Chemical potential for light by parametric coupling

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Usually photons are not conserved in their interaction with matter. Consequently, for the thermodynamics of photons, while we have a concept of temperature for energy conservation, there is no equivalent chemical potential for particle number conservation. However, the notion of a chemical potential is crucial in understanding a wide variety of single- and many-body effects, from transport in conductors and semiconductors to phase transitions in electronic and atomic systems. Here we show how a direct modification of the system-bath coupling via parametric oscillation creates an effective chemical potential for photons even in the thermodynamic limit. In particular, we show that the photonic system equilibrates to the temperature of the bath, with a tunable chemical potential that is set by the frequency of the parametric coupler. Specific implementations, using circuit-QED or optomechanics, are feasible using current technologies, and we show a detailed example demonstrating the emergence of Mott insulator–superfluid transition in a lattice of nonlinear oscillators. Our approach paves the way for quantum simulation, quantum sources, and even electronlike circuits with light.

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I. INTRODUCTION

The study of the thermodynamics of photons dates back to Planck [1]. Investigating blackbody radiation, he realized photons decay due to absorption into walls of their container, and therefore, no chemical potential appeared in his expression, in contrast to Gibbs’s thermodynamic expressions for other particles using the grand canonical ensemble. Later, it was understood that in the absence of absorbing walls, photon can acquire nonzero chemical potential, e.g., photon emission in semiconductors (LED) [2], and thus the useful concept of chemical potential can start to be applied to these systems [3–5]. Moreover, if photons are confined in a cavity and coupled to excitons, they form polaritons which also can thermalize [6–8]. More recently, it was shown that photons can thermalize with a nonzero chemical potential and form a Bose-Einstein condensate [9–12] when interacting with a nonlinear medium. However, finding a general solution to creating a chemical potential for light remains an open problem [13].

At the same time, photons provide an intriguing quantum degree of freedom for implementing quantum simulators [14–19] and observing quantum phases of matter [8]. In quantum simulation, one develops a quantum system with a controllable, known Hamiltonian, enabling simulation of problems that are exponentially difficult on a classical computer. This new paradigm covers a wide range of problems from chemistry [20] and quantum field theories [21] to strongly correlated electron systems, such as high-$T_c$ superconductors [22]. Recently, several theoretical works have shown that photonic systems can have nontrivial photonic states [23,24] and many-body effects with zero chemical potential [25–28]. In the presence of strong nonlinearity photonic system can exhibit blockade effect [29–31], which can fix the number of photons in the steady state. In particular, it was recently shown that under specific conditions (flat-band models and with an incompressibility at a certain particle number), photonic systems can be stabilized by single-photon pumping and parametric drive [32]. However, many phenomena that are interesting from a quantum simulation perspective involve thermalization in systems with a controllable chemical potential, as a key parameter in phase diagrams. Both are absent for photons.

Here, we propose a parametric scheme to address the issue of chemical potential and thermalization in photonic systems, extending preliminary concepts [33] and developing simpler approaches than current theory [34–36]. In particular, by parametrically coupling a photonic system to a thermal bath, we show that a photonic system can equilibrate to the temperature of the bath, with a tunable chemical potential given by the frequency of the parametric coupler. Therefore, this scheme makes it possible to control both the temperature and the chemical potential of a photonic system. We apply our scheme to two platforms, circuit-QED and optomechanical systems, where recent and spectacular progress has been made in controlling and using them in a few quanta regime. Finally, we conclude by considering how a photonic lattice implementing a Bose-Hubbard model can be driven through the Mott insulator–superfluid (MI-SF) transition [37] using this approach even in the presence of finite dissipation.

II. PARAMETRIC THERMALIZATION

We can understand thermalization via a system-bath picture, where the system of choice with Hamiltonian $H_S$ is coupled via $\lambda H_{SB}$ to a bath with Hamiltonian $H_B$ and initial state $\rho_B \propto \exp(-\beta H_B)$ [38,39]. Our scheme will follow this approach with one small modification: replace the coupling with a parametric coupling via $\lambda \rightarrow 2\lambda \cos(\omega_p t)$, where $\omega_p$ is the angular frequency at which the coupling is modulated. Therefore, the system-bath Hamiltonian takes the form ($\hbar = 1$),

$$H = H_S + 2\lambda \cos(\omega_p t) H_{SB} + H_B,$$

again with initial conditions $\rho_B \propto \exp(-\beta H_B)$. We assume that parametric drive can be characterized by a classical field which cannot be depleted. The parametric coupling will enable
up- and down-conversion of bath excitations to photons, which will lead to controlled chemical potentials.

