

## Research Article

# Resonant Light Absorption by Semiconductor Quantum Dots

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The cross-section of light absorption by size-quantized semiconductor quantum dots (QD) is calculated in the case of a resonance with an exciton  $\Gamma_6 \times \Gamma_7$  in cubical crystals of  $T_d$  class. The interference of stimulating and induced electric and magnetic fields is taken into account. The cross-section of light absorption is proportional to the exciton nonradiative damping  $\gamma$ .

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## 1. Introduction

Study of the resonant reflection and absorption of light by low-dimensional semiconductor objects is a simple and reliable way of the determination of exciton parameters [1–5]. When the size-quantized semiconductor objects (quantum wells (QW), quantum wires, and quantum dots (QD)) are irradiated by light, elastic light scattering and absorption intensify resonantly if the light frequency  $\omega_l$  equals to the exciton frequency  $\tilde{\omega}_0$ . The resonant peak width is determined by the exciton damping  $\Gamma = \gamma_r + \gamma$ , which consists of nonradiative  $\gamma$  and radiative  $\gamma_r$  damping. An important role of the radiative damping  $\gamma_r$  was proved for the first time in [6–8]. Light reflection by some structures, consisting of quantum wells, wires, and dots was considered in [9].

The light elastic scattering on a QD of arbitrary form and sizes in a resonance with excitons is investigated in [10], where the quantum perturbation theory is used. However, the quantum method does not allow to get out of the lowest order on the light-electron interaction and to calculate corrections to the exciton energy due to that interaction. Such approximation is acceptable only under condition  $\gamma_r \ll \gamma$ .

In the present work, a semiclassical method [9, 11–14] is applied for calculation of electric and magnetic fields, while a description of electrons remains quantum one. The semiclassical method allows to calculate precisely electric and magnetic fields on large distances from a QD, that is, to take into account all the orders on the light-electron interaction. It allows to introduce into the theory the nonradiative damping  $\gamma$ , to calculate the light absorption and the exciton

energy corrections due to the long-range exchange electron-hole interaction. At last, the method admits consideration of the monochromatic and pulse irradiation (see, e.g., [15]).

There are two variants of the semiclassical method. The first of them assumes using of boundary conditions for electric and magnetic fields on the semiconductor object boundaries (see, e.g., [11, 16]). However, using of boundary conditions in the case of spherical QDs, for instance, realizes into cumbersome calculations [11], and calculations become much more complicate for other forms. Therefore, we use here the second variant of the semiclassical method—the method of retarded potentials—allowing to avoid using of boundary conditions at all.

## 2. The Method of Retarded Potentials

First of all, we calculate the current  $\mathbf{j}(\mathbf{r}, t)$  and charge  $\rho(\mathbf{r}, t)$  densities induced by the electric field inside of the object [12, 17, 18] and averaged on the ground state of the crystal. Using of stimulating electric field  $\mathbf{E}_0(\mathbf{r}, t)$  in these expressions leads again to the limitation by the lowest order on the light-electron interaction. But substitution of the genuine fields  $\mathbf{E}(\mathbf{r}, t)$  inside of the object leads to the precise results.

The induced electric and magnetic fields are represented with the help of the vector  $\mathbf{A}(\mathbf{r}, t)$  and scalar  $\varphi(\mathbf{r}, t)$  potentials

$$\begin{aligned}\Delta\mathbf{E}(\mathbf{r}, t) &= -\frac{1}{c} \frac{\partial\mathbf{A}(\mathbf{r}, t)}{\partial t} - \frac{\partial\varphi(\mathbf{r}, t)}{\partial\mathbf{r}}, \\ \Delta\mathbf{H}(\mathbf{r}, t) &= \text{rot}\mathbf{A}(\mathbf{r}, t).\end{aligned}\tag{1}$$

The retarded potentials are [19, page 447]

$$\begin{aligned} \mathbf{A}(\mathbf{r}, t) &= \frac{1}{c} \int d^3 r' \frac{\mathbf{j}(\mathbf{r}', t - \nu|\mathbf{r} - \mathbf{r}'|/c)}{|\mathbf{r} - \mathbf{r}'|}, \\ \varphi(\mathbf{r}, t) &= \frac{1}{\gamma^2} \int d^3 r' \frac{\rho(\mathbf{r}', t - \nu|\mathbf{r} - \mathbf{r}'|/c)}{|\mathbf{r} - \mathbf{r}'|}, \end{aligned} \quad (2)$$

where  $\nu$  is the light refraction coefficient (we assume it is identical inside and outside of a QD). Having used the current and charge densities with the precise field  $\mathbf{E}(\mathbf{r}, t) = \mathbf{E}_0(\mathbf{r}, t) + \Delta\mathbf{E}(\mathbf{r}, t)$  and substituting (2) into (1), we obtain an integral equation for the induced field  $\Delta\mathbf{E}(\mathbf{r}, t)$  inside of the object. In some cases this equation may be solved. Having calculated the precise field inside of the object, we obtain the fields outside with the help of (2) and (1).

