AMORPHOUS CARBON-BORON NITRIDE NANOTUBE HYBRIDS

A method for joining or repairing boron nitride nanotubes (BNNTs). In joining BNNTs, the nanotube structure is modified with amorphous carbon deposited by controlled electron beam irradiation to form well bonded hybrid a-C/BNNT structures. In repairing BNNTs, the damaged site of the nanotube structure is modified with amorphous carbon deposited by controlled electron beam irradiation to form well bonded hybrid a-C/BNNT structures at the damage site.

18 Claims, 8 Drawing Sheets
Figure 3a

Figure 3b

Figure 3c

Figure 3d
First tensile test

Second tensile test

Figure 5d

Figure 5e

Figure 5f

Figure 5g
AMORPHOUS CARBON-BORON NITRIDE NANOTUBE HYBRIDS

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR DEVELOPMENT

The U. S. Government has a paid-up license in this invention and the right in limited circumstances to require the patent owner to license others on reasonable terms as provided for by the terms of Cooperative Agreement No. NCC-1-02043 awarded by the National Aeronautics and Space Administration.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to nanocomposites, and more particularly to amorphous carbon-boron nitride nanotube (BNNT) hybrids.

2. Description of Related Art

All references listed in the appended list of references are hereby incorporated by reference, however, as to each of the above, to the extent that such information or statements incorporated by reference might be considered inconsistent with the patenting of this/these invention(s) such statements are expressly not to be considered as made by the applicant(s). The reference numbers in brackets below in the specification refer to the appended list of references.

Utilizing the full mechanical capabilities of individual nanotubes is a primary research goal in nanotube reinforced nanocomposite materials. Most studies on structural applications of nanomaterials, such as carbon nanotubes (CNTs), have focused on attempts to improve dispersion in structural matrices to achieve or exceed the performance of state-of-the-art carbon fiber reinforced polymer (“CFRP”) composites. This approach has yet to yield mechanical properties that compete with CFRPs, the aerospace structural material of choice [10, 11], because CNTs have not demonstrated the load carrying capacity of carbon fibers due to poor intertube load transfer and physical defects created during processing and fabrication. Practical use of these nanomaterials requires creating stable and strong linkages between nanotubes without sacrificing their mechanical advantage. Cross-linking between shells via electron beam irradiation [12-15] and application of large compressive forces [16] have been studied and offer a viable approach to improve tube-to-tube load transfer and hence, mechanical properties. However, these approaches result in unwanted mechanical degradation and have limitations in scale-up for their applications to hierarchical macroscopic nanocomposite materials.

It is a primary aim of the present invention to use amorphous carbon (a-C) to form stable connections between the tubes using electron beam irradiation.

It is an object of the invention to provide a method for in-situ transmission electron microscope (TEM)-atomic force microscope (AFM) techniques which precisely position BNNT specimens and use electron beam radiation to deposit a-C to modify and join BNNTs one or more times.

It is an object of the invention to provide a method for joining BNNTs in which the a-C joint on BNNT structures is comparable with those of currently available structural fibers and films such as both CNT and BNNT yarns, carbon fibers, and carbon fiber reinforced composites, both CNT and BNNT sheets, both CNT and BNNT composites and mixed composites of CNTs, boron carbon nitrides (BCN), and BNNTs.

It is an object of the invention to provide a method for a-C welding of BNNT structures which transfers load between the tubes for structural material designs.

Finally, it is an object of the present invention to accomplish the foregoing objectives in a simple and cost effective manner.

The above and further objects, details and advantages of the invention will become apparent from the following detailed description, when read in conjunction with the accompanying drawings.

