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# Functionalization as a way to enhance dispersion of CNTs in matrices: A review

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## Abstract

The past three decades of thorough scientific scrutiny of carbon nanotubes (CNTs) revealed that, despite of their remarkable properties, some technological applications are adversely affected by the difficult in handling them, along with their tendencies, as a consequence of their graphitic structure, to agglomerates and exhibit limited or naught interaction with other materials. These issues play a crucial role when CNTs are applied as nanofillers inside matrices, in particular polar ones. In this case, unless several preliminary steps are taken, an efficient and uniform dispersion of the CNTs becomes impossible, thus the nanocomposite cannot exhibit the expected final properties. Unfortunately, a universal procedure doesn't exist since the problem of the dispersion of CNTs is very complex and its solution requires an advanced understanding of the properties of the CNTs (e.g. single or multi walled, size, length, lattice defectivity...) as well as of the matrices used. This review aims to help the reader to select the appropriate dispersion procedure by acquiring fundamental knowledge regarding: I) the synthesis and properties of pristine CNTs, II) methods of chemical functionalization and properties of functionalized CNTs, and III) methodologies for the mechanical dispersion of CNTs. A brief overview regarding chemo-physical characterization techniques is also given to enable a better evaluation of the properties of the CNTs, both before and after functionalization.

**Keywords:** Carbon nanotubes; Functionalization; Dispersion; Mechanical properties; Electrical properties; Synthesis.

## 1. Introduction

The continuous development of new/advanced materials with improved properties is a mandatory requirement to satisfy human's technological needs in several breakthrough fields, such as biomedicine, sensing, energy, environmental remediation processes, robotics, optoelectronics, nanotechnology, and so on [1–10]. In the last years it has been registered a great research effort focused on the production of nanocomposites, where the performance of polymeric, metallic or

ceramic matrices is enhanced by adding nanoscopic fillers with peculiar properties [11–19]. These fillers can be zero, one, or two-dimensional, being spherical, elongated or planar systems, respectively. In this context, graphitic nanofillers such as graphene, carbon nanotubes (CNTs) and carbon nanofibers play a very promising role due to their specific structural and functional properties (e.g., high aspect ratio, high mechanical strength, high electrical and thermal conductivity, high electron emission, high optical absorption, low energy-loss, etc...) [20–22]. Among the materials adopted as matrix in research studies, polymers are the most studied ones due to simpler processing compared to both ceramics and metals [23], moreover it is also interesting to notice that computational modeling studies largely focus on polymeric matrices as well. Among the three graphitic nanomaterials mentioned, CNTs are considered as excellent fillers due to their high tensile strength, low density and high electrical and thermal conductivity [22,24]. CNTs-reinforced composites should guarantee excellent properties exploitable in various engineering applications, such as conductive glues/inks, sensors, gas/energy storage devices, light weighted-reinforced materials for applications in extreme environments, and many others [25–30]. However, there is often a significant mismatch between theory and practice as the final properties of the nanocomposites are strongly affected by the fabrication process, and in particular by the dispersion of the nanofiller in the matrix. Thus, a greater comprehension of the effects induced by different processes is mandatory, in particular for what concern the dispersion of the CNTs in the matrix. In fact, one of the main limitations to be overcome is the tendency of CNTs to stick together into bundles (i.e., CNTs exhibit strong Van der Waals forces and little or no affinity for non-graphitic materials), thus limiting the efficiency of dispersion inside the matrix and the interfacial interaction between filler and matrix [31]. For instance, to overcome this issue different functionalization techniques have been widely adopted to improve the dispersion of CNTs in matrices [32,33]. Numerous theoretical studies are increasingly providing a deeper knowledge of the molecular dynamics related to the dispersion and interaction between CNTs and matrices. This allows to better understand its mechanisms and evaluate the beneficial effect of functionalized CNTs in composite materials[34–37].

Therefore, the aim of this review is to provide a simply organized and easy-consultable document describing the science of CNTs, their chemo-structural properties and synthesis, with a particular focus on the different methods of dispersion and/or functionalization of CNTs for the production of novel/advanced nanocomposites. In this manuscript, the principles behind the most diffuse functionalization techniques involving CNTs are critically discussed, and case studies regarding the most innovative concepts in CNTs and CNTs reinforced composites are provided.

Lastly, important guidelines are summarized to facilitate the work of novice end-users interested in this field of research and development.

## 2. CNTs: Chemistry and structure-relation property

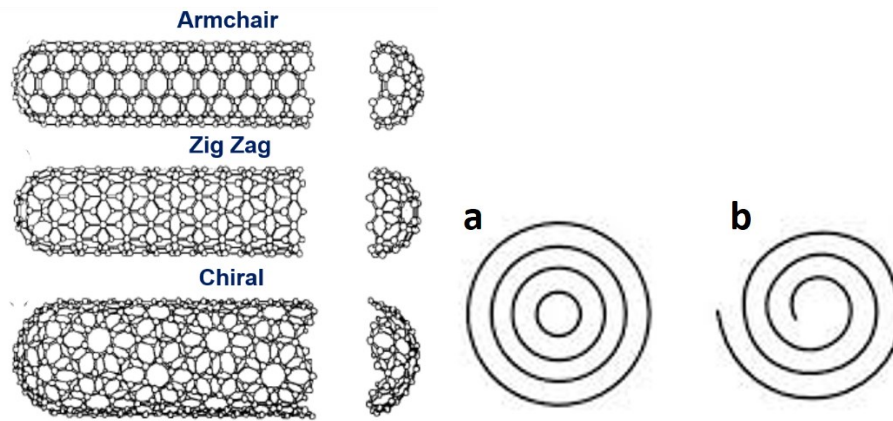
Carbon is one of the most abundant elements on earth, and it is found in nature both as native and in numerous compounds. Carbon can exhibit three hybridization forms  $sp^3$ ,  $sp^2$  and  $sp$ , coordinating with 4 atoms (bond angles  $109.5^\circ$ ), 3 atoms ( $120^\circ$ ) and 2 atoms ( $180^\circ$ ) respectively. Due to its electronic configuration, the complete ionization of carbon is thermodynamically disfavored, whereas its electronegativity pushes towards the formation of covalent bonds [32]. Carbon is naturally present under two main allotropic forms, namely: graphite (hexagonal structure with hybridization  $sp^2$ ) and diamond (cubic structure with hybridization  $sp^3$ ). Furthermore, in the last century the continuous research promoted the discoveries of other allotropic forms: carbynes [38,39], fullerenes (spherical particles) [40], graphene (assumed as single-layer graphite) [41], and CNTs (tubular structure) [42–45]. To be noticed that the curved surfaces in both carbon nanotubes and fullerenes are caused by a mixed  $sp^2$ - $sp^3$  hybridization of the carbon's orbitals.

Carbon nanotubes can be classified into two main types:

single-walled carbon nanotube (SWCNTs) consisting of a single graphene sheet wound on itself in a tubular shape;

multi-walled carbon nanotube (MWCNTs) in which two or more concentric tubes are formed by graphene sheets coaxially wound around each other.

SWCNTs usually have a diameter between 0.7 and 2 nm and a length varying between a few microns up to millimeters. Depending on the direction in which the graphene sheet is rolled up it is possible to have different chirality (defined by the chiral angle between carbon hexagons and the tube axis), which, together with the diameter, affects the electrical conductivity of the nanotube [46,47]. In figure 1 it is shown the different chirality of SWCNTs, where an armchair configuration CNTs always exhibits electrical conductivity similar to metals, whereas the chiral and zigzag configurations can be either semiconductors or metal conductors [46].



**Figure 1:** On the left the three models for SWCNTs structures with different chirality. On the right the crosssectional view of the common russian doll structure of concentric tubes of MWCNTs (a) and the rare Parchment model (b).

Multi-walled carbon nanotubes (MWCNTs) can be generally described as multiple concentric tubes (like a Russian doll) of rolled graphene layers. However, in some rare cases, it was found that some structures resemble a scroll of parchment (or a rolled newspaper) [47] where the single sheet of graphene is spirally rolled around itself in a configuration that can be explained by the Parchment model where, (the two model are shown in Figure 1). Due to Wan der Waals interaction between graphene layers, the interlayer distance between concentric tubes inside a MWCNTs is the same one found between the atomic layers in graphite, which is approximately 3.4 Å. Even though MWCNTs are formed by several concentric SWCNTs, and as mentioned earlier the electrical conductivity of each individual concentric tube is defined by chirality and diameter, all MWCNTs can be considered metal conductors as a consequence of statistical probability and restrictions on the relative diameters of the individual tubes (at least one of the shells, and thus the whole MWCNTs, is always a zero-gap metal).

**Table 1.** Electrical resistivity of different kinds of CNTs in comparison with carbon fibers and amorphous carbon [48–52]

	<b>Electrical resistivity</b> <b>[Ω m]</b>
<b>MWCNTs</b>	$8 - 20 \times 10^{-6}$
<b>SWCNTs</b>	$\leq 1 \times 10^{-6}$

<b>Carbon fibers</b>	$1 \times 10^{-6}$
<b>Amorphous carbon</b>	$5-8 \times 10^{-4}$

From a mechanical point of view, the outstanding properties of CNTs are mainly due to the covalent bonds between the carbon atoms. However, as reported in Table 1, MWCNTs tend to exhibit better performances than SWCNTs as a result of the presence of multiple concentric tubes. It is interesting to note that the Young Modulus of SWCNTs mainly depends on the diameter of the tubes, and not on the chirality, with a value of circa 1,000 GPa for a diameter of ca. 1-2 nm, as confirmed by both simulated and experimental results. Another interesting property is the ability of CNTs to resist high levels of plastic deformation without breaking (ca. 16%) [53]. These characteristics combined with a low density make CNTs better reinforcing fillers than carbon fibers [54], even considering that CNTs cost is still quite higher than CFs (Table 2)

**Table 2.** Main mechanical properties and cost of different kinds of CNTs in comparison with carbon fibers [55–57].

	<b>Young Modulus</b> [GPa]	<b>Tensile Strength</b> [GPa]	<b>Density</b> [g/cm <sup>3</sup> ]	<b>Price</b> [€/Kg]
<b>MWCNTs</b>	1,200	150	2.1-2.6	100
<b>SWCNTs</b>	1,054	75	1.33-1.94	130
<b>Carbon fibers</b>	350	2.5	1.75-2.10	21

All the properties we mentioned so far refer to the perfect CNTs which don't contain lattice defects. Defectivity can be present in the lattice for various reasons related to the synthesis of the CNTs or their reaction with harsh chemical agents. Huge number of defects can severely hinder the performance of CNTs, whereas a limited number of defects might have little or no impact of the chemo-physical performance of the CNTs (especially of the MWCNTs). Limited defectivity can be very interesting for the application of CNTs in nanocomposites, because, especially if coupled with the presence of different chemical groups, can lead to improved dispersion in matrices. Defects can

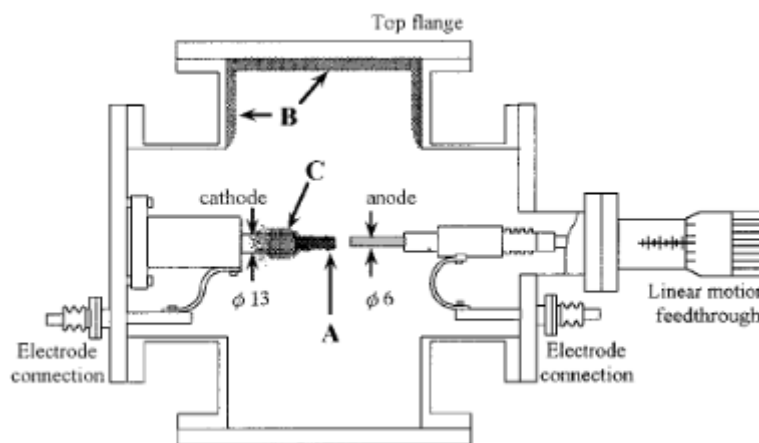
limit the agglomeration of CNTs (decrease in Van der Waals forces) enhancing their dispersion inside a matrix and their interaction with polar materials. Therefore, we can say that a limited amount of lattice defectivity can be an optimal tradeoff between the electro-mechanical performance of defect-free CNTs and their successful application in nanocomposites. It is interesting to note that, lattice defects can be removed a posteriori by a process of high temperature annealing (above 1700 °C) in vacuum or in presence of inert gas.

### **3. Methods of synthesis of CNTs**

Although the discovery of carbon nanotubes is often attributed to the Japanese Sumio Iijima for his article published in 1991 [42], in reality the history of the discovery of nanotubes is much more complex and began much earlier; first in 1954 by Radushkevich and Lukyanovich [58] and secondly in 1972 by Baker et al. [59]. The method for the synthesis of nanotubes used by Ando [60] in the article of Iijima [42] was the electric arc discharge which belongs to high temperature methodology, together with laser ablation and solar vaporization. These methodologies have in common the high temperature sublimation of graphite which is used as carbon precursor.

#### *3.1 Arc discharge*

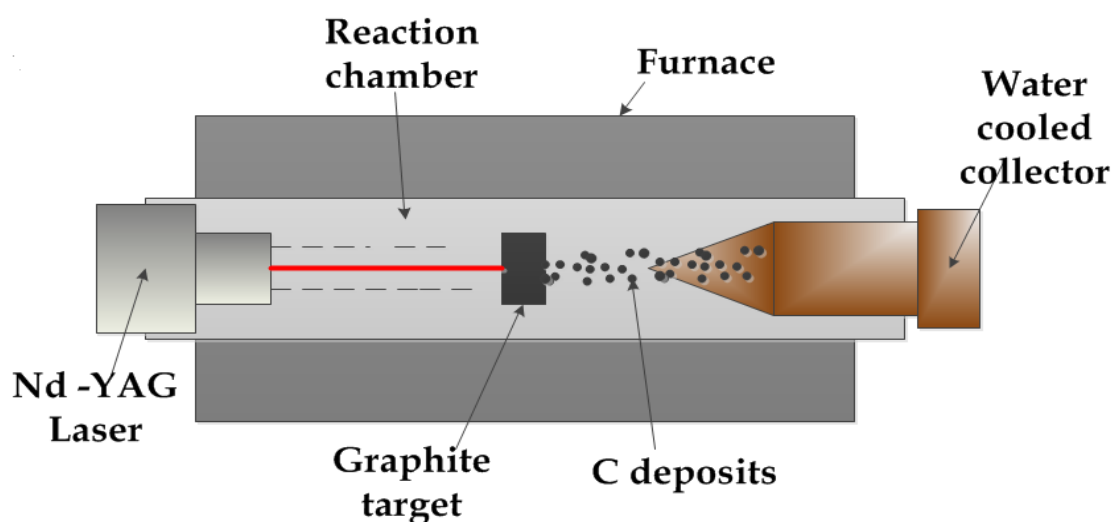
This method is based on an arc discharge generated between two graphite electrodes placed face-to-face in the instrument chamber (see Figure 2) [61] under a partial pressure of He or Ar (typically 600 mbar). The electric discharge generates a temperature (up to 6000 °C) at which the graphite sublimates, and the ejected carbon atoms form a plasma. The ejected atoms then deposit on the cathode. While this technique was developed to synthesize fullerene, it was then discovered that the presence of small amount of transition metals in the anode catalyzes the synthesis of CNTs [62]. It should be emphasized that the presence of metal catalyst, such as Fe, Ni, Co or their alloys, [63] is always necessary to catalyze the growth of the CNTs, regardless of the deposition technique. At the end of the growth the metal particles are usually lodge at the tips of the nanotube or at their base, as explained by the tip-growth mechanism and by the base-growth mechanism respectively [64].



**Figure 2:** Schematic representation of a typical arc-discharge apparatus [61]

### 3.2 Laser Ablation

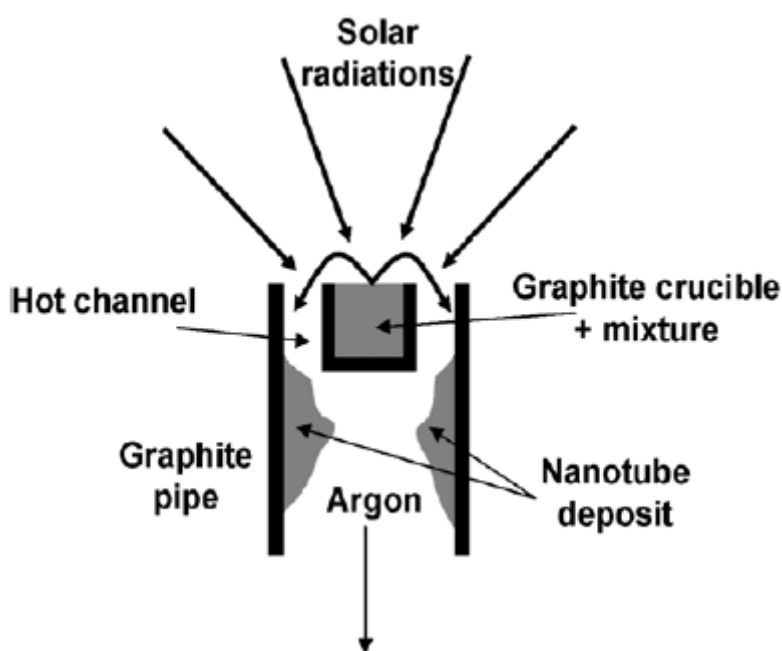
Another method investigated early on is the laser ablation process which is based on the laser-induced sublimation of a mixture of graphite and metallic catalyst (Fe, Ni, Co). The process is performed in a quartz tube reactor inserted in furnace and heated at 2700-3200 °C under He or Ar atmosphere. As depicted in Figure 3 [65], the gas flux promotes the deposition of carbon atoms and catalysts' particles on a collector at lower temperature. By varying both the cooling temperature of the collector and the gas flow rate, it possible to control the growth of the nanotubes [66]. By varying the laser, catalyst composition, growth temperature, nature of gases and gas pressure, the average nanotube diameter and size distribution can be varied [67].



