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Topical Review

Semiconductor quantum dots as an ideal source of polarization-entangled photon pairs on-demand: a review

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Abstract

More than 80 years have passed since the first publication on entangled quantum states. Over this period, the concept of spookily interacting quantum states became an emerging field of science. After various experiments proving the existence of such non-classical states, visionary ideas were put forward to exploit entanglement in quantum information science and technology. These novel concepts have not yet come out of the experimental stage, mostly because of the lack of suitable, deterministic sources of entangled quantum states. Among many systems under investigation, semiconductor quantum dots are particularly appealing emitters of on-demand, single polarization-entangled photon pairs. While it was originally believed that quantum dots must exhibit a limited degree of entanglement related to decoherence effects typical of the solid-state, recent studies have invalidated this preconception. We review the relevant experiments which have led to these important discoveries and discuss the remaining challenges for the anticipated quantum technologies.

Keywords: quantum dots, quantum optics, entangled photon pairs

(Some figures may appear in colour only in the online journal)

1. Introduction

Entanglement is one of the most fascinating aspects of quantum mechanics. The idea, originally developed from a Gedankenexperiment (see [1] and [2]), took several decades to be experimentally realized, and henceforward opened up a steadily growing field of research expanding to a manifold of sources generating 'spookily' interacting quantum states. Further,

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Original content from this work may be used under the terms of the Creative Commons Attribution 3.0 licence. Any further distribution of this work must maintain attribution to the author(s) and the title of the work, journal citation and DOI. innovative concepts of entanglement-based computation and communication protocols were developed, leading to a dramatic increase in interest in the topic [3]. If we focus our discussion on the successful distribution of locally generated quantum states to any remote destination—a fundamental requirement of quantum communication—photon states are the optimal solution due to their low environmental interaction. Nowadays, widely developed optical fiber networks could provide the communication basis of such photonic quantum states and finally pave the way towards envisioned concepts featuring secure communication via quantum cryptography [4] or even quantum computational networks [5]. However, this is far from being an easy task as fiber channels are associated with decoherence and losses of transmitted states degrading the quantum information, thus crucially limiting the communication distance [6, 7]. Likewise, a



Figure 1. The quantum repeater concept. (a) In a quantum relay the photon sources S1 and S2 emit pairs of entangled photons, where one photon of each pair is guided to the Bell state measurement apparatus enabling entanglement swapping. As a consequence, the outer photons remain in an entangled state. (b) In order to construct an entire quantum communication channel, several chained quantum relays have to be interfaced for a long-distance distribution of entangled states among individual parties A (Alice) and B (Bob).

quantum state cannot be amplified to a classical communication channel and demands more advanced non-classical technology. Therefore, several concepts of so-called quantum repeaters have been developed over the years [6, 8-12]. The very basic principle is shown in figure 1(a), where two single photons emerging from two different entangled photon-pair sources are used to perform a Bell state measurement, eventually entangling the leftover photons. The procedure of entanglement swapping allows distant and independent photons, which have never interacted before, to be entangled. Ultimately, a chain-shaped arrangement of such quantum relays principally enables the distribution of entangled states over arbitrary long distances (see figure 1(b)). We want to especially emphasize the substantial advantage of working with pairs of locally generated entangled photon-pairs in the subject of quantum repeater schemes, instead of the conditional and probabilistic NOON-state generation when relying on pure single photon sources [13].

The concept illustrated in figure 1 demands sources of high quality entangled photons, fulfilling several requirements. In particular, each source has to emit single and entangled photons deterministically, with high purity, high efficiency, high indistinguishability and a high degree of entanglement [14]. In addition, it would be desirable to have sources that are integrable on chip-like infrastructures compatible with state-of-the-art photonic technologies. In this regard, solid-state sources of entangled photons are of special interest [15]. In past years, the concept of parametric down conversion [15, 16] has been used for the majority of photon-based entanglement experiments. However, besides the obvious advantages of the source, such as its outstanding degree of entanglement and simplicity, one has to deal with severe drawbacks. First, a high degree of photon indistinguishability must be paid for by a loss of source brightness. Second, the photon generation is a probabilistic process and follows Poissonian statistics; hence, there is a nonzero probability of generating multiple photon pairs during a single excitation cycle. Envisioning a quantum repeater scheme according to [8] it would be vulnerable to errors, thus lowering its overall efficiency [17]. It is rather clear that an alternative source is necessary. In 2000, Benson et al presented the concept of a device using a single semiconductor quantum dot (QD) for the generation of polarization-entangled photon pairs via the biexciton-exciton cascade [18]. These QDs possess discrete quantum states similar to single atomic systems and, therefore, are also referred to as artificial atoms. In contrast to a single atom, QDs can be easily integrated within photonic chips and are compatible with actual semiconductor technologies. Recent research promoted QDs as emitters of single indistinguishable photons at high extraction efficiencies [16, 19]. Nonetheless, it was commonly believed for a long time that QDs are not capable of reaching a perfect degree of entanglement because of degradation effects related to the solid-state environment. In strong contrast, recent publications resolved the issue and revealed that QDs can be almost perfect emitters of polarizationentangled photon pairs [20, 21].

Here, we aim to summarize the recent developments in QDs towards the perfect generation of entangled single photon states. In section 2, we introduce the entanglement generation process—the biexciton-exciton cascade—and discuss the main degrading effects altering the quantum superposition state. In section 2.1, we provide information about the optimal growth properties of a QD, followed by promising device structures of energy tunable entangled photon sources in section 2.2. Further, in section 2.3, we review the different excitation regimes focusing on the optical population of a QD and its direct impact on the photon quality. In section 3, we present promising concepts of entanglement sources at high photon yield while simultaneously maintaining high indistinguishability, purity and ideal polarization entanglement.

2. Criteria for perfect entangled photons from a semiconductor quantum dot

The fundamental process for the generation of polarizationentangled photon pairs in a QD is the so-called biexcitonexciton cascade as illustrated in figure 2(a). Initially, the QD is prepared in the biexciton state, which consists of two electron-hole pairs. This state decays via a recombination of one electron-hole pair under the emission of a single photon and leaves the quantum dot in a single exciton state, which subsequently decays via the emission of another single photon. The two electrons in the biexciton state have a spin quantum number of $m_z = +\frac{1}{2}$ and $m_z = -\frac{1}{2}$, while the hole spins—in an ideal QD governed by heavy hole valance band states—have a spin of $m_z = +\frac{3}{2}$ and $m_z = -\frac{3}{2}$. The polarization of the emitted photons is therefore determined by the total momentum of the recombining electron-hole pair, where only transitions with a change in the total momentum of ± 1 are optically active. Hence, the cascade follows either the left or right decay channel-degenerated in energy-leading to a sequence of circular right (R) biexciton and circular left (L) exciton photons, or vice versa. The resulting two-photon state can be written as:

$$|\psi^{+}\rangle = 1/\sqrt{2} \left(|L_{\rm XX}\rangle|R_{\rm X}\rangle + |R_{\rm XX}\rangle|L_{\rm X}\rangle\right),\tag{1}$$



Figure 2. The biexciton-exciton cascade and the effect of fine structure splitting. (a) The biexciton state (XX) decays under the emission of a right circular (R) (left (L)) polarized photon to a single exciton state (X), which subsequently decays to the ground state (G) under the emission of an orthogonal-circular polarized photon. The polarization of the emitted photons is governed by the electron and hole spin configuration of the recombining electron-hole pair. (b) The anisotropic exchange interaction causes the intermediate exciton state to be energetically split by the so-called fine structure splitting (FSS). Consequently, a mixing of the exciton states appears, leading to the emission of two horizontally polarized photons (H) (vertically polarized photons (V)). (c) Polarization resolved μ -photoluminescence measurement of a QD. The biexciton and the exciton are split due to the FSS. (d) Calculated fidelity to the entangled Bell state $|\psi\rangle$ as a function of the time delay between XX and X decay. In the calculation, it is assumed that all other entanglement-reducing mechanisms, except for the FSS, are absent. The principal effect of the FSS is a time-dependent oscillation of the fidelity as depicted for varying FSS of 1 μ eV (red), 4 μ eV (blue) and 8 μ eV (green).

where $L_{\rm XX}$ ($R_{\rm XX}$) and $L_{\rm X}$ ($R_{\rm X}$) are biexciton and exciton photons in the circular left (right) polarization base, respectively. This state corresponds to a maximally entangled Bell state. By using $|H\rangle = 1/\sqrt{2}(|L\rangle + |R\rangle)$ and $|V\rangle = i/\sqrt{2}(|R\rangle - |L\rangle)$, we can rewrite equation (1) in the linear horizontal (H) and vertical (V) base as:

$$|\psi\rangle = 1/\sqrt{2} \left(|H_{XX}\rangle|H_X\rangle + |V_{XX}\rangle|V_X\rangle\right). \tag{2}$$

To take into account possible deviations of the real twophoton state from the ideal $|\psi^+\rangle$ state, a density operator formalism is commonly used. The density operator for a statistical ensemble of states is defined according to:

$$\hat{\rho} = \sum_{i} p_{i} |\phi_{i}\rangle \langle\phi_{i}|, \qquad (3)$$

where p_i is the portion of the ensemble being in the state $|\phi_i\rangle$. In the case of an ideal QD entangled photon emitter, the density operator reduces to a pure state. Using the H and V states as a basis, the matrix representation of the density operator is given by:

$$\hat{\rho}_0 = |\psi\rangle\langle\psi| = \frac{1}{2} \begin{vmatrix} 1 & 0 & 0 & 1 \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \\ 1 & 0 & 0 & 1 \end{vmatrix}.$$
(4)

The verification of photon entanglement emitted by a QD requires the reconstruction of its two-photon density matrix. This is done via polarization resolved cross-correlation measurements between biexciton and exciton photons emitted during the cascade decay. For a detailed description of the procedure and its evaluation we refer the interested reader to [22]. The degree of entanglement can be quantified via different parameters: namely concurrence, tangle, Peres criterion, negativity and fidelity (for a detailed summary see [15]). In works related to QDs, the entanglement fidelity is a widely used figure of merit to compare different sources. It can be defined as:

$$f^+ = Tr[\rho_{mes} \cdot \rho_0], \tag{5}$$

where ρ_{mes} is the reconstructed density matrix obtained from the experiment and ρ_0 is the density matrix of a pure Bell state according to equation (4). If the measured density matrix is equal to the state in equation (9), the resulting fidelity is one, while it is zero in the orthogonal case. Further, if the measured state is a mixed state, the fidelity is a measure of the probability to find an entangled photon pair in the state $|\psi^+\rangle$ in the ensemble. A classically correlated state yields a fidelity of $\frac{1}{2}$, while in the presence of entanglement $\frac{1}{2} < f^+ \leq 1$ holds. We want to point out that the fidelity can also be measured with a reduced measurement set. A full reconstruction of the density matrix is not always required if the source is unpolarized and the measurement setup is not introducing any additional phase shifts. As a consequence the fidelity can be estimated as:

$$f^{+} = \frac{1 + C_{\text{linear}} + C_{\text{diagonal}} - C_{\text{circular}}}{4}.$$
 (6)

Here C are the correlations visibilities according to

$$C_{\mu} = \frac{g_{XX,X}^2 - g_{XX,\overline{X}}^2}{g_{XX,X}^2 + g_{XX,\overline{X}}^2},$$
(7)

where $g_{XX,X}^2$ and $g_{XX,\overline{X}}^2$ are the co- and the cross-polarized correlation measurement, respectively, in the polarization basis μ . The measurement reduces to six cross-correlation measurements (see [23, 24]) instead of 16 for the full density matrix.

