

# Effect of Leucine M196 Substitution by Histidine on Electronic Structure of the Primary Electron Donor and Electron Transfer in Reaction Centers from *Rhodobacter sphaeroides*

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Received November 13, 2018

Revised December 27, 2018

Accepted December 27, 2018

**Abstract**—In our recent X-ray study, we demonstrated that substitution of the natural leucine residue M196 with histidine in the reaction center (RC) from *Rhodobacter (Rba.) sphaeroides* leads to formation of a close contact between the genetically introduced histidine and the primary electron donor P (bacteriochlorophylls (BChls) P<sub>A</sub> and P<sub>B</sub> dimer) creating a novel pigment–protein interaction that is not observed in native RCs. In the present work, the possible nature of this novel interaction and its effects on the electronic properties of P and the photochemical charge separation in isolated mutant RCs L(M196)H are investigated at room temperature using steady-state absorption spectroscopy, light-induced difference FTIR spectroscopy, and femtosecond transient absorption spectroscopy. The results are compared with the data obtained for the RCs from *Rba. sphaeroides* pseudo-wild type strain. It is shown that the L(M196)H mutation results in a decrease in intensity and broadening of the long-wavelength Q<sub>y</sub> absorption band of P at ~865 nm. Due to the mutation, there is also weakening of the electronic coupling between BChls in the radical cation P<sup>+</sup> and increase in the positive charge localization on the P<sub>A</sub> molecule. Despite the significant perturbations of the electronic structure of P, the mutant RCs retain high electron transfer rates and quantum yield of the P<sup>+</sup>Q<sub>A</sub><sup>-</sup> state (Q<sub>A</sub> is the primary quinone acceptor), which is close to the one observed in the native RCs. Comparison of our results with the literature data suggests that the imidazole group of histidine M196 forms a  $\pi$ -hydrogen bond with the  $\pi$ -electron system of the P<sub>B</sub> molecule in the P dimer. It is likely that the specific (T-shaped) spatial organization of the  $\pi$ -hydrogen interaction and its potential heterogeneity in relation to the bonding energy is, at least partially, the reason that this type of interaction between the protein and the pigment and quinone cofactors is not realized in the native RCs.

DOI: 10.1134/S0006297919050067

**Keywords:** bacterial reaction center, amino acid replacement, primary electron donor, electronic structure, electron transfer, *Rhodobacter sphaeroides*

Bacterial reaction centers (RCs) are highly specialized cofactor–protein complexes, in which the primary energy conversion of light quanta into the electrochemical energy of separated charges takes place via a series of fast electron transfer reactions. The unique feature of RCs

as natural photoconversion systems is their ability to perform charge separation with extremely high quantum yield (~100%) and high energy efficiency (tens of percent) in a molecular donor–acceptor system involving only a few pigment molecules (bacteriochlorins) embed-

**Abbreviations:**  $\Delta A$ , absorbance change; B<sub>A</sub>, monomeric bacteriochlorophyll in the active cofactors branch; BChl, bacteriochlorophyll; BPheo, bacteriopheophytin; EADS, evolution-associated decay spectra; FTIR, Fourier transform infrared spectroscopy; H<sub>A</sub> and H<sub>B</sub>, BPheo molecules in the active and inactive cofactor branches, respectively; P, primary electron donor, BChl dimer; P<sub>A</sub> and P<sub>B</sub>, BChl molecules constituting P; psWt, pseudo-wild type; Q<sub>A</sub>, primary quinone acceptor; Q<sub>B</sub>, secondary quinone acceptor; *Rba. sphaeroides*, *Rhodobacter sphaeroides*; RC, reaction center.

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ded into a specifically organized protein matrix. Obviously, the key factors for the function of this system are both basic physicochemical properties of individual pigment molecules and their interaction between themselves and with the local environment. One of the approaches to investigation of the role of intermolecular interactions in the structural and electronic configuration of pigment cofactors and their functionality is introduction of point amino acid substitutions to the RC protein in order to eliminate the existing and/or create new contacts/interactions and detailed investigation of the consequences of such mutation with X-ray diffraction analysis in combination with the methods of optical spectroscopy. In particular, one of the promising substitutions affecting intermolecular interactions in the RC is the replacement of natural amino acid residues with a histidine residue. Histidine in proteins is capable of participating in interactions of various nature: coordination bonds with metal cations, hydrogen bonds, cation- $\pi$ ,  $\pi$ -hydrogen, and  $\pi$ - $\pi$ -stacking interactions [1].

It was recently established based on the results of X-ray diffraction analysis in combination with the results of low-temperature absorption measurements and redox titration that the substitution of leucine by histidine at position M196 in the RCs from *Rhodobacter (Rba.) sphaeroides* likely resulted in formation of a new interaction between the bacteriochlorophyll (BChl) P<sub>B</sub> of the primary electron donor P and imidazole of the newly introduced histidine M196 [2]. It was suggested that such interaction could be either of  $\pi$ - $\pi$ -stacking interaction or  $\pi$ -hydrogen interaction [2].

To further elucidate the nature of the discovered interaction and its effect on the electronic properties of P and primary charge separation, in this study the isolated L(M196)H RCs were investigated with the methods of steady-state absorption spectroscopy in the near- and mid-IR (Fourier transform infrared spectroscopy; FTIR) ranges and transient absorption spectroscopy with femtosecond time resolution in the visible and near-IR ranges of the spectrum. Recording of electronic absorption spectra is widely used to obtain information on the effect of mutation on electronic properties of bacterial RCs. Light-induced FTIR difference spectroscopy demonstrates extremely high sensitivity to molecular changes accompanying charge separation providing, in particular, information on the electronic structure of the primary electron donor P in its functionally important radical cation form P<sup>+</sup> [3, 4]. Femtosecond transient absorption spectroscopy is the most appropriate method for investigation of fast electron transfer reactions occurring in bacterial RCs in the sub-picosecond time scale. The results of this work are discussed based on the analysis of key similarities and differences with the data obtained under the identical experimental conditions for the RCs from the pseudo-wild type of *Rba. sphaeroides* (psWt).

## MATERIALS AND METHODS

Construction of a recombinant *Rba. sphaeroides* strain without light-harvesting antenna but containing mutant RCs with amino acid substitution of leucine in position M196 by histidine was described previously [5]. Reaction centers isolated from the *puf*-deficient strain transformed with a plasmid containing a wild type *puf*-operon were used as psWt RCs [5]. RCs were isolated using the standard procedure by treating chromatophores with lauryldimethylamine-N-oxide (LDAO) followed by purification on DEAE-cellulose [5]. The LDAO detergent was replaced with Triton X-100 on a membrane with molecular weight cut-off of 30 kDa (Millipore, USA) in an ultrafiltration cell under applied gaseous argon pressure.

