Hot Hydrogen Exposure Degradation of the Strength of Mullite

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ABSTRACT

This study deals with the corrosion of near stoichiometric mullite (3Al₂O₃-2SiO₂) by pure dry hydrogen gas. Exposure of the mullite samples was at temperatures of 1050 and 1250 °C for times up to 500 hr. Preferential attack of the aluminosilicate glass present in the grain boundaries of the mullite occurred after 125 hr at 1250 °C. Hydrogen scrubbing of the SiO₂ from the glassy grain boundaries and the mullite grains yielded a porous alumina-rich surface. The room temperature strength increased after short exposure times at 1250 °C (up to 125 hr), then decreased by 53 percent after exposure for 500 hr. At 1050 °C, all exposure times (25 to 500 hr) decreased the strength. After 500 hr in hydrogen at 1050 °C, the room temperature strength of mullite decreased 22 percent. We also observed a rapid 25 percent strength loss after short exposure times at 1050 °C. This is attributed to the calcium/hydrogen assisted crystallization of the glassy, grain-boundary phase.

INTRODUCTION

Strength retention to high temperature combined with low thermal expansion make the monolithic form of mullite (3Al₂O₃-2SiO₂) a potentially useful material for many structural applications. At the time this study was initiated it was a candidate for the pressure vessel in the hot zone of a ceramic Stirling engine. Current technical needs emphasize the use of mullite as a matrix and/or as a reinforcing fiber for advanced fiber reinforced ceramic matrix composites. Mullite coatings are also being developed to enhance the environmental durability of the silicon-based ceramics, SiC and Si₃N₄. In some applications, the operating conditions will be highly reducing and may require significant strength retention, even after long time exposure to hydrogen at elevated temperature. Examples include the ceramic Stirling engine mentioned above and future hydrogen-fueled hypersonic vehicles. Operating temperatures in a Stirling pressure vessel could approach 1200 °C. Applications for hypersonic vehicles may require

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even higher hydrogen exposure temperatures. Under these conditions, corrosion of the mullite may occur because of uniform material loss, and/or selective pitting, which decrease the fracture strength.

The effect of high temperature hydrogen exposure on the microstructure of mullite, along with the kinetics of the reaction, has been the subject of several investigations. Wright and Wolf (Ref. 1) found that severe chemical attack of refractory mullite occurs when exposed to hydrogen above 1370 °C. Crowley (Ref. 2) further studied the effects of hydrogen on refractory mullite and measured the rate of silica loss at 1425 °C. He concluded that the weight loss was dependent on the hydrogen flow rate and that the hydrogen-mullite reaction was diffusion controlled. He also observed a decrease in compressive strength with increasing weight loss. Chen (Ref. 3), using techniques similar to those of Crowley, found the reaction of mullite with hydrogen to be a mixed control process. He said that the reaction depended on the transportation of the gaseous product species through the porous product layer and through the gaseous boundary layer. The amount of weight loss was also dependent on the flow rate of hydrogen over the mullite. The reduction of mullite by hydrogen gas was studied in more detail by Tso and Pask (Ref. 4). These investigators used a high purity stoichiometric mullite in their tests. Later, Tso (Ref. 5) found that diffusion through the porous layer was not rate limiting, but the reduction reaction was. All of the above investigators identified the loss of silica from the mullite to be the major reaction. The result of this reaction is a porous product layer of alpha alumina on the surface.

\[
\text{Al}_6\text{Si}_2\text{O}_{13}(s) + 2\text{H}_2(g) \rightarrow 3\text{Al}_2\text{O}_3(s) + 2\text{SiO}(g) + 2\text{H}_2\text{O}(g)
\]

A review of the literature reveals that the hydrogen mullite reaction has only been studied at temperatures above 1350 °C and mainly for short times, i.e., a maximum of 50 hr. In applications such as the Stirling engine several thousand hr of exposure at somewhat lower temperatures is required.

