



## Dynamic coercivity of Mo-doped FINEMETs

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### ABSTRACT

The structure and the dc magnetic behavior of FINEMET-type alloys doped with molybdenum have been recently reported. Most commercial applications of these materials are, however, not at dc but at high magnetizing frequencies. Therefore, we report a study of the frequency dependence of coercivity,  $H_c(f)$ , in amorphous and nanocrystalline ribbons of composition  $\text{Fe}_{73.5}\text{Si}_{13.5}\text{Nb}_3\text{B}_9\text{Cu}_1$  ( $x=0, 1.5$  and  $3$ ) in the frequency range from  $0.5$  to  $1.3$  kHz. The nature of  $H_c(f)$  measurements revealed the influence of eddy currents in the magnetization of samples. The frequency dependence of coercivity did not vary with the molybdenum content in the amorphous samples. All the alloys exhibited a systematic improvement in the coercivity after nanocrystallization and it was found that this improvement was better as more Nb was replaced by Mo.

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### 1. Introduction

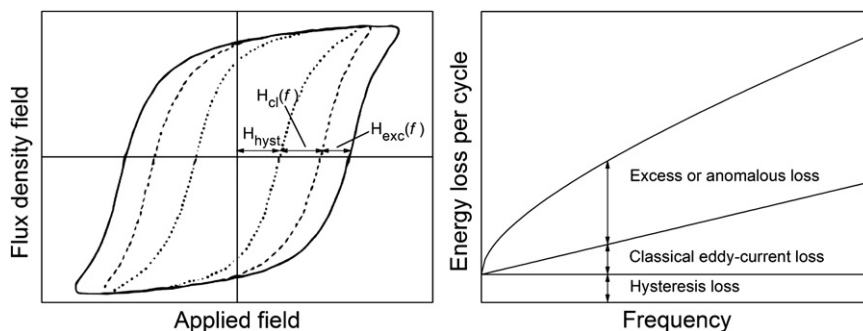
Over the past decades, there has been an immense interest in nanocrystalline soft magnetic materials due to their attractive magnetic properties compared to conventional crystalline alloys. In 1988 Yoshizawa et al. [1] developed the first nanocrystalline alloy, called FINEMET ( $\text{Fe}_{73.5}\text{Si}_{13.5}\text{Nb}_3\text{B}_9\text{Cu}_1$ ), with low coercivity, saturation magnetization and high resistivity. The outstanding properties of FINEMET are due to its special two-phase structure, i.e.,  $\alpha''\text{-Fe}_{1-x}\text{Si}_x$  crystals of  $10\text{--}20$  nm size embedded in an amorphous ferromagnetic matrix. This material is usually fabricated by a controlled annealing of an amorphous precursor, previously casted by the melt-spinning technique in the form of a ribbon. FINEMET-like alloys are suitable for a wide range of applications, particularly in ac field such as transformer cores, inductive devices, sensors, etc. [2]. It is therefore important to know the frequency dependence of their magnetic properties.

The coercive field or coercivity ( $H_c$ ) is defined as the field at which the magnetization is reduced from remanence to zero. For a magnetization loop with peak magnetization  $M_{\text{max}}$ ,  $4H_cM_{\text{max}}$  gives the order of magnitude of the loop's area. Thus,  $H_c$  is a measure of the energy that is dissipated as heat in each magnetization cycle. The losses can be conventionally divided into three categories: hysteresis, classical eddy current and anomalous or excess components (Fig. 1). This division is artificial, since they

actually do not arise from different sources, but allows treating loss mechanisms occurring on different space–time scales separately as if they were independent of each other. This subject is treated in detail in Refs. [3,4]. Hysteresis loss is due to the Barkhausen jumps between local energy minima, it is determined from the area of the static hysteresis loop and is assumed to be independent of the frequency of magnetization. The classical eddy current loss per cycle, which varies linearly with frequency, can be calculated from Maxwell's equations for a perfectly homogeneous material with no domain structure, where the boundary conditions of the problem are given by the geometry of the specimen. The so-called anomalous or excess loss increases with frequency but not linearly; it usually exhibits a rapid increase at low frequencies and then an almost constant positive slope. This component appears because the classical calculation of eddy current loss ignores the presence of domains and domain walls motion, which is damped by the eddy currents. As observed earlier, the three contributions to the energy loss have different dependences on with the magnetizing frequency. In spite of this, the total coercivity or loss has been sometimes approximated to one frequency dependent term  $\sim f^\alpha$  [5–8].

In previous works, the structure and some magnetic properties of a series of FINEMET-like alloys with a partial replacement of Nb by Mo were studied. In Ref. [9] the magnetostrictive behavior of these samples was investigated, but no influence of the refractory elements exchange on the saturation magnetostriction ( $\lambda_s$ ) was observed either for the as-quenched ( $\sim 19$  ppm) or for the annealed samples ( $\sim 2$  ppm). The phase transformations occurring during the crystallization process of the alloys were studied

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**Fig. 1.** Left: contributions to the coercivity in the hysteresis loop—static hysteresis, classical eddy currents and anomalous or excess field (modified from Ref.[3]). Right: contributions to the energy loss per cycle as a function of the frequency of the magnetizing field.

