Two-Step Sintering of Ceramics with Constant Grain-Size, I. Y₂O₃

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Two-Step Sintering of Ceramics with Constant Grain-Size, I. \(Y_2O_3\)

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Isothermal and constant-grain-size sintering have been carried out to full density in \(Y_2O_3\), with and without dopants, at as low as 40% of the homologous temperature. The normalized densification rate follows Herring’s scaling law with a universal geometric factor that depends only on density. The frozen grain structure, however, prevents pore relocation commonly assumed in the conventional sintering models, which fail to describe our data. Suppression of grain growth but not densification is consistent with a grain boundary network pinned by triple-point junctions, which have a higher activation energy for migration than grain boundaries. Long transients in sintering and grain growth have provided further evidence of relaxation and threshold processes at the grain boundary/triple point.

1. Introduction

Sintering is a complicated process of microstructure evolution, with the main outcome being porosity elimination.\(^1\)\(^4\) However, in polycrystals accelerated grain growth is also reckoned to always accompany final-stage sintering. The onset of accelerated grain growth depends on the microstructure; a uniform size and shape distribution of pores and grains postpones accelerated grain growth.\(^6\)\(^7\) In the best case the onset can be postponed to about \(\rho \approx 92\%\) which, for most ceramics with a dihedral angle of 120°, coincides with the Rayleigh transition when pore channels collapse to become isolated pores.\(^7\)\(^9\) Such collapse results in a substantial decrease in pore pinning, which triggers accelerated grain growth.\(^10\)\(^11\) The prevailing sintering paradigm in the last 40 years is to mitigate this undesirable outcome by suppressing grain boundary mobility while promoting pore pinning. The aim is to maintain pore-grain boundary contact during grain growth so that pore breakaway never occurs.

Contrary to the above paradigm, we have used a simple two-step sintering method to sinter \(Y_2O_3\), from 75% to 100% density, without grain growth.\(^1\)\(^2\) The key elements in this method are (a) reaching a higher temperature \(T_1\) to conduct first-step sintering, (b) achieving a high-density \(\rho^*\) at \(T_1\); \(\rho^* \geq 75\%\) to render pores unstable,\(^1\)\(^2\) and (c) lowering the temperature to \(T_2\) to conduct second-step sintering during which there is only densification and no grain growth. The purpose of the present paper is to analyze in detail the sintering kinetics of \(Y_2O_3\), a model system, in order to shed light on the sintering mechanism. The relevance of the present work to low-temperature kinetics and nanograin ceramics will also be explored. In addition to \(Y_2O_3\), this method has been successfully applied to BaTiO\(_3\) and Ni-Zn ferrite, which densify by solid-state diffusion, as shown in the companion paper.\(^1\)\(^4\) It also succeeded in liquid-phase-containing ceramics, \(ZnO\) and SiC, as reported in the recent literature.\(^15\)\(^16\)

Analysis of sintering data requires a sound physical model. Although theoretical models usually use idealized geometric assumptions which may not hold in real materials,\(^6\)\(^10\)\(^11\)\(^17\)\(^20\) a general dimensional argument originated by Herring\(^27\)\(^28\) states that the normalized rate of densification, via grain boundary diffusion, can be expressed as

\[
\frac{dp}{dt} = F(\rho) \frac{3\gamma \Omega D}{kT} \frac{\delta}{G^2} \quad (1)
\]

Here \(t\) is the time, \(\gamma\) the surface energy, \(\Omega\) the atomic volume, \(k\) the Boltzmann constant, \(T\) the absolute temperature, \(G\) the mean grain diameter, \(\delta\) the width of the grain boundary, \(D\) the grain boundary diffusivity, and \(F(\rho)\) an unspecified function of density \(\rho\). This argument holds for different grain size and density provided the sintering microstructure, including the pore distribution, is a function of density only once the length scale is renormalized by the grain size. For a ceramic undergoing two-step sintering, the grain boundary network is frozen, so there is no change in microstructure other than the shrinking porosity. This would seem to satisfy Herring’s assumption. Therefore, we will attempt to use Eq. (1) in our analysis.

