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MAGNETOOPTICAL PROPERTIES OF ORTHOFERRITES AND GARNETS IN INFRARED

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Résumé. — Les propriétés magnéto-optiques de YFeO₃, EuFeO₃, DyFeO₃ et DyIG sont étudiées dans l'infra-rouge. Le terme non diagonal $|\epsilon_1|$ du tenseur $|\epsilon|$ et la biréfringence des orthoferrites sont assez grands. Dans la direction de l'axe optique λ la rotation Faraday de YFeO₃ est estimée à partir de $|\epsilon_1|$. La valeur calculée de cette rotation est 700 °/cm à $\lambda = 1.15 \mu$. Le résultat expérimental est en accord avec cette estimation. La rotation Faraday du DyIG est isotrope entre 80 et 293 °K et le facteur g du Dy³⁺ est égal à la valeur pour l'ion libre. On utilise l'équation du mouvement du moment magnétique avec un facteur g anisotrope pour interpréter la rotation Faraday à 25 °K.

Abstract. — Magnetooptical properties of the YFeO₃, EuFeO₃, DyFeO₃ and DyIG were investigated in infrared. The nondiagonal components $| e_i |$ of tensor $| e_i |$ are large enough. Faraday Rotation (F. R.) along optical axis of YFeO₃ was estimated from the value $| e_i |$. The calculated value of this rotation is equal to 700 °/cm at $\lambda = 1.15 \mu$. The experimental value is — 600 °/cm. The F. R. in DyIG is isotropic at 80-293 °K and g-factor Dy³⁺ equals its free ions value. The equation of motion of the magnetic moment with anisotropic g-factor is used for interpretation F. R. at 25 °K.

Orthoferrites, like ferrite garnets, have a transparency window in the infrared [1]. The transparency of the orthoferrites has been used for observing the domain structure [1, 2]. The polar Kerr effect of orthoferrites in the visible was investigated in [3, 4]. Using the transparency of the single crystals, we have investigated magnetooptical properties of orthoferrites YFeO₃, EuFeO₃ and DyFeO₃ [5]. The single crystals orthoferrites were grown by means of the technique of zone melting with optical heating [6]. Figure 1 gives ratio $2 \delta/I_0(\lambda)$ in YFeO₃ plate 102 μ thickness cut in the {001} plane. Here 2δ is the change in intensity of radiation I passing through the system when the current in the electromagnet is switched. The electrical field of the incident radiation coincides with the axis < 100 >. The analyser was rotated by 45° from < 100 >.

For all thicknesses of the specimen the quantities $2 \delta/I_0$ are the same for the same value of wavelength. In birefringent crystals the Faraday rotation can only occur along the optical axis [7]. In the directions which do not coincide with the optic axis the angle $\chi = \delta/2 I_0$ which defines the orientation of the major axis of the ellipse with respect to direction of polarisation of the incident light, can be written in the following form [8].

$$\operatorname{tg} 2 \chi = -2 k \sin \Delta = \frac{G_{33} \varepsilon_0^{5/2}}{\Delta n} \sin \frac{2 \pi \, \mathrm{d} \Delta n}{\lambda} \qquad (1)$$

where Δn -birefringence,

$$G_{33} = \frac{\varepsilon_{12}}{\varepsilon_0^2}, \quad \varepsilon_{12} = -i(\varepsilon_1 - i\varepsilon_2),$$

 ε_{12} is the nondiagonal component of the dielectric constant tensor, ε_0 is the diagonal component, *d* is the thickness of the plate, λ is the wavelength. From the « period » of the changing χ we get

$$\begin{array}{ll} \Delta n = (3.6 \pm 0.3) \, 10^{-2} & \text{at} & \lambda = 1.15 \, \mu \,, \\ \Delta n = (3.5 \pm 0.3) \, 10^{-2} & \text{at} & \lambda = 1.8 \, \mu \,, \\ \Delta n = (2.9 \pm 0.3) \, 10^{-2} & \text{at} & \lambda = 6 \, \mu \,. \end{array}$$

Birefringence of YFeO₃ is large comparable with birefringence of the $CaCO_3$.

The inset figure 1 shows $|\varepsilon_1|$ of YFeO₃, obtained with the help of (1). The values of $|\varepsilon_1|$ are large

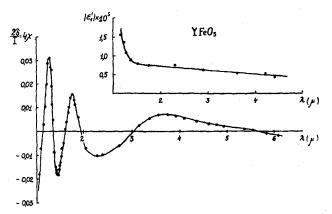


FIG. 1. — Relative change in the intensity of the radiation $2 \delta/I$ passing through the YFeO₃ of thickness 102 μ when the current in the electromagnet is switched as function of the wavelength.

