Initial Assessment of Environmental Effects on SiC/SiC composites in Helium-cooled Nuclear Systems

September 2013

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INITIAL ASSESSMENT OF ENVIRONMENTAL EFFECTS ON SiC/SiC COMPOSITES IN HELIUM-COOLED NUCLEAR REACTOR SYSTEMS

Cristian I Contescu

Date Published:

September 2013

Prepared by
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Oak Ridge, Tennessee 37831-6283
managed by
UT-BATTELLE, LLC
for the
U.S. DEPARTMENT OF ENERGY
under contract DE-AC05-00OR22725
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Impurity content in helium-cooled high temperature reactors

LIST OF ABBREVIATIONS

1D     uni-dimensional
2D     two-dimensional
3D     three-dimensional
C/C    carbon fiber / carbon matrix (composites)
CVD    chemical vapor deposition
CVI    chemical vapor impregnation
EBC    Environmental Barrier Coating
FHR    Fluoride-cooled High-temperature Reactor
GFR    Gas Fast Reactor
HCSB   Helium Cooled Solid Breeder
HTGR   High Temperature Gas-cooled Reactor
HTR    High Temperature Reactor
HTTR   High Temperature Test Reactor
LWR    Light Water Reactor
NGNP   Next Generation Nuclear Plant
NITE   nano-infiltration and transient eutectic (process)
ML     multilayer
PIP    Precursor Infiltration and Pyrolysis (method)
ppm    parts per million
PyC    pyrolytic carbon (aka pyrocarbon)
SFR    Sodium Fast Reactors
SiC    silicon carbide
VHTR   Very High Temperature Reactors
ABSTRACT

Ceramic matrix composites, in particular SiC/SiC composites, are currently being evaluated for use at high temperatures and high radiation fields in advanced concept reactors, including high-temperature gas cooled reactors (HTGR), very high temperature reactors (VHTR), gas fast reactors (GFR), sodium fast reactors (SFR) and fluoride-cooled high temperature reactors (FHR). SiC/SiC composites have many desirable properties for high temperature nuclear applications, including excellent thermal and mechanical properties and radiation resistance. However, their behavior in contact with the chemical environment in normal operation conditions and in off-normal accident conditions scenarios still needs to be evaluated. This report summarizes the information available in the literature on the chemical reactivity of SiC/SiC composites and of their components in contact with the helium coolant used in HTGR, VHTR and GFR designs. Under normal operating conditions, ultra-high purity helium will have chemically controlled impurities (water, oxygen, carbon dioxide, carbon monoxide, methane, hydrogen) that will create a slightly oxidizing gas environment. Little is known from direct experiments on the reactivity of third generation (nuclear grade) SiC/SiC composites in contact with low concentrations of water or oxygen in inert gas, at high temperature. However, there is ample information regarding the oxidation behavior of SiC/SiC composites in dry and moist air at high temperatures. This information is reviewed first in the next chapters. The emphasis is placed on the improvement in material oxidation, thermal, and mechanical properties during three stages of development of SiC fibers and at least two stages of development of the fiber/matrix interphase. The chemical stability of SiC/SiC composites in contact with oxygen or steam at temperatures that may develop in off-normal reactor conditions supports the conclusion that most advanced composites (also known as nuclear grade SiC/SiC composites) have the chemical resistance that would allow them maintain mechanical properties at temperatures up to 1200 – 1300 °C in the extreme conditions of an air or water ingress accident scenario. Further research is needed to assess the long-term stability of advanced SiC/SiC composites in inert gas (helium) in presence of very low concentrations (traces) of water and oxygen at the temperatures of normal operation of helium-cooled reactors.
1. SiC/SiC COMPOSITES AS NUCLEAR STRUCTURAL MATERIALS

Ceramic materials are strong candidates for internal core structural components in the next generation of nuclear energy systems. Several reactor concepts are currently being analyzed; instead of pressurized water, they use for coolant either molten salts, or gaseous helium, or liquid sodium. The candidate composites are continuous fiber-reinforced materials, made of either carbon or silicon carbide fibers infiltrated with a similar matrix. While carbon-fiber composites are now a mature technology and a readily-available commercial product, the technological development of SiC/SiC composites for commercial scale applications still continues. However, the third generation of SiC/SiC composites, qualified as nuclear grade, has the important advantage of being far more stable to irradiation degradation. Their high resistance to intense irradiation condition is an inherent property of the isotropic SiC crystal, which undergoes only modest property changes (swelling, hardness, elastic modulus, strength) in the range of temperatures of interest for high-temperature reactors. In contrast, carbon–fiber composites undergo densification parallel to the fiber axis and swelling on the perpendicular direction, which is caused by the growth of new basal planes from carbon atoms displaced by neutrons in interstitial positions. This difference makes nuclear grade SiC/SiC composites the preferred material over C/C composites for components designed to withstand long-time exposure to intense radiation and elevated temperatures.

1.1 TYPES, PROPERTIES, APPLICATIONS

The attractive mechanical properties of SiC/SiC composites (tensile strength, elastic modulus, creep resistance) originate from the reinforcing fibers. In addition, the temperature limits depend on the stability of component fibers in an inert atmosphere or in vacuum under high temperature conditions. Bonding of fibers to the matrix is ensured by a thin layer of compliant material, generally known as the interphase, which enhances the non-brittle mechanical behavior. Pyrocarbon is commonly used as the interphase.

The enhanced radiation resistance of SiC/SiC composites is the unique property of newly developed nuclear-grade SiC fibers, also known as the third generation fibers. Early grades of SiC fibers contained either too much oxygen (first generation) or too much carbon (second generation). Thus, the Nicalon fibers of the first generation have relatively low price, good mechanical properties, and excellent weavability, but their composition should be described as Si-Oₓ-Cᵧ rather than SiC. Second generation Hi-Nicalon has less than 1% oxygen. The third generation fibers are nearly stoichiometric, having only a low excess of carbon in the bulk and no residual oxygen, although the surface might be covered with a thin carbon layer. Some fibers contain various amounts of additives, such as aluminum (Tyranno SA) or boron (Syrlamic); Hi-Nicalon S (from Nippon Carbon) contains larger amounts of impurities. These third generation fibers have a coarse microstructure and high density, are stable up to 1800-2000 °C and maintain good mechanical properties up to 1400-1500 °C, although SiC grains may grow larger at 2100...
°C. Tyranno-SA3 and Sylramic have even higher upper temperature limit (retain strength up to 1800 °C), but also higher thermal creep rate at low temperatures.

The properties of SiC matrix depend on the fabrication route. Most common routes, compatible with the nuclear environment, are chemical vapor impregnation (CVI) and nano-infiltration and transient eutectic-phase process (NITE). Both these methods produce highly crystalline, stoichiometric matrices and can be fabricated in a variety of shapes. The CVI process, which in fact is a variety of chemical vapor deposition (CVD) of pyrolytic SiC, allows deposition of high purity, highly crystalline β-SiC phase, with excellent thermal stability and radiation resistance, but with limited matrix densification.²

The decomposition of SiC occurs at high temperatures (>2500 °C):

\[ \text{SiC(s)} \rightarrow \text{C(s)} + \text{Si(C)}(\text{l}) \]  

The gas phase in equilibrium with SiC at temperatures below the melting point contains Si(g) and SiC₂(g). Thus, exposure to high temperatures under an inert environment induces thermal degradation. However, kinetic factors limit this process, so that thermal decomposition of SiC occurs only at temperatures above 1600 °C, before melting.