Various models of intermediate and final-stage sintering have specified the function \(F(\rho)\). Typically, at \(\rho > 0.7\), the predicted \(F(\rho)\) is a gradual, decreasing function of \(\rho\), which was supported by the data of Hansen et al.\(^2\) On the other hand, Zhao and Harmer\(^2\) found, after accounting for grain-size increase, a \(F(\rho)\) that slightly increased with density in final-stage sintering. They attributed the increase to the increased number of pores that intersect a typical grain boundary as sintering proceeds. Cameron and Raj,\(^3\) however, found qualitative disagreement between their data and all the geometrical models they examined. Instead, an empirical model that depicts densification as an exhaustion process seemed to fit their data the best.

The attainment of constant grain size throughout intermediate and final-stage sintering in second-step sintering offers an excellent opportunity to verify Eq. (1) and evaluate \(F(\rho)\), which is a main objective of our work. Once Eq. (1) is confirmed, any deviation of the experimental data from the expected behavior of Eq. (1) would imply important changes in the underlying sintering kinetics.

II. Experimental Procedure

(1) Powder Preparation

The \(Y_2O_3\) powders were synthesized by a standard solution-polymerization route using yttrium nitrate (Y(NO\(_3\))\(_3\), Alfa, New York, NY) and polyethylene glycol (PEG, FW = 2000, Alfa) as raw materials. The polymeric precursor for the \(Y_2O_3\) was prepared by mixing an appropriate amount of nitrate salt and PEG into distilled water. After heating and stirring at 80°C, a transparent aqueous precursor solution was obtained. The precursor solution was then heated until the solution evaporated, leaving a soft, light brown, aerated gel. The dried gel was ground with an agate mortar and pestle, then calcined in air at various temperatures to obtain \(Y_2O_3\) powders with different particle sizes in the range of 10–60 nm.

M. Harmer—contributing editor
Mg$^{2+}$ and Nb$^{5+}$ were used as cation dopants substituting Y at 1 at.%. First, Y$_2$O$_3$ powders were dispersed in isopropyl alcohol. A solution containing Mg(NO$_3$)$_2$ was then added into the Y$_2$O$_3$ dispersion and the slurry was floced by increasing the pH using NH$_4$OH. The floced material was dried and later calcined in air at 600°C for 2 h to yield a uniform MgO coating on Y$_2$O$_3$ particles. For Nb doping, an (isopropyl) alcohol dispersion of Y$_2$O$_3$ was first mixed with a solution of niobium isopropoxide, and the mixture was next hydrolyzed by adding deionized water at room temperature. The slurry was dried and calcined at 600°C for 2 h. To avoid SiO$_2$ contamination, powder processing was conducted using plastic ware only, and sintering was performed using a dedicated "clean" furnace.

### (2) Powder Compacts and Sintering

Pellets (diameter = 10 mm, thickness = 10 mm) were prepared by die pressing followed by cold isostatic pressing. Pressure up to 310 MPa was applied to obtain green bodies with different densities. Subsequent sintering was conducted in air.

(A) **Normal Sintering:** Normal sintering up to 1600°C used a constant heating rate of 10°C/min. After reaching the desired temperature, the power was turned off to allow samples to cool naturally in the furnace. The specimen dimensions were measured in situ during heating in a lateral dilatometer using dense alumina as a reference. Isotropic shrinkage was found, which allows the density to be calculated from the current length and the final density measured by the Archimedes method. Density data are presented in fraction or percentage of the theoretical density, taken as 5.013 mg/m$^3$ for Y$_2$O$_3$.

(B) **Two-Step Sintering:** In two-step sintering, the sample was first heated at 10°C/min to $T_1$, then cooled at 50°C/min to $T_2$ and held there from 6 to 30 h. This schedule was chosen to minimize densification during heating and cooling transients. Runs interrupted after the first step or during the second step were performed in order to examine intermediate microstructure. The specimen dimensions were also measured in situ throughout two-step sintering using the same dilatometer.

