

Many-body effects in a laterally inhomogeneous semiconductor quantum well

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Many body effects on conduction and diffusion of electrons and holes in a semiconductor quantum well are studied using a microscopic theory. The roles played by the screened Hartree-Fock (SHF) terms and the scattering terms are examined. It is found that the electron and hole conductivities depend only on the scattering terms, while the two-component electron-hole diffusion coefficients depend on both the SHF part and the scattering part. We show that, in the limit of the ambipolar diffusion approximation, however, the diffusion coefficients for carrier density and temperature are independent of electron-hole scattering. In particular, we found that the SHF terms lead to a reduction of density-diffusion coefficients and an increase in temperature-diffusion coefficients. Such a reduction or increase is explained in terms of a density- and temperature-dependent energy landscape created by the bandgap renormalization.

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Our understanding of Coulomb interaction in an optically excited semiconductor has been greatly enriched recently thanks to the extensive theoretical¹⁻³ and experimental investigations⁴ over the past few decades. As far as optical properties in highly excited semiconductors (with high carrier density) are concerned, many-body effects manifest themselves in two important ways: the renormalization of the single-particle energies and the finite life time of such renormalized single-particle states. Though the quantitative calculation of these two quantities is still a topic of current research, the qualitative difference of these two features seem to be quite clear. For the sake of convenience of the presentation in this paper, we classify the two types of terms into coherent and incoherent (or scattering) parts, since the renormalization of single-particle state changes the resonance frequency, while scattering leads to a decay of coherence or oscillation. For a spatially uniform system, these two types of effects manifest themselves in the linear optical spectrum and have received extensive attention in the past decades. The manifestation of the many-body effects in a spatially non-uniform system has not received comparable attention. It is especially interesting to see, as we will show later, that these two parts of many-body interaction play different roles in the conduction and diffusion processes in a spatially non-uniform semiconductor.

In this communication, we report on our recent theoretical study on the effects of many-body interaction in a spatially inhomogeneous system. The starting point of our investigation is the set of Boltzmann-Bloch-Poisson equations which contains many-body interactions in the spirit of Refs. 1-3 and 5. Namely, the coherent part is treated within the screened Hartree-Fock (SHF) approximation, while the scatterings are treated within the second Born approximation. After following the standard moment equation approach⁶ by assuming the quasi-equilibrium distribution of carriers, a set of coupled diffusion equations for carrier densities and temperatures

can be derived⁷ and given as follows:

$$\partial_t N^\alpha + \partial_{\vec{r}} \cdot \vec{J}_N^\alpha = R_N^\alpha, \quad (1)$$

$$\begin{aligned} \partial_t T^\alpha + \partial_{\vec{r}} \cdot \vec{J}_T^\alpha &= j_W^\alpha \vec{u}^\alpha \cdot \partial_{\vec{r}} W^\alpha + \partial_{\vec{r}} j_N^\alpha \cdot \vec{J}_N^\alpha \\ &- \partial_{\vec{r}} j_W^\alpha \cdot \vec{J}_W^\alpha = R_T^\alpha, \end{aligned} \quad (2)$$

where N^α , W^α , \vec{u}^α , and T^α stand for density, thermal energy, drift velocity, and temperature of electrons ($\alpha = e$) and holes ($\alpha = h$). j_W^α and j_N^α are transform Jacobians which relate thermal energies to temperatures. They are defined in Ref. 7. For the purpose of this paper, it suffices to say that they are functions of densities and temperatures. In (1) and (2), R_N^α 's represent generation and recombination of carriers due to pumping and optical transitions, while R_T^α 's represent the corresponding heat sources or sinks. The density and thermal currents in equations (1) and (2) are defined as in Ref. 7:

$$\vec{J}_N^\alpha = \frac{\vec{P}^\alpha}{m_\alpha}, \quad (3)$$

$$\vec{J}_T^\alpha = (2j_W^\alpha \frac{W^\alpha}{N^\alpha} - j_N^\alpha) \vec{J}_N^\alpha. \quad (4)$$

