## Quantum Memory in a Microfabricated Rubidium Vapor Cell

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Scalability presents a central platform challenge for the components of current quantum network implementations that can be addressed by microfabrication techniques. We demonstrate a proof-of-principle realization of a high-bandwidth quantum memory in a warm alkali atom ensemble in a MEMS vapor cell compatible with wafer-scale fabrication. By applying an external tesla-order magnetic field, we explore a novel ground-state memory scheme in the hyperfine Paschen-Back regime, where individual optical transitions can be addressed in a Doppler-broadened medium. Working on the  $^{87}$ Rb D<sub>2</sub> line, where deterministic quantum dot single-photon sources are available, we demonstrate bandwidth-matching with 100s of MHz broad light pulses keeping such sources in mind. For a storage time of 80 ns we measure an end-to-end efficiency of  $\eta_{e2e}^{80 \text{ ns}} = 3.12(17) \%$ , corresponding to an internal efficiency of  $\eta_{\text{int}}^{0 \text{ ns}} = 24(3) \%$ , while achieving a signal-to-noise ratio of SNR = 7.9(8) with coherent pulses at the single-photon level.

Introduction.—Quantum networks [1, 2] are envisioned to enable secure quantum communication [3-5], distributed [6] and blind quantum computing [7], as well as performing precise measurements through novel forms of distributed quantum sensing [8–10]. Optical quantum memories and matched single-photon sources are the building blocks of such networks. Direct connections between individual network nodes have been realized [11], with complex systems such as cavity-trapped atoms [12] or hybrid interfaces [13, 14] regularly pushing performance records. Remote network operations have also been demonstrated both in cold ensembles [15] and diamond vacancy centers [16, 17]. As the principle feasibility of quantum networking becomes tangible, new central challenges including scalability, speed, and ease of use in deployment emerge. Platforms compatible with room-temperature operation and mass fabrication must enter any realistic vision of larger networks.

The experimental simplicity of warm alkali vapor memories is attractive for scaling quantum networks [18]. Recently, single-photon storage and retrieval has been demonstrated in broadband vapor-based memories both in ladder [19] and lambda-schemes [20]. Moreover, MEMS fabrication technology has already successfully miniaturized vapor cells for diverse applications, including quantum sensors such as atomic clocks, gyroscopes, and magnetometers [21]. However, a quantum memory has not vet been realized in a MEMS cell. Demands on vapor cell properties vary with application, and key memory demands including sufficient optical path and molecular buffer gas pressures on the order of 10s of millibar [22] are not typical features of microfabricated clock cells. Further, common methods of sealing MEMS cells such as anodic bonding involve temperatures incompatible with spin anti-relaxation coatings on the cell walls [23], which are critical to achieving long storage times in memories. Nevertheless, the state-of-the-art in cell fabrication has addressed these issues with techniques including cell-internal light routing, lower temperature bonding

methods, and techniques to achieve arbitrary buffer fillings on wafer scales [24–26]. These developments pave the way for scalability, spatial multiplexing – a wafer of suitable vapor cells potentially provides hundreds of independent memories – and could even enable satellite-borne applications [27].

In this letter we present an experimental realization of an optical quantum memory in a microfabricated vapor cell on the  ${}^{87}$ Rb  $D_2$  line. By applying a static, tesla-order, external magnetic field, degeneracy in the atomic level structure is lifted in such a way that a near-ideal threelevel lambda system can be addressed. As a proof-ofprinciple, weak coherent pulses attenuated to the singlephoton level are stored and retrieved, with operational bandwidths of hundreds of MHz. By investigating such fast signals, we demonstrate the applicability both of our novel scheme and of the miniaturized vapor cell to the task of storing single photons as they are produced by high quality solid state emitters – for instance GaAs quantum dots constitute excellent single-photon sources emitting at the Rb D<sub>2</sub> line [28]. Interfacing deterministic photon sources with vapor cell memories would enable network architectures such as the one proposed in Ref. [29]. Moreover, by evaluating memory performance and limitations, we establish a list of requirements for cell fabrication aimed at memory applications while simultaneously exploring atomic memory interactions in a hitherto untested physical parameter range.

