Photon-pair source with controllable delay based on shaped inhomogeneous broadening of rare-earth-metal-doped solids

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I. INTRODUCTION

The creation of correlated Stokes–anti-Stokes photon pairs in atomic ensembles via spontaneous Raman emission [1] plays a central role in quantum communication, starting with the implementation of quantum repeaters [2]. The basic principle requires an ensemble of lambda systems coupled to a pair of optical laser fields, see Fig. 1. An off-resonant laser pulse, the write pulse, produces a frequency-shifted Stokes photon via spontaneous Raman emission. The detection of this Stokes photon in the far field, such that no information is revealed about which atom it came from, heralds the creation of a single collective atomic spin excitation. A remarkable feature of such a collective atomic state is that it can be read out very efficiently. A resonant laser pulse, the read pulse, allows one, through a collective spontaneous Raman emission, to ideally map the collective spin excitation into an anti-Stokes photon propagating in a well-defined spatiotemporal mode. This provides a photon-pair source with a very special property: the delay between the Stokes and anti-Stokes photons can be controlled by choosing the timing between the write and read pulses. Such a source has inspired many experiments in atomic gases, including the first single-photon storage in an atomic ensemble [3,4], the first heralded creation of entanglement between atomic ensembles [5], and even the implementation of the first elementary blocks of quantum repeaters [6]. Despite this impressive body of work, there are strong motivations to use more practical systems in the solid state. Rare-earth-metal-doped solids seem naturally well suited, at least at first sight. They are widely available thanks to their use for solid-state lasers. Thanks to their particular electronic structure, they can be seen as a frozen gas of atoms, with optical and spin transitions featuring excellent coherence properties. Moreover, they have already shown excellent capability to store light for long times [7] with high efficiency [8] and negligible noise [8,9]. Last but not least, they have a large inhomogeneous absorption spectrum and narrow homogeneous lines leading to a high temporal multimode capacity [10].

All rare-earth-metal elements have in common weak dipole moments on the relevant $4f-4f$ transitions and large inhomogeneously broadened spectra due to the interaction with the host crystal. These two properties make the creation of Stokes–anti-Stokes pairs in rare-earth-metal-doped solids challenging. In hot alkali-metal gases, where there is also an inhomogeneous broadening due to the Doppler effect, spontaneous Raman processes have been successfully performed [4] with a write pulse far detuned from the resonance; the probability $p$ for the emission of a Stokes photon in a given mode being enhanced by the use of large write intensities. However, in rare-earth-metal-doped solids, where the dipole moments are typically two to three orders of magnitude weaker, the required intensities would be, at best, difficult to achieve. For resonant write pulses, atoms are transferred into the excited state, leading to a higher $p$. But their energy difference, due to the inhomogeneous broadening, makes them distinguishable. They can no longer interfere, which makes the readout of the collective excitation inefficient [11].

We propose a simple solution to this problem. It consists in shaping the spectral inhomogeneous broadening of the optical transition so that the atomic dipoles rephase at the readout step, leading to an efficient emission of the anti-Stokes photon even when resonant write pulses are used. Several shaping methods are available to force the atomic dipoles to rephase. For example, the inhomogeneous broadening could be shaped into a narrow absorption line with a reversible and controllable broadening [12]. In what follows, we focus on a shaping based on a comblike structure composed with periodic narrow peaks [13]. This provides a temporally multiplexed version of a spontaneous Raman source, similarly to the proposal of Ref. [14] in atomic gases but without the need for a cavity. Since spontaneous Raman-based protocols are well suited for entangling remote atomic ensembles in a heralded way [1] without the need for ultranarrowband pair sources, our proposal paves the way for the implementation of the first elementary link of quantum repeaters with solid-state devices.

II. WRITE STEP

Let us start by a description of the write step. We consider a medium consisting of lambda atoms (as depicted in Fig. 1) initially prepared in the state $g$. The optical transition $g-e$ is shaped into a frequency comb made of narrow peaks with a characteristic width $\gamma$, separated by $\Delta_0$, and spanning a large atomic frequency range $\Gamma$. The $g-s$ transition is considered to be homogeneous. A weak write laser pulse with the Rabi frequency $\Omega_w(t)$ is sent through the medium to transfer a small...
drives the atoms into photons. Taking the absorption into account, the write pulse maximum (FWHM) distribution made of Gaussian peaks with full width at half comb (cf. below). For concreteness, we consider an atomic length of the atomic medium. It is proportional to the ratio of the echo-photon-type re-emission (at time $2\pi/\Delta_1$) and benefiting from spontaneous emission of a Stokes photon. The detection is the total number of atoms. $\bar{N}$

$$\langle e| \hat{e}_s(z,t) = \sqrt{\frac{L}{2\pi \epsilon c}} \int d\omega \hat{a}_w(t) e^{i\omega t}.$$  

