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Neutron diffraction study of sodium nitrite in an applied electric field

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Résumé. — Le diagramme de phase de NaNO_2 sous champ électrique est exploré par diffraction neutronique. On n'observe pas de phases commensurables induites par le champ. Pour un champ de $3,4 \text{ kV cm}^{-1}$ on observe un point triple où les trois phases paraélectrique, ferroélectrique et modulée coexistent. Les résultats sont discutés sur la base d'un modèle phénoménologique à la Landau.

Abstract. — The phase diagram of NaNO_2 in an applied electric field is investigated by means of neutron diffraction. No evidence is found for field-induced commensurate phases. For an applied field of 3.4 kV cm^{-1} a triple point is observed at which the paraelectric, ferroelectric and modulated phases coexist. Results are discussed in terms of a phenomenological Landau free-energy.

1. **Introduction.** — Sodium nitrite has been the subject of a considerable number of experimental [1-5] and theoretical [6-12] studies. Between the high-temperature ($T > T_i \simeq 164^\circ\text{C}$) paraelectric phase (space group Immm) and the low temperature ($T < T_f \simeq 162.5^\circ\text{C}$) ferroelectric phase (space group Im2m), NaNO_2 exhibits a sinusoidally-modulated structure, characterized by a polarization wave of wavevector

$$\mathbf{q}_\delta = \delta \mathbf{a}^*.$$

The temperature dependence of δ , between T_i and T_f , as obtained from a recent neutron diffraction study [13], shows a smooth decrease from $\delta(T_i) \simeq 0.122$ to $\delta(T_f) \simeq 0.101$, with no evidence for intermediate « lock-in » phases at rational values of δ (e.g. $\delta = 2/17, 1/9, 2/19$). In the same study, no evidence for higher order satellite reflections could be detected, thus confirming the sinusoidal character of the polarization wave. Also, hysteresis effects were found to be limited to the vicinity of the first-order transition at T_f .

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This behaviour is in contrast to that found in thiourea, another ferroelectric compound which exhibits several intermediate modulated structures sandwiched between the high temperature paraelectric and the low-temperature ferroelectric phases. X-ray and neutron studies of thiourea [14, 15] as a function of temperature and pressure have yielded extensive evidence for the occurrence of non-sinusoidal modulations and large wavevector hysteresis.

Not surprisingly, though not necessarily for the same reasons, the two compounds appear to behave differently under an applied d.c. electric field. Birefringence measurements on thiourea [16] indicate a (E, T) phase diagram of the type shown in figure 1a. The stability region for the modulated phase is limited by a first-order transition line from T_f to T_{tr} and a second-order transition line from T_{tr} to T_i . Additional susceptibility anomalies within the modulated region, have been shown [17, 18] to correspond to a field-induced commensurate lock-in phase of wavevector $b^*/8$. On the other hand dielectric susceptibility measurements by Gesi [19] on NaNO_2 suggest a phase diagram as shown in figure 1b, i.e. with a triple point at (E_T, T_T) and a critical end point at (E_C, T_C) . No details are given in reference [19], concerning the order of the various transition lines, although the topology of the phase diagram would seem to imply the existence of a tricritical point somewhere between T_i and T_T .

The purpose of the present study is to reinvestigate the (E, T) phase diagram of NaNO_2 by means of neutron diffraction. The main advantage of the latter technique, as compared to macroscopic ones, lies in the fact that it yields useful informations about the E - and T -dependence of the amplitude, wavevector and character of the wave.

Fig. 1. — Schematic (E, T) phase diagram observed in (a) thiourea [16-18] and (b) sodium nitrite [19]. In both cases the electric field is applied along the ferroelectric axis.

2. Experimental. — The measurements were performed on the IN8 3-axis spectrometer at the Institut Laue-Langevin, using a PG-filtered incident neutron wavelength of 2.36 Å.

The specimen is a $1.5 \times 8 \times 8 \text{ mm}^3$ plate cut from a single crystal boule, grown by Professor J. P. Chapelle (SDTM, LP 3261, Université Paris-Sud, 91405 Orsay). A 1μ Cu-film has been sputtered onto the large faces ($\perp$ b) of the crystal plate. The specimen is mounted in a 30 W electrical furnace designed to provide a temperature homogeneity of a few 10^{-2} K inside the sample chamber. The temperature stability, averaged over several hours, is found to be better than 0.01 K. Electric field inhomogeneities were minimized by purposely reducing the neutron beam cross-section in such a way as to mask off the periphery of the specimen.

