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Coherent excitation of a single atom to a Rydberg stateY. Miroshnychenko,¹ A. Gaëtan,¹ C. Evellin,¹ P. Grangier,¹ D. Comparat,² P. Pillet,² T. Wilk,¹ and A. Browaeys¹¹*Laboratoire Charles Fabry, Institut d'Optique, Centre National de la Recherche Scientifique, Université Paris-Sud, Campus Polytechnique, RD 128, F-91127 Palaiseau CEDEX, France*²*Laboratoire Aimé Cotton, Centre National de la Recherche Scientifique, Université Paris-Sud, Bâtiment 505, Campus d'Orsay, F-91405 Orsay CEDEX, France*

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We present the coherent excitation of a single Rubidium atom to the Rydberg state $58d_{3/2}$ using a two-photon transition. The experimental setup is described in detail, as are experimental techniques and procedures. The coherence of the excitation is revealed by observing Rabi oscillations between ground and Rydberg states of the atom. We analyze the observed oscillations in detail and compare them to numerical simulations which include imperfections of our experimental system. Strategies for future improvements on the coherent manipulation of a single atom in our settings are given.

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I. INTRODUCTION

Rydberg atoms have attracted a lot of research interest over the last decades thanks to their exaggerated properties [1]. Among these properties is their large interaction strength, which is enhanced by many orders of magnitude compared to ground-state atoms. This long-range interaction between two Rydberg atoms has led to a wealth of studies. For example, it was suggested [2,3] and recently demonstrated [4–6] that two or more Rydberg atoms or Rydberg and ground-state atoms can be bound into exotic giant molecules using photoassociation techniques. Another example where the strong interaction plays a role is the Rydberg blockade, an effect where, in a volume smaller than the interaction range, only one atom of an ensemble can be excited into a Rydberg state.

Until recently, the excitation of atoms to Rydberg states was performed using noncoherent laser excitation, usually in multiple steps. This technique led, for example, to the observation of the Rydberg blockade in clouds of ultracold atoms [7–12]. In contrast, the coherent excitation of atoms to Rydberg states has been demonstrated only in recent years, starting with experiments using cold thermal clouds of atoms [13–15] and Bose-Einstein condensates [16–18]. It was followed by the demonstration of the coherent Rydberg excitation of individual atoms trapped in tight optical dipole traps [19,20]. Along these lines, electromagnetically induced transparency has been reported in room-temperature atomic vapor [21] and microcells [22] as well as in cold atomic clouds [23], where, also, coherent population trapping [24] has been observed recently. All those demonstrations were also made possible by the technical development of easy-to-handle laser light sources, especially in the blue region of the electromagnetic spectrum, that provide a sufficient amount of power and good stability.

This progress in the coherent manipulation of atoms between ground and Rydberg states is accompanied by an increased interest on the theoretical side. It was recognized that Rydberg atoms can play an important role in the field of quantum-information processing, using either atomic ensembles or arrays of individual atoms [25–28] (see also [29] for a recent review). In these proposals, stable ground states of single atoms are used as qubit states [30–32], and the controlled

coherent interaction between the qubits can be realized via an auxiliary Rydberg state. More generally, the long-range interaction between atoms can be tailored and used to engineer quantum states of an ensemble of atoms or to perform quantum simulations [33]. A main ingredient of those proposals is the coherent excitation of an atom to a Rydberg state and the Rydberg blockade.

Recently, the coherent excitation of individual atoms trapped in tight optical dipole traps led to the observation of controlled interactions between two individual Rydberg atoms by two groups [34,35]. This work was followed by the demonstration of the entanglement of two atoms [36] and of a controlled NOT gate [37]. The aim of the present article is to present in detail the coherent excitation of individual atoms that we used in experiments reported in Refs. [35,36]. In particular, we describe the details of the experimental setup and techniques and investigate the performance of the experiment.

The article is organized as follows. In Sec. II, we present the experimental setup and the laser system used to excite the atoms in the Rydberg state. We also detail the experimental procedure used to align the lasers and to detect the Rydberg excitation of the atom. Section III deals with the Rydberg state spectroscopy on a single atom with some emphasis on the level $58d_{3/2}$. In Sec. IV, we present Rabi oscillations between the ground state and the Rydberg state of a single atom. Moreover, we compare these measurements with numerical calculations from a model which includes our independently measured experimental imperfections.

II. EXPERIMENTAL SETUP AND TECHNIQUES

In this section, we present briefly the experimental setup and the trapping of single ^{87}Rb atoms in optical tweezers which has been described in earlier publications [35,36,38]. We then detail the laser system used for the excitation of the atom from the ground state to a Rydberg state. We present the alignment technique of the laser beams on the atom. As an aside, we use this procedure to estimate the photoionization cross section from the state $5p_{3/2}$. Finally, we discuss the detection of a successful excitation of the atom in the Rydberg state.

