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Review Article

Literature Review Paper on Mechanical Properties of Types of Carbon Nanotubes

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Abstract

Among the numerous potential uses of carbon nanotubes (CNT), its utilization to fortify polymers was given careful consideration. This reason can be because of the remarkable firmness, magnificent quality, and the low thickness of CNT. This has given various chances to the innovation of new material frameworks for applications requiring high quality and high modulus. Exact control over preparing factors, including safeguarding flawless CNT structure, uniform scattering of CNT inside the polymer grid, compelling filler– lattice interfacial communications, and arrangement/introduction of polymer chains/CNT, adds to the composite strands' unrivalled properties. Consequently, manufacture techniques assume an imperative part in deciding the composite filaments' microstructure and extreme mechanical conduct. The present best in the class of polymer/CNT elite composite filaments, particularly concerning processing– structure– execution, were looked into in this commitment. Future requirements for material by configuration approaches for handling these nano-composite frameworks were likewise examined.

Keywords:
carbon nanotubes; polymer; mechanical properties; readiness; combination; scattering; interphase; arrangement; applications

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Introduction

The materials are ruled the market as far as their flexibility for item's applications when the introduction of polymer materials science in the 1930s. These materials had been used as movies, strands, sheets, and coatings. Today, a large portion of the manufactured polymer strands being used traverse applications, for example, garments, rugs, ropes, and support materials. A portion of these filaments incorporate polyamides, for example, nylon, polyesters (e.g., polyethylene terephthalate (PET) and polybutylene terephthalate (PBT)), polyolefins (e.g., polypropylene (PP) or polyethylene (PE)), vinyl polymers (e.g., poly(vinyl liquor) (PVA) and poly(vinyl chloride) (PVC)), elastomers (e.g., polyurethane (PU) and spandex), and acrylic strands (e.g., polyacrylonitrile (PAN)) [1], [2], [3], [4], [5], [6], [7], [8], [9], [10], [11], [12], [13], [14], [15], [16], [17], [18], [19], [20], [21], [22], [23], [24], [25], [26], [27], [28], [29], [30], [31], [32], [33], [34], [35], [36], [37], [38], [39], [40], [41], [42], [43], [44], [45]. What's more, superior polymer-based strands with high solidness as well as persistence incorporate Dyneema® and Spectra® (i.e., ultra-high sub-atomic weight polyethylene (UHMWPE)- based filaments), Twaron® and Kevlar®, and Zylon® strands (i.e., sweet-smelling based polymers, for example, poly(p-phenyleneterephthalamide) (PPTA) and poly(p-phenylenebenzobisoxazole) (PBO)) [2], [3], [4], [5], [6], [7], [8]. Likewise included was PANNING, which was the overwhelming forerunner fibre for the carbon fibre industry [11], [12], [13], [14], [15], [16], [17], [18], [19], [20], [21], [22], [23], [24], [25], [26], [27], [28], [29], [30], [31], [32], [33], [34], [35], [36], [37], [38], [39], [40], [41].

General Fabrication Procedures for Polymer/CNT Fibres

This audit paper centred on the top of the line of polymer/CNT composite materials to investigate their processing-structure-property connections. The four-noteworthy fibre-turning techniques (Figure 1) utilized for polymer/CNT composites from both the arrangement and soften incorporate dry turning [51], [52], wet turning [53], dry fly wet turning (e.g., gel-turning [54]), and electro-turning [55], [56]. An old strong state turning approach has been utilized for manufacturing 100% CNT strands from the two woods and aerogels [57], [58], [59], [60]. Despite the handling system, to grow excellent strands numerous parameters should be all around controlled. As a rule, all turning systems include (I) fibre arrangement; (ii) coagulation/gelation/cementing; and (iii) drawing/arrangement. For these procedures, the even scattering of the CNT inside the polymer arrangement or dissolve was critical. Nevertheless, as far as accomplishing superb hub mechanical properties, arrangement and introduction of the polymer chains and the CNT in the composite was vital. Fibre arrangement was expert in post-preparing, for example, drawing/toughening and was critical to expanding crystallinity, rigidity, and solidness [61], [62], [63], [64], [65], [66], [67], [68], [69], [70], [71], [72], [73], [74], [75], [76], [77], [78], [79].

