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Highly tuneable hole quantum dots in Ge-Si core-shell nanowires

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We define single quantum dots of lengths varying from 60 nm up to nearly half a micron in Ge-Si core-shell nanowires. The charging energies scale inversely with the quantum dot length between 18 and 4 meV. Subsequently, we split up a long dot into a double quantum dot with a separate control over the tunnel couplings and the electrochemical potential of each dot. Both single and double quantum dot configurations prove to be very stable and show excellent control over the electrostatic environment of the dots, making this system a highly versatile platform for spin-based quantum computing. *Published by AIP Publishing.* [<http://dx.doi.org/10.1063/1.4963715>]

For spin-based quantum computing,¹ increasing research efforts have focused in recent years on C, Si, and Ge²⁻⁴ because they can be isotopically enriched to only contain nuclei with zero spin^{5,6} and thus exhibit exceptionally long spin lifetimes.^{7,8} The one-dimensional character of Ge-Si core-shell nanowires leads to unique electronic properties in the valence band, where heavy and light hole states are mixed.⁹⁻¹¹ Early experiments in Ge-Si core-shell nanowires include experiments on double quantum dots¹² and spin relaxation times.¹³ The band mixing causes an enhanced Rashba-type spin-orbit interaction (SOI),¹¹ which can be exploited for efficient spin manipulation.¹⁴ Therefore, Ge-Si core-shell nanowires are an ideal platform for future quantum computation applications.

In this letter, we define single quantum dots of several lengths in a Ge-Si core-shell nanowire. We controllably split longer quantum dots up into double quantum dots with tuneable interdot tunnel coupling. Both single and double quantum dots show an exceptional degree of measurement stability.

We will discuss measurements in two different devices *D1* and *D2* (see Fig. 1) on two different chips, which have been fabricated in the same way: A p⁺⁺-doped Si substrate is covered with 200 nm SiO₂, on which six bottom gates *g1*–*g6* with 100 nm pitch are patterned with electron beam lithography (EBL). Before metallization of the bottom gates, a 13 s buffered hydrofluoric acid dip etches 20 nm deep trenches into the SiO₂, so that the bottom gates (approximately 20 nm thick) are sunken into the SiO₂ for an improved planarity. The gates are covered with 10 nm Al₂O₃ grown with atomic layer deposition at 100 °C. Two single nanowires with a Si shell thickness of 2.5 nm and a Ge core radius of 8 nm (*D1*) and 9 nm (*D2*) are deterministically placed on top of the gate structure with a micromanipulator. Based on transmission electron microscopy studies of similar

wires, both the core and the shell are monocrystalline, and their axis is likely pointed along the ⟨110⟩ crystal axis.¹⁵ Subsequently, we define ohmic contacts to the nanowires and gate contacts made of Ti/Pd (0.5/50 nm) with EBL. The nanowire parts above the bottom gates are at no point exposed to the electron beam, preventing carbon deposition and introduction of defects into the otherwise defect-free Ge core. All measurements are performed using dc electronic equipment in a dilution refrigerator with a base temperature of 8 mK. A bias voltage V_{SD} is applied to source, and the current I is measured at the drain contact. An effective hole temperature of $T_{\text{hole}} \approx 30$ mK has been determined in one of the devices by measuring the temperature dependence of the Coulomb peak width.^{16,17}

By using different gates to induce tunnel barriers, we can form quantum dots in our nanowire with lengths varying from very long quantum dots (using *g1* and *g6*) to very short dots (using adjacent gates). This flexible quantum dot length together with a tuneable tunnel coupling between the quantum dot and the reservoirs is a great improvement compared to using lateral heterostructures,^{18,19} or Schottky barriers at the nanowire-metal interface with the contacts.^{20,21}

