

Lecture 6 – 26/03/2025

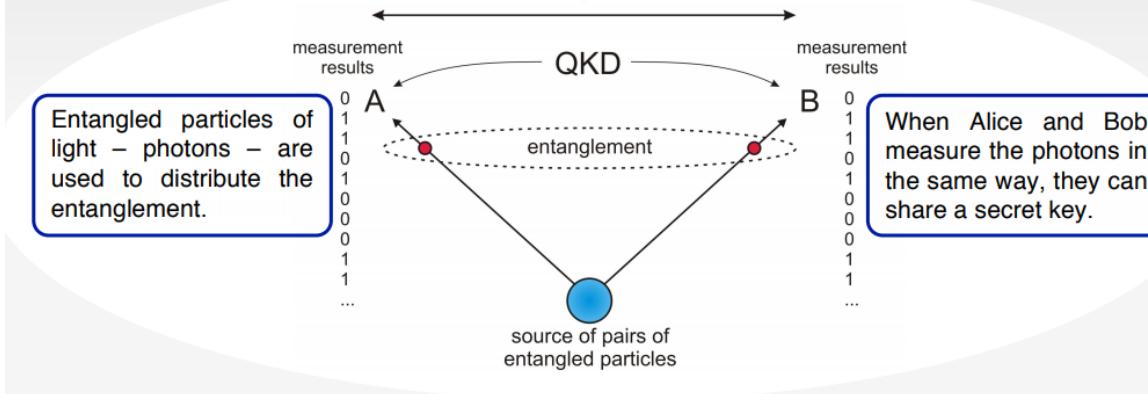
Quantum dots used as single photon emitters (SPEs)

- Role of QD electronic structure for entangled photon sources
- Second-order correlation function ($g^{(2)}(\tau)$)

Entangled photon sources for quantum communication

Quantum Key Distribution (QKD)

The Distribution of "shared" private and secret randomness – a secret key!

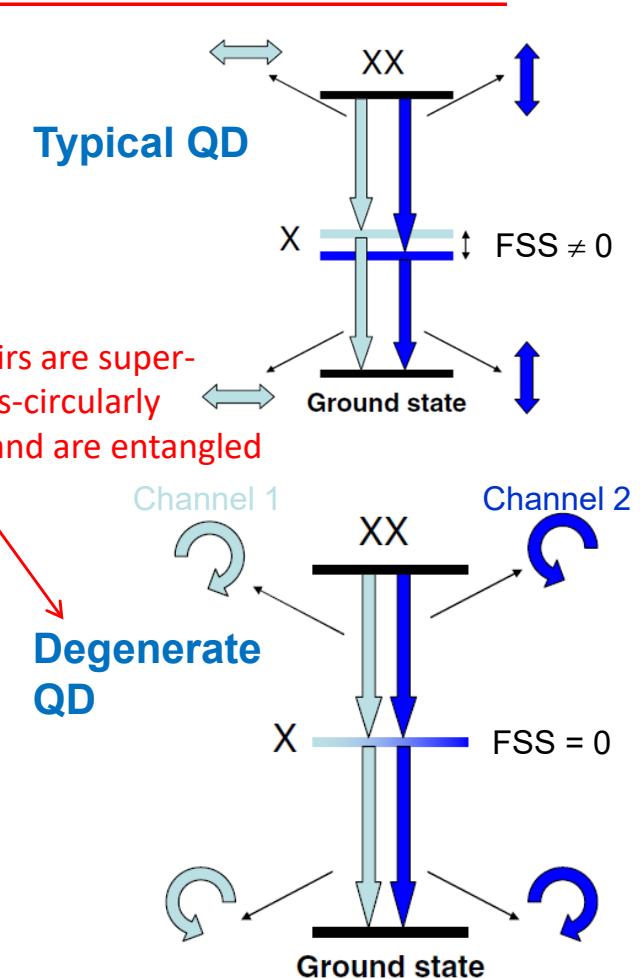


Transmitted data are encrypted and decrypted using an encryption algorithm and a secret key, which has been generated by quantum means, e.g., through the generation of entangled photons

Single QD engineered to have no fine structure splitting (FSS), i.e., same emission energy for polarization channels 1 & 2

⇒ The polarization of each photon cannot be determined by energy measurements (absence of "which-path" information)

Quantum entanglement: physical mechanism such that the quantum state of each particle of a group cannot be described independently of the state of the others



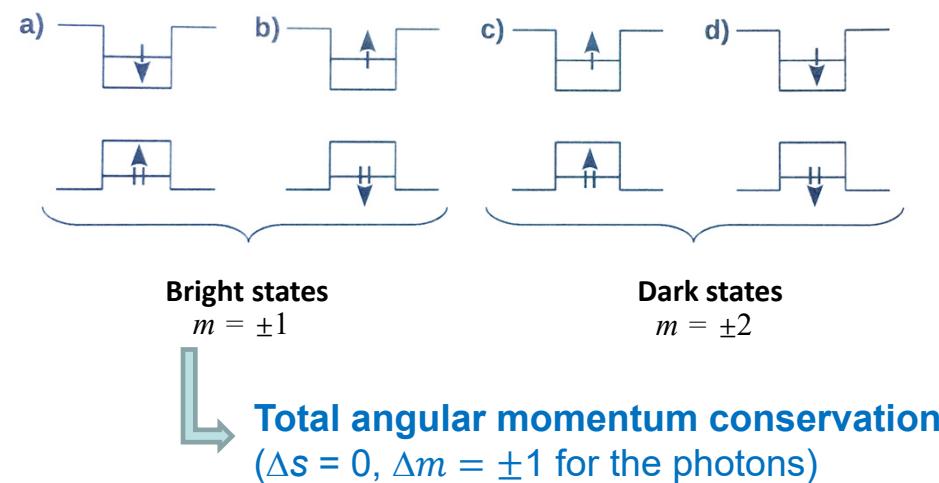
¹R. J. Young *et al.*, N. J. Phys. 8, 29 (2006). (> 240 citations)

Electronic structure of semiconductor QDs

In a QD, only the heavy hole band is considered with total angular momentum $m = \pm 3/2$ (i.e., light holes are out of the game)

Thus, an electron-hole pair is the combination of a hole ($\pm 3/2$) and an electron ($\pm 1/2$), with 4 possible states:

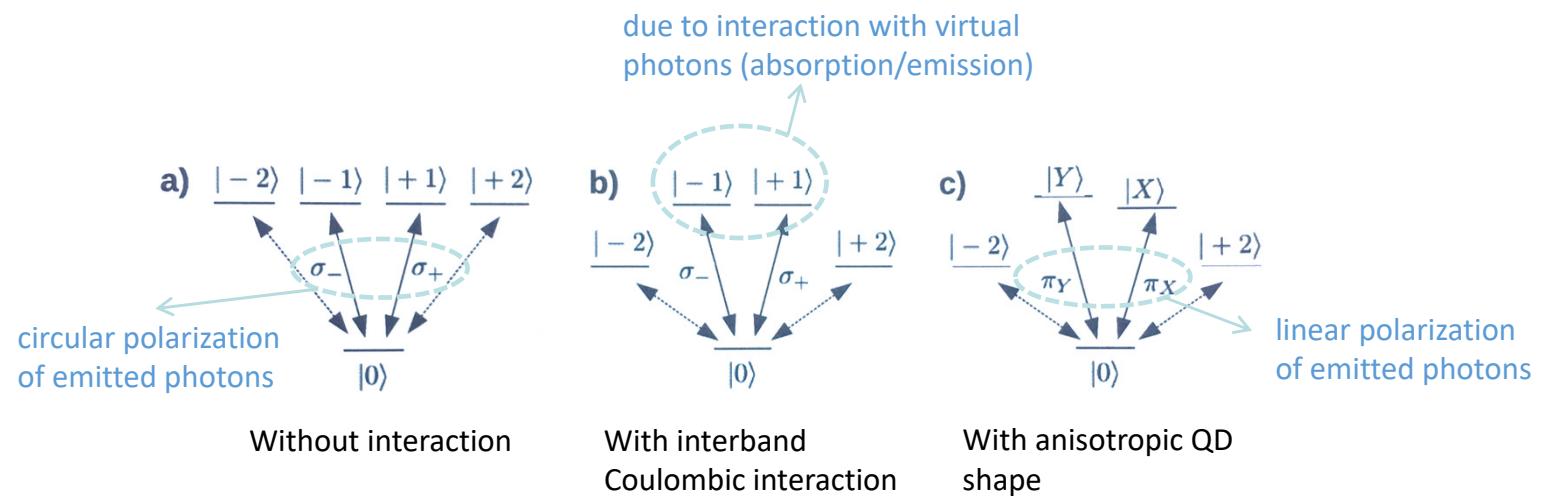
Exciton total angular momentum	
$ +2\rangle = +\frac{3}{2}; +\frac{1}{2}\rangle$	$m = +2$
$ +1\rangle = +\frac{3}{2}; -\frac{1}{2}\rangle$	$m = +1$
$ -1\rangle = -\frac{3}{2}; +\frac{1}{2}\rangle$	$m = -1$
$ -2\rangle = -\frac{3}{2}; -\frac{1}{2}\rangle$	$m = -2$



