A method is proposed to prepare $\text{Al}_2\text{O}_3$–$\text{AlN}$–Ni composites. The composites are prepared by sintering $\text{Al}_2\text{O}_3$/NiAl powder mixtures at 1600°C in a mixture of nitrogen and carbon monoxide. The presence of NiAl particles raises the green density of $\text{Al}_2\text{O}_3$/NiAl powder compacts. During sintering, NiAl reacts with nitrogen to form AlN and Ni inclusions. A volume expansion accompanies the reaction. Because of the high green density and the reaction, the volume shrinkage of the $\text{Al}_2\text{O}_3$–$\text{AlN}$–Ni composite decreases with the increase of added NiAl content.

I. Introduction

Considerable volume shrinkage accompanies the sintering of ceramic powder compacts. Shape distortion is thus frequently observed. Volume shrinkage during sintering also imposes limits on the densification of ceramic-matrix composites. The presence of differential shrinkage between the matrix and inclusion induces internal stresses. The stresses reduce the final density that can be achieved by sintering. It is therefore important to develop alternative forming processes that result in low shrinkage.

Two interesting reaction processes involve the interactions of metals and gases. One process is the directed oxidation (or nitridation) of molten metal (DMO)\(^2,3\) and another process is the reaction bonding of aluminum oxide (RBAO).\(^4,5\) In these processes, molten aluminum is reacted either with oxygen or with nitrogen to form aluminum oxide or aluminum nitride. The processes have many advantages such as low processing temperature and low shrinkage. In the present study, a process is proposed to prepare $\text{Al}_2\text{O}_3$–$\text{AlN}$–Ni composites. This process also involves the interactions of metals and gases. The metal used in this study is nickel aluminide, NiAl. NiAl is an intermetallic compound with a high melting point, 1638°C.\(^6\) Therefore, contrary to the DMO and RBAO processes, the metal used in this study involves a solid-state reaction.

II. Experimental Procedure

Alumina ($\alpha$-$\text{Al}_2\text{O}_3$, TM-DR, Taimei Chemical Co. Ltd., Tokyo, Japan, -0.2 mm and 0 to 50 vol% of nickel aluminide (NiAl, Xform Inc., New York, -5.6 mm, 68.23 wt% Ni, 31.54% Al, 0.05% Fe, 0.01% Cr, and 1300 ppm O) were attrition milled together in ethyl alcohol for 1 h using zirconia media. The mixed powder slurry was dried with a rotary evaporator. The dried lumps were crushed and sieved through a 200 mesh plastic sieve. Powder compacts were formed by pressing uniaxially at 100 MPa. By using the theoretical density of 3.98 g/cm\(^3\) for $\text{Al}_2\text{O}_3$ and 5.95 g/cm\(^3\) for NiAl, the relative density of the green compacts could be estimated. The green compacts were placed inside a closed graphite crucible. The graphite crucible was then fired in a box furnace at 1600°C for 1 h. The graphite crucible and cover were of sufficient mass to retain their shape until the end of the sintering. No gas was thus allowed to enter or escape from the closed crucible during sintering. Above 700°C, oxygen inside the crucible reacted with carbon to form carbon monoxide.\(^7\) The partial pressure of oxygen during sintering was low, $10^{-15}$ atm at 1600°C. The nitrogen inside the crucible was stable at all temperatures. Therefore, the atmosphere during sintering within the graphite crucible was mainly a mixture of CO and N\(_2\).

The apparent density after firing was determined by the water displacement method. Before the specimens were submerged in water, a wax was applied to the surface to prevent water penetration. Phase identification was performed by X-ray powder diffractometry (XRD). Several $\text{Al}_2\text{O}_3$/AlN and $\text{Al}_2\text{O}_3$/Ni powder mixtures with known composition were also prepared by the same mixing procedures. The XRD patterns for the powder mixtures were determined. Three measurements were made for each composition. Calibration curves to correlate the XRD intensities to the amounts of $\text{Al}_2\text{O}_3$, AlN, and Ni in fired composites could then be established. The standard deviation for the determination of AlN was 1 vol%, and that of Ni was 2 vol%.

Polished surfaces were prepared by grinding with diamond slurry to 6 mm and polishing with silica suspension to 0.05 mm. The microstructure was observed by scanning electron microscopy (SEM). The chemical composition of each phase after sintering was determined by electron probe microanalysis (EPMA). The residual carbon content within sintered specimens was determined by IR spectrometry (CS-244, LECO Co., U.S.A.). The interconnectivity of the nickel inclusions was determined by measuring electrical resistivity at room temperature.

III. Results and Discussion

Because of the presence of N\(_2\), CO, and a very small amount of O\(_2\) in the sintering atmosphere, NiAl can either react with N\(_2\) to form AlN and Ni

$$2\text{NiAl} + \text{N}_2 \rightarrow 2\text{AlN} + 2\text{Ni}$$

(1)

or react with O\(_2\) to form $\text{Al}_2\text{O}_3$ and Ni

$$2\text{NiAl} + \frac{3}{2}\text{O}_2 \rightarrow \text{Al}_2\text{O}_3 + 2\text{Ni}$$

(2)

The equilibrium pressures of nitrogen and of oxygen for reactions (1) and (2) are 0.1 and $10^{-17}$ atm, respectively.\(^8\) The partial pressures of nitrogen and of oxygen in the sintering atmosphere are higher than these values; the formation of either AlN/Ni or $\text{Al}_2\text{O}_3$/Ni is therefore possible.

