# Longitudinal relaxation of mechanically free KH<sub>2</sub>PO<sub>4</sub> type crystals. Piezoelectric resonance and sound attenuation

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Within the framework of proton model with taking into account the piezoelectric interaction with the shear strain  $\varepsilon_6$ , a dynamic dielectric response of KD<sub>2</sub>PO<sub>4</sub> type ferroelectrics is considered. Experimentally observed phenomena of crystal clamping by high frequency electric field, piezoelectric resonance and microwave dispersion are described. Ultrasound velocity and attenuation are calculated, peculiarities of their temperature dependence at the Curie points are described. Existence of a cut-off frequency in the frequency dependence of attenuation is predicted.

Key words: ferroelectrics, piezoelectric resonance

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## 1. Introduction

Ferroelectric compounds of the  $\text{KH}_2\text{PO}_4$  family have been studied for nearly 70 years. Extensive experimental data have been accumulated; several models of the phase transitions in these crystals have been proposed and explored (see [1–14]). At the end of 1960-s, attention of theoretical and experimental studies of the ferroelectric compounds has shifted to the problem of the dynamic phenomena. Investigation of the low-frequency dispersion of their dielectric permittivity provides an important information about mechanisms of the phase transitions or dielectric response of the crystals. Despite the success in constructing a microscopic theory of the  $\text{KH}_2\text{PO}_4$  family crystals mostly phenomenological models were used in interpreting the experimental data as regards the dynamic characteristics. Such models provide no information about the microscopic nature of the dielectric dispersion and do not allow one to adequately describe the effect of different factors on its temperature and frequency dependencies. This problem has not been solved by the Green functions method or Bloch kinetic equation method [8,9,11,12] either.

Most theoretical works on the relaxation phenomena in  $\text{KH}_2\text{PO}_4$  family ferroelectrics is based on stochastic Glauber model [16]. For the first time this approach was used to describe relaxational phenomena in  $\text{KD}_2\text{PO}_4$  in [17], where the main features of longitudinal relaxation were explored within the four-particle cluster approximation in the paraelectric phase in  $\text{KD}_2\text{PO}_4$ . In that work the long-range interactions between deuterons was not taken into account, and the theoretical results were not compared with experiment. Later [18–22] a more sophisticated model of the  $\text{KD}_2\text{PO}_4$ type ferroelectrics and  $\text{ND}_4\text{D}_2\text{PO}_4$  type antiferroelectrics was proposed, in which longitudinal dynamic properties of these crystals were calculated within the four-particle cluster approximation for short-range interactions and mean field approximation for long-range interactions. It was shown [12,15,23–25] that the proposed [18–22] theory permits a satisfactory description of longitudinal relaxation in the  $\text{KH}_2\text{PO}_4$  type ferroelectrics. An attempt to develop a more consistent theory of the  $\text{KH}_2\text{PO}_4$  family ferroelectrics within the four-particle cluster approximation with taking into account the tunneling ( $\Omega$ ) was made in [26–28]. The model was not sophisticated enough to compare its results with experimental data for the dynamic characteristics of the crystals. At the same time, in these works the fact of suppression of the dynamic characteristics of  $\text{KH}_2\text{PO}_4$  type crystals by short-range correlations was established for the first time. Instead of the tunneling frequency, an effective parameter  $\tilde{\Omega}$  ( $\tilde{\Omega} \ll \Omega$ ) renormalized by short-range interactions was obtained. It should be noted that the established [26–28] suppression explains, most likely, the Debye character of dispersion of the dielectric permittivity observed in these crystals.

It should be stressed that the ferroelectrics of the  $\rm KH_2PO_4$  family are also piezoelectric. The piezoelectric coupling is revealed particularly when the crystals are subjected to external electric fields and mechanical stresses of certain symmetry. At the ferroelectric phase transition in the  $\rm KH_2PO_4$  type crystals, a spontaneous strain arises which changes their tetragonal symmetry.

Studies of the piezoelectric coupling effect on the phase transition and on physical properties of the KH<sub>2</sub>PO<sub>4</sub> type ferroelectrics were initiated in [29], where the Slater model [30] was modified by taking into account the splitting of the lowest ferroelectric level by the strain  $\varepsilon_6$ .

Fundamental are the results obtained in [31,32], where the proton ordering model was modified for the first time by taking into account the deformational molecular field related to the strain  $\varepsilon_6$ as well as the splitting of "lateral" deuteron configurations. In [33,34], with taking into account all possible splittings of configurational energies ("upper/lower", "lateral", and single-ionized configurations) due to the strain  $\varepsilon_6$ , the phase transition and the effect of the stress  $\sigma_6$  [33] and field  $E_3$ [34] on the physical characteristics of  $K(H_{0,12}D_{0,88})_2PO_4$  crystals were explored. Tunneling was not taken into account. A satisfactory agreement of theoretical results and experimental data was obtained.

Physical characteristics of the  $KH_2PO_4$  family crystals with taking into account piezoelectric coupling and tunneling are explored theoretically in [35].

So far the model consideration of longitudinal dielectric characteristics of the  $KD_2PO_4$  type ferroelectrics has been restricted to the static limit and to the microwave region [15,18–25]. Attempts to explore the piezoelectric resonance phenomenon within a model that does not take into account the piezoelectric coupling are pointless. Conventional proton ordering model of the  $KD_2PO_4$  type ferroelectrics does not permit one to describe the effects related to the difference between the free and clamped regimes or the phenomenon of crystal clamping by high-frequency electric field.

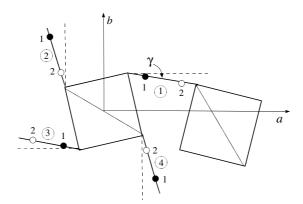
It seems natural to calculate, within the proposed in [33] proton ordering model with piezoelectric coupling, the dynamic characteristics of the  $\text{KD}_2\text{PO}_4$  type ferroelectrics in a wide frequency range from  $10^3$  Hz to  $10^{12}$  Hz, including the piezoelectric resonance region. In [36] within the framework of the four-particle cluster approximation for the proton ordering model, the thermodynamic and longitudinal dielectric, piezoelectric, and elastic characteristics of the  $\text{KD}_2\text{PO}_4$  have been calculated. It was shown that at the proper choice of the model parameters a good quantitative description of available experimental data by the proposed theory is obtained.

In this work, following the approach developed in [37], we shall calculate the longitudinal dynamic dielectric characteristics of the  $KD_2PO_4$  type ferroelectrics and study their temperature and frequency dependences. We shall investigate the effect of crystal clamping by longitudinal high-frequency external electric field. Expressions for the sound velocity and attenuation for a certain propagation direction will be obtained; their temperature and frequency dependences will be explored.

# 2. System of equations for time-dependent distribution functions of deuterons

We shall consider a system of deuterons moving on O-D...O bonds in deuterated ferroelectric of the  $KD_2PO_4$  type. A primitive Bravais cell of these crystals consists of two  $PO_4$  groups (tetrahedra) with four hydrogen bonds attached to one of them ("A" type tetrahedra), while the hydrogen bond attached to the other tetrahedron ("B" type) belongs to the four nearest structure elements surrounding it (figure 1).