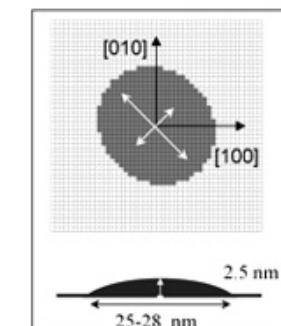
Due to the law of momentum conservation, the total angular momentum of a recombining e-h pair is equal to the angular momentum of the photon with the additional condition of opposite spins for electrons and holes

Electronic structure of semiconductor QDs

Electron-hole Coulomb interaction lifts the degeneracy of the 4 QD ground states:

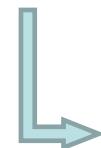
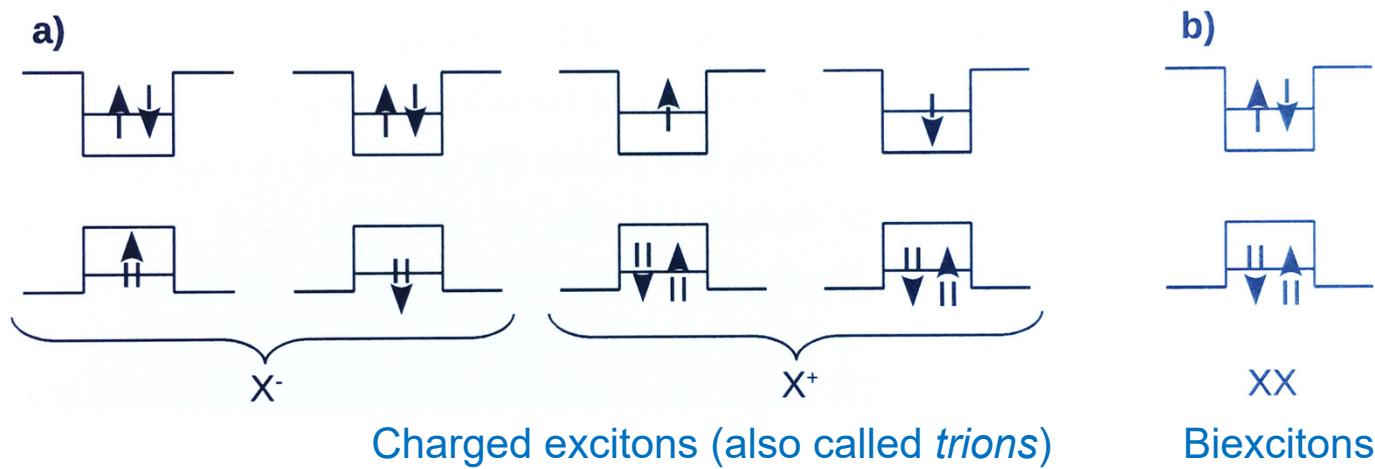


The splitting between bright states amounts to a few hundreds of μeV in InAs/GaAs QDs
⇒ This is the so-called fine-structure splitting (FSS)



Electronic structure of semiconductor QDs

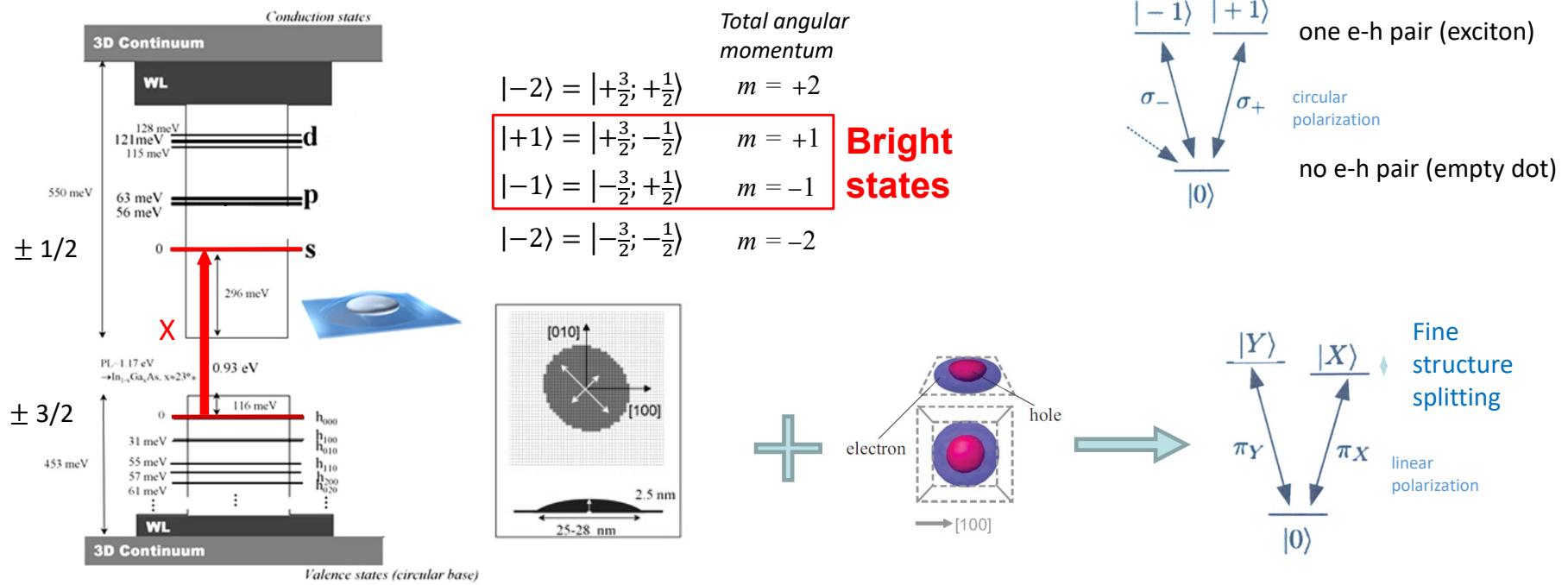
Charged states and biexciton



interband coulombic interaction vanishes \Rightarrow fine-structure splitting (FSS) = 0, i.e., the exchange splitting at the level of a neutral exciton can be switched off by injecting an additional carrier

Physical Review B **65**, 195315 (2002). (> 880 citations)

Electronic structure of semiconductor QDs



Fine-structure splitting (FSS) in QDs

PRL 95, 257402 (2005)

