Exciton-polaritons in cuprous oxide: Theory and comparison with experiment

Frank Schweiner, Jan Ertl, Jörg Main, and Günter Wunner

Institut für Theoretische Physik 1, Universität Stuttgart, 70550 Stuttgart, Germany

Christoph Uhlein

Experimentelle Physik 2, Technische Universität Dortmund, 44221 Dortmund, Germany

(Dated: July 16, 2018)

PACS numbers: 71.36.+c, 71.35.-y, 78.40.-q, 71.20.Nr

I. INTRODUCTION

Excitons are Coulomb-bound pairs of a positively charged hole in the valence band and a negatively charged electron in the conduction band of a semiconductor. Hence, these elementary excitations of a semiconductor are often regarded as the hydrogen analog of the solid state. It is now more than 80 years since Frenkel [1–3], Peierls [4], and Wannier [5] formulated the concept of excitons. After the experimental discovery of these quasi-particles in cuprous oxide (Cu$_2$O) by Gross and Karryjew in 1952 [6], excitons in bulk semiconductors became an important topic in solid-state physics from the late 1950s to the 1970s (see, e.g., Refs. [7–12] and further references therein).

Very recently, new attention has been drawn to the field of excitons by an experimental observation of the yellow exciton series in Cu$_2$O up to a large principal quantum number of $n = 25$ [13]. This discovery has opened up the research field of giant Rydberg excitons and led to a variety of new experimental and theoretical investigations [13–36]. Furthermore, it may pave the way, e.g., to a deeper understanding of inter-particle interactions in the solid [13] and to applications in quantum information technology [27].

Even though the spectrum of Rydberg excitons in Cu$_2$O can be described quite well in a first approximation by the hydrogen-like model of Wannier, one must keep in mind that excitons are complex many-body states of the solid and, hence, that there are significant limitations to the hydrogen-like model and to the atom-like description of these quasi-particles [37].

Some essential corrections to the hydrogen-like model comprise, e.g., the inclusion of the complete cubic valence band structure [38–43], which leads to a complicated fine-structure splitting, the central-cell corrections, which account for deviations from the hydrogen-like model in the limit of a small exciton radius [7, 44–50], and the exchange interaction [7, 47, 48, 51, 52]. Furthermore, interactions with other quasi-particles, like, e.g., phonons have to be considered [53–55]. All of these effects have already been discussed recently for the excitons in Cu$_2$O [15, 18, 20, 21, 25, 28].

Above all, there is another fundamental difference between atoms and excitons as regards their interaction with light. By analogy with the interaction of atoms with light one may suppose that absorption of light in a crystal can be described as the excitation of an exciton with the simultaneous disappearance of a photon [7]. Indeed, in the weak-coupling limit, the incident light acts only as a perturbation on the different energy states or excitations of the solid like, e.g., excitons [47]. However, since the excited states in the solid are connected with a polarization and since an oscillating polarization emits again an electromagnetic wave acting back onto the incident wave, there is an interplay between light and matter. If
the frequency of light is within the range of the resonance frequency of an excitation, the coupling is strong and thus anomalous dispersion can be observed [7]. Due to this coupling, excitons and photons cannot be treated as independent entities or good eigenstates, but new quasiparticles must be introduced, which are called polaritons and which represent the quanta of the mixed state of polarization and electromagnetic wave [56–64].

In this regard the 1S ortho exciton of CuO, in particular, is of interest as for this state characteristic polariton effects like propagation beats and the conservation of coherence over macroscopic distances have already been demonstrated experimentally [65]. Furthermore, the threefold degeneracy of this state is lifted for finite momentum of the center of mass \( hK \), and its oscillator strengths [66] as well as its dispersion are anisotropic [20, 67–69]. All of these effects and the number of experimental results for, e.g., the oscillator strength, the \( K \)-dependent spectra and the group velocity [66], with which theoretical results can be compared, make the 1S ortho exciton an ideal candidate for theoretical investigations.

In this paper we present the theory of exciton-polaritons in CuO. We extend the Hamiltonian of Ref. [28], which accounts for the complete valence band structure, the exchange interaction, and the central-cell corrections, for a finite momentum \( hK \) of the center of mass. The corresponding Schrödinger equation can then be solved using a complete basis. This method allows us not only to calculate dipole and quadrupole oscillator strengths but also the size of the nonanalytic exchange interaction. As the splitting due to the non-analytic exchange interaction at \( K = 0 \) is identical to the longitudinal-transverse splitting (LT-splitting) when treating polaritons, this interaction needs to be considered for a correct treatment of the complete problem. We show how to calculate the polariton dispersion for the complete exciton spectrum.

Due to the presence of the complete valence band structure, the exchange interaction, and the central-cell corrections, we obtain very complex polariton dispersions for the excitons in CuO, which by far exceed previous investigations on the polariton dispersion using a simple hydrogen-like model [26]. The main emphasis in this paper is on the only exciton state for which a pronounced polariton effect has been observed in several experiments: the 1S ortho exciton state. At first, we discuss the effects of a finite momentum \( hK \) of the center of mass on the spectrum. The \( K \)-dependent splitting, which has been discussed in Refs. [20, 67–69], is now treated using the \( K \)-dependent Hamiltonian in explicit form and including the central-cell corrections. The calculated splittings are in the same order of magnitude as the splittings observed experimentally [67–69].

Previous publications on the 1S ortho exciton polariton [26, 65, 66, 70] were confining the polariton problem to situations where the \( K \)-vector is parallel to one of the principal symmetry axes of the crystal. This ensures that the polarization of the exciton is always parallel to the electric field vector and perpendicular to the wave vector. What is still missing is a proper treatment of the general polariton problem with an arbitrary orientation of the \( K \)-vector, which considers the \( K \)-dependent splitting and the polariton effect simultaneously. The challenge is the anisotropy of the exciton-photon interaction, leading to a polarization that is neither parallel to the electric field nor orthogonal to the wave vector. Solving this very complex problem is a significant step ahead in a proper treatment of the exciton polariton problem. We are able to show that the general polariton problem can be reduced to solving an eigenvalue problem involving two photon states and three exciton states; thus giving rise to five distinct polariton branches. The general case leads to polariton states with transverse and longitudinal polarization components. These hybrid polaritons states can therefore be regarded as a mixture between a transverse polariton and a longitudinal exciton.

Using the 5 \( \times \) 5 matrix model for the anisotropic dispersion of the 1S ortho exciton-polariton we are able to calculate the group velocity, the oscillator strengths, and spectra for different orientations of \( K \) and to compare them with experimental results [65, 66, 68, 70]. We especially discuss the two cases of the polariton spectra in the planes perpendicular to [110] and [111]. In the first case the vector \( K \) is oriented in one of the six mirror planes of the cubic group \( O_h \) so that the polariton states can be classified according to two different irreducible representations of the group \( C_\gamma \). As regards the second case, there is no such symmetry left and five polariton branches are obtained independently from the polarization of light. We are able to compare our results with experimental transmission spectra. The good agreement between theory and experiment not only emphasizes the special nature of the 1S ortho exciton polariton but also proves the validity of the 5 \( \times \) 5 matrix model as a useful theoretical model for further investigations on this polariton.

Our investigations with respect to the exciton-polariton problem in CuO are gaining additional importance by recent second harmonic generation (SHG) measurements performed by J. Mund et al. [71]. The experiments were carried out with a laser generating ultra short pulses having a band width covering several meV. The resonant excitation of the 1S ortho exciton by means of a two-photon absorption process generates exciton polaritons that are giving rise to SHG radiation. The polariton nature of the quadrupole exciton explains why it is possible to generate SHG in a crystal with inversion symmetry.

The paper is organized as follows: In Sec. II we present the Hamiltonian of excitons in CuO when considering a finite momentum \( hK \) of the center of mass and show how to solve the corresponding Schrödinger equation in a complete basis. In Sec. III we present formulas to calculate dipole and quadrupole oscillator oscillator strengths. Having introduced the multi-polariton con-
cept of exciton-polaritons in Sec. IV A, we discuss the rotating wave approximation, the nonanalytic exchange interaction and criteria for the observability of polariton effects in Secs. IV B, IV C, and IV D, respectively. In Sec. V we then use the Hamiltonian, which accounts for the complete valence band structure, the exchange interaction and the central-cell corrections to calculate the complex dispersion of the polaritons connected with the excitons states of $2 \leq n \leq 4$. We also discuss the observability of polariton effects. The $K$ dependent splitting and the dispersion of the 1S ortho exciton polariton are treated separately from the other polaritons in Sec. VI.

After the discussion of the splitting in Sec. VI A, we introduce the $5 \times 5$ matrix model in Sec. VI B. The dispersion of the ortho exciton polariton, the group velocity and the oscillator strengths are compared with experimental results in Sec. VI C. We finally give a summary and outlook in Sec. VII.

II. HAMILTONIAN

In this section we shortly present the theory of excitons with a finite momentum of the center of mass in Cu$_2$O, where the cubic valence band structure, the exchange interaction and the central-cell corrections need to be considered.

The Hamiltonian of the exciton is given by

$$H = E_\text{g} + V(r_e - r_h) + H_e(p_e) + H_h(p_h) + H_{\text{CCC}}(r_e - r_h) + H_{\text{exch}}(r_e - r_h)$$

(1)

with the energy $E_\text{g}$ of the band gap. The screened Coulomb interaction $V$, the central cell corrections $H_{\text{CCC}}$, the analytic exchange exchange interaction $H_{\text{exch}}$ and the kinetic energies $H_e(p_e)$ and $H_h(p_h)$ of the electron and the hole are given in Ref. [28].

We now introduce relative and center-of-mass coordinates

$$r = r_e - r_h,$$  \hspace{1cm} (2a)

$$R = \alpha r_e + \gamma r_h.$$  \hspace{1cm} (2b)

The factors $\alpha$ and $\gamma$ are in general $3 \times 3$ matrices with $|\det(\alpha + \gamma)| = 1$ [42]. As the Hamiltonian (1) depends only on the relative coordinate $r = r_e - r_h$, electron and hole, the momentum of the center of mass

$$P = p_e + p_h = -i\hbar(\alpha + \gamma) \nabla R$$

(3)

with

$$\nabla r_e = \alpha \nabla R + \nabla r, \hspace{1cm} (4a)$$

$$\nabla r_h = \gamma \nabla R - \nabla r.$$  \hspace{1cm} (4b)

is a constant of motion, i.e., we can set $P = \hbar K$ [42, 72]. According to Ref. [42], the matrices $\alpha$ and $\gamma$ can be chosen arbitrarily, as long as $|\det(\alpha + \gamma)| = 1$ holds, but should be adapted to the problem. Note that if we insert Eqs. (2) and (4) into the Hamiltonian (1) in general a coupling term between the relative motion and the motion of center of mass appears. Only for a specific choice of $\alpha$ and $\gamma$ this coupling term vanishes. However, the correct values of $\alpha$ and $\gamma$ are difficult to find [20, 72]. In particular, the generalized relative and center of mass coordinate transformation of Ref. [20] holds only if the parameters $\eta_i$ in the kinetic energy of the hole are set to zero (cf. Ref. [20]). When assuming isotropic coefficients $\alpha = \alpha_1$ and $\gamma = (1 - \alpha_1)1$ it is not possible to find a constant value of $\alpha$ for which the relative motion and the motion of the center of mass are decoupled. This is connected to the fact that the exciton mass in Cu$_2$O is not isotropic [20]. Hence, the more complicated generalized transformation with $3 \times 3$ matrices would be needed.

Nevertheless, the results of the Hamiltonian are independent of the choice of $\alpha$ and $\gamma$. As we extend the theory of excitons in semiconductors with a cubic valence band structure for polariton effects, we simply use the coordinates and momenta of relative and center of mass motion with $\alpha = m_e/(m_e + m_h)\,1$ and $\gamma = m_h/(m_e + m_h)\,1$ known from the hydrogen atom in the following:

$$r = r_e - r_h,$$  \hspace{1cm} (5a)

$$R = (m_e r_e + m_h r_h)/(m_e + m_h),$$  \hspace{1cm} (5b)

$$p = (m_e p_e + m_h p_h)/(m_e + m_h),$$  \hspace{1cm} (5c)

$$P = p_e + p_h = \hbar K.$$  \hspace{1cm} (5d)

We can then write the Hamiltonian in the form

$$H = H_0 + (\hbar K) H_1 + (\hbar K)^2 H_2.$$  \hspace{1cm} (6)

The first part $H_0$ is exactly the Hamiltonian of relative motion presented and discussed in Ref. [28], while the last part $H_2$ describes the motion of the center of mass in the degenerate band case. The term $H_1$ depends on the relative momentum $p$ and thus couples the relative motion and the motion of the center of mass.

For the case that the wave vector $K$ is oriented along one of the directions of high symmetry, i.e., along [001], [110] or [111], one can rotate the coordinate system to make the quantization axis coincide with the direction of $K$ and then express the Hamiltonian (96) in terms of irreducible tensors [40, 76–78]. Explicit expressions for $H_0, H_1,$ and $H_2$ for these special orientations of the wave vector $K$ are given in the Supplemental Material [79].

The Schr"odinger equation corresponding to the Hamiltonian (96) is solved using the method presented in Refs. [20, 21, 28] with a complete basis. The ansatz for the exciton wave function now reads

$$|\Psi_{\nu K}\rangle = \sum_{N,LFF, M_{F_1}} c_{NLFF, M_{F_1}}^{K} |\Pi\rangle,$$  \hspace{1cm} (7a)

$$|\Pi\rangle = |N, L; (I, S_h) J; F, S_c; F_1, M_{F_1}\rangle,$$  \hspace{1cm} (7b)

with complex coefficients $c$ and the quantum numbers explained in Refs. [20, 21, 28]. Note that the coefficients
TABLE I: Material parameters used in the calculations.

| Property                        | Value          | Source   |
|---------------------------------|----------------|----------|
| band gap energy                 | $E_g = 2.17208$ eV | [13]     |
| electron mass                   | $m_e = 0.99 \, m_0$ | [73]     |
| spin-orbit coupling             | $\Delta = 0.131$ eV | [19]     |
| valence band parameters         | $\gamma_1 = 1.76$ | [19, 20] |
|                                | $\gamma_2 = 0.7532$ | [19, 20] |
|                                | $\gamma_3 = -0.3668$ | [19, 20] |
|                                | $\gamma_4 = -0.020$ | [19, 20] |
|                                | $\eta_1 = 0.0037$ | [19, 20] |
|                                | $\eta_2 = -0.0037$ | [19, 20] |
|                                | $\eta_3 = 0.0037$ | [19, 20] |
| lattice constant                | $a = 0.42696$ nm | [74]     |
| dielectric constants            | $\varepsilon_{s1} = 7.5$ | [75]     |
|                                | $\varepsilon_{s1} = \varepsilon_{s2} = 7.11$ | [75]     |
|                                | $\varepsilon_{s2} = 6.46$ | [75]     |
| energy of $\Gamma_4$-LO phonons| $\hbar\omega_{\text{LO1}} = 18.7$ meV | [48]     |
|                                | $\hbar\omega_{\text{LO2}} = 87$ meV | [48]     |
| exchange energy                 | $J_0 = 0.792$ eV | [28]     |
| central-cell correction         | $V_0 = 0.539$ eV | [28]     |

here depend on the wave vector $\mathbf{K}$, which enters the Hamiltonian (96). The index $\nu$ is a number to distinguish the different exciton states.

Inserting the ansatz (7) in the Schrödinger equation $H \Psi = E \Psi$ and multiplying from the left with another basis state $\{|\Psi\rangle\}$, we obtain a matrix representation of the Schrödinger equation of the form

$$Dc = EMCc.$$  \hfill (8)

The vector $c$ contains the coefficients of the ansatz (7). All matrix elements of $H_1$ and $H_2$, which enter the symmetric matrices $D$ and $M$ are given in the Supplemental Material [79]. The matrix elements of $H_0$ are already given in the Appendices of Refs. [20] and [28]. The generalized eigenvalue problem (8) can finally be solved using an appropriate LAPACK routine [80].

All material parameters used in the calculations are listed in Table I.

**III. OSCILLATOR STRENGTHS**

Having solved the generalized eigenvalue problem (8), one can directly calculate the relative oscillator strengths for the transitions from the ground state of the solid to the exciton states. In this section we will derive the according formulas for dipole and quadrupole transitions. The oscillator strength is strongly connected to the interaction between the excitons and photons. According to Ref. [7] the probability per unit time for a transition from the ground state $\Phi_0$ of the semiconductor to an exciton state $\Psi_{\nu, \mathbf{K}}$ is proportional to $|M|^2$ with

$$M = \int \Psi_{\nu, \mathbf{K}}^\dagger \left[ \frac{-e}{m_0} A_0 (\mathbf{K}, \xi) \sum_{l=1}^N e^{i\mathbf{k}_l \cdot \mathbf{r}_l} \right] \Phi_0 d\mathbf{r}_1 \cdots d\mathbf{r}_N.$$  \hfill (9)

Here $A_0 (\mathbf{K}, \xi)$ denotes the amplitude of the vector potential of the radiation field with the wave vector $\mathbf{k}$ and the polarization $\xi$. $N$ denotes the number of electrons with the coordinates $\mathbf{r}_l$.

