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Probing electron-phonon interaction through two-photon interference in resonantly driven semiconductor quantum dots

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We investigate the temperature dependence of photon coherence properties through two photon interference (TPI) measurements from a single QD under resonant excitation. We show that the loss of indistinguishability is only related to the electron-phonon coupling without being affected by spectral diffusion. Through these measurements, and a complementary microscopic theory, we identify two independent separate decoherence processes associated to phonons. Below 10 K, we find that the relaxation of the vibrational lattice is the dominant contribution to the loss of TPI visibility. This process is non-Markovian in nature, and corresponds to real phonon transitions resulting in a broad phonon sideband in the QD emission spectra. Above 10 K, virtual phonon transitions to higher lying excited states in the QD become the dominant dephasing mechanism, this leads to broadening of the zero phonon line, and a corresponding rapid decay in the visibility. The microscopic theory we develop provides analytic expressions for the dephasing rates for both virtual phonon scattering and non-Markovian lattice relaxation.

Many recent developments in quantum information processing rely on the use of solid-state qubits that can emit indistinguishable single photons on demand [1, 2]. However, maintaining the coherence between consecutively emitted photons remains a true challenge for realising a deterministic source of identical photons. A promising candidate for the development of such a source are self-assembled semiconductor quantum dots (QDs) embedded in photonic nanostructures [3, 4].

Despite impressive milestones in the development of these devices, a QD naturally couples strongly to its surrounding solid-state matrix, constituting an inherently open quantum system. The excitonic degrees of freedom are heavily influenced by the vibrational modes [5–8], fluctuating charges [9], and nuclear spins [10] of the host material, all of which lead to dephasing, therefore suppressing the coherence properties of the emitted photons. Decoherence may be reduced by enhancing the emission rate through the Purcell effect, or using resonant excitation to minimise laser-induced dephasing, leading to bright single-photon sources with near unity indistinguishability [11–15].

Still, there are a number of open questions regarding the role of phonon processes on the coherence properties of emitted photons. Recent experimental and theoretical work has demonstrated the importance of a microscopic model to understand the role of phonons on the emission properties of QDs [8, 10, 26]. Examples include excitation-induced dephasing of excitonic Rabi oscillations [5, 27]; sideband linewidth in resonance fluorescence (RF) [21, 23, 29]; and temperature-dependent Rabi frequency renormalisation [6, 21]. In the examples given, the excitonic degrees of the QD are assumed to couple linearly to the phonon environment, inducing thermalisation in the QD eigenbasis [7, 8], leading to a broad non-Markovian sideband in the QD emission spectra [30].

Here, we present a combined experimental and theoretical investigation of the coherence properties of photons emitted by a quantum dot, allowing us, unambiguously, to separate real and virtual decoherence processes due to phonons and their temperature dependencies. To do so, we take temperature dependent measurements of two-photon interference (TPI) in a Hong-Ou-Mandel (HOM) configuration using strictly resonant excitation conditions. We show that TPI measurements are not affected by spectral diffusion due to fluctuating charges, as this process is slow compared to the emission time interval between the two interfering photons. Therefore loss of indistinguishability is attributed only to electron-phonon scattering. Through temperature dependent TPI measurements, we demonstrate that linear electron-phonon coupling is not sufficient to capture the trend in TPI visibility. In order to describe the behaviour observed, we develop a microscopic model, based on polaron theory, to include phonon induced virtual transitions to higher lying states in the QD [31, 32]; this leads to temperature dependent broadening of the Zero Phonon Line (ZPL). Our formalism, allows us to derive analytic forms for the dephasing due to both virtual phonon processes and non-Markovian lattice relaxation. This provides novel insights into the dephasing mechanisms relevant to QD based single photon sources.

Experiment - In contrast to previous temperature dependent TPI measurements [33], our geometry (see
Fig. 1 (a) Scheme of the experimental set-up. A tunable Ti:sapphire (82 MHz) laser delivers 3-ps pulses and, for HOM experiments, pairs of pulses separated by 3 ns. The laser is focused by a microscope objective on the cleaved edge of one ridge and the RF is collected from the top surface by a second microscope objective. The sample and the two objectives are inside a closed-cycle He temperature-variable cryostat. The microscope objective. The sample and the two objectives are focused by a microscope objective on the cleaved edge of one ridge and the RF is collected from the top surface by a second microscope objective. The sample and the two objectives are inside a closed-cycle He temperature-variable cryostat. The signal is coupled to a fibered set-up for either standard spectroscopy, or Michelson interferometry \((g(1))\) or TPI experiments \((g(2))\) using a 3 ns unbalanced Mach-Zehnder interferometer. (b) Scanning electron microscopy image of one ridge with the dots schematically drawn in the layer.

Fig. 1) [34] allows us to use strictly resonant \((s\text{-shell})\) excitation removing dephasing due to relaxation from higher excited states and time-jitter [35]. Furthermore, the only filtering used in our measurements is due to a low-Q cavity which increases the collection efficiency. We present results on three different self assembled InAs/GaAs QDs from different samples (labelled QD1, QD2 and QD3 hereafter) excited with resonant \(\pi\)-pulses. Depending on the QD under study, an additional very weak (few nW) He-Ne laser is added to enhance or recover the RF as reported previously [15, 36, 37]. Experimental details can be found in the Supplemental Material [38].

The coherence time \(T_2\), corresponding to the width of the RF line, can be measured by Fourier transform (FT) spectroscopy using a Michelson interferometer. The contrast of the interference fringes is adjusted by a pseudo-Voigt profile, with an inhomogeneous contribution \(\eta\) (see [38]), usually attributed to spectral diffusion effects [39]. In Table I we give the values of the measured radiative lifetime \(T_1\), of \(\eta\) and the ratio \(T_2/2T_1\) for the three QDs. The corresponding FT spectra for the three QDs are given in Fig. 2 of the Supplemental Material [38].

