

Influence of the confinement potential on the electron-hole-pair states in semiconductor microcrystallites

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The energies of the two energetically lowest dipole-allowed electron-hole-pair states in semiconductor microcrystallites are computed variationally. Details of the quantum confinement conditions, such as the finite value of the quantum confinement potential and the different effective electron-hole masses inside and outside the crystallites, are considered explicitly. Significant deviations from the infinite-potential approximation are obtained.

I. INTRODUCTION

Three-dimensional quantum confinement effects in semiconductor microcrystallites occur when the particle size becomes comparable to the exciton Bohr radius. The confinement gives rise to interesting new effects which have attracted considerable attention recently.¹⁻¹⁴ The research has concentrated mainly on optically excited compound semiconductor clusters in glasses or other dielectrics,^{1,3,4,8-10,14} and on colloidal semiconductors precipitated in liquids,^{2,4,6,7} as well as on quantum dots in organic films or inside zeolites.¹³ The first theoretical descriptions of three-dimensional excitonic quantum confinement were given by Efros and Efros¹ and by Brus.² These authors discussed the basic optical properties of semiconductor microcrystallites as functions of crystallite size. Brus² calculated the energy of the lowest electron-hole-pair state, approximating the exciton wave function by a few configurations of the electron-hole *s* wave functions confined in a sphere. Recent variational calculations achieved some improvements in determining the energy of the lowest electron-hole-pair state.^{5,6} However, for crystallite radii comparable to the bulk exciton Bohr radius, theoretical results^{5,6} overestimate the confinement-induced blue shift in comparison with experimental results.^{3,4} We believe that one reason for this overestimation is the assumption of an infinite-potential barrier in the hard-wall model, which yields a too large kinetic energy of the electron and hole, particularly in the case of small crystallite radii. In this paper, we compute the energy of the two lowest dipole allowed quantum-confined electron-hole-pair states in semiconductor microcrystallites taking into account a finite barrier for the confining potential. To determine the wave function outside the quantum dots we need to model the environment of the microcrystallites. For the case of quantum dots in

a liquid, we assume that the electrons are mobile outside the quantum dot and use the free-electron mass m_0 for the electrons in the liquid. For the case of quantum dots in glass, we model the electron states in the glass as localized states with very low mobility and correspondingly large electron mass. We present a systematic study of the parameter dependence of the computed quantum-confined pair states.

II. VARIATIONAL CALCULATIONS

In the framework of the effective-mass approximation, the electron-hole-pair Hamiltonian is given by

$$H = -\frac{\hbar^2 \nabla_e^2}{2m_e} - \frac{\hbar^2 \nabla_h^2}{2m_h} - \frac{e^2}{\epsilon |r_e - r_h|} + V_e(r_e) + V_h(r_h) + E_g , \quad (1)$$

with the confinement potential

$$V_i(r_i) = \begin{cases} 0 & \text{for } r_i \leq R \quad (i=e,h) \\ V_{0i} & \text{for } r_i \geq R \quad (i=e,h) . \end{cases} \quad (2)$$

Here R is the crystallite radius; ϵ is the background dielectric constant of the semiconductor material; E_g is the band-gap energy of the bulk semiconductor material, m_e , r_e , and V_{0e} and m_h , r_h , and V_{0h} are the effective mass, position, and potential-well height of the electron and the hole, respectively. The contribution of the surface polarization terms is small,⁷ and we neglect it here for the sake of simplicity.

A. Ground state

In order to calculate the ground-state energy of the electron-hole pair we choose the trial wave function as

$$\psi(\mathbf{r}_e, \mathbf{r}_h) = e^{-\lambda_s |\mathbf{r}_e - \mathbf{r}_h|} \phi_{e,s}(r_e) \phi_{h,s}(r_h), \quad (3)$$

where λ_s is the variational parameter. The functions ϕ_e (ϕ_h) for the electron (hole) are the radial eigenfunctions corresponding to the lowest energy state in a spherical well of finite depth, i.e.,

