Effect of strong magnetic field on surface polaritons in ZnO

E.F. Venger¹, A.I. Ievtushenko², L.Yu. Melnichuk², O.V. Melnichuk²
¹V. Lashkaryov Institute of Semiconductor Physics, NAS of Ukraine
41, prospect Nauky, 03028 Kyiv, Ukraine
²Mykola Gogol Nezhyn State University
2, Kropyv'yansko str., 16600 Nezhyn, Ukraine; e-mail: mov310@mail.ru

Abstract. Using the attenuated total reflectance technique, we studied the effect of strong uniform magnetic field \( H \) on the main properties of surface polaritons in ZnO single crystals. The used orientations were \( C || y, \ k \perp C, \ xy || C, \ H \perp \k, \ H || y, \ k_x = k, k_{y,z} = 0 \); the free charge carrier concentration varied from \( 9.3 \times 10^{16} \) up to \( 2.0 \times 10^{18} \) cm\(^{-3} \). It has been shown that three dispersion curves exist in the zinc oxide single crystals for the above orientation. The possibility of excitation of an additional dispersion branch in optically anisotropic semiconductors placed into magnetic field is found using ZnO single crystals as an example. The damping coefficients for surface phonon and plasmon-phonon polaritons have been determined.

Keywords: ZnO single crystal, strong magnetic field, surface polariton, damping coefficient.

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1. Introduction

Zinc oxide belongs to II–VI semiconductor compounds and crystallizes in the wurtzite structure. Its unique physico-chemical properties are required for application in metallurgy, space engineering, acousto-, micro- and optoelectronics as well as other areas of science, engineering and medicine [1]. In particular, continuous growth of application of ZnO single crystals in electronic-optical engineering takes place. This promotes both theoretical and experimental investigations of electrical, magnetic and optical properties of zinc oxide.

In [2], the spectra of external reflection of light from surface of ZnO (with the electron concentration from \( 10^{16} \) up to \( 5 \times 10^{19} \) cm\(^{-3} \)) were studied experimentally and interconsistent parameters of ZnO single crystal bulk were obtained. It was shown that ZnO single crystals are characterized by strong anisotropy of properties of in the phonon subsystem, while those of the plasma subsystem demonstrate insignificant anisotropy. The authors of [3] used their experimental data to perform theoretical investigation of the coefficient of external reflection from polar uniaxial optically anisotropic single crystals of zinc oxide placed into strong uniform magnetic field. They found, for the first time, that there are some regions in the spectra of external reflection from the above single crystals where new oscillations appear. These oscillations are caused by the action of strong uniform magnetic field. The interaction between phonons and plasmons in ZnO single crystals placed into strong uniform magnetic field was studied. Similar results were obtained also for SiC single crystals (polytype 6H) that are characterized by insignificant anisotropy of properties of in the phonon subsystem, while their plasma subsystem demonstrates strong anisotropy of its properties.

In [5, 6], a possibility of excitation of various phonon and plasmon-phonon polaritons in polar optically anisotropic ZnO semiconductor was studied. It was shown in [6] that one can excite the first- and second-type surface polaritons (SP) in the region of residual rays. However, we did not find in the literature information on investigations of properties of optically anisotropic semiconductors (placed into strong magnetic field) performed using the attenuated total reflectance (ATR) technique, with allowance made for the plasmon-phonon interaction.

In this work, we studied the effect of strong uniform magnetic field on the properties of surface phonon and plasmon-phonon polaritons in optically anisotropic semiconductor (with ZnO single crystal serving as an example) for the orientations \( C || y, \ k \perp C, \ xy || C, \ H \perp \k, \ H || y, \ k_x = k, k_{y,z} = 0 \ (C \) is the optical axis of the crystal).
2. Experimental procedure