Second-order correlation measurements have been performed, allowing characterisation of the single-photon emission purity and indistinguishability. Fig. 2 shows the results obtained for QD1 at 4 K. Single-photon interferences in a Hanbury-Brown-Twiss (HBT) experiment (Fig. 2a) clearly show an antibunching with a low multiphoton probability \(g^{(2)}\), the values are given for the three QDs in Table I. We attribute the background correlations at 0 and \(\pm3\) ns delay to the remaining scattered laser. The unusual shape of the histogram of coincidences is explained by the fact that the single photons pass through the two arms of the Mach-Zehnder interferometer used for the HOM setup (see Fig. 1a and [38]). A multi-exponential decay fit (red line) is used to extract the value of \(g^{(2)}\) taking into account the overlapping of the different peaks. Fig. 2b shows the raw histogram of the TPI coincidences for QD1 at 4 K. The signature of the indistinguishability of two successively emitted photons corresponds to the small area of peak 2 compared with peaks 1 and 3. The TPI visibility \(V_{\text{TPI}}\) (see Table I) is deduced from the second-order correlation function at zero delay \(g^{(2)}\) corrected by the remaining scattered laser and by the experimental imperfections (contrast of the Mach-Zehnder interferometer and not perfect 50/50 fibered beam splitters) [38, 40]. In the literature, it is commonly accepted that the visibility of TPI experiments can be obtained from the ratio \(T_2/2T_1\), assuming random dephasing processes by phonons and charges [41], and labelled \(V\) hereafter. The values of \(V\) are given in Table I and correspond to a poor degree of indistinguishability. This is in striking contrast with the results obtained by TPI experiments where \(V_{\text{TPI}} \sim 0.80\) for all QDs at 4 K. The significant difference can be explained by recognising the distinct characteristic timescales of the two kind of experiments which probe different physical dephasing processes [11, 12, 35, 32, 43]. Indeed, because of the long

FIG. 1. (a) Scheme of the experimental set-up. A tunable Ti:sapphire (82 MHz) laser delivers 3-ps pulses and, for HOM experiments, pairs of pulses separated by 3 ns. The laser is focused by a microscope objective on the cleaved edge of one ridge and the RF is collected from the top surface by a second microscope objective. The sample and the two objectives are inside a closed-cycle He temperature-variable cryostat. The signal is coupled to a fibered set-up for either standard spectroscopy, or Michelson interferometry (\(g^{(1)}\)) or TPI experiments (\(g^{(2)}\)) using a 3 ns unbalanced Mach-Zehnder interferometer. (b) Scanning electron microscopy image of one ridge with the dots schematically drawn in the layer.

FIG. 2. Second-order correlation measurements for QD1 at 4 K for one hour acquisition time. (a) Coincidences histogram for HBT experiment, we extract \(g^{(2)}_{\text{HBT}} = 0.12 \pm 0.02\). (b) Coincidences histogram for TPI experiment. After correction by the remaining laser background, we obtain \(V_{\text{TPI}} = 0.79 \pm 0.03\).

<table>
<thead>
<tr>
<th>(T_1) (ps)</th>
<th>(\eta)</th>
<th>(T_2/2T_1)</th>
<th>(g^{(2)}_{\text{HBT}})</th>
<th>(V_{\text{TPI}})</th>
<th>(\tilde{V})</th>
</tr>
</thead>
<tbody>
<tr>
<td>QD 1</td>
<td>1100</td>
<td>0.45</td>
<td>0.35</td>
<td>0.12</td>
<td>0.79</td>
</tr>
<tr>
<td>QD 2</td>
<td>750</td>
<td>0.55</td>
<td>0.23</td>
<td>0.11</td>
<td>0.83</td>
</tr>
<tr>
<td>QD 3</td>
<td>670</td>
<td>0.10</td>
<td>0.71</td>
<td>0.07</td>
<td>0.83</td>
</tr>
<tr>
<td>Errors</td>
<td>2 %</td>
<td>±0.1-10</td>
<td>±0.02</td>
<td>±0.04</td>
<td>10-15%</td>
</tr>
</tbody>
</table>

TABLE I. Values of the experimental parameters (see text for definitions) for the three QDs under resonant excitation. \(\tilde{V}\) is the expected TPI visibility from \(T_1\) and \(T_2\) measurements assuming random dephasing processes (see text).
The measured TPI spectra. Virtual transitions give broadening of the ZPL thus significantly higher than the average phonon energies in low temperature experiments. With this in mind, any transitions from the first excited to another electronic state of the QD must be virtual in nature. Following Muljarov and Zimmerman \cite{31} we derive an effective Hamiltonian theory \cite{17}, treating the charge density operator as a perturbation, with the first- and second-order terms capturing real and virtual phonon transitions respectively, as shown schematically in Fig. 4.

We start by explicitly considering the ground state |0\rangle, and single exciton in the s-shell denoted |X\rangle, with excitonic splitting \(\omega_X\). By using an effective Hamiltonian theory \cite{33}, we can eliminate off diagonal transitions to higher lying phonon states \cite{31}, yielding the Hamiltonian, \(H = H_0 + |X\rangle\langle X| (V + V_Q)\). Here \(H_0 = \omega_X |X\rangle\langle X| + \sum_k \omega_k b_k^\dagger b_k\) is the free Hamiltonian of the QD and phonon environment, where \(b_k\) is the annihilation operator of a phonon with wave vector \(k\) and frequency \(\omega_k\).

