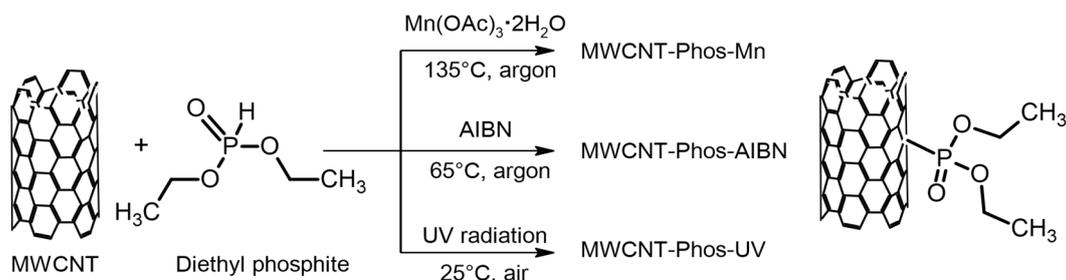
The morphologies of the samples were studied with a scanning electron microscope (ESEM Quanta Feg 250, FEI, Waltham, MA, USA). Fourier transform infrared (FTIR) spectra were collected on a Perkin Elmer Frontier spectrophotometer (Waltham, MA, USA) in the range of 500–4000 cm<sup>-1</sup>. Measurements were made in transmittance mode and the potassium bromide pellet method was used. X-ray diffraction patterns were collected on a Bruker D2 Phaser 2nd generation diffractometer with CuK<sub>α</sub> radiation ( $\lambda = 1.5404 \text{ \AA}$ ) with  $2\theta$  ranging from 5 to 70°. The Raman spectra were obtained with an integrated confocal micro-Raman system with a LabRamAraxis (Horiba Jobin Yvon, Tokyo, Japan) 460-mm spectrometer equipped with a confocal microscope. The excitation source was a diode pumped solid state (DPSS) laser emitting light at 632.8 nm. The surface chemical composition was studied using an Omicron NanoTechnology spectrometer (ScientaOmicron, Uppsala, Sweden) with Mg K<sub>α</sub> as an excitation source. The binding energies were corrected using the background C1s (285.0 eV) line as a reference. XPS spectra were analyzed with Casa-XPS software (Casa Software Ltd., ver. 2.3.23., Devon, UK) using Shirley background subtraction and Gaussian–Lorentzian curve as a fitting algorithm. Thermogravimetric analysis (TGA) was performed with a Netzsch STA 449 F1 (Netzsch, Selb, Germany). Experiments were carried out with a heating rate 10 °C/min from 40 °C to 900 °C under an argon atmosphere and under synthetic air. The N<sub>2</sub> adsorption–desorption isotherms were measured on a NOVAtouch™ 2 surface area analyser (Quantachrome Instruments, Boynton Beach, FL, USA) at 77 K. Prior to the measurements, samples were degassed under vacuum at 40 °C for 12 h. Specific surface area was calculated from the Brunauer–Emmett–Teller (BET) linear equation in the range of 0.1–0.3 relative pressure. The correlation coefficient of the linear regression was not less than 0.999.

### 3. Results and Discussion

In the present work, we present three ways of bonding of phosphorus-containing groups directly to the carbon skeleton of nanotubes by using phosphorus-centred radicals. It should be noticed, that the reaction of phosphorus-centred radicals with sp<sup>2</sup> carbon to form C–P bonds compounds is known from classical organic chemistry [42]. Homolytic cleavage of P–H bonds serves as a major approach for the generation of phosphorus-centred radicals. Phosphonyl radicals are the most popular for use in the generation of C–P bonds with alkenes, alkynes, arenes, and heteroarenes [47,48], and they can be generated from corresponding phosphonates under appropriate radical initiation conditions [49–51]. Generally, phosphorus-centred radicals can be generated via thermolysis or photolysis, and/or in the presence of well-known initiators, such as peroxides, azo compounds, or photo radical initiators. The reactions with single electron oxidants, such as cerium ammonium nitrate, Mn(OAc)<sub>2</sub>/O<sub>2</sub> or Mn(OAc)<sub>3</sub>, and AgOAc/K<sub>2</sub>S<sub>2</sub>O<sub>6</sub> have also been reported.

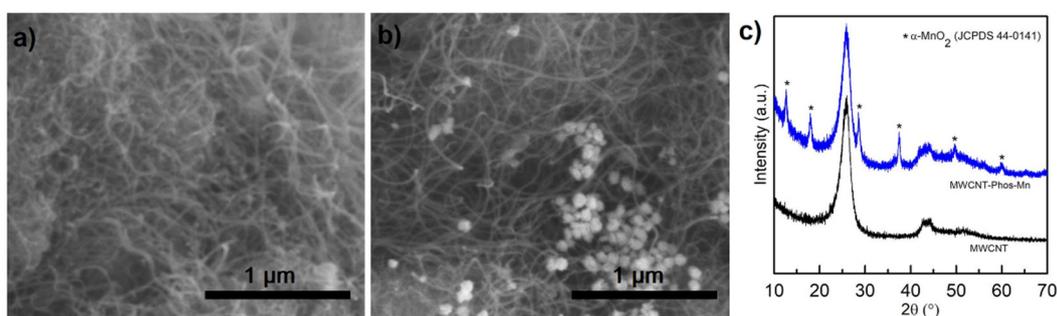
Interestingly, since the pioneering work of Jeffrey L. Bahr and James M. Tour [52], carbon-centred radicals have been frequently used for carbon nanostructures functionalization, while there is only one example in the literature describing the reaction of phosphorus-centred radicals with carbon nanomaterials. Wang and co-workers reported the Mn(OAc)<sub>3</sub>-promoted reaction of C60 with phosphonate esters or phosphine oxide.

Depending on the reaction conditions (i.e., molar ratio of the substrates) they obtained three different types of phosphorylated fullerenes, that is, singly bonded fullerene dimers, hydrofullerenes, and acetoxyated fullerenes [43,44]. Based on knowledge from organic chemistry and our experience in the free-radical functionalization of carbon nanotubes using diazonium salts [39,53–57], we proposed three synthesis pathways for covalent functionalization of MWCNTs using diethyl phosphite, which are presented in Figure 1.