Electronic absorption spectra of RCs in the range 650–1000 nm were recorded with a Shimadzu UV1800 spectrophotometer (Shimadzu, Japan).

Light-induced FTIR P<sup>+</sup>Q<sup>-</sup>/PQ difference spectra were recorded with an Equinox 55 FTIR spectrometer (Bruker, Germany) equipped with a DTGS detector and KBr beam splitter. Samples were obtained by applying a few microliters of the concentrated suspension of RCs solubilized in 20 mM Tris-HCl buffer (pH 8.0) containing 0.1% Triton X-100 onto a CaF<sub>2</sub> plate followed by partial dehydration of the sample under a gaseous argon flow and covering it with a second CaF<sub>2</sub> plate [6]. IR difference spectra were recorded under illumination of samples with constant light (1000 nm >  $\lambda$  > 620 nm). The spectral resolution was 4 cm<sup>-1</sup>. Illumination cycles were repeated hundreds of times to achieve acceptable level of signal-to-noise ratio.

Difference spectra with femtosecond time resolution were recorded using a laser spectrometer as described previously [7] with some modifications. A MaiTai titanium-sapphire laser (Spectra-Physics, USA) with a Spitfire Ace optical regenerative amplifier (Spectra-Physics) was used for generation of fundamental (800 nm) pulses with duration of 35–40 fs and energy of 3 mJ/pulse. The beam coming out of the amplifier was attenuated and split into two light beams. The first beam was used for generation of a continuum on a sapphire plate with thickness of ~2 mm. The continuum spectral components with wavelength >850 nm were separated by an RG850 light filter (Newport, USA), and the generated pulses were used for excitation of the RCs in the P absorption band. The second beam was passed over a M531.DD delay line (Physik Instrumente, Germany) and focused in a quartz cuvette with 5-mm thickness filled with water for generation of the continuum, which next was split into probing and reference beams at percent ratio 50 : 50. The spectra of probing and reference pulses were modified with spectral and spatial filtration to separate visible or near IR spectral region. Polarization of the excitation pulse was set parallel to the probing pulse. The probing and reference pulses

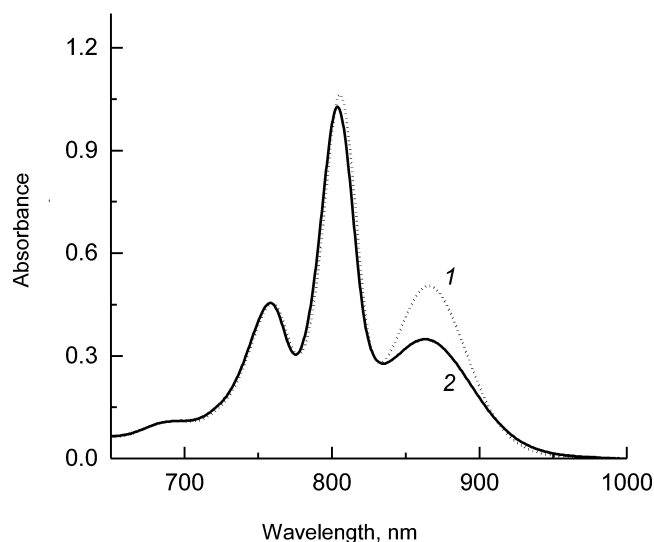
after the cuvette with the sample were directed to the input slit of a SpectraPro 2300i spectrograph (Acton Research Corporation, USA). The spectra of pulses were recorded with a Pixis 400BR CCD camera (Princeton Instruments, USA). The RC sample ( $A_{865} \sim 0.5$ ) was placed in a rotating quartz cuvette with optical pathlength of 1 mm (CDP Systems, Russia). Rotation speed and excitation frequency (10 Hz) were selected in a way so each excitation pulse would illuminate a new portion of RCs. Difference spectra were recorded in the visible (500–720 nm) or near IR (750–1000 nm) region in the time interval between 2 ps before excitation of the RCs and  $\sim 2$  ns after it; 500 spectra were averaged for each delay. The intensity of excitation pulses was attenuated to the level that provided excitation of  $<20\%$  of reaction centers in the sample.

All measurements were conducted at room temperature.

The femtosecond data were analyzed with global analysis [8] using the Glotaran program [9]. The program included mathematical description of spectral dispersion of the maximum of the instrumental function. The half width at half maximum of the instrumental function was estimated as  $\sim 35$ – $40$  fs for both investigated spectral regions.

## RESULTS

The electronic absorption spectra of RCs from *Rba. sphaeroides* psWt and L(M196)H are presented in Fig. 1; they were recorded in the region of the  $Q_y$  optical transi-

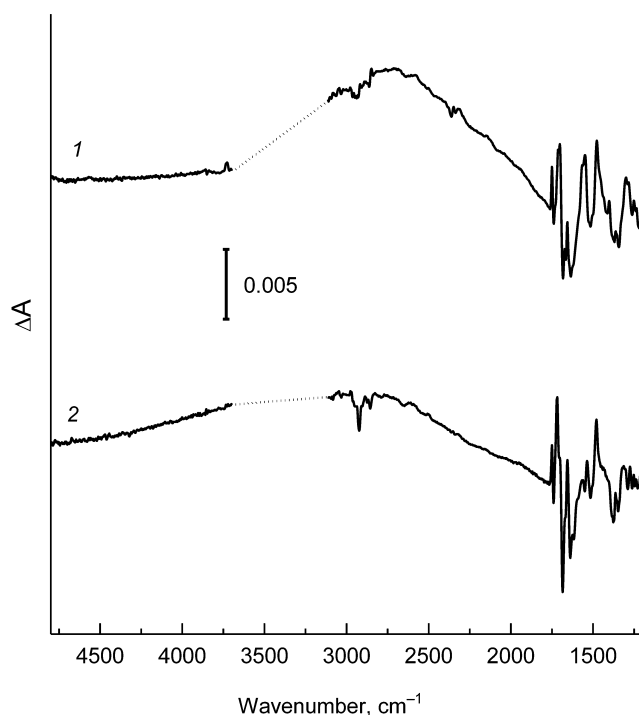


**Fig. 1.** Electronic absorption spectra of *Rba. sphaeroides* psWt RCs (1) and L(M196)H RCs (2) recorded at room temperature. The spectra are normalized at the absorption maximum of bacterio-*phycocyanin* molecules at 759 nm.

tions (650–1000 nm) at room temperature. The spectra are normalized at the maximum of the bacterio-*phycocyanin* (BPheo) band at 759 nm based on the suggestion that the mutation does not introduce significant perturbations to the electronic structure of the BPheo molecules in the RCs and does not affect significantly their optical properties as the mutation locus is located far from the binding sites of both BPheo  $H_A$  and BPheo  $H_B$ .

