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Spontaneous and electric field induced quadratic optical nonlinearity in ferroelectric crystals $AgNa(NO_2)_2$

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We demonstrate the second harmonic generation (SHG) in ferroelectric $AgNa(NO_2)_2$ crystals resulting from the spontaneous and electric field induced polarizations. Relatively high effective nonlinear optic (NLO) susceptibility is combined in this crystals with the existing several phase matching geometries of NLO interaction. Anomalously large response of SHG with respect to an applied electric field has been found in the vicinity of the paraelectric-to-ferroelectric phase transition. The behavior of NLO properties in the ferroelectric phase and especially in the region of the Curie point is discussed within the phenomenological theory. © 2010 American Institute of Physics. [doi:10.1063/1.3315941]

Ferroelectric materials are considered as very promising materials for a number of applications being usually related to their large nonlinear response with respect to the electromagnetic radiation in the optical range. Corresponding effects include electro-optic, second harmonic generation (SHG), parametric amplification and generation, or other nonlinear optic (NLO) or parametrical optic phenomena widely used for the light modulation or its frequency conversion.^{1,2}

In this letter we demonstrate the quadratic optical nonlinearity resulting from spontaneous and electric field induced polarizations in silver sodium nitride [AgNa(NO₂)₂, hereafter SSN] crystals. To probe such nonlinearity the SHG, being represented as $\chi^{(2)}$ -nonlinear frequency-conversion processes, has been chosen as the most appropriate technique. SSN crystals exhibit the phase transition (PT) at $T_c \approx 38$ °C (Refs. 3 and 4) from a paraelectric phase (space group D_{2h}^{24}) to the proper ferroelectric phase (space group C_{2v}^{19}). Comprehensive dielectric,^{3–5} specific-heat,⁶ and elastic⁷ measurements have revealed that the PT is of first order, but very close to second order one. Ferroelectricity in SSN appears due to an ordering of the NO₂⁻ dipoles which at room temperature give quite large spontaneous polarization (about 8 μ C/cm²) being oriented along [010] crystallographic direction.⁴

Single SSN crystals were grown from aqueous solution containing 9.8 wt % AgNO₂ and 37.2 wt % NaNO₂ by the slow evaporation method at constant temperature (~25 °C). We used the standard crystallographic orientation for the paraelectric phase: a=8.05 Å, b=10.77 Å, and c=10.76 Å. Crystals SSN are yellowish with the perfect cleavage plane parallel to [101]- and [101]-directions. Accordingly, the geometry of samples was adjusted to these planes, i.e., we used the slabs or plates having perfectly cleaved faces [101] (or [101]) without further polishing and with deposited silver paste electrodes on conventionally polished [010]-faces. The SHG has been excited by means of nanosecond IR Q-switched laser Vector 1064–3000–30 (λ = 1064 nm) and registered in a standard setup described earlier in Ref. 8. The sample has been set into thermostabilized optical cell and rotated by means of a step motor with angular step of 0.01 °C and rotation speed 1–5 °C/min. The accuracy of the temperature stabilization was about 0.01 °C.

In the ferroelectric phase SSN crystals exhibit only a very weak intensity SHG if the incident light propagates exactly along the [101]-direction which may be explained by a lack of phase matching between the interacting waves in this direction. Nevertheless, the phase matching indeed can be achieved in slightly tilted geometry. The inset (a) in Fig. 1 demonstrate the example of such phase matching geometry (PMG) defined by the tilt angle θ_p and azimuthal angle φ_p . Here the incident laser beam (λ =1064 nm) is polarized vertically and outgoing intense SHG (λ =532 nm) exhibits nearly horizontal polarization at small θ_p . One should be emphasized, that the measurements have been performed on



FIG. 1. (Color online) Temperature dependence of the phase matching angle φ_p being measured at the fixed tilt angle $\theta_p = 8.2^{\circ}$. Inset (a): phase matching geometry in SSN crystals. Inset (b): Maker fringe pattern in the vicinity of the phase matching angle φ_p (thickness d=0.6 mm).

