

Calhoun: The NPS Institutional Archive

# Elongational perturbations on nematic liquid crystal polymers under a weak shear 

Zhou, Hong

Physics of Fluids / Volume 19
http://hdl.handle.net/10945/25533


Calhoun is a project of the Dudley Knox Library at NPS, furthering the precepts and goals of open government and government transparency. All information contained herein has been approved for release by the NPS Public Affairs Officer.

Dudley Knox Library / Naval Postgraduate School 411 Dyer Road / 1 University Circle Monterey, California USA 93943

# Elongational perturbations on nematic liquid crystal polymers under a weak shear 

Hong Zhou<br>Department of Applied Mathematics, Naval Postgraduate School, Monterey, California 93943, USA<br>Hongyun Wang<br>Department of Applied Mathematics and Statistics, University of California, Santa Cruz, California 95064, USA

(Received 21 March 2007; accepted 11 September 2007; published online 31 October 2007)


#### Abstract

The two-dimensional Smoluchowski equation is employed to study the effect of elongational perturbations on nematic liquid crystal polymers under a weak shear. We use the multiscale asymptotic analysis to show that (1) when the elongational perturbation is small relative to the weak shear, the orientational probability density function (pdf) tumbles periodically only in an intermediate range of polymer concentration; outside this intermediate range (i.e., for very small and very large polymer concentration) the orientational pdf converges to a steady state and there is no tumbling. (2) When the elongational perturbation is about $20 \%$ of the shear rate or larger, the intermediate range of tumbling disappears and the orientational pdf always converges to a steady state regardless of the polymer concentration. Our theoretical predictions are consistent with various earlier results based on the Leslie-Ericksen theory [C. V. Chaubal and L. G. Leal, J. Non-Newtonian Fluid Mech. 82, 22 (1999)] or analogous 3D numerical simulations [M. G. Forest, R. Zhou, and Q. Wang, Phys. Rev. Lett. 93, 088301 (2004); M. G. Forest, Q. Wang, R. Zhou, and E. Choate, J. Non-Newtonian Fluid Mech. 118, 17 (2004)]. © 2007 American Institute of Physics.


[DOI: 10.1063/1.2794002]

## I. INTRODUCTION

Nematic liquid crystal polymers (LCP) are technologically important materials. ${ }^{1,2}$ For example, nematic LCPs have been spun into high strength fibers which were used as airbags that cushioned the landing of NASAs highly successful missions to Mars. ${ }^{3}$ The dynamics of nematic LCPs has been widely modeled by the kinetic Doi-Hess theory. ${ }^{4,5}$ The basic idea behind the Doi-Hess theory is to treat the LCP as a suspension of rigid rodlike nematogenic molecules and describe the ensemble with an orientational probability density function (pdf). The orientational pdf evolves according to a nonlinear Smoluchowski (Fokker-Planck) equation where the forces acting on the rodlike molecules include hydrodynamic, Brownian, and intermolecular forces. The Smoluchowski equation provides a rich variety of mathematical problems and it has been investigated theoretically for pure nematic equilibria, ${ }^{6-12}$ extensional flow-induced equilibria, ${ }^{13}$ equilibria of dipolar ensembles, ${ }^{14-16}$ effect of high ${ }^{17}$ and weak shear, ${ }^{18}$ and effect of coplanar magnetic field. ${ }^{19}$ Numerical simulations of the Smoluchowski equations and the Doi tensor model have been carried out extensively, for example, see Refs. 20-32.

The material properties of the LCPs are highly affected by the processing flows. It is practically important to understand flow effects. Marrucci and Maffettone gave the first elegant two-dimensional analysis on the effect of shear. ${ }^{33-35}$ Later Larson ${ }^{30}$ carried out three-dimensional numerical simulations using spherical harmonic expansions. Larson's results confirmed the theoretical predictions of Marrucci and Maffettone. In 1999 Chaubal and Leal ${ }^{20}$ applied a Lagrang-
ian numerical method called the smoothed particle hydrodynamics technique to explore the effect of flow perturbations on LCPs under a simple shear. They considered elongational perturbations to a simple shear to make it slightly more extensional than a pure shear and found that slight perturbations to the flow field can arrest periodic tumbling behavior. This phenomenon has been observed for both low and high shear rates even though the mechanism is different. In Ref. 20, numerical simulations were carried out in the case of finite Péclet number where the Péclet number denotes the shear rate nondimensionalized with respect to the rotational diffusivity. In the asymptotic limit of small Péclet number, the Leslie-Ericksen theory was used to explore deviations from simple shear. By introducing an effective aspect ratio and plotting the critical value of perturbation $(\alpha)$ as a function of the polymer concentration $(U)$, it was revealed that for $\alpha>0.04$, the orientational director flow-aligns for all $U$; for $\alpha<0.04$, there is an interval $\left(U_{\min }, U_{\max }\right)$ within which the orientational director displays periodic tumbling behavior; outside of the interval $\left(U_{\min }, U_{\max }\right)$, the director flowaligns again. For $U$ less than the isotropic-nematic transition 4.49, the director flow-aligns for all $\alpha \geqslant 0$.

Recently Forest's group did extensive research on various solutions of the 3D Smoluchowski equations under different kinds of conditions. ${ }^{19,22-28,36,37}$ In particular, in Ref. 36 they formulated a monodomain correspondence principle of kinetic and mesoscopic theory for arbitrary aspect ratio nematic polymers in general linear planar flow. From the principle, the monodomain response of nematic polymers for all linear flows in the plane of shear is identical to the mon-
odomain response of another nematic polymer liquid with a renormalized molecular aspect ratio parameter in pure simple shear flow. Later they extended the study in Ref. 36 to track the boundaries of the chaotic region of kinetic theory due to finite molecule aspect ratio and the addition of straining flow in the plane of shear. ${ }^{25}$ It was predicted that sheared chaotic response persists up to a threshold straining flow strength and minimum aspect ratio, beyond which chaotic behavior is arrested. Both ${ }^{25,36}$ gave predictions on how steady and unsteady attractors are modified and eventually lost or persist in the presence of a variable extensional flow contribution.

In the 2D case, it is well-known that in the absence of flow the isotropic-nematic phase transition occurs at $U=2$, where $U$ is the normalized polymer concentration. In the presence of an imposed weak shear there is a threshold $\left(U_{0}\right.$ $\approx 2.41144646$ ) for $U$ : When $U<U_{0}$, steady state solution exists; otherwise there is no steady state and the orientational pdf is temporally periodic ("tumbling") and can never reach a steady state. If one perturbs the weak shear by a small elongation, this scenario will change. In fact, a shear flow can be viewed as the superposition of a rigid body rotation and an elongational flow. So the imposed elongational flow will either increase or decrease the elongational component of the shear flow depending on how their directions are aligned with respect to each other. We will show that if the elongational component is increased by a small fraction, then there is no tumbling at large polymer concentration. This leads to the situation that the orientational pdf tumbles periodically only in an intermediate range of polymer concentrations; outside this intermediate range (below the lower end of the range or above the upper end of the range) the orientational pdf converges to a steady state and there is no tumbling. If the elongational component is increased further (by about $20 \%$ of the original elongational component of the shear flow), the intermediate range of tumbling disappears; that is, for all polymer concentrations, the orientational pdf converges to a steady state. Our theoretical results are consistent with Refs. 20, 25, and 36.