**Figure 3:** Laser ablation apparatus for the production of CNTs [65].

### 3.3 Solar beam deposition

Sublimation induced by solar beam is a particular method investigated in the past involving a solar furnace in which the sunlight is focused on a graphite sample to sublime the carbon. The soot obtained is then condensed in a cold dark zone of the reactor. The sunlight is collected by a flat tracking mirror and is reflected towards a parabolic mirror which focuses the solar radiation directly on the graphite target (as in Figure 4) [62]. Under clear-sky conditions, temperatures of ca. 2727 °C are reached at the solar station and the sublimation process can start. With this technique, initially only fullerenes were produced [68]. However, since 1998 also CNTs were obtained by changing the target composition and adjusting the experimental conditions [69].



**Figure 4:** Schematic representation of principles guiding the solar furnace technique [62]

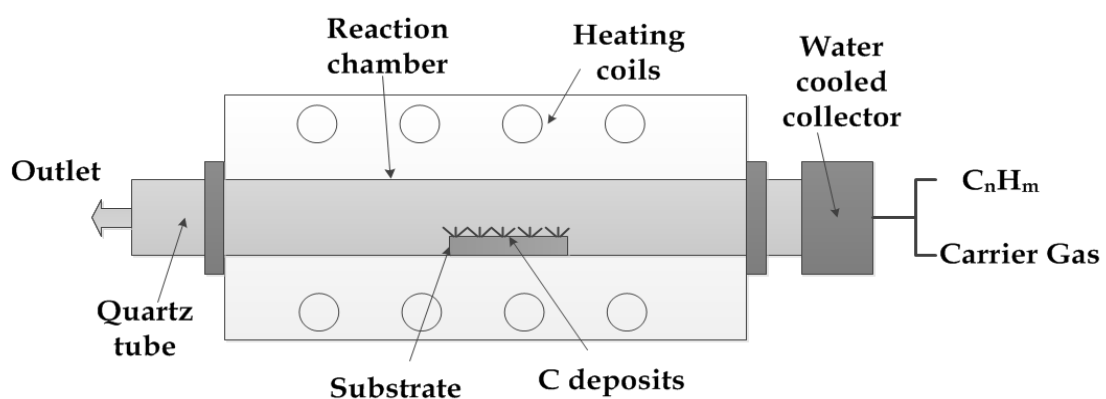
### 3.4 Chemical vapor deposition

One of the main concerns related with the high temperature synthetic route is the low production yield and the fact that such high temperature is problematic for industrial production. Nowadays, for large-scale mass production of nanotubes the methodology mostly used is chemical vapor deposition (CVD), either as thermal-CVD or plasma enhanced-CVD, which allows the production of industrial-grade nanotubes at significantly lower temperatures.

Interestingly, CVD is historically the first methodology used by two Russian scientists [58] to grow CNTs as byproducts of the synthesis of carbon nanofibers. However, since their research paper was

published in Russian in a Russian journal, its diffusion in the global scientific community was limited. As for the high temperature methods, CVD requires the presence of metal particles to catalyze the synthesis. The metal catalyst can be deposited as a thin layer directly on the surface of a substrate (usually silica), where at the temperature of synthesis of the CNTs (600-1200 °C) the thin layer will reshape into nano agglomerates of metal. Another way is to use a gas carrier to inject the catalyst, along with the carbon precursor, in the form of an organometallic compound [70–72] or a metal oxide [73–75]. At the process temperature the metal compound is reduced to metal atoms which aggregate and react with the carbon to produce CNTs. To be noticed that the diameter of the nanotube is related to the size of the metal particle catalyzing the growth and that the metal particles remain trapped inside the formed CNTs, either in the tip (far from the substrate) or at the base (meaning adjacent to the substrate), depending on the strength of adhesion between the catalyst particles and the substrate.

To start the growth of nanotubes the carbon precursor (e.g. acetylene, ethylene, ethanol, methane, camphor.....etc) [70,76] is carried into the reactor by an inert gas (nitrogen or argon), sometimes along with reactive gases such as hydrogen or ammonia to help removing amorphous carbon deposited as a byproduct. The silica substrate, where the growth of the CNTs starts, is placed in the hottest region of the deposition system and it is heated up to 600-1200 °C. The thermal cracking of the carbon precursor generates reactive species of carbon which deposit on and interact with the metal particles, thus enabling the growth of the CNTs.

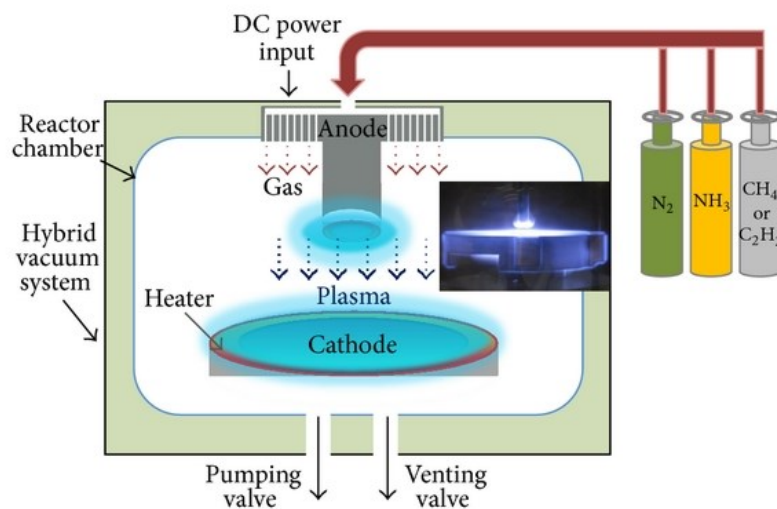


**Figure 5:** Schematic representation of a CVD apparatus [65].

The parameters influencing the growth of CNTs are several and numerous strategies have been used to implement the CVD methods towards a selective and efficient production of CNTs, including modifying the reactor set-up (i.e., horizontal furnace, fluidized bed reactor and vertical furnace) and the relative reaction parameters [77].

### 3.5 Plasma enhance chemical vapor deposition

Currently one of the most used production techniques of CNTs is plasma-enhanced CVD [78] in which the working temperature is lower than that of thermal-CVD thanks to the use of plasma. The plasma is generally generated by exciting the reacting gases between two electrodes by using either radio frequency or direct current. Figure 6 shows the schematic diagram of a typical plasma CVD apparatus. The substrate is placed on the lower electrode, while the reaction gas is fluxed in during the discharge at the upper electrode. The growth of the carbon nanotubes takes place on the nanoparticles of the metal catalyst deposited via sputtering on the substrate.



**Figure 6:** Schematic representation of principles guiding the Plasma Enhancement CVD [79].

#### 4. Dispersion of CNTs

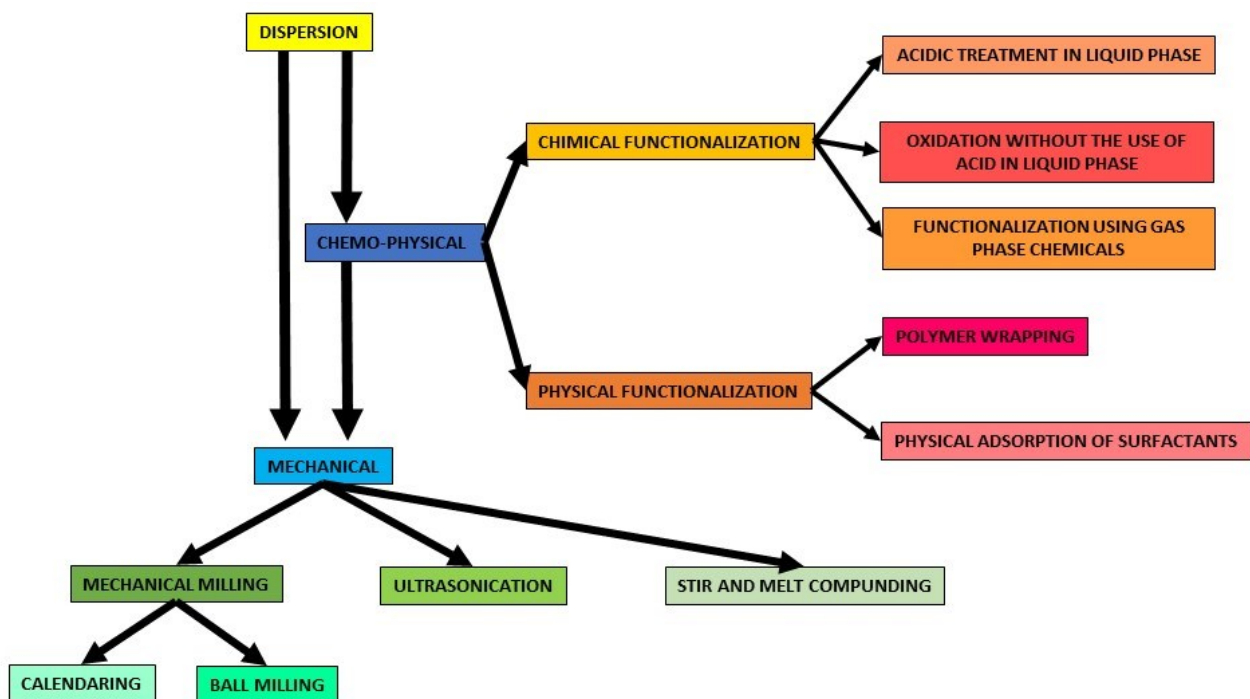
Dispersion, as defined by IUPAC, refers to a system in which distributed particles of one material are individually dispersed in a continuous phase of another material, [80,81], but the term also indicates the procedure with which such particles are dispersed. Dispersion is a key aspect for improving the performance of composite materials. The dispersion of nanotubes into matrices is difficult for three main reasons: 1) CNTs are agglomerated due to Van der Waals forces; 2) due to their chemical inertia they do not interact with other materials; 3) due to their low density they tend to occupy a large volume into the matrix. [12,82–84].

Depending on the synthesis method, length, and structure (SWCNTs or MWCNTs) the CNTs can assume different forms of agglomeration. In particular, the MWCNTs commercially synthesized in fluidized bed reactor, generally have larger dimensions, and more defects. Their higher defectivity means that they are found in the form of big agglomerates with a structure like “combed yarn” or

“bird-nests”. Instead, smaller and less defective CNTs tend to intertwine less, agglomerating into less interlocked structures which therefore make them more dispersible. On the other side, it must be considered that some evidence points toward the fact that SWCNTs are more difficult to disperse than MWCNTS, due to higher van der Waals binding of the agglomerates [85]. In any case, type of CNTs (single or multi-wall), aspect ratio, length, defectivity, synthesis procedure, all contribute to determine the agglomeration level and strength, thus rendering impossible to draw clear conclusions about the difficulty of the dispersion.

By analysing the scientific literature, several dispersion methods have been proposed by numerous researchers. They are based on two different approaches which often can be combine: mechanical dispersion, or chemo-physical treatment followed by mechanical dispersion (Figure 7). Mechanical dispersion is based on the utilization of various techniques able to disperse CNTs in solvents or solid matrices, namely: ultrasound sonication, mechanical milling, high energy stirring. On the other hand, when mechanical dispersion is preceded by chemo-physical treatment, the dispersion is enhanced by chemical or physical bond between CNTs and chemical agents.

In the following paragraphs the various types of dispersion methodologies are described.



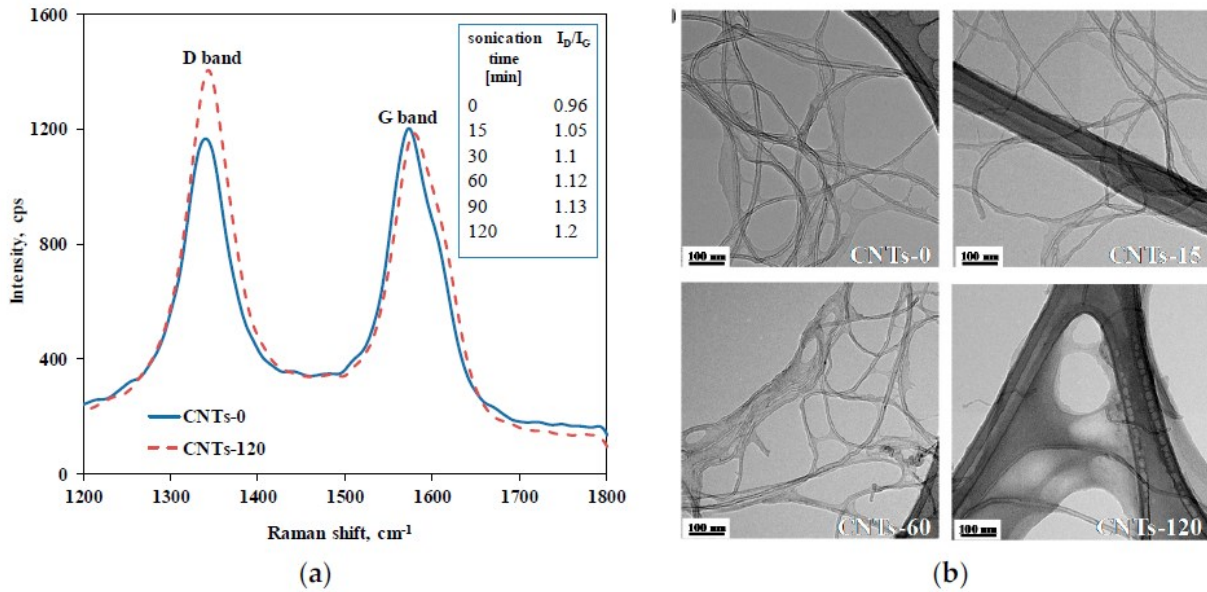
**Figure 7:** Schematic summary of the principal dispersion approaches for CNTs

## 4.1 Mechanical dispersion

### 4.1.1 Ultrasonication

Ultrasonication is an effective method widely exploited to disperse fillers (such as CNTs) in liquids having a low viscosity. Usually, this technique is commonly used to separate CNTs agglomerates in solution, but it could also be used effectively in presence of polymers, surfactants, and other compounds. Thus, for this reason, ultrasonication is commonly added to other dispersion techniques [86–90]. This technique can be performed either by using ultrasonic bath or by using an ultrasonic sonotrode (generally named probe or horn sonicator). In general, standard laboratory bath sonicators run at 20–23 kHz (power < 100 W), whereas commercial ones have an adjustable amplitude ranging from 20% to 70% (power 100–1500 W). The probe is usually made of an inert metal (e.g., titanium) and the exchangeable tips diameters are comprising between 1.6 mm and 12.7 mm.

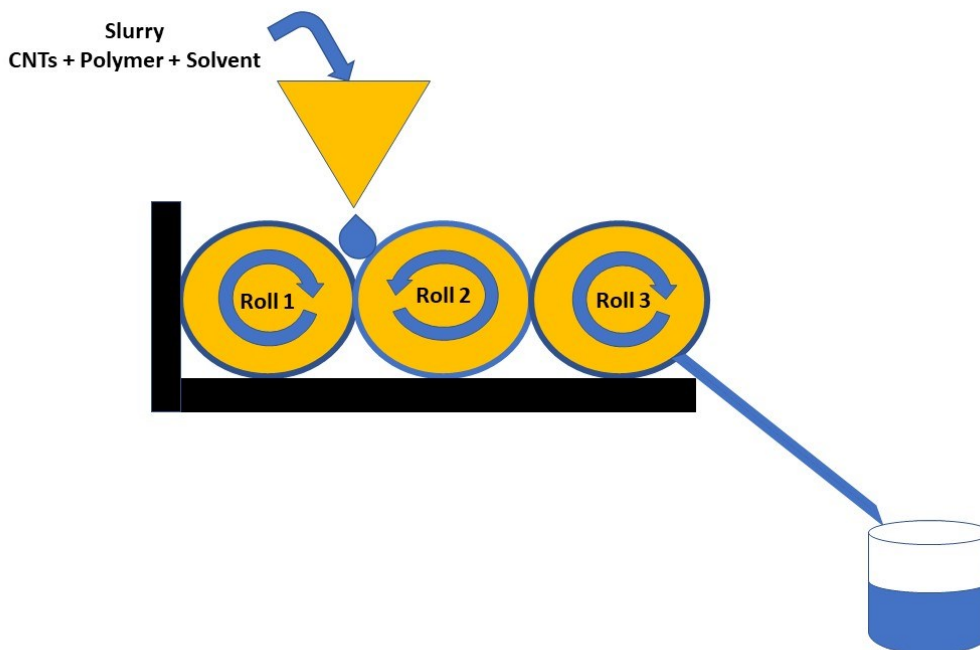
Ultrasonication is based on the cavitation mechanism, which consists in the formation of high energy bubbles which collapse within the liquid and generate a wave-shock providing high energy local shear necessary to disentangle CNTs bundles [91]. However, the main drawback of using sonotrode is that the high energy can easily damage the CNTs structure. In some cases ultrasonication can lead to a reduction in the length of the CNTs [85,92,93]. Figure 8 [94] reports the Raman spectra of CNTs before and after sonication (different sonication times are investigated). Results evidenced that after ultrasonication it has been registered the increment of signal at  $1286\text{ cm}^{-1}$  attributable to disordered  $sp^3\text{ C}$  in CNTs, suggesting the generation of defects in the CNTs lattice. In extreme cases, the graphene layers of CNTs can be completely destroyed. As aforementioned in this paper, too much structural damage can badly affected the electrical and mechanical properties of CNTs, thus negatively influencing the performances of the final composites.