The electron-phonon interaction has two contributions, the first is the standard linear electron-phonon coupling \(V = \sum_k \omega_k (b_k^\dagger + b_k)\), describing the displacement of the lattice due to the change in charge configuration of the QD \cite{8}. The coupling strength is quantified through the matrix element \(g_k = \sum_{a=e,h} M_{a,k}^{11}\), where for deformation potential coupling,

\[
M_{a,k} = \sqrt{\frac{\rho \nu}{2\mu^2 V}} D_a \int d^3r \psi_{ia}^*(\mathbf{r})\psi_{ja}(\mathbf{r}) e^{i\mathbf{k}\cdot\mathbf{r}},
\]

is the matrix element corresponding to phonon induced transition between the \(i^{th}\)- and \(j^{th}\) electronic states. Here \(\rho\) is the mass density, \(\nu\) is the speed of sound in the material, and \(V\) is the phonon normalisation volume. This matrix element is dependent on the wavefunction \(\psi_{i,e/h}(\mathbf{r})\), of the confined electron/hole, and the corresponding deformation potential \(D_a\).

The second term is quadratic in phonon operators \cite{31}

\[
V_Q = \sum_{k,k'} f_{k,k'} (b_k^\dagger + b_k)(b_{k'}^\dagger + b_{k'}),
\]

and describes virtual phonon transitions between the first exciton state and higher lying excited states. The effective coupling strength for the quadratic coupling takes the form \(f_{k,k'} = \)
the simple pure dephasing form: we may derive a master equation for the reduced density for the pure dephasing rate due to the virtual phonon quadratic Hamiltonian we obtain of the phonon environment, removing the linear electron phonon coupling. Applying this transformation to our of the excitonic degrees of freedom in which takes the form

\[ \dot{\chi}(t) = -i \omega_X [\sigma^\dagger \sigma, \chi(t)] + \Gamma L_\sigma [\chi(t)] + 2\gamma_{pd} L_{\sigma^\dagger \sigma} [\chi(t)], \]

where \( \sigma = |0 \rangle \langle X | \) is the dipole transition operator, and \( L_\sigma [\chi] = O \chi O^\dagger - \{ O^\dagger O, \chi \} / 2 \). Here \( \Gamma \) is the radiative recombination rate of the QD. If we consider only virtual transitions between the lowest exciton states and the next highest states, then, we may find an analytic form for the pure dephasing rate due to the virtual phonon transition [38]:

\[ \gamma_{pd} = \frac{\alpha^2 \mu}{\nu_c} \int_0^\infty \nu^{10} e^{-2\nu^2/\nu_c^2} n(\nu)(n(\nu) + 1) d\nu, \]

where \( n(\nu) = [e^{-\nu \mu} - 1]^{-1} \). \( \alpha \) and \( \mu \) describe the electron-phonon coupling strength and the probability of virtual phonon processes respectively. The cut-off frequency, \( \nu_c \), is directly related to the QD confinement length.

To calculate the visibility of two photon interference we must associate the excitonic degrees of freedom in the QD to the emitted field. In the polaron frame one obtains the Heisenberg picture field operator \( E(t) = \sqrt{\Gamma/2\pi} \sigma(t) B_\perp(t) \), where the standard expression for a dipole emitter is modified by the phonon displacement operator \( B_\perp(t) = e^{iS(t)} \) [21] [20]. With this expression we obtain the polaron frame first-order correlation function, \( g^{(1)}(t, \tau) = \frac{1}{\nu_c} \langle B_\perp(\tau) B_\perp(\tau) \rangle \langle \sigma^\dagger(t + \tau) \sigma(t) \rangle \). The second term in this equation describes emission through the ZPL. The first term is the phonon correlation function which takes the form \( \langle B_\perp(\tau) B_\perp(\tau) \rangle = B^2 \exp(\varphi(\tau)) \), where \( B = \exp(-\varphi(0)/2) \) is the Frank-Condon factor, and \( \varphi(\tau) = \alpha_0^{\infty \nu} \nu \exp(-\nu^2/\nu_c^2) (\coth(\beta \nu/2) \cos(\nu \tau) - i \sin(\nu \tau)) d\nu \). This function decays on a timescale related to the inverse of the cut-off frequency, which is typically on the order of picoseconds, leading to a broad phonon sideband in the spectra of the system. A detailed derivation of these expression is given in the supplement.

Following Ref. [26] we obtain an analytic form for the indistinguishability including the phonon sideband contribution [38]:

\[ I = \frac{\Gamma}{\Gamma + 2\gamma_{pd}} \left( \frac{|h(0)|^2 B^2}{|h(0)|^2 B^2 + F(1 - B^2)} \right)^2, \]

where \( |h(0)|^2 = (\kappa/2)^2 (\delta^2 + (\kappa/2)^2)^{-1} \) with \( \kappa \) the cavity width and \( \delta \) the QD-cavity detuning. The first factor gives the contribution of photons emitted through the ZPL, while the second describes the reduction of the indistinguishability from photons emitted through the phonon sideband. The temperature dependent factor, \( F \), quantifies the unfiltered fraction of the phonon sideband extracted from the low-Q cavity [50].

We use the above expression to fit the experimental data given in Fig. 3 using a least square fitted sqrt fit, we find an optimum fit for parameters \( \alpha = 0.0082 \text{ps}^{-2} \), \( \nu_c = 7.9 \text{ps}^{-1} \) and \( \mu = 4.4 \times 10^{-4} \text{ps}^2 \). \( \alpha \) and \( \mu \) depend only on the material parameters and reasonable agreement is found when compared to the theoretical values. The cut-off frequency gives a characteristic confinement length of the order of one nanometre which is the right order of magnitude for typical self assembled QD. The fit captures the qualitative and quantitative behaviour of the data. Furthermore, from the simple form of the expression given in Eq. 2 we can analyse the contributions to the indistinguishability due to the phonon sideband, and the virtual transitions to higher lying QD states. Similar results have been obtained on QD2 and are presented in the Supplemental Material [38].

