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Properties of surface Landau–de Gennes Q-tensor models

Michael Nestler,^{*a} Ingo Nitschke,^a Hartmut Löwen^b and Axel Voigt^{ib acd}

Uniaxial nematic liquid crystals whose molecular orientation is subjected to tangential anchoring on a curved surface offer a non trivial interplay between the geometry and the topology of the surface and the orientational degree of freedom. We consider a general thin film limit of a Landau–de Gennes Q-tensor model which retains the characteristics of the 3D model. From this, previously proposed surface models follow as special cases. We compare fundamental properties, such as the alignment of the orientational degrees of freedom with principle curvature lines, order parameter symmetry and phase transition type for these models, and suggest experiments to identify suitable model assumptions.

1 Introduction

Liquid crystals^{1,2} consist of particles that possess both translational and orientational degrees of freedom. If these particles are constrained to the tangent bundle of a curved surface interesting phenomena emerge, which result from the tight coupling of the elastic and bulk free energies of the liquid crystal with the topological and geometrical properties of the surface. There are various experimental realization^{3–13} and particle-based computer simulations,^{14–30} which mainly focus on the emergence and position of topological defects on spherical or more complex surfaces. However, not only defects are tightly linked to the topological and geometrical properties of the surface, but also other fundamental issues, such as the alignment of the orientational degree of freedom with principle curvature lines, order parameter symmetries, phase transition type and curvature induced phase transitions are of fundamental interest, but are much less explored. We will address these issues for uniaxial nematic liquid crystals within a field-theoretical description of a surface Landau–de Gennes Q-tensor model. Various such models have been proposed.^{31–38} They strongly differ in the coupling mechanism between the orientational ordering of the nematic liquid crystal and the geometric properties of the surface. These coupling terms strongly depend on the assumptions made in the derivation and, as will be shown, have strong consequences on the fundamental properties of the phase transition type and order parameter symmetries.

Liquid crystals on curved surfaces are in a certain sense between 2D and 3D and could thus show properties of both dimensions. Lets first compare such properties of the established Landau–de Gennes Q-tensor theory for nematic liquid crystals in 2D and 3D:¹ the nematic phase can be stable in both dimensions. However, already the isotropic-to-nematic phase transition qualitatively differs between 2D and 3D. While it is of first-order in 3D, it is controversially discussed if this generally holds in 2D. Even if it can be proven that first-order isotropic-to-nematic phase transitions are possible in 2D,^{39,40} computer simulations and experiments indicate qualitatively different behaviour including continuous, first-order and even the absence of phase transitions, see *e.g.* ref. 41–43. In ref. 44 an overview of these theoretical arguments, computer simulations and experimental observations for thin films and 2D systems is provided. The situation on curved surfaces should somehow reflect these properties. However, in some of the proposed surface models first-order phase transitions are not possible. Another aspect highlighting the differences is discussed in ref. 45 by comparing mean field theories in 2D⁴⁶ and 3D.⁴⁷ The definitions yield different eigenvalue spectra in the order tensor, which corresponds to either a symmetry under in plane rotations by 90° in 2D or a rotational symmetry (w.r.t. the average particle direction) in 3D. How a nematic liquid crystal on a curved surface fits into this picture is open. A similar situation arises for Landau–de Gennes models. Some of the proposed surface models yield an eigenvalue spectrum as in 2D, others as in 3D. In 2D uniaxial nematics is the only possible nematic state, in 3D it is the energetically most favourable state under weak assumptions on the state potential.⁴⁵ A proper classification of uniaxiality in a surface Landau–de Gennes model is open, but one might ask, if, as in 2D, only uniaxial nematics should be allowed. The third aspect considers the mean orientation. If not induced by external fields or boundary conditions the mean orientation in

^a Institut für Wissenschaftliches Rechnen, Technische Universität Dresden, 01062 Dresden, Germany. E-mail: michael.nestler@tu-dresden.de

^b Institut für Theoretische Physik II – Soft Matter, Heinrich-Heine-Universität Düsseldorf, 40225 Düsseldorf, Germany

^c Dresden Center for Computational Materials Science (DCMS), Technische Universität Dresden, 01062 Dresden, Germany

^d Center for Systems Biology Dresden (CSBD), Pfotenhauerstr. 108, 01307 Dresden, Germany



nematic liquid crystals is arbitrary in 2D and 3D. This changes on curved surfaces, where it should be preferential to align with the principle curvature lines. This aspect has been discussed previously and is identified with the influence of extrinsic curvature terms^{32,33} in surface models, which again are considered in some but not all of the proposed models.

We will review the proposed surface Landau–de Gennes Q -tensor models with the aspect of these fundamental properties. Furthermore we propose a version of a surface Landau–de Gennes Q -tensor model which effectively describes surface liquid crystals retaining the 3D phase transition type and uniaxial eigenvalue spectra. Most previously proposed models follow as special cases and we discuss under which assumptions the fundamental properties get lost. The paper is structured as follows. In Section 2 we briefly review the 3D Landau–de Gennes Q -tensor model¹ and propose its thin film limit under generic anchoring conditions. Solving this model indicates a strong influence of geometry on fundamental properties. In particular, it is established that curvature can induce biaxiality, even if parameters in the original 3D model enforce state potential minima to be uniaxial. To overcome this problem we modify the derived energy by adding a penalty term which enforces uniaxiality weakly. In Section 3 this model together with its special cases is discussed in a flat 2D scenario with respect to their fundamental properties. We demonstrate the general model to retain the 3D properties. With this established, we apply the models to curved surfaces and discuss the effects of curvature on the ordering of the liquid crystal in Section 4. We summarize our findings and discuss them in a general framework in Section 5. As all these results do not depend on specific material parameters, we only consider a one-constant approximation. Details on derivations and used numeric methods can be found in the Appendix.

2 Thin film limit of the Q -tensor model

In this section we present essential notions and properties of the Landau–de Gennes Q -tensor model in 3D¹ and derive a generic surface model as a thin film limit. Restrictions to enforce uniaxiality and special cases, which link the model to the previously proposed surface Landau–de Gennes Q -tensor models, follow.

Q -Tensor model in 3D

We consider rod like particles with a head–tail symmetry in a volume $V \subset \mathbb{R}^3$. We consider the symmetric and trace free tensorial order parameter Q . In the special case of a uniaxial eigenvalue spectrum it is defined by

$$Q = S \left(P \otimes P - \frac{1}{3}G \right), \quad \sigma(Q) = \pm \left[\frac{2}{3}, -\frac{1}{3}, -\frac{1}{3} \right] S \quad (1)$$

where P denotes the principal director, defined by the average orientation of the particles, S the scalar order parameter, encoding the degree of alignment by the particles with the average direction, G the metric of V and $\sigma(Q)$ the eigenvalue spectrum. Please note that for Cartesian coordinates the metric G coincides with the identity matrix $\mathbb{1}_3$. The phase of prevalent

liquid like material properties is characterized by an isotropic ordering of particles and $S = 0$. In the nematic phase the particles tend to preferentially align with the average direction and $S \rightarrow S^* > 0$. The Landau–de Gennes Q -tensor model is based on the free energy

$$\mathcal{F}(Q) = \int_V \frac{L}{2} \|\nabla_3 Q\|^2 + \omega \left[a \text{tr}_3 Q^2 + \frac{2}{3} b \text{tr}_3 Q^3 + c \text{tr}_3 Q^4 \right] dV. \quad (2)$$

The first term, the elastic energy, penalizes any spatial deviations from the ground state. For the sake of simplicity we consider a one-constant approximation with single elastic parameter L . ∇_3 denotes the typical ∇ -operator in \mathbb{R}^3 and $\|\cdot\|$ the Frobenius norm. For more general models see *e.g.* ref. 48–52. In the remaining terms, the state potential, we have factored out $\omega > 0$ such that classic phenomenological constants are given by $a^* = \omega a$, $b^* = \omega b$ and $c^* = \omega c$. For the sake of readability we use an index free tensor notation,⁵³ where *e.g.* \cdot denotes the inner product, regardless of the tensorial degree. We define trace notion by $\text{tr}_3 Q = Q \cdot G$. Such full contraction with the space metric coincides with the Frobenius tensor norm by $\text{tr}_3 Q^2 = \|Q\|^2 = \sum_{ij} Q_{ij}^2$. The state potential can be expressed in terms of S such that the choice of a , b and c defines the preferred ordering S^* as local minima of the state potential.

Following ref. 45 and 52 we point out that the tensorial order parameter Q in this model is not restricted to the uniaxial eigenvalue spectra $\sigma(Q)$ as defined in (1). To track such situations, a biaxiality measure is introduced⁵⁴

$$U(Q) = 1 - 6 \frac{(\text{tr}_3 Q^3)^2}{(\text{tr}_3 Q^2)^3} \quad (3)$$

for which $U(Q) = 0$ if and only if Q is uniaxial. This measure is discussed in detail in ref. 45 and it is established that for $b < 0$ state potential minima Q^* are uniaxial.

