Four-wave mixing in ultracold atoms using intermediate Rydberg states

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Ultracold Rb atoms were used to demonstrate nondegenerate four-wave mixing through a Rydberg state. Continuous $5S_1/2 - 5P_{1/2}$ two-photon excitation to the Rydberg state was combined with an $nD_{3/2}$-tuned laser in a phase-matched geometry. The angular dependence, spatial profile, and dependence on detuning were investigated, showing good agreement with theory. Under optimum conditions 50% of the radiation was emitted into the phase-matched direction. The directional emission is promising for studying quantum effects in blockaded atom clouds.

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The concept of using the strong dipole-dipole interactions between Rydberg atoms to blockade coherent excitation of single atom qubits [1,2] has recently been experimentally realized [3]. Experimental signatures of the extension of the blockade concept to mesoscopic atomic clouds [4] include observations of suppressed excitation [5–7] and evidence for quantum critical behavior [8,9]. Coherent coupling between ground and Rydberg atoms, an essential capability for exploitation of blockade concepts, has only recently begun to be realized [8,10–12].

One intriguing consequence of mesoscopic blockade is the collective emission of single photons, that when properly phase matched are predicted to be emitted into a diffraction-limited solid angle. This phenomenon could be used to produce a directional single-photon-on-demand source [13] or for fast quantum-state detection or transmission [14]. Toward this end, we recently demonstrated state-selective stimulated emission detection of Rydberg atoms [15] and showed that it can be used to study population dynamics of ultracold Rydberg clouds. The use of light for ultracold Rydberg detection has also been accomplished using electromagnetically induced transparency [10] and shown to be very sensitive to atom-atom interactions.

In this paper we present the generation of coherent light beams using nondegenerate four-wave mixing through intermediate Rydberg states in an ultracold atomic gas. In a low density gas we observe as much as 25% of the emitted light isolated in a diffraction-limited solid angle under conditions of resonant Rydberg excitation, and 50% when slightly off-resonant. The noncollinear excitation geometry used here is naturally compatible with producing the spatially localized excitations needed to realize collective coherent single-photon emission. When coupled with higher-density atom clouds [16] and exciting states with stronger dipole-dipole interactions [17], this experiment should enter the collective single-photon regime.

The four-wave mixing process occurs within a $^{87}$Rb magneto-optical trap (MOT), with density $\sim 10^{10}$ cm$^{-3}$, as described in Ref. [15]. The first photon ($k_1$) comes from a 15 mW laser locked 500 MHz above the $5S_{1/2} - 5P_{1/2}$ transition at 780 nm. 5 mW of 480 nm light from a frequency doubled, amplified 960 nm ECDL is locked to a external Fabry-Perot cavity, and used as the second step $k_2$ of the excitation process and takes the atom to (or near) an $nD_{3/2}$ Rydberg state. The third photon $k_3$ comes from a cavity-locked 75 mW 1015 nm laser, which brings the atoms from the $nD_{5/2}$ Rydberg state to the $6P_{3/2}$ state. This results in decay photons from the $6P_{3/2}$ state to the ground state at 420 nm. The relevant energy levels can be seen in Fig. 1(a).

The directions of the three laser beams are chosen to satisfy phase-matching conditions, as shown in Fig. 1(b). In principle, a wide variety of possible geometries can be chosen, but we selected this particular one to give a large angle between $k_2$ and $k_3$, allowing for a small spatial overlap volume if desired. (A 90° angle is possible, giving the smallest possible overlap volume, but is incompatible with our vacuum chamber.) This results in a phase-matched direction for the exiting 420 nm photons ($k_4$) which is 3° from $k_2$. The direction of $k_4$ is very insensitive to the direction of $k_1$. Two Hamamatsu H7360-01 photon counters are used to detect these 420 nm photons, with a series of filters placed to elimi-

FIG. 1. (Color online) (a) Energy levels for four-wave mixing. The $k_1$ and $k_2$ photons produce two-photon excitation to an $nD$ Rydberg level. The $k_3$ laser couples the Rydberg level to the $6P$ state. Atoms in the $6P$ radiate coherently to the ground state to complete the four-wave mixing process. (b) Phase-matching geometry.