Our asymptotic analysis invokes a multiscale analysis ${ }^{38}$ with the assumption that the solution is a quasisteady state. From the physical point of view, this assumption is reasonable. When the flow field is weak, it takes a long time for the flow field to change the orientation distribution significantly. In other words, the time scale of the flow field affecting the orientation distribution is large. On the other hand, in the absence of flow field, the time scale of the intermolecular force (Maier-Saupe interaction) relaxing the orientation distribution to the Boltzmann form is fixed. So in the presence of a weak flow field, it is reasonable to expect that the intermolecular interaction will be able to drive the orientation distribution approximately to the Boltzmann form before the flow field can take a significant effect on the orientation distribution. This is because the time scale of relaxing to the Boltzmann form is small in comparison to the time scale of the weak flow field. Thus, it is reasonable to assume mathematically that to the leading order the orientation distribution is of the Boltzmann form. Of course, the director of the orientation distribution may change in time driven by the weak flow field. Although this assumption is reasonable on
physical grounds, it is desirable to have a rigorous mathematical justification based on the solvability condition ${ }^{27}$ or other arguments. Currently such a rigorous mathematical justification remains an open problem. We will pursue this issue in our future work. In recent years, multiscale asymptotic analysis has become a powerful mathematical tool in many applications. In Ref. 39, Choate and Forest used it to investigate the viscoelastic response of nematic polymers to small amplitude oscillatory shear using the Doi-Hess mesoscopic orientation tensor model; Vicente Alonso et al. ${ }^{40}$ applied it to study the nonlinear dynamics of a nematic liquid crystal in the presence of a plane Couette flow using a Landau-de Gennes model; Chillingworth et al. ${ }^{41}$ employed it to describe the geometry and dynamics of a nematic liquid crystal in a uniform shear flow.

It is worthwhile to point out that other perturbation schemes have been used to study both the kinetic and tensor models for nematic liquid crystal polymers. For example, Kuzuu and Doi ${ }^{42,43}$ and Semenov ${ }^{44}$ developed a weak-flow asymptotic analysis of the molecular kinetic equation for nematic liquid crystals, which provides a molecular basis for all continuum theory parameters. Doi and Larson ${ }^{45}$ analyzed the effect of steady flow fields on the isotropic-nematic phase transition of rigid rodlike polymers by a singular perturbation method; Forest, Zhou, and Wang ${ }^{27}$ expanded the orientational distribution function in the Péclet number to find approximate steady solutions in weak shear. The works of Refs. 42, 43, 45, and 27 all solved the kinetic equation and derived rigorous "solvability conditions." The advantage of this kind of approach is that unlike multiscale asymptotic analysis, it does not posit a form of the solution. However, these earlier works focused on the persistence of steady states and did not perform constructions of asymptotic periodic solutions. Previous results on periodic solutions have been given by Lee, Forest, and Zhou ${ }^{46}$ using a mesoscopic 2D tensor model. By invoking the Poincaré-Bendixson theorem which describes the long term behavior of orbits of continuous dynamical systems on the plane, they gave an elegantly simple proof that limit cycles ("tumbling orbits") must arise beyond the parameter boundary for the steadyunsteady transition.

The purpose of this paper is to revisit the well-studied elongation-perturbed shear problem from a different theoretical point of view. Our study will provide a detailed analysis on how perturbations of flow conditions affect the system dynamics at small flow rates. We will develop a multiscale analysis directly on the Smoluchowski equation of the kinetic theory. We thus extend results of Chillingworth et al., ${ }^{41}$ Vicente Alfonso et al., ${ }^{40}$ Lee et al., ${ }^{46}$ and Choate et al. ${ }^{39}$ from a tensor model to the kinetic model. Our work differs from Refs. 25 and 36 in three parts: (1) Refs. 25 and 36 are based on numerical simulations of the 3D Smoluchowski equation, whereas ours is on the asymptotic analysis of the 2D Smoluchowski equation; (2) Refs. 25 and 36 include the effect of arbitrary aspect ratio, whereas our study is limited to the infinite aspect ratio; and (3) Refs. 25 and 36 concentrated on the persistence of steady states, whereas our focus is on both steady and unsteady solutions.

In this paper we limit our attention to the two-
dimensional case with a modified weak shear flow. More precisely, the Péclet number from shear flow ( $\varepsilon$ ) is small and the Péclet number from elongation $(q \cdot \varepsilon)$ is of the same order or smaller than that of the shear flow. Our restriction of the orientational probability density from a function on the sphere to the circle is mainly due to its mathematical simplicity, its possible relevance to experimental results on monolayer films of nematic polymers, ${ }^{47,48}$ and its potential guidance to the full 3D orientational distributions. This kind of approach has been widely adopted by others. ${ }^{33,34,46-53}$

This paper is organized as follows: We give a brief description on the mathematical formulation of the Smoluchowski equation coupled with modified shear flow for nematic liquid crystalline polymers in Sec. II. We show the effect of elongational perturbations on the weak shear behavior in Sec. III. Finally, in Sec. IV we summarize our results.

## II. MATHEMATICAL FORMULATION FOR THE TWO-DIMENSIONAL CASE

We briefly review the two-dimensional mathematical formulation of the Doi-Hess kinetic theory for homogeneous flows of rigid rodlike nematogenic molecules immersed in a viscous solvent subject to an imposed flow field. ${ }^{4,5}$ First, we denote the orientation of each polymer rod by an angle $\theta$. Then the orientational direction of a polymer rod is given by $\mathbf{u}=(\cos \theta, \sin \theta)$. To model the nematic tendency of the rodlike molecules to align with each other, it is common to use the short range Maier-Saupe potential,

$$
\begin{align*}
V_{\mathrm{MS}} & =-2 k_{B} T U \mathbf{u u}:\langle\mathbf{u u}\rangle \\
& =-k_{B} T U[\langle\cos 2 \theta\rangle \cos 2 \theta+\langle\sin 2 \theta\rangle \sin 2 \theta] \tag{1}
\end{align*}
$$

where uu and $\mathbf{A}: \mathbf{B}$ are defined as

$$
\begin{align*}
& \mathbf{u u} \equiv\left(\begin{array}{ll}
u_{1} u_{1} & u_{1} u_{2} \\
u_{2} u_{1} & u_{2} u_{2}
\end{array}\right), \\
& \mathbf{A}: \mathbf{B} \equiv \sum_{i, j} a_{i j} b_{i j} . \tag{2}
\end{align*}
$$

In Eq. (1) $U$ is the normalized polymer concentration which describes the strength of intermolecular interactions, $k_{B}$ is the Boltzmann constant, $T$ is the absolute temperature, and $\langle\mathbf{u u}\rangle$ is the second moment of the orientation distribution,

$$
\begin{equation*}
\langle\mathbf{u u}\rangle \equiv \int_{\|\mathbf{u}\|=1} \mathbf{u u} \rho(\mathbf{u}, \mathbf{x}, t) d \mathbf{u} \tag{3}
\end{equation*}
$$

where $\rho(\mathbf{u}, \mathbf{x}, t)$ is the orientational probability density function of the ensemble, i.e., the probability density that a polymer rod has direction $\mathbf{u}$ at location $\mathbf{x}$ and time $t$.