$$\phi_{i,s}(r_i) = \begin{cases} \frac{\sin(k_{i,s} r_i)}{k_{i,s} r_i} & \text{for } r_i \leq R \quad (i=e,h) \\ \frac{\sin(k_{i,s} R)}{k_{i,s} r_i} e^{-q_{i,s}(r_i - R)} & \text{for } r_i \geq R \quad (i=e,h) \end{cases} \quad (4)$$

where we defined

$$k_{i,s}^2 = \frac{2m_{i,1}E_{i,s}}{\hbar^2}, \quad (5)$$

$$q_{i,s}^2 = \frac{2m_{i,2}V_{0,i}}{\hbar^2} - k_{i,s}^2 \frac{m_{i,2}}{m_{i,1}}, \quad (6)$$

and $m_{e,1}$, $m_{h,1}$, and $m_{e,2}$, $m_{h,2}$, denote the effective mass

of the electron and hole inside and outside the microcrystallite. The effective masses inside and outside the quantum dots may be very different, describing semiconductor band electrons with an effective mass inside, and quasi-free electrons in a liquid ($m_{e,2} \approx m_0$) or electrons in more or less localized states in a glass matrix, $m_{e,2} \gg m_0$, outside, respectively. The variational wave function (3) was chosen because it approaches the hydrogenic $1s$ wave function for the relative electron-hole motion for $R \rightarrow \infty$ and $\lambda_s \rightarrow 1/a_B$, where a_B is the bulk-exciton Bohr radius.

From the boundary conditions that $\phi_i(r_i)$ and $(1/m_i) \partial \phi_i(r_i) / \partial r_i$ are both continuous at $r_i = R$, we obtain

$$\tan(k_{i,s} R) = \frac{k_{i,s} R}{1 - (1 + q_{i,s} R) m_{i,1} / m_{i,2}} \quad (i=e,h), \quad (7)$$

which determines the ground-state energy $E_{e,s}$ ($E_{h,s}$) of the noninteracting electron (hole) in the potential well.

The ground-state energy of the interacting electron-hole pair is computed from the expectation value of the Hamiltonian (1) with the trial wave function (3). After a straightforward but lengthy calculation, we obtain

$$E_s = \frac{\hbar^2}{2m_{e,1}} (\lambda_s^2 + k_{e,s}^2) + \frac{\hbar^2}{2m_{h,1}} (\lambda_s^2 + k_{h,s}^2) - \frac{e^2 C_s}{\epsilon N_s} + \frac{\hbar^2 \lambda_s^2}{4N_s} \left[\left(\frac{1}{m_{e,1}} - \frac{1}{m_{e,2}} \right) A_{e,s} + \left(\frac{1}{m_{h,1}} - \frac{1}{m_{h,2}} \right) A_{h,s} \right] - \frac{\hbar^2 \lambda_s^2}{2N_s} \left[\left(\frac{1}{m_{e,1}} - \frac{1}{m_{e,2}} \right) B_{e,s} + \left(\frac{1}{m_{h,1}} - \frac{1}{m_{h,2}} \right) B_{h,s} \right], \quad (8)$$

where we introduced the following quantities:

$$N_s = \int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,s}^2(r_e) \phi_{h,s}^2(r_h) F_1(2\lambda_s, r_e, r_h), \quad (9)$$

$$C_s = \int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,s}^2(r_e) \phi_{h,s}^2(r_h) \times (e^{-2\lambda_s |r_e - r_h|} - e^{-2\lambda_s |r_e + r_h|}), \quad (10)$$

$$A_{e,s} = \phi_{e,s}^2(R) \int_0^\infty dr_h r_h \phi_{h,s}^2(r_h) S_1(2\lambda_s, R, r_h), \quad (11)$$

$$B_{e,s} = \int_R^\infty dr_e r_e \int_0^\infty dr_h r_h \phi_{e,s}^2(r_e) \phi_{h,s}^2(r_h) F_1(2\lambda_s, r_e, r_h), \quad (12)$$

and, correspondingly, $A_{h,s} = A_{e,s}|_{e \leftrightarrow h}$, $B_{h,s} = B_{e,s}|_{e \leftrightarrow h}$. Furthermore, we defined