Within the scope of the simple band model the wave function of an exciton is given by

$$\Psi_{\nu, \mathbf{K}} = \sum_q f_{\nu, \mathbf{K}} (q) \Phi_{\nu, \mathbf{K}}^\dagger (q - \mathbf{K}, q + \alpha \mathbf{K}),$$  \hfill (10)

where $\tau$ and $-\sigma$ denote the spins of the electron and the hole, respectively. As we will discuss below, the spin orbit splitting in Cu$_2$O does not have a perceptible effect on the oscillator strength. Hence, we will keep the derivation splitting in Cu$_2$O.

The envelope function $f_{\nu, \mathbf{K}} (q)$ in Eq. (10) is the Fourier transform of the solution $F_{\nu, \mathbf{K}} (\beta)$ of the Wannier equation [5, 7],

$$f_{\nu, \mathbf{K}} (q) = \frac{1}{\sqrt{N}} \sum_{\beta} F_{\nu, \mathbf{K}} (\beta) e^{-i\mathbf{q} \beta},$$  \hfill (11)

with $\nu$ denoting the different exciton states. Note that the coordinate $\beta$ is a lattice vector which in general takes only discrete values [48]. This coordinate is identical to the relative coordinate $\mathbf{r}$ used in Sec. II in the continuum approximation.

The wave function (10) contains a Slater determinant of Bloch functions with one electron being in a Bloch state of the conduction band and $N-1$ electrons in Bloch states of the valence bands:

$$\Phi_{\nu, \mathbf{K}} = \mathcal{A} \psi_{v_1, \mathbf{k}_1, \alpha_1} \cdots \psi_{v_N, \mathbf{k}_N, \beta}.$$  \hfill (12)

Here $\mathcal{A}$ denotes the antisymmetrization operator. With the ground state of the semiconductor

$$\Phi_0 = \mathcal{A} \psi_{v_1, \mathbf{k}_1, \alpha_1} \cdots \psi_{v_N, \mathbf{k}_N, \beta},$$  \hfill (13)

we can express the exciton state (10) as

$$\langle \Psi_{\nu, \mathbf{K}} \rangle = \sum_q f_{\nu, \mathbf{K}} (q) c_{v_1, \mathbf{k}_1, \alpha_1} \cdots c_{v_N, \mathbf{k}_N, \beta} \Phi_0 \rangle,$$  \hfill (14)

using creation and annihilation operators. The operator in square brackets in Eq. (9) can be written in second quantization as

$$\sum_{nn'} \sum_{kk'} \sum_{\sigma\sigma'} \frac{-e}{m_0} A_0 (\mathbf{K}, \xi) \sum_{l=1}^N e^{i\mathbf{k}_l \cdot \mathbf{r}_l} \langle \psi_{n', \mathbf{k}', \sigma'} | e^{i\mathbf{p} \cdot \mathbf{r}} \psi_{n, \mathbf{k}, \sigma} \rangle c_{n', \mathbf{k}', \sigma'} \dagger c_{n, \mathbf{k}, \sigma}. \hfill (15)
After some transformations, using \(A_0(\kappa, \xi) = A_0(\kappa, \xi) \hat{e}_{\xi \kappa}\) and neglecting Umklapp processes, we arrive at

\[
M = -\frac{e\hbar}{m_0} A_0(\kappa, \xi) N \delta_{\tau \sigma} \delta_{\kappa, \mathbf{K}} \sum_q f_{vcv}(q) \int_{\text{WSC}} dr \times \langle u_{m0}(r) | e_{\xi \kappa} \left[ (q - \gamma \mathbf{K}) u_{v(q-\gamma \mathbf{K})}(r) - i \nabla u_{v(q-\gamma \mathbf{K})}(r) \right] \rangle \]

with an integral over the Wigner-Seitz cell (WSC) \([81]\).

To obtain expressions for the dipole and quadrupole with the components \(m = \sqrt{M k_1 N} \ldots \). We obtain \(\sum u_\gamma = \sum u_\gamma \approx U_m K_\mathbf{m} = \sum \sqrt{M k_1 N} \ldots \). We can then consider the coupling \(\sum u_\gamma = \sum U_m K_\mathbf{m}\) transforming according to \(\Gamma_\gamma^+\). With the basis functions \(|X\rangle, |Y\rangle, |Z\rangle\) of \(\Gamma_\gamma^+\) and the basis functions \(|\tilde{X}\rangle = |YZ\rangle, |\tilde{Y}\rangle = |ZX\rangle, \) and \(|\tilde{Z}\rangle = |XY\rangle\) of \(\Gamma_\gamma^+\), we see that, e.g., the \(\Gamma_\gamma^+\) like part of the products \(|X\rangle |Y\rangle_2\) and \(|Y\rangle_1 |X\rangle_2\) transforms as \(|\tilde{Z}\rangle / \sqrt{2}\). The other expressions are obtained via cyclic permutation \([84]\).

Hence, the expression requires that the operator has the symmetry \(\Gamma_\gamma^+\) \([83]\). We can then consider the coupling coefficients for the case \(\Gamma_\gamma^+ \otimes \Gamma_\gamma^+ \rightarrow \Gamma_\gamma^+\). With the basis functions \(|X\rangle, |Y\rangle, |Z\rangle\) of \(\Gamma_\gamma^+\) and the basis functions \(|\tilde{X}\rangle = |YZ\rangle, |\tilde{Y}\rangle = |ZX\rangle, \) and \(|\tilde{Z}\rangle = |XY\rangle\) of \(\Gamma_\gamma^+\), we see that, e.g., the \(\Gamma_\gamma^+\) like part of the products \(|X\rangle |Y\rangle_2\) and \(|Y\rangle_1 |X\rangle_2\) transforms as \(|\tilde{Z}\rangle / \sqrt{2}\). The other expressions are obtained via cyclic permutation \([84]\).

In Sec. II we have introduced the quasi spin \(I = 1\). If we compare the states \(|I, \mathbf{M}_I\rangle\) with the three functions \(|u_{\gamma 0}^{\pm 0}\rangle, |u_{\gamma 0}^{\pm \alpha}\rangle, \) and \(|u_{\gamma 0}^{\pm \gamma}\rangle\) transforming according to \(\Gamma_\gamma^+\), it is \([68]\)

\[
|1, +1\rangle_I = - (|u_{\gamma 0}^{\pm 0}\rangle + i |u_{\gamma 0}^{\pm \gamma}\rangle) / \sqrt{2}, \quad |1, 0\rangle_I = |u_{\gamma 0}^{\pm 0}\rangle, \quad |1, -1\rangle_I = (|u_{\gamma 0}^{\pm 0}\rangle - i |u_{\gamma 0}^{\pm \gamma}\rangle) / \sqrt{2}.
\]

In the envelope function of the exciton the angular dependence is given by the spherical harmonics \(Y_{LM}\). We know that in Eq. (21) the gradient of the envelope function at \(r = 0\) is non-zero only if \(L = 1\) holds. Furthermore, the envelope function itself vanishes at \(r = 0\) if \(L \neq 0\) holds.

Let us assume that the light is polarized in \(x\)-direction, i.e., \(e_{\xi \kappa} = e_x\). Let us furthermore write the function \(|\psi_{\nu \mathbf{K}}\rangle\) of the exciton in the form of Eq. (7), where the spins, the envelope with the angular momentum \(L\) and the function of the hole with the quasi-spin \(I\) enter. The dipole-term in Eq. (21) is then proportional to

\[
|1, +1\rangle_I = - (|u_{\gamma 0}^{\pm 0}\rangle + i |u_{\gamma 0}^{\pm \gamma}\rangle) / \sqrt{2}, \quad |1, 0\rangle_I = |u_{\gamma 0}^{\pm 0}\rangle, \quad |1, -1\rangle_I = (|u_{\gamma 0}^{\pm 0}\rangle - i |u_{\gamma 0}^{\pm \gamma}\rangle) / \sqrt{2}.
\]
\[
\lim_{r \to 0} \langle S = 0, M_S = 0 \mid \left( \langle u_{x0}^y \rangle \frac{\partial}{\partial y} + \langle u_{z0}^z \rangle \frac{\partial}{\partial z} \right) \mid \Psi_{\nu K} \rangle
\]

\[
= \lim_{r \to 0} \langle S = 0, M_S = 0 \mid \frac{\partial}{\partial r} \left( \langle I = 1, M_I = 0 \mid (-i) \sqrt{\frac{3}{8\pi}} (\langle L = 1, M_L = 1 \rangle + \langle L = 1, M_L = -1 \rangle) \right) \mid \Psi_{\nu K} \rangle
\]

\[
= -i \sqrt{\frac{3}{4\pi}} \lim_{r \to 0} \frac{\partial}{\partial r} \left( D \langle 2, 1 \mid + D \langle 2, -1 \mid \mid \Psi_{\nu K} \rangle \right)
\]

(26)

Here the state \( |F_t, M_{F_t}\rangle_D \) for the dipole transition \( (D) \) is a short notation for

\[
|S_e, S_h \rangle S, I; I + S, L; F_t, M_{F_t} \rangle = |(1/2, 1/2) 0, 1, 1, 1; F_t, M_{F_t} \rangle,
\]

in which the coupling scheme of the spins and angular momenta is different from the one of Eq. (7b) due to the requirement that \( S \) must be a good quantum number: \( S_e + S_h = S \to (I + S) + L = F_t \). As the quantization axis we choose the \( z \)-axis, which is parallel to one of the principal axes of the crystal lattice. In an analogous way, the quadrupole term can be written as

\[
\lim_{r \to 0} \langle S = 0, M_S = 0 \mid (\langle u_{x0}^y \rangle K_y + \langle u_{z0}^z \rangle K_z) \mid \Psi_{\nu K} \rangle
\]

\[
= \lim_{r \to 0} \langle S = 0, M_S = 0 \mid \frac{1}{\sqrt{4\pi}} \langle L = 0, M_L = 0 \rangle \langle I = 1, M_I = 0 \mid K_y \rangle
\]

\[
- i \sqrt{\frac{3}{2}} \langle I = 1, M_I = -1 \rangle F_{\nu K} \rangle 
\]

\[
= \lim_{r \to 0} \frac{1}{\sqrt{4\pi}} \left( Q \langle 1, 0 \mid K_y - i \sqrt{\frac{3}{2}} (Q \langle 1, -1 \rangle + Q \langle 1, 1 \rangle) \right) \mid \Psi_{\nu K} \rangle
\]

(28)

with the state \( |F_t, M_{F_t}\rangle_Q \) for the quadrupole transition \( (Q) \) being a short notation for

\[
|S_e, S_h \rangle S, I; I + S, L; F_t, M_{F_t} \rangle = |(1/2, 1/2) 0, 1, 1, 0; F_t, M_{F_t} \rangle.
\]

(29)

Note that this state is similar to the one of Eq. (27) but only \( L \) is set to zero. We finally arrive at the following expression for the relative oscillator strength:

\[
J^{rel}_{\xi K} = \lim_{r \to 0} \left[ -i \left( \hat{M}_v^* + \hat{M}_c^* \right) \frac{\partial}{\partial r} \langle T^D_{\xi K} \mid \Psi_{\nu K} \rangle + \left( -\gamma \hat{M}_v^* + \alpha \hat{M}_c^* \right) \frac{K}{\sqrt{6}} \langle T^Q_{\xi K} \mid \Psi_{\nu K} \rangle \right]^2.
\]

(30)

For the dipole term in Eq. (30) the two states \( |T^D_{\xi K} \rangle \) are given by

\[
|T^D_{\xi K} \rangle = \sum_{\ell = 1}^{3} \hat{e}_{\xi K \ell} |\pi^D_{\ell} \rangle, \quad \xi = 1, 2
\]

(31)

with the three components of the polarization vector \( \hat{e}_{\xi K} \) and three states which transform according to \( \Gamma^+_{1} \) [21]:

\[
|\pi^D_{x} \rangle = \frac{i}{\sqrt{2}} \left[ |2, -1 \rangle_D + |2, 1 \rangle_D \right],
\]

(32a)

\[
|\pi^D_{y} \rangle = \frac{1}{\sqrt{2}} \left[ |2, -1 \rangle_D - |2, 1 \rangle_D \right],
\]

(32b)

\[
|\pi^D_{z} \rangle = \frac{1}{\sqrt{2}} \left[ |2, -2 \rangle_D - |2, 2 \rangle_D \right].
\]

(32c)

The state \( |T^Q_{\xi K} \rangle \) in the quadrupole term of Eq. (30) reads

\[
|T^Q_{\xi K} \rangle = \sum_{\ell = 1}^{3} \hat{e}_{\xi K \ell} |\pi^Q_{\ell} \rangle, \quad \xi = 1, 2
\]

(33)

with the three states which transform according to \( \Gamma^+_{3} \) [21]:

\[
|\pi^Q_{x} \rangle = \hat{K}_y |1, 0 \rangle_Q,
\]

(34a)

\[
+ \hat{K}_z \frac{i}{\sqrt{2}} \left[ |1, -1 \rangle_Q + |1, 1 \rangle_Q \right],
\]

\[
|\pi^Q_{y} \rangle = \hat{K}_x |1, 0 \rangle_Q
\]
If we now set \( K = 0 \) in Eq. (30), we see that we have derived the expression for the relative oscillator strength, which has already been used in Refs. [20, 21].

We can finally make an assumption as regards the size of the parameters \( M^n \) and \( M_L \). Since in \( \text{Cu}_{2}O \) the uppermost valence bands as well as the lowermost conduction band have positive parity, we see from Eq. (22) that only bands with negative parity will contribute to the sums in Eq. (19). In \( \text{Cu}_{2}O \) there are only two bands of negative parity, which lie 449 meV above the lowest conduction band and 5.6 eV below the highest conduction band [48]. Hence, as regards the denominators of negative parity, which lie 449 meV above the lowest conduction band and 5.6 eV below the highest conduction band, the spin-orbit coupling is negligible small in comparison to the energy difference between these bands and the bands of negative parity. Thus, the spin-orbit coupling does not have a sizeable effect on the oscillator strength. Furthermore, the denominator \( \sum \) of Eq. (35), where the sum of dipole and quadrupole matrix element is squared. In contrast to the hydrogen-like model, inversion symmetry is broken when considering the more precise model of \( \text{Cu}_{2}O \). Hence, the exciton states are mixed-parity states and, thus, an interference term occurs. Especially for the 1S ortho exciton both terms in Eq. (35) are of the same magnitude. The dipole term becomes equally large as the quadrupole term for \( K \neq 0 \) due to a \( K \) dependent admixture of \( P \) excitons to the 1S ortho exciton. Even though the parameter \( \alpha \) occurs in Eq. (35), the whole expression is independent of its value. This is related to the fact that the wave function of the exciton also depends on \( \alpha \).

In this paper we choose \( \mathbf{K} \) to be oriented in [001], [110], or [111] direction and, as in Ref. [21], we also rotate the coordinate system to make the \( z \) axis of the new coordinate system coincide with the direction of \( \mathbf{K} \). The formulas for the oscillator strengths for the three orientations of \( \mathbf{K} \) [85] can be derived by analogy with Ref. [21]. The consideration of these three directions of high symmetry is sufficient to determine those parameters in Sec. VI, which are needed in the \( 5 \times 5 \) matrix model to calculate the polariton dispersion for any other direction of \( \mathbf{K} \).

## IV. EXCITON-POLARITONS

In this section we recapitulate the quantum mechanical theory of exciton-polaritons in Sec. IV A and discuss the rotating-wave approximation in Sec. IV B.