To see this explicitly, we will assume that $H_{SB}$ is bi-linear, of the form

$$H_{SB} = \sum_j (\hat{a}_j + \hat{a}_j^\dagger) \hat{B}_j,$$

(2)

where $\hat{B}_j$ is a bath operator and there exists $\hat{a}_j, n_j$ such that $[\hat{a}_j, n_j] = \delta_{j,j}$, as occurs naturally for photons. This property defines particle numbers $n_j$ and total particle number $\hat{N} = \sum_j n_j$.

Let us consider what happens when the energy scales of the bath and system are small compared to $\omega$. Specifically, we assume that the system has a low frequency cutoff, and the bath has a low-high frequency cutoff $\nu_c$. Furthermore, we will decompose $H_S$ into $H_S^+ + H_S^-$, where $H_S^\perp$ includes all terms that do not commute with the total number of particles in the photonic system, given by $\hat{N} = \sum_j \hat{a}_j^\dagger \hat{a}_j$. Therefore, $H_S^\perp$ is the part of the Hamiltonian that conserves the total number of particles. In this regime, we move to a rotating frame with the unitary transformation $U = \exp(-i t \omega_p \hat{N})$. The transformed system Hamiltonian becomes

$$U^\dagger H_S U - i U^\dagger U \approx H_S^\perp - \omega_p \hat{N},$$

(3)

where we have neglected $U^\dagger H_S^\perp U$ by making the rotating wave approximation (RWA), requiring $|H_S^\perp| \ll \omega_p$.

Meanwhile, the bath Hamiltonian remains the same, while the system bath coupling terms become

$$[\hat{a}_j + \hat{a}_j^\dagger + (e^{-2i \omega_p t} \hat{a}_j + e^{2i \omega_p t} \hat{a}_j^\dagger)] B_j \approx [\hat{a}_j + \hat{a}_j^\dagger] \hat{B}_j. \quad (4)$$

The key approximation is again the RWA to neglect $e^{2i \omega_p t} \hat{a}_j^\dagger$ type terms, consistent for a bath whose two-point bath correlation function $\langle B_i(t) B_j(t) \rangle$ has a cutoff frequency $\omega_c < \omega_p$. This provides our definition of a low frequency bath for this paper, with $H_B^\perp \equiv \sum_j [\hat{a}_j + \hat{a}_j^\dagger] \hat{B}_j$ the system-bath coupling in the RWA.

Through this set of transformations, and the rotating wave approximation, we have a new system-bath Hamiltonian which takes the traditional form

$$H = H_S^\perp - \mu \hat{N} + \lambda H_{SB} + H_B,$$

(5)

where we identity $\mu \equiv \omega_p$ as the chemical potential. For weak coupling $\lambda$ and an infinite bath at inverse temperature $\beta$, we expect the system to thermalize in the long-time limit to a density matrix

$$\rho \approx \exp[-\beta (H_S^\perp - \mu \hat{N})],$$

(6)

i.e., the distribution is exactly that of the grand canonical ensemble.

The key idea of our approach is to parametrically couple a low-temperature, low frequency bath to a set of high frequency modes. The parametric coupler up-converts bath excitations to photons and down-converts photons to bath excitations, as shown in Fig. 1. This leads to thermalization of photons, as long as the bath thermalization rate and the coupling rate between the bath and photons is faster than other photonic decay rates.

![FIG. 1. (Color online) Thermal bath with modes $\hat{b}_j$ and response functions with a cutoff $\nu_c$ can be parametrically coupled to a higher frequency (optical) system with modes $\hat{a}_j$ near the frequency $\nu_p$. Additional loss via the high frequency bath can lead transport from the parametric bath through the system to the high frequency bath.](image)

### III. IMPLEMENTATIONS

Now we show that such a scheme, which provides both thermalization and a finite chemical potential for photons, can be implemented in circuit-QED systems for microwave domain photons and using optomechanics for optical domain photons. Following the Caldeira-Leggett model, in the context of circuits [40,41], we consider the bath to be a collection of transmission lines which can be described by a quasicontinuum of harmonic oscillators. The bath Hamiltonian is given by

$$H_B = \sum_v \omega_v \left( \hat{b}_v^\dagger \hat{b}_v + \frac{1}{2} \right),$$

(7)

where $\hat{b}_v^\dagger$ is the creation operator of an electromagnetic field quantum at mode $v$ with frequency $\omega_v$. We assume that the transmission lines are in thermal equilibrium, and thus, $\langle \hat{b}_v^\dagger \hat{b}_v \rangle = \frac{\omega_v}{\omega_v + \delta_{v,nu_c}}$. We consider that each mode of the photonic system is coupled to the bath using nondegenerate parametric amplifiers, through three-wave mixing. While many configurations can implement this concept [42], we focus on the conceptually cleanest case: a Josephson parametric amplifier in a Wheatstone bridge configuration [43], as depicted in Fig. 2.

