Electric field induced inversion of the sign of half-integer disclinations in 2D nematic liquid crystals

P.P. Avelino,1,2,3 F. Moraes,3 J.C.R.E. Oliveira,1,4 and B. F. de Oliveira1,2,3

1Centro de Física do Porto, Rua do Campo Alegre 687, 4169-007 Porto, Portugal
2Departamento de Física da Faculdade de Ciências da Universidade do Porto, Rua do Campo Alegre 687, 4169-007 Porto, Portugal
3Departamento de Física, Universidade Federal da Paraíba, POB 5008 João Pessoa, PB 58051-970, Brazil
4Departamento de Engenharia Física da Faculdade de Engenharia da Universidade do Porto, Rua Dr. Roberto Frias, s/n, 4200-465 Porto, Portugal

We study the effect of the rotation of an external electric field on the dynamics of half-integer disclination networks in two dimensional nematic liquid crystals with a negative dielectric anisotropy using LICRA, a Liquid Crystal Algorithm developed by the authors. We show that a rotation of \( \pi \) of the electric field around an axis of the liquid crystal plane continuously transforms all half-integer disclinations of the network into disclinations of opposite sign via twist disclinations. We also determine the evolution of the characteristic length scale, thus quantifying the impact of the external electric field on the coarsening of the defect network.

I. INTRODUCTION

Topological defects play a fundamental role in condensed matter physics [1, 2] and cosmology [3]. Liquid crystals are an example of a very rich environment where the dynamics of topological defects can be realized experimentally at relatively low costs, making them ideal laboratories for testing different scenarios for defect formation and evolution [4–12].

External fields, such as electric and magnetic fields, can have a significant impact on the orientational order of liquid crystals [13–18] thus affecting the coarsening dynamics of disclination networks. Recently it was shown that the orientation of the applied electric field and the sign of the LC dielectric anisotropy may be used to control the type and topological charge of disclinations in two dimensional nematic liquid crystals [19]. In [17] a continuous transformation of some \(-\frac{1}{2}\) wedge disclination lines into \(\frac{1}{2}\) ones was numerically simulated on a cholesteric blue phase of a chiral liquid crystal, through the application of a constant electric field oblique to the line wedge. This provided a numerical realization of a continuous transformation between half-integer disclinations with topological charge of opposite sign [20].

In the present work we provide a numerical realization of the inversion of the sign of all half-integer disclinations on a 2D nematic LC with a negative dielectric constant induced by the rotation of an external electric field. In our implementation all half-integer disclinations of the network transform into disclinations of opposite sign via twist disclinations. We also consider other dynamical effects associated with the application of the external electric field on the nematic, by comparing the coarsening of the network under the rotation of the electric field with its evolution under a transformation of the director profile implemented by hand.

The paper is organized as follows. In Sec. [1] the equations describing the relaxational dynamics of nematic LC, in terms of a symmetric, traceless order parameter \(Q_{\alpha\beta}\), are presented. The numerical techniques are discussed in Sec. [11]. In Sec. [15] the effect of the rotation of an external electric field on the evolution of half-integer disclination networks on a two-dimensional nematic LC with a negative dielectric anisotropy is studied in detail. The conclusions and final remarks are presented in Sec. [19].

II. THE MODEL

The orientational order of a nematic LC without intrinsic biaxiality is described by a symmetric traceless tensor, \(Q_{\alpha\beta}\), at every point in space, whose components are given by [1]

\[
Q_{\alpha\beta} = \frac{3}{2} S(n_\alpha n_\beta - \frac{1}{3} \delta_{\alpha\beta}) + \frac{1}{2} T (l_\alpha l_\beta - m_\alpha m_\beta) , \tag{1}
\]

where the unit vector \(n\) is the director, determining the local average orientation of the molecules, \(l\) is the codirector, associated with the direction of orientational order perpendicular to \(n\) and \(m = n \times l\). The variables \(S\) and \(T\) represent the strength of uniaxial and biaxial ordering, respectively. The values of \(S\) and \(T\) may be found by the diagonalization of the matrix

\[
Q_{\alpha\beta} = \begin{pmatrix}
-(S + T)/2 & 0 & 0 \\
0 & -(S - T)/2 & 0 \\
0 & 0 & S
\end{pmatrix} , \tag{2}
\]

in a coordinate system where \(n = (0, 0, 1)\), \(l = (0, 1, 0)\) and \(m = (1, 0, 0)\).