A. Polariton transformation

The quantum mechanical theory of polaritons was first developed by Hopfield, Fano, and Agranovich [56–58]. According to Refs. [56, 86, 87], the second-quantized Hamiltonian for the interaction of excitons and photons

\[
H = \sum_{\mathbf{K}, \nu} \left[ \hbar \omega_{\mathbf{K}} \left( a_{\mathbf{K}}^\dagger a_{\mathbf{K}} + \frac{1}{2} \right) + \sum_{\nu} E_{\nu\mathbf{K}} \left( B_{\nu\mathbf{K}}^\dagger B_{\nu\mathbf{K}} + \frac{1}{2} \right) + i \sum_{\nu} C_{\mathbf{K}} \left( a_{\mathbf{K}}^\dagger + a_{-\mathbf{K}} \right) \left( B_{\nu\mathbf{K}} - B_{\nu-\mathbf{K}}^\dagger \right) \right.
\]

\[
+ \left. \sum_{\nu} D_{\mathbf{K}} \left( a_{\mathbf{K}}^\dagger \right) \left( a_{\mathbf{K}} + a_{-\mathbf{K}} \right) \right] (36)
\]

can be derived either from a microscopic model of excitons with the Hamiltonian describing the interaction between radiation and matter or from the equation of motion for the exciton polarization. In the Hamiltonian (36) the operators \( B_{\nu\mathbf{K}}^\dagger \) and \( B_{\nu\mathbf{K}} \) create and annihilate an exciton with energy \( E_{\nu\mathbf{K}} \), respectively, and obey Bose commutation rules in the following. Likewise, the operator \( a_{\mathbf{K}}^\dagger \left( a_{\mathbf{K}} \right) \) creates (annihilates) a photon with polarization \( \xi \) and energy \( \hbar \omega_{\mathbf{K}} = \hbar c K / \sqrt{\varepsilon b} \). The coupling coefficients in the exciton-photon and the photon-photon interaction terms of Eq. (36) are given by

\[
C_{\mathbf{K}} = \left[ \frac{\kappa_{SI} \beta_{\mathbf{K}}^2 E_{\mathbf{K}}^3}{\varepsilon b_2 \hbar \omega_{\mathbf{K}}} \right]^{\frac{1}{2}} (37)
\]

with \( \kappa_{SI} = 1/4\pi \varepsilon_0 \) and

\[
D_{\mathbf{K}} = C_{\mathbf{K}}^2 / E_{\mathbf{K}}. \quad (38)
\]

The polarizability \( \beta_{\mathbf{K}} \) is proportional to the oscillator strength of the exciton state. With our definition of the oscillator strength \( f_{\nu\mathbf{K}} \) (cf. Refs. [65, 66, 70]), this proportionality is given by [86, 88, 89]

\[
\beta_{\mathbf{K}} = \varepsilon_0 \varepsilon_0 b_2 f_{\nu\mathbf{K}}. \quad (39)
\]
The Hamiltonian (36) can be diagonalized by the Hopfield transformation [56, 57], which is similar to the Bogolyubov’s uv transformation [90, 91]: New creation and annihilation operators $p_{\mu \xi}^\dagger$ and $p_{\mu \xi}$ are introduced via

$$a_{\xi \mathbf{K}} = \sum \mu \left[ u_{\mu \xi \mathbf{K}} p_{\mu \xi} + v_{\mu \xi \mathbf{K}}^* p_{\mu \xi \mathbf{K}}^\dagger \right],$$  

$$B_{\nu \mathbf{K}} = \sum \mu \left[ u_{\mu \nu \mathbf{K}} p_{\mu \xi \mathbf{K}} + v_{\mu \nu \mathbf{K}}^* p_{\mu \xi \mathbf{K}}^\dagger \right],$$

(40a, 40b)

to obtain the polariton Hamiltonian

$$H = \sum_{\mu \xi \mathbf{K}} E_{\mu \xi \mathbf{K}} p_{\mu \xi \mathbf{K}}^\dagger p_{\mu \xi \mathbf{K}} + \text{const.} \quad \Rightarrow$$

(41)

with $\mu$ and $E_{\mu \xi \mathbf{K}}$ denoting the polariton branches and the polariton energies, respectively. The new operators must obey Bose commutation relations and the Hamiltonian shall be diagonal, i.e.,

$$[p_{\mu \xi \mathbf{K}}, H] = E_{\mu \xi \mathbf{K}} p_{\mu \xi \mathbf{K}}$$

(42)

must hold. This provides the following conditional equation for the polariton energies [58, 87, 92]:

$$\frac{\hbar^2 c^2 K^2}{E_{\mu \xi \mathbf{K}}} = \varepsilon_b + \sum \nu \frac{4\pi \kappa_S \beta_{\mu \xi \nu \mathbf{K}}}{1 - (E_{\mu \xi \mathbf{K}} / E_{\nu \mathbf{K}})^2}.$$  

(43)

Using the phase convention of Hopfield [56], the solitons for the coefficients $u$ and $v$ of the polariton transformation can be obtained (see Ref. [87]). The polariton operators can also be expressed in terms of exciton and photon operators:

$$p_{\mu \xi \mathbf{K}} = w_{\mu \xi \mathbf{K}}^{(1)} a_{\xi \mathbf{K}} + w_{\mu \xi \mathbf{K}}^{(2)*} a_{\xi \mathbf{K}}^\dagger$$

$$+ \sum \nu \left[ z_{\mu \nu \mathbf{K}}^{(1)} B_{\nu \mathbf{K}} + z_{\mu \nu \mathbf{K}}^{(2)*} B_{\nu \mathbf{K}}^\dagger \right].$$

(44)

Since all creation and annihilation operators of the three (quasi-) particles obey Bose commutation relations, we can determine the coefficients $w$ and $z$ by evaluating

$$[p_{\mu \xi \mathbf{K}}, a_{\xi \mathbf{K}}^\dagger] = + w_{\mu \xi \mathbf{K}}^{(1)} a_{\xi \mathbf{K}}^\dagger,$$

$$[p_{\mu \xi \mathbf{K}}, a_{\xi \mathbf{K}}^\dagger] = - w_{\mu \xi \mathbf{K}}^{(2)*} a_{\xi \mathbf{K}}^\dagger$$

(45a, 45b)

and

$$[p_{\mu \xi \mathbf{K}}, B_{\nu \mathbf{K}}^\dagger] = + z_{\mu \nu \mathbf{K}}^{(1)} a_{\nu \mathbf{K}}^\dagger,$$

$$[p_{\mu \xi \mathbf{K}}, B_{\nu \mathbf{K}}^\dagger] = - z_{\mu \nu \mathbf{K}}^{(2)*} a_{\nu \mathbf{K}}^\dagger.$$

(46a, 46b)

The coefficients $w_{\mu \xi \mathbf{K}}^{(i)}$ or the sum

$$W_{\mu \xi \mathbf{K}} = \sum_{i=1}^{2} |w_{\mu \xi \mathbf{K}}^{(i)}|^2$$

(47)

then allow one to determine whether the polariton is more photon-like ($W_{\mu \xi \mathbf{K}} \rightarrow 1$) or more exciton-like ($W_{\mu \xi \mathbf{K}} \rightarrow 0$).

B. Rotating-wave approximation

In the literature polaritons are often treated within the so-called rotating-wave approximation [86]. In this case the term with the coefficient $D$ and the anti-resonant terms of the form $a B$ and $a^\dagger B^\dagger$ are neglected in the Hamiltonian (36). The resulting Hamiltonian

$$H = \sum_{\xi \mathbf{K}} \left[ \hbar \omega_{\xi \mathbf{K}} a_{\xi \mathbf{K}}^\dagger a_{\xi \mathbf{K}} + \sum \nu E_{\nu \mathbf{K}} B_{\nu \mathbf{K}}^\dagger B_{\nu \mathbf{K}} \right. $$

$$+ \left. \sum \nu C_{\xi \nu \mathbf{K}} (a_{\xi \mathbf{K}}^\dagger B_{\nu \mathbf{K}} + a_{\xi \mathbf{K}} B_{\nu \mathbf{K}}^\dagger) \right]$$

(48)

is then called the Jaynes-Cummings Hamiltonian [93], where the vacuum energy of the photons has also been neglected [86] and where the operators $a_{\xi \mathbf{K}}$ have been replaced with $i a_{\xi \mathbf{K}}$. Note that this replacement does not change the physics of the problem since it only adds global phases to the occupation-number states. The occupation-number operator and the commutation relations remain unchanged.

The coefficient $C_{\xi \nu \mathbf{K}}$ can be written as [86, 89]

$$C_{\xi \nu \mathbf{K}} = \left[ \varepsilon_{\xi \nu \mathbf{K}} E_{\nu \mathbf{K}}^3 / \varepsilon_{\xi \nu \mathbf{K}} \right]^{1/2} \approx \frac{1}{2} \left( K_0 / K \right)^{3/2} \hbar \Omega_R$$

(49)

with the wave vector at the exciton-photon resonance $K_0 = E_{\nu \mathbf{K}_0} / \sqrt{\varepsilon_{\xi \nu \mathbf{K}} / \varepsilon_{\xi \nu \mathbf{K}}}$ and the Rabi frequency

$$\Omega_R = E_{\nu \mathbf{K}_0} / \sqrt{4\pi \kappa_S \beta_{\xi \nu \mathbf{K}}}. \quad \Rightarrow$$

(50)

The rotating wave approximation is generally valid if $\hbar \Omega_R \ll E_{\nu \mathbf{K}_0}$ holds. This is, e.g., the case for inorganic semiconductors and especially for Cu$_2$O [89].

Close to the resonance ($K \approx K_0$) one can assume $C_{\xi \nu \mathbf{K}} \approx \hbar \Omega_R / 2$. Note that for $K \to 0$ the coupling constant (49) diverges, which is a manifestation of the infrared catastrophe in quantum electrodynamics [86]. Hence, the simplifications made above are valid only in the vicinity of the exciton-photon resonance. Otherwise, the full Hamiltonian (36) has to be diagonalized.

In the rotating-wave approximation the polariton transformation is more simple as there is no interaction between states with different values of $\mathbf{K}$. Using the ansatz

$$p_{\mu \xi \mathbf{K}} = w_{\mu \xi \mathbf{K}} a_{\xi \mathbf{K}} + \sum_{i} z_{\mu \xi \nu \mathbf{K}} B_{\nu \mathbf{K}}$$

(51)

the Bose commutation relations of the creation and annihilation operators, and the condition (42) for the polariton operator, one ends up with the eigenvalue problem

$$P_{\xi(\nu) \mathbf{K}} z_{\xi(\nu) \mathbf{K}} = E_{\mu \xi \mathbf{K}} z_{\mu \xi(\nu) \mathbf{K}}$$

(52)

with
Knowing the energies \( E_{\nu K} \) and the Rabi frequencies \( \Omega_{R, \nu} \) of the exciton states, one can directly obtain the corresponding polariton energies by determining the eigenvalues of Eq. (52).

Finally, as the polariton is a mixed state of a photon and excitons, one can again determine the photon-like part

\[
W_{\mu \xi K} = |w_{\mu \xi K}|^2 \tag{54}
\]

of the polariton or the contribution

\[
Z_{\mu \nu K} = |z_{\mu \nu K}|^2 \tag{55}
\]

of the exciton with the energy \( E_{\nu K} \) to the polariton.

C. Nonanalytic exchange interaction

There is another interaction affecting the exciton states: the nonanalytic (NA) exchange interaction. It is well known that the splitting caused by \( H_{\text{exch}}^{\text{NA}} \) is identical to the longitudinal-transverse splitting (LT-splitting) when treating polaritons [94]. Hence, it is indispensable to include the nonanalytic exchange interaction in the theory to obtain a correct treatment of the complete problem.

In this section we will derive an expression for the nonanalytic exchange interaction. We start with the formula of Ref. [25] for the nonanalytic exchange energy between two exciton states \( \Psi_{\nu K}^\sigma, \Psi_{\nu' K'}^\tau \) in second quantization

\[
H_{\text{exch}}^{\text{NA}} = \sum_{\nu \nu' K K'} \frac{m^2}{\varepsilon_0 b^2 V_{\text{uc}}} B_{\nu K} B_{\nu' K'} \tag{56}
\]

with the volume \( V_{\text{uc}} \) of one unit cell and

\[
m_{\nu K} = \delta_{\sigma \tau} \frac{e}{\sqrt{N}} \sum_q f_{\nu \nu} (q) \left\{ \frac{\hbar}{m_0} \frac{K \cdot p_{nc}}{E_v - E_c} + \hbar^2 \sum_{n \neq v, c} \left[ \frac{(q - \gamma K) \cdot p_{en}}{(E_v - E_n)} \frac{(q + \alpha K) \cdot p_{nc}}{(E_c - E_n)} \right. \\
+ \left. \frac{(q + \alpha K) \cdot p_{en}}{(E_c - E_v)} \frac{(q - \gamma K) \cdot p_{nc}}{(E_v - E_n)} \right] \right\}. \tag{57}
\]

Here \( m_{\nu K} \) is a short notation for the function \( m_{\nu K} (K, 0) \) of Ref. [25]. For the definitions of \( \Psi_{\nu K}^{\sigma \tau}, f_{\nu \nu} (q) \), and \( p_{mn} \) see Sec. III. The exchange energy includes the term \( \delta_{\sigma \tau} \delta_{\nu \nu'} \). Introducing the total spin \( S = S_e + S_h = \tau - \sigma \) of electron and hole, this term can be written for singlet and triplet states as \( 2 \delta_{S,0} \) [82].

Using Eq. (20) and rearranging the different terms in

\[
m_{\nu K} = \frac{\hbar^2}{m_0} \frac{K}{(E_c - E_v)} \times \lim_{r \to 0} \left[ \left( \hat{N}_v + \hat{N}_c \right) \cdot (-i \nabla_r F_{\nu \nu} (r)) + \left( -\gamma \hat{N}_v + \alpha \hat{N}_c \right) \cdot (F_{\nu \nu} (r) K) \right]. \tag{58}
\]
with the matrices
\[
\begin{align*}
\hat{N}_v &= \langle u_{v,0} | p M_v (\hat{K} \cdot p) | u_{v,0} \rangle, \\
\hat{N}_c &= \langle u_{v,0} | (\hat{K} \cdot p) M_c p | u_{v,0} \rangle, 
\end{align*}
\] (59a, 59b)
and \( \hat{K} = K/K \).

Due to the similarity between Eq. (21) and Eq. (58), we can perform the same calculation as in Sec. III to obtain
\[
m_{\nu K} \sim K \lim_{r \to 0} \left[ -i \left( \hat{N}_v + \hat{N}_c \right) \frac{\partial}{\partial r} \langle L^D_K | \Psi_{\nu K} \rangle + \frac{\alpha K}{\sqrt{6}} \langle L^Q_K | \Psi_{\nu K} \rangle \right] 
\] (60)
with the states
\[
| L^D_K \rangle = \sum_{i=1}^3 \tilde{K}_i | \pi_i^D \rangle, \quad | L^Q_K \rangle = \sum_{i=1}^3 \tilde{K}_i | \pi_i^Q \rangle, 
\] (61)
where we have introduced \( \tilde{K} = K/K \).

We can see from Eq. (56) that there is no interaction between states with different values of \( K \), which is the same case as for the Hamiltonian (48) of the polariton interaction in the rotating-wave approximation. Knowing the exciton energies \( E_{\nu K} \) and the corresponding wave functions \( | \Psi_{\nu K} \rangle \), we can simultaneously diagonalize the polariton Hamiltonian and the NA-exchange Hamiltonian by solving the eigenvalue problem
\[
(\mathbf{P}_{\xi \nu} K + \mathbf{N}_{\nu K}) z_{\mu \xi \nu} K = E_{\mu \xi} K z_{\mu \xi \nu} K 
\] (63)
with the matrix
\[
N_{\nu K} = \frac{\zeta}{\varepsilon_{b2} K^2} \begin{pmatrix} 0 & 0 & \cdots & 0 & \cdots \\ 0 & m_{\nu_1}^K m_{\nu_2} K & \cdots & m_{\nu_1}^K m_{\nu_2} K & \cdots \\ 0 & m_{\nu_1}^K m_{\nu_2} K & \cdots & \vdots & \cdots \\ \vdots & \vdots & \ddots & \ddots & \cdots \\ 0 & m_{\nu_1}^K m_{\nu_2} K & \cdots & m_{\nu_1}^K m_{\nu_2} K & \cdots \end{pmatrix}, 
\] (64)
and the matrix \( \mathbf{P}_{\xi \nu} K \) and vector \( z_{\mu \xi \nu} K \) defined in Eq. (53). The constant parameter \( \zeta \) can be determined by the fact that the splitting caused by \( H_{\text{exch}}^{\text{NA}} \) is identical to the LT-splitting.

### D. Observability of polariton effects

We now shortly discuss the criteria for the observability of polariton effects, which were derived by Tait in Ref. [95]. To obtain these criteria, he included a damping term \( \Gamma \) in the model of polaritons since damping is always present in the solid due to the interaction between excitons and phonons or the leakage of photons out of the solid [89]:
\[
\frac{\hbar^2 c^2 K^2}{E_{\nu K}^2} = \varepsilon_{b2} + \sum_{\nu} \frac{4 \pi \kappa_{\text{SI}} \beta_{\nu K} E_{\nu K}^2}{E_{\nu K} - E_{\nu \xi K} - i \Gamma_{\nu \xi K}}. 
\] (65)
This equation can either be solved for a fixed wave vector \( K \) or for a fixed frequency \( \omega = E_{\nu K}/\hbar \).

The first case corresponds to nonlinear optical experiments like, e.g., two-photon absorption. For this case a criterion of temporal coherence between the photon and the exciton can be derived [86, 95]. As long as
\[
\hbar \Gamma < \hbar \sqrt{\frac{\kappa_{\text{SI}} \pi}{\varepsilon_{b2}} \beta_{\nu K} E_{\nu K}^2} = \frac{\hbar}{2} \Omega_R 
\] (66)
holds, where \( \hbar \Gamma \) is the broadening of the linewidth due to damping, the polariton splitting is observable. This criterion can be interpreted in terms of Rabi oscillations, i.e., polariton effects are observable if a coherent energy transfer between an exciton and a photon is possible at least once [89]. Spatial coherence is already provided here by keeping \( K \) fixed [86].

The second case corresponds to reflectivity or absorption experiments. Here the coupling between the photon and the exciton must remain coherent during the propagation of the polariton through the solid in the presence of damping [86]. The criterion of spatial coherence reads [86, 89, 96]
\[
\hbar \Gamma < \sqrt{\frac{16 \pi}{M c^2} \kappa_{\text{SI}} E_{\nu K}^2 \beta_{\nu K} E_{\nu K}^2} = \frac{\hbar}{2} \Omega_R \sqrt{\frac{16 \varepsilon_{b2}}{M c^2} E_{\nu K}}, 
\] (67)
and it is generally more difficult to satisfy than Eq. (73) [89, 97] since \( \sqrt{16 \varepsilon_{b2} E_{\nu K}/M c^2} \ll 1 \) holds for an exciton mass \( M \) on the order of \( m_0 \) and an exciton.
energy on the order of a few eV. The criterion (74) is equivalent to \( l \gg \lambda \) with \( \lambda \) denoting the light wavelength and \( l = v_\parallel / \Gamma \) the mean free path of the exciton [86]. Hence, polariton effects can hardly be observed in semiconductors with very shallow excitons, e.g., in GaAs [86], when using linear optical techniques. Therefore, polariton effects are often investigated using nonlinear optical spectroscopic techniques due to the much less stringent criterion (73) [98, 99].