The aim of the experiment is to monitor the intensity and position of the strong $(\pm \delta, 2, 0)$

satellite reflections, as a function of temperature and electric field, independently. In practice a major difficulty arises from the finite electrical conductivity of the specimen ($\rho = 2.4 \times 10^6 \Omega\text{m}$ at 164°C), giving rise to Joule heating under an applied electric field. The heat generated in the bulk of the specimen, a few mW at large fields, diffuse outwards and creates a temperature difference of up to 0.5 K between the specimen and the sensor. Note that thermal gradients within the specimen remain negligible because of the thin plate-like shape of the latter with large metallized faces. Since the thermal resistance between specimen and sensor can be taken as independent of field and temperature (over a range of a few K) the temperature difference between the two may be written as :

$$\Delta T = kVI, \quad (1)$$

where V and I stand for the measured voltage and current across the specimen. The value of the proportionality constant k was adjusted empirically ($k \simeq 0.13 \text{ mW}^{-1} \text{ K}$) as discussed below. The temperature scale for the data shown in figure 2, has been corrected according to equation (1) above (the correction is approximately proportional to E^2).

Fig. 2. — Observed (E, T) phase diagram of NaNO_2 . The commensurate iso- δ lines are shown as dotted lines.

3. Results. — **3.1 PHASE BOUNDARIES.** — Figure 2 summarizes our results concerning the position of the paraelectric-to-modulated transition line $T_i(E)$ and of the modulated-to-ferroelectric line $T_f(E)$. Strong critical scattering is observed near the $T_i(E)$ line, indicating a continuous transition all the way up to the triple point ($T_T = 164.8^\circ\text{C}$, $E_T = 3.4 \text{ kV/cm}$). The tran-

sition at $T_f(E)$ is clearly discontinuous as evidenced by both the abrupt vanishing of the satellite intensity and the simultaneous observation of a discontinuity in the sample's electrical resistance (V/I ratio). The initial slope of the $T_f(E)$ line is found to be :

$$\left[\frac{dE}{dT_f} \right]_{E=0} = 3.26 \text{ kV cm}^{-1} \text{ K}^{-1}. \quad (2)$$

This result is independent of the value chosen for the coefficient k in equation (1). However at finite field values the position of the $T_i(E)$ and $T_f(E)$ lines depend upon k . In fact, the value of k was chosen in such a way as to obtain a *straight line* for the $T_f(E)$ line. This criterion will be justified on the basis of the model developed in § 4.

Above E_T , no satellite reflections are observed. However, a resistivity anomaly can still be detected indicating a weakly first-order *direct* transition between paraelectric and ferroelectric phases. The location of the critical end-point could not be determined in the present study.

Our results are in quantitative disagreement with those of Gesi [19], which indicate a triple point at a lower field value ($\approx 2.6 \text{ kV cm}^{-1}$). Too few details are given in reference [19] about measuring conditions for us to make a plausible guess as to the origin of this discrepancy.

3.2 FIELD-DEPENDENCE OF MODULATION WAVEVECTOR. — Also shown in figure 2 are the values $\delta_i(E)$ and $\delta_f(E)$ of the modulation wavevector in reduced units, measured along the $T_i(E)$ and $T_f(E)$ lines, respectively. For fields up to 2.5 kV/cm we find that the quantity $\delta_i^2(E) - \delta_i^2(0)$ varies quadratically with E :

$$\delta_i^2(E) - \delta_i^2(0) \simeq \lambda E^2,$$

with

$$\lambda = -(1.6 \pm 0.2) \times 10^{-4} (\text{r.u.})^2 \text{ kV}^{-2} \text{ cm}^2. \quad (3)$$

At the triple point we have :

$$\delta_i(E_T) = \delta_f(E_T) = 0.109.$$

Within the modulated region the modulation wavevector appears to vary smoothly as a function of field and temperature. No evidence is found for a plateau at the commensurate values $\delta = 2/17, 1/9, 2/19$ (dotted lines in Fig. 2). This is in contrast to the case of thiourea where a field-induced commensurate phase at $\delta = 1/8$ is observed. This difference is readily understood on the basis of the following symmetry argument. The lowest order candidate for a lock-in energy term at, say, $\delta = 1/9$ has the form

$$V^{(9)} P_{q_\delta}^9 + \text{c.c.},$$

where P_{q_δ} is the (complex) amplitude of the modulation and $V^{(9)}$ is an appropriate anharmonic coefficient. In the present case, the term written above is forbidden for 2 reasons :

