On the Electrical Properties of AgNO₃, TlNO₃, and NH₄NO₃

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Values of the dielectric constant and resistivity measured at 1 kHz are presented for polycrystalline AgNO₃, TlNO₃, and NH₄NO₃ below room temperature. Results are also given for AgNO₃ and TlNO₃ in hitherto unexplored regions above room temperature. The data show dielectric transitions in AgNO₃, TlNO₃, and NH₄NO₃, with the critical temperatures −35, −35, and −32°C, respectively. Previously published values of the heat capacity of TlNO₃ are examined, and shown to contain an anomaly at ~−35°C which appears to be associated with the dielectric transition there. Consideration of the present and other results for NH₄NO₃ leads to the conclusion that a previously detected transition in the dielectric constant at ~−32°C on cooling is of the same character as those found at comparable temperatures in other univalent nitrates, rather than a consequence of the IV→V structural transition. A maximum in the dielectric constant of AgNO₃ is found at 110°C.

Two structural phases have been identified hitherto in solid AgNO₃ at atmospheric pressure. On heating samples from room temperature, the calorimetrically determined phase II→I transition temperature is reported as 158.9 to 160.6°C, in good agreement with that given by Bridgman,²,³ who identified four structural modifications at pressures of up to 50 kbar. The phase diagram for AgNO₃ is of quite a different form from that for TlNO₃, whose properties closely resemble those of RbNO₃ and CsNO₃, however. Measurements of specific heat from 15 to 300 K do not show any anomalies which could indicate additional phase transformations at atmospheric pressure.

The melting point of carefully dried AgNO₃ is reported to be 209.6°C.⁴ Appreciable decomposition of the melt has been reported for temperatures in the range 305 to 450°C,⁵,⁶ although the density and electrical conductivity were found to be reversible for temperature cycles with a maximum temperature of 320°C.⁷ X-Ray measurements on the solid were, however, affected by decomposition of phase I crystals.⁸

Phase I AgNO₃ has trigonal symmetry,⁹ whereas phase II is orthorhombic.¹⁰,¹¹ This phase is weakly birefringent, and has rather high values of refractive indices compared, for example, with those for KNO₃ in the orthorhombic phase II.¹²

The large refractive indices of AgNO₃ are mainly due to the large electronic polarizability of Ag⁺ compared with that for K⁺, the values of this parameter having been given as 2.4 and 1.33 Å³, respectively.¹³ This, together with the
distinctive phase diagram, suggests that the lattice bonding and hence also the electrical properties of AgNO₃ might differ considerably from those of the alkali metal nitrates. In order to test this point, it is of interest to compare the frequencies of a lattice mode of vibration for the various compounds. The frequencies of strong absorption peaks of this part of the IR spectrum are shown as a function of the cation radius for LiNO₃, NaNO₃, AgNO₃, KNO₃, TlNO₃, and NH₄NO₃ in Fig. 1. It is seen that the frequency varies regularly

![Fig. 1. A lattice mode frequency \( \nu \) for univalent nitrates at room temperature versus cation radius \( r_c \). The value of \( \nu \) for NH₄NO₃ is from Landolt-Börnstein and Newman and Halford; \( \nu \) for TlNO₃, Newman and Halford; the remaining data are taken from Ferraro and Walker.](image)

with cation radius, and therefore with the anion-cation separation. Thus it seems from this dynamic property of the lattice that the interionic bonding is of a uniform nature in the compounds, and it is to be expected that the electrical among other ionic properties of AgNO₃ will conform with those of the alkali metal nitrates.

According to dilatometric measurements, the volume expansion coefficient of AgNO₃ drops sharply at the II→I transition or a few degrees above. Sparse d.c. two-terminal resistivity data for \( t > 140^\circ C \) indicate a sharp minimum at the I→II transition. Measurements of a.c. conductivity have been obtained in the vicinity of the melting point. The dielectric constant \( \epsilon \) has been given as 9.0 at 20°C for a measuring frequency of 3 MHz.

By means of optical microscopy, it was shown that TlNO₃ adopts three modifications between room temperature and the melting point, the structural symmetries in order of increasing temperature being orthorhombic, rhombohedral, and cubic. The cubic phase I was confirmed to be optically isotropic, the melting point being 210.4°C.