V. RESULTS INCLUDING VB STRUCTURE AND CENTRAL-CELL CORRECTIONS

In this section we will treat the exciton-polaritons with \( 2 \leq n \leq 4 \) in Cu\(_2\)O using the Hamiltonian which accounts for the valence band structure, the exchange interaction, the central-cell corrections, and the finite momentum \( \hbar K \) of the center of mass. For the three orientations \([001],[110]\), and [111] of \( K \) considered here the cubic symmetry \( O_h \) is reduced to \( C_{4v},C_{2v}, \) and \( C_{3v} \), respectively [83].

Excitons of even and odd parity behave differently in dependence on the momentum \( \hbar K \). On the one hand, states of the odd series having the symmetry \( \Gamma_4^\pm \) and a component with \( L = 1 \) show a finite dipole oscillator strength at \( K = 0 \). Since the angular momentum \( L \) is not a good quantum number in Cu\(_2\)O and \( P \)-like exciton states are admixed to other states of odd parity at \( K = 0 \) [28], \( P, F, H, \ldots \) states of symmetry \( \Gamma_6^\pm \) have a finite dipole oscillator strength. Hence, for these states already a splitting at \( K = 0 \) occurs due to the nonanalytic exchange interaction and the exciton-photon coupling, which affect the states of this symmetry.

On the other hand, the even exciton states do not have an oscillator strength at \( K = 0 \) and, therefore, no splitting occurs. For those even states which have the symmetry \( \Gamma_4^\pm \) and an \( L = 0 \)-component at \( K = 0 \) the coupling to light via a finite \( K \) vector is possible. Since \( S \)-like exciton states are admixed to other states of even parity at \( K = 0 \) [28], \( S, D, G, \ldots \) states of symmetry \( \Gamma_5^\pm \) obtain a finite quadrupole oscillator strength being proportional to \( \hbar^2 \).

At first, we will compare our numerical results for the relative oscillator strengths of the 1S excitons with the absolute value from the experiment. This will allow us to calculate absolute oscillator strengths for all exciton states. Having determined the correct size of the nonanalytic exchange interaction for the \( n \) excitons and, hence, for all other exciton states, we can then investigate the dispersion of exciton-polaritons for the three orientations of \( K \) along the axes of high symmetry.

Since the formula derived in Sec. III allows us only to calculate relative oscillator strengths \( f_{\xi\nu}^{rel} \) but not absolute oscillator strengths for the different polarizations \( \xi \) and exciton states \( \nu \)\( K \), we determine the scaling factor \( \eta \) in

\[
f_{\xi\nu} = \eta f_{\xi\nu}^{rel}(\hbar K)
\]

by comparing the theoretical results for \( f_{\xi\nu}^{rel} \) for the 1S exciton state at the exciton-photon resonance \( K = K_0 \) with the experimentally obtained value of [65]

\[
f_{\xi 1S} = 3.6 \times 10^{-9}
\]

for \( \hbar K \approx 0 \). This yields \( \eta = 825.9 \). Knowing this scaling factor, we can give the absolute oscillator strengths of all exciton states in the following. With this value the oscillator strengths of, e.g., the \( n \)P excitons are given by

\[
f_{\xi nP} = 3.75 \times 10^{-4} \frac{n^2 - 1}{\eta^2}.
\]

For a complete description of the polariton problem, we also have to include the nonanalytic exchange interaction. The splitting caused by this interaction at \( K = 0 \) must exactly equal the LT-splitting due to the polariton transformation. As the rotating wave approximation does not hold for \( K = 0 \), we determine at first the polariton energies \( E_{\mu \xi} \) for the transverse exciton states via the conditional equation

\[
\frac{\hbar^2 c^2 K^2}{\epsilon_{\nu 2} E_{\mu \xi}^2} = 1 + \sum_\nu \frac{f_{\xi \nu} K^2}{1 - (E_{\mu \xi} / E_{\nu \xi})^2},
\]

which is obtained when using the complete exciton-photon interaction Hamiltonian [58, 87, 92]. The conditional equation (71) can be rewritten so that the polariton energies are the roots of the function

\[
F(E_{\mu \xi}) = 1 - \frac{\hbar^2 c^2 K^2}{\epsilon_{\nu 2} E_{\mu \xi}^2} + \sum_\nu \frac{f_{\xi \nu} K^2}{E_{\nu \xi}^2 - E_{\mu \xi}^2},
\]

where the photon energy \( \hbar \omega = \hbar K c / \sqrt{\epsilon_{\nu 2}} \) is used. Note that a root of \( F(E_{\mu \xi}) \) is always located between any neighboring pair of the exciton energies \( E_{\nu \xi} \), \( E_{\nu + 1 \xi} \) [101–103].

We calculate the effect of the nonanalytic exchange interaction on the longitudinal states for \( K \rightarrow 0 \) by diagonalizing the matrix \( N_{(\nu) \xi} \) given in Eq. (64). As the size of the nonanalytic exchange is a priori unknown, we have scaled the matrix \( N_{(\nu) \xi} \), which describes this interaction, with a parameter \( \zeta \). We vary the parameter \( \zeta \) in such a way that the energies of the longitudinal and transverse \( n \)P exciton states are identical. This yields \( \zeta = 213.5 \pm 2.0 \).

Of prime interest are now the polariton dispersions in the vicinity of the exciton phonon resonance. In this range of \( K \) the rotating wave approximation is valid and we solve the eigenvalue problem (63). Note that the errors arising due to the use of the rotating-wave approximation are smaller than the error due to uncertainties in \( \epsilon_{\nu 2} \).

The numerical result for \( K \parallel [001] \) is shown in Fig. 1. In the panels (a) we show the exciton spectrum in dependence on \( K = |K| \). Since the changes in the energy in dependence on \( K \) are in the order of tens of \( \mu \)eV, the exciton states appear as straight horizontal lines.
FIG. 1: (a) The exciton energies $E_{\nu K}$ in dependence on $K = |K|$ for $K \parallel [001]$. Due to the inclusion of the complete valence band structure and the central cell corrections, the spectrum is much more complicated than in the hydrogen-like case (cf. Refs. [26, 100]). The colorbar shows the oscillator strengths for $\xi = \sigma^\pm_p$ polarized light. For both polarizations the spectrum is identical. We denote from which states at $K = 0$ the exciton states originate (cf. Ref. [28]). For reasons of space we introduce the abbreviated notation $\Gamma_{ij}^\pm$ to replace $\Gamma_i^\pm, \Gamma_j^\pm$. The blue solid line gives the photon dispersion $\hbar \omega_{\xi K} = \hbar K_c/\sqrt{\varepsilon_b^2}$.

(b) Polariton dispersion obtained by solving the eigenvalue problem (63). Since the $P$ excitons have a large oscillator strength, large avoided crossings can be observed. They clearly shift the position of the smaller avoided crossings for the $2S, 3D$ and $4F$ excitons away from the position where the dispersion of light and the dispersion of these exciton states cross. (c) Photon-like part $W_{\mu \xi K}$ of the polariton states. In all cases the mixing of excitons and photons is strongest in the vicinity of an avoided crossing. Only for the $P$ excitons an admixture of photons far away from the resonance occurs.

The spectrum is much more complicated than when using the hydrogen-like model of excitons, which was done in Refs. [26, 100]. Even for vanishing momentum of the center of mass, the complete valence band structure already leads to a complicated fine structure splitting and to a mixing of the exciton states with even or odd parity. This explains the observability of $F$ excitons in absorption spectra due to the admixture of $P$ excitons [18, 20]. Due to the finite momentum of the center of mass $S$ and $D$ excitons also obtain a small oscillator strength.

We state in the panels (a) of Fig. 1 from which states at $K = 0$ the exciton states originate using the nomen-
clature \( nL_{y/z}(\Gamma^\pm_0) \) of Ref. [28] with the abbreviations \( y \) and \( g \) for yellow and green. In the case of the \( P \) and \( F \) excitons we do not give the symmetry of the complete exciton state but only the combined symmetry of envelope and hole [18].

In the panels (b) we show the polariton dispersion. Only for the \( P \) excitons, which have a comparatively large oscillator strength, significant deviations between the polariton spectrum and the exciton spectrum can be observed. The insets in the panels show that also for the other exciton states avoided crossings appear due to their finite oscillator strength. However, the panels (c), which show the photon-like part of the polariton states, indicate that the mixing between excitons and photons is small in these cases.

Due to the proximity of the \( nS \) and \( nP \) states as well as that of the \( nF \) states, avoided crossings are not well-separated. Hence, the multi-polariton concept has to be used and a single-polariton concept would lead to different results. The large avoided crossings of the \( P \) excitons clearly affect the other avoided crossings as they shift them away from the position where the dispersion of light and the dispersion of the other exciton states cross.

As regards the other orientations \( K \parallel [110] \) or \( K \parallel [111] \) of the momentum of center of mass the differences in the polariton spectra are slight. Since the \( K \)-dependent shift of the exciton energies is on the order of tens of \( \mu \)eV, the exciton energies are almost the same for the three orientations of \( K \) considered. The main difference between the spectra for \( K \parallel [001] \), \( K \parallel [110] \) or \( K \parallel [111] \) is the values of the oscillator strengths due to the different symmetry breaking. Therefore, also the polariton dispersions are different, however, mainly in the vicinity of the avoided crossings [104].

Note that we do not show the polariton dispersion for exciton states with \( n \geq 5 \) since the number of states with a finite oscillator strength increases rapidly. However, the oscillator strengths of the \( G, H, \ldots \) exciton states are very small so that polariton effects are likewise very small. Hence, we do not expect to observe considerably new effects for the exciton-polaritons with \( n \geq 5 \).

Using the criteria of Tait [95] for temporal and spatial coherence presented in Sec. IV D, we can estimate the possible observability of the polariton effects. If we set \( n = 2 \), the criterion of temporal coherence reads

\[
\hbar \Gamma < \frac{\hbar}{2} \Omega_R \approx 6.4 \text{ meV}.
\]

(73)

For the criterion of spatial coherence, we obtain

\[
\hbar \Gamma < \frac{\hbar}{2} \Omega_R \sqrt{\frac{16 \epsilon_{\text{k}2}}{M c^2} E_\nu K_0} \approx 0.11 \text{ meV}
\]

(74)

with the isotropic mass \( M \approx 1.57 m_0 \) of the exciton and \( m_0 \) denoting the free electron mass. In Ref. [13] the spectrum of the giant Rydberg excitons has been investigated in an absorption experiment and, thus, experimental values of the exciton linewidths \( \hbar \Gamma \) for the criterion of spatial coherence are known. For \( n = 2 \) the experimental line width is \( \hbar \Gamma = 1.58 \text{ meV} \) [13, 15], which is significantly larger than 0.11 meV. Therefore, we expect that the polariton effects for the yellow exciton states with \( n \geq 2 \) in \( \text{Cu}_2\text{O} \) may only be observed using nonlinear spectroscopy methods and high quality crystals [98, 99]. As regards the 1S ortho exciton state, the linewidth is small enough to clearly observe polariton effects [65, 66, 70].

VI. YELLOW 1S ORTHO EXCITON POLARITON

We now come to the yellow 1S ortho exciton polariton, for which a pronounced polariton effect has been proven in experiments. In Sec. VIA we will discuss at first the effect of finite momentum of the center of mass \( hK \neq 0 \) on the exciton spectrum. We will especially pay attention to the small quadrupole oscillator strength [65] and the \( K \)-dependent splitting of this state [20, 67–69]. In Sec. VIB we set up a \( 5 \times 5 \) matrix model, which allows us to calculate the anisotropic dispersion of the 1S ortho-exciton polariton for any direction of \( K \). We then present in Sec. VIC the polariton dispersion, determine the group velocity [66, 70] as well as the spectra for rotations about the \([110]\) and \([111]\) axis and compare them with experimental values [67, 68].

A. \( K \)-dependent splitting

As has already been stated in Sec. V, we have to consider the reduction of the irreducible representations of the cubic group \( O_h \) by the groups \( C_{4v}, C_{2v}, \) and \( C_{3v} \) for the three cases of \( K \) being oriented along the \([001]\), \([110]\), or the \([111]\) direction, respectively. In particular, for an exciton state having the symmetry \( \Gamma^\pm_0 \) at \( K = 0 \) degeneracies are lifted for \( K \neq 0 \).

We therefore observe a splitting of the \( 1S_o (\Gamma^\pm_0) \) ortho exciton state depending on \( K \) when solving the full \( K \)-dependent Hamiltonian of excitons in \( \text{Cu}_2\text{O} \). This splitting is shown in Fig. 2. It was observed experimentally in Refs. [67–69] and originally discussed in terms of a \( K \)-dependent exchange interaction. However, a closer examination of this interaction revealed that it is far too weak in \( \text{Cu}_2\text{O} \) to describe the observed splitting [25]. Instead, it could be shown that the effects due to the cubic valence band structure lead to a \( K \)-dependent effective mass and a \( K \)-dependent splitting of the 1S ortho exciton [20]. Hence, the directional dispersion is the true cause of the experimentally observed splitting.

Note that in Ref. [20] the splitting was treated within a perturbation approach and it was already emphasized that the complete \( K \)-dependent Schrödinger equation including the central cell corrections would have to be solved to obtain correct results. This has now been done.

As the 1S ortho exciton state exciton state has the symmetry \( \Gamma^\pm_0 \) for \( K = 0 \), we expect for \( K \parallel [001] \) and
\[ K \parallel [111] \text{ a splitting into two degenerate and one non-degenerate state. For } K \parallel [110] \text{ all degeneracies are lifted. This splitting can be described by the Hamiltonian} \]

\[
H_{\text{disp}}(K) = \Delta_1 \begin{pmatrix} K^2 & 0 & 0 \\ 0 & K^2 & 0 \\ 0 & 0 & K^2 \end{pmatrix} + \Delta_3 \begin{pmatrix} 3K_x^2 - K^2 & 0 & 0 \\ 0 & 3K_y^2 - K^2 & 0 \\ 0 & 0 & 3K_z^2 - K^2 \end{pmatrix} + \Delta_5 \begin{pmatrix} 0 & K_xK_y & K_xK_z \\ K_xK_y & 0 & K_yK_z \\ K_xK_z & K_yK_z & 0 \end{pmatrix}. \tag{75}
\]

We can prove the consistency with this formula by diagonalizing the Hamiltonian (75) for \( K \parallel [001], K \parallel [110], \text{ and } K \parallel [111] \) and fitting the resulting eigenvalues to our numerical results obtained with the full exciton Hamiltonian (96). This is shown in Fig. 2. The values of the fit parameters \( \Delta_1, \Delta_3, \text{ and } \Delta_5 \) are given in Table II. Since we have only three independent parameters \( \Delta_i \), but seven exciton states to be fitted, the consistency is proven by the fact that we obtain the same values of the parameters \( \Delta_i \) in all fits.

When performing the fit, it is not necessary to account for the \( K \)-dependent nonanalytic exchange interaction. In the next section VI B a formula for the size of the nonanalytic exchange interaction of the 1S exciton will be derived:

\[
\Delta_Q = 2f_0E_0/K_0^2. \tag{76}
\]

**TABLE II:** Results for the three coefficients \( \Delta_i \) when fitting the eigenvalues the Hamiltonian (75) to the theoretical spectra of Fig. 2 for the different orientations of \( K \). For the [001] and the [111] directions the values \( \Delta_5 \) and \( \Delta_3 \) cannot be determined, respectively. All results are given in \( 10^{-14} \mu eV m^2 \). For a comparison, we also list the experimentally determined values of Refs. [67–69]. For further information see text.

|         | [001] | [110] | [111] | exp. |
|---------|-------|-------|-------|------|
| \( \Delta_1 \) | 1.727 | 1.729 | 1.728 | 1.074 |
| \( \Delta_3 \) | -0.156 | -0.155 | -   | -0.189 |
| \( \Delta_5 \) | -0.213 | 0.213 | 0.292 |

Here \( f_0 = 3.6 \times 10^{-9} \) and \( E_0 \) denote the oscillator strength and the energy of the transverse exciton for \( K \parallel [001] \) at the exciton-photon resonance \( K_0 \), respectively. From the numerical results we obtain

\[
K_0 = 2.614 \times 10^7 \frac{1}{m}, \quad E_0 = 2.032 \text{eV} \tag{77}
\]

and, therefore,

\[
\Delta_Q = 2.135 \times 10^{-17} \mu eV m^2. \tag{78}
\]

Obviously, the nonanalytic exchange interaction is two to three orders of magnitude smaller than the anisotropic dispersion (cf. Table II).

