A Comparative Study of Double Layers Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$ Prepared By Microwave and Natural Drying

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Abstract: In this work, a comparative study of green double layers ceramic bodies prepared by natural and microwave drying techniques was investigated. The double layers ceramic with combination of Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$ samples were prepared via compaction method. Alumina powder, Al$_2$O$_3$ with particle size 90μm and <45μm, silica powder, SiO$_2$ with particle size <45μm were used in this study. Measurement of linear shrinkage of pellet shows that microwave drying contributed to lower shrinkage value compared to the natural drying method. The characterization of the microstructure of double layers ceramic body also showed that microwave technique formed a good bonding structure especially at the critical interface layer. In fact, the implementation of microwave drying also significantly reduced the drying time and produced a good and homogenous final structure.

Keywords: Ceramic pellet, microwave drying, microstructure, linear shrinkage

1. Introduction

Most of the ceramic component that used in varies application i.e membrane, mould for casting and wall consists of multilayered structure [1,2]. Basically, ceramic double layers structure is fabricated with two similar and dissimilar materials and these layers can be connected by the diffusion bonding [3]. Diffusion bonding of layered structures can be established via solid state or liquid state sintering process. This sintering stage is needed in order to connect the two similar and dissimilar materials. During diffusion stage several physical and metallurgical processes occur simultaneously, such as creep, diffusion, dislocation formation, recrystallization and formation of reaction product at the interface. Typically, bonding of the ceramic interfaces depending on the materials used such as alumina, silica, and titania.

Drying is one of the most important processes and complicated process for ceramic materials due to the fact that ceramic body will typically shrink during firing [4-7]. This shrinkage phenomenon can lead to cracking mechanism and loss the quality of the final products. A crucial task in the drying process of ceramic green bodies is the minimization of non uniform drying stresses in order to prevent cracking and non uniform deformation in the green body structure [8]. Drying can be defined as a process of removing water from unfired body or raw materials in the green body. Drying will be accomplished by supplying energy to the ceramic in order to complete evaporation and drying by evaporation is commonly applied to a large number of materials for instance ceramic [9, 15]. Drying process can change the microstructure of materials which consequently affect the characteristic of the dried materials. Some of the materials must be dried slowly and carefully to avoid crack, some have to be dried as thoroughly as possible and others need very low temperature [16]. Several methods have been utilized for the purpose of ceramic drying starting with conventional drying method such as natural air drying and oven drying. However, these methods are not always sufficient. Over the past years, the microwave drying method is becoming more and more important regarding to advance or faster drying [10]. Besides, the microwave processing of materials offers the potential of reducing production time of ceramic materials and this advantage of speed up of drying process can lead up the reduction of energy requirement in processing [11]. Basically, the conventional drying is slow in thick materials and can cause the drying process taking the longer time [12]. The natural drying is not only slow but nonuniform where the surface is hotter than inside. In the previous studies [13, 14], the microwave drying has been proved that this technique offered the advantage especially in drying acceleration and during microwave drying, the samples is heated from inside to the outside [17]. Thus, this technique is needed for the thick materials especially for double layers structure due to the microwave can penetrate internally and the interior part become hotter than its surface. This energy can heat materials faster than other drying method and the microwave drying offered possibility of predicting higher reduction in drying time.

In this study, the double layers ceramic pellet were fabricated from Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$ using powder compaction technique and then dried with different drying modes; microwave and natural. Double layers of Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$ ceramics are expected to be used in a wide range application, such as for filters or membrane for gases and liquids.
of semiconductor industries and structural products [18]. Basically, ceramic membranes consisting of several layers of one or more different ceramic materials [19]. The aim of this study was to investigate the influence of drying technique to the microstructure of the ceramic layer and properties. The results presented provide a basic understanding the effect of drying on the double layers ceramic bonding sintered structure.