PHYSICAL REVIEW LETTERS

week ending
16 DECEMBER 2005

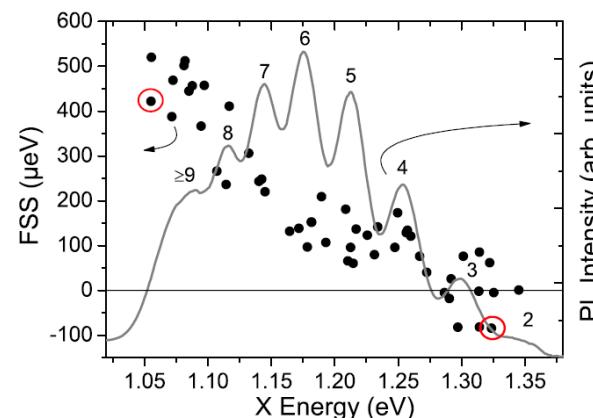
Size-Dependent Fine-Structure Splitting in Self-Organized InAs/GaAs Quantum Dots

R. Seguin, A. Schliwa, S. Rodt, K. Pötschke, U. W. Pohl, and D. Bimberg

Institut für Festkörperphysik, Technische Universität Berlin, Hardenbergstrasse 36, 10623 Berlin, Germany

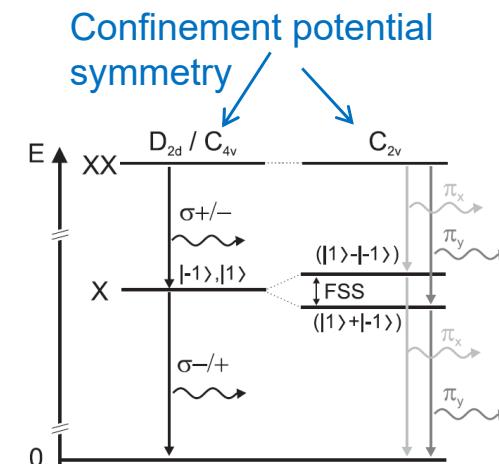
(> 270 citations)

QDs/matrix	FSS [μeV]	Setup
InAs/GaAs	110–180	μPL
InAs/GaAs	≤140	μPL
InAs/GaAs	40	FWM
InGaAs/GaAs	30–150	μPL
InGaAs/GaAs	10–42	Transmission
InGaAs/GaAs	8–36	Pump & probe
InGaAs/GaAs	6–96	FWM
InAs/AlGaAs	500–1000	μPL

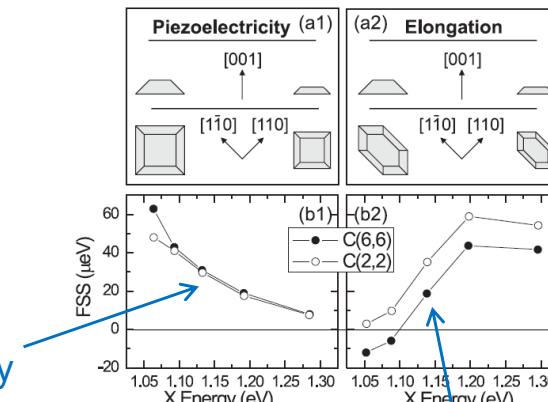


- Qualitative agreement when including piezoelectric effects
- Elongation ruled out

With piezoelectricity



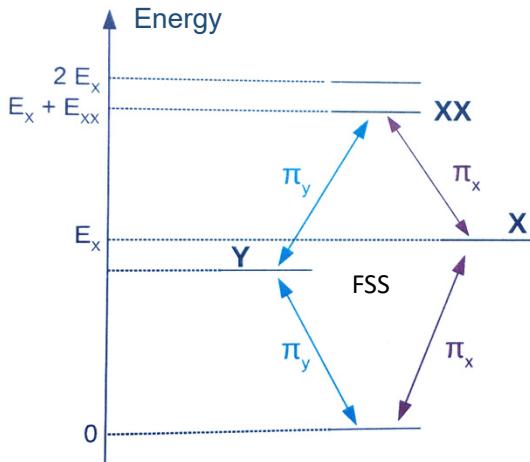
What is governing the FSS?



Without piezoelectricity

Electronic structure of semiconductor QDs

Measuring the fine structure splitting



$$X \quad |+1\rangle = \left| +\frac{3}{2}; -\frac{1}{2} \right\rangle$$

$$Y \quad | -1\rangle = \left| -\frac{3}{2}; +\frac{1}{2} \right\rangle$$

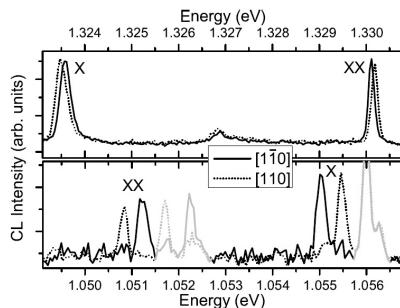
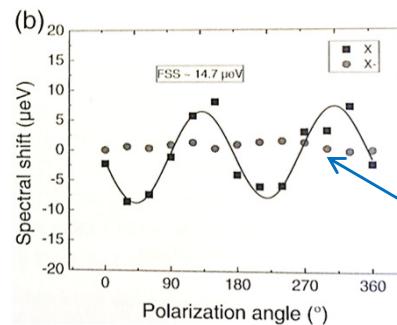
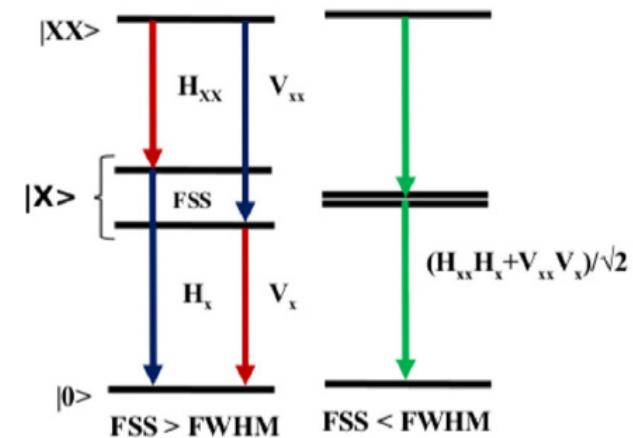


FIG. 2: Polarized spectra for two different QDs emitting at high and low energies are shown. The FSS is -80 and 420 μeV , respectively. Gray lines in the lower panel originate from charged excitonic complexes not considered in this Letter.

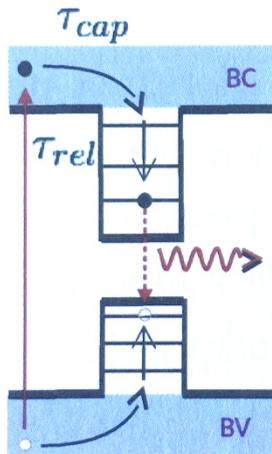
Phys. Rev. Lett. 95, 257402 (2005)



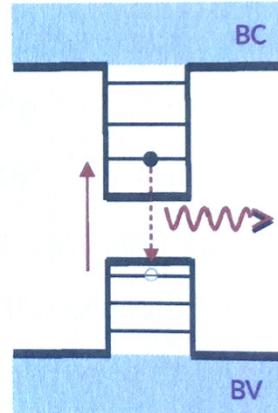
The FSS can only be measured if the emission linewidth of the QD is smaller than the energy separation

