Effect of pyrolytic carbon interface thickness on microstructure and mechanical properties of lightweight zirconium boride modified carbon-bonded carbon fiber composites

Xianghong Xu a, Baosheng Xu b, Changqing Hong c,*, David Hui d,**

a State Key Laboratory of Nonlinear Mechanics (LNM), Institute of Mechanics, Chinese Academy of Sciences, Beijing 100190, PR China
b AML, Department of Engineering Mechanics, School of Aerospace Engineering, Tsinghua University, Beijing 100084, PR China
c Science and Technology on Advanced Composites in Special Environment Laboratory, Harbin Institute of Technology, Harbin 150001, PR China
d Department of Mechanical Engineering, University of New Orleans, LA 70148, USA

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ABSTRACT

To improve the mechanical properties of carbon-bonded carbon fiber (CBCF) composites, they are firstly fabricated by chemical vapor deposited (CVD) pyrolytic carbon (PyC) coating layer on the carbon fiber surface, and then modified by zirconium boride (ZrB 2 ) using three cycles of precursor infiltration and pyrolysis (PIP) process. The effects of different PyC interface thickness on the microstructure and mechanical properties of ZrB 2 modified PyC coated CBCF (PyC–CBCF/ZrB 2 ) composites were studied and characterized. As the PyC thickness increased from 0.5 to 3.6 μm, the flexural properties of PyC–CBCF/ZrB 2 composites are noticeably enhanced in x/y and z direction, respectively. Mechanical enhancements for PyC–CBCF/ZrB 2 composites are mainly attributed to the effective interface bonding between carbon fiber and PyC, crack deflection and branching within the laminar PyC layer and carbon fiber pullout from PyC interface coating.

1. Introduction

Carbon-bonded carbon fiber (CBCF) composites possess many excellent properties such as low density, low thermal conductivity, low coefficient of thermal expansion (CTE), good resistance of irradiation and high temperature capability [1–4]. Although the deposition of external coatings on CBCF composites can prevent them from oxidation in oxidation-containing atmospheres, CBCF composites have a poor mechanical strength to meet the increasing demand of actual application [2]. Therefore, the enhancement of mechanical properties is crucial for CBCF composites used in high temperature and load-bearing environment.

To overcome these limits, many researchers have focused on the surface modification of carbon fibers to enhance the interfacial adhesion between carbon fibers and matrix. For example, Li et al. [3] studied the C/C composites with in situ grown CNFs using natural gas as carbon source. The CNFs are beneficial to the improvement of mechanical properties of C/C composites due to bridging effect which redistributes the load while the cracks extend toward fibers or CNFs leading to higher toughness as the more complex crack propagation channels dissipate more energy. Lim et al. [4] succeeded to prepare C/C composites with improved mechanical strength by using a precursor made of CNFs grafted on carbon microfilaments. The as-synthesized carbon nanofibers acted as a “solid glue” through the formation of numerous nano-and micro junctions between the microfilaments and exhibited high elasticity and strength compared to raw materials. Another method is to invite the preparation process on the mechanical properties of C/C composites carried out by Weisshaus et al. [5]. The result indicated that the duration of graphitization tended to reduce compression strength of C/C composites. It is generally accepted that effective interfacial carbon layers can be remarkable enhancement of mechanical properties for C/C composites due to good thermal coefficient matching and remarkable mechanical property of interfacial carbon layers [6–9]. Especially for porous carbon-based composites, it can form a uniform thinner carbon layer on the carbon fiber surface by chemical vapor deposition (CVD), in which the pores provide the diffusion channel for gas containing carbon in a CVD process.

* Corresponding author. Tel./fax: +86 451 86403016.
** Corresponding author.
E-mail addresses: hongcq@hit.edu.cn (C. Hong), DHui@uno.edu (D. Hui).

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In this work, pyrolytic carbon (PyC) modified porous CBCFs (PyC–CBCF) composites were fabricated by the formation of a PyC coating layer on the carbon fiber surface via a CVD method. And then, the lightweight ZrB₂-modified PyC–CBCF (PyC–CBCF/ZrB₂) composites were fabricated by polymeric precursor infiltration and pyrolysis (PIP). The main purpose of the paper is to investigate the effect of PyC interphase thickness on the mechanical strength and fracture behavior of porous PyC–CBCF/ZrB₂ composites.