Under the dipole and rotating wave approximation, the interaction between the Stokes field and the medium is governed by the Hamiltonian

$$H_{ int} = -\hbar g \sqrt{\frac{L}{2\pi \epsilon c}} \sum \int d\omega \hat{a}_w e^{i\omega t} |e\rangle \langle s| + H.c.,$$

where $g = \sqrt{\frac{\epsilon}{2\hbar \Delta_0 \Delta'}}$ is the atom-field coupling constant with $\Delta_0$ the dipole moment of the $e-s$ transition and $A$ the interaction section. The equations of motion for the Stokes field and for the atomic coherence $\sigma_{se}'(t) = \langle s| \hat{e}_s(t) \hat{e}_s^\dagger(t) s\rangle$ are given by

$$\frac{d}{dt} \sigma_{se}'(t) = -i \Delta' \sigma_{se}^j - ig \hat{e}_{st}(z,t) \sigma_{se}',$$

where $\Delta' = \omega_{se} - \omega_s$ is the frequency detuning and $\sigma_{se}^j = |e\rangle \langle e| - |s\rangle \langle s|$ is approximated by its mean value $\bar{N}_{se} \epsilon e^{-\alpha z}$ in what follows. Plugging the formal solution of Eq. (A3) into Eq. (A2), we obtain the expression of the Stokes field at $z$ (see Appendix)

$$\hat{e}_{st}(z,t) = e^{i \frac{g L}{\epsilon} \sum_{j,z,t} e^{-\alpha z} \hat{a}_s \hat{a}_t} \hat{e}_{st}(0,t) + i \frac{g L}{\epsilon} \sum_{j,z,t} e^{i \frac{g L}{\epsilon} \sum_{j,z,t} e^{-\alpha z} \hat{a}_s \hat{a}_t} e^{-i \Delta' \sigma_{se}'(0)}.$$  

We consider, for simplicity, that the transitions $g-e$ and $s-e$ have the same dipole moments. Since the state of the complete system after the write pulse is given by $\langle \Psi_w | = |\Psi_w,0\rangle$ where $|0\rangle$ is the vacuum for the electromagnetic field, the average number of Stokes photons emitted at time $t_d$ in a mode of temporal duration $\sqrt{2\pi} / \Gamma$ is given by

$$\frac{\sqrt{2\pi} \epsilon}{\Gamma L} \langle \Psi_w | \hat{e}_{st}(L,t_d) \hat{e}_{st}(L,t_d) |\Psi_w\rangle \approx \theta_{\alpha}^2 (1 - e^{-\alpha L}),$$

for $\theta_\alpha \ll 1$ (see Appendix). This formula is very useful and can easily be used in practice. Note first that the relative number of atoms transferred into the excited state by the write pulse is given by

$$\frac{1}{N} \int_0^L \rho_0^2 e^{-\alpha z} \frac{dx}{L} = \frac{\theta_\alpha^2}{\alpha} (1 - e^{-\alpha L}).$$

Therefore, the formula (6) tells us that the average number of photons in a mode with a duration corresponding to the inverse of the overall spectrum is merely the optical depth $\alpha L$ times the relative number of atoms in the state $e$. In the rest of the article we will be interested in the regime where the optical depth $\alpha L$ is large enough to get high readout efficiencies, but the write pulse is weak enough to get a high signal-to-noise ratio for the photon-echo-like re-emission for the readout of the atomic spin wave.

III. SPONTANEOUS EMISSION OF STOKES PHOTONS

Let us consider the Stokes field which propagates in the forward direction with the carrier frequency $\omega_s$. Its envelope is described by the slowly varying quantum operator [15]

$$\hat{e}_{st}(z,t) = \sqrt{\frac{L}{2\pi \epsilon c}} e^{i\omega t} \hat{a}_w(t) e^{i\omega t}.$$  

FIG. 1. (a) Basic level scheme and (b) pulse sequence for the creation of correlated Stokes–anti-Stokes pairs via spontaneous Raman emission. All atoms start in $g$. The inhomogeneously broadened optical transition is shaped into an atomic frequency comb. A resonant write pulse $\Omega_w$ excites the $g-e$ transition, making possible the spontaneous emission of a Stokes photon. The detection of a Stokes photon (at time $t_d$) is uniquely correlated with the creation of a single collective atomic excitation corresponding to one excitation in $S$ delocalized over all the atoms. This collective excitation can be mapped very efficiently into an anti-Stokes photon using a resonant read pulse $\Omega_r$ (at time $t + t_d$) and benefiting from the echo-photon-type re-emission (at time $2\pi/\Delta + \tau$) inherent to inhomogeneous transition shaped into a frequency comb.

part of the atoms in the excited state $e$ in such a way that it can spontaneously decay into the level $S$ by emitting Stokes photons. Taking the absorption into account, the write pulse drives the atoms into

$$|\Psi_w\rangle = \prod_j \left( \frac{1}{\sqrt{n_j}} \left( -\frac{1}{2} + \frac{\theta_{\alpha}^2 e^{-\alpha z_j}}{\sqrt{n_j}} \right) |e\rangle + e^{i\omega_0 z_j} \frac{\theta_{\alpha}^2 e^{-\alpha z_j}}{\sqrt{n_j}} |e\rangle \right).$$

