Preparation of sialon–transition metal silicide composites

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Abstract

A process to prepare sialon–transition metal silicide composites is proposed in the present study. The oxides of transition metals are used as the starting materials and the metal silicides are formed in situ in sialons at elevated temperature. The metal oxides, silica and alumina in starting materials first form a liquid phase on sintering to assist the densification in the graphite furnace. The transition metals then precipitate out to react with Si3N4 to form silicides. The sialon–metal silicide composites with uniform microstructures can be produced. By applying a suitable post-sintering heat-treatment, metallic fibers can also be formed in situ in the surface region on samples. Results for both sialon–Cr-silicide and sialon–Fe-silicide composites are demonstrated in the present study.

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1. Introduction

Sialon belongs to the family of silicon nitride based ceramics and is a typical structural material.1,2 There are two major forms of sialons, namely α-sialon (α', Ln4Si12−8zAl2O3N16−z) and β-sialon (β', Sis4Al3O2N6). Both sialon phases are derived from the parent silicon nitride (Si3N4) with partial substitutions of Al2O3 and AlN bonds for SiN bonds. The α-sialon is stabilized by trapping “modifying” cations such as Li, Ca, Y and selected rare-earth elements in the interstices of the [Si, Al]–[O, N] network, but β-sialon does not contain the “stabilizing” cations. Sialon ceramics are promising candidates for engineering applications because of their excellent hardness, wear resistance and corrosion resistance at room and high temperatures. However, the toughness of the material is low (α-sialon: 3–5 MPa m1/2 and β-sialon: 5–7 MPa m1/2) and densification requires high temperatures (normally above 1750 °C), which limit their wide applications as structural components.

To improve the toughness of sialons, many attempts have been made to adjust the microstructure and grain morphology. The bimodal microstructures containing large elongated grains in a fine and uniform matrix have shown obvious advantages in toughening because of crack deflection and crack bridging mechanisms introduced by the large elongated grains.6,8 However, the sintering temperature of the material is still very high and the toughness may not improve if the grain boundary is strongly bonded.8 Another toughening approach used metal additions to enhance the toughness of sialons, similar to what were achieved in Al2O3 ceramics.7,8 Several transition metals, such as Fe, Ni, Cr, Mn, have been used to improve the toughness of sialons and a two-fold increase in the toughness of sialons has been reported.7 But in these studies, the transition metals were added into sialon compositions in metallic powder forms. Several difficulties are encountered when metallic powder is used in processing of sialon–transition metal composites. Due to the flammability of fine metallic powders, the size of metallic particles is usually above 1 μm. At high temperature, the metallic phase aggregated easily and the size of the metallic inclusions in composites usually becomes far too large. The large size inho-
mogeneity could generate microcracks due to thermal expansion mismatch between the metal and ceramic phases. Furthermore, Si₃N₄ is very reactive to many transition metals directly to form metallic silicides and generate gas phases. These reactions usually take place before the densification of the composite systems, so fully dense composites are thus difficult to be achieved.

In the present study, transition metal oxides, instead of transition metals, are used in the starting compositions. Because of the transient liquid phase sintering mechanism for sialon ceramics, it may be possible to form a uniform transient liquid phase containing metal oxides at the sialon grain boundaries when sintering temperature is above the eutectic of the system, from which metallic phases may precipitate at a higher temperature by reduction in graphite furnace. In this case, the metal oxides in the starting powders would not only provide the metallic phases in final composites, but also play an important role to assist sintering. To validate these assumptions, three aspects are explored in the present study: (1) the possibility to stable α-sialon using transition metals; (2) the possibility to obtain sialon-transition metal silicides composites with a uniform microstructure; and (3) the possibility to toughen α-sialon ceramics by metallic silicides.