Memory scheme and setup.—The quantum memory is implemented in a microfabricated atomic vapor cell with 2 mm internal thickness and an aperture diameter of 5 mm (see Fig. 1(a)). The interaction is based on a ground state lambda scheme. Atoms are initially prepared in state  $|g\rangle$  by optical pumping. An incoming signal photon is stored by mapping it to a collective spinwave excitation between the two metastable ground states  $|g\rangle$  and  $|s\rangle$  by a strong control pulse resonant to the second "leg" of the lambda system. By applying a second control pulse after a variable delay, the signal is

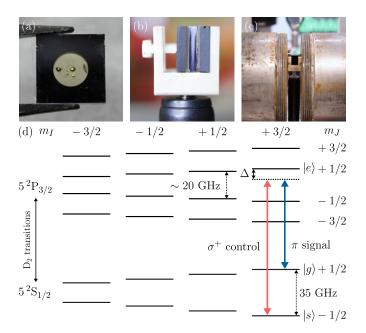


FIG. 1. Front (a) and side (b) view of the microfabricated vapor cell. Colored glass filters for absorbing the heating lasers sandwich the cell. (c) The ferromagnetic cores of the electromagnet limit the physical and optical access close to the cell. (d) Energy levels of <sup>87</sup>Rb in the external field and lambda-scheme used for the atomic memory. Energy splitting and decoupling of **J** and **I** yield a "clean" three-level system in the Doppler broadened <sup>87</sup>Rb D<sub>2</sub> line. The signal and control fields are on two-photon resonance with a detuning  $\Delta = -2\pi \times 750\,\mathrm{MHz}$  with respect to  $|e\rangle$ .

## retrieved from the memory.

Generally speaking, the energetic structure of atoms is more complex than just three levels. In order to avoid spurious noises processes, the isolation of a near-ideal three-level system is crucial. To accomplish this goal we operate the memory in the hyperfine Paschen-Back regime, reached by applying a large, static B-field. The resulting decoupling between nuclear spin manifolds, and the large energy level splittings within each manifold caused by the Zeeman effect, allow us to obtain the necessary resolution of individual transitions to address a spectrally well-isolated, "clean" three-level lambda-system for storage, as illustrated in Figure 1(d). The magnetic field is applied orthogonally to the propagation axis of the light, a configuration known as Voigt geometry, by a Bruker B-E 10 electromagnet for flexibility in testing (see Fig. 1(c)). In a future miniaturized setup we envision employing a suitably designed permanent magnet akin to what is used in [30]. For a thorough spectroscopic characterization of this regime, information on the viability of state preparation therein, and a broader overview of its usefulness in establishing quantum control in hot vapors we point to our concurrent article [31].

The memory operates on the  ${}^{87}{\rm Rb}$  D<sub>2</sub> line at  $780\,{\rm nm}$ 

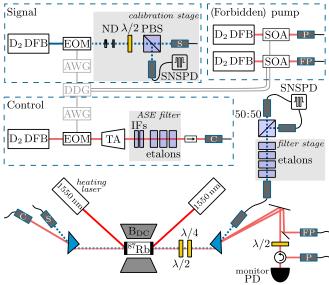


FIG. 2. Sketch of the experimental setup. The dashed boxes represent the preparation stages for the various optical pulses involved in the memory protocol. DFB, distributed feedback (laser); EOM, electro-optic modulator; AWG, arbitrary waveform generator; DDG, digital delay generator; SOA, semiconductor optical amplifier; TA, tapered amplifier; IF, interference filter;  $\lambda/2$ , half-wave plate;  $\lambda/4$ , quarterwave plate; 50:50, beam splitter; SNSPD, superconducting nanowire single-photon detector. The labels S, C, P, and FP represent the fiber links of signal, control, pump, and forbidden pump, respectively.

in a 1.06 T magnetic field. The vapor cell is filled with enriched  $^{87}{\rm Rb}$  (abundance  $\leq 90\,\%$ ) and about 11 mbar of Ar buffer gas to impede atomic motion. The fabrication process of such MEMS type cells is described in [32, 33]. The cell is sandwiched by two 2 mm-thick pieces of RG9 filter glass (see Fig. 1(b)) and heated with two multimode, telecom lasers in order to increase the Rb vapor pressure and reach moderate optical depths. With this heating technique we can achieve atomic temperatures  $> 130\,^{\circ}{\rm C}.$ 