SUMMARY OF THE INVENTION

The present invention addresses these needs by providing a method for joining or repairing boron nitride nanotubes (“BNNTs”). In the joining method, a plurality of BNNTs is provided and amorphous carbon is deposited on the surface of the BNNTs to form well bonded hybrid a-C/BNNT structures. The BNNTs are preferably fabricated by a pressurized vapor/condenser method using a CO₂ laser and are preferably highly crystalline. The BNNTs preferably consist of few-walled BNNTs having diameters of less than 7 nm. The amorphous carbon is preferably deposited on the BNNT surface by electron beam induced deposition. To repair boron nitride nanotubes (“BNNTs”), at least one damaged BNNT is provided and amorphous carbon is deposited on the surface of the at least one BNNT at the damage site to form hybrid a-C/BNNT at the damage site; the remaining steps are the same as for joining BNNTs. Finally, these steps can be used for forming hybrid a-C/BNNT boron nitride nanotubes (“BNNTs”) structures by depositing amorphous carbon on the surface of the BNNTs to form well bonded hybrid a-C/BNNT structures.

BRIEF DESCRIPTION OF THE DRAWINGS

A more complete description of the subject matter of the present invention and the advantages thereof, can be achieved by reference to the following detailed description by which reference is made to the accompanying drawings in which:

FIG. 1a shows a representative high resolution-TEM (JEOL JEM-2100F) image of BNNT bundles fabricated by the pressurized vapor/condenser method using a CO₂ laser;

FIG. 1b shows a representative high resolution-TEM (JEOL JEM-2100F) image of single-walled BNNTs fabricated by the pressurized vapor/condenser method using a CO₂ laser;

FIG. 1c shows a representative high resolution-TEM (JEOL JEM-2100F) image of double-walled BNNTs fabricated by the pressurized vapor/condenser method using a CO₂ laser;

FIG. 1d shows a representative high resolution-TEM (JEOL JEM-2100F) image of quadruple-walled BNNTs fabricated by the pressurized vapor/condenser method using a CO₂ laser.

FIGS. 2a-2f show TEM images and force displacement curves of mechanical tests conducted on pristine BNNTs in a TEM-ADF holder inside a TEM;

FIG. 2a shows a TEM image of an individual BNNT tested under compression before buckling;

FIG. 2b shows a TEM image of an individual BNNT tested under compression after buckling;

FIG. 2c shows a force displacement curve for the compressive test is shown in FIGS. 2a and 2b;

FIG. 2d shows a TEM image of a pristine BNNT under tension before breaking;
FIG. 2e shows a TEM image of a pristine BNNT under tension after breaking;

FIG. 2f shows a reconstructed stress-strain curve of the BNNT shown in FIGS. 2d and 2e;

FIGS. 3a-3g show a series of TEM images during multiple tensile tests with a-C/BNNT hybrid specimens in which:

FIG. 3a shows a TEM image before a-C/BNNT welding on the AFM tip by electron beam induced deposition (EBID);

FIG. 3b shows a TEM image after a-C/BNNT welding on the AFM tip by EBID;

FIG. 3c shows a TEM image after a first tensile test with the a-C/BNNT hybrid;

FIG. 3d shows a TEM image at the failure site after touching the broken pieces together;

FIG. 3e shows a TEM image after a second tensile test after tip-to-tip connection using a-C to repair the broken area;

FIG. 3f shows a TEM image after a second tensile test after tip-to-tip connection using a-C by EBID at the failure location of the first test to repair the broken area;

FIG. 3g shows reconstructed stress-displacement curves from the first and second tensile tests;

FIGS. 4a-4k show a series of TEM images during multiple mechanical tests with the same BNNT using a-C as a welding material;

FIG. 4a shows a TEM image of a pristine BNNT attached to the AFM tip with a-C by using electron beam irradiation and then tested under compression until it fractured, but did not completely break in two;

FIG. 4b shows a TEM image of a pristine BNNT attached to the AFM tip with a-C by using electron beam irradiation and then tested under compression until it fractured, but did not completely break in two;

FIG. 4c shows a TEM image of a pristine BNNT attached to the AFM tip with a-C by using electron beam irradiation and then tested under compression until it fractured, but did not completely break in two;