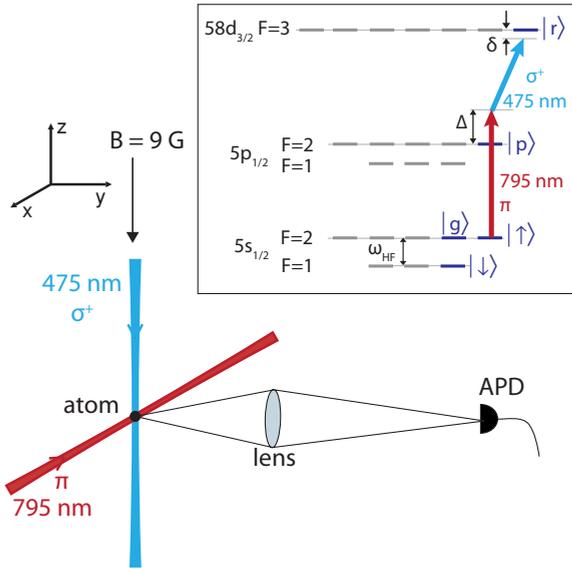


FIG. 1. (Color online) Scheme of the experimental setup. A single atom is trapped in a dipole trap and cooled by optical molasses (not shown). The atomic fluorescence light is detected on a single-photon counting avalanche photodiode (APD), allowing us to detect the presence or absence of an atom in the dipole trap. A blue σ^+ -polarized laser beam in combination with a π -polarized infrared laser beam at 795 nm is used to drive a two-photon transition to a Rydberg state. Inset shows relevant atomic levels.

A. Single-atom trapping

A single ^{87}Rb atom is held in optical tweezers which are formed by an 810 nm laser beam focused with a high numerical aperture lens to a waist of $0.9 \mu\text{m}$.¹ We operate the trap at a typical power of 0.5 mW, which corresponds to a trap depth of 0.6 mK. The trap is loaded from an optical molasses created by three retroreflected cooling beams with a power of about 3 mW per beam collimated to a waist of 1.8 mm. The detuning of the cooling laser from the ($5s_{1/2}, F=2$) to ($5p_{3/2}, F=3$) transition is $-\Delta$, where $\Gamma/2\pi = 5.75 \text{ MHz}$ is the decay rate of the $5p_{3/2}$ level. A repumping beam drives the ($5s_{1/2}, F=1$) to ($5p_{3/2}, F=2$) transition.

We detect the presence of a single atom in the dipole trap by its fluorescence light induced by the optical molasses laser beams. The fluorescence photons are collected using the same high numerical aperture lens and detected on an avalanche photodiode (APD) in single photon counting mode, as shown in Fig. 1. Owing to the small trapping volume, the number of atoms in the dipole trap is either zero or one [38]. The temperature of the atom in the optical tweezers is about $70 \mu\text{K}$, which was measured using a release-and-recapture technique [39].

B. Laser system for Rydberg excitation

We use a two-photon transition to excite a ^{87}Rb atom from its $5s_{1/2}$ ground state to the desired Rydberg level $58d_{3/2}$. We have chosen this particular state for experiments involving two atoms in neighboring traps [35,36] because of the existence of a Förster resonance [40,41] between the

two-atom states ($58d_{3/2}, 58d_{3/2}$) and ($60p_{1/2}, 56f_{5/2}$) which enhances the interaction energy between the atoms. The $5p_{1/2}$ excited state of the D_1 line of rubidium is the intermediate level. Therefore we need a laser at 795 nm and a second laser at 475 nm. As depicted in Fig. 1, the 475 nm laser is σ^+ polarized and travels along the quantization axis, which is defined by a magnetic field of 9 G along the z axis. In the chamber, we have about 35 mW of 475 nm light available, which is focused to a spot of $20 \mu\text{m}$ at the position of the atom. The π -polarized 795 nm laser has a power of about 7 mW, is applied onto the atoms from the side along the x axis, and has a waist of $130 \mu\text{m}$. The fast switching (rise time 10 ns) of the blue and red laser beams is done using electro-optic modulators.

We have chosen the intensities of the lasers such that the excitation toward the Rydberg state is fast enough ($\approx 100 \text{ ns}$) to ensure passive phase coherence between the two lasers for the duration of the excitation pulse (see more details later). The lasers are not actively phase locked, and we control only the frequencies of the lasers with respect to the atomic transitions.

The 795 nm laser is an external cavity diode laser (Toptica DL100) which is locked by a feedback loop acting on the grating piezoelectric transducer. As a reference, we use the D_1 line of ^{87}Rb , or more precisely, the transition between levels ($5s_{1/2}, F=2$) and ($5p_{1/2}, F=2$). The lock scheme relies on a saturated absorption spectroscopy. The error signal is generated using a lock-in technique by modulating the laser diode current at a frequency of 80 kHz. The frequency of the laser is then additionally shifted by a detuning Δ toward the blue of the transition with an acousto-optical modulator to avoid populating the intermediate state.

