



Filling and wetting transitions of nematic liquid crystals on sinusoidal substrates

Chi Tuong Pham, José Manuel Romero-Enrique, Nuno M. Silvestre, Pedro Patrício

► To cite this version:

Chi Tuong Pham, José Manuel Romero-Enrique, Nuno M. Silvestre, Pedro Patrício. Filling and wetting transitions of nematic liquid crystals on sinusoidal substrates. *Physical Review E* , 2011, 10.1103/PhysRevE.84.021701 . hal-03863935

HAL Id: hal-03863935

<https://universite-paris-saclay.hal.science/hal-03863935>

Submitted on 21 Nov 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Filling and wetting transitions of nematic liquid crystals on sinusoidal substrates

P. Patrício,^{1,2,*} N. M. Silvestre,² C.-T. Pham,³ and J. M. Romero-Enrique⁴

¹*Instituto Superior de Engenharia de Lisboa, Rua Conselheiro Emídio Navarro 1, P-1959-007 Lisboa, Portugal*

²*Centro de Física Teórica e Computacional, Universidade de Lisboa, Avenida Professor Gama Pinto 2, P-1649-003 Lisboa Codex, Portugal*

³*Laboratoire d'Informatique pour la Mécanique et les Sciences de l'Ingénieur, CNRS-UPR 3251, Université Paris-Sud 11, BP 133, F-91403 Orsay Cedex, France*

⁴*Departamento de Física Atómica, Molecular y Nuclear, Area de Física Teórica Universidad de Sevilla, Apartado de Correos 1065, ES-41080 Sevilla, Spain*

(Received 31 May 2011; published 1 August 2011)

Close to sinusoidal substrates, simple fluids may undergo a filling transition, in which the fluid passes from a dry to a filled state, where the interface remains unbent but bound to the substrate. Increasing the surface field, the interface unbinds and a wetting transition occurs. We show that this double-transition sequence may be strongly modified in the case of ordered fluids, such as nematic liquid crystals. Depending on the preferred orientation of the nematic molecules at the structured substrate and at the isotropic-nematic interface, the filling transition may not exist, and the fluid passes directly from a dry to a complete-wet state, with the interface far from the substrate. More interestingly, in other situations, the complete wetting transition may be prevented, and the fluid passes from a dry to a filled state, and remains in this configuration, with the interface always attached to the substrate, even for very large surface fields. Both transitions are observed only for a same substrate in a narrow range of amplitudes.

DOI: [10.1103/PhysRevE.84.021701](https://doi.org/10.1103/PhysRevE.84.021701)

PACS number(s): 61.30.Dk, 61.30.Hn, 61.30.Jf

I. INTRODUCTION

Wetting on smoothly structured substrates show a rich phenomenology. For simple fluids, the wetting behavior on structured substrates presents a variety of phenomena [1–3], for which several physical laws were theoretically proposed, as the *Wenzel* [4] or the *Cassie-Baxter* laws [5]. These laws extend *Young's law* [6] for planar substrates, including new substrate geometrical parameters. If these laws are justified in some situations [7], they may also fail in general theoretical or experimental conditions [8]. In addition to the usual wetting transition, simple fluids at bulk coexistence may present other surface transitions as the filling transition [9,10]. From a thermodynamical point of view, the filling transition in simple fluids always precedes a complete wetting transition. However, the filling transition may not exist for shallow substrates [9], but if it does exist, it always precedes the wetting transition.

The existence of long-range order in complex fluids may alter the scenario depicted above [11]. In a nematic liquid crystal, there is an orientational (continuous) long-range order. However, the presence of substrates that favor specific orientations frustrate the liquid crystal tendency to align along a given direction. As a consequence, elastic distortions emerge, altering the subtle balance between the different free-energy contributions that lead to the surface transitions on structured substrates.

In the past, we estimated the importance of the nematic elastic distortions and derived a generalized *Wenzel's law* [Eq. (12)] for very general smooth structured substrates [7]. We calculated numerically the wetting phase diagram for a periodic triangular (or sawtoothed) substrate and explained how the nucleation of defects, with noninteger topological

charges, near the singular points of the triangular substrate, leads to logarithmic contributions to the elastic part of the nematic free energies [7,13]. More recently, we reviewed and highlighted the differences and similarities of the wetting transitions of simple and nematic fluids for different types of periodic substrates, and reported some preliminary results for wetting of sinusoidal and rectangular substrates [11].

In this paper, we analyze in detail the wetting behavior of nematics on sinusoidal substrates (the rectangular substrate will be analyzed elsewhere). We show that the filling-wetting sequence that is observed in simple fluids on sinusoidal substrates may be deeply modified when we have a nematic liquid crystal instead of a simple fluid. Depending on the physical elastic parameters of the nematic, several scenarios may occur: We may have only a wetting transition; or a filling transition only, preventing the subsequent complete wetting transition; or the double sequence as in the simple fluid case. Both filling and wetting transitions are observed only for the same geometry in a narrow range of amplitudes.

This article is organized as follows: In Sec. II A we briefly review the main simple analytical results about filling and wetting transitions of simple fluids. In Sec. II B, we extend these results to nematic liquid crystal systems. In Sec. III, we use the Landau–de Gennes model of nematics to study their wetting behavior on a sinusoidal substrate. At the end, we draw some conclusions.

II. MACROSCOPIC ANALYSIS

A. Thermodynamic description of filling and wetting transitions for simple fluids

In this section, we review the simple description of filling and wetting transitions for simple fluids, following the ideas presented in Ref. [10]. To that end, let us consider a structured

*patricio@cii.fc.ul.pt

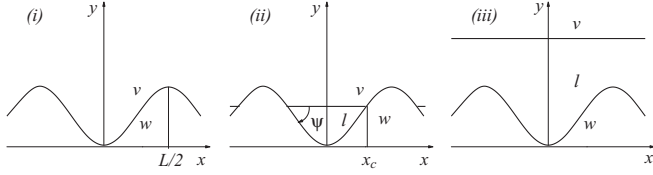


FIG. 1. Schematic representation of (i) dry, (ii) filled, and (iii) wet states of a simple fluid (l) on a sinusoidal substrate (w) of wave length L with vapor (v) above.

substrate, like the one presented in Fig. 1. The substrate is translationally invariant along z , with L_z being the total length along this direction, and periodic along the x axis with a period L . The substrate profile can be described by a simple even function $g(x)$. To simplify the analysis, we consider a monotonically increasing function for $0 < x < L/2$.