The ATR spectra of zinc oxide were measured using a spectrometer HKC-29M with an attachment HIIBO-2, according to the procedure considered in [2]. A semicylinder KRS-5 (index of refraction \( n_0 = 2.38 \)) served as ATR element. An air gap (width \( d = 2 \pm 15 \, \mu m \)) was made between the ATR prism and ZnO single crystal to be studied. The accuracy of angle adjustment in the attachment HIIBO-2 with an element KRS-5 corresponded to 0.1°. The spectral width of the slit in the course of ATR spectra recording with a polarizer was 3 cm\(^{-1}\); the polarization coefficient was 0.98. The accuracy of SP frequency determination was \( \pm 1 \, cm^{-1} \). The sample sizes were chosen in such a manner that one could obtain the ATR spectra in the \( p \)-polarized light at the above-mentioned orientations of the wave vector \( k \), crystal optical axis \( C \) and magnetic field \( \vec{H} \).

3. Theory

Let us consider the polar optically anisotropic ZnO single crystal in which excitation and propagation of various SP occur along the semiconductor surface (see Fig. 1). The \( x \)-axis lies along the direction of electromagnetic wave propagation, and \( C \parallel y \) (\( p \)-polarization).

The ATR coefficient \( I/I_0 \) in the IR spectral region (here \( I \) and \( I_0 \) are the intensities of IR radiation of a given frequency with and without a sample, respectively) was calculated with the expressions given in [2]. The contributions from the phonon and plasma subsystems of the semiconductor were taken to be additive, and allowance was made for the effect of strong uniform magnetic field on the single crystal. The agreement between the theoretical results and experimental data was achieved by usage in calculations of the components of permittivity tensor in magnetic field:

\[
\begin{align*}
\varepsilon_1 &= \frac{\varepsilon_{\omega, \omega} - \nu_1^2}{\varepsilon_{\omega, \omega} + \nu_1^2} - \frac{\nu_1^2}{\nu_1^2 - \nu_p^2} \\
\varepsilon_2 &= \frac{\varepsilon_{\omega, \omega} \nu_p^2}{\nu_p^2 - \nu_1^2} - \frac{\nu_p^2}{\nu_p^2 - \nu_1^2} \\
\varepsilon_3 &= \frac{\varepsilon_{\omega, \omega} - \nu_1^2}{\varepsilon_{\omega, \omega} + \nu_1^2} - \frac{\nu_1^2}{\nu_1^2 - \nu_p^2}
\end{align*}
\]

where \( \Omega = \frac{eH}{mc} \) is the cyclotron frequency, \( \varepsilon_{\omega, \omega} \) – high-frequency permittivity, \( \nu_{L, L} \), \( \nu_{T, L} \) – frequencies of the longitudinal and transverse optical phonons, respectively, \( \nu_p \) – plasma resonance frequency, \( \nu_{p, \omega} \) – plasmon damping frequency, and \( \nu_{f, \omega} \) – optical phonon damping frequency.

Tables 1 and 2 present the interconsistent bulk and electrophysical parameters of the optically anisotropic ZnO single crystals. The parameters were obtained with the analysis of variance of the IR reflectance spectra in the region of plasmon-phonon interaction and using polarized light at different angles of incidence [2].

Table 1. Bulk parameters of ZnO single crystals grown using the hydrothermal method.

<table>
<thead>
<tr>
<th>Sample</th>
<th>( n ), ( cm^{-3} )</th>
<th>( \nu_r ), ( cm^{-1} )</th>
<th>( \gamma_r ), ( cm^{-1} )</th>
<th>( \nu_p ), ( cm^{-1} )</th>
<th>( \gamma_p ), ( cm^{-1} )</th>
<th>( m_s )</th>
<th>( m_\perp )</th>
</tr>
</thead>
<tbody>
<tr>
<td>ZO2–3</td>
<td>9.3 x 10^16</td>
<td>90</td>
<td>150</td>
<td>11</td>
<td>0.21</td>
<td>0.258</td>
<td></td>
</tr>
<tr>
<td>ZO1–3</td>
<td>6.6 x 10^17</td>
<td>240</td>
<td>280</td>
<td>13</td>
<td>0.23</td>
<td>0.260</td>
<td></td>
</tr>
<tr>
<td>ZO6–B</td>
<td>2.0 x 10^18</td>
<td>420</td>
<td>406</td>
<td>21</td>
<td>0.22</td>
<td>0.260</td>
<td></td>
</tr>
</tbody>
</table>