The 475 nm laser is a frequency-doubling system (Toptica TA-SHG). We stabilize the fundamental wavelength of the 950 nm master laser as well as an external cavity diode laser by feedback onto its grating piezoelectric transducer. Here we use a Fabry-Perot cavity to fix the 950 nm laser relative to the 795 nm laser, which gives us more freedom in the choice of the Rydberg state we are addressing. Both the 795 nm and the 950 nm laser is coupled to the permanently scanning cavity, which has a finesse $F \approx 100$ and a free spectral range of 300 MHz. The cavity is continuously swept with a scan frequency of about 100 Hz across the transmission peaks of both lasers. Our locking scheme relies on the stabilization of the separation between these two transmission peaks in the sweep. We use a homemade circuit to convert the time difference into a voltage. When the signal on the photodiode behind the cavity exceeds for the first time a threshold value, an integrator is started and accumulates a voltage until the photodiode signal exceeds the threshold value for the second time. Subtracting this value with a set voltage generates an error signal, which is treated and fed back onto the 950 nm laser.

This locking technique is only suitable for compensating long-term drifts of the laser since the bandwidth of the correcting signal is low and depends on the frequency of the Fabry-Perot cavity sweep. Moreover, this method relies on the fact that the cavity piezo responds linearly, that the sweep has a very good stability in frequency, and that the cavity has slow drifts. Despite those constraints, we were able to stabilize the 795 nm and 950 nm lasers with respect to each other to about 4 MHz for the duration of ~ 1 hour. We also checked the intrinsic short-term stability of the 950 nm master laser using

¹All beam waists in this article are given at a $1/e^2$ -waist radius.

a self-heterodyne technique. This consists in observing a beat note of the laser with itself shifted in frequency by 80 MHz and delayed by $\sim 7 \mu\text{s}$ using a 1.3 km long fiber. We have measured a full width half maximum (FWHM) of the beat signal of 210 kHz, leading to a linewidth of the 950 nm laser of 105 kHz.² Using the same technique, we have measured a linewidth of the 795 nm laser of 600 kHz, which is larger because of the current modulation used to lock it on the atomic transition.

However, this method does not allow one to set the wavelength of the 950 nm laser absolutely since we do not know a priori which longitudinal mode of the cavity we address. We use a wavelength meter with an absolute accuracy of 10 MHz (HighFinesse-Ångstrom WS-U) to set the frequency of the laser close to the desired frequency. The locking scheme described earlier has been used for all data shown in this article and for the data discussed in [36]. For the data in [35], we used an even simpler method: the HighFinesse-Ångstrom wave meter provides a Proportional-Integral-Derivative control option that generates an error signal which can be used to stabilize the laser to any desired frequency with a precision of 2 MHz. Automatic recalibration of the wave meter every 10 s using the 795 nm laser locked to ^{87}Rb avoids long term drifts of the wave meter. The limiting factor of this method is the absolute accuracy of the wave meter, which is estimated around 5 MHz on the time scale of half an hour.

C. Alignment of the 475 nm laser

In order to reach a high two-photon Rabi frequency with the limited available 475 nm laser power, we need to align the maximum of the intensity profile of the 475 nm light, which is focused to a spot size of about $20 \mu\text{m}$, to the position of the atom. For this purpose, we take advantage of the fact that light at 475 nm photoionizes rubidium atoms from the $5p_{3/2}$ -state [see Fig. 2(a) for the excitation scheme] and that the photoionization rate is proportional to the light intensity.

After we have prepared a single atom in the tweezers, we turn off the molasses beams, and the atom is trapped in the dark. After 31 ms from the start of the sequence, we illuminate the atom with both the 475 nm laser beam and the optical molasses beams. The optical molasses creates a population in the excited $5p_{3/2}$ state from where the 475 nm light photoionizes the atom, which is consequently lost from the trap. Figure 2(b) shows a typical trace of the observed fluorescence signal averaged over 100 experimental runs. We observe a peak in the fluorescence when the lasers are switched on, which decays exponentially to a value corresponding to the background of the molasses beams and indicating that the atom is lost. A fit by an exponential decay gives the photoionization rate $1/\tau$, which depends on the applied powers and on the beam alignments. We verified that the inverse of τ is proportional to the intensity of the 475 nm laser [see the inset of Fig. 2(b)]. Therefore this method allows us to measure the beam waist of the 475 nm laser using the single atom as a probe. Keeping the laser power fixed and evaluating the photoionization rate for

