Generating single-photons “on demand” represents the ultimate control of the light emission process. Impressive progress has been made in the past several years in obtaining such sources using single quantum dot based technology [1–3] and photonic band-gap materials [4]. The intense effort is motivated by the emerging field of quantum information science [5], which requires deterministic single-photon sources. Indeed, with nearly 100% conversion efficiency [3], such single-photon sources may be used in broad fields of research and technology ranging from single-photon quantum key distributions [6], quantum circuitry for single-photon switching [7], to single uv photon imaging technology [8].

In this work, we investigate a highly efficient ultraviolet single-photon “on demand” scheme using the highly efficient single probe photon four-wave mixing (FWM) technique [9,10]. The motivation for this study is twofold. First, a highly efficient single-photon source in the uv region is not available and its entanglement properties and potentials should be explored [11]. Secondly, there has been no study on a high-efficiency scheme to entangle photons of very different frequencies, and a study that provides basic understandings of this process is desirable. Aside from these fundamental physics viewpoints, a well characterized single uv photon source is also desirable for the fast development of single-photon counting sensors working in the uv spectra region. Such a detection system, although insensitive to the entanglement properties of the uv photon, requires well characterized single-photon responsivity in the region of 180—800 nm. Furthermore, fast developing GEM-based single uv photon multiplier technology for uv imaging [8] requires a well characterized single uv photon source as a metrological calibration source to provide a well traceable metrological standard. These considerations have prompted us to investigate efficient single uv photon generation schemes and to study the fundamental physical properties of such systems.

The scheme reported here has four unique features. (1) It provides the first single-photon “on demand” source in ultraviolet spectrum region and the deterministic properties of the generated ultraviolet photons are dictated by the deterministic properties of the single-photon input source. (2) It does not require high-energy, short-pulsed pump lasers as in the case of the familiar down-conversion scheme to produce single-photons. In fact, using a well-characterized single-photon on demand source, our scheme is capable of generating single ultraviolet photons (≈300 nm) using highly stable commercial infrared (≈800 nm) diode lasers with near 100% photon flux conversion efficiency. This may significantly improve the stability and controllability of the system, both of which are critically important to applications in quantum information processing. (3) Under suitable conditions, the generated ultraviolet photons can travel in the medium with ultraslow group velocities (e.g., v_g/c ∼ 10^{-4} or less), and both the ultraviolet and infrared photon wave packets maintain a well matched temporal profile as that of the input single-photon wave packet. It has been established that such ultraslow propagation velocities significantly increase the interaction time between single-photon wave packets and may significantly enhance certain nonlinear processes [9,10,12–15]. These enhancement effects may lead to controlled interaction between photons, which is a main challenge in quantum information processing. Finally, (4) it opens the possibility of achieving very efficient quantum entanglement of photons with very large frequency differences within a small propagation distance. We will show that it is possible to achieve nearly maximum entanglement of two quantized electromagnetic fields with very large frequency differences for an extended period of time permitted by the ultraslow propagation of single-photon wave packets. This may lead to the possibility of transferring quantum entanglement properties from multiple low-frequency carriers to a single ultraviolet carrier. In addition to the above described...
aspects from a fundamental physics point of view, the system studied may also have potential as a standard single UV photon calibration source for fast developing single UV photon sensor technology. For instance, it can serve as a well-characterized single-photon source for GEM-based single UV photon multipliers as a metrological traceable calibration standard.

The scheme under consideration is a lifetime broadened four-state ladder system (Fig. 1). Here, we use a quantum dot single-photon on demand laser source ($\Omega_{QD}$) as the first step of frequency upconversion process (i.e., $|0\rangle \rightarrow |1\rangle$). We note that the following calculation is independent of the type of single-photon source as long as it is a well-characterized “on-demand” type single-photon source. Two continuous wave (cw) classical fields ($\Omega_c, \Omega_b$) complete the remaining two steps up to resonantly reach state $|3\rangle$ where a phase matched FWM field ($\Omega_F$) is originated. We show that with experimentally achievable parameters, a single-photon FWM field in the ultraviolet spectrum region can be very efficiently generated. We further show that under suitable conditions, both the ultraviolet and infrared photon wave packets can travel with well matched ultralow group velocity.

