Self-similar dynamics of order parameter fluctuations in pump-probe experiments

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Upon excitation by a laser pulse, broken-symmetry phases of a wide variety of solids demonstrate similar order parameter dynamics characterized by a dramatic slowing down of relaxation for stronger pump fluences. Motivated by this recurrent phenomenology, we develop a simple nonperturbative effective model for photoinduced dynamics of collective bosonic excitations. We find that as the system recovers after photoexcitation, it shows universal prethermalized dynamics manifesting a power-law, as opposed to exponential, relaxation, explaining the slowing down of the recovery process. For strong quenches, long-wavelength overpopulated transverse modes dominate the long-time dynamics; their distribution function exhibits universal scaling in time and space, whose universal exponents can be computed analytically. Our model offers a unifying description of order parameter fluctuations in a regime far from equilibrium, and our predictions can be tested with available time-resolved techniques.

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I. INTRODUCTION

The concept of universality plays a central role in the theory of equilibrium phase transitions because it allows us to reduce a plethora of experimentally studied systems to a few fundamental classes [1]. For systems far from equilibrium, the notion of universality [44] is relatively unexplored and has recently emerged as an active field [2–10], partially motivated by recent progress in ultracold-atom [11–14] and ultrafast pump-probe experiments [15]. In a nonequilibrium context, one of the dramatic manifestations of universality is the emergence of the self-similar evolution of correlation functions [16,53]. In particular, after a strong perturbation, the transient equal-time two-point correlation function $D(|x - y|, t)$ might depend only on a single evolving length scale $\xi(t)$ and two universal functions:

$$D(|x - y|, t) = g(t)f(|x - y|/\xi(t)).$$

(1)

Functional forms of $f(x)$ and $g(t)$ depend neither on microscopic parameters nor on initial conditions. Typical equations of motion, often a complex system of partial integro-differential equations, represent an interplay between many degrees of freedom such as quasiparticles, order parameter (OP), phonons, and/or magnons. If these equations allow for the above self-similar form, the analysis might reduce to just a few differential equations, which is particularly appealing since it eases the interpretation of the involved evolution. From a physical standpoint, the self-similarity suggests that there exists a stabilization-like mechanism responsible for this form.

Recent pump-probe experiments observed several recurrent phenomena that hint at the existence of universality in the out-of-equilibrium context. In particular, common features have been observed in the dynamics of the OP and low-energy collective excitations in charge-density-wave (CDW) compounds [15,17–27], putative excitonic insulators [28,29], magnetically ordered systems [30,31], and systems that exhibit several intertwined orders [32,33]. Common phenomenology in these materials includes the following: (i) The recovery of a photosuppressed OP takes longer at stronger pump-pulse fluence; (ii) the amplitude of the OP restores faster than the phase, exhibiting a separation of timescales; (iii) related to (ii), peaks in diffraction experiments remain broadened compared to equilibrium shape long after photoexcitation, showing prolonged suppression of long-range phase coherence. For example, all of these features have been observed in the same set of experiments [19] on LaTe$_3$, an incommensurate CDW material. Below, for concreteness, we primarily focus our otherwise generic theoretical formalism on this material, but we expect that our conclusions should qualitatively apply to other systems as well. We note, however, that quantitative results may be different depending on the details of specific systems. For example, in the case of ferromagnetic systems, our analysis should be modified to include the conserved character of the OP [44].

A common approach to describing many-body dynamics in symmetry-broken states is based on the so-called three-temperature model (3TM) [34–38]. In this framework, a nonequilibrium state is characterized by assigning different temperatures to different subsystems, such as electrons, phonons, and OP degrees of freedom [39]. Upon photoexcitation, most incoming light is absorbed by electrons, instantaneously increasing the electronic temperature, $T_e$. The introduction of $T_e(t)$ is justified provided we are only interested in phononic timescales sufficiently exceeding the fast electron-electron scattering time. Subsequent dynamics corresponds to energy exchange between hot electrons and the other two subsystems. In this process, it is often assumed

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II. THEORETICAL FRAMEWORK

Spontaneous symmetry breaking (SSB) is described by the time-dependent Landau-Ginzburg formalism (model A in Ref. [44]):

$$\frac{d\phi_\alpha(x,t)}{dt} = -\Gamma \frac{\delta F}{\delta \phi_\alpha(x,t)} + \eta_\alpha(x,t).$$  \hspace{1cm} (2)

