value is smaller by a factor of about 3 to 4 than those values given by Blinc et al. [7, 8] and Novaković [6]. It is, however, almost the value $\Omega = 1.0 \times 10^{-14}$ erg which has recently been determined by Cochran [10] from Raman-scattering

data of Kaminow and Damen [14].

Since the values of S_1 for KH₂AsO₄, KH₂PO₄, and RbH₂PO₄ differ only slightly (cf. Table 1), we used for KH₂AsO₄ and RbH₂PO₄, too, the value a $S_1 = 9.4 \times \times 10^{-3} \, \mathrm{kbar^{-1}}$ determined for KD₂PO₄. In this way, we found $\Omega/k = 43 \, ^{\circ}\mathrm{K}$ for KH₂AsO₄, and $\Omega/k = 112 \, ^{\circ}\mathrm{K}$ for RbH₂PO₄. In this estimate we assumed the same value for ζ as for KH₂PO₄, because no experimental data for ζ are available for KH₂AsO₄ and RbH₂PO₄. However, this assumption is not so important because there is only a weak dependence of the estimated values of Ω on ζ . This is shown in Fig. 5 where for the parameter value $\Omega/k = 75 \, ^{\circ}\mathrm{K}$ also curves with $\zeta = 0.15$ and 0.25 Å are given as dashed lines.

If instead of the dielectric data for the shift of T_c with pressure of KH₂PO₄ and KD₂PO₄ the neutron diffraction data of Umebayashi et al. [1], $dT_c/dp = -4.5 \text{ deg/kbar}$ and $dT_{c,D}/dp = -2.6 \times 10^{-3} \text{ deg/kbar}$, are used, the same method results in $\alpha S_1 = 6.1 \times 10^{-3} \text{ kbar}^{-1}$ and $\Omega/k = 93$ °K. The Ω -value does not differ essentially from that obtained from dielectric data. For reasons of comparison, data from dielectric-constant measurements have only been

taken in Fig. 5.

From (6), with $q^2 \sim m^{1/2} \, \zeta^2$, the ratio of the tunneling energies follows: $\Omega_{\rm D}/\Omega = 2^{-1/4} \, (\zeta_{\rm D}/\zeta) \exp \left\{-q^2 \, [\sqrt{2} \, (\zeta_{\rm D}/\zeta)^2 - 1]\right\}$, the quantities of the deuterated crystal having the index D. Assuming $\zeta_{\rm D}/\zeta = 1.0$ to 1.1, for all three substances $\Omega_{\rm D}/\Omega < 0.2$ and $\Omega_{\rm D}/kT_{\rm c,\,D} < 0.1$ result. This justifies our neglect of the influence of tunneling on the shift of $T_{\rm c}$ for the deuterated crystals as assumed above. For these crystals, therefore, the linear relation ${\rm d}T_{\rm c,\,D}/{\rm d}p \approx -0.02 \, T_{\rm c,\,D} \, {\rm kbar^{-1}}$ is expected to hold.

Contrary to our determination of the value of α from experimental data Novaković [6], and Bline and Žekš [7] determined α by a-priori assumptions which, however, resulted in very different values for α . Novaković puts $\mathrm{d}\zeta/\mathrm{d}a = \zeta/a$, i.e. $\alpha = 1$. Bline and Žekš assume that with compression the O-H···O bonds (two per lattice constant) are shortened only and that within these bonds the distance 2ζ between the potential minima is reduced only, i.e. $\mathrm{d}\zeta = \mathrm{d}a/4$ or $\alpha = 9.5$. This value is approximately the same we used ($\alpha = 7.8$ to 8.3). According to the semi-empirical model for the O-H···O bonds of Lippincott and Schroeder [15] one might expect $\mathrm{d}\zeta/\mathrm{d}a \approx 0.29$ or $\alpha \approx 11$. A direct experimental determination of α (by neutron-diffraction measurements under pressure) would be of interest because the a-priori choice of α is affected with a considerable uncertainty.

Having determined $\Omega/kT_{\rm e},~\Omega/J$ is directly obtained from (2). For the deuterated crystals, (2) simplifies to $J_{\rm D}\approx 4\,kT_{\rm c,\,D}$. In Table 1, values of $4\,\Omega/J$ and $J_{\rm D}/J$ are also listed. Obviously, the ferroelectric interaction in the deuterated crystals is stronger; this fact corresponds qualitatively to the theoretical expectations

Within the range of the applied pressures up to 1.2 kbar, no deviation from the linear dependence between $T_{\rm c}$ and p was observed. From the above mentioned dependence of the values Ω and J on ζ , according to (2), we have to expect, however, that due to tunneling, at higher pressures the transition temperature $T_{\rm c}$ decreases more rapidly, and ferroelectricity disappears com-