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nate all other wavelengths of light from the detectors. The detection efficiency of the filters and photodetectors is 3%. One detector is placed along an arbitrary or “off-axis” direction, while a second is placed along the anticipated phase-matched or “on-axis” direction. The waists of the three beams were \((w_1, w_2, w_3) = (4.0, 0.58, 0.83)\) mm, comparable to the MOT cloud size of 0.74 mm, so that most of the MOT atoms participate in the four-wave mixing process. Other important nonlinear effects, such as stimulated hyper-Raman scattering [18] and parametric four-wave mixing [19], are predicted to be relatively negligible due to the small Rydberg matrix elements and thin optical depth.

Phase-matched emission was initially seen for Rydberg principal quantum number \(n = 28\) by moving \(k_1\) far off the phase-matched direction and confirming nearly equal \(k_2\) detection rates in the on- and off-axis directions. Then as \(k_1\) was rotated in to the phase-matched direction a factor of 10 jump in the count rate on the on-axis detector was observed, with no change in the off-axis count rate. The width of acceptance for phase matching when the angle of \(k_1\) was rotated was found to be \(\delta \theta_0 = 1.6 \pm 0.2\) mrad, slightly larger than the expected 1.2 mrad.

We scanned a razor blade across the phase-matched output and recorded the count rate as a function of position. The waist is measured with a diffraction limited beam with a waist of 0.39 ± 0.06 mm at the atoms. An iris placed in front of the phase-matched detector reduced the solid angle of the on-axis detector by a factor of 11 while changing the count rate by only 9%. All further data presented here are taken with this iris in place.

When exciting to a different \(n\)-level Rydberg state, different frequencies are involved, so the phase-matched geometry will vary slightly. In fact, a change of only a few \(n\) levels at fixed laser beam angles is enough to move the phase-matching condition outside the angular bandwidth. After changing from 28D\(5\_2\) to 58D\(5\_2\), \(k_1\) was rotated until phase matching was again attained, and the rotation angle was found to be \(4.4 \pm 0.3\) mrad. This is consistent with the theoretical value of 4.1 mrad.

Neglecting propagation effects, in the absence of blockage we can consider the four-wave mixing process to result from the spatial phasing of the atomic dipole moments as a result of the three driving fields \((d_i) = (d_0(r_i)) \exp[i(k_1 + k_2 - k_3) \cdot r_i]\), where \(d_i\) is the dipole moment of the \(i\)th atom and \(d_0\) its magnitude as a function of atomic position \(r_i\). The fields from the atomic antennas constructively interfere in the far field to produce an electric field a large distance \(R\) away of

\[
E(\phi) = n(d_0)^2 k_0^2 \frac{e^{i k_0 R}}{R} \left(\frac{\pi w^2}{2}\right)^{3/2} e^{-\left(\pi w^2/\lambda^2\right)^2} \phi^2.
\]

We have assumed that the dipole moments are oriented perpendicular to the plane of the lasers, and that the effective spatial distribution of the dipoles is Gaussian with standard deviation \(w/2\) and peak density \(n\). The angle \(\phi\) is measured with respect to the phase-matched direction.

The ratio of on-axis to off-axis powers is obtained by integrating the intensity deduced from Eq. (1) over the angular distribution, and comparing to the spontaneously radiated power \(P_o\) from the 6\(P\) state

\[
\frac{P_{pm}}{P_o} = \frac{N_e^2 c^2 (d_0)^2 k_0^2 w^2}{N_i \hbar c k_0 \Gamma \rho_p} \frac{\sigma_{sp}}{w^2 \rho_p^2},
\]

where \(N_e = n(\pi w^2/2)^{3/2}\) is the effective number of participating atoms, \(\rho_p\) is the fraction of atoms in the 6\(P\) state, and \(\sigma_{sp}\) is the 6\(P\)-5\(S\) optical coherence density matrix element.

From Eq. (2) we expect the phase-matched fraction to be dependent on the number of atoms, since the phase-matched power is a nonlinear process. To verify this, the percentage of light in the phase-matched direction was measured as a function of the number of atoms in the MOT, with the results shown in Fig. 2. The expected nonlinear response of the four-wave mixing can be seen as a linear increase in the phase-matched fraction with the number of atoms at small atom numbers. As the cloud becomes optically thick, the phase-matched light is partially scattered due to the linear susceptibility of the atoms, and this scattered component, up to 24% for these conditions, is observed as additional off-axis light. This is taken into account in our analysis, resulting in slight modifications to Eqs. (1) and (2).