Planar anchoring and tensor decomposition

For the boundary of the volume V , ∂V , with outward pointing normal ν , planar anchoring of uniaxial Q -tensors is modeled by a bare surface energy as discussed in ref. 35–37,

$$\mathcal{F}_B(Q) = \int_{\partial V} \alpha [\nu Q \nu - \beta]^2 + \gamma \|(Q - \nu \otimes \nu)(Q \nu)\|^2 d\partial V, \quad (4)$$

with non-negative coefficients $\alpha, \gamma \in \mathbb{R}^+$. The formulation can be interpreted as a penalty energy enforcing ν as an eigenvector of Q with eigenvalue β . In the case of a tangential aligned uniaxial Q -tensor ($P \cdot \nu = 0$) this would translate into $\beta = -\frac{1}{3}S$. Given this concept of prescribing a specific eigenvalue in the boundary normal direction motivates the separation of Q in a normal part β and tangential parts q ($\|q\nu\| = \| \nu q \| = 0$). Analogous to ref. 35–37 we thereby choose a separation such that q is symmetric and trace free (in the boundary domain sense)

$$Q(q, \beta) = q - \frac{\beta}{2}g + \beta \nu \otimes \nu, \quad (5)$$



where g is the metric of ∂V and $\text{tr}_2 q = q \cdot g = 0$. Within this decomposition we can consider q as a two dimensional Q -tensor on ∂V with tangential principal director p ($p \cdot \nu = 0$)

$$q = S \left(p \otimes p - \frac{1}{2} g \right), \quad \sigma(q) = \pm \left[\frac{1}{2}, -\frac{1}{2}, 0 \right] S. \quad (6)$$

Thin film limit models

Lets consider $V = \mathcal{S}_h$ as a thin film with thickness h , such that $\mathcal{S}_h = \mathcal{S} \times \left[-\frac{h}{2}, \frac{h}{2} \right]$ and \mathcal{S} a regular surface. We perform the thin film limit $\lim_{h \rightarrow 0} \frac{1}{h} \mathcal{F}(Q) = \mathcal{F}^{\mathcal{S}}(Q)$ in the spirit of ref. 38 under boundary condition $\|(G - \nu \otimes \nu)(Q\nu)\| = 0$, which corresponds to $\alpha = 0$ and $\gamma \rightarrow \infty$ in eqn (4) and keeps β as a degree of freedom. For details see the Appendix. By averaging out the normal direction we yield a surface model. Inserting (5) in (2) leads to

$$\begin{aligned} \mathcal{F}^{\mathcal{S}}[q, \beta] &= \frac{1}{2} \int_{\mathcal{S}} L \|\nabla q\|^2 + \frac{3}{2} L \|\nabla \beta\|^2 \\ &\quad - 6L\mathcal{H}\beta\langle \mathcal{B}, q \rangle + L \|\mathcal{B}\|^2 \left(\text{tr}_2 q^2 + \frac{9}{2} \beta^2 \right) d\mathcal{S} \\ &\quad + \omega \int_{\mathcal{S}} \frac{1}{2} (2a - 2b\beta + 3c\beta^2) \text{tr}_2 q^2 + c \text{tr}_2 q^4 \\ &\quad + \frac{\beta^2}{8} (12a + 4b\beta + 9c\beta^2) d\mathcal{S}. \end{aligned} \quad (7)$$

In contrast to (2) all operators are defined by the Levi-Civita connection (covariant derivative) ∇ . For a definition see *e.g.* ref. 55. Furthermore, the inner products are considered at the surface. As in 3D we identify the first integral with the elastic energy and the second with the state potential. The first integral contains additional coupling terms, where \mathcal{B} denotes the shape operator and \mathcal{H} the mean curvature of \mathcal{S} , for details see the Appendix. These are extrinsic curvature contributions, the term $6\mathcal{H}\beta\langle \mathcal{B}, q \rangle$ induces an alignment of p with one of the lines of principle curvatures depending on the sign of \mathcal{H} . The second term $\|\mathcal{B}\|^2 \left(\text{tr}_2 q^2 + \frac{9}{2} \beta^2 \right)$ poses an isotropic coupling between curvature and ordering. The term is closely related to the state potential such that the curvature can locally deform the potential and can induce phase transition or violate uniaxiality even if $b < 0$, a setting guarantees state potential minima to be uniaxial in the initial 3D model.⁴⁵ An example is provided in the Appendix.

As we are only interested in uniaxial nematic liquid crystals, we require a constraint, which suppresses curvature induced biaxiality. Inserting the decomposition (5) in the biaxiality measure (3) leads to a condition for β enforcing uniaxial eigenvalue spectra of Q -tensors, for details see the Appendix. It reads

$$\beta = \pm \frac{\sqrt{2}}{3} \|q\|. \quad (8)$$

Using this constraint in (7) to eliminate β leads to numerically cumbersome terms in the first variation, which is not further

pursued. We instead add a penalty term to (7) to enforce the constraint weakly,

$$\omega_{\beta} \int_{\mathcal{S}} \frac{1}{4} \left(\beta^2 - \frac{2}{9} \|q\|^2 \right)^2 d\mathcal{S}, \quad (9)$$

with $\omega_{\beta} > 0$.

As evolution laws we propose L^2 -gradient flows of the energy (7) with (9) with independent variables q and β , which read

$$\begin{aligned} \partial_t q &= L \Delta_{\mathcal{S}}^{\text{DG}} q - (L(\mathcal{H}^2 - 2\mathcal{K}) - \omega(2a - 2b\beta + c(3\beta^2 + 2\text{tr}_2 q^2))) q \\ &\quad + 3L\mathcal{H}\beta \left(\mathcal{B} - \frac{1}{2} \mathcal{H} g \right) - \omega_{\beta} \frac{2}{9} \left(\beta^2 - \frac{2}{9} \|q\|^2 \right) q \end{aligned} \quad (10)$$

$$\begin{aligned} \partial_t \beta &= L \Delta_{\mathcal{S}} \beta - \omega(3c\beta^3 + b\beta^2) - \omega(2a + 2c \text{tr}_2 q^2) \beta \\ &\quad - 3L(\mathcal{H}^2 - 2\mathcal{K})\beta + 2L\mathcal{H}\langle \mathcal{B}, q \rangle + \frac{2}{3} b \text{tr}_2 q^2 \\ &\quad + \omega_{\beta} \left(\beta^2 - \frac{2}{9} \|q\|^2 \right) \beta \end{aligned} \quad (11)$$

where $\Delta_{\mathcal{S}}^{\text{DG}}$ denotes the surface Div-Grad (Bochner) Laplace operator, $\Delta_{\mathcal{S}}$ the Laplace-Beltrami operator and \mathcal{K} the Gaussian curvature. Details of the derivation can be found in the Appendix. Eqn (10) and (11) provide a general surface Landau-de Gennes Q -tensor model with uniaxial eigenvalue spectra with a minimum of *a priori* assumptions. This system can be solved numerically, see the Appendix.

Previously proposed models consider special choices for β . They result as a thin film limit by considering the boundary conditions $\nu Q \nu - \beta = 0$ and $\|(G - \nu \otimes \nu)(Q\nu)\| = 0$, corresponding to $\alpha \rightarrow \infty$ and $\gamma \rightarrow \infty$ in eqn (4), which specifies β . $\beta = 0$ yields a model with degenerate Q -tensors $Q = q$ as in ref. 31 and 34, for $\beta = -\frac{1}{3} S^*$ we yield the model of approximate uniaxial Q -tensors,^{32,37,56,57} where some consider a different constant value for β , and the one-constant approximation of the derived models in ref. 35, 36 and 38. The resulting evolution equations are special cases of eqn (10). In the following we will compare the three models:

1. $\beta = \pm \frac{\sqrt{2}}{3} \|q\|$ (eqn (10) and (11))
2. $\beta = -\frac{1}{3} S^*$ (eqn (10) with $\beta = -\frac{1}{3} S^*$ and $\omega_{\beta} = 0$)
3. $\beta = 0$ (eqn (10) with $\beta = 0$ and $\omega_{\beta} = 0$).

(1) ensures 3D uniaxial eigenvalue spectra, (2) approximates this condition, and (3) essentially has a biaxial eigenvalue spectrum in 3D and is only uniaxial if considered as a 2D model.

3 Fundamental properties of the surface models

To keep the discussion as simple as possible we start with planar surfaces, which already show fundamental differences.