Here $n_j$ ensures the normalization, $\theta_{\alpha} = \frac{1}{2} \int ds \Omega_w(s) \ll 1$ refers to the area of the write pulse at the entrance of the crystal, and $\omega_0$ corresponds to its wave number. $z_j$ is the position of the $j$th atom within a medium of length $L$ and $N$ is the total number of atoms. $\alpha$ is the absorption per unit length of the atomic medium. It is proportional to the ratio between the absorption per peak $\alpha$ and the finesse $F$ of the comb (cf. below). For concreteness, we consider an atomic distribution made of Gaussian peaks with full width at half maximum (FWHM) $\gamma$ so that $\alpha = \frac{\pi \epsilon}{\sqrt{2\pi} F}$ where $F = \Delta_0/\gamma$ (see Appendix). Note that this shaping, which is at the heart of memories based on the atomic frequency comb [13], is known to produce a photon echo-type of re-emission in a well-defined spatial mode at time $2\pi/\Delta_0$. Here, on the other hand, we look at the spontaneous emission of a Stokes photon at time $t_d < 2\pi/\Delta_0$ and we benefit from the efficiency of the

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readout (cf. below). In this case, the success probability for the emission of a Stokes photon (6) reduces to $p \approx \theta_0^2$.

## IV. READOUT EFFICIENCY

We now calculate the efficiency of the readout process. At time $\tau$ after the detection of the Stokes photon, a read pulse resonant with the transition s-e and associated with the Rabi frequency $\int d s \dot{Q}_s(s) = \pi$ goes through the atomic ensemble and exchanges the population of states $s$ and $e$. For $\theta_0 \ll 1$, the resulting atomic state is given by (see Appendix)

$$\ket{\psi_r} = \sum_{j \geq 1} e^{iz_j/(a_{0j}-\theta_0/\omega_c)} e^{-2a_{0j}/2} e^{-i\pi/\Delta} e^{-i\Delta/\Delta} \ket{e_j}$$

$$\times \prod_{l \neq j} (G_l e^{-i\omega_{0l}(t+\tau)} \ket{g_l} + E_l e^{i\omega_{0l}(t+\tau)} \ket{e_l}),$$

$$\text{(7)}$$

with $\zeta = (\alpha L)/\sqrt{N(1-e^{-2\alpha L})}$. Consider a re-emission associated with an anti-Stokes mode $\bar{\mathcal{E}}_{\text{AS}},(t)$ propagating in the backward direction. Following the method presented before (under the assumption that $|e\rangle\langle e| - |g\rangle\langle g|$ $\approx -1$) we find at $z = 0$

$$\bar{\mathcal{E}}_{\text{AS}}(0,t) = e^{-2zL} \mathcal{E}_{\text{AS}}(L,t)$$

$$+ i \frac{gL}{c} \sum_j e^{i\omega_j \theta z/(1-\theta_0)^2} \mathcal{E}_{\text{AS}}(L,t)$$

$$\text{(8)}$$

so that, at time $2\pi/\Delta + \tau$ where all the atoms are in phase, the efficiency of the readout process is

$$\sqrt{2\pi c \mathcal{T}} \mathcal{E}_{\text{AS}}(0,2\pi/\Delta + \tau)\mathcal{E}_{\text{AS}}(0,2\pi/\Delta + \tau) \ket{\psi_r}$$

$$= \left(1 - e^{-\frac{\pi c}{\Delta}} \right) e^{-\frac{\pi c}{\Delta}}.$$

$$\text{(9)}$$

One sees that there is a tradeoff between absorption and dephasing. However, for large enough $\alpha L$ and optimized $F$, the readout efficiency can be arbitrarily close to 100%. Let us directly note some of the advantages of spontaneous Raman protocols over other schemes. First, the proposal based on spontaneous Raman protocols is significantly more efficient than a memory where the photon has first to be absorbed before being reemitted. It leads to a much higher efficiency for small optical depths (see Appendix). Furthermore, since the readout of spontaneous Raman protocols is conditioned on the detection of a Stokes photon, the retrieved signal is not affected by the nonunit coupling efficiency of an input photon into the memory (e.g., due to imperfect spectral filters, nonunit coupling into monomode fibers, and so on, see Appendix). Contrary to the photon-pair source based on rephased amplified spontaneous emission [16], our proposal provides highly correlated pairs even for large optical depths where the retrieval efficiency is high.

## V. NOISE RATE

We now account for intrinsic noise. The conditional state (7) that we used to calculate the efficiency of the readout process corresponds to the ideal case where a single Stokes photon has been emitted and detected. However, many atoms prepared in the excited state by the write pulse can emit Stokes photons in all the spatiotemporal modes. This unwanted emission populates the state $s$ so that after the interaction with the read pulse, atoms occupy the excited state $e$ and can produce spontaneous noise in the anti-Stokes mode. To take this noise into account, we consider the worst case where all the atoms prepared in the excited state by the write pulse decay on the $e$-$s$ transition. Tracing over the Stokes photons, the atomic state after the interaction with the read pulse is well approximated by