We model the coherence and population seen in Eq. (2) using an effective three-level density matrix approach:

\[
i\dot{\rho}_r = \frac{\epsilon_s}{2} \sigma_{rg} \rho_{rg} + \frac{\epsilon_s}{2} \sigma_{pg} \rho_{pg} - \frac{\epsilon_s}{2} \sigma_{pr} \rho_{pr} - \frac{i \rho_r}{\tau_r},
\]

\[
i\dot{\rho}_g = \frac{\epsilon_s}{2} \sigma_{rg} \rho_{rg} + \frac{\epsilon_s}{2} \sigma_{pg} \rho_{pg} - \frac{i \rho_g}{\tau_g} + \frac{i \rho_p}{\tau_p},
\]

\[
i\dot{\rho}_p = \frac{\epsilon_s}{2} \sigma_{pr} \rho_{pr} - \frac{\epsilon_s}{2} \sigma_{pg} \rho_{pg} + \frac{i \rho_p}{\tau_p},
\]

\[
i\sigma_{rg} = -\left(\delta_s + \frac{i \gamma_r}{2}\right) \sigma_{rg} + \frac{\epsilon_s}{2} (\rho_g - \rho_r) + \frac{\epsilon_s}{2} \sigma_{pg},
\]

\[
i\sigma_{pr} = \left(\delta_s + \frac{i \gamma_r}{2}\right) \sigma_{pr} + \frac{\epsilon_s}{2} (\rho_r - \rho_p) - \frac{\epsilon_s}{2} \sigma_{pg},
\]

FIG. 2. (Color online) The fraction of light emitted in the on-axis direction increases with the number of atoms in the MOT, as expected from the nonlinear character of four-wave mixing. The dashed line shows the model prediction, taking rescattering at large optical depths into account.

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The $5S$ state “$g$” and the Rydberg state “$r$” are coupled by an effective two-photon Rabi frequency $\epsilon_2 = \Omega_1 \Omega_2 / 2 \Delta$ ($\sim 4$ kHz) obtained by adiabatic elimination of the $5P_{3/2}$ state. The Rydberg state is then coupled to the $6P_{3/2}$ state by $\epsilon_3$ with Rabi frequency $\epsilon_3 \sim 1$ MHz. In order to simulate the effects of the substantial linewidths of the lasers used ($\sim 1$–3 MHz), plus other possible additional broadening mechanisms, we gave the $\sigma_g$ and $\sigma_p$ coherences effective homogeneous broadening factors of $\gamma_p = 6$ MHz, $\gamma_p = 2$ MHz [20]. These were chosen to reproduce the observed linewidths for two-photon excitation and non-phase-matched four-wave mixing, respectively. In addition, the Rydberg state was assumed to have a shortened lifetime ($\tau_r = 80$ to 40 $\mu$s depending on the $n$ level) due to the effects of superradiance [15], and the $6P$ state was assumed to decay at its spontaneous rate ($\tau_s = 121$ ns). Since the hyperfine coupling in the Rydberg state is negligible [21], the effects of nuclear spin were accounted for by calculating effective Rabi frequencies assuming nuclear spin conservation in the Rydberg state. Given measured intensities and atom numbers, the model then makes absolute predictions of the on- and off-axis count rates observed, and is typically within a factor of 3 of the observations, limited mainly by uncertainty in the absolute atom number and effective excitation rate. By examining the ratio of the on- and off-axis counters information can be gathered which is less sensitive to these uncertainties.

The on- and off-axis count rates as a function of $k_3$ are shown in Fig. 3. For off-axis light the $F’=3$ count rate is slightly higher than the other $F’$ levels, with the $F’=2$ count rate being 65% of the $F’=3$ rate. The model predicts 48%. The on-axis light, on the other hand, has an observed 10:1 ratio for the two states. This is partly explained by a factor of $\sqrt{2}$ greater dipole matrix element for the $6P_{3/2}(F’=3)$ $\rightarrow 5S_{1/2}(F=2)$ transition as compared to the corresponding matrix element from the $F’=2$ state. In addition there is a competition between the coherent deexcitation from the Rydberg state and the various decoherence processes that occur in the Rydberg state. This further favors the larger Rabi coupling to the $F’=3$ state. From these effects the model predicts a ratio of 7.7, in reasonable agreement with the observations.