We choose the signal to be near resonant to the  $\pi$ -transition coupling the stretched state  $|g\rangle = |m_J = +\frac{1}{2}, m_I = +\frac{3}{2}\rangle$  to the excited state  $|e\rangle = |m_J = +\frac{1}{2}, m_I' = +\frac{3}{2}\rangle$  for its higher transition strength. Consequently, the storage state is  $|s\rangle = |m_J = -\frac{1}{2}, m_I = +\frac{3}{2}\rangle$  and the control field must be  $\sigma^+$  polarized. Initially, the  $m_I = +\frac{3}{2}$ -manifold is prepared in  $|g\rangle$  through optical pumping. A second pump laser is used to drive the singly forbiden transition  $|m_J = +\frac{1}{2}, m_I = +\frac{1}{2}\rangle \longleftrightarrow |m_J' = -\frac{3}{2}, m_I' = +\frac{3}{2}\rangle$ , to partially polarize the nuclear spin and increase the number of atoms in the addressed  $m_I$ -manifold. A working point detuning  $\Delta = -2\pi \times 750\,\mathrm{MHz}$  from the excited state  $|e\rangle$  is empirically found to optimize the memory performance under experimental operating conditions.

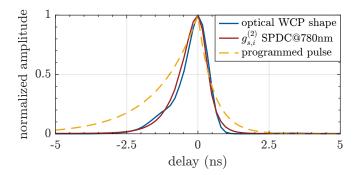


FIG. 3. Signal pulse shape. The blue trace shows the optical signal pulse, measured as input. The target shape, consisting of the cross-correlation  $g_{s,i}^{(2)}$  from the downconversion source in [20], is shown in red. For comparison, the dashed yellow line shows the voltage pulse programmed in the AWG.

The optical setup is sketched in Fig. 2. Optical pulses are generated from CW DFB lasers by amplitude modulation with EOMs (Jenoptik AM785), with their waveforms set by an AWG (PicoQuant PPG512). The signal pulses are subsequently attenuated to the single-photon level and their temporal shape, shown in Fig. 3, is matched to that of heralded photons from an SPDC source described in [20]. The Gaussian-shaped control pulses are amplified to the required intensities with a TA. Amplified spontaneous emission (ASE) from the TA is filtered with two interference filters (Laseroptik, 0.37 nm FWHM bandwidth, specified by manufacturer) and three monolithic etalons (two 1150(20) MHz FWHM bandwidth; one 280(10) MHz FWHM bandwidth). All etalons herein reach suppressions of -33(1) dB at FSR/2.

Signal and control are each coupled into single-mode (SM) fibers, and are combined at the memory stage on a polarizing calcite prism. The signal (control) is focused with the outcoupler to a  $1/e^2$  diameter at the center of the cell of  $185(6) \, \mu m$  ( $440(6) \, \mu m$ ). The beam waists are measured at an equivalent distance from the cell center with a beam profiler. The control beam is chosen to be larger for a more homogeneous Rabi frequency across the spatial mode of the signal, which is known to improve the memory efficiency at lower peak  $\Omega$  in simulation [20].

The atomic ensemble is optically pumped before each storage attempt with a dedicated pump laser tuned to resonance with  $|s\rangle \rightarrow |e\rangle$ . The pump counter-propagates with the control beam, and is coupled in through an optical circulator in the control arm. A semiconductor optical amplifier is used as a fast optical switch to turn off the pumping beam, as described in [34]. The forbidden transition is pumped by a further laser, which is aligned under a small angle (estimated trigonometrically to be 2.6(9) mrad) to the main pump using a D-shaped mirror. Both pump lasers are collimated and have a diameter of approximately 2 mm. In order to maximize depletion of  $|s\rangle$ , the pump laser is switched off 50 ns later than the

forbidden pump. The optical power on the atoms from the pump and forbidden pump lasers is  $14.8(4)\,\mathrm{mW}$  and  $16.0(5)\,\mathrm{mW}$  respectively.

A second prism filters the retrieved signal by polarization. Over eight orders of magnitude of control suppression are achieved this way. After coupling the signal into an SM fiber a stack of monolithic etalons filters spectrally. Three of these are chosen to have an FSR = 71.1 GHz, equal to twice the ground state splitting of <sup>87</sup>Rb in the external field (1.45 mm thickness, FWHM bandwidth 1.19(2) GHz), and one (4 mm thickness, FWHM bandwidth 550(10) MHz) matches the bandwidth of the retrieved photon. Finally, the signal is evenly split, fiber coupled, and detected with SNSPD (Single Quantum EOS). When the vapor cell is cold, the transmission through the whole setup at the signal frequency is measured to be slightly below 20 % for a strong CW probe.