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FIG. 2. (Color online) Temperature dependences of $\chi_{\rm eff}^{(2)}$ determined at heating and cooling in PMG. Red dashed line marks the temperature behavior of the spontaneous polarization P_s measured at heating in Ref. 4. Black solid line is the fit of $\chi_{\rm eff}^{(2)}(T)$ -dependence measured at heating.

fresh samples being cut from the crystals that were grown in the ferroelectric phase. Obtained in such a way crystals are usually single domain and for this reason we believe that we deal with the birefringent phase matching rather than with so-called quasiphase matching provided by a regular ferroelectric domain structure. The φ -dependence of SHG intensity $I_{2\omega}$ reveals Maker-fringe pattern [Fig. 1, inset (b)]. At the fixed tilt angle $\theta = \theta_p$ the other phase matching angle φ_p is only weakly temperature dependent in the ferroelectric phase (see Fig. 1). Thus the variation of the effective second order NLO susceptibility $\chi^{(2)}_{\rm eff}(T)$ scales with the temperature dependence of $I^{1/2}_{2\omega}$ measured at PMG. Comparing it with the intensity of SHG in SiO₂ (χ_{111} -component) one obtains at T=23 °C the magnitude $\chi_{eff}^{(2)}=4.0$ pm/V. Moreover, our evaluations reveal that the weight factors bonded to the partial contributions of symmetry allowed tensor components $\chi_{ijk}^{(2)}$ into the effective NLO susceptibility $\chi_{eff}^{(2)}$ either do not depend at all or change only slightly with temperature due to a weakly varying $\varphi_p(T)$ -dependence. Corresponding corrections appear thus beyond the accuracy of our measurements and will be ignored in further analysis.

Figure 2 shows the temperature dependences of $\chi_{eff}^{(2)}$ measured in PMG at heating and cooling. As one expects, the PT from the ferroelectric to paraelectric phase is characterized by vanishing of SHG, however variation of $\chi_{eff}^{(2)}(T)$ in the vicinity of Curie point appears to be gradual with a lack of a jump characteristic for the first order PTs. Both at heating and cooling the PT appears to be smeared in the temperature interval of about 1.5 °C and is characterized by the temperature hysteresis of about 0.5 °C. In the ferroelectric phase $\chi_{eff}^{(2)}(T)$ well scales with the temperature dependence of spontaneous polarization $P_s(T)$ measured in Ref. 4 (see red dashed line in Fig. 2). We interpret such behavior within the phenomenological Landau theory of second order PTs. In the case of $D_{2h}^{24} \rightarrow C_{2v}^{19}$ transition the free energy takes the form

 $F = F_P + F_{int},$

$$F_{P} = \frac{1}{2}A(T - T_{0})P^{2} + \frac{1}{4}BP^{4} + \frac{1}{6}CP^{6} - PE,$$

$$F_{int} = -\alpha_{ijmn}^{0,k,k,2k}[P_{i}(0)P_{j}(k)P_{m}(k)P_{n}(-2k) + c \cdot c] + \frac{1}{2\kappa_{ii}^{(K)}}P_{i}(K)P_{i}(-K),$$
(1)

