

229
8/28/78

LD. 404
ORNL/TM-6443

MASTER

MASTER

**Irradiation Creep by Climb-Enabled
Glide of Dislocations Resulting from
Preferred Absorption of Point Defects**

L. K. Mansur

OAK RIDGE NATIONAL LABORATORY
OPERATED BY UNION CARBIDE CORPORATION · FOR THE DEPARTMENT OF ENERGY

DISTRIBUTION OF THIS DOCUMENT IS UNLIMITED

DISCLAIMER

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency Thereof, nor any of their employees, makes any warranty, express or implied, or assumes any legal liability or 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 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. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof.

DISCLAIMER

Portions of this document may be illegible in electronic image products. Images are produced from the best available original document.

Printed in the United States of America. Available from
National Technical Information Service
U.S. Department of Commerce
5285 Port Royal Road, Springfield, Virginia 22161
Price: Printed Copy ~~\$4.50~~; Microfiche \$3.00

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor any of their employees, contractors, subcontractors, or their employees, makes any warranty, express or implied, nor assumes any legal liability or responsibility for any third party's use or the results of such use of any information, apparatus, product or process disclosed in this report, nor represents that its use by such third party would not infringe privately owned rights.

ORNL/TM-6443
Dist. Category UC-25

Contract No. W-7405-eng-26

Metals and Ceramics Division

IRRADIATION CREEP BY CLIMB-ENABLED GLIDE OF DISLOCATIONS
RESULTING FROM PREFERRED ABSORPTION OF POINT DEFECTS

L. K. Mansur

Date Published - August 1978

NOTICE
This report was prepared as an account of work sponsored by the United States Government. Neither the United States nor the United States Department of Energy, nor any of their employees, nor any of their contractors, subcontractors, or their employees, makes any warranty, express or implied, or assumes any legal liability or 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.

NOTICE This document contains information of a preliminary nature. It is subject to revision or correction and therefore does not represent a final report.

Oak Ridge National Laboratory
Oak Ridge, Tennessee 37830
operated by
Union Carbide Corporation
for the
Department of Energy

DISTRIBUTION OF THIS DOCUMENT IS UNLIMITED

THIS PAGE
WAS INTENTIONALLY
LEFT BLANK

CONTENTS

ABSTRACT	1
INTRODUCTION	1
THEORY	3
Climb-Glide Creep in Terms of Dislocation Climb Velocity .	3
The Climb Velocity of Dislocations under Irradiation. . . .	5
The Climb-Glide Creep Rate	6
The Creep Rate by Dislocation Climb	8
Relative Characteristics of Climb-Glide and Climb Components	9
Magnitudes	11
DISCUSSION	15

IRRADIATION CREEP BY CLIMB-ENABLED GLIDE OF DISLOCATIONS
RESULTING FROM PREFERRED ABSORPTION OF POINT DEFECTS

L. K. Mansur

ABSTRACT

A mechanism of irradiation creep is proposed that arises from the climb-enabled glide of dislocations due to stress-induced preferred absorption of radiation-produced point defects. This creep component we term preferred absorption glide, PAG. PAG-creep operates in addition to the previously studied components of creep from climb by stress-induced preferred absorption, PA-creep, and the climb-enabled glide due to excess absorption of interstitials on dislocations during swelling, I-creep. A formulation of the various climb and climb-enabled glide processes is presented which includes earlier results. PAG-creep is comparable in magnitude to PA-creep in the parameter range of applications. While the PA-creep rate and the I-creep rate are linear in stress, the PAG creep rate is quadratic stress and thus dominates at high stress.

INTRODUCTION

When a specimen is subjected to a stress, the resulting creep may take place by two physically distinct processes. These are the atomic transfer of material from planes more nearly parallel to those more nearly perpendicular to the stress direction and the translation of planes inclined to the stress direction by the glide of dislocations. These processes may be coupled in general since the transfer of material may be accomplished by dislocation climb, causing dislocations to sample different microscopic regions of the specimen. Within some of these regions, dislocation glide may be possible.

In the absence of irradiation, dislocation climb may be accomplished by vacancy diffusion which results from the stress induced preferred emission (PE) of vacancies from dislocations whose Burgers vectors are

nearly parallel to the stress direction. This results in a form of Nabarro-Herring creep.¹ During irradiation, a net excess of interstitials may precipitate on dislocations of all orientations resulting in dislocation climb, while the excess vacancies accumulate at cavities. This process gives rise to macroscopic swelling which has been the subject of intensive study in recent years.²⁻⁵ During irradiation, there may be, however, an additional component of dislocation climb even in the absence of swelling.⁶⁻⁸ This climb results from the stress-induced preferred absorption of radiation-produced interstitials on dislocations with Burgers vectors nearly parallel to the stress direction⁹ with the corresponding excess vacancies absorbed on dislocations with Burgers vectors nearly perpendicular to the stress direction. This results in (SI)PA-creep.

It has been shown previously that the dislocation climb resulting from swelling will give rise to a climb-enabled glide component of irradiation creep termed I-creep.¹⁰ The purpose of the present paper is to show that the dislocation climb caused by the PA process will also result in a climb-enabled glide component of creep. This previously unremarked glide creep resulting from preferred absorption, PAG-creep, is shown to be of comparable magnitude to the climb component itself, PA-creep. It is shown that these two components differ in their dependences on the stress and microstructural parameters.