The objective of this study was to determine the effect of long time hydrogen exposure on the microstructure and retained room temperature strength of near stoichiometric mullite.
EXPERIMENTAL

Mullit' test bars 6 by 3 by 25 mm were prepared by diamond grinding. The bars were ultrasonically cleaned in neat ethanol, dried, measured and weighed before each hydrogen exposure. Approximately 20 test bars were used in each hydrogen test.

The hydrogen gas was dried and purified to a moisture level of about 1 ppm. The flow rate was regulated at room temperature and held constant at 475 cc/min. Hydrogen exposures were done in a silicon carbide resistance heated horizontal tube furnace with a 99.8 percent pure Al₂O₃ tube. The specimens were placed on high purity Al₂O₃ setters in a high purity Al₂O₃ boat located in the central hot-zone of the furnace. The temperature variation within the furnace hot-zone was about ±10 °C.

Heating to the exposure temperature was in high purity helium. Once the desired temperature was achieved, the atmosphere was switched to hydrogen. At the completion of the exposure cycle, the gas was again switched to helium for the cool-down cycle. The mullite samples were exposed to hydrogen at 1050 and 1250 °C for times of 25 to 500 hr.

At the completion of the hydrogen exposure, the samples were removed from the furnace, weighed, and measured. The exposed specimens were subsequently stressed to fracture in a four-point bend fixture. The loading rate was 0.05 cm/min with an outer span of 1.90 cm and an inner span of 0.95 cm. The following equation:

\[ \text{MOR Strength} = \frac{3P(L - a)}{2bd^2} \]  

was used to calculate fracture strength. \( P \) is the load at failure, \( L \) is the outer span, \( a \) is the inner span, \( b \) is the specimen width, and \( d \) is the specimen thickness. The original sample dimensions were used to calculate the fracture stresses. Results showed that the surface recession due to material loss on the surface did not exceed 20 µm and was considered to have no significant effect (<2 percent) on the strength calculation.

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¹K-635 mullite, Kyocera International, San Diego, CA.
A number of techniques were used to characterize the exposed samples including: X-ray diffraction (XRD), scanning electron microscopy (SEM), X-ray energy dispersive spectroscopy (XEDS), transmission electron microscopy (TEM), selected area diffraction (SAD), electron microprobe, and optical microscopy.

RESULTS AND DISCUSSION

Etching the as-received mullite with a 2:1 mixture of HCl/HF:H2O for five seconds revealed the microstructure shown in Fig. 1. In the as-received sample, the grains were highly acicular with an aspect ratio approaching 8 and an average grain length of 10 μm. A silica-rich glass (marked G in Fig. 1) between the mullite grains (marked M in Fig. 1) makes up about 16 percent of the structure. This is typical of commercial mullites. The slow transport rates of the constituent species require the use of excess silica to facilitate densification. As shown in Table I, this mullite also contained calcium and magnesium for enhancement of sintering.

Table I.—Chemical Analysis of Mullite, wt%

<table>
<thead>
<tr>
<th>Element</th>
<th>Aluminum</th>
<th>Silicon</th>
<th>Calcium</th>
<th>Iron</th>
<th>Magnesium</th>
</tr>
</thead>
<tbody>
<tr>
<td>Value (wt%)</td>
<td>35.4</td>
<td>13.8</td>
<td>0.7</td>
<td>0.3</td>
<td>0.8</td>
</tr>
</tbody>
</table>

Elements not detected (<1 ppm): B, Ba, Co, Cr, Mn, Mo, Ni, P, Ti, V, Y, Zn, Zr.

Point count approximation on ceramographic sections indicated a porosity of about 3 percent. This value was confirmed by immersion density measurements. The measured bulk density was 3.05 ±0.01 g/cc.