A shear flow with velocity

$$
\begin{equation*}
\mathbf{v}=\gamma(0,-x) \tag{4}
\end{equation*}
$$

can be decomposed into the sum of a rotational flow $\mathbf{v}_{R}$ and an elongational flow $\mathbf{v}_{E}$,

$$
\begin{equation*}
\mathbf{v}=\frac{\gamma}{2}(y,-x)+\frac{\gamma}{2}(-y,-x) \equiv \mathbf{v}_{R}+\mathbf{v}_{E}, \tag{5}
\end{equation*}
$$

where $\gamma$ is the shear rate. For the elongational flow $\mathbf{v}_{E}$ $=\gamma / 2(-y,-x)$, we have

$$
\begin{align*}
\nabla \mathbf{v}_{E} & =\frac{\gamma}{2}\left[\begin{array}{cc}
0 & -1 \\
-1 & 0
\end{array}\right] \\
& =\frac{\gamma}{2}\left[\begin{array}{cc}
\frac{\sqrt{2}}{2} & \frac{\sqrt{2}}{2} \\
-\frac{\sqrt{2}}{2} & \frac{\sqrt{2}}{2}
\end{array}\right]\left[\begin{array}{cc}
1 & 0 \\
0 & -1
\end{array}\right]\left[\begin{array}{cc}
\frac{\sqrt{2}}{2} & -\frac{\sqrt{2}}{2} \\
\frac{\sqrt{2}}{2} & \frac{\sqrt{2}}{2}
\end{array}\right] \tag{6}
\end{align*}
$$

So $\nabla \mathbf{v}_{E}$ has eigenvector $[\sqrt{2} / 2,-\sqrt{2} / 2]^{T}$ associated with the eigenvalue 1 and eigenvector $[\sqrt{2} / 2, \sqrt{2} / 2]^{T}$ associated with the eigenvalue -1 .

The torque induced by the rotational flow $\mathbf{v}_{R}$ on a polymer rod is

$$
\begin{equation*}
\mathbf{F}_{R}=\frac{k_{B} T}{D r} \frac{\gamma}{2}=k_{B} T \frac{\varepsilon}{2}, \tag{7}
\end{equation*}
$$

where $D r$ is the rotational diffusivity of the polymer rod (assumed constant here), and $\varepsilon \equiv \gamma / D r$ is the Péclet number of the shear flow. Similarly, the torque induced by the elongational flow $\mathbf{v}_{E}$ is

$$
\begin{equation*}
\mathbf{F}_{E}=-\frac{k_{B} T}{D r} \frac{\gamma}{2} \cos 2 \theta=-k_{B} T \frac{\varepsilon}{2} \cos 2 \theta \tag{8}
\end{equation*}
$$

If we increase the elongation by a fraction of $q$, then the total elongation is

$$
\begin{equation*}
\mathbf{v}_{E_{T}}=\frac{\gamma}{2}(1+q)(-y,-x) \tag{9}
\end{equation*}
$$

and the torque induced by the total elongation is

$$
\begin{equation*}
\mathbf{F}_{E_{T}}=-k_{B} T \frac{\varepsilon}{2}(1+q) \cos 2 \theta \tag{10}
\end{equation*}
$$

Figure 1 depicts velocity fields of a shear flow and an elongation-perturbed shear flow, respectively.

If we assume the polymer orientation distribution is homogeneous in space, then the orientational pdf under an imposed perturbed shear evolves according to the Smolochowski equation, ${ }^{4}$

$$
\begin{equation*}
\frac{\partial \rho(\theta, t)}{\partial t}=D_{r} \frac{\partial}{\partial \theta}\left\{\left[V_{\mathrm{PS}}^{\prime}(\theta)+V_{\mathrm{MS}}^{\prime}(\theta)\right] \rho+\frac{\partial \rho}{\partial \theta}\right\} \tag{11}
\end{equation*}
$$

where $V_{\mathrm{MS}}(\theta)$ is the Maier-Saupe potential (1) and $V_{\mathrm{PS}}(\theta)$ is the potential due to the external field (i.e., perturbed shear flow)

$$
\begin{equation*}
V_{\mathrm{PS}}(\theta)=-\frac{\varepsilon}{2} \theta+\frac{\varepsilon}{4}(1+q) \sin 2 \theta \tag{12}
\end{equation*}
$$



FIG. 1. (Color online) Velocity field of (a) a shear flow $(q=0)$ and (b) a shear flow plus a $20 \%$ elongational perturbation $(q=0.2)$.

Here, for simplicity, all potentials have been normalized by $k_{B} T$. In Eq. (11), we can set $D_{r}=1$ by introducing the nondimensionalized time $t_{\text {new }}=t D_{r}$. Below we are going to do multiscale asymptotic analysis on (11) with $D_{r}=1$.

## III. MULTISCALE ASYMPTOTIC ANALYSIS

We consider Eq. (11) for small $\varepsilon$ which allows us to carry out a multiscale asymptotic analysis. First, we simplify the expression of the Maier-Saupe potential (1) by selecting an angle $\alpha$ so that $\langle\sin 2(\theta-\alpha)\rangle=0$. Then the Maier-Saupe potential can be written in a compact form,

$$
\begin{equation*}
V_{\mathrm{MS}}(\theta)=-U\langle\cos 2(\theta-\alpha)\rangle \cos 2(\theta-\alpha) . \tag{13}
\end{equation*}
$$

We restrict our application of the multiple scale method to the introduction of two time scales, $T_{0}=t$ and $T_{1}=\varepsilon \cdot t$. Assume that $\rho$ possesses an expansion of the form

$$
\begin{equation*}
\rho(\theta, t)=\rho^{(0)}\left(\theta, T_{0}, T_{1}\right)+\varepsilon \rho^{(1)}\left(\theta, T_{0}, T_{1}\right)+O\left(\varepsilon^{2}\right) \tag{14}
\end{equation*}
$$

and substitute it into Eq. (11) to obtain the leading order equation

$$
\begin{equation*}
\frac{\partial \rho^{(0)}}{\partial T_{0}}=\frac{\partial}{\partial \theta}\left[V_{\mathrm{MS}}^{(0)^{\prime}}(\theta) \rho^{(0)}+\frac{\partial}{\partial \theta} \rho^{(0)}\right] \tag{15}
\end{equation*}
$$

Here $V_{\mathrm{MS}}^{(0)}(\theta)=-U\langle\cos 2(\theta-\alpha)\rangle^{(0)} \cos 2(\theta-\alpha)$ corresponds to the leading order term of the Maier-Saupe potential and the average in the expression is taken with respect to the pdf
$\rho^{(0)}(\theta)$. Since Eq. (15) does not contain an external field effect, we have a pure nematic equation at the leading order. Also notice that Eq. (15) does not contain time scale $T_{1}$. As a result, the orientation distribution will converge to a steady state with respect to time scale $T_{0}$ (it may not be a steady state with respect to time scale $T_{1}$ ). The steady state pdf of Eq. (15) is given by the Boltzmann distribution,

$$
\begin{equation*}
\rho^{(0)}\left(\theta, T_{0}, T_{1}\right)=\frac{\exp [r \cos 2(\theta-\alpha)]}{\int_{0}^{2 \pi} \exp [r \cos 2(\theta-\alpha)] d \theta}, \tag{16}
\end{equation*}
$$

where $r$ is defined as

$$
\begin{equation*}
r=U\langle\cos 2(\theta-\alpha)\rangle^{(0)} \tag{17}
\end{equation*}
$$

Equation (17) is also the equation for determining $r$ for a given value of normalized polymer concentration $U$ in the absence of flow. In the case of weak shear plus weak elongational perturbation, the leading term of $r$ satisfies Eq. (17). We consider the simple situation where the polymer orientation starts with the steady state pdf (16). Then Eq. (15) implies that the pdf remains unchanged with respect to $T_{0}$. Mathematically, this means

$$
\begin{equation*}
\frac{\partial \rho^{(0)}\left(\theta, T_{0}, T_{1}\right)}{\partial T_{0}}=0 . \tag{18}
\end{equation*}
$$