¹F.R.N. Nabarro, *Phil. Mag.* 16: 231 (1967).

²S. D. Harkness and Che-Yu Li, *Met. Trans.* 2: 1457 (1971).

³H. Wiedersich, *Rad. Eff.* 12: 111 (1972).

⁴A. D. Brailsford and R. Bullough, *J. Nucl. Mater.* 44: 121 (1972).

⁵L. K. Mansur, *Nucl. Technol.* (in press).

⁶P. T. Heald and M. V. Speight, *Phil. Mag.* 29: 1075 (1974).

⁷R. Bullough and J. R. Willis, *Phil. Mag.* 31: 855 (1975).

⁸W. G. Wolfer and M. Ashkin, *J. Appl. Phys.* 47: 791 (1976).

⁹In principle, there also may be a stress-induced preferred absorption of vacancies on dislocations with Burgers vector nearly perpendicular to the stress direction. However, with the vacancy parameters usually employed this effect is negligible.

¹⁰J. H. Gittus, *Phil. Mag.* 25: 345 (1972).

In the next section, the formulation of the various climb and glide processes of creep is developed. The relative magnitudes and some limiting cases are explored. The results of calculations in parameter ranges of interest are presented. In the last section, the implications of this work are discussed.

THEORY

We visualize the creep along the j axis to be comprised as follows

$$\dot{\epsilon}^j = \dot{\epsilon}_C^j + \dot{\epsilon}_{CG}^j + \dot{\epsilon}_G^j \quad (1)$$

where $\dot{\epsilon}_C^j$ denotes the sum of the climb components and $\dot{\epsilon}_{CG}^j$ the sum of the climb-glide components. $\dot{\epsilon}_G^j$ includes the possibility of glide components not enabled by climb and is not discussed in this paper. $\dot{\epsilon}_C^j$ contains contributions due to preferred absorption and preferred emission of point defects

$$\dot{\epsilon}_C^j = \dot{\epsilon}_{PA}^j + \dot{\epsilon}_{PE}^j \quad (2)$$

In addition, these PA and PE processes as well as the climb due to swelling produce creep by climb-enabled glide, denoted by $\dot{\epsilon}_{CG}^j$.

Climb-Glide Creep in Terms of Dislocation Climb Velocity

The creep rate resulting from climb-enabled glide may be obtained in terms of the dislocation climb velocity by the following argument.^{10,11} Under applied stress, pinned dislocations glide until they reach a bowed-out configuration where the line tension restoring force equals the applied stress. Since the dislocations are pinned the creep by this process is limited to approximately one elastic deflection.^{12,13}

¹¹P. T. Heald, Proceedings of the Conference on *Effects of Radiation on Breeder Reactor Structural Materials*, Scottsdale, Arizona, ed. by M. L. Bleiberg and J. W. Bennett, p. 781, 1977.

¹²N. F. Mott, *Phil. Mag.* 43: 1151 (1952).

¹³J. Friedel, *Phil. Mag.* 44: 444 (1953).

However, if climb is possible the dislocations climb past the initial pinning points while the initially bowed-out segments encounter new pinning points. The released segments now between the new pinning points bow out until their line tension balances the applied stress. This cycle produces another elastic deflection and so on. This is, of course, in addition to any creep due to the climb per se. This action produces a directional glide of dislocations and a macroscopic creep while the overall configuration of the dislocation network is maintained. If the magnitude of the average dislocation climb velocity is V and the pinning points are distance λ apart, then the creep rate is

$$\dot{\epsilon}_{CG} = c/(\lambda/V) \quad , \quad (3)$$

where ϵ denotes the elastic deflection σ/E , σ is the stress, and E is Young's modulus. Where the pinning is due to the dislocation network itself, i.e., where the presence of other objects such as precipitates can be ignored, then λ must be determined by the dislocation spacing. We take λ as one-half the dislocation spacing.¹¹ However, our conclusions are not affected by the proportionality constant. Thus

$$\lambda \cong (\pi L)^{-\frac{1}{2}} \quad (4)$$

where L denotes the dislocation density. We now obtain the creep rate in terms of the dislocation density and climb velocity as

$$\dot{\epsilon}_{CG} = \epsilon (\pi L)^{\frac{1}{2}} V \quad . \quad (5)$$

Here V denotes the magnitude of the average climb velocity of a dislocation. The possibility of climb-glide creep does not depend upon the sign of the velocity, i.e., upon whether the dislocation is climbing due to a net interstitial or a net vacancy absorption. In either case the dislocation may climb past pinning points. We make use of this idea later.

The Climb Velocity of Dislocations under Irradiation

The current of excess interstitial volume per unit volume to dislocations whose Burgers vectors are aligned in direction j is

$$I^j = \Omega (Z_i^{dj} D_i C_i - Z_v^{dj} D_v C_v + Z_v^{dj} D_v C_v^{dj}) L^j, \quad (6)$$

where Ω is the atomic volume, Z_i^{dj} and Z_v^{dj} are capture efficiencies of dislocations in orientation j for interstitials and vacancies, D_i and D_v are the interstitial and vacancy diffusion coefficients where $D_{i,v} = D_{i,v}^0 \exp(-E_{i,v}^m/kT)$, $D_{i,v}^0$ is a constant, $E_{i,v}^m$ is the interstitial or vacancy migration energy, k is Boltzmann's constant, and T is temperature. C_i and C_v are the bulk-averaged physical concentrations of interstitials and vacancies and C_v^{dj} is the vacancy concentration in equilibrium with a dislocation of orientation j . For a tensile stress σ aligned with the l -axis we have