The experimentally observed splitting was described by the ansatz (75) as well [67–69]. When comparing the results, we have to note that the values of the \( \Delta_i \) are given with respect to \( K_0 \) and that the factor \( \Delta_1 \) of

FIG. 2: Energy of the 1S, (\( \Gamma_\pm \)) ortho exciton state in dependence of \( K = |K| \). We do not plot the absolute energies but the energy difference \( \Delta E = E_{1S,K} - E_{1S,0} \) to show the small increase of the energy for \( K \neq 0 \). The 1S exciton obtains a finite oscillator strength for \( K \neq 0 \), which increases quadratically with \( K \). Furthermore, a \( K \)-dependent splitting of the three ortho exciton states can be observed. The green solid lines are the fits of the eigenvalues of the Hamiltonian (75) to the numerical results. For further information see text.
Refs. [67–69] only describes the “exchange interaction,” i.e., the interaction without the spherically symmetric part of the kinetic energy. Therefore, we have to compare ∆1/K02 + ℏ2/2M, ∆3/K02, and ∆5/K02 with our results of the parameters ∆i. Here M = me + mh ≈ 1.64m0 is the value of the isotropic exciton mass used in Refs. [67–69].

Within our model the average mass of the 1S ortho exciton is

\[ M = \frac{\hbar^2}{2K^2} \frac{3}{\text{Tr} [H_{\text{disp}}(K)]} = \frac{\hbar^2}{2\Delta_1} = 2.2 m_0, \]  

(79)

which is significantly larger than the sum \( M = m_e + m_h = m_e + m_0/\gamma_1 \approx 1.56m_0 \) of the isotropic quasi-particle masses. However, there is a clear deviation from the experimentally determined value of \( M = (3.0 \pm 0.2)m_0 \) [105–107]. The disparity in the masses can be explained by the fact that the electron and the hole are not bare particles but polarize the surrounding lattice and are thus accompanied by clouds of longitudinal optical phonons [47]. Hence, in the experiment always polaron masses are measured which are larger than the bare particle masses [47]. Although we accounted for this effect via the Haken potential in the Hamiltonian, we already stated in Ref. [28] that there are some difficulties in applying this theory to Cu2O due to the existence of two optical phonons contributing to the Fröhlich interaction and the small distance between the electron and the hole in the exciton ground state, for which the Haken potential cannot describe the non-Coulomb electron-hole interaction.

For the other two coefficients ∆3 and ∆5 a good agreement is obtained (see Table II), in particular, as regards the sign of the parameters. Note, however, that in Refs. [67–69] ∆Q has been assumed to be of the same order of magnitude as the parameters ∆i and that it has been included in the fit of the experimental data. This affects the values for ∆3 and ∆5 and is, therefore, another reason for the difference between the experimental and the theoretical values in Table II.

Finally, differences between the experimental and theoretical values of the parameters ∆i could furthermore be explained by the neglectation of polaronic nature of the ortho exciton in Refs. [67–69], in terms of small strains in the crystal [68] and uncertainties in the experimental values (cf. the large error bars in the figures of Ref. [67]). In particular, the fact that the experimental spectra are not identical when changing the angle of the laser beam by the same amount in opposite directions in Ref. [67] shows the presence of strains.

We wish to note that also splittings in the same order of magnitude are obtained, e.g., for the 2P exciton state. However, these splittings cannot be observed experimentally due to the large linewidth of the 2P exciton.

### B. 5 × 5 matrix model

The yellow 1S ortho exciton is well separated from the other exciton states regarding its energy. Hence, it can be treated separately from the other ones as regards polariton effects. In this section we set up a model with a 5 × 5 matrix, which allows calculating the dispersion of the 1S ortho exciton polaron for any direction of \( K \) close to the resonance (\( K \approx K_0 \)). This model includes the two photon states with the polarization vectors \( \hat{e}_{\xi K} \) and the three ortho exciton states \( \Psi_i \), which transform according to \( yz, zx \) and \( xy \).

First we will treat the oscillator strength and the Rabi frequency. Let us consider the most simple case with \( K \parallel [001] \). Due to group theoretical reasons, the three states \( \Psi_{yz}, \Psi_{zx}, \) and \( \Psi_{xy} \) are good eigenstates of the Hamiltonian. The \( \Psi_{zx} \) exciton interacts with the photon in \( x \) polarization and the \( \Psi_{yz} \) exciton interacts with the photon in \( y \) polarization. Let us denote the oscillator strength of these exciton states at the exciton-photon resonance by \( f_0 \).

For other orientations of \( K \) superpositions of the form \( \sum_i a_i \Psi_i \) are eigenstates of the Hamiltonian. From the expression (35) or especially from the form of the states \( |T_{\xi K}^\alpha \rangle \) (33) we can see that the \( K \)-dependent oscillator strength of these exciton states is given by

\[ f_{1S \xi K} = f_0 \left| \begin{pmatrix} \hat{e}_{\xi K_y} K_z + \hat{e}_{\xi K_z} K_y \\ \hat{e}_{\xi K_x} K_y + \hat{e}_{\xi K_y} K_x \\ \hat{e}_{\xi K_z} K_y + \hat{e}_{\xi K_y} K_z \\ \hat{e}_{\xi K_x} K_y + \hat{e}_{\xi K_y} K_x \\ \hat{e}_{\xi K_z} K_y + \hat{e}_{\xi K_y} K_z \end{pmatrix} \cdot \begin{pmatrix} a_{yz} \\ a_{zx} \\ a_{xy} \end{pmatrix} \right|^2 \]  

(80)

with the components \( \hat{e}_{\xi K_i} \) of the polarization vector \( \hat{e}_{\xi K} \).

For states of symmetry \( \Gamma^+_5 \) it has been shown in Ref. [25] that the nonanalytic exchange interaction can be written as

\[ H_{\text{exch}}^{\text{NA}}(K) = \frac{\Delta Q}{K^2} \begin{pmatrix} K_y^2 K_z^2 & K_x^2 K_y & K_y^2 K_z & K_x^2 K_y \\ K_x^2 K_y & K_x^2 K_z & K_y^2 K_z & K_x^2 K_y \\ K_y^2 K_z & K_y^2 K_z & K_y^2 K_z & K_y^2 K_z \\ K_y^2 K_z & K_y^2 K_z & K_y^2 K_z & K_y^2 K_z \end{pmatrix}. \]  

(81)

Contrary to dipole allowed excitons, the nonanalytic exchange energy depends on the fourth power of the angular coordinates of \( K \). The prefactor \( \Delta Q \) is connected to the oscillator strength and can be determined for the 1S state in the following way: For the special case of \( K \) being oriented in [111] direction, the \( \Gamma^+_5 \) state splits into one longitudinal \( \Gamma_1 \) and two transverse \( \Gamma_5 \) states. The longitudinal state is an eigenstate of the operator (81) with the eigenvalue \( \Delta Q/3 \). An excitation of the longitudinal exciton leads to an oscillating longitudinal polarization. Due to the Maxwell equation \( \nabla \cdot D = 0 \), the dielectric function must be zero. Hence, we have

\[ \varepsilon(\omega, K_0)|_{\Gamma_1} = \varepsilon_{b2} + \frac{\frac{3}{2} f_0 \varepsilon_{b2}}{1 - (E_0 + \frac{1}{3} \Delta Q K_0^2)^2 / (E_0)^2} = 0 \]  

(82)
[cf. also Eq. (71)]. Here \( E_0 = \hbar k_0 / \sqrt{\varepsilon_b} \) is the energy of the \( \Gamma_1 \) exciton at \( K = K_0 \) without the nonanalytic exchange interaction. Using \( f_0 \ll 1 \) [65, 66, 70], we obtain

\[
\Delta_Q = 2 f_0 E_0 / K_0^2 = 2 f_0 h^2 c^2 / \varepsilon_b. \tag{83}
\]

Combining all the \( K \) dependent effects for the 1S ortho exciton, we arrive at the Hamiltonian for the 1S ortho exciton-polariton in the rotating-wave approximation:

\[
H = \begin{pmatrix}
    H_{\text{ph}} & H_{\text{exc} - \text{ph}} \\
    H_{\text{exc} - \text{ph}}^\dagger & H_{\text{exc}}
\end{pmatrix} \tag{84}
\]

with the \( 2 \times 2 \) matrix \( H_{\text{ph}} \) containing the photon dispersion,

\[
H_{\text{ph}} = \frac{\hbar k}{\sqrt{\varepsilon_b}} \begin{pmatrix}
    1 & 0 \\
    0 & 1
\end{pmatrix} = E_0 \frac{K}{K_0} \begin{pmatrix}
    1 & 0 \\
    0 & 1
\end{pmatrix}, \tag{85}
\]

and \( H_{\text{exc}} = E_0 \mathbf{1} + H_{\text{disp}}(K) + H_{\text{exch}}^{\text{NA}}(K) \), \tag{87}

where \( \mathbf{1} \) is the \( 3 \times 3 \) identity matrix.

Note that the Rabi energy depends on the square root of the oscillator strength [cf. Eq. (50)]. Hence, questions about the sign of the terms \((\hat{e}_1 K_y K_z + \hat{e}_1 K_z K_y)\) in Eq. (86) may arise. However, for reasons of symmetry the terms must be linear in \( K \). As the photon has negative parity, i.e., since it transforms according to \( \Gamma_4 \) in \( O_h \), the terms have to change the sign if the direction of \( K \) is reversed.

The eigenstates of the Hamiltonian (84) can be calculated using an appropriate LAPACK routine [80].

\[ H_{\text{exc} - \text{ph}} = \frac{1}{2} \hbar \Omega_{\text{ph}} \frac{1}{K_0} \begin{pmatrix}
    (\hat{e}_1 K_y K_z + \hat{e}_1 K_z K_y) & (\hat{e}_1 K_y K_z + \hat{e}_1 K_z K_y) \\
    (\hat{e}_2 K_y K_z + \hat{e}_2 K_z K_y) & (\hat{e}_2 K_y K_z + \hat{e}_2 K_z K_y)
\end{pmatrix}, \tag{86}
\]

and light being polarized along \( \hat{e}_2 K \) no exciton-photon coupling occurs. For the transverse states we present the photon-like part for the two polarizations.

In Ref. [66] the group velocity of the 1S ortho exciton-polariton has been measured as a function of the photon energy for \( K || [110] \). From the dispersion shown in Fig. 3(b) and (e), we can directly calculate the group velocity via

\[
v_g = \frac{d\omega}{dK} = \frac{1}{\hbar} \frac{dE}{dK}. \tag{90}
\]

The result is shown in Fig. 4. It can be seen that the group velocity decreases on the lower polariton branch close to the resonance for increasing values of \( K \). However, for large values of \( K \) the polariton dispersion approaches the exciton dispersion, which increases quadratically in \( K \), so that \( v_g \) is then proportional to \( K \). Hence, there must be a minimum value of the group velocity. In the experiment group velocities as low as \( 40 \text{ km/s} \) could be measured [66]. However, it was not possible to measure the complete dispersion. In particular, the region of very low group velocities is not experimentally accessible. From Eq. (90) and the theoretical results it is possible to calculate all group velocities. The minimum value of \( v_g \) obtained in our calculations is

\[
v_{g, \text{min}} = 1.5 \text{ km/s}. \tag{91}
\]

at \( K = 2.758 \text{ km/m} \). When comparing our result for the group velocity, i.e., Fig. 4(b), with the experimental results of Ref. [66], a very good agreement can be observed.

From the results of the \( 5 \times 5 \) matrix model it is also possible to calculate the polarization vector of the photon-like part of the polariton or the orientation of the electric field as \( \sum_{\mu=1}^2 W_{\mu} K^\dagger \hat{e}_\mu \). Connected with the exciton-like part of the polariton via the
FIG. 3: Dispersion of the 1S ortho exciton polariton calculated using the 5 × 5 matrix model. For the different orientations of \( \mathbf{K} \) we give the photon-like part \( W_{\mu \xi K} \) for the two polarizations \( \xi \). It can be seen that for \( \mathbf{K} \parallel [110] \) and light being polarized along \( \hat{e}_2 \) no exciton-photon coupling occurs. We do not plot the absolute energies but the energy difference \( \Delta E = E - E_0 \).

For further information see text.

symmetric cross product of \( \hat{e}_1 \) and \( \mathbf{K} \) [cf. Eq. (80)] according to the group theoretical condition \( \Gamma_5^+ \otimes \Gamma_4^+ \rightarrow \Gamma_1^- \). One obtains for \( \Delta_3 = \Delta_5 = 0 \) states with purely longitudinal or transverse polarization. However, in the general case, the states are mixed longitudinal-transverse states and the polarization is not parallel to the applied electrical field.

Another interesting point is the polariton energies if the crystal is rotated about the [1 1 0] or the [1 1 1] axis (cf. Refs. [67–69]). In the case that the crystal is rotated about the [1 ¯1 0] plane, the \( \mathbf{K} \) vector passes all three orientations of high symmetry from [0 0 1] to [1 1 1] and then to [1 1 0]:

\[
\mathbf{K} = \frac{K}{\sqrt{2}} \begin{pmatrix}
\sin \varphi \\
\sin \varphi \\
\sqrt{2} \cos \varphi
\end{pmatrix}.
\]

The polarization vectors are given by

\[
\hat{e}_{1K} = \frac{1}{\sqrt{2}} \begin{pmatrix}
\cos \varphi \\
\cos \varphi \\
-\sqrt{2} \sin \varphi
\end{pmatrix}, \quad \hat{e}_{2K} = \frac{1}{\sqrt{2}} \begin{pmatrix}
-1 \\
-1 \\
0
\end{pmatrix}.
\]

Independent of the angle \( \varphi \) the symmetry of the problem is always \( C_3 \) [84]. This group contains the identity and a reflection at the plane with the normal vector \( \mathbf{n} = (1, -1, 0)/\sqrt{2} \), which is identical to one of the six \( \sigma_d \) reflections of \( O_h \). The irreducible representations of \( C_3 \) are either symmetric (\( \Gamma_1 \)) or antisymmetric (\( \Gamma_2 \)) under reflection. Hence, the complete problem falls into two parts: The linear combinations \( \Psi_{x y} \) and \( (\Psi_{z x} + \Psi_{y z})/\sqrt{2} \) of the ortho exciton states transform according to \( \Gamma_1 \) while the linear combination \( (\Psi_{x y} - \Psi_{y z})/\sqrt{2} \) transforms according to \( \Gamma_2 \). Furthermore, photons transform in \( C_3 \) also according to \( \Gamma_1 (\hat{e}_{1K}) \) and \( \Gamma_2 (\hat{e}_{2K}) \). Therefore, the problem decouples group theoretically in a \( \Gamma_1 \) problem with three polariton branches and a \( \Gamma_2 \) problem with only two polariton branches.

We keep the amount of \( \mathbf{K} \) fixed at \( K = K_0 \) and increase the angle \( \varphi \) from 0° to 90°. The result is shown in Fig. 5(a) and (b). From the photon-like part of the polaritons we can clearly see the decoupling of the problem. Furthermore, the expected degeneracies of the polariton states occur if \( \mathbf{K} \) is oriented in [0 0 1], [1 1 1] or [1 1 0] direction.

Especially the polariton dispersion for \( \mathbf{K} \parallel [1 1 2] \) (\( \varphi = \arccos(-2/\sqrt{6}) \)) is interesting as regards second harmonic generation (SHG) measurements. This dispersion is shown in Fig. 6. SHG measurements are selective with respect to \( \omega \) and \( \mathbf{K} \) creating bulk polaritons with \( 2\omega \) and \( 2K \). This allows the excitation of polaritons on the upper polariton branch. The lower polariton branch is inaccessible since the total \( K \)-vector is too small. Due to symmetry reasons it is not possible to observe SHG for \( \mathbf{K} \) being parallel to [0 0 1] or [1 1 0]. The SHG measurements in Ref. [71] were therefore performed in a configuration where \( \mathbf{K} \) is parallel to [1 1 1] or [1 1 2]. Especially SHG measurements with \( \mathbf{K} \parallel [1 1 2] \) are interesting because there exist 3 hybrid polariton branches having a polarization within the (1, 1, 0)-plane. The polariton states on the upper and intermediate polariton branch are both accessible in SHG. Therefore, this is a situation where two exciton polaritons can get coherently excited...
FIG. 4: (a) Group velocity \( v_g \) (colorscale) of the 1S ortho exciton polariton with \( \mathbf{K} \parallel [110] \) and \( \hat{\mathbf{e}}_1 \mathbf{K} \) polarization in dependence on the energy difference \( \Delta E = E - E_0 \) and the wave vector \( \mathbf{K} \). (b) and (c) The group velocity \( v_g \) only in dependence on the energy difference \( \Delta E = E - E_0 \) for a comparison with Fig. 3 of Ref. [66]. The minimum value of \( v_g \) depends on \( f_0 \) and is given by \( v_{g,\text{min}} = 1.5 \text{ km/s} \).

by means of the same nonlinear polarization \( P(2\omega, 2\mathbf{K}) \). The interference of both exciton polaritons should give rise to a polariton beat showing up in time resolved SHG measurements.

