Shear-banding in a lyotropic lamellar phase

Part 2: Temporal fluctuations

Jean-Baptiste Salmon, Sébastien Manneville, and Annie Colin
Centre de Recherche Paul Pascal, Avenue Schweitzer, 33600 PESSAC, FRANCE
(Dated: October 31, 2018)

We analyze the temporal fluctuations of the flow field associated to a shear-induced transition in a lyotropic lamellar phase: the layering transition of the onion texture. In the first part of this work [Salmon et al., submitted to Phys. Rev. E], we have evidenced banded flows at the onset of this shear-induced transition which are well accounted for by the classical picture of shear-banding. In the present paper, we focus on the temporal fluctuations of the flow field recorded in the coexistence domain. These striking dynamics are very slow (100–1000 s) and cannot be due to external mechanical noise. Using velocimetry coupled to structural measurements, we show that such fluctuations are due to a motion of the interface separating the two differently sheared bands. Such a motion seems to be governed by the fluctuations of $\sigma^\star$, the local stress at the interface between the two bands. Our results thus provide more evidence for the relevance of the classical mechanical approach of shear-banding even if the mechanism leading to the fluctuations of $\sigma^\star$ remains unclear.

I. INTRODUCTION

Emulsions, shampoos, and paints are example of everyday life complex fluids. A mesoscopic scale located between the molecular size and the sample size is one of the most important features of these complex materials. In the case of an oil-in-water emulsion for instance, this length scale is the diameter of the oil droplets ranging from 100 nm to a few microns. During the last decade, many experimental and theoretical works have been devoted to the understanding of the effect of shear on these fluids [1, 2]. A robust fact has emerged: the flow modifies the structure of the fluid, which in turn changes the flow field.

In some cases, this strong coupling between flow and structure may induce the nucleation of a new phase or even create new structures or textures that do not exist at rest. In wormlike micelles for instance, shear may induce a nematic phase [3, 4], whereas in membrane phases, shear creates new textural organizations [5, 6]. Such Shear-Induced Structures (SIS) usually appear above a critical shear rate $\dot{\gamma}$ or shear stress $\sigma$. For some range of controlled $\dot{\gamma}$ (or $\sigma$), the SIS coexists with the unmodified structure, and then progressively fills the flow as $\dot{\gamma}$ (or $\sigma$) is further increased. Usually, the SIS flows very differently from the initial structure and large variations of the effective viscosity $\eta=\sigma/\dot{\gamma}$ are recorded near the shear-induced transition. Since the two structures have different viscosities, inhomogeneous flows or banded flows, are expected. Such an experimental picture is referred to in the literature as shear-banding [11, 12]. Shear-banding scenarii have been reported in a wide class of materials using both structural and rheological measurements [13, 14, 15, 16, 17, 18]. Moreover, a few experiments based on local velocimetry have reported the existence of such inhomogeneous flows where bands flow at different shear rates [12, 13, 14, 15, 16, 17, 18, 19]. These experiments mainly concern glassy materials and wormlike micellar systems.

In the first part of this work [20], we have started an exhaustive study of a shear-induced transition in a lyotropic lamellar phase composed of Sodium Dodecyl Sulphate (6.5% wt.) and Octanol (7.8% wt.) in Brine at 20 g.L$^{-1}$. Since the pioneering work of Roux and coworkers [21, 22], this system has been widely studied [21, 22, 23]. For $\dot{\gamma} \gtrsim 1$ s$^{-1}$, the membranes are wrapped into multiamellar vesicles called onions, that form a monodisperse disordered close-compact assembly. The characteristic size of the onions is a few microns. At higher shear rates ($\dot{\gamma} \gtrsim 15$ s$^{-1}$), onions get a long range hexagonal order and are organized on layers sliding onto each other. This shear-induced ordering transition is referred to in the literature as the layering transition [24]. Using local velocimetry, coupled to structural and rheological measurements [21], we have shown that above a critical shear rate, a highly sheared band, most probably composed of the ordered texture, is nucleated at the rotor of the Couette cell. This band coexists with the viscous disordered state thus leading to a banded flow. When $\dot{\gamma}$ is further increased, the highly sheared band grows up to finally invade the whole gap of the Couette cell. The classical picture of shear-banding thus holds for the layering transition.

However, one point remains very striking. Indeed, the flow field presents large temporal fluctuations of the local velocity in the coexistence domain. Such fluctuations reach up to 20% at the interface separating the two differently sheared bands. This phenomenon seems quite general in shear-banded flows since such temporal fluctuations have also been detected in some wormlike micellar systems using local velocimetry [10, 17]. On a more general ground, it has been shown recently that some com-

PACS numbers: 83.10.Tv, 47.50.+d, 83.85.Ei
plex fluids may also exhibit some dynamics when submitted to a shear flow in the vicinity of shear-induced transitions \cite{3,14,22,26,27}. In particular, in our lamellar phase, the effective viscosity \( \eta = \sigma/\dot{\gamma} \) presents aperiodic or relaxational oscillations under controlled \( \sigma \), suggesting deterministic behaviors in a very narrow range of parameters \cite{25,24}. For all these dynamical behaviors, referred to as *rheochaos*, it seems that spatial degrees of freedom play a relevant role \cite{14,25,24}. On the experimental point of view, it is important to determine whether the spatial organization of the flow field near such transitions is responsible for the observed dynamics. Such experimental data may then be very helpful to unveil the most relevant ingredients of the existing theoretical models \cite{31,32}.

