The fundamental role of quantized vibrations in coherent light harvesting by cryptophyte algae

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The influence of fast vibrations on energy transfer and conversion in natural molecular aggregates is an issue of central interest. This article shows the important role of high-energy quantized vibrations and their non-equilibrium dynamics for energy transfer in photosynthetic systems with highly localized excitonic states. We consider the cryptophyte antennae protein phycoerythrin 545 and show that coupling to quantized vibrations which are quasi-resonant with excitonic transitions is fundamental for biological function as it generates non-cascaded transport with rapid and wider spatial distribution of excitation energy. Our work also indicates that the non-equilibrium dynamics of such vibrations can manifest itself in ultrafast beating of both excitonic populations and coherences at room temperature, with time scales in agreement with those reported in experiments. Moreover, we show that mechanisms supporting coherent excitonic dynamics assist coupling to selected modes that channel energy to preferential sites in the complex. We therefore argue that, in the presence of strong coupling between electronic excitations and quantized vibrations, a concrete and important advantage of quantum coherent dynamics is precisely to tune resonances that promote fast and effective energy distribution.

I. INTRODUCTION

The functionality of natural multichromophoric complexes depends fully on the strength and structure of interactions between electronic states and vibrational degrees of freedom associated either to intramolecular vibrations or to intermolecular interactions with the protein and the surrounding solvent environment1-3. In these systems, the phonon environment influencing electronic excitation dynamics can be broadly divided into two distinct components.1 Firstly, a continuous distribution of modes with a characteristic cutoff frequency below or comparable to the thermal energy scale, $k_B T$. These modes, arising from the solvent and low-energy protein vibrations, induce thermal fluctuations of onsite energies which in turn drive transitions between energetically close excitonic states. Secondly, spectroscopy studies4-11 also reveal the active participation of specific vibrational modes during excitation dynamics, often described as peaks in the spectral density characterising exciton-phonon interactions. Vibrations with lower frequencies may arise from inter-molecular coupling between pigments and the local protein environment11,12 while peaks at high energies originate mostly from intramolecular vibrations that give rise to vibronic transitions13-16. We denote these high-energy vibrational modes as quantized vibrations to indicate that their quantum mechanical features can be significant even at ambient temperatures1. In contrast, vibrations with frequencies below $k_B T$ will be thermally activated. Indeed, the influence of quantized vibrations in energy transport in a variety of natural molecular systems is currently of central interest12-11,15.

In this work we discuss the important role of quantized vibrations and their non-equilibrium dynamics for electronic excitations in the cryptophyte antennae protein phycoerythrin 545 (PE545)19-20, where exciton coherence beating has been observed21. Chromophores in cryptophyte light-harvesting antenna proteins have large energy gaps and are separated by an average distance of 20 Å20, which is double the average pigment distance in proteins of plants22; hence the majority of electronic couplings in PE545 are small (<30 cm$^{-1}$)20. How can this antenna protein function efficiently with such structure? We show that a fundamental mechanism for the biological function of this complex and for supporting coherence oscillations at long-times is the coupling between excitons and vibrations quasi-resonant with excitonic energy differences. The importance of these interactions in similar light-harvesting systems has recently been highlighted by ultrafast spectroscopy experiment22-23.

Vibrations reported in spectroscopy of PE545 have high-frequencies20,22 with the largest couplings for modes between 514 cm$^{-1}$ and 1450 cm$^{-1}$. Given that these frequencies match closely energy gaps between exciton states, transport in PE545 is fundamentally activated by these vibrations giving rise to the transfer pathways and transfer times observed in experiment22. This result can be anticipated using Förster theory23 but is derived here using a non-Markovian polaron-representation master equation for multichromophoric systems24,25. This more general framework allows us to investigate the evolution of excitonic coherences and to show that non-equilibrium dynamics associated to such vibrations can contribute to the long-time oscillations of the coherences probed in recent ultrafast experiments22. Based on this analysis, we are able to elucidate two key features of
excitation transfer assisted by vibrations in cryptophytes. First, excitation dynamics follows a non-sequential order bridging large energy gaps and distances and therefore generating faster and wider spatial distribution of excitation energy. Second, vibrations that modulate the evolution of excitonic coherences also promote energy transfer to preferential sites in the complex. Both of these processes support biological function of the antennae by assuring that energy transfer to other photosynthetic complexes happens in the relevant biological time scale. On the basis of these results we argue that in complexes with highly localized excitons a fundamental role for quantum coherent contributions to dynamics is precisely to ensure that resonances with selected vibrations are tuned to guarantee faster and effective distribution of excitation energy. The interactions between excitons and quantized vibrations in close resonant with exciton gaps are therefore an important design principle for optimal light-harvesting antennae.

II. MARINE CRYPTOPHYTE ANTENNAE PROTEINS

Cryptophyte photosynthesis relies on phycobiliproteins such as PE454 and Phycocyanin 645 (PC645) to ensure photon absorption in the visible part of the spectrum, with an optimal performance in the blue (545 nm) and green (645 nm) regions. Excitation energy absorbed by cryptophyte antennae proteins is transferred towards photosystems II and II (PSI and PSII). Crystalllography reveals that PE454 contains eight light-absorbing bilin molecules covalently bound to a four subunit protein scaffold labelled A, B, C and D (see Figure 1a). The lowest-energy chromophores are 15,16-dihydroxybiliverdin chromophores and are linked to separate protein subunits A and B (DBV_A and DBV_B). The remaining six chromophores are phycerythrobilin (PEB) bilins, two of which are doubly covalently bound to the protein environment while the remaining four are singly bound. The PEB bilins are arranged such that pigments PEB50/61C, PEB82C and PEB158C occupy the subunit C, while chromophores PEB50/61D, PEB82D and PEB158D are attached to subunit D. One particular feature of this complex is that it contains a deep, water-filled slot exposing the pigments to a large contact with water. The combined pigment-protein structure is shown in Figure 1a while the relative positions of the isolated pigments is illustrated in Figure 1b.