Nuclear grade SiC/SiC composites are considered candidates for components of light-water reactor (LWR) fuel cladding and channel boxes; for fuel and in-vessel components for advanced fission reactors such as high-temperature gas-cooled reactors (HTGR); for fluoride salt-cooled high temperature reactors (FHR); for gas-cooled fast reactors (GFR), and also for fusion energy and other extreme irradiation conditions. In the U.S. the reference SiC/SiC material for the NGNP program is the composite made with Hi-Nicalon S fiber and CVI SiC matrix, with an interphase made of pyrolytic carbon and SiC.² Several U.S. companies are known to manufacture SiC/SiC composites by CVI. The list includes GE Energy, BF Goodrich, ad Hypertherm High-Temperature Composites (now Rolls-Royce), as well as other small businesses capable of manufacturing nuclear grade SiC/SiC CVI composites.

1.2 ENVIRONMENT CONDITIONS IN HELIUM-COOLED HIGH TEMPERATURE REACTORS

The coolant helium gas in high temperature reactors (HTRs) is expected to contain low level gas impurities. Of concern during normal operating conditions are oxidant species such as H₂O, O₂, and CO₂ that may cause slow gasification over the lifetime of components. Other impurities (H₂, CO) have reducing character, and some others are neutral (N₂, CH₄). Methane results from reaction of carbon with H₂, but this gasification reaction is not probable at temperatures above 600 °C. If however, methane is formed, it is a product of radiolytic processes. Other radiolytic processes may lead to formation of tritium and of some reactive ionized species.

Table 1 shows coolant gas impurities in historic high- and very high temperature gas-cooled reactors (HTGR and VHTR) and in modern reactors operated in Japan and China. Depending on design the coolant temperature (output) varies between 750 °C (Dragon, Peach Bottom) and 900-950 °C (HTTR, several NGNP design concepts). The nominal operating temperature in gas-cooled fast reactors (GFR) is even higher (about 1000 °C).³ The main technological challenge is development of materials able to resist the damage caused by fast neutron fluences at high temperatures.

SiC is an excellent candidate for these extreme requirements. SiC/SiC composites are recommended for components subjected to high fluence conditions or where anisotropic changes in dimension and
elastic properties cannot be tolerated. Such components may be control rod sheath and joints, core supports, core restraint, etc. for which the required stiffness, high temperature strength, stress rupture life, or irradiation performance are not met by the existing engineered alloys. All designs considered in the U.S. (pebble bed design by Westinghouse and the prismatic reactor designs by AREVA and General Atomics) consider the use of SiC/SiC composites for components in reactivity control system, such as the control rods and shutdown rods. This selection is motivated by the fact that SiC/SiC is the only available material that can withstand the high fluence (up to 30 dpa) and temperature (1100 °C in normal conditions, 1600 °C in off-normal conditions) required for these components.

Table 1  Impurity content in helium-cooled high temperature reactors

<table>
<thead>
<tr>
<th></th>
<th>units</th>
<th>H₂O</th>
<th>H₂</th>
<th>CO</th>
<th>O₂</th>
<th>N₂</th>
<th>CO₂</th>
<th>CH₄</th>
</tr>
</thead>
<tbody>
<tr>
<td>DRAGON, UK (1964-75)⁴</td>
<td>Pa</td>
<td>0.1</td>
<td>2</td>
<td>1.2</td>
<td>n/a</td>
<td>0.3</td>
<td>&lt;0.04</td>
<td>0.3</td>
</tr>
<tr>
<td>Peach Bottom, USA (1967-74)⁵</td>
<td>ppmv</td>
<td>0.5</td>
<td>10</td>
<td>0.5</td>
<td>n/a</td>
<td>0.5</td>
<td>&lt;0.5</td>
<td>1</td>
</tr>
<tr>
<td>AVR, Germany (1967-88)⁴</td>
<td>Pa</td>
<td>3</td>
<td>30</td>
<td>10</td>
<td>n/a</td>
<td>n/a</td>
<td>10</td>
<td>n/a</td>
</tr>
<tr>
<td>Fort St. Vrain, USA (1976-79)⁵</td>
<td>ppmv</td>
<td>&lt;1</td>
<td>2</td>
<td>7</td>
<td>1--10</td>
<td>n/a</td>
<td>n/a</td>
<td>0.5--3</td>
</tr>
<tr>
<td>HTR-10, China (2003)⁶</td>
<td>ppmv</td>
<td>&lt;0.2</td>
<td>&lt;3</td>
<td>&lt;3</td>
<td>n/a</td>
<td>&lt;1</td>
<td>n/a</td>
<td>&lt;1</td>
</tr>
<tr>
<td>HTR-PM, China (project)⁷</td>
<td>ppmv</td>
<td>2</td>
<td>30</td>
<td>30</td>
<td>0.2</td>
<td>2</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>HTTR, Japan (1998-present)⁸</td>
<td>ppmv</td>
<td>&lt;0.2</td>
<td>&lt;3</td>
<td>&lt;3</td>
<td>n/a</td>
<td>&lt;0.2</td>
<td>&lt;0.6</td>
<td>&lt;0.5</td>
</tr>
<tr>
<td>General Atomics (concept)²</td>
<td>ppmv</td>
<td>0.5</td>
<td>3</td>
<td>2</td>
<td>n/a</td>
<td>2</td>
<td>1</td>
<td>0.1</td>
</tr>
</tbody>
</table>

Different off-normal (accident) scenarios have been envisioned. Although not probable, the behavior of materials in those extreme conditions should be known. VHTR reactors are protected by passive safety features, and the most serious accident for this reactor design is air-ingress followed by a large pipe break. Fast oxidation of graphite in the reflector and moderator would generate a large amount of heat that would increase the temperature of the core. However, various analysis results show that even under severe chemical reacting conditions, the maximum core temperature might rise up to 1430 °C but would not reach the maximum temperature criteria (1600 °C).⁹,¹⁰

Based on the above information, the environment conditions that must be analyzed for assessing the stability of SiC/SiC composites in VHTR are as follows:

Normal operation conditions:
- helium coolant with ppm-range impurities (H₂O, CO₂, O₂, H₂, CH₄) – see Table 1
- temperature 700 – 950 °C

Off-normal conditions:
- air or water ingress
- temperature < 1500 °C

The general concern is about gas impurities with oxidizing character (H₂O, CO₂, O₂) that may affect the interphase layer of SiC/SiC composites. Water and oxygen, in particular, have several reaction channels with SiC, depending on temperature and gas phase composition. Reducing impurities (CO, H₂)
are of little concern for the normal operation conditions. The helium coolant in HTGR and VHTR will have a slightly oxidizing character in order to protect metallic components against carburization and/or embrittlement. However, at some extreme conditions (high pressures of H₂ or CO) gasification of SiC may also occur. A special case is the condition of low oxygen environment at high temperatures, as it is expected to be the gas coolant atmosphere in HTGR. The reactivity of various components in SiC/SiC composites is presented in the next chapter.
2. OXIDATION OF SiC/SiC COMPONENTS

Oxidation of all silicon-rich materials, either pure silicon or silicon-based ceramics, occurs according to the same basic mechanisms: in a plentiful supply of oxygen oxidation occurs with the formation of a passive, protecting silica (SiO$_2$) surface layer (*passive oxidation*); however, with a limited supply of oxidant, severe corrosion occurs through oxidative volatilization and release of SiO gas phase (*active oxidation*). This pattern will be discussed below, separately for each of the components of SiC/SiC composites: monolithic crystalline SiC material, SiC fibers, and the interphase layer.

### 2.1 OXIDATION OF MASSIVE SiC

#### 2.1.1 Passive Oxidation in Presence of Oxygen

Massive SiC oxidized in air at high temperatures produces a thin, adherent surface silica layer through the following reactions:

\[
\begin{align*}
\text{SiC(s)} + \frac{3}{2} \text{O}_2(\text{g}) &= \text{SiO}_2(\text{s}) + \text{CO(}g) \quad (2) \\
\text{SiC(s)} + 2 \text{O}_2(\text{g}) &= \text{SiO}_2(\text{s}) + \text{CO}_2(\text{g}) \quad (3)
\end{align*}
\]

The growth rate of the silica layer (in terms of the scale thickness, $x$, or mass gain) follows a linear-parabolic law with two rate constants, $K_P$ and $K_L$, for the parabolic and linear regime:

\[
\frac{K_P}{x^2} + \frac{K_L}{x} = t + \tau
\]

where $\tau$ is a constant that accounts for the initial scale thickness. The parabolic term capturing the diffusion-limited kinetics becomes the main contributor at long oxidation times while the linear term represents reaction-limited kinetics and is important during early stages. Formation of the passive SiO$_2$ layer explains the outstanding oxidation resistance of silicon-based ceramics.