FIG. 4d shows a TEM image of a pristine BNNT attached to the AFM tip with a-C by using electron beam irradiation and then tested under compression until it fractured, but did not completely break in two;

FIG. 4e shows the a-C/BNNT specimen repaired using a-C by electron beam joining;

FIG. 4f shows a second compression testing of the repaired a-C/BNNT hybrid until it fractured prior to ultimate failure;

FIG. 4g shows a second compression testing of the repaired a-C/BNNT hybrid until it fractured prior to ultimate failure;

FIG. 4h shows the new fracture which did not appear at the previous repair site, but rather developed at a different location during the second compression test shown in FIGS. 4f and 4g;

FIG. 4i shows the new fracture which did not appear at the previous repair site, but rather developed at a different location during the second compression test shown in FIGS. 4f and 4g;

FIG. 4j shows a third compression testing of the repaired a-C/BNNT hybrid;

FIG. 4k shows force-displacement data for each of the first, second and third compressions tests in FIGS. 4a-4j;

FIGS. 5a-5g show a series of TEM images and a stress-strain curve for the tensile testing of a-C/BNNT hybrid specimens;

FIG. 5a shows a TEM image of the initial tensile test performed on the twice repaired compression specimen shown in FIG. 4j;

FIG. 5b shows a TEM image of the initial tensile test performed on the twice repaired compression specimen shown in FIG. 4j after complete breakage of the sample;

FIG. 5c shows electron beam induced end-to-end joining with a-C of the severed hybrid a-C/BNNT specimen;

FIG. 5d shows electron beam induced end-to-end joining with a-C of the severed hybrid a-C/BNNT specimen in which to form an end-to-end joint bonded with a-C, one broken end was precisely aligned with another broken end and joined with a-C;

FIG. 5e shows the repaired sample subjected to tensile testing again and the failure occurring at the same location;

FIG. 5f shows the repaired sample subjected to tensile testing again and the failure occurring at the same location;

FIG. 5g shows the stress-strain curve for the test performed in FIGS. 5a through 5f;

FIG. 6a shows the formation of a lap bond, by precisely aligning the broken ends from the above discussed tensile tests (specimen image from FIG. 50 to contact;

FIG. 6b shows the formation of a lap bond, by precisely aligning the broken ends from the above discussed tensile tests (specimen image from FIG. 50 to overlap and then joining by a-C via electron beam joining;

FIG. 6c shows first lap shear test specimen prepared by side-by-side electron beam joining;

FIG. 6d shows the failure sample subjected to lap shear testing and the failure occurring at the a-C joint;

FIG. 6e shows residual a-C in the top half of the broken specimen and precisely aligning the broken ends from the above discussed first lap shear test to opposite side of the broken ends;

FIG. 6f shows the formation of a lap bond by a-C via electron beam joining and the access a-C joining via electron beam joining;

FIG. 6g shows the unbroken sample after second lap shear test; and

FIG. 6h shows lap shear stress-strain curves from two successive tests.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

The following detailed description is of the best presently contemplated mode of carrying out the invention. This description is not to be taken in a limiting sense, but is made merely for the purpose of illustrating general principles of embodiments of the invention. The embodiments of the invention and the various features and advantageous details thereof are more fully explained with reference to the non-limiting embodiments and examples that are described and/or illustrated in the accompanying drawings and set forth in the following description. It should be noted that the features illustrated in the drawings are not necessarily drawn to scale, and the features of one embodiment may be employed with the other embodiments as the skilled artisan recognizes, even if not explicitly stated herein. Descriptions of well-known components and techniques may be omitted to avoid obscuring the invention. The examples used herein are intended merely to facilitate an understanding of ways in which the invention may be practiced and to further enable those skilled in the art to practice the invention. Accordingly, the examples and embodiments set forth herein should not be construed as limiting the scope of the invention, which is defined by the appended claims. Moreover, it is noted that like reference numerals represent similar parts throughout the several views of the drawings.

