Quantum quenches and competing orders

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Abstract

We study the non-equilibrium dynamics of an electronic model of competition between an unconventional charge density wave (a bond density wave) and \(d\)-wave superconductivity. In a time-dependent Hartree-Fock+BCS approximation, the dynamics reduces to the equations of motion of operators realizing the generators of SU(4) at each pair of momenta, \((\mathbf{k}, -\mathbf{k})\), in the Brillouin zone. We also study the non-equilibrium dynamics of a quantum generalization of a O(6) non-linear sigma model of competing orders in the underdoped cuprates (Hayward et al., Science 343, 1336 (2014)). We obtain results, in the large \(N\) limit of a O(\(N\)) model, on the time-dependence of correlation functions following a pulse disturbance. We compare our numerical studies with recent picosecond optical experiments. We find that, generically, the oscillatory responses in our models share various qualitative features with the experiments.
I. INTRODUCTION

A remarkable series of recent optical experiments\(^1\)-\(^4\) have explored time-dependent non-equilibrium physics in the cuprate superconductors at the picosecond time scale. Our work is specifically motivated by the observations of Ref. \(^1\): these experiments observed terahertz oscillations in the reflectivity of underdoped \(YBCO\) in a time-domain, pump-probe experiment. The onset temperature of the reflectivity oscillations was the same as the onset temperature of charge ordering in the recent X-ray measurements,\(^5\)-\(^7\) and so the oscillations were interpreted\(^1\) as an oscillation in the amplitude of the charge order. The reflectivity oscillations also showed a remarkable \(\pi\) phase shift, and a temperature-dependent frequency, across the superconducting critical temperature \(T_c\). The authors interpreted these phenomena in a classical phenomenological model of competition between superconductivity and charge order.

Our purpose here is to develop a more microscopic and quantum model of these oscillations. We will do this by examining two distinct models.

The first is a simple ‘hot spot’ electronic model for the competition between unconventional charge density wave (CDW) order (a bond density wave) and superconductivity (SC) which was proposed in Ref. \(^8\). The CDW ordering wavevectors of this first model are \((\pm Q_0, \pm Q_0)\), where \(Q_0\) is determined by the positions of the hot spots. We will extend the equilibrium results to time-dependent phenomena using a time-dependent Hartree-Fock-BCS theory similar to that used in Ref. \(^9\) for the quench dynamics of BCS superconductors. This model has the advantage of dealing directly with the underlying fermionic degrees of freedom. However, our analysis has the disadvantage that the spatial correlations of the order parameter are treated in a mean-field manner. Our results for this model appear in Sections \(\text{II-\text{V}}\).

The second model, described in Sections \(\text{VI-\text{VIII}}\), has a more complete treatment of spatial fluctuations of the CDW and SC orders, but works instead with an effective model for these bosonic order parameters alone. Also, the CDW wavevectors can now be either along the cardinal directions or the diagonals: so they can also take the experimentally observed values of \((\pm Q_0, 0)\), \((0, \pm Q_0)\). This model for the competing order parameters has an energy functional which is drawn directly from recent work by Hayward et al.\(^10\) They argued for a non-linear sigma model for a 6-component order parameter: two of the components, \(\hat{\Psi}\), represented \(d\)-wave SC, while the remaining four, \(\hat{\Phi}\), represented the complex order parameters for CDWs along the \(x\) and \(y\) directions; we will implicitly assume that \(\hat{\Psi}\ (\hat{\Phi})\) is a 2 (4) component real vector. The thermal fluctuations in Ref. \(^10\) were restricted to be on the space constrained by \(\hat{\Psi}^2 + \hat{\Phi}^2 = 1\), and we will also impose this constraint. However, we need to extend the model of Ref. \(^10\) to include a kinetic energy term to describe the dynamic questions of interest here. In Sections \(\text{II-\text{V}}\), the dynamics is derived from the equations of motion of the underlying electrons, and so for the non-linear
sigma model the analogous procedure is to integrate out the fermionic degrees in a path-integral formulation of the Hamiltonian. While integrating out fermions is a delicate matter in a metal due to the presence of Fermi surface, we argue that for our purposes the consequences are simple. The key point is the observation that both the $\Psi$ and $\Phi$ gap out the same important portion of the Fermi surface in the anti-nodal region (near the “hot spots”). As our study is restricted to the manifold $\Psi^2 + \Phi^2 = 1$, we can always assume that the antinodal Fermi surface is gapped. Consequently, integrating out the fermions only induces analytic time derivative terms in the effective action for $\Psi$ and $\Phi$, and we will only keep terms containing up to 2 time derivatives. We will ignore the small damping that can be induced by the gapless fermions in the nodal regions because both the CDW and SC orders do not couple strongly to these fermions.

We will conclude the paper in Section IX with a summary of our results, and a comparison of the distinct methodologies employed in the paper.

II. HOT SPOT MODEL

We begin by reviewing the equilibrium properties of the simple “hot spot” model of competing orders presented in Ref. 8. The model is defined in terms of 4 species of fermions $\psi_{aa}$, $a = 1 \ldots 4$, $\alpha = \uparrow, \downarrow$ located near “hotspots” on the Fermi surface as shown in Fig. 1. Their kinetic energy is

![Figure 1](image)

**FIG. 1.** Definitions of the $\psi_{1,2,3,4}$ fermions around the Fermi surface. Each fermion resides around a curved patch of the Fermi surface shown by the thick lines. The red (green) hot spots are where the superconducting and bond density wave orders are positive (negative).
given by

\[ H_0 = \sum_k \left[ \epsilon_1(k) \psi_{1\alpha}^\dagger(k) \psi_{1\alpha}(k) + \epsilon_2(k) \psi_{2\alpha}^\dagger(k) \psi_{2\alpha}(k) + \epsilon_1(-k) \psi_{3\alpha}^\dagger(k) \psi_{3\alpha}(k) + \epsilon_2(-k) \psi_{4\alpha}^\dagger(k) \psi_{4\alpha}(k) \right]. \]

(1)

We take the origin of momentum space at the hot spots, and orient the x-axis orthogonal to the Fermi surface for the \( \psi_{1,3} \) fermions; so we can write

\[ \epsilon_1(k) = k_x + \gamma k_y^2. \]

(2)

We have taken the Fermi velocity to be unity, while \( \gamma \) measures the curvature of the Fermi surface. The dispersion \( \epsilon_2(k) \) has the form obtained by rotating \( \epsilon_1(k) \) so that the direction orthogonal to the Fermi surfaces of the \( \psi_{2,4} \) has a linear dispersion. After rescaling momenta appropriately, we can choose the convenient momentum space cutoffs \(-\pi < k_x, k_y < \pi\), and the value \( \gamma = 1/\pi \).

Next, we add interactions between these fermions. The microscopic exchange \( (J) \) interactions and Coulomb repulsion \( (V) \) when projected onto the hot spots lead to

\[ H_1 = \int d^2x \left[ -J \left( \psi_{1\alpha}^\dagger \sigma_{\alpha\beta} \psi_{2\beta} + \psi_{2\alpha}^\dagger \sigma_{\alpha\beta} \psi_{1\beta} \right) \right. \\
\left. \cdot \left( \psi_{3\gamma}^\dagger \sigma_{\gamma\delta} \psi_{4\delta} + \psi_{4\gamma}^\dagger \sigma_{\gamma\delta} \psi_{3\delta} \right) \right. \\
\left. -V \left( \psi_{1\alpha}^\dagger \psi_{2\alpha} + \psi_{2\alpha}^\dagger \psi_{1\alpha} \right) \left( \psi_{3\beta}^\dagger \psi_{4\beta} + \psi_{4\beta}^\dagger \psi_{3\beta} \right) \right]. \]

(3)

The full Hamiltonian \( H_0 + H_1 \) has an exact \( SU(2) \times SU(2) \) pseudospin rotation symmetry\(^{11}\) when \( \gamma = 0 \) and \( V = 0 \).

