# Symmetry analysis of the photorefractive electrogyration and electrogyration study in the LiNbO<sub>3</sub> crystals

R. Vlokh<sup>1,2</sup>, M. Kostyrko<sup>1</sup>, V. Kruchkevych<sup>1</sup>

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#### **Abstract**

The article is devoted to the phenomenological investigation of the possibility of the optical activity appearing under the photorefractive effect in the non-gyrotropic crystals. On the base of the symmetry analysis it is shown that in the crystals that belong to the planar and axially-inversion point groups of the symmetry ( $\bar{4}3m$ ,  $\bar{6}m2$ ,  $\bar{6}$ , 3m, 6mm, 4mm) optical activity can be induced by the electrogyration effect caused by the photorefractivity. In the crystals that belong to the point group of symmetry 6mm, 4mm, 3m,  $\bar{6}$ ,  $\bar{6}m2$  optical activity can be induced by photovoltaic and linear electrogyration effects while in the crystals that belong to the  $\bar{4}3m$  group of the symmetry - by photovoltaic and quadratic electrogyration effects. The electrogyration effect in the LiNbO<sub>3</sub> crystal was experimentally investigated. The component of electrogyration tensor was determined as  $\gamma_{41}$ =(3.39±1.3)×10<sup>-12</sup>m/V and electrogyration diffraction effectivity was calculated:  $\eta$ =1.5%. It was shown that the diffraction of light on the electrogyration photorefractive grating should be only anisotropical.

**Key words:** electrogyration, photorefraction effect, optical activity.

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## Introduction

The photorefractive effect can appear in the crystals as a result of the photovoltaic and electrooptical effects. The space charges electric field induced by electric field of the optical radiation through the photovoltaic effect caused the change of refractive indexes by the Pockels effect. But in the general case space charges electric field ( $E^{sc}_{m}$ ,  $E^{sc}_{n}$ ) that appear in the result of the charges distribution can change not only the real part of the dielectric permitivity (refractive indexes) but also the imaginary part (gyration) [1]:

$$\varepsilon_{ii} = \varepsilon_{ii}^{o} + i e_{ii} g_{lk} k_{k}, \tag{1}$$

$$\Delta \varepsilon_{ii}^{o} = r_{iim} E^{sc}_{m} + R_{iimn} E^{sc}_{m} E^{sc}_{n}, \qquad (2)$$

$$\Delta g_{lk} = \gamma_{lkm} E^{sc}_{m} + \delta_{lkmn} E^{sc}_{m} E^{sc}_{n}, \qquad (3)$$

where  $r_{ijm}$ ,  $R_{ijmn}$  - coefficients of the Pockels and

Kerr effect and  $\gamma_{lkm}$ ,  $\delta_{lkmn}$  - third and fourth rank axial tensors of the linear and quadratic electrogyration effect, respectively. It is known that due to the symmetry of the gyration tensor (second rank axial tensor  $g_{lk}$ ) optical activity can exist only in the non-centrosymmetrical crystals as well as photovoltaic effect which is described by the polar third rank tensor  $(j_n = \beta_{nrt} E_n E_t$ , where  $j_n$  is the vector of the bulk photovoltaic current that is parallel to the vector of the space charge electric field  $E^{sc}$ ). So, one can suppose that photorefractive electrogyration effect can appear only in the crystals that already possess optical activity. The present paper is devoted to analyzing the conditions in which optical activity can appear in the nongyrotropic crystalsunder the photorefractivity and experimental studying of the electrogyration in the LiNbO<sub>3</sub> crystals.

<sup>&</sup>lt;sup>1</sup>Institute of Physical Optics, 23 Dragomanov Str., 79005, L'viv, Ukraine <sup>2</sup>National University "L'viv Polytechnic", 12 S.Bandera Str., L'viv, Ukraine

