Effect of loading on recovery of internal friction of AI-16 Wt % Ag and AI-16 Wt % Ag-0.28 Wt % Fe alloys

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The phenomenon of internal friction recovery of AI-16 wt %Ag and AI-16 wt %Ag-0.28 wt % Fe alloys is studied after cold-working the samples in uniaxial tension using the free decay method. The wire samples are investigated for internal friction recovery while they were loaded-within the elastic limit-by different loads at different resting temperatures. It is found that by increasing the temporary loading the rate of recovery decreases. TEM investigations confirmed that the addition of Fe to AI-Ag alloy accelerates the formation and coarsening of G.P.zones. The results are explained on the basis of dislocation damping adopting the Granato-Lücke model for recovery of internal friction. © 2001 Kluwer Academic Publishers

1. Introduction

The basic sequence of decomposition in Al-Ag Alloy has been found as: α (f.c.c.) \rightarrow Spherical G. P. zones (tetrahedral) \rightarrow metastable γ `-phase \rightarrow equilibrium γ phase (hcp) [1]. Dubey *et al.* [2] using small angle X-ray scattering (SAXS) studies, concluded that the composition of G.P.zones (\sim Ag₂Al) remained constant over a wide temperature range from 20 to 250 °C. The experimental evidence for the formation of an ordered η state of G.P.zones below 170 °C and a disordered ε -state above this temperature was obtained by many authors [3, 4].

The transient increase of internal friction, Q^{-1} , observed in metals after plastic deformation is known to decrease more or less rapidly as a function of time. Such a phenomenon is known as the recovery of internal friction [5]. According to Koehler [6] and Granato and Lücke [7] models the increase of internal friction (IF) after plastic deformation is interpreted as being caused by the creation of a large number of new, freely moving dislocations. The subsequent disappearance of the effect (i.e., recovery of IF) is ascribed either to the pinning of dislocations by points defects and secondphase particles or rearrangements of dislocation networks resulting in an overall decrease in dislocation damping.

Reviewing the literature of previous work on the recovery of IF using torsion pendulum one can recognize - as a matter of experimental procedure-that the stress acting on the test sample is virtually zero. Now a question arises as follows: if the test sample is loaded -within the elastic limit-for certain periods of time just before measuring its IF, how such loading will affect the recovery behavior. No studies were made on the effect of loading on the recovery of IF as affected by second phase particles. Hence the present work is devoted to investigate the effect of loading on the recovery behavior of IF in samples of Al-16 wt %Ag and Al-16 wt %Ag -0.28 wt %Fe alloys strained plastically in tension immediately before measurements. Here, an ordered η -state of G.P.zones formed by aging in the appropriate temperature range represent the second phase particles to be considered.

2. Experimental procedure and technique

Two alloys each containing Al-16 wt % Ag were made by melting silver of purity 99.999% with either pure Al (99.999%) (Alloy A) or with Al containing 0.28 wt % Fe (alloy B). Each alloy was melted under vacuum in a high purity graphite crucible. After annealing for homogenization at 823 K for 4 days, the ingots were swaged and cold drawn into wires 0.35 mm in diameter for IF measurements, or rolled into strips 0.2 mm in thickness for electron microscopy investigation. Chemical analysis revealed that samples composition is very close to the weighed-in compositions.

A solution treatment was carried out by heating the samples at a temperature above solvus (\sim 823 K) for 2 h and samples were then quenched rapidly in cold water and immediately aged at 428 K for 2 h. Internal friction was determined by the free decay method by means of a low-frequency (~ 1 Hz) torsion pendulum with electromagnetic excitation and optical read out. The uniaxial tensional deformation was carried out "in situ" only at room temperature. The tensional deformation $\varepsilon = \Delta \ell / \ell$ (where $\Delta \ell$ is the increase in the length of the samples and ℓ is its original length) was 0.04 for all samples. The IF, Q^{-1} , was first measured before exerting tensional deformation upon samples, this is denoted $Q_{\rm u}^{-1}$ (i.e. Q^{-1} corresponding to the undeformed state). Immediately after deformation ($\varepsilon = 0.04$) the IF corresponding to a time t = 0, (Q_0^{-1}) is determined. Then the sample wire is set at a certain testing temperature,

and a certain load is applied for a certain time ranging from 0.5 min at the beginning of the recovery process to 3 min at its end. The load is then removed and IF, Q_t^{-1} , is measured. The above step is repeated by loading the sample for an additional period of time, then the load is removed and Q_t^{-1} is again measured. In every step Q_t^{-1} is determined where *t* is the time measured immediately after the tensional deformation. The short intervals during which the samples was unloaded for IF measurement are actually involved in *t*, thus a negligibly small error in Q_t^{-1} value exists. Calculations which took such an error into consideration proved that it has no appreciable effect on the value of Q_t^{-1} as a function of *t*. The recovery behavior under different loads 0.45, 0.95, 1.45, and 1.95 N was studied at testing temperatures of 293, 323, 343, and 363 K.