Let us now discuss the rotation about the [111] axis. The [111] axis is a threefold axis, for which reason after every 60° \( \mathbf{K} \) is oriented in a direction of the form [110]. Between each two of these cases \( \mathbf{K} \) is oriented in a direction of the form [112]. We have

\[
\mathbf{K} = \frac{\mathbf{K}}{\sqrt{6}} \begin{pmatrix}
\sqrt{3} \cos \varphi + \sin \varphi \\
-\sqrt{3} \cos \varphi + \sin \varphi \\
-2 \sin \varphi
\end{pmatrix},
\]

(94)

and the polarization vectors

\[
\hat{\mathbf{e}}_j \mathbf{K} = \frac{1}{3\sqrt{3}} \begin{pmatrix}
3j - 4 + \cos \varphi - \sqrt{3} \sin \varphi \\
3j - 4 + \cos \varphi + \sqrt{3} \sin \varphi \\
3j - 4 - 2 \cos \varphi
\end{pmatrix}, \quad j = 1, 2.
\]

(95)

The resulting spectrum when varying the angle \( \varphi \) and keeping the amount of \( \mathbf{K} \) fixed at \( \mathbf{K} = K_0 \) is shown in Fig. 5(c) and (d). In the cases with \( \mathbf{K} \) being of the form [110] only one exciton state is allowed for the polarization vector of the form [001] and all exciton states are forbidden for the other polarization vector. Note, however, that if we start with the configuration \( \mathbf{K} \parallel [010] \), \( \hat{\mathbf{e}}_1 \mathbf{K} \parallel [001] \), and \( \hat{\mathbf{e}}_1 \mathbf{K} \parallel [110] \) of Ref. [67] and perform a rotation about the [111] axis, none of the polarization vectors is oriented in the [010] direction at \( \varphi = 60° \) while \( \mathbf{K} \parallel [101] \) holds. Hence, only for \( \varphi = 0° \) all exciton states are forbidden for the polarization \( \hat{\mathbf{e}}_2 \mathbf{K} \) in Fig. 5(c) and (d).

In the cases with \( \mathbf{K} \) being of the form [112] all exciton states are allowed. Since the difference between \( \mathbf{K} \parallel [110] \) and \( \mathbf{K} \parallel [211] \) are only \( \Delta \varphi = 30° \), a rotation about the [111] axis with \( \varphi = 4° \) already shows a significant effect on the spectrum as can be seen especially from Fig. 5(d).

Exactly for this case with \( \varphi = 4° \) and the polarization \( \hat{\mathbf{e}}_2 \mathbf{K} \) the transmission spectrum of the ortho exciton-polariton is shown in Fig. 2(c) of Ref. [67]. The shape of the transmission spectrum of is clearly affected by polari-
ton dispersion. Photons with \( \mathbf{K} \parallel [112] \), i.e., for \( \varphi = \arccos(-2/\sqrt{3}) \) in Eqs. (92) and (93). In panel (a) there are three hybrid polariton branches having a polarization within the \((1, 1, 0)\)-plane. This is in contrast to the directions of high symmetry, where only two polariton branches occur. For further information see text.

We can exclude uncertainties in the parameter \( \Delta E \) of less than 1 eV. The third quadrupole-allowed state is \( \Delta E = 4 \mu eV \) is caused by polariton dispersion. If single photon spectroscopy was \( K \)-selective, this exciton polariton would show up with a similar small line width as is in the pure exciton case.

For a comparison with the experiment, we calculate the polariton dispersion for different values of \( K \) and keeping \( \varphi \) fixed to \( \varphi = 4^\circ \). The result is shown in Fig. 7. It can be seen that for the polarization \( \hat{\epsilon}_\xi K \) two states have a significant photon amplitude. Hence, these are the two states, which could be observed as very narrow lines in the experimental transmission spectrum [67]. However, while in the experiment the energy difference between the two states is about 4 \( \mu eV \), we obtain only 2.4 \( \mu eV \). We can exclude uncertainties in the parameter \( \Delta_1 \) or the angle \( \varphi \) since a variation of these parameters does not change the size of the splitting. Only a variation of \( \Delta_3 \) and \( \Delta_5 \) changes the size of the splitting. However, even if we set these parameters to the experimentally determined values listed in Table II, the splitting increases only to 3.0 \( \mu eV \). As we have already stated above, the presence of small strains in the crystal cannot be excluded. These may be the major reason for the observed discrepancy.

Of course, it is also possible that the positions of the transmission maxima in the experiment are not exactly given by the positions of the avoided crossings. In this case, one would have to calculate the transmission spectrum for the polariton dispersion using Pekar’s boundary conditions [108]. This is likewise difficult and beyond the scope of this work. Despite all that, we obtain a good qualitative agreement with the experiment.

VII. SUMMARY AND OUTLOOK

We presented the theory of exciton-polaritons in \( \text{Cu}_2O \). In the derivation of the formulas we accounted for all relevant effects which are needed to describe the spectra theoretically in an appropriate way, i.e., the complete valence-band structure, the exchange interaction, and the central-cell corrections [28]. This leads to a likewise complicated expression for the momentum-dependent Hamiltonian of excitons. Our method of solving the corresponding Schrödinger equation allows calculating dipole and quadrupole oscillator strengths, for which general formulas have been derived. The subsequent polariton transformation can be performed within the so-called rotating-wave approximation. Within this approximation it is straightforward to additionally account for the nonanalytic exchange interaction.

We have treated the dispersion of polaritons in \( \text{Cu}_2O \) using a multi-polariton concept. When considering the correct model of excitons in \( \text{Cu}_2O \), all states which have
the symmetry $\Gamma_5^+$ for $K = 0$ obtain a finite $K$-dependent oscillator strength, i.e., $S$, $D$, $G$, and higher excitons. The more complex spectrum of excitons when using the full Hamiltonian leads to a more complex polariton dispersion than in previous works [26, 100]. We also estimated that polariton effects for the yellow exciton states with $n \geq 2$ may only be observed using nonlinear spectroscopy methods and high quality crystals.

When including the complete valence band structure, a $K$-dependent splitting of the three components of the yellow $1S$ ortho exciton appears. In contrast to Ref. [20], we have solved the complete $K$-dependent Schrödinger equation numerically including also the correct values for the central-cell corrections of Ref. [28]. The splittings calculated are on the same order of magnitude as the central-cell corrections of Ref. [28]. The splittings observed experimentally [67–69].

As the exciton ground state is well separated from the other exciton states, it can be treated separately as regards polariton effects. Exploiting the symmetry properties of the $1S$ ortho exciton, we were able to set up a $5 \times 5$ matrix model, which allows for the calculation of the corresponding polariton dispersion for any direction of $K$. Using the $5 \times 5$ matrix model, we investigated the dispersion of the $1S$ ortho exciton polariton for different orientations of $K$. In a comparison of the results for the group velocity with experimental results we obtained a very good agreement. The calculations allowed us to determine the minimum value of the group velocity, which is not directly accessible in the experiment, to $v_{\text{g, min}} = 1.5 \text{ km}$. We also presented results for the two special cases that the $K$-vector is rotated in planes perpendicular to [110] and [111]. Especially in the second case, another comparison with experimental results was possible for a rotation angle of $4^\circ$. The splitting between the two allowed states obtained is smaller than in the experiment. This can, however, be explained in terms of the presence of small strains and uncertainties in the parameters $\Delta_i$.

As a next step, it is possible to calculate transmission spectra from the polariton dispersion obtained via the $5 \times 5$ matrix model by assuming Pekar’s boundary conditions [108] and applying the Fresnel equations. This would allow for an even better comparison of theoretical results with the experiment and may also lead to an understanding of the shape of the peaks in the spectra.

Furthermore, we plan to investigate exciton spectra in an external magnetic field in Voigt configuration, where, in contrast to the Faraday configuration of Ref. [21], a finite value of the wave vector $K$ must be considered.

Acknowledgments

F.S. is grateful for support from the Landesgraduiertenförderung of the Land Baden-Württemberg.

[1] J. A. Frenkel, Phys. Rev. 37, 17 (1931).
[2] J. A. Frenkel, Phys. Rev. 37, 1276 (1931).
[3] J. A. Frenkel, Phys. Z. Sowjetunion 9, 533 (1936).
[4] R. Peierls, Annalen der Physik 405, 905 (1932).
[5] G. H. Wannier, Phys. Rev. 52, 191 (1937).
[6] E. F. Gross and I. A. Karryjew, Dokl. Akad. Nauk. SSSR 84, 471 (1952).
[7] R. S. Knox, Theory of excitons, vol. 5 of Solid State Physics Supplement (Academic, New York, 1963).
[8] V. T. Agekyan, phys. stat. sol. (a) 43, 11 (1977).
[9] V. M. Agranovich and V. L. Ginzburg, Crystal Optics with spatial Dispersion and Excitons (Springer, Berlin, 1984).
[10] G. L. Rocca, in Electronic Excitations in Organic Based Nanostructures, edited by V. M. Agranovich and G. F. Bassan (Elsevier, Amsterdam, 2003), vol. 31 of Thin Films and Nanostructures, pp. 97–128.
[11] A. Stahl and I. Balslev, Electrodynamics of the Semiconductor Band Edge (Springer, Berlin, 1987).
[12] G. Czajkowski, F. Basani, and L. Silvestri, Riv. Nuovo Cimento 26, 1 (2003).
[13] T. Kazimierczuk, D. Fröhlich, S. Scheel, H. Stolz, and M. Bayer, Nature 514, 343 (2014).
[14] M. Aßmann, J. Thewes, D. Fröhlich, and M. Bayer, Nature Mater. 15, 741 (2016).
[15] F. Schweiner, J. Main, and G. Wunner, Phys. Rev. B 93, 085203 (2016).
[16] P. Grünwald, M. Aßmann, J. Heckötter, D. Fröhlich, M. Bayer, H. Stolz, and S. Scheel, Phys. Rev. Lett. 117, 133003 (2016).
[17] M. Feldmaier, J. Main, F. Schweiner, H. Cartarius, and G. Wunner, J. Phys. B: At. Mol. Opt. Phys. 49, 144002 (2016).
[18] J. Thewes, J. Heckötter, T. Kazimierczuk, M. Aßmann, D. Fröhlich, M. Bayer, M. A. Semina, and M. M. Glazov, Phys. Rev. Lett. 115, 027402 (2015), and Supplementary Material.
[19] F. Schöne, S. O. Krüger, P. Grünwald, H. Stolz, S. Scheel, M. Aßmann, J. Heckötter, J. Thewes, D. Fröhlich, and M. Bayer, Phys. Rev. B 93, 075203 (2016).
[20] F. Schweiner, J. Main, M. Feldmaier, G. Wunner, and Ch. Uihlein, Phys. Rev. B 93, 195203 (2016).
[21] F. Schweiner, J. Main, G. Wunner, M. Freitag, J. Heckötter, Ch. Uihlein, M. Aßmann, D. Fröhlich, and M. Bayer, Phys. Rev. B 95, 035202 (2017).
[22] F. Schweiner, J. Main, and G. Wunner, Phys. Rev. Lett. 118, 046401 (2017).
[23] J. Heckötter, M. Freitag, D. Fröhlich, M. Aßmann, M. Bayer, M. A. Semina, and M. M. Glazov, Phys. Rev. B 95, 035210 (2017).
[24] S. Zieilska-Raczyńska, D. Ziemkiewicz, and G. Czajkowski, Phys. Rev. B 95, 075204 (2017).
[25] F. Schweiner, J. Main, G. Wunner, and Ch. Uihlein, Phys. Rev. B 94, 115201 (2016).
[26] S. Zieilska-Raczyńska, G. Czajkowski, and D. Ziemkiewicz, Phys. Rev. B 93, 075206 (2016).
[27] S. Zielińska-Raczyńska, D. Ziemkiewicz, and G. Czajkowski, Phys. Rev. B 94, 045205 (2016).
[28] F. Schweiner, J. Main, G. Wunner, and Ch. Uihlein,
K. Cho, Phys. Rev. B 95, 195201 (2017).
[29] F. Schweiner, J. Main, and G. Wunner, Phys. Rev. E 95, 062205 (2017).
[30] F. Schweiner, P. Rommel, J. Main, and G. Wunner, Phys. Rev. B 96, 035207 (2017).
[31] F. Schöne, H. Stolz, and N. Naka, Phys. Rev. B 96, 115207 (2017).
[32] M. Kurz, P. Grünewald, and S. Scheel, Phys. Rev. B 95, 245205 (2017).
[33] D. Semkat, S. Sobkowiak, F. Schöne, H. Stolz, Th. Koch, and H. Fehske, arXiv:1705.08769 (2017).
[34] T. Stielow, S. Scheel, and M. Kurz, arXiv:1705.10527 (2017).
[35] J. Heckötter, M. Freitag, D. Fröhlich, M. Aßmann, M. Bayer, M. A. Semina, and M. M. Glazov, Phys. Rev. B 96, 125142 (2017).
[36] H. Stolz, F. Schöne, and D. Semkat, arXiv:1709.00895 (2017).
[37] R. Huber, B. A. Schmid, Y. R. Shen, D. S. Chemla, and R. A. Kaindl, Phys. Rev. Lett. 96, 017402 (2006).
[38] A. Baldreschi and N. O. Lipari, Phys. Rev. B 3, 439 (1971).
[39] A. Baldreschi and N. O. Lipari, Phys. Rev. B 9, 1525 (1974).
[40] A. Baldreschi and N. O. Lipari, Phys. Rev. B 8, 2697 (1973).
[41] N. O. Lipari and M. Altarelli, Phys. Rev. B 15, 4883 (1977).
[42] M. Altarelli and N. O. Lipari, Phys. Rev. B 15, 4898 (1977).
[43] K. Suzuki and J. C. Hensel, Phys. Rev. B 9, 4184 (1974).
[44] H. Haken, Il Nuovo Cimento 3, 1239 (1956).
[45] H. Haken, in *Halbleiterprobleme IV*, edited by W. Schottky (Vieweg, Berlin, 1957), pp. 1–48.
[46] H. Haken and W. Schottky, Z. Phys. Chem. 16, 218 (1958).
[47] C. F. Klingshirn, *Semiconductor Optics* (Springer, Berlin, 2007), 3rd ed.
[48] G. M. Kavoulakis, Y.-C. Chang, and G. Baym, Phys. Rev. B 55, 7593 (1997).
[49] C. G. Kuper and G. D. Whitefield, eds., *Polarons and Excitons* (Olive and Boyd, Edinburgh, 1963).
[50] J. T. Devreese and F. Peeters, eds., *Polarons and Excitons in Polar Semiconductors and Ionic Crystals*, vol. 108 of *NATO ASI Ser. B* (Plenum Press, New York, 1984).
[51] K. Cho, Phys. Rev. B 14, 4463 (1976).
[52] U. Rösler and H.-R. Trebin, Phys. Rev. B 23, 1961 (1981).
[53] H. Fröhlich, Advances in Physics 3, 325 (1954).
[54] J. Bardeen and W. Shockley, Phys. Rev. 80, 72 (1950).
[55] Y. Toyozawa, J. Phys. Chem. Solids 25, 59 (1964).
[56] J. J. Hopfield, Phys. Rev. 112, 1555 (1958).
[57] U. Fano, Phys. Rev. 103, 1202 (1956).
[58] V. M. Agranovich, Sov. Phys. JETP 10, 307 (1960).
[59] J. J. Hopfield, Phys. Rev. 112, 1555 (1958).
[60] J. J. Hopfield and D. G. Thomas, Phys. Rev. 132, 563 (1963).
[61] J. B. R. Claus, L. Merten, *Light Scattering by Phonon Polaritons*, vol. 75 of *Springer Tracts Mod. Phys.* (Springer, Berlin, 1975).
[62] V. M. Agranovich and V. L. Ginzburg, *Crystal Optics with Spatial Dispersion, and Excitons*, vol. 42 of *Springer Series in Solid State Sciences* (Springer, Berlin, 1984), 2nd ed.
[63] B. Hönerlage, R. Lévy, J. B. Grun, C. Klingshirn, and K. Bohnert, Phys. Rep. 124, 161 (1985).
[64] C. Klingshirn, in *Ultrafast Dynamics of Quantum Systems: Physical Processes and Spectroscopic Techniques*, edited by B. D. Bartolo (Plenum Press, New York, 1998), vol. 372 of *NATO ASI Ser. B*, pp. 143–211.
[65] D. Fröhlich, A. Kulik, B. Uebbing, A. Mysyrowicz, V. Langer, H. Stolz, and W. von der Osten, Phys. Rev. Lett. 67, 2343 (1991).
[66] D. Fröhlich, G. Dasbach, G. Baldassarri, H. von Högersthal, M. Bayer, R. Klieber, D. Suter, and H. Stolz, Solid State Commun. 134, 139 (2005).
[67] G. Dasbach, D. Fröhlich, H. Stolz, R. Klieber, D. Suter, and M. Bayer, Phys. Rev. Lett. 91, 107401 (2003).
[68] G. Dasbach, D. Fröhlich, R. Klieber, D. Suter, M. Bayer, and H. Stolz, Phys. Rev. B 70, 045206 (2004).
[69] G. Dasbach, D. Fröhlich, H. Stolz, R. Klieber, D. Suter, and M. Bayer, phys. stat. sol. (c) 2, 886 (2005).
[70] D. Fröhlich, J. Brandt, C. Sandfort, M. Bayer, and H. Stolz, phys. stat. sol. (b) 243, 2367 (2006).
[71] J. Mund, D. Yakovlev, D. Fröhlich, and M. Bayer (2017), to be published.
[72] M. A. Kanehisa, Physica B+C 117-118, 275 (1983).
[73] J. W. Hodby, T. E. Jenkins, C. Schwab, H. Tamura, and D. Trivich, J. Phys. C: Solid State Phys. 9, 1429 (1976).
[74] H. E. Swanson and R. K. Fuyat, NBS Circular 539, 112 (1953).
[75] O. Madelung and U. Rösler, eds., *Landolt-Börnstein*, vol. 17 a to i, 22 a and b, 41 A to D of *New Series, Group III* (Springer, Berlin, 1982-2001).
[76] A. R. Edmonds, *Angular momentum in quantum mechanics* (Princeton University Press, Princeton, 1960).
[77] Ch. Uihlein, D. Fröhlich, and R. Kenklies, Phys. Rev. B 23, 2373 (1981).
[78] J. Broeckx, Phys. Rev. B 43, 9643 (1991).
[79] See Supplemental Material, for explicit expressions of $H_0$, $H_1$, and $H_2$ as well as the matrix elements of $H_1$ and $H_2$, which enter the symmetric matrices $D$ and $M$.
[80] F. Schweiner, Z. Bai, S. Blackford, J. Demmel, J. Dongarra, J. D. Croz, A. Greenbaum, S. Hammarling, A. McKenney, et al., *LAPACK Users’ Guide* (Society for Industrial and Applied Mathematics, Philadelphia, PA, 1999), 3rd ed.
[81] U. Rösler, *Solid State Theory* (Springer, Berlin, 2009), 2nd ed.
[82] Y. Toyozawa, *Optical processes in solids* (Cambridge University Press, Cambridge, 2003).
[83] A. Abragam and B. Bleaney, *Electron Paramagnetic Resonance of Transition Ions* (Clarendon Press, Oxford, 1970).
[84] E. Anderson, Z. Bai, C. Bischof, S. Blackford, J. Demmel, J. Dongarra, J. D. Croz, A. Greenbaum, S. Hammarling, A. McKenney, et al., *LAPACK Users’ Guide* (Society for Industrial and Applied Mathematics, Philadelphia, PA, 1999), 3rd ed.
[85] For the detailed derivation of the formulas for the oscillator strengths for the three orientations of $K$ see the Supplemental Material.
[86] L. C. Andreani, in *Strong Light-Matter Coupling: From Atoms to Solid-State Systems*, edited by A. Auffèves, D. Gerace, M. Richard, S. Portolan, M. F. Santos, L. C. Kwek, and C. Miniatura (World Scientific, Singapore, 2014), pp. 37–82.
[87] W. C. Tait and R. L. Weiher, Phys. Rev. 178, 1404 (1969).
[88] J. Schmutzler, D. Fröhlich, and M. Bayer, Phys. Rev. B 96, 062205 (2017).
[89] F. Schweiner, P. Rommel, J. Main, and G. Wunner, Phys. Rev. E 95, 062205 (2017).
SUPPLEMENTARY INFORMATION