$$\bar{\mathcal{E}}_{\text{AS}}(0,2\pi/\Delta + \tau)\mathcal{E}_{\text{AS}}(0,2\pi/\Delta + \tau) \rho_\gamma$$

$$\approx \frac{\theta_0^2}{2} (1 - e^{-2\alpha L}),$$

where $\rho_\gamma = \mathcal{E}_N \otimes |0\rangle\langle 0|$. For large enough $\alpha L$ and optimized $F$, the signal-to-noise ratio $\mathcal{R}$ is lower bounded by

$$\mathcal{R} \geq \frac{2 \theta_0^2}{\rho_\gamma} = \frac{2}{p}.$$

One can thus conclude that despite the inhomogeneous broadening inherent to solid-state systems, our protocol achieves similar characteristics to the ones obtained in cold atomic gases; $\mathcal{R}$ can be very high provided that the number of Stokes photons per mode is low.

## VI. FEASIBILITY

For concreteness, we now discuss the experimental feasibility of spontaneous Raman processes in rare-earth-metal-doped materials. Pr:Y$_2$SiO$_5$ is a very promising material for initial experiments since excellent hyperfine coherence [17] and quantum memory efficiencies of order 70% [8] have already been demonstrated. The frequency difference between the Stokes and the anti-Stokes transitions is only of some hundreds of MHz, so no problems in relation with group velocity dispersion arise. The main drawback of praseodymium is the small hyperfine separation (a few MHz) which limits the number of peaks within the atomic comb and thus the multimode capacity (cf. below). If we consider the $3_{\text{H}} 6_{\text{H}} = 1 = D_2$ transition at 606 nm, one can realistically shape combs with individual peaks of FWHM $\nu \approx 30$ kHz [18] over a spectral range of $\Gamma \approx 2$ MHz and an absorption per peak of order $\alpha L \approx 10$ [19] (the branching ratio is approximated by 1). For $F = 5, p \approx 0.1$ can be achieved provided that the write pulse satisfies $\theta_0^2 = 0.1$ at the entrance of the crystal. This would lead to a readout efficiency of 65% and a signal-to-noise ratio larger than 10 under the assumption that the noise is dominated by the spontaneous emission. Now consider a setup involving two Pr-doped solids located 1 km apart so that the corresponding Stokes modes are combined on a beamsplitter at a central station. Further consider a fiber attenuation of 9 dB/km, corresponding to 606-nm photons [20] and assume a coupling efficiency into optical fibers of $\eta_c = 50\%$. The average time
to detect a Stokes photon after the beamsplitter and thus to entangle the two crystals is \( T = \frac{1}{2\pi \eta_{p} \eta_{d}} \) where \( \eta_{p}, \eta_{d} \) are the transmission and detection efficiencies and \( r \) is the repetition rate. Assume \( \eta_{d} = 70\% \). For \( p = 0.05 \) where the fidelity of the entanglement is \( F = 1 - 3p(1 - \eta_{p}\eta_{d}) \approx 0.85 \) [2], one finds \( T \approx 0.1 \) s for a repetition rate of \( r = 1 \) kHz so that a few hours would be sufficient for performing a full tomography. Note that the fiber lengths have to be actively stabilized to guarantee an interferometric phase stability on this time scale [2].

VII. CONCLUSION

Our approach opens an avenue toward the heralded entanglement of remote solids. Beyond that, the present scheme might be useful for quantum communications since spontaneous Raman processes lead to the production of narrowband pairs well suited for storage in atomic ensembles. Let us finally emphasize that in our protocol, spin waves created at different times, say \( t_{1}, t_{2}, t_{3}, \ldots \), are independent and if the read pulse is sent at time \( t \) after the first detection, these spin waves rephase at times \( t + 2\tau / \Delta, t + 2\pi / \Delta - (t_{2} - t_{1}), t + 2\pi / \Delta - (t_{3} - t_{1}), \ldots \). The number of spin waves that can be stored is roughly given by the number of peaks composing the comb (see Ref. [13] and the Appendix) and can be merely increased by making use of a wider range of the inhomogeneous broadening. In the framework of quantum repeaters, this temporal multiplexing has been shown to greatly enhance the distribution rate of entanglement [21].

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APPENDIX

1. Atomic distribution

For concreteness, we consider an atomic spectral distribution shaped into a frequency comb made with Gaussian peaks

\[
\Theta(\Delta) := \frac{\Delta_{0}}{2\pi \tilde{\gamma} \Gamma} e^{-\frac{\Delta^{2}}{2 \Delta_{0}^{2}}} \sum_{j=-\infty}^{\infty} e^{-\frac{(\Delta - j\Delta_{0})^{2}}{2 \Delta_{0}^{2}}},
\]

where \( \Gamma \) denotes the overall width, \( \Delta_{0} \) is the peak separation, and \( \tilde{\gamma} \) is the width of an individual peak.

We are interested in the regime \( \Gamma \gg \Delta_{0} \gg \tilde{\gamma} \) where a large number of well-separated peaks spans a large spectral bandwidth. In this regime, one can check that \( \int d\Delta \Theta(\Delta) = 1 \).