### (3) Characterization

Initial particle size was determined using transmission electron microscopy, scanning electron microscopy (SEM) and X-ray diffraction. In the Archimedes method for density measurements for samples with only closed porosity, distilled water was used as the displacement fluid and the estimated accuracy was within ±0.01 g/cm$^3$. For each experiment, the average density of three pellets was used in the analysis (their densities were within ±2% of each other). For samples with open porosity, such as after first-step sintering, the density was estimated from the weight and dimension of duplicate specimens; the estimates were consistent with the data of dilatometry. Microstructures of sintered compacts were observed in an SEM using specimens that were either fractured, or polished and then thermally etched. At lower densities, the mean grain (particle) size was estimated on the fracture surface. At higher densities, the grain size was obtained by multiplying the average linear intercept length of at least 500 grains by 1.56.

Dihedral angles were determined using samples sintered in a previous study from a coarse powder (200 nm). These samples had a density of around 85%-90%, and they were subjected to long annealing at low temperatures to develop the dihedral angles between pores and grain boundaries. At least 200 dihedral angles were measured for each material to obtain a statistical distribution.

### III. Results and Data Analysis

**1) Microstructure of Two-Step Sintering**

We have previously published the microstructure of two-step sintered Mg$^{2+}$-doped Y$_2$O$_3$ using powders with an initial particle size of 10 nm and a final dense ceramic grain size of 60 nm. The temperatures used were 1080°C for $T_1$ and 1000°C for $T_2$. To contrast this observation, we here show the microstructure of a set of coarser ceramics obtained by two-step sintering using powders of a much larger size, 200 nm. These powders were prepared by a homogeneous precipitation method using urea to assist precipitation. They were spherical in shape and doped with 1% Mg$^{2+}$. Two-step sintering was conducted using (1) 10°C/min heating to $T_1$ (1422°C) to gain a density of 84%, followed by cooling, and (2) reheating at 20°C/min to $T_2$ (800°C) for 11 h to gain full density. Microstructures of the green body after first-step sintering, and at various times during second-step sintering, are shown in Fig. 1 to demonstrate the absence of grain growth at $T_2$. Throughout the second step a constant grain size of 390 nm was maintained, which is about twice the size of the starting powder. We are not aware of any other sintering method that resulted in a less coarsened microstructure, from starting powder size to dense ceramic grain size, than that in Fig. 1. Using similar two-step sintering methods, we were able to densify a Nb$^{5+}$-doped Y$_2$O$_3$ without grain growth in the second step, starting with a 200 nm powder and reaching a final grain size of 400 nm (data not shown.)

These experiments were initially conducted to search for an appropriate ($T_2$) temperature suitable for thermal etching of partially dense Y$_2$O$_3$ in order to reveal dihedral angles between pores and grain boundaries. Many such experiments yielded ceramics that continued to densify at $T_2$, eliminating most or all the pores. From these experiments we concluded that $T_2$ sintering was feasible for Y$_2$O$_3$ of a grain size of 60-400 nm. In BaTiO$_3$ and Ni-Zn ferrite, constant grain size sintering in the 70-800 nm range was also demonstrated as shown in the companion paper.

Measured dihedral angles for undoped and doped Y$_2$O$_3$ are shown in Fig. 2. They are around 110° regardless of dopants. Therefore, doping is not likely to cause a systematic difference in the microstructure. In particular, the critical size ratio of a pore to neighboring grains that renders the pore thermodynamically unstable in Y$_2$O$_3$ is likely to be similar regardless of dopants. These results justify the use of the same $F(p)$ in the following analysis.

**2) $F(p)$ as a Universal Function**

The densification curves during $T_2$ sintering are described next. Figure 3 shows five sets of data from runs that had no grain growth, although they varied in composition, sintering temperatures ($T_1$ and $T_2$), and starting grain size at $T_2$. According to Eq. (1), if the function $F$ depends on density only, then these
data, plotted in a semi-log form, should fall on curves that are vertically parallel to each other. This is indeed the case.

By plotting normalized densification rate against grain size in a log–log form (not shown), we obtained an exponent of 3.7, which is close to four used in Eq. (1). The normalized densification rate was further used to determine the activation energy of grain boundary diffusion, by rewriting Eq. (1) into

$$\delta D = \frac{d\rho}{dt} \frac{1}{F(\rho)} \frac{kT}{3\pi\Omega} \gamma^4$$  \hspace{1cm} (2)

The obtained activation energies are listed in Table I and compared with the data in the literature. These grain size exponent and activation energies are essentially independent of the density used in the data analysis, which is consistent with the assumption that $F(\rho)$ is a universal function.