So $\rho^{(0)}\left(\theta, T_{0}, T_{1}\right)$ is independent of $T_{0}$ and thereby "quasisteady." As a result, the phase angle $\alpha$ is also independent of
$T_{0}$. But $\alpha$ may depend on $T_{1}$ and this corresponds to the slow time evolution invoked by the imposed flow field. To find $\alpha\left(T_{1}\right)$, consider the expansion of $\partial \rho(\theta, t) / \partial t$,

$$
\begin{align*}
\frac{\partial \rho(\theta, t)}{\partial t}= & \frac{\partial \rho^{(0)}\left(\theta, T_{0}, T_{1}\right)}{\partial T_{0}}+\varepsilon\left[\frac{\partial \rho^{(0)}\left(\theta, T_{0}, T_{1}\right)}{\partial T_{1}}\right. \\
& \left.+\frac{\partial \rho^{(1)}\left(\theta, T_{0}, T_{1}\right)}{\partial T_{0}}\right]+\cdots . \tag{19}
\end{align*}
$$

Both our extensive numerical simulations and the physical intuitions suggest that $\rho^{(1)}\left(\theta, T_{0}, T_{1}\right)$ is independent of time scale $T_{0}$. Although a rigorous mathematical proof is still an open problem, here in our multiscale expansion, we adopt the assumption that $\rho^{(1)}\left(\theta, T_{0}, T_{1}\right)$ is quasisteady. With this assumption, Eq. (19) becomes

$$
\begin{align*}
\frac{\partial \rho(\theta, t)}{\partial t} & =\varepsilon \frac{\partial \rho^{(0)}\left(\theta, T_{0}, T_{1}\right)}{\partial T_{1}}+O\left(\varepsilon^{2}\right) \\
& =-\varepsilon \frac{\partial \rho^{(0)}\left(\theta, T_{0}, T_{1}\right)}{\partial \theta} \cdot \frac{d \alpha\left(T_{1}\right)}{d T_{1}}+O\left(\varepsilon^{2}\right) \tag{20}
\end{align*}
$$

On substituting this into Eq. (11) and rearranging terms we obtain

$$
\begin{align*}
& \frac{\partial}{\partial \theta}\left[\left(-\frac{\varepsilon}{2}+\frac{\varepsilon}{2}(1+q) \cos 2 \theta+V_{\mathrm{MS}}^{\prime}(\theta)\right) \rho(\theta)+\frac{\partial \rho(\theta)}{\partial \theta}\right. \\
& \left.\quad+\varepsilon \rho^{(0)}(\theta) \frac{d \alpha\left(T_{1}\right)}{d T_{1}}+O\left(\varepsilon^{2}\right)\right]=0 . \tag{21}
\end{align*}
$$

This suggests

$$
\begin{align*}
& {\left[-\frac{\varepsilon}{2}+\frac{\varepsilon}{2}(1+q) \cos 2 \theta+V_{\mathrm{MS}}^{\prime}(\theta)\right] \rho(\theta)+\frac{\partial \rho(\theta)}{\partial \theta}+\varepsilon \rho^{(0)}} \\
& \quad \times(\theta) \frac{d \alpha\left(T_{1}\right)}{d T_{1}}+O\left(\varepsilon^{2}\right)=-\varepsilon J_{1} \tag{22}
\end{align*}
$$

where $\varepsilon J_{1}$ is the integration constant of the order $O(\varepsilon)$. In Eq. (22) both $d \alpha\left(T_{1}\right) / d T_{1}$ and $J_{1}$ are unknown so we need two linear equations to determine them. To do so, we first integrate Eq. (22) from 0 to $2 \pi$ to obtain

$$
\begin{align*}
-\frac{\varepsilon}{2} & +\frac{\varepsilon}{2}(1+q) \int_{0}^{2 \pi} \cos 2 \theta \rho(\theta) d \theta+\int_{0}^{2 \pi} V_{\mathrm{MS}}^{\prime}(\theta) \rho(\theta) d \theta \\
& +\varepsilon \frac{d \alpha\left(T_{1}\right)}{d T_{1}}+O\left(\varepsilon^{2}\right)=-2 \pi \varepsilon J_{1} \tag{23}
\end{align*}
$$

The second term on the left of Eq. (23) has the expression

$$
\begin{align*}
\int_{0}^{2 \pi} \cos 2 \theta \rho(\theta) d \theta= & \langle\cos 2 \theta\rangle \\
= & \langle\cos 2(\theta-\alpha) \cos 2 \alpha \\
& -\sin 2(\theta-\alpha) \sin 2 \alpha\rangle \\
= & \langle\cos 2(\theta-\alpha) \cos 2 \alpha\rangle \\
= & \langle\cos 2(\theta-\alpha)\rangle^{(0)} \cos 2 \alpha+O(\varepsilon) \\
= & \frac{r}{U} \cos 2 \alpha+O(\varepsilon) \tag{24}
\end{align*}
$$

where $r / U \equiv\langle\cos 2(\theta-\alpha)\rangle^{(0)}$ is the order parameter. The third term on the left of Eq. (23) has the expression

$$
\begin{align*}
\int_{0}^{2 \pi} V_{\mathrm{MS}}^{\prime}(\theta) \rho(\theta) d \theta & =\int_{0}^{2 \pi} \frac{d}{d \theta}[-r \cos 2(\theta-\alpha)] \rho(\theta) d \theta \\
& =\int_{0}^{2 \pi} 2 r \sin 2(\theta-\alpha) \rho(\theta) d \theta \\
& =2 r\langle\sin 2(\theta-\alpha)\rangle=0 \tag{25}
\end{align*}
$$

where we have used the fact that the angle $\alpha$ is selected to make $\langle\sin 2(\theta-\alpha)\rangle=0$. With these expressions, Eq. (23) becomes

$$
\begin{equation*}
-\frac{1}{2}+\frac{1+q}{2} \frac{r}{U} \cos \left[2 \alpha\left(T_{1}\right)\right]+\frac{d \alpha\left(T_{1}\right)}{d T_{1}}+O(\varepsilon)=-2 \pi J_{1} . \tag{26}
\end{equation*}
$$

To find a second equation for $d \alpha\left(T_{1}\right) / d T_{1}$ and $J_{1}$, we manipulate Eq. (22) by dividing it by $\varepsilon \rho(\theta)$, approximating $\rho(\theta)$ by $\rho^{(0)}(\theta)$, and integrating from 0 to $2 \pi$ to obtain

$$
\begin{align*}
\int_{0}^{2 \pi} & \left(-\frac{1}{2}\right) d \theta+\int_{0}^{2 \pi} \frac{1}{2}(1+q) \cos 2 \theta d \theta+\frac{1}{\varepsilon} \int_{0}^{2 \pi} V_{\mathrm{MS}}^{\prime} d \theta \\
& +\frac{1}{\varepsilon} \int_{0}^{2 \pi} \frac{1}{\rho^{(0)}(\theta)} \frac{\partial \rho^{(0)}(\theta)}{\partial \theta} d \theta+\int_{0}^{2 \pi} \frac{d \alpha\left(T_{1}\right)}{d T_{1}} d \theta+O(\varepsilon) \\
& =-\int_{0}^{2 \pi} \frac{J_{1}}{\rho^{(0)}(\theta)} d \theta \tag{27}
\end{align*}
$$

Note that the integrands $\cos 2 \theta, V_{\mathrm{MS}}^{\prime}$, and $1 / \rho^{(0)}(\theta) \partial \rho^{(0)}$ $\times(\theta) / \partial \theta=\partial \ln \left[\rho^{(0)}(\theta)\right] / \partial \theta$ are all periodic on $[0,2 \pi]$, so the second, third, and fourth definite integrals in Eq. (27) vanish. Using Eq. (16) and carrying out the details, we get

$$
\begin{equation*}
-\pi+2 \pi \frac{d \alpha\left(T_{1}\right)}{d T_{1}}+O(\varepsilon)=-(2 \pi)^{2} g(r) J_{1} \tag{28}
\end{equation*}
$$

where the function $g(r)$ is defined as

$$
\begin{equation*}
g(r) \equiv\left[\frac{1}{2 \pi} \int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta\right]^{2} \tag{29}
\end{equation*}
$$