Figure 1 shows that the absorption band of the low-energy  $Q_y$  exciton transition of the primary electron donor P at  $\sim 865$  nm is preserved in the mutant RCs, although it is slightly broadened and significantly lower in intensity compared to the respective band in the psWt RCs. Control experiments with the addition of reductants to the samples of mutant RCs (sodium ascorbate or sodium dithionite) revealed that the observed decrease in intensity of the P band at  $\sim 865$  nm was not due to the oxidation of P (data not shown). Hence, despite the fact that the general dimeric nature of P is maintained in the mutant RCs, the electronic structure of the dimer is likely subjected to substantial modification. The minor short wavelength shifts of the absorption bands of the monomeric BChl molecules at  $\sim 805$  nm and of the BPheo molecules at  $\sim 760$  nm are also characteristic for the mutant (Fig. 1). The decrease in intensity of the  $Q_y$ -band of P and its spectral broadening in the L(M196)H RCs in comparison with the psWt RCs was observed previously in the low-temperature (90 K) absorption spectrum [2]. It is interesting that in this case maximum of the  $Q_y$  absorption band of P in the L(M196)H RC spectrum demonstrated larger long-wavelength shift than in the spectrum of the psWt RCs (894 versus 888 nm). It is well known that the spectral position of the long wavelength  $Q_y$ -band of P depends on a number of factors including, in particular, formation of hydrogen bonds [10] and change of the energy of the charge-transfer states between the molecules  $P_A$  and  $P_B$  in the P dimer [11].

The light-induced FTIR difference spectra  $P^+Q^-/PQ$  (Q is a quinone acceptor) for reaction centers from psWt and L(M196)H are presented in Figs. 2 and 3. The positive and negative peaks in the spectra reflect an appearance of the state  $P^+Q^-$  and a disappearance of the state PQ, respectively. The FTIR spectra of psWt and L(M196)H (Figs. 2 and 3) were normalized to the amplitude of the differential signal  $(+)/(-)1750/(-)1740$   $\text{cm}^{-1}$  attributed to the shift of stretching vibrations of the  $13^3$ -ester groups of the BChl [6, 12]. In agreement with the data published previously [6, 12], the FTIR spectrum  $P^+Q^-/PQ$  for the psWt RCs (Fig. 2, curve 1; Fig. 3, curve 1) demonstrates a broad band of the low-energy electronic transition at  $\sim 2700$   $\text{cm}^{-1}$  related to the transfer of the positive charge (hole) within the dimeric  $P^+$  radical cation and three phase-phonon IR bands at  $\sim 1550$ ,  $1477$ , and  $\sim 1290$   $\text{cm}^{-1}$ . The band at  $\sim 2700$   $\text{cm}^{-1}$  is not observed for the monomeric BChl $^+$  radical cation [12], which makes it a unique characteristic of the dimeric structure



**Fig. 2.** Light-induced difference (light-minus-dark) FTIR spectra  $P^+Q^-/PQ$  for *Rba. sphaeroides* psWt RCs (1) and L(M196)H RCs (2) recorded at room temperature in the range 4800–1200  $\text{cm}^{-1}$ . The spectra are normalized at the amplitude of the differential signal (+)1750/(–)1740  $\text{cm}^{-1}$ . Region  $\sim 3700$ –3100  $\text{cm}^{-1}$  is saturated due to the strong absorption of the sample and water and is excluded from the figure.

of the oxidized primary electron donor in the RCs of purple bacteria [4]. Vibrational modes of phase-phonon type are formally symmetry-forbidden in the monomeric BChl; however, they are amplified due to coupling with the hole-transfer transition, which make them also markers of the dimeric structure of  $P^+$  [13]. The intensity of all indicated bands depends significantly on the degree of electronic coupling between the  $P_A$  and  $P_B$  molecules in the  $P$  dimer [4, 13].

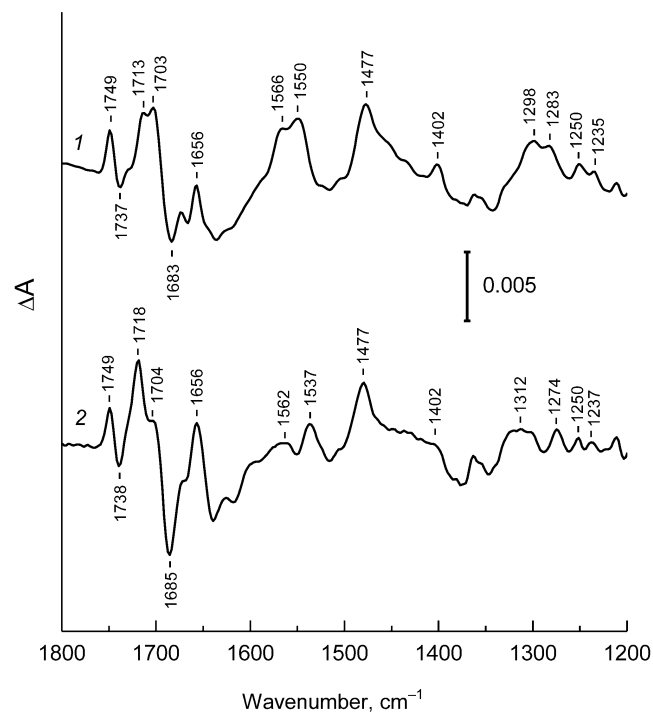
The FTIR  $P^+Q^-/PQ$  spectrum for the L(M196)H RCs also contains the hole-transfer band in the region of  $\sim 2700$   $\text{cm}^{-1}$  (Fig. 2, curve 2) and phase-phonon bands at 1537, 1477, and  $\sim 1300$   $\text{cm}^{-1}$  (Fig. 3, curve 2), which is in agreement with preservation of the dimeric structure of the electron donor  $P$  in the mutant [2] (Fig. 1). However, the pronounced decrease in intensity of the hole-transfer band and large changes in the intensity and frequencies of the phase-phonon bands in the spectrum of the mutant RCs indicate reduced electronic coupling and enhanced asymmetry in distribution of the positive charge between the molecules of BChl in the  $P^+$  radical cation [13].

The signals predominately related to the shifts of  $13^1$ -keto-C=O-groups in the  $P_A$  and  $P_B$  molecules upon  $P$  photooxidation are observed in the region 1710–

1650  $\text{cm}^{-1}$  of the IR spectrum of the psWt RCs (Fig. 3, curve 1) [12]. These groups are conjugated with the  $\pi$ -electron system of the macrocycle and are sensitive to changes in the electronic structure of the  $P^+$  dimer. The relative increase in intensity of the stretching vibrations of the  $13^1$ -keto-group of  $P_A^+$  at 1718  $\text{cm}^{-1}$  in comparison with the similar band of the  $13^1$ -keto-group of  $P_B^+$  at 1704  $\text{cm}^{-1}$  in the spectrum of the mutant L(M196)H RCs (Fig. 3, curve 2) can be interpreted as an increase in localization of the positive charge on the  $P_A$  molecule [14]. Redistribution of the positive charge towards  $P_A^+$  due to the wakening of electronic coupling between molecules  $P_A$  and  $P_B$  is likely due to the interaction of the introduced histidine M196 with the  $\pi$ -electron system of  $P$ .