We define the finesse of the comb as the ratio of the peak separation over the FWHM of a single peak \( \gamma \), that is,

\[
F := \frac{\Delta_{0}}{\gamma} = \frac{1}{\sqrt{8\ln 2}} \frac{\Delta_{0}}{\tilde{\gamma}}.
\]

Note that the Fourier transform of the atomic spectral distribution

\[
\tilde{\Theta}(\tau) := \int d\Delta \Theta(\Delta)e^{-i\Delta \tau},
\]
is also a series of Gaussian peaks with individual peak width \( 1/\Gamma \), separated by \( 2\pi/\Delta_{0} \), and spanning the overall temporal width \( 1/\tilde{\gamma} \).

2. Emission of Stokes photons

We now detail the way to find the solution of equations associated to the dynamics of the Stokes field

\[
\frac{d}{dt} \tilde{\mathcal{E}}_{s}(z,t) = igL \sum_{j} \delta(z - z_{j})\sigma_{j}^{e}, \quad (A2)
\]

\[
\frac{d}{dt} \sigma_{j}^{e} = -i\Delta_{j} \sigma_{j}^{e} - ig\tilde{\mathcal{E}}_{s}(z,t)\sigma_{j}^{s}.
\]

First, we plug the formal solution of Eq. (A3)

\[
\sigma_{j}^{e}(t) = -ig \sigma_{j}^{s}(0) \int_{0}^{t} dt' e^{-i\Delta_{j}(t-t')} \tilde{\mathcal{E}}_{s}(z,j,t') + e^{-i\Delta_{j}t} \sigma_{j}^{s}(0),
\]

into Eq. (A2) and since we consider Stokes modes with a characteristic duration \( \tau_{s} / c \), we can neglect the temporal retardation effects in the crystal [i.e., the time derivative in Eq. (A2)]. This leads to

\[
\frac{d}{dz} \tilde{\mathcal{E}}_{s}(z,t) = igL \sum_{j} \delta(z - z_{j})e^{-i\Delta_{j}t} \sigma_{j}^{s}(0)
\]

\[
+ g^{2}L \sum_{j} \delta(z - z_{j}) \sigma_{j}^{s}(0) \int_{0}^{t} dt' e^{-i\Delta_{j}(t-t')} \tilde{\mathcal{E}}_{s}(z,j,t').
\]

We then replace \( \sigma_{j}^{s}(0) \) by its mean value calculated on the state \( |\psi_{e} \rangle \) (i.e., \( \sigma_{j}^{s}(0) \approx 0_{\Theta}^{2} e^{-\Theta^{2}/2} \)) and we take the continuous limit \( \sum_{j} \rightarrow \int_{0}^{L} \frac{d\zeta}{\tilde{\gamma}} \int_{-\infty}^{\infty} d\Delta \Theta(\Delta) \), \( N \) being the total number of atoms and \( \tilde{L} \) the length of the medium. The last term of Eq. (A5) reduces to

\[
\frac{g^{2}N}{c} \theta_{0}^{2} e^{-\Theta^{2}/2} \int_{0}^{t} dt' \tilde{\Theta}(t - t') \tilde{\mathcal{E}}_{s}(z,t').
\]

Since we are interested in the emission process which has a typical duration \( \tau_{s} \), we only need to consider values of \( t - t' \) of order \( \tau_{s} \). Considering the regime where \( \tau_{s} \ll 2\pi /\Delta_{0} \), only the central peak of \( \tilde{\Theta} \) contributes and \( \tilde{\Theta}(t - t') \) is well approximated by

\[
\tilde{\Theta}(t - t') \approx e^{-(t-t')^{2}/\Gamma^{2}/2}.
\]

For \( \Gamma \tau_{s} > 1 \), \( \tilde{\Theta}(t - t') \) acts as a delta function

\[
\tilde{\Theta}(t - t') \approx e^{-(t-t')^{2}/\Gamma^{2}/2} \approx \frac{\sqrt{2\pi}}{\Gamma} \delta(t - t'),
\]
and the equation for the Stokes field reduces to
\[
\frac{d}{dz} \hat{E}_s(z,t) = i \frac{g_L}{c} \sum_j \delta(z - z_j) e^{-i \Delta t/j} \sigma_j^r(0) + \bar{\alpha} \frac{\theta_0^2}{2} e^{-\bar{\alpha}z} \hat{E}_s(z,t). \tag{A6}
\]

Here \( \bar{\alpha} \), which corresponds to the optical depth per unit length, is defined by
\[
\bar{\alpha} := \frac{\sqrt{2\pi}}{4 \Gamma} \frac{g^2 N}{c}. \tag{A7}
\]

In the main text, we introduced the optical depth per unit length associated to the central peak \( \alpha \). It is given by \( \alpha := \frac{\varepsilon N \Delta_0}{\Gamma} \), so that \( \bar{\alpha} = \sqrt{\frac{2\pi \alpha}{4 \Gamma}} \).