The sheet samples -for electron microscopy investigation- were prepared by electropolishing using a twin jet machine. The electrolyte was 10% perchloric acid and 90% ethanol at 273 K. The voltage and current values were around 15 to 25 V and 10 to 20 mA respectively. Microstructures were examined using^a, JOEL-100 transmission electron microscope operating at 100 kV.

3. Experimental results

The recovery behavior of IF is presented as a relation between the normalized IF $S(t) = (Q_t^{-1} - Q_u^{-1})/(Q_0^{-1} - Q_u^{-1})$ and the time *t*, (*S* is a fractional quantity which represents the freely moving dislocations i.e. those dislocations contributing to the damping effect). Fig. 1a and b shows the recovery of IF for both alloys A and B respectively as a relation between the normalized IF, *S*, and time *t* under the effect of elastic stressing by different loads at different testing temperatures mentioned before. From this figure it is seen that –for both alloys- the recovery behavior can be divided into three stages namely, a rapid initial stage, intermediate stage and finally a slow saturation stage. We will be only interested in the first and second stages. Generally it is clear that *S* takes larger values by increasing both testing temperatures and the loads. It is worth noting that under the same conditions, corresponding



Figure 1 (a and b) Normalized IF, (*S*), of deformed and then loaded samples as a function of time *t* for both alloys A and B respectively. Different loads applied are (\Box) 0. 45, (\triangle) 0. 95, (\bullet) 1. 45, and (\bigcirc) 1. 95 N. The testing temperatures applied are; a) 293, b) 323, c) 343, and d) 363 K.

variations in *S* are appreciably higher in alloy B than in alloy A.

4. Discussion

The Granato-Lücke [7, 8] break-away mechanism for recovery of IF has been adopted here to explain the results of the present work. It is well known that IF and its recovery are both influenced by the presence of solute atoms and second-phase particles [9-15]. In the present work, the G.P.zones are considered as the second phase-particles that affect the recovery behavior (see Fig. 2) where they act as pinners for dislocations [16–19]. Before cold-working the heat treated samples, the interaction between the pre-existing dislocations and G.P.zones gives a value of IF equal to Q_{μ}^{-1} . Immediately after deformation, the freshly created dislocations are vibrating with full network loop length, causing the highest value of IF, Q_0^{-1} . The time dependence of Q_t^{-1} (and consequently S) after straining as shown in Fig. 1 can be explained in view of the rate of interaction of G.P.zones with dislocations. According to Cottrell and Bilby [20], the number of second-phase particles (G.P.zones here) per unit volume which interacts with dislocation proceeds by a relatively higher rate immediately after deformation (rapid stage of recovery). As a result, the average dislocation loop length $L_{\rm p}$ ($L_{\rm p}$ is the average distance between zone pins and is less than the full network loop length) and consequently S are both reduced by a higher rate. In the intermediate stage of recovery, due to the relatively lower rate of interaction between zones and dislocations, a relatively lower rate of reduction of $L_{\rm p}$ results in a lower rate of recovery. Finally a slow saturation stage is reached corresponding to a minimum interaction of G.P.zones with dislocations, i.e. minimum value of L_p .

Particularly for alloy B, the higher value of S and lower rate of IF recovery as compared with corresponding values for alloy A may be explained as following. For alloy A, immediately after a minimum time of aging, the number of G.P.zones is small. As the time of aging proceeds, new zones are to be formed leading eventually to a larger number of small-sized zones. In alloy B (Al-Ag-Fe) due to binding between vacancies and Fe solutes, vacancy-impurity clusters are likely to be formed [21]. These clusters might act as nucleation sites for a large number of small zones. Thus a large number of zones is expected to be formed immediately after a minimum time of aging. In the course of aging time, growth of zones on the expense of their number occurred leading to existence of a smaller number of large-sized zones (compare Fig. 2 with Fig. 3). These large-sized zones of smaller number should be less effective as pinners for dislocation [19] and hence produce less recovery.