We suppose the substrate favors the liquid phase and consider three possible interfacial states: (1) dry, (2) filled, and (3) wet (Fig. 1). If the bulk free energies of the vapor and liquid phases are the same (so they can be set equal to zero), and if we neglect interactions between the interfaces, then the energies associated with each state will be simply the sum of surface energy terms. Let us denote $\mathcal{S}$ as the whole substrate area, $\mathcal{A}$ its projection on the x - z plane, and $\mathcal{F}$ the excess interfacial free energy. Then, the free energy of the dry state [Fig. 1(i)] is

$$\mathcal{F}^{(i)} = \mathcal{S}\sigma_{vw}, \quad (1)$$

where σ_{vw} is the vapor-wall (substrate) surface tension.

The free energy corresponding to the filled interfacial state [see Fig. 1(ii)] is

$$\mathcal{F}^{(ii)} = S(x_c)\sigma_{lw} + [\mathcal{S} - S(x_c)]\sigma_{vw} + \left(\frac{2x_c}{L}\right)\mathcal{A}\sigma_{lv}, \quad (2)$$

where σ_{lw} is the liquid-wall (substrate) surface tension, σ_{lv} is the liquid-vapor surface tension, and $S(x_c)$ is the substrate area in contact with the liquid phase, which can be obtained in terms of the abscissa x_c of the contact line of the filled region with the substrate as

$$S(x_c) = \mathcal{S} \frac{\int_{-x_c}^{x_c} \sqrt{1 + g_x^2} dx}{\int_{-L/2}^{L/2} \sqrt{1 + g_x^2} dx}, \quad (3)$$

where g_x is the derivative of g with respect to x . The value of x_c can be obtained by minimization of the free energy [Eq. (2)] with respect to it. Making use of Young's law, which relates the surface tensions mentioned above with the contact angle of a sessile drop of the liquid in a planar substrate θ_π ,

$$\sigma_{vw} - \sigma_{lw} = \sigma_{lv} \cos \theta_\pi, \quad (4)$$

it can be shown that the minimum energy condition is fulfilled when x_c satisfies [10]

$$\begin{aligned} 0 &= 2\sigma_{lv} \left[1 - \sqrt{1 + g_x^2(x_c)} \cos \theta_\pi \right] \\ &= 2\sigma_{lv} \left(1 - \frac{\cos \theta_\pi}{\cos \psi} \right), \end{aligned} \quad (5)$$

showing that the filling region must make contact with the substrate with an angle $\psi = \theta_\pi$. For smooth substrates,

this solution is a local minimum only if $g_{xx}(x_c) = d^2g/dx^2(x_c) < 0$.

Finally, the free energy for the wet state [see Fig. 1(iii)] is given by

$$\mathcal{F}^{(iii)} = \mathcal{S}\sigma_{lw} + \mathcal{A}\sigma_{lv}. \quad (6)$$

Different transitions between the different interfacial states can be observed. The transition from the dry state directly to the wet state is possible when the energies of these configurations are the same, that is,

$$\mathcal{F}^{(iii)} - \mathcal{F}^{(i)} = 0 = \mathcal{S}(\sigma_{lw} - \sigma_{vw}) + \mathcal{A}\sigma_{lv}. \quad (7)$$

Using Young's law [Eq. (4)], it is possible to deduce from Eq. (7) Wenzel's law for the dry-to-wet transition:

$$\frac{\mathcal{S}}{\mathcal{A}} \cos \theta_\pi = r \cos \theta_\pi = \cos \theta_r = 1, \quad (8)$$

where $r = \mathcal{S}/\mathcal{A} > 1$ is the roughness of the substrate. As $\theta_r \rightarrow 0$, a wetting transition may occur. However, this transition may be preempted by the presence of filled states. The filling or unbending transition between a dry and a filled state may occur if the free energies of these interfacial states are the same:

$$\mathcal{F}^{(ii)} - \mathcal{F}^{(i)} = 0 = S(x_c)(\sigma_{lw} - \sigma_{vw}) + \mathcal{A} \left(\frac{2x_c}{L} \right) \sigma_{lv}, \quad (9)$$

which leads to the equation

$$\frac{\int_{-x_c}^{x_c} \sqrt{1 + g_x^2} dx}{2x_c} \cos \theta_\pi = r_c \cos \theta_\pi = 1, \quad (10)$$

where $r_c > 1$. If this transition occurs at a lower temperature than the wetting transition predicted by the Wenzel law (i.e., for larger values of θ_π , as θ_π usually decreases with temperature), then a filling transition occurs and Wenzel's law does not hold. Under these conditions, macroscopics dictates that the wetting transition between a filled and a wet interfacial state must occur under the same conditions as in the planar substrate (i.e., $\theta_\pi = 0$). However, this prediction can be changed as interactions between the substrate and the liquid-vapor interface are taken into account. So, if there is a first-order wetting transition for the planar substrate, the wetting transition is still first order but shifted to larger values of θ_π (smaller temperatures) [10]. On the other hand, for planar continuous wetting its location and nature are unchanged [9].