$$C_v^{d1} = C_v^e \exp(\sigma\Omega/kT) \quad (7)$$

and

$$C_v^{d2} = C_v^{d3} = C_v^e \quad (8)$$

Here

$$C_v^e = \Omega^{-1} \exp(S_v^f/k) \exp(-E_v^f/kT) \quad (9)$$

is the bulk thermal equilibrium vacancy concentration, S_v^f and E_v^f are the entropy and energy of vacancy formation. In this analysis we neglect the thermal equilibrium interstitial concentration since interstitial formation energies in materials of interest are so high as to make the thermal population entirely negligible. Conservation of atoms requires the current I^j to be reflected in an accumulation recorded by the climb of dislocations

$$I^j = b L^j \nu^j, \quad (10)$$

where b is an atomic dimension and ν^j is the climb velocity of dislocations in orientation j . Using Eq. (6), Eq. (10) may be rewritten for purposes of illustration as

$$\begin{aligned} \nu^j = & \frac{\Omega}{b} \left[\left(Z_i^{dj} D_i C_i - Z_v^{dj} D_v C_v + Z_v^{dj} D_v C_v^{dj} \right) \right. \\ & - (3L^j)^{-1} \sum_{m=1}^3 \left(Z_i^{dm} D_i C_i - Z_v^{dm} D_v C_v + Z_v^{dm} D_v C_v^{dm} \right) L^m \left. \right] \\ & + \frac{\Omega}{3bL^j} \sum_{m=1}^3 \left(Z_i^{dm} D_i C_i - Z_v^{dm} D_v C_v + Z_v^{dm} D_v C_v^{dm} \right) L^m, \end{aligned} \quad (11)$$

by adding and subtracting the velocity component due to volumetric swelling. The last line in Eq. (11) is the dislocation climb velocity due to isotropic swelling and results in a climb-enabled glide component of creep, which has been identified previously as l -creep.¹⁰ The first two lines represent the dislocation climb velocity resulting from processes other than swelling, i.e., the volume conserving PA and PE processes. The climb components of velocity expressed in this term in square brackets give rise to climb-enabled glide components of creep which we label PAG- and PEG-creep. It is the climb-glide creep resulting from this term in square brackets that has not been identified previously and which is explored in this paper.

The Climb-Glide Creep Rate

According to Eq. (5) it is the *magnitude* of the climb velocity of the average dislocation which determines the climb-glide creep rate. For dislocations isotropically distributed among the three axis directions

$$L^1 = L^2 = L^3 = L/3; \quad (12)$$

this is

$$v = \frac{|v^1| + |v^2| + |v^3|}{3} \quad (13)$$

From Eq. (11) we obtain

$$v_{I+PA+PE} = \frac{\Omega}{3b} \left\{ \left| z_i^{d1} D_i C_i - z_v^{d1} D_v C_v + z_v^{d1} D_v C_v^{d1} \right| + 2 \left| z_i^{d2} D_i C_i - z_v^{d2} D_v C_v + z_v^{d2} D_v C_v^{d2} \right| \right\} \quad (14)$$

which is valid in general whether or not swelling is occurring.¹⁴ When PE and PA processes are occurring in the absence of swelling then conservation of atoms requires

$$\left(z_i^{d1} D_i C_i - z_v^{d1} D_v C_v + z_v^{d1} D_v C_v^{d1} \right) = -2 \left(z_i^{d2} D_i C_i - z_v^{d2} D_v C_v + z_v^{d2} D_v C_v^{d2} \right) \quad (15)$$

¹⁴ If we were to ignore preferred absorption, i.e., set the capture efficiencies of dislocations in different orientations equal, $z_i^{d1} = z_i^{d2} = z_i^d$; $z_v^{d1} = z_v^{d2} = z_v^d$, Eq. (14) would reduce to

$$v_{I+PE} = \frac{\Omega}{b} \left[z_i^d D_i C_i - z_v^d D_v (C_v - \bar{C}_v^d) \right]$$

where

\bar{C}_v^d is given by Eq. (17).

When inserted into Eq. (5) this gives the climb-glide creep rate due to swelling (with preferred vacancy emission when \bar{C}_v^d is non-negligible). This result is identical to the climb-glide creep rate due to swelling, I-creep, proposed earlier.^{10,11} This approximation excludes the possibility of preferred absorption enabled glide which is the focus in the present paper.

In this case Eq. (14) becomes

$$v_{PA+PE} = \frac{2}{3} \frac{\Omega}{b} \left| z_i^{d1} D_i C_i - z_v^{d1} D_v (C_v - C_v^{d1}) \right| \quad (16)$$

Finally, in the absence of irradiation, preferred vacancy emission is possible but preferred interstitial absorption and swelling are absent. In this case C_i may be replaced by zero and C_v becomes

$$\bar{C}_v^d = C_v^e \frac{z_v^{d1} \exp\left(\frac{\sigma\Omega}{kT}\right) + 2z_v^{d2}}{z_v^{d1} + 2z_v^{d2}} \quad (17)$$

In this case Eq. (16) reduces to

$$v_{PE} = \frac{4}{9} \frac{\Omega}{b} z_v^{d1} D_v C_v^e \left| \exp\left(\frac{\sigma\Omega}{kT}\right) - 1 \right| \quad (18)$$

since $z_v^{d1} \cong z_v^{d2}$.¹⁵ Equation (5) gives the creep rate due to climb enabled glide in terms of the dislocation climb velocity [Eqs. (14), (16), or (18)]. Equation (14) is the correct expression for that velocity in general. When there is no swelling occurring the climb velocity reduces to Eq. (16). In the absence of radiation the dislocation climb is due only to preferred vacancy emission and the climb velocity is given by Eq. (18).