The recovery of IF which occurs during the sequence of time after deformation could be controlled by an Avrami-type relation [22] of the form:

$$S(t) = \exp(-Kt^n) \tag{1}$$



140.000 x

Figure 2 Thin foil micrograph from Al-16 wt % Ag samples heated at 428 K for 2 h, showing G.P.zones.



140.000 x

Figure 3 Thin foil micrograph from Al-16wt % Ag-0. 28wt% Fe samples heated at 428 K for 2 h, showing G.P.zones of large sizes.

n represents the power of time dependence of damping and K is the decay rate. To calculate both n and K under the experimental conditions applied in the present work, Equation 1 can be written in the form:

$$\ln[\ln(S^{-1})] = n\ln t + \ln K.$$

Fig. 4a and b are plots of $\ln[\ln(S^{-1})]$ versus $\ln t$ for samples loaded to different values at different testing temperatures for both alloys A and B respectively. At each testing temperature the intercept of $\ln[\ln(S^{-1})]$ axis in Fig. 4 represents $\ln K$, from which the decay rate K can be obtained. From this figure, it is clear that two linear parts with different slopes n are distinguishable. They are considered to represent two stages.

4.1. Rapid stage of recovery

The power n of time dependence of damping as calculated from the slopes of the parallel straight lines in Fig. 4 has an average value of unity, for both alloys, indicating that the operating mechanism for the decay of IF in this stage (stage I) is the same irrespective of the conditions of loading and testing temperatures applied. However, this value is high as compared with the ordinarily known limits [8] which range between 0.33 to 0.7. It is clear that the relatively high value of n obtained here indicates that the responsible mechanism is highly



Figure 4 (a and b) Plot of $\ln(\ln S^{-1})$ and $\ln t$ for differently loaded samples a) 0. 45, b) 0. 95, c) 1. 45, and d) 1. 95 N. The testing temperatures applied are; (\bigcirc) 293, (\bigcirc) 323, (\triangle) 342, and (\square) 363 K.

time-dependent. In view of Granato-Lücke model, this may be attributed to the relatively large numbers of G.P.zones which act as pinners.

Under the effect of loading, the crystal is elastically strained due to the tensional forces acting on it. During loading with a load P, the elastic energy (E) stored in the crystal is proportional to P^2 [23]. Fig. 5a and b shows the dependence of decay rate K on the applied load, P, as a relation between K and P^2 . From Fig. 5, it is clear that - at a certain testing temperature- the value of K decreases linearly with the elastically stored energy E. This energy which is stored in the crystal during loading is assumed to enable moving dislocations to surmount the G.P. zones which act as pinning points for the moving dislocations, i.e. reducing the rate of pinning exerted by G.P.zones on dislocations. Thus redistribution of dislocation loop lengths will be in favor of magnification of the effective average loop length $L_{\rm p}$. Such a magnification proceeds with a higher rate as the load applied is larger, accordingly, the value of K decreases. Increasing testing temperature which leads to a decreasing in K value may explained as being due to a higher rate with which the redistribution process just mentioned takes place.

4.2. Intermediate stage of recovery

The power *n* of time dependence in this stage (stage II) was found to have an average value of 0.7 for alloy A and 0.4 for alloy B. These values is in good agreement with the ordinarily known limits mentioned before. Fig. 6a and b shows the relation between K and P^2 where straight lines are obtained as in previous stage (stage I), but with a lower value in K as compared with those having the same testing temperatures in stage I.

Since the two groups of curves in both stages are similar so, one can conclude that the operating mechanism for the recovery behavior -for both stages- is the same. The difference is that the rate with which the mechanism proceeds in stage II is lower. This is acceptable in view of the fact that recovery is basically a timedependent process i.e. during stage II the "residual" fraction of dislocations that contribute to damping being less in number should correspond to a lower rate of recovery as compared to that in stage I.

4.3. Activation energy calculation

To calculate the activation energy of the operating mechanism for the recovery behavior, the procedure





Figure 5 (a and b) Relation between the decay rate (*K*) for stage I and the square of the applied load P^2 for both alloys A and B respectively. The testing temperatures applied are; (\bigcirc) 293, (\bullet) 323, (\triangle) 342, and (\Box) 363 K.