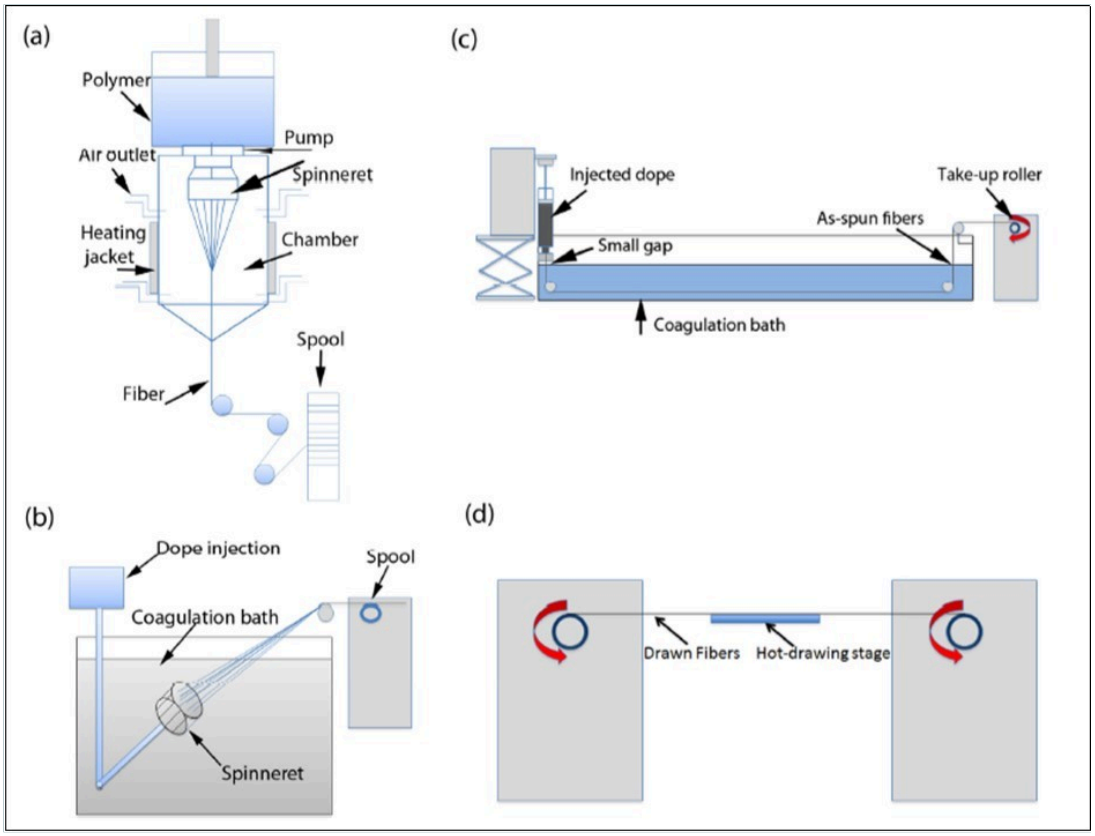


Figure 1. Schematics for the different fibre preparing techniques (a) dry-turning; (b) wet-turning; (c) dry-stream wet or gel turning; and (d) post-handling by hot-arrange drawing [10].

Smaller scale Structural Development in Polymer/CNT Fibres

The general picture of mechanical execution for polymer/CNT filaments delivered at the exploration level demonstrates a wide scope of properties (Figure 2). These strands were delivered utilizing a few manufacture strategies. As said, the revelation of CNT introduced a lot of research endeavours concentrated on using these nano-materials to make polymer composite strands to catch these outstanding properties (i.e., 1 TPa in ductile modulus and 10 to 150 GPa [21], [22], [23], [24] of every rigidity).

The natural properties of CNT accept that the structure was very much protected (i.e., substantial angle proportion and without surrenders). Going further, the initial move toward viable support of polymers utilizing nano-fillers was to accomplish a uniform scattering of the fillers inside the facilitating network, and this was additionally identified with the as-blended nano-carbon structure. Furthermore, successful interfacial communication and stress exchange amongst CNT and polymer was fundamental for enhanced mechanical properties of the fibre composite. At last, like polymer particles, the great-inborn mechanical properties of CNT can be completely misused just if a perfect uniaxial introduction was accomplished. In this way, amid the manufacture of polymer/CNT filaments, four key territories should be tended to and comprehended with a specific end goal to effectively control the small-scale auxiliary improvement in these composites. These are: (I) CNT perfect structure; (ii) CNT scattering; (iii) polymer-CNT interfacial communication; and (iv) introduction of the filler and grid particles (Figure 3). This survey will feature some key papers that had concentrated on these zones to tailor the composite structure and propel the mechanical execution of the polymer nano-composite [23], [24], [25], [26], [27], [28], [29], [30], [31].

A further examination of the distributed writing additionally demonstrates a fascinating pattern, whereby the percent expansion in mechanical properties for polymer composite filaments was identified with the inalienable polymer structure (Figure 4) [16], [17], [18], [19], [20].