The Creep Rate by Dislocation Climb

In this section we recall the formulation of the dislocation climb mechanism of irradiation creep, PA-creep, utilized by Heald.¹⁶ The deformation due to precipitation of a net excess of interstitials at dislocations and corresponding vacancies at cavities is traditionally accounted as swelling rather than volumetric creep. Thus, to find the creep rate in any direction we subtract from the total extension rate in that direction the component of extension rate due to swelling.

¹⁵L. K. Mansur and M. H. Yoo, *J. Nucl. Mater.* (in press).

¹⁶P. T. Heald, Proceedings of the Conference on *Effects of Radiation on Breeder Reactor Structural Materials*, Scottsdale, Arizona, ed. by M. L. Bleiberg and J. W. Bennett, p. 781, 1977.

$$\begin{aligned} \dot{\epsilon}_C^j = & \Omega \left(z_i^{dj} D_i C_i - z_v^{dj} D_v C_v + z_v^{dj} D_v C_v^{dj} \right) L^j \\ & - \frac{\Omega}{3} \sum_{m=1}^3 \left(z_i^{dm} D_i C_i - z_v^{dm} D_v C_v + z_v^{dm} D_v C_v^{dm} \right) L^m . \end{aligned} \quad (19)$$

If we now utilize the expressions (7), (8), and (12) for equilibrium vacancy concentrations and dislocation density, Eq. (19) becomes for the stress direction $j = 1$,

$$\dot{\epsilon}_C = \dot{\epsilon}_{PA} + \dot{\epsilon}_{PE} = \frac{2}{9} \Omega L \left\{ \left[\Delta z_{i,v}^d D_i C_i - \Delta z_{v,v}^d D_v C_v \right] + D_v C_v^e \left[z_v^{d1} \exp \left(\frac{\sigma \Omega}{kT} \right) - z_v^{d2} \right] \right\}, \quad (20)$$

where $\Delta z_{i,v}^d = z_{i,v}^{d1} - z_{i,v}^{d2}$. The first square bracket gives rise to the dislocation climb creep rate resulting from preferred absorption, PA-creep, while the second square bracket gives rise to the dislocation climb creep rate resulting from preferred emission of vacancies, PE-creep.

Relative Characteristics of Climb-Glide and Climb Components

In this section we explore the characteristics of the newly proposed climb-glide creep mechanism in relation to the dislocation climb mechanisms. We consider the case where there is no swelling. In this case Eq. (5) together with Eq. (16) gives the climb-glide creep rate due to PA and PE processes. In this case also, Eq. (15) applies and may be rewritten as

$$\bar{z}_{i,v}^d D_i C_i = \bar{z}_{v,v}^d \left(C_v - \bar{C}_v^d \right) \quad (21)$$

where

$$\bar{z}_{i,v}^d \equiv \frac{z_{i,v}^{d1} + 2z_{i,v}^{d2}}{3} = z_{i,v}^{d1} - \frac{2}{3} \Delta z_{i,v}^d \quad (22)$$

In view of expressions (17) and (22), Eq. (21) may be rewritten

$$z_v^{dl} = \frac{(z_i^{dl} - \frac{2}{3} \Delta z_i^d)}{D_v(C_v - \bar{C}_v^d)} D_i C_i + \frac{2}{3} \Delta z_v^d . \quad (23)$$

We substitute Eq. (23) into Eq. (16) to obtain

$$v_{PA+PE} = \frac{2}{3} \frac{\Omega}{b} \left| z_i^{dl} D_i C_i - \left[\frac{(z_i^{dl} - \frac{2}{3} \Delta z_i^d) D_i C_i}{D_v(C_v - \bar{C}_v^d)} + \frac{2}{3} \Delta z_v^d \right] D_v (C_v - C_v^{dl}) \right| . \quad (24)$$

We have already mentioned, however, that when we use the usual vacancy parameters that

$$\Delta z_v^d \cong 0 , \quad (25)$$

meaning that there is no stress induced preferred absorption of vacancies at dislocations.¹⁵ If for the moment we also ignore thermal emission of vacancies (we treat the thermal creep rate separately later), $C_v \gg C_v^{dl}$, $C_v \gg \bar{C}_v^d$, we obtain from Eqs. (24) and (5) the PAG-creep rate due to stress induced preferred absorption of interstitials

$$\dot{\epsilon}_{PAG} = \frac{4}{9} \frac{\epsilon}{b} \left(\pi L \right)^{\frac{1}{2}} \Omega D_i C_i \Delta z_i . \quad (26)$$

The first term in square brackets in Eq. (20) gives the PA (climb) creep rate due to stress induced preferred absorption of interstitials

$$\dot{\epsilon}_{PA} = \frac{2}{9} \Omega L D_i C_i \Delta z_i . \quad (27)$$

Thus the ratio of the climb glide creep rate to the climb creep rate caused by stress induced preferred absorption of interstitials is

$$\frac{\dot{\epsilon}_{PAG}}{\dot{\epsilon}_{PA}} = \frac{2\epsilon}{b} \left(\frac{\pi}{L} \right)^{1/2} . \quad (28)$$

In typical cases ϵ is in the range 10^{-4} to 10^{-3} , L is in the range 1×10^{10} to $5 \times 10^{11} \text{ cm}^{-2}$, and b is $\sim 2 \times 10^{-8} \text{ cm}$. Thus the ratio of Eq. (28) is in the range 0.02 to 2. Therefore, we have shown that the creep rate produced by PAG is comparable to that produced by PA which has been studied previously.