The Hamiltonian of the deuteron subsystem with taking into account short-range and longrange interactions in the presence of mechanical stress  $\sigma_6 = \sigma_{xy}$  and electric field  $E_3$  along the



**Figure 1.** Primitive Bravais cell of the  $KD_2PO_4$  type crystal. One of the possible ferroelectric configurations is shown.

crystallographic axis c, which induce contributions to polarization  $P_3$  and strain  $\varepsilon_6$  of the crystal, consists of the "seed" and pseudospin parts [33,35]:

$$\hat{H} = NH^{(0)} + \hat{H}_s \,, \tag{2.1}$$

where N is the number of primitive cells. The "seed" energy of the primitive cell corresponds to the heavy ion subsystem and does not depend explicitly on hydrogen arrangement. It is expressed via the strain  $\varepsilon_6$  and electric field  $E_3$  and includes the elastic, piezoelectric, and electric counterparts

$$H^{(0)} = \bar{v} \left( \frac{1}{2} c_{66}^{E0} \varepsilon_6^2 - e_{36}^0 E_3 \varepsilon_6 - \frac{1}{2} \chi_{33}^{\varepsilon_0} E_3^2 \right),$$
(2.2)

where  $\bar{v} = \frac{v}{k_{\rm B}}$ , v is the primitive cell volume;  $c_{66}^{E0}$ ,  $e_{36}^{0}$ ,  $\chi_{33}^{\varepsilon 0}$  are the so-called "seed" elastic constant, coefficient of the piezoelectric stress, and dielectric susceptibility, respectively. These quantities determine the temperature behavior of the corresponding observable characteristics far from the transition point  $T_{\rm c}$ .

The pseudospin part of the Hamiltonian is

$$\hat{H} = N \left[ H^0 + 2\nu_{\rm c} (\eta^{(1)})^2 \right] + \sum_q \hat{H}_{q6}^{(4)},$$

where

$$\hat{H}_{q6}^{(4)} = -\sum_{f=1}^{4} \frac{z_{6}}{\beta} \frac{\sigma_{qf}}{2} + \frac{\varepsilon_{6}}{4} \left( -\delta_{s6} + 2\delta_{16} \right) \sum_{f=1}^{4} \frac{\sigma_{qf}}{2} 
- \varepsilon_{6} \left( \delta_{s6} + 2\delta_{16} \right) \left( \frac{\sigma_{q1}}{2} \frac{\sigma_{q2}}{2} \frac{\sigma_{q3}}{2} + \frac{\sigma_{q1}}{2} \frac{\sigma_{q2}}{2} \frac{\sigma_{q4}}{2} + \frac{\sigma_{q1}}{2} \frac{\sigma_{q3}}{2} \frac{\sigma_{q4}}{2} + \frac{\sigma_{q2}}{2} \frac{\sigma_{q3}}{2} \frac{\sigma_{q4}}{2} \right) 
+ \left( V + \delta_{a6} \varepsilon_{6} \right) \left( \frac{\sigma_{q1}}{2} \frac{\sigma_{q2}}{2} + \frac{\sigma_{q3}}{2} \frac{\sigma_{q4}}{2} \right) + \left( V - \delta_{a6} \varepsilon_{6} \right) \left( \frac{\sigma_{q2}}{2} \frac{\sigma_{q3}}{2} + \frac{\sigma_{q4}}{2} \frac{\sigma_{q1}}{2} \right) 
+ U \left( \frac{\sigma_{q1}}{2} \frac{\sigma_{q3}}{2} + \frac{\sigma_{q2}}{2} \frac{\sigma_{q4}}{2} \right) + \Phi \frac{\sigma_{q1}}{2} \frac{\sigma_{q2}}{2} \frac{\sigma_{q3}}{2} \frac{\sigma_{q4}}{2} ,$$
(2.3)

and

$$z_6 = \beta(-\Delta^c + 2\nu_c \eta^{(1)} - 2\psi_6\varepsilon_6 + \mu_3 E_3), \qquad \left(\beta = \frac{1}{k_B T}\right)$$

The dynamic properties of the  $KD_2PO_4$  type ferroelectrics will be explored within the proposed dynamic model [21], based on the stochastic Glauber model [16] ideas. When calculating the

dynamic characteristics, we shall restrict our consideration to the four-particle cluster approximation, providing a successful description of thermodynamic characteristics of the crystals [8–10,15]. Using the approach, developed in [18–21], we obtain the following system of equations for the time-dependent deuteron distribution functions:

$$-\alpha \frac{\mathrm{d}}{\mathrm{d}t} \left\langle \prod_{f} \sigma_{qf} \right\rangle = \sum_{f'} \left\{ \left\langle \prod_{f} \sigma_{qf} \left[ 1 - \sigma_{qf'} \tanh \frac{1}{2} \beta \boldsymbol{\varepsilon}_{qf'}^{z}(t) \right] \right\rangle \right\},\tag{2.4}$$

where  $\varepsilon_{qf'}^{z}(t)$  is the local field acting on the f'th deuteron in the qth cell, which can be found from the Hamiltonian (2.3).

The expressions  $\tanh \frac{\beta \varepsilon_{qf'}^z(t)}{2}$  are presented in the following form

$$\tanh \frac{\beta \varepsilon_{q1}^2}{2} = P_6^z \sigma_{q3} + Q_{61}^z \sigma_{q2} + Q_{62}^z \sigma_{q4} + R_6^z \sigma_{q2} \sigma_{q3} \sigma_{q4} \\
+ M_{61}^z \sigma_{q2} \sigma_{q3} + M_{62}^z \sigma_{q3} \sigma_{q4} + N_6^z \sigma_{q2} \sigma_{q4} + L_6^z , \\
\tanh \frac{\beta \varepsilon_{q2}^2}{2} = P_6^z \sigma_{q4} + Q_{61}^z \sigma_{q1} + Q_{62}^z \sigma_{q3} + R_6^z \sigma_{q1} \sigma_{q3} \sigma_{q4} \\
+ M_{61}^z \sigma_{q4} \sigma_{q2} + M_{62}^z \sigma_{q3} \sigma_{q4} + N_6^z \sigma_{q1} \sigma_{q3} + L_6^z , \\
\tanh \frac{\beta \varepsilon_{q3}^2}{2} = P_6^z \sigma_{q1} + Q_{61}^z \sigma_{q4} + Q_{62}^z \sigma_{q2} + R_6^z \sigma_{q1} \sigma_{q2} \sigma_{q4} \\
+ M_{61}^z \sigma_{q1} \sigma_{q4} + M_{62}^z \sigma_{q1} \sigma_{q2} + N_6^z \sigma_{q2} \sigma_{q4} + L_6^z , \\
\tanh \frac{\beta \varepsilon_{q4}^2}{2} = P_6^z \sigma_{q2} + Q_{61}^z \sigma_{q3} + Q_{62}^z \sigma_{q1} + R_6^z \sigma_{q1} \sigma_{q2} \sigma_{q3} \\
+ M_{61}^z \sigma_{q2} \sigma_{q3} + M_{62}^z \sigma_{q1} \sigma_{q2} + N_6^z \sigma_{q1} \sigma_{q2} + L_6^z .$$
(2.5)

