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ENDOR Spectroscopy Reveals A Light Induced Movement of the H-Bond from Ser-L223 Upon Forming the Semiquinone ($Q_B^{\cdot-}$) in Reaction Centers from *Rhodobacter sphaeroides*

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Abstract

Proton ENDOR spectroscopy was used to monitor local conformational changes in bacterial reaction centers (RC) associated with the electron transfer reaction $DQ_B \rightarrow D^{+\cdot}Q_B^{\cdot-}$ using mutant RCs capable of photo-reducing Q_B at cryogenic temperatures. The charge separated state $D^{+\cdot}Q_B^{\cdot-}$ was studied in mutant RCs formed by either (i) illuminating at low temperature (77K) a sample frozen in the dark (ground state protein conformation) or (ii) illuminating at room temperature prior to and during the freezing (charge separated state protein conformation). The charge recombination rates from the two states differed greatly ($>10^6$ fold) as shown previously, indicating a structural change (Paddock *et al* (2006) *Biochemistry* **45**, 14032 - 14042). ENDOR spectra of $Q_B^{\cdot-}$ from both samples (35 GHz, 77K) showed three nearly identical sets of hyperfine couplings due to exchangeable protons that were similar to those for $Q_B^{\cdot-}$ in native RCs indicating that in all RCs, $Q_B^{\cdot-}$ was located at the proximal position near the metal site. In contrast, one set of H-bond couplings was observed *only* in the sample frozen under illumination in which the protein can relax prior to freezing. This H-bond was assigned to an interaction between the Ser-L223 hydroxyl and $Q_B^{\cdot-}$ based on its absence in Ser L223 $\rightarrow$ Ala mutant RCs. The Ser-L223 hydroxyl H-bond was also observed in the native RCs frozen under illumination. Thus, part of the protein relaxation in response to light induced charge separation involves the formation of an H-bond between the OH group of Ser-L223 and the anionic semiquinone $Q_B^{\cdot-}$. This proton movement serves to stabilize the charge separated state and facilitate proton transfer to reduced Q_B .

Keywords

B-branch; bacterial reaction center; quinone movement; bacterial photosynthesis; electron transfer; nanoswitch

Conformational changes in protein structure associated with electron transfer reactions play an important role in stabilizing the resultant charge separated state and determining the rates of electron transfer (1-3). In photosynthetic bacteria, reaction centers (RCs) perform the initial photochemical reactions in photosynthesis (see *e.g.* 4,5) resulting in the light induced reduction of a bound quinone, Q_B . The reduction of Q_B clearly demonstrates the influence of conformational dynamics on electron transfer. The overall reaction is shown by the solid curves in Figure 1. The first step is the initial photochemistry, a rapid ($\tau = 10^{-10}$ s) light induced electron transfer from an electron donor species D (a bacteriochlorophyll dimer) along the A-branch (one of two pseudo-symmetric pathways) through a series of acceptor molecules bacteriochlorophyll, B_A , bacteriopheophytin, H_A , to a tightly bound quinone, Q_A (Figure 1). This is followed by a

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slower ($\tau = 10^{-4}$ s) electron transfer from $Q_A^{-\bullet}$ to a substrate quinone, Q_B (for a review see 8). The rate of the electron transfer reaction $Q_A^{-\bullet}Q_B \rightarrow Q_AQ_B^{-\bullet}$, $k_{AB}^{(I)}$, is coupled to protein dynamics as shown by several experimental observations (for a review see 9). First, the rate of reaction is dependent on protein structure as illustrated by the effects of crosslinking (10), embedding in a polymer matrix (11) and hydration (12). Second, the rate is independent of driving force (13), showing that electron transfer is not the rate limiting step, indicating a conformational gating mechanism. Third, large differences in the rate of electron transfer at cryogenic temperature were found depending upon whether RCs are frozen in the light or in the dark, indicating that a structural change between the neutral and charge separated states is important for electron transfer (14,15). In the present study we use ENDOR spectroscopy (Electron Nuclear Double Resonance) (see *e.g.* 16,17) to examine the differences in local structure near Q_B between the neutral and charge separated states trapped at cryogenic temperatures in order to understand the structural changes associated with the reduction of Q_B that may contribute to the conformational gating of electron transfer.

Studies made on protein samples frozen at cryogenic temperatures served to elucidate the importance of conformational effects on reaction dynamics. In early studies, large differences in transfer rates were observed between RCs frozen in the light or in the dark (14,15,18). For RCs frozen in the dark and illuminated at cryogenic temperatures, the reaction $k_{AB}^{(I)}$, became slower as the temperature decreased and did not appreciably occur below 150K. However, if RCs were illuminated during the freezing process in the $D^{+\bullet}Q_B^{-\bullet}$ state and the charges were allowed to recombine without thawing, photochemical electron transfer at cryogenic temperatures was observed (15,18), showing that a change in a structure has occurred between the neutral state and the charge separated state. The x-ray crystal structure of RCs frozen in the light ($Q_B^{-\bullet}$) and frozen in the dark (Q_B) showed a large change in the position of the quinone head group. $Q_B^{-\bullet}$ was displaced by $\sim 5\text{\AA}$ to a proximal position from the prevalent distal position observed for Q_B . In this proximal position, $Q_B^{-\bullet}$ could form 4 H-bonds to protein groups (Figure 2) (6).

The large change in the position of the quinone head group suggested that the movement of Q_B from the distal to the proximal position is a possible mechanism for conformational gating of the electron transfer to Q_B (8,9). Arguments against this process being rate limiting were the finding that the rate was independent of the tail length of the quinone molecule (13,20) and the finding that a mutant having Q_B bound in the proximal site retained a slower gated rate (21). In addition, infrared studies suggest that at room temperature Q_B is predominantly in the proximal site (22). These results suggested that additional conformational changes besides the quinone movement must contribute to the reaction.

