Single-Molecule Spectroscopy Reveals that Individual Low-Light LH2 Complexes from *Rhodopseudomonas palustris* 2.1.6. Have a Heterogeneous Polypeptide Composition

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ABSTRACT   *Rhodopseudomonas palustris* belongs to the group of purple bacteria that have the ability to produce LH2 complexes with unusual absorption spectra when they are grown at low-light intensity. This ability is often related to the presence of multiple genes encoding the antenna apoproteins. Here we report, for the first time to our knowledge, direct evidence that individual low-light LH2 complexes have a heterogeneous $\alpha\beta$-apoprotein composition that modulates the site energies of Bchl a molecules, producing absorption bands at 800, 820, and 850 nm. The arrangement of the Bchl a molecules in the "tightly coupled ring" can be modeled by nine $\alpha\beta$-Bchls dimers, such that the Bchls bound to six $\alpha\beta$-pairs have B820-like site energies and the remaining Bchl a molecules have B850-like site energies. Furthermore, the experimental data can only be satisfactorily modeled when these six $\alpha\beta$-pairs with B820 Bchl a molecules are distributed such that the symmetry of the assembly is reduced to C_3. It is also clear from the measured single-molecule spectra that the energies of the electronically excited states in the mixed B820/850 ring are mainly influenced by diagonal disorder.

INTRODUCTION

Purple photosynthetic bacteria have evolved an elegant system of modular units that make up the light-harvesting apparatus. These modules consist of pairs of hydrophobic, low-molecular-weight polypeptides, called α and β (usually 50–60 amino acids long), that noncovalently bind a small number of bacteriochlorophyll (Bchl a) and carotenoid (Car) molecules. The modules then oligomerize to produce the native circular, elliptical, or horseshoe complexes (1). Most purple bacteria have two main types of complexes: the core complex (RC-LH1) and the peripheral complex (LH2). In the photosynthetic membranes, LH2 surrounds the RC-LH1 complexes in a two-dimensional network (2,3).

The LH2 complex from *Rhodopseudomonas (Rps.) acidophila* is a nonameric complex (nine α/β units) that contains 27 Bchl a and nine carotenoid molecules (4,5). The Bchl a molecules are arranged in LH2 into two groups. The first group consists of nine well-separated monomeric Bchl a molecules, one for each $\alpha\beta$-apoprotein pair. The bacteriochlorin rings of this group lie flat within the plane of the membrane. These Bchl a molecules absorb at 800 nm and are collectively called B800. Each $\alpha\beta$-apoprotein apoprotein pair also binds two other Bchl a molecules, whose central Mg^{2+} are complexed via His residues to the $\alpha\beta$-apoprotein helices. These Bchl a molecules form a closely interacting ring and collectively give rise to the absorption at 850 nm (B850). It is now well established that the spatial arrangement of the pigments determines, to a large extent, the

spectroscopic features of the complexes, and that in these systems collective effects (especially in the case of the B850 Bchls) have to be considered to appropriately describe their electronically excited states (6–9). This leads to so-called Frenkel excitons, which arise from the interactions of the transition-dipole moments of the individual pigments and correspond to delocalized electronically excited states (10,11). Since the interaction strength between the individual pigments can be calculated on the basis of the available structural data, information about the pigment arrangement within the LH complexes can be obtained via optical spectroscopy. From this perspective, the LH2 complex from *Rps. acidophila* has served as a cornerstone for the development of a detailed understanding of structure-function relationships in such antenna systems (12,13).

Of interest, in some species of purple bacteria, such as *Rps. acidophila* or *Rps. palustris*, the exact spectral positions of the absorption bands can vary depending on the growth conditions. When grown under low-light (LL) conditions, the light-harvesting apparatus of *Rps. acidophila* strains 7050 and 7750 features an additional peripheral LH2 complex that absorbs at 820 nm (14–16). The ability to change the type of LH2 in response to growth at different light intensities is related to the presence of multiple $\alpha\beta$-polypeptides, which are (in the case of *Rps. acidophila*) encoded by at least four different $\alpha\beta$-apoprotein gene pairs (16,17). To distinguish the two types of peripheral LH2 complexes, they are often referred to as B800–850 and B800–820 complexes, or LH2 and LH3. The origin of this spectral variation results from fine-tuning of the electronic energy levels of the B850 molecules by interaction with the protein matrix. In the B800–850 complex, the C_3-acetyl

group of the α-bound B850 Bchl a molecule is hydrogen (H)-bonded to the αTyr[44] residue and the C_3-acetyl group of the β-bound B850 Bchl a molecule is H-bonded to the αTrp[45] residue (14). In contrast, in the B800–820 complex, the respective residues in these positions, i.e., αPhe[44] and αLeu[45], are unable to form hydrogen bonds. Instead the C_3-acetyl group of the αB820 Bchl a molecules is H-bonded to the αTyr[41] residue, and this locks the acetyl group into an out-of-plane position with respect to the bacteriochlorin plane. Similarly, the β-bound B820 Bchl a molecule, lacking any H-bonds, has its acetyl tail in an out-of-plane position with respect to the bacteriochlorin plane. Rotation of the acetyl group of the Bchl a molecule into an out-of-plane position reduces the extent of π-conjugation and results in a blue shift of the BChl a site energies (18). This then is reflected in the shift of the absorption band from 850 nm to 820 nm.

