

Forming and confining of dipolar excitons by quantizing magnetic fields

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We show that a magnetic field perpendicular to an AlGaAs/GaAs coupled quantum well efficiently traps dipolar excitons and leads to the stabilization of the excitonic formation and confinement in the illumination area. Hereby, the density of dipolar excitons is remarkably enhanced up to $\sim 10^{11}$ cm⁻². By means of Landau level spectroscopy we study the density of excess holes in the illuminated region. Depending on the excitation power and the applied electric field, the hole density can be tuned over one order of magnitude up to $\sim 2.5 \times 10^{11}$ cm⁻²—a value comparable with typical carrier densities in modulation-doped structures.

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The growing interest in the spectroscopy of coupled quantum wells (CQWs) is related both to the prediction of Bose-Einstein condensation of dipolar excitons^{1,2} and to the possible applications in optoelectronic devices.³⁻⁵ In both cases control of high densities of dipolar excitons is required. As shown both theoretically and experimentally, the dynamics of photogenerated unbound electron-hole pairs prevents the desired formation and control of a dipolar exciton ensemble.⁶⁻¹⁴ In particular, excess hole densities in the illumination area of interband excitation are deduced from the formation of ring-shaped patterns resulting from dipolar exciton transitions.^{11,13,15-17} Different methods to spatially confine dipolar excitons within the two-dimensional (2D) plane have been applied, either by using aleatory traps created at the interfaces¹⁸ or by building electrostatic,¹⁹⁻²² stress,²³ or magnetic²⁴ traps. In all strategies the main goal was to create an in-plane confining potential, which would trap dipolar excitons at its minimum. Theoretical studies of the influence of an additional magnetic field on excitonic formation and transport suggest that a magnetic field applied perpendicular to the 2D plane will stabilize high densities of dipolar excitons.^{25,26}

We show experimentally that formation and trapping of dipolar excitons in the illumination spot is dramatically enhanced by a quantizing magnetic field. Under focused excitation the magnetic field confines dipolar excitons at the illumination spot and thus it acts as a trap. This method allows us to achieve dipolar exciton densities up to $\sim 10^{11}$ cm⁻² demonstrating very high trapping efficiency, which is comparable with the most effective in-plane traps.^{21,22} Additionally, for higher magnetic fields we are able to determine the density of excess holes at the excitation spot from magnetic depopulation of Landau levels, which depending on the applied electric field and excitation conditions reaches $\sim 2.5 \times 10^{11}$ cm⁻². This also places a lower bound on the excess hole density at zero magnetic field. Presumably a high density of excess holes prevents Bose-Einstein condensation of excitons. We show how this density can be decreased using the electric field.

The studied heterostructure was grown by molecular beam epitaxy on a semi-insulating GaAs (001) substrate. It consists of two 8-nm-thick GaAs coupled quantum wells separated by a 4-nm Al_{0.3}Ga_{0.7}As barrier and embedded between two Al_{0.3}Ga_{0.7}As barriers [Fig. 1(a)], which separate the CQWs from an *n*-doped GaAs back gate and the semitransparent

Schottky gate deposited on the sample surface. The resulting field-effect device allows the CQWs potential to be tuned relative to the Fermi energy pinned to the back gate. The electric field applied in the structure can be estimated from the formula $F = -(V_g + V_S)/d$, where V_g is the voltage applied between the contacts, V_S is the height of the metal Schottky barrier (-0.7 V), and d is the distance between the gates (370 nm). A detailed description of the structure can be found in Refs. 14 and 22. The microphotoluminescence (micro-PL) spectroscopy is performed using a confocal setup²⁷ in a He bath cryostat with a superconducting coil. A 2-mm focal length aspheric lens is used to focus the excitation beam from a continuous diode laser (680 nm) and to collect the PL from the sample. Our confocal scheme allows us to investigate optical properties of the emission coming from an area of ~ 1 μ m in diameter.²⁸ All experiments are performed at the temperature of 4.2 K and in magnetic fields up to 9 T in Faraday configuration.