Oxidation rates of crystalline $\alpha$-SiC are limited by the transfer of oxygen through the scale. Below about 1400 °C oxygen permeates in molecular form, and the process is independent of oxygen pressure, $P_{O_2}$ (activation energy $E_a = 120$ kJ/mol). At higher temperatures, atomic oxygen permeates the scale and the rate is proportional with ($P_{O_2}$)$^n$. A higher activation energy (200 -300 kJ/mol) reflects the energy needed to dissociate the oxygen molecules.12

Polycrystalline SiC follows a similar oxidation mechanism, but the characteristics vary depending on the SiC purity and the degree of crystallization. Activation energies reported by various authors are spread from 120 to 420 kJ/mol. Additives in SiC and impurities in the gas phase increase, in general, the rate of oxidation. Crystallization of silica changes the oxidation kinetics by changing the diffusion mechanism of CO and oxygen.13

#### 2.1.2 Active Oxidation at Low Oxygen Concentrations

In an environment with low oxidizing potential, below a critical $P_{O_2}$, silica is no longer stable, as a new oxidation reaction replaces the reaction paths illustrated by equations (2) and (3):
\[
\text{SiC}(s) + O_2(g) = \text{SiO}_2(g) + \text{CO}(g)
\] (5)

Conversion of SiC into gaseous products is recognized by the accompanying mass loss. The transition between active and passive oxidation has been discussed by several investigators.\textsuperscript{14} The rate and the transition temperature depend on factors such as SiC purity, temperature, gas flow composition and velocity, total pressure, and oxygen partial pressure.\textsuperscript{15,16,17}

### 2.1.3 Effect of Water Vapor and Other Gas Environments

The passive mode of oxidation by water vapor occurs through the following reactions:

\[
\text{SiC}(s) + 2 \text{H}_2\text{O}(g) \rightarrow \text{SiO}_2(s) + \text{CH}_4(g) \quad \text{at } T < 1400 \degree \text{C} \tag{6}
\]

\[
\text{SiC}(s) + 3 \text{H}_2\text{O}(g) \rightarrow \text{SiO}_2(s) + \text{CO}_2(g) + 3 \text{H}_2(g) \quad \text{at } T > 1400 \degree \text{C} \tag{7}
\]

Presence of water vapor in oxygen (or air) enhances the rate of passive oxidation, from about 2-3 times (when 10 % H\textsubscript{2}O is added to oxygen) to about 10-20 times (when comparing oxidation in steam versus dry oxygen).\textsuperscript{18} The rate enhancement is explained by water having higher affinity to silica than oxygen, and thus higher permeability through the scale SiO\textsubscript{2} layer. In addition, water can break Si-O bonds, which enhances diffusion. For these reasons, the linear term dominates in the linear-parabolic kinetics shown by equation (4) above.\textsuperscript{19}

In a water-rich atmosphere (steam), gasification of the protective silica layer may occur with formation of gaseous SiO(OH)\textsubscript{x} volatile species, according to the following reactions:

\[
\text{SiO}_2(s) + \text{H}_2\text{O}(g) = \text{SiO(OH)}_2(g) \tag{8}
\]

\[
\text{SiO}_2(s) + 2 \text{H}_2\text{O}(g) = \text{SiO(OH)}_4(g) \tag{9}
\]

for which the rate is proportional with \(P_{\text{H}_2\text{O}}\) and \((P_{\text{H}_2\text{O}})^2\) respectively. Therefore the higher the moisture content, the more important is the weight loss of the passive SiO\textsubscript{2} layer.\textsuperscript{20}

In a mixed \text{H}_2\text{O}/\text{H}_2\text{ environment, oxidation of SiC follows an active mechanism at } 1300 – 1400 \degree \text{C} according to the following reactions:

\[
\text{SiC}(s) + \text{H}_2\text{O}(g) = \text{SiO}(g) + \text{CH}_4(g) \tag{10}
\]

\[
\text{SiC}(s) + 2 \text{H}_2\text{O}(g) = \text{SiO}(g) + \text{CO}(g) + 2 \text{H}_2(g) \tag{11}
\]

In presence of high \(P_{\text{H}_2}\) gasification may occur through reduction of the silica layer, according to:

\[
\text{SiO}_2(s) + \text{H}_2(g) = \text{SiO}(g) + \text{H}_2\text{O}(g) \tag{12}
\]

Conditions of active oxidation at low oxygen partial pressure may also occur in presence of \text{CO}_2/\text{CO} mixtures. It was reported that active oxidation occurs at \(P_{\text{CO}_2}/P_{\text{CO}}\) ratios in the range of \(10^{-4}\) to \(10^{-1}\) and the mass loss of SiC is maximum at intermediate ratios.\textsuperscript{21} Active volatilization of SiC may also occur at low partial pressures of \text{CO}_2.\textsuperscript{22}
2.2  OXIDATION OF SiC MATRIX

In order to understand the behavior of SiC/SiC composites in an oxidative atmosphere, the knowledge of oxidation kinetics of the SiC matrix (formed under specific conditions) is as important as the knowledge of oxidation kinetics of crystalline SiC materials or of SiC fibers. A comparison between the oxidation kinetics of SiC deposits formed by CVI from CH$_2$SiCl$_3$/H$_2$ mixture and that of pure silicon (selected as standard) and other SiC ceramics showed that the behavior of CVI β-SiC under pure oxygen follows the characteristics of passive oxidation at temperatures in the 900 – 1500 °C range. The surface layer grows according to the parabolic law, and the protective silica layer is amorphous except after treatment at 1500 °C. Figure 1 shows crystobalite crystals formed from SiC (CVI) maintained at 1500 °C in a flow of pure oxygen at 100 kPa.

In conditions of VHTR or GFR, the partial pressure of oxygen in helium coolant is much lower, and the investigation of passive to active transition at much lower $P_{O_2}$ was needed for understanding of SiC behavior at the high temperatures that may occur in accident conditions. Charpentier et al. measured oxidation kinetics of sintered α-SiC and CVD β-SiC at oxygen pressures between 0.2 and 100 Pa in He (total pressure 100 kPa) and a broad temperature range. The α–SiC was obtained from Boostec (France) and was processed by pressureless sintering at ~ 2000 °C with B$_4$C as additive. The β–SiC was obtained from Rohm & Haas (USA) and was processed by CVD at 1200-1300 °C. The authors cited above found that the transition between passive and active...
oxidation regime occurs at higher temperatures for β-SiC than for α-SiC, and the mass loss rate of β-SiC is lower than the one measured for α-SiC on the common temperature range investigated (up to 1800 °C). Figure 2 show theoretical and experimental transitions observed for the two SiC allotropes. Oxygen partial pressure has no effect on the mass loss rate of β-SiC in active oxidation conditions up to $P_{O_2} = 10$ Pa in this temperature range, contrary to the behavior of α-SiC. This is an important conclusion, showing that high purity helium is not necessary to prevent β-SiC from severe oxidation up to 1800 °C. This temperature is way above the normal operation range of VHTR and GFR, and is improbable even in severe accident conditions. Severe damage of β-SiC occurred at ~ 2000 °C and $P_{O_2} = 2$ Pa caused by combined active oxidation and sublimation of SiC. At these extreme conditions, the mass loss rate was ~ 30 mg cm$^{-2}$ h$^{-1}$.