- i) The quantity $P_{q_\delta}^9$ is not rotationally invariant.
- ii) The first Umklapp vector along $\mathbf{a}^*$ is $2\mathbf{a}^*$ (due to the body-centred nature of the high-temperature structure). Thus for $\mathbf{q}_\delta = \mathbf{a}^*/9$ the above term is not translationally invariant.

Applying an electric field induces a new potential lock-in term of the form :

$$V^{(10)} P_0 P_{q_\delta}^9 + \text{c.c.},$$

where P_0 is the amplitude of the homogeneous polarization. This type of cross-term is believed to stabilize the $\delta = 1/8$ phase in thiourea [15]. Here, however, it is still *translationally* forbidden. Hence, one must resort to the next higher-order term :

$$V^{(18)} P_{q_\delta}^{18} + \text{c.c.}.$$

Such a term is allowed, with or without field, but is obviously too weak to be effective.

3.3 AMPLITUDE AND CHARACTER OF THE MODULATION. — The study of the first-order satellite intensity shows that the order parameter varies monotonically as a function of temperature and electric field, the iso-intensity curves being roughly parallel to the second-order line $T_i(E)$.

Finally we find that the applied electric field does not alter the sinusoidal character of the modulation. This point is illustrated in figure 3 which shows that at 2.3 kV/cm the second- and third-order satellite reflections are too weak to be detected. This observation is at variance with the prediction made in reference [8].

Fig. 3. — Attempt to observe higher-order satellites under field ($E = 2.3 \text{ kV cm}^{-1}$, $T = 164.65^\circ\text{C}$). The background intensity corresponds to diffuse quasielastic scattering which persists throughout the incommensurate region.

4. Model. — **4.1 FORM OF THE FREE-ENERGY.** — The phase diagram in figure 2 is best discussed in terms of a free-energy functional involving the component of the polarization along $\mathbf{b}$, P_y , and its spatial derivatives along $\mathbf{a}$:

$$G = \int \left\{ \frac{1}{2} A_0 P_y^2 + \frac{1}{4} B P_y^4 + \frac{1}{6} C P_y^6 + \frac{1}{2} \alpha \left(\frac{dP_y}{dx} \right)^2 + \frac{1}{4} \beta \left(\frac{d^2 P_y}{dx^2} \right)^2 + \gamma \left(P_y \frac{dP_y}{dx} \right)^2 - E P_y \right\} dx. \quad (4)$$

General free-energy functionals of the above type have been considered by Hornreich *et al.* [20], Michelson [21], and Coutinho-Filho *et al.* [22] and, in the specific context of uniaxial ferroelectrics, by Ishibashi *et al.* [8] and Lederer *et al.* [23].

Following the authors of reference [8] we have included a term proportional to $\left(P_y \frac{dP_y}{dx} \right)^2$. This term is essential for the overall consistency of the model, as discussed below. On the other hand our model differs from that of reference [8] in two important respects :

i) We assume $B < 0$. The opposite assumption ($B > 0$) is not consistent with our observation

of a direct para-ferro transition above E_T . Furthermore, it is not consistent with the dielectric behaviour of NaNO_2 in zero-field. For example, two different Curie laws are observed [1] in the paraelectric and ferroelectric phases :

$$\chi_{22}^{\text{p,f}} = \frac{C^{\text{p,f}}}{T - T_c^{\text{p,f}}}.$$

If the direct (virtual) para-ferro transition were continuous ($B > 0$), one would expect to find

$$T_c^{\text{p}} \simeq T_c^{\text{f}} \simeq T_f'$$

where T_f' is the temperature of the virtual direct transition

$$T_f < T_f' < T_i.$$

Instead Nomura [1] finds :

$$T_c^{\text{f}} - T_c^{\text{p}} \simeq 3.5 \text{ K}.$$

Similarly, the spontaneous polarization measured below T_f , should extrapolate to zero at T_f' (i.e. below T_i). The data of Hamano [5] show that this is not possible, except through the use of an unrealistic critical exponent ($\beta = 0.20$).