A microscopic understanding of the interplay between pigment organization and its spectral properties requires high-resolution structural information. Unfortunately, this information is only available for very few types of peripheral light-harvesting complexes, two B800–850 complexes (*Rps. acidophila* 10050 (4,5) and *Rhsp. molischianum* (19)), and one B800–820 complex (*Rps. acidophila* 7050 (14)). In each of these cases the LH2 complexes have a well-defined apoprotein composition. When cells of *Rps. palustris* are grown at HL intensity, they synthesize a standard LH2 complex. However, at LL intensities they produce an LH2 complex with a strikingly different absorption spectrum (20–23), as illustrated in Fig. S1 of the Supporting Material. The LL LH2 complex has been studied before, and indeed a low-resolution model based on crystal diffraction data at 7.5 Å resolution has been described (24). This model suggests that the complex is an $\alpha\beta$-octamer and that each of its $\alpha\beta$-apoprotein pairs binds an extra Bchl a relative to LH2 from *Rps. acidophila*. It has also been shown that the *Rps. palustris* genome contains five $\alpha\beta$-apoprotein gene pairs (25,26). Moreover, preparations of both HL and LL LH2 complexes from *Rps. palustris* contain a complex apoprotein composition, which raises the question as to whether its LH2 complexes consist of rings that each contain a mixture of apoprotein types, as opposed to the homogeneous case of, e.g., *Rps. acidophila* (22,27). However, it is difficult experimentally to distinguish between a mixture of LH2 complexes where each ring is homogeneous but there are various types of rings present, and where each individual ring has an inhomogeneous apoprotein composition. It is interesting to consider the possible effect that rings with a heterogeneous population of apoproteins might induce, especially since some of the *Rps. palustris* apoproteins have sequences that would change the site energies of the ''B850'' Bchls from the standard HL energies to ones more reminiscent of the B820 complex from *Rps. acidophila*. How would such an energetic heterogeneity affect the absorption properties of LH2? And does such heteroge-

neity actually exist? One way to address these questions is to use single-molecule spectroscopy.

The usefulness of single-molecule spectroscopy as a tool to unravel the spectroscopic complexities of light-harvesting proteins has been well demonstrated (28–35). (For reviews of single-molecule studies on bacterial light-harvesting complexes, see Cogdell et al. (36) and Berlin et al. (37).) In this study we used fluorescence-excitation spectra (i.e., fluorescence-detected absorption spectra) to investigate individual LL LH2 complexes from *Rps. palustris*. Our idea was to investigate whether different classes of single complexes exist, and to establish whether there is any evidence for spectral properties that can only be explained by rings with a heterogeneous apoprotein composition.

MATERIALS AND METHODS

Materials

Rps. palustris strain 2.1.6. was grown anaerobically in C-succinate media at 15 μW m^{-1} at 30°C. The cells were harvested after 120 h by centrifugation (1248 g, 30 min) and resuspended in 2-[N-Morpholino]ethanesulfonic acid (20 mM, pH 6.8; Sigma-Aldrich, Steinheim, Germany) containing KCl (100 mM). The harvested cells were broken by three passages through a French press (9500 psi) in the presence of a little DNAase and MgCl$_2$. The broken membranes were then sedimented by centrifugation at 184,000 g for 2 h. The pellet was resuspended and homogenized in Tris-HCl (20 mM, pH 8.0; Fisher Scientific, Leicester, UK) and its concentration was adjusted to yield an absorption at B850 of 70 cm^{-1}. The membranes were then solubilized by the addition of 1% v/v lauryldimethylamine N-oxide (LDAO; Fluka, Steinheim, Germany). After the mixture was stirred in the dark at room temperature for 30 min, any unsolubilized material was removed by centrifugation at 16,000 g for 10 min. The supernatant was then layered onto a sucrose step gradient. The gradient consisted of 0.8, 0.6, 0.4, 0.2 M sucrose prepared in Tris-HCl (20 mM, pH 8.0) containing LDAO (0.1% v/v). The gradients were then centrifuged overnight at 149,000 g at 4°C. The upper pigmented band contained the LH2 complexes, and the lower pigmented band contained the LH1-RC complexes. The band containing the LH2 complexes was collected and purified on a DE-52 cellulose column (Whatman, Maidstone, England). The purity of the LH2 complex was determined by calculating the ratio between the absorption maximum at B800 and the maximum at the protein absorption (A260). Fractions with a ratio > 3.0 were collected for spectroscopic analysis.

For single-molecule spectroscopy, the purified LL LH2 complexes were diluted to <10^{-9} M in Tris-HCl buffer (20 mM, pH 8.0) containing 0.1% LDAO. In the last dilution step, 2% (w/w) polyvinyl alcohol (PVA, mw 30,000–70,000 g mol^{-1}) was added to the LL LH2 and a drop of the solution was spin-coated onto a quartz substrate for 10 s at 500 rpm and 60 s at 2,500 rpm (model P6700; Specialty Coating System, Indianapolis, IN). This produced thin, amorphous polymer films of <1 μm thickness in which the LH2 complexes were embedded. Then the sample was immediately mounted in a helium-bath cryostat and cooled down to 1.4 K.

Low-temperature microscopy

Fluorescence-excitation spectroscopy was performed with the use of an in-house-built microscope that could be operated in either wide-field or confocal mode. The excitation source was a continuous-wave tunable titanium-sapphire laser (3900S; Spectra Physics, Mountain View, CA) pumped by a frequency-doubled continuous-wave neodynium:yttrium-vanadate (Nd:YVO4) laser (Millennia Vs; Spectra Physics). Well-defined changes in the wavelength of the Titanium:Sapphire laser were achieved by rotating

the intracavity birefringent filter with a motorized micrometer screw. For calibration purposes, a wavemeter was used, and the accuracy of the laser frequency and a reproducibility of 1 cm^{-1} were verified. First, a 50 × 50 μm^2 wide-field image of the sample was taken by exciting the sample at 800 nm and detecting the fluorescence with a back-illuminated CCD camera (512 SB; Roper Scientific Princeton Instruments, Trenton, NJ) after the light passed through suitable bandpass filters (BP893/21; Dr. Hugo Anders, Nabburg, Germany), which blocked the residual laser light. Subsequently, a spatially well-isolated complex was selected from the wide-field image and a fluorescence-excitation spectrum of this complex was obtained. The setup was switched to the confocal mode and the fluorescence was detected by a single-photon counting avalanche photodiode (APD, SPCM-AQR-16; EG&G Optoelectronics, Vaudreuil, Canada) while the laser was scanned repetitively between 784 and 872 nm. The recorded traces were stored separately in computer memory. With a scan speed of the laser of 3 nm · s^{-1} ($\approx$50 cm^{-1} s^{-1}) and an acquisition time of 10 ms per data point, this yields a nominal resolution of 0.5 cm^{-1}, ensuring that the spectral resolution is limited by the spectral bandwidth of the laser (1 cm^{-1}). To examine the polarization dependence of the spectra, a λ/2-plate was put in the confocal excitation path and rotated in steps of 3.2° between two successive scans, changing the angle of polarization of the excitation light by twice that value. The excitation intensity was ~50 W cm^{-2}.