## Symmetry analysis

Linear electrogyration can exist in the non–centrosymmetrical as well as in the centrosymmetrical medium, except crystals which belong to the m3m and  $\overline{4}$ 3m point group of the symmetry. The effect of the change of the optical activity under the photorefractive effect in the crystals that already possess gyration have been already studed yet [2-4]. From other side in the crystals that belong to the planar and axially-inversion point groups of the symmetry ( $\overline{4}$ 3m,  $\overline{6}$ m2,  $\overline{6}$ , 3m, 6mm, 4mm) optical activity could be induced by the electrogyration effect caused by the photorefractivity and in these crystals photovoltaic effect could also exist. The

appearance of the optical activity in these crystals is connected with the local lowering of the symmetry under the space charge electric field induced by the photovoltaic effect. In the cubic crystals with the symmetry  $\overline{4}3m$  optical activity can be induced only by the quadratic electrogyration. Let us consider the possibility of the appearing of the photorefractive electrogyration effect in these crystals in the case when optical radiation is propagated along the principle cystallophysical axis. The results of this analyzing are collected in the Table1. It is necessary to note, that the well known photorefractive LiNbO<sub>3</sub> crystals belong to the point group of the symmetry 3m.

Table 1

Symmetr	Components of	Change of the	Components of the	Equations for the gyration tensor
y of the	the wave vector	symmetry under	photovoltaic current	
crystals	and the light	the space charges		
	polarization	electric field		
1	2	3	4	5
6mm,	$k_1$ , $E_2E_2$	non change	$j_3 = \beta_{13} E_2 E_2$	$g_{ij}=0$
4mm	$k_1$ , $E_3E_3$	non change	$j_3=\beta_{33}E_3E_3$	$g_{ij}=0$
	$k_1$ , $E_2E_3$	$6\text{mm} \rightarrow \text{m}$	$j_3 = \beta_{13}E_2E_2 + \beta_{33}E_3E_3$	$g_{3I}$ =- $\gamma_{4I}E^{sc}_{2}$
		$4\text{mm} \rightarrow \text{m}$	$j_2 = \beta_{42} E_2 E_3$	$g_{12} = \delta_{16} E^{sc2}_{2}$
	$k_2$ , $E_1E_1$	non change	$j_3=\beta_{13}E_1E_1$	$g_{ij}$ =0
	$k_2$ , $E_3E_3$	non change	$j_3=\beta_{33}E_3E_3$	$g_{ij}$ =0
	$k_2$ , $E_3E_1$	$6\text{mm} \rightarrow \text{m}$	$j_1 = \beta_{42} E_3 E_1$	$g_{32} = \gamma_{41} E^{sc}_{l}$
		$4\text{mm} \rightarrow \text{m}$	$j_3 = \beta_{13} E_1 E_1 + \beta_{33} E_3 E_3$	$g_{12}$ =- $\delta_{l6}E^{sc2}_{l}$
	$k_3$ , $E_1E_1$	non change	$j_3=\beta_{I3}E_IE_I$	$g_{ij}$ =0
	$k_3$ , $E_2E_2$	non change	$j_3 = \beta_{13} E_2 E_2$	$g_{ij}$ =0
	$k_3$ , $E_1E_2$	non change	$j_3 = \beta_{13} E_1 E_1 + \beta_{13} E_2 E_2$	$g_{ij}$ =0
3m	$k_1$ , $E_2E_2$	$3m \rightarrow m$	$j_2 = \beta_{12} E_2 E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
			$j_3 = \beta_{13} E_2 E_2$	$g_{12} = -\gamma_{11} E^{sc}_{2} + \delta_{61} E^{sc2}_{2}$
	$k_1$ , $E_3E_3$	non change	$j_3=\beta_{33}E_3E_3$	$g_{ij}$ =0
	$k_1$ , $E_2E_3$	$3m \rightarrow m$	$j_2 = \beta_{42} E_2 E_3 + \beta_{12} E_2 E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
			$j_3 = \beta_{33}E_3E_3 + \beta_{13}E_2E_2$	$g_{12} = -\gamma_{11} E^{sc}_{2} + \delta_{61} E^{sc2}_{2}$
	$k_2$ , $E_1E_1$	$3m \rightarrow m$	$j_2 = -\beta_{12} E_1 E_1$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
			$j_3=\beta_{13}E_1E_1$	$g_{12} = -\gamma_{11} E^{sc}_{2} + \delta_{61} E^{sc2}_{2}$
	$k_2$ , $E_3E_3$	non change	$j_3 = \beta_{33} E_3 E_3$	$g_{ij}$ =0