Below 10 K, the average energy of phonons \( k_B T \) is not sufficient to induce virtual transitions (\( \gamma_{pd} \ll \Gamma \)). If we use the extracted parameters to consider only the sideband contribution (dashed curve) we observe a good fit at low temperatures, with a ~10% reduction in the indistinguishability, suggesting that emission via the phonon sideband is the principal cause for the reduction of TPI visibility. However, above 10 K the data demonstrate a rapid decrease in the indistinguishability not captured by the sideband theory. At these temperatures \( k_B T \) is sufficient to induce virtual transitions between the \( s- \) and \( p- \) states of the QD, leading to pure dephasing of the ZPL (\( \gamma_{pd} \sim \Gamma \)), and consequently a suppression of the indistinguishability.

The fact that real and virtual phonon processes occur on separate temperature scales is of vital importance to the development of solid-state single photons sources. Although the phonon sideband is a persistent problem at low temperatures, it may be easily removed through spectral filtering, or use of a high Q-cavity [20]. Though this could reduce the efficiency of the source, transform limited photons may be obtained as \( \gamma_{pd} \approx 0 \) at low temperature. In contrast, the broadening of the ZPL cannot be removed through simple filtering; instead one must rely on Purcell enhancement to reduce its influence. However, the sensitivity of virtual phonon processes to temperature will place significant limitations on the operating regimes of QD systems.

In summary, through temperature dependent TPI
measurements, we have demonstrated that both real and virtual phonon transitions, occurring on very different timescales, play a role in reducing the indistinguishability of photons emitted from QDs under resonant excitation. Using a rigorous microscopic theory, we provide analytic expressions for the dephasing due to these mechanisms, providing new insights into potential operating regimes of QD single photon sources. Furthermore, we expect such a general microscopic approach could be used to describe other specific discrete quantum systems coupled to a bosonic reservoir.

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[49] This is a manifold of three degenerate states with \( p \)-symmetry.
[50] The value of \( F \) varies between 0.19 for \( T \sim 4 \) K and 0.33 for \( T \sim 22 \) K.
InAs/GaAs self-assembled QDs were grown by molecular beam epitaxy (MBE) on a planar (001) GaAs substrate
and embedded in a planar microcavity made of unbalanced Bragg mirrors, with 24 pairs below and 12 pairs above the
QDs. To create single-mode one-dimensional waveguides, ridges ranging from 1 µm to 3 µm wide, depending on the
sample, were etched approximately 1.5 µm deep by inductively coupled plasma etching. Passivation of the surface
has further improved the quality of the heterointerfaces, leading to an important suppression of the scattered laser
and allowing the realization of resonant excitation [1, 2]. Indeed, in this geometry, the excitation laser propagates
along the 1D waveguide, while RF is collected in a perpendicular direction as shown in Fig. 1a of the main text. The
cavity has little effect as a spectral filter but rather enhances the QDs RF collection by a factor of 20 to 50. The
quality factor is rather low around $Q \sim 200 - 300$ corresponding to width $\kappa \approx 4 - 5$ meV. We show in Fig. 5
high power non resonant excitation spectra for QD1 and QD2. We clearly see that only the QDs in the cavity mode can be
observed and we extract the transfer function of the cavity by fitting the background luminescence (removing the QDs luminescence). We indicate by an arrow the luminescence lines of QD1 and QD2 on which TPI have been performed and we extract for each one the width of the cavity and the detuning between the QDs under study and the cavity mode. These values are used in the fitting procedure in Fig. 3 of the main text and in Fig. 9 in the Supplement.

![Figure 5](image.png)

**FIG. 5.** High power non resonant excitation spectra for QD1 and QD2 at 4 K. The red line is the transfer function of the cavity and is extracted by fitting the background luminescence (removing the QDs luminescence). The single exciton transition used in each sample is indicated in the figure by the arrow.

**FOURIER TRANSFORM SPECTROSCOPY MEASUREMENTS**

We determine the coherence time $T_2$ of the RF of the QDs from first-order correlation function measurements using
a Michelson interferometer. As said in the main text, the contrast of the interference fringes is fitted by a pseudo-Voigt profile (see Eq. (3)) which allows to extract $T_2$ and the inhomogeneous contribution $\eta$.

\[
f(t) = (1 - \eta)e^{-|t|/T_2} + \eta e^{-|t/T_2|^2}
\]

This inhomogeneous contribution gives rise to a Gaussian shape of the interference contrast and reflects the slight
energy shift of the RF because of fluctuating charges trapping in the vicinity of the QD [3, 4]. Figure 6 shows the
results of Fourier transform spectroscopy (FTS) for the three QDs mentioned in the main text. The errors on $T_2$
are of the order of $10 - 15$% for QD 1 and 3 of the order of $5$% for QD 2. They are mainly due to the path-length
difference between the two arms of the interferometer that should be large compared to $cT_2$, $c$ being the light velocity.
For the same reason, the errors on $\eta$ are of the order of 0.1.