Examining the details of the JJ-Wheatstone parametric coupler, we assume that each junction has a large area, and hence, a large capacitance, so that its charging energy can be ignored. In this approximation, the energy $U$ of the JJ-Wheatstone bridge is [44]

$$-4E_J \left[ \cos \left( \frac{\Phi_s}{4\phi_0} \right) \frac{\Psi_X}{2\phi_0} \cos \left( \frac{\Psi_Y}{2\phi_0} \right) \cos \left( \frac{\Psi_Z}{2\phi_0} \right) \cos \left( \frac{\Psi_Z}{2\phi_0} \right) \sin \left( \frac{\Psi_X}{2\phi_0} \right) \sin \left( \frac{\Psi_Y}{2\phi_0} \right) \sin \left( \frac{\Psi_Z}{2\phi_0} \right) \right],$$

where we have taken all four JJ’s to have the same $E_J$, and $\phi_0 = \Phi_0/(2\pi), \Phi_s = \hbar/(2e)$ being the superconducting flux quantum. Setting $\Phi_s = \Phi_0/2$ by choice of flux bias, and assuming the mode intensities $\Psi_X, \Psi_Y, \Psi_Z \ll \Phi_0,$ consistent with moderate to low characteristic impedance circuits, we can expand $U$ in $\psi_i = \Psi_i/\Phi_0, i \in \{X,S,Z\}$ to third
order [45]:

\[ U = -2\sqrt{2} E_j + M \left( \psi_1^2 + \psi_2^2 + \psi_3^2 \right) + g \psi \psi S \psi Z. \] (8)

Here \( M = \sqrt{2} E_j \pi^2 \) and \( g = -2\sqrt{2} E_j \pi^3 \).

A transmission line is connected inductively to the coupler mode \( \Psi_X = \Phi_1 - \Phi_2 \) through inductance \( L \). The modes, assumed to be in thermal equilibrium at a temperature \( T \), act as a bath. The (microwave) photonic system is coupled to the mode \( \Psi_X = \Phi_1 - \Phi_3 \), while the mode \( \Psi_Z = \Phi_1 - \Phi_3 + \Phi_2 - \Phi_4 \) is externally modulated as \( \Psi_Z = \lambda \Phi_0 \cos(\omega_0 t + \phi) \), where \( \lambda \) is the dimensionless amplitude of the modulation and controls the system-parametric bath coupling strength.

Let \( C_I = CL \) be the capacitance of the transmission line, with \( C \) being its capacitance per unit length. Because of the presence of the transmission line, \( \Psi_X = \sum \psi_i \sqrt{\frac{1}{2\pi_i}} (b_i + b_i^\dagger) \). Here, \( \psi_i \) is a dimensionless parameter that depends on the boundary conditions at \( z = L \). For our particular coupling—current-flux—we expect \( \psi_i \sim \sin(k_i L) \) and, in the weak coupling limit, \( \psi_i \propto v \). Ignoring coupling between different transmission line modes, the system Hamiltonian is

\[ H_S + H_B + \lambda \cos(\omega_0 t + \phi) \sum \psi_i (b_i + b_i^\dagger) \Psi_S, \] (9)

where \( h_v = \sqrt{\frac{1}{2\pi_i}} \psi_i \). This then directly produces our model Hamiltonian for generating a chemical potential, where the density of states \( J(v) = h(v)^2 \rho(v) \propto v \), i.e., an Ohmic bath [41].

For the optical domain, we need a different parametric process. A convenient one is the optomechanical coupling between motion of a mirror and the frequency of light in a cavity formed by the mirror. This example case has been worked in partial detail in Ref. [33]. The key idea is for a pump field to take the radiation pressure coupling \( a^\dagger a x \) to a fast oscillating coupling via \( a \rightarrow a + \alpha e^{-i\omega_0 t} \), producing a parametric coupling to the phonon “bath” with frequency \( \omega_0 \). The details and benefits of the optomechanical approach will be considered in a separate work.

Note that in any experimental implementation, one needs to filter out the pump photons from the signal system photons. This can be easily achieved by using different polarization or spatial modes of the photonic system on each site. Alternatively, in certain schemes, one can reject the pump by frequency filtering. For example, in the Mott insulator case, discussed later in this article, the pump has higher frequency than the prepared Mott state, and therefore, the pump can be filtered out by frequency selection.