A. The Electronic System

Quantum chemical calculations as well as spectroscopy have allowed the refinement of a parametrized Hamiltonian for the electronic structure of PE545 given by (ℏ = 1):

$$H_{el} = \sum_{m} \epsilon_{m} \sigma_{m}^{+} \sigma_{m} + \sum_{(m,n)} V_{mn} (\sigma_{m}^{+} \sigma_{n}^{+} + \sigma_{n}^{+} \sigma_{m}),$$

with $\sigma_{n}^{+}$ being the operator that creates an excitation on site $m$ i.e. $\sigma_{m}^{+}|0\rangle = |m\rangle$. A salient characteristic of this system is that site energy differences $|\epsilon_{m} - \epsilon_{n}|$ are very large in comparison to inter-site electronic interaction $V_{mn}$, for instance, the largest coupling is $V_{54} = 92$ cm$^{-1}$ between the central pigments PEB50C and PEB50D while their energy difference is about 1040 cm$^{-1}$ (remaining values of $V_{mn}$ can be found in the supplemental information). As a result, each excitonic eigenstate, which we denote $|\epsilon_{m}\rangle$, is highly localized to a particular site $m$. The table in Figure 1c presents the site energies of the eight chromophores in ascending order as well as the subunits to which each of them is bound. Notice the interesting relationship between the spatial arrangement of the pigments and their energies: the three highest energy chromophores are located in subunit D, the next three highest energy chromophores lie in subunit C and finally the two lowest chromophores reside in subunits B and A. Therefore, the hierarchy of pigment energies imposes also a hierarchy of energies among protein subunits. In section we illustrate how one of the functions of quantized vibrations is precisely to support efficient energy transfer between protein subunits.

B. System-Environment interaction

Optimal energy transfer in light harvesting systems depends crucially on the transitions and correlations that can arise from interaction between the electronic system and their environment. The later is usually described as a bath of harmonic modes of frequencies $\omega_{k}$, with boson creation operators $b_{k}^{+}$ and Hamiltonian $H_{B} = \sum_{k} \omega_{k} b_{k}^{+} b_{k}$. The linear interaction of this harmonic bath and the electronic system is given by

$$H_{I} = \sum_{m} \sigma_{m}^{+} \sigma_{m} \sum_{k} (g_{k,m} b_{k}^{+} + g_{k,m}^{*} b_{k}),$$

where $g_{k,m}$ is the coupling of site $m$ to mode $k$. The strength and structure of these interactions is characterized by the spectral density, $J(\omega) = \sum_{k} |g_{k}|^{2} \delta(\omega - \omega_{k})$ which has two contributions: coupling to low-frequency continuous modes denoted as $J_{CM}(\omega)$ and coupling to quantized vibrations denoted $J_{QM}(\omega)$ such that $J(\omega) = J_{CM}(\omega) + J_{QM}(\omega)$.

Guided by linear spectra studies on PE545 in this work we consider that $J_{CM}(\omega)$ is given by a super-Ohmic spectral density of the form

$$J_{CM}(\omega) = \frac{s_{1} \omega^{3}}{2T_{1}^{2}} e^{-\omega/\Omega_{1}} + \frac{s_{1} \omega^{3}}{2T_{1}^{2}} e^{-\omega/\Omega_{1}}^{\frac{1}{4}}.$$ 

Here the parameters (in units of cm$^{-1}$) for the continu-
nous modes are $s_1 = 3.46 \times 10^{-4}$, $s_2 = 2.02 \times 10^{-4}$, $\Omega_1 = 1.45 \times 10^{-3}$ and $\Omega_2 = 4.34 \times 10^{-3}$. We have chosen a super-Ohmic form for the continuous component of the spectral density taking into account that the frequency dependence of this function has been shown to appropriately reproduce vibrational sidebands\cite{30}. Moreover, the theoretical framework that we will use for describing dynamics is more accurate in this case\cite{24,25}.

Spectroscopy\cite{20} also reveals the influence of 14 vibrational components in the excitation dynamics on PE545, whose frequencies $\omega_{QM}$ and Huang-Rhys factors $s_{QM}$ are shown in Table I. Notice that the majority of the vibrations satisfy the condition $\hbar \omega_{QM} \geq k_B T \sim 200 \text{cm}^{-1}$ falling in the regime of high-energy. The largest couplings (given by $g = \omega_{QM} \sqrt{s_{QM}}$) are for modes $\omega_{QM} = 1111 \text{ cm}^{-1}$ and $\omega_{QM} = 1450 \text{ cm}^{-1}$. In this work we include the effect of these modes via a spectral density exhibiting sharp components. When combined with the non-Markovian polaron treatment (see\cite{23}), this provides a starting point to investigate the effects of nonequilibrium dynamics associated to these vibrations. In practice, these modes could be broadened by interactions with the surrounding environment. Therefore, in this work we assume broadened vibrational modes, each with a Lorentzian line shape of broadening $b$\cite{31}. The contribution of each of these modes to the spectral density is given by:

$$J_{QM}(\omega) = \frac{2s_{QM}G_{QM}}{\pi} \frac{\omega^3b}{(\omega^2 - \omega_{QM}^2)^2 + b^2\omega^2}.$$ (4)

A first insight into the relevance of the different components of $J(\omega)$ in excitation dynamics can be gained by understanding the relative position of excitonic transitions to the spectral density. Figure 2a shows the forms of $J_{CM}(\omega)$ and $J_{QM}(\omega)$ as well as superimposed bars indicating the excitonic energy differences of the electronic Hamiltonian (Eq. I). One striking feature in this figure is that the energies of the vibrations closely match a large number of the excitonic transitions. For example, the localized mode at $\omega_{QM} = 1111 \text{ cm}^{-1}$ lies close to resonance with the transition between eigenstates $|e_8\rangle$ and $|e_4\rangle$ and also to the transition between eigenstates $|e_7\rangle$ and $|e_2\rangle$. To understand the physical significance of these resonances, consider a scenario where a single localized mode is exactly on resonance with an excitonic
transition. In this case, the electronic system is able to emit a phonon into the mode, preserving energy during the transition. Therefore, quasi-resonant vibrations can activate otherwise energetically unfavorable excitonic transitions, bridging large energy gaps and, in this case, also large distances.

**Environmental Correlations:** The mechanisms supporting beating of excitonic coherences in light-harvesting systems are still under and they can indeed be system-dependent. In the case of PE545, it has been argued that non-Markovian dynamics arising from slow relaxing modes or solvent-induced correlated fluctuations of on-site energies can help supporting long-lasting oscillations of excitonic coherences. So far however none of these hypothesis has been experimentally probed. Indeed, atomistic descriptions have challenged the idea of spatially correlated fluctuations in other molecular complexes. However, solvent-induced correlated fluctuations can be expected to play a more significant role in systems where chromophores are in close proximity to water. An example is hydrated DNA where hole transport is affected by correlated fluctuations that arise from the nearby solvent. Interestingly, the structure of PE545 is such that a deep pocket of bound water occupies space between the monomers, thus allowing close contact of the pigments with the water molecules. Atomistic calculations to understand whether interaction with nearby solvent can induce correlated fluctuations in PE545 are under development. Therefore, in this work we compute line spectra and dynamics in PE545 with and without solvent-induced correlated fluctuations.