An additional model for the conformational gate is that the protein structure alters the free energy of electron transfer from $Q_A^{-\bullet}$ to Q_B . In this model the free energy is unfavorable with the protein in the “unrelaxed” structure near Q_B (RCs frozen in the dark). It becomes favorable as the protein and solvent “relax” into a conformation that stabilizes the charge separated state (reflected in the RCs frozen in the light). In order to test this hypothesis, we have studied the reduction of Q_B using RCs mutants that are modified to allow direct electron transfer to Q_B from the B-branch bacteriopheophytin located close to Q_B (see red curve in Figure 1). Using the higher driving force for electron transfer it was possible to form the charge separated state $D^{+\bullet}Q_B^{-\bullet}$ in RCs frozen in the less thermodynamically favorable “unrelaxed” conformation of the dark (7). In this way it was possible to monitor the differences between the stability of the charge separated $D^{+\bullet}Q_B^{-\bullet}$ states and measure the EPR and ENDOR spectra for RCs frozen in the “unrelaxed” (dark frozen) and “relaxed” (light frozen) conformations.

In a previous paper (7) we showed that the lifetime of the charge separated state $D^{+\bullet}Q_B^{-\bullet}$, which is dependent on the stabilization of $Q_B^{-\bullet}$, was dramatically effected by the conditions

of its formation. In a dark frozen sample (77K), charge separation occurred in a minor fraction (~30%) of the sample with a lifetime of ~6s. In a sample frozen under illumination, $D^{+\bullet}Q_B^{-\bullet}$ was formed in the entire sample with a lifetime exceeding 10^7 s (77K) (7). Thus, the protein response, which was prevented by freezing in the dark in the former sample, resulted in a $> 10^6$ -fold difference in the lifetime at 77K. For simplicity, we shall refer to samples prepared in that manner as unrelaxed (frozen in the dark) and relaxed (in which illumination occurred prior to freezing allowing the protein to respond/relax). In those studies the $Q_B^{-\bullet}$ were formed by electron transfer from BPhe_B, in a mutant designed to increase the efficiency of B-branch electron transfer (described in 23). This mutant RC carried the Ala-M260 → Trp replacement designed to remove Q_A (23,24,25), thereby creating an RC system without an admixture of $Q_A^{-\bullet}$.

In this work, we extend the previous EPR studies by using ENDOR spectroscopy at 35GHz to examine the local structure near $Q_B^{-\bullet}$ in both the relaxed and unrelaxed protein conformations. ENDOR spectroscopy can be used to determine positions of protons that are not observable with x-ray crystallography and is particularly well suited for examination of detailed interactions between a radical cofactor and its protein surrounding (see *e.g.* 26). Interactions between the magnetic moment of the unpaired electron of the radical and magnetic moments of protons (nuclei) from the surrounding protein are detected with a spatial resolution approaching 0.03Å (see *e.g.* 27). Thus, ENDOR can provide a precise measure of the binding position and details on the interactions of $Q_B^{-\bullet}$ in its binding pocket (see Figure 2). At 35GHz, the 1H ENDOR signals resulting from interactions between $Q_B^{-\bullet}$ and the protein are spectrally separated from those associated with $D^{+\bullet}$. The 1H ENDOR spectra of the relaxed and unrelaxed states were measured and differences between them discussed and the implications for the catalytic functions (*i.e.* reduction and protonation) of Q_B are discussed.

MATERIALS AND METHODS

Construction and purification of the quintuple B-branch mutant RCs

The construction and purification of the B-branch mutant was previously described (23). The mutant RC contained the following changes: GCC → TGG (Ala-M260 → Trp), CTG → CAC (Leu-M214 → His), TAC → TTC (Tyr-M210 → Phe), GGT → GAC (Gly-M203 → Asp), TTC → TAC (Phe-L181 → Tyr) and in one mutant TCG → GCC (Ser-L223 → Ala). RCs were purified as previously described (28).

Biochemical Zn²⁺ Replacement

To perform EPR and ENDOR spectroscopic studies on $Q_B^{-\bullet}$, the high spin Fe^{2+} was removed and replaced with diamagnetic Zn^{2+} as described by Debus *et al* (29) and as modified by Utschig *et al* (30). Protonated quinone/semiquinone was removed from the Q_B site by incubating for 30 minutes in 100mM Caps pH 11 and 100μM ferricyanide (to oxidize the fraction of $Q_B^{-\bullet}$ present in the dark), and 2% LDAO and 100mM o-phenanthroline (to displace UQ_{10}). The RCs were bound to a DEAE column and rinsed with 10 column volumes of 10mM Caps pH 11, 10 μM ferricyanide, 100mM o-phenanthroline and then with 10 column volumes 10mM Tris pH 8, 0.025% LDAO, 1mM EDTA. The B-branch mutant RCs were reconstituted with ~ 3-fold excess of deuterated UQ_{10} in 1% LDAO to occupy the Q_B site followed by dialysis against TMK (2mM Tris-HCl pH 8, 0.04 % β-D-maltoside, 5mM KCl). The Q_B site of the native RC was reconstituted with 10-fold excess deuterated UQ_{10} in 1% LDAO, diluted 4-fold and concentrated in an Amicon concentrator. The concentrated material was then used to make EPR and ENDOR samples with the intent of minimizing incubation that would allow exchange of protonated quinone from the Q_A site.

