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Optical observation of single-carrier charging in type-II quantum ring ensembles

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A high-purity GaSb/GaAs quantum ring system is introduced that provides both strong hole-confinement in the GaSb ring and electron confinement in its GaAs core. The latter is responsible for a reduced inhomogeneous linewidth measured in photoluminescence, in comparison to the previous measurements made on nanostructures with differing morphology in this material system. This allows the resolution of multiple peaks in the photoluminescence due to discrete charging with holes, revealing the mechanism responsible for the excitation-power-induced blueshift. © 2012 American Institute of Physics. [doi:10.1063/1.3688037]

Semiconductor nanostructures1 are appealing for applications in quantum information processing (QIP) (Ref. 2), as they provide an interface between flying- and stationary-qubits.3 Two significant physical limitations are present in most systems: shallow confinement requires low-temperature operation and strong interactions with their surroundings typically result in short coherence times.3,5 GaSb/GaAs structures6 could potentially address these limitations, as they present a deep confining potential for holes7 making room-temperature operation possible,8 and hole-spin can be robust against dephasing.9,10 The type-II nature of this system is non-ideal, however, and has led to inferior optical properties.11 The excitonic physics of type-II quantum dot systems, in which just one of the two carrier types is confined, is known to be strikingly different from that of type-I systems where both are.12 For a neutral multi-exciton state in a type-II nanostructure, the strongly confined carrier type will experience a much greater Coulomb repulsion than would be expected in a type-I system, due to the close proximity of like-, and absence of oppositely, charged carriers. In this letter, the significance of the Coulomb interaction in a type-II system is directly demonstrated by optical measurement of the discrete charging of ensembles of GaSb/GaAs quantum rings with holes.

Molecular-beam-epitaxial (MBE) growth was used to form dislocation-free rings with a very high material contrast, as shown in Fig. 1. The experimental procedure used to control the morphology of GaSb/GaAs nanostructures formed during growth by molecular beam epitaxy was introduced in Ref. 13. MBE growth tends to result in the formation of rings, rather than dots,14 with the final capping, or annealing, temperature influencing the ratio of major/minor axes of the rings that form.15 The primary sample grown for this study comprised of a single layer of GaSb quantum rings capped with 250 nm of GaAs. The ring layer was grown by depositing 2.1 monolayers (MLs) of GaSb at a pyrometer-measured temperature of 490 °C with a growth rate of 0.3 MLs s⁻¹ and a V/III ratio of approximately 10. This layer was capped with a 5 nm layer of GaAs at 430 °C prior to growing the remaining 245 nm of cap at 1 MLs⁻¹ at a temperature of 580 °C. The band-structure of GaSb/GaAs quantum rings is illustrated in Ref. 6. Samples containing layers of rings were also grown under similar conditions for cross-sectional microscopy; both high-resolution transmission electron microscopy (HRTEM) and scanning tunneling microscopy (STM) were employed to assess the structural result of growth (Fig. 1). As the latter technique requires a tunnel current to facilitate local density of state measurements, a low level of background doping (n-type ~10¹⁷ cm⁻³ using GaTe) was introduced to the growth of all layers in the sample. The image in Fig. 1(d) was taken with a tip bias of +0.44 V.

When the rings are measured in cross-section two GaSb-rich lobes are typically seen, with separations varying between 0 and ~25 nm, the value measured is dependent on the position of the section with respect to the center of the ring (Fig. 1(c)). The major radius of the rings is unlikely to influence the heavy-hole’s wave-function in this system, as it is expected to be localized in a relatively small section of the ring. Despite this, the geometry does strongly influence the excitonic properties in the system. STM measurements indicate that the GaAs in the center of the ring is purer, i.e., contains less Sb, than that surrounding it. A local electron density-of-states measurement confirms the presence of a maximum towards the center of the ring, as shown in Fig. 1(d).

Optical measurements on the quantum rings were performed at 1.4 K by placing the nominally undoped sample in a ⁴He cryostat. Stabilized 532 nm continuous-wave laser light was delivered to the sample, via an optical fiber with a large-diameter core, to excite carriers above the GaAs band-gap. Photoluminescence (PL) was collected from an area corresponding to ~2 mm² using a second optical fiber. Microscopy estimated the ring areal number density to be of the order of 1 x 10¹⁰ cm⁻². The sample’s emission was passed into a 0.3 m spectrometer equipped with a Peltier-cooled InGaAs array with which multi-channel wavelength-resolved measurements were made. PL from the rings taken at low (high) excitation power is shown in (the inset of) Fig. 2. Emission from the nanostructures is strong when compared

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to other emitters in the sample, indicating that the rings have good optical properties. The peak emission energy of the rings is below 1 eV demonstrating the very strong confining potential for this material system. The peak also contains clear oscillatory features that were investigated further by monitoring their evolution as a function of the excitation power, as shown in Fig. 3.