Figures 3 and 4 show selected SEM images$^{3,24}$ of oxidized surfaces of sintered α- and crystalline β-SiC exposed to 0.2 and 100 Pa oxygen in helium. The temperatures where active to passive oxidation
occurs vary, depending on the crystalline form and the oxygen pressure. These images correspond to several experimental data points shown in Fig. 2.

### 2.3 OXIDATION OF SiC FIBERS

Oxidation of SiC fibers follows the general behavior of pure (crystalline) SiC, and exhibits the same transition between passive and active oxidation. However, SiC fibers may also contain oxycarbide (first generation), free carbon (second generation) or additives (some of the third generation) which may influence the oxidation properties.13

Oxidation at 1500 °C of Nicalon CG fibers (Si-C-O system, first generation) was active at 100 Pa O₂ and transitioned to passive oxidation at 250 Pa. At higher P₀₂ oxidation was passive, with mass gain caused by the growth of silica layer. This impermeable silica layer formed by passive oxidation decreased thermal decomposition by preventing gaseous products from escaping the fiber.25

Passive oxidation of Hi-Nicalon fibers (second generation) followed a parabolic law, with radial silica growth. The active to passive transition occurred at 10-25 Pa at 1500 °C. The silica layer improved the high temperature behavior. Hi-Nicalon fibers did not keep any residual strength when exposed to Ar at 1500 °C, but pre-oxidation at 800-1500 °C enabled a quite high tensile strength.26 27

In the third generation fibers (stoichiometric), Hi-Nicalon S, showed the transition from passive to active oxidation at 1500 °C under very low partial pressure of oxygen, and Tyranno SA fibers did not undergo active oxidation above 1 Pa O₂. Thus, the resistance against active oxidation has continuously improved from first generation fibers to the third (Fig. 5).25

When 20 kPa water was present in O₂-Ar mixtures (20 kPa O₂), oxidation rates of Hi-Nicalon fibers increased dramatically (about 20 times at 1000 °C) as in the case of massive SiC, and the activation energy decreased. This shows that water can act as the main oxidizing agent according to reactions (8-10) above.28

![Figure 5: Oxygen partial pressure for active-to-passive oxidation transition of SiC as a function of temperatures, showing continuous improvement of oxidation resistance over three generations of SiC fibers (Nicalon, Hi-Nicalon, and Hi-Nicalon S). From reference 25.](image-url)
2.4 OXIDATION OF THE INTERPHASE

All carbons, including pyrocarbon, are sensitive to oxidation by oxygen, carbon dioxide and water, according to the following reactions:

\[ C(s) + O_{2(g)} = CO_{2(g)} \]  (13)
\[ C(s) + \frac{1}{2} O_{2(g)} = CO_{(g)} \]  (14)
\[ C(s) + CO_{2(g)} = 2 CO_{(g)} \]  (15)
\[ C(s) + H_{2O(g)} = CO_{(g)} + H_{2(g)} \]  (16)

However, pyrocarbon is known to have higher chemical resistance to oxidation than other forms of carbon. The lower oxidation rates observed by pyrocarbon are caused by kinetic factors (limited in-pore diffusion caused by extremely low porosity). The SEM image in Fig. 6 show the cross-end of a T300 carbon fiber tows (not heat-treated) infiltrated with a pyrocarbon interphase after mild oxidation at 550 °C in pure oxygen. The superior resistance of pyrocarbon is evident. However, when the same carbon fibers were heat-treated at 1600 °C to improve their structural order, they showed similar oxidation behavior as the pyrocarbon interphase.²⁸

Figure 6: SEM image of pyrocarbon-infiltrated carbon fiber tow showing that pyrocarbon has higher oxidation resistance. From reference ²⁸.
3. THERMAL AGEING AND OXIDATION OF SiC/SiC COMPOSITES

Fibers and interphase layers play a major role in the mechanical properties of composites. Because composites are designed to form micro-cracks upon excessive loading, and fibers and interphase are sensitive to oxidation, oxidizing species can access the inner components and oxidation gaseous products can escape, causing weight loss and structural damage in composites. Improving the lifetime of composites requires developing protection methods to mitigate fibers thermal instability and the interphase preferential oxidation.13

3.1 THERMAL AGEING

The thermal stability in vacuum of SiC/SiC composites prepared with chemical vapor infiltration has improved considerably from first to the third generation of SiC fibers. For example, a parallel examination of structural and mechanical properties of composites prepared from commercial grade Nicalon, Hi-Nicalon, and stoichiometric Hi-Nicalon type S after heat treatment in vacuum (1 hour at 1000 – 2000 °C) has shown important stability differences.29

Figure 7. Microstructure of SiC/SiC composites formed with SiC fibers of three generations, after ageing in vacuum at high temperatures. The SiC fibers were Nicalon commercial grade (left column), Hi-Nicalon (center column) and Hi-Nicalon-S (right column). From reference 29.
The oxygen-rich Nicalon fibers of the first generation lost volatile SiO in the gas phase according to the reaction shown below, starting from about 1400 °C:

\[(\text{SiC, SiC}_{2x\text{O}_{1-x}}) \rightarrow w \text{SiC}_{}(s) + x \text{C}_{}(s) + y \text{SiO}_{}(g) [+ z \text{C}_{}(s)] \quad (17)\]

The oxycarbide SiC\(_{2x\text{O}_{1-x}}\) decomposed completely above 1700 °C, which caused a drop in flexural strength and fracture toughness. Significant growth of SiC grains occurred simultaneously,\(^{30}\) as a side reaction between SiO in the gas phase and the carbon interphase:

\[\text{SiO}_{}(g) + 2 \text{C}_{}(s) = \text{SiC}_{}(s) + \text{CO}_{}(g) \quad (18)\]

In the same conditions, the second generation fiber Hi-Nicalon experienced just a slight degradation of bending properties,\(^{29}\) caused by the reaction

\[\text{SiC}_{1+x}(s) + \text{O}_{\text{trace}}(g,s) \rightarrow \text{SiC}_{}(s) + \text{CO}_{}(g) \quad (19)\]

In contrast, the composite with the stoichiometric Hi-Nicalon S fibers (third generation) suffered no significant change in bending properties after thermal treatment. The composites made with these latter fibers did not suffer any apparent deterioration after heat treatment up to 1900 °C in vacuum. The microstructural changes suffered by the three types of composites are illustrated by the microscopy images in Fig. 7.\(^{29}\) Composites made with Tyranno-SA fibers and CVI SiC matrix maintained high strength up to 1700 °C. Above this temperature, the strength decreased because of CVI matrix recrystallization and degradation of the Tyranno-SA fibers.\(^{31}\)

### 3.2 OXIDATION

#### 3.2.1 Effect of Oxygen

When SiC/PyC/SiC composites with machined surfaces or open cracks are exposed to environmental conditions that promote passive oxidation of SiC (e.g. 100 kPa O\(_2\), 700 °C < T < 1400 °C) the attack on the composite is initiated at the interphase layer, which is sensitive to gasification according to equations (13) – (15). It is believed\(^ {32}\) that the mechanism involves the steps of (1) oxidation of carbon interphase by direct reaction with oxygen and formation of a pore in the structure; (2) oxygen and carbon oxides diffusion along the pore; (3) reaction of oxygen with the pore walls leading to the growth of a silica layer on both the fiber and the matrix, according to reactions (2) and (3) above. If the interphase is thin enough, silica is able to seal the pore and the active oxidation process ends. However, if the pyrocarbon interphase layer is thicker, the pore formed is larger and the reactions (13) – (15) proceed until the carbon layer is entirely consumed.