35GHz EPR and ^1H ENDOR measurements

EPR and ENDOR spectroscopy was performed at a microwave frequency of 35 GHz at $T = 77$ K as previously described (31). For the B-branch mutant RC, two parallel samples were made from the same RC material (at ~ 100 - $700\mu\text{M}$ RC). One was frozen in liquid nitrogen in the dark. The other was illuminated by a projector ($\sim 1\text{W}/\text{cm}^2$, ~ 5 s) and frozen in liquid nitrogen under illumination. A similar projector was used to illuminate the sample in the magnetic resonance cavity to generate the charge separated state in the sample frozen in the dark. For the native RC, the $\text{D}^{+\bullet}\text{Q}_\text{A}^{-\bullet}$ state was produced by chemical reduction as described (31). The $\text{D}^{+\bullet}\text{Q}_\text{B}^{-\bullet}$ state was produced by illuminating with a projector ($\sim 1\text{W}/\text{cm}^2$, ~ 1 s) and frozen in liquid nitrogen under illumination.

THEORY

ENDOR spectroscopy utilizes radio-frequency radiation to excite nuclear magnetic resonance transitions that are detected by changes in the EPR signal of an interacting electron spin (16, 17). ENDOR spectra are plots of the EPR signal amplitude *versus* the frequency of the applied radio-frequency radiation used to modulate the proton spin. The simplest ENDOR spectrum of a single proton interacting with an electron spin for a molecule in a fixed orientation, such as in a crystal, consists of 2 lines centered symmetrically around the free proton frequency, ν_free where A is the hyperfine coupling due to the interaction between the nuclear spin and electron spin (Eqn. 1). The larger the coupling, the larger the interaction and most likely the smaller the distance between the proton and electron spin.

$$\nu = \nu_\text{free} \pm \frac{A}{2} \quad (1)$$

where ν is the peak frequency, ν_free the free proton frequency and A the hyperfine coupling constant.

The ENDOR spectra due to interactions between H-bonded protons and the semiquinone have been extensively described (26). In its simplest form the ENDOR spectra from H-bonded protons arise from the magnetic dipole interaction between the nuclear spin of the proton with the electron spin at the H-bonded oxygen on the quinone. The interaction is anisotropic, *i.e.* depends on the orientation of the magnetic field vector with respect to the H-bond direction. For a randomly oriented sample such as used in this study, the powder pattern ENDOR spectrum due to a single proton shows resonances at many frequencies due to the different orientations of the molecule in the magnetic field. At the magnetic field position corresponding to g_y , the spectrum corresponds to a “two-dimensional” powder-type spectrum, with weighted contributions mainly from all *in-plane* directions, providing little field selection (see *e.g.* 27). The major features of the spectra are four peaks with two peaks below the free proton frequency and two peaks above the free proton frequency symmetrically spaced around the free proton frequency. The larger coupling, $A_\parallel$, is from an orientation in which the magnetic field is parallel to the H-bond direction and the smaller coupling, $A_\perp$, is from an orientation in which the magnetic field is perpendicular to the H-bond direction. For the case where the interaction is due to a magnetic dipolar interaction alone (*i.e.* no isotropic coupling due to covalency), the principal values are related by a factor of two such that $A_\parallel = -2A_\perp$ (26). For this simple case the value of the hyperfine coupling is given by Eqn. 2:

$$A = \frac{79.2\rho}{r^3}(3\cos^2\delta - 1) [\text{MHz}] \quad (2)$$

where r is the distance in Å between proton and oxygen, ρ is the spin density on the oxygen, δ is the angle between the magnetic field and the H-bond direction. For $A_\parallel$, $\delta = 0$ and for $A_\perp$, $\delta = 90^\circ$.

$= 90^\circ$. The simple point dipole model adequately describes some systems, such as (within 5%) benzosemiquinones in solution (see *e.g.* 32). However, recent studies of the interaction of $Q_A^{\bullet-}$ with its two H-bonds in the RC showed that this simple picture must be modified due for interaction with the His-Zn²⁺ proton due to a significant isotropic coupling with the semiquinone. The point dipole approximation can also fail for the case where the direction of the H-bond significantly deviates from the plane of the quinone ring (27,33).

RESULTS

EPR spectra in the frozen state at 77K

The signals from the charge separated $D^+Q_B^{\bullet-}$ states were monitored by EPR spectroscopy (35 GHz, 77 K) using Quintuple B-branch mutant RC (Quint) RCs (containing Zn²⁺/deuterated Q₁₀) frozen in the light or frozen in the dark. In the samples frozen under illumination, an EPR signal was observed due to the charge separated state $D^+Q_B^{\bullet-}$ (Figure 3, LIGHT). At 35GHz, the signal due to $Q_B^{\bullet-}$ at low field was completely separated from the signal at higher field due to D^+ (see Figure 3) allowing ENDOR spectra of the quinone to be made with no contribution from the donor. The EPR signal was stable at 77 K and does not decay for $\tau > 10^6$ s. For the sample frozen in the dark no EPR signal was observed initially (Figure 3, DARK). However, an EPR signal due to the charge separated state $D^+Q_B^{\bullet-}$ was formed upon illumination. (Figure 3, DARK illum) The EPR signals for RCs frozen in the light and dark exhibited very similar *g* values. The light induced signal decayed in the dark with a lifetime of ~ 6 s as previously observed (7). The signal was absent in control samples (not shown) to which the Q_B site inhibitor, stigmatellin, was added showing that the signal results from $D^+Q_B^{\bullet-}$. Similar EPR signals were seen in Quint RCs having the mutation Ser L223 $\rightarrow$ Ala that were frozen in the light. However, no light induced EPR signals were seen in these mutant RCs that were frozen in the dark.

¹H ENDOR spectra of $Q_B^{\bullet-}$ in the frozen state at 77K in the B-branch mutant RC

To investigate local structural changes that occur upon charge separation, ¹H ENDOR spectra of $Q_B^{\bullet-}$ were measured in the Quint RC frozen in the light and frozen in the dark. In addition, the ENDOR spectrum of Quint RCs having a mutation to the H-bond donor residue Ser L223 $\rightarrow$ Ala. was measured in order to help identify ENDOR lines. The samples contained deuterated quinone. Thus the proton ENDOR lines due the protons on the quinone are eliminated and only ENDOR lines contributed by the protein and solvent protons are expected. The samples were monitored at the magnetic field position corresponding to *g_y* (see arrow in Figure 3), which provided the largest $Q_B^{\bullet-}$ ENDOR signals with minimal contribution from D^+ .