Since  $\sigma_{qf} = \pm 1$ , we find for the coefficients  $P_6, \ldots, L_6$ :

$$P_{6}^{z} = \frac{1}{8}(l_{1}^{z} - l_{2}^{z} + n_{1}^{z} - n_{2}^{z} + m_{1}^{z} - m_{2}^{z} + m_{3}^{z} - m_{4}^{z}),$$

$$Q_{61}^{z} = \frac{1}{8}(l_{1}^{z} - l_{2}^{z} - n_{1}^{z} + n_{2}^{z} + m_{1}^{z} + m_{2}^{z} - m_{3}^{z} - m_{4}^{z}),$$

$$Q_{62}^{z} = \frac{1}{8}(l_{1}^{z} - l_{2}^{z} - n_{1}^{z} + n_{2}^{z} - m_{1}^{z} - m_{2}^{z} + m_{3}^{z} + m_{4}^{z}),$$

$$R_{6}^{z} = \frac{1}{8}(l_{1}^{z} - l_{2}^{z} + n_{1}^{z} - n_{2}^{z} - m_{1}^{z} + m_{2}^{z} - m_{3}^{z} + m_{4}^{z}),$$

$$N_{6}^{z} = \frac{1}{8}(l_{1}^{z} + l_{2}^{z} + n_{1}^{z} + n_{2}^{z} - m_{1}^{z} - m_{2}^{z} - m_{3}^{z} - m_{4}^{z}),$$

$$M_{61}^{z} = \frac{1}{8}(l_{1}^{z} + l_{2}^{z} - n_{1}^{z} - n_{2}^{z} + m_{1}^{z} - m_{2}^{z} - m_{3}^{z} + m_{4}^{z}),$$

$$M_{62}^{z} = \frac{1}{8}(l_{1}^{z} + l_{2}^{z} - n_{1}^{z} - n_{2}^{z} - m_{1}^{z} + m_{2}^{z} + m_{3}^{z} - m_{4}^{z}),$$

$$L_{6}^{z} = \frac{1}{8}(l_{1}^{z} + l_{2}^{z} + n_{1}^{z} + n_{2}^{z} + m_{1}^{z} + m_{2}^{z} + m_{3}^{z} + m_{4}^{z}),$$

$$(2.6)$$

where we use the following notations

$$l_{\frac{1}{2}}^{z} = \tanh \frac{\beta}{2} \left[ \pm w + (\delta_{s6} + \delta_{16})\varepsilon_{6} + \frac{z_{6}^{z}}{\beta} \right], \qquad n_{\frac{1}{2}}^{z} = \tanh \frac{\beta}{2} \left[ \pm (w - w_{1}) - \delta_{16}\varepsilon_{6} + \frac{z_{6}^{z}}{\beta} \right],$$
$$m_{\frac{1}{4}}^{z} = \tanh \frac{\beta}{2} \left[ \pm (\varepsilon - \delta_{a6}\varepsilon_{6} - w) - \delta_{16}\varepsilon_{6} + \frac{z_{6}^{z}}{\beta} \right], \qquad m_{\frac{2}{3}}^{z} = \tanh \frac{\beta}{2} \left[ \mp (\varepsilon + \delta_{a6}\varepsilon_{6} - w) - \delta_{16}\varepsilon_{6} + \frac{z_{6}^{z}}{\beta} \right]$$

Taking into account the symmetry of distribution functions

$$\eta^{(1)z} = \langle \sigma_{q1} \rangle = \langle \sigma_{q2} \rangle = \langle \sigma_{q3} \rangle = \langle \sigma_{q4} \rangle,$$
  

$$\eta^{(3)z} = \langle \sigma_{q1}\sigma_{q2}\sigma_{q3} \rangle = \langle \sigma_{q1}\sigma_{q3}\sigma_{q4} \rangle = \langle \sigma_{q1}\sigma_{q2}\sigma_{q4} \rangle = \langle \sigma_{q2}\sigma_{q3}\sigma_{q4} \rangle,$$
  

$$\eta^{(2)z}_{1} = \langle \sigma_{q2}\sigma_{q3} \rangle = \langle \sigma_{q1}\sigma_{q4} \rangle, \quad \eta^{(2)z}_{2} = \langle \sigma_{q1}\sigma_{q2} \rangle = \langle \sigma_{q3}\sigma_{q4} \rangle, \quad \eta^{(2)z}_{3} = \langle \sigma_{q1}\sigma_{q3} \rangle = \langle \sigma_{q2}\sigma_{q4} \rangle, \quad (2.7)$$

from (2.4) with taking into account (2.5) and (2.7), we can obtain a closed system of equations for the time-dependent single-particle, pair, and three-particle deuteron distribution functions in the  $KD_2PO_4$  type crystals

$$\alpha \frac{\mathrm{d}}{\mathrm{d}t} \begin{pmatrix} \eta^{(1)z} \\ \eta^{(3)z} \\ \eta^{(2)z} \\ \overline{c}_{51} \quad \overline{c}_{52} \quad \overline{c}_{53} \quad \overline{c}_{54} \quad \overline{c}_{55} \end{pmatrix} \begin{pmatrix} \eta^{(1)z} \\ \eta^{(3)z} \\ \eta^{(3)z} \\ \eta^{(2)z} \\ \eta^{(2$$

Here we use the following notations

$$\begin{split} \bar{c}_{11} &= -(1-P_6^z-Q_{61}^z-Q_{62}^z), \ \bar{c}_{12} = R_6^z, \ \bar{c}_{13} = M_{61}^z, \ \bar{c}_{14} = M_{62}^z, \ \bar{c}_{15} = N_6^z, \ \bar{c}_{1} = L_6^z, \\ \bar{c}_{21} &= (2P_6^z+2Q_{61}^z+2Q_{62}^z+3R_6), \ \bar{c}_{22} = -(3-P_6^z-Q_{61}^z-Q_{62}^z), \ \bar{c}_{23} = (N_6^z+M_{62}^z+L_6^z), \\ \bar{c}_{24} &= (N_6^z+M_{61}^z+L_6^z), \ \bar{c}_{25} = (M_{61}^z+M_{62}^z+L_6^z), \ \bar{c}_{2} = (N_6^z+M_{61}^z+M_{62}^z), \\ \bar{c}_{31} &= 2(N_6^z+M_{62}^z+L_6^z), \ \bar{c}_{32} = 2M_{61}^z, \ \bar{c}_{33} = -2(1-R_6^z), \ \bar{c}_{34} = 2P_6^z, \ \bar{c}_{35} = 2Q_{61}^z, \ \bar{c}_{3} = 2Q_{62}^z, \\ \bar{c}_{41} &= 2(N_6^z+M_{61}^z+L_6^z), \ \bar{c}_{42} = 2M_{62}^z, \ \bar{c}_{43} = 2P_6^z, \ \bar{c}_{44} = -2(1-R_6^z), \ \bar{c}_{45} = 2Q_{62}^z, \ \bar{c}_{4} = 2Q_{61}^z, \\ \bar{c}_{51} &= 2(M_{61}^z+M_{62}^z+L_6^z), \ \bar{c}_{52} = 2N_6^z, \ \bar{c}_{53} = 2Q_{61}^z, \ \bar{c}_{54} = 2Q_{62}^z, \ \bar{c}_{55} = -2(1-R_6^z), \ \bar{c}_{5} = 2P_6^z. \end{split}$$

In the single-particle approximation from (2.4) we obtain the following equation for distribution functions

$$\alpha \frac{\mathrm{d}}{\mathrm{d}t} \eta^{(1)z} = -\eta^{(1)z} + \tanh \frac{1}{2} \bar{z}_6^z \,. \tag{2.10}$$

# 3. Dynamic characteristics of a mechanically free $K(H_{1-x}D_x)_2PO_4$ crystal. Piezoelectric resonance

In this section we shall consider vibrations of a thin  $l \times l$  square plate of a KD<sub>2</sub>PO<sub>4</sub> crystal, cut in the [001] plane, induced by time-dependent electric field  $E_{3t} = E_3 e^{i\omega t}$ . This field, in addition to the shear strain  $\varepsilon_6$ , also induces diagonal components of the strain tensor  $\varepsilon_i$ . For the sake of simplicity, we shall neglect the diagonal strains.