B. Thermodynamic description of filling and wetting transitions for nematic fluids

If instead of a simple fluid under saturation conditions, we have a liquid crystal at the nematic-isotropic phase coexistence, some new phenomena will inevitably arise. Let us consider that the substrate favors the nematic phase at a particular anchoring condition. From a thermodynamical point of view, if the bulk free energies of the phases are the same, the total energy of the nematic interfacial state has a new positive extra term $\mathcal{F}_d > 0$, corresponding to the Frank elastic energy of the nematic distortions imposed by the anchoring at the substrate. Figure 2 shows schematically an example of a nematic phase in contact with a substrate with homeotropic anchoring conditions (however, the theoretical

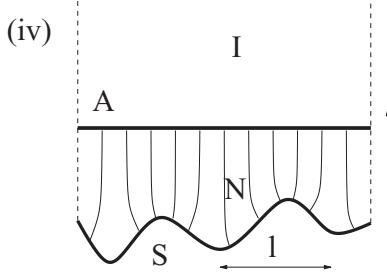


FIG. 2. Schematic representation of the nematic wet state on a rough substrate of typical l at isotropic-nematic phase coexistence.

arguments presented here are very general, and independent on the particular anchoring conditions involved). The total free energy of this interfacial state can be written as

$$\mathcal{F}^{(iv)} = S\sigma_{nw} + A\sigma_{in} + \mathcal{F}_d. \quad (11)$$

The elastic deformation creates an effective long-range repulsion between the substrate and the interface: The closer the interface is to the substrate, the more constrained the order will be, leading to higher energies. As the interface goes to infinity, the system will have only a characteristic length scale, which is imposed by the substrate (Fig. 2). If no defect arises, the elastic contribution may be simply estimated as $\mathcal{F}_d = (KS/l)\tilde{\mathcal{F}}_d$, where K is the Frank elastic constant, and $\tilde{\mathcal{F}}_d$ is a reduced elastic contribution to the interfacial free energy, dependent only on the substrate geometry, but not on its scale l [12]. In some cases, topological defects nucleate close to or on the substrate. The presence of defects imposed by the substrate introduce a $\ln(l/\xi)$ dependence on the reduced elastic contribution $\tilde{\mathcal{F}}_d$ [13]. This dependence is easily justified. If the defects are induced by the substrate, they will place themselves at a distance of order of l (the characteristic length of the substrate), usually close to its crests and/or troughs. On the other hand, the cutoff length is given by the size of the defects, which scales with the correlation length ξ .

The dry-to-nematic wet transition may be derived from the generalized Wenzel law [7]:

$$\mathcal{F}^{(iv)} - \mathcal{F}^{(i)} = 0 = S(\sigma_{nw} - \sigma_{iw}) + A\sigma_{in} + S\frac{K}{l}\tilde{\mathcal{F}}_d,$$

which can be rewritten as

$$r \cos \theta_\pi = 1 + \frac{rK\tilde{\mathcal{F}}_d}{l\sigma_{in}}. \quad (12)$$

For large enough substrate lengths, $l \gg K/\sigma_{in}$, the effects of elastic terms are unimportant, and Wenzel's law is recovered. Usually, K and σ_{in} are not independent, and their ratio scales with the correlation length ξ . In fact, using a Landau-de Gennes model for nematic liquid crystals (see next section), it is possible to show that $K/\sigma_{in} \sim 10\xi$ (at coexistence temperature).

For the simple fluid case ($\cos \theta_\pi = 1/r < 1$), the dry-to-wet transition for the rough substrate occurs always before the wetting transition for the planar substrate, as the surface field or the temperature is increased. In contrast, for the nematic case, the wetting transition for a rough substrate may occur either before (if the roughness is more important) or after the wetting transition for the planar substrate (if the elastic deformations

are more important). These deviations with respect to Wenzel's law are enhanced if defects are present, as the elastic term of the generalized Wenzel's law [Eq. (12)] will decay slower with increasing l . The sawtoothed substrate exemplifies this effect [13].

We now turn to the case of nematic filled states. Once again, it is useful to first consider configurations without defects, which do not appear if the substrate is shallow enough. Given the fact that the nematic-isotropic interface favors a particular nematic anchoring [14], if the interface is near the substrate, it will induce strong elastic deformations, because the nematic director is constrained to follow the anchoring conditions on both the interface and the substrate. Thus the elastic energy of the deformations may be large enough to prevent any filling transition, even if it would normally occur in the case of a simple fluid. It should also be pointed out that the interfacial free energies of the filled and wet states are normally very close to each other [10], and a very tiny perturbation can exchange their relative stability.

If the substrate imposes defects on the nematic matrix, a different picture emerges. The case of a sinusoidal grating is studied in detail in the next section. If the substrate is rough enough, and if the nematic-isotropic interface favors, for example, homogeneous planar anchoring, then the filled state (with the interface placed in between the crests of the sinusoidal substrate) may always have a smaller energy than the wet state, for in the latter case, a periodic array of defects is necessarily created at the top of each crest.

III. THE SINUSOIDAL SUBSTRATE: NUMERICAL RESULTS

In this section, in order to substantiate our arguments, let us consider a nematic liquid crystal in contact with a sinusoidal grating, characterized by a wave vector $q = 2\pi/L$ and an amplitude A (see also Fig. 1):

$$g(x) = A(1 - \cos qx). \quad (13)$$

At the substrate, the nematic molecules preferentially align homeotropically, i.e., perpendicularly to the substrate. The system is translationally invariant along the out-of-plane axis z and periodic along the x axis. Finally we impose that, far from the substrate, we have an isotropic ordering.

A. The Landau-de Gennes model

In the Landau-de Gennes (LdG) model, both isotropic and nematic phases can be locally represented by a traceless, symmetric order-parameter tensor with components:

$$Q_{ij} = \frac{3}{2}S\left(n_i n_j - \frac{1}{3}\delta_{ij}\right) + \frac{1}{2}B(l_i l_j - m_i m_j), \quad (14)$$

where n_i are the Cartesian components of the director field $\mathbf{n}$, S is the nematic order parameter that measures the orientational ordering along the nematic director, and B is the biaxiality parameter, which measures the ordering of the molecules on the orientations perpendicular to $\mathbf{n}$, characterized by the eigenvectors $\mathbf{l}$ and $\mathbf{m}$. In our model we will consider only in-plane deformations, although out-of-plane or twist deformations may also be important (a twist instability may

occur for particular choices of parameters [15]). In this case $\mathbf{n} = (\cos \theta, \sin \theta, 0)$, and the tensor order parameter has only three independent components (namely, Q_{11} , Q_{22} , and Q_{12}).