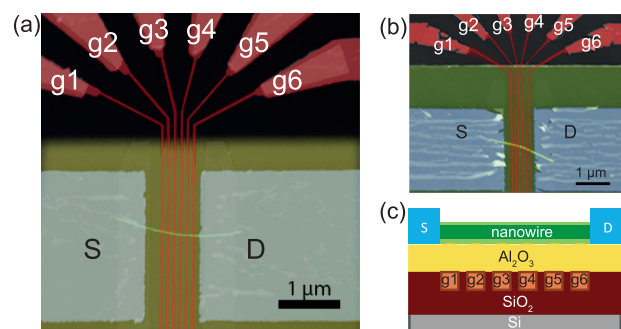


FIG. 1. False-colour AFM image of device (a) *D1*, and (b) *D2*. (c) Schematic cross-section displaying the p⁺⁺-doped Si substrate (grey) with 200 nm of SiO₂ (dark red), six bottom gates *g1*–*g6* (light red), each 35 nm wide and with a pitch of 100 nm. The bottom gates are buried under 10 nm of Al₂O₃ (yellow), on top of which the nanowire is deposited (green) and ohmic contacts (0.5/50 nm Ti/Pd, blue) are defined.

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We assume the length of our gate-defined quantum dots to be the distance between the inner edges of the barrier gates. Using a gate width of ~ 40 nm, this results in quantum dot lengths of ~ 60 nm for adjacent barrier gates, ~ 160 nm for barrier gates with one plunger gate in between, ~ 260 nm for two plunger gates, ~ 360 nm for three plunger gates, and ~ 460 nm for four plunger gates, i.e., we are able to tune the dot length over almost an order of magnitude.

In Figs. 2(a)–2(e), we plot $dI/dV \equiv dI/dV_{SD}$ versus V_{SD} and the voltage on the plunger gate V_P . The formation of quantum dots of five different lengths is reflected in the clear Coulomb diamonds. The shortest quantum dot is formed in device *D1* [Fig. 2(a)]. The quantum dots formed with one up to four plunger gates are formed in both devices *D1* and *D2*. Figs. 2(b)–2(e) display bias spectroscopies of quantum dots formed in *D2*.

We extract the respective charging energies E_C for both devices from the Coulomb diamond height and find a decreasing E_C from 18.3 meV to 4.2 meV, inversely proportional to the increasing dot length [see Fig. 2(f) and Table I]. Since E_C is linked to the total capacitance C of the quantum dot via $E_C = e^2/C$,²² C is directly proportional to the quantum dot length. E_C and C are highly consistent for the two devices.

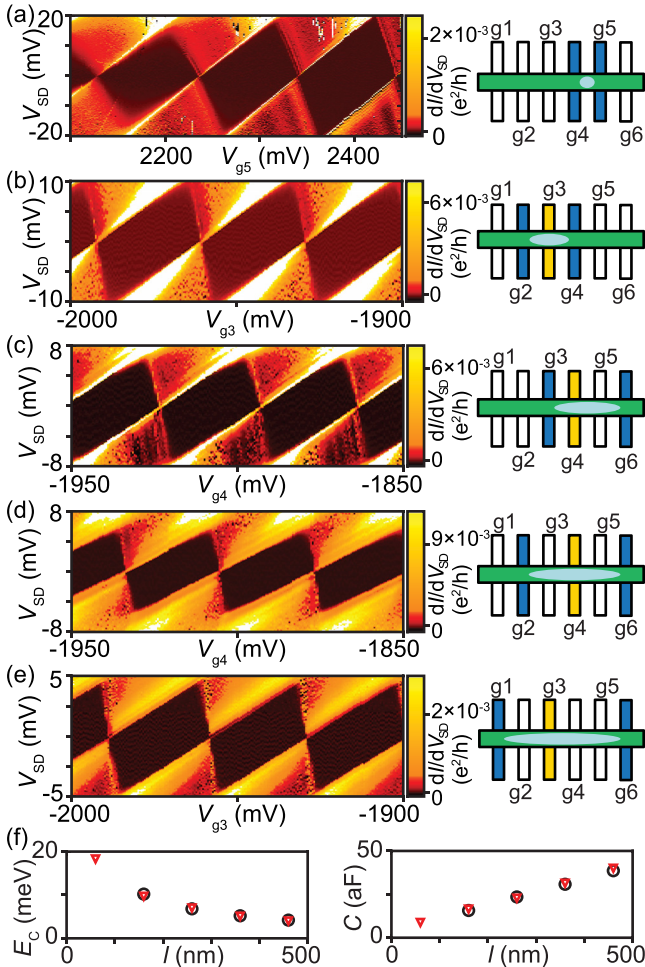


FIG. 2. Bias spectroscopy of gate-defined single quantum dots formed with (a) zero, (b) one, (c) two, (d) three, and (e) four gates between the barrier gates (indicated in blue). (a) is measured in *D1*, (b)–(e) in *D2*. (f) Charging energy E_C (left) and total capacitance C (right) of the dot plotted versus the dot length l . Red triangles measured in *D1*, black circles in *D2*.

