Indentation Creep Behavior and Microstructure of Cu-Ge Ferrites

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ABSTRACT

Cu-Ge ferrite was prepared using the standard ceramic method. The creep rate of polycrystalline Cu₁_x,GexFe₂₋₂xO₄ ferrite has been measured as a function of time at room temperature. It is found that the indentation length increases with the increase of both time and applied load. A regime of individual creep curves is observed for the first and second stages. It is not possible to record the third stage of the curve as usually happened in an ordinary creep test, because fracture of the samples does not occur. The slope is found to increase with increasing germanium and copper content in the steady state region. The high value of n (stress exponent factor) indicates that the dislocation creep is the dominate mechanism. The porosity arrangements developed within the specimens were examined using optical microscope. The results are discussed with regard to models describing the role of the steady state creep rate of metals. The morphology of the samples shows that the porosity is increased by increasing both copper and germanium ions.

Keywords: Ferrites, Indentation Creep, Microstructure

1. Introduction

Ferrites—ceramic ferromagnetic materials have been considered as highly important electronic materials for more than half a century. Ceramic like ferromagnetic materials, which are mainly composed of ferric oxide, α-Fe₂O₃, are called “ferrites”. Although the saturation magnetization for ferrites is less than half that of ferromagnetic alloys, they have advantages, such as applicability at higher frequency, greater heat resistance, higher corrosion resistance and lower cost. The practical applications of ferrites have been expanded by completely utilizing these advantages.

Individual creep curves generally show primary, secondary, and tertiary creep. The majority of the primary creep is not recoverable. The best representation of the data is one where the creep rate depends exponentially on the stress, rather than on the traditional power law.

The steady state rate equation for creep of crystalline materials is often represented by

\[ \dot{\varepsilon} = A \sigma^n d^{-p} e^{(-Q/RT)} \]  

where \( \dot{\varepsilon} \) is the strain rate, \( \sigma \) is the applied stress, \( A \) is a material parameter, \( d \) is the grain size, \( Q \) is the activation energy for creep, \( R \) is the gas constant, \( T \) is the temperature, \( n \) denotes the stress exponent factor and \( p \) is the grain size exponent.

Nishikawa et al. [1], Nishikawa and Okamoto [2] discussed the control of deformation mechanisms for creep under some conditions by analyzing stress, grain size and temperature dependency of steady state strain rate. According to their study, at intermediate stresses, power low creep is the dominant mechanism giving size of \( n = 3 \) to 5, \( p = 0 \) and \( Q = Q_L \), in which \( Q_L \) is the activation energy of the lattice diffusion. There are three distinct deformation mechanisms which exhibit the stress dependence. Two of them include the flow of vacancies from surfaces and grain boundaries under relative compression. In the case that vacancies flow through the crystal lattice, \( p = 2 \), \( Q = Q_L \) (Nabarro-Herring creep) [3,4]. In the case that the vacancies flow along grain boundaries, \( p = 3 \) and \( Q = Q_b \), where \( Q_b \) is the activation energy for grain boundary diffusion (Coble creep) [5]. The third process was reported firstly by Harper and Dorn [6] on aluminum, involves dislocation, multiplication and motion.
Nishikawa et al. [1], Nishikawa and okamoto [2] studied Mn-Zn ferrite (Mn_{0.5} Zn_{0.5} Fe_{2} O_{4}) with a wide range of temperature, stress and grain size. They showed convincing evidence for the operation of diffusional creep at low stress and small grain size accordingly the strain rate can be expressed in the form {Norton–Baley–Arrhenius (NBA) model [7,8 ]} : 

\[ \dot{\varepsilon} = A \sigma^n e^{-Q/RT} \]  

(2)

The diffusion that drives creep requires vacancies to operate. An atom can move to a site of an adjacent vacancy, provided that it has adequate thermal energy to jump from its original site (Figure 1. [9]). At any temperature, the average thermal energy of an atom is 3 kT, where k is Boltzmann’s constant (1.38 × 10^{-23} J/K). As atoms vibrate about their mean positions, they collide with their neighbors transferring energy continually from one to another. As a result, the thermal energy is not homogeneously distributed among the atoms in the crystal, and at any instant, any single atom has more or less energy than the average value. Even if the atom has sufficient energy to jump into a neighboring lattice site, it can move only if a vacancy exists on that lattice site to allow the movement to occur. The likelihood of the two independent processes above occurring simultaneously is the product of the individual probabilities. The rate of diffusion in a solid also depends on the frequency with which individual atoms move. Surfaces, grain boundaries, and dislocation cores provide “easy paths” for diffusion, since atomic arrangements are less regular there than in a perfect lattice. Activation energies are lower for these easy path processes than for diffusion through the bulk of the crystal.