Next, we review the Hartree-Fock-BCS theory of the hotspot model \( H_0 + H_1 \). The superconducting (SC) order parameter, \( \Delta \), involves pairing of particles on antipodal points on the Fermi surface, while the charge density wave (CDW) order, \( \Pi \), involves pairing of particles with holes on the antipodal point;\(^8\) consequently, the CDW ordering wavevector has the values \( \pm Q_0, \pm Q_0 \), as
is clear from Fig. 1.

\[
\Delta_1(k) = \left\langle \varepsilon_{\alpha \beta} \psi_{1\alpha}^\dagger(k) \psi_{3\beta}^\dagger(-k) \right\rangle ; \quad \Delta_1 \equiv \sum_k \Delta_1(k)
\]

\[
\Delta_2(k) = \left\langle \varepsilon_{\alpha \beta} \psi_{2\alpha}^\dagger(k) \psi_{4\beta}^\dagger(-k) \right\rangle ; \quad \Delta_2 \equiv \sum_k \Delta_2(k)
\]

\[
\Pi_1(k) = \left\langle \psi_{1\alpha}^\dagger(k) \psi_{3\alpha}(k) \right\rangle ; \quad \Pi_1 \equiv \sum_k \Pi_1(k)
\]

\[
\Pi_2(k) = \left\langle \psi_{2\alpha}^\dagger(k) \psi_{4\alpha}(k) \right\rangle ; \quad \Pi_2 \equiv \sum_k \Pi_2(k)
\]

(4)

It was found\textsuperscript{8} that optimal state has a \( d \)-wave signature for both the superconducting and charge orders, with \( \Delta_1 = -\Delta_2 \) and \( \Pi_1 = -\Pi_2 \). For the charge order, this \( d \)-wave structure implies that the charge modulation is primarily on the \emph{bonds} of the underlying lattice.\textsuperscript{12} With the above orders, the mean field Hamiltonian is

\[
H_{MF} = H_0 + \frac{(3J - V)}{2} \left( -\Delta_1 \varepsilon_{\alpha \beta} \psi_{2\alpha}(k) \psi_{4\beta}(-k) \right.
+ \Delta_2^* \varepsilon_{\alpha \beta} \psi_{1\alpha}^\dagger(k) \psi_{3\beta}^\dagger(-k) - \Delta_2 \varepsilon_{\alpha \beta} \psi_{1\alpha}^\dagger(k) \psi_{3\beta}(-k)
+ \Delta_1^* \varepsilon_{\alpha \beta} \psi_{2\alpha}^\dagger(k) \psi_{4\beta}^\dagger(-k) \right)
+ \frac{(3J + V)}{2} \left( \Pi_1 \psi_{2\alpha}(k) \psi_{4\alpha}(k) + \Pi_2^* \psi_{1\alpha}^\dagger(k) \psi_{3\alpha}(k)
+ \Pi_2 \psi_{3\alpha}^\dagger(k) \psi_{1\alpha}(k) + \Pi_1^* \psi_{2\alpha}^\dagger(k) \psi_{4\alpha}(k) \right).
\]

(5)

Ref. 8 presented the solution of the equilibrium properties of the Hartree-Fock-BCS equations for a variety of values of \( J \) and \( V \). Here, we reproduce in Fig. 2 the solution at one set of parameter values to illustrate the basic temperature dependence of the mean-field order parameters. Note that the CDW order, \( \Pi_1 \) has an onset at a higher \( T \). However, at the superconducting \( T_c \), it starts ‘competing’ for the Fermi surface with the SC order \( \Delta_1 \), and so decreases with decreasing \( T \).

III. EQUATIONS OF MOTION

We will follow the same general strategy as in Ref. 9: we will work with Heisenberg equations of motion from the Hamiltonian \( H_{MF} \), where the mean field order parameters \( \Delta_1 \) and \( \Pi_1 \) take their instantaneous average values.

An important feature of this method for the present model is that that commutators of the operators \( \Delta_1(k) \) and \( \Pi_1(k) \) with \( H_{MF} \) do not close among themselves: they produce additional operators whose equations of motion we have to also consider. By repeatedly evaluating commu-
FIG. 2. Superconducting ($\Delta_1$) and bond ($\Pi_1$) orders in the hot spot model as a function of $T$.

tators of the operators so generated, we find that we also have to consider the operators

$$N_i(k) = \psi_{i\alpha}^\dagger(k)\psi_{i\alpha}(k) , \quad P_i(k) = \varepsilon_{\alpha\beta}\psi_{i\alpha}^\dagger(k)\psi_{i\beta}^\dagger(-k)$$

(6)

where $i = 1 \ldots 4$; note

$$P_i(-k) = P_i(k) , \quad N_i^\dagger(k) = N_i(k).$$

(7)

Among all the operators introduced so far, the operator

$$N_1(k) + N_3(k) - N_1(-k) - N_3(-k)$$

(8)

commutes with all other operators. The remaining 15 operators

$$N_1(k) + N_3(-k), N_1(k) + N_1(-k) - 1, N_3(k) + N_3(-k) - 1,$$

$$\Delta_1(k), \Delta_1(-k), \Delta_1^\dagger(k), \Delta_1^\dagger(-k),$$

$$\Pi_1(k), \Pi_1(-k), \Pi_1^\dagger(k), \Pi_1^\dagger(-k),$$

$$P_1(k), P_1^\dagger(k), P_3(k), P_3^\dagger(k)$$

(9)

form the Lie algebra of SU(4). This is to be compared with the SU(2) algebra of Ref. 9 of the operators $P_1(k), P_1^\dagger(k), N_1(k) + N_1(-k) - 1$.

It is now a straightforward, but tedious, exercise to evaluate the commutators of this SU(4) algebra, and so generate the equations of motion associated with $H_{MF}$. We display the explicit form of these equations of motion in Appendix A.
FIG. 3. Oscillation of CDW order parameter $\Pi$ (left) and SC order parameter $\Delta$ (right) as a function of time in the quench case, from low temperature at the bottom to high temperature at the top, temperatures are taken from 0.025 to 0.375 with 0.05 step. Note that here we are plotting the absolute value of the order parameters. Also, we have added constants to the curves to make them evenly spaced. The initial value $J_0 = 1.2$, $V_0 = 0.9$, the quench is taken as $\Delta J = 0$, $\Delta V = -0.1$. The initial $T_c$ at equilibrium can be computed to be 0.25, the final $T_c$ to be 0.33.

IV. QUENCH

First let us consider the quench case. By quench, we mean the coupling changes abruptly, i.e.

$$V(t) = V_0 + \Delta V \theta(t), \quad J(t) = J_0 + \Delta J \theta(t)$$

(10)

where $\theta(t)$ is the step function, and $\Delta V$ and $\Delta J$ are the sizes of the steps. Similar problems have been considered in the BCS system. We take the system to be at equilibrium at the beginning with both CDW and SC order, at a fixed temperature which can be both below and above superconducting critical temperature $T_c$ in Fig. 2. The evolutions of order parameters can be obtained using Heisenberg equations of motions. We obtained oscillations of the CDW order parameter $\Pi$, and the SC order parameter $\Delta$ as a function of time at different temperatures, as shown in Fig. 3 for the parameters $J_0 = 1.2$, $V_0 = 0.9$, $\Delta J = 0$, $\Delta V = -0.1$. We find that at high temperature which is larger than $T_c$, $\Delta$ stays zero, while the oscillation of $\Pi$ is suppressed.

As in Ref. 1, we use a decaying sinusoidal function to fit the data. Unlike the experimental data, the amplitude of the oscillation does not decay to zero at long times in the present mean-field model. Later we will analyze the fourier spectrum of the oscillation, but for now we proceed with a naive fitting to the following function

$$f(t) = a e^{-b(t-t_0)} \sin(c(t-t_0) + 2\pi d) + e$$

(11)
FIG. 4. Fitting of CDW order parameters Π in the left panel of Fig. 3, the dashed lines are fitting lines using Eq. 11. We used the data after time = 5 and later fittings also obey this rule.

FIG. 5. Left to right: amplitude a, frequency c and phase d of the fit Eq. 11 fitting the data in Fig. 4 as a function of temperature. The blue line denotes initial equilibrium $T_c = 0.25$, the green line denotes after quench, the equilibrium $T_c = 0.33$.

which works quite well over the time window studied. The fit is shown in Fig. 4. We have set an onset $t_0$ in the fitting function, the fitting phase $d$ will depend on the choice of $t_0$, since the frequency $c$ does not stay constant over the temperature range. In all of the fits we choose $t_0 = 5.5$, which will give nearly $\pi$ phase shift in the later pulse case. The fitting phase $d$ here just helps us to better see the phase shift rather than simply estimating by eye.

In Fig. 5, we show the variation of the amplitude $a$, frequency $c$, phase $d$ in Eq. 11 as a function of temperature. The most important feature is that the amplitude $a$ is enhanced below the initial $T_c$ at equilibrium, which is also a key feature in the $O(6)$ field theory description that will be
discussed in later sections. This resembles the oscillatory behavior in the experiment Ref. 1. Also, the frequency varies against temperature, and there is a phase shift in the oscillations upon crossing \( T_c \). Here the phase shift is smaller than \( \pi \) by our choice of \( t_0 = 5.5 \).

We can further use spectral analysis to fit the simulation data. Using the Lomb-Scargle algorithm, we can find frequencies' spectral power as shown, for example, in the inserted figure in Fig. 6. We choose frequencies which have spectral power larger than ten percent of the largest power to fit the oscillation. Including frequencies from main peak also large side peaks, the fit result is quite good. However, the peak frequencies are quite similar as the fitting number \( c \) that we found using the simple decaying sinusoidal function as shown in the middle panel of Fig. 5.