TABLE I. Parameters for electrostatically defined quantum dots of varying length as extracted from Figs. 2(a)–2(e).

l (nm)	E_C (meV)	C (aF)	ΔV_g (mV)	C_g (aF)
60	18.3(2)	8.8(2)	104(1)	1.54(2)
160	10.2(2)	15.7(3)	31.5(2)	5.09(3)
260	6.8(2)	23.5(5)	29.6(4)	5.41(7)
360	5.2(1)	30.8(6)	28.6(4)	5.63(8)
460	4.2(1)	38.6(9)	29.7(4)	5.39(7)

For the quantum dot configurations with a dedicated plunger gate [Figs. 2(b)–2(e)], E_C as well as the shape of the Coulomb diamonds stay constant over several charge transitions, reflecting the validity of the constant interaction model. In Fig. 2(a), E_C increases significantly from 16.7 meV to 20.0 meV and also the slopes of the Coulomb diamond edges change. We attribute this to using the right barrier gate as a plunger, leading to a decreasing dot size and changing capacitive couplings to this barrier gate and the adjacent reservoir. Therefore, the constant interaction model is not valid in this configuration, and the tuneability of the quantum dot is limited compared to the longer quantum dots with a dedicated plunger gate. We extract the values for E_C and C for the zero-plunger configuration from the middle Coulomb diamond, for which they are in line with those for the longer dots.

The constant charging energies over several Coulomb diamonds in Figs. 2(b)–2(e) are accompanied by constant Coulomb peak spacings ΔV_g at $V_{SD} = 0$, indicating a constant gate capacitance C_g over several charge transitions, another indication for the validity of the constant interaction model. If we now compare the plunger gate capacitances between Figs. 2(b) and 2(e), we find them to be all very similar, (~ 5.5 aF), while the total capacitance increases linearly by ~ 7.5 aF per additional plunger gate [see Fig. 2(f)]. The discrepancy of ~ 2 aF can be explained by the finite capacitance of the global back gate which increases with the dot length and the change in the self-capacitance of the quantum dot. The linearly increasing total capacitance indicates equal coupling of all gates, consistent with the gate geometry (equal width and distance to the nanowire). In Fig. 2(f), we also plot E_C and C for quantum dots formed in *D1* with at least one plunger gate alongside the data for *D2*. The consistency between the data therefore demonstrates a high degree of control over the electrostatic environment of the gate-defined quantum dot.

Tunable double quantum dots are essential for spin readout via Pauli spin blockade.²³ For a fully tuneable double quantum dot, we need five gates: three barrier gates to form tunnel barriers, and two plunger gates to tune the electrochemical potential of each dot separately. We use device *D1* starting from a situation equivalent to Fig. 2(d) and increase the voltage on the middle gate V_{g4} . When approaching the pinch-off voltage, a tunnel barrier is formed and the single quantum dot splits up into two tunnel-coupled quantum dots.