| $\omega_{\text{QM}}$ | 207 244 312 372 438 |
| $\omega_{\text{QM}}$ | 0.0924 0.0761 0.0578 0.0313 0.0578 |
| $\omega_{\text{QM}}$ | 514 718 813 938 1111 |
| $\omega_{\text{QM}}$ | 1450 1520 1790 2090 |

**TABLE I.** Vibrational parameters for PE545 taken from Ref. Frequencies in units of cm$^{-1}$.

Exact treatment of the exciton-phonon interactions across the whole range of energy and couplings is not a straightforward task. Importantly, in the regime of strong system-environment coupling, Förster theory accounts well for the influence of high-energy vibrations in transfer rates. However, Förster theory fails to capture the non-equilibrium dynamics of such vibrations due to interactions with the electronic components. A formal treatment of quantized vibrations consider them as part of what is denoted as system such that coherent quantum interactions between electronic and vibrational modes are accounted for. We illustrate such an approach with an example in section VC but the full quantum treatment of PE545 with 14 vibrational modes per pigment is beyond the scope of this paper. Instead, here we use a non-Markovian many-site polaron master equation, which allows consideration of non-equilibrium dynamics of high-energy quantized vibrations, that is the fact that these modes take some time to thermalize. As we will see here, these non-equilibrium dynamics induce modulation of excitonic population and coherences. The treatment is particularly accurate for systems where excitonic states are highly localized and hence suitable for describing dynamics and spectroscopy of PE545. Full details of the polaron master equation for multichromophoric systems are given in Refs. Notice that this formalism allows reconstruction not only of the site-population dynamics as originally proposed but also of the evolution of excitonic coherences. For self-consistency of the paper we summarize the framework here.

The polaron formalism describes a dynamics in a framework of displaced oscillators where the original Hamiltonian $H = H_{\text{el}} + H_B + H_I$ is transformed via a small polaron-transformation i.e. $\tilde{H} = e^{S}He^{-S}$ with $S = \sum_m \sigma_m^+ \sigma_m \sum_k (h_{k,m} b_k^\dagger - h_{k,m}^\dagger b_k)$ and $h_{k,m} = g_{k,m}/\omega_k$. In this shifted-oscillators framework, the effective electronic system is described by

$$\tilde{H}_{\text{el}} = \sum_m \tilde{\varepsilon}_m \sigma_m^\dagger \sigma_m + \sum_{(m,n)} V_{mn} \beta_{mn} (\sigma_m^+ \sigma_n^- + \sigma_n^+ \sigma_m^-).$$

Here $\tilde{\varepsilon}_m$ and $V_{mn} \beta_{mn}$ are shifted onsite energies and renormalized electronic interactions respectively. The eigenstates of $\tilde{H}_{\text{el}}$ are labelled $|\sigma_m\rangle$ and are denoted as renormalized exciton states from now on. Interaction with the bath of shifted oscillators induce fluctuations of

$$\tilde{J}_{mn}^\dagger (\omega) = \tilde{J}_{mn} (\omega) = \tilde{J}_{CM} (\omega) \sin(\omega r_{\text{mn}}/v_{\text{ph}})$$

and $n$'th chromophores, while $v_{\text{ph}} = 1000 \text{ ms}^{-1}$ is comparable to the speed of sound in water. If instead $v_{\text{ph}}$ is set to zero, we recover the case of independent harmonic baths. For a detailed justification of this model of solvent-induced correlations see Nalbach et al.
electronic couplings given by

\[ \tilde{H}_I = \sum_{(m,n)} V_{mn} (\tilde{B}_{mn} \sigma^+_m \sigma^-_n + \tilde{B}^\dagger_{mn} \sigma^+_n \sigma^-_m). \]  

Bath-induced renormalisation factors are defined as \( \beta_{mn} = (B_{mn}) \) while displaced bath operators become \( \tilde{B}_{mn} = B_{mn} - \beta_{mn} \) with \( B_{mn} = e^{-\frac{1}{2} \sum_k \delta h_{k,mn} b_k^\dagger b_k} \). When a common bath is assumed, the factor \( \delta h_{k,mn} = h_{k,m} - h_{k,n} \) depends on the difference in electron-phonon couplings of sites \( m \) and \( n \). For a harmonic oscillator bath in thermal equilibrium, the renormalisation factors evaluate to \( \beta_{mn} = e^{-\frac{1}{2} \sum_k \coth(\beta \omega_k/2) \delta h_{k,mn}} \).

Perturbation theory with respect to \( \tilde{H}_I \), leads to a non-Markovian polaron master equation

\[ \frac{d}{dt} \tilde{\rho}(t) = \mathcal{R}(t)\tilde{\rho}(t) + \mathcal{I}(t)\tilde{\rho}(0), \]

The homogeneous term \( \mathcal{R}(t)\tilde{\rho}(t) \) describes the relaxation of the electronic degrees of freedom due to the interaction with a thermalised environment. The inhomogeneous term \( \mathcal{I}(t)\tilde{\rho}(0) \) is only non-zero when the initial environmental state differs from the thermal state in the polaron frame. Therefore, this term allows us to capture the non-equilibrium dynamics of all vibrational components and in particular the fact that high-energy quantized vibrations are taken out of thermal equilibrium due to interaction with the electronic component. Explicit expressions for the homogeneous and inhomogeneous terms are presented in the supplemental information.

**Original frame observables** The master equation presented in Eq. (7) describes the evolution of system observables within the polaron frame \( (\tilde{\rho}(t)) \). However, we are ultimately interested in the dynamics as seen in the original untransformed ‘lab’ frame \( (\rho(t)) \) since this will be the appropriate operator to compute expected values of physical observables. A general prescription for the transformation of system observables from the polaron to the lab frame was presented in Kolli et al.\(^{24}\). These transformations can be summarised as follows: site populations \( \langle \sigma^+_m \sigma^-_m \rangle \) remain unaffected upon transformation to the lab frame as the operator \( \sigma^+_m \sigma^-_m \) commutes with the polaron transformation \( S \). Expected values of off-diagonal operators in the site basis are given by an elaborated transformation, which up to first order in \( \tilde{H}_I \) reads:

\[ \langle \sigma^+_m \sigma^-_n \rangle = \text{tr}\{\sigma^+_m \sigma^-_n \rho(0)\} + \beta_{mn} \text{tr}\{\sigma^+_m \sigma^-_n (\tilde{\rho}(t) - \tilde{\rho}(0))\} \]

\[ + \text{tr}\{\sigma^+_m \sigma^-_n S_1(t) \tilde{\rho}(t)\} + \text{tr}\{\sigma^+_m \sigma^-_n T_1(t) \tilde{\rho}(0)\}. \]

Expression for super-operators \( S_1(t) \tilde{\rho}(t) \) and \( T_1(t) \tilde{\rho}(0) \) are presented in the supplemental information.

