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D1-S169A Substitution of Photosystem II Perturbs Water Oxidation

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KEYWORDS: cyanobacteria; mutation; oxygen evolution; photosystem II; structure; substrate water

ABSTRACT: In photosystem II (PSII), photosynthetic water oxidation occurs at the tetramanganese-calcium cluster that cycles through light-induced intermediates ($S_0 - S_4$) to produce oxygen from two substrate waters. The surrounding hydrogen-bonded amino-acid residues and waters form channels that facilitate proton transfer and substrate water delivery, thereby ensuring efficient water oxidation. The residue D1-S169 lies in the “narrow” channel and forms hydrogen bonds with the Mn_4CaO_5 cluster via waters W1 and Wx. To probe the role of the narrow channel in substrate-water binding, we studied the D1-S169A mutation. PSII core-complexes isolated from mutant cells exhibit inefficient S-state cycling and delayed oxygen evolution. The S_2 state multiline EPR spectrum of D1-S169A PSII core-complexes differed significantly from that of wild-type and FTIR difference spectra showed that the mutation strongly perturbs the extensive network of hydrogen bonds that extends at least from D1-Y161 (Y_Z) to D1-D61. These results imply a possible role of D1-S169 in proton egress or substrate water delivery.

Introduction

Water oxidation, a key step in oxygenic photosynthesis, is catalyzed by photosystem II (PSII), a protein complex embedded in the thylakoid membranes of cyanobacteria, algae and higher plants.¹⁻⁶ As seen in the 1.9 Å crystal structure,⁷ the PSII complex is composed of around 20 protein subunits including D1, D2, CP43, CP47 and cytochrome b-559. The catalytic center for water oxidation is the oxygen-evolving complex (OEC), consisting of a Mn_4CaO_5 cluster ligated by water, hydroxo/oxo species and amino-acid residues. In each turn of the catalytic cycle, the OEC oxidizes two H_2O molecules, forming O_2 via a light-driven cyclic process that cycles the metal cluster through intermediate oxidation states (the so-called storage (S) states of oxidizing equivalents, S_n states with $n = 0 - 4$). Protons released to the lumen by this process establish a pH gradient across the thylakoid membrane while electrons extracted from water are used to reduce plastoquinone to plastoquinol.¹⁻⁶

S-state cycling is initiated by light-induced charge separation between the reaction center chlorophyll molecules, P_{680} , and a nearby pheophytin molecule. P_{680}^{+*} is subsequently reduced by the redox-active tyrosine residue, Y_Z . The oxidized Y_Z drives single electron oxidations of the OEC, forming the redox intermediates of the Mn_4CaO_5 cluster.¹⁻⁶ The S-state intermediates have been characterized by different biophysical techniques, including extended X-ray absorption fine structure spectroscopy (EXAFS), electron paramagnetic resonance (EPR) spectroscopy and Fourier transform infrared (FT-IR) spectroscopy.¹⁻⁶ EPR spectroscopy has been extensively used to characterize the S-state intermediates. The dark-stable S_1 state is integer spin and shows EPR signals in parallel mode, while the paramagnetic S_2 state can exist in the form of two spin isomers with effective spins of $S_T = 1/2$ and $5/2$, respectively.⁸

Under native conditions, in His-tagged *Synechocystis* PSII core complexes, an 18-22 multiline EPR signal is observed centered at $g = 2$. This multiline signal is characteristic of the $S_T = 1/2$ isomer where the Mn1, Mn2, Mn3 and Mn4 ions are in the III, IV, IV, IV oxidation states, numbered as in Figure 1.⁹ The light-driven oxidation of the S_3 state likely forms a transient S_4 intermediate that spontaneously evolves O_2 to regenerate the S_0 state. The water-oxidation mechanism is yet to be fully elucidated because of limited knowledge of the transient intermediates generated during the S_3 to S_0 transition.

Another unresolved aspect of the water-oxidation chemistry is the identification of the two substrate water molecules. Both are already bound in the S_2 state¹⁰ and one may relocate during the S_2 to S_3 transition. Though the structures of both the S_2 and S_3 states are well characterized, there is an ongoing debate regarding the mechanism of S_3 -state formation and the identity of the substrate water molecule that relocates. The hydrogen-bonding interactions with the neighboring waters and amino-acid residues are crucial in maintaining the pK_a of the substrate waters.³ Hence, a few point mutations have been generated to study a putative substrate-water delivery channel.¹¹ The identification of the substrate water molecules will help to resolve the two most debated mechanisms of oxygen formation: water-nucleophilic attack and oxo-oxyl radical mechanisms.^{12,13}

The OEC is surrounded by a hydrogen-bonded network of water molecules and amino-acid residues that have been classified into channels referred to as the large, broad and narrow channels.^{2,14,15} These channels are thought to play crucial roles in proton transfer to the lumen and substrate water delivery to the Mn cluster. Mutational analyses have been performed to characterize the proton-transfer pathway(s). These studies include mutations of D2-K317, D1-D61, D1-E65, D1-R334, and D1-E333 that resulted in inefficient S-state

cycling, delay of oxygen evolution and/or alteration of the decay kinetics of the S states.^{11,16-22} These residues lie in the broad channel that has been designated as a proton-exit channel that functions during at least the S₃ to S₀ transition.^{14,15,23} Further, it has been observed that modulation of the chloride-binding site present in the broad channel can also influence the proton-transfer process.²⁴ Ishikita and coworkers have further proposed that the narrow channel involving O4 is involved in deprotonation during S₀-S₁ transition.²⁵