RESULTS

The fluorescence-excitation spectra of several individual LL LH2 complexes from *Rps. palustris* are shown in Fig. 1. The top trace shows, for comparison, the fluorescence-excitation spectrum taken from an ensemble of LL LH2 complexes (*gray line*) together with the spectrum that results from the summation of the spectra of 31 individual LL LH2 complexes (*black line*). The two spectra are in reasonable agreement, indicating that the selected individual LL LH2 complexes are a fair statistical representation of the ensemble. The ensemble spectrum shows two broad bands centered at 11,684 cm^{-1} (856 nm) and 12,524 cm^{-1} (798 nm) with widths (full width at half-maximum (FWHM)) of 353 cm^{-1} and 174 cm^{-1}, respectively. The peak intensity of the B850 band is about three times lower than that of the B800 band, which agrees with previously published absorption spectra obtained both at room temperature and at 77 K (23).

By measuring the fluorescence-excitation spectra of the individual complexes, as shown for three examples in the lower traces of Fig. 1, we can visualize remarkable features that are obscured in the ensemble average. Around 12,500 cm^{-1} (800 nm), the spectra show a distribution of narrow absorption bands, with linewidths (FWHM) below 10 cm^{-1}, whereas around 11,750 cm^{-1} (850 nm) a few broad bands, with linewidths (FWHM) on the order of 160 cm^{-1}, are present. This observation is reminiscent of the situation for LH2 from *Rps. acidophila*. However, in striking contrast to the single-molecule spectra from LH2 from *Rps. acidophila*, we find that ~90% of the individual LL LH2 complexes from *Rps. palustris* show an additional broad band in the B820 spectral region. The peak position of this band varies between 12,048 cm^{-1} and 12,288 cm^{-1} (center of mass at 12,206 cm^{-1}) and its linewidth covers the range between 31 cm^{-1} and 158 cm^{-1} (average 79 cm^{-1}).
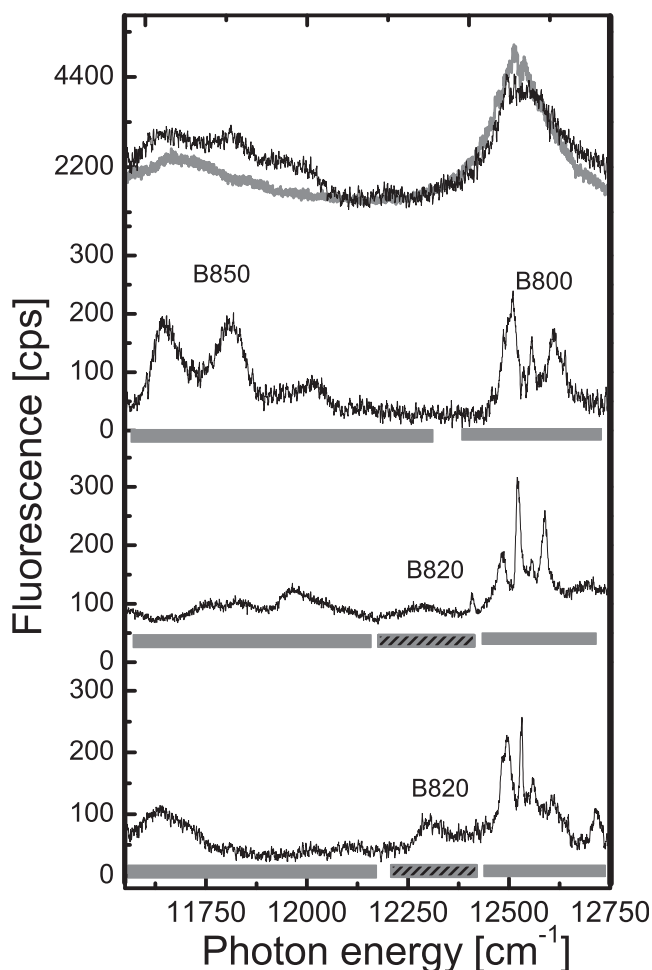

FIGURE 1 Low-temperature (1.4 K) fluorescence-excitation spectra of LH2 complexes from LL *Rps. palustris* 2.1.6. The top traces show the spectrum from an ensemble (*gray line*) and the spectrum that corresponds to the sum of 31 spectra from individual LH2 complexes (*black line*). The lower three traces display typical fluorescence-excitation spectra from individual LH2 complexes. The spectra have been averaged over all polarizations of the incident laser field. The bars indicate the spectral positions of the 800, 820, and 850 nm bands, respectively. The excitation intensity was 50 W cm^{-2}. The vertical scale is given in counted photons per second (cps).

To analyze the spectral bands from the individual LL LH2 complexes in more detail, we recorded the fluorescence-excitation spectra as a function of the polarization of the incident laser excitation. The excitation spectra were recorded in rapid succession as the polarization of the excitation light was rotated by 6.4° between two consecutive scans. An example of this protocol is shown in the top part of Fig. 2 A in a two-dimensional representation in which 410 individual scans are stacked on top of each other. The horizontal axis corresponds to photon energy, the vertical axis to the individual scans (or equivalently to the polarization of the excitation), and the detected fluorescence intensity is coded by the gray scale. The sum spectrum of these scans is presented at the bottom of Fig. 2 A. The pattern shown at the top of Fig. 2 A clearly reveals the polarization dependence of the
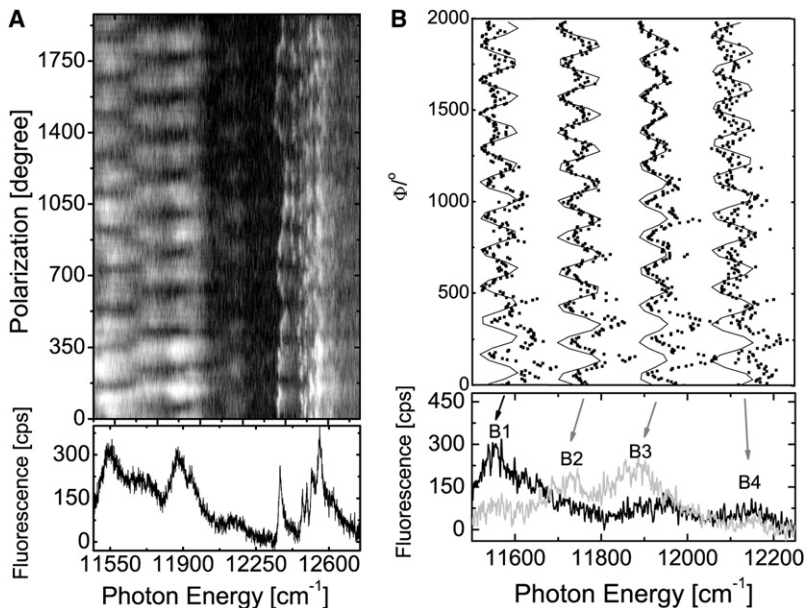
FIGURE 2 Fluorescence-excitation spectrum from an individual B800–820/850 LH2 complex from *Rps. palust-ris* as a function of the polarization of the excitation light. (*A*) (*Top*) Stack of 410 individual spectra recorded consecutively. Between two successive spectra, the polarization of the incident radiation has been rotated by 6.4°. The horizontal axis corresponds to the photon energy, the vertical axis corresponds to the scan number (or equivalently to the polarization angle), and the intensity is coded by the gray scale. The excitation intensity was 50 W cm^{-2}. (*Bottom*) Spectrum that corresponds to the average of the 410 consecutively recorded spectra. (*B*) (*Top*) Expanded view of the fluorescence intensity of the four bands (B1–B4) marked by the arrows in the lower part as a function of the polarization of the incident radiation (*dots*) together with cos^2-type functions (*black*) fitted to the data. (*Bottom*) Two fluorescence-excitation spectra from the stack that correspond to mutually orthogonal polarization of the excitation light.