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Differential thermal analysis has showed the III→II transition temperature to be 79°C, and II→I at 144°C, the II→III transition being diffuse. A calorimetric investigation gave the I→II transition temperature range as 142.5 to 143.8°C, compared with the value 144.6°C given by Bridgman. The latter author found the II→III transition to occur at 75°C and recorded no additional phases at pressures of up to 12 kbar.

The orthorhombic, room temperature phase III has been studied by several authors using X-ray diffraction. From measurements of the anisotropic IR absorption spectra for this phase, the directions of the two-fold axes of the nitrate groups, which lie in planes equidistant from those through the metal atoms, were deduced to bisect the angle between the a and b axes; the molecular planes being tilted about these directions.

A large polarizability of Tl⁺ is reflected in both the refractive index and the dielectric constant at room temperature (16.5), but as in the case of AgNO₃, this does not appear to influence the IR response, as may be seen in Fig. 1.

A determination of the electrical conductivity at 900 Hz and 30 kHz did not show the III→II transition, but the II→I transition produced an overall increase of ~1.5 orders of magnitude in σ. Later measurements showed a decrease in σ by a factor of 200 at III→II, probably due to a partial disintegration of the sample, which may have included water. An increase in σ of two orders of magnitude was noted at II→I.

The values of the specific heat C_p for TINO₃ according to two calorimetric investigations are shown in Fig. 2, where there is seen to be a small change in slope at 238 ± 5 K. Apparently undetected hitherto, this effect seems to exceed the experimental error; the two sets of data agreeing well in the region.

Fig. 2. The specific heat c_p of TINO₃ as a function of the absolute temperature T, showing a discontinuity of slope at 238 ± 5 K. The experimental points are from Latimer and Ahlberg, and the section A→B of the curve is drawn according to the equation given by Satoh.

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of overlap. The anomaly may indicate a threshold in the bonding or disorder properties of the ions, and will be shown to coincide with a dielectric transition.

By means of optical spectroscopic and dilatometric observations, it was found that orthorhombic, room temperature phase IV crystals of \( \text{NH}_4\text{NO}_3 \) transform at \( \sim -16^\circ \text{C} \) into a tetragonal low temperature phase \( \text{V}_7 \) of lesser birefringence.\(^{31}\) Samples were also found to be optically uniaxial at \(-50^\circ \text{C}\). By using a dilatometer, the same author found that on increasing the temperature, the expansion coefficient diminished at \(-16^\circ \text{C} \), and was followed by a region of contraction over the range \(-14 \) to \(-8^\circ \text{C} \), above which the crystals expanded normally. Later dilatometric experiments yielded the \( \text{V} \rightarrow \text{IV} \) transition temperature of \(-18^\circ \text{C} \).\(^{32}\) In common with several other univalent nitrate phases, the anions lie in parallel planes, in the present case parallel to \( (010) \).\(^{34,35}\)

Although X-ray diffraction measurements indicated the symmetry of phase \( \text{V} \) to be hexagonal or pseudohexagonal, with only expansion differences between the unit cell dimensions at \(-78, -33, \) and \(-18^\circ \text{C} \),\(^{36}\) the earlier suggestion of tetragonal symmetry,\(^{31}\) has been confirmed at \(-50^\circ \text{C}.\)\(^{37,38}\)

Specific heat measurements showed a sharp peak in value at \(-60.4^\circ \text{C}\);\(^{39}\) and this anomaly was later noted in addition to others at \(-160 \) and \(-18^\circ \text{C}\).\(^{40}\) The transformations at \(-160 \) and \(-60.4^\circ \text{C} \) were absent from other calorimetric investigations, however.\(^{41,42}\) The temperatures reported for the \( \text{V} \rightarrow \text{IV} \) and \( \text{IV} \rightarrow \text{III} \) transitions being \(-16.9 \) and \(-32.2^\circ \text{C} \).

