

Storage and Recall of Weak Coherent Optical Pulses with an Efficiency of 25%

M. Sabooni, F. Beaudoin, A. Walther, N. Lin, A. Amari, M. Huang, and S. Kröll

Department of Physics, Lund University, P.O. Box 118, SE-22100 Lund, Sweden

(Received 10 December 2009; published 2 August 2010)

We demonstrate experimentally an efficient coherent rephasing scheme for the storage and recall of weak coherent light pulses in an inhomogeneously broadened optical transition in a Pr^{3+} :YSO crystal at 2.1 K. Precise optical pumping using a frequency stable (≈ 1 kHz linewidth) laser is employed to create a highly controllable atomic frequency comb structure. We report single photon level storage and retrieval efficiencies of 25%, based on coherent photon-echo-type reemission in the forward direction. The high efficiency is mainly a product of our highly controllable and precise ensemble-shaping technique. The coherence property of the quantum memory is proved through interference between a super-Gaussian pulse and the emitted echo.

DOI: 10.1103/PhysRevLett.105.060501

PACS numbers: 03.67.Hk, 42.50.Ct, 42.50.Ex, 42.50.Md

Introduction.—Quantum information processing applications such as quantum networks require coherent and reversible mapping between light and matter [1,2]. Currently, this requirement is expected to be met by means of quantum repeaters, which allow temporal storage of quantum information in quantum memories and distribution of entanglement across long distances [3]. In the repeater protocol, the entanglement distribution has probabilistic behavior, and without quantum memories all probabilistic steps would have to succeed simultaneously. Quantum memories are therefore a key ingredient in any future long distance quantum communication scheme and have thus been subject to very active development in recent time. To be useful in actual processing applications, these devices must maintain fidelities close to 100% during storage times of the order of seconds and also ensure high recall output efficiency [4–6].

Quantum memory protocols [such as electromagnetically induced transparency (EIT) [7] based on stopped light [8], controlled reversible inhomogeneous broadening (CRIB) [9–11], and atomic frequency comb (AFC) [6,12], based on photon-echo experiments] use atomic ensembles to obtain the required control and strong coupling between the light field and the storage medium. These protocols can be adapted to the original Duan-Lukin-Cirac-Zoller protocol [1] to create entanglement generation via single photon detection. Therefore, it is important to demonstrate that quantum memories can be operated at the single photon level.

Protocols which enable storage of multiple temporal modes can increase the entanglement generation rate by a factor approximately equal to the number of temporal modes [13]. Storing many modes with high efficiency requires a high optical depth, which generally can be an experimental difficulty. Achieving efficient multimode storage at lower optical depth is therefore desirable. It has been shown [12,14] that the number of modes N_m that can be efficiently stored by using EIT scales as $N_m \approx \sqrt{\alpha L}$, where αL is the optical depth. CRIB offers a better

scaling since $N_m \approx \alpha L$ (to double the number of modes, it is sufficient to double αL), but in the AFC protocol the number of temporal modes that can be stored with a given efficiency is independent of optical depth.

Based on EIT, storage and retrieval efficiency of 10% for 10 μs storage time in hot atomic vapor was published by Eisaman *et al.* [15]. With the same scheme, Choi *et al.* [16] achieved 17% for 1 μs in a cold atomic ensemble. Recently, Lauritzen *et al.* [17] published a storage and retrieval efficiency of 0.25% for 600 ns in an erbium-doped crystal by using the CRIB protocol. By using the AFC protocol, de Riedmatten *et al.* [6] obtained 0.5% storage and retrieval efficiency after 250 ns in Nd^{3+} :YVO₄, and