IV. BATH DISCUSSION

We now examine our assumption of a cutoff in the bath degrees of freedom, as well as a strictly parametric system-bath coupling. For simplicity, we divide the bath modes into three, independent sets of modes, and consider coupling to a single system mode \( a \). Given a parametric coupling at frequency \( \nu \), the low frequency modes of the bath, \( b_j \), are defined as those with natural resonance frequencies \( \omega_j \leq \nu /2 \). The “natural” modes, \( c_j \), are those with frequencies \( \nu /2 < \omega_j \leq 3\nu /2 \). The “doubly rotating” modes, \( d_j \), are those with frequencies \( \omega_j > 3\nu /2 \). Thus the more general system-bath interaction is

\[ H_{SB} = [A + \lambda \cos(v t)] \sum \left( f_j (b_j + b_j^\dagger) + g_j (c_j + c_j^\dagger) + h_j (d_j + d_j^\dagger) \right). \] (10)

We now move to an appropriate rotating frame, with \( a \rightarrow ae^{-i\omega t}, b_j \rightarrow b_j, c_j \rightarrow c_j e^{-i\omega t}, \) and \( d_j \rightarrow d_j e^{-i\omega t} \). With the assumption of weak coupling \((A, \lambda \) small), we look at the rotating wave approximation for the different couplings:

\[ H_{SB,b} = [A + \lambda \cos(v t)] (ae^{-i\omega t} + a^\dagger e^{i\omega t}) \sum f_j (b_j + b_j^\dagger) \]

\[ \rightarrow \frac{1}{2} (A + a^\dagger) \sum f_j (b_j + b_j^\dagger), \] (11)

\[ H_{SB,c} = [A + \lambda \cos(v t)] \sum g_j (c_j e^{-i\omega t} + c_j^\dagger e^{i\omega t}) \] (12)

\[ \rightarrow A \sum g_j (c_j + c_j^\dagger a), \] (13)

\[ H_{SB,d} = [A + \lambda \cos(v t)] (ae^{-i\omega t} + a^\dagger e^{i\omega t}) \]

\[ \rightarrow \frac{1}{2} (A + a^\dagger) \sum h_j (d_j e^{-i\omega t} + d_j^\dagger e^{i\omega t}) \] (14)

\[ \rightarrow \frac{1}{2} (A + a^\dagger) \sum h_j (d_j e^{-i\omega t} + d_j^\dagger e^{i\omega t}) \] (15)

By breaking up the bath into three different frequency regions, we see that the natural and doubly rotating frequency regions both lead to a system-bath interaction of the quantum optics type, i.e., that of a beam splitter interaction \( a^\dagger c + c^\dagger a \). For these
portions of the system-bath interaction, we may then proceed in deriving the master equation in the usual way [46,47], and find, in appropriate limits, a decay of excitations of a rate $\kappa \sim A^2|g|^2\rho(\nu) + 4\lambda^2|J|^4\rho(2\nu)$, where $\rho(\nu)$ is the bath density of states near the parametric modulation frequency $\nu$. We now investigate the remaining portion of the system-bath to investigate this portion of the system-bath interaction in a specific setting, to illustrate the emergence of a Mott insulator-superfluid transition in a photonic lattice.

V. LATTICE MODEL AND MASTER EQUATION

We consider now what happens to a lattice of coupled, interacting photonic resonators, coupled to both a parametric bath at inverse temperature $\beta$ and nominal coupling rate $\gamma$ and to a high frequency (loss) bath with loss rate $\kappa$. For simplicity, we consider only strong on-site repulsion $\kappa$ and to a high frequency (loss) bath with loss rate $\kappa$. Formally defines an operator that reduces or keeps constant the energy, and $x_j(t) = c_j(t) + c_j^\dagger(t) + x_0$, with the last term time independent and neglected in what follows.