### 3. The Induced Current Density

The quantum theory of conductivity of low-dimensional objects is elaborated in [17, 18, 20]. The average current and charge densities induced by a weak spatially inhomogeneous electromagnetic field are calculated for the spatially inhomogeneous systems. It was shown that the averaged current and charge densities contain two contributions, first of which is expressed through the electric field, and the second is expressed through the first spatial derivatives of the electric field. Situations when the second contribution may be neglected are discussed in [20]. The main contribution into the average current density at temperature  $T = 0$  is as follows:

$$\begin{aligned} j_\alpha(\mathbf{r}, t) &= \frac{i}{\hbar} \int d^3 r' \int_{-\infty}^t dt' \\ &\times \langle 0 | [\hat{j}_\alpha(\mathbf{r}, t), \bar{d}_\beta(\mathbf{r}', t')] | 0 \rangle E_\beta(\mathbf{r}', t') + c.c., \end{aligned} \quad (3)$$

where determinations of [11] are used:  $\langle 0 | \dots | 0 \rangle$  is the average value on the ground state,  $[\hat{a}, \hat{b}]$  is the commutator of operators  $\hat{a}$  and  $\hat{b}$ ,  $\hat{\mathbf{j}}(\mathbf{r}, t)$  is the operator of current density in the interaction representation,  $\bar{\mathbf{d}}(\mathbf{r}) = \sum_i \bar{\mathbf{r}}_i \rho_i(\mathbf{r})$ ,  $\bar{\mathbf{r}}_i = \mathbf{r}_i - \langle 0 | \mathbf{r}_i | 0 \rangle$ ,  $\rho_i(\mathbf{r}) = e\delta(\mathbf{r} - \mathbf{r}_i)$ ,  $\mathbf{E}(\mathbf{r}, t)$  is the classic electric field.

We assume  $T = 0$ . It was assumed also in (3) that currents and charges are absent at infinite distances, and that the electromagnetic field equals 0 at  $t \rightarrow -\infty$ , what corresponds to adiabatic switching on of fields. We suppose an interaction between charged particles, a possible presence of an external potential and quantizing magnetic field.

We apply (3) to consideration of low-dimensional size-quantized semiconductor objects [7] and suppose that the stimulating light frequency or carrying frequency of a pulse irradiation are close to the semiconductor energy gap  $\hbar\omega_g$ . Sizes  $R$  of a semiconductor object are much more than a lattice constant  $R \gg a$ . The distances, on which the smooth multiplier from the wave function varies, are much greater  $a$  and comparable to the sizes of the object. Then the effective mass approximation is applicable and the size-quantization condition is satisfied. The last statement means that the exciton energy spectrum is discrete one. Then [7, 8],

the average density of induced current in QWs, QDs, and quantum wires is determined by the expression

$$\begin{aligned} \mathbf{j}(\mathbf{r}, t) &= \frac{ie^2}{2\pi\hbar\omega_g m_0^2} \sum_\eta \\ &\times \left\{ \mathbf{p}_{cv\eta}^* F_\eta(\mathbf{r}) \int_{-\infty}^{+\infty} d\omega \frac{e^{-i\omega t}}{\omega - \omega_\eta + i\gamma_\eta/2} \right. \\ &\times \int d\mathbf{r}' F_\eta^*(\mathbf{r}') (\mathbf{p}_{cv\eta} \mathcal{E}(\mathbf{r}', \omega)) \\ &+ \mathbf{p}_{cv\eta} F_\eta^*(\mathbf{r}) \int_{-\infty}^{+\infty} d\omega \frac{e^{-i\omega t}}{\omega + \omega_\eta + i\gamma_\eta/2} \\ &\left. \times \int d\mathbf{r}' F_\eta(\mathbf{r}') (\mathbf{p}_{cv\eta}^* \mathcal{E}(\mathbf{r}', \omega)) \right\} + c.c., \end{aligned} \quad (4)$$

where the determinations are used:  $e = -|e|$ ,  $m_0$  are the electron charge and mass, respectively,  $p_{cv\eta}$  is the interband matrix element of quasimomentum operator, corresponding to the exciton with indexes  $\eta$ ,  $F_\eta(\mathbf{r})$  is the envelope exciton wave function at  $\mathbf{r}_e = \mathbf{r}_h = \mathbf{r}$ , where  $\mathbf{r}_e(\mathbf{r}_h)$  is the electron (hole) radius-vector,  $\hbar\omega_\eta$  is the  $\eta$  exciton energy counted from the ground state energy,  $\gamma_\eta$  is the nonradiative exciton damping. The set  $\eta$  includes indexes of semiconductor valence and conduction bands, and 6 indexes, characterizing an exciton in the effective mass approximation (these 6 indexes describe the function  $F_\eta(\mathbf{r})$ ). Finally,  $\mathcal{E}(\mathbf{r}, \omega)$  is the Fourier transform of the electric field

$$\mathbf{E}(\mathbf{r}, t) = \frac{1}{2\pi} \int_{-\infty}^{+\infty} d\omega e^{-i\omega t} \mathcal{E}(\mathbf{r}, \omega) + c.c. \quad (5)$$

The main and conjugated contributions in (5) are chosen so that

$$\mathcal{E}_0(\mathbf{r}, \omega) = 2\pi E_0 \mathbf{e}_l e^{i\mathbf{k}\mathbf{r}} D_0(\omega), \quad (6)$$

where  $\mathbf{k}$  is the wave vector of stimulating light, the function  $D_0(\omega)$  describes the form of a stimulating pulse [12]. The average density of charge induced inside of a semiconductor object may be determined with the help of (4) and continuity equation

$$\text{div } \mathbf{j}(\mathbf{r}, t) + \frac{\partial \rho(\mathbf{r}, t)}{\partial t} = 0. \quad (7)$$

### 4. The Case of the Exciton $\Gamma_6 \times \Gamma_7$ in a QD

Let us consider an exciton consisting of an electron from twofold degenerated conductive band  $\Gamma_6$  and a hole from twofold degenerated valence band  $\Gamma_7$  chipped of by the spin-orbital interaction in  $T_d$  class crystals. The exciton  $\Gamma_6 \times \Gamma_7$  (see [10, 11]) is the most simple object in contrast to excitons containing light and heavy holes. All the measurable values do not depend on the direction of vectors relatively of crystallographic axes, that is, a crystal plays a role of an isotropic medium.

