

Identification of Oxidized Amino Acid Residues in the Vicinity of the Mn_4CaO_5 Cluster of Photosystem II: Implications for the Identification of Oxygen Channels within the Photosystem

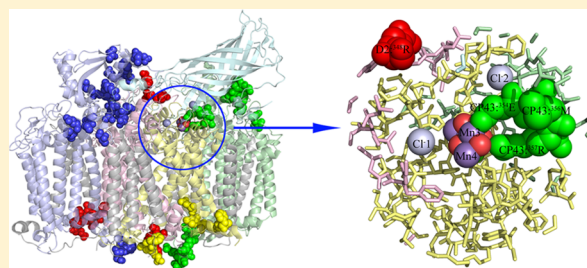
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S Supporting Information

ABSTRACT: As a light-driven water–plastoquinone oxidoreductase, Photosystem II produces molecular oxygen as an enzymatic product. Additionally, under a variety of stress conditions, reactive oxygen species are produced at or near the active site for oxygen evolution. In this study, Fourier-transform ion cyclotron resonance mass spectrometry was used to identify oxidized amino acid residues located in several core Photosystem II proteins (D1, D2, CP43, and CP47) isolated from spinach Photosystem II membranes. While the majority of these oxidized residues (81%) are located on the oxygenated solvent-exposed surface of the complex, several residues on the CP43 protein (³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R) which are in close proximity (<15 Å) to the Mn_4CaO_5 active site are also modified. These residues appear to be associated with putative oxygen/reactive oxygen species exit channel(s) in the photosystem. These results are discussed within the context of a number of computational studies which have identified putative oxygen channels within the photosystem.



Photosystem II (PS II) functions as a light-driven, water–plastoquinone oxidoreductase. In higher plants and cyanobacteria at least six intrinsic proteins appear to be required for O_2 evolution. These are CP47, CP43, D1, D2, and the α and β subunits of cytochrome b_{559} . Deletion of these subunits uniformly results in the loss of PS II function and assembly.^{1,2} Additionally, in higher plants, three extrinsic proteins, PsbO, PsbP, and PsbQ, are also required for maximal rates of O_2 evolution under physiological inorganic cofactor concentrations.³ Of these three proteins, the PsbO protein appears to play a central role in the stabilization of the manganese cluster, is essential for efficient and stable O_2 evolution, and is required, along with PsbP, for photoautotrophic growth and PS II assembly in higher plants propagated under normal growth conditions.^{4–6} Under low light growth conditions the PsbQ component is also required for photoautotrophy.⁷ Over the past 11 years, moderate resolution crystal structures of cyanobacterial PS II have significantly enhanced our understanding of the molecular organization of the constituent polypeptides of the photosystem and the active site for oxygen evolution, the $\text{Mn}_4\text{O}_5\text{Ca}$ cluster.^{8–12} Recently, a high-resolution 1.9 Å crystal structure of cyanobacterial PS II has been presented.¹³ It should be noted that no crystal structures for higher plant PS II are available. While there are differences between the higher plant and the cyanobacterial photosystems, particularly with respect to the identity and organization of the extrinsic proteins,³ the amino

acid sequences of the intrinsic components (D1, D2, CP43, and CP47) are nearly identical (>85% similarity¹⁴). Consequently, one would expect that the structural organization of these components within PS II would be highly homologous between higher plants and cyanobacteria.

The production of molecular oxygen by PS II is accompanied by the unavoidable possibility of oxidative modification of amino acid residues within the PS II complex. While one would predict that amino acid residues located at the surface of the complex which are in contact with the bulk, oxygenated solvent would be susceptible to such modification, buried amino acid residues in the vicinity of the Mn_4CaO_5 cluster, the active site for water oxidation, might also be particularly susceptible to oxidative modification. This is exacerbated by the production of reactive oxygen species (ROS), such as H_2O_2 , $\text{O}_2^{\bullet-}$, and OH^{\bullet} at, or in the vicinity of, the Mn_4CaO_5 cluster under a variety of stress conditions.¹⁵ Indeed, earlier studies have identified the double oxidation of the CP43 residue ³⁶⁵W to form an *N*-formylkynurenine derivative.^{16,17} This residue is located 17 Å from Mn2 of the Mn_4CaO_5 cluster.

