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### Abstract

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## Multimode quantum memory based on atomic frequency combs

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An efficient multimode quantum memory is a crucial resource for long-distance quantum communication based on quantum repeaters. We propose a quantum memory based on spectral shaping of an inhomogeneously broadened optical transition into an atomic frequency comb (AFC). The spectral width of the AFC allows efficient storage of multiple temporal modes without the need to increase the absorption depth of the storage material, in contrast to previously known quantum memories. Efficient readout is possible thanks to rephasing of the atomic dipoles due to the AFC structure. Long-time storage and on-demand readout is achieved by use of spin states in a lambda-type configuration. We show that an AFC quantum memory realized in solids doped with rare-earth-metal ions could store hundreds of modes or more with close to unit efficiency, for material parameters achievable today.

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### I. INTRODUCTION

The distribution of entanglement between remote locations is critical for future long-distance quantum networks and extended tests of quantum nonlocality. It is likely to rely on quantum repeaters [1,2], which require quantum memories (QMs) that can store entanglement between distant network nodes [3,4]. Recent experimental achievements in quantum state storage [5–8] demonstrate that currently investigated QMs can store a single mode. Yet, long-distance quantum repeaters having QMs that are only capable of storing one mode would only generate very limited entanglement generation rates [9]. To achieve useful rates some way of multiplexing the QM will be required [10,11]. By using time [11], spatial [10,12,13], or frequency multiplexing to store single photons in many modes  $N$ , the entanglement generation rate can be increased by a corresponding factor  $N$  [11]. Here we consider multimode QMs capable of storing  $N$  temporally distinguishable modes, which is a natural form of sending information also used in today's telecommunications networks. Time multiplexing is extremely challenging when using current QM protocols such as stopped light based on electromagnetically induced transparency (EIT) [14], photon echoes based on controlled reversible inhomogeneous broadening (CRIB) [15,16], or off-resonant Raman interactions [17]. For a single mode,  $N=1$ , the efficiency as a function of optical absorption depth  $d$  is similar for EIT, CRIB, and Raman-type memories [18,19]. For multimode storage using EIT and Raman, however,  $N$  scales as  $\sqrt{d}$  [20,21], which severely limits the number of potential modes in a practical experiment, while CRIB offers a better scaling since  $N \sim d$  [11,21]. Still, even using CRIB, efficient multimode storage requires extreme absorption depths that are currently out of reach.

In this paper we present a QM, where the number of potential modes is essentially independent of the optical depth. As we will show, efficient storage of hundreds of modes or more is then feasible with realistic material parameters. The QM is based on absorption of light by an ensemble

of atoms on an optical transition which is inhomogeneously broadened. Ensembles are in general very attractive as QMs due to strong collective enhancement of the light-matter coupling [2]. Storage of single photons using EIT (stopped light) has been demonstrated with cold alkali atoms [6,8] which can be treated as homogeneous ensembles of identical atoms. Here we will instead consider ensembles of rare-earth-metal (RE) ions in solids, which are inhomogeneously broadened. Stopped light with storage times up to 1 s has been demonstrated in RE-doped solids [22,23], where an approximate homogeneous ensemble was created by spectrally isolating a narrow absorption peak through optical pumping. In contrast to this approach, the quantum memory proposed here uses the inhomogeneous broadening as a resource in order to achieve better multimode performance. For that one needs to coherently control the dephasing which is caused by the inhomogeneous frequency distribution of the atoms. The key feature of our proposal is to achieve this control by a specific shaping of this distribution into an atomic frequency comb.

The paper is organized in the following way. In Sec. II we give an overview of the proposal using a simplified physical picture. In Sec. III we show results from an analytical analysis of the physics of atomic frequency combs, which is further detailed in Appendix. In Sec. IV the multimode storage capacity is discussed, and in Sec. V we discuss implementation of the quantum memory in rare-earth-ion-doped solids. Conclusions are given in Sec. VI.

### II. ATOMIC FREQUENCY COMBS

The storage material is an ensemble of atoms with an excited state  $|e\rangle$  optically connected to two lower states  $|g\rangle$  and  $|s\rangle$ , see Fig. 1. The states  $|g\rangle$  and  $|s\rangle$  are typically ground-state hyperfine or Zeeman states. We assume that the optical transition  $|g\rangle-|e\rangle$  has a narrow homogeneous linewidth  $\gamma_h$  and a large inhomogeneous broadening  $\Gamma_{in}$  ( $\Gamma_{in}/\gamma_h \gg 1$ ). This is the case for RE-doped solids that we will consider for physical implementation in Sec. V. The  $|g\rangle-|e\rangle$  transition is spectrally shaped such that the atomic density function consists of a series of narrow peaks spanning a large frequency range, which we call an atomic frequency comb (AFC). This

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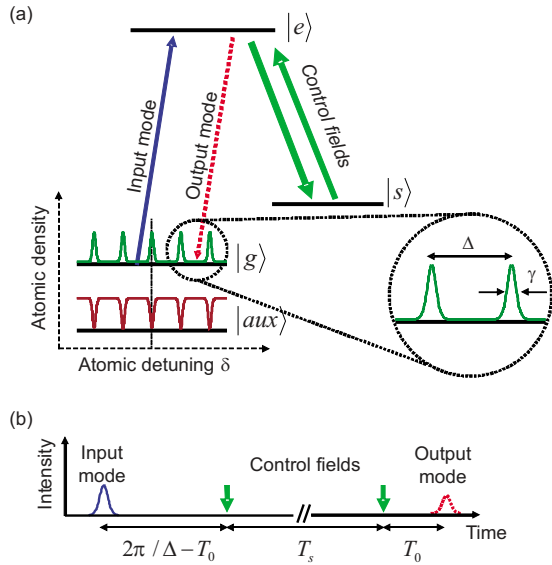


FIG. 1. (Color online) The principles of the proposed AFC quantum memory. (a) An inhomogeneously broadened optical transition  $|g\rangle\text{-}|e\rangle$  is shaped into an AFC by frequency-selective optical pumping to the  $|aux\rangle$  level. The peaks in the AFC have width (FWHM)  $\gamma$  and are separated by  $\Delta$ , where we define the comb finesse as  $F=\Delta/\gamma$ . (b) The input mode is completely absorbed and coherently excites the AFC modes, which will dephase and then rephase after a time  $2\pi/\Delta$ , resulting in a photon-echo type coherent emission. A pair of control fields on  $|e\rangle\text{-}|s\rangle$  allows for long-time storage as a collective spin wave in  $|s\rangle$ , and on-demand readout after a storage time  $T_s$ .

can be done by frequency-selectively transferring atoms from  $|g\rangle$  to a metastable state  $|aux\rangle$ , for instance a third hyperfine state, through optical pumping techniques (see Sec. V). Note that the maximum spectral width of the AFC is limited by the level spacings between the ground-state levels.