According to Hansen et al.,\textsuperscript{26} the value of $F(\rho)$ varies very slightly between $\rho = 0.75$ and 0.85 and is around 12,000 at $\rho = 0.8$. (Here we need to account for a factor three to convert length measurement to density measurement.) Accepting these values, we estimated the grain boundary diffusivity and showed it in Fig. 4. In the above estimate, we have assumed $\gamma$ to be 1 J/m\(^2\) and $\Omega$ $7.74 \times 10^{-29}$ m\(^3\) in Eq. (2). The $\delta D_0$ determined for $Y_2O_3$ is $(2.36 \pm 0.40) \times 10^{-12}$ m\(^3\)/s; for $Y_2O_3$ doped with Mg\(^{2+}\) and Nb\(^{5+}\), $\delta D_0$ are $(1.08 \pm 0.15) \times 10^{-13}$ and $(5.79 \pm 0.11) \times 10^{-17}$ m\(^3\)/s, respectively. These values are comparable with those determined from grain boundary mobilities using grain growth data at higher temperatures (see Table I)\textsuperscript{32,33}

(3) Kinetic Window for Constant Grain Size Sintering

If second-step sintering was conducted at too low a temperature ($T_2$), sintering proceeded for a while (typically for 5%), then became exhausted. The normalized densification rates in Fig. 5 (curves A–C) fall below what is expected from $F(\rho)$, starting at about $\rho = 0.85$. As there was no grain growth at $T_2$, the slowdown may be attributed to grain boundary diffusion that also became exhausted. At higher $T_2$, grain growth was observed during second-step sintering. The normalized sintering rates again fall below what is expected from $F(\rho)$, as is also shown in Fig. 5 (curves D–F), at $\rho > 0.85$. This slowdown is because of increasing grain size.

The temperature ($T_2$) and the grain size of various samples that successfully underwent second-step sintering to reach full density (at least 99% in this paper) without grain growth are marked in Fig. 6 using solid symbols. They are bordered by two lines which delineate the “kinetic window.” The open symbols below the lower boundary line are experiments that showed no grain growth but densification was exhausted; the open circles above the upper boundary line are ones in which grain growth occurred. For each of the latter data, we also draw a horizontal arrow that points to the final grain size when full density was reached. It seems reasonable that all the arrows end at the upper boundary line, as once within the “kinetic window” the grain size should stop growing. A partial list of the successful two-step sintering experiments (reaching nearly full density without grain growth) is given in Table II. It is clear that the lowest $T_2$ required to reach full density was much lower than those reported in the

<table>
<thead>
<tr>
<th>Sample</th>
<th>Activation energy (KJ/mol)</th>
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<tr>
<td>$Y_2O_3$</td>
<td>410</td>
</tr>
<tr>
<td>$Y-1%$ Mg</td>
<td>553</td>
</tr>
<tr>
<td>$Y-1%$ Nb</td>
<td>310</td>
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Table I. Activation Energy for Grain Boundary Diffusion of $Y_2O_3$

Fig. 2. Dihedral angles of undoped and doped $Y_2O_3$.

Fig. 3. Normalized densification rates during second-step sintering without grain growth. Samples were heated to $T_1$ (the first temperature given below) then cooled to $T_2$ (the second temperature given below) and held for the time specified. The grain size during $T_2$ sintering is also specified (in parentheses). A, $Y_2O_3$, 1250°C—20 h (122 nm); B, $Y_2O_3$, 1310°C—20 h (140 nm); C, $Y_2O_3$ (1% Nb\(^{5+}\)), 1450°C, 1200°C—40 h (118 nm); D, $Y_2O_3$, 1310°C, 1100°C—30 h (140 nm); E, $Y_2O_3$ (1% Mg\(^{2+}\)), 1250°C, 1050°C—30 h (116 nm).

Fig. 4. Estimated grain boundary diffusivity against reciprocal temperature.