In CQWs two types of excitonic transition can occur: a direct exciton (*D*) involving an electron and a hole from the same quantum well and an indirect exciton (*I*), called also a dipolar exciton, formed by carriers from different wells [Fig. 1(a)]. The dipolar transition can be easily manipulated with the electric field (*F*) and becomes energetically favorable for higher fields. Under focused excitation a spatial nonequilibrium of photogenerated carriers forms: electrons drift out of the focus spot much faster than much heavier holes [Fig. 1(b)]. Therefore the PL signal from the illuminated area is relatively weak. The densities of electrons, holes, and dipolar excitons can evolve quite independently in the focus area of the illumination, a particular feature of CQWs.¹¹ In order to prevent electron escape one can use a magnetic field (*B*).^{25,26} The electron drift is modified by the circular motion due to the Lorentz force, resulting in a cycloidlike trajectory [Fig. 1(c)]. At a threshold field B^* this force dominates the electron movement, and for higher *B* the electrons propagate on cyclotron orbits and stay within the excitation area [Fig. 1(d)].

Figure 2(a) shows PL spectra measured at $B = 0$ and 3.7 T. Without a magnetic field the indirect excitons are very weak and the emission is dominated by the direct transition (lower curve). Under applied magnetic field the intensity of the indirect transition increases by a factor of ~ 10 (upper curve). Additionally, we note that the spectrum splits under magnetic

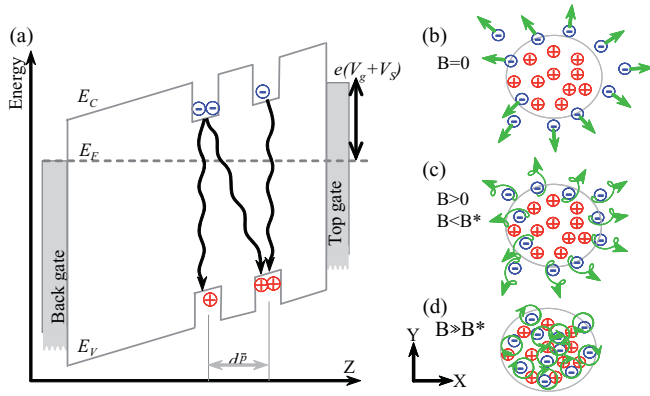


FIG. 1. (Color online) (a) Sketch of the band gap between conduction (E_C) and valence (E_V) band edges (along the growth direction Z). The applied gate voltage V_g and built-in voltage V_s sum up to an electric field between the n -doped back gate and the top gate. Two different optical transitions (direct and indirect) are marked with wavy arrows. (b)–(d) scheme of the in-plane (X - Y) distribution of electrons \ominus and holes \oplus at the excitation spot (big circle) for different magnetic fields. Arrows mark the diffusion of electrons: (b) for $B = 0$, (c) for $B < B^*$, small magnetic fields, and (d) for $B \gg B^*$.

fields, and the optical transitions form a Landau level (LL) fan chart [Fig. 2(b)]. The two first LL transitions are clearly visible for indirect excitons [Fig. 2(a)]. Comparing the spectra for two excitation powers in Fig. 2(a) (middle and upper curves), we observe that the intensity of indirect excitons increases and their spectral position shifts toward higher energies as expected for interacting systems of dipolar excitons with increased density.^{7,10,12,29} Also the relative intensities of the two indirect exciton LLs change with laser power. The difference in the filling of consecutive LLs suggests an increase of excess hole density with increasing excitation intensity. Already this first observation suggests the transition from a hole-rich to an exciton-rich system induced by a finite B . In order to better understand this process we measure systematically the dependence of PL on B under different excitation powers and electric fields.

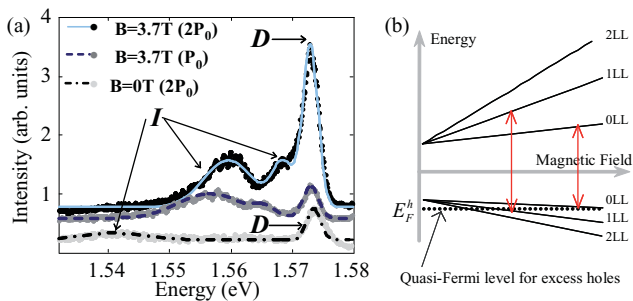


FIG. 2. (Color online) (a) PL spectra of direct (D) and indirect (I) excitons for different magnetic fields and excitation powers: ($B = 0$ T, $2P_0$), lowest; ($B = 3.7$ T, P_0), middle; ($B = 3.7$ T, $2P_0$), highest data set ($P_0 \approx 10 \mu\text{W}$), measured at $F = -30$ kV/cm. Lines, Gaussian fits. (b) Scheme of optical selection rules between electron and hole Landau levels in a perpendicular magnetic field. Dotted horizontal line marks the quasi-Fermi level (E_F^h) for holes.