Here \(\phi_\alpha\) is an \(N\)-component vector of real fields representing the OP. The free-energy functional reads

$$F[\phi] = \int d^3x \left[ \frac{r}{2} \phi_\alpha^2 + \frac{K}{2} (\nabla \phi_\alpha)^2 + u(\phi_\alpha^2)^2 \right].$$  \hspace{1cm} (3)

and \(\eta_\alpha\) represents the noise originating from the phononic bath at temperature \(T\):

$$\langle \eta_\alpha(x,t)\eta_\beta(x',t') \rangle = 2T \Gamma \delta_{\alpha,\beta} \delta(x-x')\delta(t-t').$$  \hspace{1cm} (4)

Here \(r, K, u,\) and \(\Gamma\) are the model parameters. For homogeneous quenches, without loss of generality, we assume that SSB occurs along the first direction: \(\phi(t) = \langle \phi_1(x,t) \rangle\). Associated with the OP are longitudinal (Higgs modes) \(D^\parallel_k(t) \equiv \langle \phi_1(k;t)\phi_1(-\mathbf{k};t) \rangle\) and transverse (Goldstone modes) \(D^\perp_k(t) \equiv \langle \phi_{\mathbf{a} \neq 1}(k;t)\phi_{\mathbf{a} \neq 1}(-\mathbf{k};t) \rangle\) correlation functions. The model-A formalism in Eqs. (2)–(4) can be conveniently rewritten in terms of the Fokker-Planck equation:

$$\partial_t \mathcal{P} = T\Gamma \sum_{\mathbf{k},a} \frac{\delta}{\delta \phi_{a,k}} \left[ \mathcal{P} \frac{\delta F}{\delta \phi_{a,-\mathbf{k}}} + \frac{\delta \mathcal{P}}{\delta \phi_{a,-\mathbf{k}}} \right].$$  \hspace{1cm} (5)

where \(\mathcal{P}((\phi, t)\rangle\) is the probability distribution functional of space-dependent field configurations \(\phi_\alpha(x)\). To leading order in \(1/N, \mathcal{P}((\phi, t)\rangle\) is Gaussian, implying that the OP \(\phi(t)\) and the correlators \(D^\parallel_k(t), D^\perp_k(t)\) form a closed set of dynamical variables. The self-consistent equations of motion read [45,46] (see Appendix A for a derivation)

$$\frac{d\phi(t)}{dt} = -\Gamma r_{\text{eff}} \phi,$$  \hspace{1cm} (6)

$$\frac{dD^\parallel_k(t)}{dt} = 2T\Gamma - 2\Gamma (Kk^2 + r_{\text{eff}})D^\parallel_k,$$  \hspace{1cm} (7)

$$\frac{dD^\perp_k(t)}{dt} = 2T\Gamma - 2\Gamma (Kk^2 + r_{\text{eff}} + 8u\phi^2)^2D^\perp_k.$$  \hspace{1cm} (8)

Here the self-consistent “mass” term is defined as

$$r_{\text{eff}}(t) = r(t) + 4u(\phi^2)^2 + \frac{n_{\text{tot}}}{\Lambda} + (N - 1)n_{\text{hot}},$$  \hspace{1cm} (9)

where \(n_{\text{tot}}^{(l)} \equiv \int d^k \mathbf{q} D^{(l)}_q\), and \(\Lambda\) is the UV cutoff. Note that quantities such as energy or total number of excitations are not conserved due to the external bath.

