Lawrence Berkeley National Laboratory

Recent Work

Title

CREEP DURING SINTERING OF POROUS COMPACTS

Permalink

https://escholarship.org/uc/item/38w8b7hw

Authors

Rahaman, M.N. Janghe, L.C. De Hsueh, C.H.

Publication Date

1985-03-01

LBL-19360

63

LBL-19360 Preprint RECEIVED

LAWRENCE

BERKELEY LAROPATOR

MAY 22 1985

LIBRARY AND DOCUMENTS SECTION

Submitted to the Journal of the American Ceramic Society

CREEP DURING SINTERING OF POROUS COMPACTS

M.N. Rahaman, L.C. De Jonghe and C.H. Hsueh

March 1985



TWO-WEEK LOAN COPY

This is a Library Circulating Copy which may be borrowed for two weeks.

University of California Berkeley, California 94720

Prepared for the U.S. Department of Energy under Contract DE-AC03-76SF00098

Center for Advanced Materials

DISCLAIMER

This document was prepared as an account of work sponsored by the United States Government. While this document is believed to contain correct information, neither the United States Government nor any agency thereof, nor the Regents of the University of California, nor any of their employees, makes any warranty, express or implied, or assumes any legal responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by its trade name, trademark, manufacturer, or otherwise, does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof, or the Regents of the University of California. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof or the Regents of the University of California.

CREEP DURING SINTERING OF POROUS COMPACTS

M. N. Rahaman, L. C. De Jonghe and C.H. Hsueh Center for Advanced Materials, Lawrence Berkeley Laboratory and

Department of Materials Science and Mineral Engineering University of California, Berkeley Berkeley, CA 94720

Abstract

Creep during sintering of powder compacts of cadmium oxide was studied in a loading dilatometer by applying a small, transient, uniaxial load to the compacts. After removal of the load, the axial shrinkage rate is lower but the radial shrinkage rate is actually higher than that of a compact sintered under no load. This reduction in the axial shrinkage rate is more pronounced for longer, transient loading times. The results provide further support for a mechanism of simultaneous creep and densification in which creep at constant volume occurs by diffusion controlled grain boundary sliding.

Supported by the Division of Materials Sciences, Office of Basic Energy Sciences, U.S. Department of Engergy, under Contract No. DE-ACO3-76F00098.

I. INTRODUCTION

It is now widely recognized that inhomogeneities within a powder compact can lead to non-uniform sintering rates which, in turn, produce transient (and sometimes residual) stresses.¹⁻⁵ These stresses may influence creep processes within the porous sintering compact. It is therefore important to quantify the interaction of creep and densification processes during sintering.

In previous work by Rahaman and De Jonghe^{6,7} the usefulness of a loading dilatometer to study simultaneous creep and densification was demonstrated. In the loading dilatometer, a small, measured, uniaxial load is applied to the sintering compact. This small load does not affect the volumetric densification rate but may cause extensive creep. A later paper,⁸ extending the study in both theoretical and experimental areas, showed how a number of important sintering parameters such as the sintering stress, the effective viscosity and the sintering mechanism, could be measured. Based on these results and some initial results involving transient loads, a simplified model was put forward to interpret the microstructural processes occuring during simultaneous creep and densification. In this model, creep at constant volume occurred by a mechanism of grain boundary diffusion controlled grain boundary sliding.

The purpose of this paper is to report the results of further exeriments performed to test the validity of this microstructural model, and to compare them with theoretical calculations based on this model.

II. EXPERIMENTAL PROCEDURE

Cadmium oxide powder* was uniaxially pressed at 20MPa into cylindrical compacts (6mm by 6mm) and then isostatically pressed at ~60MPa to give compacts with a green density of 0.56±0.01 of theoretical. All sintering experiments, with or without load, were performed at 1123K in flowing air (~50cm³/min) in a loading dilatometer described in detail elsewhere.⁵

In one set of experiments, different compacts were sintered under no load and under a constant load of 5N for two hours. In another set of experiments involving transient loads, compacts were sintered under 5N load for 11 min and 55 min, respectively, and then the load was removed. Continued sintering under no load then occurred, and the experiments were terminated after a total time of two hours. Finally, the (differential) radial shrinkage was measured after the removal of the transient load as follows: Two compacts, one subjected to 5N load in the axial direction and the other subjected to no load, were sintered for 25 min and the differential axial shrinkage was measured. The compacts were removed and flat and parallel surfaces were machined on their cylindrical surfaces, to allow for shrinkage measurements in the radial direction. These compacts were sintered further, both subjected to no load, and the differential radial shrinkage was measured.