where F_P is the free energy expansion on the Y-component

of the polarization P being the order parameter of this model, A, B, and C are the free energy expansion coefficients, T_0 is the Curie–Weiss temperature, $\alpha_{ijmn}^{0,k,k,2k}$ are the coupling constants, $\kappa_{ii}^{(K)} = \varepsilon_0 \chi_{ii}(K)$, ε_0 is the vacuum permittivity and $\chi_{ii}(K)$ are the components of the inhomogeneous linear dielectric susceptibility at the wavevector K. F_{int} includes α -terms representing the invariants with respect to the rotational-translational operations of D_{2h}^{24} space group and indeed describe the interaction between the static homogeneous polarization $P_i(0)$, inhomogeneous polarizations $P_i(k)$ and $P_m(k)$, being induced by the light, and its second harmonic $P_m(2k)$. In the simplest approximation one ignores the static polarization resulting from the NLO conversion. Then at E=0 the minimization of F with respect to P and $P_n(2k)$ yields $|P_n(2k)| = 2\alpha_{2jmn}^{(0,k,k,2k)} \kappa_{nn}^{(2k)} P_s|P_j(k)||P_m(k)|$ where P_s is the equilibrium value of spontaneous polarization being equal to 0 above the <u>Curie point $T_c = T_0 + 3B^2 / 16AC$ and de-</u> fined as $P_s^2 = (-B + \sqrt{B^2 - 4CA(T - T_0)})/2C$ at $T < T_c$. Accordingly, $\chi_{njm}^{(2)} \propto \alpha_{2jmn}^{(0,k,k,2)} \kappa_{nn}^{(2k)} P_s$ thus NLO susceptibility may be viewed as such which is resulted from the spontaneous polarization, i.e., one may call it as spontaneous SHG in analogy to the terminology used for the characterization of electro-optical properties in ferroelectrics (see e.g., Ref. 9). More general form of this equation ignores an origin of the polarization itself, i.e., $\chi_{njm}^{(2)} \propto \alpha_{2jmn}^{(0,k,k,2k)} \kappa_{nn}^{(2k)} P$. Here *P* implies the total polarization being presented as superposition of P_s and P_{ind} , where $P_{\text{ind}} = \varepsilon_0 (\varepsilon_{22} - 1) E \approx \varepsilon_0 \varepsilon_{22} E$ is the induced polarization caused by an applied external field $(E || \mathbf{Y})$ and ε_{22} is the dielectric constant. In the paraelectric phase and at E=0 the SHG is not observed since $P_s=0$ and all the tensor components $\chi_{njm}^{(2)}=0$. However, the SHG may be induced here by applying an external electric field along the ferro-electric axis. In this case $\chi_{njm}^{(2)} \propto \alpha_{2jmn}^{(0,k,k,2k)} \kappa_{nn}^{(2k)} \varepsilon_0 \varepsilon_{22} E$ thereby one expects its anomalous increase in the vicinity of the Curie point T_c keeping in mind that the dielectric constant ε_{22} of proper ferroelectric SSN is subjected to the Curie-Weiss law: $\epsilon_{22}(T) \propto C_0(T-T_0)^{-1}$, C_0 is the Curie–Weiss constant.³ An independent symmetry interpretation of this phenomena may be given basing on the Curie principle. Following to it the electric field applied along the Y-axis lowers the crystal symmetry from the centrosymmetric point group mmm to a noncentrosymmetric (polar) group mm_{2y} thus the SHG becomes to be symmetry allowed if $E \neq 0$.

Figure 3 demonstrates the SHG in the centrosymmetric paraelectric phase of SSN crystals induced by applied electric field $\vec{E} || \mathbf{Y}$. Far above T_c the field dependences $\chi_{\text{eff}}^{(2)}(E)$ are nearly linear, however closer to T_c they becomes more nonlinear and finally even get a saturated character. The constant β , defined as derivative $d\chi_{\text{eff}}^{(2)}/dE \propto dP_{\text{ind}}/dE \propto \varepsilon_{22}$, represents the third order nonlinear susceptibility. Figure 4 shows the constant β , being determined at $E \rightarrow 0$, and its inverse magnitude β^{-1} versus *T*. One can realize that $\beta(T)$ behaves in analogical manner as static dielectric constant $\varepsilon_{22}(T)$. By extrapolating linear dependence $\beta^{-1}(T)$ into the region below T_c one determines the temperature T_0 . The difference ΔT $= T_c - T_0$ is only 0.25 °C what can be considered as an evidence that we indeed deals with the PT of the first order being very close to the tricritical point.⁶

The anomalous behavior of $\chi_{\text{eff}}^{(2)}(T, E \neq 0)$ near T_c (see Fig. 3) may be evaluated within the phenomenological theory by a solving the electric equation of state $\partial F_P / \partial P$



FIG. 3. (Color online) Field induced SHG in the paraelectric phase. Points correspond to $\chi_{\text{eff}}^{(2)}$ vs *T* measured in PMG at different magnitudes of the external electric field *E* being applied along the *Y*-axis. Solid lines are the best fits obtained by solving the electric equation of state.