absorptions in the B850 region. This becomes even more evident in Fig. 2 *B*. At the bottom of Fig. 2 *B*, two individual spectra are shown on an expanded energy scale. The spectra were obtained for mutually orthogonal polarizations of the excitation light. For simplicity, we refer in the following to the absorptions in the B820/850 region as B1, B2, B3, and B4, in order of increasing photon energy, where B stands for band. The angle of the polarization that yields the maximum intensity for the band B1 was set arbitrarily to 0° and provides the reference point. For this particular complex, we find the absorption bands B1, B2, B3, and B4 at the peak positions of 11,562 cm^{-1}, 11,727 cm^{-1}, 11,874 cm^{-1}, and 12,150 cm^{-1}, and the linewidths are 174 cm^{-1}, 125 cm^{-1}, 187 cm^{-1}, and 52 cm^{-1}, respectively. The top part of Fig. 2 *B* shows the fluorescence intensity of the four bands as a function of the polarization of the excitation light (*dots*). This variation is consistent with a cos^2 dependence (*black line*).

From such experiments, we can determine the energetic separation (ΔE) between B1 and the other bands, as well as the relative angle ($\Delta \alpha$) between the transition-dipole moments that are associated with them. The distributions of these parameters for the full data set of 28 complexes are shown in histograms (Fig. 3). The energetic separation between the bands B1 and B2 varies between 62 cm^{-1} and 342 cm^{-1} and is centered at ~174 cm^{-1} (Fig. 3 *A*). The distribution of the mutual orientation of the transition-dipole moments that are associated with the B1 and B2 bands increases to a maximum at ~90° (Fig. 3 *D*). The energetic separation between B1 and B3 is distributed between 230 cm^{-1} and 491 cm^{-1}, with a maximum at ~361 cm^{-1} (Fig. 3 *B*). The distribution of relative orientations of the transition-dipole moments that is associated with B1 and B3 decreases from a maximum value at ~10° when going

to larger angles (Fig. 3 *E*). The distribution of the energetic separation between B1 and B4 is between 430 cm^{-1} and 670 cm^{-1} and is centered at ~550 cm^{-1}, whereas the distribution of the relative orientations of the transition-dipole moments associated with B1 and B4 shows a slow decrease from a maximum value at ~20°.

DISCUSSION

When the single-molecule spectra obtained with HL B800–850 complexes from both *Rps. palustris* and *Rps. acidophila* are compared with those recorded from LL LH2 complexes from *Rps. palustris*, both common and strikingly different features can be observed. In the spectral region around 12,500 cm^{-1} (800 nm), all three types of complexes show narrow bands. These narrow bands have been well explained and arise from essentially monomeric weakly interacting Bchl molecules (B800 manifold) (32,38–40). The single-molecule spectra in the region of ~11,600–12,250 cm^{-1} (815–860 nm) differ greatly between the LL and HL complexes. The HL complexes show two and sometimes three broad bands in the 850 nm region. Previous studies have shown that these broad bands arise from excitonic interactions among the strongly coupled B850 Bchls (7,30,41).

All of the LL complexes studied here show broad bands not only in the 11,750 cm^{-1} (850 nm) region, but also in the 12,200 cm^{-1} (820 nm) region. They appear to have a mixed composition that gives rise to exciton bands in both the 850 and 820 nm regions. In an effort to explain these differences, we performed a series of simulations to try to understand the structural features that are required to accurately reproduce the experimental data.

The general approach used to describe the electronically excited states of the B850 LH2 ring is based on a model
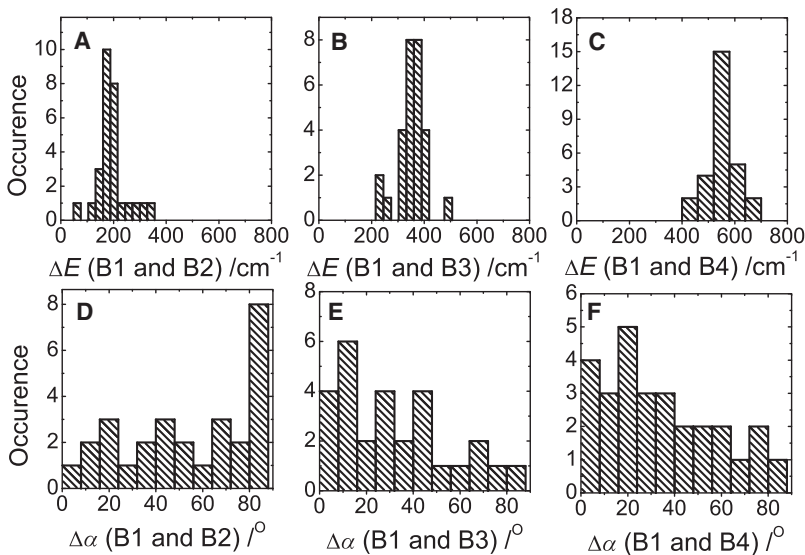
FIGURE 3 (*Top*) Distributions of the energetic separations ΔE observed between (*A*) B1 and B2, (*B*) B1 and B3, and (*C*) B1 and B4. (*Bottom*) Distributions of the mutual angles $\Delta\alpha$ between the transition-dipole moments that are associated with (*D*) B1 and B2, (*E*) B1 and B3, and (*F*) B1 and B4.