The results of investigation of the dynamics and mechanism of charge separation in the mutant RCs provided by femtosecond time-resolved difference absorption spectroscopy are presented in Figs. 4–6. The data were analyzed using global analysis [8].

The kinetics of the stimulated emission decay from the excited state of the primary donor  $P^*$  reflecting the electron transfer from  $P^*$  to BPheo  $H_A$  in the psWt RCs (curve 1) and mutant RCs (curve 2) are compared in Fig. 4a. The monoexponential analysis of the kinetic traces revealed that the lifetime of  $P^*$  (that is the time constant of the reaction  $P^* \rightarrow P^+H_A^-$ ) in the psWt RCs is 3.2 ps, which is in good agreement with the literature data for the wild type RCs. The lifetime of  $P^*$  in the

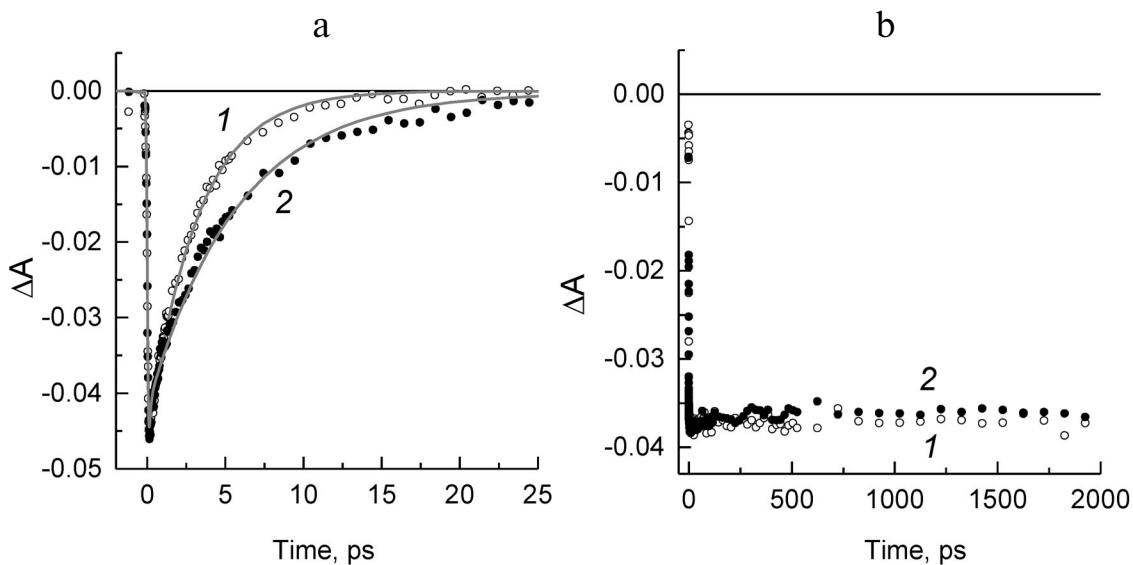


**Fig. 3.** Low-frequency region (1800–1200  $\text{cm}^{-1}$ ) of the FTIR spectra  $P^+Q^-/PQ$  for *Rba. sphaeroides* psWt RCs (1) and L(M196)H RCs (2), taken from Fig. 2.

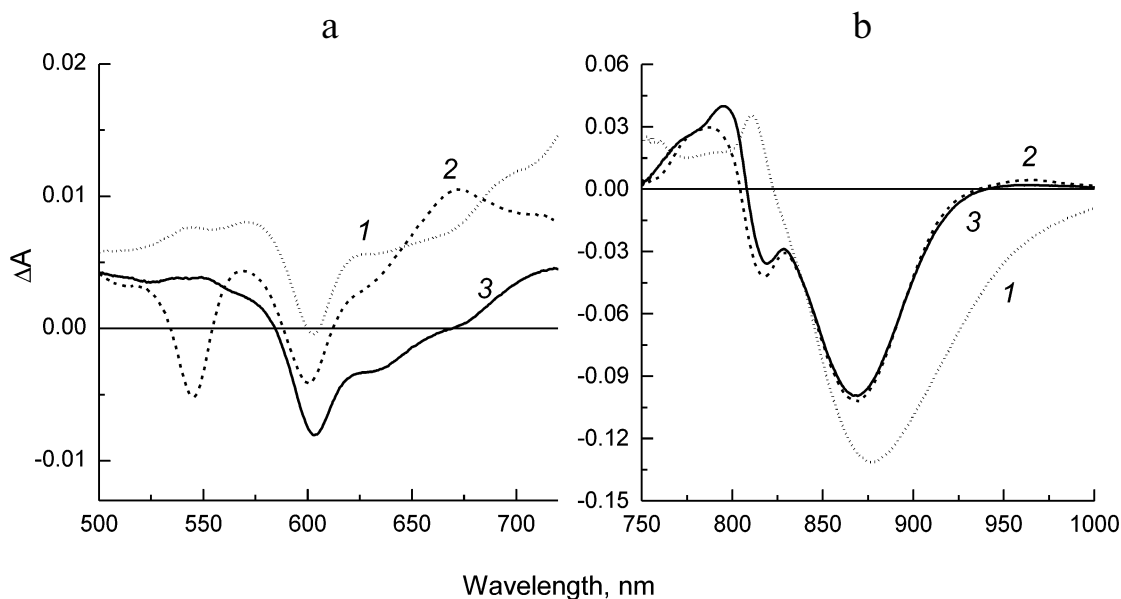
L(M196)H RCs increases to 5.7 ps. Investigation of the bleaching kinetics on the short-wavelength side of the P absorption band at ~830–840 nm (region, where contribution of the stimulated emission is minimal) provides information on the quantum yield of the charge separation in the RCs. It can be seen in Fig. 4b (curve 1) that the amplitude of bleaching in the psWt RCs remains practically unchanged in the time interval of 2 ns. This implies

that the charge separation is not accompanied by the return of  $P^*$  and  $P^+$  to the ground state, and that the quantum yield of the  $P^+Q_A^-$  state is close to 100%. Comparison of the kinetics 1 and 2 in Fig. 4b reveals that within the accuracy of our measurements a similar conclusion can be made for the L(M196)H RCs.

