Characterization of an Ultra-High Temperature Ceramic Composite

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Ultra-high temperature ceramics (UHTC) are of interest for hypersonic vehicle leading edge applications. Monolithic UHTCs are of concern because of their low fracture toughness and brittle behavior. UHTC composites (UHTCC) are being investigated as a possible approach to overcome these deficiencies. In this study a small sample of a UHTCC was evaluated by limited mechanical property tests, furnace oxidation exposures, and oxidation exposures in a flowing environment. The composite was prepared from a carbon fiber perform using ceramic particulates and a preceramic polymer.

The as-received composite plate was non-uniform from front to back surface. Plate dimensions were 150 x 150 x 6 mm. The back surface had a fibrous, uniform appearance; XRD analysis revealed the presence of SiC and C. The front surface was smooth and non-uniform in appearance with evidence of a coarse grain structure produced by a liquid phase; XRD analysis revealed the presence of HfB₂. Microcracks were present throughout the thickness as one might expect from a carbon fiber reinforced composite with attendant large thermal expansion mismatch between the matrix phases and the fibers. The HfB₂ phase on the front surface was comparable in thickness to a fiber ply or about 0.6 mm, and surface microcracks were evident. Limited four point flexural tests were carried out at span to depth ratios of approximately 14 and 16 with markedly different results. Tests were run with the front or the back surface in tension. At the shorter span to depth failures occurred under a loading pin for both orientations. At a span to depth of 16 failures occurred in the center of the span with fracture clearly initiating from a tensile failure. Ultimate flexural strength, strain at ultimate stress, stress and strain at deviation from linear elastic behavior are reported. Strains at ultimate stress ranged from about 0.6 to 0.7 % for the back surface in tension, and 0.4 to 0.6 for the front surface in tension. At constant span to depth the strain at ultimate stress was about 0.2% greater for the back surface in tension and the ultimate strength was also higher. Strengths were in line with predictions from theory.

Furnace oxidation studies were carried out at 1627 and 1927°C in a static furnace environment using ten minute cycles and one, five, and ten cycles. Limited oxidation studies were also carried out in a flowing oxyacetylene torch environment. Specimens were photographed, and weight and dimensional changes were determined. XRD and SEM characterizations were performed. Weight losses were attributed primarily to carbon fiber oxidation. The composite survived the torch test with little visible distress. Further details will be determined once metallographic studies are completed.

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Temperature Ceramic Composite Characterization of an Ultra-High

Outline

- Background
- Objectives
- UHTCC Description
- Results
- -Flexural tests
- -Furnace oxidation
- -Torch tests
- Concluding Remarks

n Temperature Ceramic Composite (UHTCC) Leading Edge	Key Issues Key Issues Fremal stress resistance Oxidation resistance Temperature capability Architecture optimization	Oxidation Resistant Coating Functionally Graded Transition UHTC Composite Core
Ultra High Te (U		

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UHTCC Processing by Starfire

Constituents

Zoltek Panex® 30 Carbon Fabric PW06

Starfire Systems' SP-Matrix Polymer (Allylhydridopolycarbosilane (AHPCS))

•HfB₂ Powder

SiC Powder

Processing of Part Number 000928-6-64

Initial Cycle:

Zoltek cloth is cut into ~6"x6" pieces. 11 plies used

 For the initial lay up the bottom 6 layers of cloth are coated with a SiC/AHPCS slurry and the top 5 layers are coated with a HfB₂ /AHPCS slurry.

The cloth is put into an AI mold and pressed to ~1800lbs. The mold is clamped and cured under inert gas to 400°C

 The plate is removed from the mold and clamped between graphite plates and fired to 850°C under inert gas to pyrolyze.

•Cycle 2:

Coat the HfB, side of the plate with more HfB, /AHPCS slurry.

•Clamp and fire directly to 850°C under inert gas.

Cycle 3: Repeat cycle 2

-Cycle 4 - 10:

Vacuum infiltrate with AHPCS only.

Pyrolyze directly to 850°C under inert gas, no clamping necessary.