For completeness we also note that the creep rate produced by PEG has the same ratio to that produced by PE as given in Eq. (28). This is shown by the same procedure as above: Obtain the PEG creep rate from Eqs. (18) and (5) and obtain the PE creep rate as the second term in Eq. (20). Taking the ratio gives

$$\frac{\dot{\epsilon}_{\text{PEG}}}{\dot{\epsilon}_{\text{PE}}} = \frac{2\epsilon}{b} \left(\frac{\pi}{L}\right)^{1/2} \quad (29)$$

Magnitudes

Equation (28) shows that the PAG- and PA-creep rates have different dependences on the stress (recalling that $\epsilon = \sigma/E$) and dislocation density. From Eq. (20) the PA-creep rate may be expressed as

$$\dot{\epsilon}_{\text{PA}} = \frac{2}{9} \Omega D_i C_i \epsilon L \Delta z_i^d \quad (30)$$

where $\Delta z_i^d = \Delta Z_i^d / \epsilon$ and does not depend on stress. Expressions for Δz_i^d have been given previously.^{16,17} The expression given by Heald is

$$\Delta z_i^d = \frac{3(1-\nu) \left\{ 2\pi / k n \left[\frac{2R_d}{3\pi(1-\nu)} \frac{(1+\nu) \mu b \Delta V_i}{KT} \right] \right\}^2 \Omega a_i}{2\pi(1+\nu) \Delta V_i} \quad (31)$$

¹⁷W. G. Wolfer, L. K. Mansur, and J. Sprague, Proceedings of the Conference on *Effects of Radiation on Breeder Reactor Structural Materials*, Scottsdale, Arizona, ed. by M. L. Bleiberg and J. W. Bennett, p. 841, 1977.

The expression given by Wolfer et al. is

$$\Delta z_i^d = - \frac{\Omega L \left(b/r_d \right)^2}{540 \pi \left[\ln(R_d/r_d) \right]^2} \frac{(1 + \nu)}{[(1 - \nu)kT]^2} (5 - 4\nu) E \alpha_i^\mu \Delta V_i \quad (32)$$

where ν is Poisson's ratio, $R_d = \left(\pi L \right)^{\frac{1}{2}}$, r_d is the dislocation core radius, E is Young's modulus, μ is shear modulus, ΔV_i the interstitial relaxation volume, and α_i^μ is the shear polarizability of the interstitial. Here $a_i = 15(1 + \nu) \Delta \mu_i / [15(1 - \nu)\mu + 2(4 - 5\nu) \Delta \mu_i]$, where $\Delta \mu_i$ is the difference in shear modulus of the matrix and the effective modulus of the interstitial. Equations (31) and (32) give similar numerical values for Δz_i^d in the stress and temperature range of interest.¹⁸

The corresponding expression for $\dot{\epsilon}_{\text{PAG}}$ is obtained from Eq. (26)

$$\dot{\epsilon}_{\text{PAG}} = \frac{4\pi^{1/2}}{9b} \Omega D_i C_i \epsilon^2 L^{1/2} \Delta z_i^d \quad (33)$$

We see from Eqs. (30) and (33) that the creep rate due to PAG is proportional to the square of the applied stress while that due to PA is linear in the stress. At high stresses, PAG-creep thus dominates. C_i in these equations also depends upon the sink strength in the specimen, one component of which is the dislocation density. C_i can be determined from chemical rate theory.

$$C_i = \frac{\left[K_i K_v + R(G_v - G_i) \right]}{2R K_i} \left\{ \left[1 + \frac{4R G_i K_v K_i}{\left[K_i K_v + R(G_v - G_i) \right]^2} \right]^{1/2} - 1 \right\} \quad (34)$$

where G_v and G_i are generation rates for vacancies and interstitials, K_v and K_i are loss rates per vacancy and interstitial to all distributed sinks, and R is the coefficient of recombination. These parameters are defined in detail by Mansur.⁵

¹⁸L. K. Mansur and W. G. Wolfer, to be published.

Figure 1 shows the creep rates due to PA and PAG. Table 1 gives the parameters used in this calculation. For these parameter values PAG becomes dominant at a stress of one to several times 10^9 dynes/cm² which is within the range of engineering application. Equations (30) and (33) also show that the PA and PAG creep rates have different dependences on the dislocation density. The predicted transition from linear (PA dominated) to parabolic (PAG dominated) stress dependence takes place at lower stress for lower dislocation density.