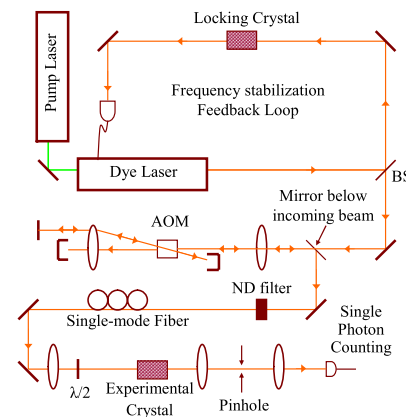


FIG. 1 (color online). Experimental setup. The frequency stabilization feedback system uses a Pr^{3+} :YSO crystal as a locking crystal. All pulses are created by using a double pass AOM. The incoming and outgoing beams in the AOM have different height, and a mirror below the incoming beam redirects the outgoing beam to the experiment. Absorptive neutral density filters stand for most of the attenuation such that the AOM can operate in a linear light intensity vs rf amplitude regime for adjusting the storage pulse intensity. After spatial cleaning in the single mode fiber, the light interacts with the Pr^{3+} :Y₂SiO₅ experimental crystal at 2.1 K. A 50 μm pinhole is used to select the center of the beam for the single photon counting detector.

Chanelière *et al.* [18] achieved 9% storage and retrieval efficiency for 1.5 μs in Tm^{3+} :YAG crystal.

In this Letter, we use the AFC protocol to store weak coherent pulses, with on the average 0.1 photons/pulse, up to 800 ns with 25% storage and retrieval efficiency in a Pr^{3+} :YSO crystal. Recently, in this material 96% transfer efficiency from an excited state to ground state spin level was achieved [19], and such transfer pulses were used to demonstrate an on-demand readout quantum memory based on a collective spin wave excitation in the hyperfine ground state levels of Pr:YSO [20]. The lifetime of such spin state storage has subsequently been extended to the 100 μs range [21]. Therefore, to achieve high efficiency on demand quantum memory operation, the key issue is to demonstrate high efficiency coherent rephasing in the low photon number regime which here is achieved through our highly controllable and precise ensemble-shaping technique.

Materials and methods.—The experimental setup is shown in Fig. 1. A 6-W Coherent Verdi-V6, 532 nm Nd:YVO₄ laser pumps a Coherent 699-21 dye laser, which emits approximately 500 mW light at 605.82 nm. This frequency matches the $^3\text{H}_4\text{-}^1\text{D}_2$ transition of praseodymium ions doped into yttrium silicate (Pr^{3+} :Y₂SiO₅). We use a frequency stabilization feedback system [22], based on a Pr^{3+} :Y₂SiO₅ crystal, which has a frequency drift below 1 kHz/s. This precise frequency control gives us ability to create and control the AFC structure and is the main reason for increasing the storage and retrieval efficiency. An acousto-optic modulator (AOM) with 200 MHz center frequency, in a double pass arrangement that cancels spatial movement when changing the frequency of the diffracted beam, controls phase, amplitude, and frequency of the light. The radio-frequency signal used to drive the AOM is created by a 1-GS/s arbitrary waveform generator (Tektronix AWG520). Two mechanisms for light attenuation to the single photon level are used. The light beam can be attenuated by reducing the amplitude of the radio-frequency field driving the AOM, but since rf amplitude versus light amplitude is highly nonlinear and difficult to calibrate due to low diffraction efficiency at low rf amplitudes, a shutter wheel with absorptive neutral density filters was added in the setup. The remaining control of the light amplitude was then handled through the AOM rf amplitude finally yielding about 0.1 photons per pulse at the sample. For cleaning up the spatial mode, after the AOM, the light was sent to a single mode fiber and then to a $\lambda/2$ plate such that the light polarization could be aligned along the transition dipole moment to yield maximum absorption. The sample is a Pr^{3+} :Y₂SiO₅ crystal with 0.05% doping concentration, and it has three perpendicular optical axes labeled D_1 , D_2 , and b , where the direction of light propagation was chosen along the b axis. The crystal dimensions are $10 \times 10 \times 20$ mm along the $D_1 \times D_2 \times b$ axis. The width of the Gaussian beam at the center of the crystal is about 100 μm , and it is imaged (1:1) onto a 50 μm pin-

hole. The pinhole gives us the opportunity to pick the light emitted by ions at the center of the beam that all have experienced roughly the same light intensity. To keep the optical coherence time (T_2) of the praseodymium $>100 \mu\text{s}$, the crystal was immersed in liquid helium at 2.1 K. After the sample crystal, a mechanical shutter was used to protect the detector from the intense AFC preparation pulses. A Hamamatsu single photon counting detector (model H8259-01) with a quantum efficiency of 7.5% and a minimum gating time of 100 ns was used.