Oxidation in air at intermediate temperatures (600 – 800 °C) was shown to consume the interphase PyC layer, and cause stronger bonding between fibers and matrix by the SiO\(_2\) layer formed at 800 °C. Both effects lead to decreased cyclic fatigue life, but the strong fiber-matrix bonding by the silica layer leads to the shortest fatigue life.\(^ {33}\)
When exposed to oxygen at 1000 °C, SiC/PyC/SiC composites experience first a mass loss attributed to oxidation of the pyrocarbon layer accompanied by recession of the interphase, followed by a parabolic weight gain that indicates development of the silica layer. The recession of thick interphase layer occurs between 800 – 1200 °C after either a linear or parabolic kinetic law, where the reaction order for oxygen is between 0.5 and 1.\textsuperscript{13} Gasification of the interphase impacts considerably the mechanical strength of the composite and cause embrittlement. Comparing TGA curves during oxidation with in situ Young’s modulus measurements of a SiC\textsubscript{f}(CG)/PyC/CVI-SiC composite revealed that interphase recession causes a sharp decrease of modulus, which may recover eventually if the newly formed silica layer closes the gap between fibers and matrix.\textsuperscript{34}

Degradation of SiC/SiC composites follows a different mechanism when exposure to high temperatures occurs in air or in vacuum.\textsuperscript{35} Figure 8 shows weight changes versus exposure time at different temperatures. In this example, the fiber produced by Ube Companies was of ZMI type, with a non-stoichiometric composition described as SiZr\textsubscript{-0.01}C\textsubscript{1.44}O\textsubscript{0.32}. The large weight loss observed in vacuum at 1200 – 1400 °C is caused by decomposition of the fiber and loss of gaseous products, SiO\textsubscript{(g)} and CO\textsubscript{(g)}. In air, a slight weight gain is observed which indicates formation of silica protective layer. The fracture mechanism was also different: the vacuum-exposed composite showed only the fiber-pullout type fracture, while the fracture mode of air-exposed composites changed with the progress of oxidation from the fiber-pullout type to the nonfiber-pullout type.\textsuperscript{35}

Nearly stoichiometric SiC fibers of the third generation are more stable than those of generation one or two. The oxidation behavior in air of Tyrannohex\textsuperscript{TM} composite containing Tyranno\textsuperscript{TM} Si-C-Al-O fibers is different from that of Al-free fibers, which is an effect of the different crystallinity of the SiO\textsubscript{2} scale.\textsuperscript{36} During heating, an amorphous SiO\textsubscript{2} scale was formed which hosted gas bubbles generated by the gaseous products of oxidation. This caused a large weight loss after 4000 h exposure to 1500 °C in spite of the conditions favoring passive oxidation. The large weight loss was caused by destruction of the oxidation-protective scale by bursting SiO\textsubscript{2} bubbles, along with higher diffusivity of oxygen in amorphous SiO\textsubscript{2} scale containing aluminum ions.

Controlling the matrix crystallinity in SiC/SiC composites made with third generation fibers (Tyranno\textsuperscript{TM} SA) by the polymer impregnation and pyrolysis (PIP) method showed good results not only on mechanical properties, but also on oxidation resistance in air.\textsuperscript{37} The improvement was obtained through reduction of impurity content in the matrix and better organization of the crystal structure through annealing in argon at 1500 – 1700 °C. Oxidation resistance was improved by the two-layered interphase (100 nm PyC and 500 nm SiC). The superior flexural properties achieved by this treatment did not significantly degrade in air up to 1000 °C and mechanical properties remained stable up to 1300 °C in air. Conventional SiC/SiC composites showed a slight reduction of flexural strength in the same conditions,
caused by oxidation of impurities in constituents, such as the Tyranno-LoxM fiber with about 10 wt % oxygen.37

3.2.2 Effect of Water

The effect of water on SiC/SiC composites was studied intensively in relation with the use of these materials in gas turbines where they are exposed to high temperature steam. Oxidation of SiC by steam according reactions (6) and (7) above forms a SiO₂ layer. In presence of H₂O, however, this layer no longer protects the underlying SiC fiber and gasification continues according to reactions (8) and (9) above, controlled by the diffusion of volatile species Si-Oₓ-Hᵧ(g) through a porous boundary layer. This is illustrated by the electron microscopy images presented in Figure 9, which also compare the surface of α– and β-SiC after exposure to high pressure steam in air, or dry air only.38

Figure 9: (Left): Typical silica scale morphology observed for CVD β-SiC after 500 h exposure to 10 bar air + H₂O at 1200 °C; (Right): Silica scale on sintered β-SiC after 100 h exposure to 10 bar of (a) air and (b) air + H₂O at 1200 °C. The arrows show the dense silica layer. Only this layer was formed when no H₂O was present. From reference 38.

The same mechanism applies to SiC/SiC composites. Oxidation of 1D SiC/SiC composites is enhanced when water is present in the oxidant gas. Compared with oxidation in dry air at 1520 °C, oxidation in air saturated with water vapor at 25 °C is up to 50 % higher. Even when oxygen is not present
in the gas phase, water alone causes oxidation. Thus, oxidation of the same 1D SiC/SiC composite in argon saturated at 25 °C with water vapor was reported to cause half of the oxidation effects observed in dry air.\textsuperscript{39}

A similar result was obtained by comparing the behavior of SiC/PyC/SiC composites (made with Hi-Nicalon\textsuperscript{TM} fiber and CVD SiC matrix) during oxidation in dry air and in a combustion gas containing 13 % H\textsubscript{2}O and 33 % CO\textsubscript{2}.\textsuperscript{40} Compared with air, the larger amount of water in combustion gas changed the oxidation mechanism from diffusion-controlled to reaction rate-controlled, and silica formation in water rich combustion gas was faster. Consequently, the oxidation channel could be sealed much earlier and the PyC interphase was oxidized less. However, because the oxidation consumed the interphase and weakened the interface with both the matrix and the fiber, the fracture displacement and the fracture work were greatly increased after oxidation in combustion gas, compared with oxidation in dry air.\textsuperscript{40} The oxidation resistance of 3D SiC/SiC composites (Nicalon\textsuperscript{TM}) was always higher than that of 2D C/SiC composites (T-300, Toray) in the combustion environment.\textsuperscript{41}

3.2.3 Environmental Effects on Fatigue Behavior

It is known that the macroscopic mechanical behavior of ceramic matrix composites is controlled not only by the individual failure of their constituents (fibers, matrix) but also by the interaction between constituents at the fiber-matrix interface. That is the reason why these composites exhibit a non-brittle mechanical behavior, although the constituents are brittle. This phenomenon is caused by the friction between fibers and matrix during sliding of fibers in the matrix. Changes in the microstructure of constituents, or in the interphase, may modify the interfacial shear stress. Such changes can be induced by thermal instability or by local chemical changes (oxidation), and they are reflected in the mechanical hysteresis during cyclic fatigue tests.\textsuperscript{42}

For example, Figure 10 compares experimental evolution of stress/strain loops under tension/compression cyclic loading at room temperatures of a cross-weave SiC/SiC composite (unprotected against oxidation (2D SiC/SiC GS4C, Nicalon fibers) provided by the Société Européenne de Propulsion (France) in two different conditions: as received and after vacuum ageing of 50 h at 800 °C. In the as-received specimen, the mechanical hysteresis increases continuously during the cyclic fatigue test, and the tensile elastic modulus decreases continuously, while the compressive mean elastic modulus is constant. After ageing in vacuum, the cyclic fatigue tests show an increase followed by decrease of the mechanical hysteresis, accompanied by an increase of tensile mean elastic modulus which approaches and reaches the value of the compressive mean elastic modulus. These changes suggest that vacuum heating lowers the initial interfacial shear stress due to chemical reactions at the interface, possibly because of residual oxygen diffusion from the Nicalon SiC fibers to the pyrocarbon interphase.\textsuperscript{42}
Mechanical stability of SiC/SiC composites under cyclic loading in oxidizing environments is important in a multitude of applications where these composites are used, such as structural components in turbine engines of hypersonic flight vehicles, or in spacecraft reentry thermal protection systems. These advanced applications in aero- and space industries motivated recent interest on tension/compression fatigue at high temperature in air and steam environments. The composites were comprised of Hi-Nicalon™ fibers, pyrolytic fiber coating (with B₄C overlay) and SiC-based oxidation inhibited matrix containing alternate layers of SiC and B₄C. The tests showed that the damage and failure of the composites at 1200 °C are caused by oxidation embrittlement in the two environments, air and steam. Fatigue from tension-compression cycles causes more extensive matrix cracking than tension-tension fatigue. Because these cracks are exposed to the oxidizing environment, oxygen (or steam) enters through the cracks and reacts with SiC to form reaction products (silica) that fuse the fibers together. But once these fused fibers in the oxidized matrix fail, larger cracks form with appreciable depth. These unbridged
matrix cracks cause the load redistribution to intact regions in the composites, and ultimately the composite fails. Damage is significantly accelerated in presence of steam, resulting in reduced fatigue performance.\textsuperscript{43,44} Figure 11 shows micrographs of the fracture surfaces in specimens tested in tension-compression cycles in different environments. They show that a large fraction of fracture surface is oxidized, supporting the above mechanism.\textsuperscript{44}