ORNL-DWG 78-9402

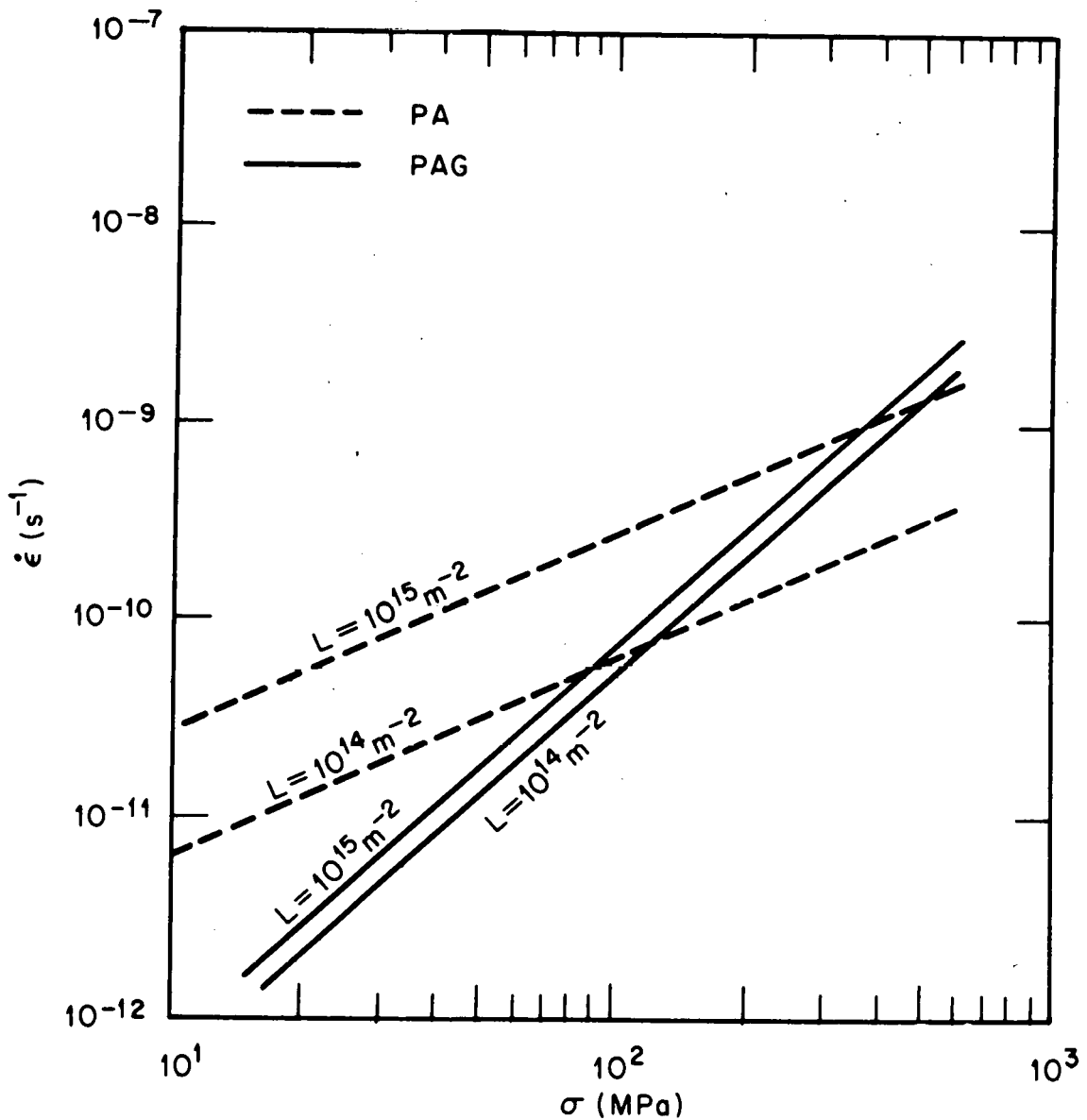


Fig. 1. Creep Rates from PA- and PAG-Mechanisms as a Function of Stress.

Table 1. Parameter Values Used in Obtaining Results Shown in Figure 1

Parameter	Value
Shear modulus, dyne/cm ²	7.75×10^{11}
Poisson's ratio	0.312
$\Delta\mu_v$, dyne/cm ²	0
$\Delta\mu_i$, dyne/cm ²	-7.75×10^{11}
Burgers vector, cm	2.1×10^{-8}
ΔV_v , cm ³	-5.06×10^{-24}
ΔV_i , cm ³	1.4×10^{-23}
V_v , cm ³	5.06×10^{-24}
V_i , cm ³	1.4×10^{-23}
T, °C	500
G, dpa/s	10^{-6}
D_v , cm ² /s	$0.014 \exp(-1.38 \text{ eV}/kT)$
D_i , cm ² /s	$0.008 \exp(-0.15 \text{ eV}/kT)$
S_v^f , k	1.5
E_v^j , eV	1.4

DISCUSSION

In this paper, three mechanisms of irradiation creep have been treated in an integrated fashion. Two mechanisms of thermal creep also are incorporated. This has been done in order to draw the correct perspective for the newly proposed mechanism of PAG-creep. I-creep arises from the climb-enabled glide of dislocations due to the net interstitial flux to dislocations associated with swelling during irradiation. PA-creep (or SIPA-creep) arises from the dislocation climb during irradiation due to preferred absorption of interstitials on dislocations whose Burgers vectors are aligned with the stress axis. The newly proposed PAG-creep arises from the climb-enabled glide of dislocations during irradiation due to the preferred absorption of interstitials on dislocations whose Burgers vectors are aligned with the stress axes. PE-creep (a form of Herring-Nabarro creep) arises from dislocation climb due to the stress-induced preferred thermal emission of vacancies from aligned dislocations. There is also a corresponding PEG-creep which arises from the climb-enabled glide of dislocations due to the stress-induced preferred thermal emission of vacancies from aligned dislocations.