Hamiltonian using the Heitler-London approximation (41,42):

$$H = \sum_{n=1}^{N}(E_0 + \Delta E_n)|n\rangle\langle n| + \frac{1}{2}\sum_{n=1}^{N}\sum_{m\neq n}V_{nm}|n\rangle\langle m|, \quad (1)$$

where N refers to the number of Bchl a molecules; $|n\rangle$ and $|m\rangle$ correspond to excitations localized on molecules n and m, respectively; and $(E_0 + \Delta E_n)$ denotes the site energy of pigment n. Due to local variations in the protein environment of the binding site, there is a separation of the individual site energies into an average, E_0, and a deviation from this average, ΔE_n, which is commonly termed diagonal disorder. This energetic disorder is usually modeled by a Gaussian distribution of site energies. As the simplest approach to evaluate the dependence of the interaction on the distance and the mutual orientation of the pigments, we used a dipole-dipole type function (43):

$$V_{nm} = V_0 \frac{\kappa_{nm}}{r_{nm}^3}, \quad (2)$$

where κ_{nm} denotes the usual orientation factor, and r_{nm} is the distance between pigments n and m. The coupling strength V_0 was set to 135,072 cm^{-1} Å^3 (43). Two limiting cases for the dipolar interaction can be distinguished: weak coupling $|V/\Delta E| << 1$, and strong coupling $|V/\Delta E| >> 1$. In the case of the B850 Bchls, there is strong coupling due to much larger interaction energy (V_{nm}) compared to the difference in their site energies (ΔE_{nm}). As a result of this, the eigenfunctions of the Hamiltonian (Eq. 1) are given by the combinations of the excited-state wavefunctions of the individual pigments.

In the case of both the 9-mers (HL LH2s from *Rps. acidophila* 10050 and *Rps. palustris*) and the 8-mer (LH2 from *Rhsp. molischianum*), the energy scheme of the excited-state manifold of the B850 ring was constructed by combining symmetric or antisymmetric linear combinations of the localized wave functions of the $\alpha\beta$-dimers in such a way that they formed the basis for the irreducible representations of the C_9 and C_8 pure-rotational point groups, respectively (7). As previously described for the C_9-symmetry case, the symmetric and antisymmetric manifolds consist of one nondegenerate ($k = 0$) and four doubly degenerate ($k = \pm1, \pm2,..., \pm4$) exciton states (7). The corresponding C_8-symmetry case has the ($k = 0, \pm1, \pm2, \pm3, 4$) exciton states. To determine which exciton states are involved in giving rise to the measured bands B1, B2, B3, and B4, we calculated the absorption spectra by Monte Carlo simulations, treating the diagonal disorder as a random variable. This simulation was carried out for both 9-mer and 8-mer possibilities. Typical examples of the simulated spectra of individual realizations of the disorder are shown in Fig. 4. The first simulated spectrum of an LL LH2 complex was obtained by introducing diagonal disorder into the circular 9- and 8-mer models (Fig. 4, *A* and *D*). For simplicity, the site energies of B850 Bchl a dimers were set according to the previously determined site energies of LH2 from *Rps. acidophila* (E_0 (αB850) = 12,300 cm^{-1} and E_0 (βB850) = 12,060 cm^{-1}) (42). The random diagonal disorder is taken from a Gaussian distribution with a width of $\Gamma_{\text{intra}} = 160$ cm^{-1}, where Γ_{intra} stands for intracomplex heterogeneity (41). Within each complex, the 18 or 16 Bchl a molecules will show variations of their site energies with respect to their spectral means. In the case of a 9-mer ring, this simulation produces a sharp band from the $k = 0$ state, broad bands from the $k = \pm2$ states, and broad bands from the $k = \pm1$ states, which are split in energy and feature mutually orthogonal transition-dipole moments (Fig. 4 *A*). In the 8-mer ring, only a sharp $k = 0$ band and the two orthogonally polarized $k = \pm1$ bands are present (Fig. 4 *D*). In Fig. 4, *B* and *E*, this simulation was extended by inserting the expression $\Delta E_n = \Delta E_n^{(0)} + E_{\text{mod}}\cos[2\phi(n + 1/2)]$ for the site energies.
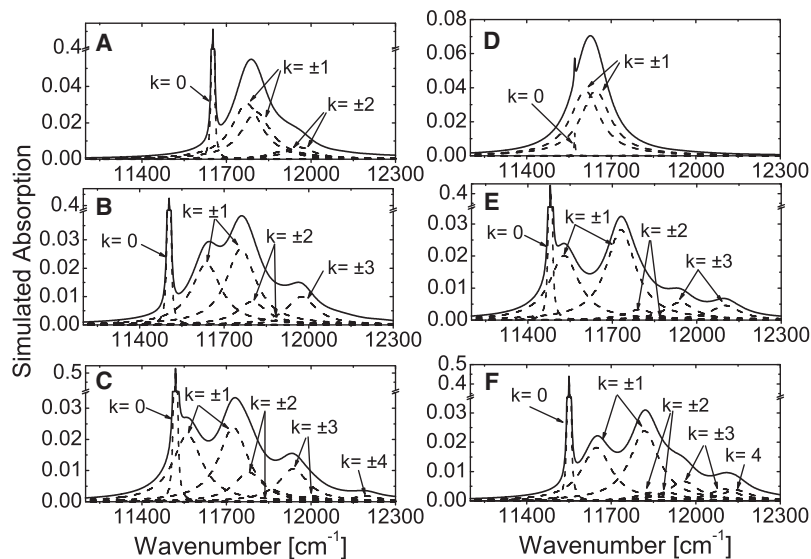
FIGURE 4 Examples for simulated absorption spectra for an individual LH2 complex (*solid line*). The spectra correspond to a single realization of the disorder. The calculated energies of the exciton states have been dressed by a Lorentzian with a width of 5 cm^{-1} for the $k = 0$ state and a width of 130 cm^{-1} for all other exciton states (*dashed lines*). Simulations A–C (*left*) are based on a nonamaric structure, and simulations D–F (*right*) are based on an octameric structure. The simulations vary with respect to the type of diagonal disorder: (A and D) random diagonal disorder taken from a Gaussian distribution of width 160 cm^{-1} (FWHM); (B and E) random diagonal disorder as in A and D, with an additionally correlated diagonal disorder; (C and F) like B and E, additionally using B820-like-pair ($E_0(\alpha B820) = 12,860$ cm^{-1}, $\Delta E = 260$ cm^{-1}) and B850-like-pair ($E_0(\alpha B850) = 12,300$ cm^{-1}, $\Delta E = 240$ cm^{-1}) randomly distributed site energies across the ring assembly. For more details see text.