The bath, cf. Eq. (4), will also always result in the thermalization of the system, in contrast to quenches in the isolated \(O(N)\) model, where, to leading in \(1/N\) order, the system does not demonstrate equilibration [47–51]. In that case, thermalization occurs only after subleading corrections are taken into consideration [52]. As such, the presence of the external bath in our case motivates us to disregard these subleading corrections here.
From the equations of motion, we obtain the equilibrium correlators:

\[ D_k^{\parallel} = \frac{T}{K k^2 + 8 \mu \phi^2 + r_{\text{eff}}} , \quad D_k^{\perp} = \frac{T}{K k^2 + r_{\text{eff}}} . \]  

(10)

This result is a manifestation of the equipartition theorem. In the symmetry-broken phase, where \( r_{\text{eff}} = 0 \) and \( \phi \neq 0 \), we observe that the OP equilibrium value \( \phi \) is affected by the thermal fluctuations; cf. Eq. (9). The transverse correlation length \( \xi_{\perp} \propto r_{\text{eff}}^{-\frac{1}{2}} \) is divergent. In the disordered phase, \( r_{\text{eff}} \neq 0 \) and \( \phi = 0 \), the transverse and longitudinal correlations are not distinguishable.

A useful point of view on the above approximations is as follows. The equations of motion (6)–(9) are equivalent to

\[ \frac{d\delta \phi_k^\parallel(t)}{dt} = -\Gamma (K k^2 + r_{\text{eff}}) \phi_k^\parallel + \eta_k^\parallel(t), \]  

(11)

\[ \frac{d\delta \phi_k^\perp(t)}{dt} = -\Gamma (K k^2 + r_{\text{eff}} + 8 \mu \phi^2) \phi_k^\perp + \eta_k^\perp(t), \]  

(12)

where \( \delta \phi_k^\parallel \) represents the fluctuating part of the corresponding Fourier mode \( \phi_k^\parallel \). We observe that each of the fluctuating modes lives in an effectively parabolic potential, \( \delta \phi_k^\parallel = 0 \), and the noise term establishes the equilibrium variances given by Eq. (10).

We now formulate the quenching protocol. For simplicity, we assume that the electronic temperature \( T_e \) cools down to the equilibrium value \( T \) with a constant rate \( \tau_{\text{QP}} \) defined by the electron-phonon coupling. This assumption seems to be not too crude for LaTe\(_4\), where the Fermi surface is only partially gapped, and the electrons excited by the laser pulse can relax through a gapless channel [19,37]. For a different situation, our framework should be straightforwardly extended. In the usual Landau-Ginzburg theory, the coefficient \( r(T_e) \) depends linearly on \( T_e \). To mimic a photoexcitation event, we therefore impose the following dynamics on \( r(t) \) [Fig. 1(b)]:

\[ r(t) = r_i + \theta(t) \exp (-t/\tau_{\text{QP}})(r_f - r_i), \]  

(13)

where \( \theta(t) \) is the Heaviside theta function, \( r_i \) is the prepulse value chosen such that \( \phi \neq 0 \), and \( (r_f - r_i) \) characterizes the pulse strength. Below we focus on time delays much beyond \( \tau_{\text{QP}} \).

In the next section, we apply the above theoretical framework to investigate the intrinsic dynamics after the arrival of a laser pulse. Our conclusions are not specific to the choice of model parameters, but in the numerical simulations below, we fix them to roughly mimic the experimental results in Ref. [19]. There, the values of \( \Gamma \) and \( \tau_{\text{QP}} \) are expected to be similar.

III. EVOLUTION AFTER PHOTOEXCITATION

Upon photoexcitation, cf. Eq. (13), the system passes through four dynamical stages [Fig. 1(c): (i) depletion, (ii) inflation, (iii) mode decoupling, and (iv) relaxation to the thermal equilibrium. We cover each of them below.

In Fig. 2(a), we show numerical results for the dynamics of the OP, \( \phi(t) \). For a weak pump, \( \phi(t) \) becomes slightly suppressed and then quickly recovers to the initial value \( \phi_0 \). This should be contrasted to the case of a strong pulse, for which initially the OP becomes strongly suppressed and then goes through a long recovery process. The recovery takes longer for stronger pulses [Fig. 2(b)]. This slowing down is due to the power-law dynamics \( \delta \phi_k(t) \equiv [\phi(t) - \phi_0] \sim t^{-d_{r}} \) with \( d_r = \frac{2}{3} \), which we further discuss in the next section.