The density currents can be written in terms of gradients of the four macroscopic variables ($X = N^e, N^h, T^e, T^h$) and the electrical potential, Φ , as⁷

$$\vec{J}_{N^\alpha} = - \sum_X D_{N^\alpha X} \partial_{\vec{r}} X - \frac{\sigma^\alpha}{q^\alpha} \partial_{\vec{r}} \Phi, \quad (5)$$

where we have introduced various diffusion coefficients and conductivities⁷ ($\alpha \neq \beta$):

$$D_{N^\alpha N^\alpha} = \mu_\alpha \left[(1 + \eta_\alpha) (C_{N^\alpha}^\alpha + H_{N^\alpha}^\alpha) + H_{N^\alpha}^\beta \right], \quad (6)$$

$$D_{N^\alpha N^\beta} = \mu_\alpha \left[(1 + \eta_\alpha) H_{N^\beta}^\alpha + H_{N^\beta}^\beta + C_{N^\beta}^\beta \right], \quad (7)$$

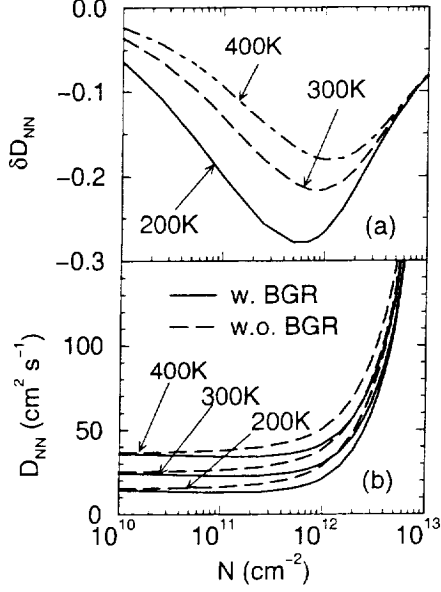


FIG. 1: Diffusion coefficient D_{NN} (b) and its relative change δD_{NN} (a) versus carrier density at three temperatures as indicated. Solid and dashed curves in Fig. 1(b) are for D_{NN} and D_{NN}^0 , respectively.

$$D_{N^\alpha T^\alpha} = \mu_\alpha \left[(1 + \eta_\alpha) (C_{T^\alpha}^\alpha + H_{T^\alpha}^\alpha) + H_{T^\alpha}^\beta \right], \quad (8)$$

$$D_{N^\alpha T^\beta} = \mu_\alpha \left[(1 + \eta_\alpha) H_{T^\beta}^\alpha + H_{T^\beta}^\beta + C_{T^\beta}^\beta \right], \quad (9)$$

$$\sigma^\alpha = \mu_\alpha e^2 [N^\alpha (1 + \eta_\alpha) - N^\beta]. \quad (10)$$

Following two shorthand notations have been introduced:

$$C_X^\alpha = \partial_X W^\alpha, \quad (11)$$

$$H_X^\alpha = N^\alpha \partial_X \delta \epsilon^\alpha. \quad (12)$$

Obviously, the C_X^α 's are specific heats of a certain kind, which represent the contribution from free electron and hole gases, while H_X^α 's are the contributions due to many-body interaction, as they are proportional to the derivatives of self-energy renormalization² ($\delta \epsilon^\alpha$) with respect to densities or temperatures. While equations (6)–(9) define diffusion coefficients in the density currents (noting the first index of the coefficients being N^α), the corresponding diffusion coefficients in the thermal currents are defined through the relation between the density currents J_N^α and thermal currents J_T^α , Eq. (4). Factors μ_α and η_α in Eqs. (6)–(10) are defined as follows:

$$\mu_\alpha = \left[\frac{\gamma_{LO}^\alpha \gamma_{LO}^\alpha}{\gamma_{eh}^\beta} (m_e + m_h) + (m_e \gamma_{LO}^\alpha + m_h \gamma_{LO}^\alpha) \right]^{-1} \quad (13)$$

$$\eta_\alpha = \frac{\gamma_{LO}^\beta}{\gamma_{eh}^\beta} \frac{m_e + m_h}{m_\alpha}, \quad (14)$$

where $\alpha, \beta \in \{e, h\} \mid \alpha \neq \beta$. Moreover, γ_{LO}^α and γ_{eh}^α are the relaxation rates of the α component momentum due