**Figure 1.** Three pathways leading to phosphonated MWCNTs.

The first synthesis was following the work of Wang; however, using carbon nanotubes in the place of fullerenes. The reaction was performed under argon, in 1,2-dichlorobenzene with  $\text{Mn}(\text{OAc})_3$  as a single electron oxidant, promoting  $(\text{EtO})_2\text{P}(\text{O})\cdot$  formation from  $(\text{EtO})_2\text{P}(\text{O})\text{H}$  at an elevated temperature (135 °C). The functionalization was successful; however, a detailed analysis of the obtained material revealed the presence of a by-product, which was impossible to remove. Figure 2a,b shows SEM images of MWCNTs before and after functionalization. It can be clearly seen that the carbon material is decorated with particles, which were not present in the pristine nanotubes. XRD study proved our assumption, that it was  $\text{MnO}_2$  formed as a by-product of  $\text{Mn}(\text{OAc})_3$  decomposition. A comparison of diffractograms recorded for pristine MWCNTs and the MWCNT-Phos-Mn sample is presented in Figure 2c. Typical (002) and (100) reflexes of MWCNT are present at 26° and 43° in both diffractograms; however, additional reflexes are visible only for the functionalized sample. The reflexes marked with asterisks refer to an  $\alpha\text{-MnO}_2$  crystal structure (Joint Committee on Powder Diffraction Standards (JCPDS)44-0141). Spectroscopic and thermal analyses revealed other differences and they will be discussed later in comparison to the material obtained using the two other proposed synthesis pathways.



**Figure 2.** SEM images of MWCNT (a) and MWCNT-Phos-Mn (b) and their XRD patterns (c).

As the obtained material, although interesting, was not what we expected, another synthetic protocol was proposed. In a second approach, thermal decomposition of azobisisobutyronitrile (AIBN) served as a free radical reaction initiator to further produce phosphonyl radicals from diethyl phosphite. The sample was heated for 5 h under argon at 65 °C in diethylphosphite, which served as a reagent and a solvent. The obtained sample was denoted as MWCNT-Phos-AIBN. The final method was based on photo-induced generation of phosphonyl radicals from diethyl phosphite in a solvent-less reaction carried

out for 1 h at RT under UV light illumination ( $P = 230 \text{ W}$ ;  $\lambda = 365\text{--}370 \text{ nm}$ ). The sample was labelled as MWCNT-Phos-UV.

Similar to the MWCNT-Phos-Mn sample, the other two were also observed under SEM. The obtained images can be seen in Figure 3. No significant structural changes were visible for the studied samples in reference to the pristine nanotubes. However, large area imaging presented a higher purity of functionalized samples in terms of a lower presence of amorphous carbon. It is frequently reported, that nanotubes are purified during a functionalization reaction and washing after synthesis [58,59].

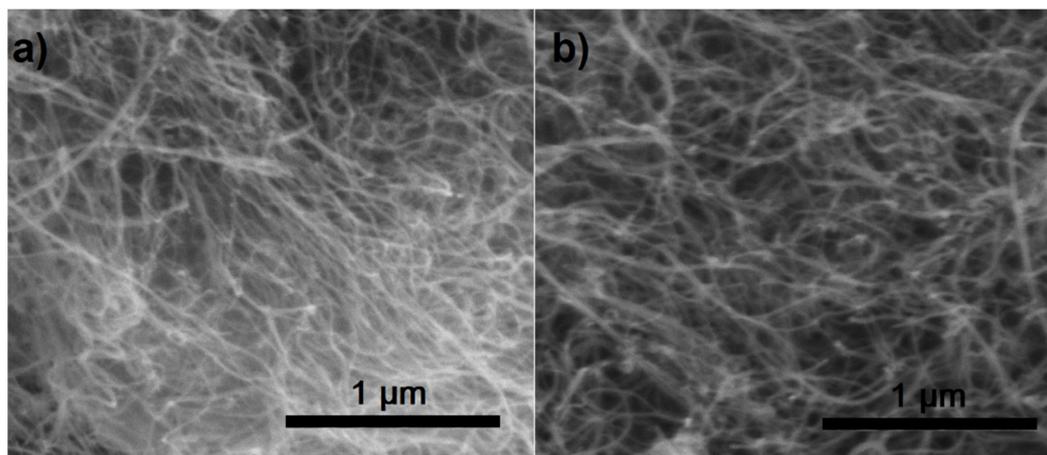


Figure 3. SEM images of MWCNT-Phos-AIBN (a) and MWCNT-Phos-UV (b).

Pristine and functionalized MWCNTs were characterized using Raman, FTIR, and XPS spectroscopy. Raman spectra of MWCNTs samples recorded at 633 nm excitation wavelength are presented in Figure 4. Raman spectra of the pristine nanotubes are in accordance with reported data for this type of material and excitation energy [60–62]. Typical D and G bands were recorded for all samples. A comparison of Raman mode positions, full width at half maximum (FWHM), and  $I_D/I_G$  ratio are presented in Table 2. The decrease in the  $I_D/I_G$  ratio was ascribed to the removal of amorphous carbon and the most defective MWCNTs during washing after functionalization [63,64]. Both the D band and G band of functionalized MWCNTs are widened, indicating the introduction of functional groups into the nanotube structures [65]. Moreover, the G bands of MWCNT-Phos-AIBN and MWCNT-Phos-UV are shifted to a lower wavenumber. Similar results were obtained by Sun et al. [66] and described as evidence of the formation of C-P bonds.

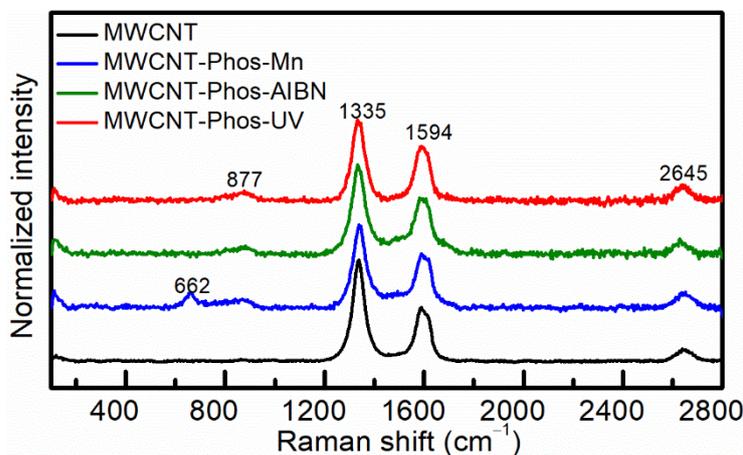


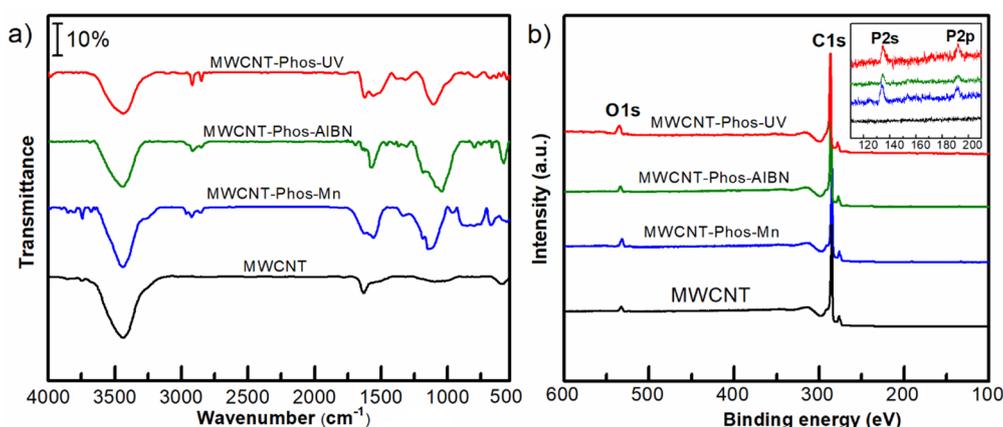
Figure 4. Raman spectra of pristine and functionalized MWCNTs.