Compound functionalization increments the between tube contacts (i.e., helpful for working up a conductive system) and gives more potential outcomes to bond the nanotubes to a framework because

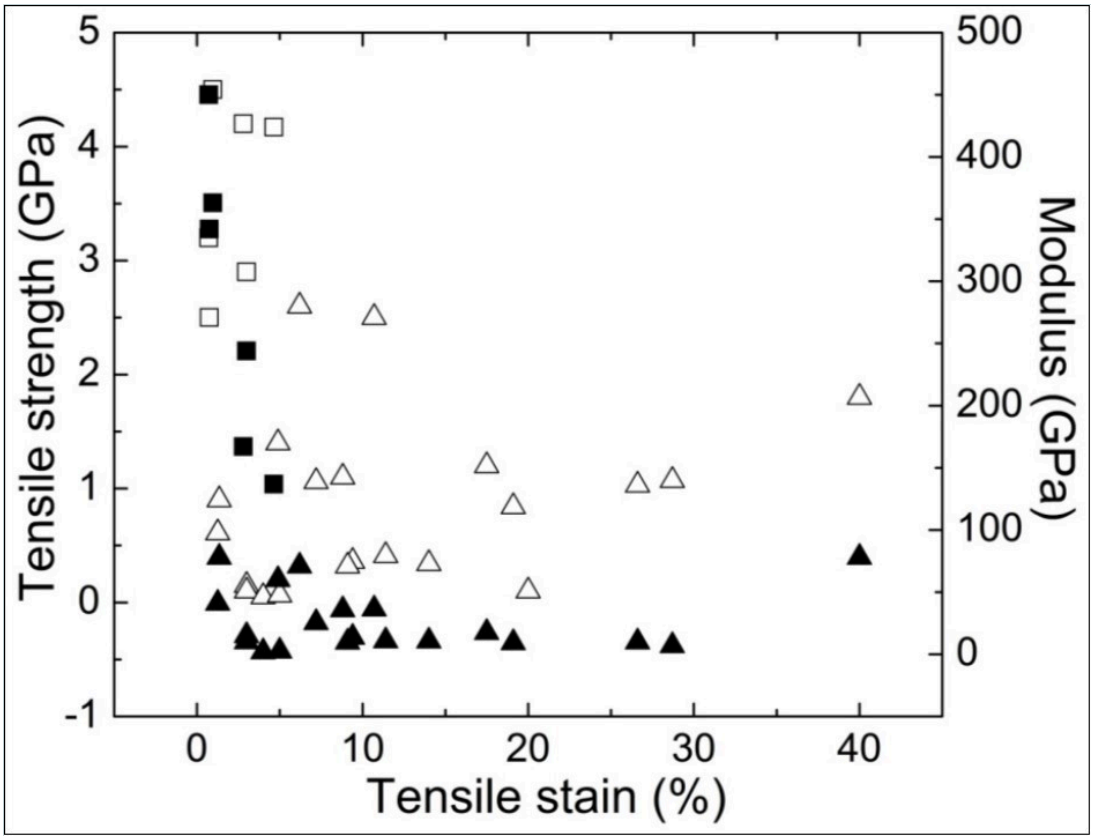


Figure 2. Rundown of Young's modulus, elasticity, and strain-to-disappointment properties for different polymer/CNT filaments delivered at the examination scale [35], [36], [38], [39], [40], [41], [42], [43], [44], [45], [46], [47], [112], [113], [114], [115], [116] (Note: □/■ images for rigidity/modulus properties for superior strands, and Δ/▲ images for rigidity/modulus properties of material review filaments).

of responsive concoction gatherings. Then again, covalent surface medications can decimate tube structure, bringing about shortening of nanotubes [55], [56], making of deformities in the graphitic structure of CNT dividers [33], [56], [57], and at times, unfastening of the tube structure. Therefore, substance functionalization will diminish the mechanical properties of CNT [58]. Non-covalent scattering techniques had additionally been created to peel SWNT groups into singular tubes in various

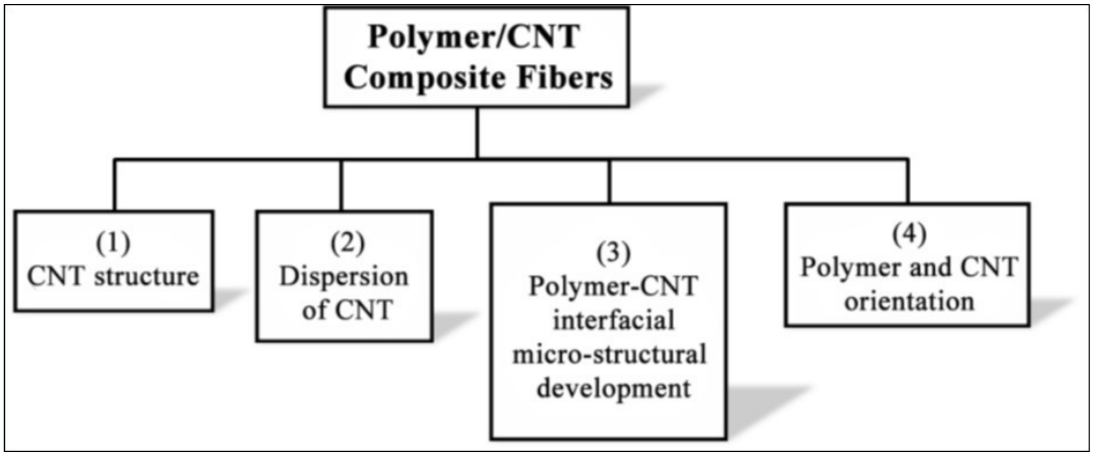


Figure 3. Four central points, which are influencing the small-scale basic advancement in polymer/CNT composite fibre amid preparing [35], [36], [38], [39], [40], [41], [42], [43], [44], [45], [46], [47], [112], [113], [114], [115], [116].