The charge stability diagrams at four different V_{g4} are plotted in Fig. 3(a). We keep the outer barrier gates at constant voltages ($V_{g2} = 2490$ mV, $V_{g6} = 2940$ mV) and plot the current at a fixed $V_{SD} = 1$ mV. For $V_{g4} = 0$ mV, we observe the typical stability diagram of a single quantum dot.²⁴ The

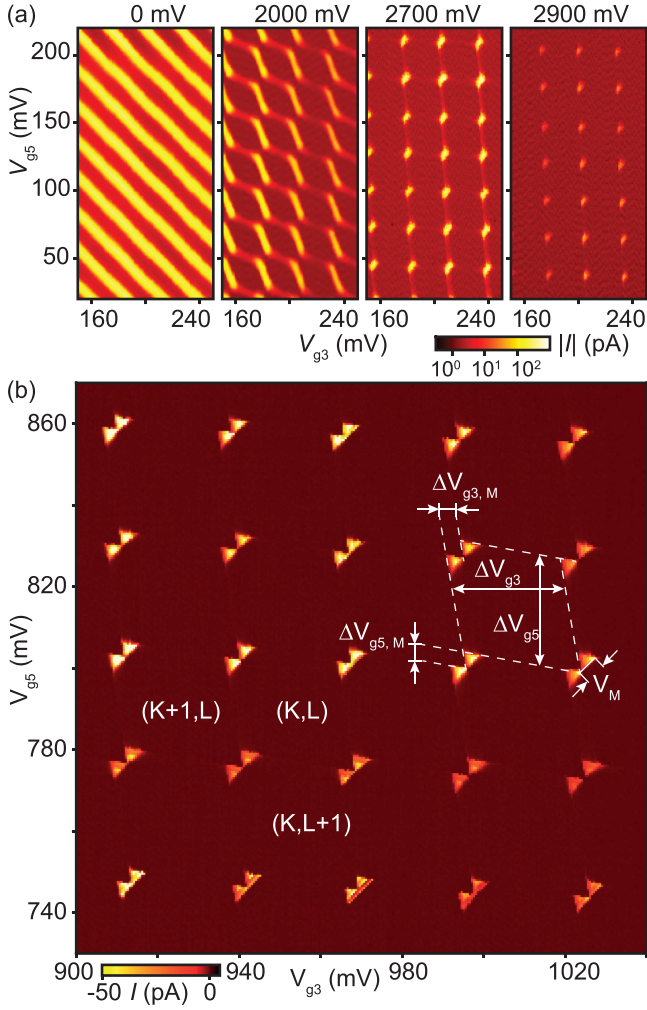


FIG. 3. (a) Charge stability diagrams with current I plotted versus V_{g3} and V_{g5} for varying voltages on $g4$ at fixed $V_{g2} = 2490$ mV and $V_{g6} = 2940$ mV. (b) Charge stability diagram of a double quantum dot at $V_{g2} = 2500$ mV, $V_{g4} = 2100$ mV, and $V_{g6} = 3180$ mV. (k, l) denote the charge occupation numbers on the left (k) and right dot (l). All measurements performed on device DI .

spacing of the diagonal, parallel lines of finite current along the respective plunger gate axis is directly related to the capacitance between the quantum dot and this gate: $C_g = e/\Delta V_g$. We observe $\Delta V_g \approx 27$ mV for both $g3$ and $g5$, i.e., both gates have the same capacitance $C_g \approx 5.8$ aF to the quantum dot. This indicates that the quantum dot indeed stretches over the whole distance between the tunnel barriers above gates $g2$ and $g6$ and is also in agreement with the gate capacitances in Table I.

At $V_{g4} = 2000$ mV, the straight lines have evolved into a regular honeycomb pattern with two distinct slopes that form the long edges of each honeycomb, indicating the formation of a strongly coupled double quantum dot.²⁴ From the distance between adjacent parallel lines, we extract the voltages needed to add a hole to the left (right) dot ΔV_{g3} , (ΔV_{g5}), and calculate the corresponding capacitances. For the left dot, we find $\Delta V_{g3} = 25.1(5)$ mV, and for the right dot, $\Delta V_{g5} = 26.1(5)$ mV, resulting in gate capacitances $C_{g3} = 6.3(2)$ aF and $C_{g5} = 6.1(2)$ aF. The sets of honeycomb edges representing the addition of a hole to either the left or the right dot are both significantly slanted because of