VIII. HAMILTONIAN AND MATRIX ELEMENTS

In the paper the Hamiltonian of excitons has been written in the form

\[ H = H_0 + (\hbar K) H_1 + (\hbar K)^2 H_2. \]  

We now give in Secs. VIII A-VIII C the expressions for \( H_0, H_1 \) and \( H_2 \) in terms of irreducible tensors for the case that the wave vector \( K \) is oriented in \([001], [110] \) or \([111] \) direction [1-4]. Note that we rotate the coordinate system to make the quantization axis or \( z \) axis coincide with the direction of \( K \). Hence, we rotate the coordinate system by the Euler angles \((\alpha, \beta, \gamma) = (\pi, \pi/2, \pi/4)\) for \( K \parallel [110] \) and by \((\alpha, \beta, \gamma) = (0, \arccos(1/\sqrt{3}), \pi/4)\) for \( K \parallel [111] \).

The first-order and second-order tensor operators used in the following correspond, as in Ref. [3], to the vector operators \( r, p, I, S_h \) and to the second-rank Cartesian operators

\[ i_{mn} = 3 \{ i_{m}, i_{n} \} - \delta_{mn} I^2, \]
\[ P_{mn} = 3 \{ p_{m}, p_{n} \} - \delta_{mn} p^2, \]

respectively. We also use the abbreviation

\[ D^2_k = \left( t^{(1)} \times S_h^{(1)} \right)^2_k. \]

The coefficients \( \gamma'_1, \mu' \) and \( \delta' \) are given by [2, 5]

\[ \gamma'_1 = \frac{m_0}{m_c}, \quad \mu' = \frac{6\gamma_3 + 4\gamma_2}{5\gamma'_1}, \quad \delta' = \frac{\gamma_3 - \gamma_2}{\gamma'_1}, \]

and we define by analogy [4]

\[ \nu = \frac{6\eta_3 + 4\eta_2}{5\eta_1}, \quad \tau = \frac{\eta_3 - \eta_2}{\eta_1}. \]

In Sec. VIII D we then give the matrix elements of the terms of the Hamiltonian in the basis \([\Pi] = [N, L; (I, S_h)] J; F, S_c, F_t, M_F] \) in Hartree units using the formalism of irreducible tensors [1]. The matrix elements of the first part \( H_0 \) of the Hamiltonian are given in the Appendices of Refs. [4, 6]. The functions of the form \( \langle R_1 \rangle_{N_L} \) and the quantities \( I_{N'} L' ; N L \) are taken from the Appendix of Ref. [4].

A. Wave vector in [001] direction

\[ H_0 = E_g - \frac{e^2}{4\pi\varepsilon_0 \varepsilon r} - \frac{e^2}{4\pi\varepsilon_0 r} \left[ \frac{1}{2\varepsilon_1} \left( e^{-r/\rho_{a1}} + e^{-r/\rho_{a1}} \right) + \frac{1}{2\varepsilon_2} \left( e^{-r/\rho_{a2}} + e^{-r/\rho_{a2}} \right) \right] \]
\[ + \frac{2}{3} \Delta \left( 1 + \frac{1}{\hbar^2} f^{(1)} \cdot S_h^{(1)} \right) + \left[ -V_0 + J_0 \left( \frac{1}{4} - \frac{1}{\hbar^2} S_h^{(1)} \cdot S_h^{(1)} \right) \right] V_{uc} \delta(r) \]
\[ H_1 = \frac{1}{2\hbar^2 m_e} \left\{ -2\sqrt{\frac{5}{3}} \mu' \left[ P^{(1)} \times I^{(2)} \right]_0^{(1)} - 4\sqrt{\frac{5}{3}} \delta' \left[ P^{(1)} \times I^{(2)} \right]_0^{(3)} \right\} + \frac{\eta_1}{\hbar^2 m_e} \left\{ \frac{2}{3} P^{(1)}_0 \left( I^{(1)} \cdot S^{(1)} \right) - 2\sqrt{\frac{5}{3}} \nu \left[ P^{(1)} \times D^{(2)} \right]_0^{(1)} - 4\sqrt{\frac{5}{3}} \tau \left[ P^{(1)} \times D^{(2)} \right]_0^{(3)} \right\}, \tag{101a} \]

\[ H_2 = \frac{\gamma_1 m_e - m_0}{2\gamma_1^2 \hbar^2 m_e^2} + \frac{m_0}{2\gamma_1^2 \hbar^2 m_e^2} \left\{ -\sqrt{\frac{5}{3}} \mu' + \frac{2}{5} \sqrt{6} \delta' \right\} I^{(2)}_0 + \frac{3\eta_1 m_0}{\gamma_1^2 \hbar^2 m_e^2} \left\{ \frac{1}{3} \left( I^{(1)} \cdot S^{(1)} \right) + \left( -\sqrt{\frac{2}{3}} \nu + \frac{2}{5} \sqrt{6} \tau \right) D^{(2)}_0 \right\}. \tag{101c} \]

**B. Wave vector in [110] direction**

\[ H_0 = E_k - \frac{e^2}{4\pi\varepsilon_0 e} \frac{1}{r} - \frac{e^2}{4\pi\varepsilon_0 r} \left[ \frac{1}{2\varepsilon_1^2} \left( e^{-r/\rho_1} + e^{-r/\rho_1} \right) + \frac{1}{2\varepsilon_2^2} \left( e^{-r/\rho_2} + e^{-r/\rho_2} \right) \right] 
+ \frac{2}{3} \Delta \left( 1 + \frac{1}{\hbar^2} I^{(1)} \cdot S^{(1)} \right) + \left[ -V_0 + J_0 \left( \frac{1}{4} - \frac{1}{\hbar^2} S^{(1)}_e \cdot S^{(1)}_h \right) \right] V_{uc} \delta(r) 
+ \frac{\gamma_1}{2\hbar^2 m_0} \left\{ \hbar^2 p^2 - \frac{\mu'}{3} \left( P^{(2)} \cdot I^{(2)} \right) + \frac{\delta'}{4} \left( \sum_{k=\pm 4} \left[ P^{(2)} \times I^{(2)} \right]_k^{(4)} \right) 
- \frac{\sqrt{5}}{6} \delta' \left( \frac{1}{10} \left[ P^{(2)} \times I^{(2)} \right]_0^{(4)} \right) \right\} 
+ \frac{3\eta_1}{\hbar^2 m_0} \left\{ \frac{1}{3} P^2 \left( I^{(1)} \cdot S^{(1)} \right) - \frac{\nu}{3} \left( P^{(2)} \cdot D^{(2)} \right) + \frac{\tau}{4} \left( \sum_{k=\pm 4} \left[ P^{(2)} \times D^{(2)} \right]_k^{(4)} \right) \right\} \]
\[ -\frac{\sqrt{7}}{6}\tau \left( \sum_{k=\pm 2} \left[ P^{(2)} \times D^{(2)} \right]^{(4)}_{k} + \sqrt{\frac{1}{10}} \left[ P^{(2)} \times D^{(2)} \right]^{(4)}_{0} \right), \]  

\[ H_1 = \frac{1}{2\hbar^2m_e} \left\{ -2\sqrt{\frac{5}{3}}\mu' \left[ P^{(1)} \times I^{(2)} \right]^{(1)}_{0} + \sqrt{3}\delta' \left( \sum_{k=\pm 2} \left[ P^{(1)} \times I^{(2)} \right]^{(3)}_{k} + \sqrt{\frac{2}{15}} \left[ P^{(1)} \times I^{(2)} \right]^{(3)}_{0} \right) \right\} + \frac{3\eta_1}{\gamma_1^2}\frac{m_0}{\hbar^2m_e} \left\{ \frac{1}{3} \left( I^{(1)} \cdot S^{(1)}_{h} \right) - \frac{1}{2}\tau \left( D^{(2)}_{-2} + D^{(2)}_{2} \right) + \left( -\sqrt{\frac{7}{3}}\mu' - \frac{1}{5}\sqrt{\frac{3}{2}}\delta' \right) I^{(2)}_{0} \right\}, \]  

\[ H_2 = \frac{\gamma'_m - m_0}{2\gamma'_m} + \frac{m_0}{2\gamma'_m} \left\{ -\frac{1}{2}\delta' \left( I^{(2)}_{-2} + I^{(2)}_{2} \right) + \left( -\sqrt{\frac{7}{3}}\mu' - \frac{1}{5}\sqrt{\frac{3}{2}}\delta' \right) I^{(2)}_{0} \right\}, \]  

\[ H_0 = E_h - \frac{e^2}{4\pi \varepsilon_0 \varepsilon r} - \frac{e^2}{4\pi \varepsilon_0 r} \left[ \frac{1}{2\varepsilon_1} \left( e^{-r/\rho_{\alpha 1}} + e^{-r/\rho_{\alpha 1}} \right) + \frac{1}{2\varepsilon_2} \left( e^{-r/\rho_{\alpha 2}} + e^{-r/\rho_{\alpha 2}} \right) \right] + \frac{2}{3} \Delta \left[ 1 + \frac{1}{\hbar^2} \left( I^{(1)} \cdot S^{(1)}_{h} \right) + \left[ -V_0 + J_0 \left( \frac{1}{4} - \frac{1}{\hbar^2} S^{(1)}_e \cdot S^{(1)}_{h} \right) \right] V_{uc}\delta(r) \right] + \frac{\gamma'_m}{2\hbar^2m_0} \left\{ \hbar^2 p^2 - \frac{\mu'}{3} \left( P^{(2)} \cdot I^{(2)} \right) \right\} + \frac{4}{27}\delta' \left( \sum_{k=\pm 3} k \left[ P^{(2)} \times I^{(2)} \right]^{(4)}_{k} - 3\sqrt{\frac{7}{10}} \left[ P^{(2)} \times I^{(2)} \right]^{(4)}_{0} \right) \right\} + \frac{3\eta_1}{\hbar^2m_0} \left\{ \frac{1}{3} p^2 \left( I^{(1)} \cdot S^{(1)}_{h} \right) - \frac{\nu}{3} \left( P^{(2)} \cdot D^{(2)} \right) + \frac{4}{27}\delta' \left( \sum_{k=\pm 3} k \left[ P^{(2)} \times D^{(2)} \right]^{(4)}_{k} - 3\sqrt{\frac{7}{10}} \left[ P^{(2)} \times D^{(2)} \right]^{(4)}_{0} \right) \right\}, \]  

\[ H_1 = \frac{1}{2\hbar^2m_e} \left\{ -2\sqrt{\frac{5}{3}}\mu' \left[ P^{(1)} \times I^{(2)} \right]^{(1)}_{0} - \frac{4}{9}\delta' \left( \sum_{k=\pm 3} k \left[ P^{(1)} \times I^{(2)} \right]^{(3)}_{k} - 6\sqrt{\frac{2}{5}} \left[ P^{(1)} \times I^{(2)} \right]^{(3)}_{0} \right) \right\} \]
\[ H_2 = \frac{\gamma_1' m_e - m_0}{2\gamma_1' m_e^2} + \frac{m_0}{2\gamma_1' m_e^2} \left\{ -\sqrt{\frac{2}{3}} \left( \mu' + \frac{4}{5} \delta' \right) I^{(2)}_0 \right\} \]

\[ + \frac{3n_1 m_0}{\gamma_1' h^2 m_e^2} \left\{ \frac{1}{3} \left( I^{(1)} \cdot S_h^{(1)} \right) - \sqrt{\frac{2}{3}} \left( \nu + \frac{4}{5} \tau \right) D^{(2)}_0 \right\}. \]

(D) Matrix elements and reduced matrix elements

\[ \langle \Pi' I^{(2)}_q | \Pi \rangle = \delta_{LL'} \cdot 3\sqrt{5} (-1)^{F_i' + F_i - M_{F_i} + 2F' + 2J + L} \]

\[ \times \left[ (2F_i + 1) (2F_i' + 1) (2F + 1) (2F' + 1) (2J + 1) (2J' + 1) \right]^\frac{1}{2} \]

\[ \times \left( \begin{array}{c} F_i' \\ -M_{F_i'} \\ q \end{array} \right) \left( \begin{array}{c} F_i' \\ F_i' \frac{1}{2} \\ F_i \end{array} \right) \left( \begin{array}{c} J' \\ F' \end{array} \right) \left( \begin{array}{c} L \\ 1 \end{array} \right) \]

\[ \times \sum_{j=-1}^{1} (R_1^j)^2 \left[ N + L + j + 1 \right]^{-1} \delta_{N',N+j}, \]

\[ \langle \Pi' D^{(2)}_q | \Pi \rangle = \delta_{LL'} \cdot 3\sqrt{5} (-1)^{F_i' + F_i - M_{F_i} + 2F' + L + J + \frac{1}{2}} \]

\[ \times \left[ (2F_i + 1) (2F_i' + 1) (2F + 1) (2F' + 1) (2J + 1) (2J' + 1) \right]^\frac{1}{2} \]

\[ \times \left( \begin{array}{c} F_i' \\ -M_{F_i'} \\ q \end{array} \right) \left( \begin{array}{c} F_i' \\ F_i' \frac{1}{2} \\ F_i \end{array} \right) \left( \begin{array}{c} J' \\ F' \end{array} \right) \left( \begin{array}{c} L \\ 1 \end{array} \right) \]

\[ \times \sum_{j=-1}^{1} (R_1^j)^2 \left[ N + L + j + 1 \right]^{-1} \delta_{N',N+j}, \]

\[ \langle \Pi' | \left( I^{(1)} \cdot S_h^{(1)} \right) | \Pi \rangle = \delta_{LL'} \cdot \delta_{M_{F_i} M_{F_i'}} \cdot 3\sqrt{3} (-1)^{F_i + 2F' + L + J + \frac{1}{2}} \]

\[ \times \left[ (2F_i + 1) (2F + 1) (2F' + 1) (2J + 1) (2J' + 1) \right]^\frac{1}{2} \]
\[
\times \left\{ \begin{array}{cc}
F' & F_t \\
F_t & F
\end{array} \right\} \times \left\{ \begin{array}{cc}
J' & F' \\
F & J
\end{array} \right\} \times \left\{ \begin{array}{cc}
\frac{1}{2} & \frac{1}{2} \\
\frac{1}{2} & \frac{1}{2}
\end{array} \right\} \\
\times \sum_{j=-1}^{1} (R_1)^j_{NL} [N + L + j + 1]^{-1} \delta_{N'L',N + j},
\]

(106)

\[
\langle \Pi' \left| P^{(1)} \times I^{(2)} \right\rangle^{(K)}_q \rangle \Pi \rangle = 3\sqrt{5} (-1)^{F_t' + F_t - M_{F_t} + F' + J + K} \langle N' L' \parallel P^{(1)} \parallel N L \rangle
\]

\[
\times \left\{ \begin{array}{cc}
F' & F_t \\
F_t & F
\end{array} \right\} \times \left\{ \begin{array}{cc}
J' & F' \\
F & J
\end{array} \right\} \times \left\{ \begin{array}{cc}
\frac{1}{2} & \frac{1}{2} \\
\frac{1}{2} & \frac{1}{2}
\end{array} \right\},
\]