^{*}Reagent grade, J.T. Baker Chemical Company, Phillipsburg, N.J.

The density of the compacts at any time was calculated from the green density and the measured shrinkage. The final density was also measured using Archimedes' principle.

III. RESULTS AND DISCUSSION

Fig. 1 shows the axial shrinkage, $\Delta L/L_O$, versus time, t, for compacts sintered under no load (Curve A), at a constant load of 5N (B) and under transient loads of 5N (C and D). For Curves C and D the load was quickly removed after times, t^* , of 11 and 55 min, respectively. (L_O = intial sample length, and ΔL = $L-L_O$, where L = instantaneous sample length). A load of 5N represents a stress of 0.22 MPa and t = 0 represents the beginning of sintering. The sintering temperature was reached after t = 8 min and each curve is reproducible to within $\pm 2\%$.

As found earlier, 7 the small applied stresses have almost no effect on the volumetric densification rate. The density of all the compacts at any sintering time agreed to within ±1%. The final density (0.86 of theoretical) calculated from the dimensional changes was also in good agreement with that measured using Archimedes' principle.

The axial shrinkage rate, $(\frac{1}{L_0}\frac{dL}{dt})$, is plotted versus t in Fig. 2. The letter by each curve represents the same conditions as stated for Fig. 1. It is seen that after the removal of the transient load, the axial shrinkage rate (Curves C and D) is lower than that of the compact sintered throughout under no load (Curve A). Moreover this reduction depends on the time, t, * at which the transient load was removed, being more pronounced at longer t*.

Fig. 3 shows the differential shrinkage in the radial direction between two compacts, one previously sintered for 25 min under 5N load in the axial direction and the other sintered under no load for the same time. These results show that the previously loaded compact shrinks <u>faster</u> in the radial direction after the load is removed.

It is important now to compare the nature of these results with those expected from the simplified microstructural model put forward earlier⁸ to account for simultaneous creep and densification in CdO. Fig. 4 shows the model in which the grain boundaries in a porous compact are subjected to an external, uniaxial stress. Depending on their orientation, boundaries may be under compression, under tension or sliding. For CdO subjected to low stresses, the rate controlling creep and densification mechanisms are the same, namely, grain boundary diffusion. 8 Creep at constant volume is controlled by diffusion of matter from boundaries under compression into neighboring pores (A), and from neighboring pores (B) into boundaries under tension. According to this model, a CdO compact will behave as it were denser in the direction of the applied stress. Thus after removal of the stress, the axial shrinkage rate should be lower than that of a compact sintered throughout under no stress. Moreover, since the effect of load on the microstructure is cumulative, this decrease in the axial shrinkage rate should be more pronounced after longer transient loading times, t. These predictions are very well substantiated by the results of Fig. 2.

On the basis of the microstructural model, a quantitative estimate may be made of the apparent densification rate in the axial direction (as measured by the dilatometer) after application or removal of the transient loads.

Consider a compact undergoing sintering under no load. Then for CdO the densification rate at time $t = t^*$, say, may be written as ⁸

$$\dot{\rho}_0 = K \Sigma_a \phi_a^{3/2}$$
 Eqn. (1)

where K is a kinetic constant that includes the grain size, Σ_a is the sintering stress and ϕ_a is the stress intensification factor.