There are differing proposals regarding the substrate water delivery pathway.³ Ammonia, a substrate water analogue, binds to Mn4 which is located near the terminus of the narrow channel.²⁶⁻²⁸ Further, another substrate water analogue, methanol binds in this channel near Mn4 and O4.²⁹⁻³² These results along with molecular dynamic simulations strongly suggest the narrow channel as the substrate water delivery channel. On the other hand, the water cavity in the large channel involving W3 was perturbed by the D1-V185N mutation.^{11,33} On the basis of these and other studies and computational analyses, the large channel has been proposed as the substrate water delivery channel with W3 acting as one of the substrate waters.^{11,33-37}

In this study, we investigate the substrate-water delivery pathway by perturbing the surrounding hydrogen-bonding network connecting the narrow channel around Mn4. The residue D1-S169 lies in the narrow channel and is hydrogen bonded to W1 (ligated to Mn4) through its backbone and Wx (ligated to O4) through its side-chain (Figure 1). Hence, to study the role of these waters and the narrow channel in the substrate-delivery process, we constructed the D1-S169A mutation. The characteristics of water oxidation exhibited by the D1-S169A mutated PSII provide new insight into the identity of the substrate waters and the mechanism of substrate water binding to the OEC during the S₂ to S₃ transition.

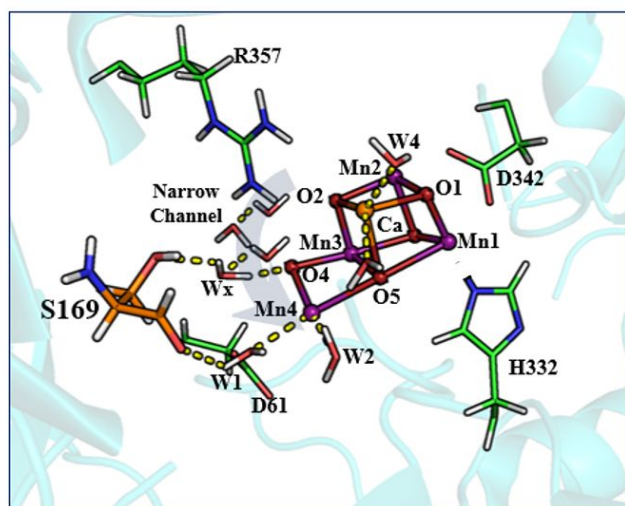


Figure 1. Structure of the OEC depicting the hydrogen-bonding network connecting D1-S169 (in orange), Wx and O4 in cyanobacterial PSII based on the crystal structure (PDB id: 3WU2).⁷ The Mn atoms are marked in purple, Ca in light orange, C atoms in green, N atoms in blue and O atoms in red.

Materials and Methods

Construction of Mutants and Isolation of PSII Core Complexes. The D1-S169A mutation was introduced into the psbA-2 gene of *Synechocystis* sp. PCC 6803 and transformed into a host strain of *Synechocystis* that lacks all three psbA genes and contains a hexa-histidine tag (His-tag) fused to the C-terminus of CP47.³⁸ The cultures were maintained and propagated as described previously.²⁰ Oxygen-evolving PSII core complexes were purified under dim green light at 4 °C with a Ni-NTA superflow affinity resin (Qiagen, Valencia, CA) as described previously.³⁹ The purified PSII core complexes were finally suspended in 1.2 M betaine, 10 % (v/v) glycerol, 50 mM MES-NaOH (pH 6.0), 20 mM CaCl₂, 5 mM MgCl₂, 50 mM histidine, 1 mM EDTA, and 0.03 % (w/v) n-dodecyl β-D-maltoside, frozen in liquid N₂, and stored at 77 K. To verify the integrity of the mutant cultures, sequencing of the relevant portion of the psbA-2 gene was performed after polymerase chain reaction amplification of genomic DNA.¹⁸ No traces of the wild-type codon were detected in any of the mutant cultures.

Steady-State Oxygen-Evolution Assays. Oxygen evolution from purified PSII core complexes of His-tagged wild-type and D1-S169A mutated PSII was monitored with a Clark-type electrode as described previously.¹⁸ The assay buffer contained 1 M sucrose, 50 mM MES-NaOH (pH 6.5), 50 mM CaCl₂ and 10 mM NaCl. 250 μM PPBQ and 1 mM K₃[Fe(CN)₆] were used as electron acceptors and 5 μg of Chl was used for each assay.