The purpose of this study is to evaluate the creep behavior of Cu-Ge ferrite and the mechanism controlling such ferrites.

2. Experimental Technique

2.1. Samples Preparation

The ferrite, Cu_{1+x}Ge_{x}Fe_{2-x}O_{4} with x =0, 0.2 and 0.4 were prepared using the standard ceramic technique. High-purity oxides of CuO, GeO_{2} and Fe_{2}O_{3} were used as starting materials. The weighted oxides with the required mole ratios were thoroughly mixed, grounded and then pre-sintered at 1023 K for 5 h soaking time. Afterwards, the fine powder was pressed under 3 tons to form disk shape of 13mm diameter. The disks were sintered at 1273 K for 10 h and then slowly cooled down to room temperature with cooling rate of 2 K/min.

The samples were prepared for creep indentation measurements by making the two opposite surfaces quite parallel using a polishing machine with different grad emery papers 300, 600, 800, 1200 and 2000, respectively. The polishing process was performed at rotating speed of 300 rpm under water cooling. Electrochemical polishing which is necessary to eliminate the surface strains was also considered during mechanical polishing.

2.2. Indentation Creep Tests

The indentation tests were performed using a Vickers hardness tester (AKASHI MVK-H2, Japan) where the applied load (F) in Newton and testing time (t) in minute are the two only variables. The applied loads were 0.49, 1.96 and 4.9 N using a 136° pyramid diamond indenter with a square base. The indentation times were 2, 5, 10, 15, 20, 25, 30, 40, 50 and 60 minutes. Each reading was an average of five measurements taken at random places on the surface of each samples. The indentation locations were separated by a distance of at least 2 mm and away from the edge of the specimen by about 2 mm.
2.3. Microstructure

The microstructure of the ferrite specimens was obtained using Nikon optical microscope model (M 001-378, Japan) provided with a computerized camera (FUJIX Digital Camera HC-300 I). An electrochemical etching is performed in an etching cell with Pt hollow cylindrical cathode, where the sample is placed as an anode in the center of cylindrical cathode. The samples were etched in a solution composed of 90% acetic acid and 10% perchloric acid for 20 sec. After that the samples were washed using deionized water and finally cleaned in acetone using ultrasound cleaner.

3. Results and Discussion

3.1. Creep Behavior

Figure 2 shows the variation of indentation diagonal length d versus time under constant loads of 0.49, 1.96 and 4.9 N for the studied ferrite with different Cu and Ge concentration. It can be seen from the figure that the indentation length increases with the increase of both time and applied load. It also shows the two stages which are similar to an ordinary creep curve. The first stage indicates a rapid increase in indentation length up to 20 min for all samples when the load is at its highest value (4.9 N) followed by the second stage with a steady or linear increase. Since compression is used as the hardness test, fracture of the samples does not occur. Therefore, it is not possible to record the third stage of the curve as usually happened in an ordinary creep test. In addition, the figure shows that the level in the steady state region and slope are increased by increasing the germanium and copper concentration instead of iron. This implies that as Ge⁴⁺ and Cu²⁺ ions increase, room temperature creep occurs more rapidly.

Figure 3 shows the rate of variation in indentation length (d’) against dwell time, which obtained by differentiation of the curves in figure 1 to show the effect of increasing copper and germanium more clearly. One can see from this figure that there is a sharp drop in the creep rates at short times corresponding to the primary creep stage, followed by a lower decreasing rate region which corresponding to the steady state stage of the studied compositions. It is further demonstrated that samples with high concentration of both copper and germanium ions have higher creep rates than that of lower concentrations. It is also observed that the almost for all compositions, there is exists of an apparent steady state.