We then consider a single-photon input field, having a spectral distribution  $\gamma_p$  larger than the AFC peak separation  $\Delta$ , but narrower than the total width of the AFC  $\Gamma$ , which is in resonance with the  $|g\rangle\text{-}|e\rangle$  transition. If the atomic density integrated over the photon bandwidth is high enough, the photon can be totally absorbed by the AFC although the spectral density of atoms is concentrated to narrow peaks. This can be understood in terms of the Heisenberg energy-time uncertainty relation. For the time scale of the absorption, which is of the order of the input pulse duration  $\tau=1/\gamma_p$ , the optical transition will have an uncertainty of the order of  $\gamma_p \gg \Delta$ . This causes a spectral averaging of the sharp AFC structure into a smooth distribution, allowing for uniform absorption over the photons bandwidth (see also Sec. III).

After absorption (at  $t=0$ ) the photon is stored as a single excitation delocalized over all the atoms in the ensemble that are in resonance with the photon. The state is described by a collective Dicke state [24]

$$|\psi\rangle = \sum_{j=1}^N c_j e^{i\delta_j t} e^{-ikz_j} |g_1 \cdots e_j \cdots g_N\rangle, \quad (1)$$

where  $z_j$  is the position of atom  $j$ ,  $k$  is the wave number of the light field (for simplicity, we only consider a single spa-

tial mode defined by the direction of propagation of the input field),  $\delta_j$  the detuning of the atom with respect to the laser frequency, and the amplitudes  $c_j$  depend on the frequency and on the spatial position of the particular atom  $j$ .

The collective state can be understood as a coherent excitation of a large number of AFC modes by a single photon. These modes are initially (at  $t=0$ ) in phase with respect to the spatial mode  $k$ . But the collective state will rapidly dephase into a noncollective state that does not lead to a strong collective emission since each term acquires a phase  $\exp(i\delta_j t)$  depending on the detuning  $\delta_j$  of each excited atom. If we consider an AFC having very sharp peaks, then the detunings  $\delta_j$  are approximately a discrete set such that  $\delta_j = m_j \Delta$ , where  $m_j$  are integers. It follows that the collective state is re-established after a time  $2\pi/\Delta$ , which leads to a coherent photon-echo [25–27] type re-emission in the forward direction. The efficiency of this process (see Sec. III for details) can reach 54% in the forward direction (limited only by reabsorption). But if the re-emission is forced to propagate in the backward direction, by a proper phase matching operation (see below), the process can reach 100% efficiency.

The process described so far only implements a QM with a fixed storage time. In order to allow for on-demand readout of the stored field (which is a necessary requirement for use in quantum repeaters) and long-term storage, the single collective excitation in  $|e\rangle$  is transferred to a ground-state spin level  $|s\rangle$ . This can be done by applying an optical control field on  $|e\rangle\text{-}|s\rangle$ , for instance a short  $\pi$  pulse. The excitation is thereafter stored as a collective spin wave, which is also the basis of storage in EIT, CRIB, and off-resonant Raman-type memories based on ensembles of lambda atoms [14,15,17]. The spin wave allows for long-lived storage since spin coherence lifetimes are generally longer than the optical coherence lifetimes. Moreover, the spin transition does not have a comb structure [28], which means that the evolution of the phases in Eq. (1) will be locked during the storage in the spin wave. If we assume that the spin transition is completely homogeneous, the storage time is only limited by the coherence lifetime of the spin wave. If there is inhomogeneous broadening, spin-echo techniques can be used to exploit the homogeneous coherence lifetime of the system (see Sec. V). To retrieve the field a second counterpropagating control field is applied after some time  $T_s$ . These operations will result in a backward propagating output mode after a total time  $T_s+2\pi/\Delta$ . As we will show below, the efficiency of the storage for backward retrieval can theoretically approach unity for high optical depths.

### III. ANALYTICAL TREATMENT

We will now discuss some important features of the AFC and give quantitative formulas relating the input field to the output field as a function of the AFC parameters. This can be used to calculate, for instance, storage efficiency. To this end we have used an approximate linearized Maxwell-Bloch (MB) model. A linearized model can be used since the population in the excited state is negligible. For a single-photon input field at most one atom in the ensemble is excited. For

multimode storage, the number of totally excited atoms should be much less than the total number of atoms, which is the case for the number modes we consider and the RE solids we discuss in Sec. V. The linearity of the analytical model also implies the equivalence between classical and single-photon dynamics [17,18,29]. The important features of the model are discussed below while the details of the calculations are given in Appendix.

The matter part was assumed to be an ensemble of two-level atoms, including levels  $|g\rangle$  and  $|e\rangle$  in Fig. 1, and the light field was treated as a one-dimensional spatial mode. This model can fully describe the mapping of light onto the atoms, the atomic evolution including the inhomogeneous dephasing and rephasing of the AFC modes, and the re-emission of light. The model did not include population relaxation ( $T_1$ ) or decoherence ( $T_2$ ). This is a valid approximation since the peaks in the AFC are assumed to be inhomogeneously broadened. Then all relevant dephasing times are much faster than  $T_1$  and  $T_2$  (as in the example given in Sec. V). In practice, however, the width of the individual peaks will be limited by the homogeneous linewidth  $\gamma_h = 1/(\pi T_2)$ . The two counterpropagating control fields were modeled by transferring the forward atomic mode, associated with the forward propagating optical mode, to a backward atomic mode [29]. This achieves the phase matching operation that results in a backward propagating optical mode. Clearly this approach assumes that the transfer efficiency of the control pulses is unity and that the spin transition  $|g\rangle\text{-}|s\rangle$  is decoherence-free. We thus only analytically treat the physics of the atomic frequency comb, which is the unique feature of this protocol as compared to previous quantum memory proposals [14,15,17]. The transfer efficiency and spin decoherence could, however, be straightforwardly included with appropriate loss factors. The equations of motion describing the light-matter dynamics is given in Appendix [see Eqs. (A1)–(A4)]. In addition to the analytical calculations, we also show results from numerical solutions of the MB model.

The atomic spectral distribution  $n(\delta)$  is a frequency comb described by a series of Gaussian functions characterized by the peak separation  $\Delta$  and peak width  $\tilde{\gamma}$ , with an overall comb width of  $\Gamma$

$$n(\delta) \propto e^{-\delta^2/(2\Gamma^2)} \sum_{j=-\infty}^{\infty} e^{-(\delta - j\Delta)^2/(2\tilde{\gamma}^2)}. \quad (2)$$

The full width at half-maximum (FWHM) peak width is then  $\gamma = \sqrt{8 \ln 2} \tilde{\gamma}$ , and we define the finesse  $F$  of the comb as  $F = \Delta/\gamma$ . We are interested in the regime where we have a significant number of well-separated peaks within the overall characteristic width of the AFC. Moreover, we assume that the spectrum of the photon is narrower than the width of the AFC but wider than the peak separation, i.e.,  $\Gamma \gg \gamma_p \gg \Delta \gg \tilde{\gamma}$ .