Multiplying Eq. (26) by a factor $2 \pi g(r)$ and subtracting it from Eq. (28) leads to

$$
\begin{align*}
-\pi+2 \pi \frac{d \alpha\left(T_{1}\right)}{d T_{1}}= & 2 \pi g(r)\left[-\frac{1}{2}+\frac{1+q}{2} \frac{r}{U} \cos \left[2 \alpha\left(T_{1}\right)\right]\right. \\
& \left.+\frac{d \alpha\left(T_{1}\right)}{d T_{1}}\right] \tag{30}
\end{align*}
$$

Solving for $d \alpha\left(T_{1}\right) / d T_{1}$ yields

$$
\begin{align*}
\frac{d \alpha\left(T_{1}\right)}{d T_{1}}= & \frac{1}{2[g(r)-1]}\left[-1+g(r)\left(1-\frac{(1+q) r}{U}\right.\right. \\
& \left.\left.\times \cos \left[2 \alpha\left(T_{1}\right)\right]\right)\right] \tag{31}
\end{align*}
$$

Applying the double angle formula $\cos \left[2 \alpha\left(T_{1}\right)\right]=1$ $-2 \sin ^{2}\left[\alpha\left(T_{1}\right)\right]$, we obtain after some manipulations

$$
\begin{equation*}
\frac{d \alpha\left(T_{1}\right)}{d T_{1}}=c_{1} \cdot\left\{\sin ^{2}\left[\alpha\left(T_{1}\right)\right]+c_{2}\right\}, \tag{32}
\end{equation*}
$$

where $c_{1}$ and $c_{2}$ are

$$
\begin{align*}
& c_{1}=\frac{g(r)}{g(r)-1} \cdot \frac{r}{U} \cdot(1+q), \\
& c_{2}=\frac{-1}{2(1+q)}[p(r)+q] . \tag{33}
\end{align*}
$$

Here the function $p(r)$ is defined as

$$
\begin{equation*}
p(r) \equiv \frac{U}{r}\left[\frac{1}{g(r)}-1\right]+1 \tag{34}
\end{equation*}
$$

Note that in the order parameter $r / U \equiv\langle\cos 2(\theta-\alpha)\rangle^{(0)}$, probability density (16) is used in calculating the average. As a function of $r, r / U$ has the expression

$$
\begin{equation*}
\frac{r}{U}=\frac{\int_{0}^{2 \pi} \cos 2 \theta \exp (r \cos 2 \theta) d \theta}{\int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta} \tag{35}
\end{equation*}
$$

As a matter of fact Eq. (35) is the equation for determining $r$ for a given value of normalized polymer concentration $U$. Here for mathematical convenience, we use $r$ (instead of $U$ ) as the independent variable. Note that, for $r>0$, the mapping between $r$ and $U$ as given in Eq. (35) is monotonic (see Appendix A for a mathematical derivation of the monotonicity). As a consequence of the monotonicity, for $r>0$, the mapping between $r$ and $U$ is one-to-one. Therefore, a result in terms of $r$ can easily be translated to the corresponding result in terms of $U$. Below we will first discuss the dynamic behavior of $\alpha\left(T_{1}\right)$ governed by Eq. (32) for various regions of $r$. Then all conclusions will be restated in terms of $U$, the normalized polymer concentration.

Equation (32) describes the dynamics of the slow time evolution of the phase angle $\alpha\left(T_{1}\right)$. In Eq. (32), the factor $c_{1}$ is always positive for $r>0$. If $c_{2}$ is negative (and $\left|c_{2}\right|<1$ ), then the phase angle $\alpha\left(T_{1}\right)$ will stop at an angle $\alpha_{0}$ satisfying $\sin ^{2} \alpha_{0}+c_{2}=0$. In other words, the polymer orientation distribution will converge to a steady state. If $c_{2}$ is positive, then the phase angle $\alpha\left(T_{1}\right)$ will keep increasing and the polymer orientation distribution will rotate periodically in time (tumbling). Thus, positive $c_{2}$ causes tumbling. The sign of $c_{2}$ is the opposite of the sign of $p(r)+q$. It follows that negative value of $p(r)+q$ corresponds to tumbling. Figure 2 shows the graph of function $p(r)$ and its asymptotic approximation as $r \rightarrow+\infty$,

$$
\begin{equation*}
p(r)=\frac{-1}{2 r}\left(1+\frac{3}{4 r}\right)+\cdots \quad \text { as } r \rightarrow+\infty . \tag{36}
\end{equation*}
$$

The asymptotic approximation of $p(r)$ as $r \rightarrow+\infty$ is derived in Appendix B. As shown in Fig. 2, the function $p(r)$ starts at $r=0$ with $p(0)=1$. As $r$ increases from 0 to $r_{\min }$, the function $p(r)$ decreases monotonically. In particular, the function $p(r)$ intersects with the $x$-axis at $r_{0}=1.3276$. At $r_{\text {min }}=2.6463$, the function $p(r)$ attains its minimum, and the minimum value is $\min _{r} p(r)=-0.18754$. For $r>r_{\text {min }}$, function $p(r)$ increases monotonically. For $r>r_{\text {min }}$, function $p(r)$ is well approxi-


FIG. 2. (Color online) Graph of function $p(r)$ and its asymptotic approximation as $r \rightarrow+\infty$. The function $p(r)$ intersects with the $x$-axis at $r_{0}$ $=1.3276$. At $r_{\text {min }}=2.6463$, the function $p(r)$ attains its minimum, and the minimum value is $\min _{r} p(r)=-0.18754$.
mated by its asymptotic expression at $r=+\infty$.
Figure 3(a) shows the tumbling region (shaded) in the $(q, r)$ space. Here the tumbling region is the largest region where $p(r)+q$ is negative. Recall from our discussion above that negative value of $p(r)+q$ corresponds to tumbling. So if the value of $(q, r)$ falls in the tumbling region, then the polymer orientation rotates periodically in time (tumbling). From the asymptotic expression of function $p(r)$ for large $r$, we see that for small $q$, the upper boundary of the tumbling region in the $(q, r)$ space is approximated by

$$
\begin{equation*}
q=\frac{1}{2 r}\left(1+\frac{3}{4 r}\right) . \tag{37}
\end{equation*}
$$

Solving $r$ in terms of $q$, we obtain

$$
\begin{equation*}
r_{\text {upper }}=\frac{1+\sqrt{1+6 q}}{4 q} \tag{38}
\end{equation*}
$$

This asymptotic expression is valid for small $q$.
Because of the one-to-one correspondence between $r$ and $U$, a region in the $(q, r)$ space can be uniquely mapped to a region in the $(q, U)$ space. Figure $3(\mathrm{~b})$ shows the corresponding tumbling region in the $(q, U)$ space, where $U$ is the normalized polymer concentration and $q$ is the magnitude of the elongational perturbation relative to the original elongational component of the shear flow. Again, for small $q$ (corresponding to large $r$ and large $U$ ), the upper boundary of the tumbling region in the $(q, U)$ space can be well approximated by using the asymptotic expression of function $p(U)$ for large $U$ (which is derived and presented in Appendix B). For small $q$, the upper boundary in $U$ satisfies approximately

$$
\begin{equation*}
q=\frac{1}{2 U}\left(1+\frac{5}{4 U}\right) . \tag{39}
\end{equation*}
$$

Solving $U$ in terms of $q$, we obtain


FIG. 3. (Color online) Tumbling regions are shown as shaded. (a) Tumbling region in the ( $q, r$ ) space. (b) Tumbling region in the ( $q, U$ ) space. The critical value for $q$ is $q_{c}=0.18754$.

$$
\begin{equation*}
U_{\text {upper }}=\frac{1+\sqrt{1+10 q}}{4 q} \tag{40}
\end{equation*}
$$

This asymptotic expression is valid for small $q$. In Fig. 3, the asymptotic approximation of the upper bound of the tumbling region for small $q$ is shown as a dashed line and it agrees well with the exact solution.