2. Experimental procedure

2.1 Sample Preparation

A commercial alumina powder (purity of 99%, Sigma Aldrich) and silica (industrial grade) were used in the preparation of double layers ceramic. Then, the alumina and silica powder with different particle size were added with 5% binder, polyvinyl alcohol (PVA) and the mixture was milled for 30 minutes by using ball mill until the mixture was homogenized. The two groups of double layers pellets Al₂O₃/Al₂O₃ and Al₂O₃/SiO₂ were fabricated with two similar and dissimilar materials see Fig.1. The particle sizes of 90 μm and below 45 μm were used for double layers Al₂O₃/ Al₂O₃ structure whereas alumina (90 μm) and silica (below 45 μm) were used for Al₂O₃/SiO₂ pellets. The pressure applied on the samples was 3 tons for 3 minutes for each of the layer.

![Fig. 1 (a) and (b) Schematic representation of ceramic double layered structures](image)

2.2 Drying Conditions

After the desired double layers samples were fabricated, the samples were dried by using microwave and natural drying. Under microwave drying, the samples were removed periodically from the microwave for every 10 minutes. This procedure was repeated until the desired moisture content (constant in weight) was achieved. The time taken for the samples to reach desired moisture content also was recorded. On the other hand, natural drying also was carried out at room temperature for 24 hours.

After drying process, the green bodies were sintered in a temperature programmable furnace. The sintering involves two stages, the first stage is 500°C in order to burn out the binder, PVA and followed by densification of ceramic bodies at 1550°C, both with the heating rates of 2°C/min. The samples were soaked for 3 hours during the densification process.

2.3 Properties Characterizations

The sintered double layers ceramic structures were characterized by physical properties and microstructural analysis. The linear shrinkage of double layered structures was determined using the following equations:

\[ \text{Shrinkage} = \frac{d_i - d_f}{d_i} \times 100\% \quad (1) \]

where \( d_i \) is the diameter of sample before drying and \( d_f \) is the diameter after drying of green and sintered double layered structure. The diameter was measured by vernier caliper (Mitutoyo) and average values were taken.

Archimedes’ method was used for calculation of the density and porosity of the sample after sintering process. The measurement of density and porosity were carried out by using Mettler Toledo Density Kit. The density and porosity of the ceramic pellets were calculated by using mass of dried sample, \( W_d \), mass of immersed and suspended specimen in liquid, \( W_i \) and mass of immersed and suspended specimen in air, \( W_w \).

\[ \text{Porosity} = \frac{W_w - W_d}{W_w - W_i} \times 100\% \quad (2) \]

\[ \text{Density} = \frac{W_i}{W_d - W_i} \times 100\%. \quad (3) \]

Scanning electron microscope, SEM (JEOL, JSM-638OLA) was used to examine the cross section of the sample. The sample micrographs were observed by using back scattered electron mode using 100 × magnifications.

3. Experimental procedure

3.1 Linear Shrinkage

Shrinkage phenomenon occurs in ceramic bodies during water removal in drying and sintering process which sometimes can result in physical defects in ceramic products such as crack and warping. Percentage of shrinkage and percentage of expansion for sintered double layers pellet between natural drying and microwave drying is shown in Fig. 2. Fig. 2 (a) shows that the double layers Al₂O₃/Al₂O₃ undergoes shrinkage while the double layers Al₂O₃/SiO₂ undergoes expansion after the sintering process. This graph shows that the microwave drying gives the lowest value of drying shrinkage for two similar materials, Al₂O₃/Al₂O₃. Typically, the drying process in microwave is more uniform due to the penetration depth [15, 20] of the microwave radiation and allows the shrinkage take place from the centre the surface of the material. As shown in Fig. 2 (a), double layers Al₂O₃/Al₂O₃ structure has a stable percentage for both drying method meanwhile the
percentage of expansion for double layer Al$_2$O$_3$/SiO$_2$ significantly different expansion between both of the techniques. The Al$_2$O$_3$/SiO$_2$ structure undergoes expansion due to the sintering temperature. This double layered structures were sintered at 1550°C (based on sintering temperature of Al$_2$O$_3$) and this causes the molten silica flow into the alumina structure. This is due to the fact that silica has slightly lower sintering temperature. In this case also the microwave drying gives a higher percentage of expansion compare to the natural drying technique due to the fact that this drying method also allows the particles stacking closed together which provide a good path for the diffusion process in the later stage of sintering process.

