Dielectric Properties of Strongly Polar Nematogens

Joanna Czub^a, Roman Dąbrowski^b, Stanisław Urban^c, and Dorota Ziobro^b

^a Faculty of Metals Engineering and Industrial Computer Science, AGH University of Technology, Al. Mickiewicza 30, 30-059 Kraków, Poland

^b Institute of Chemistry, Military University of Technology, Kaliskiego 2, 00-908 Warsaw, Poland

^c Institute of Physics, Jagiellonian University, Reymonta 4, 30-059 Kraków, Poland

Reprint requests to Dr. J. C.; Fax: 0048-12-6173190; E-mail: joanna.czub@agh.edu.pl

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Results of the static and dynamic dielectric investigations of four liquid crystalline compounds consisting of two or three phenyl rings with *n*-alkoxy and cyano terminals, carboxylic bridging groups, and fluoro lateral substituents in pure state and in the solution of low polar liquid crystalline medium, are presented. These compounds form the nematic phase being a monotropic one in the case of two ring compounds. The dielectric parameters are analyzed in relation to the dipolar structures of the molecules. The dielectric properties of pure substances are strongly influenced by conductivity and dimerization effects, while extracted from the data for solutions are consisted well with the structures. In the nematic phase the compounds exhibit a crossover of the principal permittivity components among the kilo and megahertz frequency range.

Key words: Liquid Crystals; Nematic Phase; Dielectric Properties; Dipolar Structure.

1. Introduction

Dielectric relaxation studies of liquid crystals (LC) yield important information on molecular reorientations and their interactions in mesophases [1–4]. Knowledge of the dielectric properties of LCs is also important for the technical reasons due to their connection with electrooptical display devices as they strongly influence threshold voltage and steepness of electrooptical curves [5–7].

The aim of the present work was to analyze the influence of rigid core length, terminal chain length, and dipole moment value on the dielectric properties of four substances having similar chemical structures (Table 1). All substances are nematogens, although two-ring compounds exhibit a monotropic nematic (N) phase only. They have several dipolar groups ($-\text{OC}_n\text{H}_{2n+1}$ and $-\text{CN}$ terminals, F atoms at the lateral positions, $-\text{COO}$ bridging groups) which create strong longitudinal and transverse dipole moment components. This should be reflected in the dielectric properties of the isotropic and nematic phases observed at low frequencies (static case) as well as at high frequencies (dynamical case). Such a compound structure is especially interesting for some applications [7]. They belong to the class of materials termed as dual frequency materials exhibiting a crossover in electric permittivity

constants at a certain frequency what leads to a change of sign of the dielectric anisotropy [6, 7].

It was found that such compounds, especially the three-ring ones, exhibited a large conductivity effect at low frequencies in spite of purification the chromatography column filled with silica-gel and several recrystallization procedures repeated. It is known from calculations carried out by Demus et al. [8] that for a LC medium with dielectric anisotropy $\Delta\epsilon > 10$ the resistivity starts to decrease probably because of dissociation of polar bonds and inclusion of ions. Therefore, for those substances the static permittivity components were determined also from the studies of their 10% wt. solution in the **Base 903** ester mixture showing a small negative dielectric anisotropy (Table 1). Similar studies were reported by Heppke et al. [9].

2. Experimental

All substances were synthesized in the Institute of Chemistry, Military University of Technology, Warsaw, according to the route described in [10, 11].

The dielectric relaxation spectra, $\epsilon^*(f) = \epsilon'(f) - i\epsilon''(f)$, were recorded with the aid of an impedance analyzer HP 4192A in the frequency range of 1 kHz – 30 MHz. The parallel plate capacitor was used ($C_0 \sim 50$ pF). The thickness of the samples was 0.7 mm.