To understand the fitting frequency behavior above \( T_c \), we have \( \Delta = 0 \), and the mean-field Hamiltonian Eq. (5) reduces to

\[
H_{MF} = H_0 + \frac{(3J + V)}{2} \left( \Pi_1 \psi^\dagger_{4\alpha}(k)\psi_{2\alpha}(k) + \Pi_2 \psi^\dagger_{1\alpha}(k)\psi_{3\alpha}(k) \\
+ \Pi_2 \psi^\dagger_{3\alpha}(k)\psi_{1\alpha}(k) + \Pi_1 \psi^\dagger_{2\alpha}(k)\psi_{4\alpha}(k) \right). \tag{12}
\]

From the commutation relations in Eq. (A1), we can make the identification so that these operators satisfy the \( SU(2) \) algebra:

\[
\Pi_1 \rightarrow S_+, \Pi_1^\dagger \rightarrow S_-, \frac{N_1 - N_3}{2} \rightarrow S_z \tag{13}
\]

Furthermore if we ignore the curvature of the Fermi surface, then \( \epsilon_1(k) = -\epsilon_1(-k) \), the above
Hamiltonian becomes

\[ H_{MF} = \sum_k \left[ 2\epsilon_1(k)S_z(k) + \frac{3J + V}{2} \left( -\langle S_- \rangle S_+(k) - \langle S_+ \rangle S_-(k) \right) \right] \]  

(14)

here we have used \( \Pi_2 = -\Pi_1 \) and only considered 1, 3 hotspot field (the 2, 4 channel would be similar). And this resembles the well-known pseudospin formulation of the BCS system, as studied in Ref. 9. For a small deviation, the frequency would be proportional to the order parameter \( \Pi \). This explains the fact that the oscillation frequency decrease rapidly above \( T_c \).

We have also computed the positive quench case in Fig. 22 of Appendix B, where \( J_0 = 1.2, V_0 = 0.9, \Delta J = 0, \Delta V = 0.1 \). We also get amplitude enhancement below \( T_c \), although the effect is not that big.

V. PULSE

Since in the experiment,\(^1\text{–}^4\) the disturbance is a short-time optical pulse, it should be more reasonable to consider a pulse in our time-dependent Hamiltonian, \( i.e. \)

\[ J(t) = J_0 + \Delta J \left( 1 - \tanh^2(\omega t) \right) \quad , \quad V(t) = V_0 + \Delta V \left( 1 - \tanh^2(\omega t) \right) \]  

(15)

We have chosen \( \omega = 1, \Delta V = 0.1 \) as shown in Fig. 7. Fig. 8 shows the fit using Eq. 11 and the fitting parameters are shown in Fig. 9. We have similar amplitude enhancement effect and temperature dependent frequencies, however, simply by eye, the first valley at low temperature becomes a peak upon crossing \( T_c \). In Fig. 9, we can see that there is a nearly \( \pi \) (or \(-\pi\)) phase shift crossing \( T_c \). The spectral analysis is shown in Fig. 10, the peak frequencies are quite closed to the fitting frequencies. From the inserted figure in the left panel, near \( T_c \) side peak will grow then becomes the main peak. This reflects the frequency abrupt change near \( T_c \) in the right panel.

Fig. 11 shows a direct comparison of our simulation and the experimental data. Both data indicate enhancement of CDW oscillation below \( T_c \). If we put back the real energy scale \( T_c \approx 75K \) in the simulation, the fitting frequency is of the order THz, which is also the same order found in Ref. 1. The difference is that the frequency is more dependent on temperature in the numerics than in the experiment.

We also calculated the case \( \Delta V = -0.1 \) as shown in Figs. 23–25 of Appendix B. Other than the phase shift to the opposite direction, the other results are quite similar. In the experiment,\(^1\) the pump duration is 60 fs. In the simulation above, if we put back the energy scale, the choice of \( \omega = 1 \) means that the pulse duration is of the order 100 fs. We checked the case when we set \( \omega = 4 \) (duration 25 fs) as shown in Figs. 26–28 of Appendix B. Because the duration is shorter,
FIG. 7. Oscillation of CDW order parameter $\Pi$ (left) and SC order parameter $\Delta$ (right) as a function of time in the pulse case, from low temperature at the bottom to high temperature at the top, temperatures are taken from 0.025 to 0.375 with 0.05 step. The initial value $J_0 = 1.2$, $V_0 = 0.9$, the pulse is taken as $\Delta J = 0$, $\Delta V = 0.1$, $\omega = 1$. The initial $T_c$ at equilibrium can be computed to be 0.25, at the largest derivation $V = V_0 + \Delta V$, the corresponding equilibrium $T_c$ to be 0.2.

the oscillation amplitude gets smaller, however the frequency and phase shift do not change much from the $\omega = 1$ case.

Finally, we examined the quench or pulse $J$ case, i.e. $\Delta J \neq 0$. In this case, our numerics showed no clear enhancement of oscillation below $T_c$. We showed one case when pulse $\Delta J = 0.1$, $\Delta V = 0$ in Fig. 29: no obvious enhancement or phase shift is observed crossing $T_c$. In the experiment, the pump light suppresses the superconductivity condensation, which enhances the CDW as mentioned in Ref. 1. In our model the nearest neighbor repulsion $J$ favors CDW and suppresses the condensation, while anti-ferromagnetic coupling $J$ favor both. Therefore, disturbing in $V$ mimics the pump effect in the experiment.

VI. QUANTUM NON-LINEAR SIGMA MODEL

In the sections above, we worked with a model of electrons with an underlying Fermi surface, and then examined the dynamics implied by the electron dispersion on order parameters consisting of fermion bilinears. In the following, we will work directly with the competing order parameters, via an effective Hamiltonian for the bosonic order parameters themselves. This approach will allow for spatial fluctuations of the order parameters, and hence goes beyond the mean-field treatment of the previous sections. However, our analysis will be limited here to a “collisionless” large $N$ limit in which true thermalization does not take place in the long-time limit.

As we noted in Section I, the CDW of the present model can have the experimentally observed
FIG. 8. Fitting of CDW order parameters $\Pi$ in the left panel of Fig. 7, the dashed lines are fitting lines using Eq. 11.

FIG. 9. Left to right: amplitude $a$, frequency $c$ and phase $d$ of the fit Eq. 11 fitting the data in Fig. 8 as a function of temperature. The blue line denotes initial equilibrium $T_c = 0.25$, the green line denotes at the largest derivation $V = V_0 + \Delta V$, the equilibrium $T_c = 0.2$.

wavevectors of $(\pm Q_0, 0), (0, \pm Q_0)$.

Our model for the competing order parameter has an energy functional which is drawn directly from recent work by Hayward et al.$^{10}$ as we discussed in Section I. And this is supplemented with a “relativistic” time derivative term, as also discussed in Section I, thus ignoring any damping that may arise from the gapless fermions at the nodes of the $d$-wave superconductor.
FIG. 10. Left: spectral fitting of the left panel in Fig. 7. The inserted figure is the power spectrum at $T = 0.225$ near $T_c$. Right: peak frequencies from the left panel as a function of temperature.

FIG. 11. Comparison of the numerics (dashed red line) from Fig. 9 and experiment data\textsuperscript{1}(red and blue dots). Left panel is amplitude against temperature, where dashed grey line denotes initial $T_c$ before perturbation in the experiment, the blue line denotes initial $T_c$ in the numerics, the green line denotes the equilibrium $T_c$ at the largest derivation $V = V_0 + \Delta V$ in the numerics. The right panel shows the comparison in frequency and phase.

In Lorentzian signature, the path integral of such an O(6) model takes the following form:

$$
\int D\Psi \, D\Phi \, D\sigma \, e^{iS},
$$

where

$$
S = \frac{N\rho_s}{2} \int d^3 x \left( (\partial \bar{\Psi})^2 + \lambda (\partial \Phi)^2 - (g \bar{\Phi}^2 - \mu (\Phi \Phi)^2 + \sigma (\Phi^2 + \bar{\Psi}^2) - 1) \right),
$$
where the $\sigma$ integral imposes the condition

$$\Phi^2 + \Psi^2 = 1,$$

(18)

and $\Psi$ is an $N/3$ dimensional vector, whereas $\Phi$ a $2N/3$ dimensional vector. The symmetries of the problem also allow a linear time-derivative term $\Psi_1 \partial_t \Psi_2$, which is allowed by the absence of particle-hole symmetry about the Fermi surface. However, the particle-hole asymmetry is small and we will ignore it in our analysis. Also we have chosen the velocity of ‘light’ in our relativistic formulation to be unity by rescaling the time co-ordinate.

The subsequent procedure we follow is very much the same as in Ref 13, which is further elaborated and extended in Ref. 14, accommodating more general dynamical evolution beyond a strict quantum quench. The models considered in these previous works, however, focus on the linear sigma model. Our path-integral treatment parallels that in Ref. 15 and particularly Ref. 16 where the Schwinger Keldysh formalism is employed. We note however that this is identical to the approach taken elsewhere, and that one can show that the self-consistent mean-field equation of a general linear sigma model with a $\phi^4$ coupling, considered for example in Ref. 14, reduces to a NLSM by taking the large $\phi^4$ coupling limit while holding the ratio of the $\phi^4$ and $\phi^2$ couplings constant.