In order to have an efficient memory, the probability of absorption of the input mode must be sufficiently high. But to create a high finesse AFC atoms are removed, by optical pumping, from the natural inhomogeneous spectrum. This preparation step lowers the total density of atoms and hence

the probability of absorption. The analytical calculations show that the amplitude of the forward propagating mode  $E_f(z, t)$  after the sample of length  $L$  is given by [see Eqs. (A12) and (A13)]

$$E_f(L, t) = E_f(0, t) e^{-\tilde{\alpha}L/2} = E_f(0, t) e^{-\tilde{d}L/2}, \quad (3)$$

where the effective absorption depth  $\tilde{d} = \tilde{\alpha}L$  can be expressed as

$$\tilde{d} = \frac{d}{F} \sqrt{\frac{\pi}{4 \ln 2}} \approx \frac{d}{F}. \quad (4)$$

The comb absorption is thus reduced by a factor of  $1/F$ , which is to be expected since the total density of atoms in the comb (integrated over the frequency bandwidth of the photon) is reduced by a factor of  $1/F$  in the preparation step. But if the peak absorption depth  $d$  of the comb is high enough, the total density of atoms in the comb can still be sufficiently high to absorb the input mode with close to 100% probability.

An interesting feature of the AFC is that it can absorb uniformly over the frequency spectrum of the input mode although the atomic frequency comb has large ‘‘holes.’’ This can be understood by considering that the absorption is an event well localized in time, of the order  $\tau = 1/\gamma_p$ . In that case the light-atom system reacts with a Fourier-limited resolution  $\gamma_p$ , which effectively smears out the structure of the AFC to an essentially flat distribution [see discussion preceding Eq. (A11)]. In other words the peak separation  $\Delta$  must be smaller than the photon bandwidth  $\gamma_p$ . This also implies that Eq. (3) is only valid for times  $t \ll 2\pi/\Delta$ , i.e., before the appearance of the first emission due to rephasing of modes in the comb.

By applying the two counterpropagating control fields, the spatial phase pattern of the atomic polarization changes such that it couples with the backward propagating field mode [15,16,29]. Under the assumption of unit transfer efficiency and no phase decoherence during the transfer to or storage in  $|s\rangle$ , the atomic phases will evolve to a collective state coupled to the backward propagating mode. This leads to emission in the backward mode. In Appendix [see Eq. (A18)] we show that the output field amplitude is a coherent sum over the amplitudes corresponding to all possible paths of the photon in the medium, where each path is characterized by a particular turning point in the medium. Even though there is absorption also along the path of the backward mode, a constructive interference effect between all these possible paths leads to the output amplitude potentially being arbitrarily close in magnitude to the input amplitude for large enough  $d$ . This interference effect is also present in the CRIB quantum memory scheme for backward retrieval [29]. The analytical calculation leads to the following relation between the forward input field  $E_{\text{in}} = E_f(z=0, t=0)$  and output backward field  $E_{\text{out}} = E_b(z=0, t=2\pi/\Delta)$  [see Eq. (A18)],

$$E_{\text{out}} = -E_{\text{in}} e^{-i2\pi(\Delta_0/\Delta)} (1 - e^{-\tilde{d}}) e^{-(1/F^2)(\pi^2/4 \ln 2)}. \quad (5)$$

The first factor represents a global phase shift that depends on the relative detuning  $\Delta_0$  between the carrier frequency of the light and the center of the AFC. The second factor de-



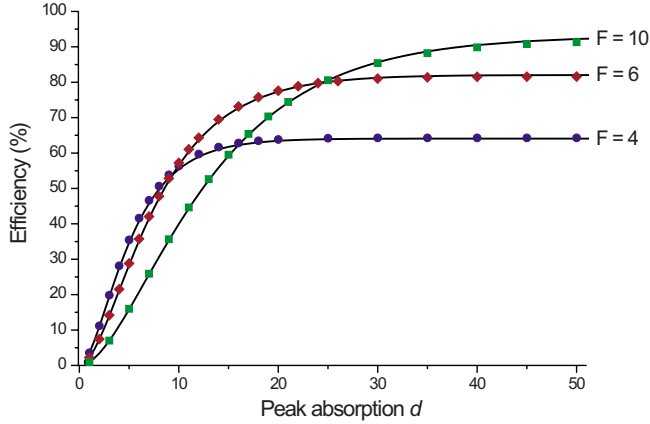


FIG. 2. (Color online) The storage efficiency is shown as a function of peak absorption depth  $d$  for AFC finesse values of  $F=4$ , 6, and 10. The solid lines represent the efficiency calculated using the analytical result, Eq. (5), whereas the symbols represent the numerical calculation. As an example, for  $d=40$  and  $F=10$ , an efficiency of 90% can be achieved. Even for an absorption depth of 25 and a modest finesse of 6, the efficiency reaches 80%.

describes the coupling between the input and output fields, which increases with increasing effective absorption depth, Eq. (4), whereas the third factor takes into account dephasing due to the finite width of the peaks.

The dephasing factor can be understood by considering the dephasing during the storage in the optically excited state  $|e\rangle$ . In the qualitative description in Sec. II we assumed that each peak was a delta function, which leads to complete rephasing of collective state Eq. (1). Clearly finite Gaussian peaks will not lead to complete rephasing, which lowers the probability of re-emission. The decay of the re-emitted field amplitude is given by the Fourier transform of a peak in the comb [see Eq. (A6) and discussion preceding Eq. (A16)], in analogy with the role of the initial peak shape in the CRIB quantum memory protocol [29,30]. In the case of Gaussian peaks, the decay is given by  $e^{-t^2\tilde{\gamma}^2/2}$ , where  $t=2\pi/\Delta$  is the storage time in the excited state. This factor can be expressed in terms of the finesse  $F$ , which leads to the last term in Eq. (5). To reduce this dephasing one should thus maximize the comb finesse  $F$ .

The efficiency, defined as  $\eta=|E_{\text{out}}|^2/|E_{\text{in}}|^2$ , is plotted as a function of optical depth for different comb finesses in Fig. 2. For a moderate finesse of  $F=4$ , an efficiency greater than 50% can be achieved for  $d=10$ . For a high finesse,  $F>10$ , the efficiency can be close to 100% for large enough  $d$ . From an experimental point of view  $d$  is often a fixed parameter. The storage efficiency can then be maximized for the given material by optimizing  $F$  using Eq. (5) (see Fig. 3). This optimum in the choice of  $F$  is due to a tradeoff between absorption probability (favoring small  $F$ ) and low dephasing during storage (favoring large  $F$ ).