Our investigation begins with the system Hamiltonian given by

$$
\hat{H}/\hbar = -\sum_{j=1}^{3} \Delta_j |j\rangle\langle j| - (\Omega_c|2\rangle|1\rangle + \Omega_b|3\rangle|2\rangle + \text{c.c.}) - (\Omega_{QD}|1\rangle|0\rangle + \Omega_F|3\rangle|0\rangle + \text{H.c.}).
$$

Using Eq. (1), the equations of motion for the atom density operator $\hat{\rho}$ and the quantized electromagnetic fields can be obtained as

$$
\frac{d\hat{\rho}_{10}}{dt} = i(\Delta_1 + i\gamma_1)\hat{\rho}_{10} + i\Omega_{QD}\hat{\rho}_{00} + i\Omega_c\hat{\rho}_{20} - i\hat{\rho}_1\hat{\Omega}_{QD} - i\hat{\rho}_2\hat{\Omega}_F.
$$

(2a)

$$
\frac{d\hat{\rho}_{20}}{dt} = i(\Delta_2 + i\gamma_2)\hat{\rho}_{20} + i\Omega_b\hat{\rho}_{30} + i\Omega_c\hat{\rho}_{10} - i\hat{\rho}_2\hat{\Omega}_{QD} - i\hat{\rho}_3\hat{\Omega}_F.
$$

(2b)

$$
\frac{d\hat{\rho}_{30}}{dt} = i(\Delta_3 + i\gamma_3)\hat{\rho}_{30} + i\Omega_b\hat{\rho}_{00} + i\Omega_c\hat{\rho}_{20} - i\hat{\rho}_3\hat{\Omega}_{QD} - i\hat{\rho}_3\hat{\Omega}_F.
$$

(2c)

$$
\frac{d\hat{\Omega}_{QD}}{dc} + \frac{1}{c}\frac{d\hat{\Omega}_F}{dt} = i\kappa_0\hat{\rho}_{10},
$$

(2d)

$$
\frac{d\hat{\Omega}_F}{dc} + \frac{1}{c}\frac{d\hat{\Omega}_Q}{dt} = i\kappa_0\hat{\rho}_{30}.
$$

Here, $2\Omega_{B(C)}$ and $\omega_{B(C)}$ are the Rabi and optical frequencies of the classical pump field $E_{B(C)}$, $2\hat{\Omega}_{QD(F)} = D_{01(03)}\hat{\mathcal{E}}_{QD(F)}^{(\pm)}/\hbar$ is the Rabi frequency operator for the quantized infrared (FWM) single-photon field $\hat{\mathcal{E}}_{QD(F)}^{(\pm)}$ with frequency $\omega_{QD}$ ($\omega_F$), $\gamma_j$ denotes the decoherence rate of $\hat{\rho}_{00}$, and $\kappa_0$ is $2N\omega_{QD(F)}|D_{01(03)}|^2/(\hbar c)$ with $N$ and $D_{01(03)}$ being the concentration and dipole moment between states $|0\rangle$ and $|1\rangle$ ($|3\rangle$), respectively. In deriving Eqs. (2a)–(2d), we have assumed that the input infrared field is in a single-photon wave packet state with a shape function $F(t)$ (see below). We have also taken the slowly varying envelope approximation and defined detunings $\Delta_1 = \omega_{QD} - \epsilon_1/\hbar$, $\Delta_2 = \omega_{QD} + \omega_c - \epsilon_2/\hbar$, and $\Delta_3 = \omega_{QD} + \omega_c + \omega_b - \epsilon_3/\hbar$ with $\epsilon_j$ being the energy of state $|j\rangle$ ($\epsilon_0 = 0$).

The task of solving Eqs. (2a)–(2d) for the quantized fields starts with the nondepleted ground-state approximation, i.e., $\hat{\rho}_{00} \approx 1$. We note that the last two terms in Eqs (2a)–(2c) are higher-order terms of small quantities $\Omega_{QD}$ and $\Omega_F$. For a small signal treatment, we neglect these higher-order terms.
By taking the time Fourier transform of Eqs. (2a)–(2d), we obtain
\begin{align}
(\omega + \Delta_1 + i \gamma_1) \hat{\alpha}_{10} + \Omega_c^* \hat{\alpha}_{20} &= - \hat{\Lambda}_{QD}, \\
\Omega_c \hat{\alpha}_{10} + (\omega + \Delta_2 + i \gamma_2) \hat{\alpha}_{20} + \Omega_p^* \hat{\alpha}_{30} &= 0, \\
\Omega_p \hat{\alpha}_{20} + (\omega + \Delta_3 + i \gamma_3) \hat{\alpha}_{30} &= - \hat{\Lambda}_F, \\
\hat{L}_d \hat{\Lambda}_{QD} = i \kappa_0 \hat{\alpha}_{10}, \quad \hat{L}_d \hat{\Lambda}_F = i \kappa_0 \hat{\alpha}_{30},
\end{align}
where \( \hat{\alpha}_{j0}, \hat{\Lambda}_{QD}, \) and \( \hat{\Lambda}_F \) are the Fourier transforms of \( \hat{\alpha}_j, \) \( \hat{\Lambda}_{QD}, \) and \( \hat{\Lambda}_F, \) respectively, \( \omega \) is the Fourier transform variable, and \( \hat{L} = \partial / \partial z - i \omega / c. \)