The complete ENDOR spectra centered around the free proton frequency are displayed in Figure 4a. Lines with larger splittings ($|A| > 3$ MHz) are due to hydrogen bonding interactions (H-bonds) between the oxygen atoms of $Q_B^{\bullet-}$ and protons from the nearby protein (26). Lines with smaller splittings ($|A| < 3$ MHz) due to weaker interactions of the quinone with more distant protons are seen in the matrix region (Matrix). A few changes occurred in the “matrix” region. In particular, the lower frequency edge of the matrix region (52.2 to 52.5 MHz) differed between the Quint (light) and Quint (dark) and between these and the mutant RC with the additional SAL223 replacement. Assignments for these lines were not attempted, since such lines are the result of interactions with numerous more distant protons. In addition, not all of the matrix protons were exchangeable indicating contributions to the spectra of non-exchangeable side chain protons.

The low frequency side ENDOR spectra in the H-bonding region for the three samples are shown in Figure 4B. In the ENDOR spectrum of Quint RC frozen in the light we tentatively assign 3 pairs of lines to *A_{||}* and *A_⊥* components of 3 H-bonds to $Q_B^{\bullet-}$. These are shown in

Figure 4B, LIGHT labeled (1, 1'), (2, 2') and (3, 3') respectively. The smaller peaks (1, 2, 3) were attributed to the parallel ($A_{\parallel}$) components of the three H-bond hyperfine coupling tensors. The higher intensity lines (1' & 3') and shoulder (2') were attributed to their respective perpendicular ($A_{\perp}$) components. The amplitude of all of the H-bonding lines were all diminished when the solvent was changed to D₂O (48 hrs) showing that the protons were exchangeable (data not shown) as expected. It is likely that the peaks 3 and 3' are composed of overlapping signals from two different protons based on the previous site-directed mutagenesis experiments on Native RCs (7).

The ENDOR spectrum of $Q_B^{\bullet-}$ in the H-bonding region in RCs frozen in the dark was similar to that of RCs frozen in the light indicating that almost all of the H-bonds are present in both cases (Figure 4b, DARK). However, peak 2 (and 2'), indicated by the arrow in Figure 4B, is present in the sample frozen in the light but is not present in the sample frozen in the dark. To help identify the missing peak the ENDOR spectrum of the mutant RC containing the additional Ser-L223 → Ala replacement (SAL223) was measured. The ENDOR spectrum of the SAL223 mutant in the H-bond region is similar to that of the Quint RCs but lacks peak 2 (and 2') (Figure 4b, SAL223). The absence of peak 2 in the SAL223 ENDOR spectrum shows that this peak is due to the H-bond from Ser L223.

¹H ENDOR spectra of $Q_B^{\bullet-}$ in the Native RC

We wished to assess the similarity of the native $Q_B^{\bullet-}$ ENDOR spectra with that measured for the B-branch mutant for samples frozen under illumination. However, it was difficult to obtain a pure ¹H ENDOR spectrum of $Q_B^{\bullet-}$ of the native RC due to contributions from $Q_A^{\bullet-}$ in the sample due to the intrinsic equilibrium between $Q_A^{\bullet-}$ and $Q_B^{\bullet-}$ and a fraction of RCs that lack Q_B . In order to remove the spectrum of $Q_A^{\bullet-}$ from the native ENDOR spectrum we focused on the matrix region which was expected to be very similar since it is composed of contributions of more distant protons that are essentially structurally the same in the native and mutant RCs (35). Due to the reconstitution procedure, the Q_A contained almost exclusively the native (non-deuterated) Q_{10} (spectrum shown in Figure 5A). A fraction of the $Q_A^{\bullet-}$ spectrum was subtracted from the native OBS spectrum giving a DIFF spectrum (Figure 5A, OBS) that resembled that of the B-branch mutant $Q_B^{\bullet-}$ in the matrix region (Figure 4A, quintuple LIGHT). In particular, the peaks in the matrix region near 52.6 and 53.8 MHz were diagnostic of the $Q_A^{\bullet-}$ contribution to the observed spectrum. The resultant difference spectrum is shown labeled as DIFF in Figure 5. This spectrum revealed peaks that were analogous to those observed in the ¹H-ENDOR $Q_B^{\bullet-}$ spectrum of the B-branch mutant RC frozen under illumination (Figure 4, LIGHT). In particular, a peak analogous to that assigned to the hydroxyl group of Ser-L223 (arrow) was observed in the native sample (Figure 5B); the slight shift in the position of the peak near 49.3MHz could be the result of a small ($< 0.1 \text{ \AA}$) increase in the length of the H-bond in the native RC.

DISCUSSION

In this work, we used ENDOR spectroscopy to investigate the local changes in the protein that occur near Q_B upon its reduction in the photosynthetic bacterial RC from *Rhodobacter (Rb.) sphaeroides*. Changes were observed that could contribute to the $>10^6$ -fold greater lifetime of the $D^{\bullet+}Q_B^{\bullet-}$ state formed by freezing under illumination compared to that formed by illumination of a sample frozen in the dark (Figure 2). The observed changes, the implications on the binding position of the quinone and its properties and the possible functional relevance for facile proton transfer of the observed changes are discussed below.