Let us now describe (see Fig. 2) in more detail how to create an AFC [12], i.e., a periodic series of narrow (FWHM = γ) and highly absorbing (optical depth αL) peaks with a periodicity Δ as a frequency grating of ions in the $|1/2g\rangle$ state (see Fig. 2). By means of optical pumping, an 18 MHz zero absorption frequency region (see Ref. [23]) is created within the 5 GHz wide Pr^{3+} :Y₂SiO₅ inhomogeneous profile [24]. This nonabsorbing frequency region is hereafter referred to as a spectral pit. Optimal pit creation needs a series of explicit pulses [23]. Afterwards, a frequency-dependent absorption grating was created inside this spectral pit. Complex hyperbolic secant optical pulses were used to transfer some ions from $|5/2g\rangle$ to $|5/2e\rangle$ and then from the $|5/2e\rangle$ to the $|1/2g\rangle$ state [25]. Changing the center frequency of the complex hyperbolic secant pulses, by an amount Δ , is a simple technique to create a second peak inside the spectral pit. By repeating this procedure, it is possible to create an atomic frequency comb in the $|1/2g\rangle$ level with a periodicity Δ . The number of peaks (N) is limited by the

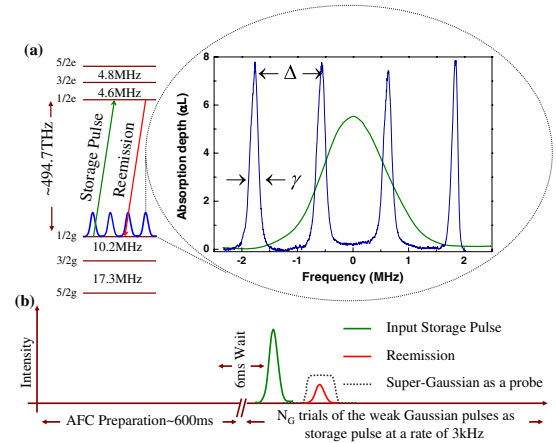


FIG. 2 (color online). (a) The $^3\text{H}_4\text{-}^1\text{D}_2$ transition of the Pr^{3+} :Y₂SiO₅ crystal is used for the experiment. An experimental AFC absorption profile with defined height (αL), peak separation (Δ), and FWHM (γ) with all ions in the state $|1/2g\rangle$ is created. A Gaussian pulse with center frequency set to the $|1/2g\rangle \rightarrow |1/2e\rangle$ transition is used as a storage pulse. (b) A sequence of pulses which can be divided into preparation and storage pulses of a total duration of approximately 1.1 s is sent into the sample. A super-Gaussian pulse is used as a probe pulse for interfering with the reemission (echo) and showing the phase preservation of the AFC interface.

$|1/2e\rangle \rightarrow |3/2e\rangle$ separation, $(\delta f_{|1/2e\rangle \rightarrow |3/2e\rangle})$, and $\Delta [\Delta \times (N-1) < \delta f_{|1/2e\rangle \rightarrow |3/2e\rangle}]$.

Results and discussions.—First, we offer a brief theoretical background to the AFC quantum memory interface. A weak light pulse is sent as a coherent state $(|\alpha\rangle_L)$, on the average containing less than a single photon, with a well-defined bandwidth $(1/\tau_p)$, to an AFC structure with well-defined peak separation (Δ) , peak width (γ) , finesse $F = \Delta/\gamma$, and peak height (αL) consisting of N ions all in the $|1/2g\rangle$ state. The objective is to transfer all the energy of the weak coherent wave packet to the distribution of ions, $|g_1 \dots g_N\rangle$, in the spectral grating with defined finesse $(F = \Delta/\gamma)$. After the light-matter interaction, the initial state of the N atoms will develop into the state [12]:

$$|\alpha\rangle_A = |g_1 \dots g_N\rangle + \sum_{j=1}^N c_j e^{i(2\pi\delta_j t - kz_j)} |g_1 \dots e_j \dots g_N\rangle, \quad (1)$$