Eigenvalue spectra and symmetries

We recall the results of the mean field modeling approach for 3D systems, see *e.g.* ref. 47,

$$Q^{\text{MF}} = \int_{S^2} \left(M \otimes M - \frac{1}{3}G \right) \phi_3(M) dM, \quad M \in \mathbb{R}^3, \|M\| = 1 \quad (12)$$

with M as the particle orientation and ϕ_3 the associated probability distribution. For planar alignment of M w.r.t. the tangential direction, of a thin film, the mean field model implies an eigenvalue spectrum $\sigma(Q^{\text{MF}}) = \left[\frac{2}{3}, -\frac{1}{3}, -\frac{1}{3} \right]$. The mean field model in 2D, *e.g.* ref. 46, reads similar

$$q^{\text{MF}} = \int_{S^1} \left(m \otimes m - \frac{1}{2}g \right) \phi_2(m) dm, \quad m \in \mathbb{R}^2, \|m\| = 1 \quad (13)$$

but implies an eigenvalue spectrum of $\sigma(q^{\text{MF}}) = \left[\frac{1}{2}, -\frac{1}{2} \right]$. Considering the eigenvalue spectrum for the decomposed Q tensor (5)

$$\sigma(Q(q, \beta)) = \pm \left[\frac{1}{2}(S - \beta), -\frac{1}{2}(S + \beta), \beta \right] \quad (14)$$

we observe for $\beta = 0$ a spectrum compatible with the 2D mean field theory, while $\beta = -\frac{1}{3}S^*$ and $\beta = \pm \frac{\sqrt{2}}{3} \|q\|$ conform to 3D theory.

Phase transition type

We now turn to the isotropic-to-nematic phase transition. In the Landau–de Gennes Q -tensor model such transitions can be accounted for by a temperature dependent coefficient $a = a(T) = a_0(T - T^*)$ where T^* denotes the critical temperature where the isotropic phase is stable for $T > T^*$. Fig. 1A shows the phase portrait of the Landau–de Gennes Q -tensor model for the 3D case w.r.t. $\frac{a}{c}, \frac{b}{c}$ with a typical transient for increasing T from the pure nematic phase, *via* phase coexistence of the nematic and

isotropic phase to the pure isotropic phase. T^c denotes the critical temperature where the nematic phase ceases to exist. In this framework the transition is discontinuous/first-order.

We transfer this investigation of the transient from 3D to the surface models. For this purpose we consider states of uniform $Q(q, \beta)$ and evaluate the minima of the state potential contribution in (7) w.r.t. to $a(T)$ and the choice of β . In Fig. 1B we have plotted the minima and their stability. Reviewing the results for

$\beta = \pm \frac{\sqrt{2}}{3} \|q\|$ we observe a behavior identical to the 3D case.

This is quite natural, since inserting the Q -tensor decomposition in 3D state potential energy density yields directly the surface counterpart. Therefore this surface model exhibits a first order phase transition type and enables phase coexistence. For the model of fixed $\beta = -\frac{1}{3}S^*$ we observe a first-order transition at T^c as in a previous model but no phase coexistence for $T \in [T^*, T^c]$. In the case of $\beta = 0$ the transition type changes to continuous/second-order and shifts to the lower temperature T^* .

Fig. 2 further highlights the qualitative differences of the three models on a torus (with two major radii r, R), which is chosen as a prototypical surface with varying curvature, which avoids the presence of topological defects.

4 Impact of curvature in surface models

To investigate the impact of β on the geometry coupling mechanisms and to demonstrate the sketched effects we consider two numerical experiments.

Distortion energy minima

To focus on the first coupling term in the surface Landau–de Gennes Q -tensor energy (7), $6\mathcal{H}\beta\langle \mathcal{B}, q \rangle$, we consider a uniform director field $P(\varphi) = \cos(\varphi)p_r - \sin(\varphi)p_R$ on a torus defined by the linear combination of two director modes p_r and p_R as shown in Fig. 3A. With this director field we define $q(\varphi) = (p(\varphi) \otimes p(\varphi) - \frac{1}{2}g)$ with fixed norm. For given β we can now variate φ and evaluate

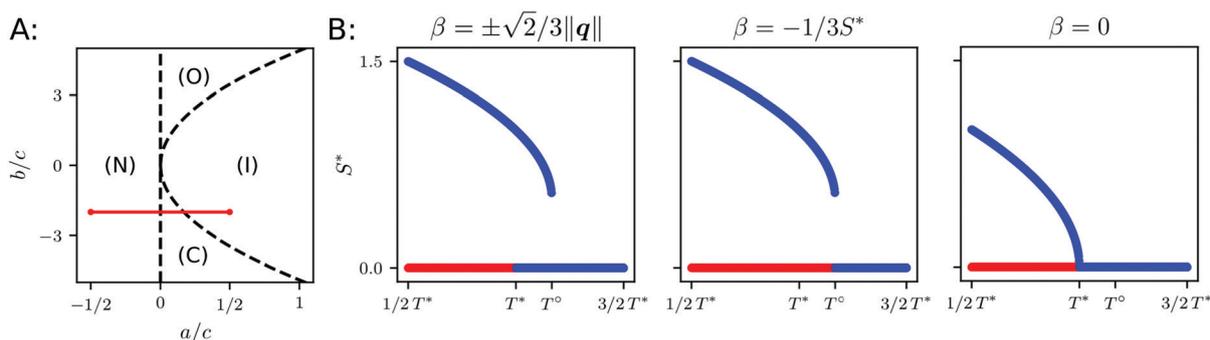


Fig. 1 Types of phase transition in surface models: (A) phase portrait for bulk Landau–de Gennes Q -tensor model as defined in (2). Four distinct phases w.r.t. state potential parameters a, b and c exist. Only the isotropic phase is stable (I), only the nematic phase is stable (N) isotropic-to-nematic phase coexistence (C) and orthogonal ordering (O). The red line indicates a typical phase transition for $a(T) = \alpha(T - T^*)$, $T \in [1/2 T^*, 3/2 T^*]$ where T^* denotes critical temperature such that the isotropic phase loses stability, (I)-to-(N) transition. Here, the temperature T^c marks the transition from only the isotropic stable regime to coexistence, (I)-(C) transition. (O) denotes also a region of coexistence but due to $\frac{b}{c} > 0$ a preferred ordering $S^* < 0$ is observed. Such a situation of orthogonal ordering (w.r.t. P) is not discussed here. (B) Stability of minima $S = \{0, S^*\}$ of state potential w.r.t. to temperature for $a(T)$ and choice of the β model. Blue lines denote stable minima while red lines indicate unstable minima.



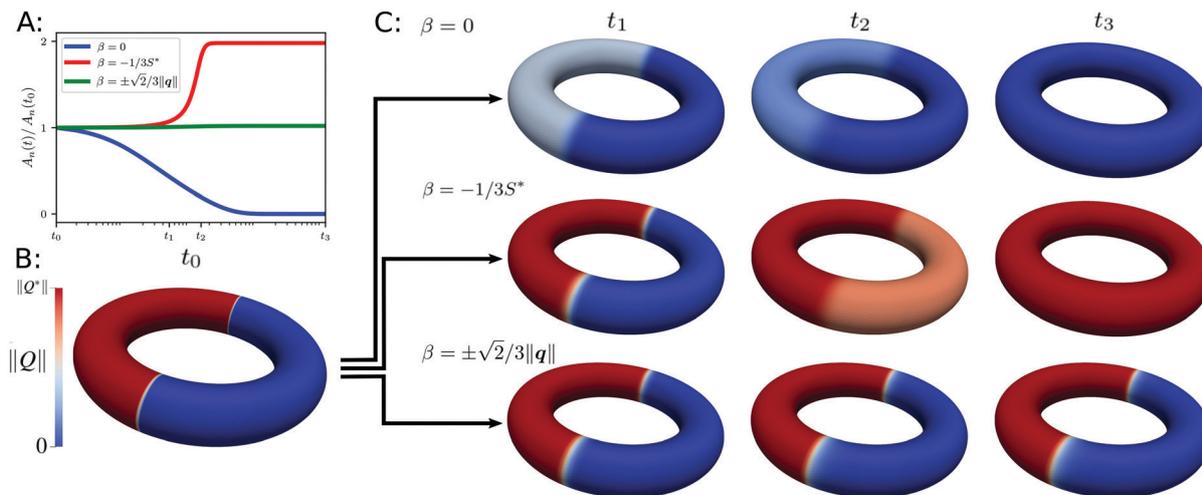


Fig. 2 Phase coexistence in surface models: (A) temporal evolution of area covered by nematic domain $A_n(t)$ vs. initial nematic domain $A_n(t_0)$. Starting at equally sized domains for $\beta = 0$ the nematic domain slowly shrinks, for $\beta = -\frac{1}{3}S^*$ the isotropic phase rapidly changes to nematic ordering, and for $\beta = \pm\frac{\sqrt{2}}{3}\|q\|$ the ratio remains almost constant. (B) Initial distribution of isotropic (blue) and nematic (red) domains on torus ($R = 2$, $r = \frac{1}{2}$). (C) Snapshots of $\|Q\| \in [0, \|Q^*\|]$ for $t_i \in [1e - 4, 1]$. Parameters: $L = 1$, $a = \frac{1}{4}$, $b = -4$, $c = 1$ and $\omega = 100$, $\omega_\beta = 10$, $\omega_\nu = 1000$.

the distortion energy contributions. In this set up the isotropic coupling term is constant as well as $\|\nabla\beta\|^2$, furthermore the models $\beta = -\frac{1}{3}S^*$ and $\beta = \pm\frac{\sqrt{2}}{3}\|q\|$ coincide. For $\beta = 0$ the directed coupling term vanishes and the minimum of distortion energy is defined by the $\|\nabla q\|^2$ contribution. As shown in Fig. 3C (top) two minima exist. The minimum $\varphi^* = \frac{\pi}{4}$ results in a surprising q configuration, shown in Fig. 3B (left). The second minimum $\varphi^* = \frac{3}{4}\pi$ yields a corresponding configuration with in-plane 90° rotated eigenvectors. In the model $\beta = \pm\frac{\sqrt{2}}{3}\|q\|$ and on the chosen geometry the $-3\beta L\mathcal{H}(\mathcal{B}, q)$ contribution dominates the distortion energy, see Fig. 3C (bottom) such that the minimum is achieved for

$\varphi^* = \frac{\pi}{2}$. The corresponding Q -tensor configuration consists of principal eigenvectors aligned with lines of minimal curvature, see Fig. 3B (right).