Quenching of the exchange splitting for the trion

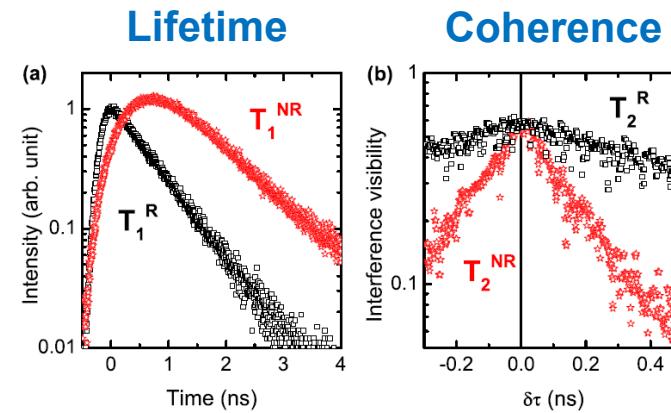
Resonant versus nonresonant excitation



Nonresonant excitation (NR):
 laser energy $> E_g$ barrier
 $\tau_{cap} \approx \tau_{rel} \approx 1-5$ ps
 Few electron-hole pairs created



Resonant excitation (R):
 laser energy $\approx X$ energy
 One electron-hole pair created
 (Pauli blocking)



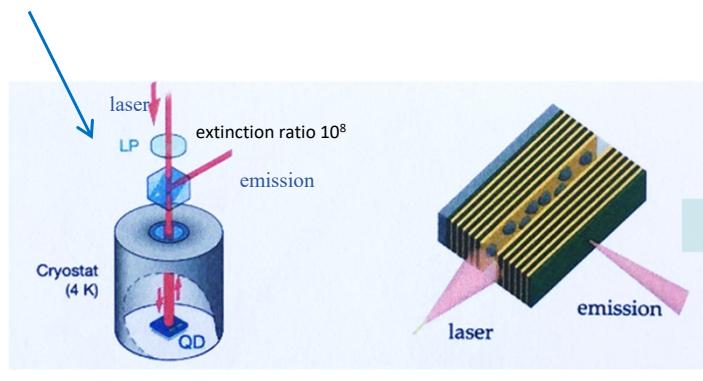
Radiative lifetime $T_1^{NR} = 850$ ps $T_1^R = 670$ ps Dephasing time
 $T_2^{NR} = 200$ ps $T_2^R = 950$ ps
 Capture and relaxation processes induce photon decoherence ($T_2 \downarrow$)

Phys. Rev. B **90**, 041303(R) (2014)

In the limit of high quality InAs/GaAs QD, at low $T(K)$, the homogeneous linewidth is given by the relationship $\Gamma = \frac{2\hbar}{T_2^R}$ with T_2^R the dephasing time of radiative origin (= 630 ps at 7 K (slide 13, Lecture 5), i.e., $\Gamma \sim 2.1$ μ eV)

Resonant excitation: experimental setup

Use of two crossed polarizers to separate the laser from the QD emission

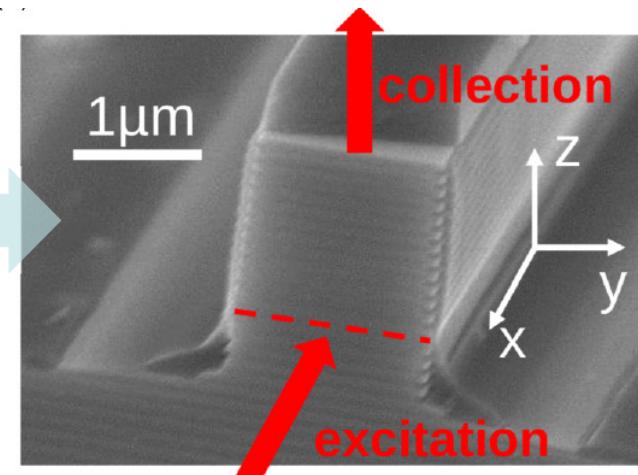


Polarization selection

Nature 525, 222 (2015)

Spatial selection

Phys. Rev. Lett. 114, 067401 (2015)



PHYSICAL REVIEW B 90, 041303(R) (2014)

Number of photons per laser pulse: $N = \lambda P / hc v_{rep}$ $\sim 10^4 - 10^5 \Rightarrow 1$ photon from the dot

Repetition rate of the laser



Need a special experimental setup to discriminate QD photons from laser ones

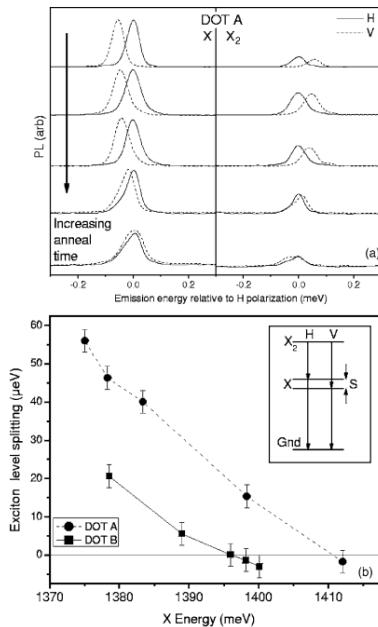
Control of the fine-structure splitting

APPLIED PHYSICS LETTERS 90, 011907 (2007)

Control of fine-structure splitting of individual InAs quantum dots by rapid thermal annealing

D. J. P. Ellis,^{a)} R. M. Stevenson, R. J. Young, and A. J. Shields
Toshiba Research Europe Limited, Cambridge Research Laboratory, 260 Science Park, Milton Road, Cambridge CB4 0WE, United Kingdom

P. Atkinson and D. A. Ritchie
Cambridge Laboratory, Cambridge University, JJ Thomson Avenue, Cambridge CB3 0HE, United Kingdom



PRL 109, 147401 (2012) Selected for a Viewpoint in Physics
 PHYSICAL REVIEW LETTERS

week ending
 5 OCTOBER 2012

Universal Recovery of the Energy-Level Degeneracy of Bright Excitons in InGaAs Quantum Dots without a Structure Symmetry

R. Trotta,^{1,2,*} E. Zallo,¹ C. Ortiz,³ P. Atkinson,^{1,4} J. D. Plumhof,¹ J. van den Brink,³ A. Rastelli,^{1,2,†} and O. G. Schmidt¹

¹Institute for Integrative Nanosciences, IFW Dresden, Helmholtzstrasse 20, D-01069 Dresden, Germany

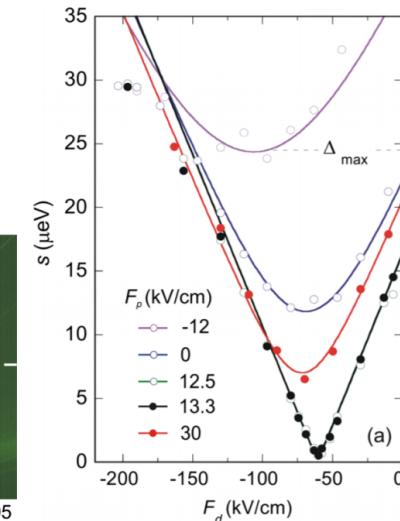
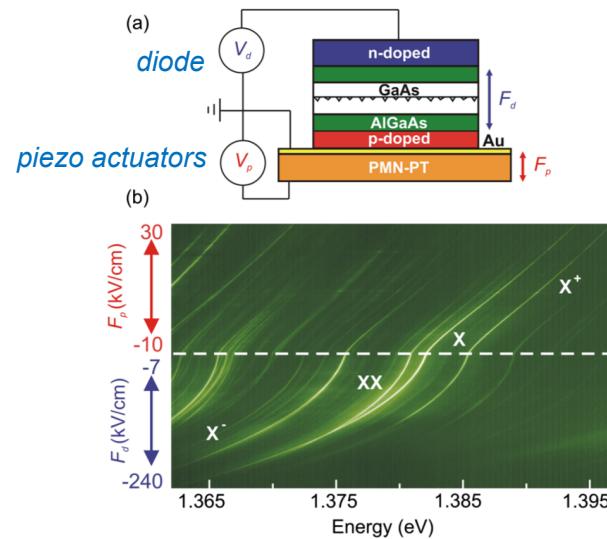
²Institute of Semiconductor and Solid State Physics, Johannes Kepler University Linz, Altenbergerstrasse 69, A-4040 Linz, Austria