Let $q_{c} \equiv-\min _{r} p(r)=0.18154 . q_{c}$ is the critical value of $q$, when the $q>q_{c}$ function $p(r)+q$ is positive for all values of $r>0$; when $0<q<q_{c}$ the function $p(r)+q$ may be negative for certain values of $r$. As shown in Fig. 3, when $0<q$ $<q_{c}$ tumbling occurs only for $r(U)$ in a finite interval. As $q$ increases toward the critical value $q_{c}$, this tumbling interval gets smaller and smaller. When $q>q_{c}$, the tumbling interval disappears and there is no tumbling. That is, with an elongational perturbation of less than $20 \% \quad\left(q_{c}=0.18154\right)$ of the original elongational component of the shear flow, the tumbling disappears for all values of $U$.

## IV. CONCLUSIONS

We have applied the multiscale asymptotic analysis to study the effect of elongational perturbations on the weak shear behavior of nematic liquid crystal polymers. It is found that when the elongational perturbation is small, the tumbling behavior occurs only in an intermediate range of polymer concentrations; when the elongational perturbation is above about $20 \%$ of the shear rate, tumbling disappears for all polymer concentrations. These findings suggest a mechanism on the control of the structure of nematic liquid crystal polymers.

## ACKNOWLEDGMENTS

The authors thank Professor M. Gregory Forest for introducing them to the field of complex fluids many years ago, and for his inspiration, support, and encouragement ever since. The authors thank the anonymous referees for their constructive and critical suggestions on improving this manuscript. This work was partially supported by the Air Force Office of Scientific Research under Grant No. F1ATA06313G003 and by the National Science Foundation.

## APPENDIX A: MONOTONICITY OF $\boldsymbol{U}(r)$

We first apply integration by parts to the numerator on the right-hand side of Eq. (35),

$$
\begin{align*}
& \int_{0}^{2 \pi} \cos 2 \theta \exp (r \cos 2 \theta) d \theta \\
& \quad=\frac{1}{2} \int_{0}^{2 \pi} \exp (r \cos 2 \theta) d(\sin 2 \theta) \\
& \quad=r \int_{0}^{2 \pi} \sin ^{2} 2 \theta \exp (r \cos 2 \theta) d \theta \\
& \quad=r \int_{0}^{2 \pi}\left(1-\cos ^{2} 2 \theta\right) \exp (r \cos 2 \theta) d \theta \tag{A1}
\end{align*}
$$

Substituting this result into Eq. (35) and writing $U$ as a function of $r$, we have

$$
\begin{equation*}
\frac{1}{U(r)}=1-\frac{\int_{0}^{2 \pi} \cos ^{2} 2 \theta \exp (r \cos 2 \theta) d \theta}{\int_{0}^{2 \pi} \exp (r \cos \theta) d \theta} \tag{A2}
\end{equation*}
$$

To show that $U(r)$ is monotonically increasing, we only need to show

$$
\begin{equation*}
w(r) \equiv \frac{\int_{0}^{2 \pi} \cos ^{2} 2 \theta \exp (r \cos 2 \theta) d \theta}{\int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta} \tag{A3}
\end{equation*}
$$

is monotonically increasing. Function $w(r)$ can be viewed as an average, $w(r)=\left\langle\cos ^{2} 2 \theta\right\rangle$ with probability density

$$
\begin{equation*}
\rho(\theta, r)=\frac{\exp (r \cos 2 \theta)}{\int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta} \tag{A4}
\end{equation*}
$$

The derivative of the probability density with respect to $r$ is

$$
\begin{equation*}
\frac{d \rho(\theta, r)}{d r}=(\cos 2 \theta-\langle\cos 2 \theta\rangle) \rho(\theta, r) . \tag{A5}
\end{equation*}
$$

The first derivative of $w(r)$ is

$$
\begin{equation*}
w^{\prime}(r)=\left\langle\cos ^{2} 2 \theta(\cos 2 \theta-\langle\cos 2 \theta\rangle)\right\rangle \tag{A6}
\end{equation*}
$$

The second derivative of $w(r)$ is

$$
\begin{align*}
w^{\prime \prime}(r)= & \left\langle\cos ^{2} 2 \theta(\cos 2 \theta-\langle\cos 2 \theta\rangle)^{2}\right\rangle \\
& -\left\langle\cos ^{2} 2 \theta\right\rangle \cdot\left(\left\langle\cos ^{2} 2 \theta\right\rangle-\langle\cos 2 \theta\rangle^{2}\right) \\
= & -2\langle\cos 2 \theta\rangle w^{\prime}(r)+\operatorname{var}\left(\cos ^{2} 2 \theta\right) \tag{A7}
\end{align*}
$$

Thus, function $w(r)$ satisfies the special property that whenever $w^{\prime}(r)=0$ we have $w^{\prime \prime}(r)>0$. At $r=0$, we have $w(0)$ $=0.5$ and $w^{\prime}(0)=0$. The special property of $w(r)$ guarantees that $w^{\prime}(r)>0$ for $r>0$. This leads immediately to the conclusion that $U(r)$ is monotonically increasing.

## APPENDIX B: ASYMPTOTIC APPROXIMATION OF $p(r)$

To derive an asymptotic approximation for $p(r)$ as $r \rightarrow+\infty$, we need to deal with integrals such as $\int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta$. First, we use the symmetry of $\cos 2 \theta$ to rewrite the integral as

$$
\begin{equation*}
\int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta=4 \int_{0}^{\pi / 2} \exp (r \cos 2 \theta) d \theta \tag{B1}
\end{equation*}
$$

To apply Watson's lemma ${ }^{38}$ on the integral, we use a change of variable,

$$
\begin{align*}
& s=(1-\cos 2 \theta) / 2 \\
& d s=\sin 2 \theta d \theta=2 \sin \theta \cos \theta d \theta=2 \sqrt{s} \sqrt{1-s} d \theta  \tag{B2}\\
& d \theta=\frac{1}{2} \frac{1}{\sqrt{s} \sqrt{1-s}} d s
\end{align*}
$$

Applying the change of variables to the integral yields

$$
\begin{align*}
\int_{0}^{2 \pi} & \exp (r \cos 2 \theta) d \theta \\
& =2 \int_{0}^{1} \exp [r(1-2 s)] \frac{1}{\sqrt{s} \sqrt{1-s}} d s \\
& =2 \exp (r) \int_{0}^{1} \exp (-2 r s) \frac{1}{\sqrt{s}}\left(1+\frac{1}{2} s+\cdots\right) d s \\
& =2 \exp (r)\left(\frac{\Gamma(1 / 2)}{\sqrt{2 r}}+\frac{1}{2} \cdot \frac{\Gamma(3 / 2)}{2 r \sqrt{2 r}}+\cdots\right) \\
& =\frac{\sqrt{2 \pi}}{\sqrt{r}} \exp (r)\left(1+\frac{1}{8} \cdot \frac{1}{r}+\cdots\right) \tag{B3}
\end{align*}
$$

Substituting this result into function $g(r)$, we have

$$
\begin{align*}
g(r) & =\left(\frac{1}{2 \pi} \int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta\right)^{2} \\
& =\frac{1}{2 \pi} \cdot \frac{\exp (r)}{r}(1+\cdots) . \tag{B4}
\end{align*}
$$