$$S_1(\lambda, x_1, x_2) = \frac{\partial}{\partial x_1} \frac{F_1(\lambda, x_1, x_2)}{x_1}, \quad (13)$$

with

$$\begin{aligned} & \frac{F_1(\lambda, x_1, x_2)}{2x_1 x_2} \\ &= \int_{-1}^1 dy y^{i-1} \exp[-\lambda(x_1^2 + x_2^2 - 2x_1 x_2 y)^{1/2}], \end{aligned} \quad (14)$$

and

$$\begin{aligned} & \frac{D_1(\lambda, x_1, x_2)}{2x_1 x_2} \\ &= \int_{-1}^1 dy y^{i-1} \exp[-\lambda(x_1^2 + x_2^2 - 2x_1 x_2 y)^{1/2}] \\ &\quad \times (x_1^2 + x_2^2 - 2x_1 x_2 y)^{1/2}, \end{aligned} \quad (15)$$

$$\frac{G_i(\lambda, x_1, x_2)}{2x_1 x_2}$$

$$= \int_{-1}^1 dy \frac{y^{i-1} \exp[-\lambda(x_1^2 + x_2^2 - 2x_1 x_2 y)^{1/2}]}{(x_1^2 + x_2^2 - 2x_1 x_2 y)^{1/2}}. \quad (16)$$

As usual, we now have to determine the variational parameter λ_s by minimizing E_s .

B. First excited state

For the first excited state we need a variational wave function that (i) can reduce to the strong-confinement product of p -state functions for the electron and hole for $R \rightarrow 0$, and (ii) can reproduce the 2s wave function for the electron-hole relative motion for large quantum dots. We choose

$$\Phi(\mathbf{r}_e, \mathbf{r}_h) = \phi_{e,p}(r_e)\phi_{h,p}(r_h)\cos(\gamma) \times (\delta - \lambda_p |\mathbf{r}_e - \mathbf{r}_h|) e^{-\lambda_p |\mathbf{r}_e - \mathbf{r}_h|}, \quad (17)$$

where

$$\phi_{i,p}(r_i) = \begin{cases} \frac{\sin(k_{i,p} r_i)}{k_{i,p}^2 r_i^2} - \frac{\cos(k_{i,p} r_i)}{k_{i,p} r_i} \\ \text{for } r_i \leq R \quad (i = e, h) \\ \frac{[\sin(k_{i,p} R) - k_{i,p} R \cos(k_{i,p} R)] q_{i,p}^2}{(q_{i,p} R + 1) k_{i,p}^2} \\ \times \frac{e^{-q_{i,p}(r_1 - R)}}{q_{i,p} r_i} \left[\frac{1}{q_{i,p} r_i} + 1 \right] \\ \text{for } r_i \geq R \quad (i = e, h). \end{cases} \quad (18)$$

With γ in Eq. (17) we denote the angle between \mathbf{r}_e and \mathbf{r}_h , and δ and λ_p are the variational parameters. One of these parameters is fixed by the condition that the wave function (17) has to be orthogonal to the ground-state wave function (3). We obtain

$$\delta = \lambda_p \frac{\int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,s}(r_e) \phi_{e,p}(r_e) \phi_{h,s}(r_h) \phi_{h,p}(r_h) F_2(\lambda_s + \lambda_p, r_e, r_h)}{\int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,s}(r_e) \phi_{e,p}(r_e) \phi_{h,s}(r_h) \phi_{h,p}(r_h) D_2(\lambda_s + \lambda_p, r_e, r_h)}. \quad (19)$$