To obtain the steady state creep exponents, the following equation of creep analysis was applied to the indentation data of the samples [10].

\[ n = \left[ \frac{\delta \ln \ln d}{\delta \ln \ln H_v} \right] \]  

where \( H_v \) is the Vickers hardness number which calcu-
Figure 3. Indentation length rate (ḋ) plotted against dwell time for 0.49 up to 4.9 N loads for Cu₁₋ₓGeₓFe₂O₄.

Figure 4. Plots of ln (ḋ) versus ln (HV) at different loads to determine the stress exponent for the studied ferrites.

is the applied load.

The rate of diagonal variation is plotted against the Vickers hardness number as shown in Figure 4 for all samples and testing loads. A straight line is resulted from the least square fitting of all data points using origin lab program for each sample and load. M. Sternitze and H. Hubner [12] studied the creep properties for Al₂O₃/ SiC nano-composite and they found that the stress exponent factor (n) ranged from 2.5 to 5.5.

Creep stress exponent can help to narrow the field...
among many theoretical possibilities [13]. It was reported that diffusion creep is associated with n value around one [14], grain boundaries sliding leads to n values close to 2 [15] and dislocation climbs responsible for n values in the range of 4 - 6 [16]. Goetze [17] and Durthan [18] indicated that the dislocation creep is the dominating mechanism when the values of the stress component n lie between 3 and 10. In our present work the relatively high values of n (3 - 14.5) imply that the operative creep mechanism in the studied samples is dislocation creep.

The variation of hardness with dwell time due to indentation creep is shown in Figure 5. It is obvious that hardness decreases with increasing the dwell time. The decrease is very sharp in the first stage followed by a linear decrease with increasing time.

Kagiarova et al. [19] stated that the low value of the stress exponent is consistent with samples with low cavity concentration. According to the morphology of the investigated samples, the porosity increases by increasing copper and germanium concentration; accordingly the exponent stress factor would be increased too. As a result of that, we can assume that a higher concentration of dopent associated with the enhancement of cavity formation. Moreover, the indentation caused by the applied stress may create new cavities. Then, as the stress increases the contribution of cavitations is expected to increase and hence the contribution to the creep rate will increase and consequently the exponent stress factor will increase.

3.2. Microstructure

In order to give an accurate view about the morphology, optical microstructure for Cu-Ge ferrites are depicted in Figure 6. The micrographs of the studied samples reveal circular pores narrowly distributed, spread throughout the entire microstructure. The interaction of grain boundary and porosity along with sintering temperature is important in determining the limiting grain size [20]. When many pores are present and the sintering temperature is not too high, grain growth is inhibited.

Figure 6 shows that the porosity of the samples are increased by increasing Ge and Cu ions content which making the density to decreases. When substituting Ge in small amount, the effect on the microstructure of Cu-ferrite can be explained in terms of the increased pore mobility. This is due to the creation of excess cation vacancy formation by doping the ferrite with high valence cations which can be argued from the site and change neutrality as well as the oxidation-reduction equilibrium of iron. It has been reported that the presence of large pores is due to the amount of metal ion vacancies caused by oxidation [21]. This structure indicates that the samples can easily exhibit absorption and condensation of water vapor as Ge$^{4+}$ ions increase. According to Hemeda [22,23] and Ecklet [24] the increase of Cu$^{2+}$ ions leads to the increase of porosity. Also Visser et al. [25] state that
the non magnetic particles and trace amount of impurities forming a non magnetic grain boundary leads to the reduction of grain size and increasing of porosity. Accurate measurements of the grain size were hard to obtain because of the difficulty to observe grain boundary by using optical microscopy.

4. Conclusions

1) The indentation length increases with the increase of both time and applied load.

2) By increasing the germanium and copper concentration on the expenses of iron, room temperature creep occurs more rapidly.

3) The samples with high concentration of both copper and germanium ions have higher creep rates than that of lower concentrations.

4) The relatively high values of, creep stress exponent (3 - 14.5) imply that the operative creep mechanism in the studied samples is dislocation creep.

The porosity of the samples is increased by increasing Ge and Cu ions content.

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