**SECOND ORDER CORRELATION FUNCTION AND TPI VISIBILITY MEASUREMENTS**

**Second order correlation function for HBT experiment**

As said in the main text, the unusual shape of the histogram of coincidences for HBT experiment is due to the fact
that the single photons pass through the two arms of the Mach-Zehnder interferometer used for the HOM setup (see
Fig. 1a in the main text). For this experiment, the laser delivers single 3-ps pulses at 82 MHz. The contribution to
FIG. 6. Fourier transform spectra for the three studied QDs. The acquisition time for each position of the delay line is 1 s which means that we average all decoherence processes that take place at shorter time scale. The red solid lines are fits using Eq. (3).

the coincidences histogram at 0 and ±3 ns is attributed to the remaining scattered laser. Indeed, the peak labelled 2 (at zero delay) in Fig 2a of the main text, corresponds to coincidences of photons taking the same path of the Mach-Zehnder interferometer. The peaks labelled 1 and 3 (at ±3 ns delay) are due to coincidences of photons taking each a different path of the Mach-Zehnder interferometer.

We measure $g^{(2)}_{\text{HOM}}$ by normalizing the three central peaks integrated intensity by the averaged integrated intensity of the two three-peaks bunches at (12, 9, and 15 ns) and (-12, -9, and −15 ns). The red solid-line corresponds to a multi-exponential decay fit which allows to take into account the overlapping of the different peaks.

TPI visibility

To extract a quantitative value of the second order correlation function at zero delay for HOM experiment $g^{(2)}_{\text{HOM}}$, we follow a procedure close to the one describes in ref. [5]. This method has the advantage to take into account the presence of the remaining scattered laser (obtained by HBT measurements) and the imperfections of the experimental setup. The coincidence histograms for HBT and HOM experiments are fitted by a multi exponential decay allowing to extract the value of $A_{i}^{\text{HBT}}$ and $A_{i}^{\text{HOM}}$ ($i \in \{1, 2, 3\}$) which are respectively the areas of the peak labeled $i$ in Figure 2a and b of the main text. In the case of an ideal experimental setup and for perfect single photon emitter, the value of the second order correlation function at zero delay for HOM experiment would be given by:

$$g^{(2)}_{\text{HOM}}(0) = \frac{A_{i}^{\text{HOM}}}{A_{1}^{\text{HOM}} + A_{3}^{\text{HOM}}}$$

The first deviation to the ideal case describes above is due to the presence of the remaining scattered laser mixed to the luminescence of the QD and send into the Mach-Zehnder interferometer. The areas of the peak 1, 2 and 3 are then corrected as follow:

$$B_{i} = A_{i}^{\text{HOM}} - 2 \frac{t_{i}^{\text{HOM}}}{t_{\text{HBT}}} A_{i}^{\text{HBT}}$$

where $t_{\text{HBT}}$ and $t_{\text{HOM}}$ are the acquisition times for HBT and HOM experiments respectively. This correction shown in Figure 7 allows to remove from the HOM histogram the laser/QD coincidences. The factor 2 comes to the fact that for HOM experiment two pulses are sent into the interferometer. This procedure is equivalent to the presence of the terms proportional to $2g$ in equation 2 of ref. [5] without assuming that the parameter $g$ characterises the two-photon emission probability but only characterising the multi-photon emission probability. Note that these two procedures neglect the coincidences due to the laser between the two pulses.

The second deviation to the ideal case is due to the contrast of the Mach-Zehnder interferometer $C \neq 1$ and not perfect 50/50 fibered beam splitter (FBS). In our case we measured, $R = 0.430 \pm 0.005$, $T = 0.570 \pm 0.005$ and $C^{2} = 0.98 \pm 0.02$. The value of the TPI visibility is then given by

$$V_{\text{TPI}} = \frac{R^{2} + T^{2}}{2RTC^{2}} (1 - g^{(2)}_{\text{HOM}})$$
FIG. 7. (a) Fits for the three central peaks for HOM (solid line) and HBT (dashed line) experiments for QD 1 at $T = 4$ K presented in Fig. 2 of the main text. For the HBT contribution, the fits have been multiplied by the factor $2t_{\text{HOM}}/t_{\text{HBT}}$. (b) Result of Eq. 5 allowing to take into account the remaining laser background.

EFFECTS OF THE PRESENCE OF AN HE-NE LASER ON THE TPI VISIBILITY

Let us notice that for QD1 we used a few nW of an additional He-Ne laser to enhance the RF while for QD2 and QD3 this non resonant laser allows to recover the RF [2, 6]. We show here that this He-Ne laser has no influence on the measured TPI visibility, by presenting results on QD1 at 4 K without the He-Ne laser (see Fig. 8). We measured $V_{\text{TPI}} = 0.80 \pm 0.05$ whereas $V_{\text{TPI}} = 0.79 \pm 0.03$ with a low power of He-Ne laser (see Fig. 2 of the main text).

FIG. 8. Second-order correlation measurements for QD1 at 4 K without He-Ne laser to enhance the RF for $t_{\text{HBT}} = 8500$ s and $t_{\text{HOM}} = 5200$ s. (a) Coincidences histogram for HBT experiment, we extract $g^{(2)}_{\text{HBT}} = 0.14 \pm 0.02$. (b) Coincidences histogram for TPI experiment. After correction by the remaining laser background, we obtain $V_{\text{TPI}} = 0.80 \pm 0.05$.

