Phase-ordering kinetics in the Allen-Cahn (Model A) class: universal aspects elucidated by electrically-induced transition in liquid crystals

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The two-dimensional (2d) Ising model is the statistical physics textbook example for phase transitions and their kinetics. Quenched through the Curie point with Glauber rates, the late-time description of the ferromagnetic domain coarsening finds its place at the scalar sector of the Allen-Cahn (or Model A) class, which encompasses phase-ordering kinetics endowed with a nonconserved order parameter. Resisting exact results sought for theoreticians since Lifshitz’s first account around a half-century ago, the central quantities of 2d Model A – most scaling exponents and correlation functions – remain known up to the level of competing approximations that, nevertheless, urge for substantial and accurate experimental confrontation. Here, we report a comprehensive study on the coarsening of 2d twisted nematic liquid crystals (TNLC) whose kinetics is induced by a super-fast electrical switching from a spatiotemporally chaotic (disordered) state to a two-phase concurrent, equilibrium one. Tracking the dynamics via optical microscopy, we firstly show the sharp evidence of well-established Model A aspects, such as the dynamic exponent $z = 2$ and the dynamic scaling hypothesis, to then move forward and: i) confirm the Bray-Humayun theory for Porod’s regime describing intradomain length scales; ii) show that Gaussian-based models, namely the Ohta-Jasnow-Kawasaki and Mazenko theories, are good descriptors of two-point spatial correlators in interdomain length scales; iii) corroborate the aging hypothesis in Model A systems which includes the collapse of two-time correlators into a master curve whose format is, actually, iv) overall best accounted for by a solution of the local scaling invariance theory: the same solution that fits the 2d nonconserved Ising model (2dIM) correlator along with the Fisher-Huse conjecture. We also v) measure a local persistence exponent with sufficient precision to distinguish between the numerically estimated 2dIM value and that exactly derived from the diffusion equation, thus suggesting the former as the true value for the Model A class. Finally, we vi) observe a fractal morphology for persistence clusters and extract their universal dimension. Due to its accuracy and possibilities, this experimental setup may work as a prototype to address further universality issues in the realm of nonequilibrium systems.

I. INTRODUCTION

Since the seminal studies $[1,2]$ by Lifshitz, Slyozov and Wagner on the Ostwald ripening, and those $[3,4]$ own to Lifshitz, Allen and Cahn on curvature-driven domain growth, investigations of phase ordering continue very alive as a centre of statistical physics and interdisciplinary sciences for a myriad of natural, social, and artificial phenomena $[5–11]$. Over the past decade, we have witnessed solid developments not only in traditional subjects, but also in the discovery on the pivotal roles that phase separation and domain coarsening play in biology and artificial active matter. The traditional road includes the unveiling of novel coarsening mechanisms such as the mechanically-driven relaxation of network-forming morphologies $[12]$, the continued interest on spinodal decomposition of alloys and glasses $[13–14]$, evolving patterns of foams $[15–16]$, growing crystalline films $[17]$, not to mention the evergreen simulations of Ising ($e.g.$ $[18–22]$), Potts $[23–24]$ and voter $[25]$ kinetics. The second path is directed to a quantitative understanding of emergent behaviour in active systems, which range from the intra-cellular assembly kinetics and cellular functionalization – notably liquid-liquid phase separations involved in the formation of membraneless cell compartments – $[11,26,27]$, to the segregation $[28]$ and phase separation $[29]$ within uni-cellular (bacteria) communities, pattern formation in groups of complex multi-cellular organisms $[30,31]$, besides experiments $[32–34]$ and theoretical $[10]$ accounts for artificial active matter undergoing motility-induced phase separation. Amid such a multidisciplinary environment, the iconic and perhaps the simplest example of ordering remains being the (kinetic) Ising model $[35,36]$ because of its simple conceptualization, theoretical and numerical tractability, both of which are accompanied by a fundamental universality that often emerge from nontrivial, real dynamics ($e.g.$, $[28,37]$).

Fascination on phase-ordering kinetics then stems from its virtual ubiquitoussness and complexity that yet bears universal aspects $[5,11]$. Among these, generically one finds a dynamic scaling property that guarantees that the domain mosaic becomes statistically time-independent whenever lengths are measured in the unit of a single, asymptotically emergent, growing scale $l(t)$. Free to relax in the absence of quenched disorder, geometric frustration and external forces, domains display an algebraic growth $l(t) \sim t^{1/z}$ where $z$ is the universal dynamic expo-
dent. For kinetics where the system’s order parameter is not conserved, viz. the Model A dynamics in the classification by Hohenberg and Halperin [38], one has [39] \( z = 2 \) [3, 4, 10]. Although simulations and experiments extensively support this dynamic scaling, rigorous derivations have been limited to the one-dimensional nonconserved kinetic Ising model (1dIM) [41, 42] and to the nonconserved \( O(n \to \infty) \) model [43], where \( n \) is the number of the components of the order parameter. In the case of scalar \((n = 1)\), two-dimensional (2d) continuum Model A systems quenched from a fully disordered regime to an equivalent zero-temperature \((T = 0)\) state, dynamic scaling with \( z = 2 \) is obtained from an exact result that relies on the assumed convergence to the critical percolation fixed point [44]. Similar conclusion is reached for quenches performed from the critical state [44], for which the cluster size distribution is exactly known [45].

Theoretically suggested [46, 47], then observed in models [48] and real systems [49] around the end of the 70’s, time-independent forms of two-point spatial correlators remain hitherto exactly known only for specific cases [41–43, 50]. These cases do not include the scalar 2d Model A whose approximate forms were proposed along a remarkable deal of effort made around the 80’s [51, 52] and in the 90’s [53, 54], where the Ohta-Jasnow-Kawasaki (OJK) theory [51] and the theory of unstable growth (TUG) by Mazenko [52, 53] played the major roles [55]. Tested against discrete [55, 56] and continuum [51, 52] models, both OJK and TUG functional forms generically succeeded [57] in fitting to data. The OJK form was tested in an notable experiment [57] back to the 80’s, although agreement was mainly confined to the short intradomain scale ruled by the Porod’s regime [58].

On the Porod’s regime, Bray and Humayun [59] derived exact formulae for the short-distance limit of the spatial correlator (or the tail of the structure factor) – hereinafter referred to as the Bray-Humayun amplitude –, so that there is no need to rely on approximate theories in this regime. Moreover, because the formulae were derived under minimal assumptions on the existence of defects, the Bray-Humayun amplitude shall be valid regardless of the conservation laws of the order parameter and whether the dynamic scaling holds [59]. Despite so, to our knowledge, no experimental measurement has turned attention to this prediction.

The memory of systems in the coarsening stages is an additional aspect of interest because corresponding two-time and infinitely-many time statistics may also display scale invariance [60, 61, 62], including aging akin to that observed in glassy materials [63]. As revealed by Fisher and Huse [70], the description of two-time correlators requires an additional nonequilibrium exponent – defined in Sec. [14], Eq. [19] \( \lambda(n, d) \) – which is affected by the presence of long-range order in initial conditions [72, 73]. For the Model A class, exact results for \( \lambda \) are known for the 1dIM [42], the 1d noiseless time-dependent Ginzburg-Landau (TDGL) equation [74], including for the \( O(n) \) model in terms of the \( 1/n \) expansion [75]. In the scalar, 2d case, Fisher and Huse conjectured \( \lambda_{FH} = 5/4 \) based on the convergence of the dynamics to the critical percolation fixed point, which then supplements their heuristic arguments [70]. Such conjecture was initially supported by 2dIM [70, 72, 73] and 2dTDGL simulations [75], but recent studies based on finite-size scaling for the very same models suggest a value closer to the prediction by TUG [76, 77], \( \lambda_{TUG} \approx 1.29 \) [78]. In parallel, the OJK theory predicts a rather different value, \( \lambda_{OJK} = 1 \) [79], coinciding with the lower bound of the inequality \( \lambda \geq 2/d \) for nonconserved dynamics [70, 80]. The OJK value has not been observed so far but for 2dIM simulations with strongly long-range interactions [22]. Noting that deviations from the \( Z_2 \)-symmetric dynamics must be taken into account when analysing the two-time correlator – otherwise strongly biasing and artificially changing its corresponding asymptotic decay – it is important to elucidate on the experimental support to \( \lambda_{FH} \) in Ref. [81] in the light of dealing with systems that are not \( Z_2 \)-symmetric, as it happens in experiments.