the mutual capacitive coupling C_M between the two dots that leads to a separation between the two triple points.²⁴ We express this shift in terms of gate voltages and find $\Delta V_{g3,M} = 9.2(5)$ mV and $\Delta V_{g5,M} = 10.9(5)$ mV. Using the expression $C_{g,M} = C_g \Delta V_{g,M} / \Delta V_g$,²⁴ we obtain $C_{g3,M} = 2.3(3)$ aF and $C_{g5,M} = 2.5(3)$ aF. A second mechanism affecting the slopes of the honeycomb edges is the finite cross capacitance between $g3$ and the right dot $C_{g3,C}$, and $g5$ and the left dot $C_{g5,C}$. This cross capacitance leads to a shift of the triple points along the $g3$ gate axis while changing the charge occupation of the right dot, and along the $g5$ gate axis while changing the number of holes on the left dot. This effect is very weak, and we extract $C_{g3,C} \approx C_{g5,C} \approx 0.1$ aF.

Increasing the voltage on the interdot barrier gate to $V_{g4} = 2700$ mV only slightly changes the gate capacitances to $C_{g3} = 5.9(2)$ aF and $C_{g5} = 5.7(2)$ aF. For the mutual gate capacitances, we find a much stronger relative change to $C_{g3,M} = 0.6(1)$ aF and $C_{g5,M} = 0.7(1)$ aF, which indicates a significantly increased separation of the charge distribution of both dots. The now only faintly visible long edges of the honeycombs also suggest a decreased tunnel coupling to the reservoirs so that cotunnelling is suppressed.²⁵ A finite, but very small cross capacitance of the plunger gates is also observed here, again on the order of 0.1 aF.

A further increase in the interdot barrier gate to $V_{g4} = 2900$ mV completely quenches the cotunnelling current at the long honeycomb edges, so that now transport is only possible at the triple point pairs. This indicates well defined charge states confined in the quantum dots weakly coupled to the reservoirs. Again, we observe a slight decrease of the gate capacitances to $C_{g3} = 5.8(2)$ aF and $C_{g5} = 5.5(2)$ aF, and also the mutual capacitances decrease further to $C_{g3,M} = 0.4(1)$ aF and $C_{g5,M} = 0.4(1)$ aF. All extracted capacitances are summarized in Table II.

In Fig. 3(b), we show a high-resolution stability diagram of a double quantum dot weakly coupled to the reservoirs at $V_{SD} = -1.5$ mV with barrier gate voltages of $V_{g2} = 2500$ mV, $V_{g4} = 2100$ mV, and $V_{g6} = 3180$ mV. Clearly visible is a very regular pattern of 25 bias triangle pairs, from which we extract the gate-to-dot capacitances in the same way as before. We obtain $C_{g3} = 5.9(2)$ aF and $C_{g5} = 5.9(2)$ aF, and mutual capacitances of $C_{M,g3} = 0.9(1)$ aF and $C_{M,g5} = 0.9(1)$ aF. The increased values for $C_{M,i}$ indicate an indeed increased capacitive coupling between the dots.

We extract the charging energies from Fig. 3(b) by relating the bias triangle size to an energy of 1.5 meV. We obtain a charging energy of the left dot $U_1 = 10.6(5)$ meV and of the right dot $U_2 = 9.3(5)$ meV. For the mutual charging energy U_M , we extract $U_M = 1.5(2)$ meV. The size and

TABLE II. Capacitances for increasing voltage on the middle barrier gate $g4$ of an electrostatically defined single ($V_{g4} = 0$ mV) or double quantum dot ($V_{g4} \geq 2000$ mV) as extracted from Fig. 3(a).

V_{g4} (mV)	C_{g3} (aF)	C_{g5} (aF)	$C_{g3,M}$ (aF)	$C_{g5,M}$ (aF)
0	5.8(3)	5.8(3)		
2000	6.3(2)	6.1(2)	2.3(3)	2.5(3)
2700	5.9(2)	5.7(2)	0.6(2)	0.7(2)
2900	5.8(2)	5.5(2)	0.4(1)	0.4(1)

shape of the bias triangles are exceptionally stable over the whole range of the measurement. This underlines the high degree of control over the electrochemical potentials of the quantum dots as well as the tunnel and capacitive couplings. Our devices are therefore exceptionally suitable for direct-transport experiments in comparison to other systems, where tunnel couplings change strongly when changing the charge occupation.^{26–29} Such experiments are relevant, because the applied bias between the two reservoirs serves as an energy scale, which, e.g., allows for the determination of the singlet-triplet splitting³¹ and the Zeeman splitting.³⁰

In summary, we demonstrate a high degree of control over the charge distribution in a double quantum dot. We have changed the mutual capacitances, a measure for the degree of separation of the dots, by a factor of six while keeping the capacitances between the left (right) dot and g^3 (g^5) almost constant. The corresponding charging energies are in agreement with the experiments on single quantum dots of the same length in $D2$.