Here $\Delta E_n^{(0)}$ refers to the random diagonal disorder chosen from the Gaussian distribution described above, and $E_{mod}\cos[2\phi(n + 1/2)]$ introduces an additional correlated modulation of C$_2$-type symmetry into the site energies of the pigments, where $\phi = 2\pi/N$ and N refers to the number of the pigments in the ring (42). When this modulation is taken into account, the 9-mer model shows a sharp $k = 0$ band, broad orthogonally polarized $k = \pm 1$ bands, and broad $k = \pm 2$ bands. Furthermore, although the $k = \pm 3$ states gain significant oscillator strength, but they have no observable splitting. Though the $k = \pm 2$ bands do contribute, it is apparent that the overall spectrum of the 9-mer model is dominated by the contributions of the $k = 0$, $k = \pm 1$, and $k = \pm 3$ states. In the case of the 8-mer model, a sharp $k = 0$ band, broad orthogonally polarized $k = \pm 1$ bands, broad $k = \pm 2$ bands, and $k = \pm 3$ bands are also present. The splitting of the $k = \pm 3$ bands is clear in the 8-mer model. The overall spectrum of the 8-mer model is dominated by the $k = 0$, $k = \pm 1$, and $k = \pm 3$ states.

There is strong evidence from mass spectroscopic analysis of LL LH2 polypeptides from *Rps. palustris* that this LL complex contains multiple types of $\alpha\beta$-polypeptides (T. H. P. Brotosudarmo, A. Gall, V. Moulisova, A. T. Gardiner, and R. J. Cogdell, unpublished results). The sequence of some of these polypeptides features changes for key amino acids shown to be involved in H-bonding to the Bchl a macrocycles of the strongly coupled ring of Bchls in LH2. This heterogeneity raises the possibility that some of these Bchl a molecules may have quite different site energies. In other words, some may have site energies characteristic of B850 and some may have ones characteristic of B820. Therefore, the simulation was extended to introduce two different site energies—E_0 (αB850) $= 12,300$ cm^{-1}, E_0 (βB850) $= 12,060$ cm^{-1} (B850-pair) (42); and E_0 (αB820) $= 12,860$ cm^{-1}, E_0 (βB820) $= 12,600$ cm^{-1}) (B820-pair) (44)—for the Bchl a-pairs in which the B850- and B820-pairs of

Bchl a are randomly distributed in the ring (Fig. 4, C and F). In this case the 9-mer model shows the gain of the oscillator strength of all of the exciton states (Fig. 4 C). The bands at 11500 cm^{-1} (870 nm) and 11700 cm^{-1} (855 nm) are due to the $k = +1$ and $k = -1$ states, respectively. The $k = \pm 2$ bands are obscured under the $k = -1$ band. Although the splitting of the $k = \pm 3$ and $k = \pm 4$ states are seen, the $k = +3$ and $k = +4$ bands are the most intense. The overall features of the simulated spectrum (*solid line*) from the 9-mer model are dominated by the $k = 0$, ± 1, $+3$ and $+4$ states (*dashed lines*) and is blue-shifted compared to the case shown in Fig. 4 B. In the case of the 8-mer model, the simulated spectrum is identical to that of the 9-mer (Fig. 4 F).

When the absorption spectra of the simulations are compared with the experimental spectrum of the LL LH2 (Fig. 2), it can be seen that the traces in Fig. 4, A, B, and D, are not consistent with the experimental data. The simulations presented in Fig. 4, C, E, and F, however, all are in reasonable agreement with the experimental spectrum. Simulations with a 9-mer model can only account for the experimental spectrum if two different site energies for Bchl a-pairs are considered (Fig. 4 C). If this model describes the situation properly, then the experimentally measured bands B1, B2, B3, and B4 can be assigned to the $k = +1, -1, +3$, and $+4$ states, respectively. Simulations with an 8-mer model produce agreement with the experimental data both when different site energies for the pairs of Bchl a molecules are taken into account and when they are not. If the model in Fig. 4 E provides a proper description of the experimentally determined situation, then the B1, B2, B3, and B4 bands in the experimental data can be assigned to the $k = +1, -1, -3$, and $+3$ states, respectively. If, on the other hand, the model that corresponds to the simulation shown in Fig. 4 F is appropriate, then the B1, B2, B3, and B4 bands can be assigned to the $k = +1, -1, +3$ and 4 states,
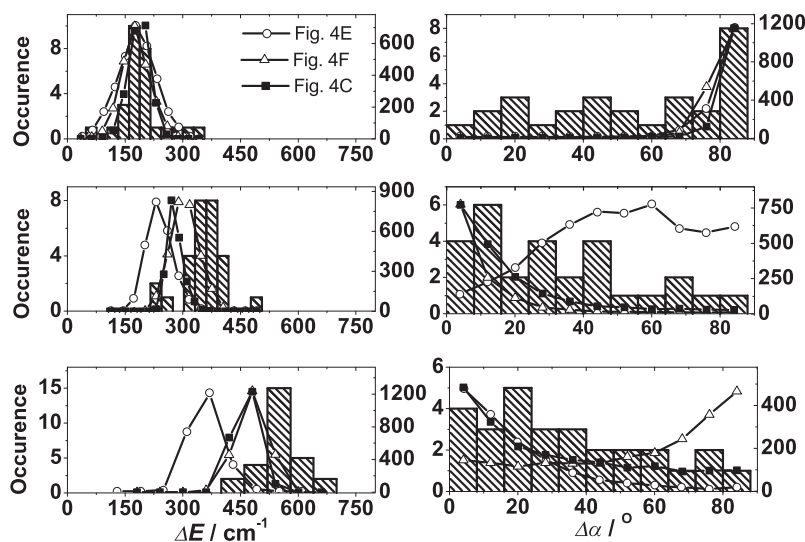
FIGURE 5 Comparison of the energetic separations of bands B1–B4 with the separations of the respective exciton states (*left*) and the mutual orientation of the associated transition-dipole moments (*right*) as predicted from Monte Carlo simulations for 2000 realizations of the disorder for the models shown in Fig. 4 *E* (*open circles*), Fig. 4 *F* (*open triangles*), and Fig. 4 *C* (*solid squares*).