Polarographic Oxygen-Evolution Measurements. Flash-induced O₂ yields of PSII core complexes of His-tagged wild-type and the D1-S169A mutant were measured using a bare platinum electrode.⁴⁰ The samples were suspended in buffer containing 1 M sucrose, 10 mM CaCl₂, 200 mM NaCl, and 50 mM MES-NaOH (pH 6.5) as described previously.⁴⁰ 500 μM DCBQ and 1 mM K₃FeCN₆ were added as electron acceptors. The oscillations resulting from 20 flashes were fit to the VZAD model⁴¹ to obtain the corresponding miss parameters, hits and double hits. An interval spacing of 1 s was used to collect data for flash oxygen kinetics.

Electron Paramagnetic Resonance Studies. EPR samples were prepared in a buffer containing 50 mM MES-NaOH (pH 6.0), 1 M sucrose, 20 mM CaCl₂, 5 mM MgCl₂, and 1 mM EDTA and concentrated to 1 mg of Chl/mL using Amicon centrifugal cells having a 100 kDa cutoff. The measurements were performed using a Bruker ELEXSYS E500 spectrometer equipped with a SHQ resonator and an Oxford ESR-900 continuous flow cryostat at 7.5 K. The EPR parameters used for recording the spectra are as follows: microwave frequency, 9.38 GHz; modulation frequency, 100 kHz; modulation amplitude, 19.95 G; microwave power, 5 mW; sweep time, 84 s; conversion time, 41 ms; time constant, 82 ms. Each spectrum is the average of two scans. The dark scan of the EPR samples, concentrated to 1 mg of Chl/mL, corresponding to the S₁ state was recorded. The S₂ state was generated by illuminating the sample with a Xe lamp in a 200 K acetone/dry ice bath for 5 min and the spectrum was recorded.

FTIR Measurements. Samples were prepared as described previously¹⁸ except that they were maintained at a relative humidity of 95 % with droplets of 40 % (v/v) glycerol in water. Spectra were obtained as described previously²¹ with a Bruker Vertex 70 spectrometer (Bruker Optics, Billerica, MA) containing a pre-amplified, midrange D317 photovoltaic MCT detector (Kolmar Technologies, Inc., Newburyport, MA). For

each sample, the absorbance at 1657 cm^{-1} (amide I band) was 0.6 - 1.1. After pre-flashing and dark adaptation, samples were illuminated at $0\text{ }^{\circ}\text{C}$ with six flashes at 13 s intervals. Two transmission spectra (each consisting of 100 scans) were recorded before the first flash and one transmission spectrum was recorded after the first and subsequent flashes. The difference spectra of the successive S-state transitions (e.g., S_{n+1} -minus- S_n difference spectra, henceforth written $S_{n+1} - S_n$), were obtained by dividing the transmission spectrum obtained after the n th flash by the transmission spectrum obtained before the n th flash, then converting the ratio to units of absorption. The baseline stability/noise level was obtained by dividing the second pre-flash transmission spectrum by the first and converting the ratio to units of absorption (these spectra are labeled dark-dark in each figure). The sample was then dark-adapted for 30 min and the cycle was repeated. For each sample, the illumination cycle was repeated up to 15 times. The spectra of 20 - 22 samples were averaged (see figure legends).

QM/MM Calculations. The QM/MM computational model of the oxygen-evolving complex (OEC) was constructed as described in previous work.^{42,43} Residues in the model included those with C_{α} within 15 \AA of the OEC. The two chloride anions and all water molecules whose oxygen atoms fall within this boundary were also included. Cut residues were capped with neutral backbone fragments (ACE/NME) whose position was dictated by the location the neighboring residues.

Residues included in the model are listed as follows, with capping residues only including the backbone atoms indicated by (parenthesis):

D1 (chain A): (57)-58-67-(68), (81)-82-91-(92), (107)-108-112-(113), (155)-156-192-(193), (289)-290-298-(299), (323)-324-344:C-terminus;

CP43 (chain C): (290)-291-(292), (305)-306-314-(315), (334)-335-337-(338), (341)-342-(343), (350)-351-358-(359), (398)-399-402-(403), (408)-409-413-(414);

D2 (chain D): (311)-312-321-(322), (347)-348-352:C-terminus.

The model was optimized as a two-layer QM/MM model using the ONIOM⁴⁴ method as implemented in the Gaussian09⁴⁵ software suite. The QM layer included the OEC, surrounding residues (D1-D170, D1-E189, D1-H332, D1-E333, D1-D342, CP43-E354, the C-terminus of D1-A344, D1-H337, CP43-R357, and D1-D61), and 10 bound water molecules. Clipped sidechains were modeled as described previously.⁴³ The QM layer was calculated at the B3LYP^{46,47} level of theory, with the LanL2DZ⁴⁸ basis set and pseudopotential used for Mn and Ca atoms, 6-31G⁴⁹ for C, N, and H, and 6-31G(d)⁵⁰ for O. The lower MM layer was calculated using the AMBER⁵¹ force field. All atoms were allowed to relax, excluding the capping residues and the chloride and oxygen atoms of waters in the MM layer.