In conclusion, we have electrostatically formed highly tunable single and double quantum dots inside Ge-Si core-shell nanowires. We can vary the length of the single quantum dots from 60 nm to 460 nm corresponding to charging energies of the quantum dots varying from ~ 18 meV down to ~ 4 meV.

Furthermore, we have split a single quantum dot into a double quantum dot in a controlled way. Our low-cross-capacitance gate design enables us to keep the voltage on the outer barriers constant while varying the interdot barrier, i.e., it is not necessary to retune all gates. All capacitances and charging energies extracted from single and double quantum dot measurements are highly consistent. 25 bias triangle pairs form a very regular pattern in the stability diagram with constant triangle sizes, indicating an exceptional degree of control over the tunnel couplings over a large range of gate voltages.

This combination of tuneability and stability makes Ge-Si core-shell nanowires an ideal platform for further experiments towards quantum computation applications.

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¹D. Loss and D. P. DiVincenzo, *Phys. Rev. A* **57**, 120 (1998).

²E. A. Laird, F. Kueemeth, G. A. Steele, K. Grove-Rasmussen, J. Nygård, K. Flensberg, and L. P. Kouwenhoven, *Rev. Mod. Phys.* **87**, 703 (2015).

³F. A. Zwanenburg, A. S. Dzurak, A. Morello, M. Y. Simmons, L. C. L. Hollenberg, G. Klimeck, S. Rogge, S. N. Coppersmith, and M. A. Eriksson, *Rev. Mod. Phys.* **85**, 961 (2013).