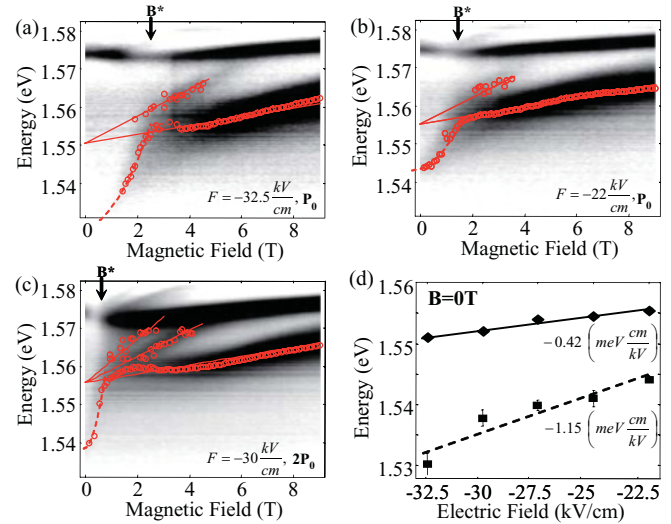


FIG. 3. (Color online) (a)–(c) Magnetic field evolution of PL spectra for different electric fields. (a), (b) measured at constant excitation power P_0 and (c) at $2P_0$. Points, fits of energy position of LL; dashed line, quadratic fit for low magnetic fields; solid lines, linear fit of Landau levels for higher fields. (d) Stark shift deduced from the extrapolation to $B = 0$ T: from the low-field evolution (squares, dashed line) and from Landau levels (diamonds, solid line).

Figures 3(a)–3(c) present the PL evolution in a perpendicular magnetic field measured for different electric fields and excitation intensities. Two field regimes are clearly distinguishable for the indirect excitons. For low fields ($B < B^*$) the signal is relatively weak and the transition energy shifts by several meV and approximately proportionally to B^2 to the blue. For higher fields, however, the emission has higher intensity, and LL splitting becomes clearly visible.

The evolution of the spectra for high magnetic fields ($> B^*$) can be described using a simple picture of excitonic transitions in the presence of carriers. The Landau quantization ($n + 1/2$) $\hbar\omega_c$ can be fitted with cyclotron frequency $\omega_c = eB/m_*$ using a reduced excitonic mass $m_* = 0.05m_0$, independent of B , V_g , and excitation intensity, and with e and m_0 the electron charge and mass, respectively. The observation of higher LLs and their magnetic depopulation above a specific magnetic field B_d enables us to assign a Landau level filling factor ν and hence a corresponding effective density of excess holes n_h via $\nu = \frac{hn_h}{eB_d}$, with h the Planck constant. We use the first excited LL ($n = 1$) to determine the dependence of n_h on the electric field F and find a systematic increase of n_h with F . In Figs. 3(a)–3(c) one can see the reduction and complete disappearance of this level with increasing magnetic field. We also studied higher LLs ($n > 1$), when they were clearly resolved in the spectra, and the values of the densities obtained were the same within the experimental error. Hence, the hole density does not increase with $B > B^*$. We wish to point out that the excess hole density obtained from the magnetic depopulation is essentially located in the surface near the QW and at least partially excitonically bound to electrons in the lower QW. This reflects the situation where the in-plane hole distance is comparable to the spacing of the QWs, which in turn is comparable to the excitonic Bohr radius.