In this second part of the article, we present a detailed study of the temporal fluctuations in our system. In the next section, after recalling briefly the experimental setup, we analyze the temporal fluctuations of the global measurements and of some time series of the local velocity. In Sec. \textbf{III}, we show that these temporal fluctuations are due to a motion of the interface separating the bands inside the gap. We then come back in Sec. \textbf{IV} to the mechanical approach used to describe the shear-banding scenario in Ref. \cite{20}. Assuming that the stress \( \sigma^* \), at which the interface between the disordered and the ordered textures is stable, fluctuates in time, we are able to describe the dynamics of the flow field recorded under controlled shear rate but also under controlled shear stress.

\section{II. GLOBAL AND LOCAL FLUCTUATIONS IN THE COEXISTENCE DOMAIN}

\subsection{A. Experimental setup}

The experimental setup has been fully described in the first part of this work \cite{20}. It allows us to perform rheological experiments, local velocity measurements and to determine the structure of the fluid at the same time. Figure 1 sketches the main features of this setup. A controlled-stress rheometer imposes a constant torque \( \Gamma \) on the axis of a Couette cell and records its rotation speed \( \Omega \) in real time. From these two *global* quantities, the rheometer indicates the following stress \( \sigma \) and shear rate \( \dot{\gamma} \):

\begin{align}
\sigma &= \frac{R_i^2 + R_o^2}{4\pi H R_i^2 R_o^2} \Gamma, \\
\dot{\gamma} &= \frac{R_i^2 + R_o^2}{R_o^2 - R_i^2} \Omega,
\end{align}

where \( H, R_i, \) and \( R_o \) are the geometrical characteristics of the Couette cell (see Fig. \textbf{1}). The values \( \langle \sigma, \dot{\gamma} \rangle \) given by the rheometer represent the spatial averages of the *local* stress \( \sigma(r) \) and of the *local* shear rate \( \dot{\gamma}(r) \) over the gap of the Couette cell, in the case of a Newtonian fluid \( \eta \) is the radial position in the flow). Temperature is controlled using a water circulation around the Couette cell. Since the system under study is very sensitive to the exact fractions of its components, great care has been taken to minimize evaporation using a thermostated plate on top of the cell.

A laser beam crosses the transparent Couette cell along the velocity gradient direction \( \nabla v \). The corresponding diffraction patterns are collected by a Charge-Coupled Device camera on a screen in the \((q_x, q_z)\) plane. Dynamic Light Scattering (DLS) experiments in the heterodyne geometry are performed to measure the local velocity. Such a method along with its spatial and temporal resolutions, has been described at length in Ref. \cite{33}. The light scattered by a small volume of the sample (denoted by \( \bullet \) in Fig. \textbf{1}) is collected using a single mode fiber. The scattered electric field is then mixed with the incident beam using an optical device coupling the two fibers. The Doppler frequency shift associated to the flow field inside the scattering volume is computed from the correlation function measured using an electronic correlator connected to a photomultiplier tube (see Fig. \textbf{1}). From the Doppler frequency shift, the mean value of the velocity inside the scattering volume is then obtained using a careful calibration of the setup \cite{33}. The characteristic size of the scattering volume is about 50 \( \mu m \).
3–5 s. A full velocity profile is then obtained in about 2–3 min.

B. Temporal fluctuations of the global rheology

The experimental procedure used to prepare the onion texture is the following. The temperature is set at $T = 30^\circ$C. A constant shear rate $\dot{\gamma} = 5 \text{ s}^{-1}$ is applied for $7200 \text{ s}$. This first step allows us to start the experiment with a well-defined stationary state of disordered onions. The shear rate is then increased from $\dot{\gamma} = 5$ to $20 \text{ s}^{-1}$ by applying different steps lasting $5400 \text{ s}$. The increment between two different steps is $5 \text{ s}^{-1}$. Finally, a constant shear rate of $\dot{\gamma} = 22.5 \text{ s}^{-1}$ is applied. Figure 2 presents the evolution of the controlled $\dot{\gamma}$ and that of the measured $\sigma$ during this last step. Let us recall that our rheometer imposes a torque $\Gamma$. A constant rotation speed $\Omega$ and thus a constant shear rate $\dot{\gamma}$ is obtained by a computer-controlled feedback loop on the applied $\Gamma$. As shown in Fig. 2(a), this procedure is very successful since the temporal fluctuations of the controlled $\dot{\gamma}$ are less than 0.05%. However, some variations greater than 0.1 s$^{-1}$ are sometimes recorded but they are very rare and probably induced by exterior mechanical noise. The measured stress $\sigma(t)$ is almost stationary: the temporal fluctuations of $\sigma(t)$ are about 0.4% [see Fig. 2(b)]. The non-equality between the fluctuations of the controlled $\dot{\gamma}$ and the measured $\sigma(t)$ raises important questions: such a difference might result from bad filling conditions of the Couette cell or may hide more fundamental points in the rheological analysis.