Measurements of dielectric constant at 100 kHz for a powdered sample showed an increase in value at \(-33^\circ \text{C} \) on cooling, and hysteresis to \(-6^\circ \text{C} \) on heating.\(^{43}\) A two-terminal conductivity measurement for \( t > -25^\circ \text{C} \), at a sample voltage of 200 V, showed an anomaly at \( \sim -6^\circ \text{C} \) on heating.\(^{57}\)

Observations of the electron paramagnetic resonance (EPR) of VO\(^{2+} \) included in the \( \text{NH}_4\text{NO}_3 \) lattice have shown that a rapid readjustment of the ion which is present at \( 25^\circ \text{C} \) is totally absent at \(-70^\circ \text{C} \). The gradual onset of the motion appears to occur over the range \(-30 \) to \(-20^\circ \text{C}.\)\(^{44}\)

An ideal study of the low frequency (or non-optical) properties of a compound would probably consist of measurements of the temperature dependence of the four-terminal conductivity and dielectric constant, virtually from d.c. to a practical upper limit in the radio frequency range. The investigation would also provide data on polarization barrier formation and the influence of substitutional impurities in single crystals; together with the detection of any non-linearity of displacement current. The effect of pressure upon the electrical parameters would also be determined. Actual investigations usually only fulfil part of these requirements, but nevertheless provide a useful addition to the information gained using other, and often more sophisticated techniques.

The present work was aimed primarily at discovering whether electrical anomalies of the type found in the alkali metal nitrates\(^{45-49}\) occur also in \( \text{AgNO}_3, \text{TiNO}_3, \) and \( \text{NH}_4\text{NO}_3 \). Satisfactory techniques for the measurement of four-terminal resistivity and dielectric constant below room temperature in these compounds are still to be fully developed, and the series of measurements is therefore restricted to a.c. two-terminal parameters for polycrystalline samples. The measurements were extended above room temperature for \( \text{AgNO}_3 \) and \( \text{TiNO}_3 \).

Of the three compounds measured, \( \text{NH}_4\text{NO}_3 \) has been most thoroughly studied below room temperature, and it is the only univalent nitrate known...
to have a structural phase transition in this region. It is not possible to reconcile all of the reported physical properties of the compound, however; and some extra complexity is to be expected when both ions are non-spherical.

EXPERIMENTAL

The p.a. grade compounds were finely divided by crushing, and dried under vacuum; AgNO₃ (E. Merck A.G.) and TlNO₃ (Fluka A.G.) at \( \sim 120^\circ \text{C} \) for 24 h, NH₄NO₃ (E. Merck A.G.) at 70°C for 1 h. Laminar polycrystalline samples were then formed by cooling the melt between silver electrode plates of \( \sim 10 \text{ cm}^2 \) area, spacers of quartz or Pyrex being used to obtain separations of \( \sim 0.3 \text{ mm} \).

The apparatus used to control the sample temperature and measure dielectric constant and dissipation at 1 kHz were as described previously.⁶³,⁶⁴

![Fig. 3. Dielectric constant \( \varepsilon \) and resistivity \( \sigma_D \) for AgNO₃ as functions of temperature. In this and the following diagrams open circles denote values obtained with decreasing temperature; filled circles with increasing temperature.](image)

![Fig. 4. Dielectric constant \( \varepsilon \) and resistivity \( \sigma_D \) for TlNO₃ versus temperature.](image)

RESULTS AND DISCUSSION

Values of the dielectric constant \( \varepsilon \) and resistivity \( \sigma_D \) at 1 kHz for AgNO₃, TlNO₃, and NH₄NO₃ below room temperature are shown in Figs. 3 to 5, and results obtained above room temperature for AgNO₃ and TlNO₃ in Figs. 6 and 7.
Fig. 5. Dielectric constant \( \varepsilon \) and resistivity \( \rho_B \) for \( \text{NH}_2\text{NO}_3 \) as functions of temperature.

Fig. 6. Dielectric constant \( \varepsilon \) and resistivity \( \rho_B \) for \( \text{AgNO}_3 \) versus temperature.