The LdG free energy may be written as

$$\mathcal{F}_{\text{LdG}} = \int_V (\phi_{\text{bulk}} + \phi_{\text{el}}) dV + \int_S \phi_{\text{surf}} dS, \quad (15)$$

where ϕ_{bulk} is the bulk free-energy density, ϕ_{el} is the elastic free-energy density, and ϕ_{surf} is the surface free energy, defined as [16]

$$\phi_{\text{bulk}} = a \text{Tr} \mathbf{Q}^2 - b \text{Tr} \mathbf{Q}^3 + c [\text{Tr} \mathbf{Q}^2]^2, \quad (16)$$

$$\phi_{\text{el}} = \frac{L_1}{2} \partial_k Q_{ij} \partial_k Q_{ij} + \frac{L_2}{2} \partial_j Q_{ij} \partial_k Q_{ik}, \quad (17)$$

$$\phi_{\text{surf}} = -\frac{2}{3} w \text{Tr} [\mathbf{Q} \cdot \mathbf{Q}_{\text{surf}}], \quad (18)$$

where a depends linearly on the temperature, b and c are positive constants, and L_1 and L_2 are positive parameters related to the elastic constants. If we rescale all the variables as follows [17]: $\tilde{\mathbf{Q}} = 6c\mathbf{Q}/b$, the positions $\tilde{\mathbf{r}} = \mathbf{r}/\xi$, where the correlation length ξ is defined as $\xi^2 = 8c(3L_1 + 2L_2)/b^2$, and $\tilde{\mathcal{F}}_{\text{LdG}} = 24^2 c^3 \mathcal{F}_{\text{LdG}} / \xi^3 b^4$, we get $\tilde{\mathcal{F}}_{\text{LdG}} = \int_{\tilde{V}} (\tilde{\phi}_{\text{bulk}} + \tilde{\phi}_{\text{el}}) d\tilde{V} + \int_{\tilde{S}} \tilde{\phi}_{\text{surf}} d\tilde{S}$, with rescaled free-energy densities:

$$\tilde{\phi}_{\text{bulk}} = \frac{2}{3} \tau \text{Tr} \tilde{\mathbf{Q}}^2 - \frac{8}{3} \text{Tr} \tilde{\mathbf{Q}}^3 + \frac{4}{9} [\text{Tr} \tilde{\mathbf{Q}}^2]^2, \quad (19)$$

$$\tilde{\phi}_{\text{el}} = \frac{1}{3 + 2\kappa} [\tilde{\partial}_k \tilde{Q}_{ij} \tilde{\partial}_k \tilde{Q}_{ij} + \kappa \tilde{\partial}_j \tilde{Q}_{ij} \tilde{\partial}_k \tilde{Q}_{ik}], \quad (20)$$

$$\tilde{\phi}_{\text{surf}} = -\frac{2}{3} \tilde{w} \text{Tr} [\tilde{\mathbf{Q}} \cdot \tilde{\mathbf{Q}}_{\text{surf}}]. \quad (21)$$

Here $\tau = 24ac/b^2$ is a dimensionless temperature, $\kappa = L_2/L_1$ is an elastic dimensionless parameter (for stability reasons, the elastic parameter is restricted to the values, $\kappa > -3/2$), and $\tilde{w} = 16wc/b^2\xi$ is the dimensionless anchoring strength. Hereafter, we will consider these rescaled expressions, so we will drop the tilde notation.

In our model we will place ourselves at coexistence temperature, $\tau = 1$, for which the bulk free-energy density has two minima, corresponding to $\phi_{\text{bulk}} = 0$ for rescaled scalar order parameters $S_I = 0$ (isotropic phase) and $S_N = 1$ (nematic phase). It is important to note that the order parameter S in the coexisting nematic phase is rescaled, so its value in real units is $b/6c$, which must be smaller than 1 (typically ≈ 0.4). If the elastic parameter κ is positive (negative), the nematic prefers to align parallel (perpendicular) to a possible nematic-isotropic interface. Finally, $\mathbf{Q}_{\text{surf}}$ defines the favored tensor at the substrate. We favor a homeotropic alignment of the nematic by setting $\mathbf{Q}_{\text{surf}} = (3\mathbf{v} \otimes \mathbf{v} - 1)/2$, with $\mathbf{v}$ the normal vector to the substrate, establishing a direct connection to previous papers [18–20].

B. Numerical procedure

For every set of model parameters, we numerically minimized the Landau–de Gennes free energy, using a conjugate-gradient method. The numerical discretization of the continuum problem was performed with a finite-element method combined with adaptive meshing in order to resolve the different length scales [21]. Due to the translational symmetry

along the z axis and the periodicity on the x axis, we restricted the minimization to a section of the system perpendicular to the z axis and with a width along the x axis equal to the period L of the substrate. We used periodic boundary conditions on the lateral sides, and the fluid on the substrate, which corresponds to the bottom boundary of our unit cell, does not have an imposed ordering or anchoring, although a nematic phase with homeotropic orientation is energetically favorable. Finally, we impose different fixed boundary conditions at the cell top (which we place at a large height H) in order to explore the different interfacial states we may observe. The value of H was varied until we found convergence in the free energy. In order to find the dry (D) or the filling (F) interfacial states, we fix a bulk isotropic phase on top of the cell. Under these conditions, different states may be obtained when we vary the initial condition (for example, by considering initial conditions with different heights for the nematic filling region). The complete wet (W) interfacial states are obtained by imposing a fixed nematic phase at the cell top with a homogeneous nematic director along either the x axis ($W^{\parallel}$) or the y axis ($W^{\perp}$) (see Fig. 4 for illustrations). The free energy for the wet states is calculated by adding to the numerically obtained free energy the contribution of an isotropic-nematic interface parallel to the x - z plane, $L\sigma_{\text{in}}$, calculated also numerically (and in some cases analytically) and corresponding to the most favored anchoring conditions (homeotropic to the interface for $\kappa < 0$, planar for $\kappa > 0$). In this way we consider the presence of a nematic-isotropic interface infinitely away from the substrate in the wet state, allowing us to neglect the elastic deformations, which may exist between the top cell and the nematic-isotropic interface. The true equilibrium state will be the state that gives the least free energy at the same thermodynamic conditions.