**Figure 8:** 1 a) Raman spectra of pristine CNTs and after 2 min and 120 min of ultrasonication and b) representative TEM micrographs of CNTs before and after ultrasonication [94]

#### 4.1.2 Mechanical milling: Calendaring

The calender, commonly known as three roll mills, is a machine tool that employs the force created by rollers to mix, disperse or homogenize viscous materials. Interestingly, the use of a three-roll machine has recently provided good results for the dispersion of nano-reinforcement in materials [95]. This dispersion method exploits shear forces rather than compression forces during the process and provides the CNTs dispersion without structural damages. A calendaring machine consists of three adjacent cylindrical rollers (i.e., the feeding and the apron rollers rotate in the same direction, whereas the center roller rotates in the opposite) running at different velocities (as in Figure 9). The material is introduced in the system by a hopper directly between the first and the second roller. The material is then transported by the second roller into the gap between second and third roller where the filler is dispersed. At the exit, the material that remains on the central roller moves through the third roller towards a metal drain tip. The third roller tends to spin faster than the other two and the material is subjected to a greater shear force. This milling cycle can be repeated several times to ensure the desired dispersion degree.

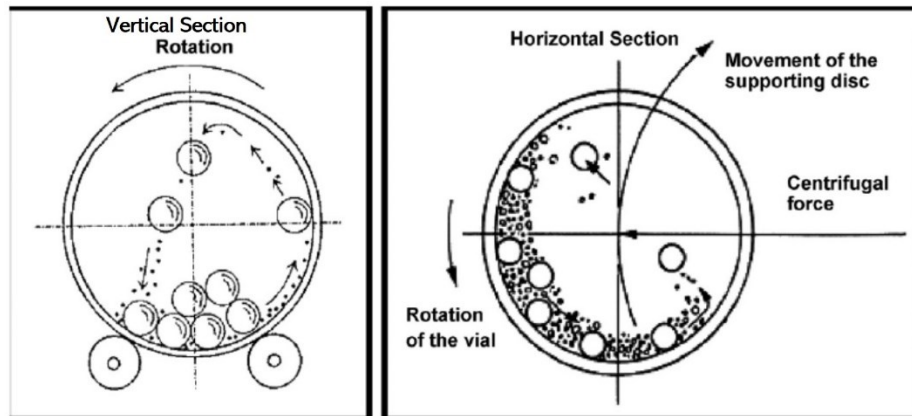


**Figure 9:** Schematic of the three roll mill machine used for the calendaring process

The employment of calendaring to disperse CNTs in a polymer matrix has become a very promising approach to achieve a relatively good carbon nanotubes distribution and dispersion [96–98]. However, there are several concerns about using this technique: for example the minimum gap is about 1–5  $\mu\text{m}$ , which is larger than the typical dimension of single CNTs. Due to this, calendaring can better disperse the large agglomerated bundles of CNTs into small ones rather than disperse individual CNTs. In addition, the feeding materials should be in a viscous state, thus this technique could be used in presence of thermoplastic matrices for heated systems[99].

#### 4.1.3 Mechanical milling: Ball milling

Mechanical milling, or grinding, is commonly used to reduce the physical size of particles to extremely fine powder. In ball milling, the materials to be ground are placed in a proper container together with milling balls. The rotation of the cylindrical container generates the kinetic energy necessary to move the sample and the milling balls, also known as charge materials. Subsequently, the collision between the sample and the milling balls generates high enough pressure to favour crack propagation and fracture, leading to the breakage of the sample [100]. The higher the impact energy, the smaller the fragments produced. The typical motion of balls inside the container is shown in Figure 10:210 [101]. Different materials, including ceramic, flint pebbles and stainless steel, are used as balls.



**Figure 10:2** Schematic representation of the balls motion during the milling process. On the left: vertical configuration (conventional ball mill). On the right: horizontal configuration (planetary ball mill) [101]

Ball milling has been used for a variety of applications related to CNTs: such as to obtain amorphous carbon or onion-like particles from the breakdown of its structure [102,103]. In some cases the nanotube length was reduced and, simultaneously, the primary agglomerate morphology was altered from loosely packed to compact [104]. With this technique it is also possible to carry out some chemo-mechanical reactions, as shown by Flahaut et al [75], where an in situ amino functionalization of CNTs was made while using ball milling. Mainly this technique is used to form metallic composites by grinding metal particles and CNTs to form composite powders but it is possible also for other matrices [104]. The mechanical alloying is used primarily for pure Al and Al alloys matrices [15,102,105–108]. Moreover, different alternative processes of ball milling could be used on the basis of either the material and/or the purpose of the process (e.g., high energy milling (HEM)) [15,109,110].

#### 4.1.4 Stir and melt compounding

Stirring is a common technique for dispersing particles in liquid systems and can also be used to disperse CNTs in a polymer matrix. The results of the dispersion depend on the size, shape of the propeller and on the mixing speed. Intense agitation of CNTs in a polymer matrix can bring to a relatively good dispersion [12,111]. However, MWCNTs are more easily dispersed than SWCNTs using stirring, even if MWCNTs tend to re-agglomerate. Indeed, Schmid and Klingenberg [112] suggested that in a fiber suspension agglomeration is mainly caused by elastic interlocking mechanisms and frictional contacts, occurring even without attractive forces between fibers.

During stirring, sliding and Van der Waals forces seem to have a smaller effect on the dispersion, however, agglomeration becomes spontaneous under static flow. In the case of heavily agglomerated

CNTs, however, to obtain a fine dispersion in the polymer matrix higher shear forces are required, that can be obtained by using a high speed shear mixer.

Extrusion is a popular technique for dispersing CNTs into polymers such as in most thermoplastics, where thermoplastic pellets mixed with CNTs are fed into the extruder hopper. The twin screws rotate at high speed creating a high shear flow, resulting in the dispersion of the CNTs agglomerates, and mixing with the molten polymer. This technique, commonly used for thermoplastic matrices is particularly useful for producing CNTs / polymer nanocomposites with both high and low filler content [113].

## **4.2 Functionalization**

Carbon nanotubes surface properties can be modified by attaching “chemical functionalities” (functionalization treatment) to their surface, which results in a good strategy to overcome the dispersion problems by improving the solubility, compatibility and processability of CNTs [32,114–117].

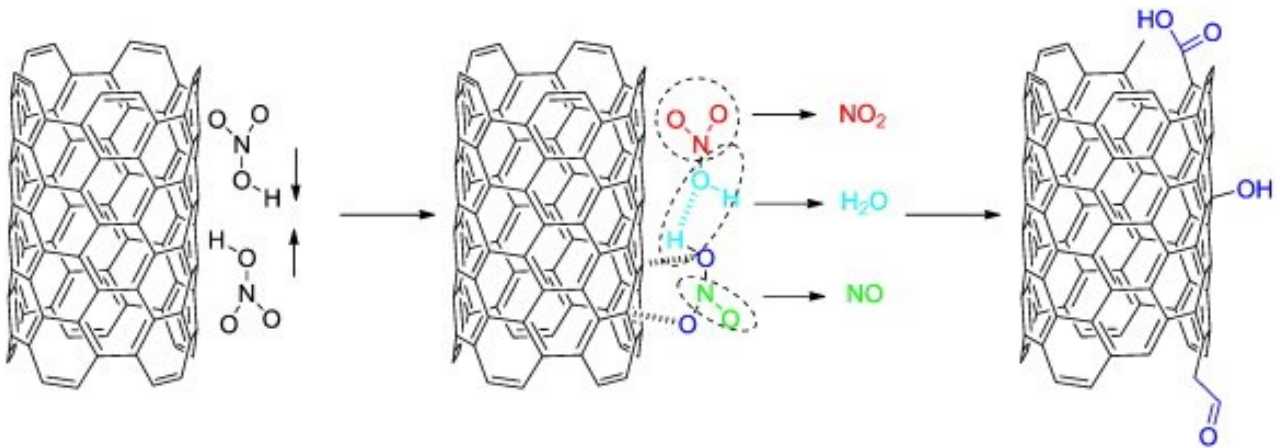
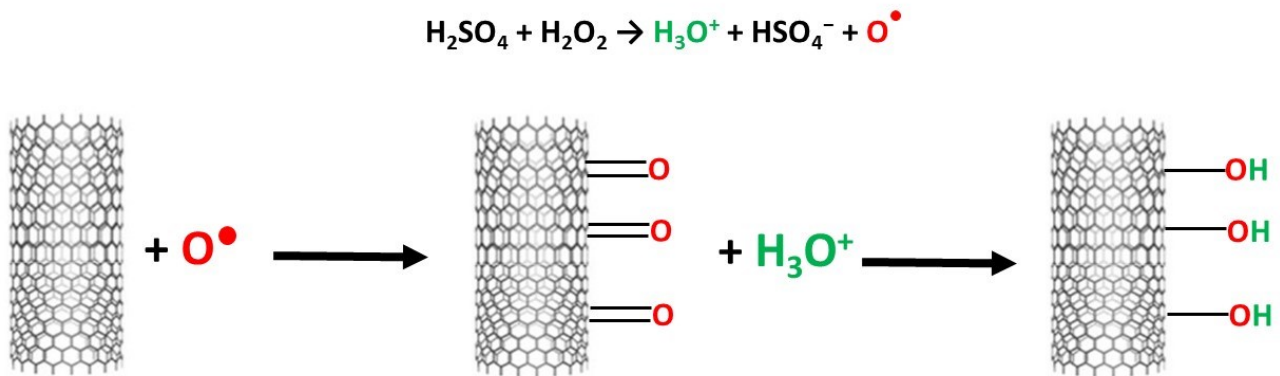
According to the literature, the functionalization of CNTs can be grouped into two main categories:

- chemical or covalent functionalization, in which new chemical groups are chemically bonded to the carbon atoms of the tubes;
- physical or non-covalent, in which various macromolecules are physisorbed or wrapped around the tubular structure.

### **4.2.1 Chemical functionalization**

#### *4.2.1.1 Acidic treatment in liquid phase*

Among the different types of chemical functionalization of CNTs, wet functionalization via acidic oxidation is one of the most commonly adopted [118]. It serves the dual goal of introducing new chemical groups and removing metal particles (trapped catalyst) and amorphous carbon. This is especially true when nitric acid is used to remove residual metals and to introduce oxygen-containing groups (mainly carboxylic acid) as reported in Figure 11a.

**A****B**

**Figure 11:** a) mechanism of oxidation of nitric acid [119]; b) mechanism of oxidation of piranha solution

Aviles *et al.* [120] evidenced that performing the process in presence of small concentration of nitric acid ( $\text{HNO}_3$ ) improves the presence of O-containing groups without damaging the structure of CNTs. Consequently, the functionalization with  $\text{HNO}_3$  enhances the dispersion of CNTs in polar solvents. It must be emphasized that process parameters, such as temperature and reaction time play significant roles. As demonstrated by Rosca *et al.* [121], for long time treatment (1-2 days) in concentrated  $\text{HNO}_3$  ( $\%_v > 60 \%$ ) could strongly damage the structure of CNTs, covering their surface with amorphous carbon. On the contrary, a temperature of  $111^\circ\text{C}$  for 6 hours leads to an efficient elimination of both metal impurities and amorphous graphitic platelets. Sulfuric acid ( $\text{H}_2\text{SO}_4$ ) is less used than nitric acid as it does not generate a significant oxidation of the external lattice. However, sulphuric acid in combination with hydrogen peroxide (i.e., piranha solution,  $3 \text{H}_2\text{SO}_4 : 1 \text{H}_2\text{O}_2$ ) exhibits a good functionalization performance by introducing both carboxylic functionalities and hydroxyl groups without causing damages to the CNTs' structure. As shown in Figure 11b the piranha solution generates an oxygen radical (reactive oxygen species) that interacts with the carbon in  $\text{sp}^2$

hybridization state leading to the formation of a double bond with oxygen, which subsequently reacts with  $\text{H}_3\text{O}^+$  to finally create hydroxyl groups on the surface of the CNTs. The functionalization with piranha solution guarantees a good dispersion in both polar and non-polar media [17,118]. Furthermore, by performing this treatment at temperature over  $70\text{ }^\circ\text{C}$  it is possible to cut the tubes [122]. The treatment with sulfonitric acid (i.e  $3\text{ H}_2\text{SO}_4 : 1\text{ HNO}_3$ ) for 8 hours at  $70\text{ }^\circ\text{C}$  can produce a high number of defect thus distorting the CNTs [123]. A study shows how the rise in temperature (from  $60\text{ }^\circ\text{C}$  to  $140\text{ }^\circ\text{C}$ ) with this treatment increase the presence of carboxylic ( $-\text{COOH}$ ) functional groups[124]. The wettability of CNTs after sulfonitric treatment was also studied in literature. One study in particular [125] was performed on vertically aligned (VA) CNTs and showed how the interaction with water was enhanced by the presence of carboxyl and sulfonic groups introduced after 6 hours of treatment. As proved by the experiment, the acidic treatment can reduce the contact angle between water and VA-CNTs from  $176^\circ$  (hydrophobic) to  $25^\circ$  (hydrophilic). Obviously, the increment in wettability is due to the extended presence of O-containing functionalities on the surface. The effect on the dispersion in water after different acid treatments is an important way to evaluate the effectiveness of the treatment and the degree of interaction with the matrix. Furthermore, the use of sonication, via ultrasonic bath, during acidic treatment of CNTs can significantly promote the grafting of a great number of polar groups, while guaranteeing a continuous disentanglement of CNTs bundles during the process[126]. At the end of acid oxidation in general it is necessary to neutralize the reaction solutions containing the treated CNTs. Once neutralized, the CNTs are washed several times with distilled water to remove acid and salt residues adsorbed on the surface and are then recovered through filtration or centrifugation.

#### *4.2.1.2 Oxidation reactions without the use of acid in liquid phase*

The literature suggests several oxidation processes not involving acids. In comparison to acidic treatments, this process allows to introduce a lower oxygen content with lesser damaging of the honeycomb lattice. In this context, Wepanick et al. [123] tried to analyze the degree of CNTs oxidation by comparing oxidizing chemical agents such as  $\text{KMnO}_4$  (potassium permanganate),  $(\text{NH}_4)_2\text{S}_2\text{O}_8$  (Ammonium persulfate) and  $\text{H}_2\text{O}_2$  (hydrogen peroxide) with  $\text{HNO}_3$  and  $\text{HNO}_3/\text{H}_2\text{SO}_4$ . The study successfully demonstrated that even a non-acidic treatment is sufficient to decorate the surface of CNTs with polar functional groups as shown in table 2. In detail, the use of  $\text{KMnO}_4$  is a good technical solution to introduce a high number of carboxylic groups and functionalities. On the other hand,  $(\text{NH}_4)_2\text{S}_2\text{O}_8$  can only create carboxyl, lactone and phenolic acid groups on pre-existing lattice defects, hence it does not guarantee a high oxygen content [123]. It should be noticed that these studies suggest that, by avoiding the use of strong acids, it would be possible to have more

“environmental-friendly” approaches. While most studies focus on low pH treatments, currently there is not much knowledge regarding the structural alteration of graphitic materials under basic conditions. Both Datsyuk et al. [118] and Kim et al. [127] have studied the treatment of MWCNTs with an  $\text{NH}_4/\text{H}_2\text{O}_2$  (ammonium hydroxide/hydrogen peroxide) mixture, which generates epoxy groups with little damage to the external lattice of CNTs. Peng and Liu [126] showed that  $\text{H}_2\text{O}_2$  in contact with CNTs for 1-6 days reacts with amorphous carbon at the surface of CNTs forming  $-\text{OH}$  groups (hydroxyl groups) while removing the disordered carbon. Table 2 reports the comparison between different oxidation processes.