Hydrogen corrosion is seen in the photomicrographs of Fig. 2. This figure shows the as-ground surface of the mullite in the as-received condition and after 500 hr exposure in hydrogen at 1050 and 1250 °C. Large surface voids were present in the as-received material, but the surface between voids was quite dense (Fig. 2(a)). The initial effects of surface corrosion are seen as material loss across the surface after 500 hr at 1050 °C (Fig. 2(b)). The grain-boundary glass phase is the first material removed from the surface, revealing clearly mullite's needle-like structure. Surface corrosion at 1250 °C is more severe. After 500 hr (Fig. 2(c)) the glass is no longer evident and actual corrosion of the mullite grains is seen.
Corrosion of the mullite was so severe at 1250 °C that formation of silica whiskers was frequently observed near the outlet end of the furnace tube. The formation of gaseous SiO and H₂O occurs when SiO₂ reacts with hydrogen according to the following equation:

\[
\text{Si}_2\text{(s)} + \text{H}_2\text{(g)} \rightarrow \text{SiO(g)} + \text{H}_2\text{O(g)}
\]  

(3)

It is assumed this same reaction is occurring to the aluminosilicate glass located in the grain boundaries. Corrosion of the mullite grains seen at 1250 °C can also contribute to the formation of SiO by the reaction shown in Eq. 1. The SiO from these reactions is removed from the reaction layer by the bulk gas and transported to the cooler outlet end of the furnace tube where conditions are favorable for reoxidation to SiO₂. Removal of SiO₂ from the glass phase and the mullite grains leaves regions of porous alpha alumina behind. The resulting alumina-rich surface was analyzed by XEDS, and Fig. 3 shows the increase in aluminum/silicon XEDS peak height ratio in going from the as-received material to the exposed material after 125 and 500 hr at 1250 °C.

Removal of the SiO₂ also contributes to a drop in density and an increase in porosity near the surface. Figure 4 shows the effect of exposure time on the density of samples heated in hydrogen at 1250 °C. The density decrease seems linear with time up to the maximum of 500 hr studied. The porous surface layer formed by hydrogen corrosion is shown in the back-scattered electron SEM micrographs of Fig. 5. Figure 5 (a) is a cross-section of an as-received mullite specimen with the top surface marked X. Several large voids (10 to 15 μm in diameter) visible in this micrograph, were also present in all the exposed samples (Figs. 5 (b to d)). During exposure to H₂, escape of SiO (g) produced by reaction (Ref. 3) leaves porous regions in the material. The extent of the porosity depends on both the time and temperature of the exposure. At 1050 °C, exposure for 500 hr produces a porous layer only about 4 μm deep marked by Y in Fig. 5 (b). At 1250 °C, exposures of 125 and 500 hr produce porous surface layers. The layer is 150 μm deep after 125 hr and increases to a depth of 375 μm after 500 hr. Chen (Ref. 3) found the reaction rate to depend on the position of the bars in the furnace and the flow rate of the hydrogen. This implies that removal

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1 Backscattered electron (BSE) imaging is used to image glass removal. During mounting for ceramographic preparation, epoxy infiltrates the sample, filling pores where the glass was removed. In BSE imaging, contrast is provided by atomic number differences. The epoxy is composed of carbon, atomic number 6, and the mullite has an average atomic number of 9.85. Therefore, the epoxy filled voids show up darker than the mullite.
of reaction products from the boundary layer, not transport through the porous reaction layer, is the rate controlling step in the corrosion process.

Fracture strength data for mullite bars exposed to hydrogen at 1050 and 1250 °C are shown in Fig. 6. Data for short time exposures in helium is also presented in the same figure. At room temperature, the fracture strength of as received mullite bars was approximately 211 MPa. All 1050 °C exposures produced a significant strength loss. Most of this strength loss occurred in the first 25 to 50 hr, and was independent of exposure atmosphere, H2 or He. After 500 hr in hydrogen the measured strength loss averaged 22 percent. The slight strength recovery, from =150 MPa to =167 MPa, observed between 125 and 500 hr exposure time is not believed to be of great significance. The loss in strength after 500 hr of exposure is attributed primarily to formation of a thin porous reaction layer due to corrosion of the grain-boundary phase between the mullite grains. Because of the different strength behavior of bars exposed at 1250 °C, to be described later in this report, an attempt was made to rationalize the sudden loss in strength for short exposure times at 1050 °C in both hydrogen and helium through microstructural characterization.