**Table 2.** Raman mode positions, linewidths (FWHM), and  $I_D/I_G$  ratios for MWCNT, MWCNT-Phos-Mn, MWCNT-Phos-AIBN, and MWCNT-Phos-UV.

	D Band		G Band		$I_D/I_G$
	Position (cm <sup>-1</sup> )	FWHM (cm <sup>-1</sup> )	Position (cm <sup>-1</sup> )	FWHM (cm <sup>-1</sup> )	
MWCNT	1337	57	1596	69	1.88
MWCNT-Phos-Mn	1340	62	1597	77	1.53
MWCNT-Phos-AIBN	1334	65	1593	81	1.64
MWCNT-Phos-UV	1333	63	1593	72	1.49

After functionalization, a new band at 877 cm<sup>-1</sup> appeared in the spectra of functionalized samples, being most evident for MWCNT-Phos-AIBN and MWCNT-Phos-UV. This can be ascribed to a superposition of bands, referring to C–O–P, C–P and C–C–O stretching. The 662 cm<sup>-1</sup> band, visible only in the MWCNT-Phos-Mn spectrum is additional evidence of the presence of MnO<sub>2</sub> in this sample. According to the literature, the spectrum of  $\alpha$ -MnO<sub>2</sub> exhibits peaks in the region of 400–800 cm<sup>-1</sup>, which are ascribed to the stretching mode of the MnO<sub>6</sub> octahedra. The highest band present in standard  $\alpha$ -MnO<sub>2</sub> around 650 cm<sup>-1</sup> could be attributed to the symmetric stretching vibration (Mn–O) of the MnO<sub>6</sub> groups. In MWCNT-Phos-Mn the band is observed at 662 cm<sup>-1</sup> and its shift to higher wavenumbers may be due to the presence of carbon material in the sample. A similar behaviour was observed for MnO<sub>2</sub>/graphene composites by Liu et al. [67].

The presence of functional groups was evidenced by FTIR and XPS spectroscopy. The FTIR spectra of functionalized nanotubes in comparison to pristine MWCNTs are presented in Figure 5a. In the spectrum of pristine MWCNTs, a small band originating from carbon–carbon bond vibrations is visible, next to well-pronounced water infrared bands at 3490 and 1630 cm<sup>-1</sup> [68]. For all functionalized samples, bands at 2914 and 2854 cm<sup>-1</sup> referring to C–H stretching in the ethoxy group of phosphonate are clearly visible. In the lower frequency region, C–P, P–O and P=O bands are present at 1450, 1180 and 1040 cm<sup>-1</sup>, respectively [69,70]. In the MWCNT-Phos-Mn sample, multiple bands in the range of 900–600 cm<sup>-1</sup>, resulting from Mn–O–Mn stretching are observed. This is additional evidence of the presence of MnO<sub>2</sub> by-product in this sample [71].

**Figure 5.** FTIR (a) and XPS survey (b) spectra of pristine and functionalized MWCNTs.

The functionalization of carbonaceous material with phosphonates was additionally proved using X-ray photoelectron spectroscopy (XPS). The survey spectra of all samples are presented in Figure 5b. It can be seen, that the C to O ratio decreased in the functionalized samples and was lowest in the case of MWCNT-Phos-UV. This indicates, that oxygen content increased due to the introduction of  $-P(O)(OEt)_2$  groups and the highest

functionalization was obtained in the UV-promoted reaction. The inset in Figure 6 shows magnified P 2s and P 2p energy regions. Again, the highest signals were obtained for the MWCNT-Phos-UV sample, while pristine MWCNT did not contain phosphorus. Moreover, grafting of phosphorous-containing groups onto the MWCNT surface was confirmed by the deconvolution of high resolution C 1s spectra and analysis of carbon binding states (Figure 6). Pristine MWCNTs revealed four components C=C (284.3 eV), C-C (285.3 eV), C-O (287.7 eV), and  $\pi-\pi^*$  (290.3 eV), indicating the presence of amorphous carbon and a small amount of oxygen contamination in the sample [72,73]. After functionalization, a new peak appeared at 286.4 eV, which can be assigned to the C-P [32] and is evidence of direct bonding between the carbon skeleton and the functional group. In addition, an increase in C-O peak was observed due to the presence of C-O-P bonds from the introduced group. The highest phosphorus content, ca. 6% at. was recorded for MWCNT-Phos-UV sample, which is in agreement with data obtained in other studies.