More fundamentally, experimental support to the dynamic scaling according to aging hypothesis [69] is surprisingly elusive for the Model A class. As consequence, functional forms for the two-time correlator given by OJK [79], TUG [78], and the local scale invariance (LSI) theory by Henkel [82], all lack confrontation with real dynamics.

Another nontrivial facet of phase-ordering kinetics is encoded in their first-passage statistics [83], specifically in the local persistence probability \( Q(t, t_0) \) [67, 68, 84]. In the language of the Ising model, \( Q(t, t_0) \) is defined as the fraction of spins at time \( t \) which has never flipped since a previous time \( t_0 \). By construction, it involves correlation of infinitely-many points. In the thermodynamic limit \((L \to \infty)\), followed by \( t \to \infty \) with fixed \( t/t_0 \), \( Q \) often decays as \( Q(t, t_0) \sim (t/t_0)^{-\theta} \) for large \( t/t_0 \) with the persistence exponent \( \theta \), which is normally independent of \( z \) and \( \lambda \) [67, 68].

The associated body of knowledge was seeded in the numerical work by Derrida et al. [59] on the 1d \( q \)-state Potts model (1dPM), then grew and took shape thanks to 1d-5d IM simulations [85, 86], analytical approaches for the noiseless 1d TDGL equation [74], reaction-diffusion schemes [87, 88], exact solutions for 1dPM [89, 90], approximations for simple diffusion (DIF) with random initial conditions (RIC) [91, 92], besides being nurtured by experiments [93, 94]. Solvability of \( Q \) in Model A systems is constrained to 1d cases [74, 95, 96]. In the scalar sector of 2d Model A, early numerics predicted \( \theta \approx 0.22 \) [89, 89] while a recent 2dIM simulation suggests \( \theta = 0.199(2) \) [93, 94]. This value is rather discernible from the exact \( \theta_{DIF} = 3/16 = 0.1875 \) value derived from the 2d DIF-RIC [96, 97] and OJK theory [92], which is also faced as a candidate value for 2d Model A. The early experiment by Yurke, Pergelis, Majumdar and Sire reported \( \theta = 0.19(3) \) [95].
2dIM [100, 101] calls for substantiation. A traditional experimental approach to nonconserved kinetics has been the twisted nematic liquid crystal (TNLC) [64, 81, 95], which can be quenched through a first-order transition between the high-temperature fluid phase and the lower-temperature nematic phase. Not forgetting to pay due tribute to these previous systems that aimed at measuring dynamic scaling properties, we however note that the inexorable slow quench rate implied in the isotropic-nematic transition and the issues of metastability, nucleation and nuclei growth stages, have conspired to blur comparisons with predictions made from genuinely sudden transitions across second-order points. It is also highly desirable to seek for a versatile setup in which different sorts of kinetics can be triggered.

Here, we exploit electrically-driven regimes in TNLC layers to induce a virtually instantaneous switch from a spatiotemporally chaotic (disordered-like) state to an equilibrium, two-phase competing state. Because of the sudden removal of the driving, problems related to first-order transitions are negligible (if existent), while thermal effects are finely controlled. In addition, this setup offers an elegant way to induce different sorts of kinetics by transiting between nonequilibrium states. We focus on the simplest transition type that is towards the equilibrium, so that we assess the several aspects on 2d Model A waiting for experimental elucidation.

We firstly show well-established Model A features such as the sharp evidence of dynamic scaling with $\gamma = 2$. Then we move to assess the Bray-Humayun amplitude and test OJK and TUG theories in length scales much larger than the Porod’s regime. Paying attention to the asymmetry in the twisted phases, an aging hypothesis for Model A systems is confirmed by a rescaled two-time correlator whose master form is used to confront the OJK, TUG and LSI theories. We find that a particular solution of LSI, which fits 2dIM data and assumes the Fisher-Huse ($\lambda$), gives the superior account for the experiment. TUG and LSI theories. We find that a particular solution of LSI, which fits 2dIM data and assumes the Fisher-Huse conjecture, gives the superior account for the experiment.

Time correlation properties are studied in Sec. [IV] Two-time correlators are presented in Sec. [IV A] it covers the measurement of $\lambda$, dynamic scaling aspects, and tests of theoretical forms. Local persistence probability and morphology of persistence clusters are presented in Sec. [IV B]. We conclude this contribution in Sec. [V].

II. EXPERIMENTAL SETUP AND ELECTRICAL SWITCHING

We exploit the electrohydrodynamic convection of nematic liquid crystal [102,106] that arises when a thin layer (typically 10 to 100 $\mu$m) of nematic liquid crystal – with negative dielectric anisotropy ($\varepsilon_{xx} - \varepsilon_{zz} < 0$)[107] and positive conductance anisotropy ($\sigma_{xx} - \sigma_{zz} > 0$) – is subjected to an electric voltage of $V \sim 10$ V or higher [108]. Upon increasing $V$, one observes a sequence of convective patterns [109,111] and finally a spatiotemporally chaotic state called the dynamic scattering mode 2 (DSM2) [112,113,116]. DSM2 stands out by the presence of a high density of disclinations; fluctuations of the director field are short-range correlated in space and time: $\approx 1-3 \mu$m and 10 ms, respectively, for a 50 $\mu$m thick layer [109]. We use DSM2 as a disordered initial condition to study relaxation towards the equilibrium state.

We built a capacitor of parallel glass plates coated with indium tin oxide. The gap was set by polymeric spacers of thickness 12 $\mu$m and enclosed an empty region $16 \times 16$ mm. Prior to assembly, a polyvinyl alcohol film was coated over each plate and rubbed to set the direction of the surface alignment. Rubbing directions were set to be orthogonal between the two plates to realize a twisted nematic liquid crystal (TNLC) cell [117]. The capacitor was then filled with $N$-4-methoxybenzylidene-4-butylaniline (purity $> 98.0\%$, Tokyo Chem. Ind., Japan) doped with 0.01 wt% of tetrabutylammonium bromide (Tokyo Chem. Ind., Japan). Director field twisted along the bulk in either left- or right-handed orientation in the cell [Fig. 1(c)].

The cell was inserted in a handmade thermocontroller [Fig. 1(b)]; the set was placed on the stage of an inverted microscope (IX73, Olympus, Japan). Green-filtered, circularly polarised light was emitted through the cell; transmitted images were recorded by a charge-coupled device [Fig. 1(a)]. We monitored an area of dimensions $L_x \times L_y = 2.9 \times 2.2$ mm in the sample by a $\times 4$ objective. The whole apparatus was inserted in a thermally isolated chamber whose inner temperature was kept roughly constant by a thermostat circulator. Peltier elements in the thermocontroller were connected to a bipolar current supply set by a proportional-integral-derivative feedback using the signal of one of the thermistors located at 1 mm to the cell. During the experiments, we kept the temperature at 25°C with fluctuations smaller than $\pm 1.5$ mK according to the reference thermistor. Prior to the experiments, we also monitored the temperature at different positions near the cell. Ther-
minal fluctuations of thermistors not used for the feedback control were at most ±10 mK, whilst the temperature gradient formed across the cell was a few tens of mK.

Ordering kinetics was triggered as follows. We applied a 100 Hz, 70 V sinusoidal voltage across the cell to generate the DSM2 state. Equilibrium twist alignment of the director field was then replaced by disordered and fluctuating configurations. The cell was kept in DSM2 during 120 s (≈10^4 correlation times of the DSM2 state) before we instantaneously removed the electric field. From this switching time t = 0 s onwards, the twisted boundary conditions for the director field induced either left- or right-handed twist; these twisted alignments were formed after complex disentangling of disclination loops. We recorded the dynamics at 3 Hz over 2000 s with pixel size a ≈ 1.82 μm; 20 independent relaxation histories were collected. Due to the polarization optics, twists of opposite handedness were distinguished in the images by darker and brighter shades of grey [Fig. 1(d)]. We attributed to each pixel a local state variable ϕ(r, t) that took ϕ = −1 and +1 at the dark- and bright-grey colour pixels, respectively. Domains of the opposite handedness were bordered by disclination lines [Fig. 1(c) and (d)] that appeared as dark lines of thickness ξ ≈ a. Because it was hard to fully detect disclinations out of dark-grey colour phases, we regarded disclinations as part of the ϕ = −1 phase: a fair approximation because ξ ≈ a and because interest is in the scaling regime where ξ/ℓ(t) → 0. Nonetheless, we warn for specific instances where the binarization effect might play some role. Data for t ≤ 2.33 s are neglected because the large fraction of disclinations blurred rational detection of domains.