where z_j is the spatial position of ion j , k is the wave number of the weak coherent wave packet, and c_j is an amplitude that depends on the absorption probability, the spatial position, and the absorption frequency of the ion. The AFC is a well-separated periodic structure with frequency detuning $\delta_j = m_j \Delta$ (m_j is an integer). Studying the time evolution of the superposition state in Eq. (1) shows that in this state the ions will start to dephase. But the key feature of the AFC structure, with its periodicity Δ , is that it gives rise to a constructive emission exactly after a time $t = 1/\Delta$. The storage and retrieval efficiency of the AFC protocol in the forward direction can be written [6] as

$$\eta = (\alpha L/F)^2 e^{-\alpha L/F} e^{-7/F^2}. \quad (2)$$

Because of the trade-off between the coherent response of the sample (first), reabsorption (second), and dephasing (third) factors, there is an optimum value for the finesse as discussed in Ref. [6].

In our measurements we have two sets of pulses. They correspond to the preparation pulses described in Ref. [23] and the storage pulses described below. The amplitude of the storage pulse was adjusted by taking into account attenuation factors due to windows, optical components, pinhole, and the quantum efficiency of the single photon detector until the mean number of photons per pulse was about $\bar{n} \approx 0.1$ at the sample. To improve measurement statistics, weak Gaussian storage pulses were sent in 2000 times at a rate of 3 kHz. To measure the efficiency, the storage pulse, which had a 200 ns FWHM, was sent through the empty spectral pit. In the empty pit there are no ions absorbing the pulse; therefore this is a good reference to compare with the recall pulse. The photon counting results fitted to a Gaussian time distribution for the empty pit are shown (circles and dashed curve) in Fig. 3.

Next, an AFC structure with peak separation $\Delta = 1.2$ MHz, peak width $\gamma = 200$ kHz, and $\alpha L \approx 6$ is created and a Gaussian weak coherent storage pulse ($\tau_{\text{FWHM}} = 200$ ns) tuned to the $|1/2g\rangle \rightarrow |1/2e\rangle$ transition [Fig. 2(a)] is sent through the medium, which has a phase

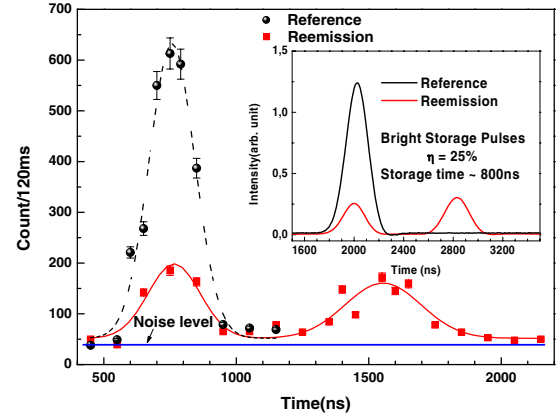


FIG. 3 (color online). Black circles (reference) are the single photon counts for the transmission of a 200 ns Gaussian storage pulse ($n = 0.1$ photon per pulse) sent through the transparent empty pit without any AFC grating. Red squares correspond to the counts for an identical storage pulse sent through the AFC spectral grating with $\Delta = 1.2$ MHz, $\gamma = 300$ kHz, and $\alpha L \approx 6$. The ratio of the areas below the echo and the reference after fitting with a simple Gaussian function and subtracting the noise level is 0.25. The same result is achieved with bright storage pulses which is shown in the inset.

relaxation time much longer than the duration of the pulse. Reemission (the echo) occurs after 800 ns ($1/\Delta$) (solid curve in Fig. 3). The measured storage and retrieval efficiency is 25%, which means that the energy contained in the collective reemission (echo) pulse relative to the energy transmitted through empty pit (dashed curve) is 0.25. Based on our knowledge this is the highest AFC echo efficiency in the single photon regime observed so far, and it is the product of our high controllable and precise ensemble-shaping technique.