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4. Results and discussion

Shown in Fig. 2 (curves 1, 2, and 3) are the calculated (curve) ATR spectra of ZnO (sample ZO2−3) for undoped zinc oxide \( (n = 9.3 \times 10^{10} \text{ cm}^{-3}) \) taken at the air gap \( d = 14.6 \text{ (1, 1')} \), 8.7 (2, 2') and 4.1 \( \mu \text{m} (3, 3') \), and angles of 30° (1, 1'), 35° (2, 2') and 50° (3, 3'). The calculations were performed with the expressions from [2] for the case of absent magnetic field. The dots show the experimental data for the sample ZO2−3 that were measured with the spectrometer HKC-29M using the procedure considered in [2]. The width of the air gap between the ATR prism and sample ZO2−3 was varied until the intensity of the absorbed wave did not exceed 20%, while the frequency of the minimum in the ATR spectrum remained constant [7]. The curves 1', 2', 3' show the calculated ATR spectra for a ZnO single crystal placed into uniform magnetic field. The calculations for the sample ZO2−3 were made for the case when magnetic field reached 100 kOe.

![Fig. 2. The ATR spectra of ZnO (sample ZO2−3): 1-3 – H=0; 1'-3' – H=100 kOe; 1, 1' – d = 14.6 \mu m, \varphi = 30°; 2, 2' – d = 8.7 \mu m, \varphi = 35°; 3, 3' – d = 4.1 \mu m, \varphi = 50°. Full and broken curves – calculation; dots – experiment.](image)

![Fig. 3. The ATR spectra of ZnO (sample ZOI–3): 1-3 – H=0; 1'-3' – H=100 kOe; 1, 1' – d = 13 \mu m; 2, 2' – d = 7.4 \mu m; 3, 3' – d = 3.4 \mu m. Full and broken curves – calculation; dots – experiment.](image)

The minima of the experimental and calculated spectra correspond to the SP frequencies \( \nu_{\text{min}} = 518 (1), 537 (2) \) and 551 \text{ cm}^{-1} (3) (without magnetic field) and \( \nu_{\text{min}} = 518 (1'), 538 (2') \) and 552 \text{ cm}^{-1} (3') (with magnetic field). The half-widths of the spectra are, correspondingly, \( \Gamma_v = 2, 3 \) and 9 \text{ cm}^{-1}.

Fig. 3 presents the ATR spectra (taken in the absence of magnetic field) for a low-doped zinc oxide single crystal \( (n = 6.6 \times 10^{17} \text{ cm}^{-3}) \), the angles of light incidence onto the ATR prism being 30° (1, 1'), 35° (2, 2') and 50° (3, 3'). The width of the air gap between the sample ZOI−3 and ATR prism varied from 13 (I) down to 3.4 \mu m (3). The minima of the spectra correspond to the SP frequencies \( \nu_{\text{min}} = 527 (1), 550 (2), \) and 563 \text{ cm}^{-1} (3), while the half-widths of the spectra are \( \Gamma_v = 5 (1), 17 (2) \) and 28 \text{ cm}^{-1} (3). The curves 1', 2', 3' are calculated with allowance for the effect of uniform magnetic field (100 kOe) on the zinc oxide single crystal (sample ZOI−3). The frequencies of minima and half-widths of the ATR spectra are, correspondingly, \( \nu_{\text{min}} = 528 (1'), 551 (2'), \) and 564 \text{ cm}^{-1} (3'), \( \nu_{\text{min}} = 7 (1'), 19 (2'), \) and 32 \text{ cm}^{-1} (3'). The dots show the experimental results for the ZnO single crystal (sample ZOI−3).