To proceed further in this analysis, we plot the probability density function of the time series $\sigma(t)$ displayed in Fig. 2(b). The continuous line is a Gaussian distribution with mean $16.21 \text{ Pa}$ and with standard deviation $0.065 \text{ Pa}$. (b) Power spectrum of the time series $\sigma(t)$ displayed in Fig. 2(b).

FIG. 2: (a) Time series of the controlled shear rate $\dot{\gamma}(t)$. (b) Corresponding temporal response of the measured shear stress $\sigma(t)$.

Let us now turn to the local velocimetry measurements. Figure 4(a) presents various velocity profiles obtained under the same applied shear rate of $\dot{\gamma} = 22.5 \text{ s}^{-1}$. We recall that it takes about 3 min to record a full velocity profile. These profiles clearly differ from each other and large variations of the velocity are observed. These variations are meaningful and greatly exceed the resolution of our experimental setup. If we decide, as in the first part of this paper [20], to average these measurements and to calculate their standard deviations, we may define errorbars on the time-averaged velocity profile that correspond to the amplitudes of the temporal fluctuations.

The time-averaged profile reveals three major issues [see Fig. 4(b)]. (i) Sliding occurs: the velocity of the fluid does not vanish at the stator and does not reach its expected value at the rotor. (ii) Moreover, the sliding velocities $v_{si}$, defined as the difference between the fluid velocity near wall number $i$ and the velocity of wall...
number \( i \), are almost constant. The lubricating layers responsible for the observed sliding, do not present any dynamics at the precision of our setup. (iii) Two bands with different shear rates coexist in the gap of the Couette cell. The highly sheared band is located near the rotor and is about 0.5 mm thick.

Finally, as shown in the inset of Fig. 4(b), the amplitude of the relative temporal fluctuations \( \frac{\delta v}{v} \) varies with the position in the gap and seems to reach a maximum at \( x \approx 0.5 \) mm where lies the interface between the two differently sheared bands. Note that \( \frac{\delta v}{v} \) seems to diverge at \( x \to 0 \): this is due to the small values of the velocities near the stator and to the intrinsic uncertainty of our setup (\( \approx 5\% \)).

### 2. Fluctuating time series \( v(x,t) \)

The variations recorded above refer to the amplitude of the temporal fluctuations. To analyze them in more details, we record the evolution of the local velocity at a given position in the flow field, by measuring successively several correlation functions accumulated over 3 s. We thus obtain the evolution of the velocity with a temporal sample rate of 3 s. This sample rate is short enough to follow the slow dynamics described above and long enough to measure precisely the velocity \( v(x,t) \). Figure 5(a) presents the measurement performed in the middle of the gap at \( x_2 = 0.46 \) mm. The signal was recorded during 11500 s. The inset clearly reveals fluctuations of the local velocity with an amplitude of about 4 mm.s\(^{-1}\) and characteristic times ranging from 100 to 500 s. The power spectrum displayed in Fig. 5(b), points out the same feature: it presents a bump centered between 0.002 and 0.01 Hz. These measurements show that the time series of the local velocity presents large fluctuations that involve long time scales. In order to check whether these behaviors occur homogeneously in the entire gap of the Couette cell, we have repeated the same measurement for three positions in the gap \( x_1 = 0.07, x_2 = 0.46, \) and \( x_3 = 0.70 \) mm.

Figure 6 reports these measurements and presents the corresponding probability density functions \( p(v) \) of the local velocities.

![FIG. 4: (a) 6 velocity profiles obtained simultaneously to the rheological data displayed in Fig. 2 (\( \eta = 22.5 \) s\(^{-1}\) and \( T = 30^\circ C \)), (b) Corresponding time-averaged velocity profile. The errorbars are the standard deviation inferred from the 6 profiles displayed in (a). Inset: temporal fluctuations (i.e. standard deviation) of the local velocity vs. the position \( x \). The horizontal lines indicate the imposed rotor velocity \( v_0 \). The vertical lines indicate the positions \( x_1 = 0.07, x_2 = 0.46, \) and \( x_3 = 0.70 \) mm where the time series \( v(x,t) \) displayed in Figs. 5 and 6 are measured.](image1)

![FIG. 5: (a) Time series \( v(x_2,t) \) measured at \( x_2 = 0.46 \) mm, simultaneously to the data presented in Figs. 2 and 4. Inset: blow-up of the dynamics. (b) Corresponding power spectrum.](image2)

![FIG. 6: Time series \( v(x_i,t) \) and corresponding probability density functions obtained simultaneously to the data displayed in Figs. 2 and 4; \( x_1 = 0.07, x_2 = 0.46, \) and \( x_3 = 0.70 \) mm. The continuous lines are the Gaussian distributions with the means and the standard deviations of the time series \( v(x_i,t) \).](image3)
tainties on the measurement. It is thus rather difficult to analyze precisely this data set. However, temporal fluctuations on about 100 s are clearly present. A Gaussian distribution centered around 15.5 mm.s\(^{-1}\) and characterized by a standard deviation of 0.6 mm.s\(^{-1}\) nicely fits the experimental distribution of the local velocity.