respectively. Therefore, it is unclear at this stage which of the underlying models that led to the simulations presented in Fig. 4 represents a realistic approximation for describing the structure of the LL LH2.

To distinguish among the three different options presented in Fig. 4, *C*, *E*, and *F*, we considered the extra experimental spectroscopic information that is available (see Fig. 3), i.e., the energetic separation of the spectral bands and the mutual orientation of their transition-dipole moments. Therefore, we compared the experimental histograms with the parameters calculated from the respective simulations for 2000 realizations of the energetic disorder (Fig. 5, *circles*, *triangles*, and *black squares*). Using the 8-mer model represented by Fig. 4 *E*, it can be seen that the simulated distributions of ΔE and the mutual angle between the transition-dipole moments of the $k = \pm 1$ states can be reproduced; however, the simulated distributions of the energetic separations between $k = +1$ and $k = -3$, as well as between $k = +1$ and $k = +3$, with maxima around 200 cm^{-1} and 400 cm^{-1} are not (Fig. 5). Moreover, the simulated distribution for the mutual angle between the transition-dipole moments of the $k = +1$ and $k = -3$ states is even worse, with a broad maximum covering the range from 40° to 60°. Simulations based on the 8-mer model represented by Fig. 4 *F* are able to satisfactorily account for the distribution of the simulated mutual angles between the transition-dipole moments of the $k = \pm 1$ states (Fig. 5). However, these simulations also do not satisfactorily reproduce the distributions of the angles between the transition-dipole moments of the $k = +1$ and $k = 4$ states. Simulations based on the 9-mer model represented by Fig. 4 *C* show quite good agreement with the distribution of all three mutual angles (Fig. 5). However, there is a significant though smaller (compared to the two 8-mer models) mismatch between this simulation and the distribution of energetic separations, ΔE ($k = +1$ and $k = +3$) and ΔE ($k = +1$ and $k = +4$).

At this stage, it appears that the 9-mer model is able to more closely account for the experimental data. The LL LH2 complexes from *Rps. palustris* have at least four types of α-polypeptides (T. H. P. Brotosudarmo, A. Gall, V. Moulisova, A. T. Gardiner, and R. J. Cogdell, unpublished results). One, the α_{d}-polypeptide, has phenylalanine and methionine residues at positions 44 and 45 (25,45). The other polypeptides have tyrosine and tryptophan residues at these positions. The presence of H-bonding residues at positions 44 and 45, such as tyrosine and tryptophan, correlates with B850-type site energies, whereas if non-H-bonding residues, such as phenylalanine and methionine, are present, it correlates with B820-like site energies. Shifting in the site energy of one α-bound Bchl *a* will also affect the nearest-neighbor β-bound Bchl *a*. Therefore, we extended the simulations using this additional information. The parameters of the coordinate arrangement of the Bchls for this simulation were taken from the crystal structures of the B800–850 and B800–820 LH2 complexes (see Table S1 and Fig. S2). The site energies of the α-bound-B820-like Bchl *a* molecules E_0 (αB820) was set to be 12,640 cm^{-1}, whereas the β-bound-B820-like Bchl *a* molecule E_0 (βB820) was 12,400 cm^{-1}. The position of the B820-like pairs was randomly distributed (Fig. 6). The simulation based on this model, though not perfect, shows closer agreement with the experimental distribution of energetic separations ΔE ($k = +1$ and $k = +3$). Again, the match to the distributions of the mutual angles of the respective transition-dipole moments between ($k = +1$ and $k = +3$) and ($k = +1$ and $k = +4$) is good. We extended this simulation further by systematically setting fewer B820-like site energy pairs. However, none of these simulations yielded an improved reproduction of the experimental data (data not shown).

Next, the effect of a symmetrical arrangement of these B820-like site energies was tested. Six B820-like site energies were distributed symmetrically with respect to the
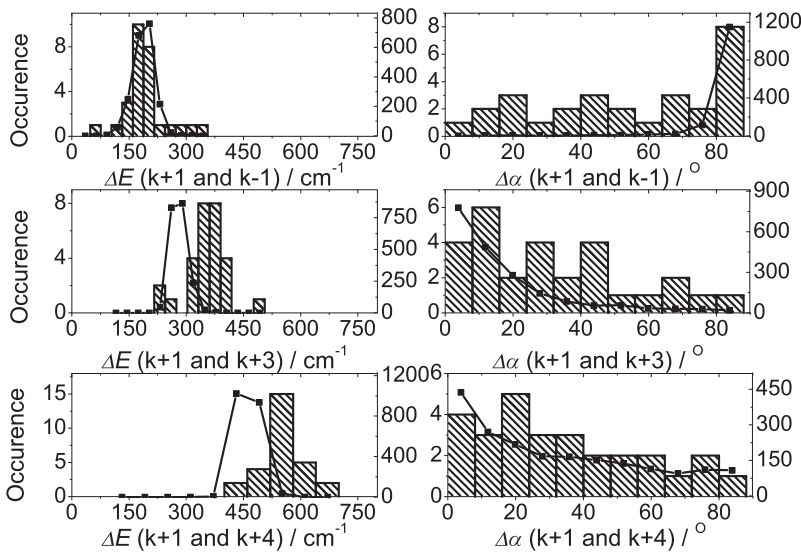
FIGURE 6 Extended simulations based on the model shown in Fig. 4 *C*, i.e., a nonameric structure with random and correlated diagonal disorder and multipolypeptide composition. The B820-like-pair site energies of $E_0(\alpha B820) = 12,640$ cm^{-1} with $\Delta E = 250$ cm^{-1} and the B850-pair site energy of $E_0(\alpha B850) = 12,300$ cm^{-1} with $\Delta E = 240$ cm^{-1}. The positions of the B820-like pairs are randomly distributed.