According to determinations of [21, page 73], the electron wave functions have the structure

$$\Psi_{c1} = iS \uparrow, \quad \Psi_{c2} = iS \downarrow, \quad (8)$$

and the hole wave functions are

$$\begin{aligned}\Psi_{h1} &= \frac{1}{\sqrt{3}}(X - iY) \uparrow - \frac{1}{\sqrt{3}}Z \downarrow, \\ \Psi_{h2} &= \frac{1}{\sqrt{3}}(X + iY) \downarrow + \frac{1}{\sqrt{3}}Z \uparrow.\end{aligned}\quad (9)$$

Having combined (8) and (9) in pairs, we obtain fourfold degenerated excitonic state for which the interband matrix elements of momentum operator result in

$$\begin{aligned}\mathbf{p}_{cv1} &= \frac{p_{cv}}{\sqrt{3}}(\mathbf{e}_x - i\mathbf{e}_y), & \mathbf{p}_{cv2} &= \frac{p_{cv}}{\sqrt{3}}(\mathbf{e}_x + i\mathbf{e}_y), \\ \mathbf{p}_{cv3} &= \frac{p_{cv}}{\sqrt{3}}\mathbf{e}_z, & \mathbf{p}_{cv4} &= -\frac{p_{cv}}{\sqrt{3}}\mathbf{e}_z,\end{aligned}\quad (10)$$

where the scalar

$$p_{cv} = i\langle S | \hat{p}_x | X \rangle \quad (11)$$

is introduced;  $\mathbf{e}_x, \mathbf{e}_y, \mathbf{e}_z$  are the unite vectors along the crystallographic axes.

We assume that the QD radius  $R$  is much less than the bulk exciton Bohr radius (strong-confinement regime). Under these conditions the binding energy of the exciton, determined by the Coulomb interaction, may be neglected. From the four exciton states, to which the interband matrix elements (10) of the quasimomentum correspond, it is easy (by drawing up the linear combinations) to obtain three bright and one dark excitonic states, about which the speech goes, for example, in [11, 22, 23]. Below we show that the corrections to the bright exciton energy, caused by long-range exchange interaction of electrons and holes, occur in our theory automatically and coincide with the results of [11, 22, 23].

## 5. Electric and Magnetic Fields on Large Distances from a QD

Let us consider light scattering by a QD near the resonance with some energy level of  $\Gamma_6 \times \Gamma_7$  exciton, when other excitons and excitonic energy levels may be neglected. Our energy level is nondegenerated, then the index  $\eta$  takes on four values from 1 to 4, and only the values  $\mathbf{p}_{cv\eta}$ , according to (10), depend on  $\eta$ . Without light-electron interaction we have

$$\begin{aligned}\omega_\eta &= \omega_0, \\ \gamma_\eta &= \gamma, & F_\eta(\mathbf{r}) &= F(\mathbf{r}),\end{aligned}\quad (12)$$

though the function  $F(\mathbf{r})$  may be chosen as a real one. After summation on  $\eta$  in (4) with using (10), we obtain

$$\mathbf{j}(\mathbf{r}, t) = \frac{ie^2 p_{cv}^2}{3\pi\hbar\omega_g m_0^2} F(\mathbf{r}) \int_{-\infty}^{\infty} d\omega e^{-i\omega t} \mathbf{T}(\omega) L(\omega) + c.c., \quad (13)$$

where

$$\begin{aligned}\mathbf{T}(\omega) &= \int d^3 r \mathcal{E}(\mathbf{r}, \omega) F(\mathbf{r}), \\ L(\omega) &= \frac{1}{\omega - \omega_0 + i\gamma/2} + \frac{1}{\omega + \omega_0 + i\gamma/2}.\end{aligned}\quad (14)$$

With the help of (7) we obtain the induced charge density

$$\rho(\mathbf{r}, t) = \frac{e^2 p_{cv}^2}{3\pi\hbar\omega_g m_0^2} \int_{-\infty}^{\infty} \frac{d\omega}{\omega} e^{-i\omega t} \left( \frac{dF(\mathbf{r})}{d\mathbf{r}} \right) \mathbf{T}(\omega) L(\omega) + c.c. \quad (15)$$

The nonresonant term from (14) is neglected below.

Using (2), we obtain electric and magnetic fields on large distances from a QD

$$\begin{aligned}\Delta \mathbf{E}_c(\mathbf{r}, t)|_{r \rightarrow \infty} &= E^+(\mathbf{r}, t) \mathbf{e}_s^+ + E^-(\mathbf{r}, t) \mathbf{e}_s^- + c.c., \\ \Delta \mathbf{H}_c(\mathbf{r}, t)|_{r \rightarrow \infty} &= H^+(\mathbf{r}, t) \mathbf{e}_s^+ + H^-(\mathbf{r}, t) \mathbf{e}_s^- + c.c.,\end{aligned}\quad (16)$$

where

$$\begin{aligned}E^\pm(\mathbf{r}, t) &= -\frac{e^2}{2\pi\hbar\omega_g m_0^2 c^2 r} \int_{-\infty}^{\infty} d\omega \omega e^{i(kr - \omega t)} Q(\mathbf{k}_s, \omega, \mathbf{e}_s^\pm), \\ H^\pm(\mathbf{r}, t) &= \mp i\nu E^\pm(\mathbf{r}, t), \\ Q(\mathbf{k}_s, \omega, \mathbf{e}_s^\pm) &= \frac{2}{3} p_{cv}^2 P(\mathbf{k}_s) (\mathbf{T}(\omega) \mathbf{e}_s^\mp) L(\omega), \\ P(\mathbf{k}) &= \int d^3 r e^{-i\mathbf{k}\mathbf{r}} F(\mathbf{r}),\end{aligned}\quad (17)$$

$\mathbf{k}_s = (\omega\nu/c)(\mathbf{r}/r)$  is the wave vector of scattered light.