The solution to Eqs. (3a)–(3c) can be found as
\begin{align}
\hat{\Lambda}_{QD}(z, \omega) &= \frac{\hat{\Lambda}_{QD}(0, \omega)[U_+ e^{i \kappa_0 z} - U_- e^{-i \kappa_0 z}] - \hat{\Lambda}_F(0, \omega)[e^{i \kappa_0 z} - e^{-i \kappa_0 z}]}{U_+ - U_-}, \\
\hat{\Lambda}_F(z, \omega) &= \frac{\hat{\Lambda}_F(0, \omega)[U_+ e^{i \kappa_0 z} - U_- e^{-i \kappa_0 z}] + U_+ U_- \hat{\Lambda}_{QD}(0, \omega)[e^{i \kappa_0 z} - e^{-i \kappa_0 z}]}{U_+ - U_-},
\end{align}
where
\begin{align}
K_\pm &= \omega / c + (\kappa_0 D_F + \kappa_0 D_{QD} \pm G) / (2D), \\
U_\pm &= (\kappa_0 D_F - \kappa_0 D_{QD} \pm G) / (2\kappa_0 \Omega_c^* \Omega_p), \quad G = \sqrt{[\kappa_0 D_F - \kappa_0 D_{QD}]^2 + 4 \kappa_0 \kappa_0 [\Omega_c^* \Omega_p]^2}.
\end{align}

We focus our attention on the adiabatic regime, where \( K_\pm \) and \( U_\pm \) can be expressed by a rapidly converging power series of \( \omega. \) This allows analytical evaluation of Eqs. (5a) and (5b) so that a clear physical picture of the process can be obtained. To ensure such a robust adiabatic process, we assume \( |\Omega_c|^2, |\Omega_p|^2 > \max(|\Delta_1|, |\Delta_2|, |\Delta_3|). \) It can be shown that under these conditions, the linearization treatment introduced before is well justified. When these conditions are satisfied, we find that \( U_\pm = W_+ + O(\omega) \) and \( K_\pm = (K_\pm)_{\omega=0} + \omega / V_{gs} + O(\omega^2) \) can accurately describe the process at hand. After applying the inverse Fourier transform, we obtain from Eqs. (5a) and (5b):
\begin{align}
\hat{\Omega}_{QD}(z, t) &= \frac{W_+ \hat{\Omega}_{QD}(\eta_+) - W_- \hat{\Omega}_{QD}(\eta_-)}{W_+ - W_-}, \\
\hat{\Omega}_F(z, t) &= \frac{W_+ [\hat{\Omega}_F(\eta_-) - W_- \hat{\Omega}_{QD}(\eta_+)] e^{i \beta_+} + W_- [W_+ \hat{\Omega}_{QD}(\eta_-) - \hat{\Omega}_F(\eta_+)] e^{i \beta_-}}{W_+ - W_-},
\end{align}
where \( \hat{\Omega}_{QD}(t) = \hat{\Omega}_{QD}(z=0, t) \) and \( \hat{\Omega}_F(t) = \hat{\Omega}_F(z=0, t) \) are the quantized infrared and FWM quantized fields at the entrance \( z=0, \) respectively, \( \eta_\pm = t - z / V_{gs}, \) \( \beta_\pm = i (K_\pm)_{\omega=0}, \) and
\begin{align}
\beta_+ &= i \frac{\kappa_0 \kappa_0 \Delta_2}{\kappa_0 |\Omega_p|^2 + \kappa_0 |\Omega_c|^2} - \frac{\kappa_0 \kappa_0 \gamma_3}{\kappa_0 |\Omega_p|^2 + \kappa_0 |\Omega_c|^2}, \\
\beta_- &= \frac{(\kappa_0 |\Omega_p|^2 + \kappa_0 |\Omega_c|^2)(B_1 + i B_2)}{B_1^2 + B_2^2}.
\end{align}
quently, after a characteristic propagation distance, the field operators take the form

\[
\hat{E}_{0D}^{(+)}(z,t) = \frac{\kappa_{01}\Omega C_0^2 \exp(z\beta_c)}{\kappa_{01}\Omega^2_{\theta l}^2 + \kappa_{03}\Omega C_0^2} \left[ \hat{E}_{0D}(t-\frac{z}{V_g}) \right] \\
+ \frac{\kappa_{01}D_{03}\Omega B}{\kappa_{03}D_{01}\Omega} \left( \hat{E}_{0D}(t-\frac{z}{V_R}) \right),
\]