III. SPATIAL SCALING AND PHASE ASYMMETRY

A. Shrinkage law

Figure 2 shows TNLC domain configurations in ordering (see also Supplemental Videos 1 and 2). Configurations at t = 2.66 s appear as a random patchwork of hundreds of domains (or clusters) that progressively coarsen afterwards. Initial irregular contours of domains become smoother in time. While the total length of boundaries tends to decrease, larger domains increase their area at the expense of smaller shrinking ones. Around ten clusters are left at t ≈ 150 s, but ordering is only completed a decade later, at a finite-size equilibration time t* ≈ 1.2 × 10^3 s.

To quantify shrinkage of interfaces (disclinations), we measure the interface density ρ(t) as the number of neighbouring pairs with the opposite handedness multiplied by a/LxLy. Based on scaling hypothesis [5]:

\[ \rho(t) \sim t^{-1/z}. \]  

(1)

Figure 3 shows that such a decay, dictated by a slope near 1/2, occurs in the experiment for nearly two decades. The decay is better quantified by the effective exponent, z^−1(t) = −d(ln ρ(t))/d ln t, whose plateau for 6 s ≤ t ≤ 60 s indicates the core of the algebraic regime (Fig. 3 inset). Time averaging in this regime, or fitting Eq. (1) to the data of each realization and averaging, we obtain:
FIG. 2. Configurations of the ordering of twisted nematic liquid crystals (TNLC) during a relaxation towards the equilibrium. The dynamics was triggered by a virtually instantaneous electrical switch (at $t = 0$ s) from a spatiotemporally chaotic called dynamic scattering mode 2 (DSM2) towards the two-phase competing, equilibrium state. Different shades of grey correspond to twists of opposite handedness in the nematic director field. See also Supplemental Videos 1 and 2.

![Configuration of the ordering of twisted nematic liquid crystals](image)

FIG. 3. Density of the interface length in time. Error bars indicate one standard deviation. Dashed black line is a guide for eyes with the slope $1/2$ expected for Model A dynamics. Inset shows the time series of the effective exponent, $1/z_{eff} = -d(\ln \rho(t))/d\ln t$, which fluctuates around $1/2$ (dashed black line) within the core of the power-law decay, $\rho(t) \sim t^{-1/2}$, for $6 \lesssim t \lesssim 60$ s.

\[ 1/z = \begin{cases} 0.505(15) & \text{(effective exponent)}; \\ 0.498(6) & \text{(power-law fit)}. \end{cases} \tag{2} \]

The number(s) into parentheses refers to the uncertainty (standard deviation for the effective exponent; standard error for power-law fit) in the last digit(s). Converting to $z$: $z = 1.98(6)$ (effective exponent) and $z = 2.008(22)$ (power-law fit). These results are in accurate agreement with the theory for curvature-driven interface dynamics \[3\] \[4\] \[10\], a hallmark of the Model A systems.

B. Phase asymmetry and domain shrinkage

The twist phases in the cell are not strictly $Z_2$-symmetric because the rubbing method inserts a nonvanishing pretilt angle for the director field at the anchoring surfaces \[119\]. Misalignment between top and bottom surfaces may add for the symmetry breaking \[120\].

Asymmetry in the kinetics is captured by the “magnetization”, $M(t) = 1 - 2A(t)$, defined from the area fraction $A(t)$ of the $\phi = -1$ (darker) phase [Fig. 4(a), inset]. Note that $M(t)$ contains a contribution from disclinations included in $\phi = -1$. To remove this effect, we carried out an extra set of 20 independent experimental runs with the cell placed upside down: by this changing, the brightness of the two competing domains is inverted, so that the sign of $\phi$ was flipped – except for disclinations which, keeping looking as dark-coloured pixels, remained regarded as $\phi = -1$ after binarization. Measuring the magnetization $M_{flip}(t)$ for this sign-flipped data set, we compute $M'(t) = (M(t) - M_{flip}(t))/2$ as the disclination-free $M'(t)$ magnetization [Fig. 4(a), main panel]. The $M'(t)$ reveals that the $\phi = -1$ phase is slightly favoured in the main data set, albeit asymmetry is small; for instance, $|M(t)|, |M'(t)| \lesssim 0.2$ for $t \lesssim 60$ s. Time average in this interval gives $\langle M \rangle = -0.17(3)$, which corresponds to $\bar{A} = 0.58(2)$.

Asymmetry is also seen in the shrinkage rate of domains of phase $\phi$ surrounded by a “sea” of $-\phi$. For curvature-driven dynamics, the squared radius $R^2(t)$ of a circular domain (or “bubble”) evolves, from a reference time $t_0$, as \[3\]:

\[ R^2(t) - R^2(t_0) = -2D_{AC}(t - t_0) + K, \tag{3} \]

where $D_{AC}$ is the Allen-Cahn diffusion parameter and $K$ is a constant. We monitored 6 independent, isolated and nearly circular bubbles; 3 of them made of the $\phi = -1$ favoured phase, and the remaining ones made of $\phi = +1$. Measuring their area and equalizing $\pi R^2(t)$ to define $R^2(t)$ for each bubble, we accompany their squared radius in time [Fig. 4(b)]. Fitting according Eq. (3) yields $D_{AC} = 117.3(17)\,\mu m^2 s^{-1}$ and $D_{AC}^+ = 126.0(6)\,\mu m^2 s^{-1}$ for $\phi = -1$ and $+1$ bubbles, respectively. The asymmetry translates in nearly $4\%$ of deviation of each diffusion constant relative to the average,

\[ D_{AC} = 122(4)\,\mu m^2 s^{-1}, \tag{4} \]
where the number in the parentheses was set to cover the interval \([D_{AC}, D_{AC}^\prime]\).

We remind that the initial state of the experiment is DSM2, which may provide an unbiased (or weakly biased) initial condition because the surface anchoring of the director field is not preserved \([121]\) and because the bulk director is strongly disturbed by the high density of disclinations. Thus, the observed asymmetry — including the one in the earlier times not captured in Fig. 4(a) — is likely developed along with the kinetics instead of being set by initial conditions.

The usual presence of asymmetry in experiments demands additional care when comparison is made with predictions for \(\mathbb{Z}_2\)-symmetric models. As we shall see, although this a simple task for some quantities such as the spatial correlator (Sec. III C), they are subtle for, e.g., the time correlator (Sec. III C).

**C. Growth law and the Bray-Humayun amplitude**

The two-point spatial correlator,

\[
C_s(r, t) = \langle (\phi(r', t)\phi(r' + r, t))_{r'} \rangle_e, \tag{5}
\]

where \(\langle \ldots \rangle_r\) denotes the spatial average and \(\langle \ldots \rangle_e\) the ensemble average, is shown as function of \(r = |r|\) in Fig. 5(a). The larger the \(t\), the slower the \(C_s\) decay because of the domain growing size \(l(t)\); here set it as the length at which \(C_s(r, t) = 0.2\) for \(|r| = l(t)\). The inset of Fig. 5(a) displays the growth law with \(z = 2\),

\[
l(t) \simeq Bt^{1/2} \quad (\xi \ll l(t) \ll L_{sys}), \tag{6}
\]

with a nonuniversal amplitude \(B\) discussed below.

A time-independent scaling function \(\mathcal{F}(\cdot)\),

\[
C_s(r, t) \simeq \mathcal{F}(x) \quad (x = |r|/l(t)), \tag{7}
\]

is defined when dynamic scaling holds, as it does in the experiment within the accuracy of \(\delta C_s(x, t) \approx 0.07\) (standard deviation) for \(t \lesssim 21\) s and \(x \leq 3\) [see Fig. 5(b)]. Moreover, the Porod’s law \([65]\),

\[
C_s(r, t) \simeq 1 - \alpha x \quad (\text{for } x \ll 1), \tag{8}
\]

is emphasized in the inset of Fig. 5(b). The angular coefficient \(\alpha\), estimated from the convergence of \((1 - C_s)/x\) for \(x \to 0\) at a late time \([t = 21.33\) s; see Fig. 5(c)] reads:

\[
\alpha = 0.92(9). \tag{9}
\]

We also remind that Bray and Humayun \([66]\) exactly derived \(C_s(r, t)\) at \(r \ll l(t)\), for \(n \leq d\):

\[
C_s(r, t) \simeq 1 + A(d, n)\rho_{\text{Euc}}(t)\xi^n \quad (\xi \ll r \ll l(t)), \tag{10}
\]

where \(\rho_{\text{Euc}}\) is the interface density measured with the Euclidian metric; \(A(d, n)\) is the universal Bray-Humayun (BH) prefactor \([66]\):

\[
A(d, n) = \pi^{(n/2) - 1} \frac{\Gamma(-n/2)\Gamma(d/2)\Gamma^2((n + 1)/2)}{\Gamma((n + d)/2)\Gamma(n/2)}, \tag{11}
\]

for odd \(n\), where \(\Gamma(\cdot)\) denotes the gamma function. For our system, \(d = 2\) and \(n = 1\), so that \(A(2, 1) = -4/\pi\).