In general a QD can underly various entanglement degrading effects, which drastically reduce the measured degree of entanglement and yield $f^+ < 1$. The dominant contributions are: (i) fine structure splitting (FSS), (ii) recapture processes, (iii) valence band mixing and (iv) exciton spin-flip processes. In the following, we would like to discuss these effects in more detail.

(i) The FSS causes a mixing of the $|\pm 1\rangle$ spin states of the intermediate exciton states (compare 2(a) and 2(b)) due to an anisotropic electron-hole exchange interaction. In turn, the broken symmetry leads to a lifting of the degeneracy of the two excitonic states (see figure 2(b)). As a result, the emission of a linear horizontal (H) (vertical (V)) polarized biexciton photon is followed by an equally polarized exciton photon during the cascade decay. The origin of the FSS can be illustrated by the exchange interaction Hamiltonian:

$$H_{\text{exch}} = -\sum_{i=x,y,z} (a_i \hat{S}_{h,i} \cdot \hat{S}_{e,i} + b_i \hat{S}_{h,i}^3 \cdot \hat{S}_{e,i}),$$
(8)

where $S_{h,i}$ ($S_{e,i}$) is the hole (electron) spin operator and a_i and b_i are the spin-spin coupling constants in the x, y, z spatial axis. The FSS arises for example, if the rotational in-plane symmetry of the wavefunction is reduced (for example, due to a geometric symmetry lower than D_{2D}) leading to $b_x \neq b_y$ ($b_x = b_y$ for FSS = 0). For a detailed description we refer the interested reader to [25, 26]. In contrast to the exciton, the biexciton does not show an FSS as the electrons and holes are in a singlet state. However, in polarization resolved μ -photoluminescence measurements (see figure 2(c)), one will observe the FSS in the exciton and biexciton spectral

lines, simply due to the fact that the biexciton decays into an exciton state.

Due to the FSS a time-dependent phase factor is introduced into the two-photon state. Therefore, equation (1) transforms into:

$$|\psi'\rangle = 1/\sqrt{2} \left(|H_{\rm XX}\rangle|H_{\rm X}\rangle + e^{\frac{iSt}{\hbar}}|V_{\rm XX}\rangle|V_{\rm X}\rangle\right),\tag{9}$$

where *S* is the FSS and *t* is the time span between the biexciton and exciton photon emission. The state is still entangled, but corresponds to $|\psi^+\rangle$ defined in equation (9) only for decays with $t = \frac{2N\pi\hbar}{s} = NT$ (with *N* an integer). In general, f^+ will exhibit an oscillatory behavior as a function of decay time [27]. This means that only a narrow subset of the decay events can be used as well-defined entanglement resources as shown in figure 2(d) for different values of the FSS. If no event selection is applied, the time-averaged fidelity will be less than unity. Assuming an exponential time distribution of the exciton photon emission, the resulting density matrix is given by:

$$\rho = \frac{1}{\tau_1} \int_0^\tau e^{-t/\tau_1} |\psi'\rangle \langle\psi'| dt, \qquad (10)$$

where τ_1 is the exciton lifetime and τ is the integration time window. Hence, in the case of a large FSS and/or long exciton lifetime one measures only classical correlations between the emitted photons for time windows $\tau \ge T$, where *T* is the oscillation period of f^+ .

As a consequence, the first attempts towards the demonstration of entangled photon generation with QDs did not show any evidence of non-classical behavior [28]. In 2006, Akopian et al [29] and Young et al [30] confirmed polarization-entangled photons from a single QD [29]. The measurement of Akopian et al relied on narrow spectral filtering of the biexciton and exciton photons to minimize the effect of the FSS. Similarly, one could also apply temporal filtering [27, 31–34], such that photon pairs with high fidelity f^+ are selected. Implementing this scheme in 2017, Huwer et al achieved fidelities to the maximal entangled Bell state up to 0.92 [34]. We want to emphasize that post-selection schemes suffer from severe photon pair losses and inherently limit the possibility of on-demand generation. In particular, temporal filtering introduces an unacceptable increase in complexity to any quantum optics system. A more favorable strategy is to directly eliminate the FSS itself. In sections 2.1 and 2.2, we present two successful strategies to eliminate the FSS without photon losses.

(ii) In a recapture process (also referred to as re-excitation), the intermediate exciton level of the biexciton-exciton cascade is re-excited to the biexciton level before it decays to the ground state. The effect can be either optically driven (see section 2.3) or related to charged carriers trapped in the surrounding QD. Evidence of recapture is, for example, visible in the cross-correlation measurements between biexciton and exciton. It produces coincidence counts at negative time delays corresponding to exciton photons detected before the biexciton, a circumstance not expected from the cascade's temporal order [35]. The detected photons obviously feature no polarization correlation. The effect is well known (see [35–37]) and particularly

dominant under non-resonant excitation (see section 2.3). An additional scenario is that a cascade is followed by re-excitation within the same excitation cycle. If the XX of the first cascade and the X of the second cascade are detected while the other photons are lost to the detection inefficiency, the measured fidelity will be lowered.

(iii) In an ideal system, as described at the beginning of the chapter, the hole state of the exciton is purely heavy. The light hole band is considered to be energetically separated due to strong quantum confinement. In reality, the validity of this assumption is restricted to truly symmetric excitonic wavefunctions and an additional valance band mixing has to be considered in common QDs: the heavy-light hole mixing has two major effects: first, it contributes to the FSS and second, it leads to a depolarization of the exciton [38, 39], with both effects degrading the measured degree of entanglement. Valence band mixing is not only related to the geometric inplane symmetry of the QD, but also depends on strain anisotropies, interface intermixing effects and composition gradients within the QD. Even in the absence of these perturbations, significant valance band mixing can be introduced by a large vertical aspect-ratio of the wavefunction $\eta = l_z/l_v$ (where l_z is the geometric parameter in growth direction and l_v is one in-plane component) as shown by Liao et al in 2012 (see [40]). Hence, a flat highly symmetric strainfree QD shape is preferred in order to avoid FSS and limit valance band mixing (see section 2.1).

(iv) Lastly, we want to discuss the effect of spin-flips affecting the intermediated exciton decay. Such an event randomizes the polarization correlation between the biexciton and exciton photon. After a scattering event equation (9) transforms to:

$$|\psi_{s}\rangle = 1/2(|H_{XX}H_{X}\rangle + |V_{XX}V_{X}\rangle + |H_{XX}V_{X}\rangle + |V_{XX}H_{X}\rangle).$$
(11)

If we first restrict ourselves to non-resonant excitation (see section 2.3), such a spin scattering can be explained by spin-flip processes due to the interaction with excess charges. The fluctuating electric field can induce an effective magnetic field altering the exciton spin via spin–orbit coupling [20, 41].

Under (quasi) resonant excitation conditions, which avoid the generation of excess charges, experiments show that spin scattering processes are still apparent [21]. Another aspect of spin scattering is the interaction with the nuclear spins of the atoms forming the QD and is completely decoupled from the excitation conditions [41-43]. Thereby, it is assumed that the confined electron of the exciton is interacting via Fermi-contact interaction with the nuclear spins, while the heavy hole dephasing is related to a dipole-dipole interaction [44]. In 2017, Fognini et al performed high resolution (30 ps) time-resolved correlation measurements on wurtzite InAsP QDs in a tapered InP nanowire under quasiresonant excitation (see section 2.3). Despite the presence of high nuclear spin In atoms in the QD and a long exciton lifetime (1 ns)—implying a large impact of the nuclei-induced spin dephasing-the authors show that these QDs are dephasing free and the model of spin scattering [43] is not experimentally supported. Further, exciton spin-flips have not been observed in InAs/GaAs QDs at relevant time scales (<30 ns) at cryogenic temperatures (<20 K) [45].

In the work of Huber *et al* [21], the authors performed a measurement of the dependence of the entanglement fidelity versus the FSS under resonant excitation on a single GaAs QD. The results indicate the deviation from perfect entanglement compatible with a residual spin scattering process with an unknown origin.

2.1. Growth of symmetric quantum dots

In the previous section, we discussed that an anisotropy of the confinement potential leads to a finite FSS and valance band mixing. Over the last decade the majority of entanglement experiments have been performed on Stranski-Krastanow InGaAs QDs. Specifically, this material system suffers from anisotropic strain fields related to the growth mode, interface diffusion processes [46] and large values of valance band mixing $(0.2 < |\beta| < 0.7)$, where $\beta = 0$ corresponds to a pure heavy hole and absence of valance band mixing [47]). As a consequence, the probability of finding InGaAs QDs with FSS smaller than 1 μ eV is of the order of a few per cent [48]. In 2013, Juska et al presented a work [49] on pyramidal sitecontrolled InGaAs_{1- δ}N_{δ} QDs, which are grown on a patterned (111)B-oriented GaAs substrate via metalorganic vapor phase epitaxy. The growth technique leads to ensemble of QDs with high structural symmetry (see also [50-52]) and up to 15% of the QDs can generate polarization-entangled photons with a fidelity up to 0.72(4) under non-resonant excitation. A different attempt at symmetrically engineered excitonic wavefunctions based on the InAs material system relys on the growth of large low-strain $In_{0.3}Ga_{0.7}$ As QDs [53] exhibiting fine structure splittings well below $10 \,\mu\text{eV}$ on average, despite structural anisotropies. Here, the shallow confinement potential lowers the sensitivity of the wavefunction with respect to structural asymmetries of the QD. Details of the QD shape practically become negligible compared to the large wavefunction extension. Other material systems, with the exception of GaAs QDs which are introduced in the following in more detail, have not yet met the demanding requirements of a high, and statistically frequent, excitonic confinement symmetry [54–56]. The results clearly highlighted that the shape of the QDs and the in-plane anisotropy of the confining potential have a dominant role in the underlying physics of the FSS. However, even in the case of high wavefunction symmetry, the InGaAs material system still suffers from the In-related large Overhauser field [44], which leads to a fluctuation of the FSS over time [57]. From this point of view, a promising material system, namely GaAs/AlGaAs QDs, is of increasing interest. The material combination is characterized by a negligible lattice mismatch, which facilitates the realization of QDs with high in-plane symmetry combined with the small nuclear spin of Ga (3/2)compared to In (9/2), thus minimizing the effective Overhauser field perturbing the FSS [58].

We want to briefly discuss three approaches, which lead to as-grown GaAs QDs emitting highly entangled photons. The first method is a droplet epitaxy method [59–61], where pure Ga is evaporated on an AlGaAs surface, via molecular beam epitaxy. The Ga forms droplets on the surface, which are subsequently crystallized by an additional As flux. After annealing, the QDs are capped with an AlGaAs layer forming the top barrier of the QD. In 2013, Kuroda et al used such highly symmetric QDs to measure a fidelity of 0.86 (the authors claim an undetectable small value of the FSS) without the aid of photon post-selection. A drawback of the growth method is an unfavorable broad size distribution due to the low crystallization temperature used [62]. In 2017, Basso Basset et al offered an alternative solution by presenting a modified approach on GaAs (111) substrates, where the Ga droplets are crystallized and the subsequent barrier layer is deposited at a high substrate temperature of around 520 °C [63]. The authors claim an improved crystallinity of the QDs by reducing the defect concentration typical for low-temperature growth eventually leading to a higher optical quality of the QDs.