Generally, the present invention relates to CNTs which have received significant attention due to their outstanding combination of mechanical, electrical and thermal properties. However, applications requiring greater thermal and chemi-
Mechanical properties of individual pristine BNNTs, and a-C/BNNT hybrids were measured inside a TEM (200 keV, Philips CM200) equipped with an integrated AFM system (TEM-AFM, Nanofactory). To minimize damage to the BNNTs prior to testing, the samples were prepared by touching an as-grown dry BNNT forest with a holey carbon-coated Cu half grid. The sample was fixed on the grid using one drop of ethanol. The assembly was vacuum-dried at 75°C before testing. Touching an individual BNNT with the conductive Au-coated AFM tip (The spring constant of AFM cantilever was 3.6-4.0 N/m) was sufficient to attach it to the tip for testing. In the case of mechanical testing using a customized sample with the TEM-AFM set-up, one end of the BNNT was attached to the AFM tip with a-C by using electron beam irradiation to ensure good contact between the tube and the tip. Development of a-C Using e-Beam Irradiation on the BNNT Surface

The structure of nanotubes can be engineered using electron beam (13, 18-25) or ion beam (13) irradiation. The a-C was grown on the BNNT surface by EBID at the contact area for a few minutes without using a liquid nitrogen (LN2) trap to condense gas phase carbon species. The a-C joins BNNTs together to form well bonded hybrid a-C/BNNT structures. The TEM-AFM set-up used in this study allows for in-situ nano-manipulation to position the BNNTs for bonding, structural modification using electron beam, and mechanical measurements such as compression, tensile, and lap shear tests, in addition to conventional TEM tasks such as morphology and chemical composition observation. It is important to note that a-C readily forms with a focused electron beam only when a LN2 trap is not used. Also, a focused electron beam can cause damage to the BNNTs when a LN2 trap is used. Therefore, during mechanical testing the electron beam was spread to minimize BNNT damage.