2. Experimental

The compositions of the starting materials, α-Si₃N₄ (M11, Stark, Germany), AlN (AT, Stark, Germany), Al₂O₃ (99.9%, AJAX, Australia) and Cr₂O₃ (99%, BDH, England) or Fe₂O₃ (99%, BDH, England) are shown in Table 1. Compositions of metal oxides are designed to form α-sialon phases according to a nominal α-sialon composition (m = 1, n = 1.7). An extra amount of transition metal oxide was added as an additive for the metal source. The starting powders were ball milled in isopropanol for 24 h and, then dried and pressed uniaxially into cylinders in a diameter of 25 mm. Hot-pressing (HP) and pressure-less sintering (PLS) were carried out in a graphite furnace with a flowing N₂. The heating rate was 30 °C/min and after sintering, the specimens were cooled down with the natural cooling rate in the furnace. Some samples embedded in graphite powder were further heat-treated in a tube furnace in a nitrogen atmosphere at 1300 °C for 6 h.

The density of the specimens was measured according to the Archimedes principle. Phase characterization was performed using X-ray diffraction method (XRD, Rigaku Geigerflex Diffractometer, Rigaku Co., Japan). The sintered samples were ground and polished with diamond paste. The microstructures were examined using back-scattered image model of scanning electron microscopy (SEM, JEOL JSM 840A). The composition of each phase was determined with an energy dispersive spectrometer. The volume fraction of the metallic phase in the fired composites was measured with a point counting technique from the SEM micrographs.

3. Results and discussion

3.1. Preparation of sialon–Cr-silicide composites

In order to observe the densification and reaction sequence of the composites, sample Cr-15 was hot-pressed in the graphite furnace at 1200, 1300, 1400, 1500, 1600 °C for 1 min and 1600 °C for 1 h, respectively. The densification curve and phase evolution for the composites are shown in Figs. 1 and 2, respectively. The density of the composites remains low after the first appearance of some metallic silicides at 1400 °C. Densification starts to pick up as the hot-pressing temperature rises to 1500 °C. As shown in Fig. 2, Cr₂O₃ disappeared and CrSi₂ formed after heating to 1300 °C. As temperature increased, the

Table 1 Compositions used to prepare sialon-metal silicide composites

<table>
<thead>
<tr>
<th>Samples</th>
<th>Nominal phases</th>
<th>α-Sialon (m = 1, n = 1.7) (vol.%)</th>
<th>Extra M₂O₃ (wt.%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cr-3</td>
<td>Cr–α-sialon + 3% Cr₂O₃</td>
<td>75.8</td>
<td>16.62</td>
</tr>
<tr>
<td>Fe-3</td>
<td>Cr–α-sialon + 3% Fe₂O₃</td>
<td>75.51</td>
<td>16.55</td>
</tr>
<tr>
<td>Cr-15</td>
<td>Cr–α-sialon + 15% Cr₂O₃</td>
<td>75.8</td>
<td>16.62</td>
</tr>
<tr>
<td>Fe-15</td>
<td>Cr–α-sialon + 15% Fe₂O₃</td>
<td>75.51</td>
<td>16.55</td>
</tr>
</tbody>
</table>

³ Cr₂O₃, ⁵ Fe₂O₃.
amount of CrSi₂ decreased and a high-temperature silicide, Cr₃Si, was then formed. The oxygen partial pressure in the graphite die can be as low as 10⁻¹⁵ atm at 1600 °C. Under this condition, Cr₂O₃ in the sample can be reduced to produce Cr metal. The presence of silicon nitride may further accelerate the reduction of Cr₂O₃. However, no Cr is detected in the sample fired at all temperatures studied, indicating that as soon as Cr was formed from the reduction of Cr₂O₃, it reacted with silicon nitride to form silicide following the reaction,

\[
\text{Si₃N₄} + 1.5\text{Cr} \rightarrow 1.5\text{CrSi₂} + 2\text{N}_2
\]  

(1)

The lowest eutectic temperature in the Cr–Si binary alloy system is 1305 °C. This eutectic is formed between Si (with the solution of 10 ppm Cr) and Cr₅Si₃. From the binary phase diagram of the Cr and Si system, a series of Cr–Si compounds can be formed, following a phase transition sequence from Cr₅Si₃, Cr₃Si, Cr₅Si₃ to Cr₃Si with the increase of Cr content. The Cr₃Si-silicide phase has the highest melting point, 1770 °C among all Cr-silicides. Thus, the final product obtained at 1600 °C containing a very refractory Cr-silicide phase.