1	2	3	4	5
3m	$k_2$ , $E_3E_1$	$3m \rightarrow 1$	$j_1 = \beta_{42} E_1 E_3$	$g_{11}=\gamma_{11}E^{sc}_{1}$
			$j_2 = -\beta_{12} E_1 E_1$	$g_{22} = -\gamma_{II} E^{sc}{}_{I} = -g_{II}$
			$j_3 = \beta_{13} E_1 E_1 + \beta_{33} E_3 E_3$	$g_{23} = \gamma_{41} E^{sc}{}_{1}$
				$g_{13} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2} + \delta_{46} E^{sc2}_{1}$
				$g_{12} = -\gamma_{11} E^{sc}_{2} - \delta_{61} E^{sc2}_{1} + \delta_{61} E^{sc2}_{2}$
	$k_3$ , $E_1E_1$	$3m \rightarrow m$	$j_2 = -\beta_{12} E_1 E_1$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
			$j_3=\beta_{13}E_1E_1$	$g_{12} = -\gamma_{11} E^{sc}_{2} + \delta_{61} E^{sc2}_{2}$
	$k_3$ , $E_2E_2$	$3m \rightarrow m$	$j_2=\beta_{12}E_2E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
			$j_3 = \beta_{13} E_2 E_2$	$g_{12} = -\gamma_{11} E^{sc}_{2} + \delta_{61} E^{sc2}_{2}$
	$k_3$ , $E_1E_2$	$3m \rightarrow 1$	$j_1=\beta_{12}E_1E_2$	$g_{11} = \gamma_{11} E^{sc}_{1}$
			$j_2 = -\beta_{12}E_1E_1 + \beta_{12}E_2E_2$	$g_{22} = -\gamma_{I1} E^{sc}{}_{I} = -g_{II}$
			$j_3 = \beta_{13}E_1E_1 + \beta_{13}E_2E_2$	$g_{23} = \gamma_{41} E^{sc}_{1}$
				$g_{13} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2} + \delta_{46} E^{sc2}_{1}$
_		_		$g_{12} = -\gamma_{11} E^{sc}_{2} - \delta_{61} E^{sc2}_{1} + \delta_{61} E^{sc2}_{2}$
$\overline{6}$	$k_1$ , $E_2E_2$	$6 \rightarrow m$	$j_1 = -\beta_{11} E_2 E_2$	$g_{23} = \gamma_{41}E^{sc}{}_{1} + \gamma_{42}E^{sc}{}_{2} + \delta_{41}E^{sc2}{}_{1}$
			$j_2 = \beta_{12} E_2 E_2$	$\delta_{42}E^{sc2}_{2}$
				$g_{13} = \gamma_{42} E^{sc}_{1}$
				$\gamma_{41}E^{sc}_{2} + \delta_{46}E^{sc2}_{1} + \delta_{46}E^{sc2}_{2}$
	$k_1, E_3E_3$	non change	$j_i=0$	$g_{ij}=0$
	$k_1, E_2E_3$	$6 \rightarrow m$	$j_1 = -\beta_{11}E_2E_2$	$g_{23} = \gamma_{41} E^{sc}_{l} + \gamma_{42} E^{sc}_{2} + \delta_{41} E^{sc2}_{l}$
			$j_2 = \beta_{12} E_2 E_2$	$\delta_{42}E^{sc2}_{2}$
				$g_{13} = \gamma_{42} E^{sc}_{1}$
		<del>-</del>		$\gamma_{41}E^{sc}_{2} + \delta_{46}E^{sc2}_{1} + \delta_{46}E^{sc2}_{2}$
	$k_2$ , $E_1E_1$	$6 \rightarrow m$	$j_1=\beta_{11}E_1E_1$	$g_{23} = \gamma_{41} E^{sc}_{1} + \gamma_{42} E^{sc}_{2} + \delta_{41} E^{sc2}_{1}$
			$j_2 = -\beta_{12} E_1 E_1$	$\delta_{42}E^{sc2}_{2}$
				$g_{13} = \gamma_{42} E^{sc}_{1}$
				$\gamma_{41}E^{sc}_{2} + \delta_{46}E^{sc2}_{1} + \delta_{46}E^{sc2}_{2}$
	$k_2$ , $E_3E_3$	non change	$j_i=0$	$g_{ij}=0$
	$k_2$ , $E_1E_3$	$6 \rightarrow m$	$j_1 = \beta_{11} E_1 E_1$	$g_{23} = \gamma_{41} E^{sc}_{1} + \gamma_{42} E^{sc}_{2} + \delta_{41} E^{sc2}_{1}$
			$j_2 = -\beta_{12} E_1 E_1$	$\delta_{42}E^{sc2}_{2}$
				$g_{13} = \gamma_{42} E^{sc}_{1}$
	1			$\gamma_{41}E^{sc}_{2} + \delta_{46}E^{sc2}_{1} + \delta_{46}E^{sc2}_{2}$
	$k_3$ , $E_1E_1$	$6 \rightarrow m$	$j_1 = \beta_{11} E_1 E_1$	$g_{23} = \gamma_{41}E^{sc}_{1} + \gamma_{42}E^{sc}_{2} + \delta_{41}E^{sc2}_{1}$
			$j_2 = -\beta_{12} E_1 E_1$	$\delta_{42}E^{sc2}_{2}$
				$g_{13} = \gamma_{42} E^{sc}_{1}$
	1 F F		. 0.55	$\gamma_{41}E^{sc}_{2} + \delta_{46}E^{sc2}_{1} + \delta_{46}E^{sc2}_{2}$
	$k_3$ , $E_2E_2$	$6 \rightarrow m$	$j_1 = -\beta_{11} E_2 E_2$	$g_{23} = \gamma_{41} E^{sc}_{1} + \gamma_{42} E^{sc}_{2} + \delta_{41} E^{sc2}_{1}$
			$j_2 = \beta_{12} E_2 E_2$	$\delta_{42}E^{sc2}_{2}$
				$g_{13} = \gamma_{42} E^{sc}_{1}$
				$\gamma_{41}E^{sc}_{2} + \delta_{46}E^{sc2}_{1} + \delta_{46}E^{sc2}_{2}$