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⁴Institut des NanoSciences des Paris, UPMC CNRS UMR 7588, 4 Place Jussieu Boite courrier 840, Paris 75252 Cedex 05, France

(Received 7 June 2012; published 1 October 2012)

(> 140 citations)



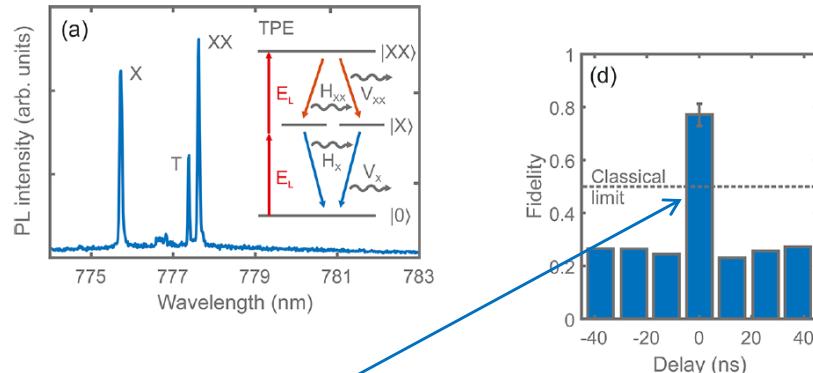
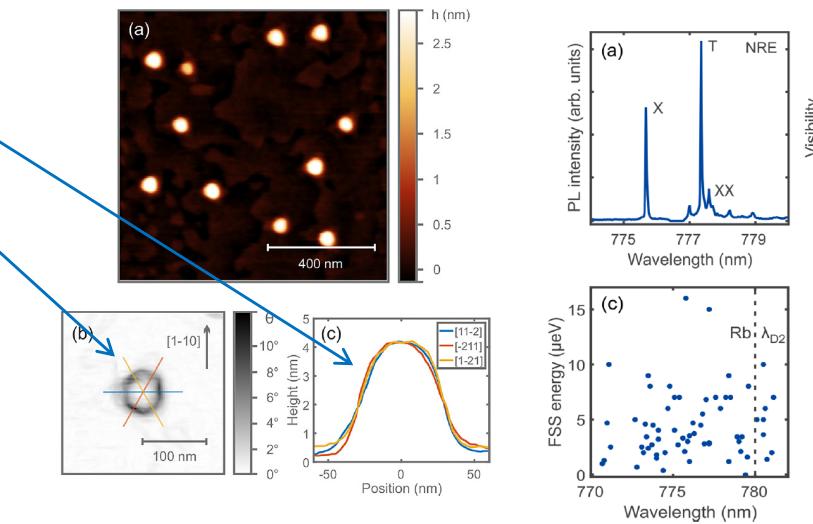
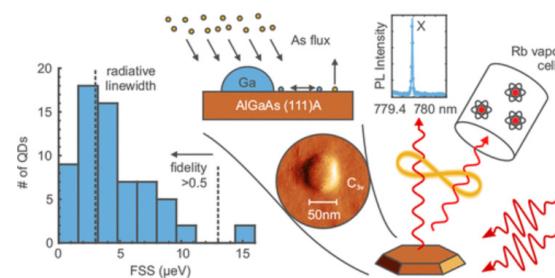
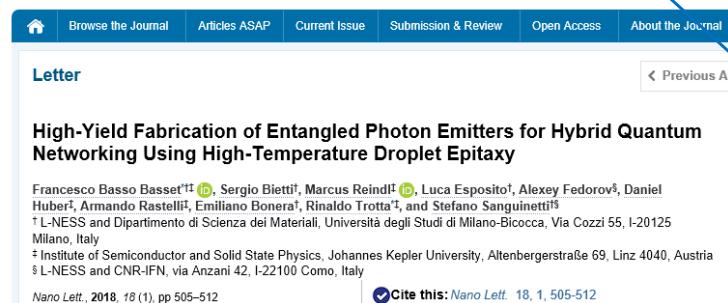
Same trend as in slide 7!

Entangled photon sources

Isotropic GaAs quantum dots obtained from liquid Ga droplets further exposed to As form GaAs islands

To cancel the FSS

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Fidelity to the expected maximally entangled Bell states \Rightarrow
important figure of merit for an entangled quantum light source

Entangled photon sources

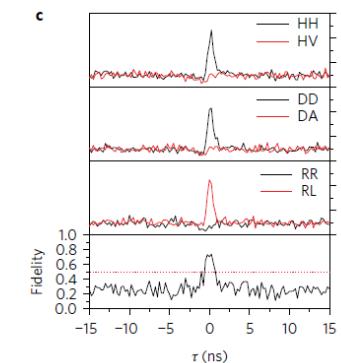
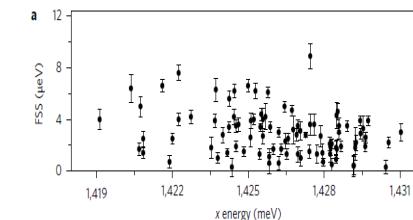
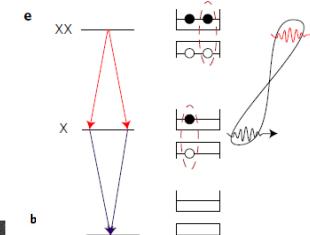
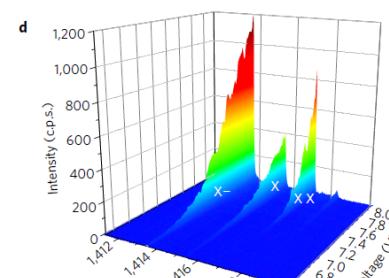
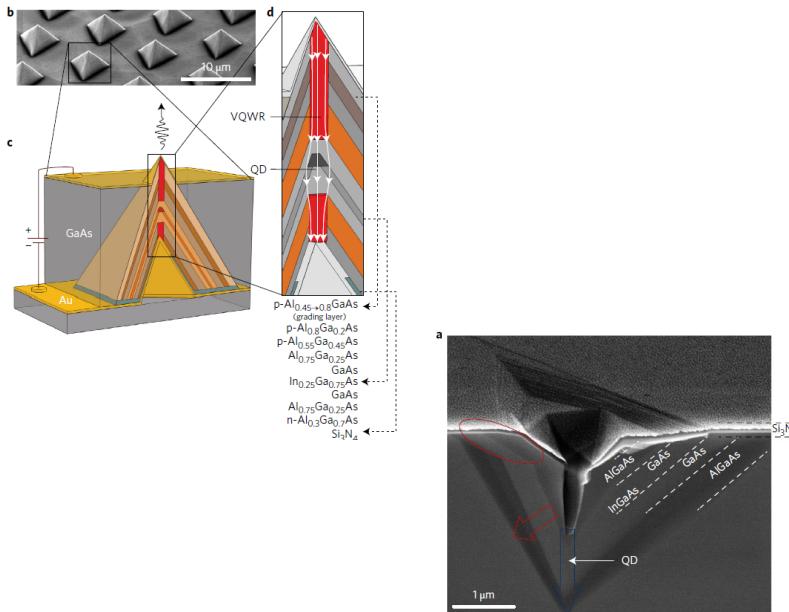
ARTICLES

PUBLISHED ONLINE: 31 OCTOBER 2016 | DOI: 10.1038/NPHOTON.2016.203

nature
photronics

Selective carrier injection into patterned arrays of pyramidal quantum dots for entangled photon light-emitting diodes