The second method is the fabrication of QDs by etching of nanoholes into a AlGaAs substrate via local droplet-etching [64]. The nanoholes are subsequently filled with GaAs and capped, after annealing, with AlGaAs [62]. In 2013, Huo *et al* demonstrated that, under optimized growth conditions, droplet-etched QDs yield an ultra small FSS with an average value of 4(2) μ eV [65]. Further, a small valance band mixing (<5%) is reported [66, 67]. In addition to the in-plane symmetry, a tall morphology is useful to reduce the FSS, as the exchange interaction depends on the strength of the electronhole wave functions overlap [68].

In 2017, Huber *et al* measured the degree of entanglement and achieved a fidelity of 0.94(1), already at a finite FSS of 1.2(5) μ eV, without any post-selection and is to date the highest value of fidelity measured on a QD without any postgrowth techniques. Similar values of entanglement were reported by Keil *et al*—where the authors claim a fidelity of 0.91 at an FSS of 2.3 μ eV—using the same type of QDs [24]. Both works rely on a resonant two-photon excitation scheme to directly populate the biexciton state (see section 2.3).

In spite of the impressive progress towards the fabrication of entanglement-ready QDs [23, 24, 34, 37], finding QDs with sub- μ eV-FSS in an ensemble is still very challenging due to unavoidable fluctuations occurring during the high-temperature growth of such nanostructures. The mentioned experiments yielding record fidelities are always based on the search of 'hero dots' with low FSS. Nonetheless, fidelities of the order of 0.94 are definitely an attractive value for ready-to-use quantum relays, including error correction protocols [69].

2.2. Energy tunable source of polarization-entangled photons

The compensation of residual asymmetries leading to a finite FSS can possibly be attributed to external perturbations, such as the optical Stark effect, electric/magnetic fields and anisotropic strain fields or a combination of them (for details, see [26, 35, 70, 71]).

When envisioning the future application of QDs in the field of quantum communication one has to deal with an additional problem. A quantum relay, as shown in figure 1(a),

relies on perfect indistinguishablility of the emitted photon pairs. It inherently demands perfect energy matching of photons emitted by different sources, basically requiring all QDs emitting fully identical entangled photons. As the wavelengths of QDs statistically differ from QD to QD the probability of finding at least two QDs under these conditions -at zero FSS and equal energy—is $\approx 10^{-9}$ [72] in standard Stranski-Krastanow QDs. Furthermore, the entangled photons are ideally interfaced with other quantum systems such as atomic clouds or N-V-centers (acting as quantum memories) adding another complex parameter to be controlled [73]. Arguably, the desired condition is not achievable with realistic growth processes. It was shown that the emission energy of a QD can be precisely engineered using the mentioned external perturbations [26, 70], but sophisticated applications demand a device structure that enables independent control of emission energy and FSS. Only a device structure featuring simultaneous control of emission energy and FSS on any emitter of investigation, a truly scalable QD-based quantum communication network is feasible.

Since the commonly used perturbation fields are of vectorial (e.g. electric field) or tensorial (strain field) nature and each of them produces a distinct effect on the electronic structure of QDs, these perturbations provide, in principle, a large enough set of independent 'tuning knobs' to control emission energy of X and XX photons as well as FSS. Two individual concepts were presented in 2015 by Wang et al (see [74]) and by Trotta et al (see [72]), both relying on properly engineered strain fields. The first concept (see figure 3(a)) uses a piezoelectric lead zirconic titanate ceramic stack with QDs glued on top. By applying two independent voltages the QDs are exposed to two independent in-plane strain components, which can be used to erase the FSS of an arbitrary QD [75]. For energy tuning, a second independent stressor is placed on top of the sample. The extraction of the photons, however, requires the technologically challenging use of a transparent strain transmitter between the sample and the top stressor.

The second concept, as shown in figure 3(b), uses a micromachined piezoelectric structure ($[Pb(Mg_{1/3} Nb_{2/3})O_3]_{0.72}$ – $[PbTiO_3]_{0.28}$), with three stress fields acting on the QDs by three pairs of 'legs', as shown in figure 3(b). Three independent voltages (V_1, V_2, V_3) applied across the legs and the top (grounded) contact allow for full control over the in-plane stress state in a nanomembrane—hosting the QDs—bonded on top of the piezoelectric actuator [77]. As a consequence, one can use two legs of the device to fully erase the FSS, while the third leg enables energy tuning at fixed FSS. As the membrane is open to the top, the emitted photons can be easily collected by microscope objectives. Furthermore, the structure enables the possibility of using QDs embedded in a diode structure to electrically pump the entangled photon source (see [33]).

In 2016, Trotta *et al* presented an experimental proof of their concept [76]. The authors erased the FSS of an InAs QD and verified photon entanglement (with resulting fidelity of 0.80(5)) at different energies of the X transition. Additionally, they interfaced the exciton photons with the D1 line of cesium vapor and at the same time kept the FSS constantly below the



Figure 3. Energy-tuneable entangled photon sources based on QDs. (a) A device structure according to Wang *et al.* Adapted figure with permission from [74], Copyright (2015) by the American Physical Society. The piezoelectric actuator (PZT) can be used to apply two independent in-plane stresses via the voltages V_z and V_y to cancel the fine structure splitting. A second stressor on top of a transparent stress transmitter would allow for energy shifting. (b) A device structure according to Trotta *et al.* Adapted by permission from Macmillan Publishers Ltd: Nature Communications [76], Copyright (2016). Six pairwise connected piezoelectric legs on a micromachined PMN-PT substrate are used to apply any arbitrary in-plane stress configuration on the membrane via the voltages V_1 , V_2 , V_3 to erase the fine structure splitting and simultaneously tune the exciton (biexciton) emission energy. (c) Excitonic degeneracy tuning of the same, single QD at arbitrarily chosen emission energies. Adapted by permission from Macmillan Publishers Ltd: Nature Communications [76], Copyright (2016). The piezoelectric actuator from (b) allows one to tune the fine structure to zero independently from the targeted emission energy. The minimum of the red curve coincides with the D2 line of a cloud of Cs atoms.

spectral resolution of the setup ($<0.2 \ \mu eV$) (see figure 3) (c). The QD-Cs interface allowed the authors to observe slow light [78], and in particular, slow entangled photons.

Other approaches enabling wavelength tunable sources of entangled photons with InAs QDs were presented in 2016 by Chen *et al* using patterned thin film piezoelectric structures fabricated throughout focused ion beam and wet chemical etching [79]. In the same year, Zhang *et al* presented an approach using a monolithic piezoelectric actuator in combination with a vertical electric field [80]. We want to emphasize that such 'two-knob' devices act as wavelength tunable sources of entangled photons only if the structural anisotropy of the QD is oriented along the main stress direction of the piezoelectric actuator, which is given for around 30% of the QDs according to [81]. The same consideration applies to approaches combining electric and magnetic fields with fixed orientation [82].

In 2018, Huber *et al* combined the patterned piezo structure with μ -membranes hosting droplet-etched GaAs/AlGaAs QDs in a planar distributed Bragg reflector cavity [21]. The authors achieved an entanglement fidelity up to 0.978(5) after taking into account setup-related entanglement deteriorating effects (phase shifts induced by the optical elements and laser background). Accordingly, the only known limit is a spin-flip process with a characteristic scattering time of 14(10) ns. The origin of the spin scattering is not yet clear. A forced reduction of the excitonic emission lifetime from around 290 ps to 100 ps via Purcell enhancement [36] would allow GaAs QDs to reach levels of near-unity entanglement compatible to parametric down conversion sources with the mentioned benefit of deterministic operation.

2.3. The role of excitation

The population of an excited state in a QD can be achieved in various ways. If we restrict our discussion to pulsed optical pumping one has three major options: (i) non-resonant excitation, (ii) quasi-resonant excitation and (iii) resonant excitation, which we will discuss in more detail:

(i) In non-resonant excitation, the laser is mainly exciting carriers in the barrier material or in the wetting layer of the QD (if available) to photo-generate electron-hole pairs. Subsequently, these electrons and holes are captured by the QDs and relax to the lowest energy levels, the so-called s-shell. Arguably, the excitation regime is not favorable for an entangled photon generation supporting the appearance of pronounced recapture processes [35–37]. Additionally, the fidelity is lowered by spin dephasing processes arising from exciton and excess charge interactions [20]. Besides the negative effects on the entanglement fidelity there are additional drawbacks; first, the possibility of achieving deterministic population of the biexciton state is inhibited [83, 84], and second, the indistinguishability of the emitted photons suffers from the carrier relaxation induced time jitter and fluctuating electric fields [19, 85] (see section 3.1).

(ii) In contrast to non-resonant excitation, quasi-resonant excitation is a more favorable approach for generating entangled photon pairs. In this regime, the electron-hole pairs are generated via excitation in a higher QD shell, e.g. p-shell, and the carriers normally undergo a fast relaxation process to the s-shell. The excitation allows for a 'near' coherent population once resonantly driving the p-shell [41]. In 2017, Fognini *et al* studied the entanglement properties in a InAsP QD under quasi-resonant excitation, where no adverse effects related to the excitation could be identified [20]. However, in a recent study by Kiršanskě *et al*, the authors showed that under quasi-resonant excitation conditions, the indistinguishability is essentially limited by the time jitter introduced via p-shell to s-shell relaxation processes [86].