**Table 3:** Comparison of different oxidative processes: different chemical groups formed on the surface in function of oxidant used

Oxidation process	O%	Functional groups introduced	Time	Temperature [°C]	Comment	References
$\text{NH}_4/\text{H}_2\text{O}_2$	2.5	$-\text{COOH}$ ; $-\text{OH}$	5 h	80	Functionalization not very effective, neutralization required at the end of the process with an acid solution (e.g. HCl)	[118]
$\text{H}_2\text{SO}_4$ ,	3.9	$-\text{SO}_3\text{H}$	15 min	60	Sulphonic groups can be detrimental in cementitious matrices. The treatment does not guarantee a uniform functionalization of the CNTs. Neutralization is achieved by rinsing with a strong base (e.g. NaOH) solution. Neutralization can lead to the formation of salts which must be removed	[120]

					by further rinsing with deionized water.	
$(\text{NH}_4)_2\text{S}_2\text{O}_8$	4.5	-OH; -COOH	4 h-24 h	RT	Long reaction times, with little yield of functional groups.	[123,128]
$\text{H}_2\text{O}_2$	4.5	-C=O; -OH	4-6 days	70-65 °C	The instability of $\text{H}_2\text{O}_2$ requires addition of fresh solutions of peroxide at least every 24 h.	[123,126]
$\text{O}_3$	4.7	-C=O	1h	25°C	The use of a gas requires a closed system to recycle ozone.	[123,129,130 ]
$\text{H}_2\text{SO}_4/$ $\text{H}_2\text{O}_2$	7.1	-OH; -COOH	15 min-9 h	RT-70°C	The piranha solution is very aggressive and gives excellent functionalization results. High reactivity of the mixture can be hazardous for the operator. As per the sulfuric acid, neutralization is needed at the end of the treatment. Neutralization can lead to the formation of salts which must be removed by further rinsing with deionized water.	[118,120,131 ]
$\text{HNO}_3$	7.9	-COOH; -C=O	15 min-48 h	RT- 150 °C	Temperature and reaction time can be tailored depending on the sample	[118,120,121 ,123,124,132 ,133]

					used and results required. Stirring or sonication can be used. When heating the acid vapor can be refluxed using a condenser. Carboxylic groups are formed on the surface of the nanotubes. Neutralization with base and several washes are required to remove acid residues from the surface of the CNTs.	
H <sub>2</sub> SO <sub>4</sub> / HNO <sub>3</sub>	8.8	-COOH; -OH; - SO <sub>3</sub> H	1-15 h	RT-60 °C	This treatment can be adapted using simultaneous dispersion techniques (e.g. sonication). Functionalization is homogeneous and severely damages the CNTs.	[120,123– 125,127,132, 134]
KMnO <sub>4</sub>	9	-COOH; -C=O	5 h	150 °C	Some authors claim this technique to be "green", however, to remove permanganate residues a washing with strong acids (H <sub>2</sub> SO <sub>4</sub> ) is often required	[123]

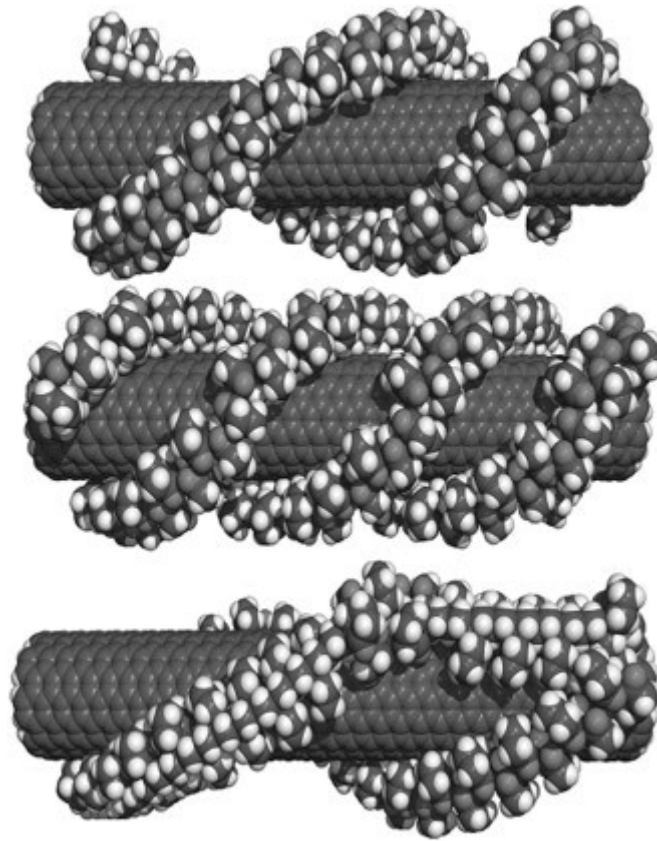
#### 4.2.1.3 Functionalization using gas phase chemicals

An alternative method to oxidize CNTs consists in fluxing an oxidant gas or vapor through CNTs powder. This method guarantees a substantial reduction in waste production and a more homogenous functionalization with respect to acidic treatments. Xia et al. [135] developed a gas-phase route for the oxidation of CNTs by means of HNO<sub>3</sub> vapor treatment. It is a more efficient way to introduce oxygen-containing functional groups on CNTs surface (21%) rather than conventional treatment with liquid HNO<sub>3</sub> (7.9 %). In addition, the use of HNO<sub>3</sub> vapor prevents secondary purification steps (e.g. filtration, washing, and drying) required with the liquid treatment. Another interesting oxidant gas is ozone (O<sub>3</sub>), which can be fluxed directly on carbon nanotubes at temperature equal or greater than RT. The reaction between O<sub>3</sub> gas and carbon produces two distinct functional groups on the surface of the CNTs (namely esters and quinones) as well as volatile byproducts such as CO<sub>2</sub> and CO [129]. The treatment with O<sub>3</sub> guarantees a stable dispersion of CNTs for days, as demonstrated by Li et al., just by simple sonication [136]. The application of ultra-violet radiation of 28 W/cm<sup>2</sup> during ozone treatment for at least 2 minutes, introduces carboxylic acids groups, whose amount is directly related with the exposure time [137]. Another interesting solution is to exploit CO<sub>2</sub> gas both to create functional groups and to eliminate the metallic residues present in the CNTs after the synthesis [138]. Wang et al. [139] explain that a treatment in flux of CO<sub>2</sub> at 900 °C for 1 hour is enough to introduce oxygen-containing functional groups that were uniformly distributed on the external surfaces of CNTs. The results revealed that the CO<sub>2</sub> oxidation thinned the CNTs walls and introduced numerous defects [140]. Another way to purify CNTs by eliminating the residual amorphous carbon and the metal particles, is via air treatment [141,142]. As evidenced by Li et al. [143], the results of the treatments with air are influenced by temperature: for a 20 minutes treatment at temperature between 480 °C and 650 °C the specific surface area of CNTs is increased, whereas at temperature greater than 700 °C surface area is decreased. The authors explained that at temperature greater than 700 °C the oxygen molecules present in the air first oxidize the amorphous carbon and then attack the CNTs capped ends. The capped ends are more reactive than the lateral walls of the CNTs because in the caps the C-C bonds are slightly weaker due to the radius of curvature (greater bond strain) and the presence of 5-fold and 7-fold rings [143,144]. Andrews et al. [145] show that at 1600 °C time plays a critical role. A 45 minutes treatment in air reduces the amount of amorphous carbon present on the surface, whereas for briefer treatment (less than 45 minutes) generated more oxidised sites on the CNTs. A recent article by some of the authors of this manuscript shows that an optimal oxidation and consequentially an optimal dispersion of the nanotubes in polar solvents, is possible when the CNTs are oxidized by a mixture of 2% O<sub>2</sub> and 98% N<sub>2</sub> at 350 °C [146]. Another method to functionalize CNTs is to use plasma functionalization in presence of different gasses (such as but not limited to O<sub>2</sub>, N<sub>2</sub>, NH<sub>3</sub>, Ar, CF<sub>4</sub>, CO<sub>2</sub>, H<sub>2</sub>, F<sub>2</sub>) [147]. Plasma can be produced via a variety of methods such as

dielectric barrier discharge, corona discharge, pulsed corona, microwave, radio frequency (RF) plasma, etc. [148]. Many studies have been carried out in the literature to modify the surface properties of carbon nanotubes using different combinations of gas and plasma parameters [149]. The main advantage of using plasma is that the functionalization temperature is low (from RT to approximately 600°C) [149], and the damage of the CNTs is also limited. Moreover, plasma allows nanotubes to be functionalized with complex molecules such as polymers [150] or biomolecules [151,152]. Main disadvantages are the low stability of the grafted functional groups over time, and the inhomogeneous functionalization of the sample's surface [153]. The latter occurs because only the face of the sample exposed to the plasma is functionalized, unless the powder sample is uniformly stirred during the treatment [154].

#### *4.2.1.4 Physical (or non-covalent) functionalization*

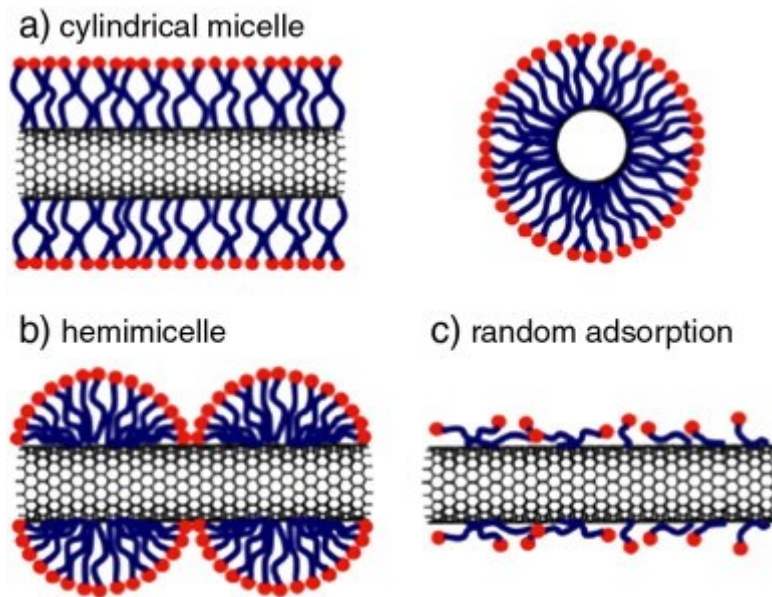
As discussed so far, it is clear that functionalization of CNTs by using covalent methods can efficiently introduce useful functional groups onto surface of carbon nanotubes. However, this kind of approach has two major drawbacks: i) the large number of lattice defects produced on the CNTs results in the depletion of their mechanical properties and the disruption of their p-electron system (thus with negative consequences on the electrical and thermal properties), and ii) the reactants adopted for these processes are dangerous and not environmental-friendly. Hence, many efforts have been put forward to develop methods able in reducing both the cost of the process and the damage to the CNTs structure. In this context, non-covalent functionalization is an alternative approach to tune the interfacial properties of CNTs. Non-covalent functionalizations are based on the physical interaction (physisorption) between CNTs and the chemicals and can be mainly classified as polymer wrapping or physical adsorption of surfactants.



**Figure 13:** Examples of polymer wrapping [155]

Polymer wrapping consists of a solution of suspended CNTs in the presence of polymers, such as poly(phenylenevinylene) [84] or polystyrene [156]. The van der Waals interactions and  $\pi$ - $\pi$  stacking between CNTs and polymer chains containing aromatic rings leads to the wrapping of the polymeric chains around the tubes to form supramolecular complexes (as in Figure 13) [155,157].

The other approach is based on the use of amphiphilic species such as surfactants (as in Figure 14). Several studies have been conducted on the effects of surfactants both on the dispersion of CNTs and their final properties [158–167]. Well-known in the literature, surfactants are usually organic amphiphilic compounds, meaning that they contain both hydrophobic groups (their tails) and hydrophilic groups (their heads). Surfactants can be divided into three categories: non-ionic surfactants such as Triton X-100 [161,162], anionic surfactants such as sodium dodecylsulfate (SDS) [164,165,168], and cationic surfactants (CTAB) [169,170]. As a result of the physical adsorption of surfactant on the surface of CNTs the surface tension of CNTs is decreased, and the inter-tube Van der Waals attraction is reduced, thus preventing the formation of agglomerates and bundles.



**Figure 14:** Models of surfactant-based physical functionalizations [171]

The efficiency of this method depends strongly on the properties of the surfactants; however it must be noted that due to the high surface area of the CNTs, a very large amount of surfactant is required to obtain a good dispersion. The use of large quantities of dispersant (in some cases the amount of dispersant is equal to that of the CNTs used) can hinder the interaction between the nanotubes and the matrix in which they are dispersed. As a consequence, the resulting mechanical, electrical and thermal performances of the composite might be much worse than expected [163,172]. Some researches, in fact, have proved that the dispersing capability of surfactant to CNTs increases with the ratio of surfactant to CNTs, but an excessive concentration level of surfactants has negative effect on the hydration of cement and will cause a marked air-entraining effect [173].

Table 4 summarizes all the main methods of dispersion and functionalization of CNTs shown in this review, with particular attention to the advantages and disadvantages of each one. As previously mentioned, functionalization and dispersion can be combined, reducing, in some cases, the disadvantages of the dispersion methods since changing the surface chemistry reduce the Van der Waals attraction between CNTs, thus preventing re-agglomeration.

**Table 4.** Dispersion technique

		Advantages	Disadvantages
DISPERSION	Ultrasonication	Excellent dispersion of nanotubes. Easy technique that can be coupled with other dispersion techniques	Risk of damaging the structure of the nanotube. If carried out for prolonged periods, it tends to heat the solution
	Calendaring	Usable for dispersion in polymeric matrices  The feeding materials should be in a viscous state	Very long times to obtain an excellent dispersion. Not as effective with low viscosity matrices
	Ball milling	Excellent for mixing nanotubes with metallic or ceramic powders	Risk of damaging the CNTs. Mainly usable for mixing with other powders
	Stir and melt compounding	Excellent for dispersing nanotubes in polymer matrix. Useful for disperse both high and low quantity of CNTs. Commonly used for thermoplastic polymer.	Possible re-agglomeration of CNTs after stir in statical condition.
FUNCTIONALIZATION	Acidic treatment in liquid phase	High surface functionalization	Risk of damage the CNTs. Not environmental friendly

	Oxidation without the use of acid in liquid phase	Eco-friendly. Nanotubes are usually not damaged	Low surface functionalization
	Functionalization using gas phase chemicals	Plasma allows to decorate CNTs with more complex functional groups.  Allows to remove amorphous carbon from the CNTs surface. Eco-friendly	Little stability of functional groups over time and lack of uniform functionalization over the surface of the sample (unless the sample is stirred). In some cases, high temperatures are required
	Physical functionalization	Allows an excellent dispersion of the nanotubes  No lattice damages	Large quantities of dispersant might be required to obtain an excellent dispersion. Reduced interaction between filler and matrix

## 5. Characterization techniques

It is important to use the most appropriate characterization technique to evaluate the effectiveness of the functionalization and to characterize the CNTs at the end of the treatment. In addition, the dispersion in polar matrices can be qualitatively evaluated by adding the CNTs in a polar solvent contained in a test tube and observing whether the CNTs aggregate and precipitate and how long it takes.

### 5.1 Raman spectroscopy

Raman spectroscopy is a non-destructive technique that allows researchers to evaluate the lattice defectivity of the nanotubes and their structure. The Raman spectrum, graphically represented as Intensity of the signal (arbitrary units) vs Raman shift ( $\text{cm}^{-1}$ ), of the CNTs can be characterized by the presence of 3 peaks. The D-band at circa  $1350 \text{ cm}^{-1}$ , arising from the disorder-induced phonon mode (A<sub>1g</sub>-band), the G-band at circa  $1590 \text{ cm}^{-1}$ , assigned to the Raman-allowed phonon mode (E<sub>2g</sub>-band) [174] and the Radial breathing mode (RBM) at circa  $248 \text{ cm}^{-1}$  which is visible only for single or double walled CNTs. The RBM frequency shifts depends on chirality and diameter [175]. The extent of the lattice defectivity can be evaluated by the ratio of the intensities of D-band over G-band (I<sub>D</sub>/I<sub>G</sub>) [176]. The higher is the ratio the greater is the amount of lattice defects and amorphous carbon. Even though this technique doesn't help when it comes to identify and quantify functional groups introduced on the CNTs, as a rule of thumb, the more chemical groups are covalently bonded to the CNTs the greater is the number of lattice defects.

## 5.2 UV-Vis-NIR spectroscopy

UV-Vis-NIR spectroscopy is based on absorption spectroscopy or reflectance spectroscopy in the ultraviolet-visible-near-infrared region collected using a UV-Vis spectrometer. UV-Vis spectroscopy is applicable to characterize CNTs [177] in liquid suspensions. As the CNTs are dispersed, the solution will become darker and will absorb more light and hence the maximum absorbance can be used as an indicator of dispersion quality. The ability to monitor liquid solution has the potential to detect the quality of dispersion of CNTs in solution [178]]. CNTs show absorption, in aqueous solution, between 262 and 230 nm respectively in the UV–visible spectrum. This is due to the  $\pi$ - $\pi^*$  transitions of aromatic C–C bonds [179]

## 5.3 X-ray diffraction (XRD)

This non-destructive characterization method is used to obtain information regarding the interlayer spacing of MWCNTs, the structural strain and the impurities contained in a CNTs sample or in a CNTs-nanocomposite. However, carbon nanotubes have multiple orientations compared to the X-ray incident beam. Diameters and chiralities distribution are also observed as well as various number of layers for MWNTs. This leads to a statistical characterization of carbon nanotubes. Due to their intrinsic nature, the main features of X-ray diffraction pattern of CNTs are close to those of graphite and presents two principal peaks: (i) a graphite-like peak (usually between  $20^\circ$  and  $30^\circ$ ) that measure the interlayer spacing of graphite layer and can be obtained from its position using the Bragg law, (ii) a family of peaks due to the honeycomb lattice of single graphene sheet. Consequently, the X-ray

diffraction profile is not useful to differentiate microstructural details between the CNTs and the graphite structure [180] but can help to determinate the sample purity (catalyst, functional groups).

#### 5.4 *Scanning Electron Microscopy (SEM)*

A scanning electron microscope is a type of electron microscope that produces images and analyzes the elemental composition (SEM-EDX) of a sample by scanning the surface with a focused beam of electrons. The electrons interact with atoms in the sample, producing various signals that contain information about the surface topography and the composition of the sample. SEM imaging cannot achieve resolution better than tens of nanometer. This technique is mostly used for the morphological characterization of CNTs or crosssections of nanocomposites containing CNTs. With this technology is possible evaluate also the state of agglomeration of CNTs and the defectivity of CNTs tube structure.

#### 5.5 *Transmission Electron Microscopy (TEM)*

Transmission electron microscopy is a microscopy technique in which a beam of electrons is transmitted through a specimen to form a see-through image. The specimen is usually either an ultrathin section of the sample (less than 10 nm thick) or a suspension on a grid. The resolving power (the minimum distance between two points for which they can be distinguished as such and not as one) is about 0.2 nm, that is about 20 times greater the best SEM. Given its great resolving power it is possible to see the different graphitic layers present in the MWCNTs [181]. It is possible to observe the metal catalyst trapped inside the tubes, and in some cases, it is also possible to distinguish functional groups anchored onto the surface of the nanotubes. This technique is used for the morphological characterization of CNTs.

#### 5.6 *X-ray photoelectron spectroscopy (XPS)*

X-ray photoelectron spectroscopy is a surface-sensitive quantitative spectroscopic technique that measures the elemental composition, the chemical state and electronic state of the elements that exist within a material. XPS is very useful to investigate functionalized CNTs by the analysis of the C1s peak in the region from 284.1 to 289.4 eV. The decomposition of the C1s peak could be used to identify and quantify the relative ratio of carbon bond types [182]. XPS technique can give information about the chemical structure of carbon nanotubes. But the most widely used data refers to the structure modification of the CNTs walls due to the chemical interaction with organic compounds or gases adsorption [183].