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was observed but densification was incomplete in A, 1310°C. The universal density function was plotted for easy comparison with the uniparametric density function. Curves are shifted vertically for easy comparison with the universal density function. Solid symbols are ones reaching full density without grain growth; open symbols did not. Data above the upper boundary had Y2O3-8 44 1400 0 88 200 1050 40 99.0 196 Y2O3-7 44 1400 0 88 200 1150 20 99.5 202 Y2O3-6 44 1350 0 85 160 1100 30 99.0 162 Y2O3-5 44 1350 0 85 160 1150 20 99.5 165 Y2O3-4 44 1310 0 80 138 1100 30 99.6 140 Y2O3-3 44 1310 0 80 138 1150 20 99.6 140 Y2O3-2 44 1250 0 78 116 1050 30 99.2 118 Y2O3-1 44 1250 0 78 116 1100 20 100 120 Y–1%Mg-1 44 1250 0 78 116 1050 30 99.6 120 Y–1%Mg-3 44 1250 0 78 116 1050 30 100 118 Y–1%Mg-4 44 1250 0 78 116 1100 20 100 118 Y–1%Mg-2 44 1250 0 78 116 1050 30 100 118

Fig. 5. Normalized densification rate of Y2O3 samples in second-step sintering. Curves are shifted vertically for easy comparison with the universal density function $F(\rho)$. Same notation as in Fig. 3. No grain growth was observed but densification was incomplete in A, 1310°C—10 h; B, 1250°C—10 h; and C, 1250°C—10 h. Grain growth was observed in D, 1310°C, 1300°C—10 h; E, 1310°C, 1250°C—10 h; and F, 1310°C, 1200°C—10 h.

Fig. 6. Temperature ($T_2$) and grain size used for second-step sintering in Y2O3. Solid symbols are ones reaching full density without grain growth; open symbols did not. Data above the upper boundary had grain growth (arrow from initial to final grain size); data below the lower boundary did not fully densify. Triangle represents a one-step sintering experiment at the temperature shown, with grain size starting at the triangle and growing to the arrowhead, when full density was reached.

Table II. Two-Step Sintering of Y2O3 (30 nm Powders)

<table>
<thead>
<tr>
<th>Sample</th>
<th>$\rho_0$ (%)</th>
<th>$T_1$ (°C)</th>
<th>$t_1$ (h)</th>
<th>$\rho_1$ (%)</th>
<th>$G_1$ (nm)</th>
<th>$T_2$ (°C)</th>
<th>$t_2$ (h)</th>
<th>$\rho_2$ (%)</th>
<th>$G_2$ (nm)</th>
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<td>Y2O3-1</td>
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<td>1200</td>
<td>0</td>
<td>75</td>
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<td>99.0</td>
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<td>0</td>
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<td>1100</td>
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thermal history, their grain size trajectories were again not bounded by the boundary lines.

IV. Discussion

To achieve densification without grain growth, grain boundary diffusion needs to be maintained while grain boundary migration is suppressed. It is well known that the kinetics of these two processes are decoupled when solute drag of the grain boundary is involved, because the latter entails lattice diffusion that has a higher activation energy. Indeed, solute drag is an effective strategy for controlling abnormal grain growth in final-stage sintering.2 In our case, however, solute drag cannot be a significant factor as the same phenomena were also observed in (Mg)-doped samples that showed enhanced grain boundary kinetics. We previously suggested that another drag mechanism, because of relatively immobile triple points (including three-grain junction lines, four-grain junction points and their equivalent involving grain/pore junctions), may provide a decoupling mechanism.10,11 This idea is schematically sketched in Fig. 10 that delineates the grain boundary mobility $M_b$ and junction mobility $M_j$, assuming the latter has a higher activation energy. The mobility of the grain boundary network, which is required to allow grain growth in a polycrystal, is given by $(M_b^{-1} + M_j^{-1})^{-1}$. Therefore, the network mobility follows the grain boundary mobility at high temperature and junction mobility at low temperature, and below the temperature when $M_b \sim M_j$, it is essentially frozen despite considerable grain boundary diffusion. Direct observation of triple point drag has been reported by Czubyko et al. in high purity zinc tri- crystals at low temperature.36,37 They found the moving grain boundaries maintained a (triangle) dihedral angle $< 120^\circ$ in order to exert a force on the grain junction to pull it forward. At higher temperatures, the dihedral angle reverted to $120^\circ$, indicating little triple-point drag. The same drag could occur in our samples at $T_2$. Meanwhile, as densification does not involve grain boundary migration, it still proceeds via grain boundary diffusion.