If a stress, σ , is applied to a freely sintering compact at t = t^* , then at $t^* + \delta t$, where δt is small, the apparent densification rate as measured by the dilatometer is given by

$$\dot{\rho}_{\sigma} = K(\Sigma_{a} + \sigma \phi_{a}) \phi_{a}^{3/2} \qquad Eqn. (2)$$

Then

$$\dot{\rho}_{\sigma}/\dot{\rho}_{o} = 1 + \sigma\phi_{a}/\Sigma_{a}$$
 Eqn. (3)

Both ϕ_a and Σ_a can be obtained from experiment.⁸

Consider now a compact undergoing sintering under a constant stress, σ . Then the apparent densification rate at t = t^* is given by

$$\dot{\rho}_{\sigma}' = K \left(\Sigma_{b} + \sigma \phi_{b} \right) \phi_{b}^{3/2}$$
 Eqn. (4)

where Σ_b and ϕ_b are the appropriate sintering stress and stress intensification factor, respectively. Σ_b and ϕ_b are somewhat different from Σ_a and ϕ_a in Eqn. 1 since the apparent density as measured by the dilatometer is higher in sintering under load, and Σ_b and ϕ_b are functions of density.

If the stress is now quickly removed from a compact at $t = t^*$, then at $t^* + \delta t$, the apparent densification rate of this unloaded compact is given by

$$\dot{\rho}_{o} = K \Sigma_{b} \phi_{b}^{3/2} \qquad Eqn. (5)$$

Then

$$\dot{\rho}_{\sigma}/\dot{\rho}_{0} = 1 + \sigma\phi_{b}/\Sigma_{b}$$
 Eqn. (6)

And

$$\dot{\rho}_{\sigma}^{\prime}/\dot{\rho}_{o} = (\Sigma_{b}/\Sigma_{a})(\phi_{b}/\phi_{a})^{3/2}$$
 Eqn. (7)

 $\Sigma_{\rm b}$ and $\phi_{\rm b}$ are derived, first by using the axial shrinkage (Fig. 1) to calculated the apparent density, as measured by the dilatometer, and then found at this density from other experiments that determined Σ and ϕ as a function of density. The values calculated for $\dot{\rho}_{\rm o}/\dot{\rho}_{\rm o}$, $\dot{\rho}_{\rm o}/\dot{\rho}_{\rm o}$ using equations 3, 6 and 7 are in good agreement with the experimental results (extrapolated to t = t*) to within $\pm 2\%$, as is evident from Table I.

Table I. Comparison of the experimental results with theoretical results calculated on the basis of the microstructural model.

RATIO	EXPT.	THEORY	COMMENT
ρ ₀ /ρ ₀	0.89	0.88	Curve C, t* = 11 min.
ρ <mark>'</mark> /ρ _ο	0.74	0.74	Curve D, t* = 55 min.
ۀ',/ۀ ₀	1.60	1.58	Curve A, t* = 20 min. (Fig. 10, Reference 8)

IV CONCLUSIONS

The small applied stresses cause constant volume creep without affecting the volumetric densification rate. Another consequence of the microstructural model is that following the removal of the stress, the radial shrinkage should be less than that of a compact sintered throughout under no load, i.e. the compact sintered under the uniaxial stress appears denser in the axial direction but less dense in the radial direction compared to a compact sintered under no load. The results of Fig. 3 clearly confirm this prediction.

It is clear that the mechanism of constant volume creep by diffusion controlled grain boundary sliding, put forward earlier to explain the microstructural changes occurring during simultaneous creep and densification of CdO, has been further supported by the present results. It should be stressed that the model has been verified so far only for the case in which the creep process is grain boundary diffusion controlled grain boundary sliding. The study of simultaneous creep and densification in other ceramic systems is currently in progress.

REFERENCES :