#### 5.7 *Small angle neutron scattering (SANS)*

Neutron scattering is used to determine the atomic and/or magnetic structure of a material. A sample to be examined is placed in a beam of thermal or cold neutrons to obtain a diffraction pattern that provides information of the structure of the material. The technique is similar to X-ray diffraction but due to their different scattering properties, neutrons and X-rays provide complementary information: X-Rays are suited for superficial analysis, strong x-rays from synchrotron radiation are suited for shallow depths or thin specimens, while neutrons having high penetration depth are suited for bulk samples [184]. This technique is mainly used for the morphological analysis of bulk samples of CNTs [185].

### 5.8 *Fourier Transform-Infrared spectroscopy (FT-IR)*

Infrared spectroscopy (IR spectroscopy) involves the absorption of infrared radiation with material. Samples may be solid, liquid, or gas. An IR spectrum can be visualized in a graph of infrared light absorbance (or transmittance) vs. frequency or wavelength. Typical units of frequency used in IR spectra are reciprocal centimeters (sometimes called wavenumbers), with the symbol  $\text{cm}^{-1}$ . Units of IR wavelength are commonly given in micrometers ( $\mu\text{m}$ ), which are related to wave numbers in a reciprocal way. A typical instrument that uses this technique is a Fourier transform infrared (FT-IR) spectrometer. In CNTs characterization, infrared spectroscopy is often used to determine impurities remaining from synthesis or molecules bonded on the nanotube surface. Numerous works are performed on organic molecules and CNTs: infrared spectroscopy exhibits all the modification of the CNTs structure and reveals the nature of compounds added to the CNTs [186]. However, the spectrum of nanotubes is difficult to visualize, since they tend to absorb most of the radiation incident on the sample. The determination of the functional groups on the surface of the nanotubes (especially if they contain long carbon chains) is complex, however polar functionalities containing oxygen can be detected [187]. This characterization technique must be coupled with one of the other characterization techniques. In particular, IR spectroscopy is complementary to Raman and they are often used together for a more complete characterization of the surface.

### 5.9 *Thermogravimetric analysis (TGA)*

Thermogravimetric analysis or thermal gravimetric analysis is a method of thermal analysis in which the mass variation of a sample is measured while the temperature changes. This measurement provides information about thermal stability, thermal decomposition, and thermal oxidation of samples [188]. This technique can be coupled to a mass or IR spectrophotometer to analyze the gases emitted during the thermal decomposition. TGA is mainly used in the study of the thermal stability

of CNTs (lattice defectivity affects the thermal stability), and differences in the thermal degradation between functionalized and pristine nanotubes can be distinguished.

**Table 5.** Different techniques used for the characterization of carbon nanotubes.

<b>Characterization technique</b>	<b>It can be used for:</b>
Raman spectroscopy	Detect single and double wall CNTs, determine the amount of lattice defects
UV-Vis-NIR spectroscopy	Evaluate the quality of CNTs dispersion in liquid solution
FT-IR and ATR-FTIR	Purity, typology and semiquantitative analysis regarding the amount of functional chemical groups
X-ray photoelectron spectroscopy (XPS)	Elemental composition, functionalization (covalent and non-covalent)
Thermogravimetric analysis	Purity, defectivity, presence of by-products, quality control of synthesis and manufacture processes
Neutron diffraction	Morphological analysis of bulk samples
X-ray diffraction	Morphological analysis of bulk samples
Scanning electron microscopy	Morphological analysis (diameter and length), agglomeration state
Transmission electron microscopy	Morphological analysis of internal structure: diameter, presence and location of metal catalyst, number of layers and distance between them

## 6. Suggestion for a good dispersion

The mechanism of dispersion is still unclear today. In the literature it is assumed that it proceeds by breaking large agglomerates of CNTs into smaller agglomerates. This phenomenon is supposed to occur through two mechanisms, namely rupture (a bulk phenomenon) and erosion (a surface

phenomenon). In rupture mechanism, the large agglomerates are broken down into smaller ones in short times, whereas in erosion mechanism, large agglomerates are eroded into smaller ones by removing single nanotubes or small nanotubes bundles from the agglomerate surface, thus needing comparatively longer time [189]. An efficient dispersion of carbon nanotubes inside a matrix plays a key role when CNTs are used as fillers in nanocomposites, hence the dispersion technique used significantly affects the final properties of the composite. Generally mechanical dispersion is used alone or in combination with chemical functionalization. A good approach to successfully disperse CNTs can be summarized in 5 points:

- Define the structure and dimension of CNTs agglomeration: this is important since the typology of agglomeration can affect the degree of CNTs dispersion. Big agglomerates require long time or high energy for dispersion, with the risk of damaging the structure of CNTs.
- Analyze the chemical properties of the solvent/matrix in which the nanotubes must be dispersed. The agglomerated CNTs can be dispersed directly in some solvents or polymers by directly using mechanical dispersion (such as ultrasonication or high shear stirring), thus obtaining good dispersion and good performance of composite.
- Perform a pre-dispersion in solution by breaking the agglomerates of nanotubes using mechanical dispersion. This step is essential for performing a homogeneous wet chemical functionalization of CNTs. Big agglomerates do not permit the penetration of chemical reactant in the middle of the agglomerate affecting the effectiveness of surface modification of CNTs in that position.
- Functionalize the CNTs to obtain a surface compatible with the medium in which they must be used. It is possible also couple the functionalization with mechanical dispersion methods, e.g. using ultrasonication in a solution with surfactant increasing the deagglomeration of CNTs and their stability in the final solution.
- Disperse the treated CNTs in the matrix by using the most appropriate mechanical dispersion method. This must be chosen not only as a function of the matrix used but also as a function of the properties researched with the use of CNTs. If the mechanical dispersion requires high energy that can damage the structure of CNTs, a loss mechanical and/or electrical properties of the final composite will occur.

It is important to remember that chemical functionalization can, as already mentioned above, alter the graphitic structure of the nanotubes. This can lead to a decrease in some properties of nanotubes (e.g. electrical conductivity). The functionalization of SWCNTs leads to a more severe reduction in electrical conductivity, while the effects are less severe for MWCNTs. If the functionalization is too

strong, the CNTs can be easily dispersed in the matrix or solvent, but this very good compatibilization with the matrix will not show beneficial effects on the final composite performance [190–192]. In addition to the methodologies presented in this paper, several new studies [193–198] are attempting to bypass the mechanical dispersion and chemical functionalization steps of the surface. These techniques plan a direct synthesis of CNTs on the powder with which the composite material is prepared.

## 7. Conclusion

In this review it was provided an overview of various aspects playing important roles in the optimal use of CNTs as nanofillers in composite materials. We underlined that it is useful to have a basic knowledge of the way CNTs are synthesized (CVD is the most used technology at industrial scale) and what CNTs properties can be expected as a consequence. For instance, as shown, single-wall and multi-wall CNTs can exhibit significant electric and mechanical dissimilarity, but differences can be also found within the same family of SWCNTs or MWCNTs due to the degree of lattice defectivity. Lattice defects can be present in the pristine CNTs because of the synthesis methodology used. But defects can also be introduced a posteriori when mechanical or chemical dispersion methods are applied to improve and facilitate the dispersion of the CNTs inside a matrix. As discussed, CNTs tend to be agglomerate in different form like bundles, combed yarn and bird-nest which must be untangled. Various mechanical dispersion methods can be used for this purpose. However, the mechanical dispersion is often not enough to disperse CNTs, especially inside polar media (CNTs have apolar graphitic structure). Thus, it is necessary to introduce a functionalization step. Several chemical and physical functionalization treatments have been illustrated in this work. All these methods change the CNTs surface, making them more polar and, consequently, more wettable in polar media. Functionalization can also help to produce a chemo-physical interaction with the matrix. However, the functionalization and mechanical dispersion techniques can also significantly damage the CNTs structure with consequent loss of the remarkable properties needed in the composite. Therefore, it is necessary a trade-off to minimize the CNTs damage as much as possible while maximizing the dispersion and interaction within the matrix. In order to help the reader with this endeavor, we have illustrated several characterization techniques to evaluate the damage and the properties of CNTs, both before and after treatments.

## References

- [1] A. Bianco, K. Kostarelos, C.D. Partidos, M. Prato, Biomedical applications of functionalised carbon nanotubes, *Chem. Commun.* (2005) 571–577. <https://doi.org/10.1039/B410943K>.

- [2] S. Singh, S. Vardharajula, P. Tiwari, Erdal Eroğlu, Komal Vig, V. Dennis, Ali, Functionalized carbon nanotubes: biomedical applications, *Int. J. Nanomedicine*. (2012) 5361. <https://doi.org/10.2147/IJN.S35832>.
- [3] A. Abate, J.-P. Correa-Baena, M. Saliba, M.S. Su'ait, F. Bella, Perovskite Solar Cells: From the Laboratory to the Assembly Line, *Chem. – Eur. J.* 24 (2018) 3083–3100. <https://doi.org/10.1002/chem.201704507>.
- [4] R.H. Baughman, A.A. Zakhidov, W.A. de Heer, Carbon Nanotubes--the Route Toward Applications, *Science*. 297 (2002) 787. <https://doi.org/10.1126/science.1060928>.
- [5] R. Nisticò, Magnetic materials and water treatments for a sustainable future, *Res. Chem. Intermed.* 43 (2017) 6911–6949. <https://doi.org/10.1007/s11164-017-3029-x>.
- [6] B.C.-K. Tee, A. Chortos, A. Berndt, A.K. Nguyen, A. Tom, A. McGuire, Z.C. Lin, K. Tien, W.-G. Bae, H. Wang, P. Mei, H.-H. Chou, B. Cui, K. Deisseroth, T.N. Ng, Z. Bao, A skin-inspired organic digital mechanoreceptor, *Science*. 350 (2015) 313. <https://doi.org/10.1126/science.aaa9306>.
- [7] P. Avouris, M. Freitag, V. Perebeinos, Carbon-Nanotube Optoelectronics, in: A. Jorio, G. Dresselhaus, M.S. Dresselhaus (Eds.), *Carbon Nanotub. Adv. Top. Synth. Struct. Prop. Appl.*, Springer Berlin Heidelberg, Berlin, Heidelberg, 2008: pp. 423–454. [https://doi.org/10.1007/978-3-540-72865-8\\_14](https://doi.org/10.1007/978-3-540-72865-8_14).
- [8] L. Lavagna, R. Nisticò, S. Musso, M. Pavese, Hydrophobic cellulose ester as a sustainable material for simple and efficient water purification processes from fatty oils contamination, *Wood Sci. Technol.* 53 (2019) 249–261. <https://doi.org/10.1007/s00226-018-1060-8>.
- [9] K.A. Williams, P.T.M. Veenhuizen, B.G. de la Torre, R. Eritja, C. Dekker, Carbon nanotubes with DNA recognition, *Nature*. 420 (2002) 761–761. <https://doi.org/10.1038/420761a>.
- [10] X. Ren, C. Chen, M. Nagatsu, X. Wang, Carbon nanotubes as adsorbents in environmental pollution management: A review, *Environ. Nanotechnol.* 170 (2011) 395–410. <https://doi.org/10.1016/j.cej.2010.08.045>.
- [11] L. Lavagna, D. Massella, M.F. Pantano, F. Bosia, N.M. Pugno, M. Pavese, Grafting carbon nanotubes onto carbon fibres doubles their effective strength and the toughness of the composite, *Compos. Sci. Technol.* 166 (2018) 140–149. <https://doi.org/10.1016/j.compscitech.2018.03.015>.
- [12] P.-C. Ma, N.A. Siddiqui, G. Marom, J.-K. Kim, Dispersion and functionalization of carbon nanotubes for polymer-based nanocomposites: A review, *Compos. Part Appl. Sci. Manuf.* 41 (2010) 1345–1367. <https://doi.org/10.1016/j.compositesa.2010.07.003>.
- [13] S. Barus, M. Zanetti, P. Bracco, S. Musso, A. Chiodoni, A. Tagliaferro, Influence of MWCNT morphology on dispersion and thermal properties of polyethylene nanocomposites, *Polym. Degrad. Stab.* 95 (2010) 756–762. <https://doi.org/10.1016/j.polymdegradstab.2010.02.013>.
- [14] A. Chiolerio, S. Musso, M. Sangermano, M. Giorcelli, S. Bianco, M. Coisson, A. Priola, P. Allia, A. Tagliaferro, Preparation of polymer-based composite with magnetic anisotropy by oriented carbon nanotube dispersion, *Proc. Diam. 2007 18th Eur. Conf. Diam. Diam.- Mater. Carbon Nanotub. Nitrides Silicon Carbide*. 17 (2008) 1590–1595. <https://doi.org/10.1016/j.diamond.2008.01.117>.
- [15] J. Liao, M.-J. Tan, Mixing of carbon nanotubes (CNTs) and aluminum powder for powder metallurgy use, *Powder Technol.* 208 (2011) 42–48. <https://doi.org/10.1016/j.powtec.2010.12.001>.
- [16] A. Saboori, M. Pavese, C. Badini, P. Fino, A Novel Approach to Enhance the Mechanical Strength and Electrical and Thermal Conductivity of Cu-GNP Nanocomposites, *Metall. Mater. Trans. A*. 49 (2018) 333–345. <https://doi.org/10.1007/s11661-017-4409-y>.
- [17] L. Lavagna, S. Musso, G. Ferro, M. Pavese, Cement-based composites containing functionalized carbon fibers, *Cem. Concr. Compos.* 88 (2018) 165–171. <https://doi.org/10.1016/j.cemconcomp.2018.02.007>.
- [18] A. Peigney, C. Laurent, E. Flahaut, A. Rousset, Carbon nanotubes in novel ceramic matrix nanocomposites, *Ceram. Int.* 26 (2000) 677–683. [https://doi.org/10.1016/S0272-8842\(00\)00004-3](https://doi.org/10.1016/S0272-8842(00)00004-3).
- [19] R. Nisticò, P. Evon, L. Labonne, G. Vaca-Medina, E. Montoneri, M. Francavilla, C. Vaca-Garcia, G. Magnacca, F. Franzoso, M. Negre, Extruded Poly(ethylene-co-vinyl alcohol) Composite Films Containing Biopolymers Isolated from Municipal Biowaste, *ChemistrySelect*. 1 (2016) 2354–2365. <https://doi.org/10.1002/slct.201600335>.
- [20] T. Ando, The electronic properties of graphene and carbon nanotubes, *NPG Asia Mater.* 1 (2009) 17–21. <https://doi.org/10.1038/asiamat.2009.1>.

- [21] J.N. Coleman, U. Khan, Y.K. Gun'ko, Mechanical Reinforcement of Polymers Using Carbon Nanotubes, *Adv. Mater.* 18 (2006) 689–706. <https://doi.org/10.1002/adma.200501851>.
- [22] L. Lavagna, D. Massella, M. Pavese, Preparation of hierarchical material by chemical grafting of carbon nanotubes onto carbon fibers, *Diam. Relat. Mater.* 80 (2017) 118–124. <https://doi.org/10.1016/j.diamond.2017.10.013>.
- [23] J. George, M.S. Sreekala, S. Thomas, A review on interface modification and characterization of natural fiber reinforced plastic composites, *Polym. Eng. Sci.* 41 (2001) 1471–1485. <https://doi.org/10.1002/pen.10846>.
- [24] S. Musso, M. Giorcelli, M. Pavese, S. Bianco, M. Rovere, A. Tagliaferro, Improving macroscopic physical and mechanical properties of thick layers of aligned multiwall carbon nanotubes by annealing treatment, *NDNC 2007 Proc. Int. Conf. New Diam. Nano Carbons 2007.* 17 (2008) 542–547. <https://doi.org/10.1016/j.diamond.2007.10.034>.
- [25] M.A. Worsley, S.O. Kucheyev, J.H. Satcher, A.V. Hamza, T.F. Baumann, Mechanically robust and electrically conductive carbon nanotube foams, *Appl. Phys. Lett.* 94 (2009) 073115. <https://doi.org/10.1063/1.3086293>.
- [26] K. Kordás, T. Mustonen, G. Tóth, H. Jantunen, M. Lajunen, C. Soldano, S. Talapatra, S. Kar, R. Vajtai, P.M. Ajayan, Inkjet Printing of Electrically Conductive Patterns of Carbon Nanotubes, *Small.* 2 (2006) 1021–1025. <https://doi.org/10.1002/sml.200600061>.
- [27] K.H. An, S.Y. Jeong, H.R. Hwang, Y.H. Lee, Enhanced Sensitivity of a Gas Sensor Incorporating Single-Walled Carbon Nanotube–Polypyrrole Nanocomposites, *Adv. Mater.* 16 (2004) 1005–1009. <https://doi.org/10.1002/adma.200306176>.
- [28] L. Sun, X. Wang, Y. Wang, Q. Zhang, Roles of carbon nanotubes in novel energy storage devices, *Carbon.* 122 (2017) 462–474. <https://doi.org/10.1016/j.carbon.2017.07.006>.
- [29] R. Lv, F. Kang, J. Gu, X. Gui, J. Wei, K. Wang, D. Wu, Carbon nanotubes filled with ferromagnetic alloy nanowires: Lightweight and wide-band microwave absorber, *Appl. Phys. Lett.* 93 (2008) 223105. <https://doi.org/10.1063/1.3042099>.
- [30] J.N. Coleman, U. Khan, W.J. Blau, Y.K. Gun'ko, Small but strong: A review of the mechanical properties of carbon nanotube–polymer composites, *Carbon.* 44 (2006) 1624–1652. <https://doi.org/10.1016/j.carbon.2006.02.038>.
- [31] P.-C. Ma, S.-Y. Mo, B.-Z. Tang, J.-K. Kim, Dispersion, interfacial interaction and re-agglomeration of functionalized carbon nanotubes in epoxy composites, *Carbon.* 48 (2010) 1824–1834. <https://doi.org/10.1016/j.carbon.2010.01.028>.
- [32] D. Tasis, N. Tagmatarchis, A. Bianco, M. Prato, Chemistry of Carbon Nanotubes, *Chem. Rev.* 106 (2006) 1105–1136. <https://doi.org/10.1021/cr050569o>.
- [33] E. Vázquez, F. Giacalone, M. Prato, Non-conventional methods and media for the activation and manipulation of carbon nanoforms, *Chem. Soc. Rev.* 43 (2014) 58–69. <https://doi.org/10.1039/C3CS60164A>.
- [34] J. Chen, L. Yan, W. Song, D. Xu, Interfacial characteristics of carbon nanotube-polymer composites: A review, *Compos. Part Appl. Sci. Manuf.* 114 (2018) 149–169. <https://doi.org/10.1016/j.compositesa.2018.08.021>.
- [35] K.Z. Milowska, M. Burda, L. Wolanicka, P.D. Bristowe, K.K.K. Koziol, Carbon nanotube functionalization as a route to enhancing the electrical and mechanical properties of Cu–CNT composites, *Nanoscale.* 11 (2019) 145–157. <https://doi.org/10.1039/C8NR07521B>.
- [36] S. Imani Yengejeh, S.A. Kazemi, A. Öchsner, Carbon nanotubes as reinforcement in composites: A review of the analytical, numerical and experimental approaches, *Comput. Mater. Sci.* 136 (2017) 85–101. <https://doi.org/10.1016/j.commatsci.2017.04.023>.
- [37] A.M.K. Esawi, M.M. Farag, Carbon nanotube reinforced composites: Potential and current challenges, *Mater. Des.* 28 (2007) 2394–2401. <https://doi.org/10.1016/j.matdes.2006.09.022>.
- [38] S.-Y. Li, H.-H. Zhou, J.-L. Gu, J. Zhu, Does carbyne really exist? — carbynes in expanded graphite, *Carbon.* 38 (2000) 934–937. [https://doi.org/10.1016/S0008-6223\(00\)00056-7](https://doi.org/10.1016/S0008-6223(00)00056-7).
- [39] J.-B. Donnet, É. Fousson, M. Samirant, T.K. Wang, M. Pontier-Johnson, A. Eckhardt, Shock synthesis of nanodiamonds from carbon precursors: identification of carbynes, *Comptes Rendus Académie Sci. - Ser. IIC - Chem.* 3 (2000) 359–364. [https://doi.org/10.1016/S1387-1609\(00\)01149-X](https://doi.org/10.1016/S1387-1609(00)01149-X).