2. Experimental

The schematic illustration of the typical microstructure of CBCF composites and manufacturing process for PyC–CBCF/ZrB₂ composites are shown in Fig. 1. As seen from Fig. 1(a)–(c), the CBCF composites consist of chopped carbon fiber network bonded together at the intersections of the fibers by discrete regions of vitreous carbon. In addition, the discontinuous carbon fibers are orientated into layers to form a 2D planar random structure at xy direction [10,11]. The schematic illustration of manufacturing process for PyC–CBCF/ZrB₂ composites is shown in Fig. 1(d)–(g). Firstly, porous PyC–CBCF composites were fabricated by the formation of a PyC coating layer on the carbon fiber surface via a CVD method (Fig. 1(d), (e)). And then, the lightweight PyC–CBCF/ZrB₂ composites were fabricated via a PIP process (Fig. 1(f), (g)).

2.1. Starting materials

Rayon-based carbon fiber (diameter of 75 μm and length of 1 mm) was purchased from Ji lin Ji yan High-tech Fibers Co., Ltd. (Jilin, China). Phenolic resin powder (PF4090, fineness >95%/200 mesh) was obtained from the Holyspring chemical Co., Ltd. (Juxian, China). Polyethylene imine (PEI) with an average molecular weight of 10,000 was obtained from Aladdin Co. Ltd. (Shanghai, China). ZrB₂-containing precursors was obtained from Institute of Process Engineering (Chinese Academy of Science, Beijing, China).

2.2. Preparation of CBCF composites

CBCF composites were prepared using the classical pressure filtration procedure [12]. A 200 ml aqueous solution consisting of 60 mg PEI was subject to a vigorous stirring in a 250 ml beaker, and 5 g short carbon fiber was then added slowly to the solution to form a water-based slurry. After stirring for 20 min, the same weight of phenolic resin powder was added to the slurry and the slurry was stirred for another 30 min to achieve good dispersion of water-based slurry. The water-based slurry was then poured into a cylindrical mould [13] to remove the water from the slurry under an appropriate pressure to obtain a desired density. The acquired wet billets were cured at 80 °C for 2 h and 150 °C for 3 h. Then the cured samples were carbonized at 1000 °C in Ar for 1 h. The CBCF composites with a density of 0.26 g/cm³ and porosity of 84.71% were prepared.

2.3. Preparation of PyC–CBCF–ZrB₂ composites

PyC was deposited on the fibers surface in CBCF framework by chemical vapor deposition (CVD) at 1200 °C using propane as carbon source. Three different PyC interphase thicknesses of 0.5, 2.6 and 3.6 μm were obtained by controlling deposition time for 30, 100 and 150 h, respectively. Ultimately, PyC–CBCF composites with different interface thicknesses were impregnated into ZrB₂ polymeric precursor to prepare ZrB₂ modified PyC–CBCF composites by a PIP method [2]. In a typical experiment, the PyC–CBCF composites were impregnated into ZrB₂-containing polymeric precursor using a vacuum infiltration devise, drying at 80 °C in a drying oven at Ar atmosphere. The pyrolysis was performed at a heating rate of 2 °C/min to the desired temperature of 1500 °C for 2 h under flowing Ar atmosphere. The composites were densification by another two cycles of ZrB₂-containing polymeric precursor infiltration and pyrolysis.

For comparison, CBCF composites without CVD PyC were fabricated by three cycles of ZrB₂-containing polymeric precursor infiltration and pyrolysis under the same condition.

Fig. 1. Schematic illustration of the typical microstructure of CBCF composites and the manufacturing process for PyC–CBCF/ZrB₂ composites.
2.4. Characterization

The phase composition of the PyC–CBCF composites was investigated by a Rigaku D/max-Rb X-ray diffraction (XRD). The morphology of the composites was investigated by a Holland FEI Sirion scanning electron microscopy (SEM). The samples for three-point bending tests were machined from bigger blocks and the effective size of the samples was $3 \times 4 \times 36 \text{ mm}^3$. The span used to be 20 mm and a crosshead speed of 0.5 mm/min was utilized during all tests. Samples were tested parallel to the x/y and z axes with the mechanical property anisotropy ratio being defined as the ratio of mechanical property values in the x/y and z planes.