Reviewing these results we conclude, once more, that degenerate ($\beta = 0$) surface Landau-de Gennes Q -tensor models describe a substantial different type of physical systems than surface models with non-degenerate Q -tensors. Retaining the 3D nature of liquid crystals in the surface model yields an intensified geometry-ordering coupling.

Balancing curvature effects for $\beta = \pm\frac{\sqrt{2}}{3}\|q\|$

To obtain an intuition on the possible interactions of the geometry-order couplings we consider a thick torus ($R = 0.55$, $r = 0.45$) where \mathcal{H} changes sign, see Fig. 4A. On this surface we

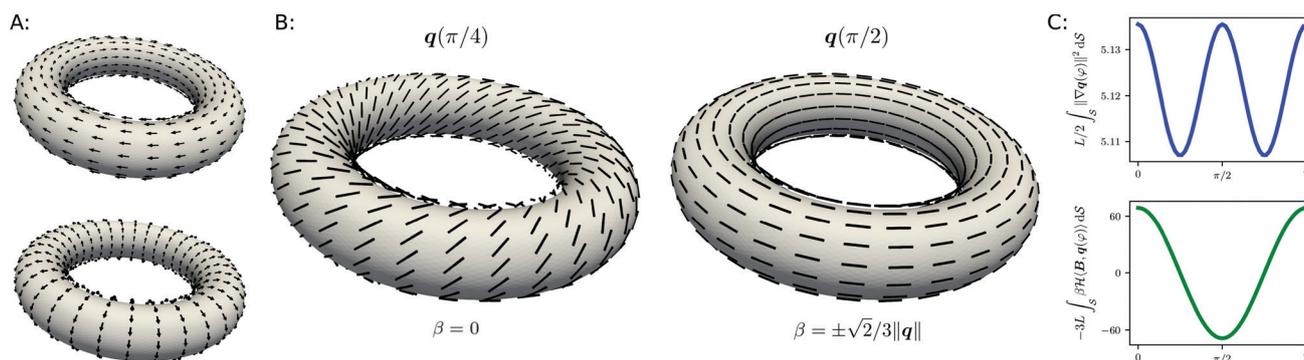


Fig. 3 Distortion energy of defect free configurations of torus: (A) director modes on torus ($R = 2$, $r = 1/2$). p_R (top) and p_r (bottom). (B) Equilibrium states of evolution equations for $\beta = 0$ (left) and $\beta = \pm\frac{\sqrt{2}}{3}\|q\|$ (right). Principal eigenvectors of minimum energy configurations $q(\varphi^*)$ for $\beta = 0$ (left): $\varphi^* = \frac{\pi}{4}$; $\beta = \pm\frac{\sqrt{2}}{3}\|q\|$ (right): $\varphi^* = \frac{\pi}{2}$. (C) Contributions of integral distortion energy for angle φ . (top) $\|\nabla q\|^2$ with minima at $\varphi = \frac{\pi}{4}, \frac{3}{4}\pi$, (bottom) $-3\beta L\mathcal{H}(\mathcal{B}, q)$ for $\beta = -\frac{\sqrt{2}}{3}\|q\|$ with minima at $\varphi = \frac{\pi}{2}$.



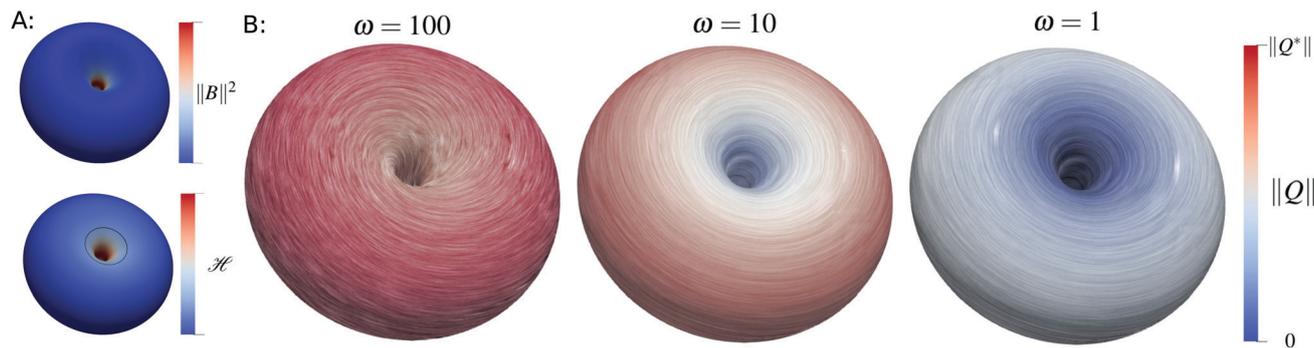


Fig. 4 Curvature impact on equilibrium configurations on thick torus: (A) curvature of thick torus ($R = 0.55$, $r = 0.45$). (top) $\|B\|^2 \in [4.7, 155]$, (bottom) \mathcal{H} changes sign, the location indicated by a black line, across geometry such that strong positive at center and mild negative at rim $\mathcal{H} \in [-3.2, 10]$. (B) Equilibrium configurations depending on the choice of ω [$\omega = 100$ (left), $\omega = 10$ (mid) and $\omega = 1$ (right)]. Lines indicate the direction of the principal director of Q , obtained by LIC. Colors denote $\|Q\| \in [0, \|Q^*\|]$, where $\|Q^*\|$ is given by a minimum value of state potential ($a = \frac{1}{4}$, $b = -4$, $c = 1$). (blue) isotropic, and (red) nematic. Simulations are performed with $L = 1$, $\omega_\beta = 10$ and $\omega_\nu = 10^3$.

evaluate the equilibrium states Q of the $\beta = \pm \frac{\sqrt{2}}{3} \|q\|$ model for several values of ω , see Fig. 4B. Recalling ω as factor weighting the distortion contribution *versus* the state potential we observe for $\omega = 100$ a dominant state potential such that a uniform nematic phase, $\|Q\| = \|Q^*\|$, as prescribed by the state potential, is enforced across the entire surface, suppressing the effects of the isotropic geometry coupling. In contrast, the directed geometry coupling is not affected and we observe a strong forcing towards the – geometry induced – preferred alignment such that a non uniform ordering of principal directors is yielded, see Fig. 4B (left). For regions of positive \mathcal{H} the alignment is with minimal curvature lines (outer part), whereas for negative \mathcal{H} the alignment is with maximal curvature lines (inner part). Similar effects have also been reported for polar liquid crystals within a surface Frank–Oseen model.^{58,59} Weakening the state potential by choosing $\omega = 10$ we observe a curvature induced phase transition with a localized isotropic phase at the inner part of the torus, where $\|B\|^2$ has its maximum. This area coincides with the area of the strongest directed geometry coupling. Finally matching the distortion contribution and state potential by $\omega = 1$ we yield a global isotropic phase since the isotropic coupling distorts the state potential such that only the isotropic $Q = 0$ phase remains stable.

The isotropic geometry coupling term, $\|B\|^2 \left(\text{tr}_2 q^2 + \frac{9}{2} \beta^2 \right)$, turns out to be the dominating effect compared to the directed geometry coupling term $6\mathcal{H}\beta\langle B, q \rangle$. Only in situations where the state potential is strong enough to suppress the isotropic coupling the effects of directed coupling become traceable. In this situation the geometry induces a preferred alignment, see also ref. 38 for similar results on ellipsoidal geometries. For geometries with strong variations in \mathcal{H} , including sign changes, this leads to nonuniform ordering. An effect is obviously possible only in models with $\beta \neq 0$.

5 Discussion and conclusions

Exploring a thin film limit of the 3D Landau–de Gennes Q -tensor model allows bridging the gap to previously proposed surface

models. Planar anchoring at the boundary of the thin film was thereby used to fix the boundary normal as eigenvector with eigenvalue β , which motivated the decomposition of the tensorial order parameter Q in tangential q and normal β parts. We will now review the central results of the models with different choices of β , discuss suitable application scenarios, assess their inherent couplings to the curvature of the surface and discuss possible experiments to confirm these results.

