Nucleation-induced transition to collective motion in active systems

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(Received 30 April 2012; revised manuscript received 30 July 2012; published 26 September 2012)

While the existence of polar ordered states in active systems is well established, the dynamics of the self-assembly processes are still elusive. We study a lattice gas model of self-propelled elongated particles interacting through excluded volume and alignment interactions, which shows a phase transition from an isotropic to a polar ordered state. By analyzing the ordering process we find that the transition is driven by the formation of a critical nucleation cluster and a subsequent coarsening process. Moreover, the time to establish a polar ordered state shows a power-law divergence.

Understanding collective motion in driven or self-propelled particle systems is a topic of recent interdisciplinary interest [1–3]. Coherently moving groups have been observed over a broad range of scales, spanning from micrometer-sized systems [4–10] over millimeter-large granules [11–13] to large cooperative animal groups [14]. The ubiquity of this phenomenon raises the questions of how these coherently moving and ordered clusters arise. Coarse-grained particle-based models [15–22] have shown that interactions favoring the alignment of the particles’ direction of motion is sufficient for the emergence of large scale order. As possible origins for the alignment, excluded volume interactions [23], dissipative collisions [24], and hydrodynamic interactions [25,26] have been discussed. Complementary to these coarse-grained agent-based models, hydrodynamic models have been derived from mesoscopic collision rules [27–31], or by means of microscopic interactions [32–36]. These approaches allow to determine the kinetic coefficients, and thereby to analyze pattern-forming instabilities. Furthermore, hydrodynamic equations based on symmetry arguments [37–42] were derived, enabling to show that collections of self-propelled particles exhibit a true, long-range ordered, spontaneously broken symmetry state. In addition, scaling exponents have been calculated analytically [37,38], and validated by agent-based simulations [39]. Moreover, these studies revealed the existence of giant number fluctuations and long-lived density correlations, which have also been found experimentally [10,11,43].

While previous approaches on propelled particle systems focused on the long-time dynamics, the assembly processes leading to collective motion remain elusive. To close this gap, we present an agent-based model of (self-)propelled particles moving at constant speed performing a persistent random walk.

Filaments of length \( L \) move on a hexagonal lattice with constant speed performing a persistent random walk.

shows a power-law divergence upon approaching the critical packing fraction.

To follow the self-assembly processes over long-time periods and for a large number of particles we consider a lattice gas automaton; cf. Fig. 1. Each of the \( N \) particles is taken as a filament of length \( L \) moving at constant speed \( v \), and with a direction performing a persistent random walk. To emulate a spatially isotropic system as close as possible we choose a hexagonal lattice, and use a simulation box of hexagonal shape to avoid possible artifacts due to symmetry breaking with the local hexagonal lattice structure; all lengths are measured in units of the lattice constant. The time scale is fixed by allowing each filament’s head to move one lattice unit per time step. The particles’ dynamics is assumed to be fully determined by its head, while the tail strictly follows the head’s trail. The persistent random walk is implemented by a stochastic process, where each filament’s head \( x_0 \) moves to one of its five neighboring sites \( x_q \) according to a fixed set of conditional probabilities \( P_{\text{prev}}(x_q|x_0) \). For a fixed filament length \( L = 10 \) we choose the conditional probabilities such that each filament performs a persistent random walk with constant unit speed and a kinetic persistence length \( \ell_p^0 = 8.7 \) comparable to \( L \); for more details see the Supplemental Material [44].

Our model includes an effective excluded volume interaction and a local alignment field. Both enter into the model by supplementing the probabilities for the unperturbed persistent random walk by appropriate factors. In contrast to recent off-lattice simulations of overdamped rods, solely interacting by means of excluded volume [23], this allows us to separately...
tune excluded volume and alignment interactions and thereby study their interplay. For a lattice site that is already \( k \)-times occupied, the probability for further occupation is reduced by a Boltzmann factor \( e^{-\epsilon x} \), where \( \epsilon \) characterizes the penalty for multiple occupations. Formally, for \( \epsilon \rightarrow \infty \), the limit of strict excluded volume is obtained [34]. The limit of weak penalties for multiple occupations (low \( \epsilon \)) is appropriate for motility assay experiments, where filament crossings occur frequently [5,6,8,9]. Moreover, these experiments also indicate that there is a local alignment interaction between filaments [5,6,8,9].