**Linear Spectra:** Here we provide a derivation of the linear spectra based on the polaron-representation of the dynamics. Our derivation of the spectra appeals to the Markov and secular approximations in the polaron frame, formulation which may still capture some non-Markovian effects in the untransformed frame (see details in Kolli et al.\(^{24}\)). The absorption spectra of a multichromophoric system is defined in terms of Fourier transform of the dipole-dipole correlation function\(^{24}\):

\[ A(\omega) = \text{Re} \int_0^\infty dt \ e^{i \omega t} C_{d-d}(t) \]

The dipole-dipole correlation function is defined as the two-time correlation function of the transition dipole op-
where \( \mu(t) = e^{-iHt} \mu e^{iHt} \) is the system transition dipole moment in the Heisenberg picture. The system transition dipole operator \( \mu \) is given by the sum of the dipole operators for the individual chromophores 
\[
\mu = \sum_m \langle m | \mu_m | 0 \rangle + \text{h.c.}
\]
The dipole-dipole correlation function after transformation to the polaron frame reads
\[
C_{d-d}(t) = \text{Tr}_{S+B} \{ \mu(t) \mu(0) \rho_B | 0 \rangle \langle 0 | \}
\]
where the polaron frame dipole operator is given by:
\[
\tilde{\mu} = \sum_m \prod_k \left( D(-h_{k,m}) \mu_m | 0 \rangle | m \rangle + \text{h.c.} \right) = \sum_{\alpha,m} \prod_k \left( D(-h_{k,m}) \mu_m u_{m\alpha} | 0 \rangle | \alpha \rangle + \text{h.c.} \right)
\]
where \( D(h_{k,m}) = e^{b_{k,m} b_{k,m}^\dagger} \) is the bath displacement operator of mode \( k \) due to interaction with site \( m \)
and \( u_{m\alpha} = \langle m | \alpha \rangle \) is the amplitude of site \( m \) on renormalized eigenstate \( | \alpha \rangle \).

Under the Markovian and secular approximations within the polaron frame, the dipole-dipole correlation function can be written as:
\[
C_{d-d}(t) = \sum_\alpha C_\alpha(t) G_\alpha(t) \exp(i \omega_\alpha t)
\]
For each renormalized exciton \( | \alpha \rangle \) there are two contributions to the dipole-dipole correlation function: one arising from the displacement of the vibrational environment, and a component due to dephasing of the electronic degrees of freedom within the polaron frame. The bath correlation function in the polaron frame \( C_\alpha(t) \) is defined as
\[
C_\alpha(t) = \sum_{mn} \mu_{m,n}^* u_{m\alpha} u_{n\alpha} \text{Tr}_B \{ D(-h_{k,n}) D(h_{k,m}(t)) \hat{\rho}_B \}
\]
The electronic correlation function is \( G_\alpha(t) = \exp(-\tilde{\Gamma}_\alpha t) \) with a dephasing rate for each transition given by
\[
\tilde{\Gamma}_\alpha = \sum_\mu \left( \Gamma^{(1)}_{\alpha\mu,\mu\alpha} + \Gamma^{(2)}_{\alpha\mu,\mu\alpha} + \Gamma^{(3)}_{\alpha\mu,\mu\alpha} + \Gamma^{(4)}_{\alpha\mu,\mu\alpha} \right)
\]
and \( \Gamma^{(i)}_{\alpha,\beta,\mu,\nu} \) Markovian polaron rates given in Kolli et al.\[24\]. Full details of the above derivation is presented in Appendix A.

The final form of the absorption spectra is given by
\[
A(\omega) = \sum_\alpha \text{Re} \int_0^\infty dt \, e^{i(\omega-\omega_\alpha)t} \, C_\alpha(t) G_\alpha(t)
\]
In a similar manner we can derive the following expression for the fluorescence spectra:
\[
F(\omega) = \sum_\alpha n(\omega_\alpha) \text{Re} \int_0^\infty dt \, e^{i(\omega-\omega_\alpha)t} \, C_\alpha^*(t) G_\alpha^*(t)
\]
where \( n(\omega_\alpha) = (\exp(\beta \omega_\alpha) - 1)^{-1} \) is the thermal occupation of eigenstate \( \alpha \).

IV. SIGNATURES OF QUANTIZED VIBRATIONS IN SPECTRA AND DYNAMICS OF PE545

A. Signatures of quantized vibrations in the linear spectra

Experimental signatures of specific vibrations participating in excitation dynamics in light-harvesting systems can be seen in photon-echo spectroscopy\[5\], resonance Raman studies\[6\], fluorescence narrowing\[7\] and hole-burning experiments\[8,9\]. Here we show that the theoretical framework considered reproduces known room-temperature spectra and we point out signatures of quan-
tized high-energy vibrations in steady state fluorescence. Absorption and fluorescence spectra are computed in the absence and presence of spatial correlations and averaged over 10000 realisations of Gaussian static disorder with FWHM= 500 cm\(^{-1}\). These results are presented in Figure 3 together with the experimental data of Novoderezhkin et al.\(^{22}\) The polaron theory gives the correct location of the absorption peak frequency, though it does tend to underestimate the absorption just beyond this maximum (Figure 3). The differences in experimental and calculated absorption spectra may indicate that full quantum treatment of selected vibrations may be needed. Calculated and experimental fluorescence spectra are presented in Figure 3. We recover the measured experimental peak fluorescence after shifting the theoretical result by 6nm to the blue. Upon this correction, there is good qualitative agreement between theory and experiment.

Importantly, the main features of fluorescence are only recovered when quantized vibrations are included. While the continuous component of the spectral density \(J_{CM}(\omega)\) determines the width of the fluorescence profile, quantized high-energy vibrations described through \(J_{QM}(\omega)\) give rise to the asymmetry observed towards the blue (around 630 nm). The linear spectra are found to be unchanged as the degree of broadening, \(b\), of the modes is varied from 1 to 50 cm\(^{-1}\), being in good agreement with the results of Novoderezhkin et al.\(^{22}\)

Finally, it is worth noting that we obtain near identical spectra in the absence and presence of spatial correlations. Consequently, the dipole-dipole correlation function and the resulting spectra are insensitive to the degree of the spatial correlations. In contrast to the linear spectra, excited state dynamics is susceptible to spatially correlated fluctuations as we discuss in the next section.

**B. Signatures of quantized vibrations in dynamics**

In this section we highlight the effects of quantized vibrations by systematically comparing dynamics with and without such vibrations as well as comparisons when correlated fluctuations are included. We shall illustrate how the presence of quantized vibrations is evidenced in transfer times and pathways of energy transport in PE545 and in the beating profile of populations and coherences. Before proceeding to the dynamics, however, let us first briefly discuss two important aspects regarding the initial state and the basis chosen to describe excitation dynamics.