- 1. A. G. Evans, "Considerations of Inhomogeneity Effects in Sintering," J. Amer. Ceram. Soc., 65 (10) 497-501 (1982).
- 2. R. Raj and R. K. Bordia, "Sintering Behaviour of Bimodal Compats," Acta Metall., 32 (7) 1003-19 (1984).
- 3. C. H. Hsueh, A. G. Evans, R. M. Cannon and R. J. Brook, "Viscoelastic Stresses and Sintering Damage in Heterogeneous Powder Compacts," Manuscript submitted to Acta Metall.
- 4. C. H. Hsueh and A. G. Evans, "The Influence of Heterogeneities on Sintering Behaviour," Manuscript submitted to J. Amer. Ceram. Soc.
- 5. L. C. De Jonghe, M. N. Rahaman and C. H. Hsueh, "Transient Stresses in Bimodal Compacts During Sintering," Manuscript submitted to Acta Metall.
- 6. L. C. De Jonghe and M. N. Rahaman, "Loading Dilatometer," Rev. Sci. Instrum., 55 (12) 2007-10 (1984).
- 7. M. N. Rahaman and L. C. De Jonghe, "Sintering of CdO Under Low Applied Stress," J. Amer. Ceram. Soc., 67 (10) C-205-7 (1984).
- 8. M. N. Rahaman, L. C. De Jonghe and R. J. Brook, "Effect of Shear Stress on Sintering," Manuscript submitted to J. Amer. Ceram. Soc.

LIST OF FIGURES

- 1. Axial shrinkage versus time for compacts sintered under no load (Curve A), a constant load of 5N (B) and transient loads of 5N (C and D).
- 2. Axial shrinkage rate versus time.
- 3. Differential radial shrinkage, after removal of the axial load, versus time.
- 4. Microstructural model showing pores and grain boundaries subjected to an externally applied load. (Reference 8).