It is then straightforward to check that the solution of Eq. (A6) is given by
\[
\hat{E}_s(z,t) = e^{\frac{g_L}{c} \int_{z_j}^z e^{\bar{\alpha}z'} dz'} \hat{E}_s(0,t) + i \frac{g_L}{c} \sum_{j: \bar{\alpha}z_j < z} e^{\frac{g_L}{c} \int_{z_j}^z e^{\bar{\alpha}z'} dz'} e^{-i \Delta t/j} \sigma_j^r(0). \tag{A8}
\]

To find the average number of Stokes photons per temporal mode, we first evaluate the ket \( \hat{E}_s(L,t_d) | \Psi_w \rangle \). Since initially there is no excitation in the Stokes mode \( | \Psi_w \rangle = | \psi_w,0 \rangle \), we obtain
\[
\hat{E}_s(L,t_d) | \Psi_w \rangle = i \frac{g_L}{c} \sum_j \left( e^{\frac{g_L}{c} \int_{z_j}^L e^{\bar{\alpha}z'} dz'} e^{-i \Delta t/j} \sigma_j^r(0) \right) \times \prod_{k \neq j} (G_k | g_k \rangle + E_k | e_k \rangle) | 0 \rangle. \tag{A9}
\]

Note that each term of the sum has only one atom in the state \( | s_j \rangle \), so when taking the scalar product with the corresponding bra, only the term with the same atom in \( | s_j \rangle \) will give a nonzero contribution. Under the approximation \( | \hat{E}_j \rangle | s_j \rangle \approx | \theta_0^2 e^{-\bar{\alpha}z} \rangle \), one gets
\[
\langle \Psi_w | \hat{E}_s^\dagger(L,t_d) \hat{E}_s(L,t_d) | \Psi_w \rangle = g^2 L^2 \frac{\Delta_0^2}{c^2} \sum_{j: \bar{\alpha}z_j < L} e^{\frac{g_L}{c} \int_{z_j}^L e^{\bar{\alpha}z'} dz'} | s_j \rangle \langle \theta_0^2 e^{-\bar{\alpha}z} | s_j \rangle. \tag{A10}
\]

Finally, we replace the sum over \( j \) by the integral expression. Since there is no term with a spectral dependence, the integral over \( \Delta \) is carried out directly and gives 1. This yields to the spatial density of photons
\[
\frac{1}{L} \langle \Psi_w | \hat{E}_s^\dagger(L,t_d) \hat{E}_s(L,t_d) | \Psi_w \rangle = g^2 L^2 \frac{\Delta_0^2}{c^2} \int_0^L d z e^{\bar{\alpha} z} | s_j \rangle \langle \theta_0^2 e^{-\bar{\alpha}z} | s_j \rangle = g^2 L^2 \frac{\Delta_0^2}{c^2} \left( e^{\frac{g_L}{c} \int_0^L e^{\bar{\alpha}z'} dz'} - 1 \right) = g^2 L^2 \frac{\Delta_0^2}{c^2} \left( e^{\theta_0^2 - 1} - 1 \right) \approx \frac{\Gamma}{\sqrt{2\pi c}} \theta_0^2 \left( 1 - e^{-\bar{\alpha}L} \right),
\]

where, at the last line, we expanded the exponential to the first order. Consequently, the number of photons per temporal mode of duration \( \sqrt{2\pi} / \Gamma \) is given by
\[
\frac{\sqrt{2\pi}}{\Gamma} \frac{c}{L} \langle \Psi_w | \hat{E}_s^\dagger(L,t_d) \hat{E}_s(L,t_d) | \Psi_w \rangle \approx \theta_0^2 \left( 1 - e^{-\bar{\alpha}L} \right). \tag{A11}
\]

Note that, in the general case, the number of photons emitted in a temporal mode \( \sqrt{2\pi} / \Gamma \) is given by \( e^{\bar{\alpha}L} - 1 \). In the particular case where all the atoms are prepared in \( e \), the average number of Stokes photons per mode is given by \( e^{\bar{\alpha}L} - 1 \). This agrees with the result presented in Ref. [16].

Further, note that the number of Stokes photons emitted per write attempt is given by
\[
\frac{2\pi c}{\Delta_0 L} \langle \Psi_w | \hat{E}_s(L,t_d) \hat{E}_s(L,t_d) | \Psi_w \rangle \approx \frac{\sqrt{2\pi}}{\Delta_0} \theta_0^2 \left( 1 - e^{-\bar{\alpha}L} \right),
\]

and is thus roughly the product of the number of peaks \( \Gamma / \Delta_0 \) composing the atomic frequency comb by the success probability for the emission of a Stokes photon per mode. It can thus be merely increased by making use of a wider range of the inhomogeneous broadening.