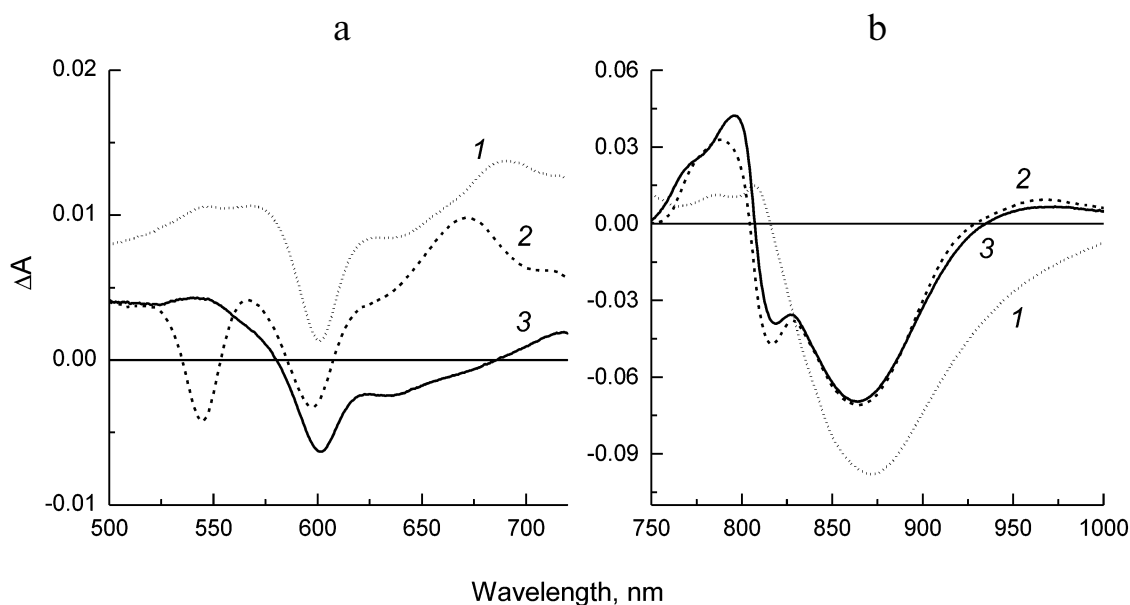
The difference spectra associated with the evolution of the states involved in the electron transfer from  $P^*$  to



**Fig. 4.** a) Normalized decay kinetics of the stimulated emission from the  $P^*$  excited state at 936 nm for the *Rba. sphaeroides* psWt RCs (1) and at 930 nm for the L(M196)H RCs (2). Monoexponential approximation of the kinetics is presented by solid lines. b) Normalized kinetics of absorption change at 836 nm for the psWt RCs (1) and at 830 nm for the L(M196)H RCs (2).



**Fig. 5.** Difference spectra associated with evolution of  $P^*$  (1),  $P^+H_A$  (2), and  $P^+Q_A^-$  (3) states obtained from global analysis of the spectral–temporal data in the  $Q_A$  (a) and  $Q_B$  (b) regions for the *Rba. sphaeroides* psWt RCs. Lifetimes of the states  $P^*$  and  $P^+H_A$  are 3.2 and 190 ps, respectively. The state  $P^+Q_A^-$  does not decay within the investigated time scale (2 ns).



**Fig. 6.** Difference spectra associated with evolution of  $P^*$  (1),  $P^+H_A^-$  (2), and  $P^+Q_A^-$  (3) states obtained from global analysis of the spectral–temporal data in the  $Q_x$  (a) and  $Q_y$  (b) regions for the *Rba. sphaeroides* L(M196)H RCs. Lifetimes of the states  $P^*$  and  $P^+H_A^-$  are 5.7 and 219 ps, respectively. The state  $P^+Q_A^-$  does not decay within the investigated time scale (2 ns).

$Q_A$  (EADS, evolution-associated decay spectra) [8] in the psWt and L(M196)H RCs are presented in Figs. 5 and 6, respectively. The EADS were obtained from the global analysis of the complete set of the experimental absorption difference spectra recorded at different time delays between the pump and probe laser pulses using the sequential kinetic model with increasing lifetimes of the states ( $1 \rightarrow 2 \rightarrow 3 \rightarrow \dots$ ). It follows from Figs. 5 and 6 that the EADS spectral profiles and corresponding lifetimes are close for the two types of RCs and are in agreement with the data published for the wild type RCs (see, for example, Sun et al. [15]). The EADS associated with lifetimes 3.2 and 5.7 ps show spectral features typical for the excited state  $P^*$  (Figs. 5 and 6, curves 1). A broad negative band is observed in the  $Q_y$ -region of both spectra that include contributions from the bleaching  $P$  absorption band at  $\sim 865$  nm and from stimulated emission from  $P^*$  in the region of  $\sim 920$  nm, as well as the characteristic positive peak at 810 nm. The negative band at 600 nm is present in the  $Q_x$  spectral regions on the background of the weakly structured absorption of  $P^*$  in the range 500–720 nm. The  $P^*$  state evolves into the charge-separated state  $P^+H_A^-$  with EADS demonstrating the preserved negative band at 600 nm, bleaching of the  $Q_x$ -absorption band of the BPheo  $H_A$  at 545 nm, and emerging of the absorption bands of the radical anion  $H_A^-$  at  $\sim 670$  and 960 nm. Bleaching of the  $P$  absorption band at  $\sim 865$  nm is observed in the  $Q_y$ -region as well as electrochromic shift in the region of  $\sim 800$  nm (Figs. 5 and 6, curves 2). The amplitude of the bleaching at  $\sim 865$  nm in the EADS of the mutant RCs is reduced relative to the amplitude in

the EADS of the psWt RCs in accordance with the differences observed in the RC absorption spectra (Fig. 1). In the course of several hundreds of picoseconds ( $\tau \sim 200$  ps), the  $P^+H_A^-$  state in both types of RCs is evolved to the terminal state  $P^+Q_A^-$ , which does not decay on the time scale of our measurements (Figs. 5 and 6, curves 3).

Hence, all the results of femtosecond measurements indicate that the sequence  $P^* \rightarrow P^+H_A^- \rightarrow P^+Q_A^-$  and dynamics of the electron transfer involved in the charge separation as well as the total yield of the process that are observed for psWt RCs are practically fully maintained in the L(M196)H RCs.

Note that formation of the  $P^+B_A^-$  intermediate state [16–18], which decays faster than forms in the native RCs causing a low level of its population and difficulties in its experimental detection [19], has not been considered in this work during the global analysis. However, preliminary analysis of the difference spectra of the L(M196)H RCs with the reduced  $Q_A$  revealed a weak absorption band at  $\sim 1025$  nm, which could be assigned to the  $B_A^-$  radical anion [16, 18] (data not shown). It is likely that the two-step mechanism of the initial electron transfer  $P^* \rightarrow P^+B_A^- \rightarrow P^+H_A^-$  typical for the native RCs (see, for example, Khatypov et al. [20]) is conserved also in the mutant preparations.