Transmission electron microscopy revealed the presence of a faulted crystalline grain-boundary phase in samples exposed at 1050 °C for 500 hr (Figs. 7 (a) and (b)). This phase is present after both helium and hydrogen exposures. Strength reductions of 22-28 percent were observed when the crystalline grain-boundary phase was present regardless of surface corrosion. Dilatometric analyses indicate that the formation of the crystalline phase does not alter the thermal expansion behavior. Differential thermal analysis traces in air and helium were featureless. In contrast to 1050 °C, samples heated to 1250 °C, even for 500 hr, contain no crystalline grain-boundary phase (Figs. 7 (c) and (d)). Figs. 8 (a) and (b) shows the corresponding XEDS for the grain-boundary and the superimposed mullite phase in K-635 mullite for each temperature. Comparison of Figs. 8 (a) and (b) suggests that the crystalline grain-boundary phase in the 500 hour/1050 °C sample was rich in calcium and also contains some sodium in addition to aluminum and silicon. To identify this grain-boundary phase, selected area diffraction (SAD) patterns were recorded from several crystalline grains in different orientations. In each case, the SAD pattern could be unambiguously indexed to a calcium aluminosilicate hydride called beidellite, [(Ca, Na)2 Al2Si4O10(OH)2]. H2O(s).

Beidellite has an orthorhombic crystal structure with unit cell parameters: a=0.518 nm, b=0.897 nm, and c=1.76 nm. The representative SAD patterns along zone axes [121], [421], and [210] are shown in Figs. 9 (a) to (c)
respectively. Beidellite is a clay mineral which belongs to the montmorillonite group and is capable of absorbing water molecules between its sheet structure (Ref. 7).

Many high density ceramic compositions contain a glassy grain-boundary phase which, when exposed to high temperatures, can devitrify to a variety of crystalline phases. In sintered alumina materials it has been found that calcium facilitates the crystallization of the glassy grain-boundary phase (Ref. 8). It was therefore logical to see if calcium was the cause of the strength reducing recrystallized grain-boundary glass observed in the sample noted above. A sample of another high density mullite material, designated mullite-I,3 which contained no detectable amounts of calcium was exposed to dry hydrogen for 50 hr at 1050 °C. TEM of this sample, Figure 10, reveals only amorphous grain-boundaries. Figure 11 shows the XEDS of such a grain-boundary. No calcium is detected. This observation led us to conclude that as with alumina, calcium plays a significant role in the recrystallization of the grain-boundary glass phase at 1050 °C. The presence of this recrystallized grain-boundary phase seems detrimental to the high temperature strength of mullite.

A different strength behavior was observed for samples exposed at 1250 °C. Short time exposure (≤ 50 hr) to both hydrogen and helium resulted in a sharp strength increase of about 38 percent. For times > 50 hr, strength of mullite dropped continuously, but bars exposed to hydrogen for 125 hr were still stronger than the baseline material. The increase in strength at short exposure times is suspected to be related to crack healing promoted by the flow of silicate glass on the surface and needs further study. Strength degradation after 500 hr at 1250 °C was approximately 53 percent. Crystallization of the glassy, grain-boundary phase that led to the strength degradation at 1050 °C was not observed. The strength loss after exposure at 1250 °C is attributed to severe corrosion and actual removal of the glassy grain boundaries as shown in Fig. 5.

**SUMMARY**

Near stoichiometric mullite was exposed to pure hydrogen at 1050 and 1250 °C for times up to 500 hr.

1. Exposure to hydrogen for 500 hr at 1050 °C reduced room temperature strength by 22 percent. At 1250 °C the corresponding loss was 53 percent.

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1Ispen Ceramics, Pecatonica, IL.
2. Strength loss after long exposure times is related to the removal of SiO₂ from the grain boundaries (and from the mullite grains at 1250 °C).

3. Strength loss after short exposures at 1050 °C is due to the formation of a crystalline grain-boundary phase, beidellite.