**Initial state.**- We consider excitation by an ultrafast laser pulse in a time scale much smaller than the timescale of the solvation dynamics. Under these conditions, the excitonic system will undergo a vertical Franck-Condon transition where the vibrational degrees of freedom remains in thermal equilibrium during the fast excitation process. Therefore we may assume that the excitation pulse can excite an excitonic state \(|e_m\rangle\) of the bare Hamiltonian (See Eq.(1)). In this work we assume that the laser pulse is shifted well into the blue end of the spectrum, such that it only excites the highest excitonic eigenstate of the \(|e_0\rangle\) corresponding to a highly localized excitation on PEB\(_{50D}\). This initial state is chosen in order to investigate the transfer pathways within the PE545 complex and to allow comparisons with previous works\(^{21,22}\). Furthermore this situation will let us evaluate whether coherent superpositions of excitonic states are evolved during the dynamics even if they were not present in the initial state.

**Basis of Relaxation.**- It is worth taking a moment to consider the correct basis in which to describe the relaxation dynamics. Assuming the single-excitation system relaxes towards a quasi-equilibrium state, a natural basis of relaxation will be that in which the density matrix becomes diagonal as the steady state is approached. For arbitrary system-bath coupling, such a basis may be constructed in terms of the eigenstates of an effective Hamiltonian describing equilibration\(^{23}\). In practice, however, the evaluation of this effective Hamiltonian is not a straightforward task and therefore we must choose an approximate basis. Our calculations indicate that the long-time density matrix in the polaron frame is diagonal in the basis of the renormalized Hamiltonian Eq.(5). We therefore choose such a basis to describe relaxation but now in the untransformed frame. In general, these renormalized states depend on the characteristics of system-bath interaction such that it will be different in the absence or presence of correlations. In the absence of solvent-induced correlations, we find that fluctuations induced by \(J_{CM}(\omega)\) lead to all bath-induced renormalisations \(\beta_{mn}\) evaluating to zero. As a result, the master equation Eq.(7) becomes a set of non-Markovian rate equations describing incoherent excitation transfer between sites. In the presence of spatial correlations corresponding to a speed of sound of 1000 ms\(^{-1}\), renormalisations are no longer zero and quantum coherent contributions to dynamics can be observed. The basis of relaxation in this case is a renormalized excitonic basis.

1. **Transfer times and pathways**

Quantized vibrations drastically change the transfer times and pathways for excitation transfer in PE545. To illustrate this we begin by presenting in Figure 4 snapshots of the distribution of the excitation in real space at different times in the presence of these vibrations (Figure 4a) and in their absence (Figure 4b). The main feature that can be appreciated from these snapshots is that in both situations excitation energy initially localized on PEB\(_{50D}\) is finally distributed to the three lowest lying chromophores PEB\(_{52C}\), DBV\(_A\) and DBV\(_B\). However, the timescale of this process is 5 ps when assisted by quantized vibrations and becomes 10 ps if they are neglected.
(a) Spatial distribution of excitation energy in presence of quantized vibrations.

(b) Spatial distribution of excitation in absence of quantized vibrations.

FIG. 4. Snapshots of population dynamics. The filling of each bar is proportional to the probability of each site to be excited, with a maximum height denoting a population of 0.5 or above.
Importantly, the timescale observed in experiments is about 5 ps agreeing with our predictions.

Comparison of the snapshots at 0.1 ps indicates that quantized vibrations activate transfer to its nearest neighbour chromophore PEB$_{50C}$ (also the most strongly interacting site). This can also be appreciated in Figure 5a depicting site population dynamics as a function of time. In contrast, when no quantized vibrations are included, excitation is distributed to the chromophore nearest in energy PEB$_{82D}$ as can be seen in Figure 5b. One striking feature of the distribution profile at 0.7 ps in Figure 4a is that excitation energy has been widely distributed over all the complex, while in the absence of fast vibrations energy is concentrated mainly on protein subunit D. Again, results at these intermediate timescales are in good agreement with experimental observations. The features observed in the presence of fast vibrations allow PE545 to perform its biological function: distribution of excitation across the antenna implies that energy can be efficiently transferred to other nearby antennae, and fast transfer to units A and B, which are those probably in contact with PSI or PSII, will aid energy conversion. This is further discussed in section 4.

Finally, it is worth noting that the participation of quantized vibrations in excitation dynamics guarantees that the general timescales of transfer and overall distribution of excitation energy remains practically unchanged as correlated fluctuations are considered (Figure 5b). This is no longer the case when fast modes are excluded. In this situation, since transport is mostly driven by thermal fluctuations, a small degree of correlations slows transfer (Figure 5b). The interplay between correlated fluctuations and quantized vibrations does exhibit interesting features when contributions of each individual mode to site populations are analyzed as shown in the next section.

2. Population dynamics

High-energy vibrations quasi-resonant with excitonic transitions manifest themselves not only in the activation of transport between widely energetically separate states but also in the oscillatory pattern of the dynamics. This arises primarily from the non-equilibrium dynamics of the vibrational components. To illustrate these features, we focus now on the population evolution of the renormalized state $|\alpha_1\rangle$, which is mostly localized on the central pigment PEB$_{50C}$. Figure 6(left) presents the population of PEB$_{50C}$ when the 14 vibrational modes are included and also when only the modes with frequencies $\omega_{QM} = 1111$ cm$^{-1}$ or $\omega_{QM} = 1450$ cm$^{-1}$ are present. While $\omega_{QM} = 1111$ cm$^{-1}$ is in close resonance with the energy detuning between the initially excited site and the central site, the mode with $\omega_{QM} = 1450$ cm$^{-1}$ has the largest coupling to each site. We find that both modes are able to activate transport in comparison to the situation when no mode is included (curve labeled (I)), with a larger rate enhancement achieved with the quasi-resonant mode as expected. The Fourier spectrum of the population dynamics with the full set of 14 modes is somewhat complicated (not shown). Hence we investigate the oscillatory behaviour when only one mode is present (see (I) to (III) in Figure 6(left)). These figures show that the subtle oscillations of these populations are due to the corresponding vibrational modes.