The center of mass (CoM) of the emission from the rings blueshifts with increasing pump intensity by more than 50 meV over the five orders of magnitude of power probed here. An excitation-power-induced blueshift is a characteristic feature that has been reported in numerous different quantum-confining type-II systems. A bending of the bandstructure at the interface between the different carrier types and increased charging by the strongly confined carrier are both thought to contribute to this shift. In quantum dots, however, the latter is thought to play a much greater role than the former. The oscillations in the PL were clearly visible at all powers probed.

The quantum ring PL data was analyzed by fitting a series of fixed-width Gaussian sub-peaks to each of the measured spectra, an example of this fitting is illustrated in the inset of Fig. 2. This fit closely matched the experimental data through-out the power-range studied. The sub-peak centers and areas are plotted as a function of power in Figs. 3 (lower panel) and 4, respectively. The width of the Gaussians used in the fitting procedure was 26 meV, and the separation between the sub-peaks was found to increase from 19 meV between the lowest energy sub-peaks to 23 meV between the high energy sub-peaks. The former, i.e., 26 meV, is believed to be an upper limit to the inhomogeneous broadening in the system, while the latter represents the energy required to overcome the repulsive Coulomb interaction when adding a further heavy-hole to the dot; justification for this statement follows.

A simulation of Ga(As)Sb/GaAs quantum dots included the Coulomb potential in an 8-band $k_p$ model. As the dot was filled with holes, the ground-state ($E_1$-$H_1$)
transition was found to blue-shift strongly as a result of the Coulomb interaction. The interpretation of our experimental data is somewhat similar, though the magnitude of the effect is several times greater. Reference 17 studied rather large \((25 \times 25 \times 15 \text{ nm})\) dilute \((25\% \text{ Sb})\) pyramidal dots, whereas, in this work, the hole-confining region is much smaller (see Fig. 1) and the Sb concentration in the ring is close to 100\%. One possible alternative explanation for the sub-peaks is recombination from the excited states of the rings. However, if this explanation were true, a highly non-linear behavior of the peak intensity with power would be expected as higher energy states become rapidly occupied at increasing excitation power; this is not observed (see Fig. 4). Recombination from excited hole states\(^{11}\) and light-hole states are likely to be present in the measured PL, but not resolved. A second possible explanation for the sub-peaks is monolayer fluctuations\(^{18}\) in the size of the rings. If this were the case, the lower energy emission from larger structures would be expected to saturate at higher powers than the higher energy emission from smaller structures. Again, this is not seen in our data where the majority of the blueshift of the CoM is a result of the lower energy states saturating at smaller laser powers than the higher energy states. The sub-peaks could also potentially be explained by Fabry–Pérot modes, but this explanation is not consistent with the power-induced shift of these peaks, and substructure is not observed in the emission from the wetting layer. We conclude that the substructure in the PL from the quantum rings is most likely due to discrete hole charging.

Concerning band-bending\(^ {16}\), emission from exciton states containing a fixed number of holes, i.e., the sub-peaks in the PL spectrum, is found to blue-shift with a magnitude that is much less than the CoMs shift (Fig. 3(b)). This is attributed to a collective band-bending effect as a result of an increase in the number of electrons that surround the rings and charging of the wetting layer.

Recent 8-band \(k\cdot p\) calculations have shown that the maximum localization energy of a pure GaSb/GaAs nanostructure with similar dimensions to those studied here is approximately 600–650 meV.\(^ {15,19}\) With a low-temperature GaAs band-gap of \(\sim 1.5\text{ eV}\), emission from the excitonic state in which just one heavy-hole occupies the dot should not occur below 0.9 eV. This allows the number of holes in each of the PL peaks to be quantified, as indicated by the numbers in circles in Figs. 3 and 4, and allows important observations to be made about this ring system. The peak corresponding to an occupancy of one hole/ring is difficult to fit due to its proximity to an absorption-dip in the optical fiber which is used to collect the emission from the sample. As the pump power tends towards zero, emission appears preferentially from the excitonic states with 4, rather than 1, holes/ring. This indicates that a degree of unintentional doping is present in the sample. A dependence of the exciton’s oscillator strength on the number of holes in a ring may contribute to this, though unintentional incorporation of carbon acceptors, as indicated by the presence of an acceptor-bound GaAs peak in the PL measurements, is thought to be the primary cause. Such background doping has been measured previously in the growth of GaSb nanostructures.\(^ {20}\)

The GaSb/GaAs quantum rings introduced in this work have unique physical properties, linked directly to their material properties, which may make them useful for applications in QIP. Unlike most zero-dimensional type-II systems, here the “unbound” carrier type does experience a confining potential that is spatially close to that of the bound carriers. This is believed to be responsible for the reduction in emission line-width of the ensemble that facilitated the measurement of discrete charging. It is also expected to lead to an increase in the exciton’s optical oscillator strength, though tools required to measure this were not available in this study. Weak oscillator strengths and broad line-widths have disincentivised research on individual type-II nanostructures; this system may help to address these issues. GaSb/GaAs has two significant advantages over widely researched InAs/GaAs dots; it has a small nuclear moment and sufficient localization energy to provide confinement well above room temperature.\(^ {8}\) It may have a promising future for applications manipulating and storing individual hole-spins.

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