3. Results and discussion

3.1. PyC coated CBCF composites

Fig. 2 shows the microstructures of PyC coated carbon fiber with different deposition time. When deposition time increases from 30 h to 150 h, the PyC interface thickness increases from 0.5 to 3.6 $\mu$m according to the cross-section morphology (Fig. 2(a), (c) and (e)). The surface of carbon fiber transforms from rough to smooth shape with the increasing of PyC thickness (Fig. 2(b), (d) and (f)). Furthermore, the PyC coated CBCF is oriented preferentially and arranged randomly in the z direction (Fig. 2(h), (j) and (l)) and perpendicular to z axis in the x/y direction (Fig. 2(g), (i) and (k)) due to the consequence of the vacuum or pressured molding process [13]. Clearly, the porosity of PyC coated CBCF both in x/y and z direction decreases with the deposition time and the PyC distributed uniformly on the surface of CBCF composites. Table 1 summarizes the density and porosity of PyC coated CBCF composites. The density for composites increases from 0.26 to 0.67 g/cm$^3$ and the open porosity decreases from 84.71 to 54.9% as the deposition time increased.

The XRD patterns of virgin CBCF and PyC coated CBCF composites are shown in Fig. 3. Due to the virgin CBCF containing low-graphitization rayon carbon fiber framework, a turbostratic carbon structure exists corresponding to the appearance of (002), (103) and (110) broad diffraction peak [14,15]. However, it is clear that the (002) peak intensity becomes sharper as the increasing of the PyC thickness, because the increase of deposition time enhances the good graphitization of PyC compared to that of virgin carbon fiber.

The ZrB$_2$ modified PyC–CBCF composites were prepared by a precursor infiltration and pyrolysis method. The densification of ZrB$_2$ modified PyC–CBCF composites increases and most pores in PyC–CBCF composites are filled and blocked by ZrB$_2$ ceramic and the ceramic are coated on the carbon fibers and PyC interface surface (shown in Fig. S1). Table 2 shows the density and porosity of

![Fig. 2. Microstructures of PyC coated CBCF with different deposition time. (a), (b), (g) and (h): PyC$_{0.5}$–CBCF; (c), (d), (i) and (g): PyC$_{2.6}$–CBCF; (e), (f), (k) and (l): PyC$_{3.6}$–CBCF; (i) and (k): x/y direction; (b), (j) and (l): z direction.](image)
PyC–CBCF/ZrB₂ composites with different PyC interface thickness. The density of ZrB₂ modified PyC–CBCF composites increases from 0.79 to 0.98 g/cm³ and the porosity decreases from 48.5 to 38.8%.

### Table 1

| Samples       | PyC0.5–CBCF | PyC1.5–CBCF | PyC2.5–CBCF |
|---------------|-------------|-------------|-------------|
| Density (g/cm³) | 0.79        | 0.79        | 0.79        |
| Porosity (%)   | 48.5        | 40.1        | 38.8        |

### 3.2. Mechanical properties and fracture behavior

The flexural properties of the ZrB₂ modified PyC–CBCF with different PyC interface thicknesses are shown in Fig. 4. It can be seen that the PyC interface can effectively improve the mechanical property of CBCF/ZrB₂ as shown in Fig. 4(a) and (b). The flexural strength of CBCF/ZrB₂ composite is 1.6 and 1.08 M for x/y and z direction, respectively. When the PyC interface is introduced on the fiber surface, the composites significantly exhibit higher values in both x/y and z direction. The flexural strength of PyC0.5–CBCF/ZrB₂, PyC1.5–CBCF/ZrB₂ and PyC2.5–CBCF/ZrB₂ are 10.2, 17.3 and 17.9 MPa for x/y direction, respectively, and 6.9, 13.6 and 16.0 MPa for z direction, respectively, which obtained 1018.75% and 1354.5% improvement compared with CBCF/ZrB₂. Moreover, the flexural modulus also has been significantly improved from 164 to 2028 MPa and from 57 to 1535 MPa for x/y direction and z direction, respectively.