To obtain precise induced electric and magnetic fields on large distances from a QD, we have to calculate the vector  $\mathbf{T}(\omega)$ , determined by the genuine electric field inside of the QD and consisting of stimulating and induced fields

$$\mathbf{E}(\mathbf{r}, t) = E_0 \mathbf{e}_l \int_{-\infty}^{\infty} d\omega e^{i(kr - \omega t)} D_0(\omega) + \Delta \mathbf{E}(\mathbf{r}, t), \quad (18)$$

where  $k = \omega\nu/c$ ,  $\mathbf{e}_l$  is the circular polarization vector. In the case of a monochromatic irradiation

$$D_0(\omega) = \delta(\omega - \omega_l), \quad (19)$$

but at a pulse irradiation the frequency  $\omega$  is widespread in the interval  $\Delta\omega$  near  $\omega_l$ , and  $\Delta\omega$  is vice inverse to the pulse duration  $\Delta t$ .

Using the formulas for retarded potentials, we obtain the induced field

$$\begin{aligned}\mathcal{E}(\mathbf{r}, \omega) &= 2\pi \mathbf{e}_l E_0 e^{i\mathbf{k}\mathbf{r}} D_0(\omega) - \frac{2e^2}{3\hbar c} \frac{p_{cv}^2 \omega}{\omega_g m_0^2 c} \\ &\times L(\omega) \left[ \mathbf{T}(\omega) + \frac{1}{k^2} \left( \mathbf{T}(\omega) \frac{d}{d\mathbf{r}} \right) \frac{d}{d\mathbf{r}} \right] \Phi(\mathbf{r}),\end{aligned}\quad (20)$$

where

$$\Phi(\mathbf{r}) = \int d^3 r' F_\eta(\mathbf{r}') \frac{e^{ik|\mathbf{r}-\mathbf{r}'|}}{|\mathbf{r}-\mathbf{r}'|}. \quad (21)$$

Equation (20) is an integral equation for  $\mathcal{E}(\mathbf{r}, \omega)$ . Then

$$\begin{aligned}\Phi(\mathbf{r}) &= \frac{1}{(2\pi)^3} \int d^3 q e^{i\mathbf{q}\mathbf{r}} P(\mathbf{q}) \left\{ \frac{4\pi P}{q^2 - k^2} + \frac{2i\pi^2}{k} \delta(q - k) \right\}, \\ \frac{P}{a - b} &= \frac{1}{2} \left( \frac{1}{a - b + i\delta} + \frac{1}{a - b - i\delta} \right), \quad \delta \rightarrow +0.\end{aligned}\quad (22)$$

Let us substitute (22) in (20), multiply both sides on  $F(\mathbf{r})$ , and integrate on  $\mathbf{r}$ . We obtain the equation for the vector  $\mathbf{T}(\omega)$ :

$$\begin{aligned} \mathbf{T}(\omega) & \left(1 + C(\omega) \int d^3q J(\mathbf{q})\right) \\ & = 2\pi E_0 \mathbf{e}_l \mathcal{D}_0(\omega) P^*(\mathbf{k}) \\ & + \frac{C(\omega)}{k^2} \int d^3q \mathbf{q} (\mathbf{q} \mathbf{T}(\omega)) J(\mathbf{q}), \end{aligned} \quad (23)$$

where determinations

$$\begin{aligned} C(\omega) & = \frac{2}{3} \left( \frac{e^2}{\hbar c} \right) \frac{p_{cv}^2 \omega}{\omega_g m_0^2 c} L(\omega), \\ J(\mathbf{q}) & = |P(\mathbf{q})|^2 \left[ \frac{1}{2\pi^2} \frac{P}{q^2 - k^2} + \frac{i}{4\pi k} \delta(q - k) \right], \end{aligned} \quad (24)$$

are introduced.

Equation (23) may be considered as a system of three equations for the components  $T_x(\omega)$ ,  $T_y(\omega)$ ,  $T_z(\omega)$ . Having solved this system, we can substitute the results into expressions for induced fields on large distances from the QD. Thus, the problem of light scattering in the resonance with an exciton  $\Gamma_6 \times \Gamma_7$  in a QD of arbitrary form is solved in principal at any relation of the light wave length and QD's sizes.

## 6. The Special Case of a Spherical Symmetrical Envelope Wave Function

Let us consider a special case when the system of three equations is reduced to one equation. It happens if the function  $P(\mathbf{k})$  depends only on the module  $k$ , that is,

$$P(\mathbf{k}) = P(k). \quad (25)$$

The condition (25) is fulfilled, if the function  $F(\mathbf{r})$  is spherically symmetrical or the QD sizes are much less than the wave length of the stimulating light when  $P(\mathbf{k}) = P(0)$ . For instance, (25) is satisfied in the case of a spherical QD limited by the infinitely high rectangular potential barrier. Then the envelope wave function

$$F(r) = \frac{1}{2\pi R} \frac{\sin^2(\pi r/R)}{r^2} \theta(R - r) \quad (26)$$

corresponds to the lowest excitonic energy level  $\omega_0$ , and

$$P(k) = \frac{2}{kR} \int_0^\pi dx \sin \frac{kRx}{\pi} \frac{\sin^2 x}{x}, \quad P(0) = 1. \quad (27)$$