in Ref. [10] by a variety of techniques, including differential scanning calorimetry (DSC), differential thermal analysis (DTA), thermogravimetric analysis (TGA) under a magnetic field, X-ray diffraction (XRD) and transmission electron microscopy (TEM). The thermal stabilities of both the ferro-paramagnetic transition of the amorphous phase and the precipitation of the  $\alpha''$ -Fe<sub>1-x</sub>Si<sub>x</sub> phase were found to deteriorate when Nb was gradually replaced by Mo. Since Mo has a smaller atomic size than Nb, the Fe–Fe interatomic distances in the as-quenched amorphous samples became shorter when Nb was replaced by Mo. This explains why the exchange interaction among Fe atoms and therefore the Curie temperature of the amorphous phase, decreased after the substitution. Furthermore, the inhibition of diffusion of Fe and Si atoms was systematically weakened and thus,  $\alpha''$ -Fe<sub>1-x</sub>Si<sub>x</sub> crystallization began at lower temperatures for the same continuous heating runs. For 1 h isothermal annealing at 813 K, a linear increase of the mean grain size was observed with the addition of Mo (from 9.1 to 11.6 nm). The samples annealed at 813 K for 1 h were also studied in Ref. [11] by means of Mössbauer spectroscopy and quasistatic magnetic measurements. In that work, the fractions of various phases were estimated and it was observed that, when Nb was replaced by Mo, the crystalline fraction increased from  $55 \pm 2$  to  $66 \pm 1$  at%, in agreement with the thermal stability deterioration reported in Ref. [10].

The aim of this work is to present a study of the frequency dependence of the coercivity in as-quenched (amorphous) and isothermally annealed (nanocrystalline) samples of composition Fe<sub>73.5</sub>Si<sub>13.5</sub>Nb<sub>3-x</sub>Mo<sub>x</sub>B<sub>3</sub>Cu<sub>1</sub> ( $x=0, 1.5$  and  $3$ ), which will be called Mo0, Mo1.5 and Mo3, respectively. Some data on the magnetic properties of the amorphous magnetic alloys have already been reported with dc [12–13] or low frequency [14] magnetizing field. However, most of the commercial applications of these materials require higher frequencies. Therefore here we present coercivity measurements in the frequency range from 0.5 to 1.3 kHz, where the frequency dependence of coercivity is almost linear.

## 2. Experimental

Ingots of homogeneous compositions were prepared in an induction furnace. The amorphous samples were obtained in the form of ribbons  $\sim 20$   $\mu$ m thick and  $\sim 10$  mm wide by planar flow casting in air at the Institute of Physics of the Slovak Academy of Sciences (Bratislava, Slovakia) with the help of RNDr. Dušan Janičkovič. The amorphous structure of the samples was checked by XRD. The composition of the samples was analyzed by inductively coupled plasma spectroscopy to confirm that the intended compositions were achieved. Finally, an isothermal annealing at 813 K for 1 h in vacuum was performed to induce the precipitation of  $\alpha''$ -Fe<sub>1-x</sub>Si<sub>x</sub> nanocrystals.

Dynamic coercivity was measured from hysteresis loops (magnetic polarization,  $J=\mu_0 M$ , vs. applied magnetic field,  $H_a$ ) with a specially designed experimental setup based on the inductive method [15]. The magnetic field was applied along the length of the samples with a sinusoidal waveform frequencies within the range 0.5–1.3 kHz and amplitude  $\sim 4200$  A/m (large enough to magnetically saturate the sample).

## 3. Results and discussion

The magnetic hysteresis loops of the alloys Mo0, Mo1.5 and Mo3 were recorded at different frequencies of the applied magnetic field in their as-quenched state (with amorphous structure) and after controlled annealing (with nanocrystalline structure). As an example, the loops measured for Mo3 at 0.5 and 1.3 kHz are shown in Fig. 2. It was observed that the as-quenched Mo3 did not saturate completely even at the maximum applied magnetic field (Fig. 2 left), whereas annealed Mo3 reached its magnetic saturation at quite low fields at all the measured frequencies, revealing the magnetic softening achieved after the controlled heat treatment (Fig. 2 right).

$H_c$  was found to be higher at 1.3 than at 0.5 kHz both in as-quenched and annealed samples (Fig. 2). The frequency dependence of  $H_c$  exhibited an almost linear increase with frequency of the magnetizing field (Fig. 3).

All the as-quenched samples had approximately the same structural disorder and this was reflected in their similar magnetic behavior. The measured saturation magnetostriction was high ( $\lambda_s \sim 19$  ppm) [9] and their domain structure was heterogeneous (it exhibited wide curved in-plane domains as well as island-like fingerprint domains with the magnetization oriented out of the plane of the ribbon [13]). In the present work, no appreciable difference among the  $H_c$  vs. frequency curves of the as-quenched samples was found (Fig. 3). The continuous increase of coercivity with increasing frequency was due to the eddy currents. When a magnetic field  $H_a$  is applied along a body, a flux density field  $B$  is induced in the material. If  $H_a$  varies with time,  $B$  and therefore the flux ( $\phi$ ) vary as well through the cross-section of the body. Thus, according to Faraday's law, an electromotive force (emf) will be induced in the specimen proportional to  $d\phi/dt$  and, if the material is a conductor, this emf will cause an eddy current. The maximum of the applied field was kept constant in our measurements, so the flux rate increased only when the frequency of  $H_a$  was increased. During each cycle, as  $H_a$  increases, the eddy currents generate a magnetic field ( $H_{ec}$ ) antiparallel to  $H_a$ . The resulting field will be the sum of both fields,  $H_a$  and  $H_{ec}$ . It follows that the true field acting on the material the interior of the specimen, where the contributions of all the eddy currents are added, becomes weaker. Hence the higher the frequency, the larger the eddy currents and the  $H_{ec}$ ,

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