To assess Eq. (10) in the experiment, we convert \(\rho_{\text{Euc}}\) to \(\rho\) in the Manhattan metric because this metric (the
FIG. 5. (a) Spatial two-point correlator $C_s(r, t)$ [Eq. (5)] versus $r = |r|$ at different times. Data were obtained by rotational average (averaging values of $C_s(r, t)$ at different $r$ with the same $r = |r|$), data binning, and ensemble average. Error bars indicate one standard deviation. Dashed line indicates $C_s(r, t) = 0.2$. Inset shows the growth of $l(t)$, where dashed red line shows $l(t) = Bt^{1/2}$ with $B = 24 \mu m \text{s}^{-1/2}$ estimated below Eq. (14). (b) Dynamic scaling hypothesis is confirmed from the time-independence of $C_s(r, t)$ versus $x = r/l(t)$. Grey dashed line represents the Porod law, $C_s(r, t) \sim 1 - \alpha x$, with $\alpha = 0.92(9)$ estimated from (c). Inset of (b) is a zoom in on the region $x \ll 1$ for the main plot. (c) Estimation of $\alpha$ by reading $(1 - C_s)/x$ for $x \to 0$; dashed line indicates the mean value, 0.92, of the ordinates in the range $x \leq 0.5$. (d) Plateau of the rescaled interface density $\rho t^{1/2}$ for 6 s $\leq t \leq 60$ s; dashed line indicates the mean value 0.0381 $\mu m^{-1/2}$. (e) Plot of $(1 - C_s)t^{1/2}/r$ versus $r$ from which the amplitude $A^*$ is estimated in the limit $r \to 0$. Dashed line indicates the average value $CA^* = 0.0377 \mu m^{-1}s^{1/2}$.

Legends in (b), (c), and (e) are the same as that in (a).

Manhattan one) was used in Fig. 3. Thus, $\rho_{\text{Enc}} = L_{\text{Enc}}^2/L_\text{Man}/L_y$, is replaced by $\rho = L_\text{Man}/L_y$, where $L_{\text{Enc}}$ ($L_\text{Man}$) is the interface length in the Euclidean (Manhattan) ruler. Noting that $L_{\text{Man}}^2/L_{\text{Enc}}^2 = 4/\pi$ holds for isotropic space [60], Eq. (10) becomes:

$$C_s(r, t) \simeq 1 - A^* \rho(t) r \quad \left(\xi \ll r \ll l(t)\right), \quad (12)$$

with the BH amplitude $A^* = 1$.

We test such prediction as follows. Defining $C$ as the prefactor of $\rho(t) \approx Ct^{-1/2}$ (Fig. 3), and recalling $B/l(t) \approx t^{-1/2}$ [Eq. (6)], we insert these in Eq. (12) to obtain:

$$C_s(r, t) \simeq 1 - CA^* r t^{-1/2}$$

$$= 1 - A^* BC \frac{r}{l(t)} \quad \left(\xi \ll r \ll l(t)\right), \quad (13)$$

and we identify $\alpha = A^* BC$.

From the plateau in the plot of $\rho(t)t^{1/2}$ versus $t$ [Fig. 5(d)], observed for 6 s $\leq t \leq 60$ s, we read $C = 0.0381(11) \mu m^{-1}s^{1/2}$. Similarly from $(1 - C_s)t^{1/2}/r$ versus $r$ in Fig. 5(e), the plateau interval $0 \leq r \leq 0.5 l(t) \approx 57 \mu m$ for $t = 21.33$ s - which was used in the measurement of $\alpha$ in Eq. (6) - gives $CA^* = 0.0377(14) \mu m^{-1}$, so that we find:

$$A^* = 0.99(7) \quad (14)$$
FIG. 6. Two-point spatial correlator $C_s(r, t)$, Eq. (9), in the experiment (symbols) compared with the forms predicted by the Ohta-Jasnow-Kawasaki (OJK) theory and the theory of unstable growth (TUG) own to Mazenko. The twisted nematic liquid crystal (TNLC) data for $t \leq 21.33\, s$ are identical to those in Fig. 5(a)(b), but the number of data points is reduced for the sake of visibility. Error bars indicate one standard deviation. (a) $C_s(r, t)$ for the experiment (symbols) at several times and the OJK theoretical form (dashed lines), Eq. (15), with $D_{OJK} = 61\, \mu m^2 s^{-1}$ obtained from the relation with the Allen-Cahn constant: $D_{OJK} = D_{AC}/2$. (b) Rescaled $C_s(r, t)$ in the experiment compared with both OJK (dashed black line) and TUG (dash-dotted blue and solid turquoise lines) functions. The two TUG curves were obtained with different values of $D_{TUG}$ as indicated in the legend. The inset shows the same data in the main plot, but the horizontal axis is displayed in logarithmic scale.

in striking accordance with the BH prediction $A^* = 1$ [60]. By passing, from $\alpha = A^*BC = 0.92(9)$ in Eq. (9) and $CA^* = 0.0377(14)\, \mu m^{-1}$ above, one has $B = 24(3)\, \mu m^{-1/2}$: a value that describes very well the experimental growth law [Fig. 5(a), inset].

D. Experimental test of Gaussian theories

In scalar 2d Model A, the order parameter field tends to change discontinuously in the scaling limit when interfaces are crossed. Because such discontinuity is hard to deal with analytically, both OJK and TUG theories replace the order parameter by an auxiliary field that smoothly varies in the whole space. OJK does so by describing interfaces as a collection of positions satisfying $u(r, t) = 0$, where $u(r, t)$ is an auxiliary field such that $\phi(r, t) = \text{sgn}(u(r, t))$ [5] [51]. Starting from the Allen-Cahn interface motion [4] and relying upon approximating $\nabla_j u \nabla_k u / |\nabla u|^2 \approx \delta_{jk}/d$ with $j, k = 1, 2, \ldots, d$, OJK reaches at the diffusion equation for $u(r, t)$. Assuming $u(r, 0)$ as a Gaussian random field [122], the form of $\mathcal{F}(\cdot)$ in Eq. (7) [5] [51] is [123]:

$$
\mathcal{F}_{OJK}(r/t^{1/2}) = \frac{2}{\pi} \sin^{-1} \left[ \exp \left( -\frac{r^2}{8 D_{OJK} t} \right) \right],
$$

with $D_{OJK} = D_{AC}/2$ for $d = 2$ [51].

We test Eq. (15) in the experiment, free from fit, using $D_{OJK} = D_{AC}/2 = 61(2)\, \mu m^2 s^{-1}$ after Eq. (4). Because $C_s(r, t) \rightarrow M(t)^2$ for $|r| \rightarrow \infty$ and $M(t) \neq 0$ in the experiment [Fig. 5(a), inset], but $M = 0$ for OJK, comparison is restricted to $C_s(r, t) \geq M(t)^2$. Figure 5(a) compares the OJK form with TNLC data in the region $C_s(r, t) \geq M(t)^2 \approx 0.04$ for $t = 42.66\, s$, and $\geq 0.02$ for shorter times. We find noteworthy agreement not only in the initial linear decay (Porod’s regime), but also in the interdomain length scale beyond it. Agreement is better for larger times. We also may look at the TNLC growth law, $l(t) \approx Bt^{1/2}$ with $B = 24(3)\, \mu m^{-1/2}$ (Sec. III C), and compare it with the OJK outcome, $l(t) = \sqrt{-SD_{OJK} \ln(\sin(\pi/10))} t^{1/2}$ from $\mathcal{F}_{OJK}(l(t)/t^{1/2}) = 0.2$ in Eq. (15): this comparison yields $D_{OJK} = 64(18)\, \mu m^2 s^{-1}$ in compatibility with $D_{AC}/2 = 61(2)\, \mu m^2 s^{-1}$.