Then

$$\mathbf{T}(\omega) = \frac{2\pi E_0 \mathbf{e}_l \mathcal{D}_0(\omega) P^*(k) (\omega - \omega_0 + i\gamma/2)}{\omega - \tilde{\omega}_0 + i(\gamma + \gamma_r)/2}, \quad (28)$$

where

$$\gamma_r = \frac{8\gamma}{9} \frac{e^2}{c\hbar} \frac{p_{cv}^2 \omega^2}{\omega_g m_0^2 c^2} |P(k)|^2, \quad (29)$$

$$\tilde{\omega}_0 = \omega_0 + \Delta\omega,$$

$$\begin{aligned} \Delta\omega & = -\frac{4e^2}{9\pi\hbar} \frac{p_{cv}^2}{\omega_g \omega m_0^2 \gamma^2} \int_0^\infty dq q^2 (P(q))^2 \\ & \times (q^2 - 3k^2) \frac{P}{q^2 - k^2}. \end{aligned} \quad (30)$$

The value (30) of an exciton energy correction coincides completely with the results of [6] and preceding results of [16–18], where it was considered as the result of the long-range exchange electron-hole interaction (see [8]). Having substituted (28) in (16) and (17), we obtain electric and magnetic fields on the large distances from the QD applicable in the cases of the monochromatic or pulse irradiation.

The precise results for the fields distinguish from the results of the lowest approximation on the light-electron interaction only by the substitution  $\omega_0$  by  $\tilde{\omega}_0$  and  $\gamma$  by  $\Gamma = \gamma + \gamma_r$ .

For the case (19) of the monochromatic irradiation we obtain

$$\begin{aligned} \Delta\mathbf{E}(\mathbf{r}, t)|_{r \rightarrow \infty} & = -\frac{3}{4} E_0 \frac{\gamma_r}{k_l r} [(\mathbf{e}_l \mathbf{e}_s^+) \mathbf{e}_s^+ + (\mathbf{e}_l \mathbf{e}_s^+) \mathbf{e}_s^-] \\ & \times \frac{e^{i(k_l r - \omega_l t)}}{\omega_l - \tilde{\omega}_0 + i\Gamma/2} + c.c., \end{aligned} \quad (31)$$

$$\begin{aligned} \Delta\mathbf{H}(\mathbf{r}, t)|_{r \rightarrow \infty} & = \frac{3i\gamma}{4} E_0 \frac{\gamma_r}{k_l r} [(\mathbf{e}_l \mathbf{e}_s^-) \mathbf{e}_s^+ - (\mathbf{e}_l \mathbf{e}_s^+) \mathbf{e}_s^-] \\ & \times \frac{e^{i(k_l r - \omega_l t)}}{\omega_l - \tilde{\omega}_0 + i\Gamma/2} + c.c., \end{aligned}$$

where  $k_l = \omega_l \nu / c$ .

## 7. The Pointing Vector on Large Distances from the QD and the Total Cross-Section of Light Scattering at the Monochromatic Irradiation

The Pointing vector on large distances from the QD equals

$$\mathbf{S}|_{r \rightarrow \infty} = \mathbf{S}_0 + \mathbf{S}_{\text{interf}} + \mathbf{S}_{\text{scat}}, \quad (32)$$

where

$$\mathbf{S}_0 = \frac{c}{4\pi} \mathbf{E}_0 \times \mathbf{H}_0 = \frac{c\nu}{2\pi} E_0^2 \mathbf{e}_z,$$

$$\mathbf{S}_{\text{interf}} = \frac{c}{4\pi} [\mathbf{E}_0 \times \Delta\mathbf{H} + \Delta\mathbf{E} \times \mathbf{H}_0], \quad (33)$$

$$\mathbf{S}_{\text{scat}} = \frac{c}{4\pi} \Delta\mathbf{E} \times \Delta\mathbf{H}.$$

With the help of (31) we obtain

$$\mathbf{S}_{\text{scat}} = \frac{9\pi}{4} S_0 \frac{\gamma_r^2}{(k_l r)^2} \frac{\mathbf{r}}{r} \frac{|\mathbf{e}_l \mathbf{e}_s^-|^2 + |\mathbf{e}_l \mathbf{e}_s^+|^2}{(\omega_l - \tilde{\omega}_0)^2 + \Gamma^2/4}. \quad (34)$$

The module of the total flux of scattered light per time unite is

$$\Pi_{\text{scat}} = \frac{3\pi}{2} S_0 \frac{\gamma_r^2}{(k_l)^2} \frac{1}{(\omega_l - \tilde{\omega}_0)^2 + \Gamma^2/4}. \quad (35)$$

Obtaining (35) from (34) we have used the relations

$$\begin{aligned} |\mathbf{e}_l^+ \mathbf{e}_s^-|^2 &= |\mathbf{e}_l^- \mathbf{e}_s^+|^2 = \frac{1}{4}(1 + \cos \theta)^2, \\ |\mathbf{e}_l^+ \mathbf{e}_s^+|^2 &= |\mathbf{e}_l^- \mathbf{e}_s^-|^2 = \frac{1}{4}(1 - \cos \theta)^2, \end{aligned} \quad (36)$$

where  $\theta$  is the scattering angle. An integration  $\theta$  is performed. Having divided  $\Pi_{\text{scat}}$  on the density  $S_0$  of the stimulating light flux and using  $k_l = 2\pi/\lambda_l$ , we obtain the total cross-section of light scattering

$$\sigma_{\text{scat}} = \frac{3}{2\pi} \lambda_l^2 \frac{\gamma_r^2/4}{(\omega_l - \tilde{\omega}_0)^2 + \Gamma^2/4}. \quad (37)$$

The differential cross-sections have been determined in [8].