PAG-creep has been formulated, its characteristics examined and compared to PA-creep. Both mechanisms operate during irradiation whether or not swelling is occurring concurrently. While the PA-creep rate (and the I-creep rate) is linear in stress, the PAG-creep rate is quadratic in stress. Thus at high stresses PAG-creep will dominate and the creep rate should approach a quadratic stress dependence. For typical parameter values it is predicted that the transition from linear to quadratic stress dependence of the creep rate will begin below a few times 10^9 dynes/cm². While it is true that the dislocation density also generally increases with stress in unirradiated materials, the dislocation density in irradiated materials is usually quite high and consequently insensitive to stress. Thus it is predicted that the second power stress dependence will be observed. The second power of the stress arises physically because the PA climb velocity is proportional to stress and the creep produced by bowing-out of dislocations

is also proportional to stress. PAG-creep combines these two processes multiplicatively and thus leads to a second power stress dependence.

The PA- and PAG-creep processes also exhibit different dependences on the total sink strength. Example calculations performed for cases where dislocations comprise most of the sink strength show that the PAG-creep rate is reduced less than the PA-creep rate for a reduction of dislocation density from 10^{11} to 10^{10} cm^{-2} . Thus the predicted transition from linear to quadratic stress dependence occurs at a lower stress for a lower dislocation density.

The PAG creep mechanism offers a possible explanation for the greater than linear stress dependence of creep rate which has recently been noted (Scottsdale Conference, 1977). Controlled experiments to better establish the dependence of this behavior on experimental conditions and properties of the material would enable a more definitive comparison.

INTERNAL DISTRIBUTION

- | | | | |
|--------|-------------------------------|--------|-----------------------------|
| 1-2. | Central Research Library | 40-49. | L. K. Mansur |
| 3. | Document Reference Section | 50. | P. J. Maziasz |
| 4-8. | Laboratory Records Department | 51. | C. J. McHargue |
| 9. | Laboratory Records, ORNL-RC | 52. | J. Narayan |
| 10. | ORNL Patent Office | 53. | T. S. Noggle |
| 11. | J. Bentley | 54. | O. S. Oen |
| 12. | E. E. Bloom | 55. | S. M. Ohr |
| 13. | D. N. Braski | 56. | N. H. Packan |
| 14. | W. H. Butler | 57. | T. C. Reiley |
| 15. | R. W. Carpenter | 58. | M. T. Robinson |
| 16. | J. C. Cathcart | 59. | A. F. Rowcliffe |
| 17. | R. E. Clausing | 60. | P. S. Sklad |
| 18. | W. A. Coghlan | 61. | J. O. Stiegler |
| 19. | J. E. Cunningham | 62. | R. W. Swindeman |
| 20. | W. P. Eatherly | 63. | P. F. Tortorelli |
| 21. | L. C. Emerson | 64. | D. B. Trauger |
| 22. | K. Farrell | 65. | T. Weerasooriya |
| 23. | J. S. Faulkner | 66. | J. R. Weir, Jr. |
| 24. | G. Gessel | 67. | C. L. White |
| 25. | M. L. Grossbeck | 68. | F. W. Wiffen |
| 26. | R. W. Hendricks | 69. | M. K. Wilkinson |
| 27-29. | M. R. Hill | 70. | M. H. Yoo |
| 30-32. | J. A. Horak | 71. | F. W. Young, Jr. |
| 33. | E. A. Kenik | 72. | A. Zucker |
| 34. | C. C. Koch | 73. | R. W. Balluffi (consultant) |
| 35. | E. Lee | 74. | P. M. Brister (consultant) |
| 36. | J. M. Leitnaker | 75. | W. R. Hibbard (consultant) |
| 37. | M. B. Lewis | 76. | R. J. Jaffe (consultant) |
| 38. | C. T. Liu | 77. | John Moteff (consultant) |
| 39. | K. C. Liu | 78. | N. E. Promisel (consultant) |

EXTERNAL DISTRIBUTION

79. A. Argon, Department of Mechanical Engineering, Massachusetts Institute of Technology, Cambridge, MA 02139
80. R. J. Arsenault, Engineering Materials Group, University of Maryland, College Park, MD 20742
81. M. Baron, Westinghouse Advanced Reactor Division, P. O. Box 158, Madison, PA 15663
82. J. R. Beeler, Jr., Department of Nuclear Engineering, North Carolina State University, Raleigh, NC 27607
83. A. L. Bement, Department of Metallurgical and Materials Science, Massachusetts Institute of Technology, Cambridge, MA 02139
84. A. Boltax, Westinghouse Advanced Reactor Division, P. O. Box 158, Madison, PA 15663