threefold axis in the ring. The other 12 Bchl *a* molecules were given B850-site energies. This simulation yielded the best agreement between the simulations and the experimental data that we found (Fig. 7), but only when we introduced a rather high intracomplex heterogeneity, Γ_{intra},

at 320 cm^{-1}. This value is significant higher than that previously reported for the B800–850 LH2 of *Rps. acidophila* (Γ_{intra} = 250 cm^{-1}) (41,42). This may be a consequence of having individual rings with a heterogeneous polypeptide composition.
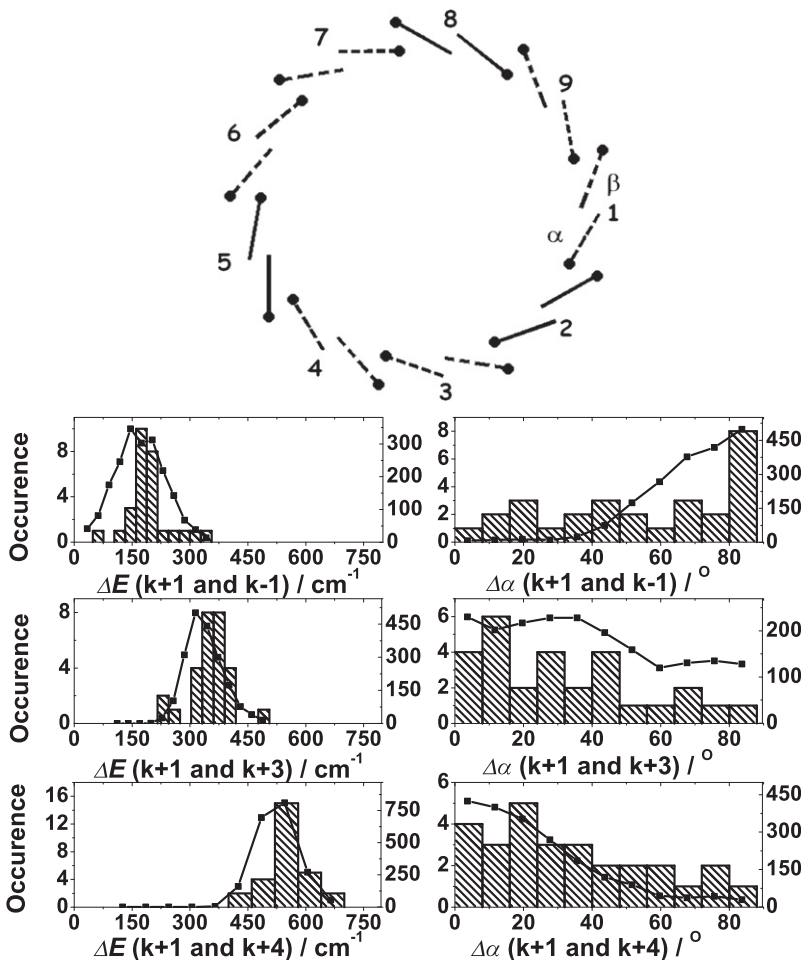

FIGURE 7 (*Top*) Model structure featuring six 820-like-pair Bchl a molecules (*dashed lines*) distributed in C$_3$ symmetry around the ring. (*Bottom*) Comparison of the experimental data with the results from Monte Carlo simulations (2000 realizations) for the nonameric structure as shown in the top part, taking random and correlated diagonal disorder into account (*black squares*). The site energies of the pigments were chosen as detailed in Fig. 6. The intracomplex heterogeneity (Γ_{intra}) is 320 cm^{-1}. The full and dashed lines in the top part of the figure indicate the B820-like (*dashed*) and B850 (*full*) site energies.

It has been suggested that some species of purple bacteria may contain LH2 complexes in which rings of multiple sizes coexist (3). To test whether various mixtures of rings of different sizes could satisfactorily account for our experimental data, we added together simulations for the best 8-mer, 9-mer, and 10-mer models in a number of different combinations. None of these ''addition'' simulations produced satisfactory agreement with the experimental data (data not shown). Though it is not possible to claim that our simulations can provide definitive proof of the structure of the LL LH2 complex from *Rps. palustris*, they do strongly favor both a nonameric model and one in which the assembly has multiple apoprotein types within a single LH ring. It is also worth pointing out that the absorption spectrum of the native LL LH2 complex from *Rps. palustris* studied in this work is not the same as that of the B800-only LH2 complex used in the structural studies mentioned above (46). We do not understand why this is so, but we have never been able to isolate an LL LH2 complex from *Rps. palustris* that has the spectrum described by Hartigan et al. (24). Photosynthetic bacteria such as *Rps. palustris* have a very complicated set of control mechanisms that can modify which of the multigene family of LH2 $\alpha\beta$-gene pairs are expressed (25,26,45). These differences, therefore, may reflect variations in the growth conditions and laboratories in which the bacteria were cultured.

CONCLUSIONS

The single-molecule spectra reported here can only be satisfactorily explained by assuming that the individual LL LH2 complexes from *Rps. palustris* contain a heterogeneous polypeptide composition. Multiple types of $\alpha\beta$ apoproteins coexist in the same individual complex, producing individual rings, with Bchl molecules having both B820-like and B850-site energies. The simulations described above strongly reinforce this conclusion. The experimental data can best be modeled by taking an arrangement of the Bchl *a* molecules in the mixed B820/850 ring of nine $\alpha\beta$ Bchls dimers, where in six $\alpha\beta$-pairs the Bchl *a* molecules have B820-like site energies and are distributed around the ring in such a way that the symmetry of the nonameric assembly is reduced to C_3. The high intracomplex disorder reflects the heterogeneity in the polypeptide composition.

SUPPORTING MATERIAL

A table and two figures are available at http://www.biophysj.org/biophysj/supplemental/S0006-3495(09)01205-3.

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