Currently, the experiment is repeated periodically and triggered by a digital delay generator (Highland Technology T564) with a repetition rate of 300 kHz, allowing for 2.8 µs of optical pumping. Nevertheless, electronic and optical switches are all suitable for asynchronous operation, as is required when paired with a probabilistic single photon source (cf. Ref. [20]).

Storage and retrieval.—Memory experiments were performed for a storage time of about 80 ns at an atomic temperature T = 90(1) °C, determined spectroscopically. Gaussian control pulses with a duration of FWHM = 3.8 ns were used, with a maximum available peak Rabi frequency of  $2\pi \times 683(15)$  MHz at the given waist. The signal pulse attenuation was calibrated right before the storage and retrieval measurement. From the counts measured on the calibration channel, within a 6.48 nswide region of interest (ROI) around the signal pulse, a mean photon number of  $|\alpha|^2 = 0.97(6)$  was set. Detection events are tagged with a time-to-digital converter (qutools quTAU). The counts from the Hanbury Brown and Twiss configured detectors after the memory are added for evaluation. All figures of merit are specified using a 6.48 ns ROI.

A photon arrival-time histogram is shown in Fig. 4(a). For an integration time of  $t_{\rm int} \approx 1 \, \rm min$ , the memory receives  $N_{\rm trig} = 1.81 \times 10^7$  triggers, each corresponding to a storage attempt, and  $N_{\rm ret} = 4.46 \times 10^5$  counts accumulate within the retrieval window. To estimate noise during read-out, the experiment is repeated with the input physically blocked. Within the same ROI,  $4.28 \times 10^4$ noise counts are detected. As this noise measurement does not account for the finite suppression of the signal EOM during the retrieval time window, we add the averaged difference of 183 counts per bin between the storage and noise traces far from signal switching to the noise counts, obtaining  $N_{\text{noise}}$ . Over the ROI we estimate that this offset adds 7325 spurious counts. The detected counts correspond to a signal-to-noise ratio of  $(N_{\text{ret}} - N_{\text{noise}})/N_{\text{noise}} = 7.9(8).$ 

From the latter quantity  $\mu_1 = \eta_{\rm det}^{\rm HBT}/{\rm SNR} = 0.09(1)$  can be computed, describing the required mean photon number at the input to reach SNR = 1. Here  $\eta_{\rm det}^{\rm HBT} = 70(4)\,\%$  is the detection probability of a weak coherent pulse in an HBT setup with non-photon-number-resolving detectors given by  $\eta_{\rm det}^{\rm HBT}(|\alpha|^2) = 2\left(1-\exp(-|\alpha\sqrt{\eta_{\rm det}/2}|^2)\right)$ , where  $\eta_{\rm det}$  is the detection efficiency of the SNSPD. The measurements yield an end-to-end efficiency of  $\eta_{e2e}^{80\,\rm ns} = (N_{\rm ret}-N_{\rm noise})/(\eta_{\rm det}^{\rm HBT}N_{\rm trig}) = 3.12(17)\,\%$ . In the region between the read-in and read-out control pulses in Fig. 4(a) unintentional retrieval can be observed. This is caused by ringing in the control EOM which is translated to an optical signal. These detrimental features account for a 0.74(4) % loss in end-to-end efficiency.

The memory lifetime is measured by the decrease in efficiency over time. This measurement was performed with a repetition rate of 100 kHz, allowing us to use the same pump pattern for all data, guaranteeing unaltered initial conditions. The end-to-end efficiency as a function of storage time is plotted in Fig. 4(b). The lifetime data are scaled by a factor 1.13 to match the measured efficiency for a storage time of 80 ns to the one obtained from the data set shown in Fig. 4(a). This discrepancy is a question of day-to-day optimization and the correction is applied to accurately capture the internal efficiency of the main result. Without any direct optimization attempt, a 1/e lifetime of 224(8) ns was achieved. The data are well fit by a decaying exponential (solid line). This lifetime is shorter than expected from simple models of atomic motion out of the interaction region [35]. A limitation due to motional dephasing of the spin wave can be ruled out as it would be described by Gaussian decay [36]. A possible explanation lies in spin destruction collisions with the cell walls, which are uncoated and always nearby as the cell is so thin. Correcting for technical losses by dividing out the signal transmission through the setup, the internal efficiency at zero storage time is estimated to be  $\eta_{\text{int}}^{0 \text{ ns}} = 24(3) \%$ .