= $A(T-T_o)P+BP^3+CP^5-E=0$ for each particular temperature *T*. Solid colored lines in Fig. 3 and black line in Fig. 2 are the best fits that have been obtained applying the set of the free energy parameters (in SI units): $A=1.6 \times 10^7$, $B=-1.0 \times 10^9$, and $C=6.0 \times 10^{12}$. One can realize that such model quite properly describes the $\chi_{\rm eff}^{(2)}(T, E \neq 0)$ -dependences at $T > T_c + 1.5$ °C but evidently



FIG. 4. (Color online) Constant β and its inverse magnitude β^{-1} vs *T* in the paraelectric phase. Solid black lines marks the Curie–Weiss behavior.

fails in close vicinity of T_c , especially at high applied voltages. The measured values of $\chi_{\text{eff}}^{(2)}$ are considerably smaller here comparing to the ones expected from the theory. Such discrepancy may have a number of reasons. The most serious and very likely one may be a coexistence of metastable polar and nonpolar regions in the PT vicinity. A smeared character of the paraelectric-to-ferroelectric PT is an evidence for such coexistence indicating on a sufficiently spatially inhomogeneous structure that occurs in the region of the Curie point.

In conclusion, we have presented here the spontaneous and electric field induced SHG in SSN. In ferroelectric phase such crystal are characterized by the effective NLO susceptibilities $\chi^{(2)}_{eff}$ being comparable or larger comparing to a number of other known NLO inorganic or organic materials (see e.g., Ref. 10). Relatively high effective NLO susceptibilities are combined in these materials with existing PMG what makes them rather perspective for a number of NLO applications such as e.g., SHG, parametric NLO generation or amplification or other applications that deal with the frequency conversion. In addition, an anomalously large response of NLO properties with respect to applied electric field has been found in the vicinity of the Curie point. This may also have a number of applications, especially there, where a high-efficient control and/or tuning of the SHG intensity are required.

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- ¹B. E. A. Saleh and M. C. Teich, *Fundamentals of Photonics* (Wiley, New York, 1991).
- ²F. Agulló-López, J. M. Cabrera, and F. Agulló-Rueda, *Electro-Optics*, *Phenomena, Materials, and Applications* (Academic, New York, 1994).
- ³K. Gesi, J. Phys. Soc. Jpn. **28**, 395 (1970).
- ⁴K. Gesi, J. Phys. Soc. Jpn. 33, 108 (1972).
- ⁵J. Petersson, E. Schneider, and R. Siems, Z. Phys. B: Condens. Matter **39**, 233 (1980).
- ⁶J. Helwig, J. Petersson, and E. Schneider, Z. Phys. B 28, 87 (1977).
- ⁷V. P. Soprunyuk, A. Fuith, H. Kabelka, K. Knorr, A. Klöpperpieper, K. Sokalski, and A. V. Kityk, Phys. Rev. B **66**, 104102 (2002).
- ⁸B. Sahraoui, J. Luc, A. Meghea, R. Czaplicki, J.-L. Fillaut, and A. Migalska-Zalas, J. Opt. Pure Appl. Opt. **11**, 024005 (2009).
- ⁹L. G. Lomova, A. S. Sonin, and T. A. Regulskaya, Sov. Phys. Crystallogr. **13**, 68 (1968).
- ¹⁰A. Keens and H. Happ, J. Phys. C **21**, 1661 (1988); W. Bi, N. Louvain, N. Mercier, J. Luc, I. Rau, F. Kajzar, and B. Sahraoui, Adv. Mater. **20**, 1013 (2008).