To proceed with the path-integral, we can linearize the action by introducing the auxiliary field $\rho$:

$$S = \frac{N\rho_s}{2} \int d^3x \left( (\partial \Psi)^2 - \sigma \Psi^2 + (\partial \Phi)^2 - (g + \rho + \sigma)/\lambda \Phi^2 - \frac{\rho^2}{4\mu} + \sigma \right)$$

(19)

Note that we have rescaled $\Phi$ to obtain a canonical kinetic term for $\Phi$ above. $\rho$ is defined accordingly.

Integrating out $\Phi, \Psi$, gives an effective action in the remaining path integral $\int D\sigma D\rho e^{iN/2S_{\text{eff}}}$

$$S_{\text{eff}} = \frac{i}{3} \text{tr} \ln(\Box + \sigma) + \frac{2i}{3} \text{tr} \ln(\Box + (g + \rho + \sigma)/\lambda) + \int d^3x (-\frac{\rho_s \rho^2}{4\mu} + \rho_s \sigma).$$

(20)

The corresponding gap equations are

$$\rho_s = \frac{1}{3} \int \frac{d^2k}{(2\pi)^2} \left( G_\Psi(k,t) + 2/\lambda G_\Phi(k,t) \right),$$

$$\rho(t) = \frac{4\mu}{3\rho_s \lambda} \int \frac{d^2k}{(2\pi)^2} G_\Phi(k,t),$$

(21)

where $G_\Psi, G_\Phi$ are spatially Fourier transformed equal time correlation functions. We have suppressed the details of the Schwinger-Keldysh time contour, whose only consequence in the leading large $N$ calculation is to determine the boundary conditions of the Green’s functions that we will
review below.

These Green’s functions can be conveniently parametrized by the (spatially Fourier transformed) time dependent field as follows: \(^{14,16}\)

\[
\Phi_k(t) = \Phi_k(t_i) \sqrt{\frac{\Omega^\Phi_k(t_i)}{\Omega^\Phi_k(t)}} \cos\left(\int_{\Omega^\Phi_k(t_i)}^{t_i} \Omega^\Phi_k(t) dt + \Pi_{\Phi_k(t_i)} \sin(\int_{\Omega^\Phi_k(t_i)}^{t_i} \Omega^\Phi_k(t) dt) \right),
\]

(22)

and similarly we can parametrize \(\Psi\) using these time dependent functions. ie

\[
\Psi_k(t) = \Psi_k(t_i) \sqrt{\frac{\Omega^\Psi_k(t_i)}{\Omega^\Psi_k(t)}} \cos\left(\int_{\Omega^\Psi_k(t_i)}^{t_i} \Omega^\Psi_k(t) dt + \Pi_{\Psi_k(t_i)} \sin(\int_{\Omega^\Psi_k(t_i)}^{t_i} \Omega^\Psi_k(t) dt) \right).
\]

(23)

\(\Pi_{a_k}\) denotes the conjugate field of \(a \in \{\Phi, \Psi\}\), and that \(t_i\) is some initial time which we could beq taken to approach \(-\infty\).

These \(\Omega^\Phi_k(t)\) satisfies the equation

\[
\frac{\ddot{\Omega}^\Phi_k}{2\Omega^\Phi_k} - \frac{3}{4} \left(\frac{\dot{\Omega}^\Phi_k}{\Omega^\Phi_k}\right)^2 + (\Omega^\Phi_k)^2 = k^2 + m^2_{\Phi}(t).
\]

(24)

Similarly,

\[
\frac{\ddot{\Omega}^\Psi_k}{2\Omega^\Psi_k} - \frac{3}{4} \left(\frac{\dot{\Omega}^\Psi_k}{\Omega^\Psi_k}\right)^2 + (\Omega^\Psi_k)^2 = k^2 + m^2_{\Psi}(t).
\]

(25)

The effective mass is given by

\[
m^2_{\Phi}(t) = (g + \rho + \sigma)/\lambda, \quad m^2_{\Psi}(t) = \sigma.
\]

(26)

From now on, we take \(\mu = 0\) and so \(\rho = 0\).

VII. EQUILIBRIUM PROPERTIES

Before delving into time-dependent scenarios, we review the properties of the theory at equilibrium.

At equilibrium, the time ordered Green’s function at finite temperature is given by

\[
G^T_\alpha(k, t_1 - t_2) = \frac{e^{-i\omega_k|t_1 - t_2|}}{2\omega_k} \text{coth}\left(\frac{\beta\omega_k}{2}\right), \quad \omega_k = \sqrt{k^2 + m^2_\alpha}.
\]

(27)
The gap equation (21) therefore becomes

\[ 6\pi \beta \rho_s = \left( \log \left[ \frac{\sinh \left( \frac{\beta}{2} \sqrt{m_{\Psi}^2 + \Lambda^2} \right)}{\sinh \left( \frac{\beta}{2} m_{\Psi} \right)} \right] + \frac{2}{\lambda} \log \left[ \frac{\sinh \left( \frac{\beta}{2} \sqrt{g + m_{\Psi}^2 + \lambda \Lambda^2} \right)}{\sinh \left( \frac{\beta}{2} \sqrt{g + m_{\Psi}^2} \right)} \right] \right) \]  

(28)

where \( \Lambda \) is the UV cutoff. We have substituted \( \lambda m_{\Psi}^2 = m_{\Psi}^2 + g \).

This expression can be compared with the classical case of Ref. 10, where only the zero Matsubara frequency is kept in the thermal Green’s function for each species \( a \) i. e.

\[ G^a(\omega_n, k) \sim \frac{T}{k^2 + m_a^2}. \]  

(29)

The gap equation in this case reduces to

\[ 6\pi \beta \rho_s = \left( \log \left[ \frac{\sqrt{m_{\Psi}^2 + \Lambda^2}}{m_{\Psi}} \right] + \frac{2}{\lambda} \log \left[ \frac{\sqrt{g + m_{\Psi}^2 + \lambda \Lambda^2}}{\sqrt{g + m_{\Psi}^2}} \right] \right) \]  

(30)

We compare the Green’s function \( G_{\Phi}(k = 0) \) plotted against temperature at fixed \( \rho_s \). In the low temperature limit the quantum \( G_{\Phi} \) falls off much slower than linearly in \( T \) because of the behavior of \( \cot(\beta m_{\Phi}/2) \) in the Green’s function. We recall that \( m_{\Phi}^2 = (m_{\Psi}^2 + g)/\lambda \).

Note that in both the quantum calculation and the classical approximation, \( G_{\Phi}(k = 0) \) exhibits a maximum at some temperature \( T_p \). The peak marks the change between the low temperature behavior where fluctuations are dominated by the superconductivity \( \Psi \) component, and the high temperature behavior which is characterized by fluctuations exploring all directions.\(^{10}\)

**VIII. PULSE-LIKE DISTURBANCE IN \( \rho_s \)**

In the experiments reported in Ref. 1–4, the system is perturbed by pulses of lasers over a short duration of the order of tens of femtoseconds. As a first brush, to mimic the effect of such a disturbance, we consider perturbing the system by a time dependent \( \rho_s \). To be precise, we take

\[ \rho_s = \rho_0 + \delta \rho (\tanh(v t)^2 - 1). \]  

(31)

To compute using the saddle point approximation, we should rescale \( \Phi, \Psi \):

\[ \tilde{\Phi} = \sqrt{\rho_s} \Phi, \quad \tilde{\Psi} = \sqrt{\rho_s} \Psi. \]  

(32)
FIG. 12. Equal-time two point function of the charge order $\Phi$ at vanishing momentum $k$ plotted against temperature at constant $\rho_s$, at cutoff $\Lambda = 5$ and $\lambda = g = 1$. Left: the quantum Green’s function. Right: the classical Green’s function. The peak position is presumed to be near the onset of superconductivity: this onset suppresses the charge order fluctuations, leading to a peak in $G_\Phi$ with decreasing temperature.

This leads to a change of the expression for the effective mass:

$$m_{\Phi}^2 = \left(\frac{\sigma + g}{\lambda} - (\Box K + (\partial K)^2)\right) \quad m_{\Phi}^2 = (\sigma - (\Box K + (\partial K)^2)),$$

(33)

where

$$K = \frac{1}{2} \ln \rho_s.$$

(34)

When $\lambda = 1$, these functions $K$ can be absorbed in the definition of $m_{\Phi}^2 = \sigma - (\Box K + (\partial K)^2)$.