Results

D1-S169A cells are photoautotrophic and evolve oxygen at $77 \pm 7\%$ the rate of wild-type cells under light-saturated conditions (Table S1). The PSII content of D1-S169A cells was estimated to be $111 \pm 5\%$ compared to wild-type cells on the basis of measurements of the maximum fluorescence yields ($F_{\max} - F_0$) of wild-type and D1-S169A cells (Figure S1 and note S1). It was estimated that 10 - 14% of the PSII reaction centers

in D1-S169A cells lack Mn_4CaO_5 clusters *in vivo* on the basis of measurements of the kinetics of charge recombination between Q_A^- and the donor side of PSII following 5 s of illumination in the presence of DCMU (Figure S2 and note S1). The presence of PSII reaction centers lacking Mn_4CaO_5 clusters implies that the cluster is assembled less efficiently or is less stable in D1-S169A cells compared to wild-type cells. The active fraction of PSII reaction centers in D1-S169A cells has a lower light-saturated rate of oxygen evolution compared to wild-type cells, especially when normalized to the higher PSII content of D1-S169A cells. If O_2 comes from 85-90% of the mutant PSII core complexes in the cells (see above), the normalized O_2 activity relative to WT would be 76-81%, implying that the efficiency of the S-state cycle is somewhat impaired in the mutant. In D1-S169A cells, the kinetics of charge recombination between Q_A^- and the donor side of PSII following a single flash given in the presence of DCMU were slightly slower than those of wild-type cells, implying a slight decrease in the S_2/S_1 midpoint potential in the mutant cells (Figure S2 and Table S1).

PSII core-complexes isolated from the D1-S169A mutant cells exhibited a maximal light-saturated oxygen-evolution rate of $1330 \pm 28\text{ }\mu\text{mol of O}_2 / (\text{mg of Chl}\cdot\text{hr})$, approximately 47% that of wild-type PSII. The lower relative activity compared to intact cells may be caused by loss of Mn_4CaO_4 clusters during purification or by a further diminished efficiency of S-state cycling under the conditions of the measurements of the D1-S169A PSII core complexes. To characterize the turnover efficiency of D1-S169A PSII core complexes, we compared the flash O_2 -yield patterns of wild-type and D1-S169A PSII core complexes in response to 20 saturating flashes given at $23\text{ }^{\circ}\text{C}$. These patterns show highly damped flash-induced oscillations for D1-S169A PSII core complexes. The miss probability (α) is 2.2 times higher than that of wild-type PSII core-complexes (Figure 2) showing decreased S-state cycling efficiency. Further, we observed a delay in the oxygen-evolution kinetics for D1-S169A PSII core-complexes (Figure 3). Taken together, these studies indicate that the D1-S169A mutation diminishes the efficiency of water oxidation.

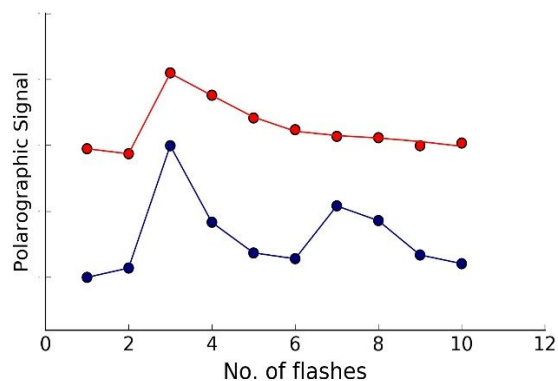


Figure 2. Comparison of the flash oxygen-yield patterns of PSII core complexes from: His-tagged wild-type *Synechocystis* (blue) and D1-S169A (red). The data from wild-type and D1-S169A PSII are offset vertically for clarity. The oscillations resulting from 20 flashes at $23\text{ }^{\circ}\text{C}$ were fit to the VZAD model to obtain the corresponding miss parameters, hits and double hits.⁴¹ Table 1 shows the Kok-model fitting parameters for wild-type and D1-

S169A PSII core complexes. Average values from 3 independent experiments are reported. The standard errors are calculated. The S-state percentages have standard errors of $\pm 5\%$.

Table 1. Kok-model fitting parameters

Parameter	Wild-type	D1-S169A
Misses, α (%)	14 \pm 1	31 \pm 2
Double hits, β (%)	3 \pm 1	4 \pm 1
S ₀ (%)	15	2
S ₁ (%)	52	64
S ₂ (%)	21	14
S ₃ (%)	13	20

FTIR characterization. The mid-frequency FTIR difference spectra produced by the first, second, third, and fourth flashes given to wild-type PSII core complexes correspond predominantly to the S₂ - S₁, S₃ - S₂, S₀ - S₃, and S₁ - S₀ FTIR difference spectra, respectively.^{22,52,53} Spectra from wild-type and D1-S169A PSII core complexes are compared in Figure 4. The amplitudes of the D1-S169A spectra produced by the first and second flashes are considerably lower than those of the corresponding spectra of wild-type (upper two sets of traces) on a protein basis (the spectra were normalized to the amplitudes of their absolute absorbances at 1657 cm⁻¹) but are recognizable as corresponding predominantly to S₂ - S₁ and S₃ - S₂ spectra, respectively. The D1-S169A spectra produced by the third and fourth flashes (lower two red traces in Figure 4) exhibit few recognizable features. The indistinct nature of the mutant spectra produced by the third and fourth flashes may reflect the much higher miss parameter measured polarographically in D1-S169A PSII core complexes.