Reducible Q_B^- bound to the proximal position

The ^1H -ENDOR spectra of Q_B^- in the H-bonding region in Quint RCs frozen in the light and in the dark (Figure 4B) and in Native RCs (Figure 5B) were very similar (with the exception of a single peak that is absent in RCs frozen in the dark). The similarity in the peak positions indicated that most of the H-bonds to the semiquinone were similar in all three RCs. Since the semiquinone Q_B^- in native RCs was found to be in the proximal position in the x-ray crystal structure for the Native RC (8), we conclude that in all cases Q_B^- was located in the proximal binding site.

Q_B^- bound to the distal position was not observed

For the Quint RC frozen in the dark, it has been shown that only a fraction (~30%) are in an active configuration capable of photochemical reduction of Q_B (7). The results from this study indicated that this active configuration is one in which Q_B is bound in the proximal site. The activity of Q_B in the proximal site is probably due to the strong effect of the H-bonds in stabilizing the negative charge on the semiquinone in the charge separated state. The X-ray crystal structure of Quint RCs frozen in the dark showed that the predominant position of Q_B was in the distal site (23). Thus, these results show that Q_B in the distal site is apparently not stably reduced. It was either not reduced efficiently by H_B^- or was reduced but rapidly recombined resulting in a small (unobserved) steady state population.

Assignment of the mobile proton upon Q_B reduction to Ser-L223

Although the majority of the ENDOR lines in the H-bonding region were the same between the Quint (light) and Quint (dark), there was one peak indicated by an arrow (labelled 2) that was absent in the Quint (dark) (Figure 4B). Most of the H-bonds (from NH protons from backbone peptides of His) to the quinone bound in the proximal site are rigidly fixed in the protein structure and their positions can be deduced from the x-ray crystal structure. In contrast, the hydroxyl group from Ser L223 can serve as an H-bond donor to either Q_B or to Asp L213 (Figure 2). Thus, the H-bond from Ser L223 cannot be determined by x-ray crystallography. The definitive assignment of the ENDOR line 2 to the hydroxyl proton of Ser-L223 came from studies on the effects of mutations on the observed spectrum. In the mutant RC with the additional Ser-L223 $\rightarrow$ Ala replacement, the peak was absent (see arrow Figure 4B). From this result we conclude that this peak is due to the interaction from the hydroxyl proton of the Ser side chain.

The absence of peak 2 assigned to Ser L223 in the un-relaxed RC was attributed to a change in the position of the proton due to the switch in the H-bond from Q_B^- to Asp L213. Since ENDOR is extremely sensitive to distance, an increase in the distance between the oxygen atom of Q_B^- and the proton of as small as $\sim 1\text{\AA}$ would shift the peak from the H-bonding region to the "matrix" region (Figure 4A). Although the ENDOR spectrum in the matrix region is very congested and difficult to analyze, there are changes that could be due to movement of the H-bonded proton, for instance the low frequency peak at 52.25 MHz. Thus, the proton responsible for the ENDOR line had a hyperfine coupling of $A_{\parallel} = 8.0\text{ MHz}$ ($\pm 5\%$) in the Quint (light) (i.e. the line is shifted $\pm 4.0\text{ MHz}$ from the free proton frequency of 53.25 MHz, to 49.25 MHz on the low frequency side shown in Figure 4B) and a smaller coupling of most likely $A_{\parallel} < 2\text{ MHz}$ in the Quint (dark).

For a proton in a typical hydrogen bond the point-dipole approximation can be used to estimate the H-bonding distance from the value of the hyperfine interaction (Eqn. 2). Using the apparent value of $A_{\parallel} = 8.0\text{ MHz}$ and previously determined values for the spin density ($\rho = 0.18$) at the oxygen atom nearest Ser-L223 (see *e.g.* 26), we obtain from Eqn. 2 a distance of $1.5\text{\AA}$ from the proton to the oxygen atom. Since this equation is known to underestimate the distance by $\sim 0.2\text{\AA}$ for the H-bonds to Q_A^- (27,33), the real distance is slightly longer and closer to $1.7\text{\AA}$.

which is essentially an optimal hydrogen bond distance (see *e.g.* 37). Thus, the magnitude of the hyperfine interaction adds further support for the assignment of the line to the H-bond from Ser L223.

The Ser-L223 hydroxyl proton ENDOR line in the Native RC

The $Q_B^{\bullet-}$ ENDOR spectrum of the native RC was determined for the relaxed protein conformation by subtracting the contribution due to $Q_A^{\bullet-}$. The similarity of the native deconvoluted $Q_B^{\bullet-}$ spectrum in the H-bonding region (Figure 5B) to that measured in the B-branch mutant (Figure 4B) showed that $Q_B^{\bullet-}$ has essentially the same H-bond interactions in both the native and mutant RCs. In particular the peak assigned to the H-bond from Ser L223 is present in both native and Quint RCs frozen in the light. The small changes in the position of the ENDOR peaks in the H-bonding region (~ 0.1 - 0.2 MHz) indicate slight changes in the H-bond juxtaposition (such as change in distance as minor as ~ 0.02 Å). These results indicate that in the relaxed protein conformation, $Q_B^{\bullet-}$ was in the proximal position with an H-bond to Ser L223 in the native RC as well as in the Quint RC.

These results extend the previous findings by Utschig *et al.* (38) who used high field ENDOR to investigate changes in $D^{+}Q_B^{\bullet-}$ states formed by freezing under a variety of light exposure conditions, resulting in a variety of partially relaxed configurations. They obtained samples in which $D^{+}Q_B^{\bullet-}$ was “inactive”, i.e. no observed recombination at 20K, or “active” in which $D^{+}Q_B^{\bullet-}$ decayed non-exponentially in 10-100 s at 20 K and could be formed by illumination at low temperature. The former state was analogous to the “relaxed” $Q_B^{\bullet-}$ state reported in this work. They reported no difference between the ENDOR spectra in the matrix regions of their active and inactive states. This observation suggests that their “active” i.e. partially relaxed, state has similar ENDOR properties as the fully relaxed state.