85. A. D. Brailsford, Ford Scientific Laboratory, P. O. Box 2053, Dearborn, MI 48120
86. J. L. Brimhall, Battelle Pacific Northwest Laboratories, Richland, WA 99352
87. R. Bullough, Theoretical Physics Division, Bldg. B.9, Atomic Energy Research Establishment, Harwell, Berkshire, England
88. L. T. Chadderton, Physics Lab II, H. G. Orsted Institute, University of Copenhagen, Universitetsparken 5, DK-2100, Copenhagen Φ , Denmark
89. J. W. Corbett, Physics Department, State University of New York at Albany, Albany, NY 12203
90. A. G. Crocker, Department of Physics, University of Surrey, Guildford GU2 5XH, United Kingdom
91. D. DeFontaine, Materials Department, UCLA School of Engineering, Los Angeles, CA 90024
92. J. Dienes, Department of Physics, Brookhaven National Laboratory, Upton, NY 11973
93. D. G. Doran, Hanford Engineering Development Laboratory, P. O. Box 1970, Richland, WA 99352
94. A.J.E. Foreman, Metallurgy Division, UKAEA Research Group, AERE Harwell, Didcot, Oxon, England
95. J. Gittus, UKAEA, Salwick, Preston, England
96. A. Goland, Brookhaven National Laboratory, Upton, NY 11973
97. D. R. Harries, Metallurgy Division, UKAEA Research Group, AERE Harwell, Didcot, Oxon, England
98. J. E. Harris, Berkeley Nuclear Laboratory, Berkeley, Gloucestershire, England GL13 9 PB
99. J. Hillairet, Département de Recherche Fondamentale, Centre d'Etudes Nucléaires, 85 X, 38041 Grenoble Cedex, France
100. R. A. Johnson, Department of Materials Science, University of Virginia, Charlottesville, VA 22903
101. W. G. Johnston, General Electric, Research and Development Center, P. O. Box 1, Schenectady, NY 12301
102. Adam Jostsons, Australian Atomic Energy, Commission Research Establishment, Lucas Heights, New South Wales, Australia
103. P. Jung, Institut für Festkörperforschung der Kernforschungsanlage, Jülich GmbH, D-5170 Jülich 1, Postfach 1913, Germany (BRD)
104. S. D. Harkness, Materials Science Division, Argonne National Laboratory, Argonne, IL 60439
105. M. Kiritani, Department of Material Physics, Faculty of Engineering Science, Osaka University, Toyonaka, Osaka, Japan
106. G. Kulcinski, Nuclear Engineering Department, University of Wisconsin, Madison, WI 53706
107. J. J. Laidler, Hanford Engineering Development Laboratory, P. O. Box 1970, Richland, WA 99352
108. V. Levy, Centre d'Etudes Nucleaires de Saclay, Boite Postale No. 2, 91190 Gif-sur-Yvette, France
109. Che-Yu Li, Department of Materials Science and Engineering, Cornell University, Ithaca, NY 14850
110. G. Martin, Centre d'Etudes Nucleaires de Saclay, Boite Postale No. 2, 91190 Gif-sur-Yvette, France

111. D. J. Michel, Naval Research Laboratory, Code 6390, Washington, DC 20390
112. T. E. Mitchell, Division of Metallurgy and Materials Science, Case Western Reserve University, University Circle, Cleveland, OH 44106
113. J. Moteff, Department of Materials Science and Metallurgical Engineering, University of Cincinnati, Cincinnati, OH 45221
114. F. A. Nichols, Materials Science Division, Argonne National Laboratory, Argonne, IL 60439
115. G. R. Odette, Department of Nuclear Engineering, University of California, Santa Barbara, CA 93107
116. W. Schilling, Institut für Festkörperforschung der Kernforschungsanlage, Jülich GmbH, D-5170 Jülich 1, Postfach 1913, Germany (BRD)
117. A. Seeger, Max-Planck-Institut für Metallforschung, Institut für Physik, D7000 Stuttgart 80, Büsnauer Strasse 171, Germany (BRD)
118. D. N. Seidman, Department of Materials Science, Bard Hall, Cornell University, Ithaca, NY 14850
119. E. P. Simonen, Battelle Pacific Northwest Laboratories, Richland, WA 99352
120. F. A. Smidt, Fuels Systems Branch, Department of Energy, Washington, DC 20545
121. B. N. Singh, Metallurgy Division, Danish Atomic Energy Commission, Research Establishment Risø, Roskilde, Denmark
122. R. E. Smallman, Department of Physical Metallurgy and Science of Materials, University of Birmingham, P. O. Box 363, Birmingham B15 2TT, England
123. M. V. Speight, Central Electricity Generating Board, Berkeley Nuclear Laboratories, Gloucestershire GL 13 9 rB, England
124. J. A. Sprague, Naval Research Laboratory, Code 6395, Washington, DC 20375
125. J. T. Stanley, College of Engineering Science, Arizona State University, Tempe, AZ 85281
126. J. L. Straalsund, Hanford Engineering Development Laboratory, P. O. Box 1970, Richland, WA 99352
127. H. Ullmaier, Institut für Festkörperforschung der Kernforschungsanlage, Jülich GmbH, D-5170 Jülich 1, Postfach 1913, Germany (BRD)
128. M. S. Wechsler, Department of Materials Science and Engineering, Iowa State University, Ames, IA 50010
129. H. Weidersich, Materials Science Division, Argonne National Laboratory, Argonne, IL 60439
130. P. Wilkes, Department of Nuclear Engineering, Engineering Research Building, University of Wisconsin, Madison, WI 53706
131. W. G. Wolfer, Department of Nuclear Engineering, Engineering Research Building, University of Wisconsin, Madison, WI 53706
- 132-133. DOE, Division of Materials Sciences, Washington, DC 20545
L. C. Ianniello
D. K. Stevens
134. DOE, Oak Ridge Operations Office, P. O. Box E, Oak Ridge, TN 37830
Director, Research and Technical Support Division
- 135-361. For distribution as shown in TID-4500 Distribution Category, UC-25 (Materials)