Dynamics of deformational processes in  $\mathrm{KD}_2\mathrm{PO}_4$  will be described using classical Newtonian equations of motion of an elementary volume

$$\rho \frac{\partial^2 u_i}{\partial t^2} = \sum_k \frac{\partial \sigma_{ik}}{\partial x_k},\tag{3.1}$$

where  $\rho$  is the cell volume,  $u_i$  are the displacements of an elementary volume along the axis  $x_i$ ,  $\sigma_{ik}$  is the mechanical stress. The shear strain  $\varepsilon_6$  is determined by the displacements  $u_x = u_1$  and  $u_y = u_z$ , that is

$$\varepsilon_6 = \varepsilon_{xy} = \frac{\partial u_1}{\partial y} + \frac{\partial u_2}{\partial x}.$$

In our case a shear strain  $\sigma_{xy} = \sigma_6$  is different from zero and [36]

$$\sigma_6 = c_{66}^{E0} \varepsilon_6 - e_{36}^0 E_3 + \frac{4\psi_6}{v} \frac{m^z}{D_6^z} + \frac{2\delta_{a6}}{\bar{v}D_6^z} M_{a6} - \frac{2\delta_{s6}}{vD_6^z} M_{s6} + \frac{2\delta_{16}}{vD_6^z} M_{16} , \qquad (3.2)$$

where

$$M_{a6} = aa_6 - \frac{a}{a_6}$$
,  $M_{s6} = \sinh(2z_6^z + \beta\delta_{s6}\varepsilon_6)$ ,  $M_{16} = 4b\sinh(z_6^z - \beta\delta_{16}\varepsilon_6)$ 

Thus,

$$\rho \frac{\partial^2 u_1}{\partial t^2} = c_{66}^{E0} \frac{\partial \varepsilon_6}{\partial y} + \frac{4\psi_6}{v} \frac{\partial \eta^{(1)}}{\partial y} + \frac{2\delta_{a6}}{v} \frac{\partial}{\partial y} \left(\frac{M_{a6}}{D_6}\right) - \frac{2\delta_{s6}}{v} \frac{\partial}{\partial y} \left(\frac{M_{s6}}{D_6}\right) + \frac{2\delta_{16}}{v} \frac{\partial}{\partial y} \left(\frac{M_{16}}{D_6}\right), \quad (3.3)$$
$$\rho \frac{\partial^2 u_2}{\partial t^2} = c_{66}^{E0} \frac{\partial \varepsilon_6}{\partial x} + \frac{4\psi_6}{v} \frac{\partial \eta^{(1)}}{\partial x} + \frac{2\delta_{a6}}{v} \frac{\partial}{\partial x} \left(\frac{M_{a6}}{D_6}\right) - \frac{2\delta_{s6}}{v} \frac{\partial}{\partial x} \left(\frac{M_{s6}}{D_6}\right) + \frac{2\delta_{16}}{v} \frac{\partial}{\partial x} \left(\frac{M_{16}}{D_6}\right).$$

At small deviations from the equilibrium we can separate in the systems (2.6) and (2.8) the static and time-dependent parts, presenting the distribution functions, effective displacement fields  $u_1$ ,  $u_2$ , and the strain  $\varepsilon_6$  as sums of equilibrium values and of fluctuational deviations

$$\eta^{(1)} = \tilde{\eta}^{(1)} + \eta_t^{(1)}, \qquad \eta^{(3)} = \tilde{\eta}^{(3)} + \eta_t^{(3)}, \qquad \eta_i^{(2)} = \tilde{\eta}_i^{(2)} + \eta_t^{(2)}, \qquad (i = 1, 2, 3),$$
  

$$\varepsilon_6 = \tilde{\varepsilon}_6 + \varepsilon_{6t}, \qquad u_{1,2} = \tilde{u}_{1,2} + u_{1,2t}, \qquad (3.4)$$
  

$$z_6^z = \tilde{z}_6 + z_{6t} = -\beta \tilde{\Delta} + 2\beta \nu_c \tilde{\eta}^{(1)} - 2\beta \psi_6 \varepsilon_6 - \beta \Delta_t + 2\beta \nu_c \eta_t^{(1)} - 2\beta \psi_6 \varepsilon_{6t} + \beta \mu_3 E_{3t}.$$

We substitute the expressions (3.4) into the system of equations (2.8), (2.10), expand into a series over the time dependent terms the coefficients  $\bar{c}_{ij}$ ,  $\bar{c}_i$ , limit oneself to linear approximation. We exclude parameter  $\Delta_t$  and obtain the system of equations for fluctuating parts of distribution functions:

$$\frac{\mathrm{d}}{\mathrm{d}t} \begin{pmatrix} \eta_{t}^{(1)} \\ \eta_{t}^{(3)} \\ \eta_{2t}^{(2)} \\ \eta_{2t}^{(2)} \\ \eta_{3t}^{(2)} \\ \eta_{3t}^{(2)} \end{pmatrix} = \begin{pmatrix} c_{11} & c_{12} & c_{13} & c_{14} & c_{15} \\ c_{21} & c_{22} & c_{23} & c_{24} & c_{25} \\ c_{31} & c_{32} & c_{33} & c_{34} & c_{35} \\ c_{41} & c_{42} & c_{43} & c_{44} & c_{45} \\ c_{51} & c_{52} & c_{53} & c_{54} & c_{55} \end{pmatrix} \begin{pmatrix} \eta_{t}^{(1)} \\ \eta_{t}^{(2)} \\ \eta_{2t}^{(2)} \\ \eta_{3t}^{(2)} \end{pmatrix} - \frac{1}{2}\beta\mu_{3}E_{3t} \begin{pmatrix} c_{1} \\ c_{2} \\ c_{3} \\ c_{4} \\ c_{5} \end{pmatrix} \\ +\beta\psi_{6}\varepsilon_{6t} \begin{pmatrix} c_{1} \\ c_{2} \\ c_{3} \\ c_{4} \\ c_{5} \end{pmatrix} - \beta\delta_{s6}\varepsilon_{6t} \begin{pmatrix} c_{1s} \\ c_{3s} \\ c_{21s} \\ c_{22s} \\ c_{23s} \end{pmatrix} + \beta\delta_{a6}\varepsilon_{6t} \begin{pmatrix} c_{1a} \\ c_{3a} \\ c_{21a} \\ c_{22a} \\ c_{23a} \end{pmatrix} - \beta\delta_{16}\varepsilon_{6t} \begin{pmatrix} \tilde{c}_{11} \\ \tilde{c}_{21} \\ \tilde{c}_{31} \\ \tilde{c}_{41} \\ \tilde{c}_{51} \end{pmatrix}. \quad (3.5)$$

The expressions for coefficients of the system (3.5) are given in [36].

Taking into account (3.4), equations for the displacements (3.3) can be rewritten as follows:

$$\rho \frac{\partial^2 u_{1t}}{\partial t^2} = c_{16} \frac{\partial \varepsilon_{6t}}{\partial y} + c_{26} \frac{\partial \eta_t^{(1)}}{\partial y}, \qquad \rho \frac{\partial^2 u_{2t}}{\partial t^2} = c_{16} \frac{\partial \varepsilon_{6t}}{\partial x} + c_{26} \frac{\partial \eta_t^{(1)}}{\partial x}, \tag{3.6}$$

where we use the following notations

$$\begin{aligned} c_{16} &= c_{66}^{E0} + \frac{4\beta\psi_6}{vD_6} f_6 - \frac{2\beta}{vD_6} \Big[ \delta_{s6}^2 \cosh(2\tilde{z}_6 + \beta\delta_{s6}\tilde{\varepsilon}_6) + \delta_{16}^2 4b \cosh(\tilde{z}_6 - \beta\delta_{16}\tilde{\varepsilon}_6) \\ &+ \delta_{a6}^2 2a \cosh\beta\delta_{a6}\tilde{\varepsilon}_6 \Big] + \frac{2\beta}{vD_6^2} (-\delta_{s6}M_{s6} + \delta_{16}M_{16} + \delta_{a6}M_{a6})^2, \\ c_{26} &= \frac{4}{v} \left( \psi_6 - \frac{\varphi_6^{\eta}}{D_6} f_6 \right), \end{aligned}$$