The mid-frequency S₂ - S₁ spectrum of D1-S169A showed substantial changes compared to wild-type throughout the mid-frequency region. In the amide I region, a negative feature appeared at 1641 cm⁻¹ and the 1680(-) cm⁻¹ and 1622(+) cm⁻¹

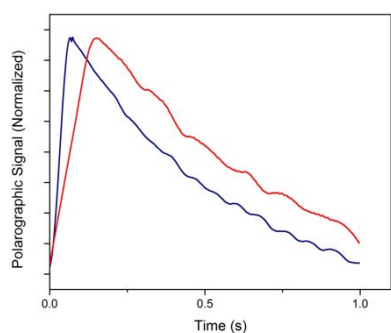


Figure 3. Comparison of the O₂-release kinetics of D1-S169A (red) and wild-type (blue) PSII core complexes. The peak values for oxygen evolution are normalized to facilitate comparison. The S169A core complexes exhibited a maximal oxygen-evolution rate of 1330 \pm 28 $\mu\text{mol of O}_2 / (\text{mg of Chl} \cdot \text{hr})$, approximately 47 % that of wild-type PSII.

features were eliminated or nearly eliminated, respectively. These features can be assigned to amide I modes because they are sensitive to global ¹³C labeling but not to global ¹⁵N labeling.^{54,55} In the asymmetric carboxylate stretching [$\nu_{\text{asym}}(\text{COO}^-)$] and amide II region, the 1587(+) cm⁻¹ feature was nearly eliminated, the 1544(-) cm⁻¹ feature was diminished, and the 1530(+)/1523(-) cm⁻¹ derivative feature was abolished. The 1587 cm⁻¹ feature corresponds to a $\nu_{\text{asym}}(\text{COO}^-)$ mode because it is sensitive to global ¹³C labeling^{54,56-58} but not to global ¹⁵N labeling.^{16,54,56,58,59} The 1544(-) and 1530(+)/1523(-) cm⁻¹ features correspond to amide II modes because they are sensitive to both global ¹³C labeling^{54,56-58} and global ¹⁵N labeling.^{16,54,56,58,59} In the symmetric carboxylate stretching [$\nu_{\text{sym}}(\text{COO}^-)$] region, the 1415(-)/1410 (+) cm⁻¹ derivative feature was eliminated and the 1400(-) cm⁻¹ feature was upshifted by 5 cm⁻¹. Finally, in the carbonyl stretching [$\nu(\text{C=O})$] region, the 1746(-) cm⁻¹ feature was diminished. The mid-frequency S₃ - S₂ spectrum of D1-S169A also showed changes throughout the mid-frequency region. The 1675(-)/1666(+) cm⁻¹ derivative feature in the amide I region was eliminated, the 1538(+) cm⁻¹ feature in the $\nu_{\text{asym}}(\text{COO}^-)$ /amide II region was eliminated, the 1422(-)/1413(+) cm⁻¹ derivative feature in the $\nu_{\text{sym}}(\text{COO}^-)$ region was eliminated and the 1363(-) cm⁻¹ and 1344(+) cm⁻¹ features were upshifted slightly.

The O-H stretching vibrations of strongly H-bonded OH groups can be observed as very broad positive features between 3200 and 2500 cm⁻¹.^{21,22,60,61} These regions of the S_n - S_{n-1} difference spectra of D1-S169A PSII core complexes are compared with wild-type PSII core complexes in Figure 5A. In the S₂ - S₁ spectrum, the broad feature is overlain with positive features that have been attributed to the N-H stretching vibration of the protonated D1-H337 side chain.⁶² The broad feature of the S₂ - S₁ spectrum was substantially altered by the D1-S169A mutation: much of the amplitude in the 3000-2700 cm⁻¹ region appeared downshifted to between 2700-2400 cm⁻¹ (upper pair of traces in Figure 5A).

In contrast, the broad feature of the S₃ - S₂ spectrum was largely unaltered by the mutation. The apparent mutation-induced alterations to the broad features produced by the third and fourth flashes may arise from significant contributions of the mutant S₂ - S₁ and S₃ - S₂ spectra to these spectra because of the high miss parameter in D1-S169A PSII core complexes.

The O-H stretching vibrations of weakly hydrogen-bonded OH groups of water molecules can be observed between 3700 and 3500 cm⁻¹.^{22,52,53,61} In the S₂ - S₁ spectrum, the D1-S169A mutation eliminated the small 3663(-) cm⁻¹ and large 3617(+) cm⁻¹ features and shifted the 3584(-) cm⁻¹ feature to 3579 cm⁻¹ (Figure 5B, upper traces). The mutation had no apparent effect on the broad negative feature in the S₃ - S₂ spectrum (Figure 5B, second set of traces). Any apparent mutation-induced alteration to the broad features produced by the third and fourth flashes may arise from significant contributions of the mutant S₂ - S₁ and S₃ - S₂ spectra to these spectra because of the mutated sample's high miss parameter.