QUADRATIC ELECTRON-PHONON COUPLING

We start by consider the general electron-phonon coupling Hamiltonian $H_{\text{e-\text{ph}}} = H_0 + H_I$. Here $H_0 = \sum_m \omega_m |m\rangle\langle m| + \sum_k \nu_k b_k^\dagger b_k$ is the energy of the uncoupled QD and phonon environment, where $|m\rangle$ is the $m^{th}$ electron-hole states of the QD and $\omega_m$ is its energy. The $k^{th}$-mode of the phonon environment is described by the creation and annihilation operators, $b_k^\dagger$ and $b_k$ respectively, with frequency $\nu_k$. For linear electron-phonon coupling [7], the interaction Hamiltonian is given by:

$$H_I = \sum_k M_k \rho(k)(b_k^\dagger + b_k),$$

where $M_k = k\nu(k)/\sqrt{2\rho V\nu_k}$ is the phonon matrix element, here $\rho$ is the mass density of the material, $V$ is the phonon quantisation volume, and $\nu(k)$ is the deformation potential energy of the phonons. We have also defined $\rho(k) = \int d^3r e^{ik\cdot r} \rho(r)$ as the Fourier transform of the particle density operator $\rho(r) = \sum_{jj'} \psi_j^\dagger(r) \psi_{j'}(r)c_{j'}^\dagger c_j$, where $c_j^\dagger$ is the fermionic creation operator of the $j^{th}$-electronic state, and $\psi_j(r)$ is its corresponding wavefunction.

This Hamiltonian couples all electronic states together through phonon transitions. To simplify our dynamical
description of the system we follow Muljarov and Zimmerman [8]. This involves removing the off-diagonal transitions to higher lying excited states in the QD through an effective Hamiltonian approach [9], restricting our description to a TLS system with ground state $|0\rangle$ and the single exciton state $|X\rangle$, with splitting $\omega_X$. This results in a Hamiltonian that is quadratic in the electron-phonon coupling [8]:

$$
H = \omega_X |X\rangle \langle X| + |X\rangle \langle X| \left[ \sum_k g_k \left( b_k^\dagger + b_k \right) + \sum_{k,k'} f_{k,k'} \left( b_k^\dagger + b_k \right) \left( b_{k'}^\dagger + b_{k'} \right) \right] + \sum_k \nu_k b_k^\dagger b_k. \tag{8}
$$

The coupling constants are given by:

$$
g_k = \langle X|M_k g(k)|X\rangle \quad \text{and} \quad f_{k,k'} = \sum_{m>1} \frac{\langle X|M_k g(k)|m\rangle \langle m|M_k g(k')|X\rangle}{\omega_X - \omega_m}, \tag{9}
$$

where $f_{k,k'}$ contains a sum over the states $|m\rangle$ describing the electronic configurations of the QD. We can gain analytic expressions for these coupling constants by assuming the electron/holes in the QD are in an isotropic harmonic potential, and the QD has mirror symmetry (i.e. the electron and hole experience the same confinement potential).

The linear coupling takes the standard form $g_k = (2\nu c_s)^{-1/2} \sqrt{\kappa}(D_e - D_h)e^{-c_s k^2/2\nu c_s^2}$, where we have assumed linear dispersion for the phonons with speed of sound $c_s$, and we have introduced the cut-off frequency of the phonon environment $\nu_c$.

For the quadratic coupling, we consider only virtual transition between the lowest excited state and the first manifold of next excited states. This yields:

$$
f_{k,k'} = -\sum_{j \in \{x,y,z\}} \frac{c_s}{2\nu \nu_c^2} \left( \frac{D_e^2}{\Delta_e} + \frac{D_h^2}{\Delta_h} \right) \frac{\sqrt{k}\sqrt{k'}}{\nu_c^2} e^{\frac{-c_s^2 k^2 + k'^2}{\nu_c^2}}, \tag{10}
$$

where $k_j$ are the Cartesian components of the wavevector.

### POLARON TRANSFORMATION AND MASTER EQUATION

To derive an equation of motion for the above Hamiltonian, we make use of the polaron transformation, $U_p = |X\rangle \langle X| e^S + |0\rangle \langle 0|$, where $S = \sum_k \nu_k^{-1} g_k (b_k^\dagger - b_k)$ [10]. Applying this to the Hamiltonian in Eq. 8 we obtain:

$$
H_V = \left( \omega_X - \sum_k \frac{g_k^2}{\nu_k} \right) |X\rangle \langle X| + \sum_k \nu_k b_k^\dagger b_k + |X\rangle \langle X| \sum_{k,k'} f_{k,k'} (b_k^\dagger + b_k)(b_{k'}^\dagger + b_{k'}), \tag{11}
$$

where any residual linear coupling is zero due to spherical symmetry. We now wish to use this Hamiltonian to derive a second-order master equation for the density operator of the QD, $\chi(t)$, which simply takes a pure dephasing form:

$$
\frac{\partial \chi(t)}{\partial t} = -i [\tilde{\omega}_X |X\rangle \langle X|, \chi(t)] + \Gamma \mathcal{L}_o[\chi(t)] + 2\gamma_{pd} \mathcal{L}_{o^\dagger o}[\chi(t)], \tag{12}
$$

where $\tilde{\omega}_X = \omega_X - \sum_k g_k^2/\nu_k$ is the polaron shifted resonance of the TLS, and $\mathcal{L}_o[\chi] = \mathcal{O} \mathcal{O}^\dagger - \{ \mathcal{O}^\dagger \mathcal{O}, \chi \}/2$. Notice that have introduced a second dissipator to account for the optical transition in the QD with lifetime $\Gamma$. The pure dephasing rate is then given by $\gamma_{pd} = \text{Re} \sum_{k,k'} \int_0^\infty d\tau f_{k,k'} |\mathcal{B}_k(\tau)\mathcal{B}_{k'}(\tau)\mathcal{B}_0(0)\mathcal{B}_{k'}(0)|$, where $\mathcal{B}_k(\tau) = b_k^\dagger e^{i\omega_{kk} \tau} + b_k e^{-i\omega_{kk} \tau}$.