To emulate such an interaction, each particle is assigned an alignment field \( u(x) \) at its occupied and neighboring lattice sites (see the arrows in Fig. 1); it is directed along the particles’ contour, and is of unit length. Overlapping alignment fields of different particles are averaged. In a collision event the alignment field modifies the transition probability to move from \( x_0 \) to \( x \) by the Boltzmann factor \( e^{\alpha \phi} \), where \( \phi \) is the relative angle between the alignment field and the direction of motion of the respective collision partner, \( \phi = \angle[\mu(x),x-x_0] \). The parameter \( \alpha > 1 \) characterizes the strength of the polarization, where stronger interactions correspond to increasing values of \( \alpha \). Assuming that all these contributions to the filament’s dynamics are statistically independent, we arrive at the following update rule: Given a configuration of \( N \) filaments, we use random sequential updating and move a chosen head from position \( x_0 \) to a target position \( x \) with probability \( P(x|x_0,u) \propto e^{-\epsilon P_\text{pen}(x|x_0)}\cdot e^{-\phi \alpha P_{\text{rots}}(x|x_0)} \).

We have performed extensive numerical simulations for a broad range of reduced densities \( \rho = NL/A \), where \( A \) is the area of the simulation hexagon. For low densities an initially statistically homogeneous distribution of filaments remains isotropic both in filament position and orientation. However, for densities above some threshold, an initially disordered state evolves into a state with small coherently moving clusters that spread perpendicular to their direction of motion and form bands [see Figs. 2(a)–2(c), and videos in the Supplemental Material [44]]. The clusters move coherently embedded in an isotropic background of randomly oriented particles. These observations resemble similar results found previously in Vicsek-like models [17] and other agent-based lattice gas models [16,21].

The pattern formation process can readily be quantified by introducing the time-dependent mean polarization \( \mathcal{P}(t) \), i.e., the magnitude of the polarization averaged over all single-particle polarizations \( n_i(t) \) contained in a region of interest (ROI) or the entire simulation box (SIM), respectively. The polarization of the \( i \)th filament is defined as the difference vector of the position of the filament’s head between two successive time steps, \( n_i(t) := x_i(t) - x_i(t-1) \). These two measures of polarity provide complementary information about the pattern-forming process; cf. Fig. 2(d). \( \mathcal{P}_{\text{ROI}} \) illustrates the formation of small polarized clusters that develop into bands: The sequence of spikes in the time traces indicate polar-moving clusters passing through the ROI. In contrast, \( \mathcal{P}_{\text{SIM}} \) characterizes the evolution of global order in the system: The mean value slowly grows in time and fluctuations are much less pronounced. These latter fluctuations originate from changes in relative polarization of just a few large polar-moving structures with time.

The temporal evolution of coherent motion can be accessed by quantifying the directional correlations in the filament’s comoving frame by means of the Lagrangian orientational correlation function

\[
C(\tau;t) := \frac{\sum_i n_i(t) \cdot n_i(t+\tau) - \left[ \sum_i n_i(t) \right]^2}{\sum_i n_i^2(t) - \left[ \sum_i n_i(t) \right]^2}. \tag{1}
\]

The sum extends over all particles in the system, and \( t \) denotes the waiting time passed since the preparation of the system in a state with a uniform distribution in both position and orientation. Starting from a disordered state at \( t = 0 \), the directional correlation function decays exponentially in time \( \tau \) with a decay length corresponding to the persistence length of the undisturbed persistent random walk \( \ell_p^0 \). As we increase the waiting time \( t \) a clear shoulder in \( C(\tau;t) \) builds up until it finally becomes stationary; for the parameters used in Fig. 2(e) this occurs at \( t \approx 5000 \). Then, in the stationary regime, there are clearly two decay processes with well-separated time scales: an initial fast decay and an extended time window with a significantly slower but again exponential decay. This indicates a phase separation where a fraction of the filaments is in a low-density isotropic phase and the remainder is organized in coherently moving polar clusters. Filaments in the low-density phase are responsible for the fast initial decay which has the same slope as the initial decay starting from an initially fully disordered filament configuration. In contrast, orientational correlations of filaments in clusters are significantly enhanced since they preferentially move into the same direction as their neighbors due to the impact of the alignment field. This results in an increased decorrelation time of \( C \). Since filaments are moving at unit velocity this time scale can also be