(ii) In the middle of the gap at \(x_2 = 0.46\) mm, the fluctuations are much larger and reach 12%. Moreover, the fluctuations are not symmetric: more events occur at high velocity than at low velocity. The distribution is thus not Gaussian.

(iii) Near the stator at \(x_3 = 0.70\) mm, this behavior is even more pronounced: the velocity seems to fluctuate between two states. Most of the time, the velocity remains equal to about 3.5 mm.s\(^{-1}\), but sometimes the velocity increases rapidly to 5–6 mm.s\(^{-1}\). The most probable velocity is now clearly smaller than the mean velocity. The probability density function is highly disymmetric and shows a long tail of rare events at high velocities.

These measurements show that (i) long time scales ranging between 100 and 1000 s are involved in the dynamics of the local velocities; (ii) the amplitude of the fluctuations varies in the gap and reaches a maximum at the interface between the bands; (iii) the probability density functions are Gaussian near the rotor and become highly disymmetric near the interface in the weakly sheared band.

### III. EVIDENCE FOR A MOTION OF THE INTERFACE BETWEEN THE TWO BANDS

#### A. Local velocimetry measurements

To proceed deeper into the analysis, a dynamical picture of the flow in the entire gap would be very helpful. Unfortunately, our experimental setup does not provide an excellent temporal resolution: about 2–3 min are necessary to obtain a full velocity profile by moving the rheometer along the velocity gradient direction \(\nabla v\). This time is comparable to the time scales involved in the temporal fluctuations. Thus, the velocity profile is not frozen during the measurement.

In the first part of this article, we have analyzed the time-averaged velocity profiles obtained using a careful protocol to study the layering transition (see Ref. 21). Keeping in mind the time needed to obtain a full profile (≈3 min), we now focus on individual profiles, i.e. without averaging.

Figure 7(a) presents two individual profiles obtained at \(\dot{\gamma} = 15\) s\(^{-1}\) and the time-averaged profile calculated over 13 successive measurements is shown in Fig. 7(b). For this applied shear rate, the time-averaged velocity profile is almost linear. Sliding is observed both at the rotor and at the stator. The inset of Fig. 7(b) presents the relative fluctuations \(\delta v / v\) vs. the position \(x\). \(\delta v / v\) reaches a maximum of 10% at \(x \approx 200\) \(\mu m\). Moreover, among the 13 individual profiles, we can distinguish two kinds of measurements: some are linear while others present bands supporting different local shear rates [see Fig. 7(a)]. In this last case, the thickness of the highly sheared band is roughly equal to 200 \(\mu m\). It seems that these two families of profiles correspond to frozen pictures of nearly stationary flow fields.

At this stage, an interpretation of the temporal fluctuations begins to emerge. These fluctuations could be correlated to the motion of the interface between the two bands. If the position of the interface moves, then different kinds of velocity profiles will be measured at this
low shear rate: linear ones when the highly sheared band is absent, and shear-banded ones when the two textures coexist. Fluctuations of the position of the interface separating the two bands will also lead to a spatial localization of the fluctuations of the local velocity. This is indeed the case as can be seen in the inset of Fig. 4(b).

In order to further check this idea, we performed the same experiments at higher shear rates. Since the temporal fluctuations are more important, frozen pictures of the flow field are more difficult to capture. However, some profiles reveal the same results as for \( \dot{\gamma} = 15 \text{ s}^{-1} \). Figure 7(c) shows two different velocity profiles averaged over some well-chosen profiles among 13 successive measurements at \( \dot{\gamma} = 22.5 \text{ s}^{-1} \). In both cases, two differently sheared bands coexist in the gap. However, the position of the interface varies a lot. For the first group of velocity profiles, the interface lies at \( x \approx 0.25 \text{ mm} \) and it has moved to \( x \approx 0.55 \text{ mm} \) for the second group. In both cases, the velocity of the fluid near the rotor is nearly constant. This implies that the shear rate in the highly sheared band has changed from 35 to 23 \( \text{s}^{-1} \). Note that the shear rate in the weakly sheared band does not seem to vary. On average, the interface between the two textures lies at \( x \approx 0.40 \text{ mm} \) and the amplitude of the fluctuations is maximal at this position [see Fig. 7(d)].