(iii) In resonant excitation, the electron-hole pairs are directly created in the s-shell of the QD. The resonant population of the biexciton state requires a resonant two-photon absorption process due to dipole selection rules [87, 88]. The concept is shown in the inset of figure 4(a), where the laser is tuned between the biexciton and exciton spectral line. The energy spacing—governed by the biexciton binding energy—is small (typically around 1–4 meV for InAs and GaAs QDs, respectively [23, 24, 84]), such that the laser is energetically close to,



Figure 4. Different coherent excitation schemes. (a) A resonantly excited GaAs quantum dot. Adapted by permission from Macmillan Publishers Ltd: Nature Communications [23], Copyright (2017). The exciton (X) and biexciton (XX) of similar intensity are visible. Two weaker, charged states appear (C1 and C2). Inset: For resonant two-photon excitation, the laser energy E_L is tuned to half the energy of the biexciton state. Two photons are absorbed to directly populate the biexciton state. (b) An exciton (X_0) population under resonant two-photon excitation (TPE) (red), white light assisted TPE (blue) and phonon-assisted excitation (green) versus pulse area on the same QD. The envelope of the Rabi oscillation is modeled with a single exponential damping. As a consequence of the cascade process, the exciton exhibits the same population evolution as the resonantly driven biexciton (see [90]). Adapted with permission from [90], Copyright (2017) American Chemical Society. (c) Population of the X state as a function of the laser detuning for varying excitation power. While the traditional TPE (blue) suffers from a steep drop in inversion efficiency, a stable plateau can be observed exploiting the QD phonon sideband (green). Adapted with permission from [90], Copyright (2017) American Chemical Society. (d) A biexciton population versus power and chirp parameters in order to identify the adiabatic rapid passage excitation condition. Reprinted with permission from [92], Copyright (2013) by the Amiercan Physical Society.

albeit not overlapped, with the QD photons. A sophisticated rejection of the scattered laser light is unavoidable, as already minor contributions of the laser can significantly lower the single photon purity, indistinguishability and the degree of entanglement [21, 23]. This is particularly relevant when measuring the degree of entanglement of the source, as common techniques relying on cross-polarization between excitation and detection cannot be adapted [83]. According to current research, however, the resonant and quasi-resonant excitation do not show appreciable differences in the degree of entanglement, but only on the indistinguishability of the emitted photons. In fact, studies by Ding et al and Somaschi et al proved that QDs are only exhibiting near-perfect degrees of photon indistinguishability under resonant conditions (indistinguishabilities of 0.996(5) [19] and 0.985(4) [89] are reported). Further, the resonant excitation allows for almost near-unity population probability (figure 4(b)) of the biexciton state after application of a single π -pulse [90, 84, 91]—a condition of high importance in terms of quantum efficiency of the entangled photon source. Hence, to simultaneously optimize the degree of entanglement, photon indistinguishability, single photon purity and on-demand generation, a resonant excitation regime is obligatory [23]. The only disadvantage is the severe QD-dependent sensitivity of the resonant condition. Small fluctuations in the laser pulse area, energy and QD environment result in a strong variation of the excited state population probability (see figure 4(c)). Therefore, two alternative excitation schemes based on resonant excitation are introduced, as discussed below:

First, we would like to introduce the adiabatic rapid passage (ARP), an excitation technique well known for single atomic systems. Here, the resonant condition of the QD twolevel system is established utilizing a chirped laser pulse, in contrast to the commonly used transform-limited π -pulse. The duration of the frequency-swept pulse should be significantly shorter than the spontaneous emission time ('rapid') but still slow enough in comparison to the Rabi frequency such that the excited state can be tracked adiabatically. The result of the light–matter interaction Hamiltonian is a traversed anticrossing of the two possible eigenstates [93] causing the system to switch into the excited state at sufficient excitation power. The robustness of the scheme is theoretically demonstrated in figure 4(d) and unambiguously shows the pulse area independent maximum state inversion as a function of the pulse chirp, strikingly different from the very sensitive Rabi oscillations of transform-limited excitation conditions. Nonetheless, the quantum dynamics of the ARP strongly relies on a complex control of the excitation conditions as well as minimized dephasing mechanisms of the coherent superposition of the two-level system such as carrier-phonon interactions [94].

An alternative excitation scheme which combines the simplicity of strictly resonant excitation with the robustness of the ARP is the so-called phonon-assisted two-photon excitation. It was proposed [95] to tune a transform-limited, two-photon excitation laser slightly above the biexciton energy in order to dress a vibrational quasicontinuum and, in the following, enable on-demand preparation of the biexciton state despite the presence of strong carrier-phonon interactions. These interaction terms are the main obstacle to achieving high preparation fidelity, and are particularly pronounced in semiconductor QDs [96, 97]. The proposal was verified in several works [98–100] and its comparison to standard resonant schemes is depicted in figure 4(b). While traditional Rabi oscillations suffer from strong phonon-induced damping for increasing driving strength, the two-photon phonon-assisted excitation unfolds its real potential. At high excitation powers it is not only possible to prepare the QD with comparable levels of state inversion; in addition, the excited state population is independent of laser intensity fluctuations. Most importantly, the phonon sideband dressing of a QD biexciton state is allowed for a certain range of laser energies (figure 4(c)) and presents a feasible way to address complex systems of dissimilar QDs at the same time by stable, frequencylocked laser systems: a condition of high practical advantage impossible to realize with strict resonant excitation. Careful studies revealed that all advantages of π -pulse driven systems regarding their emission quality, in particular polarization entanglement, hold up for the two-photon phonon-assisted condition due to the negligible time jitter introduced by the ultrafast phonon relaxation times [90].

Besides optical excited QDs, electrically driven sources of entangled photons are of special interest as the absence of an excitation laser would drastically reduce the complexity of the system [33, 101]. QDs can be embedded into light-emitting diode structures [102] and several approaches concerning pulsed entangled light-emitting-diodes have been presented [52, 81]. The highest fidelity under pulsed electrical pumping was achieved by Chung *et al* in 2016 featuring $f^+ = 0.68(2)$ at a FSS of $0.2(2) \mu eV$ (see [52]) without any post-selection. The results are significantly lower with respect to optical excitation [23, 24] and suggest entanglement degrading effects, apart from the FSS, are strikingly dominant (especially re-population [52]). The full potential of these devices might only be accessible via an electrically driven resonant population process.

3. Towards a perfect entangled photon source

QD-based sources already approach the state-of-the-art set by parametric down-converters [103] in terms of near-unity entangled-state fidelities. However, besides a high degree of



Figure 5. Two-photon interference. (a) Two photons impinge on a beam splitter and experience the Hong-Ou-Mandel effect, where indistinguishable photons always leave through the same output port. The outputs are equipped with detectors (D) to measure the correlation between the detected photons. (b) A two-photon interference experiment with photons emitted by a single QD under pulsed resonant two-photon excitation. Adapted by permission from Macmillan Publishers Ltd: Nature Communications [23], Copyright (2017). If the photons are indistinguishable (co-polarized), the height of the central peak is reduced with respect to the peak of cross-polarized photons (fully distinguishable photons).

entanglement and the capability of generating photons at predefined energies, additional criteria must be met by the 'perfect' source. For truly on-demand operation, one and only one photon pair should be emitted after each excitation cycle and, in addition, the photons emitted in subsequent cascades should be indistinguishable in all degrees of freedom apart from their polarization. Among these criteria, scalable applications are only feasible if near-unity fractions of the generated photons are extracted from the semiconductor matrix.

3.1. Single-photon purity and photon indistinguishability

The single photon purity of a photon source is characterized using a Hanbury Brown and Twiss (HBT) experiment. It allows extraction of the second-order autocorrelation function $g^{(2)}(\tau)$ versus the time delay τ . A perfect single photon source gives a $g^{(2)}(0) = 0$ and a two-photon source gives $g^{(2)}(0) = 0.5$ [41]. In view of sources of entangled photon pairs, unprecedented values of purity alongside high entanglement fidelity have been achieved, mainly relying on the resonant two-photon excitation of droplet-etched GaAs QDs [23, 24].

The photon indistinguishability can be verified by measuring the two-photon interference (TPI) visibility via the Hong-Ou-Mandel effect [104]. The concept is presented in figure 5(a). Two photons impinge simultaneously on a 50/50beam splitter. If the two photons are indistinguishable in all degrees of freedom (energy, polarization, temporal- and spatial overlap), they will always leave the beam splitter through the same output port. Hence, a measured histogram will show no coincidence counts between the detectors at zero time delay.

The two-photon interference visibility of a single source can be measured by the generation of two single photons using two delayed laser pulses and a respective compensation of the excitation time gap with the aid of an unbalanced Mach-Zehnder interferometer [105]. The result of such a measurement is shown in figure 5(b). The co-polarized (indistinguishable in polarization) photons show a drop in the central peak with respect to the cross-polarized photons (fully distinguishable). The visibility is given by:

$$V = 1 - \frac{g_{\parallel}}{g_{\perp}},\tag{12}$$

where g_{\parallel} and g_{\perp} are the integrated peak areas around the zero time delay for co-polarized and cross-polarized photons, respectively. A visibility equal to one indicates perfect indistinguishability, while V = 0 denotes fully distinguishable photons. Bearing in mind the actual applications of these sources in quantum networks [5, 73], the interference visibility of remote, entangled sources is particularly relevant. In the past several experiments have been performed using remote QDs [90, 106, 107]. The best remote visibility to date was achieved by Reindl *et al* in 2017 with V = 0.51(5) using GaAs QDs embedded in a planar distributed Bragg reflector (DBR) cavity under phonon-assisted two-photon excitation (see [90]).

3.2. Source brightness

In spite of this progress, a severe limitation towards desired applications remains the difficulties in extracting the entangled photons generated by QDs from the host matrix. However, whereas sources based on parametric down conversion suffer from an intrinsic trade-off between source brightness and entanglement fidelity [108] due to the probabilistic nature of the generation process, the deterministic QD-based sources could, in principle, reach a close-to-unity efficiency without negatively influencing the entanglement fidelity. So far this issue has been mainly addressed from the perspective of single-photon sources.

The efficiency of a single-photon source can be classified by its brightness (*B*), which denotes the probability of collecting a single photon with the first lens of the collection optics upon an excitation pulse. Following a common definition [109] the brightness $B = p_s \eta_{qe} \eta_{ee}$ is composed of the probability that the QD is excited in the desired target state (p_s), the quantum efficiency accounting for losses due to nonradiative recombination processes (η_{qe}) and the extraction efficiency (η_{ee}).

The low efficiency of QD-based sources arises mainly from a small extraction efficiency. To achieve a proper 3D confinement, QDs have to be capped and are therefore embedded in a host matrix like GaAs or AlGaAs with a high refractive index (e.g. $n(GaAs) \approx 3.5$). For planar as-grown samples the extraction efficiency is therefore limited by total internal reflection (TIR), which allows only a small fraction $(1/(4n^2))$ of the emitted light to exit the sample. Taking into account the finite numerical aperture (NA) of the collection optics, generally only less than 2% of the emitted light can be collected from the top surfaces of planar unprocessed samples [110]. Over the years, several paths have been followed in order to improve the brightness of QD-based sources whereby the principle strategies are evident from the definition of the extraction efficiency [111] given by

$$\eta_{\rm ee} = \eta_{\rm ce}\beta = \frac{\eta_{\rm ce}\Gamma_{\rm tm}}{\Gamma_{\rm tm} + \Gamma_{\rm om}} \tag{13}$$

where η_{ce} describes the fraction of the emitted light collectible via the objective lens (with a given NA) and the β -factor accounts for the spontaneous emission (SE) rates in a specific target mode Γ_{tm} or into all other modes Γ_{om} , respectively. Consequently, three main approaches to increase the extraction efficiency were followed: (i) enhancement of the SE rate Γ_{tm} into a (cavity) target mode exploiting the Purcell effect (cavity approach), (ii) inhibition of the emission into modes that cannot be collected Γ_{om} (waveguide approach) or (iii) an increase in η_{ce} e.g. by levering the TIR limitation (geometrical approach). In the following, we want to review several concepts possibly leading to bright entangled photon sources.

3.2.1. Cavity approach: micropillars. The most extensively used approach for increasing the source brightness is to embed the QDs into a planar λ -cavity either defined by DBRs or metal mirror(s) [110]. The mirrors induce a vertical confinement of the light field leading to a local change in the optical mode density. If the OD emission is properly coupled to such a cavity mode a significant acceleration of the SE rate $\Gamma_{\rm tm}$ (quantified by the Purcell factor $F_{\rm p}$) can be achieved [112]. Furthermore, the diminished transition lifetime limits the sensitivity of the QDs to various dephasing mechanisms induced by the interaction with the solid-state environment. Hence, these structures yield excellent values of singlephoton purity, two-photon interference visibility and entanglement fidelity. However, the achieved (pair) extraction efficiencies are still modest (see comparison in [113]) even if additionally furnished with a solid immersion lens [21].