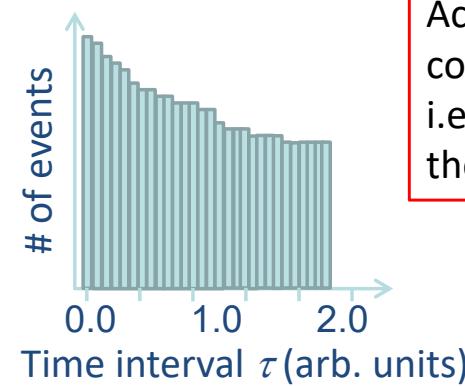
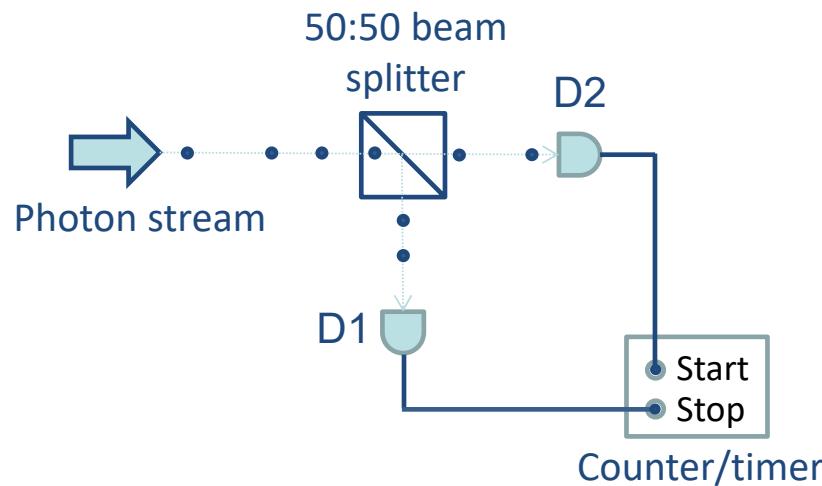
T. H. Chung[†], G. Juska^{*†}, S. T. Moroni, A. Pescaglini, A. Gocalinska and E. Pelucchi



$(1/\sqrt{2})(|DD\rangle + |AA\rangle) = (1/\sqrt{2})(|RL\rangle + |LR\rangle)$, where $|R\rangle = (1/\sqrt{2})(|H\rangle + i|V\rangle)$, $|L\rangle = (1/\sqrt{2})(|H\rangle + i|V\rangle)$, $|D\rangle = (1/\sqrt{2})(|H\rangle + |V\rangle)$ and $|A\rangle = (1/\sqrt{2})(|H\rangle + |V\rangle)$ are the right/left-hand circular, diagonal and antidiagonal polarization states, respectively.

Photon statistics as a tool to assess coherence

Hanbury Brown-Twiss experiment



Access to 2nd-order correlation function, $g^{(2)}(\tau)$, i.e., intensity fluctuations in the photon field

- Bunched (chaotic) light: $g^{(2)}(0) > 1$   
- Thermal light
- Coherent light: $g^{(2)}(0) = 1$  (random)
Laser light
- Antibunched light: $g^{(2)}(0) < 1$ 
SPEs (ideal SPE, $g^{(2)}(0) = 0$)

The Nobel Prize in Physics 2005

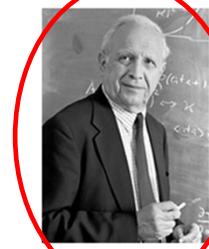


Photo: J. Reed
Roy J. Glauber
Prize share: 1/2



Photo: Sears.P.Studio
John L. Hall
Prize share: 1/4



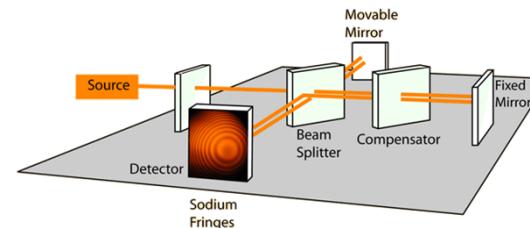
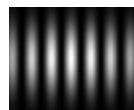
Photo: F.M. Schmidt
Theodor W. Hänsch
Prize share: 1/4

Some insights into the 2nd order coherence function

1st order coherence function: *quantifies electric field fluctuations in time*

$$g^{(1)}(\tau) = \frac{\langle \mathcal{E}^*(t)\mathcal{E}(t+\tau) \rangle}{\langle |\mathcal{E}(t)|^2 \rangle}$$

$$\text{visibility} = \frac{I_{max} - I_{min}}{I_{max} + I_{min}} = |g^{(1)}(\tau)|$$



2nd order coherence function: *quantifies intensity fluctuations in time*

$$g^{(2)}(\tau) = \frac{\langle \mathcal{E}^*(t)\mathcal{E}^*(t+\tau)\mathcal{E}(t+\tau)\mathcal{E}(t) \rangle}{\langle \mathcal{E}^*(t)\mathcal{E}(t) \rangle \langle \mathcal{E}^*(t+\tau)\mathcal{E}(t+\tau) \rangle} = \frac{\langle I(t)I(t+\tau) \rangle}{\langle I(t) \rangle \langle I(t+\tau) \rangle}$$

Quantum state describing here a set of quantum emitters

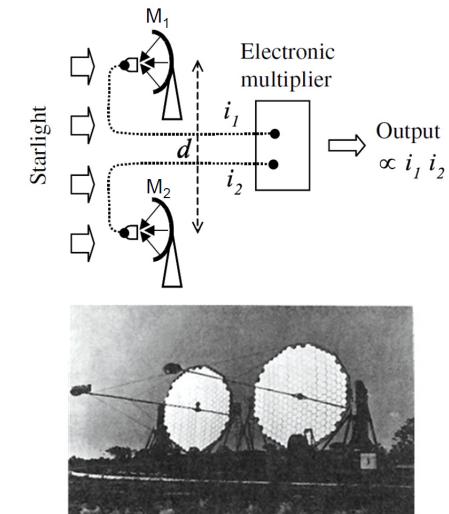
For a given number *Fock state* $|n\rangle$ satisfying the bosonic commutation, the zero time-delay correlation becomes¹

$$g^{(2)}(0) = \frac{n(n-1)}{n^2}$$

¹I. Aharonovich *et al.*, Rep. Prog. Phys. **74**, 076501 (2011).

# n of quantum emitters	$g^{(2)}(0)$
1	0
2	0.5
3	0.667

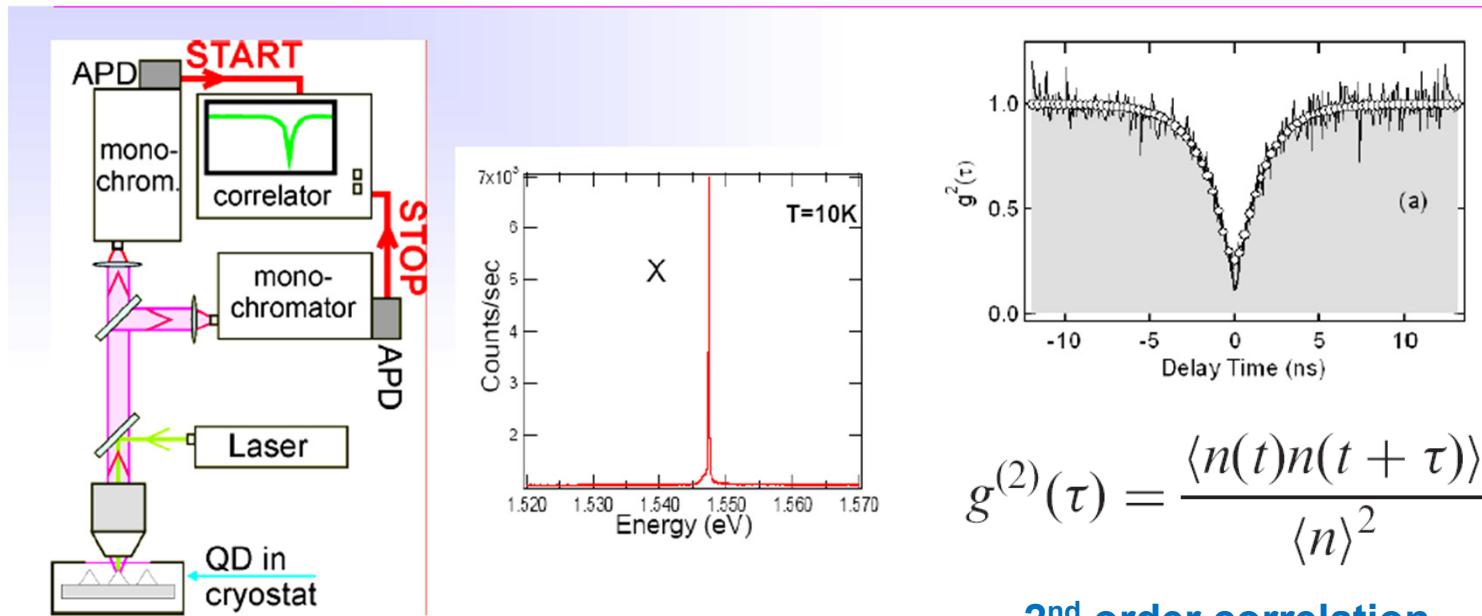
Hanbury Brown and Twiss, Nature **177**, 27 (1956). (> 2030 citations)
 Hanbury Brown and Twiss, Nature **178**, 1046 (1956). (> 850 citations)