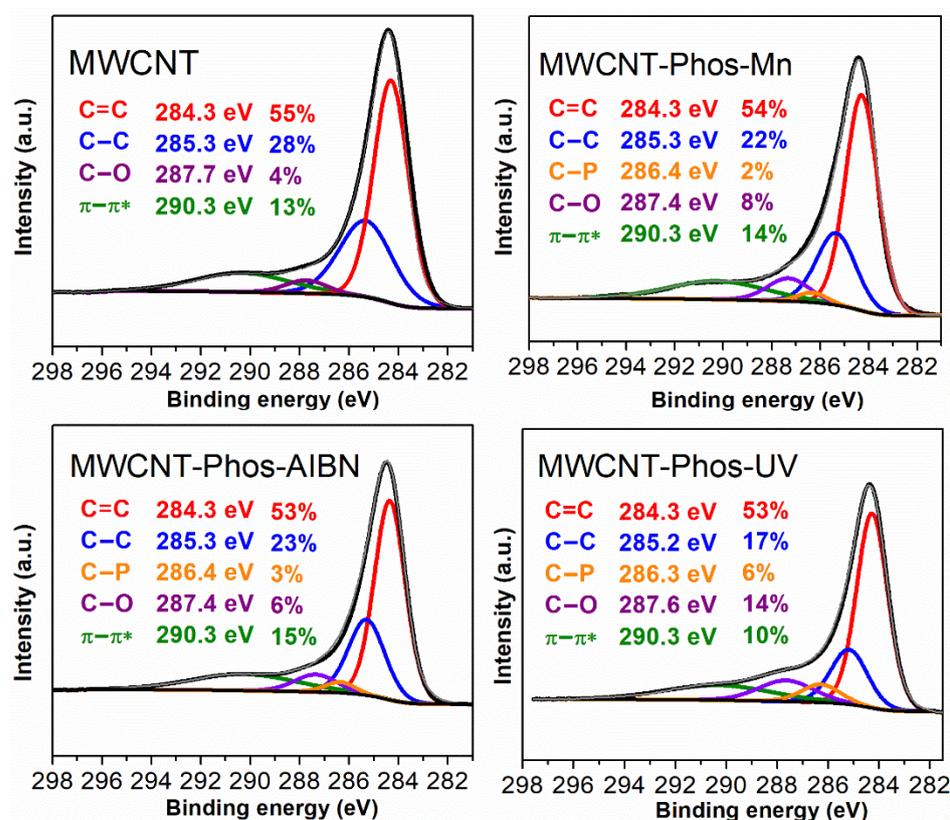
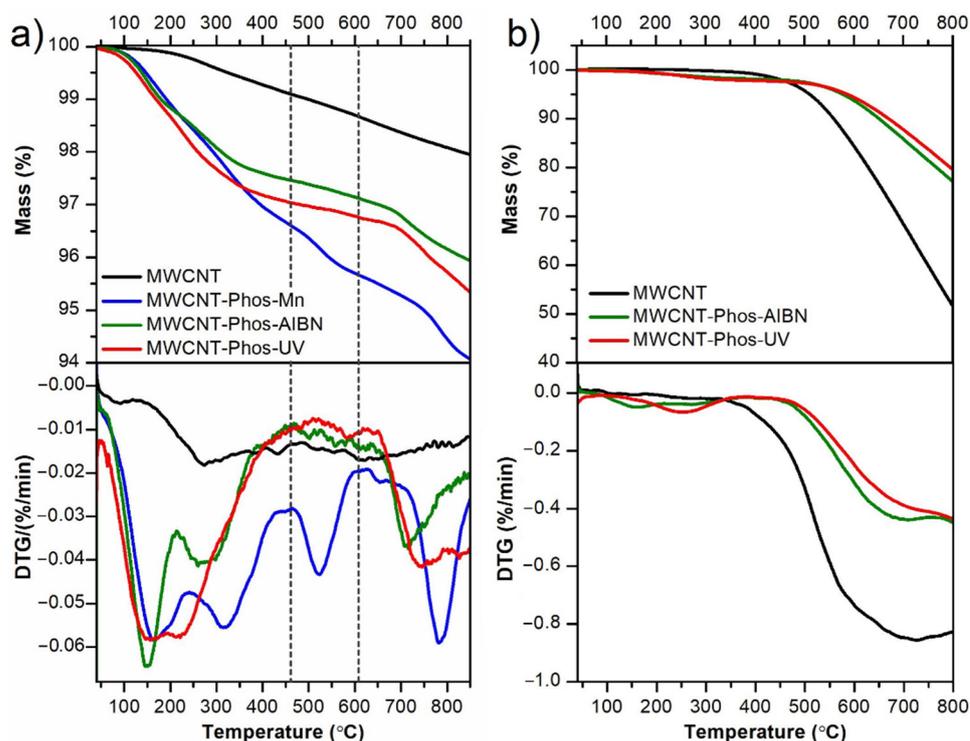


Figure 6. High resolution XPS spectra of C1s with peak positions and calculated atomic percentages.

Thermogravimetric analysis conducted in an argon environment revealed differences in the thermal stabilities of the studied samples (Figure 7a). The highest mass loss in the studied temperature range was observed for the MWCNT-Phos-Mn sample. The characteristic steps for the decomposition cascade:  $\text{MnO}_2 \rightarrow \text{Mn}_2\text{O}_3 \rightarrow \text{Mn}_3\text{O}_4$  were observed in the thermogravimetric (TG) curve, together with the corresponding minima (520, 788 °C) in the derivative thermogravimetric (DTG) curve [74]. Unfortunately, because of the presence of  $\text{MnO}_2$  in this functionalized MWCNT sample, it was not possible to determine the number of functional groups introduced during synthesis. The TG curves for the two other samples were similar, showing, however, a higher functionalization degree for MWCNT-Phos-UV. The lowest contamination of the MWCNT-Phos-UV sample with amorphous carbon was also proved by this study. The comparison of the observed mass loss for all samples is given in Table 3. The obtained mass loss for the MWCNT-Phos-UV, due to the decomposition of the functional groups, slightly exceeded 3%. This quite-low mass loss can be justified by the flame-retarding properties reported for carbon materials

with phosphorus-containing groups [59,75,76]. When phosphonated, carbon nanotubes are heated and a char layer is formed on the surface, which shields the material, preventing the formation of volatile moieties. Therefore, the recorded mass loss during TG analysis cannot be directly connected with the number of phosphonic groups attached to the carbon skeleton. To further prove this assumption, MWCNT-Phos-UV and MWCNT-Phos-AIBN were also studied under oxidative conditions. TG curves recorded in air are presented in Figure 7b. Compared to the pristine MWCNTs, functionalized samples show better thermal stability and higher char yield. It is worth noting, that the experiments were conducted in partial oxygen to be closer to environmental conditions. Therefore, incomplete combustion of CNTs and functionalized CNTs can be read out of the TG curves (Figure 7b). A similar behavior was observed by Mahajan et al. [77]. They observed ca. 50%-mass loss of CNTs heated under partial oxygen with a heating rate of 5 °C/min, which is in line with our observations. Incorporation of phosphorus-containing groups results in an increase in the decomposition onset temperature, from 527 °C for MWCNTs to 639 °C and 652 °C for MWCNT-Phos-AIBN and MWCNT-Phos-UV, respectively. In addition, the first derivative of the TG curves (DTG, Figure 7b) revealed that the decomposition rate is almost two times slower for functionalized MWCNTs. The above results indicate that even a low percentage of functionalization significantly improves the thermal properties of MWCNTs and makes them promising materials as flame retarding additives.

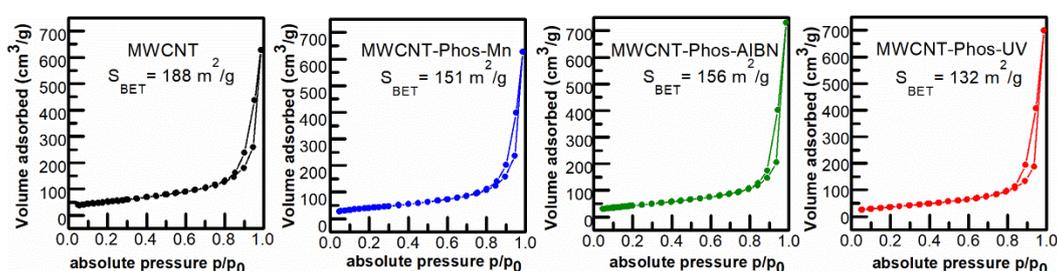


**Figure 7.** (a) TG (upper) and DTG (lower) curves recorded for pristine and functionalized MWCNTs in argon; (b) TG curves of MWCNT, MWCNT-Phos-AIBN and MWCNT-Phos-UV registered in synthetic air.