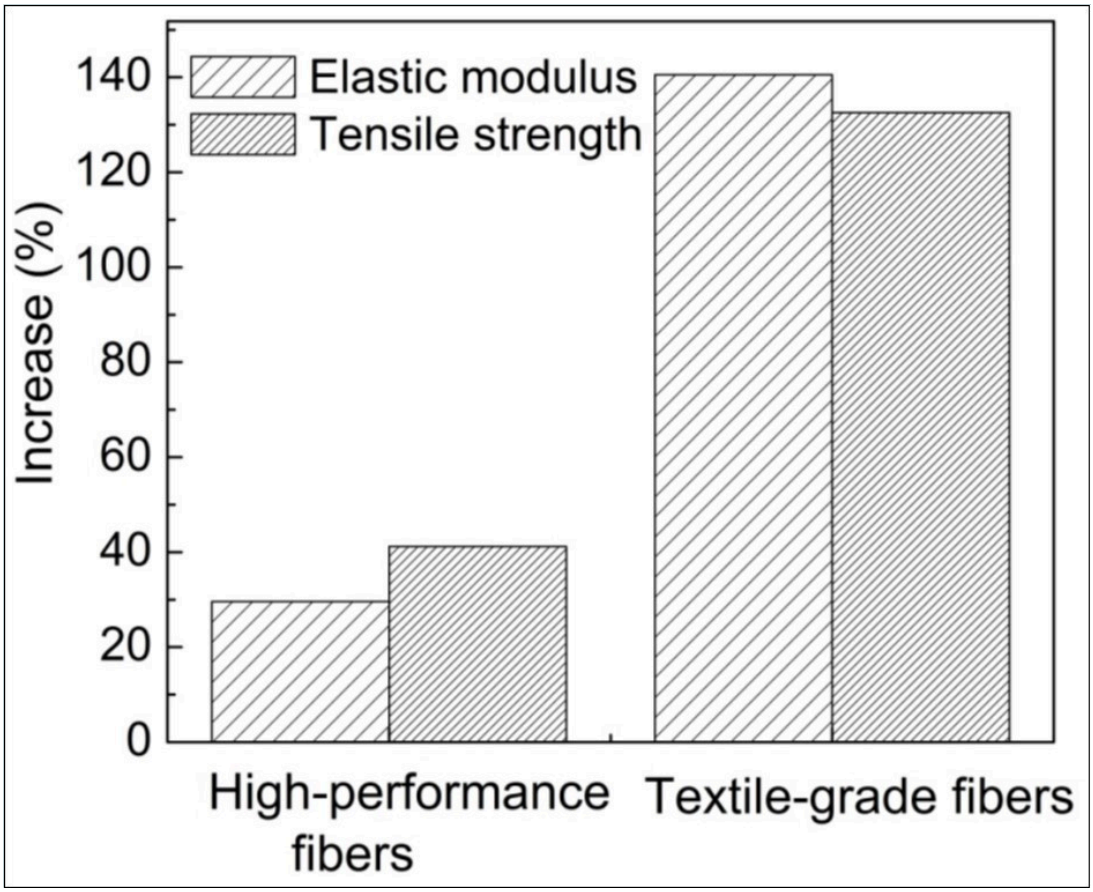


Figure 4. Normal percent expansion correlation between control strands (no fillers) and composite filaments for both the Young's modulus and rigidity properties [35], [36], [38], [39], [40], [41], [42], [43], [44], [45], [46], [47], [112], [113], [114], [115], [116].