The same conclusions may be drawn by looking at two groups of profiles obtained at \( \dot{\gamma} = 37 \text{ s}^{-1} \) and displayed in Fig. 4(c). In this case, the highly sheared band has invaded the gap on a time-averaged point of view since the velocity profile averaged over 20 successive measurements is almost linear [see Fig. 7(f)].

### B. Temporal fluctuations of the structure

The previous measurements clearly show a displacement of the interface between the two bands in the gap. As the two bands present different microscopic structures, we believe that this motion may also be detected on the diffraction patterns. The patterns associated with the disordered texture are homogeneous rings, whereas six peaks modulate the ring at the onset of the layering transition. These peaks are associated with the hexagonal long range order of the onions on planes oriented along the flow [21]. Let us recall that the diffraction patterns correspond to a measure of the structure integrated along the velocity gradient direction \( \nabla v \). During the layering transition, the diffraction pattern evolves continuously from an homogeneous ring to six well-defined peaks [21].

A series of images were taken every 10 s at \( \dot{\gamma} = 22.5 \text{ s}^{-1} \) and \( T = 30^\circ\text{C} \) in the coexistence domain. To enhance the signal-to-noise ratio, each image corresponds to an average over five successive pictures separated by 40 ms. From the intensity of the peaks relative to that of the ring, we can compute a degree of organization of the sample [25]. Such a parameter \( \phi \) is defined as the difference between the maximal intensity and the minimal intensity measured on the ring. \( \phi \) is significantly larger than 0 when peaks modulate the ring, whereas \( \phi \approx 0 \) when rings are uniform. Figure 8 presents the evolution of \( \phi \) as a function of time for \( \dot{\gamma} = 22.5 \text{ s}^{-1} \). First, note that \( \phi(t) \) is always positive since peaks are always present on the ring: the intensity modulation of the ring is due to the presence of a highly sheared band most probably corresponding to the layered state of onions. Indeed, as shown in Fig. 4(b), the width \( \delta \) of the ordered band is about 0.5 mm at the considered \( \dot{\gamma} \).

However, the time series \( \phi(t) \) fluctuates with characteristic periods of 100–500 s. The amplitude of these fluctuations is rather small but still meaningful since some well-chosen patterns [see Figs. 8(a)–(d)] clearly show the variation of the contrast of the peaks on the ring. These structural fluctuations support the idea that the proportion of ordered onions relative to disordered onions constantly changes, which is consistent with a motion of the interface between the bands. Moreover, it is reasonable to assume that such an ordering/disordering process, leading to the displacement of the highly sheared band, involves time scales longer than the characteristic times of hydrodynamics. Such long time scales correspond to those observed in the dynamics of \( \sigma(t) \) and of \( v(x_i, t) \).

### IV. Analysis using a simple mechanical approach

#### A. Constitutive equations of the mechanical approach

In the first part of this article [20], we have shown that the time-averaged velocity profiles may be well fitted by a simple mechanical approach. Using the rheological data \( \sigma \) vs. \( \dot{\gamma} \) together with the assumption that the interface between the bands always lies at the same local stress \( \sigma^* \), we were able to reproduce the velocity data. Let us recall briefly such a mechanical approach. The flow curve obtained from the rheological data where the contribu-
tion due to wall-slip are removed is sketched in Fig. 9. The shear rate plotted on this flow curve is the effective shear rate in the bulk material once slip effects are removed. The homogeneous branches corresponding to the disordered and layered states, yield the local rheological behaviors \( \sigma = f_i(\dot{\gamma}) \), where \( i = 1 \) (\( i = 2 \) resp.) denotes the disordered (layered resp.) texture.

Due to the the curvature of the Couette cell, the local stress \( \sigma(r) \) varies in the gap as:

\[
\sigma(r) = \frac{\Gamma}{2\pi H r^2} = \sigma^* \frac{R_1^2}{r^2},
\]

where \( r \) is the radial position in the gap, \( \Gamma \) is the imposed torque, and \( \sigma_i = \Gamma / (2\pi H R_1^2) \) is the stress at the rotor. In our mechanical picture, the interface lies at a given stress \( \sigma^* \). The width \( \delta \) of the highly sheared band is thus given by:

\[
\delta = R_1 \left( \sqrt{\frac{\sigma_i}{\sigma^*}} - 1 \right) .
\]

It is then straightforward to compute the theoretical profiles from an arbitrary value of \( \Gamma \). If \( \sigma(r) < \sigma^* \) (\( \sigma(r) > \sigma^* \) resp.) everywhere in the gap, the flow is homogeneous and composed of the disordered (layered resp.) state of onions. In that case, the velocity profiles are given by the following integration of the rheological behavior:

\[
v(r) = \frac{v_2}{R_2} + \int_{R_2}^{r} \frac{\dot{\gamma}(u)}{u} \, du ,
\]

where \( \dot{\gamma}(r) \) is found by solving:

\[
\sigma(r) = \frac{\Gamma}{2\pi H r^2} = f_i(\dot{\gamma}(r)) ,
\]

and \( i \) denotes the considered branch. When there exists one particular position in the gap where \( \sigma(r) = \sigma^* \), the flow displays two bands supporting different shear rates.