We will also briefly discuss the possibility of re-emission in the forward direction. This will occur if no control fields are applied or if the control fields are applied copropagating. As is shown in Appendix [see Eq. (A19)], this re-emission is limited in efficiency to 54% for an optimal optical depth of  $\tilde{d}=2$ . The optimum represents a tradeoff between strong

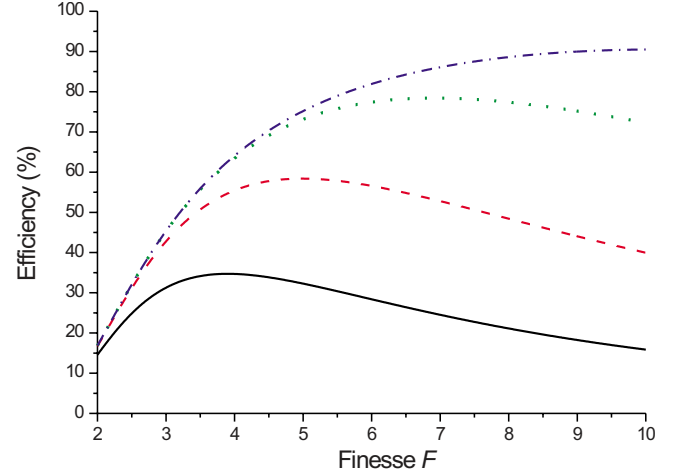


FIG. 3. (Color online) Efficiency as a function of comb finesse  $F$  for peak absorption depths  $d=5$  (black solid line),  $d=10$  (red dashed line),  $d=20$  (green dotted line), and  $d=40$  (blue dashed-dotted line). The curves were calculated using Eq. (5).

light-matter coupling (favoring high optical depth) and reabsorption of the re-emitted light as it propagates through the sample (favoring low optical depth).

#### IV. MULTIMODE STORAGE CAPACITY

The excellent multimode properties of the AFC based QM can be understood qualitatively as follows. We imagine that the input is a series of temporally distinguishable modes containing, with a certain probability, a single photon in each mode. The storage efficiency is uniform over all the modes since each mode spends the same time in the memory. Note that the effect of spin decoherence is also the same on each mode. If the control fields are applied as late as possible, the total duration of the input can be close to  $T=2\pi/\Delta$ , the time it takes to rephase the first mode that was absorbed. The shortest duration  $\tau$  of one mode, however, is only limited by the total frequency bandwidth  $\tau\sim 1/(N_p\Delta)$ , where  $N_p$  is the number of peaks in the comb. The number of modes  $N=N/\tau\sim N_p$  is then proportional to  $N_p$ , such that adding more peaks in the AFC allows one to store more modes. Thus, in materials with inherently large inhomogeneous broadenings (see Sec. V), one can increase the multimode capacity by making use of a wider range of the inhomogeneous broadening without having to increase the peak absorption depth by, for instance, increasing the atomic density in the sample.

We can make a more detailed calculation of the multimode capacity by assuming that the input modes have Gaussian temporal shapes, where each mode occupies a temporal slot of duration  $\tau$ . It is important that the modes do not overlap too much, which in the end would produce errors. The amount of overlap that can be accepted depends on the particular application. In Ref. [11] an analysis of the error due to overlap was carried out in the context of a quantum repeater protocol in which the AFC QM could be used. There it was shown that  $\tau$  should be chosen as  $\tau=12\pi/\Gamma$ , where  $\Gamma$  is the total memory bandwidth (note that here  $\Gamma$  is expressed in radians), where the width  $\Gamma$  is taken to be that of a square

distribution, as in Ref. [11]. This also guarantees that the spectral content of the input modes can be fitted inside the AFC, which in turn results in negligible temporal distortion of the output modes. The width  $\Gamma$  can be related to the number of AFC peaks  $N_p = \Gamma/\Delta$ . The total duration  $T$  of the input is  $T = 2\pi/\Delta - T_0$ , where  $T_0$  is the time necessary for applying the two control fields. In the limit where  $T_0 \ll 2\pi/\Delta$ , we have  $T \approx 2\pi/\Delta$ . The number of total temporal modes then amounts to  $N = T/\tau \approx N_p/6$ .

The number of peaks that can be created will depend strongly on the inhomogeneous  $\Gamma_{\text{in}}$  and homogeneous  $\gamma_h$  linewidths of the material. The width of each peak cannot be narrower than the homogeneous linewidth, i.e.,  $\gamma > \gamma_h$ . Since an efficient memory (>90%) requires a finesse  $F > 10$ , it follows that  $\Delta > 10\gamma_h$ . The number of peaks will depend on the ratio of the maximum AFC bandwidth  $\Gamma$  to the peak separation  $\Delta$ . The bandwidth will ultimately be limited by the inhomogeneous broadening of the material. In most materials, however, it will instead be limited by hyperfine or Zeeman transition spacings in order to avoid exciting multiple transitions that would induce quantum beat effects. In Sec. V we will give a detailed example where the number of peaks is of the order of hundreds with known material parameters.

The AFC QM dramatically reduces the necessary absorption depth for multimode storage as compared to CRIB or EIT QMs since the multimode capacity is independent of the peak absorption depth of the material. For CRIB,  $d = 30N$  is required for storing  $N$  modes with an average efficiency of 90% [11], so for instance  $N = 100$  requires  $d = 3000$ . In the case of EIT even larger  $d$  would be necessary due to worse scaling as mentioned in Sec. I. The AFC only requires  $d = 40$  to reach the same efficiency, independently of  $N$ . This makes efficient multimode QMs more realistic with current material properties, as we will discuss in the next section.

## V. IMPLEMENTATION USING RARE-EARTH-ION-DOPED SOLIDS

As storage material we here consider RE-doped crystals, which have inherently large inhomogeneous broadenings (0.1–10 GHz) and very narrow homogeneous linewidths (0.1–100 kHz) at low temperatures (typically less than 4 K) [31]. Note that the typical homogeneous linewidths correspond to optical coherence times ranging from  $\mu\text{s}$  to ms. Still RE ensembles have relatively high absorption coefficients at normal doping concentrations (10–100 ppm), typically in the range of 1–40  $\text{cm}^{-1}$ . RE crystals are therefore good candidates for the high-resolution spectral shaping we are discussing here. Reference [31] contains an extensive review of optical properties of different RE-doped crystals and glasses.

The creation of the periodic narrow structure is the crucial enabling step for the AFC QM. Narrow absorbing peaks (<50 kHz) have been isolated within the inhomogeneous broadening by optical pumping (or spectral hole burning) [32–35]. In all stopped-light [22,23] and CRIB experiments [30,34] in RE solids to this date, a narrow absorption peak within a larger transmission hole has been created in order to have an approximate homogeneous distribution of atoms.