- ⁴M. Amato, M. Palumbo, R. Rurali, and S. Ossicini, *Chem. Rev.* **114**, 1371 (2014).
- ⁵K. M. Itoh, J. Kato, M. Uemura, A. K. Kaliteevskii, O. N. Godisov, G. G. Devyatych, A. D. Bulanov, A. V. Gusev, I. D. Kovalev, P. G. Sennikov, H.-J. Pohl, N. V. Abrosimov, and H. Riemann, *Jpn. J. Appl. Phys.* **42**, 6248 (2003).
- ⁶K. Itoh, W. L. Hansen, E. E. Haller, J. W. Farmer, V. I. Ozogin, A. Rudnev, and A. Tikhomirov, *J. Mater. Res.* **8**, 1341 (1993).
- ⁷J. T. Muhonen, J. P. Dehollain, A. Laucht, F. E. Hudson, T. Sekiguchi, K. M. Itoh, D. N. Jamieson, J. C. McCallum, A. S. Dzurak, and A. Morello, *Nat. Nanotechnol.* **9**, 986 (2014).
- ⁸M. Veldhorst, J. C. C. Hwang, C. H. Yang, A. W. Leenstra, B. de Ronde, J. P. Dehollain, J. T. Muhonen, F. E. Hudson, K. M. Itoh, A. Morello, and A. S. Dzurak, *Nat. Nanotechnol.* **9**, 981 (2014).
- ⁹D. Csontos and U. Zülicke, *Phys. Rev. B* **76**, 073313 (2007).
- ¹⁰D. Csontos, P. Brusheim, U. Zülicke, and H. Q. Xu, *Phys. Rev. B* **79**, 155323 (2009).
- ¹¹C. Kloeffel, M. Trif, and D. Loss, *Phys. Rev. B* **84**, 195314 (2011).
- ¹²Y. Hu, H. O. H. Churchill, D. J. Reilly, J. Xiang, C. M. Lieber, and C. M. Marcus, *Nat. Nanotechnol.* **2**, 622 (2007).
- ¹³Y. Hu, F. Kueemeth, C. M. Lieber, and C. M. Marcus, *Nat. Nanotechnol.* **7**, 47 (2012).
- ¹⁴C. Kloeffel, M. Trif, P. Stano, and D. Loss, *Phys. Rev. B* **88**, 241405 (2013).
- ¹⁵A. Li, S. Conesa-Boj, S. Koelling, M. Brauns, J. Ridderbos, M. A. Verheijen, P. M. Koenraad, F. A. Zwanenburg, and E. P. A. M. Bakkers, “Boosting hole mobility in coherently strained [110]-oriented Ge-Si core-shell nanowires” (unpublished).
- ¹⁶F. Mueller, R. N. Schouten, M. Brauns, T. Gang, W. H. Lim, N. S. Lai, A. S. Dzurak, W. G. van der Wiel, and F. A. Zwanenburg, *Rev. Sci. Instrum.* **84**, 044706 (2013).
- ¹⁷D. Goldhaber-Gordon, J. Göres, M. A. Kastner, H. Shtrikman, D. Mahalu, and U. Meirav, *Phys. Rev. Lett.* **81**, 5225 (1998).
- ¹⁸A. Fuhrer, L. E. Fröberg, J. N. Pedersen, M. W. Larsson, A. Wacker, M.-E. Pistol, and L. Samuelson, *Nano Lett.* **7**, 243 (2007).
- ¹⁹S. Roddaro, A. Pescaglioni, D. Ercolani, L. Sorba, and F. Beltram, *Nano Lett.* **11**, 1695 (2011).
- ²⁰F. A. Zwanenburg, A. A. Van Loon, G. A. Steele, C. E. W. M. Van Rijmenam, T. Balder, Y. Fang, C. M. Lieber, and L. P. Kouwenhoven, *J. Appl. Phys.* **105**, 124314 (2009).
- ²¹H. A. Nilsson, M. T. Deng, P. Caroff, C. Thelander, L. Samuelson, L.-E. Wernersson, and H. Q. Xu, *IEEE J. Sel. Top. Quantum Electron.* **17**, 907 (2011).
- ²²L. P. Kouwenhoven, C. M. Marcus, P. L. McEuen, S. Tarucha, R. M. Westervelt, and N. S. Wingreen, in *Proceedings of the NATO Advanced Study Institute on Mesoscopic Electron Transport* (1997), pp. 105–214.
- ²³K. Ono, D. G. Austing, Y. Tokura, and S. Tarucha, *Science* **297**, 1313 (2002).
- ²⁴W. G. van der Wiel, S. De Franceschi, J. M. Elzerman, T. Fujisawa, S. Tarucha, and L. P. Kouwenhoven, *Rev. Mod. Phys.* **75**, 1 (2002).
- ²⁵S. De Franceschi, S. Sasaki, J. M. Elzerman, W. G. van der Wiel, S. Tarucha, and L. P. Kouwenhoven, *Phys. Rev. Lett.* **86**, 878 (2001).
- ²⁶C. Fasth, A. Fuhrer, M. T. Björk, and L. Samuelson, *Nano Lett.* **5**, 1487 (2005).
- ²⁷H. Liu, T. Fujisawa, Y. Ono, H. Inokawa, A. Fujiwara, K. Takashina, and Y. Hirayama, *Phys. Rev. B* **77**, 073310 (2008).
- ²⁸T. Sand-Jespersen, M. Aagesen, C. B. Soerensen, P. E. Lindelof, and J. Nygard, *J. Vac. Sci. Technol. B* **26**, 1609 (2008).
- ²⁹R. Li, F. E. Hudson, A. S. Dzurak, and A. R. Hamilton, *Nano Lett.* **15**, 7314 (2015).
- ³⁰M. Brauns, J. Ridderbos, A. Li, E. P. A. M. Bakkers, and F. A. Zwanenburg, *Phys. Rev. B* **93**, 121408(R) (2016).
- ³¹M. Brauns, J. Ridderbos, A. Li, E. P. A. M. Bakkers, and W. G. V. D. Wiel, *Phys. Rev. B* **94**, 041411(R) (2016).