Acronym	Substance	Phase transitions [°C]
1		Cr 57.8 I (N 21.5) [10]
2		Cr 62.5 I (N 39.4) [11]
3		Cr 122 N 260 I [10]
4		Cr 117.2 N 188.1 I [11]
Base 903 (B)		Cr -10 N 63.1 I

B1	10% wt. mixture of 1 with B	Cr < 20 N 62.7 I
B2	10% wt. mixture of 2 with B	Cr < 20 N 64.0 I
B3	10% wt. mixture of 3 with B	Cr < 20 N 75.0 I
B4	10% wt. mixture of 4 with B	Cr < 20 N 74.7 I

Table 1. Chemical formulae and phase transition temperatures of the substances studied.

The samples in the N phase were oriented by a magnetic field of 0.8 T. Two experimental geometries were applied: $\mathbf{E} \parallel \mathbf{B} \equiv \mathbf{n}$ and $\mathbf{E} \perp \mathbf{B} \equiv \mathbf{n}$ ($\mathbf{n}$ – director, $\mathbf{E}$ – measuring electric field), enabling the measurement of the $\epsilon_{\parallel}$ and $\epsilon_{\perp}$ permittivity tensor components, respectively. The temperature was stabilized within ± 0.1 K in the range of 20 °C–90 °C and ± 0.2 K in the range of 90 °C–160 °C. In the isotropic phase a time domain spectrometer (TDS) was employed that covers the frequency range of 10 MHz – 3 GHz [12]. All measurements were carried out with decreasing temperature which caused supercooling of the N phase. In substances **1** and **2** a monotropic N phase appeared.

3. Results

3.1. Two-Ring Compounds

Figure 1 presents a comparisons of the tensor permittivity components, $\epsilon_{\parallel}$, $\epsilon_{\perp}$, and $\bar{\epsilon} = (\epsilon_{\parallel} + 2\epsilon_{\perp})/3$, obtained for two-ring compounds. The monotropic N phase could be supercooled by c. six degrees only

Fig. 1. Static permittivity components for compounds **1** and **2**. The inset shows the dielectric anisotropy $\Delta\epsilon = \epsilon_{\parallel} - \epsilon_{\perp}$ versus the shifted temperature $\Delta T = T_{NI} - T$.

before a rapid crystallization. The relaxation spectra collected in the isotropic and nematic phases of both substances are shown in Figures 2 and 3 in form of the Cole-Cole plots [13]. The superposition of the Debye-

Fig. 2. Cole-Cole plots for the isotropic phase of two-ring substances **1** and **2**. The spectra in the upper row were collected at temperatures close to the clearing point with the aid of an impedance analyzer, in the bottom row by the TDS method in the high temperature region of the isotropic phase.

Fig. 3. Cole-Cole plots for two-ring substances in the nematic phase orientated parallel (upper row) and perpendicular (lower row) to the measuring field. The spectra were collected with the aid of an impedance analyzer.

type processes well fits the experimental points,

$$\varepsilon^*(\omega) = \varepsilon'(\omega) - i\varepsilon''(\omega) = \sum_k \frac{\varepsilon_s - \varepsilon_{\infty k}}{1 + i\omega\tau_k} - i \frac{\sigma(\omega)}{\varepsilon_0\omega}, \quad (1)$$

where ε_s and ε_{∞} stand for the static and high frequency permittivities, respectively, τ is the relaxation time, $\sigma(\omega)$ the ionic conductivity, $\omega = 2\pi f$ is a circular frequency, k numbers the relaxation processes, $\varepsilon_s - \varepsilon_{\infty}$ is called the strength of the relaxation process or the dielectric increment.

3.2. Three-Ring Compounds

Due to high clearing temperatures in both three-ring compounds (Table 1), the relaxation spectra could be collected only in a temperature region far below the clearing points. Both substances exhibited a strong conductivity at low frequencies, as shown in Figure 5. Several re-crystallization procedures were applied which did not suppressed the ionic conductivity considerably, especially in the case of compound **3**.

Fig. 4. Results of detailed analysis of the dielectric relaxation spectra presented in Figures 2 and 3 for the substance 1: a) dielectric parameters, b) relaxation times.