To calculate the resulting velocity profile, one should separate the previous integration between \( [R_i; R_i + \delta] \) and \( [R_i + \delta; R_i] \) (see Ref. 20 for details).

From this set of equations, one can compute a theoretical profile \( v(x) \) from an arbitrary value of \( \Gamma \). It is then straightforward to infer a global shear rate \( \dot{\gamma} \) indicated by the rheometer. Such a procedure has been applied to our data in Ref. 20; since Eqs. 9 and 10 lead to satisfactory fits of both the flow curve and of the velocity profiles, the assumption that the interface lies at a given stress \( \sigma^* \), holds for the layering transition.

**B. Temporal fluctuations of \( \sigma^* \)**

One point is particularly important: for a given value of torque \( \Gamma \), there exists only one profile \( v(x) \). Thus, under controlled shear rate, these equations predict that the stress \( \sigma \) indicated by the rheometer is well-defined and that the position of the interface is fixed. Moreover, one can easily check that if \( \sigma^* \) is fixed, very small variations of the applied shear rate cannot induce large fluctuations of the measured \( \sigma \). Thus, the experimental data displayed in Fig. 2 cannot be understood simply by these equations.

However, since Eqs. 9 and 10 describe nicely the time-averaged velocity profiles, we believe that if a crucial parameter of the previous equations possesses its own hidden dynamics, the mechanical approach may still help us understand the observed fluctuations. Since the interface between the two bands fluctuates in the gap and since its position is fixed by \( \sigma^* \) through Eq. 4, we can reasonably assume that some fluctuations of \( \sigma^* \) could explain our data. Figure 10 presents a sketch of this scenario. Consider an initial state A in the shear-banding region. The flow profile presents two differently sheared bands [see Fig. 10(b)]. Let us now assume that \( \sigma^* \) slightly increases. The position of the stress plateau thus changes, as well as the position in the gap that corresponds to \( \sigma^* \). The interface does not lie on a stable position anymore. Under controlled shear rate, the final stable state will lie at point B on the flow curve, and the global stress will vary from \( \sigma \) to \( \sigma^* \). This simple picture helps us understand why fluctuations of \( \sigma \) could be observed without any significant fluctuations of \( \dot{\gamma} \) and with a motion of the interface in the gap.

To proceed further into the analysis, one should compare such a scenario to the experimental data. Since the global stress \( \sigma \) fluctuates as \( \sigma^* \), we assume that the fluctuations of \( \sigma^* \) have the same shape as those of \( \sigma(t) \). Figure 11(a) displays a time series \( \sigma^*(t) \) inferred from a typical time series \( \sigma(t) \) obtained under controlled shear rate at \( \dot{\gamma} = 22.5 \, \text{s}^{-1} \). The mean value of \( \sigma^* \) is set at 16.15 Pa, the value leading to the best fits of the experimental data in Ref. 20.

We have presented in Fig. 11(b), the resulting theoretical velocity profiles computed from Eqs. 3 and 4 with \( \sigma^*(t) \) displayed in Fig. 11(a). Figure 11(b) also presents
the experimental time-averaged velocity profile obtained at $\dot{\gamma} = 22.5$ s$^{-1}$ for comparison. Figure 12 displays a time-space plot of the same theoretical velocity profiles. The time series $\delta(t)$ of the position of the interface computed from Eq. 11 has also been added on this time-space plot.

Our crude approach succeeds to describe many experimental points. (i) The fluctuations of $\sigma^*$ induce a large fluctuating motion of the interface (see Fig. 12). (ii) The amplitude of the fluctuations of the flow field is thus localized [see inset of Fig. 11(b)]: they reach a maximum of about 10% at the position of the interface. (iii) A quantitative agreement may be found between the theoretical fluctuations and the measured data within this crude approach: fluctuations of $\sigma^*$ by 2% induce fluctuations of the measured $\sigma$ by 1%, and fluctuations of the local velocities reaching 10% at the interface. Let us outline that fluctuations of the others parameters of the mechanical equations such as the viscosities of the two different textures will not be able to reproduce these dynamics.

**C. Validation under controlled shear stress**

To fully validate our mechanical approach, one should perform experiments under controlled shear stress, and show that the fluctuations of $\sigma^*$ may also induce a fluctuating motion of the interface and some variations of the measured $\dot{\gamma}(t)$.

However, for temperatures $T \geq T_c = 27^\circ$C and under controlled stress, the lyotropic lamellar phase under study displays well-defined oscillating behaviors of $\dot{\gamma}(t)$ with a period of about 500 s and an amplitude reaching about 20 s$^{-1}$ in the vicinity of the transition. Unfortunately, our setup is not suitable to follow such dynamics of the flow field, since the involved periods are of the order of minutes and the amplitudes are large. Using DLS velocimetry, it is thus rather difficult to determine whether...
the shear rate oscillations observed for \( T \geq T_c \) are due to an oscillating motion of the interface. We thus decide to perform experiments for temperatures \( T < T_c \), where the shear rate only displays fluctuating behaviors rather than sustained oscillations. Indeed, in this region of parameters, the amplitude of the dynamics is rather small and the dynamics is slow enough to be captured by our setup.