Now we turn attention to TUG. Developed from the $d$-dimensional TDGL equation, TUG introduces an auxiliary field $m(r, t)$ interpreted as the shortest distance between $r$ and the nearby interface [124]. Assuming $m(r, t)$ as a Gaussian field, TUG predicts that $\mathcal{F}(\cdot)$ in Eq. (7) is given by $\mathcal{F}(r/t^{1/2}) = \mathcal{F}_{TUG}(g)$, which satisfies the following differential equation [52] [53]:
with \( \mathcal{F}_{TUG}(0) = 1, \ g = r/(4D_{TUG}t)^{1/2} \), and \( D_{TUG} \) is a constant. The value \( \mu(d) \) is \( \mu(2) \approx 1.104 \) [52-53] for the case of interest here [125].

To test TUG form, we numerically integrated Eq. (16) by a Euler-Heuns discretization with a uniform step \( \delta g = 10^{-6} \) in \( d = 2 \), and boundary conditions \( \mathcal{F}_{TUG}(0) = 1, \ \frac{d\mathcal{F}_{TUG}(g)}{dg}|_{g=0} = -(2/\pi)^{1/2} \). After, to determine \( D_{TUG} \), we rely on the Porod’s regime \( g \ll 1 \) of the TUG correlator [53-54]:

\[
\mathcal{F}_{TUG}(g) \approx 1 - \left( \frac{D_{TUG}2\pi(d-1)}{r} \right)^{1/2} t^{1/2}.
\]

Relating Eq. (17) with \( C_s(r, t) \approx 1 - (\alpha/B)(r/t)^{1/2} \) [from Eqs. (3) and (8)], we find \( D_{TUG} = (B/\alpha\sqrt{2} \pi)^2 = 1/2(\pi(CA^*)^2 = 112(8) \mu m^2 s^{-1} \).

Figure 6(b) compares the experimental data with \( \mathcal{F}_{TUG}(g) \) for \( D_{TUG} = 112 \mu m^2 s^{-1} \) (the most probable value), as well as with \( \mathcal{F}_{OJK}(r/t)^{1/2} \) for \( D_{OJK} = 61 \mu m^2 s^{-1} \). For the sake of visibility, we show only data for \( t = 21.33 s \) and 42.66 s because they are in the core of the \( z = 2 \) scaling regime (see Fig. 3). Figure 6(b) suggests that \( \mathcal{F}_{TUG}(g) \) describes the experiment better than the TUG counterpart; however, by adjusting the value of \( D_{TUG} \) within the uncertainty, specifically by setting \( D_{TUG} = 104 \mu m^2 s^{-1} \), the form \( \mathcal{F}_{TUG}(g) \) can also be made reasonably close to the data [Fig. 6(b), inset]. Thus, both the OJK and TUG theories are good descriptors (within the statistical accuracy) for the TNLC correlator and this includes the description of length scales larger than those in the Porod’s regime. Unlike \( D_{OJK} \), \( D_{TUG} \) does not seem to have a trivial connection with \( D_{AC} \), albeit it can also be reliably evaluated by fitting the experimental correlator to the TUG Porod’s regime, Eq. (17), as we have done. It is overall noteworthy that OJK performs remarkably well without such fitting.

### IV. TIME STATISTICS

#### A. Two-time autocorrelation

The two-time autocorrelator,

\[
C_t(t, t_0) = \langle (\phi(r, t_0)\phi(r, t)) \rangle_c \quad (t \geq t_0),
\]

is anticipated [126] to decay in the coarsening stages (for most of the cases [127]) as [5]:

\[
C_t(t, t_0) \approx \mathcal{H}(t/t_0), \quad \mathcal{H}(y) \sim y^{-\lambda/z} \quad (y \gg 1),
\]

for large \( t_0 \) and \( t \), where \( \mathcal{H}(y) \) is a universal scaling function and \( \lambda \) is the autocorrelation exponent introduced by Fisher and Huse (FH) [70]. As described in Sec. 1 2d Model A theories suggest: \( \lambda_{FH} = 5/4 \) [70]; \( \lambda_{OJK} = 1 \) [79]; \( \lambda_{TUG} \approx 1.2887 \) [78].

The scaling function \( \mathcal{H}(y) \) in the OJK model is [5-79]:

\[
\mathcal{H}_{OJK}(y) = \frac{2}{\pi} \sin^{-1}\left(\frac{4y}{(1+y)^2}\right)^{d/4}.
\]

For TUG, the function \( \mathcal{H}_{TUG}(g, \tau(y)) \), analogously defined from the two-point spatiotemporal correlator \( \langle \phi(r', t_0)\phi(r', t_0) \rangle_c \), with \( g = r/(4D_{TUG}t)^{1/2} \) and \( \tau(y) = (\ln y)/4 \), satisfies the differential equation [78]:

\[
\frac{\partial \mathcal{H}_{TUG}(g, \tau(y))}{\partial \tau} = \frac{1}{\mu} \nabla^2 g \mathcal{H}_{TUG} + 2g \cdot \nabla_g \mathcal{H}_{TUG} + \frac{1}{\mu} \tan\left(\frac{\pi}{2} \mathcal{H}_{TUG}\right).
\]

This equation can be integrated numerically from the initial condition \( \mathcal{H}_{TUG}(g, 0) \), which is just the vectorial analogue of \( \mathcal{F}_{TUG}(g) \) that we also numerically obtained in Eq. (16). Here, we are interested in \( \mathcal{H}_{TUG}(y) = \mathcal{H}_{TUG}(0, \tau(y)) \).

The LSI theory is based on the hypothesis that correlators transform covariantly under the conformal group (that is, under space-time scale transformations set by the dynamic exponent \( z \) [82]. For 2d scalar Model A, LSI theory yields [128]:

\[
\mathcal{H}_{LSI}(y) = y^{\lambda/2}(y-1)^{-\lambda} \Phi\left(\frac{y+1}{y-1}\right),
\]
with

\[
\Phi(q) = E_1 q^{1-\lambda} \left[ \left( 1 + \frac{1}{q} \right)^{3-2\lambda} - \frac{1}{E_3 q} \gamma(\lambda, E_3 q) \right] + E_2 \left[ E_3^{-\lambda} q^{-\lambda} \gamma(\lambda, E_3 q) + \frac{e^{-E_3 q} - 1 + 2 F_1(\lambda - 1, 1; 3 - \lambda; -1/q)}{q} \right],
\]

(23)

where \(E_1, E_2, E_3\) are constants, \(\Gamma\) is the gamma function, \(\gamma\) is the lower incomplete gamma function, and \(2F_1\) is the hypergeometric function. Constants were determined from 2dIM simulations after a quench from the paramagnetic state to the lower-temperature \(T = 0\) state, with \(\lambda = \lambda_{FH}\); they read [129]: \(E_1 = -0.601, E_2 = 3.94\) and \(E_3 = 0.517\). Hereafter, \(\mathcal{H}_{LSI}(y)\) with this set is referred to as the LSI-2dIM correlator.

Before comparing experiment with theories, it is crucial to subtract effects from a non-vanishing \(M(t)\) in the kinetics, otherwise, the asymptotic decay of the two-time correlator \(C(t, t_0)\) becomes dependent on which phase equilibrates the system. Evaluating the magnetization \(\hat{M}(t) = \langle \phi(r, t) \rangle_r\), and the bare correlator \(\hat{C}(t, t_0) = \langle \phi(r, t) \phi(r, t_0) \rangle_r\) for each realization, we define a modified correlator \(C'(t, t_0)\):

\[
C'(t, t_0) = \left( \frac{\hat{C}(t, t_0) - \hat{M}(t)\hat{M}(t_0)}{1 - \hat{M}(t)\hat{M}(t_0)} \right)_e.
\]

(24)

This correlator satisfies \(C'(t_0, t_0) = 1\) and \(C'(t \rightarrow \infty, t_0) \rightarrow 0\), since \(\hat{C}(t \rightarrow \infty, t_0) \rightarrow M(t)M(t_0)\). Note that \(C'(t, t_0) \approx C(t, t_0)\) as long as \(\hat{C}(t, t_0) \gg \hat{M}(t)\hat{M}(t_0)\), which is roughly \(\approx 0.03\) for \(t, t_0 \lesssim 60\) s for instance.