The strong blueshift of indirect PL when $B < B^*$ exceeds 15 meV and is much larger than the expected diamagnetic shift of less than 1 meV.³⁰ However, it can be explained by the increase of the dipole-dipole interaction with increasing exciton density, in analogy to the spectral shift with the excitation power in Fig. 2(a).^{7,10,12,29} At a critical magnetic field B^* a maximum density is reached [see the arrows in Figs. 3(a)–3(c)] and the population of the excitons stabilizes within the excitation focus as schematically sketched in Figs. 1(c) and 1(d). This increase in density is also reflected in an increase in the PL intensity. To quantify this quadratic increase in exciton density n_X with B ($B < B^*$) we subtract both the cyclotron energy and the calculated diamagnetic shift to obtain the bare B -dependent and interaction-caused blueshift of the indirect emission ΔE_X , related to Δn_X by³¹

$$\Delta E_X = \frac{ed_{\bar{p}}}{\epsilon_b \epsilon_0} \Delta n_X \quad (1)$$

where $d_{\bar{p}}$ is the dipole length (approximately the distance between the center of the two QWs) and ϵ_b is the background dielectric constant. Since the exciton density at $B = 0$ is negligibly small, the magnitude of the blueshift reflects the total exciton density at $B > B^*$, which independent of B ($B > B^*$) varies from $n_X \sim 0.66 \times 10^{11} \text{ cm}^{-2}$ to $n_X \sim 1.22 \times 10^{11} \text{ cm}^{-2}$ for the electric field range -22 to -32.5 kV/cm . The increase of B^* with the applied electric field is clearly visible in Figs. 3(a) and 3(b). Stronger F induces larger separation of photoexcited carriers, reducing exciton binding and n_X . In order to suppress this effect higher B is required to localize carriers, which allow formation of excitons.²⁵ In contrast, strong excitation power increases the initial exciton density and reduces B^* , as seen by comparing Figs. 3(b) and 3(c). The n_X enhancement in the magnetic field is a profound effect if it compares with typical densities achieved in CQWs.¹⁴

In Fig. 3(d) we compare the $B = 0$ PL energy of the indirect excitons extrapolated from the low- B data with that extrapolated from the Landau level fan chart discernible at $B > B^*$. For a given electric field and excitation power their energy difference reflects the blueshift induced by the B -dependent change in exciton density (1) discussed above. More striking is the change in slope in Fig. 3(d), reflecting a difference in the Stark shift. The slope of the Stark shift obtained from the low- B data of -1.15 meV cm/kV is in good agreement with the value -1.26 meV cm/kV obtained from numerical simulations in such a heterostructure³² and comparable with results of other experiments.^{9,22} The Stark shift obtained from the extrapolation of LLs is notably reduced by about a factor of 3 to -0.42 meV cm/kV . This suggests screening of the applied electric field, most likely caused by a magnetic-field-assisted accumulation of the dipolar indirect excitons as shown in Fig. 3 ($B < B^*$) and discussed above.

In Fig. 4(a) we compare the excess hole n_h and exciton n_X densities. Note that n_h was obtained from the depopulation of LLs ($B > B^*$), whereas n_X from the low-field ($B < B^*$) blueshift of PL. The changes of n_h with the electric field can be qualitatively explained by slower escape rate of holes from the QW, which, using a simple tunneling model developed in the WKB approximation³³ and the simulated level structure in CQWs,³² is about three times smaller than for electrons.

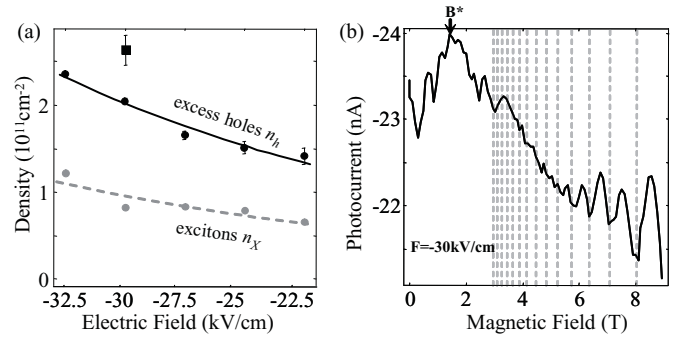


FIG. 4. (a) Densities vs electric field: for excess holes (black points, series taken at excitation power P_0 ; square, at $2P_0$) and for the exciton density (gray points). Solid and dashed lines are guides for the eyes. (b) Photocurrent vs magnetic field for $F = -30 \text{ kV/cm}$. Dashed vertical lines show the resonance of the photon energy with interband LL transition ($n = \dots, 9, 8, 7$).