As for the ground-state calculation, we use the boundary conditions for the wave function (17) to determine the excited-state energies of the noninteracting electrons and holes from the relation

$$\tan(k_{i,p} R) = \frac{k_{i,p} R}{(k_{i,p} R)^2} \frac{1 + \frac{m_{i,1}}{m_{i,2}} \frac{2 + 2q_{i,p} R + q_{i,p}^2 R^2}{1 + q_{i,p} R} - 2}{1 + q_{i,p} R} \quad (20)$$

where $k_{i,p}$ and $q_{i,p}$ are defined as in Eqs. (5) and (6) with the index s being replaced by p . The expectation value of the Hamiltonian (1) with the wave function (17) gives the excited-state energy of the interacting electron-hole pair as

$$E_p = \frac{\hbar^2}{2m_{e,1}} \left[\lambda_{e,p}^2 \frac{D_{e,p}}{N_p} + k_{e,p}^2 \right] + \frac{\hbar^2}{2m_{h,1}} \left[\lambda_{h,p}^2 \frac{D_{h,p}}{N_p} + k_{h,p}^2 \right] - \frac{e^2 C_p}{\epsilon N_p} + \frac{\hbar^2 \lambda^2}{4! N_p} \left[\left(\frac{1}{m_{e,1}} - \frac{1}{m_{e,2}} \right) A_{e,p} + \left(\frac{1}{m_{h,1}} - \frac{1}{m_{h,2}} \right) A_{h,p} \right] - \frac{\hbar^2 \lambda^2}{2N_p} \left[\left(\frac{1}{m_{e,1}} - \frac{1}{m_{e,2}} \right) B_{e,p} + \left(\frac{1}{m_{h,1}} - \frac{1}{m_{h,2}} \right) B_{h,p} \right], \quad (21)$$

where

$$D_{e,p} = \int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,p}^2(r_e) \phi_{h,p}^2(r_h) \{ [(1+\delta)^2 + \lambda_p^2 (r_e^2 + r_h^2)] F_3(2\lambda_p, r_e, r_h) - 2\lambda_p^2 r_e r_h F_4(2\lambda_p, r_e, r_h) - 2\lambda_p(\delta+1) D_3(2\lambda_p, r_e, r_h) \}, \quad (22)$$

$$A_{e,p} = \phi_{e,p}^2(R) \int_0^\infty dr_h r_h \phi_{h,p}^2(r_h) S_2(2\lambda_p, R, r_h), \quad (23)$$

$$B_{e,p} = \int_R^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,p}^2(r_e) \phi_{h,p}^2(r_h) \{ [(1+\delta)^2 + \lambda_p^2 (r_e^2 + r_h^2)] F_3(2\lambda_p, r_e, r_h) - 2\lambda_p^2 r_e r_h F_4(2\lambda_p, r_e, r_h) - 2\lambda_p(\delta+1) D_3(2\lambda_p, r_e, r_h) \}, \quad (24)$$

and, correspondingly, $A_{h,p} = A_{e,p}|_{e \leftrightarrow h}$, $B_{h,p} = B_{e,p}|_{e \leftrightarrow h}$. We also defined

$$C_p = \int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,p}^2(r_e) \phi_{h,p}^2(r_h) [\delta^2 G_3(2\lambda_p, r_e, r_h) - 2\lambda_p \delta F_3(2\lambda_p, r_e, r_h) + \lambda_p^2 D_3(2\lambda_p, r_e, r_h)], \quad (25)$$

$$N_p = \int_0^\infty dr_e \int_0^\infty dr_h r_e r_h \phi_{e,p}^2(r_e) \phi_{h,p}^2(r_h) \{ [\delta^2 + \lambda_p^2 (r_e^2 + r_h^2)] F_3(2\lambda_p, r_e, r_h) - 2\lambda_p^2 r_e r_h F_4(2\lambda_p, r_e, r_h) - 2\lambda_p \delta D_3(2\lambda_p, r_e, r_h) \}, \quad (26)$$

and

$$\begin{aligned} S_2(\lambda, x_1, x_2) = & \frac{\partial}{\partial x_1} \frac{1}{x_1} \{ [\delta^2 + \lambda^2(x_1^2 + x_2^2)] F_3(\lambda, x_1, x_2) \\ & - 2\lambda D_3(2\lambda, x_1, x_2) \\ & - 2\lambda^2 x_1 x_2 F_4(2\lambda, x_1, x_2) \}, \quad (27) \end{aligned}$$

where F_i and D_i are given by Eqs. (14) and (15), respectively.