- [40] H.W. Kroto, J.R. Heath, S.C. O'Brien, R.F. Curl, R.E. Smalley, C<sub>60</sub>: Buckminsterfullerene, *Nature*. 318 (1985) 162–163. <https://doi.org/10.1038/318162a0>.
- [41] W.S. Hummers, R.E. Offeman, Preparation of Graphitic Oxide, *J. Am. Chem. Soc.* 80 (1958) 1339–1339. <https://doi.org/10.1021/ja01539a017>.
- [42] S. Iijima, Helical microtubules of graphitic carbon, *Nature*. 354 (1991) 56–58. <https://doi.org/10.1038/354056a0>.
- [43] D.S. Bethune, C.H. Kiang, M.S. de Vries, G. Gorman, R. Savoy, J. Vazquez, R. Beyers, Cobalt-catalysed growth of carbon nanotubes with single-atomic-layer walls, *Nature*. 363 (1993) 605–607. <https://doi.org/10.1038/363605a0>.
- [44] S. Iijima, T. Ichihashi, Single-shell carbon nanotubes of 1-nm diameter, *Nature*. 363 (1993) 603–605. <https://doi.org/10.1038/363603a0>.
- [45] M. Monthieux, V.L. Kuznetsov, Who should be given the credit for the discovery of carbon nanotubes?, *Carbon*. 44 (2006) 1621–1623. <https://doi.org/10.1016/j.carbon.2006.03.019>.
- [46] X. Lu, Z. Chen, Curved Pi-Conjugation, Aromaticity, and the Related Chemistry of Small Fullerenes (, *Chem. Rev.* 105 (2005) 3643–3696. <https://doi.org/10.1021/cr030093d>.
- [47] A. Bhatt, A. Jain, E. Gurnany, R. Jain, A. Modi, A. Jain, 17 - Carbon Nanotubes: A Promising Carrier for Drug Delivery and Targeting, in: A.M. Holban, A.M. Grumezescu (Eds.), *Nanoarchitectonics Smart Deliv. Drug Target.*, William Andrew Publishing, 2016: pp. 465–501. <https://doi.org/10.1016/B978-0-323-47347-7.00017-3>.
- [48] H.O. Pierson, 3 - Graphite Structure and Properties, in: H.O. Pierson (Ed.), *Handb. Carbon Graph. Diam. Fuller.*, William Andrew Publishing, Oxford, 1993: pp. 43–69. <https://doi.org/10.1016/B978-0-8155-1339-1.50008-6>.
- [49] P.J. de Pablo, E. Graugnard, B. Walsh, R.P. Andres, S. Datta, R. Reifengerger, A simple, reliable technique for making electrical contact to multiwalled carbon nanotubes, *Appl. Phys. Lett.* 74 (1999) 323–325. <https://doi.org/10.1063/1.123011>.
- [50] D. Dasgupta, F. Demichelis, A. Tagliaferro, Electrical conductivity of amorphous carbon and amorphous hydrogenated carbon, *Philos. Mag. B.* 63 (1991) 1255–1266. <https://doi.org/10.1080/13642819108205558>.
- [51] H. Dai, E.W. Wong, C.M. Lieber, Probing Electrical Transport in Nanomaterials: Conductivity of Individual Carbon Nanotubes, *Science*. 272 (1996) 523. <https://doi.org/10.1126/science.272.5261.523>.
- [52] A. Lekawa-Raus, J. Patmore, L. Kurzepa, J. Bulmer, K. Koziol, Electrical Properties of Carbon Nanotube Based Fibers and Their Future Use in Electrical Wiring, *Adv. Funct. Mater.* 24 (2014) 3661–3682. <https://doi.org/10.1002/adfm.201303716>.
- [53] P. Zhang, P.E. Lammert, V.H. Crespi, Plastic Deformations of Carbon Nanotubes, *Phys. Rev. Lett.* 81 (1998) 5346–5349. <https://doi.org/10.1103/PhysRevLett.81.5346>.
- [54] V.N. Popov, Carbon nanotubes: properties and application, *Mater. Sci. Eng. R Rep.* 43 (2004) 61–102. <https://doi.org/10.1016/j.mser.2003.10.001>.
- [55] Ch. Laurent, E. Flahaut, A. Peigney, The weight and density of carbon nanotubes versus the number of walls and diameter, *Carbon*. 48 (2010) 2994–2996. <https://doi.org/10.1016/j.carbon.2010.04.010>.
- [56] M. Miao, Electrical conductivity of pure carbon nanotube yarns, *Carbon*. 49 (2011) 3755–3761. <https://doi.org/10.1016/j.carbon.2011.05.008>.
- [57] W.D. Callister Jr, D.G. Rethwisch, *Fundamentals of materials science and engineering: an integrated approach*, John Wiley & Sons, 2020.
- [58] L.V. Radushkevich, V.M. Lukyanovich, O strukture ugljeroda, obrazujucesosja pri termiceskom razlozenii okisi ugljeroda na zeleznom kontakte, *Zurn Fis. Chim.* 26 (1952) 88–95.
- [59] R.T.K. Baker, P.S. Harris, R.B. Thomas, R.J. Waite, Formation of filamentous carbon from iron, cobalt and chromium catalyzed decomposition of acetylene, *J. Catal.* 30 (1973) 86–95. [https://doi.org/10.1016/0021-9517\(73\)90055-9](https://doi.org/10.1016/0021-9517(73)90055-9).
- [60] Y. Ando, The Preparation of Carbon Nanotubes, *Fuller. Sci. Technol.* 2 (1994) 173–180. <https://doi.org/10.1080/15363839408009542>.

- [61] Y. Saito, K. Nishikubo, K. Kawabata, T. Matsumoto, Carbon nanocapsules and single-layered nanotubes produced with platinum-group metals (Ru, Rh, Pd, Os, Ir, Pt) by arc discharge, *J. Appl. Phys.* 80 (1996) 3062–3067. <https://doi.org/10.1063/1.363166>.
- [62] A. Loiseau, P. Launois, P. Petit, S. Roche, J.-P. Salvetat, Understanding carbon nanotubes, *Lect Notes Phys.* 677 (2006) 495–543.
- [63] J.J. Schneider, N.I. Maksimova, J. Engstler, R. Joshi, R. Schierholz, R. Feile, Catalyst free growth of a carbon nanotube–alumina composite structure, *Protag. Chem. Piero Zanello.* 361 (2008) 1770–1778. <https://doi.org/10.1016/j.ica.2006.10.025>.
- [64] J. Gao, J. Zhong, L. Bai, J. Liu, G. Zhao, X. Sun, Revealing the Role of Catalysts in Carbon Nanotubes and Nanofibers by Scanning Transmission X-ray Microscopy, *Sci. Rep.* 4 (2015) 3606. <https://doi.org/10.1038/srep03606>.
- [65] Jay P. Gore, Flame Synthesis of Carbon Nanotubes, in: Anup Sane ED1 - Siva Yellampalli (Ed.), *Carbon Nanotub.*, IntechOpen, Rijeka, 2011: p. Ch. 7. <https://doi.org/10.5772/21012>.
- [66] A.M. Morales, C.M. Lieber, A Laser Ablation Method for the Synthesis of Crystalline Semiconductor Nanowires, *Science.* 279 (1998) 208. <https://doi.org/10.1126/science.279.5348.208>.
- [67] M. Yudasaka, R. Yamada, N. Sensui, T. Wilkins, T. Ichihashi, S. Iijima, Mechanism of the Effect of NiCo, Ni and Co Catalysts on the Yield of Single-Wall Carbon Nanotubes Formed by Pulsed Nd:YAG Laser Ablation, *J. Phys. Chem. B.* 103 (1999) 6224–6229. <https://doi.org/10.1021/jp9908451>.
- [68] P. Bernier, D. Laplaze, J. Auriol, L. Barbedette, G. Flamant, M. Lebrun, A. Brunelle, S. Della-Negra, Production of fullerenes from solar energy, *Synth. Met.* 70 (1995) 1455–1456.
- [69] D. Laplaze, P. Bernier, W.K. Maser, G. Flamant, T. Guillard, A. Loiseau, Carbon nanotubes: The solar approach, *Eur. Mater. Res. Soc. 1997 Meet. Symp. Fullerenes Carbon Based Mater.* 36 (1998) 685–688. [https://doi.org/10.1016/S0008-6223\(98\)00025-6](https://doi.org/10.1016/S0008-6223(98)00025-6).
- [70] P. Nikolaev, M.J. Bronikowski, R.K. Bradley, F. Rohmund, D.T. Colbert, K.A. Smith, R.E. Smalley, Gas-phase catalytic growth of single-walled carbon nanotubes from carbon monoxide, *Chem. Phys. Lett.* 313 (1999) 91–97. [https://doi.org/10.1016/S0009-2614\(99\)01029-5](https://doi.org/10.1016/S0009-2614(99)01029-5).
- [71] L. Ci, S. Xie, D. Tang, X. Yan, Y. Li, Z. Liu, X. Zou, W. Zhou, G. Wang, Controllable growth of single wall carbon nanotubes by pyrolyzing acetylene on the floating iron catalysts, *Chem. Phys. Lett.* 349 (2001) 191–195. [https://doi.org/10.1016/S0009-2614\(01\)01098-3](https://doi.org/10.1016/S0009-2614(01)01098-3).
- [72] H.W. Zhu, C.L. Xu, D.H. Wu, B.Q. Wei, R. Vajtai, P.M. Ajayan, Direct Synthesis of Long Single-Walled Carbon Nanotube Strands, *Science.* 296 (2002) 884. <https://doi.org/10.1126/science.1066996>.
- [73] Y. Ning, X. Zhang, Y. Wang, Y. Sun, L. Shen, X. Yang, G. Van Tendeloo, Bulk production of multi-wall carbon nanotube bundles on sol–gel prepared catalyst, *Chem. Phys. Lett.* 366 (2002) 555–560. [https://doi.org/10.1016/S0009-2614\(02\)01647-0](https://doi.org/10.1016/S0009-2614(02)01647-0).
- [74] R.R. Bacsa, Ch. Laurent, A. Peigney, W.S. Bacsa, Th. Vaugien, A. Rousset, High specific surface area carbon nanotubes from catalytic chemical vapor deposition process, *Chem. Phys. Lett.* 323 (2000) 566–571. [https://doi.org/10.1016/S0009-2614\(00\)00558-3](https://doi.org/10.1016/S0009-2614(00)00558-3).
- [75] E. Flahaut, A. Govindaraj, A. Peigney, Ch. Laurent, A. Rousset, C.N.R. Rao, Synthesis of single-walled carbon nanotubes using binary (Fe, Co, Ni) alloy nanoparticles prepared in situ by the reduction of oxide solid solutions, *Chem. Phys. Lett.* 300 (1999) 236–242. [https://doi.org/10.1016/S0009-2614\(98\)01304-9](https://doi.org/10.1016/S0009-2614(98)01304-9).
- [76] A.M. Cassell, J.A. Raymakers, J. Kong, H. Dai, Large Scale CVD Synthesis of Single-Walled Carbon Nanotubes, *J. Phys. Chem. B.* 103 (1999) 6484–6492. <https://doi.org/10.1021/jp990957s>.
- [77] C.H. See, A.T. Harris, A comparison of carbon nanotube synthesis in fixed and fluidised bed reactors, *Chem. Eng. J.* 144 (2008) 267–269. <https://doi.org/10.1016/j.cej.2008.06.002>.
- [78] Y.C. Choi, Y.M. Shin, Y.H. Lee, B.S. Lee, G.-S. Park, W.B. Choi, N.S. Lee, J.M. Kim, Controlling the diameter, growth rate, and density of vertically aligned carbon nanotubes synthesized by microwave plasma-enhanced chemical vapor deposition, *Appl. Phys. Lett.* 76 (2000) 2367–2369. <https://doi.org/10.1063/1.126348>.
- [79] H.K. Jung, H.W. Lee, Effect of Catalytic Layer Thickness on Diameter of Vertically Aligned Individual Carbon Nanotubes, *J. Nanomater.* 2014 (2014) 1–6. <https://doi.org/10.1155/2014/270989>.
- [80] S. Slomkowski, J.V. Alemán, R.G. Gilbert, M. Hess, K. Horie, R.G. Jones, P. Kubisa, I. Meisel, W. Mormann, S. Penczek, R.F.T. Stepto, Terminology of polymers and polymerization processes in

dispersed systems (IUPAC Recommendations 2011), *Pure Appl. Chem.* 83 (2011) 2229–2259. <https://doi.org/10.1351/PAC-REC-10-06-03>.