solvents utilizing different anionic, cationic, non-ionic surfactants [34], [59] or polymers [35], [60]. The SWNT modulus, quality, and interfacial shear quality were taken to be 1 TPA, 50 GPa, and 100 MPa (i.e., in view of computational forecasts) [20], [64], [65], separately. To exhibit the significance of the length commitment in the composite, which was plotted by utilizing polymer framework modulus esteems running from 1 to 100 GPa, and quality qualities extending from 0.01 to 5 GPa. These qualities compare to the commonplace properties revealed for polymers utilized as a part of CNT composite preparing [35], [36], [38], [39], [40], [41], [42], [43], [44], [45], [46], [47], [112], [113], [114], [115], [116]. The modulus and quality increment as for viewpoint proportion can be seen that both solidness and quality of the filaments scale with viewpoint proportion. A comparable pattern has additionally been accounted for composite movies [26], [27], [28], [29], [30], [31], [32], [33], [34]. Plainly, the scattering of the CNT as far as shedding, circulation, and length safeguarding were exceptionally critical angles influencing the advancement of the composite microstructure. Each factor was subject to the next and finding the correct adjust remains a test. Albeit a few strategies for scattering had been talked about, it was critical to perceive that without great polymer nanotube communication; even very much scattered CNT may not give viable support of the framework. To enhance polymer-CNT collaborations, interfacial advancement was essential. The accompanying Section 3.2 examines a portion of the systems for the improvement of interfacial structures in the polymer composite strands. To do this will require further major comprehension of the nano-composite framework as far as morphology development amid handling. The interfacial connection happens through a few instruments: (I) mechanical coupling, smaller scale mechanical interlocking and polymer chain-CNT ensnarement; (ii) physical collaboration, including van der Waals powers, electrostatic powers, or epitaxial precious stone development; and (iii) substance associations. As said in the past segment, these substance connections incorporate covalent holding and physical holding, for example, surfactant-helped scattering of CNT [33], plasma polymerization [67], and polymer wrapping [68], [69]. A few examinations had concentrated on understanding

the quality of the interface for polymer/CNT materials. For PVA/CNT composites, it was discovered that the shearing brought about crack of the grid before the breakage of the interphase polymer [70]. The shear pressure was resolved to associate with 40 MPa, which was in sensible concurrence with anticipated estimations of ~50 MPa [70]. Other computational works had moreover been done to foresee the interfacial shear pressure (IFSS). Polymer frameworks, for example, polystyrene (PS) [71], epoxy [72], poly (m-phenylenevinylene-co-2, 5-dioctyloxy-p-phenylenevinylene) (PmPV), what was more, poly (phenyl acetylene) (PPA) [73] had been figured utilizing atomic progression, where the computed IFSS was subject to both the polymer and CNT. In such cases, the IFSS esteems ran from 18 to 186 MPa. Aside from the figurings and re-enactments, coordinate estimations had likewise been accounted for. The systems and gadgets for these estimations incorporate checking electron microscopy (SEM) [20], transmission electron microscopy (TEM) [74], nuclear power microscopy (AFM) [64], [75], and filtering test microscopy (SPM) [76]. These announced qualities extend from 0.02 to 500 MPa [39], [16], [26], [65], [74], [75], [76]. The bigger IFSS esteems were reliable with composites where covalent holding was available at the interphase (i.e., functionalized CNT). Estimations of 0.5 GPa assessed by Wagner et al. [74], and 0.35 GPa estimated by Cooper et al. [76] were estimated using the AFM and were ascribed to covalent holding amongst CNT and polymer. To date, the lion's share of interphase estimations and expectations had concentrated on either perfect CNT or functionalized CNT installed in shapeless polymer dissolves. Less was thought about the interfacial mechanical properties of crystalline polymer at the CNT interphase, particularly in situations where the polymer can frame requested stages along the CNT length. A few on-going papers had featured the significance of crystalline interphase arrangement in these composites [39], [42], [43], [77], [78], [79]. It has been watched that CNT can nucleate and format the development of requested polymer gems in a few polymer frameworks including PE [80], [81], [82], [83], [84], [85], nylon 6,6 [82], PVA [86], PAN [87], poly(butylene terephthalate) (PBT) [88], [89], [90], isotactic polypropylene (iPP) [91], poly(L-lactide) (PLLA) [91], poly(e-caprolactone) (PCL) [92], and polyethylene-b-poly(ethylene oxide) (PE-b-PEO) square copolymer [93]. One of the overwhelming support components in polymer/CNT composites has been recommended to be the nearness of requested polymer interfacial covering structure close CNT [94]. This arranged structure can frame because of the capacity of CNT to communicate particularly with the polymer grid. Requested or crystalline polymer structure in polymer nano-composites was mechanically more grounded than shapeless structure due to the nearness of fewer imperfections or less scattered areas. Along these lines, it was critical to ponder CNT-actuated polymer crystallization to control these systems amid the arrangement of the interphase in the polymer/CNT composites. On an atomic level, a diminished interpenetration/snare of chains close to a strong interface causes chain arrangement, the configuration-change energies, and rehash unit-surface association energies to change [95]. Likewise, changes in response energy and interfacial versatility (i.e., due to crosslink thickness) can likewise influence the framework [95]. Glass progress, polymer dispersion, nanotube dissemination, crystalline structure, crystallization energy, and properties can likewise be adjusted [95]. This marvel was not seen with other usually utilized small-scale fillers [95]. Extra work has demonstrated that the interphase polymer morphology was totally unique in relation to the mass polymer in the composite, and this means high modulus and elasticity esteems (i.e., modulus about 5 and 400 GPa and quality >1 GPa). Examination of these interphase areas by microscopy demonstrates that they show crystalline flawlessness [42], [43], [44], [78]. As already said, a few works had likewise demonstrated the capacity of the nanotube to nucleate polymer gem development at the interphase [82], [96], [97], [98], [99], [100], [101]. In addition, layout gem development and introduction in polymers [42], [43], [77], [78], [81], [85], [97]. This templating impact of CNT in polymer composites has been demonstrated to have a successful commitment toward the pressure exchange component of load between the polymer grid and filler [42], [49], [50], [51], [52], [53], [54], [55], [56], [57], [58], [59], [60], [61], [62], [63], [64], [65], [66], [67], [68], [69], [70], [71], [72], [73], [74], [75], [76], [77], [78], [79]. In such situations where templated interphase structure was observed to be available at the interphase, the mechanical properties for the composite were essentially expanded. It was additionally intriguing that the general crystallinity esteem for the composite when contrasted with the control strands was generally the same. This suggests while a bit of the grid polymer frames a much-arranged interphase structure, the mass framework remains semi-crystalline and moderately disarranged. It was additionally worth specifying that the expansion in mechanical properties does not take after governing of-blend expectations. This