In Fig. 2(c), we plot the evolution of \( r_{\text{eff}}(t) \). Upon arrival of a laser pulse, \( r_{\text{eff}}(+0) = (r_f - r_i) \). This large initial value first decreases due to the time evolution of the “bare value” of \( r(t) \), cf. Eq. (13), and later, at \( t \gtrsim \tau_{\text{QP}} \), due to the dynamics of the OP and collective modes described by Eqs. (6)–(8). Even though \( r(t) \) returns to its equilibrium value \( r_i \) during a relatively short time \( \tau_{\text{QP}} \), the dynamics of \( r_{\text{eff}} \) occurs over much longer time scale where it even changes sign [Fig. 2(c)]. We find that long-time evolution of \( r_{\text{eff}} \sim t^{-d_{r}} \) is power-law-like with \( d_r = \frac{2}{3} \). For the fluctuating modes \( \delta \phi_k^\parallel \), a large value of \( r_{\text{eff}} \) implies that each of the effective parabolic potentials becomes initially steeper, and, as such, the noise term in Eq. (4) tends to depopulate these modes [Fig. 2(d)]. Therefore, the first stage—depletion—is characterized by suppression of the OP and correlations \( D_k^{\parallel} \) and \( D_k^{\perp} \).

The second stage— inflation—starts when \( r_{\text{eff}} \) changes its sign. A negative \( r_{\text{eff}} \) implies that each of the effective parabolic potentials becomes shallower, or, as in the case for the low-momenta transverse modes, can even become inverted. Therefore, during the inflation, population in each of the modes proliferates, most dramatically for the low-momenta modes [Fig. 2(d)].
For a strong quench and at the time when the OP becomes completely suppressed, the longitudinal and transverse correlations are no longer distinguishable [Fig. 2(d)]. This parallels the disordered phase in an equilibrium situation. As the OP develops, these modes start to separate. We will associate the end of the inflation stage with the time when $D_{k=0}^\perp$ reaches its maximum value [see the inset in Fig. 2(d), dashed curve].

Because of the additional correction to the quadratic term for the longitudinal correlations in Eq. (8), the subsequent evolution—mode decoupling—is very different for the longitudinal and transverse modes; see Fig. 3. The longitudinal correlations start to relax back to the thermal equilibrium value in Eq. (10), while the transverse modes continue to proliferate, resulting in the exponent $\alpha$, cf. Eq. (15), being positive during the third dynamical stage. Moreover, by the time $n_{\text{tot}}^\parallel$ is sufficiently recovered, $n_{\text{tot}}^\perp$ is about to reach its maximum. Strong experimental evidence of this separation of timescales was reported in Refs. [19,24,25].

Just after the mode decoupling, $n_{\text{tot}}^\perp$ starts to slowly decrease, cf. Eq. (16), suggesting that the system enters the final relaxation stage. Note that even though lowest-momenta modes $D_k^\perp$ continue to proliferate at very long times, their relative contribution to $n_{\text{tot}}^\perp$ is suppressed by the reduced phase space of these modes, which is proportional to $k^2$. The underlying dynamics is reminiscent of an inverse particle cascade in the theory of turbulence [8,53,54]. The main difference is that in our system the dynamics is overdamped.

In the next section, we focus on the last two dynamical stages, where we find that the system exhibits self-similar scalings in time and space, manifesting power-law-like, as opposed to exponential, behavior.

IV. UNIVERSALITY IN THE INTRINSIC DYNAMICS

Our discovery of self-similarity can be summarized in the following equations. The distribution function of the Goldstone modes follows

$$\delta D_k^\perp(t) \simeq \frac{g(t)}{k^2} f(k/k_\perp^*(t)).$$

(14)

where $\delta D_k^\perp(t) \equiv [D_k^\perp(t) - D_{k=0}^\perp]$ and $D_{k=0}^\perp$ is the prepulse equilibrium distribution given by Eq. (10). The form in Eq. (14) is similar to the one in Eq. (1), though written in momentum space; $\xi_k(t)$ [Eq. (15)] represents the emergent time-dependent length scale. We also identify the scaling relations

$$g(t) \sim t^\alpha, \ k_\perp^* \sim t^{-\beta}.$$

(15)

Both power-law exponents $\alpha$, $\beta$ and the function $f(x)$ are universal. We find that $\beta = \frac{1}{2}; \alpha \approx 0.7$ at early times while $\alpha = -1$ in the final relaxation stage. The scaling functions $f(x)$, $k_\perp^*(t)$, and $g(t)$ are shown in Fig. 4. In a model where one retains dissipation but neglects the noise coming from the bath, an exponential behavior is expected instead [55].