## DISCUSSION

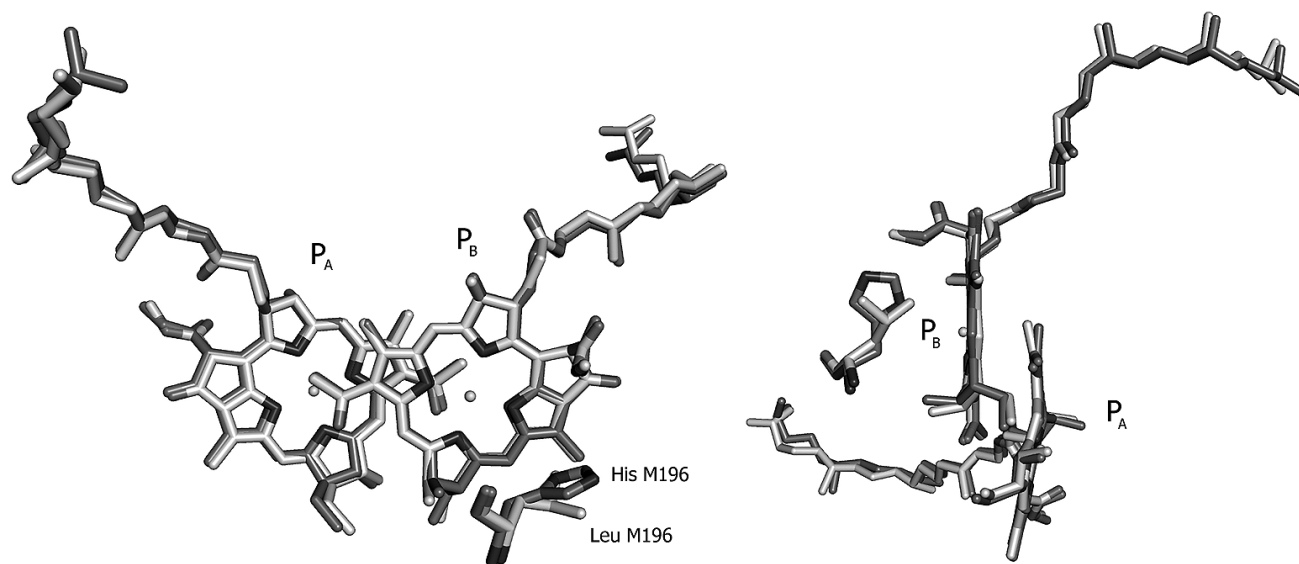
According to the X-ray diffraction data for the mutant L(M196)H RCs [2], the angle between the imida-

zole group of histidine M196 and the plane of the  $P_B$  macrocycle is estimated as  $103 \pm 10^\circ$ , which corresponds to the T-shaped geometry of the mutual arrangement of the histidine residue and BChl. The distance between the histidine nitrogen atom ND1 and ring C in the  $P_B$  molecule is  $3.8 \pm 0.15 \text{ \AA}$ . Hence, the introduced histidine and the  $P_B$  molecule are located quite close to each other, ensuring their interaction.

It can be seen from the crystal structure of the L(M196)H RCs (Fig. 7) that the mutation does not cause global changes in the geometry of the P dimer. The observed minor deviations between the RC structures of the mutant and psWt are likely beyond the limit of resolution. Hence, it can be concluded that the experimentally observed changes in the electronic structures of P and  $P^+$  (Figs. 1-3) are caused by the interaction between histidine M196 and the  $\pi$ -electron system of the  $P_B$  molecule. According to the literature data [1], the T-shaped orientation of the imidazole ring relative to the plane of the aromatic molecule is characteristic for  $\pi$ -hydrogen interaction. Formation of the  $\pi$ -hydrogen interaction in the mutant RCs would be in agreement with the fact that the L(M196)H mutation increases the redox potential of the  $P^+/P$  pair by  $\sim 60 \text{ mV}$  in comparison with the psWt RCs [2]. Apparently, formation of the hydrogen bond between the histidine and the conjugated system of P destabilizes the  $P^+$  state in comparison with the neutral state, thus hindering P oxidation and shifting the redox potential of the  $P^+/P$  pair towards the positive region as has been observed in the mutant RCs during formation of hydrogen bonds between the carbonyl groups of P and histidines [23, 24]. In this connection, it is worth men-

tioning that the long wavelength shift of the P band in the low temperature spectrum of the L(M196)H RCs [2] is in agreement with the theoretically estimated effect of formation of the hydrogen bond on the energy of the  $Q_y$ -transition in P [10].

It is interesting in this context that a similar effect on the low-temperature absorption spectrum of RCs, redox potential of P,  $P^+$  electronic structure, and primary photochemistry was observed previously for the *Rba. sphaeroides* L(M160)H mutant RCs, where a common hydrogen bond was introduced between the histidine M160 and  $13^1$ -keto-carbonyl group of the BChl  $P_B$  molecule [12, 25]. The shift of the absorption band of P towards longer wavelengths and decrease of its intensity was observed in the low-temperature (20 K) absorption spectrum of the L(M160)H RC mutant [25]. The redox potential of the  $P^+/P$  pair in the mutant RCs increased by 55 mV in comparison with the wild type RCs [25]. The hole-transfer band and one of the phase-phonon bands in the  $P^+Q_A^-/PQ_A$  FTIR spectrum of the L(M160)H mutant were shifted towards the high-frequency region to  $\sim 2800$  and  $\sim 1300 \text{ cm}^{-1}$ , respectively, and the intensity of both bands decreased [12]. It is significant that the decrease in intensity of the IR bands in the spectrum of L(M160)H [12] was pronounced to a considerably lower degree than in the L(M196)H RCs (Figs. 2 and 3, curves 2), not affecting, in particular, the phase-phonon band at  $1550 \text{ cm}^{-1}$ . These data can be explained by the fact that during formation of the common hydrogen bond in the L(M160)H RCs the hydrogen atom of the N-H-group in imidazole interacts with the electron density of oxygen in the  $13^1$ -keto-carbonyl group of  $P_B$ , while in the case of



**Fig. 7.** Overlay of crystal structures of the *Rba. sphaeroides* psWt RCs (PDB: 3V3Y [21]) and of the L(M196)H mutant [2] in the region of the primary electron donor P dimer in two projections. The respective amino acid residues at position M196 are marked. Crystal structure resolution for the psWt and L(M196)H RCs was 2.8 and 2.4  $\text{\AA}$ , respectively. The PyMOL program was used for generating this figure [22].