Shown in Fig. 4 (curves 1, 2, and 3) are the calculated ATR spectra for heavily doped ZnO \( (n = 2.0 \times 10^{18} \text{ cm}^{-3}) \). The angles of incidence were 30° (1, 1'), 35° (2, 2') and 50° (3, 3'); the widths of the air gaps between the ATR prism and zinc oxide were, correspondingly, 10.2 (1, 1'), 5.6 (2, 2') and 2.6 \mu m (3, 3'). The curves 1–3 correspond to the case when magnetic field is absent. The curves (dotted) 1', 2', 3' show the ATR spectra calculated with allowance made for the effect of strong magnetic field (100 kOe) on the ZnO single crystal, the orientation being \( \vec{H} \perp \vec{k} \), \( \vec{H} \parallel \vec{y} \). The spectral minima correspond to the SP frequencies \( \nu_{\text{min}} = 557 (1'), 574 (2'), \) and 582 \text{ cm}^{-1} (3'). The half-widths of the spectra are, correspondingly, \( \Gamma_v = 58 (1'), 92 (2'), \) and 99 \text{ cm}^{-1} (3').

Fig. 5 shows the ATR spectra for undoped zinc oxide (sample ZO2−3). The concentration of free charge carriers was \( n = 9.3 \times 10^{10} \text{ cm}^{-3} \); the angle of incidence of IR radiation onto the ATR prism was 50°. The air gap between the ATR prism and ZnO single crystal was 4 \mu m. The magnetic field strength was varied: it took the values 0 (I), 30 (2), 65 (3), and 100 kOe (4). The spectral minimum was at the frequency \( \nu_{\text{min}} = (551 \pm 1) \text{ cm}^{-1} \) for all the values of magnetic field strength; \( \Gamma_v = 9 \text{ cm}^{-1} \) for the curves 1–3 and 10 \text{ cm}^{-1} for the curve 4.
Fig. 4. The ATR spectra of ZnO (sample ZO6−B): 1–3 – \( H = 0 \); 1′–3′ – \( H = 100 \) kOe; 1, 1′ – \( d = 10.2 \) μm, \( \varphi = 30^\circ \); 2, 2′ – \( d = 5.6 \) μm, \( \varphi = 35^\circ \); 3, 3′ – \( d = 2.6 \) μm, \( \varphi = 50^\circ \).

Fig. 5. The ATR spectra of ZnO (sample ZO2−3): 1–4 – \( H = 0, 30, 65, 100 \) kOe; \( d = 4 \) μm, \( \varphi = 50^\circ \). Full and broken curves – calculation; dots – experiment.

Fig. 6. The ATR spectra of ZnO (sample ZOI−3): 1–4 – \( H = 0, 30, 65, 100 \) kOe; \( d = 5 \) μm, \( \varphi = 40^\circ \). Full and broken curves – calculation; dots – experiment.

Fig. 6 demonstrates the effect of magnetic field on the ATR spectra. In our calculations, we used the data from Tables 1 and 2 for the low-doped sample ZOI−3, with the angle of incidence 40° and air gap width 5 μm. The calculations were performed for magnetic fields of 0 (1) and 30 (2), 65 (3) and 100 kOe (4), respectively. From the ATR spectra, one can see that growth of magnetic field strength is accompanied with increase of both the ATR spectra half-width and SP damping. The position of the spectral minima remains the same: it corresponds to the SP frequency \( \nu_{\text{min}} = 558 \text{ cm}^{-1} \). \( \Gamma \) is 20, 22, 25 and 28 cm\(^{-1}\) for the curves 1–4, respectively.