The maximum n_h exceeds $2.6 \times 10^{11} \text{ cm}^{-2}$. We note that such high carrier density was previously observed only for intentionally doped structures. The density n_h can exceed the density of excitons as their charge is partially compensated by the negative gate charge.

Complementary information about the onset of Landau quantization can be deduced from the photocurrent measurements. Without illumination our field-effect devices possess typical diode characteristics: current starts to flow at gate voltages above the flatband voltage $+0.7 \text{ V}$ and at voltages below -0.6 V at which breakdown occurs. The illumination-induced photocurrent starts to flow for negative voltages and it grows with decreasing biases up to several nanoamps at $V_g = -0.5 \text{ V}$ ($F = -32.5 \text{ kV/cm}$). Figure 4(b) shows the changes of the photocurrent through the sample under applied magnetic fields. We observe a small increase of the current amplitude for $B < 1.8 \text{ T}$ and a slow decrease for higher fields. A characteristic oscillation pattern occurs for higher B and it is more pronounced for bigger excitation powers. The period of the oscillations depends weakly on E . Resonance of the LL fan chart with the laser energy (1.82 eV) is marked in Fig. 4(b) by the dashed lines. When the magnetic field is increased, LLs with decreasing index cross the excitation energy, e.g., the crossing of the seventh LL occurs at $\sim 8 \text{ T}$. Because the excitation energy comes to resonance only with the empty levels and it exceeds over 20 times the hole quasi-Fermi energy, this effect is only seen in photocurrent measurements, but not in nonresonantly excited PL. The increase of the photocurrent at low B implies charge separation of photoexcited carriers, most likely occurring from within the individual QWs as reflected in the simultaneously observed decrease in the intensity of the direct PL [see Fig. 3(a)–3(c)]. Note that charge separation in the respective quantum well prevents direct recombination and thus may also increase the interwell tunneling and formation of indirect excitons. Together with the magnetic field stabilization this is likely to be responsible for the observed intensity increase in the PL of indirect excitons.

In conclusion, we showed that under focused excitation a magnetic field stabilizes the formation and confinement of indirect excitons in CQWs. The confining ability of the magnetic field allows enhancement of the exciton densities

up to $n_X \approx 10^{11} \text{ cm}^{-2}$. Our explanation is confirmed by measurements at variable excitation intensities and electric fields. PL spectroscopy also enables us to determine the excess hole density, which can be tuned by the electric field and the excitation power from optically nonmeasurable values up to $n_h = 2.6 \times 10^{11} \text{ cm}^{-2}$.

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¹L. V. Keldysh and A. N. Kozlov, *Zh. Eksp. Teor. Fiz.* **54**, 978 (1968) [*Sov. Phys.-JETP* **27**, 521 (1968)].

²J. M. Blatt, K. W. Böer, and W. Brandt, *Phys. Rev.* **126**, 1691 (1962).

³J. Krauß, J. P. Kotthaus, A. Wixforth, M. Hanson, D. C. Driscoll, A. C. Gossard, D. Schuh, and M. Bichler, *Appl. Phys. Lett.* **85**, 5830 (2004).

⁴S. Zimmermann, A. Wixforth, J. P. Kotthaus, W. Wegscheider, and M. Bichler, *Science* **283**, 1292 (1999).

⁵A. A. High, E. E. Novitskaya, L. V. Butov, M. Hanson, and A. C. Gossard, *Science* **321**, 229 (2008).

⁶A. L. Ivanov, P. B. Littlewood, and H. Haug, *Phys. Rev. B* **59**, 5032 (1999).

⁷V. Negoita, D. W. Snoke, and K. Eberl, *Phys. Rev. B* **61**, 2779 (2000).

⁸Z. Vörös, R. Balili, D. W. Snoke, L. Pfeiffer, and K. West, *Phys. Rev. Lett.* **94**, 226401 (2005).

⁹A. Gärtner, A. W. Holleitner, J. P. Kotthaus, and D. Schuh, *Appl. Phys. Lett.* **89**, 052108 (2006).

¹⁰R. Rapaport, G. Chen, and S. H. Simon, *Phys. Rev. B* **73**, 033319 (2006).

¹¹R. Rapaport and G. Chen, *J. Phys. Condens. Matter* **19**, 295207 (2007).

¹²R. Zimmermann and C. Schindler, *Solid State Commun.* **144**, 395 (2007).