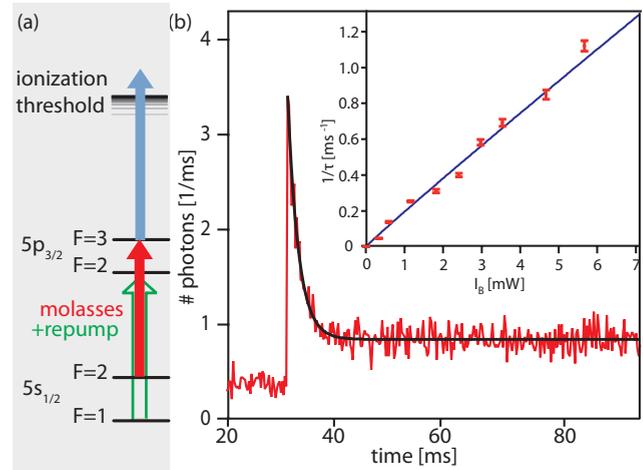


FIG. 2. (Color online) Procedure used to align the 475 nm laser onto the single atom. (a) To create population in the $5p_{3/2}$ excited state, the atom is driven by the molasses beams and the repumper. From there, the 475 nm laser photoionizes the atom with a rate proportional to its intensity. (b) Fluorescence signal of a single atom averaged over 100 experimental runs. At $t = 31 \text{ ms}$, the atom is exposed to the optical molasses and the 475 nm laser simultaneously. The atom starts fluorescing on the D_2 line, is photoionized from the $5p_{3/2}$ state by the 475 nm laser, and is consequently lost from the trap. The solid line is an exponential fit to the data and gives the photoionization rate $1/\tau$. The inset demonstrates the linear dependence of the inverse of the decay time constant as the function of the blue laser intensity.

different positions of the laser beam, we find $w_x = 22(1) \mu\text{m}$ and $w_y = 19(1) \mu\text{m}$, which correspond well to the values we expect for our optical setup. The laser beam is then centered onto the atom using the same method.

D. Estimate of the photoionization cross section

The photoionization rate can also be used to estimate the photoionization cross section σ of ^{87}Rb from the $5p_{3/2}$ state. The relation between σ and $1/\tau$ is given by [42]

$$\frac{1}{\tau} = f \frac{I_B \sigma}{h\nu_B}, \quad (1)$$

where I_B and ν_B are the intensity and the frequency of the 475 nm laser and f is the population in the $5p_{3/2}$ state. The population f depends strongly on the parameters of the optical molasses and is given by

$$f = \frac{1}{2} \frac{c_1^2 \Omega_{\text{mol}}^2 / 2}{\delta_{\text{mol}}^2 + \Gamma^2 / 4 + c_2^2 \Omega_{\text{mol}}^2 / 2}, \quad (2)$$

where the coefficients c_1 and c_2 are connected to Clebsch-Gordan coefficients [43] and have values of $c_1^2 = c_2^2 = 0.73$.³ Including the light shift due to the dipole trap, the detuning of the optical molasses is $\delta_{\text{mol}}/2\pi = 100 \text{ MHz}$. The total Rabi frequency on the cooling transition of the molasses is

²We assume that the corresponding linewidth of the light generated at 475 nm is twice larger, although we could not measure it directly.

³Although these values were originally measured for Cs atoms, we use them for ^{87}Rb because for the stretched states, the Clebsch-Gordan coefficients are the same as for Cs.

given by $\Omega_{\text{mol}} = \frac{d}{\hbar} \sqrt{\frac{2I_{\text{mol}}}{c\epsilon_0}}$, where $d = 3.58 \times 10^{-29}$ C m is the transition dipole matrix element [44] and I_{mol} is the total intensity because of the optical molasses beams at the position of the atom. In our case, the estimation of this intensity can be done only approximately since it is difficult to determine the position of the single atom within the beams. We estimate the population in the $5p_{3/2}$ state f to be between 1% and 8%. This uncertainty contributes the most to the final uncertainty of the ionization cross section.

From the fit in Fig. 2(b), we extract $\tau = 2.03(9)$ ms, which was taken for a power of 7.4 mW of the 475 nm laser. The values of the ionization cross section obtained for the two extreme populations f of the $5p_{3/2}$ state are

$$0.2 \times 10^{-17} \text{ cm}^2 \leq \sigma \leq 1.6 \times 10^{-17} \text{ cm}^2. \quad (3)$$

Despite our uncertainty of f , these values agree with theoretically predicted cross sections, which are between $1.25 \times 10^{-17} \text{ cm}^2$ and $1.4 \times 10^{-17} \text{ cm}^2$ from $5p$ states to the ionization threshold [45], and with the experimentally measured values from the literature: $1.48(22) \times 10^{-17} \text{ cm}^2$ at 476.5 nm (see [42], and references therein).

E. Detection of Rydberg atoms

We detect a successful excitation toward the Rydberg state by a loss of the atom. Our detection method is based on the fact that Rydberg atoms are not trapped in our dipole trap, the light at 810 nm being even slightly antitrapping for them. For a Rydberg atom, the electron can almost be considered as a free particle, and therefore the Thomson model can be applied to calculate the polarizability of the atom [46]:

$$\alpha = -\frac{e^2}{m_e \epsilon_0 \omega^2} < 0, \quad (4)$$

where e is the charge of the electron, m_e is the mass of the electron, and $\omega/2\pi$ is the frequency of the applied laser. Since the light shift is proportional to $-\alpha$, and in our case $\alpha < 0$, the light shift is positive for all Rydberg atoms, as long as we are far from any atomic resonance between the Rydberg state and other levels. Since the positive light shift is very small (only ≈ 1 MHz), the Rydberg atoms leave the trapping region on a time scale that corresponds to their velocities at the temperature of 70 μK . We have estimated the lifetime of a Rydberg atom in the dipole trap by exciting it to the Rydberg state (see Sec. IV) and measuring the probability to drive it back to the ground state after a given time using a second laser pulse. After 10 μs , the atom has left the trapping region, which is faster than the radiative decay time back to the ground state, which is on the order of 200 μs [1]. Moreover, an atom can be photoionized by blackbody radiation (85 μs). Photoionization by the dipole trap beam (1.4 ms [46]) is negligible on these time scales.