III. NUMERICAL RESULTS AND DISCUSSION

We numerically evaluate the expressions derived in Sec. II and minimize E_s and E_p by variation of λ_s and λ_p , respectively. For all our calculations we choose material parameters appropriate for CdS, i.e., effective electron mass inside the quantum dot $m_{e,1}=0.235m_0$, effective hole mass $m_{h,1}=1.35m_0$, band gap $E_g=2.583$ eV, exciton binding energy $E_R=27$ meV, and exciton Bohr radius $a_B=30.1$ Å. For simplicity, we neglect tunneling effects of the hole and choose $V_{0h}=\infty$, i.e., the hole is confined inside the microcrystallites. To describe hole tunneling would require additional information about the occupied electron states outside the quantum dots, which is not available at the present time.

In Fig. 1 we plot the energies of the two quantum-confined states as function of quantum-dot radius for a confinement potential $V_{0e}=40E_R$ and $m_{e,2}=m_0$. This situation is more or less appropriate for quantum dots in a liquid solution, where we assume that the electrons can move freely once they have tunneled out of the crystallites. The solid lines are the results obtained for infinite confinement potential, i.e., ideal quantum confinement, and the dashed lines are the results of our variational cal-

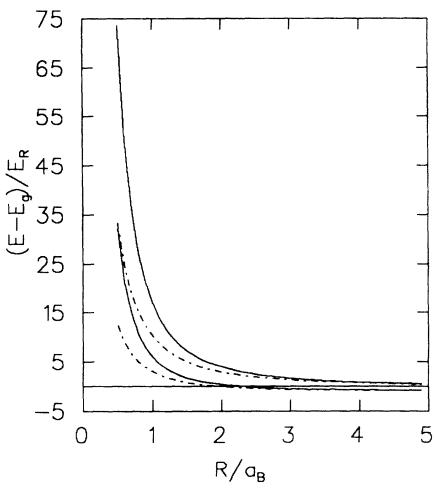


FIG. 1. Computed energies of the two lowest dipole-allowed quantum-confined states are plotted as functions of crystallite radius. The solid lines show the results for an infinite confinement potential and the dashed lines are for a confinement potential $V_0=40E_R$. The other material parameters are $m_{e,1}=0.235m_0$, $m_{e,2}=m_0$, $m_{h,1}=1.35m_0$, $E_g=2.583$ eV, $E_R=27$ meV, and $a_B=30.1$ Å.

culations. As expected, the lower potential barrier reduces the quantum-confined energy levels. The relative changes are larger for the excited state. This is due to the reduced confinement energy contribution for the case of a finite well height. This reduction is more pronounced for energetically higher states. For radii R smaller than $\approx 0.4a_B$ our calculations predict the disappearance of an excited state that is bound inside the quantum dot. In the limit $R \gg a_B$ our results approach the correct bulk semiconductor values of $-E_R$ for the ground state and $-E_R/4$ for the excited state, respectively.

To demonstrate the dependence of the energy states on the strength of the confinement potential we plot in Fig. 2 the computed energy levels as function of V_{0e} assuming a crystallite radius $R=0.5a_B$ and otherwise the same conditions as in Fig. 1. Figure 2 shows that both energy levels increase with increasing quantum confinement. The increase of the excited state is clearly more pronounced than that of the ground-state level. For a confinement potential which is less than $\approx 30E_R$ the first excited state is no more bound inside the quantum dot for the chosen parameters.

In Fig. 3 we plot the dependence of the energy levels on the electron mass outside the semiconductor material for $R=0.5a_B$ and $V_{0e}=80E_R$. We see that the energies decrease with increasing outside mass. This indicates that for quantum dots in glass, where the electrons outside are most likely in localized states simulated by a very heavy mass, the confined energy levels are lower than those for the same size quantum dots in a liquid solution. We verified the overall energy variation as function of mass ratio by solving the Schrödinger equation for a one-dimensional potential well with different masses inside and outside, using the same boundary conditions discussed above, i.e., $\phi_i(r_i)$ and $1/m_i \partial \phi_i(r_i) / \partial r_i$ are continuous at $r_i=R$. We always find that the energy of the confined states decreases with increasing outside mass.