Discussion

The first class of surface Landau–de Gennes Q -tensor models considers $\beta = 0$ and is usually labeled as planar degenerate Q -tensors.^{31,34} We observed the models to exhibit essentially 2D characteristics like eigenvalue spectra of Q matching 2D mean field theory. These models also show a continuous/second-order isotropic-to-nematic phase transition, which contradictory results on the existence of first-order isotropic-to-nematic phase transitions in 2D.^{39,40} Concerning the coupling with geometric properties the model in ref. 31 and 34 did only account for intrinsic curvature effects.

For the second class, where $\beta \neq 0$, the obtained models retain the characteristics of the 3D Landau–de Gennes Q -tensor model. As the resulting thin film limit in the general case of variable β shows a distortion of uniaxiality by curvature, we use a 3D biaxiality measure to fix $\beta = \pm \frac{\sqrt{2}}{3} \|q\|$, which enforces uniaxial Q also in the surface model. Further investigations confirmed that in this modified model for uniaxial Q -tensors the fundamental properties of eigenvalue spectra, first order isotropic-to-nematic phase transition and phase coexistence stay preserved. The model $\beta = -\frac{1}{3}S^*$, as proposed in ref. 36 and 38, has been proven to reproduce the 3D characteristics, under the assumption of $\|Q\| = \|Q^*\|$, except for phase coexistence. It can be considered as a simplified surface Landau–de Gennes Q -tensor model suitable for uniaxial nematic liquid crystals far from phase transition temperature. Furthermore $\beta \neq 0$ introduces additional curvature coupling terms. In addition to isotropic geometry coupling, the alignment of the director field with the principle



curvature lines is considered, which had already been shown in ref. 32, 36, 38 and 60. Parameter studies for a torus indicate that, depending on the strength of the curvature, phase transitions can be enforced, leading to phase coexistence and locally confined isotropic regions within a nematic phase or even a uniform isotropic phase if the curvature effect is strong enough. The derived thin film limit (without the introduced penalization to enforce uniaxiality) shows even stronger curvature coupling, e.g. curvature induced biaxiality, see the Appendix. Other implications of this model are not explored.

These discrepancies in response to the curvature of the $\beta = 0$ and $\beta \neq 0$ models provide a motivation for *in vitro* experiments to assess the prevalent 2D or 3D nature of liquid crystals which are confined to curved surfaces.

Conclusions

With the presented derivations and arguments we have provided a comprehensive study unifying recent approaches for surface Landau–de Gennes Q -tensor theories for uniaxial nematic liquid crystals confined to curved surfaces. By introducing a surface parameter β we have classified different surface limits for nematic phases on curved manifolds. Essentially, β measures the ability of the orientational degrees of freedom to fluctuate in the direction perpendicular to the curved surface while the particle centers are constrained on the manifold in the thin-film limit. In terms of physics, these are the imposed anchoring conditions at the surface. In particular, we have identified two classes of models as special limits which could be related to liquid crystals with prevalent 2D or 3D characteristics. The distinct response to curvature of these two model classes enables a path to determine suitable models for the liquid crystal systems by *in vitro* experiments. In the future it would be interesting to link the classes of surface Landau–de Gennes Q -tensor models studied in this paper to particle-resolved models where anisotropic apolar particles are bound to curved interfaces. It remains to be understood how different anchoring conditions of the particles at the surface can be mapped and described effectively by the coarse-grained mean-field-like approach. In principle, varying the anchoring conditions should result in different coupling parameters used in the surface free energy (4). In particle-resolved computer simulations, different anchoring conditions can just be implemented by an explicit orientational coupling to the curved interface. In actual experiments on colloids bound to curved interfaces (see e.g. ref. 61), on Pickering emulsion droplets⁶² the anchoring conditions can be conveniently changed by the pH⁶³ or by changing the thermodynamic parameters.

Conflicts of interest

There are no conflicts to declare.

A Appendix

A.1 Details on the uniaxiality condition

To obtain the surface uniaxiality condition (8) we can insert the Q -tensor decomposition (5) into the biaxiality measure (2).

Another elegant approach is to consider the eigenvalue spectrum (14) and insert it into the biaxiality measure: Q uniaxial $\Leftrightarrow U(Q) = 0 \Leftrightarrow 6(\text{tr}_3 Q^3)^2 = (\text{tr}_3 Q^2)^3$. We obtain $\text{tr}_3 Q^2 = \frac{1}{2}S^2 + \frac{3}{2}\beta^2$ and $\text{tr}_3 Q^3 = -\frac{3}{4}\beta S^2 + \frac{3}{4}\beta^3$ such that the uniaxiality condition $U(Q) = 0$ can be expressed by, assuming $S \neq 0$, $\beta^2(18S^2 - 81\beta^2) = S^4$. This holds if and only if $\beta^2 = \frac{1}{9}S^2$ which translates to $\beta = \pm \frac{\sqrt{2}}{3} \|q\|$.

A.2 Derivation of the thin film limit

Thin film limits require a reduction of the degrees of freedom. We deal with this issue by setting Dirichlet boundary conditions for the normal parts of Q and postulate *a priori* a minimum of the free energy on the boundary of the thin film. This is achieved by considering the natural boundary condition of the weak Euler–Lagrange equation. In this setting we restrict the density of \mathcal{F} to the surface and integrate in the normal direction to obtain the surface energy $\mathcal{F}^{\mathcal{S}}$. We closely follow ref. 38, use the notation introduced there and only point out differences. Following this notation we define the shape operator by $\mathcal{B} = -[(G - \nu \otimes \nu) \nabla_3] \nu$.

The free energy (2) in the thin film \mathcal{S}_h in index notation reads

$$\mathcal{F}[Q] = \int_{\mathcal{S}_h} \frac{L}{2} Q_{IJ;K} Q^{IJ;K} dV + \omega \int_{\mathcal{S}_h} \left(a Q_{IJ} Q^{IJ} + \frac{2}{3} b Q_{IJ} Q^{JK} Q^I_K + c Q_{IJ} Q^{JK} Q_{KL} Q^{LI} \right) dV.$$

For the choice of essential boundary conditions, we require that Q has to have two eigenvectors in the boundary tangential bundle and the remaining eigenvector has to be the boundary normal, i.e., for $P \in T\partial\mathcal{S}_h$ a pure covariant representation of Q at the boundary is

$$Q = S_1 P^b \otimes P^b + S_2 \nu^b \otimes \nu^b - \frac{1}{3}(S_1 + S_2)G \quad (15)$$

with scalar order parameters S_1 and S_2 . Hence, it holds $Q_{i\xi} = Q_{\xi i} = 0$ and $Q_{\xi\xi} = \frac{1}{3}(2S_2 - S_1)$ at the boundaries. The remaining boundary conditions are considered as natural boundary conditions $0 = Q_{ij;\xi}$ at $\partial\mathcal{S}_h$. According to the normal eigenvalue β of Q at \mathcal{S} , we extend this scalar field to the thin film \mathcal{S}_h , i.e., $Q_{\xi\xi} = \beta$ with $\hat{\beta}|_{\mathcal{S}} = \beta$.

We can relate the anchoring conditions to surface identities by sums and differences of Taylor expansions at the upper and lower boundaries, see ref. 38. This results in

$$Q_{i\xi}|_{\mathcal{S}} = Q_{\xi i}|_{\mathcal{S}} = \mathcal{O}(h^2) \quad \partial_{\xi} Q_{i\xi}|_{\mathcal{S}} = \partial_{\xi} Q_{\xi i}|_{\mathcal{S}} = \mathcal{O}(h^2) \\ Q_{\xi\xi;\xi}|_{\mathcal{S}} = \mathcal{O}(h^2) \quad Q_{ij;\xi}|_{\mathcal{S}} = \mathcal{O}(h^2).$$

The restricted Q -tensor $\{Q_{ij}|_{\mathcal{S}}\} \in T^{(2)}\mathcal{S}$ is not a Q -tensor. We have $\text{tr}_2\{Q_{ij}|_{\mathcal{S}}\} = \text{tr}_3 Q|_{\mathcal{S}} - Q_{\xi\xi}|_{\mathcal{S}} = -\beta$. To ensure Q -tensor



properties we introduce the projection $\Pi_{\underline{2}}: t \rightarrow \frac{1}{2}(t + t^T - (\text{tr}_2 t)g)$, and define

$$q := \Pi_{\underline{2}} \left\{ Q_{ij} \Big|_{\mathcal{S}} \right\} = \left\{ Q_{ij} \Big|_{\mathcal{S}} \right\} + \frac{\beta}{2} g. \quad (16)$$