We first explore the implications of the self-similarity in Eq. (14) on the experimental phenomenology. Prior to the arrival of the pump pulse, the system possesses long-range coherence manifested in the macroscopic homogeneous OP $\phi$ and divergent transverse correlation length $\xi_\perp = \infty$. The laser pulse depletes this coherence. Equation (14) suggests that as the system evolves toward equilibrium, it develops a finite correlation length $\xi_\perp(t)$ that slowly grows in a diffusive manner [56], $\xi_\perp(t) \sim \sqrt{t}$, consistent with recent experiments [24,25]. This physical picture explains the broadening of diffraction peaks observed long after the arrival of the pulse. The slowing down of the OP recovery can also be deduced from Eq. (14). The system enters the final dynamical stage with $g(t) \simeq A_Q t^{-1}$, where $A_Q$ is a constant of proportionality that increases monotonically with the quench strength. By contrast, as shown in Fig. 4(c), $k_\perp^*(t)$ does not depend on the quench. Therefore, the cumulative effect, expressed in the change of the population of transverse modes $\delta n_{\text{tot}}$,
behaves as
\[
\delta n_{\text{tot}} \equiv \int \frac{d^3k}{(2\pi)^3} \delta D_k^1(t) \sim A_0 t^{-3/2},
\]  
(16)

i.e., as a power-law. Since the transverse modes dominate the long-time dynamics, from Eq. (16) it follows that the characteristic recovery time \( r_{\text{eff}} \sim A_0^{2/3} \) is a monotonically increasing function of the quench strength [Fig. 2(b)].

In the following, we offer a formal derivation of all the long-time power-law exponents: \( \beta = \frac{1}{2}, \alpha = -1, d_\phi = \frac{3}{2}, \) and \( d_r = \frac{5}{2}. \) Reestablishing the long-range coherence, which is depleted by the laser pulse, is the slowest process that happens in the system, \( k_0^* \sim t^{-\beta}. \) Motivated by the numerical results in Figs. 2 and 4, let us assume that \( d_r > 2\beta, \) which will turn out to be consistent with the subsequent derivation. We note that the most relevant transverse modes are the ones with wave vectors close to \( k_0^*; \) cf. Fig. 4. For these modes, we can safely neglect fast \( r_{\text{eff}} \sim t^{-d_r}, \) in Eq. (7) compared to slow \( (k_0^*)^2 \sim t^{-2\beta}, \) resulting in a simple diffusion-like equation with the solution \( \delta D_k^1 = A_k \exp(-2\Gamma k_0^* t), \) where \( A_k \) is yet an unknown function of \( k \). As supported by Fig. 2(d), \( \delta D_k^1(t) \) does not diverge for \( k \to 0. \) One may then Taylor-expand \( A_k \) as \( A_k = A_0 + A_2 k^2 + A_4 k^4 + \cdots \). The relevant \( k \) vectors, the ones in the vicinity of \( k_0^* \), are small at long times, and thus it is safe to leave only the dominant harmonic \( A_0 \) in this expansion, i.e., \( \delta D_k^1 \sim \exp(-2\Gamma k_0^* t), \) consistent with \( \beta = \frac{1}{2} \) and \( \alpha = -1; \) cf. also Fig. 4(b).

To extract the value of \( d_r \), we need to consider the interplay between the OP and transverse correlations (longitudinal correlations are discussed in Appendix B). Assuming that at long times \( r_{\text{eff}} \sim t^{-d_r}, \) the equation of motion (6) reads
\[
\frac{d\phi}{dt} = -\Gamma r_{\text{eff}}(t) \phi \sim t^{-d_r},
\]
where we implied that \( \phi(t) = \phi_0 + \delta\phi(t) \) is already close to its equilibrium value \( \phi_0. \) By integrating the above equation, we obtain \( \phi^2(t) \approx \phi_0^2 + C t^{-d_r}, \) where \( C \) is some constant and \( d_\phi = d_r - 1. \) Note that since \( \phi^2(t) \) enters the definition of \( r_{\text{eff}}(t), \) cf. Eq. (9), the more dominant scaling \( t^{-d_r+1} \) from the OP must be compensated by the transverse correlations. On the other hand, from the previous paragraph, we deduce \( \delta n_{\text{tot}} \sim t^{\alpha - \beta}, \) and therefore
\[
\alpha - \beta = -d_r + 1 \Rightarrow d_r = 1 + \beta - \alpha = \frac{5}{2},
\]  
(17)