III. RYDBERG STATE SPECTROSCOPY OF A SINGLE ATOM

In this section, we discuss how the desired Rydberg line is initially localized in the spectrum. Then we explain the optical pumping in state $|5s_{1/2}, F = 2, m_F = 2\rangle$. From this initial state, a two-photon spectroscopy of the $58d_{3/2}$ level is performed on a single atom.

A. Localizing the Rydberg lines

In order to initially locate the $58d$ line and to have a proof of principle of the detection of Rydberg atoms, we perform a spectroscopy of Rydberg levels using a two-step excitation of the atoms via the $5p_{3/2}$ level. For this particular measurement, we use a two-step excitation at 780 nm, and the blue laser is tuned to 480 nm. We run the optical tweezers with the optical molasses (780 nm) continuously on and observe the fluorescence on the APD from single atoms entering and leaving the trap continuously on the APD while we scan the frequency of the blue laser. When its frequency hits one of the Rydberg lines, the loading of the dipole trap is strongly perturbed, and the atomic fluorescence signal on the APD drops down to the background level.

We have observed s , p , d , and f lines for n ranging from 54 to 68, which have a typical separation of 5–30 GHz. The positions of the lines found with this method coincide within ~ 100 MHz with respect to the calculated values, which were calculated by the quantum defect theory [1] using the values of the quantum defects measured in Refs. [47,48]. According to the selection rules, only s and d lines should be observable in a two-step excitation. The presence of p and f lines indicates the existence of a residual electric field at the position of the atom. With this spectroscopy method, we identified the $58d_{3/2}$ line, which we use in further experiments.

B. Initial state preparation: Optical pumping

In all further experiments, we perform a direct two-photon excitation of a single atom trapped in the tweezers from its ground state $|5s_{1/2}, F = 2, m_F = 2\rangle$. The atom is optically pumped into this particular state for 600 μs using two laser beams: σ^+ -polarized light resonant with the atomic transition $|5s_{1/2}, F = 2\rangle$ to $|5p_{3/2}, F = 2\rangle$ (taking into account the light shift of the ground state) and repumping light resonant with the transition $|5s_{1/2}, F = 1\rangle$ to $|5p_{3/2}, F = 2\rangle$. To ensure that there is no population left in the $F = 1$ ground state, the repumping light is switched off 1 μs later than the pumping light. The efficiency of the optical Zeeman pumping to the state $|5s_{1/2}, F = 2, m_F = 2\rangle$ is about 95%, which was checked by measuring the visibility of Rabi oscillations driven by Raman transitions between the $|5s_{1/2}, F = 1, m_F = 1\rangle$ and $|5s_{1/2}, F = 2, m_F = 2\rangle$ hyperfine ground states [32].

C. Spectroscopy of the $58d_{3/2}$ level

Figure 3(a) shows the time sequence used for this experiment. About 1 μs after the optical pumping, we switch on the 475 nm laser beam. We also switch off the dipole trap for 600 ns to avoid light shifts because of the dipole trap laser light during the Rydberg excitation. During this time window, the 795 nm laser is switched on for $T = 60$ ns to coherently excite the free atom to the Rydberg state, as will be detailed in Sec. IV. After this excitation, the dipole trap is switched on again, which results in the loss of the atom if it was in the Rydberg state and in its recapture in the optical tweezers otherwise. The frequency scan is performed by changing the absolute frequency of the 475 nm laser. For every point, we repeat the experiment 100 times.