Discussion.—The microfabricated vapor cell design and its filling were not optimized for memory application. leaving much room for future improvement. A pumpprobe relaxation in the dark measurement shows that an atomic polarization of 88(1)% in  $|g\rangle$  is reached within the  $m_I = +\frac{3}{2}$ -manifold [37]. Currently, the polarization is most likely limited by radiation trapping due to the high Rb number density n at the temperature required for efficient storage. At 90 °C and accounting for 90 % enrichment, the number density in the considered nuclear spin manifold is  $n = 5.5 \times 10^{11} \, \mathrm{cm}^{-3}$ . A molecular buffer gas, such as N2, would quench the excited state and improve the atomic polarization [38]. By more fully depleting the storage state in this manner, the contribution of spontaneous Raman scattering to the read-out noise would be significantly reduced.

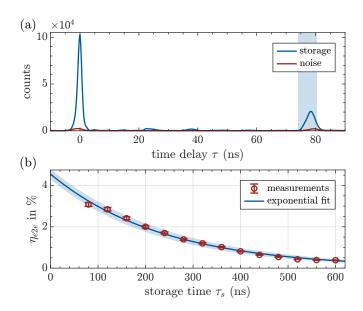


FIG. 4. (a) Arrival-time histogram for storage and retrieval experiments and corresponding noise measurements. Zero time delay corresponds to the time at which signal photons leaking through the memory in failed storage attempts reach the detector. The shaded area corresponds to the 6.48 ns-wide ROI. The features between 20 ns to 40 ns constitute unintentional read-out and are induced by insufficient suppression of the control EOM. (b) Memory lifetime measured as the decrease in efficiency. Each data point is integrated over 2 min and the data are scaled by a factor 1.13 to match the performance of the data set shown in (a). A fitted exponential decay describes the data well. The shaded area corresponds to the 95 % confidence interval of the fit.

Low optical depth limits the total internal efficiency of the memory. In the explored temperature range, an OD of 1 to 2.5 is reached on the unpumped signal transition. According to the authors of Ref. [39], for OD = 5 (full atomic polarization) the maximal total efficiency achievable with forward retrieval is limited to < 30 %. Given this, our memory is remarkably efficient. Considering the magnetic field homogeneity, characterized in [37], a longer cell could be used to increase the OD at constant Rb number density. For a 10 mm-long vapor cell, frequency shifts of at most a few megahertz are expected. By internally routing the light within the cell, longer optical paths can be obtained exploiting the capabilities of wafer-based fabrication techniques [40–42]. Furthermore, by designing the transverse profile of the cell to match the interaction region and by applying anti-relaxation coatings to the cell walls the lifetime of the memory could be extended.

The discrepancy between the measured efficiency and the theoretical expectation can be fully resolved by effects we account for. The largest contribution is caused by unintentional read-out occurring during storage, constituting nearly a quarter of the achieved end-to-end efficiency. This eliminates the discrepancy within experimental uncertainty. In order to address this issue an investigation into improving the extinction ratio of the control field is ongoing. First promising results were obtained by directly switching the TA's operating current. Over six orders of magnitude of suppression of the control field are reached almost immediately upon switching [37], which should prevent perturbation of the spinwave during storage. Additionally, it would be interesting to investigate how the cell's flat form-factor affects the remission mode, as it does not appear to remain perfectly matched to the SM fiber mode defined at the input. We also note that for efficient optical pumping its aspect ratio is particularly inconvenient.

In summary, our proof-of-principle implementation of a ground-state quantum memory in a microfabricated vapor cell opens up a route towards realistic scaling of quantum networks. Moreover, we have shown that our novel high magnetic field scheme isolates a near-ideal three-level system, even on the <sup>87</sup>Rb D<sub>2</sub> line, promising highly efficient and low noise performance in an optimized cell where the necessary optical depths can be reached at lower number densities. Once the atomic state preparation is further improved, interfacing the memory with a compatible single-photon source, e.g. based on down-conversion [20] or a GaAs quantum dot [28] will be attempted. In fact, we note that the presented memory is already built for asynchronous operation, capable of on-demand read-in and read-out.

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