Briefly, a large transmission hole is first created by spectral hole burning on the optical transition while sweeping the laser frequency. The pumped atoms are stored in another hyperfine level with long population lifetime. A narrow ensemble of atoms is then created in the hole by repumping atoms from the storage state to the initial state using a highly coherent laser source [36]. By repumping at different frequencies this method can be extended in order to create a series of peaks. One can also use other techniques in order to achieve frequency-selective optical pumping. For instance, one can use pairs of coherent pulses where each pair coherently transfers atoms to the excited state via a Ramsey interference [37]. Numerical simulations show that, by accumulating many pairs, narrow periodic structures can be created. This time-domain approach is interesting because in principle a time sequence can be created which has the appropriate frequency spectrum for creating an AFC with a certain finesse. Moreover a time-domain approach can be used for creating more complicated structures in frequency space, for instance multiple AFCs having different periodicity.

The spin coherence times have the potential of being significantly longer than the optical ones since it is free of decoherence due to spontaneous emission. In  $\text{Pr}^{3+}$ -doped  $\text{Y}_2\text{SiO}_5$  crystals, one of the most studied RE-doped material, the spin coherence time is 500  $\mu\text{s}$  at zero magnetic field [38], which can be extended to 82 ms by applying an orientation-specific magnetic field [39]. Fraval *et al.* [40] has also shown that it is possible to use dynamic decoupling methods developed in NMR spectroscopy to further extend the coherence time to 30 s. Yet another example is  $\text{Eu}^{3+}:\text{Y}_2\text{SiO}_5$  where 15 ms spin coherence time was measured at zero field [41], and  $\text{Tm}^{3+}$ : yttrium aluminum garnet (YAG) where 300  $\mu\text{s}$  was obtained in a magnetic field [42]. Efficient transfer of population between the optically excited state and ground hyperfine states has been demonstrated in  $\text{Pr}^{3+}:\text{Y}_2\text{SiO}_5$ , and manipulation of coherence between the optically excited state and the two spin ground states has recently been demonstrated in a state tomography experiment of an ensemble-based spin qubit in  $\text{Pr}^{3+}:\text{Y}_2\text{SiO}_5$  [43]. This shows the potential of spin states for long-term storage states for quantum information.

We finally give a more detailed example based on europium-doped  $\text{Y}_2\text{SiO}_5$  crystals to show the potential for efficient multimode storage using an AFC based memory. Europium ions absorb at 580 nm, and have an appropriate energy structure with three ground-state levels and an optically excited state [44], as well as excellent optical [45] and hyperfine [41] coherence times. Given the very narrow homogeneous linewidth in this material (122 Hz [45]), one could realistically create an AFC with  $\gamma = 2$  kHz and  $\Delta = 20$  kHz (i.e.,  $F = 10$ ). The total AFC bandwidth we take is 12 MHz, limited by hyperfine transition spacings, which results in  $N_p = 600$  peaks. The maximum number of modes is then roughly  $N = N_p/6 \approx 100$ . The absorption coefficient is about 3–4  $\text{cm}^{-1}$  [45] so that  $d = 40$  is within technical reach today using a multipass arrangement. Hence, one could then store 100 temporal modes in one single QM with an efficiency of 90%, which we verified by numerically simulating storage of 100 modes using the parameters above. We note that the efficiency stated above relates to the physics of the

atomic frequency comb (see the model assumptions in Sec. III). A complete storage experiment including transfer to the spin state will be affected by the limitation in spin coherence time, for  $\text{Eu}^{3+}:\text{Y}_2\text{SiO}_5$   $T_2^{\text{spin}}=36$  ms [41] has been observed so far, and by the transfer efficiency of the control fields (cf. discussion above).

We conclude this section by observing that less studied RE-doped crystals could potentially store more modes due to larger hyperfine state separations, for instance erbium-doped crystals absorbing at the telecommunication wavelength  $1.5 \mu\text{m}$  or neodymium-doped crystals absorbing at  $880$  nm.

## VI. CONCLUSIONS

We have presented a protocol for a quantum memory that can store efficiently a large number of temporal modes in a single atomic ensemble by creating an atomic frequency comb.

In materials with large optical inhomogeneous and narrow homogeneous broadenings, this allows for high multimode capacity at moderate optical absorption depths. We argue that RE-doped materials have the necessary properties for implementing the proposed protocol. The inherent multimode property of the atomic frequency comb quantum memory promises to speed up the entanglement generation rates in long-distance quantum networks by several orders of magnitude.

Recently, we have experimentally demonstrated a light-matter interface at the single-photon level [46] using an atomic frequency comb. This constitutes a proof-of-principle experiment of the idea presented in this paper.

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## APPENDIX: ANALYTICAL CALCULATIONS

### 1. Equations of motion

In this Appendix we present the analytical calculations leading to the quantitative formulas in the paper. The basic equations describing the dynamics of the light-atom system are the same as in Sec. II of Ref. [29]. Both the field and the atomic polarization are described by (slowly varying) forward and backward modes, denoted as  $E_f(z, t)$  and  $E_b(z, t)$  for the field, and  $\sigma_f(z, t; \delta)$  and  $\sigma_b(z, t; \delta)$  for the atomic polarization corresponding to atoms with resonance frequency  $\delta$ . They satisfy the following equations, cf. Eqs. (5a) and (5b), and Eqs. (6a) and (6b) of Ref. [29],

$$\frac{\partial}{\partial t} \sigma_f(z, t; \delta) = -i\delta \sigma_f(z, t; \delta) + i\varphi E_f(z, t), \quad (\text{A1})$$

$$\left( \frac{\partial}{\partial t} + c \frac{\partial}{\partial z} \right) E_f(z, t) = i\tilde{\varphi} \int_{-\infty}^{\infty} d\delta n(\delta) \sigma_f(z, t; \delta), \quad (\text{A2})$$

$$\frac{\partial}{\partial t} \sigma_b(z, t; \delta) = -i\delta \sigma_b(z, t; \delta) + i\varphi E_b(z, t), \quad (\text{A3})$$

$$\left( \frac{\partial}{\partial t} - c \frac{\partial}{\partial z} \right) E_b(z, t) = i\tilde{\varphi} \int_{-\infty}^{\infty} d\delta n(\delta) \sigma_b(z, t; \delta). \quad (\text{A4})$$

Here  $\varphi$  is the dipole moment of the atomic transition, and  $\tilde{\varphi} = g_0^2 \varphi$ , with  $g_0 = \sqrt{\omega_0 / (2\epsilon_0 V)}$ ,  $\omega_0$  is the optical frequency, and  $V$  is the quantization volume;  $n(\delta)$  is the atomic spectral distribution, with  $\int_{-\infty}^{\infty} d\delta n(\delta) = N_a$ , where  $N_a$  is the total number of atoms. See Ref. [29] for more details. Note that  $n(\delta) = N_a G(\delta)$  and  $\tilde{\varphi} = \beta / N_a$  in the notation of Ref. [29].