- [81] D.H. Everett, *Manual of Symbols and Terminology for Physicochemical Quantities and Units, Appendix II: Definitions, Terminology and Symbols in Colloid and Surface Chemistry*, *Pure Appl. Chem.* 31 (1972) 577–638. <https://doi.org/10.1351/pac197231040577>.
- [82] X.-L. Xie, Y.-W. Mai, X.-P. Zhou, Dispersion and alignment of carbon nanotubes in polymer matrix: A review, *Mater. Sci. Eng. R Rep.* 49 (2005) 89–112. <https://doi.org/10.1016/j.mser.2005.04.002>.
- [83] E.T. Thostenson, Z. Ren, T.-W. Chou, Advances in the science and technology of carbon nanotubes and their composites: a review, *Compos. Sci. Technol.* 61 (2001) 1899–1912. [https://doi.org/10.1016/S0266-3538\(01\)00094-X](https://doi.org/10.1016/S0266-3538(01)00094-X).
- [84] B. Mc Carthy, J.N. Coleman, R. Czerw, A.B. Dalton, D.L. Carroll, W.J. Blau, Microscopy studies of nanotube-conjugated polymer interactions, *Proc. Int. Conf. Sci. Technol. Synth. Met.* 121 (2001) 1225–1226. [https://doi.org/10.1016/S0379-6779\(00\)00906-1](https://doi.org/10.1016/S0379-6779(00)00906-1).
- [85] Y.Y. Huang, E.M. Terentjev, Dispersion of Carbon Nanotubes: Mixing, Sonication, Stabilization, and Composite Properties, *Polymers*. 4 (2012) 275–295. <https://doi.org/10.3390/polym4010275>.
- [86] B. Krause, M. Mende, P. Pötschke, G. Petzold, Dispersability and particle size distribution of CNTs in an aqueous surfactant dispersion as a function of ultrasonic treatment time, *Carbon*. 48 (2010) 2746–2754. <https://doi.org/10.1016/j.carbon.2010.04.002>.
- [87] C. Tang, T. Zhou, J. Yang, Q. Zhang, F. Chen, Q. Fu, L. Yang, Wet-grinding assisted ultrasonic dispersion of pristine multi-walled carbon nanotubes (MWCNTs) in chitosan solution, *Colloids Surf. B Biointerfaces*. 86 (2011) 189–197. <https://doi.org/10.1016/j.colsurfb.2011.03.041>.
- [88] H. Li, J.C. Nie, S. Kunsági-Máté, Modified dispersion of functionalized multi-walled carbon nanotubes in acetonitrile, *Chem. Phys. Lett.* 492 (2010) 258–262. <https://doi.org/10.1016/j.cplett.2010.04.053>.
- [89] X. Xu, Z. Wang, Non-covalent dispersed carbon nanotube–benzocyclobutene composites as a bonding interface material for three-dimensional integration, *Microelectron. Eng.* 91 (2012) 33–38. <https://doi.org/10.1016/j.mee.2011.10.010>.
- [90] H. Sato, M. Sano, Characteristics of ultrasonic dispersion of carbon nanotubes aided by antifoam, *Colloids Surf. Physicochem. Eng. Asp.* 322 (2008) 103–107. <https://doi.org/10.1016/j.colsurfa.2008.02.031>.
- [91] J. Hilding, E.A. Grulke, Z. George Zhang, F. Lockwood, Dispersion of Carbon Nanotubes in Liquids, *J. Dispers. Sci. Technol.* 24 (2003) 1–41. <https://doi.org/10.1081/DIS-120017941>.
- [92] J. Zhou, J. Cheiftz, R. Li, F. Wang, X. Zhou, T.-K. Sham, X. Sun, Z. Ding, Tailoring multi-wall carbon nanotubes for smaller nanostructures, *Carbon*. 47 (2009) 829–838. <https://doi.org/10.1016/j.carbon.2008.11.032>.
- [93] X. Gao, A.I. Isayev, C. Yi, Ultrasonic treatment of polycarbonate/carbon nanotubes composites, *Polymer*. 84 (2016) 209–222. <https://doi.org/10.1016/j.polymer.2015.12.051>.
- [94] R. Arrigo, R. Teresi, C. Gambarotti, F. Parisi, G. Lazzara, N. Dintcheva, Sonication-Induced Modification of Carbon Nanotubes: Effect on the Rheological and Thermo-Oxidative Behaviour of Polymer-Based Nanocomposites, *Materials*. 11 (2018) 383. <https://doi.org/10.3390/ma11030383>.
- [95] C. Schilde, H. Nolte, C. Arlt, A. Kwade, Effect of fluid–particle-interactions on dispersing nano-particles in epoxy resins using stirred-media-mills and three-roll-mills, *Compos. Sci. Technol.* 70 (2010) 657–663. <https://doi.org/10.1016/j.compscitech.2009.12.021>.
- [96] E.T. Thostenson, T.-W. Chou, Processing-structure-multi-functional property relationship in carbon nanotube/epoxy composites, *Carbon*. 44 (2006) 3022–3029. <https://doi.org/10.1016/j.carbon.2006.05.014>.
- [97] M. Sánchez, M. Campo, A. Jiménez-Suárez, A. Ureña, Effect of the carbon nanotube functionalization on flexural properties of multiscale carbon fiber/epoxy composites manufactured by VARIM, *Compos. Part B Eng.* 45 (2013) 1613–1619. <https://doi.org/10.1016/j.compositesb.2012.09.063>.
- [98] F.H. Gojny, M.H.G. Wichmann, U. Köpke, B. Fiedler, K. Schulte, Carbon nanotube-reinforced epoxy-composites: enhanced stiffness and fracture toughness at low nanotube content, *Dev. Carbon Nanotube Nanofibre Reinf. Polym.* 64 (2004) 2363–2371. <https://doi.org/10.1016/j.compscitech.2004.04.002>.

- [99] P. Pötschke, B. Krause, S.T. Buschhorn, U. Köpke, M.T. Müller, T. Villmow, K. Schulte, Improvement of carbon nanotube dispersion in thermoplastic composites using a three roll mill at elevated temperatures, *Compos. Sci. Technol.* 74 (2013) 78–84. <https://doi.org/10.1016/j.compscitech.2012.10.010>.
- [100] S. Budin, I.P. Almanar, S. Kamaruddin, N.C. Maideen, A.H. Zulkifli, Modeling of vial and ball motions for an effective mechanical milling process, *J. Mater. Process. Technol.* 209 (2009) 4312–4319. <https://doi.org/10.1016/j.jmatprotec.2008.11.016>.
- [101] C. Suryanarayana, Mechanical alloying and milling, *Prog. Mater. Sci.* 46 (2001) 1–184. [https://doi.org/10.1016/S0079-6425\(99\)00010-9](https://doi.org/10.1016/S0079-6425(99)00010-9).
- [102] H.J. Choi, J.H. Shin, D.H. Bae, The effect of milling conditions on microstructures and mechanical properties of Al/MWCNT composites, *Compos. Part Appl. Sci. Manuf.* 43 (2012) 1061–1072. <https://doi.org/10.1016/j.compositesa.2012.02.008>.
- [103] Y.A. Kim, T. Hayashi, Y. Fukai, M. Endo, T. Yanagisawa, M.S. Dresselhaus, Effect of ball milling on morphology of cup-stacked carbon nanotubes, *Chem. Phys. Lett.* 355 (2002) 279–284. [https://doi.org/10.1016/S0009-2614\(02\)00248-8](https://doi.org/10.1016/S0009-2614(02)00248-8).
- [104] K. Menzer, B. Krause, R. Boldt, B. Kretzschmar, R. Weidisch, P. Pötschke, Percolation behaviour of multiwalled carbon nanotubes of altered length and primary agglomerate morphology in melt mixed isotactic polypropylene-based composites, *Compos. Sci. Technol.* 71 (2011) 1936–1943. <https://doi.org/10.1016/j.compscitech.2011.09.009>.
- [105] A. Esawi, K. Morsi, Dispersion of carbon nanotubes (CNTs) in aluminum powder, *Compos. Part Appl. Sci. Manuf.* 38 (2007) 646–650. <https://doi.org/10.1016/j.compositesa.2006.04.006>.
- [106] Z.Y. Liu, S.J. Xu, B.L. Xiao, P. Xue, W.G. Wang, Z.Y. Ma, Effect of ball-milling time on mechanical properties of carbon nanotubes reinforced aluminum matrix composites, *Compos. Part Appl. Sci. Manuf.* 43 (2012) 2161–2168. <https://doi.org/10.1016/j.compositesa.2012.07.026>.
- [107] R. Pérez-Bustamante, J.L. Bueno-Escobedo, J. Jiménez-Lobato, I. Estrada-Guel, M. Miki-Yoshida, L. Licea-Jiménez, R. Martínez-Sánchez, Wear behavior in Al<sub>2024</sub>-CNTs composites synthesized by mechanical alloying, *Wear*. 292–293 (2012) 169–175. <https://doi.org/10.1016/j.wear.2012.05.016>.
- [108] Y. Wu, G.-Y. Kim, A.M. Russell, Mechanical alloying of carbon nanotube and Al6061 powder for metal matrix composites, *Mater. Sci. Eng. A.* 532 (2012) 558–566. <https://doi.org/10.1016/j.msea.2011.10.121>.
- [109] N. Darsono, D.-H. Yoon, J. Kim, Milling and dispersion of multi-walled carbon nanotubes in texanol, *Appl. Surf. Sci.* 254 (2008) 3412–3419. <https://doi.org/10.1016/j.apsusc.2007.11.028>.
- [110] A.M.K. Esawi, K. Morsi, A. Sayed, A.A. Gawad, P. Borah, Fabrication and properties of dispersed carbon nanotube–aluminum composites, *Mater. Sci. Eng. A.* 508 (2009) 167–173. <https://doi.org/10.1016/j.msea.2009.01.002>.
- [111] J. Sandler, M.S.P. Shaffer, T. Prasse, W. Bauhofer, K. Schulte, A.H. Windle, Development of a dispersion process for carbon nanotubes in an epoxy matrix and the resulting electrical properties, *Polymer*. 40 (1999) 5967–5971. [https://doi.org/10.1016/S0032-3861\(99\)00166-4](https://doi.org/10.1016/S0032-3861(99)00166-4).
- [112] C.F. Schmid, D.J. Klingenberg, Mechanical Flocculation in Flowing Fiber Suspensions, *Phys. Rev. Lett.* 84 (2000) 290–293. <https://doi.org/10.1103/PhysRevLett.84.290>.
- [113] T. Villmow, P. Pötschke, S. Pegel, L. Häussler, B. Kretzschmar, Influence of twin-screw extrusion conditions on the dispersion of multi-walled carbon nanotubes in a poly(lactic acid) matrix, *Polymer*. 49 (2008) 3500–3509. <https://doi.org/10.1016/j.polymer.2008.06.010>.
- [114] A. Hirsch, Functionalization of Single-Walled Carbon Nanotubes, *Angew. Chem. Int. Ed.* 41 (2002) 1853–1859. [https://doi.org/10.1002/1521-3773\(20020603\)41:11<1853::AID-ANIE1853>3.0.CO;2-N](https://doi.org/10.1002/1521-3773(20020603)41:11<1853::AID-ANIE1853>3.0.CO;2-N).
- [115] A. Hirsch, O. Vostrowsky, Functionalization of Carbon Nanotubes, in: A.D. Schlüter (Ed.), *Funct. Mol. Nanostructures --*, Springer Berlin Heidelberg, Berlin, Heidelberg, 2005: pp. 193–237. <https://doi.org/10.1007/b98169>.
- [116] K. Balasubramanian, M. Burghard, Chemically Functionalized Carbon Nanotubes, *Small*. 1 (2005) 180–192. <https://doi.org/10.1002/sml.200400118>.
- [117] S. Banerjee, M.G.C. Kahn, S.S. Wong, Rational Chemical Strategies for Carbon Nanotube Functionalization, *Chem. – Eur. J.* 9 (2003) 1898–1908. <https://doi.org/10.1002/chem.200204618>.

- [118] V. Datsyuk, M. Kalyva, K. Papagelis, J. Parthenios, D. Tasis, A. Siokou, I. Kallitsis, C. Galiotis, Chemical oxidation of multiwalled carbon nanotubes, *Carbon*. 46 (2008) 833–840. <https://doi.org/10.1016/j.carbon.2008.02.012>.
- [119] Ş.S. Bayazit, İ. İnci, Adsorption of Cu (II) ions from water by carbon nanotubes oxidized with UV-light and ultrasonication, *J. Mol. Liq.* 199 (2014) 559–564. <https://doi.org/10.1016/j.molliq.2014.10.001>.
- [120] F. Avilés, J.V. Cauich-Rodríguez, L. Moo-Tah, A. May-Pat, R. Vargas-Coronado, Evaluation of mild acid oxidation treatments for MWCNT functionalization, *Carbon*. 47 (2009) 2970–2975. <https://doi.org/10.1016/j.carbon.2009.06.044>.
- [121] I.D. Rosca, F. Watari, M. Uo, T. Akasaka, Oxidation of multiwalled carbon nanotubes by nitric acid, *Carbon*. 43 (2005) 3124–3131. <https://doi.org/10.1016/j.carbon.2005.06.019>.
- [122] J. Zhang, H. Zou, Q. Qing, Y. Yang, Q. Li, Z. Liu, X. Guo, Z. Du, Effect of Chemical Oxidation on the Structure of Single-Walled Carbon Nanotubes, *J. Phys. Chem. B*. 107 (2003) 3712–3718. <https://doi.org/10.1021/jp027500u>.
- [123] K.A. Wepasnick, B.A. Smith, K.E. Schrote, H.K. Wilson, S.R. Diegelmann, D.H. Fairbrother, Surface and structural characterization of multi-walled carbon nanotubes following different oxidative treatments, *Carbon*. 49 (2011) 24–36. <https://doi.org/10.1016/j.carbon.2010.08.034>.
- [124] T.A. Saleh, The influence of treatment temperature on the acidity of MWCNT oxidized by HNO<sub>3</sub> or a mixture of HNO<sub>3</sub>/H<sub>2</sub>SO<sub>4</sub>, *Appl. Surf. Sci.* 257 (2011) 7746–7751. <https://doi.org/10.1016/j.apsusc.2011.04.020>.
- [125] M. Pavese, S. Musso, S. Bianco, M. Giorcelli, N. Pugno, An analysis of carbon nanotube structure wettability before and after oxidation treatment, *J. Phys. Condens. Matter*. 20 (2008) 474206. <https://doi.org/10.1088/0953-8984/20/47/474206>.
- [126] Y. Peng, H. Liu, Effects of Oxidation by Hydrogen Peroxide on the Structures of Multiwalled Carbon Nanotubes, *Ind. Eng. Chem. Res.* 45 (2006) 6483–6488. <https://doi.org/10.1021/ie0604627>.
- [127] Y.J. Kim, T.S. Shin, H.D. Choi, J.H. Kwon, Y.-C. Chung, H.G. Yoon, Electrical conductivity of chemically modified multiwalled carbon nanotube/epoxy composites, *Carbon*. 43 (2005) 23–30. <https://doi.org/10.1016/j.carbon.2004.08.015>.
- [128] C. Moreno-Castilla, M.V. López-Ramón, F. Carrasco-Marín, Changes in surface chemistry of activated carbons by wet oxidation, *Carbon*. 38 (2000) 1995–2001. [https://doi.org/10.1016/S0008-6223\(00\)00048-8](https://doi.org/10.1016/S0008-6223(00)00048-8).
- [129] D.B. Mawhinney, V. Naumenko, A. Kuznetsova, J.T. Yates, J. Liu, R.E. Smalley, Infrared Spectral Evidence for the Etching of Carbon Nanotubes: Ozone Oxidation at 298 K, *J. Am. Chem. Soc.* 122 (2000) 2383–2384. <https://doi.org/10.1021/ja994094s>.
- [130] J.M. Simmons, B.M. Nichols, S.E. Baker, M.S. Marcus, O.M. Castellini, C.-S. Lee, R.J. Hamers, M.A. Eriksson, Effect of Ozone Oxidation on Single-Walled Carbon Nanotubes, *J. Phys. Chem. B*. 110 (2006) 7113–7118. <https://doi.org/10.1021/jp0548422>.
- [131] K.J. Ziegler, Z. Gu, H. Peng, E.L. Flor, R.H. Hauge, R.E. Smalley, Controlled Oxidative Cutting of Single-Walled Carbon Nanotubes, *J. Am. Chem. Soc.* 127 (2005) 1541–1547. <https://doi.org/10.1021/ja044537e>.
- [132] A.G. Osorio, I.C.L. Silveira, V.L. Bueno, C.P. Bergmann, H<sub>2</sub>SO<sub>4</sub>/HNO<sub>3</sub>/HCl—Functionalization and its effect on dispersion of carbon nanotubes in aqueous media, *Appl. Surf. Sci.* 255 (2008) 2485–2489. <https://doi.org/10.1016/j.apsusc.2008.07.144>.
- [133] M.N. Tchoul, W.T. Ford, G. Lolli, D.E. Resasco, S. Arepalli, Effect of Mild Nitric Acid Oxidation on Dispersability, Size, and Structure of Single-Walled Carbon Nanotubes, *Chem. Mater.* 19 (2007) 5765–5772. <https://doi.org/10.1021/cm071758l>.
- [134] Y. Xing, L. Li, C.C. Chusuei, R.V. Hull, Sonochemical Oxidation of Multiwalled Carbon Nanotubes, *Langmuir*. 21 (2005) 4185–4190. <https://doi.org/10.1021/la047268e>.
- [135] W. Xia, C. Jin, S. Kundu, M. Muhler, A highly efficient gas-phase route for the oxygen functionalization of carbon nanotubes based on nitric acid vapor, *Carbon*. 47 (2009) 919–922. <https://doi.org/10.1016/j.carbon.2008.12.026>.
- [136] M. Li, M. Boggs, T.P. Beebe, C.P. Huang, Oxidation of single-walled carbon nanotubes in dilute aqueous solutions by ozone as affected by ultrasound, *Carbon*. 46 (2008) 466–475. <https://doi.org/10.1016/j.carbon.2007.12.012>.