The XRD pattern for the 50% $\text{Al}_2\text{O}_3$/50% NiAl powder mix-
Figs after sintering is shown in Fig. 1. There is no NiAl detected within the detection limit of the instrument (about 1% for NiAl). Aluminum nitride, nickel, and graphite are found instead. The microstructure for the composite containing 50% added NiAl sintered at 1600°C for 1h is shown in Fig. 2. Approximately 8 at.% Al is detected in the Ni inclusions by the EPMA analysis. This corresponds to the solubility of Al in Ni. The amount of each phase in the composites measured by quantitative XRD analysis is shown in Table I. The XRD analysis suggests that reaction (1) is taking place during sintering. AlN and Ni are not found in composites containing less than 10 vol% added NiAl. It may be that the amounts of AlN and Ni are too small to be detected.

Carbon can be dissolved in the nickel melt during sintering. The solubility of carbon in nickel melt is higher than that in solid nickel. Owing to the different solubility, carbon inclusions are precipitated out upon cooling, particularly as the nickel is solidified. Therefore, graphite inclusions are observed within the nickel inclusions or at the interface. The carbon content within the composites is also shown in Table I. As carbon is dissolved in nickel, the amount of carbon increases with nickel content.

The newly formed AlN is not stable, as it may react with O₂ to form Al₂O₃

\[ \text{AlN} + \frac{3}{4}\text{O}_2 = \frac{1}{2}\text{Al}_2\text{O}_3 + \frac{1}{2}\text{N}_2 \]  

(3)

Table I. Measured Amount of Each Phase Presented in the Composites after Sintering at 1600°C for 1 h

<table>
<thead>
<tr>
<th>NiAl added (vol%)</th>
<th>Al₂O₃ (vol%)</th>
<th>AlN (vol%)</th>
<th>Ni (vol%)</th>
<th>Carbon (wt%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>ND†</td>
<td>ND†</td>
<td>0.20</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>ND†</td>
<td>ND†</td>
<td>0.3</td>
<td></td>
</tr>
<tr>
<td>20</td>
<td>94</td>
<td>3</td>
<td>3</td>
<td>1.5</td>
</tr>
<tr>
<td>40</td>
<td>66</td>
<td>15</td>
<td>19</td>
<td>3.8</td>
</tr>
<tr>
<td>50</td>
<td>54</td>
<td>26</td>
<td>20</td>
<td>3.8</td>
</tr>
</tbody>
</table>

†Not detectable.

For this reaction, the equilibrium pressure ratio of N₂ to O₂ is

\[ \frac{P_{N_2}^{1/2}}{P_{O_2}^{3/4}} = 3.5 \times 10^{12} \]  

(4)

at 1600°C. Based on the calculation from the reaction of C and O₂ to form CO, the oxygen partial pressure at 1600°C is 10⁻¹⁵ atm and the nitrogen partial pressure is 0.79 atm. The ratio of N₂ to O₂ is 1.6 × 10¹¹. The value is less than the equilibrium ratio. It thus is thermodynamically possible for AlN to react with O₂ to form Al₂O₃ and N₂. However, AlN is still detected after sintering. Therefore, even though reaction (3) is possible thermodynamically, the reaction is slow under the sintering conditions used.

The green density and fired density of the composites are shown as a function of added NiAl content in Fig. 3. The difference in particle sizes for the NiAl versus the Al₂O₃ powder results in more efficient particle packing. The green density of the composites is thus higher than that of alumina. The fired density increases with increasing NiAl content because the density of Ni is higher than that of Al₂O₃ (8.90 vs 3.98 g/cm³). Through careful microstructural observation, the porosity in fired composites increases with the increase of added NiAl content. For the Al₂O₃/50% NiAl compact, the porosity is less than 10%. An example is given in Fig. 2. It indicates that the Al₂O₃–AlN–Ni composites with relative density higher than 90% can be prepared with a pressureless sintering technique.

During the reaction between of NiAl and N₂, a volume expansion accompanies the reaction. The volume shrinkage of the composite is thus smaller than that of alumina alone (Fig. 4). Therefore, the process developed in the present study has the advantage of low shrinkage. The electrical resistivity of the composites is shown as a function of NiAl content in Fig. 5. Because particulate NiAl is used as a starting material, the resulting Ni inclusions are isolated from each other until nickel
content is higher than 20 vol%. When the added amount of NiAl is less than 40 vol%, the resulting Al₂O₃–AlN–Ni composites are electrical insulators. When the NiAl content is greater than 50 vol%, the resulting composite is an electrical conductor.

IV. Conclusions

A reaction process to prepare Al₂O₃–AlN–Ni composites is proposed in the present study. The starting material used is powder mixtures of Al₂O₃ and 5–50 vol% NiAl. As the powder mixtures are sintered in the presence of nitrogen, AlN and Ni inclusions are formed. Al₂O₃–AlN–Ni composites with relative densities higher than 90% can be prepared by pressureless sintering at 1600°C for 1 h. The green density is increased by adding NiAl particles to Al₂O₃ powder. Furthermore, volume expansion accompanies the reaction during sintering. The process developed in the present study thus has the advantage of low shrinkage.

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References