Figure 13(a) presents two well chosen groups of velocity profiles measured in the coexistence domain under controlled \( \sigma \) and at \( T = 26^\circ C \). The velocity profile averaged over 14 successive measurements is shown in Fig. 13(b). These data clearly show a very large motion of the interface in the gap (from \( x = 0.25 \) to \( 0.50 \) mm). In this experiment under controlled stress, the velocity of the fluid near the rotor and thus the measured \( \dot{\gamma} \) vary a lot. Following our crude mechanical approach, the situation is the one sketched in Fig. 13. The flow field is initially stable at point A. A small variation of \( \sigma^\star \) changes the position of the plateau on this flow curve. Under controlled stress, the final stable configuration lies at point C: the shear rate has varied from \( \dot{\gamma} \) to \( \dot{\gamma}^\prime \) and the interface has moved in the gap.

\[
\begin{align*}
\text{FIG. 13: (a) Two averaged groups of velocity profiles obtained at } & \sigma = 12.5 \text{ Pa and } T = 26^\circ C. \text{ The vertical dashed lines lie at } x = 0.25 \text{ and } x = 0.50 \text{ mm. The dotted lines help to guide the eye in the highly sheared band. (b) Velocity profile averaged over 14 consecutive measurements at the same stress. The errorbars are the standard deviations of these estimates. Inset: temporal fluctuation of the local velocity inferred from the standard deviation of the different measurements vs. } x.
\end{align*}
\]

V. CONCLUSIONS, DISCUSSIONS, AND PERSPECTIVES

The present work has provided an exhaustive study of a shear-induced transition in a specific complex fluid: the onion texture of a lyotropic lamellar phase. Under controlled shear rate and above some critical value, onions get a long range hexagonal order into planes sliding onto each other. In the vicinity of this ordering transition, one observes a spatial organization of the flow field: a band corresponding to the ordered texture is nucleated near the rotor and coexists with the disordered texture.

As pointed out by our velocity measurements, these two structures flow with different shear rates and the highly sheared band invades the gap as \( \dot{\gamma} \) is increased. The proportion of the two differently sheared bands in the flow field is governed by the controlled \( \dot{\gamma} \) and by \( \sigma^\star \), the local stress at the interface.

Our experiments have also revealed the presence of large fluctuations of the flow field localized at the interface between the two bands. Using DLS velocimetry coupled to rheology and structural measurements, we have shown that these temporal fluctuations are due to a fluctuating motion of the interface in the gap. The characteristic times of these puzzling dynamics range between 100 and 1000 s and do not correspond to external noise. Using the classical mechanical picture of shear-banding, we have demonstrated that the fluctuations of \( \sigma^\star \) may explain the observed dynamics. Indeed, under controlled shear rate, small variations of \( \sigma^\star \) may induce small variations of \( \sigma \), but also large fluctuations of the interface between the bands and thus localized fluctuations of the flow field. Under controlled stress, we have also shown that for \( T < T_c = 27^\circ C \), fluctuations of \( \sigma^\star \) may describe correctly the observed displacement of the interface.

This crude approach describes very nicely the dynamics of the fluctuations but raises a crucial question: why is \( \sigma^\star \) a fluctuating parameter? Clearly, this parameter does not vary because of some external mechanical vibrations. Following Lu’s theoretical approach [34], this parameter depends on the mathematical form of the model which describes the transition and its origin is due to the presence of gradient terms. In the phenomenological model of Ref. [35] for instance, \( \sigma^\star \) depends on the rate of change between the two structures of the differently sheared bands. Our data suggest that a dynamical equation describing the evolution of \( \sigma^\star \) is missing. Such an equation is difficult to derive since it must describe precisely the microscopic structure of the system. Other theoretical approaches such as those of Ref. [36], based on a dynamical equation for the motion of the interface, may also fully reproduce the observed fluctuations if other dynamical equations concerning the structure are added. Understanding the origin of the fluctuations of \( \sigma^\star \) from a microscopic point of view remains a major challenge in the study of shear-banding.

Finally, another point is left unclear: above the critical temperature \( T_c \) and under controlled stress, why do the dynamics seem to become deterministic or even seem to display chaotic behaviors? To answer such a crucial question, it is necessary to increase the temporal resolution of our setup. We plan to use a spectrum analyzer to reduce the acquisition time of the velocity measurements or to turn to other velocimetry methods such as ultrasonic techniques [37]. These experimental improvements will certainly help to get a more precise spatio-temporal picture of rheochaos in complex fluids.
Acknowledgments

The authors are deeply grateful to D. Roux, L. Bécu, and C. Gay for many discussions and comments on this work. The authors would like to thank the Cellule Instrumentation at CRPP for the realization of the heterodyne DLS setup. We also wish to thank A. Ajdari, A. Aradian, C. Barentin, L. Bocquet, L. Chevillard, F. Molino, G. Picard, L. White-Benon and C. Ybert for many helpful conversations.