We can determine all remaining covariant derivatives restricted to the surface by

$$\begin{aligned} Q_{i\bar{\zeta};\bar{\zeta}} \Big|_{\mathcal{S}} &= Q_{\bar{\zeta}i;\bar{\zeta}} \Big|_{\mathcal{S}} = \partial_{\bar{\zeta}} Q_{i\bar{\zeta}} \Big|_{\mathcal{S}} - \Gamma_{\bar{\zeta}i}^K Q_{K\bar{\zeta}} \Big|_{\mathcal{S}} = \mathcal{O}(h^2) \\ Q_{\bar{\zeta}\bar{\zeta};k} \Big|_{\mathcal{S}} &= \partial_k Q_{\bar{\zeta}\bar{\zeta}} \Big|_{\mathcal{S}} - 2\Gamma_{k\bar{\zeta}}^L Q_{L\bar{\zeta}} \Big|_{\mathcal{S}} = \beta|_k + \mathcal{O}(h^2) \\ Q_{i\bar{\zeta};k} \Big|_{\mathcal{S}} &= Q_{\bar{\zeta}i;k} \Big|_{\mathcal{S}} = \partial_k Q_{i\bar{\zeta}} \Big|_{\mathcal{S}} - \Gamma_{ki}^L Q_{L\bar{\zeta}} \Big|_{\mathcal{S}} - \Gamma_{ki}^{\bar{\zeta}} Q_{\bar{\zeta}\bar{\zeta}} \Big|_{\mathcal{S}} - \Gamma_{k\bar{\zeta}}^l Q_{il} \Big|_{\mathcal{S}} \\ &= [q\mathcal{B}]_{ik} - \frac{3}{2}\beta\mathcal{B}_{ik} + \mathcal{O}(h^2) \\ Q_{ij;k} \Big|_{\mathcal{S}} &= \partial_k Q_{ij} \Big|_{\mathcal{S}} - \Gamma_{ki}^l Q_{lj} \Big|_{\mathcal{S}} - \Gamma_{ki}^{\bar{\zeta}} Q_{\bar{\zeta}j} \Big|_{\mathcal{S}} - \Gamma_{kj}^l Q_{il} \Big|_{\mathcal{S}} - \Gamma_{kj}^{\bar{\zeta}} Q_{i\bar{\zeta}} \Big|_{\mathcal{S}} \\ &= \partial_k Q_{ij} \Big|_{\mathcal{S}} - \Gamma_{ki}^l Q_{lj} \Big|_{\mathcal{S}} - \Gamma_{kj}^l Q_{il} \Big|_{\mathcal{S}} + \mathcal{O}(h^2) \\ &= \left(q_{ij} - \frac{\beta}{2} g_{ij} \right) \Big|_k + \mathcal{O}(h^2) = q_{ij|k} - \frac{1}{2}\beta|_k g_{ij} + \mathcal{O}(h^2). \end{aligned} \quad (17)$$

The contributions of the energy density read

$$\begin{aligned} \|\nabla Q\|_{\mathcal{G}}^2 \Big|_{\mathcal{S}} &= Q_{ij;k} Q^{jk} + 2Q_{i\bar{\zeta};k} Q^{i\bar{\zeta};k} + 2Q_{\bar{\zeta}\bar{\zeta};k} Q^{\bar{\zeta}\bar{\zeta};k} \Big|_{\mathcal{S}} + \mathcal{O}(h^2) \\ &= \|\nabla q\|_g^2 + \frac{3}{2}\|\nabla\beta\|_g^2 + 2\left\| q\mathcal{B} - \frac{3}{2}\beta\mathcal{B} \right\|_g^2 + \mathcal{O}(h^2) \\ \text{tr}_3 Q^2 \Big|_{\mathcal{S}} &= \text{tr}_2 q^2 + \frac{3}{2}\beta^2 + \mathcal{O}(h^2) \\ \text{tr}_3 Q^3 \Big|_{\mathcal{S}} &= \frac{3}{2}\beta \left(\frac{\beta^2}{2} - \text{tr}_2 q^2 \right) + \mathcal{O}(h^2) \\ \text{tr}_3 Q^4 \Big|_{\mathcal{S}} &= \text{tr}_2 q^4 + \frac{3}{2}\beta^2 \text{tr}_2 q^2 + \frac{9}{8}\beta^4 + \mathcal{O}(h^2). \end{aligned} \quad (18)$$

Using

$$2\left\| q\mathcal{B} - \frac{3}{2}\beta\mathcal{B} \right\|^2 = \|\mathcal{B}\|^2 \left(\text{tr}_2 q^2 + \frac{9}{2}\beta^2 \right) - 6\mathcal{H}\beta\langle \mathcal{B}, q \rangle \quad (19)$$

which follows from ref. 38 (Corollary A.4.), adding all contributions up and denoting the free energy densities by F and $F^{\mathcal{S}}$, we obtain for $h \rightarrow 0$

$$\begin{aligned} \frac{1}{h} \mathcal{F} &= \frac{1}{h} \int_{\mathcal{S}_h} F dV = \frac{1}{h} \int_{-\frac{h}{2}}^{\frac{h}{2}} \int_{\mathcal{S}} (1 - \xi \mathcal{H} + \xi^2 \mathcal{K}) F d\mathcal{S} d\xi \\ &= \int_{\mathcal{S}} F^{\mathcal{S}} d\mathcal{S} + \mathcal{O}(h^2) = \mathcal{F}^{\mathcal{S}} + \mathcal{O}(h^2) \rightarrow \mathcal{F}^{\mathcal{S}}. \end{aligned} \quad (20)$$

A.3 Variational derivatives

The first variation of

$$\begin{aligned} \mathcal{F}^{\mathcal{S}}[q, \beta] &= \frac{1}{2} \int_{\mathcal{S}} L \|\nabla q\|^2 + \frac{3}{2} L \|\nabla\beta\|^2 \\ &\quad - 6L\mathcal{H}\beta\langle \mathcal{B}, q \rangle + L \|\mathcal{B}\|^2 \left(\text{tr}_2 q^2 + \frac{9}{2}\beta^2 \right) d\mathcal{S} \\ &\quad + \omega \int_{\mathcal{S}} \frac{1}{2} (2a - 2b\beta + 3c\beta^2) \text{tr}_2 q^2 + c \text{tr}_2 q^4 \\ &\quad + \frac{\beta^2}{8} (12a + 4b\beta + 9c\beta^2) d\mathcal{S}, \end{aligned}$$

w.r.t. surface Q -tensor ψ and scalar φ perturbations for q and β reads

$$\begin{aligned} \delta \mathcal{F}^{\mathcal{S}} &= L \int_{\mathcal{S}} \langle \nabla q, \nabla \psi \rangle + \langle \|\mathcal{B}\|^2 q - 3\mathcal{H}\beta\Pi_{\underline{2}}\mathcal{B}, \psi \rangle \\ &\quad + \frac{3}{2} \langle \nabla\beta, \nabla\varphi \rangle - 3 \left(\mathcal{H}\langle \mathcal{B}, q \rangle - \frac{3}{2} \|\mathcal{B}\|^2 \beta \right) \varphi d\mathcal{S} \\ &\quad + \omega \int_{\mathcal{S}} (2a - 2b\beta + 3c\beta^2) \langle q, \psi \rangle + 2c \text{tr}_2 q^2 \langle q, \psi \rangle \\ &\quad + \left(\frac{9}{2} c \beta^3 + \frac{3}{2} \beta^2 + 3(c \text{tr}_2 q^2 + a)\beta - b \text{tr}_2 q^2 \right) \varphi d\mathcal{S}, \end{aligned}$$

where β is independent. If β is constant, or prescribed generally, the associated perturbation φ vanishes. To enforce uniaxiality (8), we add the penalty energy

$$\mathcal{F}_{\text{uni}}^{\mathcal{S}} = \omega_{\beta} \int_{\mathcal{S}} \frac{1}{4} \left(\beta^2 - \frac{2}{9} \|q\|^2 \right)^2 d\mathcal{S}$$

to $\mathcal{F}^{\mathcal{S}}$, with $\omega_{\beta} > 0$. Its first variation reads

$$\delta \mathcal{F}_{\text{uni}}^{\mathcal{S}} = \omega_{\beta} \int_{\mathcal{S}} -\frac{2}{9} \left(\beta^2 - \frac{2}{9} \text{tr}_2 q^2 \right) \langle q, \psi \rangle + \left(\beta^2 - \frac{2}{9} \text{tr}_2 q^2 \right) \beta \varphi d\mathcal{S}.$$

From this, the L^2 -gradient flows lead to the considered evolution equations for q and β .