We can make a technical simplification by assuming that the dominant contribution to the virtual phonon scattering occurs due to phonons scattering into different modes, that is only scattering processes where $k \neq k'$. This allows us to factorise the $k$ and $k'$ components such that:

$$
\gamma_{pd} = \frac{\pi}{c_s} \frac{V^2}{(2\pi)^6} \int d^3k \int d^3k' |f(k,k')|^2 \left\{ n(k') n(k+1) \delta(k-k') + n(k) n(k'+1) \delta(k-k') \right\}, \tag{13}
$$

where we have taken the continuum limit and used the definition of the $\delta$-function, $\delta(x) = \pi^{-1} \text{Re}[\int_0^\infty d\tau e^{i\omega \tau}]$. We have also defined the bosonic occupation number $n(k) = [\exp(-\beta \nu_k) - 1]^{-1}$, where $\beta^{-1} = k_B T$ is the thermodynamic
where we have assumed linear dispersion to write the above in terms of frequency, and we have defined the constants $\alpha = [4\pi^2 \gamma_0^5]^{-1}(D_e - D_h)^2$ and $\mu = \pi (D_e - D_h)^{-4}(\Delta_e^{-1} D_e^2 + \Delta_h^{-1} D_h^2)^2$.

**INDISTINGUISHABILITY IN THE POLARON FRAME**

We now wish to calculate the indistinguishability/visibility of two photon interference. To do so, we use the definition of the indistinguishability in frequency space \[20\]:

$$\mathcal{I} = P^{-2} \int_{-\infty}^{\infty} d\omega \int_{-\infty}^{\infty} d\nu |S(\omega, \nu)|^2,$$  

(15)

where $S(\omega, \nu) = \langle \hat{E}^\dagger(\omega)\hat{E}(\nu) \rangle = \int_0^\infty dt_1 \int_0^\infty dt_2 g^{(1)}(t_1, t_2) e^{-i\omega t_1} e^{i\nu t_2}$ is the generalised two-colour spectrum, $g^{(1)}(t_1, t_2) = \langle \hat{E}^\dagger(t_1)\hat{E}(t_2) \rangle$ is the first-order correlation function, and $P = \int_{-\infty}^{\infty} \langle \hat{E}^\dagger(\omega)\hat{E}(\omega) \rangle d\omega$ is the total light emitted from the QD. $\hat{E}(t_1)$ and $\hat{E}(\omega)$ are the Heisenberg picture field operators written in the time- and frequency-domain respectively.

In order to relate the detected field operators to the emission of the QD, we must account for emission via the phonon environment \[11, 12\]. With this definition the first order correlation function for the QD becomes

$$\langle \hat{E}^\dagger(t_1, \tau) = 2\pi B(\tilde\Gamma_{c}(\tau)B_{c})(\sigma^\dagger(t + \tau)\sigma(t)), \rangle$$  

which the first term describes the phonon sideband, and the second the zero phonon line contribution to the spectra \[11\]. Using the quantum regression theorem, in addition to the master equation given in Eq. \[12\] we can write an analytic form for the first order correlation function $g^{(1)}(t, \tau) = 2\pi B\tilde{\Gamma}_c^2 G(\tau) \exp(-\Gamma t - (1 + 2\gamma_{pd})\tau)/2$, where $G(\tau) = \exp(\varphi(\tau))$ is the phonon correlation function, $B = \exp(-\varphi(0)/2)$ is the Franck-Condon factor, and $\varphi(\tau) = \int_0^\infty d\nu\nu^2 J(\nu)$ (coth $\nu/2k_BT \cos(\nu\tau) - i \sin(\nu\tau)$). We have introduced the spectral density $J(\nu) = \sum_k |g_k|^2 (\nu - \nu_k) = \alpha/n^2 \exp(-\nu^2/\sigma^2)$.

By recognising that phonon processes occur on a time scale much faster than optical transitions, we can separate the phononic and photonic contributions \[11\] in the generalised two colour spectrum such that $S(\omega, \nu) = S_{ZPL}(\omega, \nu) + S_{SB}(\omega, \nu)$, where $S_{ZPL}(\omega, \nu) = B^2 h(\omega) h(\nu) \Gamma(i(\omega-\nu) + \Gamma + 2\gamma) \left[ \left( \frac{1}{2} \Gamma + \gamma \right) - i\nu \right] \left[ \left( \frac{1}{2} \Gamma + \gamma \right) + i\omega \right]$, which reduces to a Lorentzian filtered Lorentzian line shape when $\omega = \nu$. The term associated with the sideband is given by $S_{SB}(\omega, \nu) = S_{SB}(\omega, \nu) + S_{SB}^G(\omega, \nu)$, with

$$S_{SB}(\omega, \nu) \approx B^2 h^2(\omega) h(\nu) \int_0^\infty dt\rho^{(1)}(\nu, \nu) g^{(1)}(t, 0) \int_0^\infty d\tau (\tilde{\Gamma} - 1)e^{-i\omega \tau} = \frac{\Gamma B^2 h^2(\omega) h(\nu)}{\Gamma - i(\nu - \omega)} S_{PH}(\omega)$$  

(16)

where we have defined the sideband spectrum $S_{PH}(\omega) = \int_{-\infty}^{\infty} d\tau (\tilde{\Gamma} - 1)e^{-i\omega \tau}$.