The main phases in the composition Cr-15 after hot-pressing at 1600 °C for 1 h are β-sialon and Cr₃Si. A small amount of α-Si₃N₄ also exists, but no α-sialon is detected. If Cr₂O₃ was replaced by Me₂O₃ (e.g. Me = Yb) and other compositions kept the same as that in sample Cr-15, Cr₃Si-silicide should have been the major phase after fired at this temperature. The result suggests that transition metal ion Cr is not an effective stabilizer for the α-sialon structure. The ionic radii of Cr³⁺ and Cr²⁺ are 0.63 and 0.89 Å, respectively. The ionic size of Cr³⁺ is similar to that of Yb³⁺ (0.658 Å) and Cr²⁺ similar to Mg²⁺ (0.66 Å). Both Yb³⁺ and Mg²⁺ are good α-sialon stabilizers. Thus, the ionic radius appears not to be a critical factor for α-sialon stabilization. The formation temperature (1300 °C) of Cr-silicide is lower than that for α-sialon (>1500 °C). Once the Cr-silicide phase is formed, it may prohibit chromium from entering the α-sialon structure. As a result, the solubility of transition metal ion in α-sialon may be too low to form a stable α-sialon phase.

Microstructural features of the sialon-Cr-silicide system at different temperatures are shown in Fig. 3. A typical micrograph of the composite after hot-pressed at 1600 °C for 1 h is also shown in the figure. EDS results confirm that the bright particles in the back scattered SEM images are Cr-silicide which is in the range of 200–300 nm in size when sintered at 1300 and 1400 °C. This particle size is much smaller than that of the starting Cr₂O₃ powder, which was in the range of several micrometers, suggesting that the Cr-silicide particles resulted from a process of solution and precipitation. It is thought that the added Cr₂O₃ dissolved in a transient liquid phase below 1300 °C, and then Cr metal precipitated from the liquid due to the reducing environment and reacted with the surrounding silicon nitride to form chromium silicide when temperature further increased. On the other hand, the transient liquid phase could also facilitate the mass transportation and assist densification. Grain growth of the Cr-silicide phase appears not significant when sample Cr-15 was fired between 1400 and 1600 °C for 1 min only. But the Cr-silicide phase underwent considerable coarsening after heating for 1 h at 1600 °C.

The microstructures of Cr–α-sialon compositions containing extra 3 and 15 wt% Cr₂O₃ (Cr-3 and Cr-15) hot-pressed at 1600 °C for 1 h are shown in Fig. 4. As indicated, the Cr-silicide particles distributed uniformly within the sialon matrix in both compositions. The microstructures of Cr-3 and Cr-15 are very similar except the difference in the volume and particle size of the metal silicide. In sample Cr-3, the silicide phase is about 4 vol.% and the average particle size around 1 µm. In sample Cr-15, the silicide is about 12 vol.%
and the average particle size around 2.5 μm. A significant increase in apparent density is noted for the increase of Cr₂O₃ content in the samples from 3 wt.% (ρ_{Cr-3} = 2.12 g/cm³) to 15 wt.% (ρ_{Cr-15} = 2.90 g/cm³). A denser microstructure can also be found for sample Cr-15 compared to Cr-3 (Fig. 4). It indicates that the introduction of chromium oxide has assisted densification of sialon–silicide composites through the formation of more liquid in the system.

3.2. Preparation of sialon–Fe-silicide composites

The phase and microstructure evolution of sialon–Fe-silicide composites are very similar to those of the sialon–Cr-silicide composites. As demonstrated in Fig. 5, the disappearance of Fe₂O₃ took place at 1200 °C, which is about 100 °C lower than the disappearance of Cr₂O₃. There was no metallic Fe phase detected in the sample when sintering temperature...
further increased. The resulting phases after fired at 1600 °C were α-Si₃N₄, β-sialon, FeSi and Fe₅Si₃. The microstructure of the sialon–Fe-silicide composite (as shown in Fig. 6) is very similar to that of sialon–Cr-silicide composite, except that the Fe-silicide particles are smaller in size. The similarity of the two systems suggests that the design concept of the present study may be applicable to other sialon–transition metal silicide systems.