The crossover depicted in Fig. 10 and the kinetic windows shown in Figs. 6 and 9 provide helpful guides to experimental design but are probably not unique in view of the evidence of history dependence and thermal hysteresis/transient in our study. Such evidence includes (a) slowdown and exhaustion of diffusion at low temperature, (b) incubation time for grain growth to initiate after constant grain-size sintering, and (c) thermal history dependence of the kinetic window. This may be caused by the slow equilibration of defect configurations on the grain boundaries, pore surface, and pore/grain junctions at low temperatures. Therefore, at intermediate times, long transients and possibly metastable states may reign. Meanwhile, pre-firing at $T_1$ may have locked pore/grain junctions into stable configurations that are not kinetically accessible at $T_2$, and the grain

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Fig. 7. Density and grain size of Y$_2$O$_3$ (with 1% Mg$^{2+}$) sintered in the second step at 1000°C for 60 h. Sample previously preheated to 1120°C ($T_1$) without holding.

Fig. 8. Density and grain size during heating (10°C/min) of undoped and doped Y$_2$O$_3$. Initial powder size = 30 nm. See text for highest temperature reached.

Fig. 9. Maps for second-step sintering for Y$_2$O$_3$ with 1% Mg$^{2+}$ (squares) and Y$_2$O$_3$ with 1% Nb$^{5+}$ (diamonds). Same notation as in Fig. 6.
boundary movement may entail the transition states of these configurations that have activation barriers insurmountable at T₂. As a result, pre-firing at T₁ followed by sintering at T₂ may enhance the pinning effects and make possible constant-grain-size sintering.

Another possible reason that a higher T₁ is required for two-step sintering is the different sintering mechanisms below and above the critical density. Below the critical density, particle coarsening and repacking (not unlike that in liquid-phase sintering) are the mechanisms responsible for densification, because otherwise the pores should grow as expected from the theory of Kingery and Francois. Although particle coarsening is achieved by surface diffusion, which typically has a lower activation energy than grain boundary diffusion, some repacking events could require more robust kinetics available only at a higher temperature. If so, a higher T₁ would be needed. On the other hand, once the critical density is reached, grain boundary diffusion alone is sufficient to cause densification even if the microstructure is frozen, as illustrated by our experiments. Therefore, a lower T₂ will suffice as long as it is not too low to exhaust diffusion.

The temperature range in the kinetic windows depicted in Figs. 6 and 9 widens with grain size. This shape has also been observed in BaTiO₃ and Ni–Zn ferrite, suggesting that it is a general feature for T₂ sintering. If so, we may conclude that a larger grain size may postpone the exhaustion of diffusion at lower T₂ and the non-parabolic grain growth at higher T₂. As the driving force for grain growth decreases with grain size, the delay of grain growth by increasing grain size is reasonable. Meanwhile, the delay of diffusion exhaustion by increasing grain size is consistent with the common notion that grain boundary diffusion may be limited by source/sink actions (e.g., kinks and ledge required to provide and accommodate atoms and point defects) at small grain sizes. As diffusion proceeds, some of these kinks and ledge are eliminated by migrating atoms/point defects; after that diffusion may be exhausted if sources/sinks are not replenished by fresh nucleation. The size effects on nucleation are well known; for example, a source/sink in the form of a screw dislocation loop entails a nucleation energy that is inversely proportional to the grain size, so that a threshold stress on the order of 2γ/G, amounting to 20 MPa for a grain size of 100 nm, is obtained. Such sluggish kinetics for source/sink nucleation may be eventually overcome after a long incubation time. According to Fig. 7, this requires 10 h at 1000°C with Mg as ‘‘sintering aid.’’ Such a long incubation time is not entirely surprising given that Y₂O₃ has a melting point of 2440°C, so most of our T₂ experiments fell below or were around 0.5 of the homologous temperature.