Taking independent, Ohmic baths for each coupling term, we have

$$S_{ij}(t) = \frac{\delta_{ij}}{\pi} \int_0^\infty dv J(v)(N_{ah}(v) + 1)e^{-i\nu t} + N_{ah}(v)e^{i\nu t}.$$  

with the effective spectral density $J(v) = v e^{-v/\gamma}$, $N_{ah}(v) = 1/[\exp(\beta v) - 1]$, where $\beta$ is the inverse temperature of the parametric bath and $\gamma \gg U$ is a high frequency cutoff that is irrelevant to the rest of our calculation. At this point, we get terms in the master equation of the form $S_{ij}(t)(c_i(\tau)c_j(\tau - t)\rho_S)$ and terms of the form $S_{ij}(t)(c_i(\tau)c_j^\dagger(\tau - t)\rho_S)$. The former will have phase evolution at a zero frequency as a function of $\tau$, and will be neglected in a rotating frame approximation. The latter will also have such terms, except for those with $\omega_{kl} = \omega_{1l}$, i.e., energy-degenerate transitions. Keeping only these transitions immediately takes us to the usual golden rule result: transitions with a positive energy difference $\nu$ occur with a rate $J(v)N_{ah}(v)$ and transitions with a negative energy difference have the rate $J(v)[N_{ah}(v) + 1]$. Thus, when the energy levels of the system are well resolved, we can derive a super operator describing both photon loss and coupling to the parametric bath. Using the commutation of $H_S$ with $N$ (the total photon number), we get transitions from $k$ to $l$ with rates that depend on whether the total photon number of the two states differs by $+1$ or $-1$ as

$$\Gamma_{k\rightarrow l}^+ = \gamma(N_{ah}(|\epsilon_k - \epsilon_l|) + \Theta(\epsilon_k - \epsilon_l))\sum_i |\langle l|a_i|k\rangle|^2,$$

$$\Gamma_{k\rightarrow l}^- = [\gamma(N_{ah}(|\epsilon_k - \epsilon_l|) + \Theta(\epsilon_k - \epsilon_l)) + \kappa]\sum_i |\langle l|a_i|k\rangle|^2,$$

where $\gamma = \gamma_0[|\epsilon_k - \epsilon_l|]$ for the Ohmic bath case, $\gamma_0$ represents the overall strength of the coupling, and $\Theta$ is the Heaviside step function. We have gone back to the physical couplings $a_i$ rather than the many-body energy lowering operator $c_j$ in order to make clear the special role loss via the high frequency bath plays in Eq. (20).

The superoperator takes Lindblad form with these rates leading to a rate equation in the energy eigenbasis. Solving this numerically for a case of four coupled sites (Fig. 3), we can immediately see an intuitive understanding of the two types of decay processes. The first type, which increases photon number, corresponds to the decay of holes (if the energy of the higher photon number state is lower in the rotating frame) or the creation of particles (if otherwise). The second type decreases photon number, and includes both creation of
holes via loss and via the parametric bath; consequently, we expect a greater rate for the second process, which will lead to a particle-hole temperature asymmetry as shown below. The simulations themselves correspond to fixing a maximum total particle number per site, finding the eigenenergies of the dissipation-free model, calculating the decay rates in Eqs. (19) and (20), determining the steady state of the master equation, and for that steady state, finding the probability of each state (shown in the inset to Fig. 3), and estimating the Mandel $Q = \sqrt{\langle \delta n^2 \rangle - \langle n \rangle^2} / \langle n \rangle$ parameter and the average hopping $\langle a \rangle \equiv \sqrt{\langle \langle a_i^\dagger a_i \rangle \rangle}$ (shown in Fig. 4).

VI. STRONG INTERACTION EXPANSION

We now take a simpler form of the superoperator describing both photon loss and coupling in the case of a single resonator site ($J = 0$) to get an analytical handle on the process. That is, we evaluate Eqs. (19) and (20) in the single site case. Specifically, defining $E_0(n) = J/2n(n - 1) - \mu n$, the sign of $\Delta E(n) = E_0(n + 1) - E_0(n) = nU - \mu$ determines both the direction of decay and the thermal bosonic enhancement factor $N_{th}(\Delta E(n))$. Thus $\Gamma_{n \rightarrow n+1} = \gamma f_+(n), \Gamma_{n+1 \rightarrow n} = (n+1)\kappa + \gamma f_-(n)$ with

$$f_+(n) = (n + 1)[N_{th}(\Delta E(n))] + \Theta(-\Delta E(n)), \quad f_-(n) = (n + 1)[N_{th}(\Delta E(n))] + \Theta(\Delta E(n)),$$

and $\gamma = \gamma_0|\Delta E(n)|/U$ for the Ohmic bath case. The change from $N_{th}$ to $N_{th} + 1$ that occurs in these two factors with the change in sign of $\Delta E(n)$ arises from having both co- and counter-rotating terms in the system bath coupling.