We shall look for solutions of the system (3.5) and (3.6) in the form of harmonic waves

$$\eta_t^{(1)} = \eta^{(1)}(6, x, y) e^{i\omega t}, \qquad \eta_t^{(3)} = \eta^{(3)}(6, x, y) e^{i\omega t}, \eta_{1t}^{(2)} = \eta_1^{(2)}(6, x, y) e^{i\omega t}, \qquad \eta_{2t}^{(2)} = \eta_2^{(2)}(6, x, y) e^{i\omega t}, \eta_{3t}^{(2)} = \eta_3^{(2)}(6, x, y) e^{i\omega t}, \qquad \varepsilon_{6t} = \varepsilon_6(x, y) e^{i\omega t}, u_{1t} = u_1(y) e^{i\omega t}, \qquad u_{2t} = u_2(x) e^{i\omega t}.$$
(3.7)

Finally, solving the system (3.5) with taking into account (3.7), we find that

$$\eta^{(1)}(6, x, y) = \frac{\beta \mu_3}{2} F^{(1)}(\alpha \omega) E_3 + \left[ -\beta \psi_6 F^{(1)}(\alpha \omega) + \beta \delta_{16} F_1^{(1)}(\alpha \omega) \right] \varepsilon_6(x, y), \qquad (3.8)$$

where we use the notations

$$F^{(1)}(\alpha\omega) = \frac{p^{(4)}(i\alpha\omega)^4 + p^{(3)}(i\alpha\omega)^3 + p^{(2)}(i\alpha\omega)^2 + p^{(1)}(i\alpha\omega) + p^{(0)}}{(i\alpha\omega)^5 + p_4(i\alpha\omega)^4 + p_3(i\alpha\omega)^3 + p_2(i\alpha\omega)^2 + p_1(i\alpha\omega) + p_0},$$

$$F^{(1)}_s(\alpha\omega) = \frac{p^{(4)}_s(i\alpha\omega)^4 + p^{(3)}_s(i\alpha\omega)^3 + p^{(2)}_s(i\alpha\omega)^2 + p^{(1)}_s(i\alpha\omega) + p^{(0)}_s}{(i\alpha\omega)^5 + p_4(i\alpha\omega)^4 + p_3(i\alpha\omega)^3 + p_2(i\alpha\omega)^2 + p_1(i\alpha\omega) + p_0},$$

$$F^{(1)}_a(\alpha\omega) = \frac{p^{(4)}_a(i\alpha\omega)^4 + p^{(3)}_a(i\alpha\omega)^3 + p^{(2)}_a(i\alpha\omega)^2 + p^{(1)}_a(i\alpha\omega) + p^{(0)}_a}{(i\alpha\omega)^5 + p_4(i\alpha\omega)^4 + p_3(i\alpha\omega)^3 + p_2(i\alpha\omega)^2 + p_1(i\alpha\omega) + p_0},$$

$$F^{(1)}_1(\alpha\omega) = \frac{p^{(4)}_1(i\alpha\omega)^4 + p^{(3)}_1(i\alpha\omega)^3 + p^{(2)}_1(i\alpha\omega)^2 + p^{(1)}_1(i\alpha\omega) + p^{(0)}_a}{(i\alpha\omega)^5 + p_4(i\alpha\omega)^4 + p_3(i\alpha\omega)^3 + p_2(i\alpha\omega)^2 + p_1(i\alpha\omega) + p_0}.$$
(3.9)

Expressions for  $p_4, \ldots, p_0, p^{(4)}, \ldots, p^{(0)}, p_s^{(4)}, \ldots, p_s^{(0)}, p_a^{(4)}, \ldots, p_a^{(0)}, p_1^{(4)}, \ldots, p_1^{(0)}$  in (3.9) are given in [36].

Taking into account the relations (3.6) and (3.7), we obtain the following wave equations for  $u_1, u_2$ :

$$c_{66}^{E}(\alpha\omega)\frac{\partial^{2}u_{1}}{\partial y^{2}} + \rho\omega^{2}u_{1} = 0, \qquad c_{66}^{E}(\alpha\omega)\frac{\partial^{2}u_{2}}{\partial x^{2}} + \rho\omega^{2}u_{2} = 0,$$
(3.10)

where

$$\begin{aligned} c_{66}^{E}(\alpha\omega) &= c_{66}^{E0} + \frac{4\beta\psi_{6}}{vD_{6}}f_{6} + \frac{2\beta}{vD_{6}^{2}}(-\delta_{s6}M_{s6} + \delta_{16}M_{16} + \delta_{a6}M_{a6})^{2} \\ &+ \frac{4\beta\psi_{6}}{v}\Big[-\psi_{6}F^{(1)}(\alpha\omega) + \delta_{s6}F_{s}^{(1)}(\alpha\omega) + \delta_{16}F_{1}^{(1)}(\alpha\omega) - \delta_{a6}F_{a}^{(1)}(\alpha\omega)\Big] \\ &- \frac{4\varphi_{3}f_{6}}{vD_{6}}\beta\Big[-\psi_{6}F^{(1)}(\alpha\omega) + \delta_{s6}F_{s}^{(1)}(\alpha\omega) + \delta_{16}F_{1}^{(1)}(\alpha\omega) - \delta_{a6}F_{a}^{(1)}(\alpha\omega)\Big] \\ &- \frac{2\beta}{vD_{6}}\Big[\delta_{s6}^{2}\cosh(2\tilde{z} + \beta\delta_{s6}\tilde{\varepsilon}_{6}) + 4b\delta_{16}^{2}\cosh(\tilde{z} - \beta\delta_{16}\tilde{\varepsilon}_{6}) + \delta_{a6}^{2}2a\cosh\beta\delta_{a6}\tilde{\varepsilon}_{6}^{2}\Big].\end{aligned}$$

Equation (3.10) can be written as

$$\frac{\partial^2 u_1}{\partial y^2} + k_6^2 u_1 = 0, \qquad \frac{\partial^2 u_2}{\partial x^2} + k_6^2 u_2 = 0, \tag{3.11}$$

where  $k_6$  is the wavenumber

$$k_6 = \frac{\omega\sqrt{\rho}}{\sqrt{c_{66}^E(\alpha\omega)}}.$$
(3.12)

We shall look for the solutions of (3.11) in the form

$$u_1 = A_1 \cos k_6 y + B_1 \sin k_6 y,$$
  $u_2 = A_2 \cos k_6 x + B_2 \sin k_6 x$ 

As a result

$$\varepsilon_6(x,y) = k_6 \Big[ -(A_1 \cos k_6 y + A_2 \cos k_6 x) + (B_1 \sin k_6 y + B_2 \sin k_6 x) \Big].$$
(3.13)

The boundary conditions are set as follows:

$$\varepsilon_6(0,0) = \varepsilon_6(l,l) = \varepsilon_6(0,l) = \varepsilon_6(l,0) = \varepsilon_0. \tag{3.14}$$

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The values of  $\varepsilon_0$  are determined from (3.2), using the relation (3.10)

$$\varepsilon_0 = \frac{e_{36}(\alpha\omega)}{c_{66}^E(\alpha\omega)} E_3.$$
(3.15)

where

$$e_{36}(\alpha\omega) = e_{36}^0 + \frac{\beta\mu_3}{v} \Big[ -\psi_6 F^{(1)}(\alpha\omega) + \delta_{s6} F_s^{(1)}(\alpha\omega) + \delta_{16} F_1^{(1)}(\alpha\omega) - \delta_{a6} F_a^{(1)}(\alpha\omega) \Big].$$
(3.16)