A micromechanics model of the thermomechanical behavior of minicomposites based on the matrix cracking mechanism in oxidizing environments was proposed by Pailler and Lamon.\textsuperscript{45} Matrix cracking is considered as a prerequisite for interphase oxidation. Matrix cracking involves multiple cracks that develop under load from microstructural heterogeneities in the matrix. As the cracks are arrested at the fiber/matrix interphase, debonding cracks are created. The latter allow oxygen ingress, which causes degradation of the fiber/matrix interphase at high temperatures. As the pyrocarbon interphase is consumed, the loads are progressively transferred to the fibers. Failure may occur as a result of fiber overloading. The model treats statistically the rate of interface degradation and crack healing induced by oxidation at high temperatures. Model predictions were found to be in good agreement with the room temperature behavior of SiC/SiC minicomposites reinforced with Hi-Nicalon fibers multifilament tows.
4. ADVANCED SiC/SiC COMPOSITES

4.1 ENGINEERED COMPOSITES WITH IMPROVED OXIDATION STABILITY

After being established that degradation of mechanical properties in oxidizing atmospheres at high temperatures is caused by oxidation of the interface (usually, pyrolytic carbon), efforts were recently directed at developing advanced composites with improved interface oxidation resistance.

Multilayer interfaces formed by repeated deposition sequences of PyC and SiC were reported to provide higher oxidation resistance and enhanced mechanical properties in comparison to those involving only one homogeneous PyC layer. For example, composites with four layers of PyC/SiC coatings (Fig. 12) exhibit high strength and high toughness, and multiple crack deflection features that are interesting for high temperature applications.

The composites were prepared by alternate sublayers of PyC and SiC deposited on Nicalon fibers by CVI. They were tested under tensile static loading (60-140 MPa) at high temperatures (700 – 1200 °C) in air. The use of the multilayered interface improved considerably the lifetime and the fatigue behavior, in comparison with the equivalent composite having a single PyC layer (50 nm thick). In addition, the multilayered composite showed interesting oxidation resistance below 1000 °C under stress lower than 120 MPa, even in presence of small and large densities of cracks.

Two generations of multilayered interfaces, composed of carbon and silicon carbide, have been developed to act as a mechanical fuse in SiC/SiC composites with improved oxidation resistance. From the mechanical point of view, pyrocarbon is the ideal interface material, but it is sensitive to oxidation. On the other hand, SiC is oxidation resistant and acts as a protective layer for PyC. In the multilayered interface the carbon mechanical fuse is split into thin sublayers, each being protected against oxidation by the neighboring SiC-based glass former layers. The first generation of multilayers was synthesized by isobaric CVI with sublayers of micrometer thickness, and used early generation SiC fibers (Nicalon NL202). The second generation of multilayer composites was developed with nanometer-size sublayers formed by pressure-pulsed CVI. They were reinforced with Hi-Nicalon SiC fibers. Minicomposites with a single two of fibers showed improved mechanical properties and oxidation resistance compared with their counterparts with a single layer of pyrocarbon coating. A great advantage of multilayer interface coatings is the deflection of matrix cracks at the fiber surface, as shown in Fig. 13 for composites with multilayered interphases of the first and second generation.
Another great advantage of multilayered interphase coatings is improvement of oxidation resistance. When sublayers have very low thickness, typically a few nanometers as obtained by the pressure-pulsed CVI, they display the self-healing behavior in oxidizing atmospheres. This is the effect of crack sealing oxides formed by reactions such as (2) and (3) in presence of oxygen and reactions (6) and (7) in presence of water, as shown above. The most common self-healing oxide phase examples are SiO₂, B₂O₃, and mixed compositions B₂O₃-SiO₂. Unfortunately, boron is incompatible with nuclear applications. However, the solution of boron-free hybrid interphases, such as PyC-SiC, and multilayer interphases, such as (PyC-SiC)_n remains. For example, the lifetime in air at 700 °C under static loading of 1D SiC (Hi-Nicalon)/SiC minicomposite is improved when the classical PyC₁₀₀ single layer interphase is replaced by a (PyC₂₀-SiC₅₀)₁₀ multilayer hybrid interphase (the subscripts stands for layer thickness in nm).

A different approach for improved oxidation stability of SiC/SiC composites is using environmental barrier coatings (EBC). It is known for example that SiC-based systems are not suitable for long term use at high temperatures (above 1200 °C) in high steam environments, such as that found in gas turbines. A research program conducted by NASA investigated other possible approaches for mitigation of accelerated oxidation caused by water vapor. Environmental barrier coatings for SiC developed under this program and the tests performed led to recognition of three efficient EBC formulations: calcium aluminosilicate, yttrium silicate, and barium-strontium aluminosilicate. These compositions may protect the SiC surface and reduce fiber recession in active oxidation conditions, but they might not be suitable for nuclear applications. A different method was proposed, which does not use other components but SiC. The double layer coating of SiC composite surface using the CVD-SiC process offered some protection in the 4000 h oxidation test of SiC-fiber boded composites (TyrannoHex™) at 1500 °C in air. However, some SiO₂ scale spalling could not be avoided because of large volume changes occurring during α − β phase transformation in cyclic heating. More recently, it was reported that the mechanical properties and the oxidation resistance of SiC/CVI-SiC composites was improved (compared with
SiC/SiC composites with the PyC interphase) when the precursor infiltration and pyrolysis (PIP) method was used for coating SiC fibers with SiC interphase.\textsuperscript{53,54} In contrast with the PyC interphase, that is consumed on oxidation and allows silica to bond strongly with the fibers and matrix, the PIP-SiC interphase can effectively protect SiC fibers against thermal oxidation, based on tests conducted at 900 °C for 6 and 12 h. However, PIP type composites cannot be used for nuclear applications.

4.2 ENGINEERED COMPOSITES FOR USE IN NUCLEAR APPLICATIONS

Because of their low induced radioactivity, excellent high temperature mechanical properties, and excellent radiation resistance, SiC/PyC/SiC composites fabricated from stoichiometric SiC fibers and pure SiC matrix are potential structural materials for gas-cooled high temperature nuclear reactors, of both fusion and fission type.\textsuperscript{55} This new and extremely demanding application raises specific constraints on the fibers, matrix, and interphase. The environment that SiC/SiC composites would see in gas-cooled nuclear reactors is not radically different than that of their classical field of application, gas turbines. The conditions of temperature, gas pressure, and lifetime are about the same. However, the continuous exposure to strong oxidizing medium is not expected to occur in normal operation conditions of high temperature reactors. What is different is the continuous exposure to intense radiation by fast neutrons, α particles and electromagnetic radiation. Such severe radiation conditions are known to change the structure of materials, and hence their properties.\textsuperscript{51}