We can evolve the system beginning at $vt < -1$, where the time dependence is negligible, and allow the system to react to the shaking. The equal time Green’s function in this case, using also the parameterizations (22,23), and the initial conditions

$$\langle \phi^a(t_i)\phi^a(t_i) \rangle = \frac{1}{2\Omega^a(t_i)}, \quad \langle \Pi^a(t_i)\Pi^a(t_i) \rangle = \frac{\Omega^a(t_i)}{2},$$

(35)

takes the form$^{16}$

$$G^a(k, t) = \frac{1}{2\Omega_k^a(t)} \coth\left(\frac{\beta_0\Omega_k^a(t)}{2}\right), \quad a \in \{\Psi, \Phi\}$$

(36)

and $\Omega^a(t_i \to -\infty) = \sqrt{k^2 + m_a(-\infty)^2}$ where $m_a(-\infty)$ is the initial mass of each field before the application of the disturbance, and that it is set by the initial temperature $\beta_0$ and $\rho_0$ by solving
the gap equation self-consistently at $t = -\infty$. The gap equation (21) then becomes

$$\lambda m^2_{\Phi}(t) - m^2_{\Psi}(t) = g,$$
$$m^2_{\Psi}(t) = \frac{A}{2B},$$
$$A = -3\ddot{\rho}_s + \int \frac{dk}{2\pi} k \left( 2(\Omega^\psi_k(t))^2 - k^2 \right) + \frac{1}{2} \left( \frac{\dot{\Omega}^\psi_k(t)}{\Omega^\psi_k(t)} \right)^2 G^\psi(k, t) + \frac{4}{\lambda} \left( \Omega^\psi_k(t)^2 - k^2 - \frac{g}{\lambda} \right) + \frac{1}{\lambda} \left( \frac{\dot{\Omega}^\psi_k(t)}{\Omega^\psi_k(t)} \right)^2 G^\Psi(k, t),$$
$$B = \int \frac{dk}{2\pi} k \left( G^\Psi(k, t) + \frac{2}{\lambda^2} G^\Phi(k, t) \right).$$

(37)

The above is obtained by replacing the Green’s function in the gap equation by the explicit forms (36), and then differentiating the gap equation with respect to time twice.

A natural regularization scheme would be to place the system on a lattice with lattice constant $a$. To do so, we make the replacement

$$k^2 \rightarrow \frac{(4 - 2\cos(ak_x) - 2\cos(ak_y))}{a^2},$$

(38)

and that $k_x, k_y$ take values between $-\pi/a$ to $\pi/a$.

A. Numerical Results

We consider dynamical oscillations of the system at various different choice of parameters $g, \lambda$ and initial temperatures $T = 1/\beta_0$, subjected to different disturbances applied for different durations. We plot the oscillations of the self-consistent effective mass $m^2_{\Psi}(t)$ as a function of time. These results are presented in Figs. 13, 14, 17 and 18. For each set of parameters $g, \lambda$ we obtain the time evolution at 10 different initial temperatures, and we indicate the position of these initial conditions in the equilibrium $G_{\Phi}(k = 0) - T$ plot. We look particularly at the vicinity of the peak, and observe the changes in the oscillations as temperature is increased across the peak.

In all these cases, the self-consistent mass $m_{\Psi}(t)$ displays a large peak while the disturbance is applied, and exhibits oscillatory behavior after the time-dependent disturbance is withdrawn.

The disturbances applied in the cases in figures 13 and 14 are relatively slow compared to the initial values of $m_{\Psi}$ and that $m_{\Psi}(t = -\infty) \sim m_{\Psi}(t = -\infty)$. In these cases, the subsequent oscillations are sinusoidal with a distinct frequency and a decaying amplitude. In these cases, we fit the oscillations by the function

$$f(t) = \exp(-at)b\sin(ct + 2\pi d) + e + ft.$$  

(39)
FIG. 13. Left Panel: Oscillations of $m_\Psi^2$ as a function of time at 10 different initial temperatures, from low temperatures at the bottom of the picture to high temperatures at the top, at constant $\rho_0 = 0.0756$ (corresponding to choosing $m_\Psi = 1/10$ at $T = 1/100$, $\lambda = 1$, $g = 0.2$, and $a = 1$). Integral along $k_x$ and $k_y$ is each divided into 90 steps. The pulse parameter is taken as $v = 1/5, \delta \rho = 1/500$. These 10 initial temperatures correspond to 10 points on the equilibrium plot of $G_\Psi(k = 0)$ against $T$, as shown on the right panel. The color of the markers match the color of the curves on the left. Note that variation of the mass before $t = 10$ is a huge peak resulting from the disturbance which is not shown in the picture.

An infinitesimal, approximately linear downward drift of the oscillations appears at sufficiently high temperatures (i.e. $f < 10^{-3}$ at $a = 1$). This is most likely numerical error since the oscillation amplitudes are also extremely small. We include in the fit function the linear term in order to remove the effect of this drift to improve accuracy for the frequency fit. In all cases, $e$ is very close to the original value of $m_\Psi(t = -\infty)$. The results of the fit for $a, b, c, d$ corresponding to the data presented in figures 13 and 14 are presented in figures 15 and 16 respectively.

A most distinctive feature is that there is a large suppression in the oscillation amplitude as the temperature $T$ increases across $T_p$, the temperature corresponding to maximal $G_\Psi(k = 0)$ at equilibrium. This strongly resembles the experimental results\textsuperscript{1} where oscillations are enhanced below the critical temperature $T_c$ of superconductivity, which is also observed in the electron ‘hotspot’ model considered in the previous section. A second feature is that the characteristic frequency of the oscillations increase with temperature at a rate faster than linearly, and the rate of increase does not appear to level off at high temperatures. One can inspect the ratio of the oscillation frequencies and peak temperature $T_p$. Consider say the results from figure 14. The oscillation frequency is roughly $\omega \sim 0.3 - 1$, whereas $T_p \sim 0.3$, giving a ratio of order between $1 - 3$. In the experiments, the oscillation frequencies are of order $2\pi \times 2/\text{ps}$ and the critical temperature is of order $\sim 50\text{K}$, giving a ratio of

$$\frac{\hbar \omega}{(k_B T)} \sim 2\pi \times 0.3,$$

(40)
FIG. 14. Left Panel: Oscillations of $m^2_{\Psi}$ as a function of time at 15 different initial temperatures, from low temperatures at the bottom of the picture to high temperatures at the top, at constant $\rho_0 = 0.1067$ (corresponding to $m_{\Psi} = 1/5$ at $T = 1/100$, $g = 0.1$, $\lambda = 6/10$, and $a = 1$). Integral along $k_x$ and $k_y$ is each divided into 90 steps. The pulse parameter is taken as $\nu = 1/5$, $\delta \rho = 1/500$. These 15 initial temperatures correspond to 15 points on the equilibrium plot of $G_{\Phi}(k = 0)$ against $T$, as shown on the right panel. The color of the markers match the color of the curves on the left. Note that variation of the mass before $t = 10$ is a huge peak resulting from the disturbance which is not shown in the picture.

which is very close to our data.

We note that the accuracy we can achieve for the value of $a$, the rate of exponential decay of the amplitude is much lower than frequency $c$ and the amplitude $b$ itself. This is particularly true at higher temperatures, where the oscillation amplitudes are very small, which explains the apparent larger fluctuations. However, it is clear that the point at which oscillatory behavior begins is shifted toward later times as temperature increases, a trend most apparent as we inspect figure 14, where the first trough has a reducing depth until it disappears altogether as initial temperature is increased. For that matter, there is not an obvious definition of a relative phase between oscillations with different initial temperatures, although simply by eye-balling the oscillations, it is very suggestive of a phase shift with temperatures.

The data presented in figure 17 corresponds to parameters chosen at $g = 0.4$, $\lambda = 0.6$, $\rho_0 = 0.1067$. At very low temperatures, the onset of oscillatory behavior appears to begin even before the time dependent disturbance is withdrawn, a feature that eventually disappears as temperature is increased.

Another feature demonstrated most clearly in figure 17 is that the height of the first peak responding to the time dependent disturbance increases as temperature is increased. We have checked that this is true for all our data sets.

In the left panel of figure 18 we show oscillations at 10 different temperatures with the same
FIG. 15. Coefficients $a, |b|, c, d$ of the fit function $f(t) = \exp(-at)b\sin(ct + 2\pi d) + e + ft$ fitting the data presented in figure 13 are plotted against temperature.

initial conditions as in figure 13, except that the pulse disturbance is more abrupt, set at $v = 1/3$. We note that the waveform looks much less regular at low temperatures corresponding to a regime in which the mass scale $m \ll v$. At higher temperatures corresponding to higher $m$, the waveform returns to sinusoidal. Suspecting that there are more than one Fourier component with significant amplitude, we inspect the power spectrum of these oscillations. A plot of the power spectrum of data obtained using initial temperature $T$ close to $T_p$ is shown in the right panel of figure 18. (i.e. The set of data corresponding to the magenta curve in figure 18). There is a distinct second peak in all the Fourier transforms, and we can track the variation of the frequencies of the two significant peaks with initial temperatures, shown in figure 19.