Figure 7(a) displays \(C'(t, t_0)\) in the experiment for \(t_0 = 2.66\) s, 5.33 s, and 10.66 s against \(t\) (inset) and \(y = t/t_0\) (main panel) in the double logarithmic scales. From the collapse of the three data sets shown in the main panel, we confirm the dynamic scaling hypothesis as prescribed in Eq. [19]. These data are then compared with the asymptotic power law decay \(C(t, t_0) \sim (t/t_0)^{-\lambda/z}\) with \(\lambda/z = 1/2\) (OJK), \(= 5/8\) (FH), and \(\approx 0.644\) (TUG). After the initial non-algebraic region, the data at intermediate times \(2 \lesssim y \lesssim 10\) tend to decay with a OJK-like value prior to their crossover to a faster, asymptotic decay. Using \(t_0 = 2.66\) s for which the asymptotic power law is most prominent, and \(z = 2\) shown in Eq. [2], we find

\[
\lambda = \begin{cases} 
1.03(5) & (2 \leq y \leq 10); \\
1.28(11) & (15 \leq y \leq 75),
\end{cases}
\]

(25)

whose values for the asymptotic TNLC decay are compatible with both the FH conjecture and TUG theory within uncertainty (uncertainties are evaluated from the effective exponents).
Figure 7(b) compares $C_f(t, t_0)$ in the experiment with the OJK [Eq. (20)], TUG [Eq. (21)], and LSI-2dIM [Eq. (23)] functions. OJK outcome is well above the TNLC data, a result similarly seen in tests with 2dIM simulations [22]. The TUG form, which to our knowledge has not been confronted with simulations or experiments, exhibits pronounced deviations from the experiment for $y \lesssim 25$ while it approaches to the collapsed correlator from below. This suggests that neither OJK or TUG theory may perform well to describe the universal autocorrelator form in scalar, 2d Model A dynamics. Interestingly, the experimental correlator is best accounted for by the LSI-2dIM theory, except for $y \lesssim 2$ since LSI solutions cannot describe such regime [82, 129]. The nice agreement with LSI-2dIM can also be appreciated in the log × linear plot shown in the inset of Fig. 7(b).

In an overall view, individual theories cannot account simultaneously for all aspects (or regimes) of the experimental two-time autocorrelator. Indeed, although both the FH conjecture and TUG theory are good descriptors for the slope of the power-law decay, the autocorrelator functional form itself is not captured by TUG, nor OJK. Instead, this form is overall best accounted for by the LSI-2dIM correlator, except for the $y \lesssim 2$ regime; note that LSI-2dIM incorporates the FH assumption, which is also in harmony with our experimental observations.

**B. Local persistence**

The local persistence probability $Q(t, t_0)$ is the probability that $\text{sgn}(\phi(r, \tilde{t}))$ does not change during $\tilde{t} \in [t_0, t]$. It typically shows nontrivial algebraic decay [67, 68],

$$Q(t, t_0) \sim (t/t_0)^{-\theta} \quad (t \gg t_0),$$  \hspace{1cm} (26)

with the persistence exponent $\theta$, as we can here directly appreciate in the experiment for several values of $t_0$ [Fig. 8(a)]. The plot $Q(t, t_0)$ versus $y = t/t_0$ yields an excellent data collapse within $\Delta Q \approx 0.04$ (standard deviation) for $y \lesssim 10^2$ [Fig. 8(b)]. The decay is quantified by the exponent, $\theta_{\text{eff}}(t, t_0) = -d[\ln Q(t, t_0)]/d[\ln (t)]$, shown in the inset of Fig. 8(b) for $t_0 = 2.66$ s and 5.33 s. Both of the $\theta_{\text{eff}}(t, t_0)$ curves fluctuate around their mean values for $8 \leq y \leq 50$. Averaging $\theta_{\text{eff}}$ data in this interval gives 0.1938(14) and 0.1976(27) for $t_0 = 2.66$ s and 5.33 s, respectively. We also evaluate $\theta$ by applying a least-squares fit individually to $Q(t, t_0)$ data of each realization, constrained to the interval $8 \leq y \leq 50$, and then ensemble averaging; the results are: $\theta = 0.1941(26)$ and 0.199(4) for $t_0 = 2.66$ s and 5.33 s. Integrating these results, we report

$$\theta = \begin{cases} 
0.196(3) & \text{(effective exponent);} \\
0.196(4) & \text{(power-law fit),} 
\end{cases}$$  \hspace{1cm} (27)

where uncertainties include (sum up) both the uncertainty from each estimate and the deviation between the two estimates for $t_0 = 2.66$ s and 5.33 s. We remark that these results do not significantly change when the interval $8 < y < 60$ s is used, since it corresponds to the interval $t \lesssim 60$ s predominantly used in Sec. III.

Results in Eq. (27) agree with systematic 2dIM outcomes [88] ($\theta \approx 0.195$ and $\theta = 0.199(2)$ for free and periodic boundary conditions, respectively), but they significantly differ from $\theta_{\text{DIF}} = 3/16 = 0.1875$ [99] and, hence, from the OJK theory as well [92]. This may suggest, based on a concrete experience, the former as the true $\theta$ for 2d Model A systems.

To probe morphological aspects of persistence, we define an index $\chi(r, t)$ such that $\chi = 1$ if $\text{sgn}(\phi(r, \tilde{t}))$ does not change during $\tilde{t} \in [t_0, t]$, with $t_0$ hereafter pinned at $t_0 = 2.66$ s, and $\chi = 0$ otherwise. The index $\chi(r, t)$ can only change from 1 to 0; such change happens only when an interface crosses the position $r$ for the first time since $t_0$. 

![Figure 8](image-url)
\[ \chi(r, t) \]

FIG. 9. Configurations of the local persistence index \( \chi(r, t) \) (see text) computed in the ordering of twisted nematic liquid crystals (TNLC) for a reference time \( t_0 = 2.66 \text{s} \). The black and white pixels correspond to \( \chi = 0 \) (nonpersistence) and 1 (persistence), respectively. See also Supplemental Videos 3 and 4.

\[ G(\zeta, t) \]

FIG. 10. Statistics of the local persistence in the ordering kinetics of TNLC phases. In all the panels, dash-dotted and dashed lines are guide to eyes; the values of their slopes or ordinates are indicated in the respective plots. (a) Mean counting \( G(\zeta, t) \) of \( \chi = 1 \) (persistence) pixels found inside a square of size \( \zeta \) that glides over the configurations of the persistence index \( \chi(r, t) \) (reference time \( t_0 = 2.66 \text{s} \)). (b) Effective dimension \( d_{f, \text{eff}}(\zeta, t) \) for the morphology of persistence clusters as a function of \( \zeta \). (c) Covariance \( C_{\chi}(r, t) \) [Eq. (29)] of the index \( \chi \) versus \( r = |r| \). Data were obtained by rotational average (averaging over different \( r \) with the same \( r = |r| \)), data binning, and ensemble average. The legend of the symbols is given in (b). Left inset depicts the effective exponent \( \eta_{\text{eff}}(r) \) related to a curve, \( t = 1365.33 \text{s} \), in the main panel. Right inset shows \( C_{\chi}(r, t) \) in time for a length \( (r = 500 \mu\text{m}) \) much larger than the crossover length \( \zeta^*(t) \). (d-e) \( d_{f, \text{eff}}(\zeta, t) \) and \( C_{\chi}(r, t) \) versus \( \zeta/t^{1/2} \) and \( r/t^{1/2} \), respectively, following the dynamic scaling hypothesis. Legends in (c),(d),(e) are as that shown in (b).

Snapshots of \( \{\chi(r, t)\} \) are presented in Fig. 9; they reveal how a single \( \chi = 1 \) cluster is progressively fragmented because of the interface motion in real space. Surviving clusters develop ragged edges in both internal and external contours resembling fractal objects (see Supplemental Videos 3 and 4). The fractal dimension \( d_f \) of this morphology is estimated by combining the gliding-box method [130] and the method in [131]. Here, a square
of side $\zeta$ glides over each $\{r(t),r\}$ image while counting the number of $\chi=1$ pixels inside the $i$th box, $g_i$, with $i=1,2,\ldots,n(r)$; $n(r)$ is the total number of distinct boxes that contain at least one $\chi=1$ pixel. The mean counting is $G(\zeta,t) = \langle |g_i| \rangle$, where $\langle \ldots \rangle$ is the average over $n(r)$ boxes glided on a snapshot at $t$. For fractal objects \cite{131,132}:

$$G(\zeta,t) \sim \zeta^{d_f} \quad (a \ll \zeta \ll L_{sys}). \quad (28)$$

In the experiment, $G$ exhibits a crossover from fractal to Euclidian length scales at a certain correlation length $\zeta^*(t)$ \cite{10}. The fractal regime is progressively built in time since it is confined to emerge for $a \ll \zeta \ll \zeta^*(t)$, with a growing length, $\zeta^*(t)$, that starts from $\zeta^*(t_0) = 0$. This progressive building is captured by $d_{f,\text{eff}}(\zeta,t) = \mu \ln(G(\zeta,t))/\mu \ln(\zeta)$ \cite{10} which shows a concave dependence with $\zeta$ and forms a plateau around a $t$-dependent minimum. For $\zeta \gg \zeta^*(t)$, $d_{f,\text{eff}}$ crossovers to $d = 2$. Finite-size and finite-time effects preclude observation of clear $t$-independent plateau for $\zeta \ll \zeta^*$; however, from the plateau at the minimum of $d_{f,\text{eff}}$ for large times, $t = 1365\,s$, we roughly read $d_f \approx 1.65(3)$ \cite{10}. The number in the parentheses indicate one standard deviation. Although $t = 1365\,s$ is far from the time interval where $M(t)$ is constant \cite{4,a},\cite{10},\cite{13}, properties of persistence clusters at long times are reminiscent of those in the early dynamics: this is revealed by the inner structures of persistence domains in the panel at $t = 1365\,s$ of Fig. \cite{9} which are mostly frozen and already present since $t = 43\,s$. Note also that the rough estimate 1.65(3) is consistent with $d_f = 1.61(5)$ reported for persistence clusters in the 2dIM \cite{10}. To evaluate $d_f$ from shorter times, for which $|M(t)|$ is small and $z = 2$ is clearly identified, we appeal to the correlator $C_X$:

$$C_X(r) = \frac{\langle \chi(r,t)\chi(r+r',t) \rangle_r}{\langle \chi(r)\rangle_r}, \quad (29)$$

which behaves as $\chi = 101,131,133$:

$$C_X \sim \begin{cases} \frac{r^{-\eta}}{t^{-\theta}} & (r \ll \zeta^*(t)), \\
\frac{r^{-\theta}}{t^{-\eta}} & (r \gg \zeta^*(t)) \end{cases}, \quad (30)$$

where $\eta$ is a universal exponent. At the crossover length, $C_X \sim \langle \chi^* \rangle^{-\eta} \sim t^{-\theta}$. Thus, if $\zeta^*(t) \sim t^{1/2}$ holds \cite{134}, then $\eta = z\theta$. Because $G(\zeta,t) \sim \int_{box} C_X(r,t) d^2r$ \cite{131}, where the integration is carried out over a box of size $\zeta$ centered at $r = 0$, integration after Eq. \cite{30} yields:

$$G(\zeta,t) = \begin{cases} \zeta^{-\eta} & (a \ll \zeta \ll \zeta^*(t)); \\
\zeta^{-\theta} \zeta^d & (\zeta^*(t) \ll r \ll L_{sys}) \end{cases}. \quad (31)$$

Comparing with Eq. \cite{28}, we identify:

$$d_f = d - \theta z, \quad (32)$$

and, consequently, $z\theta \leq d$.

Before using Eq. \cite{32} to evaluate $d_f$, we test Eq. \cite{30} in the experiment. As shown in Fig. \cite{10}c, we detect a nearly time-independent decay $C_X(r,t) \sim r^{-\eta}$ (arguably from $t = 43\,s$) within a region $20\,\mu m \lesssim r \lesssim \zeta^*(t)$ that extends in time. The exponent $\eta = -\frac{d(\ln C_X(r,t))/\ln r}{d(\ln r)}$ measured at $t = 1365.3\,s$ provide us with $\eta = 0.394(4)$ after averaging data for $38\,\mu m \leq r \leq 150\,\mu m$ \cite{10}. Note that this value is consistent with $\eta = 0.428(7)$ obtained for persistent 2dIM clusters \cite{10}. We also observe $C_X(r,t) \sim t^{-\theta}$ for $r \gg \zeta^*$ in the right inset of Fig. \cite{10}c, which displays $C_X(r,t)$ at $|r| = 500\,\mu m$. Finally, the assumption $\zeta^*(t) \sim t^{1/2}$ is verified by the collapse of $d_{f,\text{eff}}(\zeta,t)$ \cite{10} and $C_X(r,t)$ \cite{10} when plotted versus $\zeta/t^{1/2}$ and $r/t^{1/2}$, respectively. Deviations from collapsed forms for small lengths are due to effects of microscopic scales, such as $a$ and $\zeta$.

Now we safely rely on Eq. \cite{32} to evaluate $d_f$. Using $\theta$ in Eq. \cite{27} and $z$ in Eq. \cite{2}, we find:

$$d_f = \begin{cases} 1.613(14) \quad \text{(effective exponent)}; \\
1.609(9) \quad \text{(power-law fit).} \end{cases} \quad (33)$$

where the notes in the parentheses indicate the methods used to evaluate $1/z$ and $\theta$ in Eqs. \cite{2} and \cite{27}, respectively. Final estimates are slightly lower than the rough estimate $d_f = 1.65(3)$ (as may be expected), but, more importantly, they sharply agree with 1.61(5) from 2dIM simulations \cite{10}.

Altogether, experimental results in this Section provide firm evidence that the statistics of the local persistence probability correspond to a universal feature of 2d Model A systems. In addition, one can realize that the presence of a weak phase asymmetry has negligible, or very small, effect on those quantities.

V. CONCLUSIONS

We have studied the nonconserved phase-ordering kinetics of two-dimensional (2d) twisted nematic liquid crystals (TNLC) as a model experimental system to elucidate universal aspects in the Allen-Cahn (Model A) universality class. Exploiting electrically-driven hydrodynamical regimes of liquid crystals, we triggered a genuinely sudden transition between a disordered-like, spatiotemporally chaotic state, called dynamic scattering mode 2 (DSM2), towards a two-phase competing, equilibrium one. The sudden removal of the driving avoids the inexorable slow quench rate of thermally-induced transitions, besides it avoids the issues of metastability, nucleation and nuclei growth, implied in first-order transitions. In addition to its accuracy, the system offers an elegant way of inducing different types of kinetics by transiting between nonequilibrium states while finely tuning thermal effects.

Because spatiotemporal correlations of the director field are negligible in the DSM2 state \cite{109}, DSM2 essentially generates random initial conditions. When
models.

by the shrinking rate of $\phi$ count when one aims at testing theories for usual in experiments [95, 135] – must be taken into ac-

throughout the study, this weak asymmetry – that is $D$ slightly different values (4% deviation from the mean

fests in the Allen-Cahn diffusion coefficients, which take namics of disclinations (interfaces) [Eq. (3); Fig. 4(b)],

(“bubbles”) [Fig. 4(b)]. Noting the curvature-driven dy-

lation” for the director field, in left- and right-handed twists ensuing competition between two possible conformations

electrically-switched ($t = 0$ s) towards equilibrium, the ensuing competition between two possible conformations for the director field, in left- and right-handed twists (here noted $\phi = \pm 1$), resembles that between Ising spins when quenched from the higher-temperature ($T \gg T_c$, $T_c$ is the critical temperature) regime to the lower-

temperature ($T \ll T_c$) phase [see Fig. 2]. Nonetheless, the experiment develops a weakly asymmetry likely along its dynamics: this is revealed by the “magneti-

zation” $|\mathbf{M}| \lesssim 0.2 \neq 0$, for $t \lesssim 60$ s [Fig. 3(a)], and by the shrinking rate of $\phi = \pm 1$ spherical domains (“bubbles”) [Fig. 3(b)]. Noting the curvature-driven dy-

amics of disclinations (interfaces) [Eq. 2; Fig. 3(b)], the asymmetry between the twisted phases also mani-

fests in the Allen-Cahn diffusion coefficients, which take slightly different values (4% deviation from the mean $D_{AC} = 122(4) \mu m^2 s^{-1}$ for $\phi = \pm 1$. As we discussed throughout the study, this weak asymmetry – that is usual in experiments [95, 135] – must be taken into ac-

count when one aims at testing theories for $Z_2$-symmetric models.

After observing that the experiment firmly shows well-

known Model A features – such as the dynamic exponent $z = 2$ [see Fig. 3; Eq. 2] and dynamic scaling [Fig. 3(b)] – we move forward to comprehensively elucidate aspects that remain either controversial or not experimentally assessed yet for 2d Model A. Main results are summarized in Tables I and II.