C. Numerical results

1. Case of $\kappa > 0$

As an example, in Fig. 3 we plot the free energy per projected unit area $f = \mathcal{F}/A$ corresponding to the most

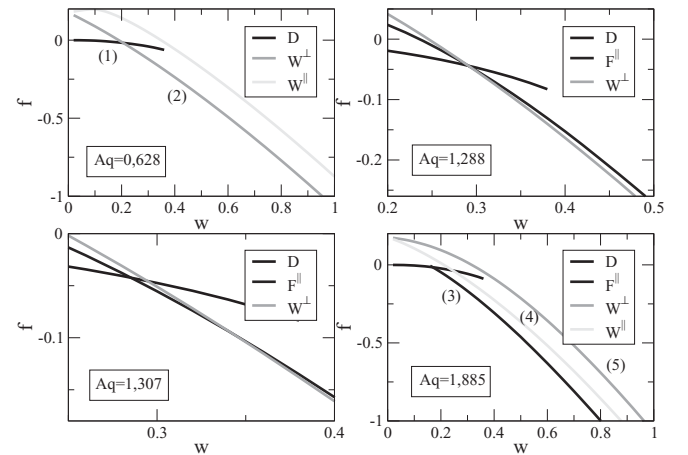


FIG. 3. Plot of the free energy densities (per projected unit area) of different branches of (meta)stable states, as a function of the anchoring strength w , for a sinusoidal substrate with $L = 10\xi$, $\kappa = 2$, and $Aq = 0.628, 1.288, 1.307$, and 1.885 . Numbers correspond to the typical configurations shown in Fig. 4. See text for explanation.

relevant interfacial states for $L = 10\xi$, $\kappa = 2$ (favoring a parallel anchoring at the nematic-isotropic interface) and different values of $Aq = 0.628, 1.288, 1.307$, and 1.885 . For the shallowest case $A = \xi$ we plot the branches corresponding to D states (which is the most stable for small w), $W^\perp$ states, and $W^\parallel$. No F states are observed. As it is expected, the $W^\parallel$ branch has always larger free energy than the others, since for shallow substrates the elastic deformations are stronger for this state than for the $W^\perp$ state. On the other hand, the D and $W^\parallel$ branches intersect at a first-order wetting transition for $w_W = 0.2056$, so $W^\perp$ branch is the most stable for $w > w_W$ [typical configurations for both D and $W^\perp$ states are shown in Figs. 4(1) and 4(2)].

For rougher substrates, the scenario is completely different (Fig. 3). For $Aq = 1.885$ four different (locally) stable branches are observed: D states (again the most stable for small w), $W^\perp$ states, $W^\parallel$ states, and a filled state where the nematic is planar at the nematic-isotropic interface. This new state will be denoted by $F^\parallel$ [a typical configuration is shown in Fig. 4(3)]. For small w , the nematic-isotropic interface is almost parallel to the x - z plane. As w increases, the position of the nematic-isotropic interface for the $F^\parallel$ state is lifted and curved slightly, and it eventually pins at the crests of the substrate. Regarding the complete wet states, the free energy of the $W^\perp$ branch is always larger than those of the $W^\parallel$ states (unlike for shallow wedges). Typical configurations are shown in Figs. 4(4) ($W^\parallel$) and 4(5) ($W^\perp$). Now roughness induces larger deformations in $W^\perp$, leading to a higher free energy. In both cases topological defects may nucleate either close to the substrate troughs ($W^\perp$ states) or the crests ($W^\parallel$ states). We can

see that the $F^\parallel$ branch has always a smaller energy than the wetting states, so there is a first-order filling transition between a D and $F^\parallel$ state at $w_F = 0.1693$. However, we do not observe a wetting transition, even for large w . This latter feature can be rationalized by the fact that the $F^\parallel$ state does not present topological defects as does the $W^\parallel$ state.

The crossover from the scenarios described above occurs at intermediate values of Aq . Above some value of Aq the branch of $F^\parallel$ emerges as a metastable branch. At $Aq \approx 1.288$ a triple point occurs, since the D , $F^\parallel$, and $W^\perp$ branches intersect at the same value of $w = 0.291$ (Fig. 3). Above this triple point, the $F^\parallel$ states branch crosses both the D and $W^\perp$ branches, so filling and wetting transitions are observed for the same geometry. However, the value of w for the wetting transition has a steep increase with Aq , so the wetting transition is no longer observable for Aq larger than 1.33 .

The global adsorption phase diagram for $\kappa = 2$ is shown in Fig. 5. The domains for the different possible stable phases (D , $W^\perp$, or $F^\parallel$) are divided by first-order transition lines that meet at a triple point. Note that the $F^\parallel$ - $W^\perp$ transition line is almost vertical, so the range in which both filling and wetting transitions are observed is quite narrow. As the substrate wavelength L increases, the importance of the elastic term of the nematic free energy diminishes, and the D - $W^\perp$ and D - $F^\parallel$ transition lines shift toward lower values of w , approaching the values obtained for simple fluids within the macroscopic theory outlined in Sec. II. However, due to the presence of elastic distortions, larger deviations than for simple fluids can be observed in all the cases,

For the D - $W^\perp$ wetting transition we can estimate this deviation through the modified Wenzel law prediction [Eq. (12)]. The different terms in this expression can either be evaluated exactly or estimated numerically. First, the nematic order parameter for the LdG model close to a planar substrate in the