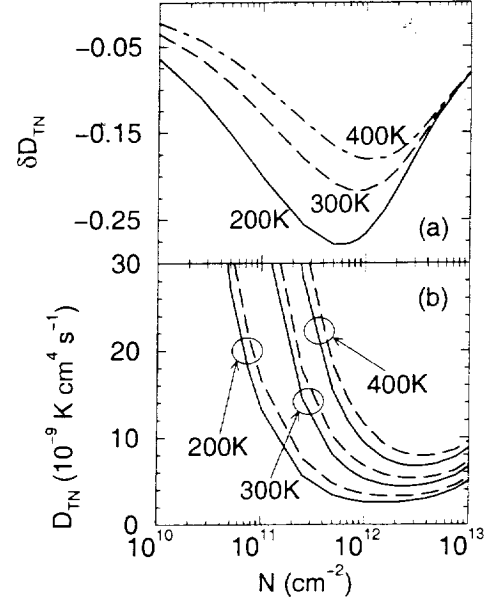


FIG. 2: Diffusion coefficient D_{TN} (b) and its relative change δD_{TN} (a) versus carrier density at three temperatures as indicated. Solid and dashed curves in Fig. 2(b) are for D_{TN} and D_{TN}^0 , respectively.

to carrier-LO (c-LO) phonon and due to electron-hole (e-h) scattering, respectively. These rates are defined microscopically in Ref. 7.

Several general features can be readily observed from the expressions in Eqs. (6)–(10): First, all diffusion coefficients and conductivities depend on momentum relaxation rates due to e-h scattering through factors μ_α and η_α . On the other hand, the coherent part of the many-body interaction (through H_X^α) only enters the diffusion coefficients, but not the conductivities. Second, it is interesting to consider the ambipolar diffusion coefficients, as commonly defined by setting equal the density currents of electrons and holes. The ambipolar currents are now written as⁷

$$\tilde{J}_N = -D_{NN} \partial_{\bar{r}} N - D_{NT} \partial_{\bar{r}} T, \quad (15)$$

$$\tilde{J}_T = -D_{TN} \partial_{\bar{r}} N - D_{TT} \partial_{\bar{r}} T, \quad (16)$$

with

$$D_{NX} = \frac{C_X^e + C_X^h}{m_e \gamma_{LO}^e + m_h \gamma_{LO}^h} + \frac{H_X^e + H_X^h}{m_e \gamma_{LO}^e + m_h \gamma_{LO}^h} \equiv D_{NX}^0 + \Delta D_{NX}, \quad (17)$$

$$D_{TX} = \left[2j_w \frac{W}{N} - j_N \right] D_{NX} \equiv D_{TX}^0 + \Delta D_{TX}, \quad (18)$$

where $X = N, T$ as we also assumed the densities and temperatures to be the same for the two components.

The identity in equation (17) defines the free-carrier diffusion coefficient D_{NX}^0 and the corresponding many-body correction ΔD_{NX} . We observe that the e-h scattering rate γ_{eh}^α (the incoherent part of the many-body effects) disappears from the ambipolar diffusion coefficients completely. Only c-LO phonon scattering rates γ_{LO}^α 's remain. This means that the ambipolar diffusion coefficients depend only on the weighted sum of the scattering rates of electrons and holes with LO phonons. The coherent part of the many-body interactions, however, remains present in the ambipolar diffusion coefficients, as expressed by the second term ΔD_{NX} and ΔD_{TX} in (17) and (18), respectively. The absence of the e-h scattering and the remaining presence of the coherent part of the Coulomb interaction in the ambipolar diffusion coefficients clearly illustrate different roles played by the two aspects of the same Coulomb interaction. We note that the absence of the e-h scattering in the ambipolar diffusion coefficients is also implied in Ref. 8. But unlike the situation here, all the Coulomb interaction terms drop out from the diffusion coefficient in Ref. 8 completely including the coherent part under the ambipolar diffusion approximation. The absence of the e-h scattering can be easily understood, since such scattering represents collisions of the electrons and holes, which now are parts of integral entities diffusing together under the ambipolar diffusion approximation. However, the coherent part creates a new density- and temperature-dependent environment (energy landscape) for the original quasi-particles (with unrenormalized energies). They diffuse in the modified energy-landscape, resulting in an effective change in diffusion coefficients, as we will explain in more detail in the following.

