Ultrafast Excitation Exchange in a Maxwell Fish-Eye Lens

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The strong coupling of quantum emitters to a cavity mode has been of paramount importance in the development of quantum optics. Recently, also the strong coupling to more than a single mode of an electromagnetic resonator has drawn considerable interest. We investigate how this multimode strong coupling regime can be harnessed to coherently control quantum systems. Specifically, we demonstrate that a Maxwell fish-eye lens can be used to implement a pulsed excitation exchange between two distant quantum emitters. This periodic exchange is mediated by single-photon pulses and can be extended to a photon-exchange between two atomic ensembles, for which the coupling strength is enhanced collectively.

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Introduction.—The coherent transfer of an excitation between different quantum systems is an essential process in quantum optics, with applications in quantum communication [1], quantum computation [2], and quantum networks [3]. At the heart of many implementations lies the coupling to monochromatic light fields, permitting Rabi oscillations between the electromagnetic field and quantum emitters [4].

Going beyond the paradigmatic Jaynes-Cummings model, the coupling of quantum emitters to more than a single mode has recently drawn considerable interest. A strong coupling to several modes is reached when the emitter-mode coupling strength becomes comparable to the free spectral range [5]. This multimode strong coupling regime (also known as the superstrong coupling regime) has been at the focus of recent theoretical studies [5–10] and experimental realizations, using photonic [11–18] or phononic modes [19,20].

The profound impact of single-mode effects on quantum technology raises the question of whether mechanisms of similar scope can also be engineered making use of many modes in a cavity. In contrast to the single-mode case, the dynamics in multimode cavities are, in general, more complex as they are dictated by a complex interference of all involved modes, whose individual properties are determined by the geometry of the underlying resonator. This, in turn, suggests that a suitable choice for the cavity geometry is the key to tame the dynamics and harness it for the coherent control of quantum systems.

In this Letter, we demonstrate that, with a Maxwell fisheye (MFE) lens [21,22] as a resonator, it is possible to coherently transfer an excitation between two distant quantum emitters when making use of their strong coupling to multiple modes. As a gradient index lens, the MFE lens comprises a radially changing refractive index and comes with the remarkable property that any point in the lens has a corresponding focal point mirrored at the lens's center. From the viewpoint of geometrical optics, light rays propagating in the lens form circular arcs, and all rays emitted from one point converge at the antipodal point after reflecting from the lens's circular boundary mirror [see Fig. 1(a)]. The presence of a continuum of pairs of focal points provides decisive advantages when considering distributed emitters or emitters that cannot be considered pointlike. Because of their extraordinary properties, MFE lenses have been studied, e.g., for imaging [23-27] and



FIG. 1. (a) 2D Maxwell fish-eye lens with two opposing quantum emitters. Within the lens, all rays emerging from one atom form circular arcs due to the radial refractive index gradient and converge at the second atom [23]. All light paths that connect the two atoms with one reflection at the bounding mirror have the same optical length. (b) Illustration of relevant coordinates and parameters in the lens. (c) Radial refractive index gradient within the lens [cf. Eq. (2)].

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coherent perfect absorption [28], as well as for radiation emission [29]; also generalizations of the MFE concept have been proposed [30–33]. In the context of quantum optics, the MFE lens has recently come into focus for the creation of entanglement by the dipole-dipole interactions it may mediate [34]. In contrast, we consider the MFE lens in a regime, where the light field in the cavity exhibits strong memory effects.