Given the possibility that oxidative modification of PS II could lead to damage of the photosystem, several groups have

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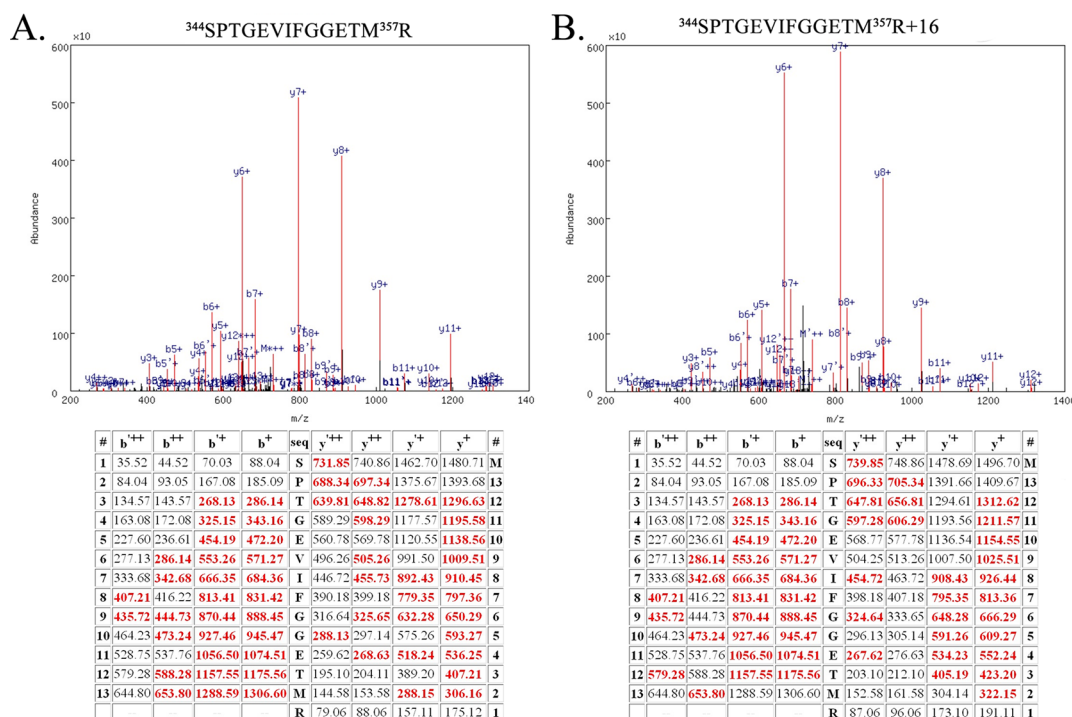


Figure 1. Example mass spectrometry data from the unmodified peptide $^{344}\text{SPTGEVIFGGETM}^{357}\text{R}$ and the oxidatively modified peptide $^{344}\text{SPTGEVIFGGETM}^{357}\text{R}+16$. (A) Top: spectrum of the CID dissociation of the unmodified peptide $^{344}\text{SPTGEVIFGGETM}^{357}\text{R}$. Various identified ions are labeled. Bottom: table of all predicted masses for the y- and b-ions generated from this peptide sequence. Ions identified in the CID spectrum (above) are shown in red. The b^{+2} , b^{+1} y^{+2} , and y^{+1} ions are generated by the neutral loss of water. (B) Top: spectrum of the CID dissociation of the modified peptide $^{344}\text{SPTGEVIFGGETM}^{357}\text{R}+16$. Various identified ions are labeled. Bottom: table of all predicted masses for the y- and b-ions generated from this peptide sequence. Ions identified in the CID spectrum are shown in red. The b^{+2} , b^{+1} y^{+2} , and y^{+1} ions are generated by the neutral loss of water. Note the additional mass of 16 amu (in comparison with part A) arising from modification of ^{357}R . This modification is evident in all identified y-ions species but is absent in all identified b-ions. Both of these spectra are from Biological Replicate #1.