One consequence of the strong interaction (sometimes called strong coupling in the Mott insulator literature) limit ($J \rightarrow 0$) is an analytical form for the steady state. Specifically, we recover a form of detailed balance, where the probability of a transition on a site from photon number $n$ to $n + 1$ is given by $f_+(n)$, while the transition from $n + 1$ to $n$ is $\gamma f_-(n) + (n + 1)\kappa$. This gives, in steady state, a set of ratios

$$p_1 = \frac{f_+(0)}{f_+(0) + \kappa/\gamma},$$

$$p_2 = \frac{f_+(1)}{f_+(1) + \kappa/\gamma},$$

...\hspace{1cm} (25)

where the correction from a thermal distribution arises from the term $\kappa/\gamma$, which depends on the energy difference via $\gamma$. We can characterize this for two regimes. First, when $\Delta E(n)$ is positive (it costs energy to add a photon), we expect the ratio $p_{n+1}/p_n = N_{th}^{(p)}(n)/N_{th}^{(p)}(n+1)$. This defines the bosonic occupation as seen by particle addition as

$$N_{th}^{(p)} = \frac{N_{th}(\Delta E(n))}{1 + \kappa/\gamma}.$$

Thus, when particles cost energy, photon loss reduces the effective temperature of the system.
Similarly, when \( \Delta E(n) \) is negative, we expect \( p_{n+1}/p_n = (N^{(h)}_{\text{eff}} + 1)/N^{(h)}_{\text{eff}} \), which defines the bosonic occupation as seen by hole addition:

\[
N^{(h)}_{\text{eff}} = \frac{[N_{\text{th}}((\Delta E(n)) + \kappa/\gamma)]}{1 - \kappa/\gamma}.
\]

Here, photon loss increases the energy, and thus increases the effective temperature of the system. Furthermore, any hope of a thermal description will necessarily break down for \( \kappa/\gamma \geq 1 \).

Having established that a Mott insulator-like phase emerges from the single site picture, we can ask how the asymmetry of particles and holes changes the standard picture of the edges of the Mott lobes, by using a picture of free particles and holes above the \( n_0 \) particle-per-site Mott state \( |\Psi_M \rangle \propto \prod_i (a_i^\dagger p_i \text{vac}) \), i.e., using small \( J \) perturbation theory in the strong interaction limit. A crucial difference from the standard treatment \cite{48} is the use of an implicit finite lifetime to such excitations due to the coupling to both parametric and high frequency baths.

When \( J \) exceeds damping and dephasing, we can no longer use a master equation appropriate to a single site. Specifically, as we want the parametric bath to resolve the kinetic terms in the Hamiltonian, we require \( \gamma = \gamma_0(J/U) \ll J \). We can, however, characterize the particle or hole occupation for a wave vector \( k \) in the dilute limit (where particle hole collisions are neglected) by using our \( N^{(ph)}_{\text{eff}} \), and we can ask over what domain of parameter space is the combined occupation of particles and holes small compared to one per site. Here we rely upon the standard picture of particle and hole energies to order \( J^2/U \), neglecting loss-induced changes to the energy differences, consistent with \( \kappa \ll \gamma \ll J \). The energy of a particle(hole) of wave vector \( k = 0 \) above the Mott state is given by Ref. \cite{48} and reproduced here to order \( J^2/U \):

\[
\Delta E^{(p)} = -z J (n_0 + \frac{1}{2}) + n_0 U + \mu + \frac{z J^2}{2U} n_0 (5n_0 + 4) - \frac{z^2 J^2}{U} n_0 (n_0 + 1),
\]

\[
\Delta E^{(h)} = -z J (n_0) - (n_0 - 1) U + \mu + \frac{z J^2}{2U} (n_0 + 1)(5n_0 + 1) - \frac{z^2 J^2}{U} n_0 (n_0 + 1),
\]

where \( z \) is the number of nearest neighbors.

We can then calculate the average particle and hole expectation values including both the parametric bath and the high frequency (photon loss) bath, and find that these lowest energy modes have just \( N^{(p)}_{\text{eff}} \) and \( N^{(h)}_{\text{eff}} \) with the above \( \Delta E^{(ph)} \). The boundary of the phase would then correspond to this effective occupation approaching unity (at which point we may expect a macroscopic occupation of particles and/or holes in the system, taking us far from the Mott state). This boundary is shown for two different values of \( \kappa/\gamma \) in Fig. 4; as \( \kappa \) increases, the lobes become asymmetric, consistent with additional hole creation via particle losses.