Gating of electron transfer

One of the unresolved questions related to the reduction of Q_B is the nature of the conformational gate that limits its reduction *via* the prevalent A-branch. The sensitivity of ENDOR spectroscopy enables one to observe small changes in proton positions that are otherwise difficult to detect but possibly important for the energetics of enzymatic reactions. In this work, we observe a change in the H-bonding of $Q_B^{\bullet-}$ in response to its reduction using the quintuple mutant in which Q_B was reduced *via* the B-branch. Electron transfer *via* the B-branch has a greater driving force thereby allowing access to states that are otherwise inaccessible. We assigned the responsive H-bond to the hydroxyl group of Ser-L223. The overall similarity with the native ENDOR spectrum suggests that these results are applicable to the native RC. An increase in the stabilization of $Q_B^{\bullet-}$ could result from the formation of an H-bond to the Ser-L223 hydroxyl group and could explain (in part) the increased lifetime of the relaxed $D^{+}Q_B^{\bullet-}$ state. However, the Ser-L223 H-bond formation is not by itself the conformational gate that limits Q_B reduction *via* the A-branch, since the observed electron transfer rate remains very similar in a mutant in which Ser-L223 is replaced with Ala (39). Additional changes observed in the matrix region indicated that more distant protons have also responded to the charge separation. Further studies are needed to assign these lines to specific protons.

Effects of protein response on functional properties of Q_B

A major functional difference between the Quint RCs frozen in the dark and frozen in the light are the lifetimes of the charge separated states which were found to be 6 s and $> 10^7$ s, respectively (7). Similar long lived states have been observed by other authors for native RCs frozen under different illumination conditions (15,18,38,41,42). In principal the longer lifetime of the long lived state frozen in the light may be due to a reduced electron transfer rate for the back electron transfer reaction. This could result from either a change in electronic coupling

or in the driving force (43). Since the ENDOR spectra indicate that the location of $Q_B^{\bullet-}$ is the same for RCs frozen in the light or in the dark, it is unlikely that a change in the electronic coupling alone could be large enough to account for the change in rate. A change in distance of greater than 5 Å would be required, which is inconsistent with ENDOR results. Thus, it seems that the stabilization of the charge separated state is due to a lowering in energy of the relaxed state.

Although it is tempting to ascribe the lowering in energy in the relaxed state to the switch of the Ser L223 H-bond, the H-bond energy is not sufficient to account for the observed change in the rate since we observed a long lifetime for the Quint mutant lacking Ser L223 (Figure 4). Instead it is likely that there are many changes that contribute to the charge stabilization. Alexov and Gunner (44), Ishikita and Knapp (45) and Zhu and Gunner (46) have provided possible molecular responses and a starting point for discussion of molecular changes resulting from the reduction of Q_B . In addition to the stabilizing effect of forming an H-bond to $Q_B^{\bullet-}$ from Ser L223, protonation of a cluster of acids near the Q_B site could provide additional stabilizing to the charge separated state. Furthermore, protonation of a water molecule in the Q_B region was suggested based on FTIR spectroscopy (47). The overall stabilization is likely to be due to the sum of many contributions.

Our ENDOR results indicate the formation of an H-bond between $Q_B^{\bullet-}$ and Ser L223 at cryogenic temperatures. In contrast, FTIR studies suggest that the $Q_B^{\bullet-}$ is not H-bonded to Ser L223 at higher temperatures (> 280K) (48). A possible resolution to this seemingly contradictory result is that the H-bonding is temperature dependent. If the Ser L223 does not form a H-bond to $Q_B^{\bullet-}$ then it should be H-bonded to the nearby Asp-L213 (Figure 2). The H-bond to Asp L213 should be especially favorable if Asp L213 is negatively charged as suggested by kinetic measurements (40,49) and FTIR studies (50). The formation of an H-bond between Ser L223 and $Q_B^{\bullet-}$ at low temperature suggests that under these conditions Asp L213 is protonated. We suggest that the protonation of Asp L213 may also be present in the relaxation process (Figure 6). The protonation of Asp L213 and other carboxylic acid residues in the region may contribute to the stabilization of $Q_B^{\bullet-}$ in the charge separated state at low temperature.

Role of Ser L223 in Proton Transfer

The H-bond from Ser L223 plays a crucial role in proton transfer to $Q_B^{\bullet-}$ associated with electron transfer. The mutation of Ser L223 to Ala drastically reduces the rate of proton coupled electron transfer that has been proposed to be due blocking of proton transfer to $Q_B^{\bullet-}$ (39). The mechanism of proton transfer in biological systems is believed to occur through a chain of H-bonded groups in series via a Grothuss mechanism (6,51). The Grothuss mechanism consists of two steps 1) the transfer of protons across the H-bonds and 2) the rotation of the H-bond from one H-bond acceptor group to another to re-establish the starting H-bond pattern. The proton transfer to $Q_B^{\bullet-}$ can be seen as an extension of the proposed relaxation process in Figure 6. The rotation of the H-bond from Asp L213 to $Q_B^{\bullet-}$ is followed by proton transfer producing the protonated semiquinone $Q_BH^{\bullet}$. In the mechanism for proton coupled electron transfer to reduced Q_B (52,53), the protonated semiquinone is not a stable species but is an intermediate state, present with low probability, that serves as the acceptor for electron transfer from $Q_A^{\bullet-}$. The protonation step is crucial to establish a favorable driving force for electron transfer. Overall, the proton equilibrates rapidly and is not rate limiting. The rapid rate of proton transfer to $Q_B^{\bullet-}$ is greatly facilitated by the switching capability of the H-bond from Ser L223 (6,44, 54,55).