The form of the samples yielding these data is both convenient and suitable when the temperature variation of parameters rather than their absolute values are of primary interest. It is clear, however, that when a compound is allowed to freeze in contact with metal electrode plates, there is little control over the growth of the crystallites, which is governed by such factors as rate of cooling, degree of purity of the sample, condition of the electrode surface, etc. With anisotropic compounds the samples may include preferred directions of crystal growth, which are not easily ascertained. It may also be difficult to measure the sample geometry with high precision. Variations in absolute value from sample to sample have been found, and the results are therefore not necessarily those for a bulk sample of small, randomly oriented grains. The temperature dependent forms of the anomalies seen in the results are reproducible, however.

Below room temperature, the curves for \( \text{AgNO}_3 \), \( \text{TlNO}_3 \), and \( \text{NH}_2\text{NO}_3 \) show dielectric transitions which are similar in form to those found in the alkali metal nitrates.\textsuperscript{46-49} The temperatures marked on the curves are taken as being characteristic of the anomalies for the purpose of comparison. The forms of the anomalies suggest the presence of structural or order-disorder
Fig. 7. Dielectric constant $\varepsilon$ and resistivity $\rho_D$ for TiNO$_3$ as functions of temperature.

transformations in the compounds, and in the case of TiNO$_3$ the dielectric transition temperature of $-35^\circ$C (Fig. 4) agrees well with that of the discontinuity in slope of the specific heat curve of Fig. 2.

The maximum at $-32^\circ$C in the values of $\varepsilon$ obtained on cooling for NH$_4$NO$_3$ (Fig. 5) is very similar to the effect found by Kamiyoshi and Yamakami at the same temperature. The latter authors employed a frequency of 100 kHz, and a powdered sample, and the differences between the two sets of data are no greater than may be expected from the techniques used. On heating, the present results show less hysteresis than those of Kamiyoshi and Yamakami.

The resistivity $\rho_D$ of NH$_4$NO$_3$ (Fig. 4) reflects the course of $\varepsilon$, showing that the increased polarizability is accompanied by relaxation. In addition, the $\rho_D$ curves show slight, but important minima at $\sim -20^\circ$C on cooling and $\sim -3^\circ$C on heating. The latter effect corresponds to the d.c. conductance anomaly found by Brown and McLaren at $\sim -6^\circ$C on heating, and identified as being due to the V$\rightarrow$IV transformation. From a comparison of these results, it appears that the dielectric transitions do not coincide with the IV$\equiv$V transformations, and that negligible changes in $\varepsilon$ are associated with the structural

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transitions, although variations in a.c. and d.c. resistivity do occur. This interpretation is supported by the large discrepancy between the dielectric transition temperature (−32°C) and that detected by other means (−18°C)* on cooling.

On the other hand, the anomalous variations of ε and ρD for NH₄NO₃ below room temperature are clearly of the same nature as those found in the other univalent nitrates, implying that the effect is not a disorder property of the NH₄⁺ ion, although the latter could be involved in the V→IV transformation. It may be significant that the EPR measurements on NH₄NO₃ containing VO³⁺ showed the onset of rapid reorientation of the ion to occur over the hysteresis interval of the dielectric transition. This is not the case for NaNO₃, KNO₃, and CsNO₃, however.⁴⁴

The results for AgNO₃ above room temperature (Fig. 6) show a maximum in ε at 110°C, succeeded by a plateau region at higher temperatures. The increase in ε of ~2.5 times the room temperature value is not associated with an increase in dissipation, the values of ρD being substantially linear, especially on cooling. Thus the increased polarizability involves purely elastic displacements or rotations. It is not known whether the anomaly is reflected in the d.c. resistivity, the measurements of Davis et al.¹⁸ commencing above 110°C. The dielectric anomaly does not appear to be directly connected with the II→I transformation at ~160°C.

The data obtained for TiNO₃ above room temperature (Fig. 7) show a gradual increase in ε above ~70°C, i.e. in the phase II region. The onset of the II→I transition is gradual, and the change completed at 147°C. The rather steeply increasing value of ε in phase I resembles those of the isostructural phase III of RbNO₃.⁴⁹ No appreciable change in ρD occurs at III→II, in accordance with the use of a dry sample (cf. Ref. 22). Apart from the effects of the II→I transition, which has a gradual onset, the variation in resistivity is substantially linear. Of all of the known structural transformations in univalent nitrates, the III→II transition in TiNO₃ has the least effect upon the electrical parameters.

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* See introductory section.

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