As-Received UHTCC Plate



150 mm







As -Received UHTCC

UHTCC Cross-section







Center of UHTCC









UHTCC Specimen Layout



Flexural Strength Tests

Calculated Load	Based on	Beam	Theory*	Z	0011				
Ultimate Load				z	972	757	1025	811	855
Ition	side	down			C/SiC	UHTC	C/SiC	UHTC	UHTC
Orienta	fracture				under pin	under pin	center	center	center
ure	span/	depth			13.8	13.5	15.9	15.8	15.8
st Fixt	outer	span		mm	80	80	96	96	96
Te	inner	span		шш	40	40	48	48	48
cimen	thickness			mm	5.79	5.92	6.04	6.06	6.07
Spe	width			mm	12.7	12.7	12.7	12.7	12.7
					∢	ш	C	Δ	Ш

 Sample B remained intact. All other samples fractured into 2 pieces Samples B, D, and E retained an obvious permanent set

*Calculated load at 0.7% strain
Panex 30 minimum property: E = 193 GPa
Rule of mixtures with no matrix contribution

Flexural Strength Results

	Te	st			Result	S		
-	5	Tensile	ultimate	ultimate	deviation	strain	deviation	strain
Δ	Span /	Side	flexural	strain	from	at dfl	from	at dfl
	Depth		strength	(crossh'd)	linearity		linearity	
					crossheac	l strain	deflector	neter
			MPa	%	MPa	%	MPa	%
۲	4	C/SiC	137.0	0.59	26.4	0.105		
Ш	4	UHTC	102.3	0.39	31.4	0.113		
C	16	C/SiC	161.8	0.73	28.2	0.078	23.0	0.058
	16	UHTC	136.7	0.56	24.7	0.056	26.0	0.059
Ш	16	UHTC	131.8	0.55	26.1	0.090	26.1	0.063







UHTCC After 1627°C Furnace Oxidation









UHTCC Oxy-Acetylene Torch Test

 Sample O: One 4 min. cycle Photos on cool down Temps with Ircon 2 color py 980-1760°C range Weight change Initial weight 41.66 g Final weight 40. 87 g Weight loss 0.79 g, or 1.99 Sample N: Three ~4 min. cyt Cycle 1 max temp 1815°C Cycle 2 max temp 1915°C Cycle 3 max temp 2015°C Cycle 3 max temp 2015°C Weight change Photos on heat up Photos on heat up Initial weight 41.99 Final weight 40.04 	to 1805°C Sample O	Time, Temp, °C	rometer, min.	0.5 1720	1.0 1750	1.5 1750	6 2.0 1755	cies 2.5 1765	3.0 1775	3.5 1790	4.0 1805			0	
	Sample O: One 4 min. cycle	 Photos on cool down 	 Temps with Ircon 2 color py 	980-1760°C range	 weight unange Initial weight 41.66 g 	Final weight 40. 87 g	• Weight loss 0.79 g, or 1.9%	Sample N: Three ∼4 min. cy(1 max temp 1815°C	 Cycle 2 max temp 1915°C 	 Cycle 3 max temp 2015°C 	Photos on heat up	 Weight change Initial weight 41.99 	 Final weight 40.04 	 Weight loss 1.95 g, or 4.6% 	

UHTCC Torch Test Video



UHTCC Torch Test Cool Down



30 sec

UHTC Surface After Three 4-Minute Torch Cycles to 1815 to 2025°C





SEM of Center of UHTC Surface After Three Torch Cycles



1 mm



	Conclusions
•	 Processing Uniform and through thickness graded microstructure achieved
	 Matrix cracking due to thermal expansion mismatch between C fibers and matrix constituents is a concern
٠	Mechanical Properties
	 Flexural strength was close to expected values based on rule of mixtures with no matrix contribution
	 Some evidence of composite behavior
٠	Furnace Oxidation
	 Based on weight loss, carbon fiber oxidation occurred rapidly
•	Torch Test
	 Material withstood ~2000°C (~3600°F), severe heat-up and thermal gradients with no major visible distress
	 Based on observed temperature spikes during test, adherence of the HfO₂-rich scale is an area of concern

Recommendations

- The thermal stress response of this early UHTCC makes the concept worthy of further study
- Fiber coatings need to be incorporated to address fiber oxidation issues
- Advanced SiC fibers need to be evaluated to address oxidation and thermal expansion mismatch issues

Future Work

- Continue UHTCC development
- Continue UHTCC evaluation
- Complete metallography on Starfire specimens
- Evaluate other NASA and industry developed materials

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- Thanks to Terry R. McCue for scanning electron microscopy support and Ronald E. Phillips for assistance with testing.
- Thanks to David Glass of NASA Langley for providing the Starfire UHTCC plate.
- providing processing details and for permission to Thanks to Walt Sherwood of Starfire Inc. for present this study.