A decisive step towards the development of bright QD single-photon sources exploiting cavity quantum electrodynamics was the establishment of an additional lateral optical confinement by etching micropillars out of planar cavity samples [114]. Initially, the randomness of self-assembled QDs in terms of spatial position and spectral properties had been a major hurdle to fully exploiting the capabilities of these structures as any spectral and spatial mismatch drastically degrades the brightness of the source. The essential breakthrough for this approach was accomplished in 2008 by Dousse et al [115]. In the related work, a novel far-field optical lithography technique has been demonstrated capable of accurate spatial alignment (50 nm accuracy) between QD location and micropillar. The lithography is performed in situ at low temperature and further allows a spectral preselection of the QD and according customization of the pillar dimension to provide optimal spectral matching between QD emission and fundamental cavity mode. Using this technique, extraction efficiencies as high as 79% have been demonstrated [116] whereby additional spectral fine tuning is performed using temperature. This has been achieved while simultaneously preserving excellent values for single-photon purity $g^{(2)}(0) = 0.002 8(12)$ [19] and indistinguishability in terms of (corrected) TPI visibility V = 98.5(1)% [89], both measured under strict resonant excitation (excitation power corresponding to π -pulse) and additional stabilization of the charge environment via electric fields. To date, QDs embedded in micropillar cavities show the best performance in terms of high brightness, high single-photon purity and high photon indistinguishability and hence, are already applied in scalable multi-photon experiments such as boson sampling [117].

However, these structures are less suitable for entangled photon pair generation, as the energy difference between biexciton and exciton (relative binding energy) typically exceeds the width of the cavity resonance. We note here that the relative binding energy of the biexciton strongly depends on the QD structure [118] and that QDs with small binding energy can be found by careful preselection [119] or postgrowth tuning via electric or strain fields [118, 120]. Unless one resorts to the concept of time-reordering [121]—which is limited to non-unity values of the entanglement fidelity [122] -achieving vanishing FSS, binding energy and mode matching at the same time is challenging, however. In addition resonant two-photon excitation becomes impracticable under these conditions. In 2010, Dousse et al [36] presented the concept of a photonic molecule, using two micropillar cavities coupled to a single QD. The authors achieved an entanglement fidelity of $f^+ = 0.68$ at a photon pair extraction efficiency of 12%. The entanglement was essentially limited by the FSS ($<3 \mu eV$) and recapture processes, as pulsed, non-resonant excitation were applied. The two-photon interference visibility or single photon purity were not addressed in this work. However, the authors claim that the fine structure could be tuned via an electrical field and resonant excitation could, in principle, be implemented. Unfortunately, no additional improvements in terms of entanglement on this structure exist (a fact that probably arises from the complexity of its fabrication). However, this is to date the brightest known entangled photon source using a QD emitter [113]. Since it is a challenging task to use narrowband cavity designs for entangled photon pair extraction, we foresee that cavity structures featuring broadband extraction efficiency enhancement and a modest Purcell factor are more suitable for the purpose.

In this context, Sapienza *et al* [123] recently presented a promising alternative approach based on circular Bragg grating 'bullseye' cavities. Hereby, a two-color photoluminescence imaging technique has been used to precisely align the customized cavities to preselected QDs. Sizeable extraction efficiencies of up to 48.5% (0.4NA) over a moderate spectral bandwidth of a few nm have been obtained whereby calculations indicate possible values of up to 80% when using objectives with higher NA. Furthermore, autocorrelation measurements performed under pulsed quasi-resonant excitation yielded an excellent single photon purity of 99.1% ($g^{(2)}(0) < 0.009(5)$).

3.2.2. Waveguiding approach: nanowires. A superior concept for a single photon source which promises extraction efficiencies even over 90% along a 70 nm wide

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spectral range was proposed in 2009 by Friedler et al [124]. Here, a single QD is integrated in a vertical tapered semiconductor nanowire (NW) waveguide, where, instead of using the Purcell effect for enhancement of the spontaneous emission into a cavity mode, the coupling over all modes, except for the fundamental waveguide (WG) mode, is suppressed. The optimal device layout obtained is shown in figure 6 along with the calculated SE rates and extraction efficiencies for variable emitter wavelengths. The first experimental realization of such a device was pioneered by Claudon et al [125] in 2010. In this work, 2.5 μ m high and 200 nm thick NWs were engineered from GaAs wafer hosting (high-density) self-assembled In(Ga)As QDs using a top-down approach for fabrication. Autocorrelation measurements of the saturated exciton transition using a HBT setup revealed a very pure single-photon emission of $g^{(2)}(0) < 0.008$ for pulsed non-resonant excitation. For the presented device design, an excellent extraction efficiency of $\eta_{ee} = 72(9)\%$ (using a 0.75 NA) was obtained. The extraction efficiency was found to be strongly NA-dependent indicating that the far-field emission pattern was not entirely collected. The discrepancy of the obtained source efficiency compared to the theoretically predicted values was attributed to the sensitivity to deviations from the optimal taper angle. This issue has been addressed in a subsequent work dealing with a novel device called a 'photonic trumpet' [126]. The revised design is more tolerant against deviations from the optimal taper angle, provides a higher directionality of the far-field emission and shows an improved extraction efficiency of 75(10)%.

Although the radial alignment of the QD with respect to the NW axis has been found to demand comparable low position accuracy [127], a more accurate alignment between the QD and the photonic structure is required for a further improvement of the source brightness and fabrication yield. This could be either achieved by employing deterministic processing technologies [115] or by using a bottom-up growth approach for the NWs. In the latter case, wurtzite In(As)P QDs in [111]-oriented (tapered) InP NWs in particular have been studied extensively. This approach turned out to be highly promising for a bright source of polarization-entangled photon pairs without the necessity of post-growth engineering of the QD properties or temporal post-selection, a major source of photon losses.

These NWs are actually grown using a combination of selective-area and vapor-liquid-solid (VLS) epitaxy [128]. Hereby, the (111)B InP substrate is initially coated with a SiO₂ film. Circular apertures are created in the oxide at predefined positions by means of electron-beam lithography (EBL) and (isotropic) under-etching of the resist pattern with hydrofluoric (HF) acid. Subsequently, the Au catalysts for VLS growth of the NW core are deposited in the center of the oxide openings and the resist is removed. As VLS growth occurs exclusively at the Au/InP interface, the lateral dimensions of the NW core can be controlled via the hole sizes of the EBL pattern (± 2 nm precision). After growth of the InP NW core containing a thin In(As)P segment, axial growth is suppressed by increasing the



Figure 6. Nanowire waveguide. (a) Color-coded intensity distribution, (b) (normalized) SE rate and (c) extraction efficiency for a QD embedded in a NW WG. Hereby, an optimal coupling to the fundamental WG mode (HE₁₁) has been obtained for a ratio $D/\lambda \approx 0.22$ between the waveguide thickness (D) at the QD position and the emission wavelength (λ) supporting large β -factors > 0.95. To achieve high extraction efficiencies the NW WG features a planar Ag bottom mirror coated with a thin silica dielectric layer to suppress plasmonic effects at the metal/semiconductor interface. Furthermore, a taper with a small angle of around 1.5° towards the NW tip is used to reduce scattering of the guided mode, increasing transmission at the top facet and reducing the beam divergence. The QD is located at the NW axis in an anti-node of the electric field to benefit from constructive interference. The calculations of the spontaneous emission rate (b) have been performed using different theoretical models (see [124]) and the values obtained for the extraction efficiency are presented for two different NAs corresponding to collection angles of $\theta = 60^{\circ}$ (NA 0.87) and $\theta = 45^{\circ}$ (NA 0.7). Reproduced with permission from [124].

temperature. This in turn supports the radial growth of the NW shell which confines the In(As)P QD and defines the WG. As the radial growth of the NW shell is limited to the diameter of the oxide openings [129], both lateral QD and WG dimensions can be controlled independently whereby the QD is exactly aligned on the NW axis. This enables an optimal device design and coupling between QD dipole and guided mode ensuring directional emission and efficient light extraction.

In 2012, Reimer *et al* [130] presented a single-photon source based on a bottom-up grown InP NW with embedded In(As)P QDs. As growth on a metallic mirror is not possible, the NWs were harvested using a flexible and fully transparent polydimethylsiloxane (PDMS) polymer film followed by evaporation of the Au-mirror at the bottom side of the NWs. To evaluate the extraction efficiency, photoluminescence studies under pulsed non-resonant excitation were performed. At saturation an extraction efficiency of 42% was obtained for a 0.75 NA, whereby the moderate value was mainly attributed to a low modal reflectivity of the Au-mirror of only 30%.

Although growth was performed on a (111) substrate an FSS of around 30 μ eV was observed for the measured QDs. However, subsequent improvements in the crystal quality of the NWs [131] and optimization of the QD dimensions finally

allowed the demonstration of polarization-entangled photon pairs in 2014 by Huber et al [132] and Versteegh et al [133]. In the latter work, half of the measured QDs had a very low FSS of $<2 \mu eV$. Quantum state tomography revealed a fidelity to the entangled state $(|JJ\rangle + |WW\rangle)/\sqrt{2}$ (J and W are orthogonal elliptic polarizations) of f = 0.76(2) for the full time window of 6.02 ns. Applying temporal postselection the value could be improved to a fidelity of f = 0.85(6) for the narrowest time window of 0.13 ns. The observation of the $(|JJ\rangle + |WW\rangle)/\sqrt{2}$ two-photon state instead of the common, maximally entangled $(|HH\rangle + |VV\rangle)/\sqrt{2}$ has been attributed to NW anisotropy (elongated cross-section) formed during shell growth (not related to QD shape) giving rise to birefringence in the NW WG, rotating the polarization state of the emitted photon pairs. Using a quasi-resonant excitation scheme Jöns et al [134] improved the fidelity further, up to $f^+ = 0.82(2)$ ($f^+ = 0.85(9)$), without (with) temporal postselection using time windows of 4.48 ns (0.13 ns).

So far, bottom-up grown NW WG with embedded QDs represent the best compromise between source brightness and achieved entanglement fidelity. The weak-point of the NW approach remains the limited visibility in TPI experiments [135] despite sub-Kelvin temperatures, an issue which may be solved by resonant excitation.



Figure 7. Quantum dot microlenses. (a) An illustration of a single QD grown above a bottom DBR and deterministically embedded in a monolithic microlens allowing for an enhanced photon extraction efficiency. Reproduced from [119]. CC BY 4.0. (b) Comparison of the numerically calculated extraction efficiency versus the numerical aperture (NA) of the collection optics for the use of a bottom DBR (red) or a gold mirror (black). In both cases a hemispheric-section lens shape has been assumed but the structure dimensions have been optimized for the respective mirror (black: height 400 nm, base width 2 μ m, gold mirror 150 nm below the QD; red: height 400 nm, base width 2.4 μ m with a DBR 65 nm below the QD). For large NAs the calculations reveal that the design with the Au-mirror outperforms the DBR as a consequence of the DBR's finite acceptance angle of $<20^{\circ}$ whereas the reflectivity of the Au-mirror is almost independent of the angle of incidence. Reproduced from [136]. CC BY 4.0.