### 2. Atomic spectral distribution and its Fourier transform

Let us suppose that

$$n(\delta) \propto e^{-\delta^2 / (2\Gamma^2)} \sum_{n=-\infty}^{\infty} e^{-(\delta - n\Delta)^2 / (2\tilde{\gamma}^2)}, \quad (\text{A5})$$

i.e., the atomic spectral distribution is a frequency comb with overall width  $\Gamma$ , peak separation  $\Delta$ , and individual peak width  $\tilde{\gamma}$ . Note that in the paper we talk about the FWHM peak width  $\gamma = \sqrt{8 \ln 2} \tilde{\gamma}$ .

We are interested in the regime where  $\Gamma \gg \Delta \gg \tilde{\gamma}$ , i.e., we have a significant number of well-separated peaks within the overall characteristic width. We define the finesse  $F$  of the atomic distribution as the ratio of the peak separation over the FWHM of the individual peak, which gives  $F = \Delta / \gamma = \Delta / (\sqrt{8 \ln 2} \tilde{\gamma})$ .

In the following the Fourier transform  $\tilde{n}(t)$  of  $n(\delta)$  will play an important role. One finds

$$\tilde{n}(t) \propto e^{-t^2 \tilde{\gamma}^2 / 2} \sum_{n=-\infty}^{\infty} e^{-(t - n2\pi/\Delta)^2 \Gamma^2 / 2}, \quad (\text{A6})$$

which is also a series of peaks, with overall temporal width  $1/\tilde{\gamma}$ , peak separation  $2\pi/\Delta$ , and individual peak width  $1/\Gamma$ . Under the above assumptions, the peaks are again numerous (within the characteristic overall width) and well separated. To summarize, both  $n(\delta)$  and  $\tilde{n}(t)$  are pulse trains, with inversely proportional separations between the individual pulses, and the *global* shape of  $n(\delta)$  determines the *local* shape of  $\tilde{n}(t)$  and vice versa.

### 3. Absorption

The absorption process is described by forward Eqs. (A1) and (A2) coupling  $E_f(z, t)$  and  $\sigma_f(z, t; \delta)$ . Plugging the solution of Eq. (A1) into Eq. (A2) one has the following equation for the field

$$\begin{aligned} & \left( \frac{\partial}{\partial t} + c \frac{\partial}{\partial z} \right) E_f(z, t) \\ &= -\varphi \tilde{\varphi} \int_{-\infty}^{\infty} d\delta n(\delta) e^{-i\delta t} \times \int_{-\infty}^t dt' e^{i\delta t'} E_f(z, t'). \end{aligned} \quad (\text{A7})$$

This can be rewritten as

$$\left( \frac{\partial}{\partial t} + c \frac{\partial}{\partial z} \right) E_f(z, t) = -\varphi \tilde{\varphi} \int_{-\infty}^t dt' \tilde{n}(t-t') E_f(z, t'), \quad (\text{A8})$$

with

$$\tilde{n}(t) = \int_{-\infty}^{\infty} d\delta n(\delta) e^{-i\delta t}. \quad (\text{A9})$$

Let us consider an incoming light pulse of temporal length  $\tau$ . Let us note right away that we are here considering the regime  $\tau \gg L/c$ , where  $L$  is the length of the medium. This allows us to neglect temporal retardation effects, i.e., we can neglect the temporal derivative in Eqs. (A7) and (A8). The absorption process then has a duration that is essentially given by  $\tau$ . After this time the field is very close to zero until the coherent photon-echo type re-emission at  $t=2\pi/\Delta$ . If we are interested only in the absorption process, we only need to consider values of  $t-t'$  of order  $\tau$  in Eq. (A8). Assuming  $\tau \ll 2\pi/\Delta$ , i.e., a pulse duration much shorter than the waiting time for the re-emission, only the central peak of  $\tilde{n}$  from Eq. (A6) will contribute, i.e.,

$$\tilde{n}(t-t') \propto e^{-(t-t')^2 \Gamma^2/2}, \quad (\text{A10})$$

for  $t-t'$  of order  $\tau$ . Assuming that the overall width of the spectral distribution  $\Gamma$  is much larger than the spectral width of the incoming pulse, i.e.,  $\Gamma\tau \gg 1$ ,  $\tilde{n}(t-t')$  will act essentially like a delta function in the integral in Eq. (A8), giving an approximate equation of the form

$$\frac{\partial}{\partial z} E_f(z, t) = -\frac{\tilde{\alpha}}{2} E_f(z, t), \quad (\text{A11})$$

with a constant absorption coefficient  $\tilde{\alpha}$ . This is easily solved, giving

$$E_f(z, t) = E_f(0, t) e^{-(\tilde{\alpha}/2)z}. \quad (\text{A12})$$

Concerning the absolute strength of the absorption, one can show that, under the above assumptions  $\tilde{\alpha} = \alpha \frac{\sqrt{2\pi}\tilde{\gamma}}{\Delta}$ , where  $\alpha$  is the absorption coefficient corresponding to the (central) peaks of  $n(\delta)$ . This can be rewritten as

$$\tilde{\alpha} = \frac{\alpha}{F} \sqrt{\frac{\pi}{4 \ln 2}} = 1.064 \frac{\alpha}{F}. \quad (\text{A13})$$

Hence, for the absorption process, the AFC is indistinguishable from a smooth distribution that would be obtained by a spectral averaging that does not resolve the individual peaks. Note that in the paper we frequently discuss the absorption depth  $d$ , here defined as  $d = \alpha L$ . The effective absorption

depth of the AFC is then  $\tilde{d} = \tilde{\alpha} L$ , such that Eq. (A13) above corresponds to Eq. (4) in the paper.