That is, as $r \rightarrow+\infty$, function $g(r)$ is exponentially large and consequently $1 / g(r)$ is exponentially small. In a similar way, we obtain

$$
\begin{align*}
\int_{0}^{2 \pi} & (1-\cos 2 \theta) \exp (r \cos 2 \theta) d \theta \\
& =4 \int_{0}^{1} \exp [r(1-2 s)] \frac{s}{\sqrt{s} \sqrt{1-s}} d s \\
& =4 \exp (r) \int_{0}^{1} \exp (-2 r s) \sqrt{s}\left(1+\frac{1}{2} s+\cdots\right) d s \\
& =4 \exp (r)\left(\frac{\Gamma(3 / 2)}{2 r \sqrt{2 r}}+\frac{1}{2} \cdot \frac{\Gamma(5 / 2)}{(2 r)^{2} \sqrt{2 r}}+\cdots\right) \\
& =\frac{\sqrt{2 \pi}}{\sqrt{r}} \exp (r)\left(\frac{1}{2} \cdot \frac{1}{r}+\frac{3}{16} \cdot \frac{1}{r^{2}}+\cdots\right) \tag{B5}
\end{align*}
$$

Combining Eqs. (B3) and (B5) gives us

$$
\begin{align*}
1-\langle\cos 2 \theta\rangle & =\frac{\int_{0}^{2 \pi}(1-\cos 2 \theta) \exp (r \cos 2 \theta) d \theta}{\int_{0}^{2 \pi} \exp (r \cos 2 \theta) d \theta} \\
& =\frac{\frac{1}{2} \cdot \frac{1}{r}+\frac{3}{16} \cdot \frac{1}{r^{2}}+\cdots}{1+\frac{1}{8} \cdot \frac{1}{r}+\cdots}=\frac{1}{2 r}\left(1+\frac{1}{4 r}+\cdots\right) \tag{B6}
\end{align*}
$$

and

$$
\begin{equation*}
\langle\cos 2 \theta\rangle=1-(1-\langle\cos 2 \theta\rangle)=1-\frac{1}{2 r}+\cdots \tag{B7}
\end{equation*}
$$

To find an asymptotic approximation for function $p(r)$, we rewrite it as

$$
\begin{align*}
p(r) & =\frac{1}{\langle\cos 2 \theta\rangle}\left(\frac{1}{g(r)}-1\right)+1 \\
& =\frac{1}{\langle\cos 2 \theta\rangle} \cdot \frac{1}{g(r)}-\frac{1-\langle\cos 2 \theta\rangle}{\langle\cos 2 \theta\rangle} . \tag{B8}
\end{align*}
$$

The first term on the right-hand side is exponentially small. The second term can be expanded using results (B6) and (B7) obtained above,

$$
\begin{align*}
p(r) & =-\frac{1-\langle\cos 2 \theta\rangle}{\langle\cos 2 \theta\rangle}+\cdots=-\frac{\frac{1}{2 r}\left(1+\frac{1}{4 r}+\cdots\right)}{1-\frac{1}{2 r}+\cdots}+\cdots \\
& =\frac{-1}{2 r}\left(1+\frac{3}{4 r}+\cdots\right) . \tag{B9}
\end{align*}
$$

Now we derive the expansion of function $p(r)$ in terms of $U$. As $r \rightarrow+\infty, U$, as a function of $r$, has the expansion

$$
\begin{equation*}
U(r)=\frac{r}{\langle\cos 2 \theta\rangle}=r\left(1+\frac{1}{2 r}\right)+\cdots . \tag{B10}
\end{equation*}
$$

Expressing $r$ in terms of $U$ yields

$$
\begin{equation*}
r(U)=U\left(1-\frac{1}{2 U}\right)+\cdots \tag{B11}
\end{equation*}
$$

Substituting into Eq. (B7), we arrive at

$$
\begin{align*}
p(U) & =p(r(U))=\frac{-1}{2 r(U)}\left(1+\frac{3}{4 r(U)}+\cdots\right) \\
& =\frac{-1}{2 U}\left(1+\frac{5}{4 U}+\cdots\right) \tag{B12}
\end{align*}
$$