Static equilibrium can only be reached for a minimum value of the free energy \((\delta F/\delta Q_{\alpha\beta} = 0)\). However, the evolution of the order parameter \(Q_{\alpha\beta}\) from a given set of initial conditions is not fully specified by the free energy functionals, and further assumptions have to be made on how the minimization process will take place. In the absence of thermal fluctuations and hydrodynamic flow, the time evolution of the order parameter is given by [19]

\[
\dot{Q}_{\alpha\beta}(r, t) = -\Gamma_{\alpha\beta\mu\nu} \frac{\delta F}{\delta Q_{\mu\nu}} . \tag{3}
\]

Here the dot represents derivative with respect to the physical time, \(t\), and the tensor

\[
\Gamma_{\alpha\beta\mu\nu} = \Gamma \left( \delta_{\alpha\mu} \delta_{\beta\nu} + \delta_{\alpha\nu} \delta_{\beta\mu} - \frac{2}{3} \delta_{\alpha\beta} \delta_{\mu\nu} \right) , \tag{4}
\]

satisfies \(\Gamma_{\alpha\beta\mu\nu} = \Gamma_{\beta\alpha\mu\nu} = \Gamma_{\mu\alpha\beta\nu} \) and \(\Gamma_{\alpha\alpha\mu\nu} = 0\) thus ensuring that the order parameter \(Q_{\alpha\beta}\) remains symmetric and traceless. In the following we shall assume that the kinetic coefficient, \(\Gamma\), is a constant.
The order parameter form a local expansion in rotationally invariant powers of the nematic-isotropic phase transition and it may be obtained from a grid point, at \( t = 499 \), only half-integer disclinations remain due to application of an external electric field along the \( z \) direction.

The free energy can be written as

\[
F = \int d^3r \left( F_b + F_{el} + F_E \right).
\]

(5)

The first term, \( F_b \), is the bulk free energy density. It describes the nematic-isotropic phase transition and it may be obtained from a local expansion in rotationally invariant powers of the order parameter

\[
F_b = \frac{A}{2} \text{Tr} Q^2 + \frac{B}{3} \text{Tr} Q^3 + \frac{C}{4} (\text{Tr} Q^2)^2.
\]

(6)

The second term, \( F_{el} \), is the elastic free energy density. Using the one elastic constant approximation the elastic free energy density is given by

\[
F_{el} = \frac{R}{2} \partial_\alpha Q_{\beta\gamma} \partial_\alpha Q_{\beta\gamma},
\]

(7)

where the constant \( R \) is the single elastic constant. The last term in the free energy functional, \( F_E \), is the contribution of the effect of an external electric field \( E \),

\[
F_E = -\frac{\tau_E}{2} Q_{\alpha\beta} E_\alpha E_\beta,
\]

(8)

where \( \tau_E = 2 \Delta \epsilon \) and \( \Delta \epsilon \) is the dielectric anisotropy. The molecules of a LC with positive dielectric constant, \( \Delta \epsilon > 0 \), tend to orient parallel to the external electric field, \( E \). On the other hand, if the LC has a negative dielectric constant, \( \Delta \epsilon < 0 \), then the molecules tend to be align perpendicularly to \( E \).