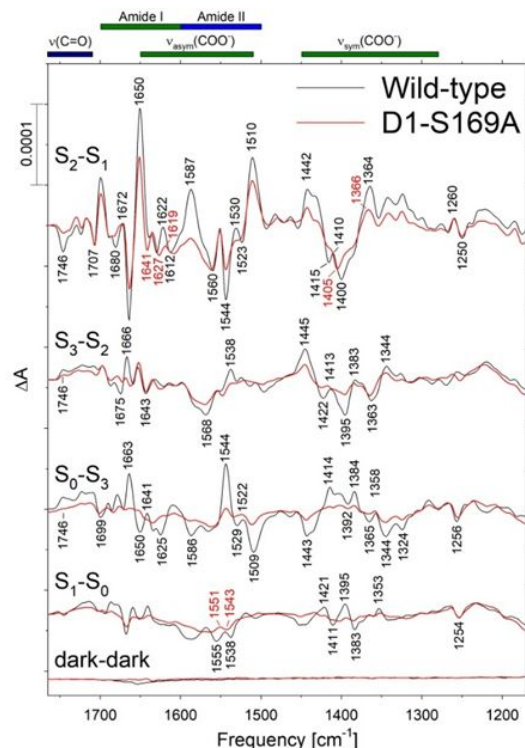


Figure 4. Comparison of the mid-frequency FTIR difference spectra of wild-type (grey) and D1-S169A (red) PSII core complexes in response to four successive flash illuminations applied at 0 °C. The data (plotted from 1770 cm^{-1} to 1170 cm^{-1}) represent the averages of 22 wild-type and 20 D1-S169A samples (33,000 and 29,600 scans, respectively). The spectra have been normalized to the average absolute amplitudes of the samples at the amide I peak at 1657 cm^{-1} . Dark-dark traces show the noise level and the stability of the baseline.

The fraction of PSII containing functional Mn_4CaO_5 clusters in PSII core-complexes was estimated from the $S_2 - S_1$ FT-IR difference spectra. The lower amplitudes of the peaks at 1664(-), 1650(+), 1510(+), 1442(+), and 1400(-) in the D1-S169A spectrum suggest that approximately 60% of the mutant PSII core complexes contain Mn_4CaO_5 clusters capable of reaching the S_2 state. On the basis of the amplitude of the O_2 flash yield on the third flash (Figure 2), we estimate that 67% of mutant PSII core complexes contain Mn_4CaO_5 clusters capable of making O_2 . The steady-state O_2 forming rate of the mutant PSII cores was 47% compared to WT. Consequently, the normalized O_2 activity of D1-S169A PSII core-complexes relative to WT would be 70-78%, similar to the 76-81% estimate in D1-S169A cells (see above) suggesting that the lower efficiency of O_2 evolution is similar in both cells and PSII core complexes.

EPR characterization To analyze the structure of the OEC, cw-EPR spectra of the S_2 states of both D1-S169A and wild-type PSII core complexes were collected (Figure S3, Figure 6). The S_2 -state spectrum of wild-type PSII exhibits an 18 - 22 multilines EPR signal centered at $g = 2$ corresponding to the $S_T = 1/2$ spin state isomer. This multilines signal has an average hyperfine line spacing of ~ 87.5 G consistent with previously reported values in the literature.⁶³

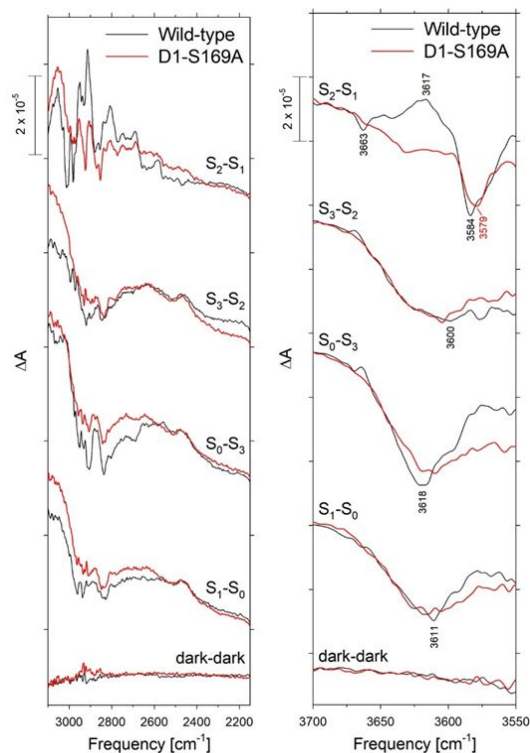


Figure 5. (A) Comparison of the strongly hydrogen bonded O-H stretching region of wild-type (grey) and D1-S169A (red) PSII core complexes in response to four successive flash illuminations applied at 0 °C. The data were collected simultaneously with those in Figure 4. (B) Comparison of the weakly hydrogen-bonded O-H stretching region of wild-type and D1-S169A PSII core complexes in response to four successive flash illuminations applied at 0 °C. The data were collected simultaneously with those in Figure 4.