enough, as in visible [3, 4]. The orthoferrites crystallize in an orthorombically distorted perovskite structure belonging to the two-axis crystals. The optical axis of YFeO₃ lies in the { 100 } plane and makes with the axis < 001 > the angle of 40°. That is very convenient to receive large F. R. The value of F. R. may be estimated from $|\varepsilon_1|$ and the angle between the optical axis and < 001 >. For example, F. R. is equal to 700 °/cm at $\lambda = 1.15 \mu$. The experimental value is - 600 °/cm. This F. R. is higher than in YIG. In direction that doesn't coincide with the optical axis, the value χ is small; $\chi = 0.4^{\circ}$ and 1.3° at $\lambda = 1.15 \mu$ and 0.63 μ respectively. Curves similar to those of figure 1 were obtained for other orthoferrites. Birefringence of the orthoferrites is proportional to its orthorombical distortions. For example, in EuFeO₃ at $\lambda = 1.15 \ \mu \ \Delta n = (2.2 \pm 0.2) \ 10^{-2}$ and $\chi = 0.6^{\circ}$. The curves $\chi(\lambda)$ in the vicinity of the absorption bands have some peculiarities [9].

In papers [10] was found the frequency independent Faraday effect α_{F_0} which exists due to the magnetic permeability in the optical frequence. It was shown that the F. R. in DyIG at 80-293 °K is isotropic [11] and the values $\alpha_{\rm F}(\lambda)$ may be written in the form [10-12]:

$$\alpha_{\rm F} = \frac{\pi \sqrt{\varepsilon} e}{mc^2} (g_{\rm Fe} I_{\rm Fe} - g_{\rm R} I_{\rm R}) + \frac{k}{\lambda^2}. \qquad (2)$$

Here the first member is α_{F_0} ; g_{F_e} , g_R , I_{F_e} , I_R are the g-factor and the magnetizations of Fe³⁺ and Dy³⁺, $\sqrt{\varepsilon} = n$. The g-factor of Dy³⁺ ion obtained from the value α_{F_0} equals its free ion value. The result of the investigation of the F.R. of oriented DyIG single crystals at 25 °K in magnetic field H = 12 kOe is represented in figure 2. From these results and (2) we obtains :

$$\begin{aligned} \alpha_{F_0}^{<111>} &= 180 \pm 10^{\text{ o}/\text{cm}}; \\ \alpha_{F_0}^{<110>} &= 120 \pm 10^{\text{ o}/\text{cm}}; \\ \alpha_{F_0}^{<100>} &= 140 \pm 10^{\text{ o}/\text{cm}}. \end{aligned}$$

Along the < 110 > direction was found the birefringence [13, 14] which varies from $\Delta n = 1.1 \times 10^{-3}$ at $\lambda = 3.5 \mu$ up to $\Delta n = 0.95 \times 10^{-3}$ at $\lambda = 6.6 \mu$. In this case the value $\alpha_{F_0} < 110 >$ was calculated using (1). From the values α_{F_0} and (2), using the magnetization data [15], may be determined $g_R < 111 > = 1.55 \pm 0.1$, $g_R < 100 > = 1.8 \pm 0.1$, $g_R < 110 > = 1.4 \pm 0.1$. To obtain these values α_{F_0} the values $\cos \alpha_i = 1, \sqrt{2/3}$, $\sqrt{1/3}$ were introduced in (2) for the < 111 >, < 110 >and < 100 > directions respectively. It is not excluded that anisotropic α_{F_i} is due to the difference of values of $\cos \alpha_i$ from those mentioned above. In this case it must vanish in the high magnetic fields.

The interpretation α_{F_0} at 80-293 °K mentioned above is based on using the Landau-Lifshitz equations with isotropic g-factor. At low temperature one must take into account that local symmetry of rare-earth

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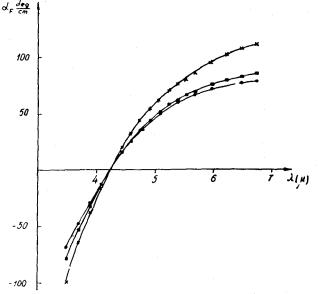


FIG. 2. — The Faraday Rotation $\alpha_{\rm F}(\lambda)$ in oriented single crystals DyIG at 25 °K in magnetic field H = 12 kOe. ($\bullet - \alpha_F < 110 >$; $\blacksquare - \alpha_{\rm F} < 100 > ; \times - \alpha_{\rm F} < 111 >).$

ions is not cubic. In this case we must employ the equations of motion of magnetic moment with anisotropic g-factor. Using the Vlasov's and Yshmuhametov's equations [16] and taking account of the six non-equivalent positions of the rare-earth ions in the unit cell, one obtains

$$g_{\rm R} = \frac{g_{11}^2 g_{22}^2 + g_{22}^2 g_{33}^2 + g_{11}^2 g_{33}^2}{3 g_{11} g_{22} g_{33}}$$
(3)

along over all directions of the crystal. In (3) the g_{11}, g_{22}, g_{33} are the principal values of g-tensor. When $g_{11} = g_{22} = g_{33} = g_J$, then $g_R = g_J$. If $g_{11} \neq g_{22} \neq g_{33}$, $g_R \neq g_J$ and this is observed on the experiment for < 111 >direction.

Allowance for non-colinear sublattices [17] within the Wolf's model [18] doesn't lead to anisotropic α_{F_0} under condition of the equal magnetic moments of the rare-earth ions in unequivalent positions.

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