[1] R. G. Larson, The Structure and Rheology of Complex Fluids (Oxford University Press, 1999).
[2] M. E. Cates and M. R. Evans, eds., Soft and Fragile Matter: Non Equilibrium Dynamics Metastability and Flow (Institute of Physics Publishing (Bristol), 2000).
[3] J.-F. Berret, D. C. Roux, and G. Porte, J. Phys. II 4, 1261 (1994).
[4] V. Schmitt, F. Lequeux, A. Pousse, and D. Roux, Langmuir 10, 955 (1994).
[5] O. Diat and D. Roux, J. Phys. II (France) 3, 9 (1993).
[6] D. Roux, F. Nallet, and O. Diat, Europhys. Lett. 24, 53 (1993).
[7] E. Cappelaere, J.-F. Berret, J.-P. Decruppe, R. Cressel y, and P. Lindner, Phys. Rev. E 56, 1869 (1997).
[8] Y. T. Hu, P. Boltenhagen, and D. J. Pine, J. Rheol. 42, 1185 (1998).
[9] E. Eiser, F. Molino, G. Porte, and O. Diat, Phys. Rev. E 61, 6759 (2000).
[10] P. Panizza, P. Archambault, and D. Roux, J. Phys. II 5, 303 (1995).
[11] L. Ramos, F. Molino, and G. Porte, Langmuir 16, 5846 (2000).
[12] G. Debrégeas, H. Tabuteau, and J.-M. di Meglio, Phys. Rev. Lett. 87, 178305 (2001).
[13] P. Coussot, J. S. Raynaud, F. Bertrand, P. Moucheront, J. P. Guilbaud, H. T. Huynh, S. Jarny, and D. Lesueur, Phys. Rev. Lett. 88, 218301 (2002).
[14] F. Pignon, A. Magnin, and J.-M. Piau, J. Rheol. 40, 573 (1996).
[15] R. W. Mair and P. T. Callaghan, Europhys. Lett. 36, 719 (1996).
[16] M. M. Britton and P. T. Callaghan, Phys. Rev. Lett. 78, 4930 (1997).
[17] E. Fischer and P. T. Callaghan, Phys. Rev. E 64, 011501 (2001).
[18] Y. T. Hu, P. Boltenhagen, E. Matthys, and D. J. Pine, J. Rheol. 42, 1209 (1998).
[19] J.-B. Salmon, A. Colin, S. Manneville, and F. Molino, Phys. Rev. Lett. 90, 228303 (2003).
[20] J.-B. Salmon, S. Manneville, and A. Colin (2003), submitted to Phys. Rev. E.
[21] O. Diat, D. Roux, and F. Nallet, Phys. Rev. E 51, 3296 (1995).
[22] P. Sierr and D. Roux, Phys. Rev. Lett. 78, 1496 (1997).
[23] J. Leng, F. Nallet, and D. Roux, Eur. Phys. J. E. 4, 77 (2001).
[24] B. J. Ackerson and P. N. Pusey, Phys. Rev. Lett. 61, 1033 (1988).
[25] R. Bandyopadhyay and A. K. Sood, Europhys. Lett. 56, 447 (2001).
[26] D. Lootens, H. Van Damme, and P. Hébraud, Phys. Rev. Lett. 90, 178301 (2003).
[27] E. K. Wheeler, P. Fischer, and G. G. Fuller, J. Non-Newtonian Fluid Mech. 75, 193 (1998).
[28] A.-S. Wunenburger, A. Colin, J. Leng, A. Arnéodo, and D. Roux, Phys. Rev. Lett. 86, 1374 (2001).
[29] J.-B. Salmon, A. Colin, and D. Roux, Phys. Rev. E 66, 031505 (2002).
[30] D. A. Head, A. Ajdari, and M. E. Cates, Phys. Rev. E 64, 061509 (2001).
[31] M. E. Cates, D. A. Head, and A. Ajdari, Phys. Rev. E 66, 025202 (2002).
[32] G. Picard, A. Ajdari, L. Bocquet, and F. Lequeux, Phys. Rev. E 66, 051501 (2002).
[33] J.-B. Salmon, S. Manneville, A. Colin, and B. Pouligny, Eur. Phys. J. AP 22, 143 (2003).
[34] C.-Y. D. Lu, P. D. Olmsted, and R. C. Ball, Phys. Rev. Lett. 84, 642 (2000).
[35] J. L. Goveas and P. D. Olmsted, Eur. Phys. J. E 6, 79 (2001).
[36] A. Ajdari, Phys. Rev. E 58, 6294 (1998).
[37] S. Manneville, L. Bécu, and A. Colin (2003), in preparation.