It is important to point out that off-resonance modes such as $\omega_{QM} = 1450$ cm$^{-1}$ are able to activate transport to PEB$_{50C}$ due to the interplay with thermal-induced fluctuations induced by $J_{CM}(\omega)$: fluctuations in excitonic energy splittings cause excitonic transitions to be close to an otherwise off-resonance vibration. Equally, fluctuations bring out of resonance transitions which are otherwise quasi-resonant with a fast mode. The interplay between thermal fluctuations and quantized vibrations bring us to the discussion of the interesting behaviour observed when a small degree of correlated fluc-
FIG. 6. Population dynamics of the excitonic state localized on PEB50C in the absence of spatial correlations (left) and in the presence of solvent-induced correlations (right). Lines labeled with (IV) denote population when all modes are included. Panels (I) to (III) depict Fourier transforms of populations when no LM is included, or when only $\omega_{QM} = 1111\text{cm}^{-1}$ or $\omega_{QM} = 1450\text{cm}^{-1}$ are present.

The population dynamics exhibits frequency components of both excitonic and vibrational origin. This can be seen in Figures 6(right)(I)-(III) which show the Fourier spectrum of populations when there is a small degree of solvent-induced correlations. When only the mode with $\omega_{QM} = 1450\text{cm}^{-1}$ is included the spectrum exhibits three peaks. One peak at $\sim 1450\text{cm}^{-1}$ due to the localized vibration assisting transport and a low-amplitude peak at $\sim 1040\text{cm}^{-1}$ which compares very well with the energy difference between eigenstates $|\alpha_8\rangle$ and $|\alpha_4\rangle$. Hence we may attribute this frequency component to a quantum-coherent excitonic oscillation and indicates that, in the basis chosen to describe the dynamics, populations can witness coherences. The feature at $400\text{cm}^{-1}$ in Figures 6(III)(right) does not correspond exactly to any transition frequency, therefore we believe that this peak arises from a beating between a frequency of excitonic coherence ($\omega \sim 1040\text{cm}^{-1}$) and the vibration-induced oscillation of frequency $\omega_{QM} = 1450\text{cm}^{-1}$. For comparison, we present the Fourier spectrum when no mode is present (see Figures 6(right)(I)) and when only the mode $\omega_{QM} = 1111\text{cm}^{-1}$ is included (Figures 6(II)(right)).
3. Coherence dynamics

As discussed in the previous section, quantum-coherent contributions to dynamics appear when solvent-induced correlated fluctuations are included. In this situation, bath-induced renormalizations do not vanish and they range between $\beta_{28} \sim 0.11$ and $\beta_{45} \sim 10^{-3}$ such that quantum coherent dynamics associated to $\hat{H}_{el}$ is at play. We therefore now investigate how non-equilibrium vibrations manifest themselves in the oscillatory profile of excitonic coherences. For simplicity, we consider the situation where only the mode $\omega_{QM} = 1111$ cm$^{-1}$ is present and show in Fig. 7(a) the evolution of the coherences between the two highest energy renomalised eigenstates i.e. $\rho_{08,07} = \langle \alpha_8 | \rho(t) | \alpha_7 \rangle$ as well as $\rho_{08,06} = \langle \alpha_8 | \rho(t) | \alpha_6 \rangle$. These off-diagonal matrix elements correspond to coherences investigated in room-temperature two-dimensional spectroscopy.\(^2\)

Large-amplitude oscillations are observed in the dynamics of both $\rho_{08,07}$ and $\rho_{08,06}$ but they last only for about 50-100 fs. On longer timescales, there is also a small-amplitude oscillatory component -a hundred times smaller than the oscillations in the short time scale. To investigate the origin of the two different oscillatory features, we once again calculate their Fourier transforms. The frequency spectrums over the 1 ps timescale are presented in Figure 7(b). Two features should be noticed. First, there is a very broad component due to the fast decaying oscillations seen at short times. For the matrix element $\rho_{08,07}$ this broad feature occurs at 500 cm$^{-1}$, which corresponds to the energy difference between eigenstates $|\alpha_8\rangle$ and $|\alpha_7\rangle$. Similarly for $\rho_{08,06}$ this broad features is observed around $\omega = 600$ cm$^{-1}$, which corresponds to the energy difference between eigenstates $|\alpha_8\rangle$ and $|\alpha_7\rangle$. Therefore, we may attribute these frequencies to quantum coherent evolution associated to $\hat{H}_{el}$. In contrast, the second feature within the Fourier spectra is much subtler and occurs in both cases at roughly 1100 cm$^{-1}$.

To investigate the fine structure of the spectrum of long-lived oscillations we perform the Fourier transform only over the time interval 250 fs to 1 ps (Figure 7(c)). Both off-diagonal elements exhibit a peak at 1100 cm$^{-1}$, which can be attributed to vibration-induced oscillations due to the localized mode with $\omega_{QM} = 1111$ cm$^{-1}$. Interestingly, however, the matrix element $\rho_{08,06}$ exhibits an additional peak at 1040 cm$^{-1}$, an energy scale comparable to the transition frequency between eigenstates $|\alpha_8\rangle$ and $|\alpha_4\rangle$. This is an indication of coherence-coherence coupling during the evolution and shows that excitonic coherences are generated during the dynamics even if they were not initially created.

In summary, we have shown that in the presence of a small degree of correlations supporting coherent evolution of electronic excitations, the beating pattern of excitonic coherences in the presence of vibrations out of equilibrium has a dual origin: short-time oscillations lasting for about 50-100 fs reflecting the time scale associated to the electronic Hamiltonian and a superimposed
FIG. 8. Total population within each protein subunit of PE545. For subunit D we have neglected the contribution of the initially excited chromophore PEB50D.

V. ROLE OF QUANTIZED VIBRATIONS

The results of the preceding section allow us to clearly point out three key roles of the strong-coupling between highly localized electronic excitations and quantized vibrations in the biological function of PE545.

A. Promotion of non-cascaded energy transfer

In the absence of quantized vibrations exciton-phonon interactions are dominated by the continuous distribution of modes that induce transitions between states near in energy. Given the local nature of excitonic states, transfer dynamics follows the hierarchy of energies of the chromophores. In light of the intimate connection between the hierarchy of energies and the physical arrangement of the pigments in the protein subunits (Figure 1c), we conclude that when quantized vibrations are neglected, electronic excitation moves through PE545 in a sequential manner, from D onto subunit C and finally to subunits B and A as shown in Figure 8b. Clearly this would be a very unfavourable dynamics for the biological function of transferring energy to other antennae.