This result also gives \( d_\phi = d_r - 1 = \frac{3}{2}. \) The above analysis has explained all long-time scalings. Note, however, that the self-similarity in Eq. (14) settles much earlier than the final relaxation stage. It is striking that the functional form of \( f(x), \) cf. Eq. (14), is the same for the last two dynamical stages (Fig. 4), an interesting feature that warrants further investigation.

It is worth mentioning that there are two well-known scenarios: (i) critical ageing dynamics [57–60] and (ii) phase-ordering kinetics [45,46,56]—where self-similar dynamics is well established in the overdamped \( O(N) \) model. Scenario (i) focuses on a temperature quench specifically to a critical point, which is not applicable here. By contrast, scenario (ii) shares some similarities with the above physical picture. The usual phase-ordering kinetics is concerned with a quench from the disordered phase to a temperature below \( T_c, \) which parallels our quenching protocol, where the OP becomes initially suppressed and then slowly recovers to its equilibrium value. Another resemblance is that the two scenarios share the same exponent \( \beta = \frac{1}{2}. \) However, in the phase-ordering kinetics, the classical field expectation value cannot develop, as trivially follows from Eq. (6), and the origin of the self-similarity is due to the growth of large phase-coherent competing domains. By contrast, the self-similarity in our case is due to the interplay between the nonzero OP \( \phi \) and proliferating fluctuations, resulting in new exponents \( d_\phi = \frac{3}{2} \) and \( d_r = \frac{5}{2}. \)

V. DISCUSSION AND OUTLOOK

This section is devoted to addressing the applicability of our results to real experiments. In Sec. I, we motivated our study with recurring phenomenology observed in a wide variety of materials. We believe that our framework describes the underlying physics of these systems qualitatively; however, due to several features specific to each of these materials, it may alter the quantitative description. Here are some sources for potential quantitative disagreement. First, our theory is based on \((3+1)\) dimension, but most of those materials are either quasi-one-dimensional, such as \( \text{Ko}_3\text{MoO}_4 \), or quasi-two-dimensional, including rare-earth tritellurides [18–22], \( \text{TiSe}_2 \), and cuprates [15]. Second, in multiple cases, such as \( \text{Ni}_2\text{MnGa} \), there is some constant and \( d_\phi = d_r - 1. \) Note that since \( \phi^2(t) \) enters the definition of \( r_{\text{eff}}(t), \) cf. Eq. (9), the more dominant scaling \( t^{-d_r+1} \) from the OP must be compensated by the transverse correlations. On the other hand, from the previous paragraph, we deduce \( \delta n_{\text{tot}} \sim t^{\alpha - \beta}, \) and therefore
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The initial conditions also seem to be very different: In the phase-ordering case, one often starts from high temperatures with profound thermal fluctuations, cf. Eq. (10); here, at the time when the OP becomes suppressed, fluctuations become depleted instead of proliferated.

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Our work suggests that behind this common phenomenology is a simple, intuitive interpretation where slow over-populated Goldstone modes dominate the system evolution after photoexcitation. We make a few observations to address the complications noted in the previous paragraph. First, in quasi-1D or -2D systems, fluctuations are expected to be much more significant compared to 3D. Hence, we anticipate a similar, if not more pronounced, proliferation of Goldstone modes. The strong spatial anisotropy can lead, though, to a different scaling exponent, as was recently observed in Ref. [15]. Second, one can straightforwardly extend the present framework to account for several orders [61,62], and it would be intriguing to see how enhanced fluctuations enter the interplay between different orders. Third, in ferromagnetic systems, conservations of the OP should lead to additional slowing-down of the long-wavelength excitations. We leave the discussion of the conserved OP dynamics for future work. Fourth, coherent dynamics [63,64], such as Higgs-like oscillations, is not expected to dictate the evolution for a strong pulse because these oscillations are expected to be overdamped. From a practical point of view, the model-A dynamics should also be a good
approximation to describe experiments that have an insufficient time resolution to detect oscillatory behavior. On the experimental side, it is essential to verify our interpretation.

