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Dielectric properties of Betaine Arsenate crystal near the phase transition

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Dielectric measurements for single crystal of betaine arsenate $(CH_3)_3NCH_2COO \cdot H_3AsO_4$ connected with the ferroelectric phase transition at 119 K were performed. The temperature dependence of electric permittivity was measured at dc electric fields up to 700 kV/m . The results show significant suppression of the dielectric constant by the application of dc field. Deviation from the classical behavior was observed. The electric permittivity was also measured in the paraelectric phase at constant temperature as a function of electric field intensity up to 700 kV/m . The electric permittivity might be well described by the classical relation with additional term including contribution to permittivity coming from clusters. The fit parameters indicate that the polar-clusters carries polarization $P_0 = 0.7 - 4.9 \cdot 10^{-3}\text{ C m}^2$ with the clusters size of $L = 12 - 20\text{ nm}$.

1. Introduction

Ferroelectric crystal of betaine arsenate $((CH_3)_3NCH_2COO \cdot H_3AsO_4)$ abbreviated as BA) has been known since 1982 [1]. According to this first paper the crystal undergoes two phase transitions, one at $T_{C1} = 411\text{ K}$ and another one at $T_{C2} = 119\text{ K}$. The first phase transition was identified as a para-ferroelastic phase transition, while the second was identified as a para-ferroelectric one [2,3]. The high frequency dielectric measurements and Raman spectroscopy indicate that the phase transition at T_{C2} appears to be of the order-disorder type [4,5]. The shape of the heat capacity curve measured near T_{C2} by adiabatic calorimeter [6] suggests that the phase transition is of the second order. The phase transition at T_{C1} seems to be of the first order [7]. The symmetry of the BA crystal above 411 K is $Pbnm$ [8] and in the range of $119\text{ K} < T < 411\text{ K}$ is $P121/n1$ [2]. There are some evidence for a third phase transition occurring at the temperature about 110 K [9,10]. The latest X-Ray studies suggested the appearance of the third phase transition at the temperature 107 K and the space group $P1n1$ was proposed for temperature below 119 K [10]. Most of the experimental investigations of BA show deviations from what is considered as a typical ferroelectric behaviour. The pyroelectric coefficient along the $[001]$ direction reveals two anomalies in the temperature range $110 - 123\text{ K}$ [11]. The hysteresis loops in BA exhibit striking S-shaped curves in the interval of a few K above T_{C2} [1]. Using a model of weakly coupled one-dimensional Ising chains, the temperature dependence of dielectric constant in paraelectric phase was described as referred by Müser [12]. The same authors observed a cross-over of effective critical exponent γ to a value

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typical for short-range forces in paraelectric phase [13]. Recent measurements also show that in BA exists short range interaction above $119K$ [5,10], which are called clusters.

The aim of our studies was to measure and describe dielectric properties of BA under influence of electric field. The description was aimed to check the possibility to use dielectric state equation with two or three terms and with additional term coming from clusters to get the best fitting of parameters, which describe phase transition.

2. Experimental

Polycrystals of BA were obtained from water solution of stoichiometric quantities of arsenic acid and betaine. Single crystals BA were grown from saturated water solution of dissolved polycrystals of the substance by slow evaporation method at constant temperature of $302K$.

The c-plates approximately 0.7 mm in thickness were cut out from the crystal. Silver paste films were used as electrodes. The sample was placed in a nitrogen gas exchange chamber of a cryostat and it was mechanically free. The sample capacitance was measured using a precision LCR-meter HP 4284A with a measured field of 1 V/cm in amplitude 1.2 kHz , allowing us to calculate the electric permittivity. The electric permittivity as a function of temperature was measured on a cooling run with a rate of 0.5 K/min with different values of dc electric field from 0 to 700 kV/m . Applying an electric field of amplitude 700 kV/m and the frequency 10^{-3} Hz , the field dependence of the electric permittivity at constant temperatures was measured. The temperature in the chamber was controlled with an accuracy not worse than $5 \cdot 10^{-3}\text{ K}$.

3. Results and Discussion

The dielectric constant and its reciprocal value are presented as a function of temperature in figure 1,2. In the paraelectric phase, the Curie-Weiss law :

$$\varepsilon' = \frac{C_0}{T - T_0}, \quad (1)$$

where C_0 and T_0 are Curie-Weiss constant and temperature, respectively. Equation (1) is well fulfilled in the temperature range of $120\text{-}150\text{K}$ (see dotted line in the figure 2) with $C_{01} = 3.9 \pm 0.01 \cdot 10^4\text{K}$. However, this fit gives Curie-Weiss temperature different than the one in which maximum of dielectric constant appears. Another fit was done in the temperature range of $154 - 190\text{ K}$ (see solid line in the figure 2) and gave practically the same Curie-Weiss temperature as the temperature of the maximum of dielectric constant. The Curie-Weiss constant is equal to $C_{02} = 3.6 \pm 0.01 \cdot 10^4\text{K}$. However, below temperature about 154K experimental data show deviation from this fit.