Fig. 4(c) and (d) shows the typical flexural stress/strain curves for CBCF/ZrB₂ and PyC–CBCF/ZrB₂ with different PyC interface thicknesses. It is interesting to note that the PyC interface increases the flexural strain, especially for PyC interface thickness of 0.5 μm which increases the strain from 0.98% to 1.24% as shown in Fig. 4(c). Moreover, it is notable for the z direction that flexural strain increases with the increasing the PyC interface thickness as shown in Fig. 4(d). The mismatch between the thermal expansion coefficient of ceramic coatings and carbon fibers results in cracking along the interface between fibers and ceramic. Hence, the brittle ceramic coatings are easily peeled off from fibers as preparation of test samples. Therefore, it is weaker surface strength of CBCF/ZrB₂ as a result of fractures prior to reaching the maximum stress value (see insert in Fig. 4(c)) even at the starting stage of loading (insert in Fig. 4(d)). However, the PyC–CBCF/ZrB₂ does not present this fracture behavior, suggesting that the PyC interface enhances the CBCF composites framework. On the other hand, it is indicated that the introduction of ZrB₂ ceramic leads to the increase of brittle fracture of the sample as seen from Fig. 4(c) and (d).

It is clear that PyC interface plays some role in the mechanical reinforcement process [16]. So, the fracture surfaces of different samples failed in flexural tests are present to understand this fracture behavior as shown in Fig. 5. The ceramic coated fiber bundles in the different samples exhibit pulling-out behavior, presenting “gaps” or “peaks” feature on the fractured surface (Fig. 5(a), (c), (e) and (g)). The “gaps” tends to bigger for the PyC3.5–CBCF/ZrB₂ due to the strong adhesive force among fibers with a thicker PyC interface as seen in Fig. 5(g), Fig. 5(h), (d) and (b) shows the fracture surfaces of the coated fibers with different thickness of PyC interface. As seen from Fig. 5(b), the ZrB₂ ceramic layer shows a brittle fracture [17,18], and the carbon fiber present irregular fracture surface which increases the impact damage tolerance [19]. In addition, interfaces between fiber and ceramic layer exhibit debonding characteristic. It is interestingly found that the coated layer is visible plastic deformation when the PyC thickness is 0.5 μm (Fig. 5(d)), indicating that the coated layer appears plastic tension strain before the failure of the composites. It is very likely that a similar failure mechanism reported by Della Bona and co-workers [20] who pointed out that the ceramic core surface was placed under tensile stress during the flexural testing of multilayer structures. The carbon fiber with a thin PyC interface (0.5 μm) also presents an irregular fracture surface which indicates that the most of the energy from the load transmit through the thin PyC interface to carbon fiber and generate and expanse of cracks within carbon fiber. The effect of PyC with a thin thickness of 0.5 μm can increase the strain as shown in Fig. 4(c). When the PyC thickness increases to 2.6 and 3.6 μm, the carbon fiber pulls out from the coated PyC layer as shown in Fig. 5(f) and (h). Moreover, a very rigorous delamination within PyC interface can be seen in the insert image in Fig. 5(f) and (h). It is worthy to note that the PyC is present a laminar structure along the carbon fiber surface with deposition time during the CVD process [21–23]. Accordingly, the laminar PyC pullout from PyC layers, and the pullout increases with increasing of PyC thickness, which can enhance the mechanical properties of composites. Crack deflection and branching also occur within the laminar PyC interface as shown in insert image in Fig. 5(f) and (h), because the driving force for interfacial crack growth equals the fracture energy of the interface at a lower load than that required for the driving force of the penetrating crack to the fracture energy of the matrix [24]. For the PyC interface with thickness of 2.6 and 3.6 μm coated carbon fibers, the PyC interfacial crack release energy might increase and the cracks propagate through the adjacent PyC layers, leading to a fast and high energy fracture of carbon fibers, resulting in smooth and flat of fiber surface (Fig. 5(f) and (h)). As seen from Fig. 5(g) the mismatch of CET between PyC and ceramic result in forming interspaces which may influence the mechanical property of composites. Moreover, the strong interface bonding between carbon fiber and PyC due to better compatibility and obstinate PyC layer may lead to the brittle fracture for CBCF modified by PyC.