### 3. Atomic state prepared by the Stokes photon detection

Consider the successful event where a Stokes photon is detected at time \( t_d \). The conditional atomic state \( \psi_d \) is obtained by calculating the evolution of the initial state \( \Psi_w \) until the time \( t_d \) and by projecting the resulting state into \( | 0 \rangle | \hat{E}_s(L,t_d) \rangle \). First, let us directly note that the initial state can be written as \( | \psi_w \rangle = \prod_{j=1}^N \left( G_j | g_j \rangle + E_j | e_j \rangle \right) | 0 \rangle \) where \( \sigma_j^r \) is an eigenstate associated to the eigenvalue 0, it stays unchanged in time. Furthermore, the operator \( \sigma_j^r \) merely acquires the phase term \( e^{-i \Omega j t} \). The evolution of \( \sigma_j^r \) is obtained by plugging the field solution (A8) back into the atomic solution (A4) and then taking the adjoint. For \( \theta_0 \ll 1 \), we get
\[
| \psi_d \rangle = \xi \sum_{j \geq 1} e^{i \varphi_j} \left( e^{\varphi_j - \frac{\Theta_0}{\sqrt{1 - e^{-\bar{\alpha}L}}} / 2} e^{-i \Delta t/j} | s_j \rangle \right) \times \prod_{j \neq j} (G_j e^{-i \varphi_j | g_j \rangle} + E_j e^{-i \varphi_j | e_j \rangle} | e_j \rangle).
\]

with \( \xi = \frac{1}{\sqrt{N (1 - e^{-\bar{\alpha}L})}_2} \). Furthermore, at time \( t \) after the detection of the Stokes photon, a read pulse resonant with the transition \( s \rightarrow e \) and associated with the Rabi frequency \( f \Delta \Omega_j(s) = \pi \) goes through the atomic ensemble and exchanges the population of states \( s \) and \( e \). Therefore, the atomic state becomes
\[
| \psi_r \rangle = \xi \sum_{j \geq 1} e^{i \varphi_j} \left( e^{\varphi_j - \frac{\Theta_0}{\sqrt{1 - e^{-\bar{\alpha}L}}} / 2} e^{-i \Delta t/j} | e_j \rangle \right) \times \prod_{j \neq j} (G_j e^{-i \varphi_j | g_j \rangle} + E_j e^{i \varphi_j | e_j \rangle} | e_j \rangle).
\]

We now have all the necessary ingredients to calculate the efficiency of the readout process.
4. Readout efficiency

The operator corresponding to the envelope of the anti-Stokes field propagating in the backward direction is given by

$$\hat{\xi}_{\text{AS}}(z,t) = \sqrt{\frac{L}{2\pi c}} e^{i\omega_{\text{AS}} z + \frac{c}{2} z^2} \int d\omega \hat{a}_\omega(t) e^{i\omega z / c}. \quad (A13)$$

Its evolution is given by the equations of motion

$$\left( \frac{d}{dt} - c \frac{d}{dz} \right) \hat{\xi}_{\text{AS}}(z,t) = -i gL \sum_j \delta(z - z_j) \sigma_j^g, \quad (A14)$$

$$\frac{d}{dt} \sigma_j^g = -i \Delta_j \sigma_j^g + i g \hat{\xi}_{\text{AS}}(z_j,t) \sigma_j^g, \quad (A15)$$

where $\sigma_j^g = \{|e\rangle \langle e| - |g\rangle \langle g|\}$ is approximated by $\sigma_{j,g} \approx -1$ in what follows. Using the methods presented before, we find at $z = 0$

$$\hat{\xi}_{\text{AS}}(0,t) = e^{-\frac{zL}{c}} \hat{\xi}_{\text{AS}}(L,t) + \frac{i gL}{c} \sum_{z_j} e^{-\frac{zL}{c}} e^{-i\Delta_j (t - t_d)} \sigma_j^g (t + t_d). \quad (A16)$$

This solution allows one to evaluate the readout efficiency. For a perfect spatial phase matching (all the wave vectors sum to zero) the ket $\hat{\xi}_{\text{AS}}(0,t) |\Psi_r\rangle$ is given by (up to a global phase factor)

$$i \frac{gL}{c} \sum_j e^{-\frac{zL}{c}} e^{-i\Delta_j (t - t_d)} |g_j\rangle \prod_{\ell \neq j} (G_\ell e^{-i\omega_\ell (t + t_d)} |g_\ell\rangle$$

$$+ E_i e^{\bar{z} e_\omega e^{-i\omega (t + t_d)} |e_\omega\rangle} \langle 0 |. \quad (A17)$$

Projecting on the corresponding bra, one can show that the spatial density of photons is well approximated by

$$\frac{1}{L} \langle \Psi_r | \hat{\xi}_{\text{AS}}(0,t) \hat{\xi}_{\text{AS}}(0,t) |\Psi_r\rangle \approx \frac{g^2 L}{c^2} \sum_j e^{-\frac{zL}{c}} e^{-i\Delta_j (t - t_d)} G_j$$

$$\times \left( \sum_{\ell} e^{-\frac{zL}{c}} e^{i\Delta_\ell (t - t_d)} G_\ell^* \right). \quad (A18)$$