Figure 1. Temperature dependence of dielectric constant measured in c-axis direction.

Figure 2. Temperature dependence of the reciprocal value of the dielectric constant.

3.1. Temperature dependence of electric permittivity in the paraelectric phase

Figure 3 shows temperature variations of the dielectric constant for various dc electric fields. As it can be seen the maximum value of ϵ' decreases with intensity of applied electric field and the position of maximum shifts to higher temperatures. It looks like classical behaviour.

Figure 3. Temperature variation of the dielectric constant at various dc electric field.

However, when we take equation for maximum value of permittivity obtained from Landau theory [14]:

$$(\epsilon_{\max})^{-1} = \frac{3}{2^{1/3}} B^{1/3} E^{1/a}, \quad (2)$$

where B is nonlinear coefficient in dielectric state equation, we do not get classical value of the exponent $a = 1.5$. The experimental data are well described with the value of the exponent $a = 1.66$. The results are presented in figure 4.

Figure 4. Maximum value of electric permittivity versus dc electric field for different temperatures.

3.2. Field dependence of permittivity at constant temperatures in the paraelectric phase

The further study was focused on the dc electric field dependence of dielectric constant at constant temperature in the paraelectric phase. The measurements were done in the range of temperature (T_{C2} , $T_{C2}+30K$). In order to describe experimental data we started with the well-known classical dielectric state equation in a form:

$$E = AP + BP^3 + CP^5, \quad (3)$$

where $A = A_0(T - T_C)$ is linear coefficient and B, C are nonlinear one, which are assumed to be temperature independent. P is an induced polarization.

Neglecting the last term of equation (3) and using relation:

$$(\epsilon_0 \epsilon')^{-1} = \frac{\partial E}{\partial P} \quad (4)$$

one can rewrite (4) in the form:

$$\epsilon' = 1/(\epsilon_0(A + 3BP^2)), \quad (5)$$

where the explicit relation between P and E

$$P = \left[\frac{E}{2B} + \sqrt{\left(\frac{A}{3B}\right)^3 + \left(\frac{E}{2B}\right)^2} \right]^{1/3} - \left[-\frac{E}{2B} + \sqrt{\left(\frac{A}{3B}\right)^3 + \left(\frac{E}{2B}\right)^2} \right]^{1/3} \quad (6)$$

was obtained analytically from equation (3) with only two first terms.

On the other hand the field dependence of dielectric constant in the paraelectric state can be expressed as [15,16]

$$\epsilon' = \frac{1}{\epsilon_0 A} - \frac{3B}{\epsilon_0 A^4} E^2 + \frac{15B^2}{\epsilon_0 A^7} E^4 \quad (7)$$

This equation is commonly used to describe the behaviour of dielectric constant in dc field at constant temperature. However, in this work, fits of ϵ' vs. E involving even powers of E^4 are insufficient to explain experimental data as it can be seen in figure 5.

Figure 5. Field dependence of the dielectric constant and its best fit to the equation (7) (solid line).

Therefore, in next step the fits were done according to equation (5) and (6) as it was propose for some systems by Dec et al. [17]. Plot in figure 6 (see the dotted line) shows that fitting to equation (5) is in better agreement with data for temperature higher than 131K. From these fits the parameters A and B were obtained in the temperature range 132 – 152K. However, below 131K fitting curves are in disagreement with experimental data as it can be seen in figure 7.

Additionally, to explain experimental data we take into account another term (E^6) in equation (7). We performed fitting with extended equation and result for $T = 138.5K$ is shown in figure 6 (see the solid line). Plots in the figure 8 show temperature dependence of fitting parameters A and B . One might notice that in this case parameters A and B were in quite good agreement with the results from the equation (4). Nevertheless at the temperature below 126K, adding next term was insufficient to explain the semi-bell shape curves (as an example see figure 9).

Following results from [5,10] we considered the appearance of clusters and according to method proposed by Bianchi et al. [18,19] and used by others authors [20], we added a cluster term to equation (5). By means of Langevin-type approach[21], the total polarization of a cluster system is expressed as

$$P = P_0 \tanh \left(\frac{P_0 L^3 E}{2k_B T} \right). \quad (8)$$

Figure 6. Field dependence of the dielectric constant and its best fit:

- solid line according to the equation (5)
- stars according to the equation (7) with additional term E^6 .

here P_0 denotes effective polarization of clusters, L the size of cluster, k_B the Boltzmann's constant, and E electric field. Therefore contribution to electric permittivity from clusters takes a form:

$$\varepsilon'_K = \frac{P_0^2 L^3}{\varepsilon_0 2k_B T} \left[\cosh \left(E \frac{P_0 L^3}{2k_B T} \right) \right]^{-2}. \quad (9)$$

It can be seen in figure 10 that term represented as equation (9) added to equation (5) improves fits and describes experimental data in temperature range from 119 to 153K. It is interesting to notice that the contribution to electric permittivity from polar clusters at the temperature 121K is about 27% without dc field (see an example in figure 11) and at the temperature about 130K it is 18%. The contribution from clusters decreases with temperature.