#### 4. Re-emission

The pair of  $\pi$  pulses, which are used for storing the excitations in a long-lived ground state and then bringing them back to the excited state, also have the effect of changing the phase pattern of the atomic polarization from forward ( $\sigma_f$ ) to backward ( $\sigma_b$ ), such that it now couples to the backward field  $E_b$ , cf. Ref. [29]. We will see that this is essential for achieving potentially perfect readout efficiency, in analogy with the case of CRIB memories [29]. We assume that the time spent in the ground state does not lead to any relative phases between the different frequency components (no inhomogeneous broadening in the ground state). From the point of view of the equations of motion, one can thus treat the two  $\pi$  pulses as being applied essentially simultaneously. We will refer to the corresponding time as  $t=0$ . Absorption thus occurred at negative times, and re-emission will occur at positive times. The backward polarization created at  $t=0$  is

$$\sigma_b(z, 0; \delta) = \sigma_f(z, 0; \delta) = \int_{-\infty}^0 dt' e^{i\delta t'} E_f(z, t'), \quad (\text{A14})$$

where  $\sigma_b$  ( $\sigma_f$ ) is to be understood after (before) the two  $\pi$  pulses. Equation (A14) gives the initial condition for Eqs. (3) and (4). We assume that the field  $E_b$  is zero at  $t=0$  (the absorption process is completed; if there was a transmitted field it has left the medium). This gives the following equation for the backward field

$$\begin{aligned} \left( \frac{\partial}{\partial t} - c \frac{\partial}{\partial z} \right) E_b(z, t) &= -\varphi \tilde{\varphi} \int_{-\infty}^0 dt' \tilde{n}(t-t') E_f(z, t') \\ &\quad - \varphi \tilde{\varphi} \int_0^t dt' \tilde{n}(t-t') E_b(z, t'). \end{aligned} \quad (\text{A15})$$

The first term on the right-hand side is the source term which gives rise to the coherent re-emission; the second term describes the absorption of the backward field in analogy with the case treated in the previous section, see Eq. (A8). The re-emission occurs because  $\tilde{n}(t-t')$  has a peak not only for  $t-t'$  around zero, but also for  $t-t'$  around  $2\pi/\Delta$ . Note that the height of this peak is reduced by a factor  $e^{-(2\pi/\Delta)^2 \tilde{\gamma}^2/2}$  with respect to the central peak, cf. Eq. (A6). Around the corresponding point in time, one can apply arguments fully analogous to the ones in the previous subsection. Equation (A15) then simplifies to

$$\frac{\partial}{\partial z} E_b(z, t) = \tilde{\alpha} e^{-\tilde{\gamma}^2 (2\pi/\Delta)^2/2} E_f \left( z, t - \frac{2\pi}{\Delta} \right) + \frac{\tilde{\alpha}}{2} E_b(z, t). \quad (\text{A16})$$

The  $E_f$  term has a factor  $\tilde{\alpha}$  instead of  $\frac{\tilde{\alpha}}{2}$  because the  $t'$  integral in Eq. (A15) is over all of the effective delta function (given by the corresponding peak of  $\tilde{n}$ ) for the  $E_f$  (source) term, whereas it is only over one half of the delta function



for the  $E_b$  (absorption) term, as it was in the previous subsection for the absorption of  $E_f$ .

The solution of Eq. (A16) is

$$E_b(z, t) = e^{-\tilde{\gamma}^2(2\pi/\Delta)^2/2} \int_L^z dz' \tilde{\alpha} e^{\tilde{\alpha}/2(z-z')} E_f\left(z', t - \frac{2\pi}{\Delta}\right). \quad (\text{A17})$$

Note that  $E_b$  propagates backward so the output field is at  $z=0$ . Using Eq. (A12) one has

$$\begin{aligned} E_b(0, t) &= -E_f\left(0, t - \frac{2\pi}{\Delta}\right) e^{-\tilde{\gamma}^2(2\pi/\Delta)^2/2} \int_0^L dz' \tilde{\alpha} e^{-\tilde{\alpha}z'/2} e^{-\tilde{\alpha}z'/2} \\ &= -E_f\left(0, t - \frac{2\pi}{\Delta}\right) e^{-\tilde{\gamma}^2(2\pi/\Delta)^2/2} (1 - e^{-\tilde{\alpha}L}). \end{aligned} \quad (\text{A18})$$

It is interesting to note that the integral over  $z'$  has the same form as in the case of the (backward) CRIB memory [29]. It corresponds to the coherent sum over all possible paths of the photon in the medium, where each path is characterized by its turning point  $z'$ . Even though there is absorption along the path, there is a constructive interference effect which leads to the output amplitude potentially being arbitrarily close in magnitude to the input amplitude (for large enough  $\tilde{\alpha}L$ ).

This interference is not present if the output field is emitted in the forward direction, which would be the case if the two control fields were not applied. Then one would have an integral

$$\int_0^L dz' \tilde{\alpha} e^{-\tilde{\alpha}z'/2} e^{-\tilde{\alpha}(L-z')/2} = \tilde{\alpha}L e^{-\tilde{\alpha}L/2}, \quad (\text{A19})$$

replacing the last factor in Eq. (A18). If one then calculates the efficiency, one finds that it can at most attain 54% for  $\tilde{\alpha}L=2$ , cf. [29].

For the backward propagating mode the achievable memory efficiency is given by, from Eq. (A18),

$$\eta = (1 - e^{-\tilde{\alpha}L})^2 e^{-\tilde{\gamma}^2(2\pi/\Delta)^2}, \quad (\text{A20})$$

which can be rewritten in terms of the finesse  $F$  and peak absorption  $\alpha$  of the atomic distribution  $n(\delta)$  as [cf. Eq. (5) in paper]

$$\begin{aligned} \eta &= (1 - e^{-(\alpha L/F)(\sqrt{\pi/4 \ln 2})})^2 e^{-(1/F^2)(\pi^2/2 \ln 2)} \\ &\approx (1 - e^{-(\alpha L/F)})^2 e^{-(7/F^2)}. \end{aligned} \quad (\text{A21})$$

There is a tradeoff between absorption (favoring small  $F$ ) and dephasing (favoring large  $F$ ); however, for large enough  $\alpha L$  and optimized  $F$  the efficiency can be made arbitrarily close to 100% (cf. Figs. 2 and 3 in paper).

### 5. Phase due to frequency offset

Here we will comment on the phase of the output field as a function of the relative frequency detuning between the carrier frequency and the frequency of the peak in the AFC (any of them), cf. Eq. (5) in the paper. If the atomic distribution  $n(\delta)$  is replaced by  $n(\delta - \Delta_0)$ , corresponding to a small frequency offset, then the Fourier transform  $\tilde{n}(t)$  acquires a phase factor  $e^{i\Delta_0 t}$ , which leads to a supplementary phase factor  $e^{i2\pi\Delta_0/\Delta}$  for the re-emitted field, cf. Eq. (A18). Note that this phase is modulus the peak separation  $\Delta$ .