Fig. 5. Dispersion and absorption spectra for three-ring compounds in the nematic phase (parallel orientation).

The common feature of these spectra is the fact that the dispersion curve does not exhibit a plateau at low frequencies so that the static value could not be determined directly. Moreover, the ϵ' values seem to be too large which especially concerns sample 3. Similar effects were observed in other strongly polar LC substances as well [14]. The absorption spectra of sample 4 consist of one Debye-type relaxation process whereas for sample 3 an additional low frequency process is present which can be attributed to the double-layer relaxation caused by free charges. In that situation we decided to establish the static permittivity values of all four substances in an indirect way by measuring their 10% wt. solutions in the **Base 903** mixture (Table 1) which exhibits the small negative dielectric

anisotropy $\Delta\epsilon = \epsilon_{||} - \epsilon_{\perp} < 0$, see Figure 6a, bottom. In Figure 6a the results of the measurements for all mixtures are presented. Their clearing points are given in Table 2. Assuming that both components of the mixtures contribute additively to the permittivity, the values ϵ_l , $\epsilon_{||}$, and $\epsilon_{\perp}$ characterizing the pure substances were calculated, see Figure 6b.

Relatively low concentrations of strongly polar molecules in the low polar solvent allow one to assume that the permittivity values displayed in Figure 6b correspond to the non-associated state of substances. In real substances a strong association of molecules can be expected which, besides of the conductivity effects, should change the permittivity values considerably. This is seen well in Figure 7 where compar-

Table 2. Dipole moment components and the angles between different molecular axes as explained in the text and Figure 10.

No	μ [D]	μ_x [D]	μ_y [D]	μ_z [D]	β_{chem}	$\Delta\beta$	β_{cal}	β_{Obs}	μ_l [D]	μ_t [D]
1	8.880	−8.258	−3.142	−0.882	21.56	4.13	25.69	26.34	8.00	3.85
2	8.959	−8.340	−3.155	−0.866	21.42	4.94	26.36	27.60	8.03	3.98
3	11.826	−10.704	−4.579	2.074	25.16	3.78	28.94	—	10.35	5.72
4	11.893	−10.739	−4.718	1.968	25.45	2.49	27.95	—	10.51	5.57

Fig. 6. (a) Tensor permittivity components measured for **Base 903** and its four 10% wt. mixtures, **B1** to **B4**, with particular substances (corresponding to 0.09577, 0.08532, 0.07560, and 0.06234 of mole fraction for compounds **1** to **4**, respectively). (b) Tensor permittivity components calculated from the measurements in mixtures for the four substances.

isons of the experimental and calculated permittivity values are presented. (In the case of compound **4** the static permittivity values were obtained from the analysis of the relaxation spectra by fitting the semicircle to the points measured above c. 100 kHz as is shown in Figure 8a). Only for compound **2** the agreement of both sets of data is fairly good. Relations between the measured and the calculated permittivities for compounds **1** and **4** are opposite which indicates the presence of parallel (**1**) and antiparallel (**4**) correlations in pure compounds. More information about those effects can be achieved if the measurements are made for several concentrations of mixtures.

4. Discussion

4.1. Static Permittivities

Several dipolar groups constitute the molecules under study. Figure 10 shows the molecule **1** simu-

lated with the aid of HyperChemTM Release 7.51 and CS Chem3D Pro with CS MOPAC ProTM software, based on the semi-empirical AM1 method [15]. These enabled us to calculate the dipole moment components (gathered in Table 2), to determine the principal inertia moments axes (marked as 1, 2, 3 in Fig. 9) and to estimate the angle β_{cal} between the net dipole moment and the long molecular axis (the 1-axis), as well as the angle β_{chem} between the net dipole moment and the *x*-axis defined as the *para*-axis of the phenyl ring with the butyloxy chain. As can be seen in Figure 9, the *para*-axis differs from the 1-axis by the angle $\Delta\beta$.