The experimental spectrum (Fig. 6, dots 1) was taken for the low-doped sample ZOI−3 at frequency scanning; both the angle of IR radiation incidence onto the ATR prism (\( \varphi = 40^\circ \)) and air gap width (\( d = 5 \) μm) remained constant. The increase of the concentration of free charge carriers up to \( 2.0 \times 10^{18} \text{ cm}^{-3} \), at constant values (30, 65 and 100 kOe) of the strength of uniform magnetic field, angle of incidence and air gap width (4 μm), is accompanied with a shift of the SP frequency towards the low-frequency spectral region (see Fig. 7). The spectral minima correspond to the SP frequencies \( \nu_{\text{min}} = 581 \) (1), 580 (2) and 578 cm\(^{-1}\) (3), respectively, and \( \Gamma = 83 \) (1, 2), 88 (3) and 98 cm\(^{-1}\) (4). The curve 1 is calculated for the case of absent magnetic field.

One can see from Fig. 5 that magnetic field affects the absorption coefficient in the region of minimum of ATR spectrum and does not change the frequency of the minimum (within experimental errors). As the level of ZnO doping is being increased, the effect of magnetic field on the character of ATR spectrum becomes more substantial (see Figs 4 and 7).

Shown in Fig. 8 are the dispersion curves of zinc oxide single crystals when there is no magnetic field and allowance is made for the damping coefficients of the phonon and plasma subsystems. The concentration of free charge carriers (electrons) in ZnO varied from \( 3.9 \times 10^{16} \) cm\(^{-3}\) (1 and 1′) up to \( 6.6 \times 10^{17} \) cm\(^{-3}\) (3 and 3′). The curves 2 and 2′ were taken at \( n = 6.6 \times 10^{17} \) cm\(^{-3}\). The curves 1–3 correspond to the high-frequency dispersion branches with the limit frequency values \( \nu^{+}(k) = 561 \) (1), 578 (2) and 627 cm\(^{-1}\) (3). The curves 1′–3′ are calculated low-frequency dispersion branches with the limit frequency values \( \nu^{-}(k) = 59 \) (1′), 152 (2′) and 246 cm\(^{-1}\) (3′). The SP dispersion curves were calculated with the expressions from [2, 6-9] and data from Tables 1 and 2. The dots represent the experimental values obtained using the ATR technique. The limit frequency values \( \nu^{+} \) obtained from the ATR spectra of the samples under investigation are in agreement with the results of [6].
Table 3. The limit frequency values for SP of different types in ZnO single crystals placed into magnetic field.

<table>
<thead>
<tr>
<th>Sample</th>
<th>$\nu^–$, cm$^{-1}$</th>
<th>$\nu^+$, cm$^{-1}$</th>
<th>$\nu^–$, cm$^{-1}$</th>
<th>$\nu^f$, cm$^{-1}$</th>
<th>$\nu^+$, cm$^{-1}$</th>
<th>$\nu^–$, cm$^{-1}$</th>
<th>$\nu^f$, cm$^{-1}$</th>
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<tbody>
<tr>
<td>ZO1−3</td>
<td>59</td>
<td>561</td>
<td>34</td>
<td>108</td>
<td>561</td>
<td>21</td>
<td>171</td>
</tr>
<tr>
<td>ZO6−B</td>
<td>246</td>
<td>627</td>
<td>221</td>
<td>287</td>
<td>620</td>
<td>196</td>
<td>316</td>
</tr>
</tbody>
</table>

Fig. 7. The ATR spectra of ZnO (sample ZO6−B ). 1-4 – $H = 0, 30, 65, 100$ kOe; $d = 4$ $\mu$m, $\varphi = 40^\circ$.

One can see from Fig. 8 that increase in the concentration of free charge carriers (electrons) in ZnO is accompanied with a shift of the frequency of surface plasmon-phonon polaritons (SPPP) towards higher values. The SPPP high-frequency branches, whatever the degree of sample doping, begin from the frequency $\nu = \nu_T$ and exist at $k \gg 2\pi\nu/c$, approaching asymptotically the SP limiting frequency [2].