ii) Following the discussion in § 3.3, we assume a purely sinusoidal polarization wave, i.e. we write :

$$P_y(x) = P_0 + 2 P_\delta \cos(2 \pi \delta x). \quad (5)$$

Inserting (5) into (4) and neglecting some unessential sixth-order terms, we get :

$$G = \frac{1}{2} A_0 P_0^2 + \frac{1}{4} B P_0^4 + \frac{1}{6} C P_0^6 + A_\delta P_\delta^2 + \frac{3}{2} B_\delta P_\delta^4 + \frac{3}{2} (B + B_\delta) P_0^2 P_\delta^2 - P_0 E \quad (6)$$

with :

$$A_\delta = A_0 + \alpha \delta^2 + \frac{1}{2} \beta \delta^4, \quad (7)$$

and :

$$B_\delta = B + \frac{4}{3} \gamma \delta^2. \quad (8)$$

In addition, we assume :

$$A_0 = a(T - T_0); \quad B < 0; \quad C > 0; \quad \alpha < 0; \quad \beta > 0; \quad \gamma > 0.$$

4.2 SINUSOIDAL PHASE IN ZERO-FIELD. — Setting $P_0 = 0$ in (6), the free-energy reduces to :

$$G_s = A_\delta P_\delta^2 + \frac{3}{2} B_\delta P_\delta^4. \quad (9)$$

The para-sinusoidal transition temperature T_i and the modulation wavevector at T_i , δ_i , are obtained from :

$$A_\delta = \frac{\partial A_\delta}{\partial \delta} = 0,$$

which yields :

$$\left. \begin{aligned} \delta_i^2 &= -\frac{\alpha}{\beta}, & (a) \\ a(T_i - T_0) &= \frac{\alpha^2}{2\beta} & (b) \end{aligned} \right\}. \quad (10)$$

In order to have a continuous transition at T_i we have implicitly assumed $B_{\delta_i} (\equiv B_i) > 0$, which in turn implies :

$$\frac{4}{3} \gamma \delta_i^2 > |B|. \quad (11)$$

Below T_i , (9) may be minimized firstly with respect to the amplitude P_δ of the modulation, yielding :

$$G_s = -\frac{A_\delta^2}{6 B_\delta}, \quad (12)$$

and secondly with respect to the value of the modulation wavevector δ :

$$\frac{\partial G_s}{\partial \delta} = 0. \quad (13)$$

With the help of (7), (8) and (10), equation (13) yields :

$$\delta^2 - \delta_i^2 = \frac{1}{2} \left\{ \left[\frac{B_i^2}{\gamma^2} + \frac{8a}{3\beta} (T - T_i) \right]^{1/2} - \frac{B_i}{\gamma} \right\}. \quad (14)$$

Close to T_i we have :

$$\delta^2 - \delta_i^2 \simeq \mu (T - T_i).$$

With :

$$\mu = \frac{2\gamma a}{3\beta B_i}. \quad (15)$$

Finally the zero-field free-energy of the sinusoidal phase is obtained as

$$G_s(T) = -\frac{a^2(T - T_i)^2}{6 B_i} + \mathcal{O}(T - T_i)^3. \quad (16)$$

4.3 PARA-SINUSOIDAL TRANSITION LINE ($T_i(E)$). — Assuming $B_{\delta_i}(E) > 0$, the equation of the transition line is obtained by setting to zero the coefficient of P_δ^2 in (6) :

$$A_\delta + \frac{3}{2}(B + B_\delta) P_0^2 = 0. \quad (17)$$

The value of $\delta_i(E)$ is obtained by taking the derivative of (17) with respect to δ . Whence :

$$\delta_i^2(E) - \delta_i^2 = -\frac{2\gamma}{\beta} P_0^2. \quad (18)$$

In the non-ferroelectric state we have :

$$P_0 = \frac{E}{A_0} \left(1 - \frac{BE^2}{A_0^3} \right) + \mathcal{O}(E^5).$$

For $E \ll \left(\frac{A_0^3}{|B|} \right)^{1/2}$ we may take

$$P_0 \simeq \frac{E}{A_0},$$

whence :

$$\delta_i^2(E) - \delta_i^2 = \lambda E^2 + \mathcal{O}(E^4),$$

with :

$$\lambda = -\frac{2\gamma}{\beta a^2(T_i - T_0)^2}. \quad (19)$$

Finally, the equation of the transition line is obtained as :

$$a(T_i(E) - T_i) = -\frac{3}{2}(B + B_i) \frac{E^2}{a^2(T_i - T_0)^2} + \mathcal{O}(E^4). \quad (20)$$