Irradiation of monolithic SiC causes moderate swelling. Irradiation of SiC fibers causes different effects, depending on the composition and structure. Stoichiometric SiC fibers undergo moderate swelling like SiC phase. Fibers of the first and second generation (Nicalon and Hi-Nicalon) experience different degrees of swelling. The mismatch of volume changes under neutron irradiation is destructive to the composite. This is one important reason why third generation stoichiometric fibers are preferred for SiC/SiC composites engineered for nuclear applications. Irradiation of PyC is known to induce anisotropic dimensional changes, which would induce residual thermal stress and might affect the fiber-matrix bonding upon prolonged irradiation exposure.\textsuperscript{51} To avoid unwanted effects, the thickness of PyC interphase should be as low as possible, but large enough to allow matrix crack deflection. This can be achieved by using hybrid multilayered (PyC-SiC)\textsubscript{n} interphase with thin (only a few nanometers) PyC sublayers. Indeed, the effect of neutron irradiation on composites with different type of interphase (either single layer PyC or multilayer PyC-SiC) showed better performance of multilayer composites with thin innermost PyC layers.\textsuperscript{56,57} As mentioned above, composites with multilayered interphases have the advantage of higher oxidation resistance over those with one single PyC layer.\textsuperscript{47,49}

The oxidation resistance of irradiated SiC/SiC composites has not been reported yet, and is currently an active direction of research. Given that at high temperature, the microstructure of irradiated SiC quickly recovers toward its original unirradiated state, significant deviation in oxidation behavior is unlikely.

The low concentration of oxygen in the helium coolant of HTR may cause active oxidation and gasification of SiC at high temperatures, which might limit the use of SiC/SiC composites in high temperature reactors. A recent study reported the behavior of SiC/SiC composites at high temperatures in helium with low concentrations of oxygen. The β-SiC-SiC/PyC/SiC composite was made from Hi-Nicalon fibers (second generation) with a PyC interphase and β-SiC matrix produced by CVI. A β-SiC coating was deposited by CVD on one face of the composite to act as an environmental barrier by closing the porosity on that face. Oxidation tests at 0.2 Pa and 2 Pa O\textsubscript{2} in helium were conducted up to 2000 °C by the gravimetric method. Water was introduced as the impurity in helium gas of different qualities (1 ppm
H₂O and respectively 2 and 20 ppm O₂). It was assumed that these conditions simulate the environment of gas-cooled fast reactors (GFR). Weight changes versus temperature at the two oxygen concentrations showed the existence of three domains in the zone of active oxidation (Fig. 14). For example, at low O₂ partial pressure (0.2 Pa) the weight loss increases slowly with the temperature between 1400 and 1700 °C; the weight loss continues with four time faster rate up to 1800 °C, and then accelerates 3.6 more times between 1800 and ~ 1925 °C. At higher O₂ partial pressure (2 Pa) the three-domain behavior is reproduced, but the transition temperatures are higher and the corresponding weight losses increase even further (Fig. 13). The differences reflect the damage introduced by oxidation of the β-SiC coating at 1270 and 1570 °C. Even more damaging is sublimation of SiC(s) into Si(g), Si₂C(g), and SiC₂(g) that starts above 1800 °C at \( P_{O_2} = 0.2 \) Pa and 1890 °C at \( P_{O_2} = 2 \) Pa. Similar results were obtained on the face of the composite that was not coated by an environmental barrier of β-SiC, and was concluded that coating was probably too thin and did not show a great influence on the oxidation behavior of β-SiC-SiC/PyC/SiC in case of accident conditions leading to active oxidation. However, the coating remains essential for closing the porosity and preventing direct contact of oxidizing gas with the body of the composites at temperatures below about 1500-1900 °C. The partial pressure of oxygen (0.2 – 2 Pa) did not have a strong impact on the mass loss rate. Unfortunately, the fibers used were second generation Hi-Nicalon; no information is available about the behavior of third generation SiC fibers in similar conditions.
The β-SiC coated SiC/PyC/SiC composite is a candidate for the cladding gas cooled fast reactor (GFR) and its chemical resistance in other environments that may develop in case of severe accident conditions was recently studied.\textsuperscript{59} Figure 15 compares active oxidation weight loss rates of β-SiC and β-SiC-SiC/PyC/SiC composite in air, helium (with 2 Pa O\textsubscript{2}) and in nitrogen (with 0.5 Pa O\textsubscript{2}) at atmospheric pressure. The same three temperature domains were observed, which is not surprising. However, degradation of the composite was always faster than of monolithic β-SiC above 1700 °C because of the damage caused to β-SiC coating which enables oxidizing species to penetrate inside the composite. At very high temperature, oxidation and sublimation of SiC occurred deep inside the composites, while only a superficial layer of β-SiC was affected. This is caused by the presence of porosity that gives access to the pyrocarbon interphase inside the composite.

The conclusion of these studies\textsuperscript{3,24,58,59} is that in order to withstand high temperatures for prolonged durations that may occur in severe accident scenarios, the SiC/SiC composites must be protected by thicker layers of β-SiC. Substitution of He (with 2 Pa O\textsubscript{2}) by N\textsubscript{2} (with 0.5 Pa O\textsubscript{2}) did not cause important differences in oxidation behavior of β-SiC (Fig. 14). When a sudden temperature increase was applied to β-SiC samples preoxidized in passive conditions, transition from passive to active oxidation occurred and surface damage was identified on the β-SiC monolith. The nature of the gas atmosphere had an effect on the final temperature achieved in these experiments. Helium, with better thermal conductivity than nitrogen, allowed a lower final temperature (1725 °C) than nitrogen (2000 °C), but since nitrogen did not
reduce the damage of β-SiC, the authors did not recommend replacing helium by nitrogen in the case of a sudden accidental loss of coolant.\textsuperscript{59}

The behavior of SiC/SiC composites in helium gas with higher oxygen content was studied experimentally to evaluate the suitability of SiC/SiC composites as candidate structural materials for fusion reactor blanket helium gas-cooled solid breeder blanket (HCSB) reactors. One recent study\textsuperscript{60} addressed the difference in oxidation behavior between conventional composites with pyrolytic carbon interphase (PyC-SiC/SiC) and advanced multilayer interphase (ML-SiC/SiC) in He with 150 Pa O\textsubscript{2} at 1000 – 1200 °C. The advanced ML-SiC/SiC composite showed small weight change at both 1000 and 1200 °C, while the conventional PyC-SiC/SiC composite behaved similarly only at 1200 °C. The good stability was due to SiO\textsubscript{2} which formed with relatively high rate and was able to seal the specimen surface and to protect the PyC interphase (passive oxidation). In contrast, the conventional PyC-SiC/SiC composite showed significant weight loss at 1000 °C, where surface sealing was not efficient and significant degradation of the pyrolytic carbon layer occurred (Fig. 16).

At high O\textsubscript{2} partial pressures (10 – 150 Pa) the weight loss caused by oxidation of SiC/SiC composites at 1100 °C depends on the oxygen pressure. The kinetics is governed by the linear-parabolic law, with the rate constants of the linear region being correlated with the O\textsubscript{2} partial pressure.\textsuperscript{61} This behavior contrasts with the report cited above\textsuperscript{58} according to which oxidation is not sensitive to $P_{O_2}$ in the range of 0.2 – 2 Pa. The difference of behavior may possibly be caused by the quality of SiC fibers used (Nicalon first generation in one report\textsuperscript{61} versus Hi-Nicalon in another\textsuperscript{58}). Gasification rates of the same Nicalon-based composite in helium containing 1 % and 0.1 % hydrogen were 6 to 26 times slower than the rates in oxygen, based on the linear kinetics approximation.\textsuperscript{61}
5. CHEMICAL COMPATIBILITY OF SiC/SiC COMPOSITES IN THE ENVIRONMENT OF ADVANCED NUCLEAR REACTORS

Ceramic matrix composites are being evaluated for use at high temperatures and high radiation fields in advanced reactor concepts, including high temperature gas-cooled reactor (HTGR), very high temperature reactor (VHTR), gas fast reactor (GFR), sodium fast reactor (SFR), and fluoride-cooled high-temperature reactor (FHR). SiC/SiC ceramic matrix composites have excellent thermal properties, mechanical strength, and radiation resistance, all desirable properties for nuclear applications. However, their qualification as component materials in each of the above reactor concepts depends critically on the stability of SiC/SiC composites in contact with the chemical environment of the coolant system.