![FIG. 2.](image-url)
interacted as a persistence length of the trajectories traced out by the filaments, $\ell_{\text{coll}}$. We attribute the loss of correlations for filaments in clusters mainly to reorientations of the clusters as a whole and to a lesser extent to collisions between clusters. The latter and collisions of clusters with the confining boundary of the simulation box are responsible for the steep drop in the correlation function at very large times. The collective persistence length of the clusters $\ell_{\text{coll}}$ depends on the density $\rho$ as well as the strength of excluded volume interaction $\epsilon$; see the Supplemental Material [44].

To further analyze the self-assembly process and to characterize the ensuing stationary state we consider the fraction of occupied lattice sites $\Gamma$, or equivalently the free volume fraction $\Phi := \rho - \Gamma$. In the absence of polar alignment processes between the filaments, we find that $\Gamma$ fluctuates around a constant value depending on the strength of the excluded volume interaction $\epsilon$ [see Fig. 3(a), inset]. For a noninteracting system ($\epsilon = 0, \alpha = 1$), the stationary value of $\Gamma$ as a function of reduced density $\rho$ is given by $\Gamma_{\text{hom}}(\rho) = 1 - e^{-\rho}$; see Fig. 3(b). This result has previously been obtained in continuum percolation theory for the fraction of lattice sites occupied by a statistically homogeneous distribution of overlapping rods [45]. In the presence of excluded volume interaction, the actual value of $\Gamma$ is only slightly smaller than $\Gamma_{\text{hom}}$ and shows the same functional form. Adding polar alignment interactions leads to qualitatively different behavior. As can be inferred from Fig. 3(b), there is now a threshold density $\rho_c$, where the free volume gain $\Phi$ jumps to a value much larger than the corresponding value for a noninteracting system. Moreover, measuring the polar order parameters it turns out that this jump actually coincides with the onset of polar order; cf. Fig. 2(e). The jump in $\Phi$ may, therefore, be taken as a signature to map out the phase diagram as shown in Fig. 3(c); for a more detailed discussion see the Supplemental Material [44]. It also clearly indicates that filaments must have formed some clusters much denser than expected for a purely statistical overlap of filaments. Figure 3(a) shows that clusters form by a nucleation process: There is a lag phase during which $\Gamma$ stays high close to a value obtained in the absence of polar alignment processes. Subsequently, there is a time window $[\Gamma_{\text{nuc}}, \Gamma_{\text{stat}}]$, where the available free volume fraction $\rho - \Gamma$ increases towards a higher stationary value; cf. Fig. 3(b). The time to reach the final polar steady state diverges as a power law $\Gamma_{\text{stat}} \propto (\rho - \rho_c)^{-\zeta}$ with $\zeta \approx 1$.

To further scrutinize the nucleation processes, we analyzed the cluster mass distribution $P(m)$ and the size of the critical nucleation cluster as a function of density; cf. Figs. 4(b) and 4(c). Here clusters are defined as connected areas on the lattice with local occupation numbers $k \geq k_c = 5$; see the Supplemental Material [44]. For an illustration, Fig. 4(a) shows time traces for the maximum of the cluster mass distribution $m_{\text{max}}$, the mean $\langle m \rangle$, and standard deviation $\delta m = \sqrt{\langle m^2 \rangle} - \langle m \rangle^2$ close to the phase transition from an isotropic to an ordered state. We observe that during the lag phase clusters are continually created and destroyed such that all cluster characteristics remain constant on average and there is no gain in available free volume. The corresponding distribution of cluster sizes $P(m)$ shows power-law tails with an exponent close to $\delta = 5$; cf. Fig. 4(b). For the representative trace shown in Fig. 4(a), there is a spontaneous nucleation of a large cluster at $t \approx 2800$ that, albeit it partially disassembles, still seems to trigger the transition to cluster formation. It defines the onset of a steady increase in $\langle m \rangle$ and also a change in the power law of the cluster size distribution; cf. Fig. 4(b). In conclusion, close to the critical point the system is waiting for a large enough density fluctuation that acts as a nucleus for cluster formation, which in turn triggers the transition to the polar ordered state; as shown above, the waiting time diverges upon approaching the transition; cf. Fig. 3(b).