1	2	3	4	5
<u></u>	$k_3$ , $E_1E_2$	$\overline{6} \rightarrow m$	$j_1 = \beta_{11} E_1 E_1 - \beta_{11} E_2 E_2$	$g_{23} = \gamma_{41} E^{sc}_{1} + \gamma_{42} E^{sc}_{2} + \delta_{41} E^{sc2}_{1}$
			$-\beta_{12}E_1E_2$	$\delta_{42}E^{sc2}{}_2$
			$j_2 = -\beta_{12}E_1E_2 + \beta_{12}E_2E_2 -$	$g_{13} = \gamma_{42} E^{sc}_{l}$ -
			$-\beta_{11}E_1E_2$	$\gamma_{41}E^{sc}{}_2 + \delta_{46}E^{sc2}{}_1 + \delta_{46}E^{sc2}{}_2$
<del>6</del> m2	$k_1$ , $E_2E_2$	$\overline{6}$ m2 $\rightarrow$ mm2	$j_2 = \beta_{12} E_2 E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
	$k_1$ , $E_3E_3$	non change	$j_i=0$	$g_{ij}$ =0
	$k_1$ , $E_2E_3$	$\overline{6}$ m2 $\rightarrow$ mm2	$j_2 = \beta_{12} E_2 E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
	$k_2$ , $E_1E_1$	$\overline{6}$ m2 $\rightarrow$ mm2	$j_2 = -\beta_{12} E_1 E_1$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
	$k_2$ , $E_3E_3$	non change	$j_i=0$	$g_{ij}$ = $0$
	$k_2$ , $E_1E_3$	$\overline{6}$ m2 $\rightarrow$ mm2	$j_2 = -\beta_{12} E_1 E_1$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
	$k_3$ , $E_1E_1$	$\overline{6}$ m2 $\rightarrow$ m	$j_2 = -\beta_{12} E_1 E_1$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
	$k_3$ , $E_2E_2$	$\overline{6}$ m2 $\rightarrow$ m	$j_2 = \beta_{12} E_2 E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} - \delta_{46} E^{sc2}_{2}$
	$k_3$ , $E_1E_2$	$\overline{6}$ m2 $\rightarrow$ m	$j_1 = -\beta_{12} E_1 E_2$	$g_{32} = \gamma_{41} E^{sc}_{1}$
			$j_2 = -\beta_{12}E_1E_1 + \beta_{12}E_2E_2$	$g_{31} = -\gamma_{41} E^{sc}_{2} + \delta_{46} E^{sc2}_{1} - \delta_{46} E^{sc2}_{2}$
<del>4</del> 3m	$k_1$ , $E_2E_2$	non change	$j_i=0$	$g_{ij}$ =0
	$k_1$ , $E_3E_3$	non change	$j_i=0$	$g_{ij}$ = $0$
	$k_1$ , $E_2E_3$	$\overline{4}3\text{m} \rightarrow \overline{4}2\text{m}$	$j_1 = \beta_{41} E_2 E_3$	$g_{22} = -\delta_{12} E^{sc2}{}_{1}$
				$g_{33} = \delta_{12} E^{sc2}{}_{1}$
	$k_2$ , $E_1E_1$	non change	$j_i=0$	$g_{ij}$ =0
	$k_2$ , $E_3E_3$	non change	$j_i=0$	$g_{ij}$ =0
	$k_2$ , $E_1E_3$	$\overline{4}3\text{m} \rightarrow \overline{4}2\text{m}$	$j_2 = \beta_{41} E_1 E_3$	$g_{11} = \delta_{12} E^{sc2}_{2}$
				$g_{33}$ =- $\delta_{l2}E^{sc2}_{2}$
	$k_3$ , $E_1E_1$	non change	$j_i=0$	$g_{ij}$ =0
	$k_3$ , $E_2E_2$	non change	$j_i=0$	$g_{ij}$ =0
	$k_3$ , $E_1E_2$	$\overline{4}3\text{m} \rightarrow \overline{4}2\text{m}$	$j_3 = \beta_{41} E_1 E_2$	$g_{11} = -\delta_{12} E^{sc2}_{3}$
				$g_{22} = \delta_{12} E^{sc2}_{3}$