4. In hydrogenous atmospheres, the presence of calcium promotes the crystallization of the grain-boundary phase, beidellite at 1050 °C.

5. Strength enhancement after short exposures at 1250 °C is suspected to be due to healing of surface flaws due to flow of the silicate glass phase on the surface.

CONCLUSIONS

Room temperature fracture strength of the mullite used in this study is severely degraded by long time exposure to hydrogen at 1050 and 1250 °C. Corrosion of the grain boundaries occurs at both temperatures and at 1250 °C, even the mullite grains are slightly attacked. The presence of calcium promotes the intermediate temperature (1050 °C in this case) recrystallization of the glassy, grain-boundary phase which degrades the strength of mullite even at short exposure times. These results indicate that the calcium content of mullite intended for use at temperatures ca 1050 °C should be as low as possible. Pure mullite without the presence of a secondary glass phase may be suitable for use in hydrogen below 1250 °C. Production of mullite without a glass phase is, however, extremely difficult.

REFERENCES


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Figure 1.—Ceramograph of as-received mullite K-635, etched with 2:1 HCl:H2O.
Figure 2.—SEM micrographs of mullite K-635 bar surfaces after exposure in hydrogen. (a) As-received. (b) 500 hr, 1050 °C. (c) 500 hr, 1250 °C.
Figure 3.—Surface Al/Si ratios (XEDS) of mullite exposed to hydrogen at 1250 °C.

Figure 4.—Density of mullite K-635 as a function of hydrogen exposure at 1250 °C.
Figure 5.—BSE micrographs reveal porous reaction layer of mullite K-635 after hydrogen exposure. (a) As-received. (b) 500 hr, 1050 °C; $Y = 4 \mu m$. (c) 125 hr, 1250 °C; $Y = 150 \mu m$. (d) 500 hr, 1050 °C; $Y = 300 \mu m$. 
Figure 6.—Room temperature fracture strength of mullite K-635 as a function of exposure time.
Figure 7.—TEM micrographs of grain-boundary phase in mullite K-635 after hydrogen exposure. M = mullite, CG = crystallized glass, G = glass. (a) 500 hr, 1050 °C; bright field. (b) 500 hr, 1050 °C; dark field. (c) 500 hr, 1250 °C; bright field. (d) 500 hr, 1250 °C; dark field.
Figure 8.—XEDS of grain-boundary phase in mullite K-635 after hydrogen exposure. (a) 500 hr, 1050 °C. (b) 500 hr, 1250 °C.
Figure 9.—Representative selected area diffraction patterns of crystalline phase formed in mullite K-635 at 1050 °C. Simulated zone axis patterns verify match with beidellite [(Ca, Na)₂Al₂Si₄O₁₀(OH)₂]·H₂O.

(a) Zone axis [121]. (b) Zone axis [421]. (c) Zone axis [210].
Figure 10.—TEM micrographs of mullite I after hydrogen exposure at 1050 °C. (a) Bright field. (b) Dark field.

Figure 11.—XEDS of grain-boundary phase in mullite I after hydrogen exposure at 1050 °C.
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**Abstract:**
This study deals with the corrosion of near stoichiometric mullite (3Al₂O₃–2SiO₂) by pure dry hydrogen gas. Exposure of the mullite samples was at temperatures of 1050 and 1250 °C for times up to 500 hours. Preferential attack of the aluminosilicate glass present in the grain boundaries of the mullite occurred after 125 hours at 1250 °C. Hydrogen scrubbing of the SiO₂ from the glassy grain boundaries and the mullite grains yielded a porous alumina-rich surface. The room temperature strength increased after short exposure times at 1250 °C (up to 125 hours), then decreased by 53 percent after exposure for 500 hours. At 1050 °C, all exposure times (25 to 500 hours) decreased the strength. After 500 hours in hydrogen at 1050 °C, the room temperature strength of mullite decreased 22 percent. We also observed a rapid 25 percent strength loss after short exposure times at 1050 °C. This is attributed to the calcium/hydrogen assisted crystallization of the glassy, grain-boundary phase.