Within the multimode strong coupling regime, it has been shown both theoretically and experimentally that a *single* excited atom may emit its excitation into the electromagnetic field and (partially) reabsorb the corresponding singlephoton pulse when it returns to the atom [7,10,16,35]. Hence, when such a pulse is emitted by an excited atom in a MFE lens, one may expect the pulse to be refocused at the opposite point. In the following, we consider a second atom placed at this opposite point and study the dynamics of the two quantum emitters. Taking into account the quantized electromagnetic field inside the cavity, we demonstrate explicitly the emission of a single-photon pulse, its propagation through the cavity, and its refocusing at the second atom. In the most straightforward implementation, the pulse does not get fully absorbed by the second atom, which we attribute to a mismatch in the temporal pulse shape [36–39]. By spectrally engineering the mode-emitter coupling, we can increase the efficiency of this exchange to a value near unity, thus implementing a pulsed passive swap operation [40] that relies on the interaction with many modes. On that basis, we then show that the dynamics of two emitters is readily translated to two atomic ensembles, where the coupling strength is collectively enhanced [12]. Our results demonstrate how concepts from classical photonics can be translated to the quantum regime when emitters couple strongly to many modes.

The model.—We model the above system using the Glauber-Lewenstein quantization [34,41] where the dynamics is described by the Hamiltonian

$$H = \hbar \omega_{\rm a} \sum_{i} |\mathbf{e}_{i}\rangle \langle \mathbf{e}_{i}| + \hbar \sum_{lm} \omega_{l} a_{lm}^{\dagger} a_{lm} - \sum_{i} \boldsymbol{d}_{i} \boldsymbol{E}(\boldsymbol{r}_{i}), \quad (1)$$

comprising the two-level atoms, the quantized field, and the coupling between them, respectively. Here, ω_a is the atoms' transition frequency and the excited (ground) state of the *i*th emitter is denoted by $|e_i\rangle$ ($|g_i\rangle$). Since we will extend the discussion from two atoms to two atomic ensembles, we keep the number of atoms general at this point. The frequencies of the cavity modes are denoted by ω_l and the modes' creation and annihilation operators obey the usual commutation relations. We note that, while the above interaction Hamiltonian is conventionally used, there has been an ongoing debate regarding its gauge invariance for multimode systems [42,43]. The key results of our present work are, however, left unchanged by this issue.

The radially varying refractive index in the MFE lens is given by

$$n(\mathbf{r}) = \frac{2n_0}{1 + (r/R)^2},$$
(2)

where *r* is the in plane distance from the lens's center and *R* is the radius of the mirror, which forms the circular lens's boundary (see Fig. 1). Without loss of generality, we set the minimal refractive index $n_0 = 1$ throughout the Letter and assume the transverse thickness *b* of the lens to be far below the lens's radius *R*. The relevant transverse eigenmodes $f_{lm}(\mathbf{r})$ in this refractive index profile are hence polarized transversally to the plane and known analytically [34]. The modes are indexed by l = 1, 2, 3, ... with angular index m = -(l-1), -(l-3), ..., (l-1) and the modes' eigenfrequencies are $\omega_l = c\sqrt{l(l+1)}/(Rn_0)$, i.e., the *l*th mode is *l*-fold degenerate. The emitters couple to the modes via the transverse electric field,

$$\boldsymbol{E}(\boldsymbol{r}) = \boldsymbol{E}^{+}(\boldsymbol{r}) + \boldsymbol{E}^{-}(\boldsymbol{r}), \qquad \boldsymbol{E}^{-}(\boldsymbol{r}) = \left[\boldsymbol{E}^{+}(\boldsymbol{r})\right]^{\dagger},$$
$$\boldsymbol{E}^{+}(\boldsymbol{r}) = i \sum_{lm} \sqrt{\frac{\hbar\omega_{l}}{2\varepsilon_{0}}} a_{lm} \boldsymbol{f}_{lm}(\boldsymbol{r}), \qquad (3)$$