$\pi$ -hydrogen bond the hydrogen atom interacts directly with the  $\pi$ -electron cloud of the  $P_B$  BChl macrocycle, causing greater perturbation of the  $P^+$  electronic structure in the L(M196)H RCs. The time constants of electron transfer from  $P^*$  to  $H_A$  and from  $H_A^-$  to  $Q_A$  in the mutant L(M160)H RCs (5.7 and 210 ps, respectively) [25] are practically identical to the ones observed for the L(M196)H RCs (5.7 and 219 ps, respectively; this study).

It was shown previously using quantum-chemical calculations and EPR data that the semiquinone form of the quinone acceptor A1 could be stabilized in complexes of photosystem 1 (PS1) reconstructed with modified quinones with short side chains via re-orientation of the quinone molecule followed by formation of a  $\pi$ -hydrogen bond with tryptophan [26]. It was also suggested based on the quantum-chemical calculations that the  $CH-\pi$ -interactions between  $\beta$ -carotene and chlorophyll together with the  $\pi-\pi$ -stacking played an important role in the binding of a  $\beta$ -carotene molecule in the PS1 from *Synechococcus elongatus* [27]. The mutant *Rba. sphaeroides* L(M196)H RCs are likely to be the first example of formation of a  $\pi$ -hydrogen bond between the protein and one of the electron transfer cofactors in bacterial RCs. To the best of our knowledge, cofactor-protein bonds of this type have not been detected in the native photosynthetic RCs, while the formation of common hydrogen bonds between the protein and redox-active pigment and quinone cofactors plays an important role in optimization of the function (and, likely, maintenance of the structure) of the native RC complexes [28-30]. According to the literature data, the most favorable orientation of the  $\pi$ -hydrogen bond is the perpendicular arrangement of the X-H bond of the donor (X is atom N, O, or C) with respect to the plane of  $\pi$ -acceptor [1, 31, 32]. It was also shown that unlike for the common hydrogen bond, a "softer" geometry is characteristic for the  $\pi$ -hydrogen interaction, which makes formation of a larger number of conformations in a wider range of bonding energy more probable [32]. It is plausible that the  $\pi$ -hydrogen bonds do not provide noticeable functional advantages in comparison with the common hydrogen bonds (with respect to the effect on the redox properties of cofactors, for example), while the quite specific spatial organization of the  $\pi$ -hydrogen interaction and its potential heterogeneity with regards to bonding energy could be at least partial reason for the fact that this type of interaction between the protein and pigment and quinone cofactors is not realized in native RCs.

### Funding

This work was financially supported by State Task No. AAAA-A17030110140-5 and partially supported by the Russian Foundation for Basic Research (grant No. 17-00-00207 KOMFI, FTIR measurements, femtosec-

ond measurements; grant No. 17-44-500828, preparation of RC samples).

### Conflict of Interest

The authors declare no conflict of interest.

### Ethical Compliance

This work does not involve any experiments using humans and animals as study objects.

### REFERENCES

1. Liao, S. M., Du, Q. S., Meng, J. Z., Pang, Z. W., and Huang, R. B. (2013) The multiple roles of histidine in protein interactions, *Chem. Cent. J.*, **7**, 44-55, doi: 10.1186/1752-153X-7-44.
2. Fufina, T. Y., Vasilieva, L. G., Gabdulkhakov, A. G., and Shuvalov, V. A. (2015) The L(M196)H mutation in *Rhodobacter sphaeroides* reaction center results in new electrostatic interactions, *Photosynth. Res.*, **125**, 23-29, doi: 10.1007/s11120-014-0062-0.
3. Lutz, M., and Mantele, W. (1991) Vibrational spectroscopy of chlorophylls, in *Chlorophylls* (Scheer, H., ed.) CRC Press, Boca Raton, FL, pp. 855-902.
4. Breton, J., Nabedryk, E., and Parson, W. W. (1992) A new infrared electronic transition of the oxidized primary electron donor in bacterial reaction centers: a way to assess resonance interactions between the bacteriochlorophylls, *Biochemistry*, **31**, 7503-7510, doi: 10.1021/bi00148a010.
5. Gabdulkhakov, A. G., Fufina, T. Y., Vasilieva, L. G., Mueller, U., and Shuvalov, V. A. (2013) Expression, purification, crystallization and preliminary X-ray structure analysis of wild-type and L(M196)H mutant *Rhodobacter sphaeroides* reaction centres, *Acta. Crystallogr. Sect. F*, **69**, 506-509, doi: 10.1107/S1744309113006398.
6. Zabelin, A. A., Fufina, T. Y., Vasilieva, L. G., Shkuropatova, V. A., Zvereva, M. G., Shkuropatov, A. Y., and Shuvalov, V. A. (2009) Mutant reaction centers of *Rhodobacter sphaeroides* I(L177)H with strongly bound bacteriochlorophyll *a*: structural properties and pigment-protein interactions, *Biochemistry (Moscow)*, **74**, 68-74, doi: 10.1134/S0006297909010106.
7. Khatypov, R. A., Khristin, A. M., Fufina, T. Yu., and Shuvalov, V. A. (2017) An alternative pathway of light-induced transmembrane electron transfer in photosynthetic reaction centers of *Rhodobacter sphaeroides*, *Biochemistry (Moscow)*, **82**, 692-697, doi: 10.1134/S0006297917060050.
8. Van Stokkum, I. H. M., Larsen, D. S., and van Grondelle, R. (2004) Global and target analysis of time-resolved spectra, *Biochim. Biophys. Acta*, **1657**, 82-104, doi: 10.1016/j.bbabo.2004.04.011.
9. Snellenburg, J. J., Liptonok, S. P., Seger, R., Mullen, K. M., and van Stokkum, I. H. M. (2012) Glotaran: a Java-based graphical user interface for the R package TIMP, *J. Stat. Soft.*, **49**, 1-22, doi: 10.18637/jss.v049.i03.