With this free energy the equation of motion can be written more explicitly as

\[
\dot{Q}_{\alpha\beta} = -\Gamma \left[ (A + C \text{Tr} Q^2) Q_{\alpha\beta} + B Q_{\alpha\gamma} Q_{\beta\gamma} - R Q_{\alpha\beta,\gamma\gamma} - \frac{\tau_E}{2} E_\alpha E_\beta \right],
\]

(9)

where a comma denotes a partial derivative and \( \overline{X_{\alpha\beta}} \) setting an arbitrary real matrix, \( X_{\alpha\beta} \), to be a symmetric traceless matrix by the following procedure

\[
\overline{X_{\alpha\beta}} = \frac{1}{2} \left( X_{\alpha\beta} + X_{\beta\alpha} \right) - \frac{1}{3} \delta_{\alpha\beta} X_{\gamma\gamma}.
\]

(10)

III. NUMERICAL IMPLEMENTATION

All the numerical simulations were performed with LICRA (LIquid CRystal Algorithm), a publicly available set of C codes and MATLAB/OCTAVE routines used to solve Eq. (9) using a standard second-order finite difference algorithm for the spatial derivatives and a second order Runge-Kutta method for the time integration. This software is free and it is available at http://faraday.fc.up.pt/licra [18].

The order parameter \( Q \) has 5 degrees of freedom associated with \( S, T, \mathbf{n} \) and \( I \) (\( \mathbf{n} \) accounts for two degrees of freedom). The initial conditions for \( S \) and \( T \) are randomly generated, at every grid point, from uniform distributions in the intervals \([0, 2/3]\) and \([0, 3S]\), respectively. The director, \( \mathbf{n} \), is also randomly generated, at every grid point, using the spherical vector distributions routines in the GNU Scientific Library (GSL) and the codirector, \( I \), was calculated by randomly choosing a direction perpendicular to \( \mathbf{n} \). The eigenvalues and eigenvectors in LICRA are computed using the library GSL.

In order to determine the evolution of the characteristic scale of the network LICRA numerically calculates the correlation function in Fourier space

\[
P(k, t) = \frac{Q_{\alpha\beta}(k, t) Q_{\beta\alpha}(-k, t)}{\int d^3k Q_{\alpha\beta}(k, t) Q_{\beta\alpha}(-k, t)},
\]

(11)

after setting the infinite wavelength mode \( Q_{\alpha\beta}(0) \) to zero. Here \( Q_{\alpha\beta}(k) \) is the Fourier transform of \( Q_{\alpha\beta}(r) \) (\( k \) is the
After that, three different simulations have been performed: only half-integer disclinations remain in the simulation \cite{18}. The electric field was applied at \(t = 500\) along the \(x\) direction and then rotates around the \(x\) direction and then rotates around the \(x\) direction by an angle of \(\varphi\), on the \(yz\) plane. At each time step (\(\Delta t = 0.1\)) the angle, \(\varphi\), was increased by \(\pi/300\) radians until a total angle of \(\pi\) (at \(t_{\text{off}} = 800\)). The negative dielectric anisotropy, \(\Delta \epsilon < 0\), ensures that the director remains perpendicular to the field ensuring that \(\mathbf{n} \cdot \mathbf{E} = 0\). If the defect centers were static then, at each time-step, the components of the director would be equal to

\[
\begin{align*}
\varphi &= \frac{\pi}{300} (t - 500) .
\end{align*}
\]

The electric field was applied at \(t_{\text{on}} = 500\) along the \(z\) direction and then rotates around the \(x\) direction by an angle of \(\varphi\), on the \(yz\) plane. At each time step (\(\Delta t = 0.1\)) the angle, \(\varphi\), was increased by \(\pi/300\) radians until a total angle of \(\pi\) (at \(t_{\text{off}} = 800\)). The negative dielectric anisotropy, \(\Delta \epsilon < 0\), ensures that the director remains perpendicular to the field ensuring that \(\mathbf{n} \cdot \mathbf{E} = 0\). If the defect centers were static then, at each time-step, the components of the director would be equal to

\[
\begin{align*}
n_x &= n_{x_0} , \\
n_y &= n_{y_0} \cos(\varphi) , \\
n_z &= -n_{y_0} \sin(\varphi) ,
\end{align*}
\]

where \((n_{x_0}, n_{y_0}, 0)\) are the components of the director at \(t = 500\) in simulation A. In practice this result must be complemented with the coarsening dynamics of the network.