3.2.3. Geometrical approach: microlenses. A geometrical approach to increase the TIR-limited extraction efficiency by shaping the sample surface at the emitter position was reported by Gschrey et al [136] in 2015. In the presented work, monolithic microlenses have been processed deterministically onto self-assembled In(Ga)As QDs grown above a 23-pair Al_{0.9}Ga_{0.1}As/GaAs DBR (see figure 7(a)). Spectral preselection of the QDs as well as accurate positioning and dimensioning of the microlenses have been achieved using a novel technique called low-temperature cathodoluminescence lithography (CLL) [137], which represents a combination of cathodoluminescence (CL) spectroscopy and in situ 3D EBL, both performed at liquid He temperature. Due to the weak directionality of the emission, the extraction efficiency achievable with microlenses strongly depends on the NA of the used collection system (see figure 7(d)). Until now, only moderate values of up to $\eta_{ee} = 29(3)\%$ [138] could be obtained using this device concept whereby most reports refer to a relatively low NA = 0.4. However, improvement of the extraction efficiency has been achieved while maintain ing ideal values of single-photon purity ($g^{(2)}(0) \leq 0.01$). Furthermore, two-photon interference experiments yielded competitive visibility values of up to V = 94(6)% [139] using a quasi-resonant two-pulse excitation scheme with a pulse separation of 2 ns. Both values indicate that the lens processing has no adverse effect on the quantum properties (i.e single photon purity and indistinguishability) of light emitted by the embedded QD. The enabled spectral preselection of QDs also facilitates the application in experiments with spatially separated sources [107].

Various paths have been followed to achieve a further increase in the extraction efficiency using the microlens approach. Recently, a corresponding value of $\eta = 40(4)\%$ has been reported [140] for the remarkable combination of a deterministic QD microlens with a 3D-printed multi-lens micro-objective [141] (with an NA ≈ 0.7): a concept which is especially interesting for coupling to low-NA fibers. For high-NA collection systems, anti-reflection coatings [142] and in particular the use of an Au-mirror instead of the bottom DBR [143] promise a significant increase in the practically achievable extraction efficiency (see figure 7(d)). In the latter case, the reported application of a flip-chip process easily enables integration with existing strain-tuning technology [70] allowing for post-growth engineering of various QD properties [144]: in particular a cancellation of the FSS necessary for the generation of strongly entangled photon pairs. In this regard, although a demonstration of entangled photon pair emission is still pending, microlenses provide a highly promising alternative due to their broadband enhancement with a bandwidth of about 50 nm [145], technological compatibility [146] and flexibility with respect to application to various QD systems.

3.3. Sources at the telecommunication wavelengths

Sources compatible with the existing fiber-based photonic infrastructure would strongly facilitate the widespread use of quantum communication. Therefore, it is important to develop entangled photon sources operating within the low loss regime of silica fibers. The ongoing progress in fabrication of QD-based entanglement-emitters matching the telecom wavelength (1260-1625 nm) started in 2005 with the first demonstrations of QD light emission in the C-band by Miyazawa et al using InAs/InP QDs [147] and in the O-Band by Alloing et al with InAs/GaAs QDs [148]. In the following, we summarize the results of some recent works. In 2016, Miyazawa showed high single-photon purity $(g^{(2)}(0) =$ $4.4(2) \times 10^{-4}$) for InAs/InP QDs emitting at 1.5 μ m under quasi-resonant excitation [149]. Further, in 2016, Kim et al studied two-photon interference with an InAs/InP QD embedded in a nanophotonic cavity. The resulting visibility was 18% (67%) without (with) post-selection at a pulse separation of 5 ns under non-resonant excitation [150]. Although the visibility is significantly lower than reported for non-telecom wavelength QDs [19, 89], the authors achieved an impressive outcoupling efficiency >36%, which is among the highest ever observed at this emission wavelength. The fabrication of entanglement-ready QDs at telecom wavelength is a challenging task as non-zero FSS values are not easy to achieve. In principle, InAs/InP QDs are proposed to have an intrinsic FSS one order of magnitude lower than standard InGa/GaAs QDs [151]. In 2014, Liu et al showed that InAs/ InAlAs droplet QDs grown on (111)A surfaces combine the advantages of a broad emission spectra covering the O, C and L telecom bands with small FSS $<4 \mu eV$ (on average 25 μ eV), but the authors showed no entanglement measurements [152]. In 2017, Olbrich et al presented QDs emitting in the C-Band where a fraction of one third has an FSS below 5 μ eV [153]. The authors used InAs quantum dots embedded in InGaAs barriers and shifted the emission by inserting an additional InGaAs metamorphic buffer. Entanglement measurements revealed a fidelity of 0.61(7) at an FSS of approximately 6.20(7) µeV. In 2017, Skiba-Szymanska et al presented a different growth strategy to realize low FSS QDs -emitting in the telecom C-band-by droplet epitaxy of InAs on (001) InP substrate [154] using metalorganic vapor phase epitaxy. The authors claim a mean FSS 4x smaller than for Stranski-Krastanow QDs. Furthermore, in 2018, Müller et al presented a quantum light-emitting diode based on InP QDs emitting at 1550 nm [155]. The authors showed a peak entanglement fidelity of 0.87(4) by measuring its time evolution (the time evolution of the fidelity is indicated in figure 2 (d)). Further, the authors observed entangled photon generation up to device temperatures of 93 K, which is an advantage compared to standard InAs/GaAs and GaAs/ AlGaAs QDs typically working at temperatures < 10 K. In 2017, Huwer et al presented a quantum relay based on a QD emitting in the O-band combined with an O-band diode laser for encoding the polarization states (for details on the relay scheme we refer the interested reader to [34]). The used QD has an FSS of 9.05(1) μ eV and gives a peak fidelity of 0.920 (2) (0.963(3)) to a maximally entangled Bell state (the exact time-evolving state), which is the highest value ever presented under these conditions at telecom wavelength. The quantum relay itself achieves an overall fidelity of 0.95(2). In 2018, Höfer et al integrated InGaAs/GaAs quantum dots emitting at the telecom O-band on an uniaxial piezoelectric actuator. This allowed the authors to tune the FSS of the QDs-with an average value of $10(6) \mu eV$ —to values below the setup resolution ($\langle 2 \mu eV \rangle$ [156]. However, the authors did not report on the achievable entanglement. Considering the rapid progress in telecom wavelength QDs, we believe that the combination of these dots with tuning techniques presented in section 2.2 may soon enable the realization of highly entangled photon sources compatible with existing fiber networks.

4. Outlook

Polarization-entangled photon pairs are now seen as the fundamental building block in quantum information sciences and in particular for quantum communication. It was long believed that there is no real application of quantum entanglement to be harnessed, alongside fundamental research, until the first proposals [157, 158] to prevent eavesdropping

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of information transfer utilizing Bell's theorem. From this point, an explosion of technological concepts emerged to distribute entangled photons used as message encrypting keys. The basic underlying mechanism is the already mentioned quantum teleportation [159], providing an instantaneous transfer of quantum information from one location to another without violating the no-cloning theorem [160]. It ultimately allows relocation of entangled photon pairs at arbitrary distances in terms of the teleportation of mixed states involving independent entanglement resources, also known as entanglement swapping, and requires the efficient use of quantum repeater schemes [161–165]. In view of such quantum relays exploiting QD polarization entanglement, experimental realizations are limited at present but are promising. Preliminary investigations of the teleportation of a single photon polarization state was implemented on an entangled light-emitting diode [166], and consequently expanded within a laser-heralded teleportation protocol [58]. Remarkably, such a protocol relying on laser generated input states was also carried out at telecom wavelength [58], making it potentially compatible with the existing global communication infrastructure, thereby supplying the necessary scalability. A major hurdle to be considered in this context is the high vulnerability of single photon polarization states to environmental dephasing effects, particularly pronounced in long-distance networks. A robust and feasible solution to account for the polarization degradation was established on QDs through time-bin entanglement [167]. Here, the QD XX-X cascade is excited in a coherent superposition of two laser pulses arriving subsequently in time. The photons then pass an equally unbalanced interferometer with respect to the excitation pulses. The information regarding the creation time is canceled and results in an entangled state encoded in the respective time-bins. Indeed, it is of practical advantage to have polarization entanglement to hand at any time in order to perform unitary transformations, as required for complex communication networks, and to solely use timebin entanglement for faithful, decoherence-free propagation between quantum relay nodes. In fact, polarization-time-bin interfaces allowing for arbitrary conversion among the different quantum superpositions without loss of entanglement fidelity are possible for QDs [168] by creating time-bins: but only after entangled photon pair emission, based on unbalanced, polarization-selective Mach-Zehnder interferometers. In summary, QDs as sources of polarization-entangled photon pairs are on the verge of revealing their full potential with respect to actual, practical applications. Once all individually achieved results on QDs are finally merged into a single, semiconductor-based entangled photon pair source, it will pave the way towards sophisticated and fully deterministic quantum networks.

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References

- [1] Einstein A, Podolsky B and Rosen N 1935 Phys. Rev. 47 777
- [2] Schrödinger E 1935 Math. Proc. Camb. Phil. Soc. 31 555
- [3] Zeilinger A 2017 Phys. Scr 92 072501
- [4] Gisin N, Ribordy G, Tittel W and Zbinden H 2002 *Rev. Mod. Phys.* 74 145
- [5] Kimble H J 2008 Nature 453 1023
- [6] Sangouard N, Simon C, de Riedmatten H and Gisin N 2011 *Rev. Mod. Phys.* 83 33
- [7] Pirandola S, Laurenza R, Ottaviani C and Banchi L 2017 Nat. Commun. 8 15043
- [8] Duan L-M, Lukin M D, Cirac J I and Zoller P 2001 Nature 414 413
- [9] Simon C, de Riedmatten H, Afzelius M, Sangouard N, Zbinden H and Gisin N 2007 Phys. Rev. Lett. 98 190503
- [10] Briegel H-J, Dür W, Cirac J I and Zoller P 1998 Phys. Rev. Lett. 81 5932
- [11] Kok P, Williams C P and Dowling J P 2003 Phys. Rev. A 68 022301
- [12] Munro W J, Stephens A M, Devitt S J, Harrison K A and Nemoto K 2012 Nat. Photon. 3 777
- [13] Fattal D, Diamanti E, Inoue K and Yamamoto Y 2004 Phys. Rev. Lett. 92 037904
- [14] Lu C-Y and Pan J-W 2014 Nat. Photon. 8 174
- [15] Orieux A, Versteegh M A M, Jöns K D and Ducci S 2017 *Rep. Prog. Phys.* 80 076001
- [16] Kwiat P G, Mattle K, Weinfurter H, Zeilinger A,
- Sergienko A V and Shih Y 1995 *Phys. Rev. Lett.* **75** 4337
 [17] Scarani V, De Riedmatten H, Marcikic I, Zbinden H and Gisin N 2005 *Eur. Phys. J.* D **32** 129
- [18] Benson O, Santori C, Pelton M and Yamamoto Y 2000 *Phys. Rev. Lett.* 84 2513
- [19] Somaschi N et al 2016 Nat. Photon. 10 340
- [20] Fognini A et al 2017 Preprint: arXiv:1710.10815
- [21] Huber D, Reindl M, Covre da Silva S F, Schimpf C, Martín-sánchez J, Piredda G, Edlinger J, Rastelli A and Trotta R 2018 Preprint: arXiv:1801.066557
- [22] James D F V, Kwiat P G, Munro W J and White A G 2001 *Phys. Rev.* A 64 052312
- [23] Huber D, Reindl M, Huo Y, Huang H, Wildmann J S, Schmidt O G, Rastelli A and Trotta R 2017 Nat. Commun. 8 15506
- [24] Keil R, Zopf M, Chen Y, Höfer B, Zhang J, Ding F and Schmidt O G 2017 Nat. Commun. 8 15501
- [25] Bayer M et al 2002 Phys. Rev. B 65 195315
- [26] Plumhof J D, Trotta R, Rastelli A and Schmidt O G 2012 Nanoscale Res. Lett. 7 336