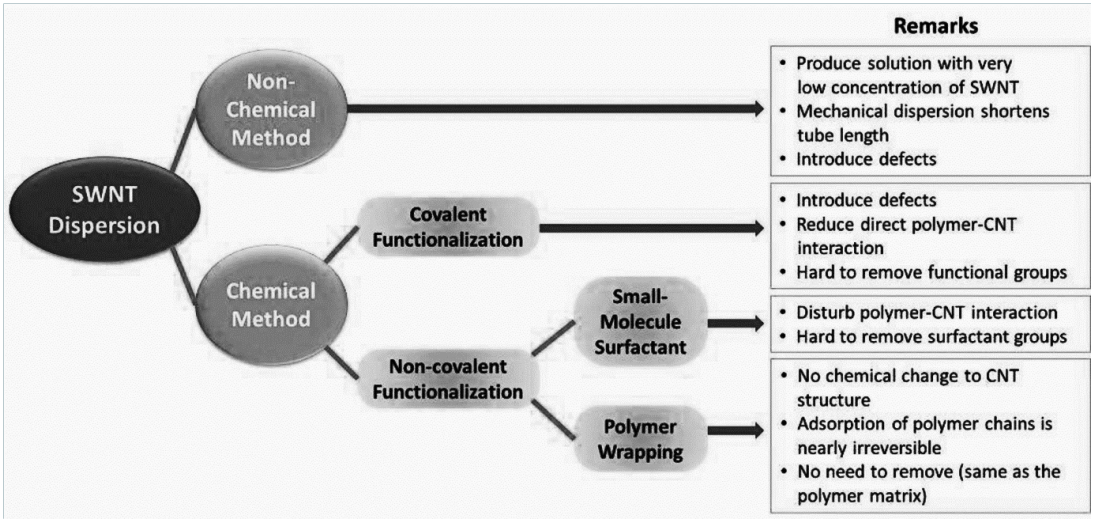


Figure 5. Utilised strategies for SWNT scattering towards creation of polymer/CNT nano-composites [35], [36], [38], [39], [40], [41], [42], [43], [44], [45], [46], [47], [112], [113], [114], [115], [116].

was because of the commitment from the interphase polymer, which was regularly unaccounted for. A few late works had endeavoured to incorporate this commitment for better comprehension of the composite small-scale basic commitment to the mass properties [40], [61]. It was additionally imperative to take note of that in some CNT-polymer frameworks where CNT templating was discovered, the crystallinity was regularly much higher in the composite versus the control framework. In such cases, the impact of templating alone was hard to evaluate. Here, the attention was on two frameworks, which show comparative crystallinity keeping in mind the end goal to comprehend the part of the format situated polymer interphase commitment. It has likewise been perceived that in situations where the interphase areas were not format or arranged (i.e., demonstrating chain issue), the mechanical upgrade was not that critical [98]. Interfacial push exchange was a basic part/parameter controlling the execution of the composite. Finish pressure/stack exchange from the polymer to the nano-filler was achievable if there was solid grip. In view of these high-determination transmission electron microscopy (HR-TEM) thinks about, better chain pressing was likewise appeared to exist at the interphase [42], [43], [81].