Tensors  $\beta_{ijk}$   $\delta_{ijkl}$  and  $\gamma_{ijk}$  sre presented in matrix form.

## **Experimental**

For the calculation of the value of the photorefractive induced optical activity it is necessary to know the electrogyration coefficients in the chosen crystals. The one of the photorefractive crystals is well investigated lithium-niobate that belongs to the planar group of the symmetry 3m and does not possess optical activity. Unfortunately the electrogyration effect in these crystals is not studied yet. The reason of this fact, perhaps is connected with the difficulties of such experiment. Electrogyration should not appear in the crystals with the

symmetry 3m if the electric field is applied along three-fold axis and optical radiation is propagated along optical axis. If the electric field is applied in the direction perpendicular to the three-fold axis optical birefringence will always appear in the Z - direction. It means that electrogyration effect should be investigated under conditions of the presence of the linear birefringence.

Let us consider the peculiarities of the electrooptical effect in the LiNbO<sub>3</sub> crystals in the case if the electric field is applied in the X and Y - directions. According to the Curie prin-

ciple the application of the electric field along Y axis should cause the lowering of the symmetry  $3m\rightarrow m$  and along X- axis -  $3m\rightarrow 1$ . The both groups of the symmetry (m and 1) permit existing of the optical activity and crystals that belong to this groups are optically biaxial. Under the electric field  $E_2$  the LiNbO<sub>3</sub> crystals become optically biaxial and the optical axes will belong to the ZY or ZX planes, dependently of the sign of the field (Fig.1,a). At the same time gyration tensor should possess two nonzero components  $g_{13}$  and  $g_{23}$  (Fig.1,a). It means that optical activity can appear in the direction of the one of optical axes only in the case if optical axes plane will be perpendicular to the YZ-plane. The orientation of the optical axes plane depends on the sign of the  $E_2$  field. Then if optical beam will propagate along one of the optical axis which belong to the XZ-plane one could observe the rotation of the polarization plane caused by the electrogyration effect. At the application of the  $E_1$  electric field the crystal should become optically biaxial,

indicatrix will turn around Z axis on the  $45^{\circ}$  and all components of the gyration tensor should be non-zero (Fig.1,b). In both cases it is possible to investigate the electrogyration effect under conditions when optical radiation will propagate along one of the optical axes by measuring of the rotation the polarization plane.