Compression and Tensile Tests on Pristine BNNTs

The mechanical tests were conducted on pristine BNNTs in a TEM-AFM holder inside a TEM. Representative TEM images and force displacement curves are presented in FIGS. 2a-2f. The applied force for direct calculation of Young’s modulus, tensile strength and strain of an isolated tube were obtained from the forward and reverse movement of the piezo-driven positioning component of the AFM holder. FIGS. 2a and 2b are TEM images of an individual BNNT tested under compression before and after buckling, respectively. In FIG. 2e, a force displacement curve for the compressive test is shown. Buckling of the BNNT samples occurred in the middle of the tube where the diameter is smallest. The Young’s modulus of the individual BNNT from the buckling was calculated to be 0.84 TPa, using Euler’s formula, which agrees with values discussed in the literature [2, 4]. The Euler’s formula was derived for a compression test of a cylinder where the column buckling is the predominant failure mode: $F_{cylinder}=(\pi^2 E I)/L^2$, $L=\pi(d_i^4-d_o^4)/64$ with $E$ the Young’s modulus, $I$ the moment of inertia, $L$ the length of BNNT between the two contact points, $d_o$ the external nominal BNNT diameter, and $d_i$ internal nominal BNNT diameter. In this specific case, the external ($d_o$) and internal ($d_i$) diameters and the length ($L$) of the tube were 3.1, 1.7, and 52.3 nm, respectively, measured from the TEM image and the buckling stress was around 1.7 GPa. Herein, we used the two-dimensional projection of the external diameter ($d_o$) to calculate the buckling stress due to uncertainty of exact cross-sectional area of the tube.
Tensile tests were also performed with few other BNNTs. FIGS. 2d and 2e are representative TEM images of a pristine BNNT under tension before and after breaking, and FIG. 2f is a reconstructed stress-strain curve. The measured tensile strength was 20.7 GPa and the strain was 7.3%. The Young’s modulus obtained from linear fitting of the corresponding stress-strain curve (FIG. 2f) was 0.61 TPa, which is close to the value from the compression test. The tensile strength is calculated by assuming that only the outermost shell of the tube carries the tensile load; thus, the cross-sectional area is estimated by \( A = \pi d^2 \), where \( d \) is the external nominal BNNT diameter. For the BNNT shown in FIGS. 2d and 2e, the external diameter (\( d_2 \)) of the BNNT was 2.7 nm, and the interlayer distance (\( d_1 \)) of the tube walls was about 0.34 nm [1]. The energy-to-failure of this individual BNNT was calculated to be 350 J/g using 1.4 g/cm³ as the estimated density of BNNT. This value of the energy at failure is slightly lower than that with isolated double-walled CNT bundles (~500 J/g) [24]. However, the value is much higher than that of state-of-the-art CNT yarns (below 100 J/g) [24]. In the case of an individual BNNT, there is no interaction between tubes which is in contrast to double-walled CNT bundles, nor are there any interactions with polymeric binder and CNTs as is the case with CNT yarns [24]. Note that the measured tensile strain and the energy-to-failure are overestimated due to pre-applied compressive forces necessary to ensure contact during the joining process. The pre-compression applied during tube joining by EBID caused some alignment and stretching of the tron beam irradiation and then tested under compression until reversed to perform tensile testing. The tensile stress was applied to the pristine BNNT using a-C as a welding material. A pristine BNNT (The ratio of cross-sectional area \( A_{\text{BNNT}} \) is 6.1 with the diameters of 6.4 and 17.1 nm prior and after the a-C deposition, respectively. This value is about four times less than the measured value of the individual pristine BNNT (0.84 TPa) discussed in the previous repair site, but rather developed at a different location during multiple mechanical tests with the same BNNT using a-C as a welding material. A pristine BNNT was attached to the AFM tip with a-C by using electron beam irradiation and then tested under compression until it fractured, but did not completely break in two (see FIGS. 3a-3f for TEM images of this test, and FIG. 3d, first test, for force-displacement data). The specimen was subsequently subjected to repair via electron beam joining with the beam focused to fill or cover the fractured area with a-C. The electron beam was subsequently spread to irradiate the entire tube and build up a-C along the tube surface to form an a-C/BNNT hybrid structure. After attaching to the AFM tip and performing a-C growth on the BNNT shown in FIG. 4a, the Young’s modulus of the a-C coated BNNT under compression was determined to be 0.20 TPa (see FIGS. 4a-4e). Here, the ratio of cross-sectional area between the a-C \( (A_{\text{a-C}}) \) and BNNT \( (A_{\text{BNNT}}) \) is 6.1 with the diameters of 6.4 and 17.1 nm prior and after the a-C deposition, respectively. This value is about four times less than the measured value of the individual pristine BNNT (0.84 TPa) discussed in the previous experiment. The reduction in Young’s modulus is likely due to an increase in overall contribution of a-C as a result of the a-C deposition. After the first compression test (crack location marked with a black arrow), the same a-C/BNNT specimen was repaired using a-C by electron beam joining (see FIGS. 4d and 4e). The repaired a-C/BNNT hybrid was subjected to compression testing again until it fractured prior to ultimate failure (see FIG. 4f-4g). The new crack did not appear at the previous repair site, but rather developed at a different location during the compression test shown in FIG. 4f (marked with a red arrow). The Young’ modulus of the repaired a-C/BNNT hybrid structure (The ratio of cross-sectional area \( A_{\text{a-C}}/A_{\text{BNNT}} \)) is 17.2) decreased to 0.10 TPa (FIG. 4f, red line). The hybrid structure did not break completely due to the resistance from the BNNT core which has a higher elastic modulus.