So far, we have considered the efficiency properties of our interface, but for quantum memory applications it is vital that the interface also conserve the phase of the storage pulse with high fidelity. To show the phase preservation, a super-Gaussian ($n = 7$) pulse with controllable phase, overlapping the reemission pulse temporally, but with a 2.3 MHz frequency offset is prepared. This frequency offset allows us to send the super-Gaussian pulse through a transparent frequency window inside the pit between $|1/2g\rangle \rightarrow |1/2e\rangle$ and $|1/2g\rangle \rightarrow |3/2e\rangle$ transition peaks [23]. To have enough frequency space between these two transitions, the peak separation (Δ) is reduced to 1 MHz, but the rest of the AFC parameters are the same as before. In these measurements the τ_{FWHM} of the Gaussian pulse is 420 ns, whereas the super-Gaussian full width at half maximum is 840 ns.

Because of the interference between the two pulses, a beating pattern is expected at the detector at the time of the reemission (the echo) pulse (Fig. 4). The measured visibility after subtraction of the noise level is 83%. To show that the phase is conserved, the phase of the super-Gaussian pulse was changed by π and the beating pattern is reversed (Fig. 4).

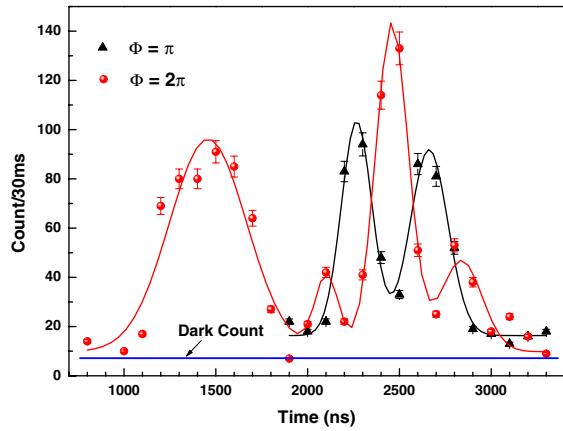


FIG. 4 (color online). Beating between reemission (echo) and the super-Gaussian ($n = 7$) pulses. The first pulse to the left is the transmitted part of the input storage pulse, which is independent of the phase of the super-Gaussian pulse. The visibility after subtracting the noise level is 83%. The interference patterns is inverted when the phase factor of the super-Gaussian pulse is changed by π , demonstrating phase conservation of the weak coherent storage pulse by the AFC spectral grating.

The 25% storage and retrieval efficiency obtained in this Letter is independent of the storage pulse intensity. Furthermore, all results with the bright storage pulses [23] are reproducible in the single photon regime. The experiment is based on coherent photon-echo-type reemission experiments in the forward direction. Theoretically, as discussed in Ref. [6], the maximum achievable efficiency for emission in the forward direction is 54%. This is basically limited by the reabsorption of the emitted echo. From an experimental point of view, the most critical issues for reaching this efficiency are first to have control over the AFC peak shape and second to obtain a high effective absorption ($\alpha L/F$). There is a good agreement between the effective absorption and obtained storage and recall efficiency which is discussed in Ref. [23]. To improve the efficiency beyond 54%, counterpropagating control fields [9] or external control of the ion frequency [26] could be used; alternatively, the crystal could be placed inside a cavity [27]. The efficiency could also be enhanced by using optimum input pulses [28–30].

This work was supported by the Swedish Research Council, the Knut and Alice Wallenberg Foundation, the Crafoord Foundation, and the European Commission through the integrated project QAP.

Note added in proof.—A quantum memory storage efficiency of 69% has been reported after the submission of this manuscript [31].

-
- [1] L. M. Duan, M. D. Lukin, J. I. Cirac, and P. Zoller, *Nature (London)* **414**, 413 (2001).
 - [2] H. J. Kimble, *Nature (London)* **453**, 1023 (2008).
 - [3] H. J. Briegel, W. Dur, J. I. Cirac, and P. Zoller, *Phys. Rev. Lett.* **81**, 5932 (1998).