In our experiments lithium-niobate crystal was prepared with the faces perpendicular to the principle crystallophysical axis with dimensions  $10 \times 10 \times 10 \text{mm}^3$  and the face that is perpendicular to the Z-axis was polished by the diamond paste. Electrical voltage was applied along X and Y axes up to the 13kV. Sample with the electrical field application was placed on the optical micropozicioner with the angle scale, which could be turned around direction of the optical beam propagation ( $\lambda$ =632.8nm). The measurements of the electrogyration effect were held with the help of the Faraday cell at the room temperature ( $T=20^{\circ}$ C). The polarization of the incident beam was parallel or perpendicular to the incident plane.

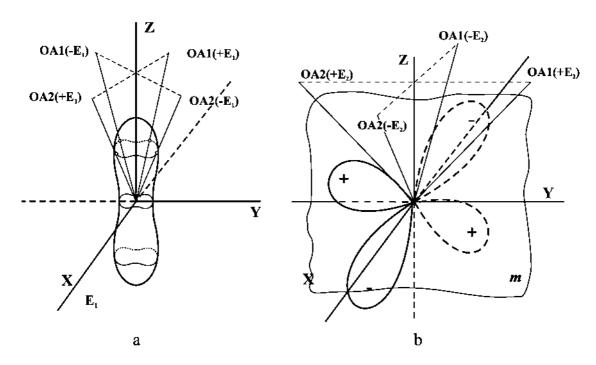


Fig.1.Gyration surfaces and optical axes in the LiNbO<sub>3</sub> crystals under the application of the electrical field a) $E_2$  and b) $E_1$ .

## Results and discussion

Let us consider the equation for the scalar gyration parameter at the application of the electric field along X and Y axis of the LiNbO<sub>3</sub> crystals. At the  $E_2$  field the relation for the gyration can be written as:

$$G = g_{ij} l_i l_j = \rho \lambda n / \pi = 2g_{12} l_1 l_2 +$$

$$+ 2g_{13} l_1 l_3 = -2 \gamma_{11} E_2 l_1 l_2 - 2 \gamma_{41} E_2 l_1 l_3,$$
(4)

where G - is scalar gyration parameter,  $g_{ij}$  - are components of the gyration tensor , n - is the refractive index in the direction of the optical axis,  $\rho$  - is the polarization plane rotary power,  $\lambda$  - is the wavelength of the optical radiation,  $\gamma_{ij}$  - are the components of the electrogyration tensor and  $l_i l_j$  - are direction cosines of the wave vector. It is suitable to rewrite the equation (4) in the spherical coordinate:

$$G=g_{ij} l_i l_j = \rho \lambda n / \pi = -\gamma_{41} E_2 \sin 2\theta$$
 (5) and

$$\gamma_{41} = -\rho \lambda n / \pi E_2 \sin 2\theta, \tag{6}$$

where  $\theta$  - is the angle between the optical axis in the optically biaxial LiNbO<sub>3</sub> crystals and Z - axis. At the  $E_I$  field the relation for the gyration can be written as:

$$\begin{aligned} G = & g_{ij} \ l_i l_j = \rho \lambda n / \pi = g_{11} l_1 l_1 + g_{22} l_2 l_2 + 2 g_{12} l_1 l_2 = \\ = & \gamma_{11} E_1 l_1 l_1 - \gamma_{11} E_1 l_2 l_2 + 2 \gamma_{41} E_1 l_3 l_2 \end{aligned} \tag{7}$$

In the spherical coordinate (7) can be rewritten as

$$G{=}g_{ij}\;l_il_j{=}\rho\lambda n/\pi{=}(2)^{1/2}\gamma_{41}E_1sin2\theta$$
 and   
 (8)

$$\gamma_{41} = \rho \lambda n/(2)^{1/2} \pi \gamma_{41} E_1 \sin 2\theta$$
.