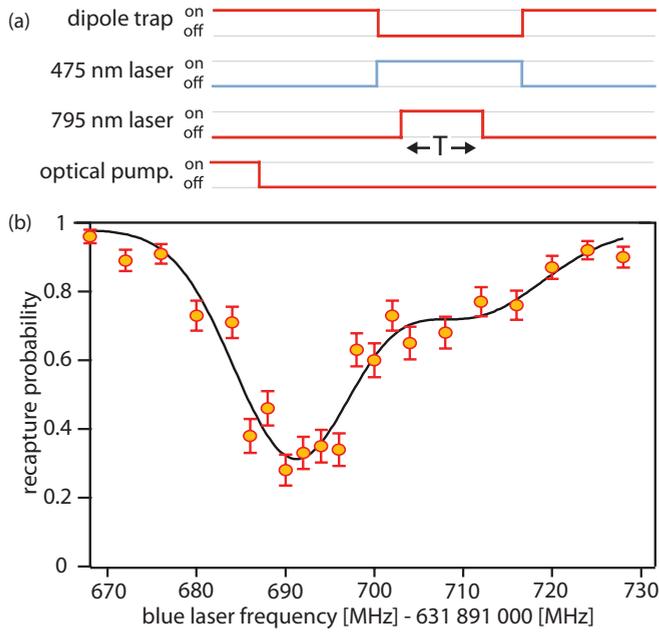


FIG. 3. (Color online) Two-photon spectroscopy of the $58d_{3/2}$ line. (a) The blue laser is switched on for 600 ns. The precise timing of the two-photon transition is controlled by the duration T of the infrared square pulse with a rise time less than 10 ns. The frequency scan is done by varying the frequency of the blue light. (b) The corresponding spectrum taken for $T = 60$ ns and $(\Omega_R, \Omega_B, \Delta)/2\pi = (255, 24, 400)$ MHz (see Sec. IV). Each point corresponds to 100 repetitions of the experiment. The solid line is a double-Gaussian fit to the experimental points.

Figure 3(b) presents the resulting two-photon spectrum. We observe a double-peak structure. The two peaks have a separation of 20(1) MHz. We attribute the more pronounced dip to the transition to the $|58d_{3/2}, F = 3, m_F = 3\rangle$ state. The width (FWHM) of this main peak is 16(1) MHz. It is compatible with the expected width (13.3 MHz) using a square pulse with duration $T = 60$ ns convoluted with the contribution from experimental imperfections (6 MHz; see Sec. IV). These are mainly fluctuations of the excitation laser frequencies and intensities as well as the rise and fall times of the square pulse. The center of the line is at $\nu_B^{\text{exp}} = 631891691(10)$ MHz, where the error bar results from the absolute precision of the wave meter. We calculate the position of this line to be at $\nu_B^{\text{calc}} = 631891657(6)$ MHz using the quantum defects reported in Ref. [47] and taking into account the light shifts induced by the 795 nm and 475 nm lasers and the Zeeman shifts. The difference between the calculated and measured frequencies of 34(12) MHz can have several reasons: an uncertainty of the light shift induced by the 795 nm laser beam itself during the excitation (≈ 10 MHz) or the absolute precision of the wave meter (≈ 10 MHz). Moreover, stray electric fields, which we cannot control on our setup, would also induce a shift. However, the calculated Stark shift $[-240 \text{ MHz}/(\text{V}/\text{cm})^2]$ does not have the correct sign to explain the observed frequency difference.

The origin of the satellite dip is not entirely clear at this stage, and further investigation is required. A possible candidate is the imperfect optical pumping in the Zeeman state

that leaves population in $|5s_{1/2}, F = 2, m_F = 1\rangle$. From that state, the $|58d_{3/2}, F = 2, m_F = 2\rangle$ and $|58d_{3/2}, F = 3, m_F = 2\rangle$ levels can be excited by the two-photon transition. Since the $|5s_{1/2}, F = 2, m_F = 1\rangle$ state is light shifted by the 795 nm laser by a smaller amount than the $|5s_{1/2}, F = 2, m_F = 2\rangle$ state, because of a smaller Clebsch-Gordan coefficient connecting it to the corresponding Zeeman state in level ($5p_{1/2}$), we expect the side dip to be at a higher frequency of the 475 nm laser, as we observe on the measured spectrum. This is also confirmed by the fact that changing the detuning Δ from the intermediate state from positive to negative detuning changes the position of the side dip from the blue to the red side of the Rydberg line. However, the area of this side dip seems to be too large to be explained only by about 5% imperfect optical pumping. Another possible effect is the nonperfect polarization of the excitation laser at 475 nm.

IV. RABI OSCILLATION BETWEEN GROUND AND RYDBERG STATES

Once the state $|58d_{3/2}, F = 3, m_F = 3\rangle$ is localized in the spectrum, we set the frequency of the blue laser to the center of the line. By varying the pulse duration T of the 795 nm laser, we observe Rabi oscillations between the ground state $|5s_{1/2}, F = 2, m_F = 2\rangle$ and the Rydberg state $|58d_{3/2}, F = 3, m_F = 3\rangle$, demonstrating the coherence of the two-photon excitation. We discuss the experimental results of measurements with different Rabi frequencies of the red (795 nm) and the blue (475 nm) laser and for different detunings from the intermediate state Δ . The measurements are compared to simulations that we performed, which include intensity and frequency fluctuations present in our system.

A. Experimental results

We have recorded the Rabi oscillations between ground and Rydberg states for different values of the red and blue Rabi frequencies Ω_R and Ω_B and the detuning Δ . In Fig. 4, the recapture probability of the single atom is plotted versus the duration T of the excitation pulse. Each point corresponds to 100 repetitions of the experiment. Figure 4(a) shows the measurement for $(\Omega_R, \Omega_B, \Delta)/2\pi = (255, 24, 400)$ MHz. For these parameters, we reach a maximal excitation probability toward the Rydberg state of $\approx 70\%$, and the oscillation is strongly damped. A fit to the data by the function $A - B e^{-T/\tau} \cos \Omega T$ (not shown) yields a two-photon Rabi frequency of $\Omega/2\pi = 7.0(1)$ MHz and $\tau = 480$ ns. The solid and the dotted line are results of simulations that will be discussed in the next section.