## 8. The Light Absorption: The Role of the Nonradiative Damping

The results for the cross-sections of light scattering on QD, obtained with the help of the quasiclassic method, coincide with the lowest approximation on the light-electron interaction with the results of the quantum perturbation theory [10]. However, the quasiclassic method allows to obtain also the cross-section of light absorption by a QD. In the case of a monochromatic irradiation the absorption is stipulated by the nonradiative damping  $\gamma$  of excitons, and it equals 0 at  $\gamma = 0$ . The same result was obtained at a monochromatic irradiation of a QW [7, 24]. The cause is that at  $\gamma = 0$  a dissipation of energy, spent on an exciton creation, is absent. And this energy returns at an exciton annihilation. In the case of a pulse irradiation the integral absorption equals 0 at  $\gamma = 0$  [15, 25–27].

It was shown at calculations of light absorption by a QW that the interference of stimulating and induced electromagnetic fields must be taken into account (see, e.g., [25]). The same is true for the case of QDs.

## 9. The Interference Contribution into Energy Fluxes

Let us calculate an interference contribution into the Pointing vector at  $r \rightarrow \infty$  in the case of a monochromatic irradiation. Using (31) and

$$\begin{aligned} \mathbf{E}_0^\pm(\mathbf{r}, t) &= E_0 \mathbf{e}_l^\pm e^{i(\mathbf{k}_l \mathbf{r} - \omega_l t)} + c.c., \\ \mathbf{H}_0^\pm(\mathbf{r}, t) &= E_0 \gamma \left[ \mathbf{e}_z \times \mathbf{e}_l^\pm \right] e^{i(\mathbf{k}_l \mathbf{r} - \omega_l t)} + c.c., \end{aligned} \quad (38)$$

we obtain

$$\mathbf{S}_{\text{interf}} = \mathbf{S}_z + \mathbf{S}_\perp, \quad (39)$$

$$\mathbf{S}_z = -\frac{3}{4} \frac{\gamma_r}{k_l r} S_0 \mathbf{e}_z \left| \mathbf{e}_l^+ \mathbf{e}_s^- \right|^2 \left( \frac{e^{i(\mathbf{k}_l \mathbf{r} - k_l r)}}{\omega_l - \tilde{\omega}_0 - i\Gamma/2} + c.c. \right), \quad (40)$$

$$\mathbf{S}_\perp = \frac{3}{4} \frac{\gamma_r}{k_l r} S_0 \left( \mathbf{e}_l^+ (\mathbf{e}_l^- \mathbf{e}_s^+) (\mathbf{e}_s^- \mathbf{e}_z) \frac{e^{i(\mathbf{k}_l \mathbf{r} - k_l r)}}{\omega_l - \tilde{\omega}_0 - i\Gamma/2} + c.c. \right), \quad (41)$$

where  $+$ ( $-$ ) correspond to the right (left) circular polarization of the stimulating light.

Since (40) and (41) are applicable at  $r \rightarrow \infty$ , it is obviously that due to the factor  $e^{i(\mathbf{k}_l \mathbf{r} - k_l r)}$  only angles  $\theta \rightarrow 0$  can contribute into the constant energy flux. However, there is the factor  $(\mathbf{e}_l^- \mathbf{e}_z)$  in the RHS of (41), which equals 0 at  $\theta = 0$ . Therefore, the constant energy flux on the large distances from a QD corresponds only to the vector  $\mathbf{S}_z$ .

Let us calculate the energy flux

$$\mathbf{\Pi}_z = \int ds \mathbf{S}_z, \quad (42)$$

going through a plane perpendicular to the stimulating light direction  $z$  in a time unite on large distances behind of a QD. A surface element equals  $ds = \rho d\rho d\varphi$ , and  $\rho = z \tan \theta$ ,  $r = z/\cos \theta$ , where  $\theta$  is the angle between  $\mathbf{k}_l$  and  $\mathbf{r}$ . Having integrated on the angle  $\theta$  which gives  $2\pi$ , using (36) and going from the variable  $\rho$  to the variable  $\theta$ , we obtain

$$\begin{aligned} \mathbf{\Pi}_z &= -\frac{3\pi}{8} \frac{\gamma_r z}{k_l} \mathbf{e}_z \frac{S_0}{\omega_l - \tilde{\omega}_0 - i\Gamma/2} \int_0^{\pi/2} d\theta \sin \theta \\ &\times \left( \frac{1 + \cos \theta}{\cos \theta} \right)^2 e^{ik_l z(1-1/\cos \theta)} + c.c. \end{aligned} \quad (43)$$

Further we substitute variable  $\theta$  by the variable  $t = (\cos \theta)^{-1} - 1$  and obtain

$$\begin{aligned} \mathbf{\Pi}_z &= -\frac{3\pi}{8} \frac{\gamma_r z}{k_l} \mathbf{e}_z \frac{S_0}{\omega_l - \tilde{\omega}_0 - i\Gamma/2} \\ &\times \int_0^\infty dt \left( \frac{2+t}{1+t} \right)^2 e^{-ik_l z t} + c.c. \end{aligned} \quad (44)$$

At  $z \rightarrow \infty$

$$\begin{aligned} &z \int_0^\infty dt \left( \frac{2+t}{1+t} \right)^2 e^{-ik_l z t} \\ &\rightarrow 4z \int_0^\infty dt e^{-ik_l z t} = -\frac{4i}{k_l} (1 - e^{-ik_l z}), \end{aligned} \quad (45)$$

and we obtain

$$\begin{aligned} \mathbf{\Pi}_z &= \frac{3\pi}{2} \frac{\gamma_r}{k_l^2} \mathbf{e}_z S_0 \left( \frac{i}{\omega_l - \tilde{\omega}_0 - i\Gamma/2} + c.c. \right) \\ &= -\frac{3\pi}{2} \frac{\mathbf{e}_z}{k_l^2} S_0 \frac{\gamma_r \Gamma}{(\omega_l - \tilde{\omega}_0)^2 + \Gamma^2/4}. \end{aligned} \quad (46)$$

In (46), the terms going to 0 at  $z \rightarrow \infty$ , and rapidly oscillating with  $z$  are neglected. Analogically we calculate the contribution  $\mathbf{\Pi}_\perp = \int ds \mathbf{S}_\perp^\pm$  into the energy flux and find that at  $z \rightarrow \infty$  the value  $\mathbf{\Pi}_\perp \rightarrow 0$ .