- [27] Stevenson R M, Hudson A J, Bennett A J, Young R J, Nicoll C A, Ritchie D A and Shields A J 2008 *Phys. Rev. Lett.* **101** 170501
- [28] Santori C, Fattal D, Pelton M, Solomon G S and Yamamoto Y 2002 Phys. Rev. B 66 045308
- [29] Akopian N, Lindner N H, Poem E, Berlatzky Y, Avron J, Gershoni D, Gerardot B D and Petroff P M 2006 *Phys. Rev. Lett.* 96 130501
- [30] Young R J, Stevenson M, Atkinson P, Cooper K, Ritchie D A and Shields A J 2006 New J. Phys. 8 29
- [31] Huber T, Predojevic A, Khoshnegar M, Dalacu D, Poole P J, Majedi H and Weihs G 2014 Nano Lett. 14 7107
- [32] Ward M B, Dean M C, Stevenson R M, Bennett A J, Ellis D J P, Cooper K, Farrer I, Nicoll C A, Ritchie D A and Shields A J 2014 Nat. Commun. 5 3316
- [33] Salter C L, Stevenson R M, Farrer I, Nicoll C A, Ritchie D A and Shields A J 2010 Nature 465 594–7
- [34] Huwer J, Stevenson R M, Skiba-Szymanska J, Ward M B, Shields A J, Felle M, Farrer I, Ritchie D A and Penty R V 2017 Phys. Rev. Appl. 8 024007
- [35] Trotta R, Wildmann J S, Zallo E, Schmidt O G and Rastelli A 2014 Nano Lett. 14 3439–44
- [36] Dousse A et al 2010 Nature 466 217
- [37] Kuroda T et al 2013 Phys. Rev. B 88 041306
- [38] Lin C-H, You W-T, Chou H-Y, Cheng S-J, Lin S-D and Chang W-H 2011 *Phys. Rev.* B **83** 075317
- [39] Plumhof J D, Trotta R, Křápek V, Zallo E, Atkinson P, Kumar S, Rastelli A and Schmidt O G 2013 *Phys. Rev.* B 87 075311
- [40] Liao Y-H, Liao C-C, Ku C-H, Chang Y-C, Cheng S-J, Jo M, Kuroda T, Mano T, Abbarchi M and Sakoda K 2012 *Phys. Rev.* B 86 115323
- [41] Michler P 2009 Single Semiconductor Quantum Dots (Berlin: Springer) 227–65
- [42] Chekhovich E A, Krysa A B, Hopkinson M, Senellart P, Lemaitre A, Skolnick M S and Tartakovskii A I 2013 Nat. Phys. 9 74–8
- [43] Hudson A J, Stevenson R M, Bennett A J, Young R J, Nicoll C A, Atkinson P, Cooper K, Ritchie D A and Shields A J 2007 Phys. Rev. Lett. 99 266802
- [44] Urbaszek B, Marie X, Amand T, Krebs O, Voisin P, Maletinsky P, Högele A and Imamoglu A 2013 *Rev. Mod. Phys.* 85 79
- [45] Langbein W, Borri P, Woggon U, Stavarache V, Reuter D and Wieck A D 2004 Phys. Rev. B 70 033301
- [46] Heyn C 2001 Phys. Rev. B 64 165306
- [47] Belhadj T et al 2010 Appl. Phys. Lett 97 051111
- [48] Gong M, Hofer B, Zallo E, Trotta R, Luo J-W, Schmidt O G and Zhang C 2014 *Phys. Rev.* B 89 205312
- [49] Juska G, Dimastrodonato V, Mereni L O, Gocalinska A and Pelucchi E 2013 Nat. Photon. 7 527–31
- [50] Treu J, Schneider C, Huggenberger A, Braun T, Reitzenstein S, Höfling S and Kamp M 2012 Appl. Phys. Lett. 1011 22102
- [51] Juska G, Murray E, Dimastrodonato V, Chung T H, Moroni S T, Gocalinska A, Pelucchi E and Murray E 2015 *Appl. Phys. Lett.* **117** 134302
- [52] Chung T H, Juska G, Moroni S T, Pescaglini A, Gocalinska A and Pelucchi E 2016 Nat. Photon. 10 782–7
- [53] Musiał A, Gold P, Andrzejewski J, Löffler A, Misiewicz J, Höfling S, Forchel A, Kamp M, Sek G and Reitzenstein S 2014 Phys. Rev. B 90 045430
- [54] Furis M, Htoon H, Petruska M A, Klimov V I, Barrick T and Crooker S A 2006 Phys. Rev. B 73 241313
- [55] An J M, Franceschetti A and Zunger A 2007 Nano Lett. 7 2129
- [56] Kindel C, Kako S, Kawano T, Oishi H, Arakawa Y, Hönig G, Winkelnkemper M, Schliwa A, Hoffmann A and Bimberg D 2010 Phys. Rev. B 81 241309

- [57] Welander E, Hildmann J and Burkard G 2015 Preprint: arXiv:1409.6521
- [58] Stevenson R M, Salter C L, Boyer De La Giroday A, Farrer I A, Nicoll C A, Ritchie D A and Shields A J 2011 Preprint: arXiv:1103.2969
- [59] Mano T, Watanabe K, Tsukamoto S, Koguchi N and Gotoh Y 2000 Japan. J. Appl. Phys. 39 79
- [60] Sanguinetti S, Watanabe K, Tateno T, Gurioli M, Werner P, Wakaki M, Koguchi N and Schieber M 2003 J. Cryst. Growth 253 71
- [61] Mano T, Kuroda T, Yamagiwa M, Kido G, Sakoda K and Koguchi N 2006 Appl. Phys. Lett. 89 183102
- [62] Heyn C, Stemmann A, Köppen T, Strelow C, Kipp T, Grave M, Mendach S and Hansen W 2009 Appl. Phys. Lett. 94 18
- [63] Basso Basset F, Bietti S, Reindl M, Esposito L, Fedorov A, Huber D, Rastelli A, Bonera E, Trotta R and Sanguinetti S 2018 Nano Lett. 18 505
- [64] Wang Z M, Liang B L, Sablon K A and Salamo G J 2007 Appl. Phys. Lett. 90 113120
- [65] Huo Y H, Rastelli A and Schmidt O G 2013 Appl. Phys. Lett. 102 152105
- [66] Huo Y H et al 2014 Nat. Phys. 10 46-51
- [67] Huo Y H, Křápek V, Schmidt O G and Rastelli A 2017 Phys. Rev. B 95 165304
- [68] Huo Y H, Křápek V, Rastelli A and Schmidt O G 2014 Phys. Rev. B 90 041304
- [69] Chau H F 2002 Phys. Rev. A 66 060302
- [70] Martín-Sánchez J et al 2018 Semicond. Sci. Technol. 33 013001
- [71] Trotta R, Zallo E, Ortix C, Atkinson P, Plumhof J D, van den Brink J, Rastelli A and Schmidt O G 2012 *Phys. Rev. Lett.* **109** 147401
- [72] Trotta R, Martín-Sánchez J, Daruka I, Ortix C and Rastelli A 2015 Phys. Rev. Lett. 114 150502
- [73] Northup T and Blatt R 2014 Nat. Photon. 8 356
- [74] Wang J, Gong M, Guo G-C and He L 2015 Phys. Rev. Lett. 115 067401
- [75] Wang J, Gong M, Guo G-C and He L 2012 Appl. Phys. Lett. 101 063114
- [76] Trotta R, Martín-Sánchez J, Schimpf C, Zallo E, Stroj S, Edlinger J and Rastelli A 2016 Nat. Commun. 7 10375
- [77] Martín-sánchez J et al 2016 Adv. Opt. Mater. 4 682
- [78] Akopian N, Wang L, Rastelli A, Schmidt O G and Zwillier V 2011 Nat. Photon. 5 230
- [79] Chen Y, Zhang J, Zopf M, Jung K, Zhang Y, Keil R, Ding F and Schmidt O G 2016 Nat. Commun. 7 10387
- [80] Zhang J, Zallo E, Höfer B, Chen Y, Keil R, Zopf M, Böttner S, Ding F and Schmidt O G 2017 Nano Lett. 17 501
- [81] Zhang J, Wildmann J S, Ding F, Trotta R, Huo Y, Zallo E, Huber D, Rastelli A and Schmidt O G 2015 Nat. Commun. 6 10067
- [82] Pooley M A, Bennett A J, Stevenson R M, Shields A J, Farrer I and Ritchie D A 2014 Phys. Rev. Appl. 1 024002
- [83] Jayakumar H, Predojević A, Huber T, Kauten T, Solomon G S and Weihs G 2013 Phys. Rev. Lett. 110 135505
- [84] Muller M, Bounouar S, Jöns K D, Glässl M and Michler P 2014 Nat. Photon. 8 224
- [85] Huber T, Predojević A, Föger D, Solomon G and Weihs G 2015 New J. Phys 17 123025
- [86] Kiršanskė G et al 2017 Phys. Rev. B 96 165306
- [87] Stufler S, Machnikowski P, Ester P, Bichler M, Axt V M,
- Kuhn T and Zrenner A 2006 *Phys. Rev.* B **73** 125304 [88] Brunner K, Abstreiter G, Böhm G, Tränkle G and
- Weimann G 1994 Phys. Rev. Lett. 73 1138
- [89] Ding X et al 2016 Phys. Rev. Lett. **116** 020401
- [90] Reindl M, Jöns K D, Huber D, Schimpf C, Huo Y, Zwiller V, Rastelli A and Trotta R 2017 Nano Lett. 17 4090