Combining (15) and (19) the quantity B_i appearing in (20), is expressed in terms of observable quantities as :

$$B_i = -\frac{a^3}{3}(T_i - T_0)^2 \frac{\lambda}{\mu}. \quad (21)$$

4.4 DIRECT PARA-FERRO TRANSITION LINE ($T'_f(E)$). — The stability of the ferroelectric phase with respect to the paraelectric phase is discussed using equation (6) with $P_\delta = 0$. The coexistence line originates at the point ($E = 0, T = T'_f$) and ends at the critical point (E_c, T_c), with

$$a(T'_f - T_0) = \frac{3B^2}{16C},$$

$$a(T_c - T_0) = \frac{9B^2}{20C}; \quad E_c = \frac{8C}{3} \left(\frac{-3B}{10C} \right)^{5/2}.$$

The slope of the line at $E = 0$ and $E = E_c$ is given by

$$\left(\frac{dE}{dT} \right)_{E=0} = \frac{1}{2} a \left(\frac{-3B}{4C} \right)^{1/2},$$

and

$$\left(\frac{dE}{dT} \right)_{E_c} = a \left(\frac{-3B}{10C} \right).$$

The transition at $T'_f(E)$ is virtual below the triple point and real above.

4.5 FERRO-SINUSOIDAL TRANSITION LINE ($T_f(E)$). — The difference $T'_f(E) - T_f(E)$ is largest at $E = 0$ and vanishes at E_T . Furthermore, a simple thermodynamic argument shows that if, as is the case here, the transition at $T_i(E)$ remains continuous all the way up to E_T , the two curves $T_f(E)$ and $T'_f(E)$ should have a common tangent at E_T .

The zero field transition temperature T_f is obtained by equating the free energies of the ferro and sinusoidal phases :

$$G_s(T_f) = G_f(T_f), \quad (22)$$

where G_s is given in (12), while G_f is obtained from (2) :

$$G_f(T) = \frac{1}{2} a(T - T_0) P_0^2 + \frac{1}{4} B P_0^4 + \frac{1}{6} C P_0^6. \quad (23)$$

Minimizing (23) with respect to the spontaneous polarization P_0 we get :

$$P_0^2 = \frac{|B| + \sqrt{B^2 - 4aC(T - T_0)}}{2C}.$$

For realistic values of the coefficients a , T_0 , B , B_i and C (cf. § 5), the numerical solution of (22) yields :

$$T'_f - T_f \simeq 0.1 \text{ K}.$$

This small temperature difference reflects the large entropy difference between the ferroelectric state on one hand and the paraelectric and sinusoidal states on the other hand.

5. Discussion. — The phenomenological model developed in the previous section is readily applied to the case of NaNO_2 . Starting from the known values [4] of the paraelectric Curie constant and Curie temperature ($C = 5 \text{ 130 K}$; $T_i - T_0 = 2.45 \text{ K}$) and from the observed value of δ_i ($= 0.122 \text{ red. units}$), we obtain, with the help of (10) :

$$\begin{aligned} a &= 2.2 \times 10^7 \text{ MKS}, \\ \alpha &= -7.3 \times 10^9 \text{ MKS (r.u.)}^{-2}, \\ \beta &= 4.9 \times 10^{11} \text{ MKS (r.u.)}^{-4}. \end{aligned}$$

From the observed values of λ and μ ($\lambda = 1.6 \times 10^{-4} \text{ (r.u.)}^2 \text{ kV}^{-2} \text{ cm}^2$; $\mu = 0.22 \times 10^{-2} \text{ (r.u.)}^2 \text{ K}^{-1}$) and using (19) and (21) we get :

$$\begin{aligned} \gamma &= 1.15 \times 10^{13} \text{ MKS (r.u.)}^{-2}, \\ B_i &= 1.55 \times 10^{11} \text{ MKS}, \end{aligned}$$

whence :

$$B = B_i - \frac{4}{3} \gamma \delta_i^2 = -0.70 \times 10^{11} \text{ MKS}.$$

Fig. 4. — Calculated (E, T) phase diagram. The model is constrained to yield the correct value of $T_i - T_f$. The virtual transition line $T'_f(E)$ is shown as a dotted line.