Figure 4 shows the on-axis fraction of the emitted light as the intensity of beam $k_3$ is varied, for two different hyperfine levels of the $6P$ state. The different dipole matrix elements for the two hyperfine levels are accounted for by scaling the $F’=2$ intensities by a factor of 0.48 and the on-axis fraction by 2. The model prediction is sensitive to the assumed broadening of the $r$-$p$ coherence, with the data favoring a narrower ($\sim 1.25$ MHz) linewidth for this coherence than deduced from the off-axis spectroscopy. This discrepancy could suggest a narrower transition linewidth than observed, a larger number of atoms or smaller interaction region than calculated, or a fault of the model in its treatment of the broadening mechanisms. Despite this, the increase of the on-axis fraction with increased intensity still shows that the four wave mixing process is competing with decoherence. At higher $k_3$ powers, the atoms are removed from the Rydberg state before they can undergo any decoherent process.

In order to look for another sign of decoherence, the population in the Rydberg state was reduced by detuning $k_3$ from the Rydberg-$6P_{3/2}$ resonance. Then the $k_3$ frequency is varied and two resonances result. The first occurs when the excitation lasers are on resonance with the Rydberg state, giving significant Rydberg populations. The second resonance occurs when the three-photon process to the $6P$ state is on resonance, but two-photon excitation to the Rydberg state is off resonance. An example of these data with $k_3$ tuned 32 MHz above the Rydberg-$6P_{3/2}$ $F’=3$ resonance is shown in Fig. 5. As the figure shows, the count rate on the “off-axis” counter is very small for off-resonant Rydberg excitation. The fraction of phase-matched light reaches 50% of the total $6P$-$5S$ emission. When the Rydberg states are resonantly produced, the fraction of phase-matched light drops to 20%.

Figure 6 shows the ratio of off-axis to on-axis light as a function of two-photon detuning, with $k_3$ adjusted to maintain three-photon resonance. Again, the fraction of on-axis light is about 20% on resonance, increasing to 50% off resonance. This trend is accounted for by the model as seen in the figure.
conditions Rydberg-Rydberg and superradiant decoherence which we do not fully understand. However, under blockade indicating that there is an additional source of decoherence not provide the amount of decoherence we have observed,ance will cause additional decoherence should be minimal. While it has been shown that superradiant effects weak excitation conditions where Rydberg-Rydberg effects linewidths. The data presented here are all obtained under state excitation with off-resonant deexcitation to the 6P state. The k1 laser was held 32 MHz above the Rydberg-6P transition. The inset scales the off-axis count rate by the solid angle to show the total amount of light emitted. In this case the amount of on-axis light is approximately equal to the total amount of light in all other directions.

The decoherence in our data is represented in the model by γr and γ’re, which are somewhat larger than our laser linewidths. The data presented here are all obtained under weak excitation conditions where Rydberg-Rydberg effects should be minimal. While it has been shown that superradiance will cause additional decoherence [15], this still does not provide the amount of decoherence we have observed, indicating that there is an additional source of decoherence which we do not fully understand. However, under blockade conditions Rydberg-Rydberg and superradiant decoherence mechanisms are predicted to be virtually eliminated, minimizing these decoherent processes.

We have demonstrated continuous wave phase-matched four-wave mixing using Rydberg states. The directional emission is promising for studying quantum effects in blocked atom clouds. The dependence of the phase-matched light on geometry, power, and detuning is well explained by a simple density matrix model, demonstrating a good understanding of the system.

As a step toward using phase-matched four-wave mixing as a single photon source, we have focused the 480 and 1015 nm beams to 12 μm. This results in a effective volume of \( \sim 10^{-6} \text{ cm}^3 \), which could be blocked for reasonable n levels [17]. When using the focused beams, phase matching was again achieved for the various n levels, with the same dependence on F’ level, k3 intensity, and detuning from the Rydberg state. The maximum percentage of light achieved in the phase-matched direction was reduced to below 1%, as expected from Eq. (2) since the relevant factor \( N_p w^2 \) was reduced by a factor of 40. In the future, this could be improved by increasing the density of the sample [16], allowing both efficient phase matching and dipole blockade and making a single-photon source possible.

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