The cw-EPR spectrum of the S_2 state of D1-S169A PSII also exhibits a multilines signal centered at $g = 2$. However, the average hyperfine line spacing of the multilines signal is reduced to ~ 72 G, which suggests alterations in the structure of the OEC.

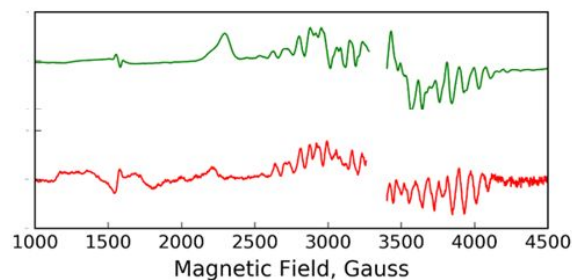


Figure 6. Comparison of the $S_2 - S_1$ state difference EPR spectra of wild-type (green) and D1-S169A PSII core complexes (red). The unsubtracted spectra are reported in the Supplementary Information (Figure S3).

QM/MM calculations on the S_1 state of D1-S169A. The D1-S169 residue is part of the narrow channel, and therefore

substitution of this residue can affect the substrate water delivery to OEC. The QM/MM minimized structure of the D1-S169A mutant in the S_1 state revealed that the elimination of the hydrogen bond between D1-S169 and Wx caused Wx to shift by 0.5 Å and W195 by 0.3 Å towards the narrow channel. We also see slight movement in CP43-R357 and D1-D61 (0.18 Å and 0.12 Å, respectively) as the two residues equilibrate to the new location of Wx (Figure 7). A disruption in the substrate-water delivery is consistent with the observed impairment of S-state cycling in D1-S169A PSII as well as delayed oxygen evolution.

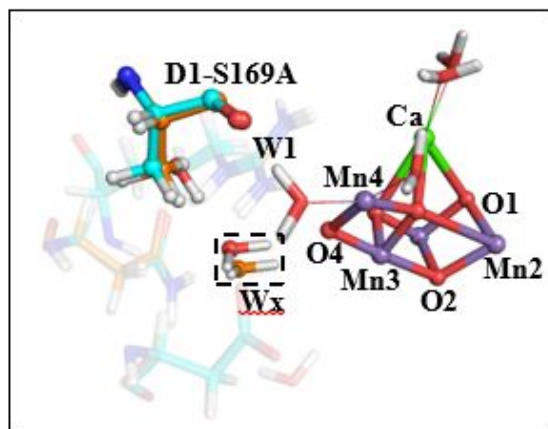


Figure 7. Superposition of the QM/MM optimized structures of wild-type (blue) and D1-S169A PSII (orange) in the S_1 state. The dashed box indicates the shift in Wx (red, in wild-type PSII).

Discussion

The narrow-channel has been proposed previously to be the substrate-water delivery channel²⁹ and to play an important role in proton egress.²⁵ Hence, we substituted a narrow channel residue D1-S169 with A169 to study the effect of perturbation of the hydrogen-bonding network involving Wx, W1, O4 and Mn4. PSII core complexes isolated from D1-S169A cells exhibit reduced oxygen evolution and impaired S-state cycling. Similar characteristics have often been displayed by PSII core-complexes of secondary-shell mutants including D2-K317A,¹⁶ D1-D61A,²¹ CP43-R357K⁶⁴ and D1-N87D.²⁴ These perturbations were caused by alterations to the hydrogen-bonding network around the OEC, presumably resulting in hindered proton transfer.^{19-22,29,33}

The substantial D1-S169A mutation-induced changes to the mid-frequency $S_2 - S_1$ FTIR difference spectrum show that the D1-S169A mutation alters the response of carboxylate groups and the protein backbone to the positive charge that develops on the Mn_4CaO_5 cluster during the S_1 to S_2 transition. Similar alterations are produced by mutations of numerous residues that are located in the networks of hydrogen bonds that surround the Mn_4CaO_5 cluster. Examples include D1-D61A,²¹ D1-E65A,¹⁹ D1-N87A,²⁴ D1-Q165E,²⁰ D1-N181A,¹⁸ D1-V185N,³³ D1-N298A,⁶⁵ D1-R334A,²⁰ D2-E312A,¹⁹ and D2-K317A.¹⁶ The diminished amplitudes of the carbonyl, carboxylate, and amide II features at 1746(-) cm^{-1} , 1587(+) cm^{-1} and 1544(-) cm^{-1} , respectively, in the $S_2 - S_1$ FTIR difference spectrum are also produced by the D1-D61A, D1-E65A, D1-V185N, D1-N298A, D2-E312Q, and D1-R334A mutations.^{19-21,33,65} These spectral

changes may reflect similar perturbations of the polypeptide backbone that are caused by disruption of the same network of hydrogen bonds. The 1530(+)/1522(-) cm^{-1} feature in the $S_2 - S_1$ difference spectrum that is eliminated by the D1-S169A mutation is also eliminated by the D1-D61A,²¹ D1-E65A,¹⁹ D1-N181A,¹⁸ D1-V185N,³³ D1-N298A,⁶⁵ D1-R334A,²⁰ D2-E312A,¹⁹ and D2-K317A mutations.¹⁶ Elimination of the same feature by mutations constructed at these eight residues implies that each mutation partly disrupts a common network of hydrogen bonds that includes the nearby Cl^- ion. The spectral changes produced by the D1-S169A mutation in the mid-frequency region are, therefore, consistent with the participation of D1-S169 in a network of hydrogen bonds that forms a water entry channel leading to the Mn_4CaO_5 cluster and/or a proton exit pathway.