SUMMARY AND CONCLUSIONS

In this work we investigated the structural changes near the reduced quinone $Q_B^{\bullet-}$ that are associated with electron transfer to Q_B in RCs. ENDOR spectroscopy was used to study the H-bond environment near the reduced quinone $Q_B^{\bullet-}$ generated by B-branch electron transfer in the mutant Quint RC that are frozen in the dark (neutral, un-relaxed state) and in the light (charge-separated, relaxed state). The ENDOR spectra in the H-bonding region of $Q_B^{\bullet-}$ formed by B-branch electron transfer in mutant RCs were mostly similar to that in native RCs formed via the conventional A-branch, indicating that $Q_B^{\bullet-}$ was in the proximal active binding position in all cases. We observed one notable difference between the ENDOR spectra of RCs frozen in the light or in the dark. A peak in the H-bonding region of the ^1H -ENDOR spectrum was observed in Quint RCs frozen in the light but absent in Quint RCs frozen in the dark. The peak was assigned to an H-bond from Ser L223 based on its absence in Quint RCs carrying the additional mutation Ser L223 $\rightarrow$ Ala. From these results we conclude the H-bond from Ser L223 to $Q_B^{\bullet-}$ is formed upon Q_B reduction.

Although this switch in the H-bond position serves to stabilize the resulting semiquinone, other studies on mutant RCs lacking Ser L223 indicate that the formation of this H-bond is not solely responsible for stabilizing the charge separated state nor does its absence change the rate limiting conformational gating step of the Q_B reduction. Rather, the H-bond formation is one of many changes, including protonation events and protein rearrangements that contribute to charge stabilization and conformational gating. However, the switch of the H-bond from Ser L223 does play a key role in the proton transfer to Q_B . The movement of the Ser L223 H-bond from Asp L213 to $Q_B^{\bullet-}$ serves as the final step in the proton transfer process leading to the protonated semiquinone. Several unanswered questions arise from this study. These include - What is the rate of the fluctuation of the H-bond? Can the H-bond fluctuation be correlated with the gating of electron transfer to Q_B ? Future experiments using ENDOR spectroscopy can help to answer these questions.

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Abbreviations

RC	reaction center
D	primary donor, dimer of bacteriochlorophylls
Q_A	primary quinone electron acceptor
Q_B	secondary quinone electron acceptor
$k_{AB}^{(1)}$	rate of <u>first</u> electron transfer from $Q_A^{\bullet-}$ to Q_B
$k_{AB}^{(2)}$	rate of <u>second</u> electron transfer from $Q_A^{\bullet-}$ to $Q_B^{\bullet-}$

Figure 1.

Structure of cofactors imbedded within the RC protein matrix (PDB ID code 1AIG,8). The normal path for electron transfer to Q_B occurs predominantly along the A-branch from D to Q_A then Q_B (solid black arrows). The nearly symmetric cofactors along the B-branch are normally inactive in electron transfer. However, direct electron transfer to Q_B along the B-branch (red lines) can be observed in Quint RCs (B-branch mutant used in this work) which lack Q_A and have mutations that bias the electron transfer through the B-branch (7).

Figure 2.

Partial structure of the RC showing the catalytic Q_B site and the proton transfer pathways connecting it to the surface (PDB ID code 1AIG, 8) showing Q_B molecule (red); residues within H-bonding distance – His-L190 & Ser-L223 (yellow); residues that form part of the proton transfer pathways – Asp-L213 & Asp-L210 (yellow), Asp-M17 (blue), and His-H126 & His-H128 (green); and internal bridging water molecules (red spheres) identified in the DQ_B and/or $D^+Q_B^-$ structures (PDB ID codes 1AIG and 1AIJ). Also shown are part of the subunit backbones of the L subunit (yellow), M subunit (blue) and H subunit (green) and the non-heme Fe^{2+} (rust sphere) that is replaced with the diamagnetic Zn^{2+} in this work. Closest H-bonding partners are connected via dotted lines. The potential H-bonds to the backbone are labeled with the corresponding amino acid position, L224 and L225. Arrows represent the general flow of protons (H^+) through the pathways with the colors distinguishing different steps that occur during the photocyclic reduction of Q_B to Q_BH_2 . Ser-L223 and Asp-L213 (names highlighted in magenta) are key residues discussed in this work.

Figure 3.

35 GHz EPR spectra ($d\chi''/dH$) of $D^{+\bullet}Q_B^{-\bullet}$ in the dark frozen sample before and after illumination ($T=77K$). The EPR peak positions (principal g -values) are the same as frozen under illumination (not shown); the amplitude was decreased to 20% due to the minor fraction in which the $D^{+\bullet}Q_B^{-\bullet}$ state can be generated (see ref. 7). The RCs were chemically treated to replace the Fe^{2+} with Zn^{2+} to allow observation of $Q_B^{-\bullet}$ which would otherwise be broadened by the spin of the Fe^{2+} (34). At 35GHz, the low field region of the EPR spectrum arises from $Q_B^{-\bullet}$ (as labeled on left side of traces) and is spectrally separated from that of $D^{+\bullet}$ (as labeled in middle of traces); a Si marker line is located on far right. The lifetime of charge separated state was the same as observed in Fe^{2+} containing RCs and the same as observed at 9GHz (7). (Conditions: [RC] $\sim 100\mu M$, ID of EPR tube = 0.2cm, $\nu = 35GHz$, 77K, 16 traces were averaged.)

Figure 4.