We now consider what near equilibrium picture can emerge, and in particular focus on a picture with two reservoirs (particles and holes) at different temperatures due to loss into the high frequency bath. In the limit of \( \kappa \to 0 \), we recover the usual picture of an equilibrium system, and get a critical temperature defined as

\[
T_c^{(0)} = \frac{1}{k_B \log 2} \min[\Delta E^{(h)}, \Delta E^{(p)}].
\]

However, including the nonequilibrium effects, we instead have for the parametric bath temperature the requirement

\[
T \leq T^{(ne)}_c = \frac{1}{k_B} \min \left[ \frac{\Delta E^{(h)}}{\log \left( \frac{2(\gamma^{(h)}_0 - 1)}{\gamma_p} \right)}, \frac{\Delta E^{(p)}}{\log \left( \frac{2(\gamma^{(p)}_0 - 1)}{\gamma_p} \right)} \right],
\]

where \( \gamma^{(h,p)}_0 \) depends on \( \Delta E^{(h,p)} \) via \( J(\gamma) \).

Further analysis of the particle-hole picture at finite temperature will no doubt elucidate additional physics for this nonequilibrium system, following perhaps the efforts of Refs. [49,50]. In addition, an appropriate mean field theory including modifications of the system-bath coupling could provide insight into the applicability of such theories for describing nonequilibrium systems.

**VII. CONCLUSION**

Providing a robust chemical potential for light allows for classical and quantum systems to access a wide variety of heretofore forbidden domains. Crucially, our approach allows one to build from well established theoretical tools for nonequilibrium problems with chemical potential imbalances, such as occurs in circuits and cold atom systems, rather than the thornier problems associated with driven steady-state systems more typical to the quantum optical domain. From a quantum simulation perspective, this simplification makes the state preparation problem much more straightforward than existing approaches, and yields a mechanism for robust quantum simulation of condensed matter and chemistry problems with light. In addition, our parametric coupling scheme has a wide range of potential implementations, all of which are accessible with current technology, and enables a variety of practical applications in the context of nonclassical sources in the microwave and optical domain that operate more in analogy to a diode than to a pumped dissipative steady-state system.

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**APPENDIX**

As a simple test of these concepts, we implement numerically a model for the intermediate time behavior of a two-level system (qubit) coupled to a bosonic bath via a parametric coupling. The usual picture of quantum Brownian motion \cite{51} has a set of bath modes coupled linearly through their position variables \( x_\omega \) with constant \( \bar{\omega} \) and mass \( m_\omega \),

This leads to the effective spectral density \( J(\omega) = \rho(\omega) \bar{\omega}^2/m_\omega \), where \( \rho(\omega) \) is the density of states. Our goal will be to well approximate such a bath with a discrete set of modes. Before engaging in that, we mention some rescaling of
the problem appropriate to simulation. First, we rewrite the system-bath coupling \( \tilde{g}_\omega x_\omega (a + a^\dagger) \) in terms of bath creation and annihilation operators with \( x_\omega = \sqrt{\frac{1}{2\omega_0}} (b_\omega + b_\omega^\dagger) \). This defines \( g_\omega = \tilde{g}_\omega \sqrt{\frac{1}{2\omega_0}} \), and \( J(\omega) = 2 \rho(\omega) g_\omega^2 \).

Our goal is to approximate the bath such that the correlation function and the commutation relation of the bath are as close to the desired approximate bath as possible. Specifically, for our quantum Brownian motion bath with a cutoff function \( f(\omega) \) considered in this work, we assume the time-ordered spectral function used in Sec. V:

\[
S(\tau) = \frac{1}{\pi} \int_0^\infty dv J(v) [(N_{ih}(v) + 1)e^{-iv\tau} + N_{ih}(v)e^{iv\tau}] \\
\approx \sum_{j=1}^N \frac{w_j J(\omega_j)}{\pi f(\omega_j)} [(N_{ih}(\omega_j) + 1)e^{-i\omega_j \tau} + N_{ih}(\omega_j)e^{i\omega_j \tau}],
\]

where the approximation of the integral as a finite sum arises from Gaussian quadrature over \( N \) orthogonal polynomials under the function \( f(\omega) \) to find the set \( \{\omega_j\} \) and the associated weights \( w_j \), and \( N_{ih}(\nu) = 1/|\exp(i\beta \nu) - 1| \). We remark that in the case of the Ohmic bath with exponential cutoff function, the appropriate choice is the Laguerre polynomials.

A simple reinterpretation of this formula is that of a discrete harmonic oscillator (quantum Brownian motion) bath with frequencies \( \omega_j \) and a coupling constant

\[
g_j = \sqrt{\frac{w_j J(\omega_j)}{f(\omega_j)}}.
\]

This approximation is immediately amenable to numerical techniques via direct integration of the Schrödinger equation.