**Table 3.** Comparison of the mass loss and differential thermal analysis of studied samples.

Sample Name	Temperature Range (°C)		
	40–450	450–600	600–850
Mass Loss in % (DTG Peak Position (°C))			
MWCNT	0.87 (100, 270)	0.43 (-)	0.75 (-)
MWCNT-Phos-Mn	3.33 (166, 317)	0.97 (520)	1.63 (780)
MWCNT-Phos-AIBN	2.51 (148, 272)	0.34 (-)	1.20 (710)
MWCNT-Phos-UV	2.93 (158, 220)	0.29 (-)	1.43 (740)

The surface area of the MWCNTs and functionalized MWCNTs were evaluated by  $N_2$  isothermal adsorption (Figure 8).

**Figure 8.**  $N_2$  adsorption–desorption isotherms and corresponding surface area of pristine and functionalized MWCNTs.

The adsorption–desorption isotherms represent a type IV isotherm, typical for multi-walled carbon nanotubes [78,79]. The observed hysteresis loop (0.8–0.99  $p/p_0$ ) is associated with the mesoporous structure and corresponds well to the diameter of the used MWCNTs. After modification, the Brunauer–Emmett–Teller (BET) surface area of the MWCNT samples decreased from 188 to 132  $m^2/g$  for MWCNT–Phos–UV. A smaller decrease was observed for the two other samples, namely MWCNT–Phos–Mn and MWCNT–Phos–AIBN, of which the surface areas were equal to 151 and 156  $m^2/g$ , respectively. The results obtained by all methods described above confirmed the highest functionalization level of MWCNT–Phos–UV sample, and the presence of contaminant in the form of  $MnO_2$  in the MWCNT–Phos–Mn sample. Therefore, the surface area of MWCNT–Phos–Mn cannot be compared with pristine MWCNTs or other samples as it resulted from an addition of two different materials. For the MWCNT–Phos–UV sample, Raman spectroscopy and TG analysis results demonstrated the most effective removal of amorphous carbon and probably the shortest and most defective MWCNTs, which caused a decrease in surface area.

#### 4. Conclusions

Multiwalled carbon nanotubes were successfully functionalized in a free-radical phosphonylation reaction. Thermolysis of the P–H bond in diethyl phosphite was enhanced by two promoters: azo compound and single electron oxidant  $Mn(OAc)_3$ . Photolysis of diethyl phosphite without any promoters also succeeded in homolytic cleavage of P–H bonds. It should be noted, that usage of  $Mn(OAc)_3$  was not suitable for the reactions of carbon nanotube functionalization, as the product of its decomposition contaminated the functionalized carbon material. Two other methods resulted in phosphonated carbon nanotubes, with a higher functionalization degree for the MWCNT–Phos–UV sample. UV illumination was found to be the most effective method for free-radical functionalization of the carbon nanotubes. Moreover, as compared to the other two protocols, the reaction time was substantially shorter and the temperature of the reaction medium did not exceed 40 °C. Therefore, this method can be used for temperature-sensitive materials. Functionalization with phosphonates significantly improve the thermal properties of MWCNTs and makes them promising materials as flame retardants additives.

The novel approach described here complements existing protocols for carbon nano-material functionalization. Moreover, it opens new possibilities for surface functionalization and the application of carbon nanotubes and other carbon nanostructures.

**Author Contributions:** Conceptualization, K.S. and M.N.; methodology, M.N., M.P.-W. and M.L.; investigation M.N., M.P.-W. and M.L.; writing—original draft preparation, K.S. and M.N.; visualization, M.N.; supervision, K.S.; project administration, K.S.; funding acquisition, K.S. and M.N. All authors have read and agreed to the published version of the manuscript.

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## References

1. Camilli, L.; Passacantando, M. Advances on Sensors Based on Carbon Nanotubes. *Chemosensors* **2018**, *6*, 62. [[CrossRef](#)]
2. Schroeder, V.; Savagatrup, S.; He, M.; Lin, S.; Swager, T.M. Carbon Nanotube Chemical Sensors. *Chem. Rev.* **2019**, *119*, 599–663. [[CrossRef](#)]
3. Wen, L.; Li, F.; Cheng, H.-M. Carbon Nanotubes and Graphene for Flexible Electrochemical Energy Storage: From Materials to Devices. *Adv. Mater.* **2016**, *28*, 4306–4337. [[CrossRef](#)] [[PubMed](#)]
4. Antiohos, D.; Romano, M.; Chen, J.; Razal, J.M. Carbon Nanotubes for Energy Applications. In *Syntheses and Applications of Carbon Nanotubes and Their Composites*; Suzuki, S., Ed.; IntechOpen: Rijeka, Croatia, 2013; Volume 22, pp. 1–13. [[CrossRef](#)]
5. Yanhui, Y.; Miao, J.; Yang, Z.; Xiao, F.-X.; Bin Yang, H.; Liu, B.; Yang, Y. Carbon nanotube catalysts: Recent advances in synthesis, characterization and applications. *Chem. Soc. Rev.* **2015**, *44*, 3295–3346. [[CrossRef](#)]
6. MabenaSuprakas, L.F.; Ray, S.S.; Mhlanga, S.D.; Coville, N.J. Nitrogen-doped carbon nanotubes as a metal catalyst support. *Appl. Nanosci.* **2011**, *1*, 67–77. [[CrossRef](#)]
7. Sarkar, B.; Mandal, S.; Tsang, Y.F.; Kumar, P.; Kim, K.-H.; Ok, Y.S. Designer carbon nanotubes for contaminant removal in water and wastewater: A critical review. *Sci. Total Environ.* **2018**, *612*, 561–581. [[CrossRef](#)]
8. Ihsanullah; Abbas, A.; Al-Amer, A.M.; Laoui, T.; Al-Marri, M.J.; Nasser, M.S.; Khraisheh, M.; Atieh, M.A. Heavy metal removal from aqueous solution by advanced carbon nanotubes: Critical review of adsorption applications. *Sep. Purif. Technol.* **2016**, *157*, 141–161. [[CrossRef](#)]
9. Karimi, M.; Solati, N.; Ghasemi, A.; Estiar, M.A.; Hashemkhani, M.; Kiani, P.; Mohamed, E.; Saeidi, A.; Taheri, M.; Avci, P.; et al. Carbon nanotubes part II: A remarkable carrier for drug and gene delivery. *Expert Opin. Drug Deliv.* **2015**, *12*, 1089–1105. [[CrossRef](#)] [[PubMed](#)]
10. Bianco, A.; Kostarelos, K.; Partidos, C.D.; Prato, M. Biomedical applications of functionalised carbon nanotubes. *Chem. Commun.* **2005**, *5*, 571–577. [[CrossRef](#)]
11. Ferreira, F.V.; Cividanes, L.; Brito, F.S.; de Menezes, B.R.C.; Franceschi, W.; Simonetti, E.A.N.; Thim, G.P. *Functionalizing Graphene and Carbon Nanotubes*, 1st ed.; Springer Nature: Basingstoke, UK, 2016; pp. 31–61.
12. Balasubramanian, K.; Burghard, M. Chemically Functionalized Carbon Nanotubes. *Small* **2005**, *1*, 180–192. [[CrossRef](#)]
13. Mallakpour, S.; Soltanian, S. Surface functionalization of carbon nanotubes: Fabrication and applications. *RSC Adv.* **2016**, *6*, 109916–109935. [[CrossRef](#)]
14. Saka, C. Overview on the Surface Functionalization Mechanism and Determination of Surface Functional Groups of Plasma Treated Carbon Nanotubes. *Crit. Rev. Anal. Chem.* **2018**, *48*, 1–14. [[CrossRef](#)] [[PubMed](#)]
15. Deline, A.R.; Frank, B.P.; Smith, C.L.; Sigmon, L.R.; Wallace, A.N.; Gallagher, M.J.; Goodwin, J.D.G.; Durkin, D.P.; Fairbrother, D.H. Influence of Oxygen-Containing Functional Groups on the Environmental Properties, Transformations, and Toxicity of Carbon Nanotubes. *Chem. Rev.* **2020**, *120*, 11651–11697. [[CrossRef](#)] [[PubMed](#)]
16. Ali, S.; Rehman, S.A.U.; Luan, H.-Y.; Farid, M.U.; Huang, H. Challenges and opportunities in functional carbon nanotubes for membrane-based water treatment and desalination. *Sci. Total Environ.* **2019**, *646*, 1126–1139. [[CrossRef](#)] [[PubMed](#)]
17. Park, S.K.; Mahmood, Q.; Park, H.S. Surface functional groups of carbon nanotubes to manipulate capacitive behaviors. *Nanoscale* **2013**, *5*, 12304–12309. [[CrossRef](#)]
18. Mao, J.; Wang, Y.; Zhu, J.; Yu, J.; Hu, Z. Thiol functionalized carbon nanotubes: Synthesis by sulfur chemistry and their multi-purpose applications. *Appl. Surf. Sci.* **2018**, *447*, 235–243. [[CrossRef](#)]