Fig. 3 presents the order parameter \(S\) and the projection of the director, \(\mathbf{n}\), onto the two dimensional grid for a snapshot taken from simulation \(B\), at \(t = 650\), when \(\varphi = \pi/2\). In this case, the director is given by \(\mathbf{n} = (n_{x_0}, 0, -n_{y_0})\) and twist disclinations are formed on the \(xz\) plane. Note that the presence of the electric field increases the size of the defect cores.

Fig. 4 shows a snapshot of simulation \(B\) for \(t = 900\) after the electric field has been rotated by an angle of \(\pi\) and then switched off when at \(t = 800\). In this case, the components of the director are given by \(\mathbf{n} = (n_{x_0}, -n_{y_0}, 0)\) and all the half-integer disclinations are back into the \(xy\) plane, with an opposite topological charge to the one they had in Figure.
A similar realization of the exchange of sign of all half-
integer disclinations network is possible through the rotation
of the electric field around any other axis of the simulation
plane. Besides this transformation of the topological charge,
the electric field produces other effects on the network coars-
ening that need to be addressed in more detail.

In simulation $C$ we have implemented by hand the modi-
fication of the director and co-director profiles corresponding
to a rotation of $\pi$ of an external electric field. This transforma-
tion was performed with the purpose of isolating the effect of
the inversion of the sign of the half-integer disclinations from
the modification of the coarsening dynamics associated with
the presence of an external electric field. We have modified
the components of $n$ and $I$ of the configuration at $t = 500,$
shown in Fig. 1, so that

$$
\begin{align*}
n_x &= n_{x0}, & n_y &= -n_{y0}, \\
l_x &= n_{y0}, & l_y &= n_{x0},
\end{align*}
$$

(16)

where $(n_{x0}, n_{y0}, 0)$ are the components of the director $n$
presented in Fig. 1. The results for the projection of the director
onto the lattice plane are presented in Fig. 4 along with the
order parameter $S$ for a snapshot of simulation $C$ at $t = 900.$
Fig. 4 is very similar to Fig. 1 except for the inversion of the
sign of the topological charges and the coarsening which took
place between $t = 499$ and $t = 900.$

The evolution of the characteristic length, $L(t),$ of simu-
lations $A,$ $B$ and $C$ is presented in Figure 5. Given that the
curves for simulations $A$ and $C$ are almost indistinguishable
we intercalate the corresponding datapoints in order to im-
prove the visualization. The effect of the rotation of the elec-
tric field is visible in simulation $B,$ between $t = 500$ and
$t = 800.$ An acceleration of the coarsening, reflected in
the larger slope of $L^2(t)$ may be observed between $t = 500$
($\varphi = 0$) and $t = 650$ ($\varphi = \pi/2$). It is also clear that the mem-
ory of the connection of the field is preserved by the coarsen-
ing dynamics in simulation $B,$ even after the electric field is
disconnected. This is reflected in the larger value of the char-
acteristic length scale of the network at late time, compared to
simulations $A$ and $C.$

The comparison of the evolution of $L^2(t)$ for simulations
$A$ and $C$ demonstrates that modifying the director and co-
director profiles by hand using Eq. (16) does not modify the
coarsening dynamics. This happens because the bulk and elas-
tic components of the free energy defined by Eqs. (6) and
(7), are invariant by a rotation of the directors around an ar-
bitrary axis. Such rotation originates a rotated director and
co-director network profile with the same free energy at every
point in space. This result can be confirmed in Fig. 6 which
shows the total free energy, given by Eq. (5), for a snapshot of simulation $A$ at $t = 900$ (the corresponding image for sim-
ulation $C$ is identical). As expected, the free energy profile is
axially symmetric around the defect centers. Hence, the ac-
celeration of the defect coarsening due to the application of
an external electric field is associated with the modification
of the degree of alignment of the molecules, reflected in the
modification of the values of the order parameters $S$ and $T,$
and it is therefore insensitive to the inversion of the sign of the
topological charges.