The operating temperature in VHTR reactors may reach 1000 °C over sustained periods of time, and the gas atmosphere is expected to be helium with small content of impurities (CO, CO₂, CH₄, H₂, H₂O, O₂) at pressures that may reach 75 bar. The slightly oxidizing environment and the presence of water vapor may be quite aggressive to SiC/SiC composites. Additional factor operating under high gas flow rates is the mechanical corrosion caused potentially by particles transported by the gas. Although comprehensive studies of the environmental effects on SiC/SiC composites under these special conditions are missing, the current technical progress and the continuous improvement of materials properties allows for drawing several educated conclusions on the compatibility issues.

The latest generation of SiC/SiC composites for nuclear applications is based on stoichiometric SiC fibers, thin pyrolytic carbon interphase or multilayered interphase with alternating layers of SiC and pyrolytic carbon, and a crystalline high purity β-SiC matrix obtained by chemical vapor infiltration (CVI). An environmentally barrier coating may also be required to seal the surface porosity. This design is the result of continuous properties improvement over three generations of SiC fibers and two generations of interphase compositions. Historically, composites with improved mechanical properties, such as fatigue resistance under cyclic operation, were needed for high temperature operation in the oxidizing environment encountered in gas turbines. Now it is known that the combination of high-purity stoichiometric SiC fibers with thin interphase layers or multilayers and crystalline β-SiC matrix imparts superior chemical resistance against oxidation by air and steam with high oxidation potential. However, very little is known about the behavior of these same composites in the helium coolant with low oxidation potential encountered in gas-cooled nuclear reactors.

A few studies were recently published on the effect of different accident condition scenarios on the environmental resistance of SiC/SiC composites at low oxygen concentrations in helium. Unfortunately the SiC fibers used were Hi-Nicalon of the second generation; no information was found about the behavior in similar conditions of composites obtained from third generation, stoichiometric SiC fibers. Even so, the few existing data demonstrate good oxidation stability at low oxygen concentrations in the temperature range of normal operation conditions of VHTR. For example, Fig. 13 (above) shows that β-SiC/PyC/SiC composites experience passive oxidation up to 1300 °C at 0.2 and 2 Pa O₂. In these composites, the fiber was second generation Hi-Nicalon. The third generation Hi-Nicalon S fibers have even better oxidation resistance, with active to passive transition at lower oxygen pressure, as shown in Fig. 5 (above). According to that figure, SiC fibers (Nicalon-S) would suffer passive oxidation limits in the range of temperatures (below 1300 K or about 1000 °C) and oxygen partial pressures (< 1 Pa) that may be encountered in normal operation conditions. This is an encouraging result, but it should be noted that the transition ranges reported in Fig. 5 (above) are based on calculations and are not compared
against direct experimental data. Moreover, little or nothing is available about the oxidation resistance and microstructural changes suffered by SiC/SiC composites in the same conditions.

Most of information available was obtained from mechanical testing of composites exposed at high temperatures in conditions where gasification by decomposition, volatilization, or active oxidation is of concern (vacuum or inert gas with low concentrations of oxidant gas). For example, it has been reported that composites fabricated with Hi-Nicalon S fibers maintain high strength in vacuum up to 1700 °C and do not show structure deterioration up to 1900 °C in vacuum.29 Consistent with that, composites fabricated with Tyranno SA fibers were reported to maintain stable mechanical properties in air up to 1300 °C, due to the engineered interphase composed of SiC and PyC sublayers and the high crystallinity of the β-SiC matrix.37 According to indirect information, SA-TyrannoHex composites fabricated with Tyranno-SA fibers and protected by a secondary CVD SiC surface layer maintained their mechanical strength after 4000 h exposure in air at 1500 °C.36 At these conditions, carbon based materials and metal alloys fail.

Although experimental data about the behavior of nuclear grade SiC/SiC composites in conditions relevant to either normal operation or off-normal conditions in VHTR are scarce, all results available so far point to the conclusion that SiC composites engineered for nuclear applications have a great environmental resistance and maintain mechanical properties up to 1500 - 1700 °C in vacuum and 1300 – 1500 °C in air. These temperatures are way above the range of normal operation regime (750 – 900 °C). Extrapolated behavior of stoichiometric fibers of the third generation shows that they should be stable against passive oxidation at low oxygen concentration at the temperatures of normal operation conditions.

The control of coolant chemistry is needed to protect the core components. A pure helium coolant should not be of concern, but in reality the coolant may have a slight oxidizing character either because traces of O2, CO2, or H2O are expected to be always present, and also because slightly oxidizing potential is required to protect metallic parts against carburization and embrittlement. At low oxygen concentrations, as expected for the He coolant, SiC may experience gasification by active oxidation and volatilization at high temperatures. These reactions would totally damage the composites. It appears however that sublimation of Hi-Nicalon fibers occurs at higher temperatures (1800 °C) than those that might hypothetically develop in severe accident conditions. On the other hand, high oxygen supply is to be avoided because oxidation of the pyrolytic carbon interphase occurs even at intermediate temperatures. Other chemical factors than must be controlled in the coolant are water and methane concentrations, because they interfere with silica growth.13

The available sources contain very little information about the long term behavior of SiC/SiC composites in conditions simulating normal operation conditions of gas-cooled reactors. Recently published data58,59 are targeted to GFR-relevant environments rather than VHTR. Long-term oxidation behavior studies in impure helium, both in irradiation-free and irradiation active conditions, are still needed. Although SiC/SiC composites have been considered for fusion reactor gas-cooled blanket, there still is a huge knowledge gap between the needs to select and qualify these composites in gas-cooled high-temperature fission reactors and the available information on the behavior of composites in other high-temperature impure helium conditions, such as the gas turbine environment.

Last, it is noted that most of the experimental results available today have been measured in short time experiments, a few hours at most. One notable exception is the 4000 h long experiment at low oxygen pressures reported above.58 More long-term experiments are needed with advanced SiC/SiC composites, using generation III fibers, engineered interphases, protecting coatings etc. These studies are needed to obtain quantitative results and validate the lifetime projections based on the limited corpus of
data existing today. The methodology might be oriented towards determination of long time oxidation behavior of each constituent in the composite (advanced fibers, engineered interphase, matrix) and then of nuclear grade composites themselves. Investigation of the oxidation mechanisms in extreme environment conditions should enable development of predictive models and standardization of qualification procedures.\textsuperscript{13}
6. CONCLUSIONS

The information available in the literature on chemical reactivity of SiC/SiC composites in contact with the helium coolant of gas-cooled reactors was reviewed in this literature report. Special emphasis was placed on collecting available information regarding chemical reactivity of different components (SiC fibers and monoliths, and fiber/matrix interphase) of SiC/SiC composites in contact with various environments, containing oxygen, water vapor and other components, and on the effect of high temperature exposure to these chemically aggressive conditions. The recent progress in development of nuclear grade SiC fibers and composites led to an increased chemical stability of SiC fibers and composites from the third generation (nuclear grade) of materials. The most advanced composites with engineered structures and added environmental barrier protection have chemical resistance that would allow them maintain mechanical properties at the highest temperatures (1200-1300 or higher °C) that might be projected in the extreme conditions of an air- or water ingress scenario. At these extreme conditions carbon-based materials and metal alloys fail. However, further research is required to assess the long-term stability of these advanced SiC/SiC composites in contact with traces of water or oxygen that might be present in the helium coolant during normal operation. Given that the microstructure of irradiated SiC recovers quickly at high temperatures to the original, non-irradiated form, significant deviations of oxidation behavior caused by irradiation with fast neutrons are unlikely.
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