As of late shear crystallization thinks about in half-breed polymer/SWNT scattering as of late shear crystallization considers in cross breed polymer/SWNT scattering were utilized to initiate arranged polymer crystallization within the sight of the SWNT. These investigations were particularly engaged on building up a method for delivering requested interphase structure on the CNT. Figure 5, Figure 6 demonstrates an HR-TEM picture for a PAN-SWNT interphase, where the polymer broadened chain

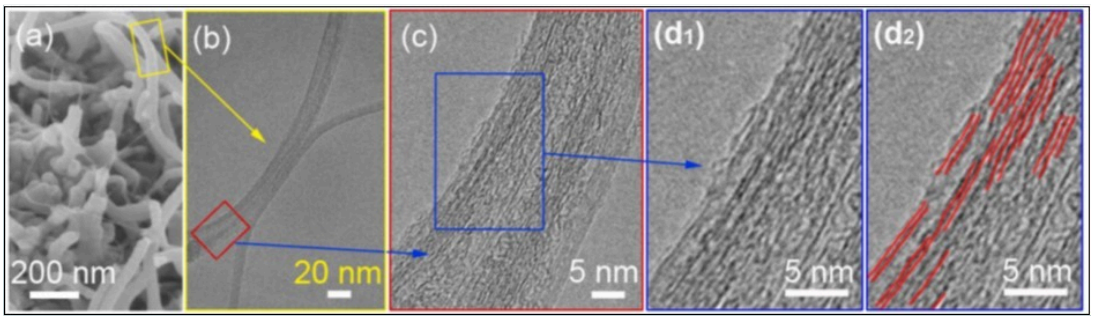


Figure 6. (a) Scanning electron micrograph (SEM) of PAN tubular covering on SWNT. High-determination transmission electron micrograph (HR-TEM) of tubular covered Dish/SWNT tests; (b) at the beginning of electron bar presentation; (c and d1) demonstrate a territory of the PAN/SWNT test where the PAN grid of ~0.52 nm is watched; and (d2) a schematic featuring the PAN grid perceptions in (d1) [87].

morphology has been templated by the nanotube [87]. This principal crystallization thinks about give great understanding at the morphological abilities of the polymer affected by this system. As far as handling polymer/CNT composite materials, these crystallization procedures may even be fused into creation methodology [109], [110], [111], [112], [113].

These nucleation, crystallization, and introduction impacts were particularly seen in composites with low nano-carbon stacking (<one wt %) and significantly affect the general structure and properties of the composite material [42], [43]. Arrangement of CNT or CNT ropes was another critical factor in deciding the mechanical properties of composites containing them. As indicated by the continuum mechanics computations, the moduli of both SWNT filler and polymer chains along the hub course drop suddenly for just slight mis-introduction regarding the fibre hub. For SWNT materials, this impact was less articulated as the SWNT package measurement diminishes [105], [106], [107], [108], [109].

What was instantly clear was that in the polymer/CNT composite fibre, the full arrangement of the polymer chain and the CNT was foremost. This was not a simple undertaking. To date, just a bunch of polymer-based elite filaments exists (i.e., Kevlar®, Spectra®, Zylon®), and this was because of the high chain arrangement in the small-scale structure either managed by the innate polymer conformational structure (i.e., pole like particles—Kevlar® and Zylon®) or uncommon preparing of low focus polymer answers for diminish chain trap (i.e., gel turning of polyethylene—Spectra®). Nevertheless, in later work, the similitudes amongst polymers and CNT, CNT templating impacts, CNT fluid crystalline nature, and the capacity of nano-carbons materials to grease up polymers amid arrangement had been perceived. These components all had huge ramifications toward significantly progressing polymer chain arrangement amid handling of the composite [66], [67], [68], [69], [70], [71], [72], [73], [74], [75], [76], [77], [78], [79], [80], [81], [82], [83], [84], [85], [86], [87], [88], [89], [90], [112], [113], [114], [115], [116], [117], [118], [119], [120].

By looking at the structure, properties, stage conduct, rheology, preparing, and applications amongst SWNT and unbending bar polymers, SWNT were considered as polymeric materials [109], [110]. As said, the likeness between CNT (particularly SWNT) and polymers will permit the polymer chains to communicate with SWNT all the more promptly and nucleate on SWNT surfaces because of epitaxy. For this reason, SWNT were conceivably ready to adjust the chains parallel to the pivot course and layout polymer crystallization with expanded chain compliance. For polymeric materials extensional power (normally directed through shear streams in dissolve or arrangement) was required for actuating the broadened chain crystallization and the ensuing developing of the package like fibrils or shish-kebab structures [81], [82], [83], [84], [85], [86], [87], [88], [89], [90], [91], [92], [93]. This shearing instrument was likewise expected to develop fibrillar (expanded chain) precious stones in polymer/CNT crossover frameworks [42], [43], [78], [81]. The handling of expanded chain polymer precious stones in CNT frameworks was troublesome and not as normal as the perception of collapsed chain gem structures in these composites [82], [83], [84], [85].