→ $g^{(2)}(0) < 0.5$ for ensuring true single photon emission

Single photon emission from QDs

Hanbury Brown and Twiss interferometry



- Single QDs are readily selected and probed
- Photon antibunching observed at X lines

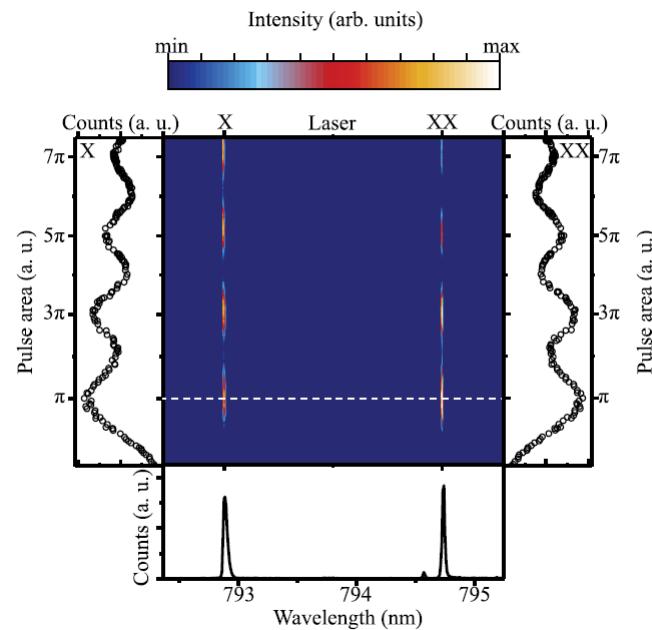
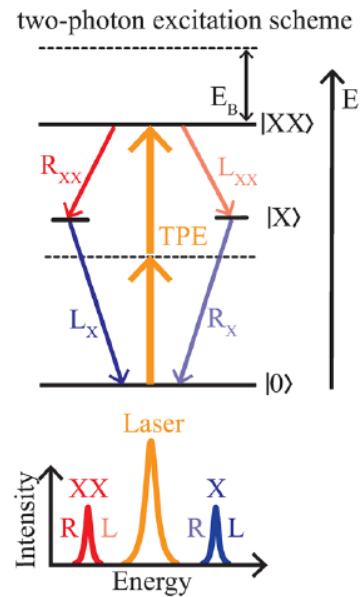
$$g^{(2)}(\tau) = \frac{\langle n(t)n(t + \tau) \rangle}{\langle n \rangle^2}$$

2nd order correlation function

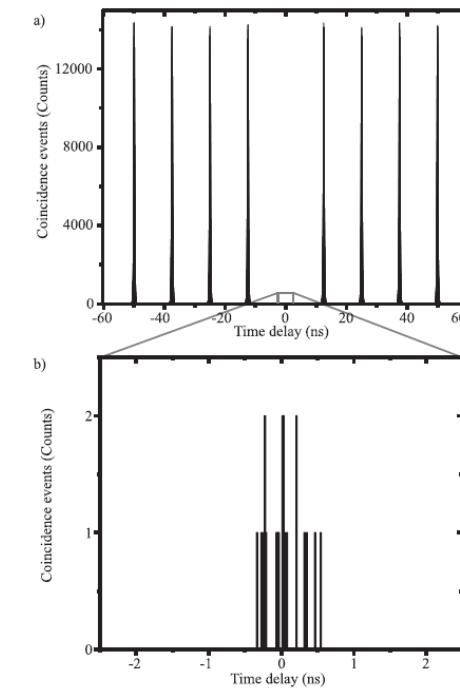
Photon antibunching

Two-photon resonant excitation of the biexciton

Reduce decoherence and multi-photon emission¹



Laser spectral width of $260 \mu\text{eV}$ and pulse length of 7 ps



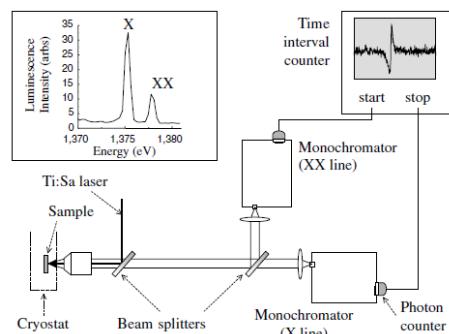
$$g^{(2)}(0) = (7.5 \pm 1.6) \times 10^{-5}$$

Low background noise

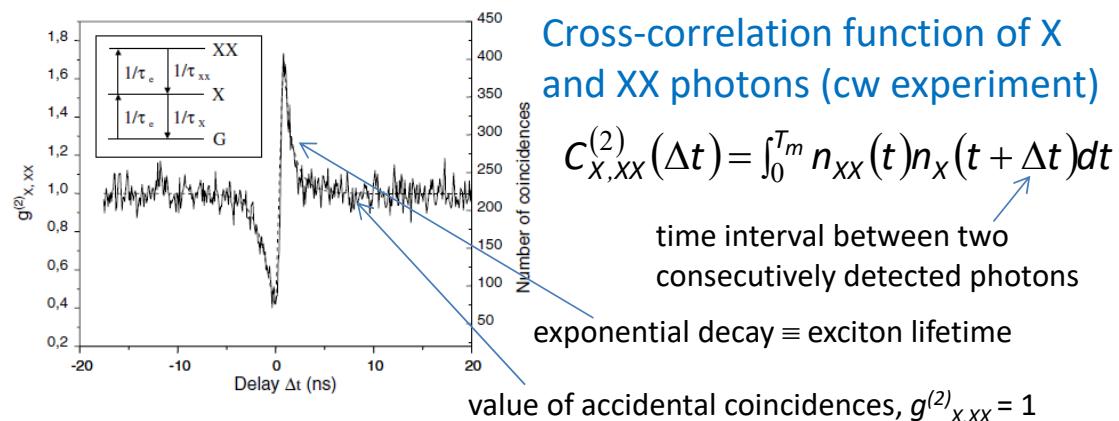
¹L. Schweickert *et al.*, Appl. Phys. Lett. **112**, 093106 (2018). (> 210 citations)

Quantum cascade of photons in quantum dots¹

Experiment highlighting the potential of single dots as an efficient source of single photons (quantum information applications)