Figure 6 (a and b) Relation between the decay rate (*K*) for stage II and the square of the applied load P^2 for both alloys A and B respectively. The testing temperatures applied are; (\bigcirc) 293, (\bullet) 323, (\triangle) 342, and (\Box) 363 K.

used by Graiss *et al.* [23] is followed. Fig. 7a and b shows a relation between the slopes $(\partial K/\partial p^2)$ of the straight lines of Figs 5 and 6 and 1000/*T* for both alloys A and B respectively, where *T* is the testing temperature in K. For both stages, the activation energy yielded a mean value of about 14 ± 0.7 kJ/mol for alloy A and a mean value of nearly 18 ± 0.9 kJ/mol for alloy B. In view of such a low values for the activation energy we assume the recovery mechanism not to be diffusion. A similar conclusion was reported by Kenawy *et al.* [13] as a result of their early work on Al-Fe alloy. Adopting the Granato-Lücke model [7, 8] for the recovery of IF as being the loop length redistribution due to the change in concentration of pinning points, we may assume that the mechanism responsible for the recovery process is the pinning imposed to vibrating dislocations by G.P.zones. When the dislocation line acquires an energy higher than 14–18 kJ/mol it can be able to surmount the G.P. zones which act as pinning points for the moving dislocations. Depending on the amount of energy supplied to the sample either thermally and/or by loading, we may except different concentrations of pinning points on the dislocation line leading to different network lengthes, L_p . At higher testing temperatures and/or applying larger loads, large values of L_p are progressively obtained corresponding to lower values of decay rate K. In view of this, the values of the slopes $(\partial K/\partial p^2)$ at different testing temperatures in Figs 5 and 6 are considered to reflect the rate of pinning of dislocations by G.P. zones. Since the values of



Figure 7 (a and b) Arrhenius plot of $\ln(\partial K/\partial p^2)$ as a function of $10^3/T$ for both alloys A and B respectively. Stage type is indicated.

activation energy for both alloys are very close, so the same pinning mechanism for the recovery process is applied.

References

- 1. D. B. WILLIAMS, Metals Trans. 19A (1983) 1745.
- 2. PH. A. DUBEY, B. SCHONFELD and G. KOSTORZ, Acta Metall. mater. **39** (1991) 1161.
- 3. G. DLUBEK, G. WENDROCK and K. PAWELZYK, *Phys. Stat. Sol. (a)* **140** (1993) 311.
- 4. A. MALIK, B. SCHONFELD and G. KOSTORZ, Z. Metalle. 88 (1997) 625.

- 5. A. S. NOWICK, J. Appl. phys. 25 (1954) 1129.
- J. S. KOEHLER, "Imperfections in Nearly Perfect Crystals" (J. Wiley, 1952) Ch. 7.
- 7. A. V. GRANATO and LÜCKE, J. Appl. Phys. 27 (1956) 583.
- R. DE. BATIST, "Internal Friction Of Structural Defects In Crystalline Solids" (North-Holland Publ. Co., 1972) p. 367.
- 9. T. MORI, M. OKABE and T. MURA, *Acta. Metall.* **28** (1980) 319.
- 10. M. OKABE, T. MORI and T. MURA, *Phil. Mag. A* 44 (1981) 1.
- 11. R. MONZEN, K. SUZUKI, A. SATO and T. MORI, *Acta. Metall.* **31** (1983) 519.
- 12. G. GRAISS and G. SAAD, *Phys. Stat. Sol.* (a) **92** (1985) 489.
- 13. M. A. KENAWY, G. GRAISS, G. SAAD and A. FAWZY, *ibid.* **107** (1988) 71.
- 14. G. GRAISS, G. SAAD, L. N. MAKAR, Solid State Commenication 70 (1988) 375.
- 15. G. GRAISS, G. SAAD, L. N. MAKAR and M. A. MAHMOUD, *Egypt. J. Sol.* **14** (1991) 1.
- 16. T. KANADANI and A. UMADA, *Phys. Stat. Sol. (a)* **148** (1995) k23.
- 17. Idem., ibid. 151 (1995) k29.
- G. H. DEAF, S. B. YOSSEF, M. A. MAHMOUD, G. GRAISS and M. A. KENAWY, *ibid.* 158 (1996) 471.
- 19. G. H. DEAF, S. B. YOSSEF and M. A. MAHMOUD, *ibid.* **168** (1998) 389.
- 20. A. H. COTTRELL and BILBY, *Proc. Phys. Soc. A.* **62** (1949) 49.
- 21. M. DOYAMA, J. Nucl. Mater. 70 (1978) 350.
- G. GRAISS, G. SAAD, L. N. MAKAR; Journal of Material Science 25 (1990) 1033.
- 23. I. G. GRAISS, G. SAAD and A. FAWZY, *Phys. Stat. Sol. (a)* **110** (1988) 481.

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