Finally, we note that a variety of time-resolved experiments could be performed to test our predictions. Examples include electron or x-ray diffuse scattering [65–67], resonant inelastic x-ray scattering [15], and Brillouin scattering [68]. These experiments give access to momentum- and/or energy-resolved dynamics of bosonic excitations related to OP, so one may specifically search for signatures of (i) nonthermal resolved dynamics of bosonic excitations related to OP, so these experiments give access to momentum- and/or energy-resolved dynamics of bosonic excitations related to OP, so one may specifically search for signatures of (i) nonthermal population of the transverse modes, (ii) the self-similarity encoded in Eq. (14), and (iii) different dynamical stages after photoexcitation [Fig. 1(c)].

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APPENDIX A: DERIVATION OF THE EQUATIONS OF MOTION

Here we provide details of the derivation of the main Eqs. (6)–(9).

1. Dynamics of $\phi$

Evolution of the field $\phi = \frac{1}{\sqrt{V}} \phi_{\alpha, q=0}$ can be obtained from

$$\partial_t \langle \phi_{\alpha, q} \rangle_t = \int D[\phi] \phi_{\alpha, q} \partial_t P(\phi) = -\Gamma \left( \frac{\delta F}{\delta \phi_{\alpha, -q}} \right).$$

(A1)

where in the last equality we used the Fokker-Planck Eq. (5), and integration by parts. The latter derivative can be calculated from Eq. (3):

$$\frac{\delta F}{\delta \phi_{\alpha, -q}} = (r + K q^2) \phi_{\alpha, q} + \frac{4 i t}{V} \sum_{k_1, k_2} \phi_{\beta, k_1} \phi_{\beta, k_2} \phi_{\alpha, q - k_1 + k_2}.$$\hspace{1cm}

(A2)

Using Wick’s theorem and leaving only terms up to the leading order in $1/N$, we obtain

$$\left( \sum_{k_1, k_2} \phi_{\beta, k_1} \phi_{\beta, k_2} \phi_{\alpha, q - k_1 + k_2} \right) \approx \phi_{1, q=0}^3 + \phi_{1, q=0} \sum_{k} [D_k^1 + (N - 1) D_k^2].$$\hspace{1cm}

(A3)

Combining Eqs. (A1) and (A3), we arrive at Eq. (6) of the main text.

2. Dynamics of the correlators

Applying the same trick as above, we derive

$$\partial_t \langle \phi_{\alpha, k} \phi_{\alpha, -k} \rangle_t = 2 T \Gamma - 2 \Gamma \times \left[ \langle \delta F / \delta \phi_{\alpha, k} \rangle - \langle \delta F / \delta \phi_{\alpha, k} \rangle \right].$$ (A4)

For the case of the transverse component, in leading in $1/N$ order we obtain

$$\langle \phi_{\alpha, k} \sum_{k_1, k_2} \phi_{\beta, k_1} \phi_{\beta, k_2} \phi_{\alpha, -k, k_1 - k_2} \rangle \approx D_k^2 \left[ \phi_{1, q=0}^2 + \sum_{q} [D_q^1 + (N - 1) D_q^2] \right].$$ (A5)

Combining Eqs. (A4) and (A5), we arrive at Eq. (7) of the main text. For the case of the longitudinal component, similar to the above discussion, we get

$$\sum_{k_1, k_2} \langle \phi_{1, k} \phi_{\beta, k_1} \phi_{\beta, k_2} \phi_{1, -k_1 - k_2} \rangle - \langle \phi_{1, k_1} \rangle \langle \phi_{\beta, k_1} \phi_{\beta, k_2} \phi_{1, -k_1 - k_2} \rangle \approx D_k^2 \left[ 3 \phi_{1, q=0}^2 + \sum_{q} [D_q^1 + (N - 1) D_q^2] \right].$$ (A6)

This equation leads to Eq. (8).