The angle β was also estimated on the basis of the experimental data. From the TDS spectra of compounds **1** and **2** presented in Figure 2 the parameters ϵ_s and ϵ_{∞} for two relaxation processes were calculated using formula (1). By applying the Onsager equa-

Fig. 7. Comparison of the tensor permittivity values for three substances obtained experimentally with those calculated from their 10% wt. mixtures in **Base 903**.

tion [16],

$$\frac{(\varepsilon_s - \varepsilon_\infty)(2\varepsilon_s + \varepsilon_\infty)}{\varepsilon_s(\varepsilon_\infty + 2)^2} = \frac{N_0}{3\varepsilon_0} \frac{\mu^2}{3kT}, \quad (2)$$

and taking into account the fitting parameters $\varepsilon_{s,lf}$ and $\varepsilon_{\infty,lf}$, governed by the longitudinal component μ_l , and $\varepsilon_{s,hf}$ and $\varepsilon_{\infty,hf}$, governed by the transverse component μ_t separately, we were able to calculate the angle $\beta_{Ons} = \tan^{-1}(\mu_t/\mu_l)$. The obtained values β_{Ons} are in good agreement with β_{cal} (Table 2).

As it is seen in Figure 1 and Table 3 the permittivity values differ considerably between substances **1** and **2** in spite of close dipole moments values (Table 2). The compound **2** may be treated as internal diluted medium because of its longer alkoxy chain. In the solution this effect is compensated by the low polar solvent. Therefore, the sequences of permittivities in the isotropic and N phases obtained from the measurements in solutions (Figure 6b) seem to be in accordance with the dipole moment values of particular substances.

Those behaviours of the permittivities may be discussed on the basis of the Onsager equation (2) (isotropic phase) or the Maier and Meier equations (3)–(5) [17] (nematic phase),

$$(\varepsilon_{||} - 1) = \varepsilon_0^{-1} N F h \left\{ \bar{\alpha} + \frac{2}{3} \Delta \alpha S + F \frac{\mu^2_{eff}}{3kT} [1 - (1 - 3 \cos^2 \beta) S] \right\}, \quad (3)$$

$$(\varepsilon_{\perp} - 1) = \varepsilon_0^{-1} N F h \left\{ \bar{\alpha} - \frac{1}{3} \Delta \alpha S + F \frac{\mu^2_{eff}}{3kT} \left[1 + \frac{1}{2} (1 - 3 \cos^2 \beta) S \right] \right\}, \quad (4)$$

$$\Delta \varepsilon = (\varepsilon_{||} - \varepsilon_{\perp}) = \varepsilon_0^{-1} N \cdot F \cdot h \cdot \left[\Delta \alpha - F \frac{\mu^2_{eff}}{2kT} (1 - 3 \cos^2 \beta) \right] S, \quad (5)$$

where $N = N_A \rho / M$ with N_A the Avogadro's number, ρ the density, M the molar mass, ε_0 is the permittivity of free space, $\Delta \alpha = \alpha_l - \alpha_t$ and $\bar{\alpha} = (\alpha_l + 2\alpha_t)/3$, S is the nematic order parameter. The local field parameters F and h are expressed by the mean polarizability $\bar{\alpha}$ and mean permittivity $\bar{\varepsilon} = (\varepsilon_{||} + 2\varepsilon_{\perp})/3$. $\mu^2_{eff} = g\mu^2$ is the effective dipole moment, and g is the dipole-dipole correlation factor. The differences in the observed ε -values (Fig. 1) cannot be attributed to the differences in the density values that change between the members of the homologous series rather slightly: the ratio $\rho_4/\rho_8 \sim 1.04$ [18], whereas the ratio $\Delta \varepsilon_4/\Delta \varepsilon_8 \sim 1.6$ (see in the inset of Fig. 1) and $\varepsilon_{l4}/\varepsilon_{l8} \sim 1.4$ in the isotropic phase. The change in the length of the butyl and octyl tails yields reduction in the ε_l and $\Delta \varepsilon$ values by factor ~ 1.2 [18]. A possible difference in

Fig. 8. Cole-Cole plots for the spectra of compound **4** in the nematic phase measured for: (a) the pure substance, (b) the 10% wt. mixture of compound **4** in the **Base 903**. The static $\epsilon_{||s}$ values presented in Figure 7 were taken from fitting the Debye semi-circle to the spectra as indicated in (a).