We attribute the damping mainly to spontaneous emission from the intermediate $5p_{1/2}$ state, which destroys the coherence. A second measurement, shown in Fig. 4(b), with about the same Rabi frequencies of the lasers $(\Omega_R, \Omega_B)/2\pi = (250, 28)$ MHz but with a larger detuning $\Delta/2\pi = 600$ MHz, already improved the excitation probability up to 80%. The fit yields $\Omega/2\pi = 5.8(1)$ MHz and a longer damping time $\tau = 560$ ns. The damping rate could even be lowered further by reducing the Rabi frequency of the red laser significantly to $\Omega_R/2\pi = 80$ MHz and by increasing one of the blue lasers to $\Omega_B/2\pi = 70$ MHz (stronger focusing of the blue laser

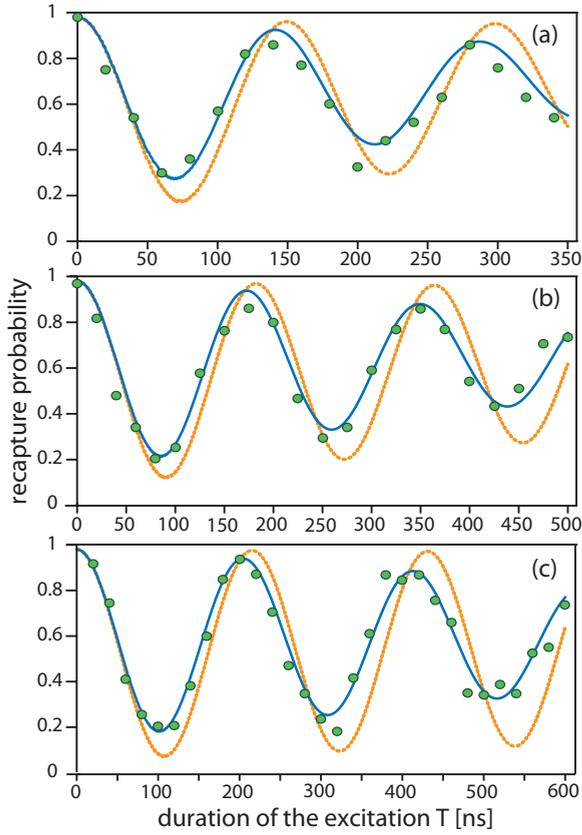


FIG. 4. (Color online) Coherent Rydberg excitation of a single atom to the level $|58d_{3/2}, F=3, m_F=3\rangle$ for different experimental parameters: (a) $(\Omega_R, \Omega_B, \Delta)/2\pi = (255, 24, 400)$ MHz, (b) $(\Omega_R, \Omega_B, \Delta)/2\pi = (250, 28, 600)$ MHz, and (c) $(\Omega_R, \Omega_B, \Delta)/2\pi = (80, 70, 600)$ MHz. Each point corresponds to 100 repetitions of the experiment. The blue line is the result of a Monte Carlo simulation of the dynamics of a five-level system, which includes a decay from the intermediate state, fluctuations of the power and of the frequency of the lasers, and imperfection of the optical pumping. The dotted orange line shows for comparison the results of simulations with the same parameters of Ω_R , Ω_B , and Δ but without fluctuations.

and improvement of the optical setup). The fit gives $\Omega/2\pi = 4.9(1)$ MHz and $\tau = 912$ ns. But even at these parameters, the excitation probability to the Rydberg state does not exceed 85%. To understand better the underlying mechanisms for the damping and the limited excitation probability, we have modeled the system including experimental imperfections, as will be discussed in the following section.

B. Comparison to numerical simulations

In our numerical calculation, we include five atomic states, which are labeled in the level scheme of Fig. 1: in the $5s_{1/2}$ ground state, we define $|\downarrow\rangle = |F=1, m_F=1\rangle$, $|\uparrow\rangle = |F=2, m_F=2\rangle$, and $|g\rangle = |F=2, m_F=1\rangle$. We call the intermediate state $|p\rangle = |5p_{1/2}, F=2, m_F=2\rangle$, and the Rydberg level is $|r\rangle$. We take into account the spontaneous emission from the intermediate level $|p\rangle$ to the ground state with a rate of $\Gamma/2\pi = 5.75$ MHz and the corresponding branching ratios of $1/2$ to state $|\downarrow\rangle$, $1/3$ to $|\uparrow\rangle$, and $1/6$ to

$|g\rangle$ [44] as well as the decay from the Rydberg state to the intermediate level with rate $\gamma/2\pi = 4.8$ kHz.