3.3. Effect of post-sintering heat-treatment

The microstructure of the sialon–Cr-silicide composite (Cr-3) prepared by pressureless sintering at 1600 °C for 2 h is shown in Fig. 7. The microstructure of the sintered composite after post-sintering heat-treatment at 1300 °C for 6 h is also shown in Fig. 7. After the post-sintering heat-treatment, fiber-like particles with bright contrast were observed within...
a zone of a thickness around 5–10 μm from the surface. Results of EDX analysis on the bright phases in the two samples are shown in Fig. 8. The Cr-silicide particles in the as-sintered sample contain a relatively high concentration of silicon. But the fiber-like phase in the heat-treated sample has a very little Si content, indicating that the fibers are a metallic Cr phase rather than Cr-silicide. This suggests that silicon may have diffused out of the CrSi phase during post-sintering heat-treatment. Sample Cr-3 has a relatively low density and contains continuous pores. Chromium has a high vapor pressure at 1300 °C, close to 1 Pa, and thus can evaporate at the heat-treatment temperature. The appearance of the needle shaped Cr phase may be a result of evaporation of Cr-silicide through continuous pore channels and condensation of metallic Cr fibers on the sample surface. This phenomenon requires further investigation for an ascertain explanation.

Experimental results of both Cr– and Fe–sialon systems have demonstrated that transition metals are not effective sialon stabilizer and they promote the formation of composites containing β-sialon (z = 1.0) and silicides.

3.4. Mechanical properties

The mechanical properties of the composites prepared by hot-pressing at 1600 °C for 2 h are determined. To facilitate densification, 4 wt.% CaO was added in the samples. The compositions and mechanical performance of the composites are listed in Table 2. Hardness and toughness of these two composites are above 15 GPa and 6 MPa m$^{1/2}$, which are very similar to the normal β-sialon ceramics. But the samples in the present study were sintered at 1600 °C, much lower than that for β-sialon (usually around 1750 °C). Thus, there is potential for applications of the sialon–silicide composites in structural, refractory and catalyst areas.

Fig. 9 shows the interactions between cracks and metallic silicide particles in both Cr- and Fe-silicide–sialon systems. Crack deflection and crack bridging due to the presence of the metal silicide are observed (denoted with arrows). The presence of these mechanisms should enhance the toughness of sialons. Nevertheless, the low toughness of the silicide itself

| Table 2 Compositions and mechanical properties of β-sialon + 4% CaO + 15% M$_2$O$_3$ samples |
|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| Samples        | Compositions (wt.%) | Density (g/cm$^3$) | HV$_{100}$ (GPa) | K$_{IC}$ (MPa m$^{1/2}$) |
|                | Si$_3$N$_4$ | AlN | Al$_2$O$_3$ | CaO | M$_2$O$_3$ |                |                  |                  |
| CrCa-15        | 67.06   | 4.07 | 9.87    | 4   | 15$^a$ | 3.28  | 15.74  | 6.96 |
| FeCa-15        | 67.06   | 4.07 | 9.87    | 4   | 15$^b$ | 3.34  | 15.75  | 6.17 |

$^a$ Cr$_2$O$_3$

$^b$ Fe$_2$O$_3$
may trade off the benefits induced from the crack bridging. Not many pores are observed in the samples shown in Fig. 9, suggesting that dense sialon–transition metal silicide can be prepared.

4. Conclusions

(1) Sialon–transition metal silicides composites with uniform microstructure can be prepared by reaction sintering of silicon nitride and transition metal oxides. A transition oxide-containing liquid forms first, then the transition metal precipitates out with increasing temperature and reacts with surrounding Si₃N₄ to form metal silicides.

(2) Transition metal oxides can be used as sintering aids for sialon ceramics; however, they are not effective stabilizers for α-sialon structures.

(3) By applying suitable post-sintering heat-treatment, metallic Cr fibers can be formed in the surface region of porous sialon body.

(4) Dense sialon–silicide composites can be prepared by adding a small amount of CaO as a sintering aid.

(5) Typical mechanical properties of dense sialon–silicide composites are HV₀ > 15 GPa and KIC > 6 MPa√m. These values are comparable to those of β-sialon; nevertheless, the composites can be prepared at a relatively lower sintering temperature than β-sialon.

References