The tested sample was subsequently repaired a second time using electron beam joining with a-C. The crack was refilled with a-C using focused electron beam irradiation as shown in FIGS. 4h and 4i and then the repaired a-C/BNNT hybrid (The ratio of cross-sectional area \( A_{\text{a-C}}/A_{\text{BNNT}} \)) is 23.7) was re-tested (FIG. 4j). In this round, the Young’s modulus dropped...
further to 0.086 TPa (FIG. 4k, blue line). The force displace-
ment curves for the three tests are shown in FIG. 4k. Interest-
ingly, upon the third compression test, the sample did not
buckle or fracture up until 150 nm of displacement was
reached, behaving in a more ductile manner. The maximum
buckling stresses calculated from FIG. 4k were increased to
152, 213, and 277 MPa, respectively, with each successive
compression test. The structural and mechanical properties
of each BNNT and a-C/BNNT hybrid tested in this study are
summarized in Table 1.

**TABLE 1**

<table>
<thead>
<tr>
<th>Characteristics summary of each BNNT and a-C/BNNT hybrid tested in this study.</th>
</tr>
</thead>
<tbody>
<tr>
<td>External Diameter (nm)</td>
</tr>
<tr>
<td>------------------------</td>
</tr>
<tr>
<td>pristine BNNT, compression</td>
</tr>
<tr>
<td>pristine BNNT, tension</td>
</tr>
<tr>
<td>a-C/BNNT Hybrid</td>
</tr>
<tr>
<td>1st tensile break</td>
</tr>
<tr>
<td>2nd tensile break</td>
</tr>
<tr>
<td>a-C/BNNT Hybrid</td>
</tr>
<tr>
<td>1st compression</td>
</tr>
<tr>
<td>2nd compression</td>
</tr>
<tr>
<td>3rd compression</td>
</tr>
<tr>
<td>Repaired a-C/BNNT Hybrid</td>
</tr>
<tr>
<td>(tip-to-tip joint)</td>
</tr>
<tr>
<td>1st tensile break</td>
</tr>
<tr>
<td>2nd tensile break</td>
</tr>
<tr>
<td>(side-by-side joint)</td>
</tr>
<tr>
<td>1st lap-shear break</td>
</tr>
</tbody>
</table>

*External and internal diameters of individual BNNTs and a-C/BNNT hybrids were determined using TEM images of
the failure site during tensile tests and from kinked sites during compression tests.

Tensile Tests on Repaired a-C/BNNT Hybrid

A series of TEM images and a stress-strain curve are pre-
sented in FIGS. 5a-5g for the tensile testing of a-C/BNNT hybrid specimens. Using the same set-up, the force direction
was reversed to perform tensile testing. The tensile stress
was calculated with the measured AFM force divided by the cross-
sectional area of the hybrids. In this specific case, the external
(d4) diameters of the hybrids were 31.8, and 39.8 nm for each
test as measured from the TEM images. The observation of the
tip movement and the measured elongation of the a-C/
BNNT hybrid structure under tensile load were used to cal-
culate strain. The TEM images in FIGS. 5a and 5b are of the initial
tensile test performed on the twice repaired compression
specimen (FIG. 4f). The complete breakage of the sample is seen clearly in FIG. 5b. The tensile strength exhibited
by this sample was 1.2 GPa. Two different slopes were
apparent from the stress-strain curve for the first test shown in
FIG. 5g. The tensile failure occurred at a location previously
repaired by a-C (second repair, marked by a red arrow), and
the broken section exhibited a clean break. The lower slope
from the first tensile test is attributed to the tensile stress of the
a-C layer while the higher slope above 4% strain is from the
damaged BNNT beneath the a-C layers.

Electron beam induced end-to-end joining with a-C of the
severed hybrid a-C/BNNT specimen is shown in FIGS. 5e and
5f. To form an end-to-end joint bonded with a-C, one
broken end was precisely aligned with another broken end
values of a-C cell with 50% of sp3 sites by the tight-binding
molecular dynamics simulation [27]. However, the mechanical
properties of a-C joints could possibly be enhanced
through graphitization of a-C by further heat treatment [21].