In Fig. 4 we compare our ground-state energies for

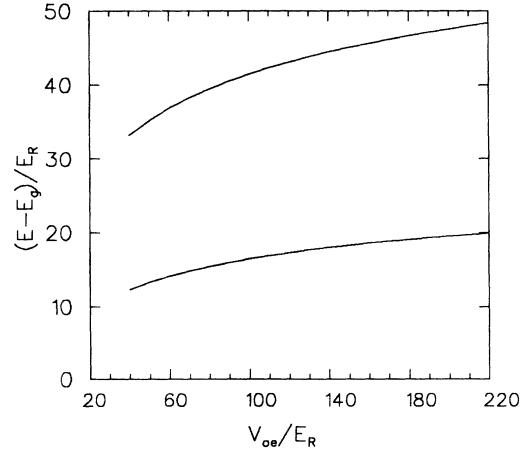


FIG. 2. Computed energies of the two lowest quantum confined states as functions of the confinement potential for a quantum dot of radius $R=0.5a_B$. The other parameters are the same as in Fig. 1.

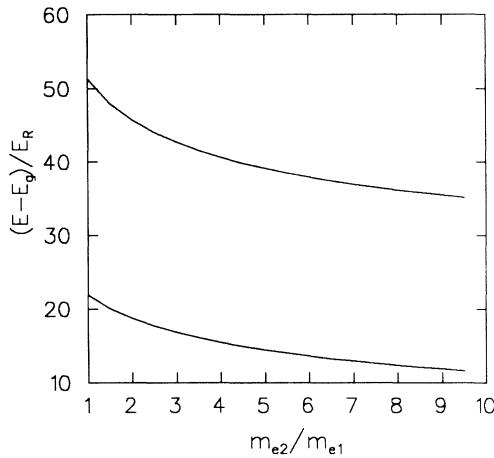


FIG. 3. Computed energies of the two lowest quantum confined states as functions of the electron mass outside the quantum dot for $R = 0.5 a_B$, $V_{0e} = 80 E_R$, and otherwise the same parameters as in Fig. 1.

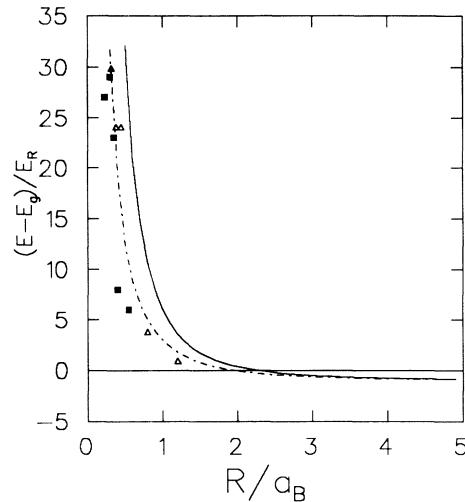


FIG. 4. Computed ground-state energy for a well depth $V_{0e} = 40 E_R$ (dashed line) and $V_{0e} = \infty$ (solid line) in comparison with the experimental results of Weller *et al.* (Ref. 4). The parameters are the same as in Fig. 1.

$V_{0e} = 40 E_R$ (dashed line) and $V_{0e} = \infty$ (solid line) to the experimental results for CdS colloids obtained by Weller *et al.*⁴ The radius of the CdS crystallites was determined by electron microscopy¹¹ (triangles) or by the fluorescence quenching method (squares).^{4,12} Our results for a finite-potential barrier show good agreement with the experimental observations, whereas the infinite-potential-barrier calculation clearly overestimates the confinement energy.

In conclusion, we have calculated the energy of the two lowest quantum-confined states of an electron-hole pair in semiconductor clusters including the tunneling effects of the electron. We found reasonably good agreement between our theoretical results and experimental data indicating the importance of the confinement conditions. Recent calculations^{15,16} have shown that the effective-mass approximation especially for the holes, may not be very good due to the mixing between the heavy- and light-hole states in small quantum dots. Clearly, for a truly quanti-

tative analysis these modifications must be taken into account, in addition to the effects considered in the present paper.

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