We now return to the universal sintering function F(ρ). Unlike the previous reports of Zhao and Harmer (p > 90%) and Hansen et al. (75% < ρ < 95%), our F(ρ) decreases much more rapidly, by a factor of about 4.5 from ρ = 0.75 to 0.9, and by another factor of about 10 from ρ = 0.9 to 0.99. In contrast, Hansen et al. found a decrease of no more than a factor of three in total, and Zhao et al. reported a slight increase. Interestingly, our data in Fig. 3 are empirically consistent with an exponential decay, i.e., F(ρ) = exp(−cρ) where c is a rather large constant, around 24 for ρ > 0.88. This form of F(ρ) has been previously noticed by Cameron and Raj. To understand these differences, we note that in normal sintering, the microstructure is constantly refreshed by grain growth, bringing the remaining pores to coalesce and then to relocate at the newly formed four-grain junctions. As a result, the diffusion distance in normal sintering could remain commensurate with the characteristic grain size even as densification progresses. In our experiments, however, the grain boundary network is frozen. Therefore, as pores around smaller grains that have a shorter diffusion distance are removed first, the remaining pores around larger grains that have a longer diffusion distance remain and become dominant. The effective pore spacing thus constantly moves toward the larger-size end of the grain size distribution, which may be responsible for the exponential-like decrease of F(ρ). This decrease is very steep, because an increase in the pore spacing by a factor of two lowers the normalized sintering rate by a factor of 16 according to Herring.

Despite the rapid decrease of F(ρ), constant-grain-size sintering at T₂ still offers the advantage of lower sintering temperature. Using powders of the same initial particle size, the lowest T₂ achieved in our experiments was always lower than the temperature of conventional isothermal sintering. For example, for 30 nm powders, the lowest T₂ successfully used for Y₂O₃ was 1050–1100°C, whereas isothermal sintering at 1200°C for 10 h was required to reach 99% density. Likewise, for Mg-doped Y₂O₃ at the same initial powder size, the lowest T₂ successfully used was 1000–1050°C, whereas isothermal sintering at 1100°C for 10 h was required to reach full density. This illustrates the advantage of lowering the sintering temperature to maintain a constant grain size once the critical density is reached. It is also clear that further advances in sintering dense nanograin ceramics may come from using finer powders and/or other processing methods to achieve the critical density with an even finer microstructure. In the companion paper, we provide one such example using high green density (61%) to lower T₁.

Lastly, we acknowledge that the discrepancy in F(ρ) between our data and the previous studies makes our earlier estimate of δDₑ₀ based on the model of Hansen et al. inaccurate. However, the uncertainty here is of the order of 10–50, which is well within the scatter of typical diffusion data. Therefore, our conclusion that second-step sintering proceeds by grain boundary diffusion should remain valid.

V. Conclusions

1. Two-step sintering in Y₂O₃ with and without a dopant proceeds by grain boundary diffusion. The suppression of grain growth may be attributed to triple point immobility, which is facilitated by high-temperature (T₁) pre-firing.

2. Constant grain-size sintering of Y₂O₃ to nearly full density was achieved at 800°C, which is less than 0.4 of the homologous temperature. The kinetic window for second-step sintering has a characteristic shape, spanning a wider temperature range with increasing grain size. At higher temperatures, grain growth occurs, for which the driving force diminishes as grain size increases. At lower temperatures, diffusion manifests long transients and eventually comes to exhaustion especially at smaller grain sizes. Dopants may enhance or suppress the overall kinetics and shift the temperature range of the kinetic window, but not affect the above general features.

3. The pre-factor in the Herring’s scaling law in constant grain size sintering follows an exponential decay. It implies a gradual increase of pore spacing. This is because pores around...
larger grains are more difficult to sinter, and they become dominant later.

4. Aways two-step sintering to $Y_2O_3$, a coarsening ratio of two, from powder particle size to dense ceramic grain size, was achieved using 200 nm powders. This ratio was 6 for 10 nm powders, which is still much lower than in conventional sintering. Therefore, the two-step method offers a promising approach for fabrication of bulk nanograin ceramics.

References