A.4 L^2 -Gradient flows

Instead of relating the energies \mathcal{F} and $\mathcal{F}^{\mathcal{S}}$, also the evolution equations in \mathcal{S}_h and on \mathcal{S} can be related. Similar calculations as in Section A.2 give

$$\langle \nabla_{L^2} \mathcal{F}^{\mathcal{S}_h}, \Psi \rangle \Big|_{\mathcal{S}} = \langle \nabla_{L^2}^q \mathcal{F}^{\mathcal{S}}, \psi \rangle + \langle \nabla_{L^2}^{\beta} \mathcal{F}^{\mathcal{S}}, \varphi \rangle + \mathcal{O}(h^2),$$

with appropriate bulk and surface Q -tensors Ψ and ψ and scalar φ perturbations, s.t. $\Psi_{\bar{\zeta}\bar{\zeta}} \Big|_{\mathcal{S}} = \varphi$. This condition is the principal difference to the calculations in ref. 38. Note that we used (19) in the derivation, weakly in the Q -tensor direction ψ , i.e.

$$\int_{\mathcal{S}} \left\langle q\mathcal{B} - \frac{3}{2}\beta\mathcal{B}, \psi \right\rangle d\mathcal{S} = \int_{\mathcal{S}} \langle \|\mathcal{B}\|^2 q - 3\mathcal{H}\beta\Pi_{\underline{2}}\mathcal{B}, \psi \rangle d\mathcal{S}.$$



Moreover, as $\partial_t g = 0$ for a stationary surface, we obtain $\langle \partial_t Q, \Psi \rangle_G|_{\mathcal{S}} = \langle \partial_t q, \psi \rangle_g + \frac{3}{2} \langle \partial_t \beta \rangle \phi + \mathcal{O}(h^2)$ and as in (20)

$$\begin{aligned} & \frac{1}{h} \int_{\mathcal{S}_h} \langle \partial_t Q + \nabla_{L^2} \mathcal{F}, \Psi \rangle dV \\ &= \int_{\mathcal{S}} \langle \partial_t q + \nabla_{L^2}^q \mathcal{F}^{\mathcal{S}}, \psi \rangle + \left(\frac{3}{2} \partial_t \beta + \nabla_{L^2}^{\beta} \mathcal{F}^{\mathcal{S}} \right) \phi d\mathcal{S} + \mathcal{O}(h^2). \end{aligned} \quad (21)$$

Therefore, the relaxation velocity parameter, for the L^2 -gradient flow w.r.t. β , has to be $\frac{2}{3}$ for consistency of the time-dependent thin film and surface problems w.r.t. h . The surface evolution equations thus read

$$\begin{aligned} \partial_t q &= L\Delta_{\mathcal{S}}^{\text{DG}} q - (L(\mathcal{H}^2 - 2\mathcal{H}) - \omega(2a - 2b\beta + c(3\beta^2 + 2\text{tr}_2 q^2)))q \\ &+ 3L\mathcal{H}\beta \left(\mathcal{B} - \frac{1}{2}\mathcal{H}g \right) + \omega\beta \frac{2}{9} \left[\beta^2 - \frac{2}{9} \|q\|^2 \right] q, \end{aligned} \quad (22)$$

$$\begin{aligned} \partial_t \beta &= L\Delta_{\mathcal{S}} \beta - \omega(3c\beta^3 + b\beta^2) - \omega(2a + 2c\text{tr}_2 q^2)\beta + 2L\mathcal{H} \langle \mathcal{B}, q \rangle \\ &- 3L(\mathcal{H}^2 - 2\mathcal{H})\beta + \frac{2}{3} b\text{tr}_2 q^2 - \omega\beta \frac{2}{3} \left[\beta^2 - \frac{2}{9} \|q\|^2 \right] \beta. \end{aligned} \quad (23)$$

A.5 Numeric solution procedure

To numerically solve the tensor- and scalar-valued surface PDEs (22) and (23), we use the surface FEM approaches of ref. 64 and 65, respectively. The approach in ref. 64 extends previous ideas for vector-valued surface PDEs^{59,66,67} to tensors of arbitrary degree. The idea of these approaches is to reformulate the problems in Cartesian coordinates and to penalize normal components. This allows for a componentwise solution using tools for scalar-valued surface PDEs, e.g. ref. 65. The penalty term added to $\mathcal{F}^{\mathcal{S}}$ reads $\omega_{\nu} \int_{\mathcal{S}} \frac{1}{2} \|q\nu\|^2 d\mathcal{S}$ with $\omega_{\nu} > 0$. This leads to an additional term in (22) reading $\omega_{\nu}(\nu \otimes \nu)q$.

To address the nonlinearity of the system of PDEs we consider a Newton method. We solve the temporal discretized problem as a sequence of time steps $[\hat{q}, \hat{\beta}] \rightarrow q, \beta$, where $\partial_t q \approx (q - \hat{q})/\tau$ and $\partial_t \beta \approx (\beta - \hat{\beta})/\tau$. We denote with $L\mathbb{Q}$, $L\mathbb{B}$ and $N\mathbb{Q}$, $N\mathbb{B}$ the collections of linear and nonlinear operators of the time step problems, where \mathbb{Q} and \mathbb{B} refer to the q and β state equations. The single Newton iteration $k \rightarrow k+1$ reads

$$\begin{aligned} & \delta_q N\mathbb{Q}(q^k, \beta^k)[q^{k+1}] + \delta_{\beta} N\mathbb{Q}(q^k, \beta^k)[\beta^{k+1}] + L\mathbb{Q}(q^{k+1}, \beta^{k+1}) \\ &= \delta_q N\mathbb{Q}(q^k, \beta^k)[q^k] + \delta_{\beta} N\mathbb{Q}(q^k, \beta^k)[\beta^k] - N\mathbb{Q}(q^k, \beta^k) \\ & \delta_q N\mathbb{B}(q^k, \beta^k)[q^{k+1}] + \delta_{\beta} N\mathbb{B}(q^k, \beta^k)[\beta^{k+1}] + L\mathbb{B}(q^{k+1}, \beta^{k+1}) \\ &= \delta_q N\mathbb{B}(q^k, \beta^k)[q^k] + \delta_{\beta} N\mathbb{B}(q^k, \beta^k)[\beta^k] - N\mathbb{B}(q^k, \beta^k) \end{aligned}$$

To evaluate $[q, \beta]$ we solve the Newton iterations until

$$\left(\int_{\mathcal{S}} \|q^k - q^{k+1}\|^2 + \|\beta^k - \beta^{k+1}\|^2 d\mathcal{S} \right)^{1/2} < \theta.$$

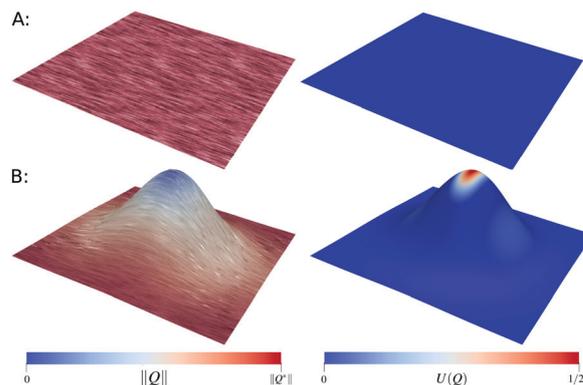


Fig. 5 Curvature distorts uniaxiality in unconstrained surface Landau-de Gennes Q -tensor model: (A) in 3D energetic minima of unconstrained surface Q -tensors $\mathcal{F}(Q(q, \beta))$ are uniaxial (for $b < 0$, see e.g. ref. 45), which also holds for the thin film limit for a flat surface. (left) 2D yields uniform alignment of the principal director (lines, obtained by line integral convolution (LIC) and constant $\|Q\|$ (color scale)). (right) Uniform values of biaxiality measure $U(Q) \approx 0$. (B) Curvature in domain distorts $\|Q\|$ and also uniaxiality. (left) Norm deficiency in high curvature areas. (right) Uniaxiality is substantially violated in high curvature regions where $U(Q) \approx \frac{1}{3}$. The geometry is defined by a Gaussian bump $h(x, y) = h_0 \exp(-(x^2 + y^2)/w_0^2)$ where $h_0 = w_0 = \frac{1}{2}$ and $(x, y) \in [-1, 1]^2$ and $\omega = 10$.

The resulting linear surface PDEs are solved by the surface FEM methods^{64,65} which are implemented in the adaptive FEM toolbox AMDiS.^{68,69}

To assess the quality of enforcing tangentiality and uniaxiality by the introduced penalty terms, simulations are performed with $\omega_{\nu} \in [10^1, 10^3]$ and $\omega_{\beta} \in [10^1, 10^3]$ on a torus ($R = 2, r = 0.5$) for $\omega = 10, a = \frac{1}{4}, b = -4$ and $c = 1$. Across the studied parameters we obtain for tangential alignment $\|q\nu\| < 10^{-3}$ and for uniaxiality $\left| \beta^2 - \frac{2}{9} \|q\|^2 \right| / \|Q^*\| < 10^{-3}$ at each point of \mathcal{S} .

A.6 Curvature induced biaxiality

Without the penalization (9) the solution space for the tensorial order parameter $Q(q, \beta)$ is not restricted to the uniaxial eigenvalue spectrum $\sigma(Q)$ as defined in (1). To demonstrate this we solve (10) and (11) without the penalization term ($\omega_{\beta} = 0$). Fig. 5A and B shows the equilibrium solution on a flat (2D) and a curved surface, respectively, demonstrating the distortion of uniaxiality by curvature.

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References

- 1 P. G. de Gennes and J. Prost, *The Physics of Liquid Crystals*, Oxford Science Publications, 2nd edn, 1993.