To study more quantitatively the effects of the coherent part of the many-body interaction, we examine the relative change in diffusion coefficients as defined by $\delta D_{XY} = \Delta D_{XY}/D_{XY}^0$. In Figs. 1-4, we plot δD_{XY} (a) and D_{XY} (b) with respect to density for all four ambipolar diffusion coefficients. As model material system, we choose GaAs of 8 nm in width. All the material parameters are standard and will not be listed. Fig. 1 shows the familiar density-diffusion coefficient D_{NN} . In Fig. 1(b), coefficients D_{NN} (solid lines) and D_{NN}^0 (dashed lines) are plotted versus carrier density at three temperatures. The overall feature of the diffusion coefficient is explained in detail in a separate paper⁹. We see that the coherent many-body effects result in a reduction in the diffusion coefficient. The relative change of diffusion coefficient is plotted in Fig. 1(a), where we see a diffusion coefficient reduction of over 25 percent at 200K. This reduction decreases as temperature increases. Similar behavior is also observed in Fig. 2 for the mutual-diffusion coefficient D_{TN} , which relates carrier density gradient to thermal flux \bar{J}_T . Fig. 3 shows the temperature-diffusion coefficient D_{TT} and D_{TT}^0 (a) and the corresponding relative change δD_{TT} . Contrary to the reduction of diffusion coefficients shown in Figs. 1 and 2, we see an increase in D_{TT} , i.e., $D_{TT} > D_{TT}^0$. Furthermore, the relative in-

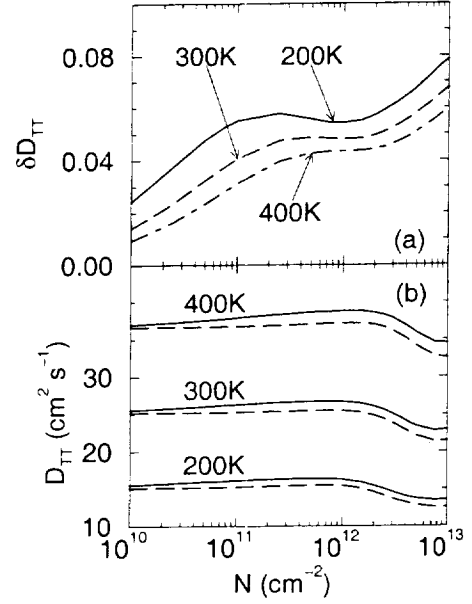


FIG. 3: Diffusion coefficient D_{TT} (b) and its relative change δD_{TT} (a) versus carrier density at three temperatures as indicated. Solid and dashed curves in Fig. 3(b) are for D_{TT} and D_{TT}^0 , respectively.

crease δD_{TT} is much smaller in magnitude than δD_{NN} and δD_{TN} in Figs. 1 and 2. The change is less than 10 percent. Similar behavior is shown in Fig. 4 for the mutual-diffusion coefficient D_{NT} , which describes carrier density flux induced by temperature gradient.

Let us now explain in more detail these figures. We begin with the reduction in diffusion coefficients, D_{NN} and D_{TN} , in Figs. 1 and 2, both of which are determined by the derivatives with respect to density [see Eqs. (11), (12), (17), and (18)]. This reduction can be explained by the bandgap renormalization. We note that D_{NN} describes a carrier density flux from high density region to low density region. Due to bandgap renormalization which increases with carrier density, the high density region has a smaller total bandgap than the low density region. This means that a diffusing particle from high density region to low density region will have to climb an uphill energy landscape due to many-body effects, thus leading to a reduction in the effective diffusion coefficients. The reduction in the mutual-diffusion coefficient D_{TN} needs slightly different explanation. First, we note that D_{TN} , by definition, describes the *thermal* flux from high *density* region to low density region. Due to the bandgap renormalization, an energy bandgap profile is created. A thermal flux is therefore induced from the high bandgap (lower density) region to lower bandgap (higher density) region to counteract the thermal flux and to equilibrate the total energy (bandgap plus the thermal energy) profile. The situation described in Figs. 3 and 4 are exactly the reverse of that in Figs. 1 and 2. The increase in the diffusion coefficients are due to the bandgap renormalization that decreases with plasma

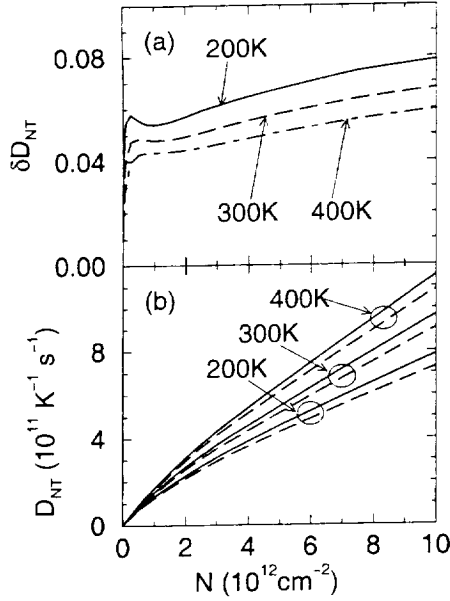


FIG. 4: Diffusion coefficient D_{NT} (b) and its relative change δD_{NT} (a) versus carrier density at three temperatures as indicated. Solid and dashed curves in Fig. 4(b) are for D_{NT} and D_{NT}^0 , respectively.