![Fig. 3. XRD spectra of original CBCF composites and PyC–CBCF composites with different deposition time.](image-url)
To understand the effect of the ceramic layer on the mechanical enhancement of composites, the microstructures of fiber surfaces after three-point flexural tests were carried out by SEM as shown in Fig. 6. Fig. 6 shows the ceramic layer coated on the surface of the carbon fiber framework via a PIP process. The appearance of microcracks is found in Fig. 6, attributing to the shrinking of ZrB2 during PIP process and the mismatch of CET between carbon and ceramic. It is worthy to note that the ceramic layers peel from the fiber surface and PyC interface, which indicates the ceramic layers present a lower strength and brittle fracture behavior. The ceramic layer firstly can peel off from the carbon surface due to the weak bonding strength, and it makes a little contribution to the strength of PyC$_3$ CBCF/ZrB$_2$ composites. Moreover, compared to Fig. 6(c) and (d), the modified carbon fiber surface (Fig. 6(b)) presents the appearance of large area peeling, showing that the PyC$_3$ CBCF/ZrB$_2$ composites with thinner PyC has larger plastic deformation than that with thicker PyC.

In addition, except for the fracture of single carbon fiber in the frameworks of composites as discussed former, the fiber bundles in the intersections of carbon fiber framework with different thickness of PyC interface also present different fracture behavior. The PyC interface strongly dominates mechanical properties and fracture behaviors for ZrB$_2$-modified PyC$_3$ CBCF composites.

(1) The mechanical properties of PyC$_3$ CBCF/ZrB$_2$ increase with the thickness of PyC interface. As the PyC thickness increased from 0.5 to 3.6 µm, the flexural strength noticeably elevated from 1.6 to 17.9 MPa and 1.10–16.00 MPa in $x/y$ direction and $z$ direction, respectively.

(2) The PyC interface presents laminar structure along the carbon fiber surface with different thickness and shows different fracture behavior. The PyC interface with 0.5 µm thickness shows plastic deformation during flexural test. However, for PyC layer with 2.6 and 3.6 µm thickness show pullout behavior due to the thicker of the PyC layers. Additionally, the lower porosity is helpful to the improvement of mechanical properties (as shown in Table 2), for example, the examined mechanical strength of PyC$_{3.6}$ CBCF/ZrB$_2$ is higher than that of PyC$_{2.6}$ CBCF/ZrB$_2$.

4. Conclusions

Carbon-bonded carbon fiber frameworks were modified by PyC coating layer by a CVD process and then the ZrB$_2$ modified PyC$_3$ CBCF was prepared by a PIP method. The PyC interface with a thickness of 0.5, 2.6 and 3.6 µm was obtained by controlling deposition time. Induced PyC interface strongly dominates mechanical properties and fracture behaviors for ZrB$_2$-modified PyC$_3$ CBCF composites.

(1) The mechanical properties of PyC$_3$ CBCF/ZrB$_2$ increase with the thickness of PyC interface. As the PyC thickness increased from 0.5 to 3.6 µm, the flexural strength noticeably elevated from 1.6 to 17.9 MPa and 1.10–16.00 MPa in $x/y$ direction and $z$ direction, respectively.

(2) The PyC interface presents laminar structure along the carbon fiber surface with different thickness and shows different fracture behavior. The PyC interface with 0.5 µm thickness shows plastic deformation during flexural test. However, for PyC layer with 2.6 and 3.6 µm thickness show pullout behavior, and crack deflection and branching occur within the laminar PyC interface.

(3) The carbon fiber has a good compatibility with PyC resulting in strong interaction between fibers and PyC layer, leading to brittle fracture behavior of PyC$_3$ CBCF/ZrB$_2$ composite compared to CBCFs/ZrB$_2$ composite.
Fig. 5. SEM images of fracture surfaces after three-point bending tests. (a) and (b) CBCF/ZrB$_2$; (b) and (c) PyC$_{0.5}$–CBCF/ZrB$_2$; (d) and (e) PyC$_{2.6}$–CBCF/ZrB$_2$; (g) and (h) PyC$_{3.6}$–CBCF/ZrB$_2$. 
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Appendix A. Supplementary material

Supplementary material associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.compositesb.2016.03.094.

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