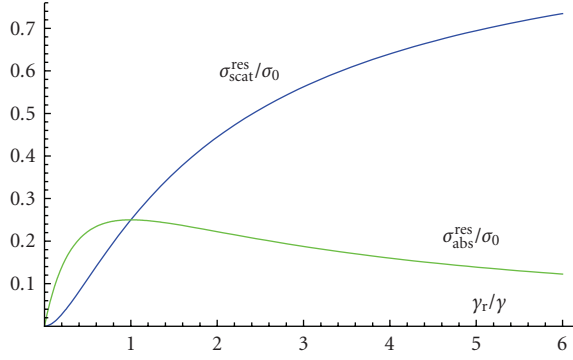


FIGURE 1:  $\sigma_{\text{scat}}^{\text{res}}$  and  $\sigma_{\text{abs}}^{\text{res}}$  as functions of  $\gamma_r/\gamma$  in the resonance  $\tilde{\omega}_0 = \omega_l$ .  $\sigma_0 = (3/2\pi)\lambda_l^2$ .

## 10. The Cross-Section of Light Absorption

Since  $\Gamma = \gamma + \gamma_r$ , the flux  $\Pi_z$  may be represented as two parts

$$\Pi_z = -\mathbf{e}_z \Pi_{\text{scat}} - \mathbf{e}_z \Pi_{\text{abs}}, \quad (47)$$

where  $\Pi_{\text{scat}}$  is determined by (35),

$$\Pi_{\text{abs}} = \frac{3\pi}{2} S_0 \frac{\gamma_r \gamma / k_l^2}{(\omega_l - \tilde{\omega}_0)^2 + \Gamma^2/4}. \quad (48)$$

Obviously that the energy flux  $-\mathbf{e}_z \Pi_{\text{scat}}$  compensates the total flux of the scattered energy, and  $\Pi_{\text{abs}}$  corresponds to the energy, absorbed by the QD per time unite at the monochromatic irradiation. Having divided  $\Pi_{\text{abs}}$  on the density  $S_0$  of the stimulating energy flux, we obtain the total cross-section of light absorption

$$\sigma_{\text{abs}} = \frac{3}{2\pi} \lambda_l^2 \frac{\gamma_r \gamma / 4}{(\omega_l - \tilde{\omega}_0)^2 + \Gamma^2/4} \quad (49)$$

and reduces to 0 at  $\gamma = 0$ , as it was supposed in advance.

Comparing (49) and (37) we find that in the lowest approximation on the light-electron interaction the absorption cross-section is of the first order on that interaction (it contains the factor  $e^2/c\hbar$ ), and the scattering cross-section is of the second order on that interaction (it contains the factor  $(e^2/c\hbar)^2$ ). The ratio of the scattering and absorption cross-sections equals  $\gamma_r/\gamma$ . The absorption cross-section has the maximum value at comparable values  $\gamma_r$  and  $\gamma$ . At  $\gamma_r = \gamma$  in the resonance  $\omega_l = \tilde{\omega}_0$

$$\sigma_{\text{scat}}^{\text{res}} = \sigma_{\text{abs}}^{\text{res}} = \frac{3}{8\pi} \lambda_l^2. \quad (50)$$

$\sigma_{\text{scat}}^{\text{res}}$  and  $\sigma_{\text{abs}}^{\text{res}}$  as functions of  $\gamma_r/\gamma$  are represented in Figure 1.

## 11. Conclusion

Thus, the electric and magnetic fields induced at the light irradiation of QDs are calculated in the resonance of the stimulating light and excitons with the help of the semiclassical method of the retarded potentials. The fields on the large distances are calculated exactly.

The concrete calculations are performed for the excitons  $\Gamma_6 \times \Gamma_7$  in cubic crystals of  $T_d$  class. It is shown that in the case of small or spherically symmetric QDs the light-electron interaction realizes into the substitution of the nonradiative excitonic damping  $\gamma$  by the damping  $\Gamma = \gamma + \gamma_r$  and the exciton energy  $\omega_0$  by the energy  $\tilde{\omega}_0 = \omega_0 + \Delta\omega$ , where  $\gamma_r$  is the radiative damping,  $\Delta\omega$  is the correction to the exciton energy due to the long-range exchange interaction of electrons and holes.

The light scattering and absorption cross-sections by QDs are obtained for arbitrary QDs under condition  $R \ll \lambda_l$  and excitonic spherical envelope wave functions.

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## Special Issue on Polymer Nanocomposite Processing, Characterization, and Applications

### Call for Papers

Polymers reinforced with nanoparticles, such as carbon nanotubes, are of great interest due to their remarkable mechanical, thermal, chemical properties as well as optical, electronic, and magnetic applications. In the general research area of polymer nanocomposites, a number of critical issues need to be addressed before the full potential of polymer nanocomposites can actually be realized. While a number of advances have recently been made in the area of polymer nanocomposites, the studies on understanding of the effects of processing parameters on the structure, morphology, and functional properties of polymer nanocomposites are deficient. There is a need for characterization techniques to quantify the concentration and distributions of nanoparticles as well as to assess the strength at the interface between the polymer and nanoparticles. Also, there is a need for the development of better models able to predict the mechanical properties of the polymer nanocomposites as functions of myriad factors including nanoparticle orientation, the type of functional groups, and the molecular weight of polymer chain. The relationships between the structural distributions and the ultimate properties of the polymer nanocomposites also need to be elucidated.