Fig. 9 presents the dispersion curves of low-doped ZnO single crystal (sample ZO1−3) obtained under action of uniform magnetic field (from 0 up to 100 kOe); the orientation was stated above. One can see from Fig. 9 (inset) that the high-frequency dispersion curve remains practically the same as magnetic field varies from zero up to 100 kOe, while the lower dispersion branch shifts towards lower frequencies as magnetic field grows. Besides, as it follows from Fig. 8, action of magnetic field on the ZnO single crystal within the frequency range from 190 up to 350 cm$^{-1}$ leads to appearance of one more dispersion branch (limited with the wave vector value) that shifts towards higher frequencies as the external magnetic field grows from 30 kOe. The reason is that, as the magnetic field strength increases, the cyclotron frequency $\Omega$ grows. In this case, coupling between plasmons and phonons breaks, and an additional dispersion branch appears [10]. The initial points for the lower and upper dispersion branches correspond to the frequencies $\nu = 0$ and $\nu_T$ that are solutions of the equations $\varepsilon_1 = 1$ or $\varepsilon_1 = 0$. Asymptotes to the additional branch are specified by the equations $1 + \varepsilon_1 \pm \varepsilon_2 = 0$. 3 gives the limit frequencies of the additional dispersion branches for ZnO single crystals with allowance made for the above requirements.

Table 4. The half-width of the minimum $\Gamma_i$ in the ATR spectra and SP damping coefficient $\Gamma_\phi$ in ZnO single crystals at $\nu_p = 0$, $\nu_T = 0$; $H = 0$ and 100 kOe.

<table>
<thead>
<tr>
<th>$\varphi$, deg.</th>
<th>$H = 0$</th>
<th>$H = 100$ kOe</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$\nu_{\text{min}}$, cm$^{-1}$</td>
<td>$\chi$</td>
</tr>
<tr>
<td>ZO1-3</td>
<td>30</td>
<td>527</td>
</tr>
<tr>
<td></td>
<td>35</td>
<td>550</td>
</tr>
<tr>
<td></td>
<td>50</td>
<td>564</td>
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<tr>
<td>ZO2-3</td>
<td>30</td>
<td>518</td>
</tr>
<tr>
<td></td>
<td>35</td>
<td>537</td>
</tr>
<tr>
<td></td>
<td>50</td>
<td>551</td>
</tr>
<tr>
<td>ZO6-B</td>
<td>30</td>
<td>555</td>
</tr>
<tr>
<td></td>
<td>35</td>
<td>576</td>
</tr>
<tr>
<td></td>
<td>50</td>
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</tbody>
</table>

Fig. 8. The dispersion curves for ZnO single crystal. Curves – calculation; dots – experiment.

One can see from Fig. 8 that increase in the concentration of free charge carriers (electrons) in ZnO is accompanied with a shift of the frequency of surface plasmon-phonon polaritons (SPPP) towards higher values. The SPPP high-frequency branches, whatever the degree of sample doping, begin from the frequency $\nu = \nu_T$ and exist at $k \gg 2\pi\nu/c$, approaching asymptotically the SP limiting frequency [2].

Fig. 9 presents the dispersion curves of low-doped ZnO single crystal (sample ZO1−3) obtained under action of uniform magnetic field (from 0 up to 100 kOe); the orientation was stated above. One can see from Fig. 9 (inset) that the high-frequency dispersion curve remains practically the same as magnetic field varies from zero up to 100 kOe, while the lower dispersion branch shifts towards lower frequencies as magnetic field grows. Besides, as it follows from Fig. 8, action of magnetic field on the ZnO single crystal within the frequency range from 190 up to 350 cm$^{-1}$ leads to appearance of one more dispersion branch (limited with the wave vector value) that shifts towards higher frequencies as the external magnetic field grows from 30 kOe. The reason is that, as the magnetic field strength increases, the cyclotron frequency $\Omega$ grows. In this case, coupling between plasmons and phonons breaks, and an additional dispersion branch appears [10]. The initial points for the lower and upper dispersion branches correspond to the frequencies $\nu = 0$ and $\nu_T$ that are solutions of the equations $\varepsilon_1 = 1$ or $\varepsilon_1 = 0$. Asymptotes to the additional branch are specified by the equations $1 + \varepsilon_1 \pm \varepsilon_2 = 0$. 3 gives the limit frequencies of the additional dispersion branches for ZnO single crystals with allowance made for the above requirements.
The effect of magnetic field on the dispersion curves of undoped and heavily doped ZnO single crystals (samples ZO2–3 and ZO6–B) is similar. Fig. 10 presents the calculated dispersion curves of ZnO placed into magnetic field of 100 kOe. One can see that, under action of magnetic field, all three dispersion branches shift towards higher frequencies as the concentration of free charge carriers grows.