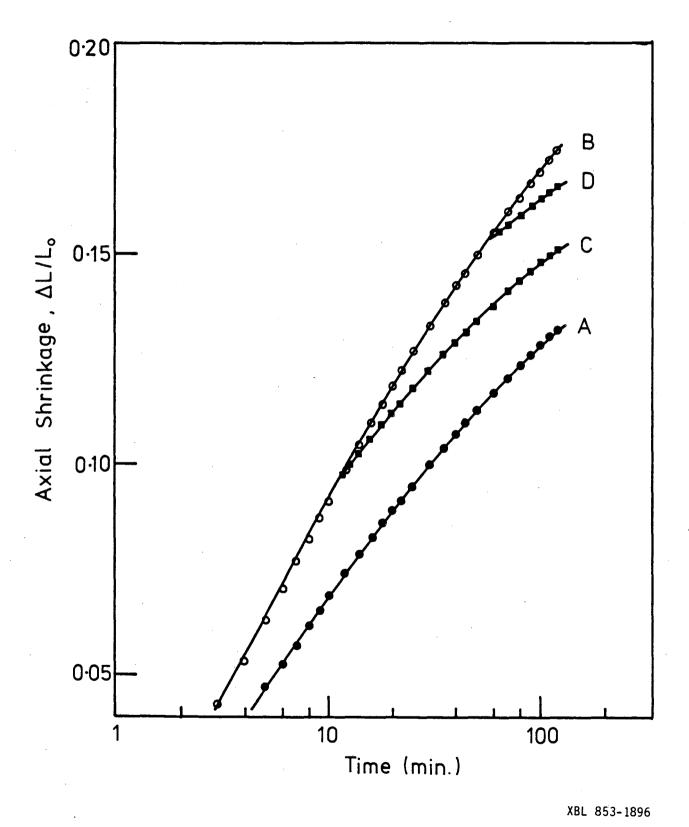


Figure 1

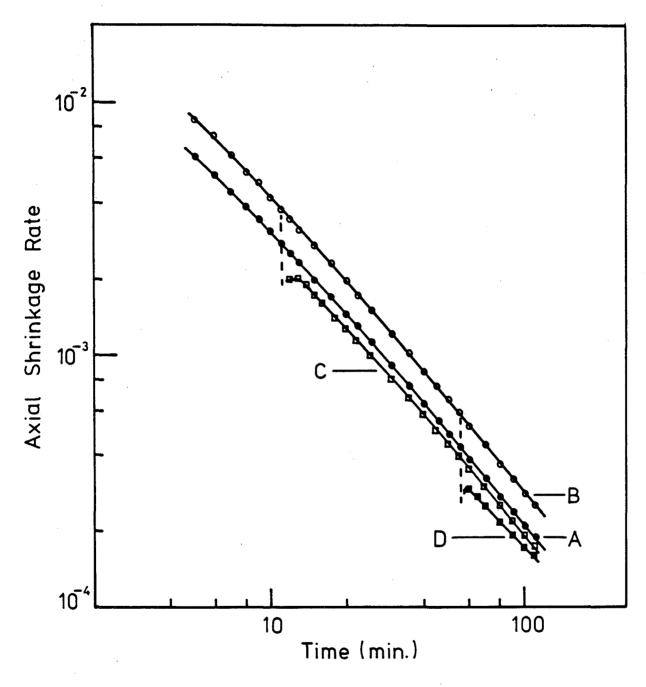


Figure 2

XBL 853-1897

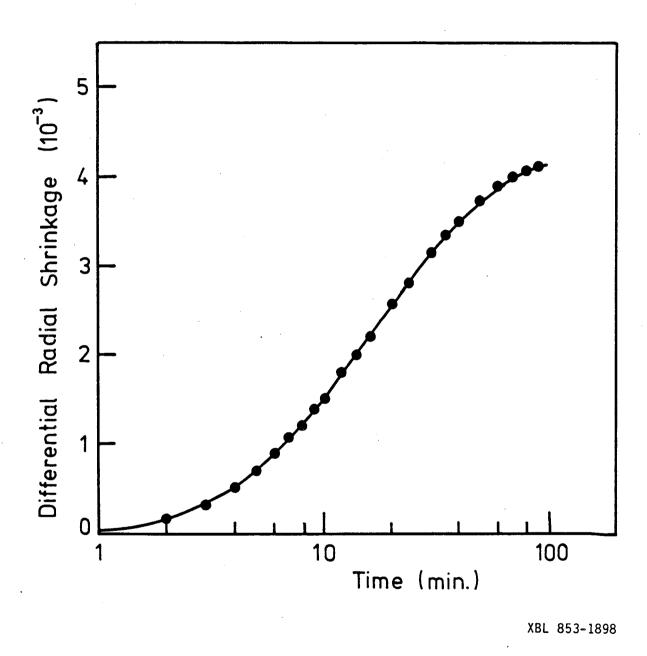


Figure 3

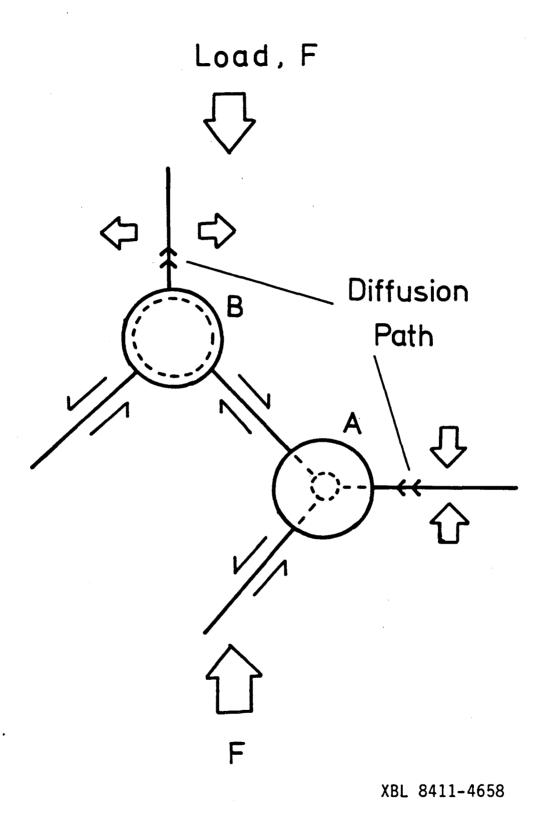


Figure 4

This report was done with support from the Department of Energy. Any conclusions or opinions expressed in this report represent solely those of the author(s) and not necessarily those of The Regents of the University of California, the Lawrence Berkeley Laboratory or the Department of Energy.

Reference to a company or product name does not imply approval or recommendation of the product by the University of California or the U.S. Department of Energy to the exclusion of others that may be suitable.

4- * ->

TECHNICAL INFORMATION DEPARTMENT LAWRENCE BERKELEY LABORATORY UNIVERSITY OF CALIFORNIA BERKELEY, CALIFORNIA 94720