Each sum is then replaced by an integral and in the limit where $\theta_0 \ll 1$

$$\sum_j e^{-\frac{zL}{c}} e^{-i\Delta_j (t - t_d)} G_j \approx \frac{N}{L} \int_0^L d\omega \sum_j e^{-\frac{zL}{c}} \int_{-\infty}^{\infty} d\Theta \Theta e^{-i\Delta\Theta (t - t_d)}.$$

$$\quad (A19)$$

The spatial integral is carried out directly, while the spectral integral is related to the Fourier transform of the atomic distribution. We obtain

$$\frac{1}{L} \langle \Psi_r | \hat{\xi}_{\text{AS}}(0,t) \hat{\xi}_{\text{AS}}(0,t) |\Psi_r\rangle \approx \frac{\Gamma}{\sqrt{2\pi c}} (1 - e^{-\bar{z} L}) |\bar{\Theta}(t - \tau)|^2. \quad (A20)$$

By evaluating the Fourier transform of the chosen atomic distribution $(A1)$ for the first revival time (i.e., around $t = 2\pi / \Delta_0$) we end up with

$$\frac{\sqrt{2\pi} c}{\Gamma} \langle \Psi_r | \hat{\xi}_{\text{AS}}(0,2\pi / \Delta_0 + \tau) \hat{\xi}_{\text{AS}}(0,2\pi / \Delta_0 + \tau) |\Psi_r\rangle \approx (1 - e^{-\bar{z} L}) e^{-\frac{\tau^2}{4\Delta_0^2}}.$$ 

The success probability to find an anti-Stokes photon within a temporal mode of duration $\sqrt{2\pi} / \Gamma$ centered at $2\pi / \Delta_0 + \tau$ approaches 1 for large enough optical depth provided that the comb finesse is optimized.

Note that for a quantum memory based on an atomic frequency comb, the efficiency has a similar expression but the term $(1 - e^{-\bar{z} L})$ is squared because the photon has first to be absorbed before being reemitted. For weak optical depths, a spontaneous Raman scheme is thus more efficient than the corresponding quantum memory. For example, for a $L = 0.1$ and optimized finesses, the efficiency of the spontaneous Raman scheme approaches 1% whereas it is limited to 0.1% for a quantum memory based on the atomic frequency comb.

Further, note that for an emission in the forward direction, the readout efficiency is given by

$$\frac{(\bar{a} L)^2 e^{-\bar{a} L}}{1 - e^{-\bar{a} L} e^{-\frac{\tau^2}{4\Delta_0^2}}}, \quad (A21)$$

and is limited to 65% by reabsorption instead of 54% for a quantum memory [22].

5. Noise

The atoms excited at the write level can produce spontaneous noise in the anti-Stokes mode of interest. To get a lower bound on the signal-to-noise ratio, we assume that all the atoms transferred to the excited state at the write level decays on the $e-s$ transition. The evolution of the resulting noise is similar to the one associated to the collective emission, except that the atomic state is now given by $\rho_n$ [see Eq. (10) in the main text]. By applying $\hat{\xi}_{\text{AS}}(0,t)$ on $\rho_n$, we obtain

$$i \frac{gL}{c} \sum_j \theta_0^2 e^{-\frac{zL}{c}} e^{-i\Delta_j (t - t_d)} e^{i\omega_j (t + t_d)} |g_j\rangle \langle e_j|$$

$$\otimes (1 - \theta_0^2 e^{-\bar{z} L}) |g\rangle \langle g| + \theta_0^2 e^{-\bar{z} L} |e\rangle \langle e| \otimes |0\rangle \langle 0|. \quad (A22)$$

We then apply the adjoint $\hat{\xi}_{\text{AS}}^\dagger(0,t)$ before taking the trace. One sees that for the trace to be nonzero, each term $e^{i\omega_j (t + t_d)} |g_j\rangle \langle e_j|$ has to be multiplied by the corresponding operator $\sigma_j^g (t + t_d)$ from $\hat{\xi}_{\text{AS}}^\dagger(0,t)$. Since the part of the state on the second line has a trace equal to 1, we get

$$\frac{1}{L} \text{tr}[\hat{\xi}_{\text{AS}}^\dagger(0,2\pi / \Delta_0 + \tau) \hat{\xi}_{\text{AS}}(0,2\pi / \Delta_0 + \tau) |\Psi_r\rangle \rho_n]$$

$$= \frac{g^2 L}{c^2} \sum_j \theta_0^2 e^{-2\bar{z} L},$$

and we conclude that the noise is bounded by $\frac{\theta_0^2}{2} (1 - e^{-2\bar{z} L})$ by replacing the discrete sum by its integral expression and by taking the duration of the anti-Stokes mode into account.
Therefore, the signal-to-noise ratio $R$ is bounded by

$$R \geq \frac{2(1 - e^{-\bar{\alpha}L})}{\bar{\theta}_0^2(1 - e^{-2\bar{\alpha}L})} e^{-\frac{\pi^2}{2n^2 F^2}},$$

that is, for large optical depth and optimized finesse

$$R \geq \frac{2}{\bar{\theta}_0^2}.$$ 

Note that, in the case of a quantum memory, the photon to be stored is subject to many defective manipulations (e.g., to the nonunit coupling efficiency into monomode fibers or to imperfect spectral filters). This reduces the signal and hence strongly limits the signal-to-noise ratio. This is a significant drawback with respect to spontaneous Raman-based sources where the readout efficiency is conditioned on the successful detection of a Stokes photon.

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