19. Zhao, R.; Rupper, P.; Gaan, S. Recent Development in Phosphonic Acid-Based Organic Coatings on Aluminum. *Coatings* **2017**, *7*, 133. [[CrossRef](#)]
20. Prabakaran, M.; Vadivu, K.; Ramesh, S.; Periasamy, V. Corrosion protection of mild steel by a new phosphonate inhibitor system in aqueous solution. *Egypt. J. Pet.* **2014**, *23*, 367–377. [[CrossRef](#)]
21. Nowack, B. Environmental chemistry of phosphonates. *Water Res.* **2003**, *37*, 2533–2546. [[CrossRef](#)]
22. Hoang, D.; Kim, J.; Jang, B.N. Synthesis and performance of cyclic phosphorus-containing flame retardants. *Polym. Degrad. Stab.* **2008**, *93*, 2042–2047. [[CrossRef](#)]
23. Liu, Y.-L.; Hsiue, G.-H.; Lee, R.-H.; Chiu, Y.-S. Phosphorus-containing epoxy for flame retardant. III: Using phosphorylated diamines as curing agents. *J. Appl. Polym. Sci.* **1997**, *63*, 895–901. [[CrossRef](#)]
24. Yudaev, P.; Kolpinskaya, N.; Chistyakov, E. Organophosphorous extractants for metals. *Hydrometallurgy* **2021**, *201*, 105558. [[CrossRef](#)]
25. Li, D. Development course of separating rare earths with acid phosphorus extractants: A critical review. *J. Rare Earths* **2019**, *37*, 468–486. [[CrossRef](#)]
26. Galezowska, J.; Gumienna-Kontecka, E. Phosphonates, their complexes and bio-applications: A spectrum of surprising diversity. *Co-ord. Chem. Rev.* **2012**, *256*, 105–124. [[CrossRef](#)]
27. Sevrain, C.M.; Berchel, M.; Couthon, H.; Jaffrès, P.-A. Phosphonic acid: Preparation and applications. *Beilstein J. Org. Chem.* **2017**, *13*, 2186–2213. [[CrossRef](#)]
28. Zheng, M.; Li, P.; Fu, G.; Chen, Y.; Zhou, Y.; Tang, Y.; Lu, T. Efficient anchorage of highly dispersed and ultrafine palladium nanoparticles on the water-soluble phosphonate functionalized multiwall carbon nanotubes. *Appl. Catal. B Environ.* **2013**, *129*, 394–402. [[CrossRef](#)]
29. Li, P.; Liu, H.; Ding, Y.; Wang, Y.; Chen, Y.; Zhou, Y.; Tang, Y.; Wei, H.; Cai, C.; Lu, T. Synthesis of water-soluble phosphonate functionalized single-walled carbon nanotubes and their applications in biosensing. *J. Mater. Chem.* **2012**, *22*, 15370–15378. [[CrossRef](#)]
30. Sainsbury, T.; Fitzmaurice, D. Templated Assembly of Semiconductor and Insulator Nanoparticles at the Surface of Covalently Modified Multiwalled Carbon Nanotubes. *Chem. Mater.* **2004**, *16*, 3780–3790. [[CrossRef](#)]
31. Zhao, B.; Hu, H.; Mandal, A.S.K.; Haddon, R. A Bone Mimic Based on the Self-Assembly of Hydroxyapatite on Chemically Functionalized Single-Walled Carbon Nanotubes. *Chem. Mater.* **2005**, *17*, 3235–3241. [[CrossRef](#)]
32. Maho, A.; Detriche, S.; Fonder, G.; Delhalle, J.; Mekhalif, Z. Electrochemical Co-Deposition of Phosphonate-Modified Carbon Nanotubes and Tantalum on Nitinol. *ChemElectroChem* **2014**, *1*, 896–902. [[CrossRef](#)]
33. Kannan, R.; Bipinlal, U.; Kurungot, S.; Pillai, V.K. Enhanced electrocatalytic performance of functionalized carbon nanotube electrodes for oxygen reduction in proton exchange membrane fuel cells. *Phys. Chem. Chem. Phys.* **2011**, *13*, 10312–10317. [[CrossRef](#)] [[PubMed](#)]
34. Oki, A.; Adams, L.; Khabashesku, V.; Edigin, Y.; Biney, P.; Luo, Z. Dispersion of aminoalkylsilyl ester or amine alkyl-phosphonic acid side wall functionalized carbon nanotubes in silica using sol-gel processing. *Mater. Lett.* **2008**, *62*, 918–922. [[CrossRef](#)]
35. Dehghani, F.; Sardarian, A.R.; Doroodmand, M.M. Preparation and characterization of multi-walled carbon nanotubes (MWCNTs), functionalized with phosphonic acid (MWCNTs-C-PO<sub>3</sub>H<sub>2</sub>) and its application as a novel, efficient, heterogeneous, highly selective and reusable catalyst for acetylation of alcohols, phenols, aromatic amines, and thiols. *J. Iran. Chem. Soc.* **2013**, *11*, 673–684. [[CrossRef](#)]
36. Dehghani, F.; Sardarian, A.R.; Doroodmand, M.M. An efficient method for synthesis of acylals from aldehydes using multi-walled carbon nanotubes functionalized with phosphonic acid (MWCNTs-C-PO<sub>3</sub>H<sub>2</sub>). *Chin. Chem. Lett.* **2014**, *25*, 1630–1634. [[CrossRef](#)]
37. Muleja, A.A.; Mbianda, X.; Krause, R.; Pillay, K. Synthesis, characterization and thermal decomposition behaviour of triphenylphosphine-linked multiwalled carbon nanotubes. *Carbon* **2012**, *50*, 2741–2751. [[CrossRef](#)]
38. Muleja, A.A.; Mbianda, X.Y.; Pillay, K.; Krause, R.W. Phosphine functionalised multiwalled carbon nanotubes: A new adsorbent for the removal of nickel from aqueous solution. *J. Environ. Sci.* **2012**, *24*, 1133–1141. [[CrossRef](#)]
39. Sadowska, K.; Jabłonowska, E.; Stolarczyk, K.; Wisner, R.; Bilewicz, R.; Roberts, K.P.; Biernat, J. Chemically modified carbon nanotubes: Synthesis and implementation. *Pol. J. Chem.* **2008**, *82*, 1309–1313.
40. Żelechowska, K.; Sobota, D.; Cieślak, B.; Przeźniak-Welenc, M.; Łapiński, M.; Biernat, J.F. Bis-phosphonated carbon nanotubes: One pot synthesis and their application as efficient adsorbent of mercury. *Full Nanotub. Carbon Nanostruct.* **2018**, *26*, 269–277. [[CrossRef](#)]
41. Maho, A.; Detriche, S.; Delhalle, J.; Mekhalif, Z. Sol-gel synthesis of tantalum oxide and phosphonic acid-modified carbon nanotubes composite coatings on titanium surfaces. *Mater. Sci. Eng. C* **2013**, *33*, 2686–2697. [[CrossRef](#)] [[PubMed](#)]
42. Pan, X.-Q.; Zou, J.-P.; Yi, W.-B.; Zhang, W. Recent advances in sulfur- and phosphorous-centered radical reactions for the formation of S-C and P-C bonds. *Tetrahedron* **2015**, *71*, 7481–7529. [[CrossRef](#)]
43. Wang, G.-W.; Wang, C.-Z.; Zhu, S.-E.; Murata, Y. Manganese(III) acetate-mediated radical reaction of [60]fullerene with phosphonate esters affording unprecedented separable singly-bonded [60]fullerene dimers. *Chem. Commun.* **2011**, *47*, 6111–6113. [[CrossRef](#)]
44. Wang, G.-W.; Wang, C.-Z.; Zou, J.-P. Radical Reaction of [60]Fullerene with Phosphorus Compounds Mediated by Manganese(III) Acetate. *J. Org. Chem.* **2011**, *76*, 6088–6094. [[CrossRef](#)] [[PubMed](#)]