Fig. 9. Molecular structure of the substance **1** obtained with the aid of the HyperChemTM software. The reference axes and the angles listed in Table 2 are shown.

the nematic order parameter values cannot be responsible for that because the effect is pronounced in the isotropic phase as well. Taking the above into consideration, as well as the relations between the data obtained from the mixtures (Fig. 7), one can suppose that the observed effects must be attributed to the dipole-dipole correlations which seem to be parallel ($g > 1$) for substance **1** and antiparallel ($g < 1$) for substance **4**, with no correlations, perhaps, for substance **2**. The models describing different molecular correlations in LC phases can be found in [19–23]. In many works it was established that antiparallel pairing (dimerization) strongly increases with alkyl (hydrocarbon part) length of molecule, see example in [24, 25]. In the case of compounds having smectic A phase, transformation of smectic A_1 subphase to smectic A_d subphase is observed with growing alkyl chain [26]. In the near future the dielectric measurements for several concentrations of substances **1** and **4** in a non-polar nematogen will be performed in order to analyze these intermolecular interactions.

4.2. Dielectric Relaxation

In the isotropic phase of two-ring compounds two relaxation modes were observed by the impedance

Fig. 10. Longitudinal relaxation times vs. inverted temperature determined for pure substances (full symbols) and their mixtures with the **Base 903** (open symbols).

(low temperatures) and TDS (high temperatures) methods, see Figure 2. These modes can be attributed to the molecular rotations around the principal inertia axes. Superposition of two Debye-type formulae (1) fits the spectra well (especially for substance **1**). It is rather unusual that these spectra are shifted to the

Fig. 11. Dispersion spectra recorded for two alignments of the nematic phase in pure compounds **1** (a) and **4** (b), and compounds **1–4** in the **Base 903** mixtures (c) at $\Delta T = 30$ K. The crossover frequencies are indicated by arrows. (d) Temperature dependence of the cross-over frequency in one mixture in two scales; linear dependence f_{co} vs. $1/T$ indicates its exponential character.

that in the substance **1** strong intermolecular interactions hinder the molecular motions around the long axes as well.

4.3. Cross-Over Effect

Figure 11 shows the dispersion spectra plotted for two orientations of the nematic phase in two pure substances (a, b) and in four mixtures (c). At a certain frequency, called the crossover frequency f_{co} , the dielectric anisotropy changes sign from positive at low frequencies to negative above f_{co} . The values of both anisotropies are listed in Table 3. This effect enables designing the dual addressing displays. The here investigated alkoxy compounds have higher f_{co} value than recently studied analogous alkyl derivatives [8,33]. Figure 11d shows that the crossover frequency exponentially depends on the temperature with the activa-

tion enthalpy slightly smaller than the value characterizing the flip-flop molecular motion in the N phase.

5. Conclusions

The four studied nematogenic compounds possess large longitudinal and transverse dipole components which make them interesting for the formulation of mixtures useful for some application. However, very large dielectric anisotropy in the nematic phase causes a dissociation of polar bonds and a distinct ionic conductivity appears which influences the dielectric properties of pure substances considerably. It was shown that the dielectric measurements of such substances diluted in low-polar medium can bring valuable data on the tensor permittivity components in the nematic phase. Differences in the dielectric properties of substances in the pure and diluted states were interpreted

in terms of strong dipole-dipole interactions, especially in the case of two-ring compounds. Also the rotational dynamic properties of molecules provide evidence of such interactions in the isotropic and nematic phases of these compounds. More information about those inter-

actions can bring further investigations performed for several concentrations of mixtures.

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