In practice, exponential cutoffs at the relevant frequencies are unlikely as the superconductors work well into the GHz domain for our implementation. Thus we consider a polynomial cutoff function induced by filtering the Ohmic bath with a low-pass filter, such as a capacitor in parallel with the resistor forming the Ohmic bath for our circuit case. The impedance of this system becomes \( Z = R/(1 + i\tau_{RC} \omega) \), where \( \tau_{RC} = RC \) is the characteristic time of the RC circuit. Therefore, the real part of the impedance, which appears in the spectral noise [41], leads to a natural modification of the effective spectral density: \( J(\omega) \rightarrow J(\omega) \frac{1}{1 + \tau_{RC}^2 \omega^2} \). We neglect imaginary contributions to the circuit by assuming they are renormalized in the system Hamiltonian. We note that for an optomechanical implementation such cutoff functions arise from the cavity Lorentzian and can have a similar functional form—quadratic suppression at high frequency.

We simulate the following simple case numerically to illustrate our system. Working with the discrete bath approximation and a photon-blockade-regime cavity with an effective two-level system description with Pauli matrices \( \sigma_z \), etc., we write

\[
H = \omega_0 \sigma_z/2 + [A + \lambda \cos(\mu t)] \sum_j g_j \sigma_z (b_j + b_j^\dagger) + \omega_j b_j^\dagger b_j,
\]

with \( g_j = \sqrt{\frac{w_j J(\omega_j)}{1 + \omega_j^2 \tau_{RC}^2}} \). The parameters \( A \) and \( \lambda \) represent the relative strength of the regular exponential decay bath and the additional parametric bath terms oscillating at frequency \( \mu \).

We take \( \tau_{RC} = 4/\omega_0 \) and \( \omega_{cutoff} = 2.5\omega_0 \), and find that decreasing or increasing \( \omega_{cutoff} \) by even a factor of two does not appreciably change the results presented below. We also work in units of time given by \( 1/\omega_0 \). For improved computation speed, we truncate the bath Hilbert space to a maximum of two bosonic excitations, and confirm post-facto that simulations produce only slightly more than one bath excitation, consistent with the truncation.

We first test the purely Ohmic case, taking \( \tau_{RC} \rightarrow 0, A = 0.5/\sqrt{17} \), and \( \lambda = 0 \) (no parametric bath). We find exponential decay with a time scale \( \gamma^{-1}_0 = 16.8(4) \). Furthermore, this decay is well approximated (with around \(< 1\% \) errors) up to times \( t \gtrsim \{30, 50, 65, 100\} \) for \( N = \{35, 50, 70, 100\} \). The fitted decay rate is independent of \( N \) in this range of values, consistent with our approximation scheme.

We then consider \( A = 0.5 \) with the filter on \( \tau_{RC} \rightarrow 4 \), leading to a slightly reduced decay rate due to the non-Ohmic nature of the bath near \( \omega_0 \) from the cutoff filter. Still, exponential decay is observed over two decades with \( \gamma^{-1}_a = 18.7(5) \), and residuals are at the 3\% level or less, largely due to corrections to exponential decay at long times from non-Ohmic bath behavior. The decay from an initially excited state \( |\uparrow\rangle \) into the zero temperature bath is shown in red in Fig. 5.

After these simple tests of our model system, we move to the more complicated regime of a parametrically coupled bath. Taking \( \lambda = 0.5 \) and \( \omega_{cutoff} = 0.5 \) (no parametric bath), we start the system in the lower-energy spin state, \( |\downarrow\rangle \). We calculate over 100 time units three different values of our chemical potential parameter, \( \mu = \{0.9, 1.0, 1.1\} \), and plot the resulting \( \langle \sigma_z \rangle \) as a function of time.

**FIG. 5.** (Color online) Time-dependent simulation of a two level system with natural frequency \( \omega_0 \) interacting normally (red) and parametrically (green, yellow, blue) with a high-frequency filtered bath. For an initial excited state \( \langle \sigma_z \rangle = 1 \), coupling to the bath leads to exponential decay when the oscillating frequency of the bath is set to zero, as shown in red. However, for an initial ground state \( \langle \sigma_z \rangle = -1 \), turning on the parametric coupling to the bath such that it oscillates at frequencies \( \mu = \{0.9 \text{ (blue)}, 1.0 \text{ (yellow)}, 1.1 \text{ (green)\}ω_0} \) leads to inversion of the spin when \( \mu > \omega_0 \), as predicted by our more general theoretical model.
We expect the system to approach spin up for $\mu > 1$ according to the chemical potential derivation given in the first part of this paper, and find these expectations confirmed in this simple numerical experiment (Fig. 5).