- 2 P. M. Chaikin and T. C. Lubensky, *Principles of condensed matter physics*, Cambridge University Press, 1995.
- 3 A. Fernandez-Nieves, V. Vitelli, A. S. Utada, D. R. Link, M. Marquez, D. R. Nelson and D. A. Weitz, *Phys. Rev. Lett.*, 2007, **99**, 157801.
- 4 T. Lopez-Leon, A. Fernandez-Nieves, M. Nobili and C. Blanc, *J. Phys.: Condens. Matter*, 2012, **24**, 284122.
- 5 H. L. Liang, J. Noh, R. Zentel, P. Rudquist and J. P. F. Lagerwall, *Philos. Trans. R. Soc., A*, 2013, **371**, 20120258.
- 6 H. L. Liang, S. Schymura, P. Rudquist and J. Lagerwall, *Phys. Rev. Lett.*, 2011, **106**, 247801.
- 7 K. May, K. Harth, T. Trittel and R. Stannarius, *EPL*, 2012, **100**, 16003.
- 8 J. P. F. Lagerwall, C. Schütz, M. Salajkova, J. H. Noh, J. H. Park, G. Scalia and L. Bergström, *NPG Asia Mater.*, 2014, **6**, e80.
- 9 L. Jia, A. Cao, D. Lévy, B. Xu, P.-A. Albouy, X. Xiangjun, M. J. Bowick and M.-H. Li, *Soft Matter*, 2009, **5**, 3446.
- 10 B. Xu, R. Piñol, M. Nono-Djamen, S. Pensec, P. Keller, P.-A. Albouy, D. Lévy and M.-H. Li, *Faraday Discuss.*, 2009, **143**, 235.
- 11 L. Jia, D. Lévy, D. Durnad, M. Imperor-Clerc, A. Cao and M.-H. Li, *Soft Matter*, 2011, **7**, 7395.
- 12 L. B. G. Cortes, Y. Gao, R. P. A. Dullens and D. G. A. L. Aarts, *J. Phys.: Condens. Matter*, 2017, **29**, 064003.
- 13 R. E. Guerra, C. P. Kelleher, A. D. Hollingsworth and P. M. Chaikin, *Nature*, 2018, **554**, 346.
- 14 J. Dzubiella, M. Schmidt and H. Löwen, *Phys. Rev. E: Stat. Phys., Plasmas, Fluids, Relat. Interdiscip. Top.*, 2000, **62**, 5081–5091.
- 15 M. A. Bates, G. Skacej and C. Zannoni, *Soft Matter*, 2010, **6**, 665.
- 16 H. Shin, M. J. Bowick and X. Xing, *Phys. Rev. Lett.*, 2008, **101**, 037802.
- 17 S. Dhakal, F. J. Solis and M. Olvera de la Cruz, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2012, **86**, 011709.
- 18 V. Koning, T. Lopez-Leon, A. Fernandez-Nieves and V. Vitelli, *Soft Matter*, 2013, **9**, 4993.
- 19 S. Paquay, G.-J. Both and P. van der Schoot, *Phys. Rev. E*, 2017, **96**, 012611.
- 20 N. D. Bade, R. Kamien, R. Assoian and K. Stebe, *Biophys. J.*, 2017, **112**, 536a.
- 21 H. Stark, *Phys. Rep.*, 2001, **351**, 378.
- 22 P. Prinsen and P. van der Schoot, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2013, **68**, 021701.
- 23 A. Martinez, M. Ravník, B. Lucero, R. Visvanathan, S. Zumer and I. I. Smalyukh, *Nat. Mater.*, 2014, **13**, 259.
- 24 T. Geigenfeind, S. Rosenzweig, M. Schmidt and D. de las Heras, *J. Chem. Phys.*, 2015, **142**, 174701.
- 25 D. Matsunaga, F. Meng, A. Zöttl, R. Golestanian and J. M. Yeomans, *Phys. Rev. Lett.*, 2017, **119**, 198002.
- 26 R. Sknepnek and S. Henkes, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2015, **91**, 022306.
- 27 F. Alaimo, C. Köhler and A. Voigt, *Sci. Rep.*, 2017, **7**, 5211.
- 28 E. Allahyarov, A. Voigt and H. Löwen, *Soft Matter*, 2017, **13**, 8120–8135.
- 29 C. E. Sitta, F. Smallenburg, R. Wittkowski and H. Loewen, *Phys. Chem. Chem. Phys.*, 2018, **20**, 5285–5294.
- 30 E. Allahyarov and H. Löwen, *Soft Matter*, 2018, **14**, 8962–8973.
- 31 S. Kralj, R. Rosso and E. G. Virga, *Soft Matter*, 2011, **7**, 670–683.
- 32 G. Napoli and L. Vergori, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2012, **85**, 061701.
- 33 G. Napoli and L. Vergori, *Phys. Rev. Lett.*, 2012, **108**, 207803.
- 34 D. Jesenek, S. Kralj, R. Rosso and E. G. Virga, *Soft Matter*, 2015, **11**, 2434–2444.
- 35 D. Golovaty, J. A. Montero and P. Sternberg, *J. Nonlinear Sci.*, 2015, **25**, 1431–1451.
- 36 D. Golovaty, J. A. Montero and P. Sternberg, *J. Nonlinear Sci.*, 2017, **27**, 1905–1932.
- 37 M. R. Novack, *SIAM J. Math. Anal.*, 2018, **50**, 6007–6048.
- 38 I. Nitschke, M. Nestler, S. Praetorius, H. Löwen and A. Voigt, *Proc. R. Soc. A*, 2018, **474**, 20170686.
- 39 A. C. Van Enter and S. B. Shlosman, *Phys. Rev. Lett.*, 2002, **89**, 285702.
- 40 A. C. Van Enter, S. Romano and V. A. Zagrebnov, *J. Phys. A: Math. Gen.*, 2006, **39**, L439.
- 41 H. H. Wensink and R. L. C. Vink, *J. Phys.: Condens. Matter*, 2007, **19**, 466109.
- 42 R. L. C. Vink, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2014, **90**, 062132.
- 43 R. Wittmann, C. E. Sitta, F. Smallenburg and H. Löwen, *J. Chem. Phys.*, 2017, **147**, 134908.
- 44 J. Fish and R. Vink, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2010, **81**, 021705.
- 45 A. Majumdar, *Eur. J. Appl. Math.*, 2010, **21**, 181–203.
- 46 J. Straley, *Phys. Rev. A: At., Mol., Opt. Phys.*, 1971, **4**, 675.
- 47 M. J. Stephen and J. P. Straley, *Rev. Mod. Phys.*, 1974, **46**, 617.
- 48 K. Schiele and S. Trimper, *Phys. Status Solidi B*, 1983, **118**, 267–274.
- 49 D. W. Berreman and S. Meiboom, *Phys. Rev. A: At., Mol., Opt. Phys.*, 1984, **30**, 1955–1959.
- 50 T. C. Lubensky, *Phys. Rev. A: At., Mol., Opt. Phys.*, 1970, **2**, 2497–2514.
- 51 L. Longa, D. Monselesan and H.-R. Trebin, *Liq. Cryst.*, 1987, **2**, 769–796.
- 52 J. M. Ball and A. Majumdar, *Mol. Cryst. Liq. Cryst.*, 2010, **525**, 1–11.
- 53 M. E. Gurtin and A. I. Murdoch, *Arch. Ration. Mech. Anal.*, 1975, **57**, 291–323.
- 54 P. Kaiser, W. Wiese and S. Hess, *J. Non-Equilib. Thermodyn.*, 1992, **17**, 153–169.
- 55 W. Kühnel, *Differential geometry*, American Mathematical Soc., 2015, vol. 77.
- 56 I. Nitschke, S. Reuther and A. Voigt, *Phys. Rev. Fluids*, 2019, **4**, 044002.
- 57 I. Nitschke, S. Reuther and A. Voigt, 2019, arXiv:1911.11859.
- 58 A. Segatti, *Math. Models Methods Appl. Sci.*, 2016, **26**, 1865.
- 59 M. Nestler, I. Nitschke, S. Praetorius and A. Voigt, *J. Nonlinear Sci.*, 2018, **28**, 147–191.
- 60 G. Napoli and L. Vergori, *Int. J. Non Linear Mech.*, 2013, **49**, 66–71.
- 61 I. B. Liu, N. Sharifi-Mood and K. J. Stebe, *Annu. Rev. Condens. Matter Phys.*, 2018, **9**, 283–305.
- 62 F. Lou, L. Ye, M. Kong, Q. Yang, G. Lia and Y. Huang, *RSC Adv.*, 2016, **6**, 24195.
- 63 S. Qin and X. Yong, *Soft Matter*, 2019, **15**, 3291–3300.



- 64 M. Nestler, I. Nitschke and A. Voigt, *J. Comput. Phys.*, 2019, **389**, 48–61.
- 65 G. Dziuk and C. M. Elliott, *Acta Numer.*, 2013, **22**, 289–396.
- 66 T. Jankuhn, M. A. Olshanskii and A. Reusken, *Interface. Free Bound.*, 2018, **20**, 353–377.
- 67 P. Hanbo, M. G. Larson and K. Larsson, *IMA J. Num. Anal.*, 2019, DOI: 10.1093/imanum/drz018.
- 68 S. Vey and A. Voigt, *Comput. Visualization Sci.*, 2007, **10**, 57–67.
- 69 T. Witkowski, S. Ling, S. Praetorius and A. Voigt, *Adv. Comput. Math.*, 2015, **41**, 1145.