Notwithstanding, a couple of past works had demonstrated that SWNT can prompt nucleation of expanded chain crystallization and layout the arrangement of polymer chains in PE [81], PBT [14], poly (ethylene terephthalate) (PET) [77], PAN [43], [44], [45], and PVA [42], [78] frameworks. The nearness of CNT was considered to add to the polymer core measure in the cross-breed framework, which stifles the vitality boundary for fibrillar crystallization by giving adequate heterogeneous nucleation destinations due to epitaxial connection [85]. Under calm conditions, the last crystalline structure and morphology were controlled by the filler attributes (i.e., fixation, synthesis, filler size, and shape) and by the cooperation between the filler and the polymer network. Within the sight of the shear stream, the affecting impacts reach out to shear rate, shear length, and the cooperation amongst shear and fillers [13]. In a polymer/nano-particles half and half framework, the presentation of nano-fillers and polymers into shear stream has been appeared to make a synergistic impact for advancing crystallization, because of the adjustments in the nearby feelings of anxiety and introduction of chains encompassing the nano-particles upon the use of shear [13], [85], [96]. Hence, the pole like CNT can enormously incite anisotropic nucleation destinations at the interphase and advance the resulting precious stone development in the stream bearing. Under fitting shear stream at a crystallization temperature, PE and PAN had been appeared to take shape into broadened chain shish straightforwardly on SWNT [81], [87] surface, trailed by nucleation of collapsed chain lamellae. In view of the little point X-beam dissemi-

nating (SAXS) investigation for the unadulterated PBT framework and PBT/SWNT composites, it was demonstrated the simple low SWNT stacking (0.2 wt. %) can format the morphology of crystallization amid stream, giving a strategy to get an exceedingly attractive fibre-like morphology [114]. Patil et al. have inferred that inside the sheared PE/CNT nano-composite framework, the nearness of CNT essentially advances the polymer chain introduction, the length increment, what's more, the steadiness of the half breed shish-kebab structures, because of CNT templating chain arrangement as contrasted with the sheared unadulterated PE framework [77], [78], [79], [80], [81], [82], [83], [84], [85], [86], [87], [88], [89], [90], [91], [92], [93], [94], [95], [96], [97], [98], [99]. Wide-edge X-beam diffraction (WAXD) thinks about on drawn PET/SWNT composite demonstrated that arranged crystallization of PET was initiated by adjusted SWNT in a randomized PET dissolve [77]. This introduction of the PET survived even after re-dissolving [78], [79], [80], [81]. No introduction was seen in the re-liquefying process in the flawless PET framework, showing the templating part of SWNT upon shear for polymer crystallization [60], [61], [62], [63], [64], [65], [66], [67], [68], [69], [70], [71], [72], [73], [74], [75], [76], [77], [78], [79], [80], [81], [82], [83], [84], [85]. These examinations exhibit the synergistic impacts of the nearness of SWNT and shear stream on advancing polymer broadened chain crystallization at the interphase in the nano-composites. Notwithstanding templating, the utilization of unbending nano-carbons in polymer lattices may likewise empower expanded polymer chain arrangement amid handling [61]. Change in chain arrangement has been detailed where an introduction factor (f) increment from 0.5 to 0.8 was found. This along these lines prompted an intense increment in the mechanical execution of the composite when contrasted with the control fibre. This work exhibits the capacity to utilize one-of-a-kind nano-fillers to go about as an ointment amid attracting to encourage polymer chain augmentation and introduction. A few examinations had demonstrated that the polymer chains shape special arrangement within the sight of CNT, and this was not the situation in their nonattendance [61], [77], [78], [81], [94]. What was required now was the comprehension of how to exploit such a marvel amid handling of the composite. The extraordinary likenesses between the CNT and polymer [110] may bear the cost of chances to grow new extraordinary handling systems that can exploit such parallels to create superior polymer/CNT filaments with all around controlled miniaturized scale structures [90], [91], [92], [93], [94], [95], [96], [97], [98], [99].

Without great connection between the segments of the framework, the commitment from each was decreased. To date, the presentation of nano-materials and their utilization in composite frameworks had demonstrated that these filler materials can have a colossal effect on the lattice segments even with no advancement. Nevertheless, the larger parts of these changes had so far been incremental. Taking full favourable position of the CNT material requires more outline in accordance with the association between the filler and the network, scattering forms, and arrangement of this half and half framework amid fibre turning. Hence, future-handling methodologies of polymer/CNT materials should consolidate some demonstrating/computational angles to foresee what sort of impacts these parameters may really have on the polymer and Nano-filler [120], [121], [122], [123].

Conclusions

This audit condenses examines on the different parameters that influence the reinforcing components in polymer/CNT fibre composite frameworks as a component of preparing. CNT containing polymeric filaments had shown enhanced mechanical and physical properties, for example, elasticity, Young's modulus, strain-to-disappointment, strength, and protection from particle changes from both dissolvable and warmth medicines. Trial factors influencing composite handling incorporate CNT structure, scattering, interfacial cooperation, and arrangement/introduction of polymer chains and CNT. The mix of these elements should be very much controlled keeping in mind the end goal to enhance the resultant mechanical properties of the mass composite fibre. A comprehension of these elements was overwhelming and an awesome test in the field of nano-composite preparing. Nevertheless, expanding essential test knowledge combined with computational and "materials by configuration" methodologies will prompt more productive utilization of CNT in composites and better improvement of creation systems.

Competing Interests

The authors have declared that no competing interests exist.

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