ENDOR spectra of $Q_B^{\bullet-}$ in the B-branch mutant (QUINTUPLE) samples frozen under illumination (LIGHT), frozen in the dark and illuminated (DARK) and in a mutant containing the additional Ser-L223 $\rightarrow$ Ala (SAL223) replacement. (A) At the magnetic field position corresponding to g_y , proton nuclei that couple to the spin of $Q_B^{\bullet-}$ result in a two pairs of lines symmetrically shifted from the free proton frequency. The principle values of the hyperfine coupling interaction, $A_{\parallel}$ and $A_{\perp}$, is obtained from the frequency difference between symmetrically related lines (see text). At the bottom, the lime colored bars indicate the H-bonding region where peaks associated with H-bonds between $Q_B^{\bullet-}$ and the protein are observed; the blue-grey bar indicates the matrix region, which is composed of numerous more weakly coupled protons; an arrow indicates the free proton frequency, this correspond to the position at which a line resulting from an uncoupled proton would appear (i.e. with no hyperfine interaction). Note the overall similarity of the traces, in particular in the region in which peaks due to H-bonds are observed. The amplitude of the dark sample was increase 5-fold, since only a fraction of the sample undergoes charge separation, to allow a better comparison of the ENDOR signals. (B) Expansion of the ENDOR spectra of $Q_B^{\bullet-}$ focusing on left H-bonding region from part A. Note that the majority of the peaks (labeled in pairs 1 & 1', 2 & 2' and 3 & 3') are at the same position in all of the samples; the peaks are slightly broader in the sample frozen in the dark. This indicates that the binding position of $Q_B^{\bullet-}$ with respect to the local protein is the same in all of the samples. There is one exception indicated by an arrow and labeled 2 that is present only in the sample frozen under illumination (LIGHT); it is not present in the sample frozen in the dark (DARK) nor in the sample with the additional Ser- L223 $\rightarrow$

Ala replacement (SAL223). (Conditions: [Zn-RC(dQ)] ~ 0.7mM, ID of EPR tube = 0.2cm, ν = 35GHz, 77K, up to 80,000 traces were averaged.)

Figure 5.

ENDOR spectra of $Q_B^{\bullet-}$ and $Q_A^{\bullet-}$ in the native (R26) RC frozen under illumination. (A) Shown are the complete ENDOR spectra within 12 MHz of the free proton frequency. The top trace shows that observed (OBS) for a native sample excited with a single saturating laser flash. The Q_B site was occupied with deuterated UQ_{10} . This trace includes an admixture of some $Q_A^{\bullet-}$. The spectrum of $Q_A^{\bullet-}$, measured in a sample lacking Q_B , is shown in the second trace; due to the reconstitution procedure, the Q_A site contained almost exclusively the native (non-deuterated) UQ_{10} . The contribution of $Q_A^{\bullet-}$ was subtracted from the observed trace to yield the spectrum of $Q_B^{\bullet-}$ shown as the third trace (DIFF ($Q_B^{\bullet-}$)). The scaling factor for the amount of $Q_A^{\bullet-}$ in the observed trace was determined by matching the matrix region of the DIFF ($Q_B^{\bullet-}$) with that of the quintuple $Q_B^{\bullet-}$ (QUINT) obtained by freezing under illumination (from Fig. 4). Since the matrix peaks result from protons located further from the semiquinones, they should be less sensitive to possible local structural changes. For example, the peak near 52.5 MHz, which is a prominent matrix peak of $Q_A^{\bullet-}$, was used to determine the relative amount of $Q_A^{\bullet-}$ to subtract from the observed spectrum. The difference spectrum better matched other parts of the matrix region indicating that the subtraction was appropriate. Without knowledge of the quintuple $Q_B^{\bullet-}$, the scaling factor for the amount of $Q_A^{\bullet-}$ in the observed spectrum would have been difficult (if not impossible) to determine. (B) Expanded view of the left side emphasizing the H-bonds. Note the similarity of the peaks in the native $Q_B^{\bullet-}$ shown in the third trace from the top (DIFF ($Q_B^{\bullet-}$)) and those from the Quintuple mutant in which $Q_A^{\bullet-}$ cannot be formed. Highlighted with an arrow is the peak attributed to the hydroxyl group of Ser-L223 which is also present in the spectrum of the native sample. The small shift in position may

indicate a slight difference in distance of the magnitude of $\sim 0.02 \text{ \AA}$. Small shifts are also observed in the position of many of the smaller amplitude peaks. Overall, the native $\text{Q}_\text{B}^{\bullet-}$ is within uncertainty the same as the quintuple $\text{Q}_\text{B}^{\bullet-}$. In particular, the H-bond from Ser-L223 (labeled 2) is present in the native $\text{Q}_\text{B}^{\bullet-}$ spectrum. (Conditions: $[\text{Zn-RC(dQ)}] \sim 0.7 \text{ mM}$, ID of EPR tube = 0.2cm, $\nu = 35 \text{ GHz}$, 77K, up to 30,000 traces were averaged.)

Figure 6.

Model of the Q_B site to explain the observed low temperature (77K) ENDOR spectra. On the left is the model for active fraction of the frozen in the dark sample in which Q_B is bound in the proximal site with the protein in the ground state conformation, i.e. prior to charge separation. Upon illumination the D⁺Q_B^{•-} state is formed and detected with EPR and ENDOR spectroscopy. Since the sample is frozen, the normal protein relaxation to the formation of Q_B^{•-} is inhibited and is thus termed the *unrelaxed* conformation. On the right is the model for the sample frozen under illumination in which Q_B^{•-} is bound in the proximal site with the protein in the excited state conformation, i.e. following charge separation. Since the charge separated state is formed at higher temperatures (298K) the protein can respond to the charge separation prior to freezing and consequently the protein is in the *relaxed* conformation. The major difference observed in the ENDOR spectra is the H-bond between Ser-L223 and Q_B^{•-} which is present only in the relaxed conformation at 77K. Calculations suggest that this H-bond occurs when Asp-L213 becomes protonated (44,45). These changes indicate the formation of an H-bond network that could promote proton transfer as indicated by the arrows.