45. Luboch, E.; Wagner-Wysiecka, E.; Szulc, P.; Chojnacki, J.; Szwarc-Karabyka, K.; Łukasik, N.; Murawski, M.; Kosno, M. Photochemical Rearrangement of a 19-Membered Azoxybenzocrown: Products and their Properties. *ChemPlusChem* **2020**, *85*, 2067–2083. [[CrossRef](#)]
46. Łukasik, N.; Chojnacki, J.; Luboch, E.; Okuniewski, A.; Wagner-Wysiecka, E. Photoresponsive, amide-based derivative of embonic acid for anion recognition. *J. Photochem. Photobiol. A Chem.* **2020**, *390*, 112307. [[CrossRef](#)]
47. Van der Jeught, S.; Stevens, C.V. Direct Phosphorylation of Aromatic Azaheterocycles. *Chem. Rev.* **2009**, *109*, 2672–2702. [[CrossRef](#)] [[PubMed](#)]
48. Wiemer, D.F. Synthesis of nonracemic phosphonates. *Tetrahedron* **1997**, *53*, 16609–16644. [[CrossRef](#)]
49. Leca, D.; Fensterbank, L.; Lacôte, E.; Malacria, M. Recent advances in the use of phosphorus-centered radicals in organic chemistry. *Chem. Soc. Rev.* **2005**, *34*, 858–865. [[CrossRef](#)] [[PubMed](#)]
50. Marque, S.; Tordo, P. Reactivity of Phosphorus Centered Radicals. In *New Aspects in Phosphorus Chemistry V*; Majoral, J.-P., Ed.; Springer: Berlin, Germany, 2005; pp. 43–76.
51. Bentrude, W.G. Free-Radical Reactions of Organophosphorus (3) Compounds. In *The Chemistry of Organophosphorous Compounds*; Hartley, F.R., Ed.; Wiley: Chichester, UK, 1990; Volume 1, pp. 117–125.
52. Bahr, J.L.; Tour, J.M. Highly Functionalized Carbon Nanotubes Using in Situ Generated Diazonium Compounds. *Chem. Mater.* **2001**, *13*, 3823–3824. [[CrossRef](#)]
53. Sadowska, K.; Roberts, K.; Wiser, R.; Biernat, J.; Jabłonowska, E.; Bilewicz, R. Synthesis, characterization, and electrochemical testing of carbon nanotubes derivatized with azobenzene and anthraquinone. *Carbon* **2009**, *47*, 1501–1510. [[CrossRef](#)]
54. Stolarczyk, K.; Sepelowska, M.; Lyp, D.; Żelechowska, K.; Biernat, J.F.; Rogalski, J.; Farmer, K.D.; Roberts, K.N.; Bilewicz, R. Hybrid biobattery based on arylated carbon nanotubes and laccase. *Bioelectrochemistry* **2012**, *87*, 154–163. [[CrossRef](#)]
55. Karaškiewicz, M.; Nazaruk, E.; Żelechowska, K.; Biernat, J.F.; Rogalski, J.; Bilewicz, R. Fully enzymatic mediatorless fuel cell with efficient naphthylated carbon nanotube–laccase composite cathodes. *Electrochem. Commun.* **2012**, *20*, 124–127. [[CrossRef](#)]
56. Żelechowska, K.; Stolarczyk, K.; Lyp, D.; Rogalski, J.; Roberts, K.P.; Bilewicz, R.; Biernat, J.F. Aryl and N-arylamide carbon nanotubes for electrical coupling of laccase to electrodes in biofuel cells and biobatteries. *Biocybern. Biomed. Eng.* **2013**, *33*, 235–245. [[CrossRef](#)]
57. Matyszewska, D.; Nopora, E.; Żelechowska, K.; Biernat, J.F.; Bilewicz, R. Synthesis, characterization, and interactions of single-walled carbon nanotubes modified with doxorubicin with Langmuir–Blodgett biomimetic membranes. *J. Nanopart. Res.* **2018**, *20*, 1–16. [[CrossRef](#)] [[PubMed](#)]
58. Khazaee, M.; Xia, W.; Lackner, G.; Mendes, R.G.; Rummeli, M.; Muhler, M.; Lupascu, D.C. Dispersibility of vapor phase oxygen and nitrogen functionalized multi-walled carbon nanotubes in various organic solvents. *Sci. Rep.* **2016**, *6*, 26208. [[CrossRef](#)]
59. Guo, M.; Huang, J.; Kong, X.; Peng, H.; Shui, H.; Qian, F.; Zhu, L.; Zhu, W.; Zhang, Q. Hydrothermal synthesis of porous phosphorus-doped carbon nanotubes and their use in the oxygen reduction reaction and lithium-sulfur batteries. *New Carbon Mater.* **2016**, *31*, 352–362. [[CrossRef](#)]
60. Singh, D.K.; Iyer, P.K.; Giri, P.K. Distinguishing defect induced intermediate frequency modes from combination modes in the Raman spectrum of single walled carbon nanotubes. *J. Appl. Phys.* **2012**, *111*, 064304. [[CrossRef](#)]
61. Che, B.D.; Nguyen, B.Q.; Nguyen, L.-T.T.; Nguyen, H.T.; Nguyen, V.Q.; Van Le, T.; Nguyen, N.H. The impact of different multi-walled carbon nanotubes on the X-band microwave absorption of their epoxy nanocomposites. *Chem. Cent. J.* **2015**, *9*, 10. [[CrossRef](#)] [[PubMed](#)]
62. Choi, Y.C.; Min, K.-I.; Jeong, M.S. Novel Method of Evaluating the Purity of Multiwall Carbon Nanotubes Using Raman Spectroscopy. *J. Nanomater.* **2013**, *2013*, 615915. [[CrossRef](#)]
63. Ye, B.; Kim, S.-I.; Lee, M.; Ezazi, M.; Kim, H.-D.; Kwon, G.; Lee, D.H. Synthesis of oxygen functionalized carbon nanotubes and their application for selective catalytic reduction of NO<sub>x</sub> with NH<sub>3</sub>. *RSC Adv.* **2020**, *10*, 16700–16708. [[CrossRef](#)]
64. Wepasnick, K.A.; Smith, B.A.; Schrote, K.E.; Wilson, H.K.; Diegelmann, S.R.; Fairbrother, D.H. Surface and structural characterization of multi-walled carbon nanotubes following different oxidative treatments. *Carbon* **2011**, *49*, 24–36. [[CrossRef](#)]
65. Osswald, S.; Havel, M.; Gogotsi, Y. Monitoring oxidation of multiwalled carbon nanotubes by Raman spectroscopy. *J. Raman Spectrosc.* **2007**, *38*, 728–736. [[CrossRef](#)]
66. Sun, J.; Zheng, G.; Lee, H.-W.; Liu, N.; Wang, H.; Yao, H.; Yang, W.; Cui, Y. Formation of Stable Phosphorus–Carbon Bond for Enhanced Performance in Black Phosphorus Nanoparticle–Graphite Composite Battery Anodes. *Nano Lett.* **2014**, *14*, 4573–4580. [[CrossRef](#)]
67. Liu, H. Facile Preparation of 1D a-MnO<sub>2</sub> as Anode Materials for Li-ion Batteries. *Int. J. Electrochem. Sci.* **2016**, *11*, 8964–8971. [[CrossRef](#)]
68. Țucureanu, V.; Matei, A.; Avram, A.M. FTIR Spectroscopy for Carbon Family Study. *Crit. Rev. Anal. Chem.* **2016**, *46*, 502–520. [[CrossRef](#)]
69. Raghu, A.V.; Karuppanan, K.K.; Pullithadathil, B. Highly Surface Active Phosphorus-Doped Onion-Like Carbon Nanostructures: Ultrasensitive, Fully Reversible, and Portable NH<sub>3</sub> Gas Sensors. *ACS Appl. Electron. Mater.* **2019**, *1*, 2208–2219. [[CrossRef](#)]
70. Niu, F.; Tao, L.-M.; Deng, Y.-C.; Wang, Q.-H.; Song, W.-G. Phosphorus doped graphene nanosheets for room temperature NH<sub>3</sub> sensing. *New J. Chem.* **2014**, *38*, 2269–2272. [[CrossRef](#)]

