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Regular article Magnetostriction enhancement of Fe₇₃Ga₂₇ alloy by magnetic field annealing

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ARTICLE INFO

ABSTRACT

Article history: Received 3 December 2017 Received in revised form 2 January 2018 Accepted 4 January 2018

Keywords: Fe-Ga alloy Magnetostriction Nanoparticles Magnetic field annealing

Magnetostrictive materials are playing an increasingly important role in actuators, transducers, and torque sensors, among others [1]. Terfenol-D materials possess giant magnetostriction, but they are extremely brittle and their saturation magnetic fields are too high, limiting practical applications [2]. Meanwhile, Fe-Ga alloys are promising magnetostrictive materials since they show high magnetostriction under low magnetic field [3], high mechanical properties [4], and a high Curie temperature (Tc > 650 °C) [5]. The introduction of Ga into the body-centered cubic (bcc) α -Fe may result in several crystal structures such as disordered bcc A2, ordered bcc B2 and D0₃, ordered fcc L1₂ and hcp D0₁₉ [6,7]. So the magnetostriction of Fe-Ga alloys varies with Ga concentration. There are two peaks in the curve of magnetostriction vs Ga concentration in Fe-Ga alloys [8]. The first peak at near Ga-19% is thought to be attributed to the short range order of Ga atoms pairs along the [100] axis of the A2 structure [3], and the second peak at Ga-27% due to the softness of shear elastic constant $\frac{1}{2}(C_{11}-C_{12})$ [8,9]. Previous investigations were mainly focused on the near Ga-19 at.% alloy [10–15]. The magnetostriction of near 27 at.% Ga alloys sharply decreases when the large-area fcc ordered phase appears in the A2 matrix [16]. However, the researches show that the heterogeneities resulted from the nanoscale precipitations can result in local strain in the surrounding of A2 matrix and is thought to be helpful to magnetostrictive property [17-21]. This indicates that the magnetostriction of Fe-Ga alloys is sensitive to structure and heterogeneity of phases. These facts give us clues for reviewing the important role of nanoprecipitations in Fe-Ga magnetostrictive materials. Given that nanoparticles are

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generally precipitated by aging or annealing for a relatively long time, in this work, numerous multiphase nanoparticles precipitated when magnetic field annealing was applied to the Fe₇₃Ga₂₇ alloy in a very short time. Moreover, the saturation magnetostriction λ s (λ s = $\lambda_{//}$ –

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In this work, on the basis of multiphase characterics of near Ga-27 at.% alloy, magnetic field annealing was applied

in a directional solidification (DS) $\langle 001 \rangle$ oriented Fe₇₃Ga₂₇ polycrystalline alloy, and the saturation magnetostric-

tion of 370 ppm under low magnetic field has been achieved. Large numbers of multiphase nanoparticles are pre-

cipitated in this process; also the magnetic heterogeneities are used to enhance magnetostriction. Meanwhile,

this method provides an approach for developing excellent magnetostrictive materials.

 λ_{\perp}) of 370 ppm under a low magnetic field has been achieved in DS (001) oriented Fe₇₃Ga₂₇ polycrystalline alloy. The directional solidification (DS) (001) Fe₇₃Ga₂₇ alloy was prepared from high pure Fe (99.9 wt%), Ga (99.99 wt%) by induction melting under the protection of argon gas and grown by the DS processes at a growth rate of 720 mm/h. The ingot was annealed at 1100 °C for 3 h in an argon atmosphere. Samples were cut from the ingot. Each sample was cut into 5 mm length and 4 mm wide for the strain gauge and magnetic field annealing, and its thickness was about 0.3 mm. The length direction (LD) of all samples was along the (001) direction of columnar crystals. The specific heat treatments methods of samples (shown in Table 1) were based on the differential scanning calorimetry (DSC) curve, as shown in Fig. 1(a). The sample was rapidly heated to 720 °C, and held for 2 min under a magnetic field of 1200 Oe along LD, which was higher than the value of saturation magnetic field measured from the magnetostriction curves of the Fe₇₃Ga₂₇ alloy (shown in Fig. 4). Finally, the sample was cooled at a rate of 100 °C/min under a magnetic field. Fig. 1(b) showed the crystal orientation information of the DS Fe73Ga27 alloy. As can be seen, the DS Fe73Ga27 sample consisted of several large columnar grains with a (100) preferred orientation. Electron backscatter diffraction (EBSD, Zeiss, z5) was implemented to obtain the grain orientation information, and FEI Tecnai G² F30 equipment was employed to get TEM images free of oxidation. The magnetostric-

Fig. 2 shows the selected area electron diffraction (SAED) patterns of samples in different states with the beam direction parallel to the [110] direction of A2 and B2 phases. The SAED patterns of samples a1, a2, a3

tions were measured using a standard strain gauge technique.







Table 1The specific heat treatments methods of samples.

Sample	Treatment	Sample	Treatment
a1 a2 a3	860 °C/2 h 860 °C/2 h + 720 °C/2 min 860 °C/2 h + 720 °C/1200 Oe/2 min	b1 b2 b3	700 °C/20 h 700 °C/20 h + 720 °C/2 min 700 °C/20 h + 720 °C/1200 Oe/2 min

are shown in Fig. 2(a1), (a2), (a3), respectively. As can be seen in Fig. 2(a1) and (a2), diffraction spots indexed to $(01\overline{1})$, $(\overline{2}00)$ and $(\overline{2}1\overline{1})$ planes correspond to bcc A2 phase indicated with white lines, and

 $(\overline{111})$, $(\overline{111})$ and $(\overline{200})$ planes correspond to bcc D0₃ phase indicated with red lines. In Fig. 2(a1) and (a2), there are also two uncalibrated light spots pointed by two black arrows in Fig. 2(a2) symmetrically distributing on both sides of the [$\overline{100}$] direction of A2 matrix, and this fact indicates that the matrix lattice is not that purity. In Fig. 2(a3), besides A2 and weakened D0₃ phases, the diffraction spots indexed to (100), $(01\overline{1})$ and $(11\overline{1})$ planes correspond to fcc L1₂ phase indicated with blue lines, and $(\overline{111})$, $(\overline{311})$ and $(\overline{200})$ planes correspond to fcc A1 phase indicated with yellow lines (referring to fcc Aluminum phase, group number: 225, a = b = c = 4.0494 Å). Furthermore, a variant of L1₂ phase appears in Fig. 2(a3) indicated with blue lines, two uncalibrated



Fig. 1. (a) DSC curve of the DS Fe₇₃Ga₂₇ polycrystalline alloy with a (001) preferred direction, (b) Crystal orientation information along LD.



Fig. 2. SAED patterns of samples in different states with the beam parallel to the [110] direction of A2 and B2 phases. Images of (a1), (a2), (a3), (b1), (b2), (b3) represent the SAED image of corresponding state sample.

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