The time evolution of the density matrix ρ is governed by the master equation

$$\dot{\rho} = -\frac{i}{\hbar}[\hat{H}, \rho] + \mathcal{L}, \quad (5)$$

where the Hamiltonian \hat{H} and the Liouville \mathcal{L} operators are given in the basis $\{|\downarrow\rangle, |g\rangle, |\uparrow\rangle, |p\rangle, |r\rangle\}$ by [49]

$$\hat{H} = \hbar \begin{pmatrix} -\omega_{\text{HF}} & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & \Omega_R/2 & 0 \\ 0 & 0 & \Omega_R/2 & -\Delta & \Omega_B/2 \\ 0 & 0 & 0 & \Omega_B/2 & -\delta \end{pmatrix},$$

$$\mathcal{L} = \begin{pmatrix} \frac{\Gamma}{2}\rho_{pp} & 0 & 0 & -\frac{\Gamma}{2}\rho_{\downarrow p} & -\frac{\gamma}{2}\rho_{\downarrow r} \\ 0 & \frac{\Gamma}{6}\rho_{pp} & 0 & -\frac{\Gamma}{2}\rho_{gp} & -\frac{\gamma}{2}\rho_{gr} \\ 0 & 0 & \frac{\Gamma}{3}\rho_{pp} & -\frac{\Gamma}{2}\rho_{\uparrow p} & -\frac{\gamma}{2}\rho_{\uparrow r} \\ -\frac{\Gamma}{2}\rho_{p\downarrow} & -\frac{\Gamma}{2}\rho_{pg} & -\frac{\Gamma}{2}\rho_{p\uparrow} & \gamma\rho_{rr} - \Gamma\rho_{pp} & -\frac{\Gamma+\gamma}{2}\rho_{pr} \\ -\frac{\gamma}{2}\rho_{r\downarrow} & -\frac{\gamma}{2}\rho_{rg} & -\frac{\gamma}{2}\rho_{r\uparrow} & -\frac{\Gamma+\gamma}{2}\rho_{rp} & -\gamma\rho_{rr} \end{pmatrix}.$$

Here $\omega_{\text{HF}}/2\pi = 6.834$ GHz is the hyperfine ground-state splitting frequency. We include the power and frequency fluctuations of the two lasers by applying a Monte Carlo simulation. We assume that the power fluctuations of the red and the blue lasers are both Gaussian distributed and are 2.5% and 5% (FWHM), respectively, as measured independently. We note that these intensity fluctuations also lead to a fluctuation of the frequency of the transition due to the light shift induced by the lasers on this two-photon transition, respectively, $\Omega_R^2/4\Delta$ on the ground state and $\Omega_B^2/4\Delta$ on the Rydberg state. To take into account the frequency fluctuations of the lasers, we assume a Gaussian distribution of the two-photon detuning with respect to the transition δ . This procedure would also include a potential variation in the Rydberg transition frequency by the Stark effect because of a fluctuating stray electric field. We take into account the efficiency of the optical pumping of 95% by assuming that the remaining 5% are in state $|g\rangle$. Finally, we multiply the results of the simulation by 0.98 to account for the measured probability to recapture the atom at the end of the sequence in the absence of laser excitation. We average over 100 evolutions of the master equation, and the solution is shown for different values of Ω_R , Ω_B , and Δ in Fig. 4 as a blue solid line. In order to have an idea of how the fluctuations influence the result, we have added to the graphs of Fig. 4 the dotted orange line, which corresponds to the solution of the model, without including the frequency and power fluctuations of the lasers.

The simulations are in good agreement with the measured data. For the simulations shown in blue in Fig. 4, we assumed fluctuations (FWHM) of the two-photon detuning δ of 6 MHz in Fig. 4(a), 4.5 MHz in Fig. 4(b), and 4 MHz in Fig. 4(c). Comparing them to the results without fluctuations, we identify two main factors limiting the population transfer to the Rydberg state: the spontaneous emission from state $|p\rangle$ and the fluctuations of laser frequency and intensity. The influence

of the spontaneous emission can be reduced by decreasing the ratio Ω_R^2/Δ^2 , resulting in an increase of the contrast of the Rabi oscillations, as shown in Figs. 4(b) and 4(c). Moreover, we estimate from our simulations that if we would reduce the laser frequency fluctuations to 1 MHz, we could increase the Rydberg excitation efficiency to 93%. We finally note that the frequency fluctuations used in the model are compatible with the one estimated from measurements on the laser system (see Sec. II B). At the present stage of the experiment, we therefore have no evidence for a fluctuation of the transition frequency because of stray electric fields.

As a last comment, we conclude from the good agreement between the data and the model that the result of the simulation is compatible at 100% efficiency with the Rydberg state detection we use in the experiment.

V. CONCLUSION

We perform coherent excitation of single atoms to the Rydberg state $|58d_{3/2}, F = 3, m_F = 3\rangle$ using a two-photon transition and observe Rabi oscillations with a high contrast.

The observed Rabi frequency is about 7 MHz, which relaxes partially the requirements for the frequency stability of the laser system. The largest population transfer to the Rydberg state observed is 80%. The good agreement between the model and the data leads us to conclude that this maximal transfer efficiency is limited by the spontaneous emission from the intermediate state and the frequency stability of the excitation lasers. In future experiments, the population of the intermediate level can be reduced by increasing the detuning from this level and using higher power in the blue beam to keep the two-photon Rabi frequency high.

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