71. Ghosh, D.; Bhandari, S.; Khastgir, D. Synthesis of MnO<sub>2</sub> nanoparticles and their effective utilization as UV protectors for outdoor high voltage polymeric insulators used in power transmission lines. *Phys. Chem. Chem. Phys.* **2016**, *18*, 32876–32890. [[CrossRef](#)] [[PubMed](#)]
72. Okpalugo, T.; Papakonstantinou, P.; Murphy, H.; McLaughlin, J.; Brown, N. High resolution XPS characterization of chemical functionalised MWCNTs and SWCNTs. *Carbon* **2005**, *43*, 153–161. [[CrossRef](#)]
73. Zhang, Y.; Liu, G.; Yu, S.; Zhang, J.; Tang, Y.; Li, P.; Ren, Y. Kinetics and Interfacial Thermodynamics of the pH-Related Sorption of Tetrabromobisphenol A onto Multiwalled Carbon Nanotubes. *ACS Appl. Mater. Interfaces* **2014**, *6*, 20968–20977. [[CrossRef](#)]
74. Tinsley, D.M.; Sharp, J.H. Thermal analysis of manganese dioxide in controlled atmospheres. *J. Therm. Anal. Calorim.* **1971**, *3*, 43–48. [[CrossRef](#)]
75. Kim, M.-J.; Jeon, I.-Y.; Seo, J.-M.; Dai, L.; Baek, J.-B. Graphene Phosphonic Acid as an Efficient Flame Retardant. *ACS Nano* **2014**, *8*, 2820–2825. [[CrossRef](#)] [[PubMed](#)]
76. Wu, X.; Radovic, L. Inhibition of catalytic oxidation of carbon/carbon composites by phosphorus. *Carbon* **2006**, *44*, 141–151. [[CrossRef](#)]
77. Mahajan, A.; Kingon, A.; Kukovecz, Á.; Konya, Z.; Vilarinho, P.M. Studies on the thermal decomposition of multiwall carbon nanotubes under different atmospheres. *Mater. Lett.* **2013**, *90*, 165–168. [[CrossRef](#)]
78. Rather, S.-U.; Nahm, K.S. Hydrogen uptake of high-energy ball milled nickel-multiwalled carbon nanotube composites. *Mater. Res. Bull.* **2014**, *49*, 525–530. [[CrossRef](#)]
79. Xiong, W.; Zeng, G.; Yang, Z.; Zhou, Y.; Zhang, C.; Cheng, M.; Liu, Y.; Hu, L.; Wan, J.; Zhou, C.; et al. Adsorption of tetracycline antibiotics from aqueous solutions on nanocomposite multi-walled carbon nanotube functionalized MIL-53(Fe) as new adsorbent. *Sci. Total Environ.* **2018**, *627*, 235–244. [[CrossRef](#)] [[PubMed](#)]