From the spatial correlator at the Porod’s regime, we measured the universal amplitude $A^*$ (see Table I) in sharp agreement with the Bray-Humayun theory [69]. Beyond the Porod’s regime, the correlator form is well captured by both the Otha-Jasnow-Kawasaki (OJK) theory [51] and the theory of unstable growth (TUG) own to Mazenko [52, 53] [Fig. 6(b)]. Such conclusion emerges from comparisons done with no free parameters: the constant $D_{OJK}$ was found from its direct relation with $D_{AC}$ [51], which we measured independently from the “bub-

ble” experiments; $D_{TUG}$ was determined from the func-

tional form in the Porod’s regime. We have seen that although the agreement between experiment and theory is more pronounced for OJK (considering the most prob-

TABLE I. Universal quantities measured in the TNLC experiment

| Quantity          | Experiment (this work) | Theory | References |
|-------------------|------------------------|--------|------------|
| $1/z$             | 0.505(15), 0.498(6)$^a$| Fig. 3, Eq. (2) | 1/2 | exact predictions [3, 4, 10] |
| $A^*$             | 0.99(7)                | Eq. (14) | 1 | BH prediction [26] |
| $\lambda$         | 1.03(5) for short times ($2 \leq y \leq 10$) | Fig. 7(a), Eq. (25) | 1 | OJK [29] |
|                   | 1.28(11) for late times ($15 \leq y \leq 75$) | $5/4 = 1.25$ | ≈ 1.2887 | FH [20] |
| $\theta$          | 0.196(3), 0.196(4)$^b$| Fig. 8, Eq. (27) | ≈ 0.195, 0.199(2)$^c$ | 2dIM simulations [98] |
| $d_f$             | 1.65(3), 1.613(14), 1.609(9)$^d$ | Fig. 10 | 1.61(5) | 2dIM simulations [101] |

$^a$ Both the estimates were obtained from the decay of the interface density, $\rho(t) \sim t^{1/z}$. The time averaged effective exponent and the least-square fit of the power law give 0.505(15) and 0.498(6), respectively. Please see Eq. (2) and text for detail.

$^b$ The estimate $\theta = 0.196(3)$ was obtained from the time average of the effective exponent; 0.196(4) from the least-square fit of the power law. See Eq. (27) and text for detail.

$^c$ The two values $\theta \approx 0.195$ and 0.199(2), from Ref. [98], were obtained from simulations with free and periodic boundary conditions, respectively.

$^d$ The estimate $d_f = 1.65(3)$ was obtained directly from Eq. (28) by using data at $t = 1365$ s, which is however far from the interval where $M(t)$ is constant and far from the core of the $z = 2$ scaling regime. The other values $d_f = 1.613(14)$ and 1.609(9) are our final estimates obtained through the scaling relation [32]. Please see Eq. (33) and text for detail.

TABLE II. Universal scaling functions measured in the TNLC experiment and compared with theoretical predictions.

| Scaling function | Experimental result | Conclusion |
|------------------|---------------------|------------|
| Spatial correlator $F(g)$ | Fig. 6 | consistent with both OJK and TUG predictions |
| Temporal correlator $H(y)$ | Fig. 7(b) | best described by LSI-2dIM prediction for $y \geq 2$ |

was determined from the func-

tional form in the Porod’s regime.
able values for $D_{OJK}$ and $D_{TUG}$). TUG also may account for the TNLC correlator if the lower limit for $D_{TUG}$, set by its uncertainty, is used [Fig. 3(b)]. To our knowledge, this is the first experimental evidence [135] that theoretical spatial correlators, such as OJK and TUG, describe experimental data in the nontrivial scale beyond the Forod’s regime, for a curvature-driven ($z=2$) system.

Regarding the time correlations, we measured the two-time correlator after proposing its due modification to account for the weak asymmetry in the kinetics [Eq. (24)]. These correlators, $C(t, t_0)$, collapse on a master curve when rescaled by $y = t/t_0$, thus offering evidence of dynamic scaling hypothesis for 2d Model A systems as demanded by aging phenomenology [69] – see Fig. 7. The exponent $\lambda$ (Eq. (19)) displays a crossover from an initial OJK-like value to an asymptotic one which is compatible with both the FH conjecture and the TUG prediction (values in Table I); this reinforces the difficult matter, witnessed by simulations, of distinguishing between the FH and TUG predictions. Light is shed by comparisons with the functional form itself, which elucidate that neither OJK or TUG theory can account for the experimental, 2d Model A, correlation function. Instead, the local scale invariance (LSI) theory own to Henkel [82], that describes the 2dIM correlator along with the FH conjecture [129], provides the best description for the experiment in the regime, $t/t_0 \gtrsim 2$, that LSI deals with [Fig. 7(b)]. These results might indicate support to the FH conjecture from a global perspective, in harmony with many numerical studies [22, 70, 72, 73, 129]. The agreement with LSI also raises up the question whether the paradigmatic 2d Model A systems may display conformal invariance in some stage of their kinetics [82].

We have also assessed the local persistence statistics. Thanks to the accuracy of our data, we measure a local persistence exponent $\theta$ (Fig. 5, Table I) in accordance with the recent estimates from 2dIM [88], but in significant disagreement with the candidate 2d Model A value obtained from the diffusion equation [99] and the OJK theory [82]. This experimentally-based observation thus points out the former as the genuine value in the 2d Model A class. Additionally, we have observed that the mosaic of persistence clusters, defined by the persistence index $\chi(r, t)$, develops a fractal morphology up to the scale of a correlation length $\xi(t) \sim t^{1/2}$ (see Fig. 9 and Fig. 10). The fractal dimension $d_f \approx 1.61$ of such morphology is measured experimentally (Table I) and substantiated as universal [101]. Correlations among persistence clusters respect the dynamic scaling hypothesis [Fig. 10(c)].

Having a versatile experimental model system in the Allen-Cahn (Model A) class opens novel paths for experimental investigation. The many interesting electro-conductive patterns in the liquid crystals [109–111], for instance, can be used to assess unusual sorts of ordering kinetics by transiting between nonequilibrium states, with the advantage of the genuinely sudden electrical switching. Moreover, the independent control on the temperature allows one to tune thermal effects by adjusting the distance from the nematic-isotropic transition point [108]. Such control can be particularly important for experimentally assessing the fundamental theoretical hypothesis that, from the viewpoint of renormalization group [4–5–44], the bath temperature merely factorizes non-universal amplitudes, while it preserves universal features – scaling exponents and correlation functions – for a subcritical dynamics. Another appealing direction is suggested by the numerically observed [44] self-tuning of 2d Model A systems to the critical percolation fixed point [137–139], which theoretically has boosted derivations of exact solutions for the distributions of hull areas [44–135], perimeters [140], domain crossing probabilities [141], besides proposing a generalization of the dynamic scaling hypothesis [142–143]. Access to response functions [73–52] is an additional and attractive avenue that calls for, and it is in principle amenable to, experimentation.

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$$\phi(x) = \frac{1}{2} \left( \sin \left( \frac{\pi x}{L} \right) + 1 \right),$$

for $0 \leq x \leq L$. The value of $\phi(x)$ at the midpoint $x = L/2$ is $0$, which is the equilibrium condition for the TDGL equation, with $\phi(x)$ playing the role of the coordinate normal to the interface.

[122] More specifically, the relation between the continuous order parameter field $\phi$ and the auxiliary field $\mu$ is obtained by requiring $d\phi/d\mu = \mu^{-1} \frac{dV(\phi)}{d\phi}$, where $V(\phi)$ is a symmetric double-well potential, with the boundary conditions $\lim_{\phi \to \pm \infty} \phi(\mu) = \pm 1$ and $\phi(0) = 0$. This is the equilibrium condition for the TDGL equation, with $\mu$ playing the role of the coordinate normal to the interface.

[123] Equation (16) admits two linearly independent solutions $\phi_g \gg 1$, one with a Gaussian decay and the other with an algebraic one. The general solution is a linear combination of both: $\phi_{\text{TUG}}(\phi_g \gg 1) = F_0 g^{1-\frac{d}{2}} \exp \left( -\frac{F_1 g^{1-\frac{d}{2}}}{2} \right)$, where $F_0$ and $F_1$ are constants [53]. The power-law solution in TUG should be guaranteed to maintain the integrability of the associated power-spectrum. The value of $\mu(\phi)$ in TUG is then given in such a way that $\phi_{\text{TUG}}(\phi_g \gg 1) \geq 1$ yields a solution with $F_1 = 0$.

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Interestingly, OJK-like form was also seen to describe the spatial correlations of polycrystals formed by colloids and whose overall kinetics do not follow $z = 2$ \cite{155}. It is also the first time that the TUG two-point spatial correlator [see Eq. (16)] is confronted with experiment after Prof. Mazenko called for experimental tests in \cite{54}.

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