Taking into account the boundary conditions (3.14), we find from (3.13)

$$\varepsilon_6(x,y) = \frac{\varepsilon_0}{2} \left[ -\frac{\cos k_6 l - 1}{\sin k_6 l} (\sin k_6 y + \sin k_6 x) + (\cos k_6 y + \cos k_6 x) \right].$$
(3.17)

Using the expression, relating polarization  $P_3$  to the order parameter  $\eta^{(1)}$  and strain  $\varepsilon_6$ , as well as relation (3.8), we find that

$$P_3(x, y, t) = P_3(x, y) e^{i\omega t}, (3.18)$$

where

$$P_3(x,y) = e_{36}(\alpha\omega)\varepsilon_6(x,y) + \chi_{33}^{\varepsilon}(\alpha\omega)E_3$$

and

$$\chi_{33}(\alpha\omega) = \chi_{33}^{\varepsilon 0} + \frac{\beta\mu_3^2}{2v}F^{(1)}(\alpha\omega).$$

Now we can calculate the dynamic dielectric susceptibility of a free crystal  $\chi^{\sigma}_{33}(\alpha\omega)$ 

$$\chi_{33}^{\sigma}(\alpha\omega) = \frac{1}{l^2} \frac{\partial}{\partial E_3} \int_0^l \int_0^l P_3(x,y) \mathrm{d}x \mathrm{d}y.$$
(3.19)

Taking into account (3.17), we find that

$$\frac{1}{l^2} \int_0^l \int_0^l \mathrm{d}x \mathrm{d}y \varepsilon_6(x, y) = \frac{2\varepsilon_0}{k_6} \tan \frac{k_6 l}{2} = \frac{\varepsilon_0}{R(\omega)}, \qquad (3.20)$$

where

$$\frac{1}{R(\omega)} = \frac{2}{k_6 l} \tan \frac{k_6 l}{2} \,.$$

With (3.18) and (3.20) from (3.19) we obtain that

$$\chi_{33}^{\sigma}(\alpha\omega) = \chi_{33}^{\varepsilon}(\alpha\omega) + \frac{1}{R(\omega)} \frac{e_{36}^2(\alpha\omega)}{e_{66}^E(\alpha\omega)}.$$
(3.21)

# 4. Attenuation and velocity of ultrasound in KD<sub>2</sub>PO<sub>4</sub> crystals

Pulsed ultrasonics provide a useful method for the investigation of crystal behavior. The ultrasound wavelength is usually much smaller than the sample dimensions. Therefore, the dynamical variables, such as elementary displacements and order parameter, depend only on the spatial coordinate which is the direction of sound propagation.

If thin bars of the crystal are cut along [001], then we shall consider a transverse sound wave propagating along the bar and polarized along [010]. Among  $\frac{\partial u_i}{\partial x_j}$  the only nonzero derivative is

 $\frac{\partial u_2}{\partial x}$ ; therefore, instead of the systems (3.5) and (3.6), we can write

$$\frac{\mathrm{d}}{\mathrm{d}t}\begin{pmatrix}\eta_{t}^{(1)}\\\eta_{t}^{(3)}\\\eta_{1t}^{(2)}\\\eta_{2t}^{(2)}\\\eta_{3t}^{(2)}\end{pmatrix} = \begin{pmatrix}c_{11} & c_{12} & c_{13} & c_{14} & c_{15}\\c_{21} & c_{22} & c_{23} & c_{24} & c_{25}\\c_{31} & c_{32} & c_{33} & c_{34} & c_{35}\\c_{41} & c_{42} & c_{43} & c_{44} & c_{45}\\c_{51} & c_{52} & c_{53} & c_{54} & c_{55}\end{pmatrix}\begin{pmatrix}\eta_{t}^{(1)}\\\eta_{t}^{(2)}\\\eta_{2t}^{(2)}\\\eta_{3t}^{(2)}\end{pmatrix} + \beta\psi_{6}\varepsilon_{6t}\begin{pmatrix}c_{1}\\c_{2}\\c_{3}\\c_{4}\\c_{5}\end{pmatrix} -\beta\delta_{s6}\varepsilon_{6t}\begin{pmatrix}c_{1s}\\c_{3s}\\c_{21s}\\c_{23s}\end{pmatrix} + \beta\delta_{a6}\varepsilon_{6t}\begin{pmatrix}c_{1a}\\c_{3a}\\c_{21a}\\c_{22a}\\c_{23a}\end{pmatrix} - \beta\delta_{16}\varepsilon_{6t}\begin{pmatrix}\tilde{c}_{11}\\\tilde{c}_{21}\\\tilde{c}_{31}\\\tilde{c}_{41}\\\tilde{c}_{51}\end{pmatrix}, \rho\frac{\partial^{2}u_{2t}}{\partial t^{2}} = c_{16}\frac{\partial\varepsilon_{6t}}{\partial x} + c_{26}\frac{\partial\eta_{t}^{(1)}}{\partial x}.$$
(4.1)

Solving the system (4.1), we obtain the same wavenumber as found above

$$k_6 = \frac{\omega\sqrt{\rho}}{\sqrt{c_{66}^E(\alpha\omega)}}.$$
(4.2)

Using (4.2), we can find the sound wave velocity

$$v_{66}(\omega) = \frac{\omega}{\operatorname{Re}|k_6|} = \operatorname{Re}\frac{\sqrt{c_{66}^E(\alpha\omega)}}{\sqrt{\rho}}$$
(4.3)

and the contribution of the pseudospin subsystem to the sound attenuation

$$\alpha_6(\omega) = \alpha_{60} - \operatorname{Im}|k_6| = \alpha_{60} - \operatorname{Im}\left|\frac{\omega\sqrt{\rho}}{\sqrt{c_{66}^E(\alpha\omega)}}\right|,\tag{4.4}$$

where  $\alpha_{60}$  is a constant frequency and temperature independent contribution of the other mechanisms to the experimentally observable attenuation.

# 5. Discussion

Let us evaluate the above found dynamic characteristics of mechanically free  $K(H_{1-x}D_x)_2PO_4$ crystals, cut as  $l \times l$  square plates (l = 1 mm) in the [0,0,1] plane. It should be noted that the developed theory is valid, strictly speaking, for highly deuterated  $KD_2PO_4$  crystals, only. However, the tunneling in undeuterated crystals is weakened due to short-range interactions [26], which is indicated by the experimentally established relaxational character of  $\varepsilon_{33}^*(\nu, T)$  dispersion see [15,23– 25] in  $KH_2PO_4$ . Therefore, proton tunneling can be neglected as well as the obtained expressions used for undeuterated  $KH_2PO_4$ . In numerical calculations we shall use the values of the model parameters determined in describing the static and dynamic permittivity of a mechanical free crystal [36] and given in table 1.

Unfortunately, no experimental data are available to perform a quantitative comparison of the theoretically obtained temperature and frequency dependences of the dynamic characteristics of mechanically free crystals in the piezoelectric resonance region.