- [91] Bounouar S, Müller M, Barth A M, Glässl M, Axt V M and Michler P 2015 Phys. Rev. B 91 161302
- [92] Debnath A, Meier C, Chatel B and Amand T 2013 *Phys. Rev.* B 88 201305
- [93] Wu Y, Piper I M, Ediger M, Brereton P, Schmidgall E R, Eastham P R, Hugues M, Hopkinson M and Phillips R T 2011 Phys. Rev. Lett. 106 067401
- [94] Glässl M, Barth A M, Gawarecki K, Machnikowski P, Croitoru M D, Lüker S, Reiter D E, Kuhn T and Axt V M 2013 Phys. Rev. B 87 085303
- [95] Glässl M, Barth A M and Axt V M 2013 Phys. Rev. Lett. 110 147401
- [96] Förstner J, Weber C, Danckwerts J and Knorr A 2003 Phys. Rev. Lett. 91 127401
- [97] Ramsay A J, Gopal A V, Gauger E M, Nazir A, Lovett B W, Fox A M and Skolnick M S 2010 Phys. Rev. Lett. 104 017402
- [98] Ardelt P-L et al 2014 Phys. Rev. B 90 241404
- [99] Quilter J H, Brash A J, Liu F, Glässl M, Barth A M, Axt V M, Ramsay A J, Skolnick M S and Fox A M 2015 Phys. Rev. Lett. 114 137401
- [100] Bounouar S, Müller M, Barth A M, Glässl M, Axt V M and Michler P 2015 Phys. Rev. B 91 161302
- [101] Stevenson R M, Salter C L, Nilsson J, Bennett A J, Ward M B, Farrer I, Ritchie D A and Shields A J 2012 Phys. Rev. Lett. 108 040503
- [102] Yuan Z, Kardynal B E, Stevenson R M, Shields A J, Lobo C J, Cooper K, Beattie N S, Ritchie D A and Pepper M 2002 Science 295 102
- [103] Shalm L K et al 2015 Phys. Rev. Lett. 115 250402
- [104] Hong C K, Ou Z Y and Mandel L 1987 *Phys. Rev. Lett.*59 2044
- [105] Gazzano O and Solomon G S 2016 J. Opt. Soc. Am. B 33 C160
- [106] Gold P, Thoma A, Maier S, Reitzenstein S, Schneider C, Höfling S and Kamp M 2014 Phys. Rev. B 89 035313
- [107] Thoma A, Schnauber P, Böhm J, Gschrey M, Schulze J-H, Strittmatter A, Rodt S, Heindel T and Reitzenstein S 2017 *Appl. Phys. Lett.* **110** 011104
- [108] Wang X-L et al 2016 Phys. Rev. Let 117 210502
- [109] Senellart P, Solomon G and White A 2017 Nat. Nanotechnol. 12 1026
- [110] Barnes W L, Björk G, Gérard J M, Jonsson P, Wasey J A E, Worthing P T and Zwiller V 2002 Eur. Phys. J. D 18 197–210
- [111] Gazzano O and Solomon G S 2016 JOSA B 33 C160
- [112] Purcell E M 1946 Phys. Rev. 69 681
- [113] Jöns K D, Schweickert L, Versteegh M A M, Dalacu D, Poole P J, Gulinatti A, Giudice A, Zwiller V and Reimer M E 2017 Sci. Rep. 7 1700
- [114] Gérard J, Sermage B, Gayral B, Legrand B, Costard E and Thierry-Mieg V 1998 Phys. Rev. Lett. 81 1110
- [115] Dousse A, Lanco L, Suffczyński J, Semenova E, Miard A, Lemaître A, Sagnes I, Roblin C, Bloch J and Senellart P 2008 Phys. Rev. Lett. 101 267404
- [116] Gazzano O, De Vasconcellos S M, Arnold C, Nowak A, Galopin E, Sagnes I, Lanco L, Lemaître A and Senellart P 2013 Nat. Commun. 4 1425
- [117] Loredo J, Broome M, Hilaire P, Gazzano O, Sagnes I, Lemaitre A, Almeida M, Senellart P and White A 2017 *Phys. Rev. Lett.* **118** 130503
- [118] Trotta R, Zallo E, Magerl E, Schmidt O G and Rastelli A 2013 Phys. Rev. B 88 155312
- [119] Heindel T et al 2017 Nat. Commun. 8 14870
- [120] Ding F et al 2010 Phys. Rev. Lett. 104 067405
- [121] Avron J E, Bisker G, Gershoni D, Lindner N H, Meirom E A and Warburton R J 2008 Phys. Rev. Lett. 100 120501
- [122] Troiani F and Tejedor C 2008 Phys. Rev. B 78 155305

- [123] Sapienza L, Davanço M, Badolato A and Srinivasan K 2015 Nat. Commun. 6 7833
- [124] Friedler I, Sauvan C, Hugonin J-P, Lalanne P, Claudon J and Gérard J-M 2009 Opt. Express 17 2095
- [125] Claudon J, Bleuse J, Malik N S, Bazin M, Jaffrennou P, Gregersen N, Sauvan C, Lalanne P and Gérard J-M 2010 *Nat. Photon.* 4 174–7
- [126] Munsch M, Malik N S, Dupuy E, Delga A, Bleuse J, Gérard J-M, Claudon J, Gregersen N and Mørk J 2013 Phys. Rev. Lett. 110 177402
- [127] Bleuse J, Claudon J, Creasey M, Malik N S, Gérard J-M, Maksymov I, Hugonin J-P and Lalanne P 2011 *Phys. Rev. Lett.* **106** 103601
- [128] Dalacu D, Mnaymneh K, Wu X, Lapointe J, Aers G C, Poole P J and Williams R L 2011 Appl. Phys. Lett. 98 251101
- [129] Dalacu D, Kam A, Austing D G, Wu X, Lapointe J, Aers G C and Poole P J 2009 Nanotechnology 20 395602
- [130] Reimer M E, Bulgarini G, Akopian N, Hocevar M, Bavinck M B, Verheijen M A, Bakkers E P, Kouwenhoven L P and Zwiller V 2012 Nat. Commun. 3 737
- [131] Dalacu D, Mnaymneh K, Lapointe J, Wu X, Poole P J, Bulgarini G, Zwiller V and Reimer M E 2012 Nano Lett.
- 12 5919
 [132] Huber T, Predojevic A, Khoshnegar M, Dalacu D, Poole P J, Majedi H and Weihs G 2014 Nano Lett. 14 7107
- [133] Versteegh M A, Reimer M E, Jöns K D, Dalacu D, Poole P J, Gulinatti A, Giudice A and Zwiller V 2014 Nat. Commun. 5 5298
- [134] Jöns K D, Schweickert L, Versteegh M A, Dalacu D, Poole P J, Gulinatti A, Giudice A, Zwiller V and Reimer M E 2017 Sci. Rep. 7 1700
- [135] Reimer M E et al 2016 Phys. Rev. B 93 195316
- [136] Gschrey M et al 2015 Nat. Commun. 6 7662
- [137] Gschrey M, Gericke F, Schüßler A, Schmidt R, Schulze J-H, Heindel T, Rodt S, Strittmatter A and Reitzenstein S 2013 *Appl. Phys. Lett.* **102** 251113
- [138] Schlehahn A et al 2015 Appl. Phys. Lett. 107 041105
- [139] Thoma A et al 2016 Phys. Rev. Lett. 116 033601
- [140] Fischbach S et al 2017 ACS Photonics 4 1327
- [141] Gissibl T, Thiele S, Herkommer A and Giessen H 2016 Nat. Photon. 10 554
- [142] Schnauber P et al 2015 Technologies 4 1
- [143] Fischbach S, Kaganskiy A, Tauscher E B Y, Gericke F, Thoma A, Schmidt R, Strittmatter A, Heindel T, Rodt S and Reitzenstein S 2017 Appl. Phys. Lett. 111 011106
- [144] Aberl J, Klenovsky P, Wildmann J S, Martín-Sánchez J, Fromherz T, Zallo E, Humlíček J, Rastelli A and Trotta R 2017 Phys. Rev. B 96 045414
- [145] Jakubczyk T et al 2016 ACS Photonics 3 2461
- [146] Schlehahn A, Schmidt R, Hopfmann C, Schulze J-H, Strittmatter A, Heindel T, Gantz L, Schmidgall E,

Gershoni D and Reitzenstein S 2016 Appl. Phys. Lett. 108 021104

- [147] Miyazawa T, Takemoto K, Sakuma Y, Hirose S, Usuki T, Yokoyama N, Takatsu M and Arakawa Y 2005 Japan. J. Appl. Phys. 44 L620
- [148] Alloing B, Zinoni C, Zwiller V, Li L H, Monat C, Gobet M, Buchs G, Fiore A, Pelucchi E and Kapon E 2005 Appl. Phys. Lett. 86 101908
- [149] Miyazawa T et al 2016 Appl. Phys. Lett. 109 132106
- [150] Kim J-H, Cai T, Richardson C J K, Leavitt R P and Waks E 2016 Optica 3 577
- [151] He L, Gong M, Li C-F, Guo G-C and Zunger A 2008 Phys. Rev. Lett. 101 157405
- [152] Liu X, Ha N, Nakajima H, Mano T, Kuroda T, Urbaszek B, Kumano H, Suemune I, Sakuma Y and Sakoda K 2014 *Phys. Rev.* B **90** 081301
- [153] Olbrich F, Höschele J, Müller M, Kettler J, Portalupi S L, Paul M, Jetter M and Michler P 2017 Appl. Phys. Lett. 111 133106
- [154] Skiba-Szymanska J et al 2017 Phys. Rev. Appl. 8 014013
- [155] Müller T, Skiba-Szymanska J, Krysa A B, Huwer J, Felle M, Anderson M, Stevenson R M, Heffernan J, Ritchie D A and Shields A J 2018 Nat. Commun. 9 862
- [156] Höfer B, Olbrich F, Kettler J, Paul M, Höschele J, Jetter M, Portalupi S L, Ding F, Michler P and Schmidt O G 2018 Preprint: arXiv:1802.03660
- [157] Bennett C H and Brassard G 1984 Proc. IEEE Int. Conf. on Computers, Systems and Signal Processing 9, p 175
- [158] Ekert A K 1991 Phys. Rev. Lett. 67 661
- [159] Bennett C H, Brassard G, Crepeau C, Jozsa R, Peres A and Wootters W K 1993 Phys. Rev. Lett. 70 1895
- [160] Wootters W K and Zurek W H 1982 Nature 299 802
- [161] Briegel H, Dür W, Cirac J I and Zoller P 1998 Phys. Rev. Lett. 81 5932
- [162] Duan L, Lukin M D, Cirac J I and Zoller P 2001 Nature 414 413
- [163] Riedmatten H D, Marcikic I, Tittel W, Zbinden H, Collins D and Gisin N 2004 Phys. Rev. Lett. 92 047904
- [164] Childress L, Taylor J M, Sørensen A S and Lukin M D 2006 Phys. Rev. Lett. 96 070504
- [165] Simon C, Riedmatten H D, Afzelius M, Sangouard N, Zbinden H and Gisin N 2007 Phys. Rev. Lett. 98 190503
- [166] Nilsson J, Stevenson R M, Chan K H A, Lucamarini M, Ward M B, Bennett A J, Salter C L, Farrer I, Ritchie D A and Shields A J 2013 Nat. Photon. 7 311
- [167] Jayakumar H, Predojevic A, Kauten T, Huber T, Solomon S G and Weihs G 2014 Nat. Commun. 5 4251
- [168] Versteegh M A M, Reimer M E, Berg A A V D, Juska G, Dimastrodonato V, Gocalinska A, Pelucchi E and Zwiller V Phys. Rev. Appl. 92 033802