${ }^{1}$ A. D. Rey and M. M. Denn, "Dynamical phenomena in liquid-crystalline materials," Annu. Rev. Fluid Mech. 34, 233 (2002).
${ }^{2}$ B. Bird, R. C. Armstrong, and O. Hassager, Dynamics of Polymeric Liquids (Wiley, New York, 1987), Vol. 1.
${ }^{3}$ A. M. Donald, A. H. Windle, and S. Hanna, Liquid Crystalline Polymers, 2nd ed. (Cambridge University Press, Cambridge, 2006).
${ }^{4}$ M. Doi and S. F. Edwards, The Theory of Polymer Dynamics (Oxford University Press, New York, 1986).
${ }^{5}$ S. Z. Hess, "Fokker-Planck-equation approach to flow alignment in liquid crystals," Z. Naturforsch. A 31, 1034 (1976).
${ }^{6}$ P. Constantin, I. Kevrekidis, and E. S. Titi, "Asymptotic states of a Smoluchowski equation," Arch. Ration. Mech. Anal. 174, 365 (2004).
${ }^{7}$ P. Constantin, I. Kevrekidis, and E. S. Titi, "Remarks on a Smoluchowski equation," Discrete Contin. Dyn. Syst. 11, 101 (2004).
${ }^{8}$ P. Constantin and J. Vukadinovic, "Note on the number of steady states for a 2D Smoluchowski equation," Nonlinearity 18, 441 (2005).
${ }^{9}$ I. Fatkullin and V. Slastikov, "Critical points of the Onsager functional on a sphere," Nonlinearity 18, 2565 (2005).
${ }^{10}$ H. Liu, H. Zhang, and P. Zhang, "Axial symmetry and classification of stationary solutions of Doi-Onsager equation on the sphere with MaierSaupe potential," Commun. Math. Sci. 3, 201 (2005).
${ }^{11}$ C. Luo, H. Zhang, and P. Zhang, "The structure of equilibrium solution of 1D Smoluchowski equation," Nonlinearity 18, 379 (2005).
${ }^{12}$ H. Zhou, H. Wang, M. G. Forest, and Q. Wang, "A new proof on axisymmetric equilibria of a three-dimensional Smoluchowski equation," Nonlinearity 18, 2815 (2005).
${ }^{13}$ Q. Wang, S. Sircar, and H. Zhou, "Steady state solutions of the Smoluchowski equation for rigid nematic polymers under imposed fields," Optim. 3, 605 (2005).
${ }^{14}$ G. Ji, Q. Wang, P. Zhang, and H. Zhou, "Study of phase transition in homogeneous, rigid extended nematics and magnetic suspensions using an
order-reduction method," Phys. Fluids 18, 123103 (2006).
${ }^{15}$ H. Zhou, H. Wang, Q. Wang, and M. G. Forest, "Characterization of stable kinetic equilibria of rigid, dipolar rod ensembles for coupled dipole-dipole and Maier-Saupe potentials," Nonlinearity 20, 277 (2007).
${ }^{16}$ H. Zhou, H. Wang, and Q. Wang, "Nonparallel solutions of extended nematic polymers under an external field," Discrete Contin. Dyn. Syst., Ser. B 7, 907 (2007).
${ }^{17}$ A. Zarnescu, "The stationary 2D Smoluchowski equation in strong homogeneous flow," Nonlinearity 19, 1619 (2006).
${ }^{18} \mathrm{H}$. Zhou and H. Wang, "Steady states and dynamics of 2-D nematic polymers driven by an imposed weak shear," Commun. Math. Sci. 5, 113 (2007).
${ }^{19}$ M. G. Forest, S. Sircar, Q. Wang, and R. Zhou, "Monodomain dynamics for rigid rod and platelet suspensions in strongly coupled coplanar linear flow and magnetic fields. II. Kinetic theory," Phys. Fluids 18, 103102 (2006).
${ }^{20}$ C. V. Chaubal and L. G. Leal, "Smoothed particle hydrodynamics techniques for the solution of kinetic theory problems. Part 2. The effect of flow perturbations on the simple shear behavior of LCPs," J. NonNewtonian Fluid Mech. 82, 25 (1999).
${ }^{21}$ G. Sgalari, G. L. Leal, and J. J. Feng, "The shear flow behavior of LCPs based on a generalized Doi model with distortional elasticity," J. NonNewtonian Fluid Mech. 102, 361 (2002).
${ }^{22}$ M. G. Forest, R. Zhou, and Q. Wang, "Symmetries of the Doi kinetic theory for nematic polymers of arbitrary aspect ratio: At rest and in linear flows," Phys. Rev. E 66, 031712 (2002).
${ }^{23}$ M. G. Forest, Q. Wang, and R. Zhou, "The flow-phase diagram of DoiHess theory for sheared nematic polymers II: Finite shear rates," Rheol. Acta 44, 80 (2004).
${ }^{24}$ M. G. Forest, R. Zhou, and Q. Wang, "The weak shear phase diagram for nematic polymers," Rheol. Acta 43, 17 (2004).
${ }^{25}$ M. G. Forest, R. Zhou, and Q. Wang, "Chaotic boundaries of nematic polymers in mixed shear and extensional flows," Phys. Rev. Lett. 93, 088301 (2004).
${ }^{26}$ M. G. Forest, R. Zhou, and Q. Wang, "Kinetic structure simulations of nematic polymers in plane Couette cells, I: The algorithm and benchmarks," SIAM J. Math. Anal. 3, 853 (2005).
${ }^{27}$ M. G. Forest, R. Zhou, and Q. Wang, "Scaling behavior of kinetic orientational distributions for dilute nematic polymers in weak shear," J. NonNewtonian Fluid Mech. 116, 183 (2004).
${ }^{28}$ M. G. Forest and Q. Wang, "Monodomain response of finite-aspect-ratio macromolecules in shear and related linear flows," Rheol. Acta 42, 20 (2003).
${ }^{29}$ S. Hess and M. Kroger, "Regular and chaotic orientational and rheological behaviour of liquid crystals," J. Phys.: Condens. Matter 16, S3835 (2004).
${ }^{30}$ R. G. Larson, "Arrested tumbling in shearing flows of liquid crystal polymers," Macromolecules 23, 3983 (1990).
${ }^{31}$ V. Faraoni, M. Grosso, S. Crescitelli, and P. L. Maffetone, "The rigid-rod model for nematic polymers: An analysis of the shear flow problem," Macromolecules 43, 829 (1999).
${ }^{32}$ F. Cocchini, C. Aratari, and G. Marruucci, "Tumbling of rodlike polymers in the liquid crystalline phase under shear flow," Macromolecules 23, 4446 (1990).
${ }^{33}$ G. Marrucci and P. L. Maffettone, "Description of the liquid-crystalline phase of rodlike polymers at high shear rates," Macromolecules 22, 4446 (1989).
${ }^{34}$ G. Marrucci and P. L. Maffettone, "Nematic phase of rodlike polymers. I. Prediction of transient behavior at high shear rates," J. Rheol. 34, 1217 (1990).
${ }^{35}$ P. L. Maffettone and G. Marrucci, "A two-dimensional approach to the constitutive equation of nematic polymers," J. Non-Newtonian Fluid Mech. 38, 273 (1991).
${ }^{36}$ M. G. Forest, Q. Wang, R. Zhou, and E. Choate, "Monodomain response of arbitrary aspect ratio nematic polymers in general linear planar flows," J. Non-Newtonian Fluid Mech. 118, 17 (2004).
${ }^{37}$ M. G. Forest, Q. Wang, and R. Zhou, "Monodomain dynamics for rigid rod and platelet suspensions in strongly coupled coplanar linear flow and magnetic fields," J. Rheol. 51, 1 (2007).
${ }^{38}$ A. W. Bush, Perturbation Methods for Engineers and Scientists (CRC, Boca Raton, 1992).
${ }^{39}$ E. P. Choate and M. G. Forest, "A classical problem revisited: Rheology of nematic polymer monodomains in small amplitude oscillatory shear," Rheol. Acta 46, 83 (2006).
${ }^{40}$ E. Vicente Alonso, A. A. Wheeler, and T. J. Sluckin, "Nonlinear dynamics
of a nematic liquid crystal in the presence of a shear flow," Proc. R. Soc. London, Ser. A 459, 195 (2003).
${ }^{41}$ D. R. J. Chillingworth, E. Vicente Alonso, and A. A. Wheeler, "Geometry and dynamics of a nematic liquid crystal in a uniform shear flow," J. Phys. Soc. Jpn. 34, 1393 (2001).
${ }^{42}$ N. Kuzuu and M. Doi, "Constitutive equation for nematic liquid crystals under weak velocity gradient derived from a molecular kinetic equation," J. Phys. Soc. Jpn. 52, 3486 (1983).
${ }^{43}$ N. Kuzuu and M. Doi, "Constitutive equation for nematic liquid crystals under weak velocity gradient derived from a molecular kinetic equation. II. Leslie coefficients for rodlike polymers," J. Phys. Soc. Jpn. 53, 1031 (1984).
${ }^{44}$ A. N. Semenov, "Rheological properties of a liquid crystal solution of rod-like molecules," Sov. Phys. JETP 66, 321 (1983).
${ }^{45}$ H. See, M. Doi, and R. G. Larson, "The effect of steady flow fields on the isotropic-nematic phase transition of rigid rod-like polymers," J. Chem. Phys. 92, 792 (1990).
${ }^{46}$ J. H. Lee, M. G. Forest, and R. Zhou, "Alignment and rheo-oscillator criteria for sheared nematic polymer films in the monolayer limit," Dis-
crete Contin. Dyn. Syst., Ser. B 6, 339 (2006).
${ }^{47}$ T. Maruyama, G. G. Fuller, M. Grosso, and P. L. Maffetone, "The dynamics of two dimensional polymer nematics," J. Non-Newtonian Fluid Mech. 76, 233 (1998).
${ }^{48}$ K. S. Yim, G. G. Fuller, A. Datko, and C. Eisenbach, "Isotropic-nematic phase transitions of lyotropic, two-dimensional liquid crystalline polymer solutions," Macromolecules 34, 6972 (2001).
${ }^{49}$ D. Grecov and A. D. Rey, "Transient rheology of discotic mesophases," Rheol. Acta 42, 590 (2003).
${ }^{50}$ P. L. Maffettone and S. Crescitelli, "The rigid rod model for nematic polymers: Testing closure approximations with bifurcation analysis," J. Rheol. 38, 1559 (1994).
${ }^{51}$ P. L. Maffettone and S. Crescitelli, "Bifurcation analysis of a molecular model for nematic polymers in shear flow," J. Non-Newtonian Fluid Mech. 59, 73 (1995).
${ }^{52}$ G. Rienäcker and S. Hess, "Orientational dynamics of nematic liquid crystals under shear flow," Physica A 267, 294 (1999).
${ }^{53}$ Q. Wang, "Biaxial steady states and their stability in shear flows of liquid crystal polymers," J. Rheol. 41, 943 (1997).