and the dipole moment, $d_i = d(\sigma_i^{\dagger} + \sigma_i)\hat{z}, \sigma_i = |g_i\rangle\langle e_i|$, of the atoms. We assume a Gaussian cutoff in frequency for the atomic dipole moments $d^2 = d_0^2 \exp[-(\omega - \omega_a)^2/(2\omega_c^2)]$, with cutoff frequency ω_c [7,35,44]. Further, we employ the rotating wave approximation, i.e., we neglect nonexcitation-preserving terms $a\sigma$, $a^{\dagger}\sigma^{\dagger}$. Note that this approximation is not necessarily guaranteed to be compatible with the multimode dynamics [45,46], since ultrastrong coupling effects may eventually become relevant [11,17,35,47–49] (an explicit justification for the validity of the approximation for our work will thus be provided at the end of the Letter). The mode-independent coupling prefactors are summarized in the constant $g = d_0 \sqrt{\omega_a^3/(\varepsilon_0 b c^2 \hbar)}$. While g parametrizes the coupling strength between modes and atoms, it does not represent the couplings' actual values, which are reduced by mode specific prefactors [see Eq. (3)]. We further note that, for coupling strengths far below the free spectral range, Rabi oscillations between two emitters can be recovered in accordance with [34].

To study a pulsed excitation transfer, we consider a single excitation only, i.e., we can represent the wave function as $|\psi(t)\rangle = \sum_i c_i(t)|0...01_i0...0\rangle_a|0...0\rangle_{ph} +$ $\sum_{lm} c_{lm}^{ph}(t)|0...0\rangle_a|0...01_{lm}0...0\rangle_{ph}$, comprising atomic and photonic degrees of freedom, and the dynamics can be calculated by standard tools (see the Supplemental Material [50]). Using this wave function, the observable electric field intensity $\langle E^-(r)E^+(r)\rangle$ is directly accessible and subsequently serves as an observable for the field in the cavity [51,52].



FIG. 2. Atom and multimode dynamics in a Maxwell fish-eye lens. The excited state populations of the two emitters are shown as a function of time. The insets display the intensity expectation value of the quantized electric field $\langle E^-(\mathbf{r})E^+(\mathbf{r})\rangle$ for five different times, as indicated by the links to the curves. Note that peaks of exceeding height were clipped in the insets. A video illustrating the dynamics is provided in the Supplemental Material [50].

Single-photon pulses.-We consider a MFE lens of radius $R = 3\lambda_a = 6\pi/\omega_a$ with two emitters placed opposite to one another at $r_a = 0.6R$ (see Fig. 1). The coupling strength is $g = 0.5\omega_a$ and the cutoff frequency is set to $\omega_{\rm c} = 2\omega_{\rm a}$; i.e., the cutoff function is flat around the atomic transition frequency [7], leading to about 10⁵ modes participating in the dynamics. Figure 2 shows the resulting excitation probabilities of the two atoms as a function of time when the left emitter is initially excited. Over some time, the emitter transfers the excitation to the cavity and returns to its ground state. Subsequently, the excitation probability of the right-hand emitter starts to increase when the time $t = l_{opt}/c = R\pi n_0/c$ is reached. This time precisely corresponds to the propagation duration of light from one emitter to the other one, including one reflection at the cavity boundary [50].

Here, since we have full access to the quantized photonic degrees of freedom, we can directly access the dynamics inside the cavity between the initial emission and the following absorption. The insets in Fig. 2 show the expectation value of the intensity at different times. During the initial emission, a radially symmetric pulse emerges. This single-photon pulse propagates and is reflected wherever it strikes the cavity boundary. Notably, due to the refractive index gradient, the wavefront deforms during propagation. Since the emitters' positions constitute a pair of focal points, the pulse eventually refocuses on the second emitter and is partly absorbed. Note that the spot size of the focused pulse may be smaller than the atomic wavelength λ_a since the emitter also interacts with high frequency modes and acts as a drain for the light field [53,54]. Before the absorption by the right emitter, both emitters are close to their ground state. The absorption thus witnesses significant memory effects in the light field that would not be compatible with a Born-Markov approximation [34].