In contrast, quantized vibrations take advantage of electronic interactions to promote non-sequential transfer by bridging the energy gap between subunits D and C and spreading out energy from each of them to A and B as shown in Figure 8a. In short, quantized vibrations in close resonance with excitonic transitions provide a built-in mechanism to control energy transfer pathways. In general, exploitation of environment and system resonances is an important design principle for directing energy transfer.

vibration-induced modulation that survives close to the picosecond timescale. Importantly, the two timescales identified here compare very well with recent experimental results on a similar complex.[12]
The position of the excitation defined as follows:

\[ \sigma(t) = \sum_m p_m(t)(|r_m|^2 - r^2(t)), \quad (17) \]

where \( p_m(t) \) denotes the population on chromophore \( m \) at a given time \( t \), \( r_m \) is the position vector of chromophore \( m \) and \( r(\omega) \) is the mean position of the excitation. The dynamics of this measure of spreading is shown Figure 9b. Using generalized Förster theory (details of which are not presented here) we have estimated that transfer times between adjacent PE545 complexes is of the order of 0.7 to 1 ps. In this biological timescale, quantized vibrations guarantee efficient inter-complex transfer by promoting a wider distribution of excitation energy (dotted vertical line in Figure 9b) and faster transfer to acceptor states (see dotted vertical line Figure 9a). These conclusions also hold when a small degree of correlated fluctuations is included as shown in insets of Figures 9a and 9b.

C. Exploitation of quantum coherent dynamics to tune useful resonances

We have discussed how a small degree of correlated fluctuations supports quantum-coherent dynamics of excitonic states and ensures tuning of a resonance with a preferred vibrational mode that enhances energy transport to a targeted site in the complex. For clarity this is summarized in Figure 10a showing the enhancement of the population of the central pigment PEB50C when coherent dynamics (correlations) is turned on and only the mode \( \omega_{QM} = 1111 \text{cm}^{-1} \) is considered. These results lead us to argue, that in the presence of quantized vibrations, any mechanism supporting coherent evolution of excitonic states, will also ensure that resonances enhancing transport are activated. The rationale behind this idea is simple: coherent evolution effectively implies that excitonic gaps during dynamics do not fluctuate much and hence frequencies associated to coherent evolution can be synchronized with quasi-resonant vibrations for longer times, which in turn enhances population transport.

VI. CONCLUDING REMARKS

The role of vibrations for adjusting and enhancing transport in wide variety of biological and chemical systems is currently of central interest. In this article we have illustrated the importance of quantized vibrations for efficient energy transport in photosynthetic systems where excitonic states are highly localized such as the PE545 antennae protein present in cryptophyte algae. The main feature allowing these systems to take full advantage of fast vibrations is the near-resonance be-
antennae. We are currently pursuing such study and our preliminary results confirm the features presented in this article and indicate that rich nonclassical and non-equilibrium behavior arise in these systems. Atomic calculations identifying the structural origin of such vibrations are therefore crucial. From the experimental viewpoint it is important to develop more direct measurements of the influence of such vibrations in channeling and/or enhancing excitation energy transfers as well as conclusive signatures of their quantum properties in conditions of biological relevance. The integration of physical descriptions as the ones presented in this article, together with atomic calculations and experimental characterization will provide useful insights for the design of antennae prototypes that exploit electronic interactions and promoting vibrations for controlled and enhanced performance.

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Appendix A: Linear Spectra Within the Polaron Formalism

In this Appendix, we derive an expression for the dipole correlation function within the polaron formalism used in section III. The dipole-dipole correlation function is defined as

\[ C_{d-d}(t) = \text{Tr}_{S+B}\{\mu(t)\mu(0)\rho_B|0\rangle\langle0|\} \] (A1)

where \( \mu(t) = e^{-iHt} \mu e^{iHt} \) and the system dipole operator \( \mu \) is defined as the sum of the dipole operators for the individual chromophores as follows:

\[ \mu = \sum_m \mu_m |0\rangle\langle m| + \mu^*_m |m\rangle\langle 0|. \] (A2)

Let us now transform into the polaron frame defined by the transformation \( H = e^{S}He^{-S} \), where \( S = \sum_m \sigma^+_m \sigma^-_m \sum_k (\hat{n}_{k,m} \hat{b}_k + \hat{n}^*_{k,m} \hat{b}^+_k) \), with \( \hat{n}_{k,m} = g_{k,m}/\omega_k \). Within the polaron frame, the dipole-dipole correlation function becomes:

\[ C_{d-d}(t) = \text{Tr}_{S+B}\{\tilde{\mu}(t)\tilde{\mu}(0)\tilde{\rho}_B|0\rangle\langle0|\} \] (A3)

while the dipole operator in the polaron frame is given by:

\[ \tilde{\mu} = e^{S}\mu e^{-S} \]
\[ \dot{\mu} = \sum_m \prod_K D(-h_{k,m}) \mu_m |0\rangle \langle m| + \text{h.c.} \]

\[ = \sum_{\alpha, m} \prod_K D(-h_{k,m}) \mu_m u_{m\alpha} |0\rangle \langle \alpha| + \text{h.c.} . \quad (A4) \]

where \( D(h_{k,j}) = e^{i h_{k,j} - \frac{\hbar}{2} h_{k,j}^2} \) is the bath displacement operator of mode \( k \) due to interaction with site \( j \).

Let us now proceed to calculate the dipole-dipole correlation function in steps. Firstly consider:

\[ \dot{\mu}(0) \hat{\rho}_B |0\rangle \langle 0| = \sum_{\alpha, m} \prod_K D(-h_{k,m}) \mu_m \mu_{m\alpha} |0\rangle \langle \alpha| + \text{h.c.} |0\rangle \langle 0| \hat{\rho}_B \]

\[ = \sum_{\alpha, m} \prod_K D(h_{k,m}) \mu_m^* u_{m\alpha} |0\rangle \hat{\rho}_B . \quad (A5) \]

Then the dipole-dipole correlation function is:

\[
C_{d-d}(t) = \sum_{\alpha, m, \beta, n} \prod_K \text{Tr}_{S+B} \{ e^{-i \hat{H} t} (D(-h_{k,n}) \mu_n u_{n\beta} |0\rangle \langle \beta| + D(h_{k,n}) \mu_n^* u_{n\beta} |0\rangle \hat{\rho}_B ) e^{i \hat{H} t} \}
\times D(h_{k,m}) \mu_m^* u_{m\alpha} |0\rangle \hat{\rho}_B \}

\[ = \sum_{\alpha, m, \beta, n} \prod_K \text{Tr}_{S+B} \{ D(-h_{k,n}) \mu_n u_{n\beta} |0\rangle \langle \beta| + D(h_{k,n}) \mu_n^* u_{n\beta} |0\rangle \hat{\rho}_B e^{-i \hat{H} t} \}
\times e^{i \hat{H} t} D(h_{k,m}) e^{-i \hat{H} t} e^{i \hat{H} t} \mu_m^* u_{m\alpha} |0\rangle \hat{\rho}_B e^{-i \hat{H} t} . \quad (A6) \]