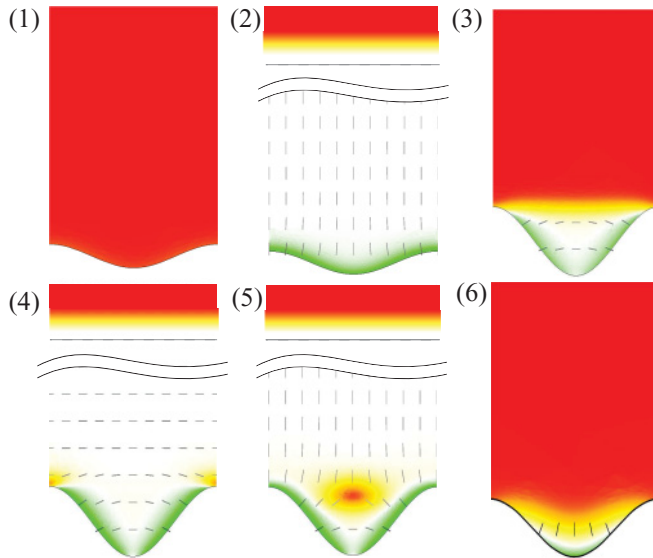


FIG. 4. (Color online) Typical configurations for a sinusoidal substrate with $L = 10\xi$: from left to right and top to bottom: (1) a D state ($\kappa = 2$, $Aq = 0.628$, and $w = 0.2$); (2) a $W^\perp$ state for shallow substrates ($\kappa = 2$, $Aq = 0.628$, and $w = 1.0$); (3) a $F^\parallel$ state ($\kappa = 2$, $Aq = 2.199$, and $w = 0.4$); (4) a $W^\parallel$ state for rough substrates ($\kappa = 2$, $Aq = 2.199$, and $w = 1.0$); (5) a $W^\perp$ for rough substrates ($\kappa = 2$, $Aq = 2.199$, and $w = 1.0$); and (6) a $F^\perp$ state ($\kappa = -1/2$, $Aq = 1.257$, and $w = 0.4$). Numbers correspond to those shown in Fig. 3.

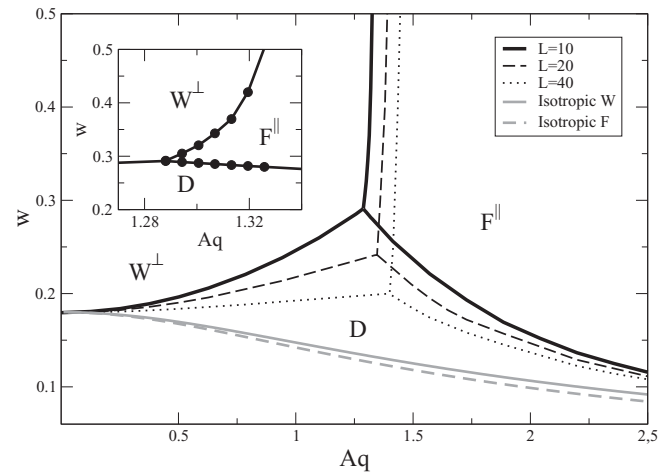


FIG. 5. Global phase diagram for nematic adsorption on sinusoidal substrates with $\kappa = 2$. Black lines correspond the phase boundaries between the D , $W^\perp$, and $F^\parallel$ states for $L = 10\xi$ (solid lines), $L = 20\xi$ (dashed lines), and $L = 40\xi$ (dotted lines). Gray lines correspond to the transition from dry to filled state (dashed line) and from filled to wet state (solid line) in the case of isotropic fluid. Inset: Zoom of the phase boundaries around the triple point for $L = 10\xi$.

presence of a bulk nematic or isotropic phase can be solved analytically [13]. The exact solutions of the corresponding Euler-Lagrange equations for the order parameter profiles, $d^2S/dy^2 = 2S - 6S^2 + 4S^3$, are given by

$$S_{\pm}(y) = \frac{S_{\pm}(0)}{S_{\pm}(0) + [1 - S_{\pm}(0)] \exp(\mp \sqrt{2}y)}, \quad (22)$$

where $S_+(y)$ and $S_-(y)$ are, respectively, the nematic order parameter profiles, as a function of the distance to the planar substrate y , corresponding to a bulk nematic or isotropic phase in bulk, and $S_{\pm}(0)$ are their values in contact with the substrate. The surface tensions between the planar substrate and the nematic or isotropic phase are given by

$$\begin{aligned} \sigma_{nw} &= \int_0^{\infty} [\phi_{el}(S_+) + \phi_{bulk}(S_+)] dy - wS_+(0) \\ &= \frac{\sqrt{2}}{6} [1 + 2S_+(0)][1 - S_+(0)]^2 - wS_+(0), \end{aligned} \quad (23)$$

$$\begin{aligned} \sigma_{iw} &= \int_0^{\infty} [\phi_{el}(S_-) + \phi_{bulk}(S_-)] dy - wS_-(0) \\ &= \frac{\sqrt{2}}{6} [3 - 2S_-(0)]S_-(0)^2 - wS_-(0). \end{aligned} \quad (24)$$

The nematic order parameter values at contact with the substrate can be obtained from the boundary condition $(dS_{\pm}/dy)(y=0) = -w$ as

$$S_{\pm}(0) = \frac{1}{1 \mp (\frac{1}{\sqrt{2}w} \pm 1) + \sqrt{(\frac{1}{\sqrt{2}w} \pm 1)^2 - 1}}. \quad (25)$$

The planar contact angle θ_{π} is obtained from Young's law:

$$\cos \theta_{\pi} = \frac{\sigma_{iw} - \sigma_{nw}}{\sigma_{in}}, \quad (26)$$

where $\sigma_{IN} \approx 0.178$ for $\kappa = 2$.

The roughness r of a sinusoidal substrate, is given in terms of the amplitude A and the wave number q as

$$r = \frac{2E[-(qA)^2]}{\pi}, \quad (27)$$

where $E(x)$ is the complete elliptic integral of second kind.