It is also of interest to inspect the Green’s function $G_\phi(k)$ at different times. A typical plot is shown in figure 20. At $t = -\infty$ the Green’s function is a thermal Green’s function. As the time-dependent disturbance sets in, one can see that all the departure from the thermal Green’s function occurs at low momenta. At late times long after the withdrawal of the disturbance, the Green’s function appears to approach the original thermal value.
FIG. 16. Coefficients $a, |b|, c, d$ of the fit function $f(t) = \exp(-at)b\sin(ct + 2\pi d) + e + ft$ fitting the data presented in figure 14 are plotted against temperature.

B. Remark: pulses in $g$

Let us also remark on the response of the system upon shaking the parameter $g$.

Consider $g$ as a function of time given by

$$g(t) = g_0 + \delta g(\tanh^2(vt) - 1).$$ (41)
FIG. 17. Left Panel: Oscillations of $m^2_\Psi$ as a function of time at 15 different initial temperatures, from low temperatures at the bottom of the picture to high temperatures at the top, at constant $\rho_0 = 0.1067$ (corresponding to $m_\Psi = 0.0041$ at $T = 1/100$, $g = 0.4$, $\lambda = 6/10$, and $a = 1$). Integral along $k_x$ and $k_y$ is each divided into 90 steps. The pulse parameter is taken as $v = 1/5$, $\delta \rho = 1/500$. These 15 initial temperatures correspond to 15 points on the equilibrium plot of $G_\Psi(k=0)$ against $T$, as shown on the right panel. The color of the markers match the color of the curves on the left. Note that in this picture we display the entire oscillations including the large peak. This is because at sufficiently low temperatures there are extra higher frequency oscillations that begin earlier.

The gap equation is then modified to

$$\lambda m^2_\Psi(t) - m^2_\Psi(t) = g(t),$$

$$m^2_\Psi(t) = \frac{A}{2B},$$

$$A = \int \frac{dk}{2\pi} k \left( 2(\Omega^\Psi_k(t))^2 - k^2 \right) + \frac{1}{2} \left( \frac{\Omega^\Psi_k(t)}{\Omega^\Psi(t)} \right)^2 G^\Psi(k,t) +$$

$$\frac{4}{\lambda} \left( \left( \frac{\Omega^\Psi_k(t)}{\Omega^\Psi(t)} \right)^2 - k^2 - \frac{g(t)}{\lambda} + \frac{1}{\lambda} \left( \frac{\Omega^\Psi_k(t)}{\Omega^\Psi(t)} \right)^2 \right) G^\Phi(k,t),$$

$$B = \int \frac{dk}{2\pi} k \left( G^\Psi(k,t) + \frac{2}{\lambda^2} G^\Psi(k,t) \right)$$

(42)

Below we find the plots at different $T$ in figure 21. The oscillations following a kick in $g$ are qualitatively the same as what happens when $\rho_s$ is taken as the time-dependent disturbance instead. In particular, the oscillations are sinusoidal with a decaying amplitude, and that the amplitude falls off as temperature increases. What is interesting however is that contrary to a time-dependent $\rho_s$, the initial large response to the disturbance before oscillatory behavior sets in has a large amplitude at low temperatures which decreases with increasing temperatures.
FIG. 18. Left Panel: 10 oscillations taking the same initial conditions as in figure 2 except that the pulse is more abrupt, with \( v = 1/3 \). At low temperatures corresponding to a regime \( m_\Psi \ll v \) the oscillation waveforms look much less regular. Right Panel: a typical plot of the power spectrum of the oscillations, obtained using the magenta curve on the left. The fit makes use of 11 significant frequencies in the power spectrum.

FIG. 19. Frequency variation of the higher and lower frequency peaks in the power spectrum with initial temperature.

IX. CONCLUSIONS

In the first part if the paper, we studied the non-equilibrium dynamics of SC and CDW order parameters in the t-J-V model, using a time-dependent Hartree-Fock computation. We examined
FIG. 20. A plot of $G_\Psi (k)$ as a function of $k$ at different times. The parameters take values $g = 0.4, \lambda = 0.6, \rho_0 = 0.1067$, and the pulse is characterized by $v = 1/5, \delta \rho = 1/5000$. In fact it corresponds to the pulse leading to oscillations displayed in the red curve (the third curve from the bottom of the left panel in figure 17).

FIG. 21. Oscillations of $m_\Psi^2$ as a function of time at 10 different initial temperatures. The parameters $g_0, \lambda$ and $T$ take the same values as the 10 plots in figure 13. Here $v = 1/5$ and $\delta g = 1/50$. 

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two setups: quench and pulse in the interaction parameters, and compared with a recent optical experiment.\textsuperscript{1} We used a decaying sinusoidal function to fit the oscillation which gives qualitatively good fitting parameters compared with the spectral analysis, other than the fact that near $T_c$, there are relatively large side peaks in the spectrum. When perturbing with the nearest neighbor Coulomb interaction $V$, we found an enhanced oscillation amplitude of the CDW order below the superconducting critical temperature in both setups. We interpret this enhancement as a competition between the charge order and superconducting order. The oscillation frequency is of the order of terahertz if we plug in the energy scale from the information of $T_c$. The frequency of the oscillations has a strong dependence on temperature, which is however different from that in the experiments. The change in frequency makes it subtle to define the relative phase between oscillations at different temperatures. But in the pulse case crossing $T_c$, if we choose a particular phase starting point in the fit function, we find a nearly $\pi$ phase shift, as is evident by eye from Fig. 7. When perturbing the exchange interaction $J$, in both setups, there is no obvious enhancement in the oscillation amplitude crossing $T_c$ because of the similar enhancement effects of $J$ on both SC condensation and charge density waves.

In the second part of the paper, we considered the dynamical evolution of a large $N$ NLSM inspired by Ref. 10 subjected to a short pulse like disturbance to mimic the effect of a pulse of laser radiation on the underdoped cuprates. We considered in detail oscillations of the system upon sending a short pulse in the coupling $\rho_s$ at different initial temperature $T$, helicity moduli $\lambda$ and relative energetic cost of superconductivity and charge density wave order $g$. We find that for disturbance with a rate of change $v$ taken at the same order as the effective mass scales of the system, the subsequent oscillation after the withdrawal of the disturbance is, to a very good approximation, sinusoidal with an exponentially decaying amplitude. The amplitude is greatly suppressed as initial temperature is increased, particularly beyond the maximal point in the equilibrium $G_{\Phi}(k = 0) - T$ plot. The characteristic oscillation frequency increases faster than linearly with increasing temperature. The onset of oscillatory behavior begins at a later time at large temperatures making a comparison of relative phases between different oscillations ambiguous, even though the data is suggestive of a phase shift as temperature changes. The increase in oscillation frequency, the presence of a phase shift and a strongly suppressed oscillation amplitude across the critical temperatures are features qualitatively consistent with the experiments.\textsuperscript{1} For more abrupt disturbances or at very low temperatures such that $m_{\Phi}$ is several orders of magnitudes less than $m_{\Phi}$, however, the oscillatory behavior is characterized by more than one frequency.
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Appendix A: Equations of motion

It is a simple matter to evaluate the commutators of the operators in Eq. (9):

\[
\begin{align*}
[N_1(k), \Delta_1(k)] &= \Delta_1(k) \\
[N_1(k), \Delta^+_1(k)] &= -\Delta^+_1(k) \\
[N_1(k), \Pi_1(k)] &= \Pi_1(k) \\
[N_1(k), \Pi^+_1(k)] &= -\Pi^+_1(k) \\
[N_1(k), P_1(k)] &= P_1(k) \\
[N_1(k), P^+_1(k)] &= -P^+_1(k) \\
[N_3(k), \Delta_1(-k)] &= \Delta_1(-k) \\
[N_3(k), \Delta^+_1(-k)] &= -\Delta^+_1(-k) \\
[N_3(k), \Pi_1(k)] &= -\Pi_1(k) \\
[N_3(k), \Pi^+_1(k)] &= \Pi^+_1(k) \\
[N_3(k), P_3(k)] &= P_3(k) \\
[N_3(k), P^+_3(k)] &= -P^+_3(k) \\
[\Delta_1(k), \Delta^+_1(k)] &= N_1(k) + N_3(-k) \\
[\Delta_1(k), \Pi_1(-k)] &= -P_1(k) \\
[\Delta_1(k), \Pi^+_1(k)] &= -P^+_3(k) \\
[\Delta_1(k), P^+_1(k)] &= \Pi^+_1(-k) \\
[\Delta_1(k), P_3(k)] &= \Pi_1(k) \\
[\Pi_1(k), \Pi^+_1(k)] &= N_1(k) - N_3(k) \\
[\Pi_1(k), P^+_1(k)] &= -\Delta^+_1(-k) \\
[\Pi_1(k), P_3(k)] &= \Delta_1(k) \\
P_1(k), P^+_1(k) &= N_1(k) + N_1(-k) - 1 \\
P_3(k), P^+_3(k) &= N_3(k) + N_3(-k) - 1
\end{align*}
\]

(A1)

and some others that follow under \( k \to -k \) and/or Hermitian conjugates.