- [137] M.-L. Sham, J.-K. Kim, Surface functionalities of multi-wall carbon nanotubes after UV/Ozone and TETA treatments, *Carbon*. 44 (2006) 768–777. <https://doi.org/10.1016/j.carbon.2005.09.013>.
- [138] I.W. Chiang, B.E. Brinson, R.E. Smalley, J.L. Margrave, R.H. Hauge, Purification and Characterization of Single-Wall Carbon Nanotubes, *J. Phys. Chem. B*. 105 (2001) 1157–1161. <https://doi.org/10.1021/jp003453z>.
- [139] D. Wang, K. Wang, H. Wu, Y. Luo, L. Sun, Y. Zhao, J. Wang, L. Jia, K. Jiang, Q. Li, S. Fan, J. Wang, CO<sub>2</sub> oxidation of carbon nanotubes for lithium-sulfur batteries with improved electrochemical performance, *Carbon*. 132 (2018) 370–379. <https://doi.org/10.1016/j.carbon.2018.02.048>.
- [140] M.R. Smith, S.W. Hedges, R. LaCount, D. Kern, N. Shah, G.P. Huffman, B. Bockrath, Selective oxidation of single-walled carbon nanotubes using carbon dioxide, *Carbon*. 41 (2003) 1221–1230. [https://doi.org/10.1016/S0008-6223\(03\)00054-X](https://doi.org/10.1016/S0008-6223(03)00054-X).
- [141] Y.S. Park, Y.C. Choi, K.S. Kim, D.-C. Chung, D.J. Bae, K.H. An, S.C. Lim, X.Y. Zhu, Y.H. Lee, High yield purification of multiwalled carbon nanotubes by selective oxidation during thermal annealing, *Carbon*. 39 (2001) 655–661. [https://doi.org/10.1016/S0008-6223\(00\)00152-4](https://doi.org/10.1016/S0008-6223(00)00152-4).
- [142] P.-X. Hou, C. Liu, H.-M. Cheng, Purification of carbon nanotubes, *Carbon*. 46 (2008) 2003–2025. <https://doi.org/10.1016/j.carbon.2008.09.009>.
- [143] C. Li, D. Wang, T. Liang, X. Wang, J. Wu, X. Hu, J. Liang, Oxidation of multiwalled carbon nanotubes by air: benefits for electric double layer capacitors, *Powder Technol.* 142 (2004) 175–179. <https://doi.org/10.1016/j.powtec.2004.04.037>.
- [144] R. Sen, S.M. Rickard, M.E. Itkis, R.C. Haddon, Controlled Purification of Single-Walled Carbon Nanotube Films by Use of Selective Oxidation and Near-IR Spectroscopy, *Chem. Mater.* 15 (2003) 4273–4279. <https://doi.org/10.1021/cm0342997>.
- [145] R. Andrews, D. Jacques, D. Qian, E.C. Dickey, Purification and structural annealing of multiwalled carbon nanotubes at graphitization temperatures, *Carbon*. 39 (2001) 1681–1687. [https://doi.org/10.1016/S0008-6223\(00\)00301-8](https://doi.org/10.1016/S0008-6223(00)00301-8).
- [146] L. Lavagna, S. Musso, M. Pavese, A facile method to oxidize carbon nanotubes in controlled flow of oxygen at 350 °C, *Mater. Lett.* 283 (2021) 128816. <https://doi.org/10.1016/j.matlet.2020.128816>.
- [147] Z.N. Utegulov, D.B. Mast, P. He, D. Shi, R.F. Gilland, Functionalization of single-walled carbon nanotubes using isotropic plasma treatment: Resonant Raman spectroscopy study, *J. Appl. Phys.* 97 (2005) 104324. <https://doi.org/10.1063/1.1913801>.
- [148] B.M. Penetrante, S.E. Schultheis, Non-thermal plasma techniques for pollution control: part b: electron beam and electrical discharge processing, Springer Science & Business Media, 2013.
- [149] C. Saka, Overview on the Surface Functionalization Mechanism and Determination of Surface Functional Groups of Plasma Treated Carbon Nanotubes, *Crit. Rev. Anal. Chem.* 48 (2018) 1–14. <https://doi.org/10.1080/10408347.2017.1356699>.
- [150] I.-H. Chen, C.-C. Wang, C.-Y. Chen, Preparation of Carbon Nanotube (CNT) Composites by Polymer Functionalized CNT under Plasma Treatment, *Plasma Process. Polym.* 7 (2010) 59–63. <https://doi.org/10.1002/ppap.200900067>.
- [151] T. Abuzairi, M. Okada, Y. Mochizuki, N.R. Poespawati, R.W. Purnamaningsih, M. Nagatsu, Maskless functionalization of a carbon nanotube dot array biosensor using an ultrafine atmospheric pressure plasma jet, *Carbon*. 89 (2015) 208–216. <https://doi.org/10.1016/j.carbon.2015.03.015>.
- [152] T. Abuzairi, M. Okada, R.W. Purnamaningsih, N.R. Poespawati, F. Iwata, M. Nagatsu, Maskless localized patterning of biomolecules on carbon nanotube microarray functionalized by ultrafine atmospheric pressure plasma jet using biotin-avidin system, *Appl. Phys. Lett.* 109 (2016) 023701. <https://doi.org/10.1063/1.4958988>.
- [153] C. Boaretti, M. Roso, R. Bonora, M. Modesti, A. Lorenzetti, Investigation of Plasma-Assisted Functionalization of Graphitic Materials for Epoxy Composites, *Nanomaterials*. 10 (2019) 78. <https://doi.org/10.3390/nano10010078>.
- [154] M. Garzia Trulli, E. Sardella, F. Palumbo, G. Palazzo, L.C. Giannossa, A. Mangone, R. Comparelli, S. Musso, P. Favia, Towards highly stable aqueous dispersions of multi-walled carbon nanotubes: the effect of oxygen plasma functionalization, *J. Colloid Interface Sci.* 491 (2017) 255–264. <https://doi.org/10.1016/j.jcis.2016.12.039>.

- [155] M.J. O'Connell, P. Boul, L.M. Ericson, C. Huffman, Y. Wang, E. Haroz, C. Kuper, J. Tour, K.D. Ausman, R.E. Smalley, Reversible water-solubilization of single-walled carbon nanotubes by polymer wrapping, *Chem. Phys. Lett.* 342 (2001) 265–271. [https://doi.org/10.1016/S0009-2614\(01\)00490-0](https://doi.org/10.1016/S0009-2614(01)00490-0).
- [156] D.E. Hill, Y. Lin, A.M. Rao, L.F. Allard, Y.-P. Sun, Functionalization of Carbon Nanotubes with Polystyrene, *Macromolecules*. 35 (2002) 9466–9471. <https://doi.org/10.1021/ma020855r>.
- [157] L. Lavagna, S. Marchisio, A. Merlo, R. Nisticò, M. Pavese, Polyvinyl butyral-based composites with carbon nanotubes: Efficient dispersion as a key to high mechanical properties, *Polym. Compos.* 41 (2020) 3627–3637. <https://doi.org/10.1002/pc.25661>.
- [158] E.A. Whitsitt, A.R. Barron, Silica Coated Single Walled Carbon Nanotubes, *Nano Lett.* 3 (2003) 775–778. <https://doi.org/10.1021/nl034186m>.
- [159] T.-H. Kim, C. Doe, S.R. Kline, S.-M. Choi, Water-Redispersible Isolated Single-Walled Carbon Nanotubes Fabricated by In Situ Polymerization of Micelles, *Adv. Mater.* 19 (2007) 929–933. <https://doi.org/10.1002/adma.200601991>.
- [160] X. Gong, J. Liu, S. Baskaran, R.D. Voise, J.S. Young, Surfactant-Assisted Processing of Carbon Nanotube/Polymer Composites, *Chem. Mater.* 12 (2000) 1049–1052. <https://doi.org/10.1021/cm9906396>.
- [161] S. Cui, R. Canet, A. Derre, M. Couzi, P. Delhaes, Characterization of multiwall carbon nanotubes and influence of surfactant in the nanocomposite processing, *Carbon*. 41 (2003) 797–809. [https://doi.org/10.1016/S0008-6223\(02\)00405-0](https://doi.org/10.1016/S0008-6223(02)00405-0).
- [162] L. Vaisman, G. Marom, H.D. Wagner, Dispersions of Surface-Modified Carbon Nanotubes in Water-Soluble and Water-Insoluble Polymers, *Adv. Funct. Mater.* 16 (2006) 357–363. <https://doi.org/10.1002/adfm.200500142>.
- [163] Y. Geng, M.Y. Liu, J. Li, X.M. Shi, J.K. Kim, Effects of surfactant treatment on mechanical and electrical properties of CNT/epoxy nanocomposites, *Compos. Part Appl. Sci. Manuf.* 39 (2008) 1876–1883. <https://doi.org/10.1016/j.compositesa.2008.09.009>.
- [164] M.F. Islam, E. Rojas, D.M. Bergey, A.T. Johnson, A.G. Yodh, High Weight Fraction Surfactant Solubilization of Single-Wall Carbon Nanotubes in Water, *Nano Lett.* 3 (2003) 269–273. <https://doi.org/10.1021/nl025924u>.
- [165] J. Yu, N. Grossiord, C.E. Koning, J. Loos, Controlling the dispersion of multi-wall carbon nanotubes in aqueous surfactant solution, *Carbon*. 45 (2007) 618–623. <https://doi.org/10.1016/j.carbon.2006.10.010>.
- [166] L. Vaisman, H.D. Wagner, G. Marom, The role of surfactants in dispersion of carbon nanotubes, *Honor Profr. Nissim Gartis 60th Birthd.* 128–130 (2006) 37–46. <https://doi.org/10.1016/j.cis.2006.11.007>.
- [167] N. Grossiord, J. Loos, O. Regev, C.E. Koning, Toolbox for Dispersing Carbon Nanotubes into Polymers To Get Conductive Nanocomposites, *Chem. Mater.* 18 (2006) 1089–1099. <https://doi.org/10.1021/cm051881h>.
- [168] M.S. Strano, V.C. Moore, M.K. Miller, M.J. Allen, E.H. Haroz, C. Kittrell, R.H. Hauge, R.E. Smalley, The Role of Surfactant Adsorption during Ultrasonication in the Dispersion of Single-Walled Carbon Nanotubes, *J. Nanosci. Nanotechnol.* 3 (2003) 81–86. <https://doi.org/10.1166/jnn.2003.194>.
- [169] V.C. Moore, M.S. Strano, E.H. Haroz, R.H. Hauge, R.E. Smalley, J. Schmidt, Y. Talmon, Individually Suspended Single-Walled Carbon Nanotubes in Various Surfactants, *Nano Lett.* 3 (2003) 1379–1382. <https://doi.org/10.1021/nl034524j>.
- [170] X. Zhang, K. Song, J. Liu, Z. Zhang, C. Wang, H. Li, Sorption of triclosan by carbon nanotubes in dispersion: The importance of dispersing properties using different surfactants, *Colloids Surf. Physicochem. Eng. Asp.* 562 (2019) 280–288. <https://doi.org/10.1016/j.colsurfa.2018.11.037>.
- [171] H. Wang, Dispersing carbon nanotubes using surfactants, *Curr. Opin. Colloid Interface Sci.* 14 (2009) 364–371. <https://doi.org/10.1016/j.cocis.2009.06.004>.
- [172] S. Siljander, P. Keinänen, A. Rätty, K. Ramakrishnan, S. Tuukkanen, V. Kunnari, A. Harlin, J. Vuorinen, M. Kanerva, Effect of Surfactant Type and Sonication Energy on the Electrical Conductivity Properties of Nanocellulose-CNT Nanocomposite Films, *Int. J. Mol. Sci.* 19 (2018) 1819. <https://doi.org/10.3390/ijms19061819>.

- [173] B. Han, X. Yu, E. Kwon, J. Ou, Effects of CNT concentration level and water/cement ratio on the piezoresistivity of CNT/cement composites, *J. Compos. Mater.* 46 (2012) 19–25. <https://doi.org/10.1177/0021998311401114>.
- [174] N. Melanitis, P.L. Tetlow, C. Galiotis, Characterization of PAN-based carbon fibres with laser Raman spectroscopy, *J. Mater. Sci.* 31 (1996) 851–860. <https://doi.org/10.1007/BF00352882>.
- [175] S. Musso, G. Fanchini, A. Tagliaferro, Growth of vertically aligned carbon nanotubes by CVD by evaporation of carbon precursors, *Proc. Diam. 2004 15th Eur. Conf. Diam. Diam.- Mater. Carbon Nanotub. Nitrides Silicon Carbide.* 14 (2005) 784–789. <https://doi.org/10.1016/j.diamond.2004.12.030>.
- [176] F. Tuinstra, J.L. Koenig, Raman Spectrum of Graphite, *J. Chem. Phys.* 53 (1970) 1126–1130. <https://doi.org/10.1063/1.1674108>.
- [177] R. Rastogi, R. Kaushal, S.K. Tripathi, A.L. Sharma, I. Kaur, L.M. Bharadwaj, Comparative study of carbon nanotube dispersion using surfactants, *J. Colloid Interface Sci.* 328 (2008) 421–428. <https://doi.org/10.1016/j.jcis.2008.09.015>.
- [178] F. Catania, E. Marras, M. Giorcelli, P. Jagdale, L. Lavagna, A. Tagliaferro, M. Bartoli, A Review on Recent Advancements of Graphene and Graphene-Related Materials in Biological Applications, *Appl. Sci.* 11 (2021) 614. <https://doi.org/10.3390/app11020614>.
- [179] J. Njuguna, O.A. Vanli, R. Liang, A Review of Spectral Methods for Dispersion Characterization of Carbon Nanotubes in Aqueous Suspensions, *J. Spectrosc.* 2015 (2015) 463156. <https://doi.org/10.1155/2015/463156>.
- [180] W.Z. Zhu, D.E. Miser, W.G. Chan, M.R. Hajaligol, Characterization of multiwalled carbon nanotubes prepared by carbon arc cathode deposit, *Mater. Chem. Phys.* 82 (2003) 638–647. [https://doi.org/10.1016/S0254-0584\(03\)00341-9](https://doi.org/10.1016/S0254-0584(03)00341-9).
- [181] Kelvii Wei Guo, TEM Morphology of Carbon Nanotubes (CNTs) and its Effect on the Life of Micropunch, in: Hon-Yuen Tam ED1 - Khan Maaz (Ed.), *Transm. Electron Microsc.*, IntechOpen, Rijeka, 2015: p. Ch. 7. <https://doi.org/10.5772/60713>.
- [182] M. Smith, L. Scudiero, J. Espinal, J.-S. McEwen, M. Garcia-Perez, Improving the deconvolution and interpretation of XPS spectra from chars by ab initio calculations, *Carbon.* 110 (2016) 155–171. <https://doi.org/10.1016/j.carbon.2016.09.012>.
- [183] T.I.T. Okpalugo, P. Papakonstantinou, H. Murphy, J. McLaughlin, N.M.D. Brown, High resolution XPS characterization of chemical functionalised MWCNTs and SWCNTs, *Carbon.* 43 (2005) 153–161. <https://doi.org/10.1016/j.carbon.2004.08.033>.
- [184] P.M.B. Piccoli, T.F. Koetzle, A.J. Schultz, SINGLE CRYSTAL NEUTRON DIFFRACTION FOR THE INORGANIC CHEMIST – A PRACTICAL GUIDE, *Comments Inorg. Chem.* 28 (2007) 3–38. <https://doi.org/10.1080/02603590701394741>.
- [185] A. Burian, J.C. Dore, H.E. Fischer, J. Sloan, Structural studies of multiwall carbon nanotubes by neutron diffraction, *Phys. Rev. B.* 59 (1999) 1665–1668. <https://doi.org/10.1103/PhysRevB.59.1665>.
- [186] T. Belin, F. Epron, Characterization methods of carbon nanotubes: a review, *Mater. Sci. Eng. B.* 119 (2005) 105–118. <https://doi.org/10.1016/j.mseb.2005.02.046>.
- [187] T. Saito, K. Matsushige, K. Tanaka, Chemical treatment and modification of multi-walled carbon nanotubes, *Proc. Tsukuba Symp. Carbon Nanotube Commem. 10th Anniv. Its Discov.* 323 (2002) 280–283. [https://doi.org/10.1016/S0921-4526\(02\)00999-7](https://doi.org/10.1016/S0921-4526(02)00999-7).
- [188] A.W. Coats, J.P. Redfern, Thermogravimetric analysis. A review, *Analyst.* 88 (1963) 906–924. <https://doi.org/10.1039/AN9638800906>.
- [189] G.R. Kasaliwal, S. Pegel, A. Gödel, P. Pötschke, G. Heinrich, Analysis of agglomerate dispersion mechanisms of multiwalled carbon nanotubes during melt mixing in polycarbonate, *Polymer.* 51 (2010) 2708–2720. <https://doi.org/10.1016/j.polymer.2010.02.048>.
- [190] S. Musso, J.-M. Tulliani, G. Ferro, A. Tagliaferro, Influence of carbon nanotubes structure on the mechanical behavior of cement composites, *Compos. Sci. Technol.* 69 (2009) 1985–1990. <https://doi.org/10.1016/j.compscitech.2009.05.002>.
- [191] L. Lavagna, D. Massella, E. Priola, M. Pavese, Relationship between oxygen content of graphene and mechanical properties of cement-based composites, *Cem. Concr. Compos.* 115 (2021) 103851. <https://doi.org/10.1016/j.cemconcomp.2020.103851>.

- [192] M. Giorcelli, M. Bartoli, Development of Coffee Biochar Filler for the Production of Electrical Conductive Reinforced Plastic, *Polymers*. 11 (2019) 1916. <https://doi.org/10.3390/polym11121916>.
- [193] A.G. Nasibulin, T. Koltsova, L.I. Nasibulina, I.V. Anoshkin, A. Semench, O.V. Tolochko, E.I. Kauppinen, A novel approach to composite preparation by direct synthesis of carbon nanomaterial on matrix or filler particles, *Acta Mater.* 61 (2013) 1862–1871. <https://doi.org/10.1016/j.actamat.2012.12.007>.
- [194] F. Ghaharpour, A. Bahari, M. Abbasi, A.A. Ashkarran, Parametric investigation of CNT deposition on cement by CVD process, *Constr. Build. Mater.* 113 (2016) 523–535. <https://doi.org/10.1016/j.conbuildmat.2016.03.080>.
- [195] L. Lavagna, G. Meligrana, C. Gerbaldi, A. Tagliaferro, M. Bartoli, Graphene and Lithium-Based Battery Electrodes: A Review of Recent Literature, *Energies*. 13 (2020) 4867. <https://doi.org/10.3390/en13184867>.
- [196] W. Ma, D. Tian, Direct electron transfer and electrocatalysis of hemoglobin in ZnO coated multiwalled carbon nanotubes and Nafion composite matrix, *Bioelectrochemistry*. 78 (2010) 106–112. <https://doi.org/10.1016/j.bioelechem.2009.08.002>.
- [197] J. Cho, A.R. Boccaccini, M.S.P. Shaffer, Ceramic matrix composites containing carbon nanotubes, *J. Mater. Sci.* 44 (2009) 1934–1951. <https://doi.org/10.1007/s10853-009-3262-9>.
- [198] L. Cao, T.K. Sinha, X. Zhang, X. Zhai, C. Wang, C. Zong, J.K. Kim, Graphene/carbon nanotubes-supported Ziegler-Natta catalysts for in situ synthesis of mechanically strong, thermally and electrically conductive trans-polyisoprene nanocomposite, *J. Polym. Res.* 26 (2019) 36. <https://doi.org/10.1007/s10965-018-1688-y>.