Finally, for the elastic term we have to resort to some numerical calculations. As a first estimate, we may use Berreman's approximation $\mathcal{F}_d \approx AK(Aq)^2q/4$ [12]. Figure 6 shows the comparison between the numerical and the modified Wenzel law predictions for the D - $W^{\perp}$ transition line when Berreman's approach for the elastic contribution is used. They are in qualitative agreement: For small L the wetting transition is shifted toward larger values of w , and as L increases, the transition line moves down, approaching Wenzel's prediction [Eq. (8)]. A similar behavior was observed for the wetting transition of the sawtoothed substrate [7,11]. However, there are quantitative discrepancies even for the largest system considered. The explanation for these discrepancies is that Berreman's expression is accurate only for small Aq and wL very large (strong anchoring conditions), but it overestimates the exact result of Frank theory [22]. So we have estimated the elastic energy obtained by using a Frank-Oseen (FO) model with a Rapini-Papoular surface contribution, where the interaction strength (which depends on w) is obtained from

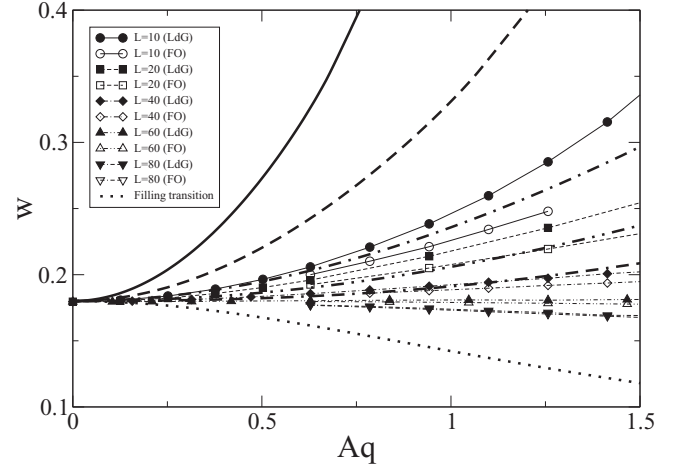


FIG. 6. Location of the D - $W^{\perp}$ wetting transitions for $\kappa = 2$ and different values of $L/\xi = 10$ (circles and solid lines), 20 (squares and dashed lines), 40 (rhombi and dot-dashed lines), 60 (upward triangles and dashed-double dot lines), and 80 (downward triangles and double dashed-dot lines). Symbols correspond to the numerical results from full minimization of the LdG model (filled symbols) and the FO model (open symbols). Thick lines indicate the predictions from the modified Wenzel law [Eq. (12)], using Berreman's approach for the elastic term. Thin lines serve only as guides for the eyes for the numerical results using the same style as the thick lines for the same parameter L . For comparison, the prediction from usual Wenzel's law [Eq. (8)] is shown (thick dotted line).

a constrained minimization of the LdG model in presence of a planar wall and a given nematic director orientation at a distance 1.5ξ [13]. We used a similar adaptative-mesh finite-element method to minimize the FO model to that explained above for the LdG model. Our results show that, for $Aq \sim 1$ and assuming strong anchoring conditions, the elastic contribution is reduced, taking a value of around 75% of that obtained from Berreman's expression. However, if weak anchoring is considered, a further reduction is observed, even for $L = 80\xi$. This fact contrasts with our previous results in the sawtoothed substrate, in which the strong anchoring regime is reached for relatively smaller values of L . When the numerically obtained values of $\mathcal{F}_d$ are used in Eq. (12), the predictions for the wetting transition from the generalized Wenzel law are in very good agreement with the full numerical values from the LdG model for $L \geq 60\xi$ (Fig. 6).

2. Case of $\kappa = 0$ or $\kappa < 0$

Now we turn to the effect of κ in the adsorption phase diagram. As was already stated, for $\kappa = 2$ the nematic fluid has a preferred planar anchoring at the interface. As κ decreases the free-energy cost to anchor the nematic in a parallel or perpendicular orientation approach each other and become identical for $\kappa = 0$. If we further decrease κ , the system starts to prefer a perpendicular orientation at the interface. As a consequence, new interfacial states emerge, as the filled state $F^{\perp}$ [Fig. 4(6)]. These new states appear at intermediate values of Aq for large w , and the NI interface shows a pronounced curvature following the substrate shape. The global adsorption phase diagrams for $L = 10\xi$ are shown in Fig. 7. Again, the

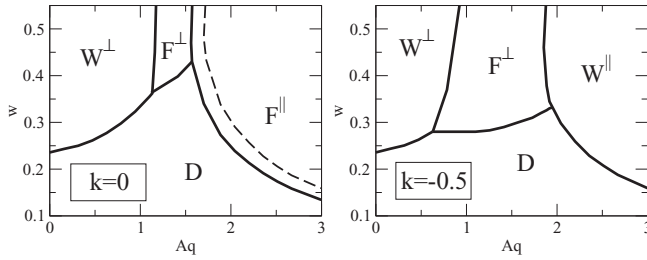


FIG. 7. Global phase diagram for $L = 10\xi$ and (left) $\kappa = 0$; (right) $\kappa = -1/2$. The dashed line on the left-hand side is the $D - W^\parallel$ metastable transition line.

continuous lines correspond to first-order transition lines that meet at triple points. For $\kappa = 0$, the region of the phase diagram corresponding to $F^\perp$ states is located between the regions corresponding to $W^\perp$ and $F^\parallel$ states, with almost vertical boundaries between them. The $D - F^\perp$ transition is weaker than the other transitions, so it is almost observed as a crossover for $L = 10\xi$. Finally, the $W^\parallel$ states' free-energy branch approach the $F^\parallel$ free-energy branch. The dashed line corresponds to the (metastable) $D - W^\parallel$ transition. For $\kappa = -1/2$, the $W^\parallel$ states become the stable states for large Aq and w . So the phase diagram is reminiscent of that for $\kappa = 0$, by swapping the $F^\parallel$ states with the $W^\parallel$ states. It is interesting to note that it is possible to observe reentrance of $F^\perp$ states as w is increased: The fluid undergoes a wetting transition from the $F^\perp$ state to the $W^\parallel$ state, and for larger values of w there is a reverse dewetting transition from the $W^\parallel$ to the $F^\perp$ state. However, this sequence of three transitions is limited to a very narrow range of values of Aq .