A similar set of relations follow from \( 1 \to 2 \) and \( 3 \to 4 \), yielding a second SU(4) algebra. However, we will not need these because we will always assume \( \Delta_2 = -\Delta_1 \) and \( \Pi_2 = -\Pi_1 \).

Then we can use \( H_{MF} \) in Eq. (5) to obtain the equations of motion of the average values of the
operators in Eq. (9):

\[
\frac{d\Delta_1(k)}{dt} = -i \left[ -\epsilon_1(k)\Delta_1(k) - \epsilon_1(k)\Delta_1(k) + \frac{(3J - V)}{2} \Delta_2(N_1(k) + N_3(-k)) + \frac{3J + V}{2} (-\Pi_2 P_1(k) - \Pi_2 P_3(k)) \right]
\]

\[
\frac{d\Delta_1(-k)}{dt} = -i \left[ -\epsilon_1(-k)\Delta_1(-k) - \epsilon_1(-k)\Delta_1(-k) + \frac{(3J - V)}{2} \Delta_2(N_1(-k) + N_3(k)) + \frac{3J + V}{2} (-\Pi_2 P_1(k) - \Pi_2 P_3(k)) \right]
\]

\[
\frac{d\Delta_1^\dagger(k)}{dt} = -i \left[ \epsilon_1(k)\Delta_1^\dagger(k) + \epsilon_1(k)\Delta_1^\dagger(k) - \frac{(3J - V)}{2} \Delta_2^\dagger(N_1(k) + N_3(-k)) - \frac{3J + V}{2} (-\Pi_2 P_1^\dagger(k) - \Pi_2 P_3^\dagger(k)) \right]
\]

\[
\frac{d\Delta_1^\dagger(-k)}{dt} = -i \left[ \epsilon_1(-k)\Delta_1^\dagger(-k) + \epsilon_1(-k)\Delta_1^\dagger(-k) - \frac{(3J - V)}{2} \Delta_2^\dagger(N_1(-k) + N_3(k)) - \frac{3J + V}{2} (-\Pi_2 P_1^\dagger(k) - \Pi_2 P_3^\dagger(k)) \right]
\]

\[
\frac{d\Pi_1(k)}{dt} = -i \left[ -\epsilon_1(k)\Pi_1(k) + \epsilon_1(-k)\Pi_1(k) + \frac{(3J - V)}{2} (-\Delta_2 P_3(k) + \Delta_2 P_1(k)) + \frac{3J + V}{2} \Pi_2(N_1(k) - N_3(-k)) \right]
\]

\[
\frac{d\Pi_1^\dagger(k)}{dt} = -i \left[ \epsilon_1(k)\Pi_1^\dagger(k) - \epsilon_1(-k)\Pi_1^\dagger(k) - \frac{(3J - V)}{2} (-\Delta_2 P_3^\dagger(k) + \Delta_2 P_1^\dagger(k)) - \frac{3J + V}{2} \Pi_2^\dagger(N_1(-k) - N_3(k)) \right]
\]

\[
\frac{dP_1(k)}{dt} = -i \left[ -(\epsilon_1(k) + \epsilon_1(-k)) P_1(k) + \frac{(3J - V)}{2} \Delta_2(\Pi_1(-k) + \Pi_1(k)) - \frac{3J + V}{2} \Pi_2(\Delta_1(-k) + \Delta_1(k)) \right]
\]

\[
\frac{dP_1^\dagger(k)}{dt} = -i \left[ (\epsilon_1(k) + \epsilon_1(-k)) P_1^\dagger(k) - \frac{(3J - V)}{2} \Delta_2(\Pi_1^\dagger(-k) + \Pi_1^\dagger(k)) + \frac{3J + V}{2} \Pi_2^\dagger(\Delta_1^\dagger(-k) + \Delta_1^\dagger(k)) \right]
\]

\[
\frac{dP_3(k)}{dt} = -i \left[ -(\epsilon_1(k) + \epsilon_1(k)) P_3(k) + \frac{(3J - V)}{2} \Delta_2(\Pi_1(k) + \Pi_1^\dagger(-k)) - \frac{3J + V}{2} \Pi_2(\Delta_1^\dagger(k) + \Delta_1(k)) \right]
\]

\[
\frac{dP_3^\dagger(k)}{dt} = -i \left[ (\epsilon_1(-k) + \epsilon_1(k)) P_3^\dagger(k) - \frac{(3J - V)}{2} \Delta_2(\Pi_1(-k) + \Pi_1^\dagger(k)) + \frac{3J + V}{2} \Pi_2^\dagger(\Delta_1^\dagger(k) + \Delta_1^\dagger(-k)) \right]
\]

\[
\frac{dN_1(k)}{dt} = -i \left[ \frac{3J - V}{2} (\Delta_2^\dagger \Delta_1(k) - \Delta_2 \Delta_1^\dagger(k)) + \frac{3J + V}{2} (\Pi_2 \Pi_1(k) - \Pi_2 \Pi_1^\dagger(k)) \right]
\]

\[
\frac{dN_1(-k)}{dt} = -i \left[ \frac{3J - V}{2} (\Delta_2^\dagger \Delta_1(-k) - \Delta_2 \Delta_1^\dagger(-k)) + \frac{3J + V}{2} (\Pi_2 \Pi_1(-k) - \Pi_2 \Pi_1^\dagger(-k)) \right]
\]

\[
\frac{dN_3(k)}{dt} = -i \left[ \frac{3J - V}{2} (\Delta_2^\dagger \Delta_1(k) - \Delta_2 \Delta_1^\dagger(k)) + \frac{3J + V}{2} (\Pi_2 \Pi_1(k) - \Pi_2 \Pi_1^\dagger(k)) \right]
\]

\[
\frac{dN_3(-k)}{dt} = -i \left[ \frac{3J - V}{2} (\Delta_2^\dagger \Delta_1(-k) - \Delta_2 \Delta_1^\dagger(-k)) + \frac{3J + V}{2} (\Pi_2 \Pi_1(-k) - \Pi_2 \Pi_1^\dagger(-k)) \right]
\]

\[\tag{A2}\]

Also note that the operators $P_1 + P_2$ and $P_3 + P_4$ (and their Hermitian conjugates) commute with the original Hamiltonian $H$ for $V = 0$; these operators generate the pseudospin symmetry between the SC and CDW order parameters.

One quality that remains constant during the oscillation is the mean field energy $\langle H_{MF} \rangle$, if the interaction parameters are constant with respect to time, like during the time after the quench. This can be easily verified using Eq. (A2). And this can be used to check the validity of the numerics.

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\[ \langle H_{MF} \rangle = \sum_k \left[ \epsilon_1(k) N_1(k) + \epsilon_1(-k) N_3(k) + \epsilon_2(k) N_2(k) + \epsilon_2(-k) N_4(k) \right] \\
+ \frac{3J - V}{2} (\Delta_1 \Delta_2^* + \Delta_2 \Delta_1^*) + \frac{3J + V}{2} (\Pi_1 \Pi_2^* + \Pi_1^* \Pi_2) \tag{A3} \]

Notice that we have added back some subtractions terms to Eq. (5). Then the time derivative of \( \langle H_{MF} \rangle \) becomes

\[
\frac{1}{2} \frac{d \langle H_{MF} \rangle}{dt} = \sum_k -i \epsilon_1(k) \left[ \frac{3J - V}{2} (\Delta_2 \Delta_1(-k) - \Delta_2 \Delta_1^*(-k)) + \frac{3J + V}{2} (\Pi_2 \Pi_1(-k) - \Pi_2 \Pi_1^*(-k)) \right] \\
- i \epsilon_1(-k) \left[ \frac{3J - V}{2} (\Delta_2^* \Delta_1(-k) - \Delta_2 \Delta_1^*(-k)) + \frac{3J + V}{2} (-\Pi_2 \Pi_1(k) + \Pi_2 \Pi_1^*(k)) \right] \\
- i \frac{3J - V}{2} \Delta_2 \left[ -\epsilon_1(k) \Delta_1(k) - \epsilon_1(k) \Delta_1(k) + \frac{3J - V}{2} \Delta_2 (N_1(k) + N_3(k)) + \frac{3J + V}{2} (-\Pi_2 \Pi_1(k) - \Pi_2 \Pi_1^*(k)) \right] \\
- i \frac{3J + V}{2} \Pi_2 \left[ -\epsilon_1(k) \Pi_1(k) + \epsilon_1(-k) \Pi_1(k) + \frac{3J - V}{2} (-\Delta_2 \Delta_1^*(k) + \Delta_2 \Pi_1(k)) + \frac{3J + V}{2} \Pi_2 (N_1(k) - N_3(k)) \right] \\
- i \frac{3J + V}{2} \Pi_2 \left[ -\epsilon_1(k) \Pi_1^*(k) + \epsilon_1(-k) \Pi_1^*(k) - \frac{3J - V}{2} (-\Delta_2 \Delta_1^*(k) + \Delta_2 \Pi_1^*(k)) - \frac{3J + V}{2} \Pi_2^*(N_1(k) - N_3(k)) \right] \tag{A4} \\
= 0
\]

where we have assumed time-independence of \( J \) and \( V \), which is true in the quench case.
Appendix B: More simulations of the hot spot model

Here we describe multiple additional initial conditions for the hot spot model as a supplement to the main text.