In figures 2 and 3 we plot the frequency dependences of the real and imaginary parts of the dynamic dielectric permittivity of mechanically free KH<sub>2</sub>PO<sub>4</sub>, KD<sub>2</sub>PO<sub>4</sub> crystals in the paraelectric phase at different temperatures  $\Delta T$ . For the MH<sub>2</sub>XO<sub>4</sub> crystals, a resonance dispersion takes place in the frequency range,  $3 \cdot 10^5 - 3 \cdot 10^8$  Hz, and for KD<sub>2</sub>PO<sub>4</sub> in the  $5 \cdot 10^5 - 5 \cdot 10^8$  Hz range. At  $\omega \to 0$ 

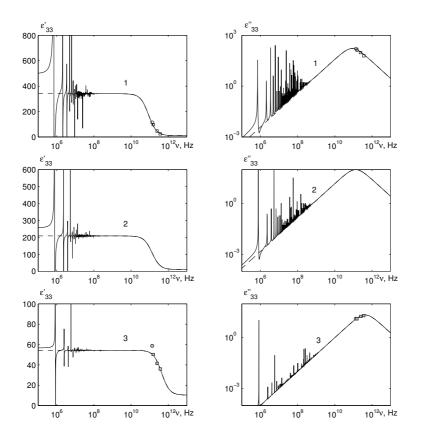
x	$T_{\rm c}$	$T_0$	$\frac{\varepsilon}{k_B}$	$\frac{w}{k_B}$	$\frac{\nu_3(0)}{k_B}$	$\mu_{3-}, 10^{-18}$	$\mu_{3+}, 10^{-18}$	$\chi^0_{33}$			
	(K)	(K)	$(\vec{K})$	(K)	$(\vec{K})$	$(esu \cdot cm)$	$(esu \cdot cm)$				
0.00	122.5	122.5	56.00	422.0	0 17.91	1.46	1.71	0.73			
0.81	205.6	204.8	85.82	781.5	5 33.44	1.76	2.02	0.42			
1.00	220.1	219.0	93.05	868.6	35.76	1.84	2.10	0.34			
x	$\frac{\psi_6}{k_B}$	$\frac{\delta_{s6}}{k_P}$	$\frac{\delta}{k}$	<u>a6</u> P	$\frac{\delta_{16}}{k_P}$	$c_{66}^0 \cdot 10^{-10}$	$e_{36}^{0}$	$P_{-}$	$R_{-}$	$P_+$	$R_+$
x	$ \begin{array}{c} \frac{\psi_6}{k_B} \\ (K) \end{array} $	$\frac{\frac{\delta_{s6}}{k_B}}{(K)}$		$\binom{a6}{B}{K}$	$\frac{\delta_{16}}{k_B}$ (K)	$\frac{c_{66}^0 \cdot 10^{-10}}{(dyn/cm^2)}$			$\begin{array}{c} R_{-} \\ \left(\frac{s}{K}\right) \end{array}$	$P_+$ $(s)$	$\begin{array}{c} R_+ \\ \left(\frac{s}{K}\right) \end{array}$
x 0.00		(K)	(1						$\left(\frac{s}{K}\right)$		
	(K)	$ \begin{array}{c c} (K) \\ \hline 0 82.0 \end{array} $	$\begin{array}{c c} & (1) \\ \hline 0 & -50 \end{array}$	K) 0.00	(K)	$\frac{(dyn/cm^2)}{7.10}$	$(esu/cm^2)$	(s) 0.35	$\left(\frac{s}{K}\right)$	(s)	$\left(\frac{s}{K}\right)$

**Table 1.** The optimal set of the model parameters for the  $K(H_{1-x}D_x)_2PO_4$  crystals.

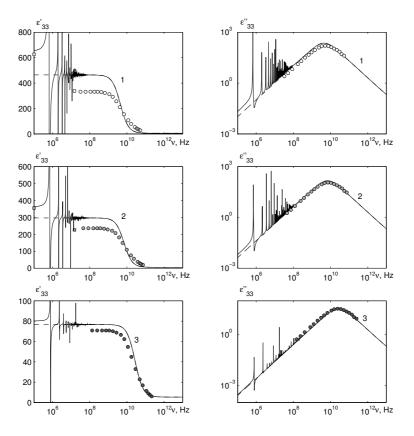
we obtain a static dielectric permittivity of a free crystal. Taking into account the dispersion (3.12), we find the equation for the resonance frequencies

$$\omega_n = \frac{\pi(2n+1)}{l} \sqrt{\frac{c_{66}^E}{\rho}},$$

where taking into account the fact that in the  $5 \cdot 10^5 - 5 \cdot 10^8$  Hz frequency range  $c_{66}^E(\omega)$  is practically frequency independent. The resonance frequencies are inversely proportional to sample dimensions.



**Figure 2.** Frequency dependences of the real and imaginary parts of the dynamic dielectric permittivity of a mechanically free and clamped KH<sub>2</sub>PO<sub>4</sub> crystal at different  $\Delta T$ , K: 1 – 5,  $\circ$  [38],  $\Box$  [39]; 2 – 10; 3 – 50  $\bullet$  [38],  $\Box$  [39].

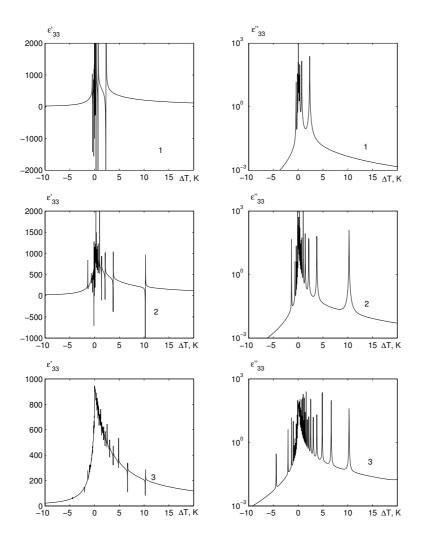


**Figure 3.** Frequency dependences of the real and imaginary parts of the dynamic dielectric permittivity of a mechanically free and clamped KD<sub>2</sub>PO<sub>4</sub> crystal at different  $\Delta T$ , K: 1 – 5, 2 – 10, 3 – 50.  $\circ$ ,  $\bullet$ ,  $\bullet$  [40].

The dashed lines in figures 2–3 describe the low-frequency permittivity of a clamped crystal. With increasing frequency and temperature  $\Delta T$ , the amplitudes of the resonance peaks decrease. With increasing temperature  $\Delta T$  the last resonance peak shifts to higher frequencies. Analogous multi-peak resonance dispersion is also observed in the ferroelectric phase. Above the resonance frequency, the crystal is clamped by the high-frequency field; above 10<sup>9</sup> Hz the clamped permittivity has a relaxational dispersion. The theoretical frequency curves  $\varepsilon'_{33}(\omega)$  and  $\varepsilon''_{33}(\omega)$  well accord with experimental data. The resonance dispersion in KH<sub>2</sub>PO<sub>4</sub> is schematically presented in [43] for the ferroelectric phase at  $10^4 - 10^6$  Hz.

In figure 4 we show the temperature curves of the real part  $\varepsilon'_{33}(\omega, T)$  of the free dielectric permittivity of KH<sub>2</sub>PO<sub>4</sub> at different frequencies. Below the frequency of the first resonance peak, the temperature variation of  $\varepsilon^*_{33}(\omega, T)$  essentially coincides with that of the static permittivity of a free crystal. Near the resonance frequencies, the sharp peaks in the temperature curve of permittivity appear, the number of which increases with an increase of frequency, whereas the magnitudes decrease. Upon further increase of frequency, numerous resonance peaks of small amplitude arise around the curve of clamped permittivity. At even higher frequencies the peaks disappear. The resonance peaks for the real and imaginary parts of the permittivity for a given frequency are observed at the same temperature  $\Delta T$ . The character of piezoelectric resonance curves is different for different crystals considered here.