While the pulse emitted by the left atom is successfully refocused on the right atom, the excitation is not fully transferred there. This can be understood by considering that the absorption process is the time reverse of the emission process, which has been studied in detail in free-space setups [36-39]. Since the emission from the left atom is of an exponential-like shape, the resulting pulse has an extended decaying tail. For perfect absorption, however, the time-reversed pulse, i.e., an increasing tail followed by the wavefront peak, would need to arrive at the right emitter. For the dynamics in Fig. 2, the mismatch in the temporal pulse shape results in an overlay between the absorption and the reemission dynamics for the right atom; i.e., when the excitation probability of the right atom reaches its peak, a new pulse is already propagating toward the left atom (see rightmost inset in Fig. 2). This process leads to a fast degradation of the pulsed excitation exchange between the two atoms.

Coherent exchange by spectral engineering.—Previous work suggests that a viable path to overcome the restrictions of the pulse shape is to artificially modify the pulse emission and absorption [36-39,55]. In the following, we show that in our setup we can achieve a nearly coherent transfer of the excitation by spectrally engineering the coupling strength between the emitters and the modes of the resonator. Specifically, we modify the couplings such that the atoms couple to a reduced number of about 200 modes (amounting to about 15 frequencies with degeneracies) close to the resonance frequency ω_a . Such an engineered coupling strength may be reached, e.g., by off resonantly driving Raman transitions with higher lying states [55,56]. Alternatively, by employing Bragg mirrors [57–59], the coupling may effectively be reduced to modes lying in the band gap, thus realizing the coupling to the restricted range of modes.

We parametrize the engineered coupling strength by reducing the cutoff frequency to $\omega_{\rm c} = 0.1\omega_{\rm a}$ [see Fig. 3(b)], which results in a symmetrically distributed Gaussian coupling. In Fig. 3(a), we consider the corresponding dynamics for the left emitter initially excited. We now find that, after the propagation, the excitation is almost completely reabsorbed by the second emitter. This greatly enhanced absorption efficiency can be understood when considering the shape of the emitted pulse shown in the intensity distributions in Fig. 3(a). Because of the spectral engineering, the emitted pulse is now symmetric around its peak and, therefore, time-reversal symmetric. Consequently, the pulse is absorbed as efficiently by the right emitter as it was emitted by the left emitter. Without the spectral engineering, the coupling constants showed a pronounced asymmetry, since, first, the relevant modes' frequencies were bounded by zero from below, but



FIG. 3. Coherent excitation exchange in the Maxwell fish-eye lens. (a) Structured in analogy to Fig. 2. All parameters but the cutoff frequency $\omega_c = 0.1\omega_a$ are left unchanged as compared to Fig. 2. Because of the narrower range of modes participating in the dynamics, a high exchange efficiency is achieved. (b) Illustration of the spectral engineering ($\omega_c = 0.1\omega_a$) in comparison with the cutoff used in Fig. 2. A video illustrating the dynamics is provided in the Supplemental Material [50].

extended far beyond the atomic transition frequency, and, second, the coupling constants obeyed their natural frequency dependence $\sim \sqrt{\omega_l}$ [see Eq. (3)]. Even with the spectral engineering, we see that the excitation probability at the right atom peaks slightly below unity, which can be shown to be due to an imperfect time-reversal symmetry of the emitted pulse. This remaining infidelity can be reduced by optimizing the coupling, the cutoff and the geometrical parameters, together with the details of the coupling constants' frequency dependence (see the Supplemental Material [50]).

Note that, while the spectral engineering reduced the number of modes, the multimode nature of the dynamics remains apparent by the pulsed emission and absorption. Further, the emitter-cavity system remains strongly non-Markovian, as in between the emission and absorption both emitters reach their ground state and the complete excitation is reversibly stored in the electromagnetic field.

Since both atoms emit pulses that can be absorbed efficiently, an autonomous periodic excitation swapping is established. Remarkably, since any point is a focal point in the MFE lens, the pulses between the emitters do not suffer from dispersion during propagation. The mechanism studied in Fig. 3 further works universally also for MFE lenses with larger diameters.