10. Thompson, M. A., Zerner, M. C., and Fajer, J. (1991) A theoretical examination of the electronic structure and excited states of the bacteriochlorophyll *b* dimer from *Rhodospseudomonas viridis*, *J. Phys. Chem.*, **95**, 5693-5700, doi: 10.1021/j100167a058.
11. Parson, W. W., and Warshel, A. (1987) Spectroscopic properties of photosynthetic reaction centers. 2. Application of the theory to *Rhodospseudomonas viridis*, *J. Am. Chem. Soc.*, **109**, 6152-6163, doi: 10.1021/ja00254a040.
12. Naberdyk, E., Allen, J. P., Taguchi, A. K. W., Williams, J. C., Woodbury, N. W., and Breton, J. (1993) Fourier transform infrared study of the primary electron donor in chromatophores of *Rhodobacter sphaeroides* with reaction centers genetically modified at residues M160 and L131, *Biochemistry*, **32**, 13879-13885, doi: 10.1021/bi00213a017.
13. Reimers, J. R., and Hush, N. S. (2003) Modeling the bacterial photosynthetic reaction center. VII. Full simulation of the intervalence hole-transfer absorption spectrum of the special-pair radical cation, *J. Chem. Phys.*, **119**, 3262-3277, doi: 10.1063/1.1589742.
14. Malferrari, M., Turina, P., Francia, F., Mezzetti, A., Leibl, W., and Venturoli, G. (2015) Dehydration affects the electronic structure of the primary electron donor in bacterial photosynthetic reaction centers: evidence from visible-NIR and light-induced difference FTIR spectroscopy, *Photochem. Photobiol. Sci.*, **14**, 238-251, doi:10.1039/c4pp00245h.
15. Sun, C., Carey, A.-M., Gao B.-R., Wraight, C. A., Woodbury, N. W., and Lin, S. (2016) Ultrafast electron transfer kinetics in the LM dimer of bacterial photosynthetic reaction center from *Rhodobacter sphaeroides*, *J. Phys. Chem. B*, **120**, 5395-5404, doi: 10.1021/acs.jpcc.6b05082.
16. Arlt, T., Schmidt, S., Kaiser, W., Lauterwasser, C., Meyer, M., Scheer, H., and Zinth, W. (1993) The accessory bacteriochlorophyll: a real electron carrier in primary photosynthesis, *Proc. Natl. Acad. Sci. USA*, **90**, 11757-11761, doi: 10.1073/pnas.90.24.11757.
17. Shkuropatov, A. Ya., and Shuvalov, V. A. (1993) Electron transfer in pheophytin *a*-modified reaction centers from *Rhodobacter sphaeroides* (R-26), *FEBS Lett.*, **322**, 168-172, doi: 10.1016/0014-5793(93)81561-D.
18. Kennis, J. T. M., Shkuropatov, A. Ya., van Stokkum, I. H. M., Gast, P., Hoff, A. J., Shuvalov, V. A., and Aartsma, T. J. (1997) Formation of a long-lived P<sup>+</sup>B<sub>A</sub><sup>-</sup> state in plant pheophytin-exchanged reaction centers of *Rhodobacter sphaeroides* R26 at low temperature, *Biochemistry*, **36**, 16231-16238, doi: 10.1021/bi9712605.
19. Zinth, W., and Wachtveitl, J. (2005) The first picoseconds in bacterial photosynthesis – ultrafast electron transfer for the efficient conversion of light energy, *Chem. Phys. Chem.*, **6**, 871-880, doi: 10.1002/cphc.200400458.
20. Khatypov, R. A., Khmel'nitskiy, A. Yu., Khristin, A. M., Fufina, T. Yu., Vasilieva, L. G., and Shuvalov, V. A. (2012) Primary charge separation within P870\* in wild type and heterodimer mutants in femtosecond time domain, *Biochim. Biophys. Acta*, **1817**, 1392-1398, doi: 10.1016/j.bbabi.2011.12.007.
21. Vasilieva, L. G., Fufina, T. Y., Gabdulkhakov, A. G., Leonova, M. M., Khatypov, R. A., and Shuvalov, V. A. (2012) The site-directed mutation I(L177)H in *Rhodobacter sphaeroides* reaction center affects coordination of P<sub>A</sub> and B<sub>B</sub> bacteriochlorophylls, *Biochim. Biophys. Acta*, **1817**, 1407-1417, doi: 10.1016/j.bbabi.2012.02.008.
22. DeLano, W. L. (2002) The PyMOL molecular graphics system (<http://www.pymol.org>).
23. Lin, X., Murchison, H. A., Nagarajan, V., Parson, W. W., Allen, J. P., and Williams, J. C. (1994) Specific alteration of the oxidation potential of the electron donor in reaction centers from *Rhodobacter sphaeroides*, *Proc. Natl. Acad. Sci. USA*, **91**, 10265-10269, doi: 10.1073/pnas.91.22.10265.
24. Allen, J. P., and Williams, J. C. (1995) Relationship between the oxidation potential of the bacteriochlorophyll dimer and electron transfer in photosynthetic reaction centers, *J. Bioenerg. Biomembr.*, **27**, 275-283, doi: 10.1007/BF02110097.
25. Williams, J. C., Alden, R. G., Murchison, H. A., Peloquin, J. M., Woodbury, N. W., and Allen, J. P. (1992) Effects of mutations near the bacteriochlorophylls in reaction centers from *Rhodobacter sphaeroides*, *Biochemistry*, **31**, 11029-11037, doi: 10.1021/bi00160a012.
26. Kaupp, M. (2002) The function of photosystem I. Quantum chemical insight into the role of tryptophan-quinone interactions, *Biochemistry*, **41**, 2895-2900, doi: 10.1021/bi0159783.
27. Wang, Y., Mao, L., and Hu, X. (2004) Insight into the structural role of carotenoids in the photosystem I: a quantum chemical analysis, *Biophys. J.*, **86**, 3097-3111, doi: 10.1016/S0006-3495(04)74358-1.
28. Allen, J. P., and Williams, J. C. (2006) The influence of protein interactions on the properties of the bacteriochlorophyll dimer in reaction centers, in *Chlorophylls and Bacteriochlorophylls: Biochemistry, Biophysics, Functions and Applications* (Grimm, B., Porra, R. J., Rudiger, W., and Scheer, H., eds.) Springer, The Netherlands, pp. 283-295.
29. Bylina, E. J., Kirmaer, C., McDowell, L., Holten, D., and Youvan, D. C. (1988) Influence of an amino acid residue on the optical properties and electron transfer dynamics of a photosynthetic reaction centre complex, *Nature*, **336**, 182-184, doi: 10.1038/336182a0.
30. Flores, M., Isaacson, R., Abresch, E., Calvo, R., Lubitz, W., and Feher, G. (2007) Protein-cofactor interactions in bacterial reaction centers from *Rhodobacter sphaeroides* R-26: II. Geometry of the hydrogen bonds to the primary quinone Q<sub>A</sub><sup>-</sup> by <sup>1</sup>H and <sup>2</sup>H ENDOR spectroscopy, *Biophys. J.*, **92**, 671-682, doi: 10.1529/biophysj.106.092460.
31. Steiner, T., and Koellner, G. (2001) Hydrogen bonds with  $\pi$ -acceptors in proteins: frequencies and role in stabilizing local 3D structure, *J. Mol. Biol.*, **305**, 535-557, doi: 10.1006/jmbi.2000.4301.
32. Du, Q.-S., Wang, Q.-Y., Du, L.-Q., Chen, D., and Huang, R.-B. (2013) Theoretical study on the polar hydrogen- $\pi$  (Hp- $\pi$ ) interactions between protein side chains, *Chem. Cent. J.*, **7**, 92-99, doi: 10.1186/1752-153X-7-92.