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- [1] H.-J. Briegel, W. Dür, J. I. Cirac, and P. Zoller, *Phys. Rev. Lett.* **81**, 5932 (1998).
- [2] L.-M. Duan, M. D. Lukin, J. I. Cirac, and P. Zoller, *Nature (London)* **414**, 413 (2001).
- [3] C. W. Chou, J. Laurat, H. Deng, K. S. Choi, H. de Riedmatten, D. Felinto, and H. J. Kimble, *Science* **316**, 1316 (2007).
- [4] Z.-S. Yuan, Y.-A. Chen, B. Zhao, S. Chen, J. Schmiedmayer, and J.-W. Pan, *Nature (London)* **454**, 1098 (2008).
- [5] B. Julsgaard, J. Sherson, J. I. Cirac, J. Fiurášek, and E. S. Polzik, *Nature (London)* **432**, 482 (2004).
- [6] T. Chanelière, D. N. Matsukevich, S. D. Jenkins, S.-Y. Lan, T. A. B. Kennedy, and A. Kuzmich, *Nature (London)* **438**, 833 (2005).
- [7] M. D. Eisaman, A. André, F. Massou, M. Fleischhauer, A. S. Zibrov, and M. D. Lukin, *Nature (London)* **438**, 837 (2005).
- [8] K. S. Choi, H. Deng, J. Laurat, and H. J. Kimble, *Nature (London)* **452**, 67 (2008).
- [9] N. Sangouard, C. Simon, B. Zhao, Y.-A. Chen, H. de Riedmatten, J.-W. Pan, and N. Gisin, *Phys. Rev. A* **77**, 062301 (2008).
- [10] O. A. Collins, S. D. Jenkins, A. Kuzmich, and T. A. B. Kennedy, *Phys. Rev. Lett.* **98**, 060502 (2007).
- [11] C. Simon, H. de Riedmatten, M. Afzelius, N. Sangouard, H. Zbinden, and N. Gisin, *Phys. Rev. Lett.* **98**, 190503 (2007).
- [12] K. Surmacz, J. Nunn, K. Reim, K. C. Lee, V. O. Lorenz, B. Sussman, I. A. Walmsley, and D. Jaksch, *Phys. Rev. A* **78**, 033806 (2008).
- [13] D. V. Vasylyev, I. V. Sokolov, and E. S. Polzik, *Phys. Rev. A* **77**, 020302(R) (2008).
- [14] M. Fleischhauer and M. D. Lukin, *Phys. Rev. Lett.* **84**, 5094 (2000).
- [15] S. A. Moiseev and S. Kröll, *Phys. Rev. Lett.* **87**, 173601 (2001).
- [16] B. Kraus, W. Tittel, N. Gisin, M. Nilsson, S. Kröll, and J. I. Cirac, *Phys. Rev. A* **73**, 020302(R) (2006).
- [17] J. Nunn, I. A. Walmsley, M. G. Raymer, K. Surmacz, F. C. Waldermann, Z. Wang, and D. Jaksch, *Phys. Rev. A* **75**, 011401(R) (2007).
- [18] A. V. Gorshkov, A. André, M. D. Lukin, and A. S. Sørensen, *Phys. Rev. A* **76**, 033806 (2007).
- [19] A. V. Gorshkov, T. Calarco, M. D. Lukin, and A. S. Sørensen, *Phys. Rev. A* **77**, 043806 (2008).
- [20] M. D. Lukin, *Rev. Mod. Phys.* **75**, 457 (2003).
- [21] 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).
- [22] A. V. Turukhin, V. S. Sudarshanam, M. S. Shahriar, J. A.

- Musser, B. S. Ham, and P. R. Hemmer, *Phys. Rev. Lett.* **88**, 023602 (2001).
- [23] J. J. Longdell, E. Fraval, M. J. Sellars, and N. B. Manson, *Phys. Rev. Lett.* **95**, 063601 (2005).
- [24] R. H. Dicke, *Phys. Rev.* **93**, 99 (1954).
- [25] T. Mossberg, A. Flusberg, R. Kachru, and S. R. Hartmann, *Phys. Rev. Lett.* **42**, 1665 (1979).
- [26] N. W. Carlson, Y. S. Bai, W. R. Babbitt, and T. W. Mossberg, *Phys. Rev. A* **30**, 1572 (1984).
- [27] M. Mitsunaga, R. Yano, and N. Uesugi, *Opt. Lett.* **16**, 264 (1991).
- [28] For the rare-earth-doped materials considered here the optical and spin inhomogeneous broadenings are generally uncorrelated. If the physical system would contain correlations between these broadenings, however, there could be a comb structure also on the spin transition.
- [29] N. Sangouard, C. Simon, M. Afzelius, and N. Gisin, *Phys. Rev. A* **75**, 032327 (2007).
- [30] A. L. Alexander, J. J. Longdell, M. J. Sellars, and N. B. Manson, *Phys. Rev. Lett.* **96**, 043602 (2006).
- [31] R. M. Macfarlane, *J. Lumin.* **100**, 1 (2002).
- [32] F. de Sèze, V. Lavielle, I. Lorgeré, and J.-L. Le Gouët, *Opt. Commun.* **223**, 321 (2003).
- [33] M. Nilsson, L. Rippe, S. Kröll, R. Klieber, and D. Suter, *Phys. Rev. B* **70**, 214116 (2004).
- [34] G. Hétet, J. J. Longdell, A. L. Alexander, P. K. Lam, and M. J. Sellars, *Phys. Rev. Lett.* **100**, 023601 (2008).
- [35] B. Lauritzen, S. R. Hastings-Simon, H. de Riedmatten, M. Afzelius, and N. Gisin, *Phys. Rev. A* **78**, 043402 (2008).
- [36] L. Rippe, M. Nilsson, S. Kröll, R. Klieber, and D. Suter, *Phys. Rev. A* **71**, 062328 (2005).
- [37] W. H. Hesselink and D. A. Wiersma, *Phys. Rev. Lett.* **43**, 1991 (1979).
- [38] B. S. Ham, M. S. Shahriar, M. K. Kim, and P. R. Hemmer, *Phys. Rev. B* **58**, R11825 (1998).
- [39] E. Fraval, M. J. Sellars, and J. J. Longdell, *Phys. Rev. Lett.* **92**, 077601 (2004).
- [40] E. Fraval, M. J. Sellars, and J. J. Longdell, *Phys. Rev. Lett.* **95**, 030506 (2005).
- [41] A. L. Alexander, J. J. Longdell, and M. J. Sellars, *J. Opt. Soc. Am. B* **24**, 2479 (2007).
- [42] A. Louchet, Y. Le Du, F. Bretenaker, T. Chanelière, F. Goldfarb, I. Lorgeré, J.-L. Le Gouët, O. Guillot-Noël, and Ph. Goldner, *Phys. Rev. B* **77**, 195110 (2008).
- [43] L. Rippe, B. Julsgaard, A. Walther, Yan Ying, and S. Kröll, *Phys. Rev. A* **77**, 022307 (2008).
- [44] M. Nilsson, L. Rippe, N. Ohlsson, T. Christiansson, and S. Kröll, *Phys. Scr.*, T **102**, 178 (2002).
- [45] F. Könz, Y. Sun, C. W. Thiel, R. L. Cone, R. W. Equall, R. L. Hutcherson, and R. M. Macfarlane, *Phys. Rev. B* **68**, 085109 (2003).
- [46] H. de Riedmatten, M. Afzelius, M. U. Staudt, C. Simon, and N. Gisin, *Nature (London)* **456**, 773 (2008).