Hanbury-Brown & Twiss setup with λ selective arms



Biexciton/exciton as sources of pairs of correlated single photons

Spectral filtering allows isolation of a single photon by a single dot (antibunching \rightarrow single photon nature of exciton emission)

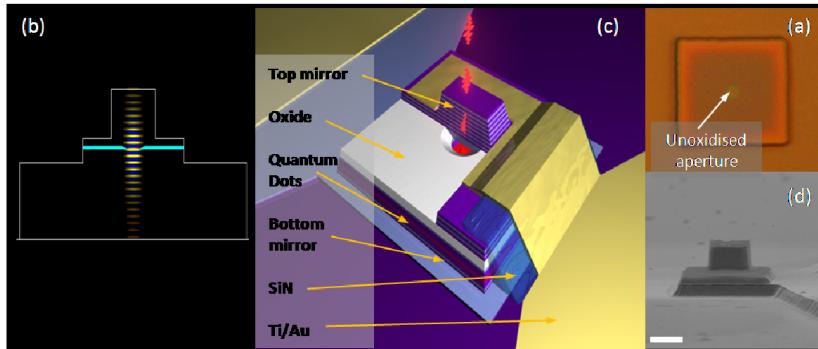
Cross-bunching ($\Delta t > 0$, i.e. XX 1st, X 2nd) and **cross-antibunching** ($\Delta t < 0$, when X photon is detected, system \equiv ground state, probability of detecting XX photon $\rightarrow 0$)

Quantum efficiency for photon pair production in QDs \gg efficiencies atomic cascades and parametric fluorescence (but operation limited to liquid He temperature)

Traditional sources of entangled photons

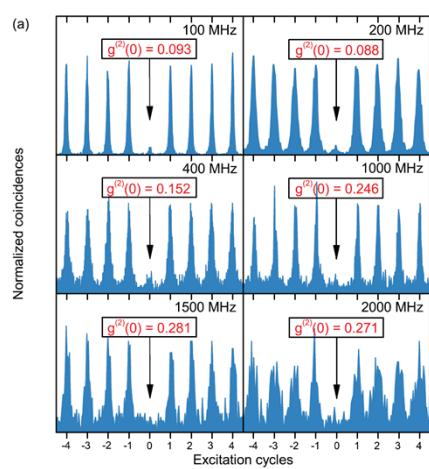
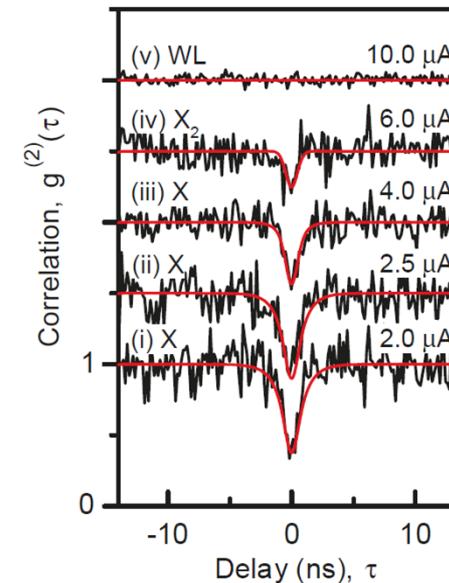
¹E. Moreau *et al.*, Phys. Rev. Lett. **87**, 183601 (2001). (> 300 citations)

Electrically-driven QD-based single photon emitters



Cambridge + Toshiba UK

Z. L. Yuan *et al.*, Science **295**, 102 (2002). (> 1030 citations)



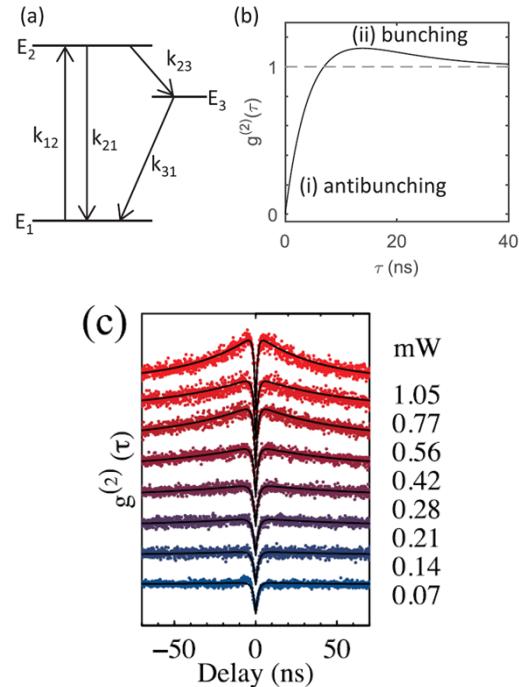
Single photon emission up to excitation repetition rates of 2 GHz

F. Hargart *et al.*, Appl. Phys. Lett. **102**, 011126 (2013).

Alternative solid-state single photon emitters

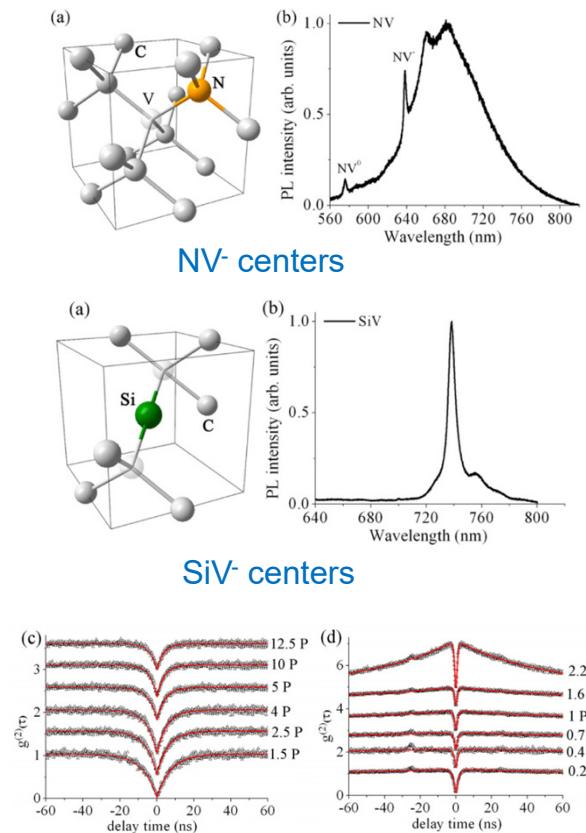
Deep-level defects in:

SiC¹



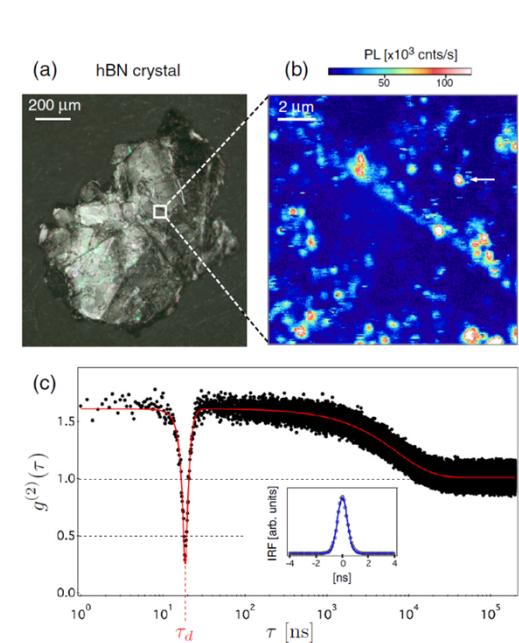
¹A. Lohrmann *et al.*, Rep. Prog. Phys. **80**, 034502 (2017).

Diamond²



²I. Aharonovich *et al.*, Rep. Prog. Phys. **74**, 076501 (2011).

hBN³



³L. J. Martínez *et al.*, Phys. Rev. B **94**, 121405(R) (2016).