- [4] W. Tittel, M. Afzelius, R. L. Cone, T. Chanelière, S. Kröll, S. A. Moiseev, and M. Sellars, *Laser Photon. Rev.* **4**, 244 (2009).
- [5] N. Sangouard, C. Simon, H. de Riedmatten, and N. Gisin, *arXiv:0906.2699*.
- [6] H. de Riedmatten, M. Afzelius, M. U. Staudt, C. Simon, and N. Gisin, *Nature (London)* **456**, 773 (2008).
- [7] S. E. Harris, *Phys. Today* **50**, No. 7, 36 (1997).
- [8] J. J. Longdell, E. Fraval, M. J. Sellars, and N. B. Manson, *Phys. Rev. Lett.* **95**, 063601 (2005).
- [9] S. A. Moiseev and S. Kröll, *Phys. Rev. Lett.* **87**, 173601 (2001).
- [10] M. Nilsson and S. Kröll, *Opt. Commun.* **247**, 393 (2005).
- [11] B. Kraus, W. Tittel, N. Gisin, M. Nilsson, S. Kröll, and J. I. Cirac, *Phys. Rev. A* **73**, 020302(R) (2006).
- [12] M. Afzelius, C. Simon, H. de Riedmatten, and N. Gisin, *Phys. Rev. A* **79**, 052329 (2009).
- [13] C. Simon, H. de Riedmatten, M. Afzelius, N. Sangouard, H. Zbinden, and N. Gisin, *Phys. Rev. Lett.* **98**, 190503 (2007).
- [14] J. Nunn, K. Reim, K. C. Lee, V. O. Lorenz, B. J. Sussman, I. A. Walmsley, and D. Jaksch, *Phys. Rev. Lett.* **101**, 260502 (2008).
- [15] M. D. Eisaman, A. Andre, F. Massou, M. Fleischhauer, A. S. Zibrov, and M. D. Lukin, *Nature (London)* **438**, 837 (2005).
- [16] K. S. Choi, H. Deng, J. Laurat, and H. J. Kimble, *Nature (London)* **452**, 67 (2008).
- [17] B. Lauritzen, J. Minár, H. de Riedmatten, M. Afzelius, N. Sangouard, C. Simon, and N. Gisin, *Phys. Rev. Lett.* **104**, 080502 (2010).
- [18] T. Chanelière, J. Ruggiero, M. Bonarota, M. Afzelius, and J. L. Le Gouët, *New J. Phys.* **12**, 023025 (2010).
- [19] L. Rippe, B. Julsgaard, A. Walther, Y. Ying, and S. Kröll, *Phys. Rev. A* **77**, 022307 (2008).
- [20] M. Afzelius, I. Usmani, A. Amari, B. Lauritzen, A. Walther, C. Simon, N. Sangouard, J. Minár, H. de Riedmatten, N. Gisin, and S. Kröll, *Phys. Rev. Lett.* **104**, 040503 (2010).
- [21] L. Rippe *et al.* (to be published).
- [22] B. Julsgaard, L. Rippe, A. Walther, and S. Kröll, *Opt. Express* **15**, 11 444 (2007).
- [23] A. Amari, A. Walther, M. Sabooni, M. Huang, and S. Kröll, *J. Lumin.* **130**, 1579 (2010).
- [24] R. W. Eqsall, R. L. Cone, and R. M. Macfarlane, *Phys. Rev. B* **52**, 3963 (1995).
- [25] L. Rippe, M. Nilsson, S. Kröll, R. Klieber, and D. Suter, *Phys. Rev. A* **71**, 062328 (2005).
- [26] A. Kalachev and S. Kröll, *Phys. Rev. A* **78**, 043808 (2008).
- [27] M. Afzelius, in *Proceedings of the Workshop on Rare-Earth-Ion-doped Solids for Quantum Information* (Lund University, Lund, Sweden, 2009).
- [28] A. Kalachev, *Phys. Rev. A* **76**, 043812 (2007).
- [29] I. Novikova, A. V. Gorshkov, D. F. Phillips, A. S. Sørensen, M. D. Lukin, and R. L. Walsworth, *Phys. Rev. Lett.* **98**, 243602 (2007).
- [30] N. B. Phillips, A. V. Gorshkov, and I. Novikova, *Phys. Rev. A* **78**, 023801 (2008).
- [31] M. P. Hedges, J. J. Longdell, Y. Li, and M. Sellars, *Nature (London)* **465**, 1052 (2010).