To quantify the size of the critical nucleation cluster and further substantiate that the nucleation phenomenon is the mechanism driving the transition, we have performed a detailed analysis of the largest cluster formed during the nucleation process and thereby quantified the heuristic observations made in Fig. 4(a). In the isotropic phase, one expects that the largest clusters formed over the whole observation window $M_{\text{max}} := \max_t m_{\text{max}}(t)$ are typically too small to act as a nucleus for the emergence of polar order. However, in a finite system, there will still be rare occasions where fluctuations are strong enough to form a critical nucleus. The fraction $S$...
FIG. 4. (Color online) (a) Time traces of \( m_{\text{max}} \) (red solid), \( \langle m \rangle \) (green dashed), and \( \delta m \) (blue dotted) of the \( P(m) \) for \( \rho = 0.6 \) during the lag phase and in the stationary regime for the polar ordered state sampled over more than \( 10^4 \) realizations. For reference, power-law distributions \( P(m) \propto m^{-\delta} \) with \( \delta = 4.5 \) are indicated by dashed lines. (c) Mean values of the largest cluster, \( \langle M_{\text{max}} \rangle \) (red squares) and \( \langle M_{\text{max}} \rangle \) (blue dots), serve as lower and upper bounds for critical nucleation cluster mass. In the transition window (shaded gray area) from the ISO to the PO phase, the fraction of realizations to 1. Parameters for all graphs are \( \alpha = 5 \) and \( \epsilon = 0.5 \).

of realizations which develop polar order sharply increases as the density is raised throughout the transition window [see Fig. 4(c)], with the critical density \( \rho_c \approx 0.4 \). Below the critical density, the values of \( M_{\text{max}} \) for those realizations which did not lead to polar order are a measure of cluster sizes which are too small to trigger the pattern-forming process, and can thereby be used as lower bounds for the size of the critical nucleus. Figure 4(c) shows the average value of \( M_{\text{max}} \) for those realizations which did not lead to polar order. We observe that this quantity increases upon approaching the critical density from below. Above the critical density, most realizations will develop polar order and one may estimate the critical nucleus size as the average size of the largest cluster at the nucleation time \( M_{\text{nuc}} := m_{\text{max}}(t = T_{\text{nuc}}) \). The corresponding average value \( \langle M_{\text{nuc}} \rangle \) gives the typical cluster mass that has been large enough to trigger a transition to polar order, and hence serves as an upper bound for the critical nucleus.

While previous numerical studies on propelled particle systems focused on the long-time dynamics, we investigated the dynamics of the self-assembly process leading to the emergence of polar ordered states. We find that the ordering process is driven by the formation of a critical nucleation cluster: Once a cluster of sufficiently large mass has assembled, the entire system builds up collective motion. This is reflected in the existence of a lag time, where the system does not succeed to assemble a cluster of sufficient mass. The signatures of the ensuing phase transitions are a jump in the free volume and a power-law divergence in the characteristic time for the formation of the polar ordered state. We suggest that exploring the principles underlying the self-assembly process of polar clusters is a promising route to understand ordering phenomena in a broader class of active systems.

The authors would like to thank Fernando Peruani for helpful discussions. Financial support from the DFG in the framework of the SFB 863 and the German Excellence Initiative via the program Nanosystems Initiative Munich (NIM) is gratefully acknowledged.
NUCLEATION-INDUCED TRANSITION TO COLLECTIVE ... Physical Review E 86, 030901(R) (2012)