Lap Shear Tests on Repaired a-C/BNNT Hybrid

In order to form a lap bond, the broken ends from the above
discussed tensile tests (specimen image from FIG. 50 were
precisely aligned to overlap and then joined by a-C via elec-
tron beam joining (FIGS. 6a and b). This specimen configu-
ration was tested to obtain lap shear strength under tension.
The lap shear strength of the a-C bonded BNNT specimen
was 0.25 GPa (FIGS. 6c and 6d), which is two orders of
magnitude higher than what is found between the outermost
shell and the neighboring inner shell of multi-walled CNTs
(0.3 MPa) [25]. The contact area on the lap shear test was
assumed to be rectangular in shape with a contact length (86.5
nm) determined from the amount of overlap and a contact
width equal to the a-C coated tube diameter (55.8 nm). FIG.
6f shows lap shear stress-strain curves from two successive
tests. The lap shear specimen failed during 400 nm of piezo-
actuator driven movement in the first test (FIG. 6d). Some
residual a-C can be seen in the top half of the broken specimen
(FIG. 6e), which indicates that the failure occurred predomi-
nately at the a-C joined site under the shear load. This speci-
men was manipulated so that the opposite surfaces which had
not been previously bonded were aligned (FIG. 6e) and sub-
sequently bonded with a-C (FIG. 6f). In this case, the speci-
Herein and defined in the following claims. Obviously, many modifications may be made without departing from the basic spirit of the present invention. Accordingly, it will be appreciated by those skilled in the art that within the scope of the appended claims, the invention may be practiced other than has been specifically described herein. Many improvements, modifications, and additions will be apparent to the skilled artisan without departing from the spirit and scope of the present invention as described herein and defined in the following claims.

LIST OF REFERENCES

[18] Banhart F 2001 *Nano Lett.* 1, 329-332

What is claimed is:

1. A method for joining boron nitride nanotubes, comprising: providing a plurality of BNNTs; precisely aligning the BNNTs to form a contact area; and depositing amorphous carbon on the surface of the BNNTs at the contact area to form bonded hybrid a-C/BNNT structures in which load is transferred between the BNNTs by amorphous carbon disposed between the BNNTs.

2. The method of claim 1 wherein the BNNTs are fabricated by a pressurized vapor/condenser method using CO2 laser.

3. The method of claim 1 wherein the BNNTs are crystalline.

4. The method of claim 1 wherein the BNNTs consist of at least one of single-walled and multi-walled BNNTs.

5. The method of claim 1 wherein the BNNTs have diameters of less than 7 nm.

6. The method of claim 1 wherein the amorphous carbon is deposited on the BNNT surface by electron beam induced deposition.

7. A method for repairing boron nitride nanotubes, comprising: providing at least one damaged BNNT having a damaged site; depositing amorphous carbon on the surface of the at least one damaged BNNT at the damage site to form hybrid a-C/BNNT at the damage site.

8. The method of claim 7 wherein at least one BNNT is fabricated by a pressurized vapor/condenser method using a CO2 laser.

9. The method of claim 7 wherein at least one BNNT is crystalline.

10. The method of claim 7 wherein at least one BNNT is a few-walled BNNT.

11. The method of claim 7 wherein at least one BNNT has diameters of less than 7 nm.

12. The method of claim 7 wherein the amorphous carbon is deposited on the surface of the at least one BNNT by electron beam induced deposition.

13. A method for forming hybrid a-C/BNNT structures, comprising: providing a plurality of BNNT; precisely aligning the BNNTs to form a contact area; and depositing amorphous carbon on the surface of the BNNTs at the contact area to form bonded hybrid a-C/BNNT structures in which load is transferred between the BNNTs by amorphous carbon disposed between the BNNTs.

14. The method of claim 13 wherein the BNNTs are fabricated by a pressurized vapor/condenser method using a CO2 laser.

15. The method of claim 13 wherein the BNNTs are crystalline.

16. The method of claim 13 wherein the BNNTs consist of at least one of single-walled and multi-walled BNNTs.

17. The method of claim 13 wherein the BNNTs have diameters of less than 7 nm.

18. The method of claim 13 wherein the amorphous carbon is deposited on the BNNT surface by electron beam induced deposition.

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