This special issue of the Journal of Nanomaterials will be devoted to emerging polymer nanocomposite processing techniques and call for new contributions in the field of characterization and applications of multifunctional nanocomposites. It intends to cover the entire range of basic and applied materials research focusing on rheological characterization, nanoparticle dispersion, and functional properties of polymer nanocomposites for sensors, actuators, and other applications. Fundamental understanding of the effects of processing and nanoparticles on the polymer structure and morphology, their optical, electrical, and mechanical properties as well as novel functions and applications of nanocomposite materials will be the highlights of this special issue.

Papers are solicited in, but not limited to, the following areas:

- Solution and melt processing of polymer nanocomposites
- Rheological and thermal characterization of nanocomposites

- Generation of nanofibers using extrusion and electrospinning of nanocomposites
- Processing-induced orientation of nanoparticles
- Quantification of nanoparticle dispersion
- Effect of nanoparticle incorporation on polymerization
- In situ nanoparticle formation in polymer matrix
- Noncovalent functionalization techniques and characterization of properties at polymer-nanoparticle interface
- Novel applications of polymer nanocomposites

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## Special Issue on Nanomaterials for Cancer Diagnosis and Therapy

### Call for Papers

Integration of nanotechnology into the treatment of various diseases such as cancers represents a mainstream in the current and future research due to the limitations of traditional clinical diagnosis and therapy. The early detection of cancer has been universally accepted to be essential for treatment. However, it remains challenging to detect tumors at an early stage. For instance, the goal of molecular imaging in breast cancer is to diagnose the tumor with approximately 100–1000 cells, compared to the traditional techniques which may require more than a million cells for accurate clinical diagnosis. On another hand, anticancer drugs are designed to simply kill cancer cells, and their entrance into healthy organs or tissues is undesirable due to the severe side effects. In addition, the rapid and widespread distribution of anticancer drugs into nontargeted organs and tissues requires a lot of drugs with high cost. These difficulties have largely limited the successful therapy of cancer.

Nanomaterials are anticipated to revolutionize the cancer diagnosis and therapy. The development of multifunctional polymeric nanoparticles allows for the early detection of cancers. The construction of intelligent polymeric nanosystems can be used as controlled delivery vehicles to improve the therapy efficiency of anticancer drugs, that is, such vehicles are capable of delivering drugs to predetermined locations and then releasing them with preprogrammed rates in response to the changes of environmental conditions such as pH and temperature. Besides polymers, these nanomaterials can also be composed of supraparamagnetic iron oxide, carbon nanotube, metallic nanoshell, core-shell aggregate, or composites. These nanomaterials represent new directions for more effective drug administration in cancer.

This special issue of the Journal of Nanomaterials will cover a wide range of nanomaterials for cancer diagnosis and therapy. It will mainly focus on the preparations, characterizations, functionalizations, and properties of nanoparticles, nanostructured coatings, films, membranes, nanoporous materials, nanocomposites, and biomedical devices. Fundamental understanding of the basic mechanisms on material and biological processes related to the unique nanoscale properties of the materials will be the highlights of this special issue.

Papers are solicited in, but not limited to, the following areas:

- Synthesis and functionalization of polymer nanoparticle/nanomicelle/nanocomplex
- Polymer nanoparticle/hydrogel for drug delivery
- Synthesis of intelligent nanogel
- Hydrogel in nanoscale sensing
- Supraparamagnetic nanoparticle for magnetic resonance imaging applications
- Carbon nanotube-based devices for drug delivery
- Core-shell nanoparticle for molecular imaging
- Metallic nanoshell for drug delivery
- Nanoporous and nanoscaled materials for drug delivery
- Nanotechnologies for targeted delivery
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## Special Issue on Phonons and Electron Correlations in High-Temperature and Other Novel Superconductors

### Call for Papers

It has been now over 20 years since the discovery of the first high-temperature superconductor by Georg Bednorz and Alex Müller and yet, despite intensive effort, no universally accepted theory exists about the origin of superconductivity in cuprates. The absence of consensus on the physics of cuprate superconductors and the recent discovery of iron-based compounds with high transition temperatures have re-emphasized the fundamental importance of understanding the origin of high-temperature superconductivity. First-principles calculations based on density functional theory (DFT) often predict a rather weak electron-phonon interaction (EPI) insufficient to explain high transition temperatures. A number of researchers are of the opinion that the EPI may be considerably enhanced by correlation effects beyond DFT. Others maintain that the repulsive electron-electron interaction in novel superconductors is pairing and provides high transition temperatures without phonons. On the other hand, some recent studies using numerical techniques cast doubt that simple repulsive models can account for high-temperature superconductivity. Therefore, it seems plausible that the true origin of high-temperature superconductivity could be found in a proper combination of strong electron-electron correlations with a significant EPI.

This Special Issue will become an international forum for researchers to summarize recent developments in the field, with a special emphasis on the results in high-temperature superconductors and some other related materials that combine sizeable electron-phonon coupling with strong correlations. We invite authors to present original research papers as well as summarizing overviews stimulating the continuing efforts to understand high-temperature and other unconventional superconductors. Potential topics include, but are not limited to:

- Phonon spectroscopies of cuprates and related compounds
- Experimental evidence for electron-phonon interactions
- First-principles calculations of EPIs
- Strong-coupling extensions of the BCS-Eliashberg theory including polarons and bipolarons

- Interplay of electron-electron and electron-phonon interactions (Holstein-Hubbard, Fröhlich-Hubbard, tJ-Holstein, and similar models)
- Phase separation of correlated electrons
- EPI effects in fullerenes, MgB<sub>2</sub>, ruthenates, ferropnictides, and other novel noncuprate superconductors
- Routes to higher temperature superconductivity

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