In the strongly hydrogen-bonded O-H stretching region, the broad positive feature observed in the $S_2 - S_1$ spectrum between 3100 cm^{-1} and 2400 cm^{-1} has been assigned to the coupled O-H stretching vibrations of strongly hydrogen-bonded water molecules in an extended network of hydrogen bonds that links D1-D61 with the Ca^{2+} ion and Y_Z .⁶¹ This feature is dominated by the O-H stretching vibrations of W1 and W2⁶¹ and is eliminated by the D1-D61A mutation.²¹ That D1-S169A also alters this feature implies that D1-S169 participates in the same hydrogen-bonding network. Such participation would be consistent with the 1.9 Å X-ray crystallographic structures; in these structures, D1-S169 forms a hydrogen bond with a water molecule that also forms hydrogen bonds with D1-D61 and the O4 oxo bridge of the Mn_4CaO_5 cluster.^{7,66,67}

In the weakly hydrogen-bonded O-H stretching region, the features observed in the $S_2 - S_1$ spectrum between 3700 cm^{-1} and 3500 cm^{-1} have been assigned to the coupled O-H stretching vibrations of weakly hydrogen-bonded water molecules in the same extended network of hydrogen bonds mentioned above.⁶¹ The small 3663(-) cm^{-1} feature that is eliminated by the D1-S169A mutation is also eliminated by the D1-D61A,²¹ D1-N298A,⁶⁵ and D1-E333Q⁶⁸ mutations. The feature is also diminished by the D1-N181A mutation.¹⁸ The larger 3617(+) cm^{-1} feature that is eliminated by the D1-S169A mutation is also eliminated by the D1-D61A,²¹ D1-N87A,²⁴ and D1-V185N³³ mutations and is diminished by the D1-N181A¹⁸ mutation and also by the substitution of Sr^{2+} for Ca^{2+} .³⁴ Consequently, the spectral features altered by the D1-S169A mutation in the strongly and weakly hydrogen bonded O-H stretching regions are consistent with D1-S169 participating in an extensive network of hydrogen bonds involved in water entry and/or proton egress.

To understand the changes in the structure of the OEC, EPR studies were carried out. EPR studies demonstrated that the S_2 state multiline EPR signal was altered in D1-S169A PSII. The lower intensity of the multiline EPR signal in D1-S169A PSII is a consequence of both the lower fraction of PSII centers having assembled/functional metal clusters in the mutant and the altered hyperfine splittings in the mutant. Similar alterations to the S_2 -state multiline EPR signal have been observed in PSII depleted of Ca^{2+} ,⁶⁹ in PSII having Ca substituted by Sr,^{70,71} in ammonia-treated PSII,^{26-28,72} in PSII from the D1-H332E mutant,⁷³ and in PSII from the D1-D61A mutant PSII,²⁶ another secondary shell mutation. Pulsed EPR studies have shown that these spectral changes are caused by slight alterations to the

environment of the S₂ state's single Mn(III) ion, caused in turn by treatment/mutation-induced alterations to the magnetic couplings between the individual Mn ions.^{28,69,71,73} The alterations to the multiline signal caused by the D1-D61A mutation can be attributed to perturbations of hydrogen bonds involving Mn⁴. The alterations caused by the D1-S169A mutation likely have the same origin. Such an interpretation is in good agreement with the QM/MM analyses which show that the waters in the narrow-channel ligated to the cluster (W_x) are affected as a result of the D1-S169A site-directed mutation. It has been shown in a recent publication that the waters in the narrow channel in wild-type PSII remain unperturbed, except W20 (here the water hydrogen bonded to W_x) which disappears in the S₂ and S₃ states.⁷⁴ However, in the D1-S169A mutant, we do observe movement of waters in the narrow channel in the S₁ state which is possibly a consequence of the mutation. Hence, based on these results it will be interesting to further probe the disappearance of W20 and understand its role in the substrate water delivery mechanism. In our previous studies, we have observed intricate connections between the channels,²⁴ and thereby we do not negate the possibility that the mutations in the narrow channel result in hindrance of substrate water delivery through the large channel. The impaired S-state cycling and the delayed-oxygen kinetics can be the consequence of a hindered substrate water channel, changes in the substrate water binding mechanism, or partial disruption of proton egress. Further studies involving pulsed EPR techniques will be required to provide deeper insight into the nature of the perturbation of the Mn₄CaO₅ cluster by the D1-S169A mutation.

ASSOCIATED CONTENT

Supporting Information. Measurement of chlorophyll fluorescence and EPR spectra of the S₂ state and dark spectra, S₂ state decay curves. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Author Contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

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