In figure 12 we showed parameters A and B as functions of temperature. The parameter B varies with temperature from $0.5 \cdot 10^{11}$ for higher temperature and rises up to $12 \cdot 10^{11} m^5 V C^{-3}$ for temperature near the temperature of the phase transition. This increase of B can be connected with existence of the clusters. The parameter A_0 was found from temperature dependences of parameters A . We found that the value of A_0 is equal to $2.1 \cdot 10^6 V m (CK)^{-1}$, it has the same order as the one calculated from Curie-Weiss constant ($A_0 = 2.8 \cdot 10^6 V m (CK)^{-1}$). The lower value of A_0 can be connected with the relative volume of cluster, which is comparable with the contribution from clusters to total electric permittivity under zero electric field. In next step according to parameters from formula, which was used for fitting $\varepsilon(E)$ and was obtained by summing equations (4) and (9) we got the clusters size, and polar clusters carried polarization. The following

Figure 7. Field dependence of the dielectric constant and its best fit to the equation (5). Dotted line shows fit for an entire range of field, while solid line shows fit for values of field up to $4 \cdot 10^5 \text{ V/m}$.

part of equation (9) is connected with clusters. We can rewrite equation (9) in the form:

$$\epsilon'_K = P_1 [\cosh(P_2 \cdot E)]^{-2}, \quad (10)$$

where parameters P_1 and P_2 are:

$$P_1 = \frac{P_0^2 L^3}{\epsilon_0 2 k_B T}, \quad (11)$$

$$P_2 = \frac{P_0 L^3}{2 k_B T}. \quad (12)$$

Since ϵ_0 , k_B and T are constant the value of variable L and P_0 can be calculated from P_1 and P_2 . We got the clusters size $L = 12 - 20 \text{ nm}$, and polar clusters carried polarization $P_0 = 0.7 - 4.9 \cdot 10^{-3} \text{ Cm}^2$.

Figure 8. Temperature dependence of fitting parameters A and B . In both figures plot with squares denotes dependences according equation (5), while plot with circles denotes dependences according to equation with term E^6 .

Figure 9. Field dependence of the dielectric constant and its best fit to the equation with E^6 term (solid line).

Figure 10. The dc electric field dependence of dielectric constant for two values of temperature. Solid lines represent fits to equation with cluster term.

Figure 11. The dc electric field dependence of dielectric constant. Plot labeled as 1 is the contribution from the polar cluster term (according equation (9)), and plot labeled as 2 is the contribution from the conventional polarization term (according equation (5)).

Figure 12. Temperature dependence of parameters A and B of equation with cluster's term.

3.3. Polarization in the paraelectric phase

The polarization was measured using an electrometer at frequency of $8.3 \cdot 10^{-3} \text{ Hz}$. Klöpperpier et al. observed that the hysteresis loops in BA exhibit striking S-shaped curves in the interval of a few K above T_{C2} [1]. In our analysis hysteresis were observed in paraelectric phase even up to temperature $167 K$. The correction of polarization obtained by subtracting the part connected with resistance of the sample gives disappearance of hysteresis above $150 K$ (see an example figure 13). However, for temperature below $150 K$ this method was insufficient to remove hysteresis loops. Character of hysteresis loops after correction is as for ferroelectric one (see figure 13) and one can conclude that presence of these loops confirmed the existence of clusters below $150 K$.

Figure 13. Polarization after correction versus dc electric field.

4. Summary

The results obtained in the present study are summarized as follows:

- From relation for $\varepsilon'_{max}(E)$ we found exponent $a = 1.66$, which is different than classical one, so the classical equation can not be used in the temperature below $130K$,
- $\varepsilon'(E)$ was described using classical state equation with the term P^3 in the range of temperature $132 - 152K$,
- $\varepsilon'(E)$ was described using classical relation with term E^6 in the temperature range $126 - 152K$,
- $\varepsilon'(E)$ was described by relation with clusters' term added to classical equation in the temperature range $121.5 - 132K$,
- The existence of clusters with $L = 12 - 20nm$ and $P_0 = 0.7 - 4.9 \cdot 10^{-3} Cm^2$ was confirmed in BA crystal,
- The parameters A and B were obtained by few methods in the broad temperature range; A_0 is equal to $2.1 \cdot 10^6 Vm(CK)^{-1}$ and the parameters B varies with temperature from $0.5 \cdot 10^{11}$ for higher temperature and rises up to $12 \cdot 10^{11} m^5 VC^{-3}$ for temperature near the temperature of the phase transition.

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