![Fig. 9](image1.png)

Fig. 9. The dispersion curves for ZnO single crystal (sample ZOI–3) with allowance made for the effect of magnetic field: 1–4 – $H = 0, 30, 65, 100$ kOe.

![Fig. 10](image2.png)

Fig. 10. Different types of dispersion curves for ZnO single crystals placed into magnetic field of 100 kOe: 1 – ZO2–3; 2 – ZOI–3; 3 – ZO6–B.

![Fig. 11](image3.png)

Fig. 11. The frequency dependence of SP damping coefficient, $\Gamma_{sp}(\nu)$, for ZnO single crystal. The calculations were made for single crystals with ideally flat surface [2]. The curves 1–3 were calculated according to [2] for zinc oxide single crystals with different doping levels and without action of magnetic field. The curves 1’–3’ show $\Gamma_{sp}(\nu)$ dependence for the heavily doped sample ZO6–B placed into magnetic field $H \perp k$, $H \parallel y$ of 30 ($1'$), 65 ($2'$) and 100 kOe ($3'$).

The SP damping coefficient was determined with a graphical method in [2, 9]. Fig. 12 shows the procedure used for determination of the SP damping coefficient $\Gamma_{sp}(\nu)$ for optically anisotropic ZnO single crystals [9]. The curve 1 is the dispersion branch, while the dashed curves 2 and 3 determine the half-width $\Gamma_{p}$ of the ATR spectrum. The “true” spectrum width, according to [2], is equal to the SP damping coefficient and is determined by the difference of the frequencies located at the intersection of the perpendicular to the abscissa axis in the system “dashed curves – dispersion point”. Table 4 presents the calculated data $\Gamma_{sp}(\nu)$ and $\Gamma_{p}$ for ZnO single crystals with different doping levels under the conditions $\gamma_p = 0$, $\gamma_f = 0$, without and with magnetic field of 100 kOe. The angles of incidence onto the ATR prism were 30°, 35° and 50°.

One can see from Table 4 that, as the angle of incidence and concentration of free charge carriers in...
ZnO single crystal grow, the SP damping coefficient increases. The effect of plasmon-phonon damping and external magnetic field on $\Gamma_s$ and $\Gamma_{sp}$ in ZnO is similar.

5. Conclusion

Thus, action of strong uniform magnetic field on optically anisotropic ZnO single crystals (orientations $C \parallel y$, $\mathbf{k} \perp C$, $xy \parallel C$, $\mathbf{H} \perp \mathbf{k}$, $H \parallel y$, $k_x=k$, $k_y,z=0$) is accompanied by variation of the main properties of phonon and plasmon-phonon SP. For the first time, it has been shown that, in optically anisotropic ZnO single crystals subjected to action of uniform magnetic field, an additional dispersion branch may be excited. As the strength of the external magnetic field grows, that branch shifts towards higher frequencies. The initial frequency and character of the additional dispersion branch depend on the free charge carrier concentration and strength of magnetic field. The lower dispersion branch is deformed under action of magnetic field on the optically anisotropic single crystal, while the high-frequency branch of the dispersion curve remains the same (within experimental errors). As to the SP damping coefficient for ZnO, this problem needs additional studies. However, from the analysis performed it follows that the effect of external magnetic field is unquestionable.

References