At this point, we make a number of approximations to simplify the expression for the dipole-dipole correlation function. Firstly, we shall assume that the interaction Hamiltonian \( \hat{H}_I \) does not perturb the bath. This amounts to assuming that the bath is in the stationary, thermal state within the polaron frame (\( \hat{\rho}_B \)) at all times. Under this assumption we may write

\[ e^{i \hat{H} t} D(h_{k,m}) e^{-i \hat{H} t} = e^{i \hat{H}_0 t} D(h_{k,m}) e^{-i \hat{H}_0 t} = D(h_{k,m}(t)) . \quad (A7) \]

The second assumption we shall make is that all system coherence terms decouple. This amounts to making the secular approximation for the system dynamics. Therefore, we may write:

\[ e^{i \hat{H}_0(t) t} \mu_m^* u_{m\alpha} |0\rangle \hat{\rho}_B e^{-i \hat{H}_0(t) t} = \mu_m^* u_{m\alpha} \hat{\rho}_{0\alpha}(t) . \quad (A8) \]

Under these assumptions, the dipole-dipole correlation function has the following very simple form:

\[ C_{d-d}(t) = \sum_{\alpha, \beta, m, n} \text{Tr}_S \{ \mu_m^* \mu_n u_{m\alpha} u_{n\beta} |0\rangle \langle \beta| \hat{\rho}_{0\alpha}(t) \}
\times \prod_K \text{Tr}_B \{ D(-h_{k,n}) D(h_{k,m}(t)) \hat{\rho}_B \}

\[ = \sum_{\alpha} C_{\alpha}(t) \hat{\rho}_{0\alpha}(t) . \quad (A9) \]

where we have defined the dressed bath correlation function \( C_{\alpha}(t) \) as

\[ C_{\alpha}(t) = \sum_{m, n} \mu_m^* \mu_n u_{m\alpha} u_{n\alpha} \text{Tr}_B \{ D(-h_{k,n}) D(h_{k,m}(t)) \hat{\rho}_B \} . \quad (A10) \]

To complete the calculation of the linear spectra we need to know how each of the off-diagonal element \( \hat{\rho}_{0\alpha} \) evolves with time. To do so, let us consider the Markovian secular polaron master equation in the interaction picture:

\[ \frac{d\hat{\rho}_{0\alpha}(t)}{dt} = -\sum_{\mu \nu} \left( \Gamma^{(1)}_{\nu \mu} [S_{\mu \nu}, S_{\nu \mu} \hat{\rho}(t)] + \Gamma^{(2)}_{\nu \mu} [S^\dagger_{\mu \nu}, S_{\nu \mu} \hat{\rho}(t)] + \Gamma^{(3)}_{\nu \mu} [S_{\mu \nu}, S^\dagger_{\nu \mu} \hat{\rho}(t)] + \Gamma^{(4)}_{\nu \mu} [S^\dagger_{\mu \nu}, S^\dagger_{\nu \mu} \hat{\rho}(t)] + \text{h.c.} \right) . \quad (A11) \]

Details of this master equation can be found in Kolli et al.\cite{Kolli2013}. The equation of motion for the off-diagonal element \( \hat{\rho}_{0\alpha} \) is then given by:

\[ \dot{\hat{\rho}}_{0\alpha}(t) = -\sum_{\mu} \left( \Gamma^{(1)}_{\alpha \mu} + \Gamma^{(2)}_{\mu \alpha} + \Gamma^{(3)}_{\alpha \mu} + \Gamma^{(4)}_{\mu \alpha} \right) \hat{\rho}_{0\alpha}(t) . \quad (A12) \]

The general solution for the evolution of the coherence terms, in the Schrödinger picture, is then simply given by
\[ \tilde{\rho}_{\alpha\alpha}(t) = \exp(i\omega_{\alpha}t) \exp(-i\tilde{\Gamma}_{\alpha}t)\tilde{\rho}_{\alpha\alpha}(0), \]
where we have defined the total dephasing rate as \( \tilde{\Gamma}_{\alpha} = \sum_{\mu} \left( \Gamma_{\alpha\mu,\mu\alpha}^{(1)} + \Gamma_{\alpha\mu,\mu\alpha}^{(2)} + \Gamma_{\alpha\mu,\mu\alpha}^{(3)} + \Gamma_{\alpha\mu,\mu\alpha}^{(4)} \right) \). Therefore, the final expression for the dipole-dipole correlation function becomes:

\[ C_{d-d}(t) = \sum_{\alpha} C_{\alpha}(t) \exp(-\tilde{\Gamma}_{\alpha}t) \exp(i\omega_{\alpha}t). \] (A13)

Appendix B: Dimer model

This model considers the central dimer in PE545 formed by pigments PEB$_{50D}$ and PEB$_{50C}$. The excitonic Hamiltonian is given by

\[ H_e = \epsilon_1 \sigma_1^+ \sigma_1^- + \epsilon_2 \sigma_2^+ \sigma_2^- + V(\sigma_1^+ \sigma_2^- + \sigma_2^+ \sigma_1^-) \]

where the strength of the electronic interaction is \( V = 92 \text{ cm}^{-1} \) and the exciton eigenstates are labeled X and Y. Each pigment is coupled to a localized mode with frequency \( \omega_{QM} = 1111 \text{ cm}^{-1} \). The free Hamiltonian for the modes is

\[ H_{\text{modes}} = \omega_{QM} (b_1^\dagger b_1 + b_2^\dagger b_2) \]

and the linear interaction between the dimer coupled dimer is given by \( H_{e-modes} = g \sigma_1^+ \sigma_1^- (b_1^\dagger + b_1) + g \sigma_2^+ \sigma_2^- (b_2^\dagger + b_2) \). Here the strength of the site-mode coupling is \( g = 267 \text{ cm}^{-1} \). The initial state of the full dimer plus modes system \( \rho(0) = \rho_{\text{X}}(0) \otimes \rho_{\text{modes}}(0) \) is propagated under the total Hamiltonian \( H = H_e + H_{\text{modes}} + H_{e-modes} \).

The initial state of the dimer is chosen to be the highest energy eigenstate \( \text{X} \) such that in the absence of the harmonic modes no transport occurs. The modes are assumed to be initially in thermal equilibrium at room temperature (300K). For the purposes of calculations, the Fock space of the modes is truncated to a maximum occupation \( M = 8 \) which fully captures the dynamics of the high-energy modes for the parameters considered. We also introduce pure dephasing of onsite energies which results in a dissipator of the form

\[ D(\rho) = \gamma \sum_{\alpha} \left( \sigma_{\alpha}^+ \sigma_{\alpha}^- - \frac{1}{2} \sigma_{\alpha}^2 \right) \]

with \( \gamma = 50 \text{ cm}^{-1} \).

See supplementary material for details on the polaron formalism and a table of renormalization factors of electronic couplings when solvent-induced correlations have been included.
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