In figure 5 we plot the calculated temperature dependences of the sound attenuation  $\alpha_6$  for KH<sub>2</sub>PO<sub>4</sub>, K(H<sub>0,195</sub>D<sub>0,805</sub>)<sub>2</sub>PO<sub>4</sub> at different frequencies along with the experimental points  $\alpha_6(T)$  taken from [41,42]. A good quantitative description of experimental data is obtained, especially



**Figure 4.** Temperature dependences of the real and imaginary parts of the dynamic dielectric permittivity of a mechanically free KH<sub>2</sub>PO<sub>4</sub> crystal at different frequencies  $\nu$ , MHz: 1 – 3, 2 – 10, 3 – 30.

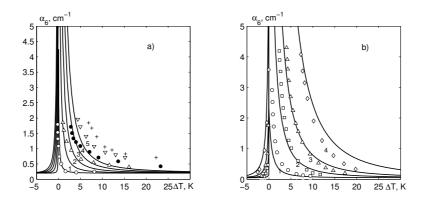
for  $K(H_{0.195}D_{0.805})_2PO_4$ .

Near the transition temperature  $T_c$ , a sharp increase of attenuation is obtained. In the ferroelectric phase, the attenuation decreases much faster when moving away from the transition than in the paraelectric phase.

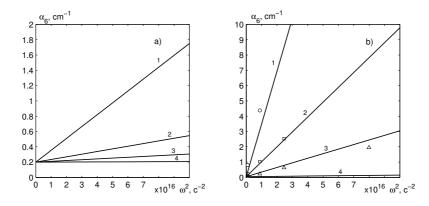
The dependence of attenuation  $\alpha_6$  on the square of frequency  $\omega^2$  for KH<sub>2</sub>PO<sub>4</sub>, K(H<sub>0,195</sub>D<sub>0,805</sub>)<sub>2</sub>PO<sub>4</sub> crystals at different temperatures  $\Delta T$  is shown in figure 6. As one can see, attenuation varies proportionally to the square of frequency. The closer the temperature is to the Curie point, the larger is the rate of this variation.

With increasing frequency, starting from ~  $10^7$  Hz, the theoretical attenuation sharply increases and then reaches a saturation at about  $10^{10}$  Hz (figure 7). In the paraelectric phase, the cut-off frequency decreases with approaching the transition temperature  $T_c$ . The cut-off frequency also decreases with increasing deuteration x. Such high values of attenuation at saturation, in fact, mean the absence of sound propagation.

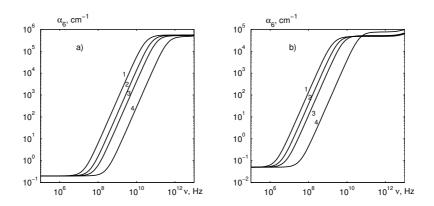
In figure 8 we depicted the calculated temperature dependence of sound velocity  $v_{66}$  for KH<sub>2</sub>PO<sub>4</sub>, K(H<sub>0,195</sub>D<sub>0,805</sub>)<sub>2</sub>PO<sub>4</sub>.



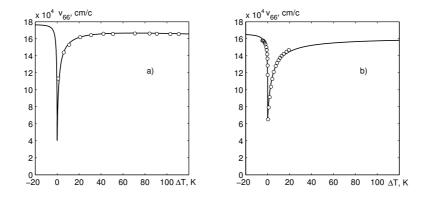
**Figure 5.** Temperature dependences of sound attenuation for KH<sub>2</sub>PO<sub>4</sub> (a), at different frequencies  $\nu$ , 10<sup>6</sup> Hz: 1 - 10,  $\circ$  [41], 2 - 30,  $\diamond$  [41], 3 - 50,  $\bullet$  [41], 4 - 70,  $\nabla$  [41], 5 - 90,+[41] and K(H<sub>0.195</sub>D<sub>0.805</sub>)<sub>2</sub>PO<sub>4</sub> (b) at  $\nu$ , 10<sup>6</sup> Hz: 1 - 5,  $\circ$  [42], 2 - 15,  $\Box$  [42], 3 - 25,  $\diamond$  [42], 4 - 45,  $\diamond$  [42].



**Figure 6.** Frequency dependences of attenuation for KH<sub>2</sub>PO<sub>4</sub> (a),  $K(H_{0.195}D_{0.805})_2PO_4$  (b) at different temperatures  $\Delta T$ ,K: 1 – 2, 2 – 5, 3 – 10, 4 – 50.  $\circ$ ,  $\Box$ ,  $\Delta$  – [42].



**Figure 7.** Frequency dependences of attenuation for  $KH_2PO_4$  (a),  $K(H_{0.195}D_{0.805})_2PO_4$  (b) at different temperatures  $\Delta T$ , K: 1 – 2, 2 – 5, 3 – 10, 4 – 50.



**Figure 8.** Temperature dependences of sound velocity for  $KH_2PO_4$  (a),  $K(H_{0.195}D_{0.805})_2PO_4$  (b).

It has a minimum at  $T = T_c$ . Below 10<sup>10</sup> Hz, the magnitude of  $c_{66}^E(\omega)$  is frequency independent; therefore, we calculated the velocity using (8.3) and experimental data for  $c_{66}^{TE}$  ([44] for KH<sub>2</sub>PO<sub>4</sub> and [42] for K(H<sub>0.195</sub>D<sub>0.805</sub>)<sub>2</sub>PO<sub>4</sub>) and crystal density  $\rho$ . The results are shown in figure 8 by symbols  $\circ$ .

At the frequency of microwave dispersion of permittivity, a sharp increase of sound velocity  $v_{66}$  should be observed, after which the frequency curve of velocity saturates at about ~  $10^{11}$  Hz. The saturation values of sound velocity are temperature independent.

## 6. Conclusions

Within the proton ordering model with taking into account the shear strain  $\varepsilon_6$  we explored a dynamic response of the KD<sub>2</sub>PO<sub>4</sub> type crystals to an external harmonic electric field  $E_3$ . Dynamics of the pseudospin subsystem is described within the stochastic Glauber approach. Dynamics of the strain  $\varepsilon_6$  is obtained from the Newtonian equations of motion of an elementary volume, with taking into account the relations between the order parameter of the pseudospin subsystem and the strain in the static limit. Expressions for the longitudinal dynamic dielectric permittivity of KD<sub>2</sub>PO<sub>4</sub> crystals are obtained. Evolution of dynamic permittivity with increasing frequency via the piezoelectric resonances  $(10^5-10^8 \text{ Hz})$  to the clamped crystal with the microwave  $(10^9-10^{12} \text{ Hz})$  relaxational dispersion is obtained. Temperature and frequency dependences of sound attenuation  $\alpha_6$  and velocity  $v_{66}$  in  $K(H_{1-x}D_x)_2PO_4$  are obtained. An anomalous increase of attenuation in the phase transition region and the temperature curve of sound velocity are described. A presence of cut-off frequency for sound propagation (an anomalous increase of attenuation at frequencies corresponding to the microwave dispersion of the dielectric permittivity) is predicted.

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# Поздовжня релаксація механічно вільних кристалів типу КН<sub>2</sub>РО<sub>4</sub>. П'єзоелектричний резонанс та поглинання звуку

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В рамках протонної моделі з врахуванням п'єзоелектричної взаємодії зі зсувною деформацією  $\varepsilon_6$  розглянуто динамічний діелектричний відгук сегнетоелектриків типу KD<sub>2</sub>PO<sub>4</sub>. Враховано динаміку п'єзоелектричної деформації. Явно описано явища затискання кристалу високочастотним електричним полем, п'єзоелектричного резонансу і НВЧ дисперсії, що спостерігаються на експерименті. Розраховано коефіцієнт поглинання звуку. Описано особливості коефіцієнта поглинання в околі точок переходу. Передбачено наявність обрізаючої частоти у частотній залежності коефіцієнта поглинання звуку.

Ключові слова: сегнетоелектрики, п'єзоелектричний резонанс

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