APPENDIX B: LONGITUDINAL CORRELATIONS AT LONG TIMES

During the evolution, the longitudinal correlation function $D_k^1$ remains bell-shaped with a maximum at $k = 0$ suggesting to define $\langle \tilde{g}(t) = D_k^1(t) \rangle$ and $k^*_1(t)$ to be the wave vector corresponding to the half-width at half-maximum in $D_k^1$. Notably, both functions at long times behave as $\langle \tilde{g}(t) \rangle \sim t^{-d_q}$—see Fig. 5. We also observe that this power-law exponent implies that the longitudinal correlation functions exhibit the leading scaling, i.e., these modes should not be entirely ignored.

To explain the above observation, we note that at long times, when the order parameter $\phi(t) = \phi_0 + \delta \phi$ is already close to being recovered, the equation of motion (8) can be
Integration of this equation gives

$$\frac{dD_k}{dt} \approx -32\Gamma_{\phi 0} \partial\phi D_{k, eq}^1 - 2\Gamma (k^2 + 8\omega_0^2) D_k^1, \quad (B1)$$

where $D_k^1(t) = D_{k, eq}^1 + \delta D_k^1(t)$ and we disregarded fast $r_{\text{eff}}(t) \sim t^{-d_k}$ compared to slow $\delta\phi(t) \sim t^{-d_\phi} \ (0 < d_\phi < d_r)$. The above equation can be solved analytically. Indeed, substituting $\delta D_k^1(t) = e^{-2(\Gamma k^2 + 8\omega_0^2)/\nu} h_k(t)$, we obtain the following equation on $h_k(t)$:

$$\frac{dh_k}{dt} = -32\Gamma_{\phi 0} \partial\phi D_{k, eq}^1 e^{2(\Gamma k^2 + 8\omega_0^2)/\nu}. \quad (B2)$$

Integration of this equation gives

$$h_k(t) = h_k(t_0) + \frac{C}{k^2 + 8\omega_0^2} \int_{t_0}^t dt' e^{2(\Gamma k^2 + 8\omega_0^2) t'/\nu} (t')^{d_\phi}, \quad (B3)$$

where $C$ is some constant. We therefore conclude that

$$\delta D_k^1 = \delta D_k^{1(1)} + \delta D_k^{1(2)}, \quad (B4)$$

where $\delta D_k^{1(1)}(t) = h_k(t_0) e^{-2(\Gamma k^2 + 8\omega_0^2)/\nu}$ decays exponentially in time, whereas

$$\delta D_k^{1(2)}(t) \sim e^{-2(\Gamma k^2 + 8\omega_0^2)/\nu} \int_{t_0}^t dt' e^{2(\Gamma k^2 + 8\omega_0^2) t'/\nu} (t')^{d_\phi} \quad (B5)$$

is potentially important.

At long times $t \to \infty$, we observe that

$$F(t) \equiv \int_{t_0}^t dt' \frac{e^{a(t')^{d_\phi}}}{(t')^b}, \quad a, b > 0. \quad (B6)$$

Indeed, by differentiating $F(t)$ we note that it satisfies

$$\frac{dF}{dt} = \frac{e^{a t^{d_\phi}}}{t^b}. \quad (B7)$$

By substituting $F(t) = e^{a t^{d_\phi}}$ we separate rapid exponential growth from slow power-law-like dynamics encoded in $p(t)$:

$$\frac{dp}{dt} + ap = \frac{1}{t^b}. \quad (B8)$$

From this equation, we finally see that $p \sim t^{-b}$ (as long as $a \neq 0$). Combining Eqs. (B5) and (B6), we conclude that

$$\delta D_k^{1(2)} \sim \frac{t^{-d_\phi}}{k^2 + 8\omega_0^2}, \quad (B9)$$

i.e., indeed $\delta D_k^{1(2)}$ gets a power-law-like contribution with the leading exponent. For completeness, we also note that

$$\delta n_{\text{tot}}^{1(2)} \equiv \int \frac{d\mathbf{k}}{(2\pi)^3} \delta D_k^{1(2)} \sim t^{-d_\phi} \quad (B10)$$

also exhibits the same scaling.
state by assigning different temperatures to each subsystem. (ii) the rest of the lattice. One then describes a nonequilibrium lattice into two groups: (i) phonons associated with the OP, and (ii) the rest of the lattice. Therefore, in CDW systems one sometimes splits the phonons, the introduction of three subsystems might be confus-

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