temperature¹⁰ instead of increasing with density as in Figs. 1 and 2. The energy landscape reverses from the cases of Figs. 1 and 2 and leads to increase of diffusion coefficients D_{NT} and D_{TT} . In short, the many-body effects on diffusion coefficients lead to a decrease in those diffusion coefficients that are related to the density gradient (D_{NN} and D_{TN}) and an increase in those that are related to temperature gradient (D_{TT} and D_{NT}). Finally, the smaller change in D_{TT} and D_{NT} (Figs. 3 and 4) than in D_{NN} and D_{TN} (Figs. 1 and 2) is due to the weaker dependence of the bandgap renormalization with respect to plasma temperature than to density.

Another feature of Figs. 1 and 2 is the decrease of the relative change δD_{NN} and δD_{TN} at lower density until the carrier density reaches the critical value near $1 \times 10^{12} \text{ cm}^{-2}$, where electrons become degenerate. This decrease is a result of the reduction of the bandgap due

to bandgap renormalization and to an almost constant value of D_{NN} and D_{TN} . The relative change δD_{NN} reaches a minimum around 28 percent for 200K. With the further increase of carrier density, δD_{NN} and δD_{TN} start to increase as the carriers become strongly degenerate and D_{NN}^0 and D_{TN}^0 begin to rise dramatically⁹. At high density for lasing over $1 \times 10^{12} \text{ cm}^{-2}$, the diffusion reduction is still over 20 percent. The larger values of δD_{NN} and δD_{TN} at higher temperature are mainly due to the increase of the D_{NN}^0 and D_{TN}^0 with temperature.

In summary, many-body effects are investigated in a semiconductor quantum well where spatial nonuniformity of densities and temperatures exists along the quantum well plane. Different roles played by the coherent and incoherent parts of Coulomb interaction are analyzed. While both coherent and incoherent parts contribute to the diffusion coefficients of the general two component system, the conductivities depend only on the scattering part. Even though e-h scattering plays an important role in legitimating the ambipolar diffusion approximation⁷, we show that the diffusion coefficients of the established composite system do not depend on the e-h scattering rate. Instead the ambipolar diffusion coefficient depends only on the coherent part of the interaction, the bandgap renormalization. We found that the coherent many-body interaction leads to a significant reduction of the ambipolar diffusion coefficients D_{NN} and D_{TN} and an increase in coefficients D_{TT} and D_{NT} . We note that this quite significant change in diffusion coefficients, especially in D_{NN} and D_{TN} , should be important in describing optoelectronic devices where spatial inhomogeneities of densities, or plasma temperatures occur. Such non-uniformities are quite ubiquitous in high power and ultrafast devices, such as lasers and photodetectors. Simulation of such devices using the microscopically calculated diffusion coefficients will be reported elsewhere.

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¹ H. Haug and S. Koch, *Theory of the Electrical and Optical Properties of Semiconductors* (World Scientific Singapore, 1994), 3rd ed.

² W. W. Chow, S. W. Koch, and M. Sargent, *Semiconductor Laser Physics* (Springer Verlag, Berlin, 1994).

³ R. Binder and S. W. Koch, *Prog. Quant. Electr.* **19**, 307 (1995).

⁴ D.S.Chemla and J.Shah, *Proceed. Nat. Acad. Sci.* **97**, 2437 (2000).

⁵ O.Hess and T.Kuhn, *Phys.Rev.A* **54**, 3347 (1996).

⁶ M. Lundstrom, *Fundamentals of Carrier Transport* (Addison-Wesley, 1990).

⁷ J.Li and C.Z.Ning, A Hydrodynamic Approach to Spatially Inhomogeneous Semiconductor Lasers: I. Theoretical Aspects (to be published) (2002).

⁸ J.Meyer, *Phys.Rev.B* **21**, 1554 (1980).

⁹ J.Li and C.Z.Ning, A Hydrodynamic Approach to Spatially Inhomogeneous Semiconductor Lasers: II Numerical Results (to be published) (2002).

¹⁰ R.Zimmermann, *Phys. Status Solidi B* **146**, 371 (1988).