Finally, fixing the value of T_f ($T_i - T_f = 1.6$ K) we determine the value of C :

$$C = 4.3 \times 10^{13} \text{ MKS.}$$

Figure 4 shows the phase diagram calculated using the above numerical values. The agreement with the observed phase diagram is only semi-quantitative. In particular the model underestimates the downward bending of the $T_i(E)$ line. This discrepancy is only partly due to our neglect of terms of order E^4 in deriving (20).

Finally, since none of the observables used in the calculation depend upon the value of k in equation (1), we can use the calculated phase diagram to estimate the uncertainty on the temperature scale arising from our identification of the $T_f(E)$ line to a straight line : we find an uncertainty of about 0.1 K at $E = E_T$.

In conclusion we find that the simple phenomenological model presented here gives a good semi-quantitative account of the observed phase diagram. In this model the term $\propto \left(P_y \frac{\partial P_y}{\partial x} \right)^2$ plays a central role in so far as it accounts for the temperature and field dependence of the modulation wavevector. At the same time it explains why the transition at T_i is found to be continuous, while the direct para-ferro transition would be discontinuous. Obviously, more elaborate treatments are possible and are probably warranted on the basis of the present data. Further experimental work is underway in order to determine the position of the critical point.

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References

- [1] NOMURA, S., *J. Phys. Soc. Japan* **16** (1961) 2440.
- [2] YAMADA, Y., SHIBUYA, I. and HOSHINO, S., *J. Phys. Soc. Japan* **18** (1963) 1594.
- [3] SAKURAI, J., COWLEY, R. A. and DOLLING, G., *J. Phys. Soc. Japan* **28** (1970) 1426.
- [4] HATTA, I., *J. Phys. Soc. Japan* **28** (1970) 1266.
- [5] HAMANO, K., *J. Phys. Soc. Japan* **35** (1973) 157.
- [6] YAMADA, Y. and YAMADA, T., *J. Phys. Soc. Japan* **21** (1966) 2167.
- [7] LEVANYUK, A. P. and SANNIKOV, D. G., *Sov. Phys. Solid State* **18** (1976) 1122.
- [8] ISHIBASHI, Y. and SHIBA, H., *J. Phys. Soc. Japan* **45** (1978) 409.
- [9] MICHEL, K. H., *Phys. Rev. B* **24** (1981) 3998.
- [10] HEINE, V. and MCCONNELL, J. D. C., *Phys. Rev. Lett.* **46** (1980) 1092.
- [11] KLEIN, M. L., McDONALD, I. R. and OZAKI, Y., *Phys. Rev. Lett.* **48** (1982) 1197.
- [12] AIZU, J., *J. Phys. Soc. Japan* **51** (1982) 515 ; **51** (1982) 2885.
- [13] DURAND, D., DENOYER, F., LAMBERT, M., BERNARD, L. and CURRAT, R., *J. Physique* **43** (1982) 149.
- [14] MOUDDEN, A. H., DENOYER, F. and LAMBERT, M., *J. Physique* **39** (1978) 1323.
- [15] DENOYER, F., MOUDDEN, A. H., CURRAT, R., VETTIER, C., BELLAMY, A. and LAMBERT, M., *Phys. Rev. B* **25** (1982) 1697.
- [16] JAMET, J. P., *J. Physique-Lett.* **42** (1981) L-123.
- [17] JAMET, J. P., LEDERER, P. and MOUDDEN, A. H., *Phys. Rev. Lett.* **48** (1982) 442.
- [18] MOUDDEN, A. H., SVENSSON, E. C. and SHIRANE, G., *Phys. Rev. Lett.* **49** (1982) 557.
- [19] GESI, K., *Japan J. Applied Phys.* **4** (1965) 818.
- [20] HORNREICH, R. M., LUBAN, M. and SHTRICKMAN, S., *Phys. Rev. Lett.* **35** (1975) 1678.
- [21] MICHELSON, A., *Phys. Rev. B* **16** (1977) 577.
- [22] COUTINHO-FILHO, M. E. and DE MOURA, M. A., *J. Magn. Magn. Mat.* **15-18** (1980) 433.
- [23] LEDERER, P. and CHAVES, C. M., *J. Physique-Lett.* **42** (1981) L-127.