(107)

\[
\langle \Pi' \left| P^{(1)} \times D^{(2)} \right\rangle^{(K)}_q \rangle \Pi \rangle = 3\sqrt{5} (-1)^{F_t' + F_t - M_{F_t} + F' + \frac{1}{2} + K} \langle N' L' \parallel P^{(1)} \parallel N L \rangle
\]

\[
\times \left\{ \begin{array}{cc}
F' & F_t \\
F_t & F
\end{array} \right\} \times \left\{ \begin{array}{cc}
J' & F' \\
F & J
\end{array} \right\} \times \left\{ \begin{array}{cc}
\frac{1}{2} & \frac{1}{2} \\
\frac{1}{2} & \frac{1}{2}
\end{array} \right\},
\]

(108)

\[
\langle \Pi' \left| P^{(1)}_0 \left( I^{(1)} \cdot S^{(1)}_{h} \right) \right\rangle^{(K)}_q \rangle \Pi \rangle = \delta_{M_{F_t},M_{F_t}'} 9 (-1)^{F_t' + F_t - M_{F_t} + F' + \frac{1}{2}} \langle N' L' \parallel P^{(1)} \parallel N L \rangle
\]

\[
\times \left\{ \begin{array}{cc}
F' & 1 \\
-1 & M_{F_t} F_t & 0
\end{array} \right\} \times \left\{ \begin{array}{cc}
\frac{1}{2} & \frac{1}{2} \\
\frac{1}{2} & \frac{1}{2}
\end{array} \right\},
\]

(109)

\[
\langle N' L' \parallel P^{(1)} \parallel N L \rangle = \delta_{L',L+1} (-i) \left[ \frac{(2L + 3)(2L + 1)}{L + 1} \right]^{\frac{1}{2}}
\]

\[
\times \left[ \sum_{j=-1}^{1} \sum_{k=-2}^{0} (L_1)^{k,j}_{N',L+1} \left\{ (RP_1)^j_{N,L} (N_1)^1_{L,0} + \delta_{0,j} (D_1)^j_{L,0} \right\} I_{N' + k,L+2;N+j,L} \right]
\]

\[
+ \delta_{L',L-1} (i) \left[ \frac{(2L + 1)(2L - 1)}{L} \right]^{\frac{1}{2}}
\]
the cubic group $O_h$ by the groups $C_{4v}$ ($K \parallel [001]$), $C_{2v}$ ($K \parallel [110]$), and $C_{3v}$ ($K \parallel [111]$) [8]. Note that only the irreducible representation $\Gamma_5$ of $C_{4v}$ and the irreducible representation $\Gamma_3$ of $C_{3v}$ are two-dimensional. Hence, almost all degeneracies in the exciton spectrum are lifted for $\Gamma_5$ (see Table III). Especially for $\Gamma_5$ the irreducible representations of $O_C$ reduced to $C_{4v}$ and $C_{3v}$ in the [111] direction. For these directions we have to consider $\mathbf{K}z$, as in Ref. [7], we also rotate the coordinate system more, as in Ref. [7], we also rotate the coordinate system.

\[ \lim_{r \to 0} \left[ -i \frac{\partial}{\partial r} (T_{\xi K}^D |\Psi_{\nu K}) + \frac{\alpha K}{\sqrt{6}} (T_{\xi K}^Q |\Psi_{\nu K}) \right] \]

(111)

for the three different orientations of $\mathbf{K}$ in [001], [110], or [111] direction. For these directions we have to consider the reduction of the irreducible representations of the cubic group $O_h$ by the corresponding subgroups. Furthermore, as in Ref. [7], we also rotate the coordinate system to make the $z$ axis of the new coordinate system coincide with the direction of $\mathbf{K}$.

### IX. OSCILLATOR STRENGTH

In this section we give the formulas for the oscillator strength

\[ f_{\xi K} = \eta \lim_{r \to 0} \left[ -i \frac{\partial}{\partial r} (T_{\xi K}^D |\Psi_{\nu K}) + \frac{\alpha K}{\sqrt{6}} (T_{\xi K}^Q |\Psi_{\nu K}) \right]^2 \]

(112)

determine the correct linear combinations of the states

\[ |\pi_x^D \rangle = \frac{i}{\sqrt{2}} [2, 1)_{D} + |2, 1)_{D} \]

(113a)

\[ |\pi_y^D \rangle = \frac{1}{\sqrt{2}} [2, 1)_{D} - |2, 1)_{D} \]

(113b)

\[ |\pi_z^D \rangle = \frac{i}{\sqrt{2}} [2, 2)_{D} - |2, 2)_{D} \]

(113c)

and

\[ |\pi_x^Q \rangle = K_y |1, 0)_{Q} \]

\[ + \hat{K}_z \frac{i}{\sqrt{2}} [1, 1)_{Q} + |1, 1)_{Q} \]

(114a)

\[ |\pi_y^Q \rangle = K_x |1, 0)_{Q} \]

\[ + \hat{K}_z \frac{1}{\sqrt{2}} [1, 1)_{Q} - |1, 1)_{Q} \]

(114b)

\[ |\pi_z^Q \rangle = K_y \frac{1}{\sqrt{2}} [1, 1)_{Q} - |1, 1)_{Q} \]

\[ + \hat{K}_z \frac{i}{\sqrt{2}} [1, 1)_{Q} + |1, 1)_{Q} \]

(114c)

which transform according to the irreducible representations of $C_{4v}$. On the other hand, it is instructive that the correct linear combinations are given according to the direction of $\mathbf{K}$ and the polarization vectors $\hat{e}_{\xi K}$ transverse to $\mathbf{K}$:

\[ \hat{K} = (0, 0, 1)^T \]

(115a)

\[ \hat{e}_{1K} = (1, 0, 0)^T \]

(115b)

\[ \hat{e}_{2K} = (0, 0, 1)^T \]

(115c)

Since light is always transversely polarized, only states of symmetry $\Gamma_5$ are allowed. The correct linear combinations of the states (113) are

\[ \Gamma_1 : |L_{\mathbf{K}K}^D \rangle = |\pi_x^D \rangle \]

(116a)

\[ \Gamma_5 : |T_{1 \mathbf{K}K}^D \rangle = |\pi_y^D \rangle \quad \text{and} \quad |T_{2 \mathbf{K}K}^D \rangle = |\pi_y^D \rangle \]

(116b)

and likewise for the states of Eq. (114). If we assume the incident light to be circularly polarized, the oscillator strength is given by

\[ f_{\xi K} = \eta \lim_{r \to 0} \left[ -i \frac{\partial}{\partial r} (T_{\xi K}^D |\Psi_{\nu K}) + \frac{\alpha K}{\sqrt{6}} (T_{\xi K}^Q |\Psi_{\nu K}) \right]^2 \]

(117)
with

\[ |T^D_{+K}\rangle = -\frac{i}{\sqrt{2}} \left[ T^D_{1K}\rangle + iT^D_{2K}\rangle \right] = |2, -1\rangle_D, \quad (118a) \]

\[ |T^D_{-K}\rangle = \frac{i}{\sqrt{2}} \left[ T^D_{1K}\rangle - iT^D_{2K}\rangle \right] = -|2, 1\rangle_D \quad (118b) \]

and

\[ |T^Q_{+K}\rangle = -\frac{i}{\sqrt{2}} \left[ T^Q_{1K}\rangle + iT^Q_{2K}\rangle \right] = |1, -1\rangle_Q, \quad (119a) \]

\[ |T^Q_{-K}\rangle = \frac{i}{\sqrt{2}} \left[ T^Q_{1K}\rangle - iT^Q_{2K}\rangle \right] = -|1, 1\rangle_Q. \quad (119b) \]

Note that the sign \( \pm \) is defined by the direction of rotation of the polarization with respect to \( K \).

### B. Wave vector \( K \parallel [110] \)

For \( K \parallel [110] \) the symmetry is reduced to \( C_{2v} \) and it is

\[ \Gamma_4^- \rightarrow \Gamma_1 \oplus \Gamma_2 \oplus \Gamma_4. \quad (120) \]

The corresponding vectors are

\[ \hat{K} = (1, 1, 0)^T / \sqrt{2}, \quad (121a) \]

\[ \hat{e}_{1K} = (1, -1, 0)^T / \sqrt{2}, \quad (121b) \]

\[ \hat{e}_{2K} = (0, 0, 1)^T, \quad (121c) \]

so that the correct linear combinations of the states (113) read

\[ \Gamma_1 : |L^D_{1K}\rangle = \frac{1}{\sqrt{2}} \left[ |\pi^D_x\rangle + |\pi^D_y\rangle \right], \quad (122a) \]

\[ \Gamma_2 : |T^D_{1K}\rangle = \frac{1}{\sqrt{2}} \left[ |\pi^D_x\rangle - |\pi^D_y\rangle \right], \quad (122b) \]

\[ \Gamma_4 : |T^D_{2K}\rangle = |\pi^D_z\rangle. \quad (122c) \]

The correct linear combinations of the states \( |T^D_{\xi K}\rangle \) as well yields

\[ |L^D_{1K}\rangle = |\pi^D_{\xi'}\rangle = \frac{1}{\sqrt{2}} \left[ |2, -1\rangle_D - |2, 1\rangle_D \right], \quad (123a) \]

\[ |T^D_{1K}\rangle = |\pi^D_{\nu'}\rangle = \frac{i}{\sqrt{2}} \left[ |2, 2\rangle_D - |2, -2\rangle_D \right], \quad (123b) \]

\[ |T^D_{2K}\rangle = |\pi^D_{\nu'}\rangle = \frac{\sqrt{3}}{2} |2, 0\rangle_D \]

\[ + \frac{1}{\sqrt{8}} \left[ |2, -2\rangle_D + |2, 2\rangle_D \right], \quad (123c) \]

and

\[ |L^Q_{1K}\rangle = |\pi^Q_{\xi'}\rangle = \frac{1}{\sqrt{2}} \left[ |1, -1\rangle_Q - |1, 1\rangle_Q \right], \quad (124a) \]

\[ |T^Q_{1K}\rangle = |\pi^Q_{\nu'}\rangle = 0, \quad (124b) \]

\[ |T^Q_{2K}\rangle = |\pi^Q_{\nu'}\rangle = |1, 0\rangle_Q. \quad (124c) \]

The labels \( \xi' \), \( \nu' \) and \( \nu' \) are meant to indicate that the states are given in the rotated coordinate system. Finally, we calculate the oscillator strengths by evaluating

\[ f_{\nu\xi K} = \eta \left| \lim_{r \to 0} \left[ -\frac{i}{\sqrt{2}} \frac{\partial}{\partial r} \langle T^D_{\xi K}\rangle |\Psi_{\nu K}\rangle + \frac{\alpha K}{\sqrt{6}} \langle T^Q_{\xi K}\rangle |\Psi_{\nu K}\rangle \right]^2 \right| \quad (125) \]

for light which is polarized in [001] or in [1\( \overline{1} \)0] direction.

### C. Wave vector \( K \parallel [111] \)

For \( K \parallel [111] \) the symmetry is reduced to \( C_{3v} \) and we have

\[ \Gamma_4^- \rightarrow \Gamma_1 \oplus \Gamma_3 \quad (126) \]

with the vectors

\[ \hat{K} = (1, 1, 1)^T / \sqrt{3}, \quad (127a) \]

\[ \hat{e}_{1K} = (1, 1, -2)^T / \sqrt{6}, \quad (127b) \]

\[ \hat{e}_{2K} = (-1, 1, 0)^T / \sqrt{2}. \quad (127c) \]

The correct linear combinations of the states in Eqs. (113) are therefore

\[ \Gamma_1 : |L^D_{1K}\rangle = \frac{1}{\sqrt{3}} \left[ |\pi^D_x\rangle + |\pi^D_y\rangle + |\pi^D_z\rangle \right], \quad (128a) \]

\[ \Gamma_3 : |T^D_{1K}\rangle = \frac{1}{\sqrt{6}} \left[ |\pi^D_x\rangle + |\pi^D_y\rangle - 2 |\pi^D_z\rangle \right] \quad (128b) \]

\[ |T^D_{2K}\rangle = \frac{1}{\sqrt{2}} \left[ -|\pi^D_x\rangle + |\pi^D_y\rangle \right] \quad (128c) \]

We now choose the quantization axis parallel to \( K \), i.e., we rotate the coordinate system by the Euler angles \( (\alpha, \beta, \gamma) = (\pi, \pi/2, \pi/4) \). Rotating the states \( |L^D_{K}\rangle \) and
FIG. 8: (a) The exciton energies $E_{\nu K}$ in dependence on $K = |K|$ for $K \parallel [110]$. The colorbar shows the oscillator strengths for $\xi = \pi \nu$ polarized light. We denote from which states at $K = 0$ the exciton states originate (cf. Ref. [6]). For reasons of space we introduce the abbreviated notation $\Gamma_{i/j}^\pm$ to replace $\Gamma_i^\pm$, $\Gamma_j^\pm$. The blue solid line gives the photon dispersion $\hbar \omega_{\xi K} = \hbar K c / \sqrt{\epsilon_{b2}}$.

(b) Polariton dispersion obtained by solving the eigenvalue problem in Sec. IVC. (c) Photon-like part $W_{\mu K}$ of the polariton states. For further information see text.

Here we rotate the coordinate system by the Euler angles $(\alpha, \beta, \gamma) = (0, \arccos(1/\sqrt{3}), \pi/4)$. Rotating the states of Eq. (128) and assuming circularly polarized light yields

$$|L''_{D K}^\nu\rangle = |2, 0\rangle_D, \quad (129a)$$

$$|T''_{+K}^\nu\rangle = \frac{i}{\sqrt{3}} \left[ \sqrt{2} |2, -2\rangle_D + |2, 1\rangle_D \right]$$

and

$$|T''_{-K}^\nu\rangle = \frac{i}{\sqrt{3}} \left[ \sqrt{2} |2, 2\rangle_D + |2, -1\rangle_D \right], \quad (129c)$$

$$|L''_{Q K}^\nu\rangle = \frac{2}{\sqrt{3}} |1, 0\rangle_Q, \quad (130a)$$

$$|T''_{+K}^\nu\rangle = \frac{-i}{\sqrt{3}} |1, 1\rangle_Q$$

and

$$|T''_{-K}^\nu\rangle = \frac{-i}{\sqrt{3}} |1, -1\rangle_Q. \quad (130c)$$
We finally calculate the oscillator strengths by evaluating

\[
\mathcal{F}_{\xi\nu} = \lim_{r \to 0} \left| \eta - \frac{\alpha K}{\sqrt{6}} \left( \langle T'_{\pm K} | \Psi_{\nu K} \rangle \right)^2 \right|^2 - \frac{\partial}{\partial r} \left( T'_{\pm K} | \Psi_{\nu K} \rangle \right) \right| .
\]

(131)

**X. POLARITON DISPERSIONS FOR \( K \parallel [110] \) AND \( \xi = \pi \)**

In this section we present the dispersions of the exciton-polaritons with \( K \parallel [110] \) and \( K \parallel [111] \) using the Hamiltonian which accounts for the valence band structure, the exchange interaction, the central-cell corrections, and the finite momentum \( hK \) of the center of mass.

As we already stated in the manuscript, the differences in the dispersion relations for the different orientations \( K \parallel [001] \) or \( K \parallel [110] \) or \( K \parallel [111] \) are only slight. A change in the energies of the exciton states cannot be observed in Figs. 8, 9, and 10, since it is on the order of tens of \( \mu eV \). However, there is an important difference between the spectra: The values of the oscillator strengths significantly change due to the different symmetry breaking. For example, for \( K \parallel [110] \) the states which have the symmetry \( \Gamma_5^+ \) for \( K = 0 \) are dipole-forbidden in this case. For further information see text.
FIG. 10: Same spectra as in Fig. 8 but for $\mathbf{K} \parallel [111]$ and $\xi = \sigma^{\pm}_{zz}$ polarized light. For both polarizations the spectrum is identical. The oscillator strength of the $4F(\Gamma_7^-)$ state is too weak to be seen here. For further information see text.

The polariton dispersions differ, however, mainly in the vicinity of the avoided crossings.

[1] A. R. Edmonds, *Angular momentum in quantum mechanics* (Princeton University Press, Princeton, 1960).
[2] A. Baldereschi and N. O. Lipari, Phys. Rev. B 8, 2697 (1973).
[3] J. Broeckx, Phys. Rev. B 43, 9643 (1991).
[4] F. Schweiner, J. Main, M. Feldmayer, G. Wunner, and Ch. Uihlein, Phys. Rev. B 93, 195203 (2016).
[5] Ch. Uihlein, D. Fröhlich, and R. Kenkiles, Phys. Rev. B 23, 2731 (1981).
[6] F. Schweiner, J. Main, G. Wunner, and Ch. Uihlein, Phys. Rev. B 95, 195201 (2017).
[7] F. Schweiner, J. Main, G. Wunner, M. Freitag, J. Heckötter, Ch. Uihlein, M. Alßmann, D. Fröhlich, and M. Bayer, Phys. Rev. B 95, 035202 (2017).
[8] G. F. Koster, J. O. Dimmock, R. G. Wheeler, and H. Statz, *Properties of the Thirty-Two Point Groups* (M.I.T. Press, Cambridge, MA, 1963).
[9] A. Abragam and B. Bleaney, *Electron Paramagnetic Resonance of Transition Ions* (Clarendon Press, Oxford, 1970).