Collective excitation exchange.—Experimentally, it may be hard to reach the multimode strong coupling regime with single emitters. It is thus worthwhile to point out that this regime has recently been reached through collective enhancement of the coupling between an ensemble of atoms and a cavity [12].



FIG. 4. Collective excitation exchange between two atomic ensembles in a Maxwell fish-eye lens. (a) Spatial distribution of the emitters in the lens. The two ensembles opposing each other consist of N = 100 atoms each. The atoms are normal distributed with a standard deviation of $\sigma = 0.02R$ around the two centers at $r_a = 0.6R$ at opposite positions in the cavity. (b) Collective excitation probabilities of the ensembles $|C_{\text{left}(\text{right})}|^2 = \sum_i |c_{i,\text{left}(\text{right})}|^2$ as a function of time. Initially, the excitation is distributed among the left ensemble as a symmetric Dicke state. The individual coupling strength of each atom of the ensemble is $g = 0.05\omega_a$, while the other parameters are analogous to Fig. 3.

To showcase that the pulsed excitation exchange by means of dielectric cavities also applies to collective multimode strong coupling, we replace each emitter by a cloud of atoms. As for the single emitters, we place the centers of the ensembles opposite to one another in the MFE lens. Within each ensemble, the atoms are normal distributed around the ensemble center [see Fig. 4(a)].

In Fig. 4, we consider N = 100 atoms per ensemble and reduce the coupling strength by a factor of $\sqrt{100}$ as compared to Fig. 3. When the left ensemble is initially prepared in a symmetric Dicke state, $|\psi(0)\rangle =$ $\sum_{i=1}^{N} |0...01_i 0...0\rangle_{a,left} |0...0\rangle_{a,right} |0...0\rangle_{ph} / \sqrt{N}$, we can clearly observe a pulsed excitation swapping between the two ensembles comparable to Fig. 3. This shows that the exchange mechanism is robust when a distributed ensemble is considered. Even when the distribution of the atoms is further broadened, the pulsed dynamics remains intact, although with reduced efficiency. On the other hand, the dynamics in Fig. 4(b) can be brought arbitrarily close to the one in Fig. 3(a) by narrowing the ensemble distribution.

Rotating wave approximation.—Since the relevant mode frequencies may deviate significantly from the atomic transition frequency ω_{av} , it is not a priori clear whether counterrotating terms can be neglected in the interaction between atoms and modes. In the following, we therefore explicitly verify the applicability of the rotating wave approximation to the setup of Fig. 3, which represents the principal result of our Letter. In first order, the counterrotating terms couple the single-excitation subspace to the three-excitation subspace. Hence, we take into account in our simulation all states that contain at most three excitations and monitor the cumulative population $P(|3\rangle)$ of the states with exactly three excitations as a function of time.



FIG. 5. Population of the three-excitation subspace $P(|3\rangle)$ (blue) when counterrotating terms are taken into account. The dynamics of the atomic populations (gray) is shown in the background for comparison to Fig. 3. All parameters are identical to Fig. 3.

Figure 5 shows that $P(|3\rangle)$ remains around 5×10^{-3} ; i.e., the dynamics is largely unaffected by counterrotating terms.

Conclusions.—We demonstrated that a Maxwell fish-eye lens can be used to transfer an excitation between distant quantum emitters in the multimode strong coupling regime. Our analysis shows that this exchange relies on the emission of single-photon pulses and can be highly efficient with the application of judicious spectral engineering of the atom-cavity coupling. In the absence of quantum emitters, the Maxwell fish-eye lens is well understood in classical photonics; our study shows that, in the multimode strong coupling regime, this intuition also applies to the resonator's quantum properties, thus providing a promising platform for multimode quantum physics. Interesting extensions to more than two emitters could be implemented with generalizations of the traditional MFE lens [30–33].

The following numerical packages have been employed in this work: [60,61].

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