In the Fig.2 the dependence of the optical rotary power on the strength of electrical field  $E_2$  is shown. According to the formula (6) the calculated value is  $\gamma_{41}$ =(3.39±1.3)×10<sup>-12</sup>m/V. In the case when electrical field  $E_2$  leads to the splitting of optical axis in YZ plane the deviation of the rotation of polarization plane was in order of the experimental error. Taking into account the largest value of the space charge electrical field ( $E_{sc}$ ~10<sup>7</sup>V/m) induced by optical radiation through photorefractivity [5] one can estimate

the amplitude of the photorefractive periodical distribution of the gyration tensor will be  $\Delta g_{ij}=3.9\times10^{-5}$ . In this case the diffraction effectivity can be written as [6]

$$\eta = \sin^2 \left( \frac{\pi \Delta nd}{\lambda \cos \theta} \right) \tag{9}$$

where d - is the interaction length,  $\theta$  - is the incident angle of regording beam. In the case of the electrooptical and electrogyration grating recording the amplitude of the birefringence can be written as

$$\Delta n = (\Delta n_1^2 + \Delta n_c^2)^{1/2},$$
 (10)

where  $\Delta n$  - is the total birefringence,  $\Delta n_l$  - is the linear birefringence induced by the electrooptical effect and  $\Delta n_c = \Delta g_{ij} / n$  - is the circular birefringence induced by the electrogyration ( $n = (n_e + n_o)/2 = 2.2864 + 2.2025 = 2.2445$  - the average value of the refractive indexes (according to [7] for the wavelength 632.8nm)). So, photo-refractive  $\Delta n_c = 1.76 \times 10^{-5}$ .

One can note that the electrogyration kind of diffraction can be only anisotropic because

$$E^{\omega}_{i} = (\chi_{ii} + ie_{iil} g_{lk}k_k)D^{\omega}_{i}, \qquad (11)$$

(where  $P^{\omega}_{i}$  is the polarization of diffracted wave and  $E^{\omega}_{j}$  - electrical field of incident wave) and in Levi-Civit tensor -  $e_{ijl}$  all indexes should be different. It means that polarization of incident and diffracted light should be mutually perpendicular and electrogyration diffraction could be simply separated from electrooptical one. Though it is reasonable to consider only the diffraction efficiency  $\eta$  of the diffraction on the electrogyration photorefractive grating.

$$\eta = \sin^2 \left( \frac{\pi \Delta n_c d}{\lambda \cos \theta} \right) \tag{12}$$

For the  $\lambda$ =632.8nm, d=1mm and  $\theta$ ≈45° [8]  $\eta$ ≈1.5×10<sup>-2</sup> or 1.5%. Comparing with diffraction effectivity obtained on the electrooptical grating for example in [8] the electrogyration diffraction effectivity it is of order smaller than electrooptical.

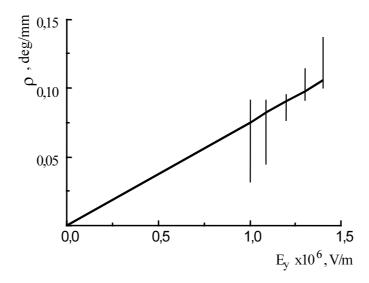


Fig.2.The dependence of the optical rotary power on the strength of electrical field  $E_2$  in the LiNbO<sub>3</sub> crystals ( $\lambda$ =632.8nm).

## **Conclusions**

1. The phenomenological investigation of the possibility of the optical activity appearing under the photorefraction effect in the nongyrotropic crystals is made. On the base of the symmetry analysis it is shown that in the crystals that belong to the planar and axiallyinversion point groups of the symmetry (43m, 6m2, 6, 3m, 6mm, 4mm) the optical activity could be induced by the electrogyration effect caused by the photorefractivity. In the crystals that belong to the point group of symmetry 6mm, 4mm, 3m,  $\overline{6}$ ,  $\overline{6}$ m2 the optical activity could be induced by photovoltaic and linear electrogyration effects while in the crystals that belong to the  $\overline{43}$ m group of the symmetry - by and quadratic electrogyration photovoltaic effects.

2.The electrogyration effect in the LiNbO<sub>3</sub> crystal was experimentally investigated. The component of electrogyration tensor was determined as  $\gamma_{41}$ =(3.39±1.3)×10<sup>-12</sup>m/V and diffractive effectivity at the diffraction on the photorefractive electrogyration grating is estimated as  $\eta$ =1.5%. It was shown that diffraction of light on the electrogyration photorefractive grating should be only anisotropical.

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