**V. CONCLUSION**

In this work we have numerically demonstrated that a ro-
rotation by $\pi$ of an external electric field around an axis of the
plane of a 2D nematic liquid crystal induces an inversion of the
sign of all half-integer disclinations, when a negative di-
electric anisotropy is considered. We have further analysed
the the impact of the external electric field on the coarsening
dynamics.

The inversion of the topological charge investigated in the
present paper may be realized experimentally in a simple
setup. The topological arguments do not depend on the model
parameters and apply to the different phases of a liquid crys-
Although we have assumed that the inversion was induced by the rotation of an external electric field, a magnetic field induced transition is also possible in the case of negative diamagnetic anisotropy.

ACKNOWLEDGMENTS

We thank CAPES, CNPq, REDE NANOBIOTEC BRASIL, INCT-FCx (Brazil) and FCT (Portugal) for partial financial support.

[1] P. G. de Gennes and J. Prost, *The Physics of Liquid Crystals* (Clarendon Press, Oxford, 1995), 2nd ed.
[2] M. Kleman and O. D. Lavrentovich, *Soft Matter Physics: An Introduction* (Springer, 2003).
[3] A. Vilenkin and E. P. S. Shellard, *Cosmic Strings and Other Topological Defects* (Cambridge University Press, Cambridge, 1994).
[4] I. Chuang, R. Durrer, N. Turok, and B. Yurke, Science 251, 1336 (1991), ISSN 0036-8075.
[5] I. Chuang, B. Yurke, A. N. Pargellis, and N. Turok, Phys. Rev. E 47, 3343 (1993).
[6] M. Zapotocky, P. M. Goldbart, and N. Goldenfeld, Phys. Rev. E 51, 1216 (1995).
[7] S. Digal, R. Ray, and A. M. Srivastava, Phys. Rev. Lett. 83, 5030 (1999).
[8] C. Denniston, E. Orlandini, and J. M. Yeomans, Phys. Rev. E 64, 021701 (2001).
[9] S. Dutta and S. K. Roy, Phys. Rev. E 71, 026119 (2005).
[10] A. de Lózar, W. Schöpf, I. Rehberg, D. Svenšek, and L. Kramer, Phys. Rev. E 72, 051713 (2005).
[11] H. Mukai, P. R. G. Fernandes, B. F. de Oliveira, and G. S. Dias, Phys. Rev. E 75, 061704 (2007).
[12] A. K. Bhattacharjee, G. I. Menon, and R. Adhikari, Phys. Rev. E 78, 026707 (2008).
[13] H. S. Kitzerow, Mol. Cryst. Liq. Cryst. 202, 51 (1991).
[14] I. Dierking, O. Marshall, J. Wright, and N. Bulleid, Phys. Rev. E 71, 061709 (2005).
[15] G. P. Alexander and D. Marenduzzo, EPL 81, 66004 (2008).
[16] J.-i. Fukuda, M. Yoneya, and H. Yokoyama, Phys. Rev. E 80, 031706 (2009).
[17] J.-i. Fukuda, Phys. Rev. E 81, 040701 (2010).
[18] B. F. de Oliveira, P. P. Avelino, F. Moraes, and J. C. R. E. Oliveira, Phys. Rev. E 82, 041707 (2010).
[19] P. G. de Gennes, Mol. Cryst. Liq. Cryst. 12, 193 (1971).