

⁵⁷Fe AND ⁶¹Ni MÖSSBAUER EFFECT INVESTIGATIONS OF NICKEL FERRITE-ALUMINATES

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In this paper we report on some new results of ⁵⁷Fe and ⁶¹Ni Mössbauer effect investigations of NiFe_{2-x}Al_xO₄ series [1]. Low temperature ⁵⁷Fe spectra of all investigated samples are interpreted as composed of tetrahedral (A) and octahedral (B) Zeeman patterns. The influence of the random distribution of Al³⁺, Ni²⁺ and Fe³⁺ ions in the A and B sites on the hyperfine fields corresponding to these sites, manifests itself in the broadening of Zeeman components. At elevated temperatures relaxation processes begin to play an important role. They appear at lower temperatures for highly diluted samples than for weakly diluted ones. The shapes of relaxation spectra indicate that the magnetic type of relaxation process, with only a small contribution of the superparamagnetic process, is involved. Some spectra of each of the investigated samples (Fig. 1) show relaxation features over temperature regions below appropriate Curie temperatures T_c. These regions are narrow for weakly diluted samples and relatively large for highly diluted ones (Fig. 2). In region A static Zeeman patterns are observed. Relaxation spectra are found in region B, while in region C quadrupole

doublets are registered. For each of the

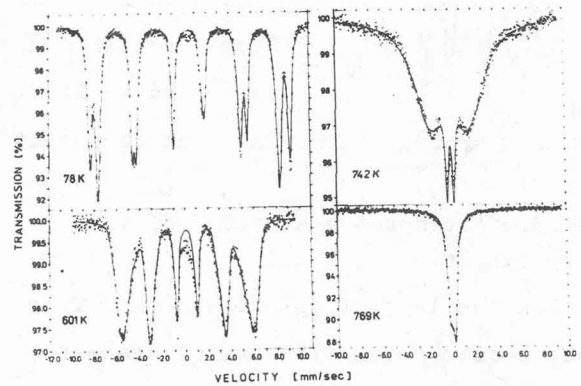


FIG. 1.—Selected ⁵⁷Fe spectra of the x=0.2 sample.

investigated samples a quadrupole doublet is observed below the magnetometric Curie temperature. This seems to be clear evidence of the influence of the external

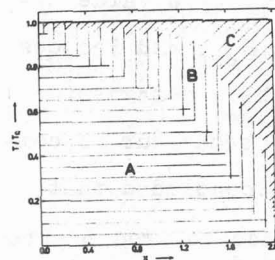


FIG. 2.—The diagram of the types of ⁵⁷Fe Mössbauer spectra.

relaxation processes. From the spectra of region A, the parameters characterizing the supertransferred hyperfine fields, while from those of region B, the relaxation times, are being studied.

⁵⁷Fe spectra for the x=1.5 sample were

taken at 4.2 K in 0, 4, 10, 30 and 60 kGs longitudinal external magnetic fields. Some of them are shown in Fig. 3. The spectrum in $H_{ext}=0$ is decomposed into three

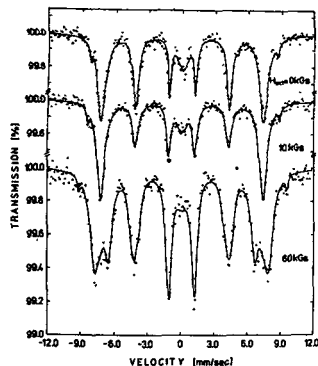


FIG. 3.—⁵⁷Fe spectra at 4.2 K of the x=1.5 sample.

components. From the linewidth of the most intense one it is concluded that both A and B Zeeman patterns are contained in it. The weak component II with the largest splitting comes from those B-site ferric ions whose nearest-neighbour coordination shell is more rich in Fe and Ni atoms than on the average. The non-resolved component III comes from Fe impurities in the beryllium sample holder. The external magnetic field splits component I into two parts, increases the splitting of component II by a value directly corresponding to H_{ext} and induces a Zeeman splitting of the non-magnetic component III. From the behaviour of component II in H_{ext} it is concluded that this component corresponds to iron B sites in which Fe magnetic moments are collinearly ordered and aligned antiparallel to the H_{ext} vector. The intensity of $\Delta m=0$ lines of the component I at first decreases with the increase in H_{ext} and then increases for still higher H_{ext} . This may be explained by the influence of H_{ext} on the ordering of the A and B sublattice magnetization vectors, similarly as is observed

for antiferromagnets [2], since the x=1.5 composition is very near the compensation point composition [1]. Similar phenomenon was observed earlier for another spinel [3].

⁶¹Ni Mössbauer spectra taken at 4.2 K in $H_{ext}=60$ kGs for samples up to x=1.25 (Fig. 3) are interpreted as being due to Ni ions occupying only the B sites, with a negative hyperfine field. For x=1.5 and 1.9

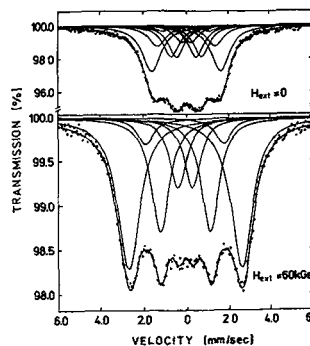


FIG. 4.—⁶¹Ni spectra at 4.2 K of the x=1.25 sample.

the signs of hyperfine fields at the A and B sites are both negative.

A more detailed analysis with the new results included will be published elsewhere.

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