where an increase in shrinkage results in the reduction of porosity in the sintered ceramic bodies. As expected, the different double layers structure give different porosity and density. It showed that the natural drying produced the higher porosity and lower density for double layers Al$_2$O$_3$/SiO$_2$. Density and porosity are interrelated properties in ceramic materials. Theoretically, ceramic bodies with higher porosity will have a low density. The decrease of the bulk density is due to the increase of the porosity in sintered ceramic bodies. The finding reveals that shrinkage phenomenon is strongly related to the porosity and density.

Fig. 2 (a) Percentage of shrinkage for double layers Al$_2$O$_3$/Al$_2$O$_3$ and (b) percentage expansion for double layers Al$_2$O$_3$/SiO$_2$

3.2 Porosity and Density

Table 1 displays the percentage of apparent porosity and bulk density for sintered double layers Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$ structure. Table 1 shows that microwave drying produced higher porosity for sintered double layers Al$_2$O$_3$/Al$_2$O$_3$ structure compared to natural drying. Basically, shrinkage has an influence to the porosity

Table 1 Porosity and density of double layers Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$

<table>
<thead>
<tr>
<th>Property</th>
<th>Al$_2$O$_3$/Al$_2$O$_3$</th>
<th>Al$_2$O$_3$/SiO$_2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drying</td>
<td>Porosity (%)</td>
<td>Density (g/cm$^3$)</td>
</tr>
<tr>
<td>Microwave</td>
<td>33.47</td>
<td>3.60</td>
</tr>
<tr>
<td>Natural</td>
<td>31.85</td>
<td>3.69</td>
</tr>
</tbody>
</table>

3.3 Microstructural Analysis

SEM (Fig. 3 and 4) micrographs with EDS analysis (Fig.5) present the microstructure of sintered double layers structure Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$. In natural drying, the particles are less agglomerated to each other and obviously create two separated layers for both cases while microwave offered uniform drying which caused the particle stacked closely together. This is clearly shown in Fig. 3(b) and 4(b) where the layer obviously separated for both double layers (Al$_2$O$_3$/Al$_2$O$_3$ and Al$_2$O$_3$/SiO$_2$) with the natural drying due to the uncontrolled drying process. Theoretically, the ceramic materials need a slow and controlled drying stage to avoid the cracking mechanism in ceramic green bodies. The structure obtained in Fig. 3 (a) and 4(a) clearly shown that both double layers structure have good linkage between layers with no crack and deformation in the sample. In natural drying, larger grains were obtained in the sintered double layered ceramic structure and this is opposite from the microwave drying where the application of microwave has successfully produced smaller grain in the sintered double layers structure. These result indicated that the microwave absorption characteristics vary depending on the microstructure where the finer and better microstructure can be produced by the microwave processing.
Fig. 3 Cross sectional images of sintered double layers ceramic structure (Al$_2$O$_3$ / Al$_2$O$_3$) that dried using different technique; (a) microwave, (b) natural.

Fig. 4 Cross sectional images of sintered double layers ceramic (Al$_2$O$_3$ / SiO$_2$) that dried using different technique; (a) microwave (b) natural.
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![Fig. 5 EDS mapping for double layers ceramic Al₂O₃ / SiO₂](image)

4. Conclusion

The application of microwave technique in drying double layers ceramic structure has potential to improve the properties of the structure. This is due to the several advantages of microwave drying such as shorter drying time which is good for thick samples especially for double layers structure and more uniform over the cross section part than other natural drying method. This is due to the penetration depth of the microwave radiation that can cause the water evaporated more uniform either at surface or centre. Meanwhile the uniform drying can lead to a homogenous temperature distribution and causes the shape of the sample maintains in the most of the structure. The finding of this study revealed that the microwave drying improves the properties of Al₂O₃/Al₂O₃ and Al₂O₃/SiO₂ double layered structures.

REFERENCES


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