In Fig. 22, we considered the quench case, where $\Delta V = 0.1, \Delta J = 0$. We still have amplitude enhancement below $T_c$, although the enhancement is not that big. Besides, there is also a phase shift upon crossing $T_c$.

![Fig. 22. Oscillation of CDW order parameter $\Pi$ (left) and SC order parameter $\Delta$ (right) as a function of time in the quench case, from low temperature at the bottom to high temperature at the top, temperatures are taken from 0.025 to 0.375 with 0.05 step. The initial value $J_0 = 1.2, V_0 = 0.9$, the quench is taken as $\Delta J = 0, \Delta V = 0.1$. The initial $T_c$ at equilibrium can be computed to be 0.25, the final $T_c$ to be 0.2.]

In Figs. 23–25, we showed the pulse case, where $\Delta V = -0.1, \Delta J = 0$. We have used both decayed sinusoidal fitting and spectral analysis to characterize oscillation properties. Fitting parameters are found to be similar to those in Figs. 7–10, although the phase shift has the opposite sign. In the spectral analysis, near $T_c$ the side peaks gets larger and become the main peak: that is why we observe frequency change against temperature in this region.
FIG. 23. Oscillation of CDW order parameter $\Pi$ (left) and SC order parameter $\Delta$ (right) as a function of time in the pulse case, from low temperature at the bottom to high temperature at the top, temperatures are taken from 0.025 to 0.375 with 0.05 step. The initial value $J_0 = 1.2$, $V_0 = 0.9$, the pulse is taken as $\Delta J = 0, \Delta V = -0.1$, $\omega = 1$. The initial $T_c$ at equilibrium can be computed to be 0.25, at the largest derivation $V = V_0 + \Delta V$, the corresponding equilibrium $T_c$ to be 0.33.

FIG. 24. Left to right: amplitude $a$, frequency $c$ and phase $d$ of the fit Eq. 11 fitting the data in Fig. 23 as a function of temperature. The blue line denotes initial equilibrium $T_c = 0.25$, the green line denotes at the largest derivation $V = V_0 + \Delta V$, the equilibrium $T_c = 0.33$.  

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FIG. 25. Left: spectral fitting of the left panel in Fig. 23. The inserted figure is the spectral power at $T = 0.225$ near $T_c$. Right: peak frequency from the left panel as a function of temperature.

In Fig. 26 - 28, we made the pulse sharper with $\omega = 4$. If we put energy scale $T_c \approx 75K$ in the simulation, this gives the pulse duration is about 25 fs. We also found similar amplitude enhancement and phase shift crossing $T_c$, except that here the overall oscillation amplitude gets smaller because of the shorter disturbance. With more simulations, we found that as long as the pulse duration does not exceed the oscillation period, these effects are quite robust.

FIG. 26. Oscillation of CDW order parameter $\Pi$(left) and SC order parameter $\Delta$(right) as a function of time in the pulse case, from low temperature at the bottom to high temperature at the top, temperatures are taken from 0.025 to 0.375 with 0.05 step. The initial value $J_0 = 1.2$, $V_0 = 0.9$, the pulse is taken as $\Delta J = 0$, $\Delta V = 0.1$, $\omega = 4$. The initial $T_c$ at equilibrium can be computed to be 0.25, at the largest derivation $V = V_0 + \Delta V$, the corresponding equilibrium $T_c$ to be 0.2.
FIG. 27. Left to right: amplitude $a$, frequency $c$ and phase $d$ of the fit Eq. 11 fitting the data in Fig. 26 as a function of temperature. The blue line denotes initial equilibrium $T_c = 0.25$, the green line denotes at the largest derivation $V = V_0 + \Delta V$, the equilibrium $T_c = 0.2$.

FIG. 28. Left: Fitting Fig. 26 using spectral analysis. The inserted figure is the power spectrum at $T = 0.225$ near $T_c$. Right: Peak frequency as a function of temperature.

Finally, we considered quench $J$ case in Fig. 29. However, now the behavior is quite different, as we discussed in the main text; $V$ rather than $J$ affects SC and CDW differently, which mimics the effect of the pump in the experiment.
FIG. 29. Oscillation of CDW order parameter $\Pi$ (left) and SC order parameter $\Delta$ (right) as a function of time in the quench case, from low temperature at the bottom to high temperature at the top, temperatures are taken from 0 to 0.5 with 0.1 step. The initial value $J_0 = 1.2$, $V_0 = 0.9$, the quench is taken as $\Delta J = 0.1, \Delta V = 0$. 

\[ \text{time} \]

\[ |\Pi| \]

\[ T \]

\[ 0 \]

\[ 5 \]

\[ 10 \]

\[ 15 \]

\[ 20 \]

\[ 0.0 \]

\[ 0.2 \]

\[ 0.4 \]

\[ 0.6 \]

\[ 0.8 \]

\[ 1.0 \]

\[ 0 \]

\[ 5 \]

\[ 10 \]

\[ 15 \]

\[ 20 \]

\[ 0.0 \]

\[ 0.2 \]

\[ 0.4 \]

\[ 0.6 \]

\[ 0.8 \]

\[ 1.0 \]
Appendix C: A different corner in parameter space for the O(6) model

In the main text, we have been inspecting a region of parameter space where $g$ is of the order of 1/10 of the lattice cutoff $a$. We present here results from a vastly different parameter space, in which we reduce $g$ by a factor of 10. At equilibrium, the correlation functions are plotted in figure 30, which has a much stronger resemblance to the classical results in.$^{10}$

![Equilibrium equal time correlation functions of \( \Phi \) at zero momentum vs temperatures at three different values of \( \lambda \) and \( g \) and \( m_\Phi = 1/500 \) at \( T = 1/100 \).](image)

In figure 31 below we present the oscillations at \( \lambda = 1, g = 0.04 \). We note that comparing with figure 17, we find that this feature where the initial large response to the disturbance changes sign close to the peak temperature \( T_p \) becomes a completely generic feature in this regime of parameter space. Note that right across the peak temperature (see blue and green curves in the middle of the right panel) the initial trough disappears, and this skipped trough leads to a more convincing \( \pi \) phase shift actually happening independently how the data is fitted. We note that in all our data sets collected this always happens close to \( T_p \), in this case and also \( g = 0.01 \), just above \( T_p \), and whereas in the case in figure 17, just below. Another feature is that ultimately the frequency still stays at roughly the scale set by \( T_p \) despite dramatically reducing \( g \) while keeping \( T_p \) roughly the same. The oscillation amplitude is highest close to \( T_p \) but decreases again as temperature is further lowered, although not as steeply as in the high temperature regime. The fit of the exponential decay at very high temperature is a gross underestimate, because the oscillation amplitude is
extremely small and that it is strongly affected by the starting point of the fit. We also present in figure 33 results of numerical fits of the oscillations by the fit function in equation 39. To improve the results of the fits, we note that close to the peak temperature, there are in fact two major Fourier components with similar amplitudes (see figure 32). Therefore, between $T = 0.5 - 0.6$ we modify the fit function to

$$f_{\text{modified}}(t) = e^{-at}(b \sin(\omega_1 t + d_1) + b_2 \sin(\omega_2 t + d_2))$$

We take the higher frequency peak and its amplitude in the plot of frequencies and amplitudes verses temperature in figure 33.

![Graph](image.png)

**FIG. 31.** Left Panel: Oscillations of $m^2_\Psi$ as a function of time at 13 different initial temperatures, from low temperatures at the bottom of the picture to high temperatres at the top, at constant $\rho_0 = 0.1058$ (corresponding to $m_\Psi = 0.002$ at $T = 1/100$, $g = 0.04$, $\lambda = 1$, and $a = 1$). Integral along $k_x$ and $k_y$ is each divided into 90 steps. The pulse parameter is taken as $\nu = 1/5$, $\delta \rho = 1/500$. These 13 initial temperatures correspond to 13 points on the equilibrium plot of $G_\Phi(k = 0)$ against $T$, as shown on the right panel. The color of the markers match the color of the curves on the left. Note that in this picture we display the entire oscillations including the large peak.
FIG. 32. Power spectrum at $T = 0.47$.

FIG. 33. Coefficients $a, |b|, c, d$ of the fit function $f(t) = \exp(-at)b\sin(ct + 2\pi d) + e$ fitting the data presented in figure 31 are plotted against temperature.