IV. CONCLUSIONS

In this article we have shown that the filling and wetting transition sequence of an isotropic fluid on a sinusoidal substrate is deeply changed for nematic liquid crystals. Substrates favoring perpendicular anchoring in contact with nematic liquid crystals, which favor parallel anchoring at the nematic-isotropic interface, exhibit only a wetting (and not filling) transition for small Aq , while for large Aq only filling (and not wetting) transitions occur. In the latter case, the complete wetting transition is prevented by the pinning of the nematic-isotropic interface at the crests of the sinusoidal substrate, thus hindering the nucleation of topological defects that would otherwise be present in the wet configuration at large enough anchoring strengths. When perpendicular

anchoring is favored at both the substrate and the interface, the filling and wetting transition lines change places. In this case the wetting transition may occur for larger Aq , while the filling transition occurs for intermediate Aq . Finally, in conditions such that the anchoring at the nematic-isotropic interface may be either planar or homeotropic, the phase diagram shows mixed features of the two previous cases. The analysis of these situations is far more complex than in the previous cases, as any perturbation can change the delicate balance between the different contributions to the free energy.

In order to check the validity of the phenomenological generalized Wenzel's law [Eq. (12)], we have compared its predictions with the numerical transition values for the one-step $D - W^\perp$ wetting transition. We see that they are in excellent agreement for $L \geq 60\xi$. This fact shows that elastic effects are important in the range of values of wavelength L considered in our work. In particular, they stabilize the D state with respect to the $W^\perp$ state, as elastic deformations are a positive contribution to the surface free energy present only in the latter state. On the other hand, the location of the wetting transition approaches the value predicted by the typical Wenzel's law [Eq. (8)] when L increases.

It is interesting to compare these results to those obtained for other substrates [7,11]. Unlike the sawtoothed or rectangular substrates, defects do not nucleate on the surface but may emerge in the nematic. The selection of defect-free states is always favored (the $W^\perp$ state for shallow wedges, the $F^\parallel$ state for rough substrates if $\kappa = 2$) and leads to the suppression of interfacial transitions that may exist for simple fluids. Due to the absence of topological defects, the finite-size effects on the wetting transitions are less important than for sawtoothed or rectangular substrates. Nevertheless, a Berreman-like elastic contribution to the free energy is present, and, consequently, wetting obeys a generalized Wenzel's law [Eq. (12)]. This is expected to be true in general for smooth substrates.

ACKNOWLEDGMENTS

We acknowledge the support from FCT (Portugal) through Grants No. PEst-OE/FIS/UI0618/2011, PTDC/FIS/098254/2008, and SFRH/BPD/40327/2007 (NMS) and Acção Integrada Luso-Espanhola Ref. E 17/09. JMR-E also acknowledges financial support from Spanish Ministerio de Ciencia e Innovación through Grants No. FIS2009-09326 and HP2008-0028, and Junta de Andalucía through Grant No. P09-FQM-4938.

- [1] C. Rascón and A. O. Parry, *Nature (London)* **407**, 986 (2000).
- [2] M. Calliès and D. Quéré, *Soft Matter* **1**, 55 (2005).
- [3] D. Bonn, J. Eggers, J. Indekeu, J. Meunier, and E. Rolley, *Rev. Mod. Phys.* **81**, 739 (2009).
- [4] R. N. Wenzel, *Ind. Eng. Chem.* **28**, 988 (1936).
- [5] A. B. D. Cassie and S. Baxter, *Trans. Faraday Soc.* **40**, 546 (1944).

- [6] P. G. de Gennes, *Rev. Mod. Phys.* **57**, 827 (1985).
- [7] P. Patrício, C.-T. Pham, and J. M. Romero-Enrique, *Eur. Phys. J. E* **26**, 97 (2008).
- [8] P. S. Swain and R. Lipowsky, *Langmuir* **14**, 6772 (1998).
- [9] C. Rascón, A. O. Parry, and A. Sartori, *Phys. Rev. E* **59**, 5697 (1999).
- [10] K. Rejmer and M. Napiórkowski, *Phys. Rev. E* **62**, 588 (2000).

- [11] P. Patricio, J. M. Romero-Enrique, N. M. Silvestre, N. R. Bernardino, and M. M. Telo da Gama, *Mol. Phys.* **109**, 1067 (2011).
- [12] D. W. Berreman, *Phys. Rev. Lett.* **28**, 1683 (1972).
- [13] J. M. Romero-Enrique, C.-T. Pham, and P. Patricio, *Phys. Rev. E* **82**, 011707 (2010).
- [14] P.-G. de Gennes, *Mol. Cryst. Liq. Cryst.* **12**, 193 (1971).
- [15] P. Patrício, M. M. Telo da Gama, and S. Dietrich, *Phys. Rev. Lett.* **88**, 245502 (2002).
- [16] P. G. de Gennes and J. Prost, *The Physics of Liquid Crystals*, 2nd ed. (Clarendon Press, Oxford, 1995).
- [17] D. Andrienko, M. Tasinkevych, P. Patricio, and M. M. Telo da Gama, *Phys. Rev. E* **69**, 021706 (2004).
- [18] P. Sheng, *Phys. Rev. Lett.* **37**, 1059 (1976).
- [19] P. Sheng, *Phys. Rev. A* **26**, 1610 (1982).
- [20] F. N. Braun, T. J. Sluckin, and E. Velasco, *J. Phys. Condens. Matter* **8**, 2741 (1996).
- [21] P. Patrício, M. Tasinkevych, and M. M. Telo da Gama, *Eur. Phys. J. E* **7**, 117 (2002).
- [22] G. Barbero, A. S. Gliozzi, M. Scalerandi, and L. R. Evangelista, *Phys. Rev. E* **77**, 051703 (2008).