¹³M. Stern, V. Garmider, E. Segre, M. Rappaport, V. Umansky, Y. Levinson, and I. Bar-Joseph, *Phys. Rev. Lett.* **101**, 257402 (2008).

¹⁴X. P. Vögele, D. Schuh, W. Wegscheider, J. P. Kotthaus, and A. W. Holleitner, *Phys. Rev. Lett.* **103**, 126402 (2009).

¹⁵R. Rapaport, G. Chen, D. Snoke, S. H. Simon, L. Pfeiffer, K. West, Y. Liu, and S. Denev, *Phys. Rev. Lett.* **92**, 117405 (2004).

¹⁶A. T. Hammack, L. V. Butov, J. Wilkes, L. Mouchliadis, E. A. Muljarov, A. L. Ivanov, and A. C. Gossard, *Phys. Rev. B* **80**, 155331 (2009).

¹⁷A. V. Paraskevov and S. E. Savel'ev, *Phys. Rev. B* **81**, 193403 (2010).

¹⁸L. V. Butov, C. W. Lai, A. L. Ivanov, A. C. Gossard, and D. S. Chemla, *Nature (London)* **417**, 47 (2002).

¹⁹S. Zimmermann, A. O. Govorov, W. Hansen, J. P. Kotthaus, M. Bichler, and W. Wegscheider, *Phys. Rev. B* **56**, 13414 (1997).

²⁰M. Hagn, A. Zrenner, G. Böhm, and G. Weimann, *Appl. Phys. Lett.* **67**, 232 (1995).

²¹A. T. Hammack, N. A. Gippius, S. Yang, G. O. Andreev, L. V. Butov, M. Hanson, and A. C. Gossard, *J. Appl. Phys.* **99**, 066104 (2006).

²²A. Gärtner, L. Prechtel, D. Schuh, A. W. Holleitner, and J. P. Kotthaus, *Phys. Rev. B* **76**, 085304 (2007).

²³V. Negoita, D. W. Snoke, and K. Eberl, *Appl. Phys. Lett.* **75**, 2059 (1999).

²⁴P. C. M. Christianen, F. Piazza, J. G. S. Lok, J. C. Maan, and W. van der Vleuten, *Physica B* **249**, 624 (1998).

²⁵A. O. Govorov and W. Hansen, *Phys. Rev. B* **58**, 12980 (1998).

²⁶Y. E. Lozovik, I. V. Ovchinnikov, S. Y. Volkov, L. V. Butov, and D. S. Chemla, *Phys. Rev. B* **65**, 235304 (2002).

²⁷B. Alén, F. Bickel, K. Karrai, R. J. Warburton, and P. M. Petroff, *Appl. Phys. Lett.* **83**, 2235 (2003).

²⁸K. Kowalik-Seidl, X. Vögele, B. Rimpfl, S. Manus, D. Schuh, W. Wegscheider, A. W. Holleitner, and J. P. Kotthaus, *Appl. Phys. Lett.* **97**, 011104 (2010).

²⁹B. Laikhtman and R. Rapaport, *Phys. Rev. B* **80**, 195313 (2009).

³⁰M. Bugajski, W. Kuzsko, and K. Regiński, *Solid State Commun.* **60**, 669 (1986).

³¹Recently several publications were devoted to the problem of the exciton-exciton interaction in CQWs and correct recalculation of the blueshift into the exciton density (Refs. 12, 13, and 34). The latest studies (Refs. 29, 35, and 36) show that for typical GaAs CQWs with exciton densities 10^{11} – 10^{12} cm^{-2} the simple capacitorlike formula (Refs. 6, 10, and 37) is still a good approximation.

³²See [<http://www.nextnano.de>].

³³W. Heller, U. Bockelmann, and G. Abstreiter, *Phys. Rev. B* **57**, 6270 (1998).

³⁴C. Schindler and R. Zimmermann, *Phys. Rev. B* **78**, 045313 (2008).

³⁵A. L. Ivanov, E. A. Muljarov, L. Mouchliadis, and R. Zimmermann, *Phys. Rev. Lett.* **104**, 179701 (2010).

³⁶A. L. Ivanov, *Europhys. Lett.* **59**, 586 (2002).

³⁷S. B.-T. de Leon and B. Laikhtman, *Phys. Rev. B* **63**, 125306 (2001).