The phonon sideband is purely incoherent, thus it does not contribute to the numerator in the indistinguishability. Furthermore, making the approximation that the cavity is flat across relevant ZPL features, we can set $h(\omega) \approx h(0)$, and we therefore obtain $\mathcal{I} \approx 4\pi^2 h(0)^2 P_{ZPL}^2 \mathcal{F}^2 \Gamma[1 + 2\gamma_{pd}]^{-1}$. Using again that the cavity is much broader than the ZPL, we approximate $P \approx |h(0)|^2 P_{ZPL} + 2P_{SB}$, where $P_{ZPL} = \int_{-\infty}^{\infty} S_{ZPL}(\omega, \omega) d\omega = 2\pi B^2$ is the power in the ZPL, and $P_{SB} = 2\pi(1 - B^2)$ is the power in the phonon sideband in the absence of a filter, while $\mathcal{F} = P_{SB}^{-1} \int_{-\infty}^{\infty} S_{SB}(\omega, \omega) d\omega$ is the fraction of the phonon sideband not removed by the filter or cavity. Putting this together, we obtain:

$$\mathcal{I} = \frac{\Gamma}{\Gamma + 2\gamma_{pd}} \left( \frac{|h(0)|^2 B^2}{|h(0)|^2 B^2 + 2\mathcal{F}[1 - B^2]} \right)^2,$$  

(17)

which is Eq. \(2\) in the main text.
FIG. 9. Plot of the measured TPI visibility for QD2 as a function of temperature (black stars). The data are fitted with Eq. 2 of the main text. The full expression is used to produce the solid red curve and the dash purple curve shows only the effect of the phonon sideband on the TPI visibility. The phonon parameters extracted from the fitting procedure are \( \alpha = 0.0071 \, \text{ps}^2 \), \( \nu_c = 11.9 \, \text{ps}^{-1} \), and \( \mu = 5.6 \times 10^{-4} \, \text{ps}^2 \).

For the fitting procedure, we can simplify this expression further by assuming weak electron–phonon coupling. Here, the phonon correlation function \( G(\tau) = \exp(\phi(\tau)) \approx 1 - \phi(\tau) \), where we have considered only terms first-order in \( \alpha \). The fraction of sideband unfiltered is given by

\[
F = P_{SB}^{-1} \int_{-\infty}^{\infty} d\omega |h(\omega)|^2 \text{Re}[S_{PH}(\omega)].
\]

We can now carry out the Fourier transform of the phonon correlation function analytically, such that

\[
\text{Re}[S_{PH}(\omega)] \approx \text{Re} \left[ \int_0^\infty \phi(\tau) e^{-i\omega \tau} d\tau \right] = \pi \alpha e^{-\omega^2/\nu_c^2} \left( \coth \left( \frac{\beta \omega}{2} \right) - 1 \right),
\]

where we have invoked the definition of the \( \delta \)-function. Using this expression, the filtered fraction becomes:

\[
F = P_{SB}^{-1} \int_{-\infty}^{\infty} |h(\omega)|^2 \omega e^{-\omega^2/\nu_c^2} \coth \left( \frac{\beta \omega}{2} \right) d\omega.
\]

TPI VISIBILITY FOR QD2

We present in Fig. 9 temperature dependent measurement of the TPI visibility for QD2. These experiments were realized in another cryostat with a minimum temperature of 7 K which explain the lack of data at low temperature. As for QD1 we use Eq. 2 of the main text to fit the experimental data with the same fitting procedure. Again, we observe very good agreement between the experimental data and the microscopic model. We extract the following parameters for QD2: \( \alpha = 0.0071 \, \text{ps}^2 \), \( \nu_c = 11.9 \, \text{ps}^{-1} \), \( \mu = 5.6 \times 10^{-4} \, \text{ps}^2 \) and \( F \sim 0.1 \) at \( T \sim 7 \, \text{K} \). We notice the importance of taking into account virtual transitions to higher lying QD states in order to fit the rapid loss of the visibility for temperature higher than 10 K.

The blue squares represent the expected TPI visibility using the measured values of \( T_1 \) and \( T_2 \) and random dephasing processes by phonons and charges [13]. We clearly see that these values do not reproduce the observed behaviour of the TPI visibility, confirming that TPI and FTS using a Michelson interferometer do not probe the same dephasing processes.

\[ \text{Re} \left[ S_{PH}(\omega) \right] \approx \text{Re} \left[ \int_0^\infty \phi(\tau) e^{-i\omega \tau} d\tau \right] = \pi \alpha e^{-\omega^2/\nu_c^2} \left( \coth \left( \frac{\beta \omega}{2} \right) - 1 \right), \]

\[ F = P_{SB}^{-1} \int_{-\infty}^{\infty} |h(\omega)|^2 \omega e^{-\omega^2/\nu_c^2} \coth \left( \frac{\beta \omega}{2} \right) d\omega. \]

\[ F \sim 0.1 \text{ at } T \sim 7 \, \text{K}. \]

\[ \text{We present in Fig. 9 temperature dependent measurement of the TPI visibility for QD2. These experiments were realized in another cryostat with a minimum temperature of 7 K which explain the lack of data at low temperature. As for QD1 we use Eq. 2 of the main text to fit the experimental data with the same fitting procedure. Again, we observe very good agreement between the experimental data and the microscopic model. We extract the following parameters for QD2: } \alpha = 0.0071 \, \text{ps}^2, \nu_c = 11.9 \, \text{ps}^{-1}, \mu = 5.6 \times 10^{-4} \, \text{ps}^2 \text{ and } F \sim 0.1 \text{ at } T \sim 7 \, \text{K}. \text{ We notice the importance of taking into account virtual transitions to higher lying QD states in order to fit the rapid loss of the visibility for temperature higher than 10 K.}

\[ \text{The blue squares represent the expected TPI visibility using the measured values of } T_1 \text{ and } T_2 \text{ and random dephasing processes by phonons and charges [13]. We clearly see that these values do not reproduce the observed behaviour of the TPI visibility, confirming that TPI and FTS using a Michelson interferometer do not probe the same dephasing processes.} \]