(8a)

\[
\hat{E}_{0D}(z,t) = \left( \frac{D_{03}\Omega B}{D_{01}\Omega} \right) \hat{E}_{0D}(z,t),
\]

(8b)

\[\kappa_{01}D_{03}/\kappa_{03}D_{01} = \sqrt{\omega_{QD}/\omega_{01}} \sqrt{\omega_{01}/\omega_{03}}.\]

We note that Eq. (8) indicates that the two quantized fields travel with the same group velocity \(V_g\). With appropriate and experimentally achievable parameters (see below), it can be shown that the matched group velocity can be substantially smaller than the speed of light in vacuum. Under the input condition, there exists only an infrared quantum dot single photon with the pulse shape function \(F(t)\) of single photons. It is straightforward from Eq. (8) [16] to obtain the state of the total quantized field at the exit \(z=L\) as follows:

\[
|\Psi\rangle = CF \left( t - \frac{L}{V_g} \right) |1_{QD}, 0_F\rangle \\
+ \left( \frac{\kappa_{01}D_{03}\Omega B E_{0D}(z,t)}{\kappa_{03}D_{01}\Omega} \right) |0_{QD}, 1_F\rangle \exp(i\theta - \alpha L),
\]

(9)

where

\[
\theta = \kappa_{01}D_{03}L/\kappa_{03}\Omega C_0^2 + \kappa_{03}\Omega C_0^2, \quad \alpha = \kappa_{01}D_{03}L/\kappa_{03}\Omega C_0^2 - \kappa_{03}D_{01}\Omega C_0^2,
\]

and \(C\) is a normalized constant. Equation (10) shows that with appropriate choice of intensities of the two classical fields so that \(\kappa_{01}D_{03}\Omega B \sim \kappa_{03}D_{01}\Omega C\), we obtain a nearly maximum entangled photon pair (the entanglement of two frequency modes) with very large frequency difference. Furthermore, it is possible to adjust the probability amplitude of the ultraviolet photon wave packet by varying the intensities of the control fields. This is a remarkable and unique advantage for a single photon on demand source.

We have carried out extensive numerical calculations to establish the validity of the above described analytical treatment by using experimentally achievable parameters. A possible experimental candidate for the proposed system is ultracold \(^{85}\)Rb atoms. We take, for instance, \(|0\rangle = |5S_{1/2}\rangle, |1\rangle = |5P_{1/2}\rangle, |2\rangle = |5D_{3/2}\rangle, \text{ and } |3\rangle = |nP_{3/2}\rangle\) with \(n > 10\). The respective transitions are \(|0\rangle \rightarrow |1\rangle\) at 795 nm \((\gamma_1 = 5.9 \text{ MHz})\), \(|1\rangle \rightarrow |2\rangle\) at 762 nm \((\gamma_2 = 0.8 \text{ MHz})\), and \(|2\rangle \rightarrow |3\rangle\) at 1.3–1.5 \(\mu\text{m} \((\gamma_3 = 0.09 \text{ MHz})\), all accessible with diode lasers. From Eq. (10) it is easily verified that the same set of parameters as those in Fig. 2 leads to a maximum entanglement between the infrared and the ultra violet photon wave packets. In addition, by choosing intensities of driving fields properly, one can adjust the probability amplitude of the ultraviolet photon wave packet. These are remarkable performances in multi-wave frequency upconversion processes.

The ultra violet single photon on demand using highly efficient single probe photon FWM technique and the newly developed efficient quantum dot single photon on demand source may have wide applications in quantum information processing and photo-sensor metrology. It is a novel scheme for achieving maximum entanglement of two photons with very large frequency difference. The flexibility of being able to adjust the probability amplitude of the ultra violet photon wave packet may lead to new research opportunities in quantum information processing. The well matched group velocity and temporal profile may also find applications in quantum computing, quantum cryptography, entanglement, and single photon quantum key distribution schemes.

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[11] Most, if not all, of single-photon on demand schemes using quantum dot or photonic band-gap materials as the core technology, however, can only generate required photons in the near-infrared region. Currently, there is no single photon on demand source that works in the ultraviolet spectral region.
[16] One operates the complex conjugate of Eq. (8) on the left-hand side of a vacuum state vector. After some algebra, one arrives at Eq. (9).