hypothesized that channels present in PS II would function to vector molecular oxygen and, by extension, ROS away from the Mn_4CaO_5 cluster toward the surface of the photosystem. Murray and Barber¹⁸ used the CAVER Program¹⁹ to examine the 3.5 Å crystal structure of *T. elongatus*,¹⁰ later examining similar PS II crystals perfused with xenon.²⁰ Gabdulkhakov et al.²¹ used noble gas and dimethyl sulfoxide cocrystallization studies in combination with CAVER to examine the 2.9 Å structure of *T. elongatus*,¹² and Ho and Styring calculated solvent-accessible surfaces for the 3.0 Å *T. elongatus* structure of Loll et al.¹¹ We hypothesized that amino acid residues in contact with such oxygen channels in the interior of PS II would be significantly more susceptible to oxidative amino acid modification than residues not exposed to molecular oxygen and/or ROS. Consequently, the identification of such oxidatively modified residues in the interior of PS II should serve to complement and extend the largely computational studies mentioned above.

MATERIALS AND METHODS

PS II membranes were isolated from market spinach by the method of Berthold et al.²² modified by Ghanotakis and Babcock.²³ After isolation, the PS II membranes were suspended at 2 mg chl/mL in 50 mM Mes-NaOH, pH 6.0, 300 mM sucrose, 15 mM NaCl buffer, and frozen at -80°C . The proteins in the samples were resolved on a 12.5–20% acrylamide gradient by LiDS-PAGE using a nonoxidizing gel system.²⁴ This was required, as standard PAGE is known to introduce numerous protein oxidation artifacts.^{24,25} In the

nonoxidizing system the gels are polymerized with riboflavin (in the presence of diphenyliodonium chloride + toluenesulfonate) followed by exposure to UV light. The upper reservoir contains thioglycolate. Preliminary experiments indicated that proteins resolved in this system exhibited extremely low levels of artifactual protein oxidation when compared to proteins resolved under standard PAGE conditions (Figure S1). After electrophoresis, the gels were stained with Coomassie Blue and destained, and protein bands containing CP47, CP43, D1, and D2 were excised. These were then processed for trypsin digestion using standard protocols. In some cases, the tryptic peptides were processed using a C18 ZipTip prior to mass analysis.

Chromatography was performed using a Finnigan Surveyor MS pump and a Finnigan Micro AS autosampler. The column used was a Water's X-Bridge C18 3.5 μm 2.1 \times 100 mm with a flow rate of 200 $\mu\text{L}/\text{min}$. The mobile phases consist of a 95:5 water:acetonitrile with 0.1% formic acid aqueous phase and a 95:5 acetonitrile:water with 0.1% formic acid organic phase. The gradient was as follows: The organic phase composition was 10% for the first 5 min, ramped to 20% for the next 10 min (total time 15 min), ramped to 50% for the next 25 min (total time 40 min), ramped to 80% in the next 35 min (total time 75 min), and held at 80% for 10 min followed by a quick ramp to 10% in 5 min and a 10 min hold to equilibrate the column (total time 100 min).

Mass spectrometry was performed on a Thermo Scientific LTQ-FT, a hybrid instrument consisting of a linear ion trap and a Fourier transform ion cyclotron resonance mass spectrom-

Table 1. Oxidatively Modified Residues in the D1, D2, CP43, and CP47 Proteins^a

D1	² T ^b + stcb or go, ³ A ^b + go, ⁵ L ^b + go, ⁶ E ^b + gad, ¹⁷ F + go, ¹⁸ C + cysh, ¹³⁰ E ^c + go, ¹³³ L ^c + go, ¹³⁵ F ^c + go, ²³⁹ F + go, ²⁴¹ Q + ca, ²⁴² E + gad
D2	¹³ K + go, ¹⁴ D + gad, ¹⁵ L + ca, ¹⁶ F + go, ¹⁹ M + go, ²⁰ D + ca, ²¹ D + gad, ²³⁸ P + ca, ²³⁹ T + go, ²⁴² E + gad, ²⁴⁷ M + go, ³⁴⁵ E + gad, ³⁴⁷ L + go, ³⁴⁹ R + ca
CP43	²⁷ D + go, ²⁸ Q + ca, ²⁹ E + gad, ¹⁹⁵ D + gad, ¹⁹⁶ V + ca, ³⁵⁴ E ^d + gad, ³⁵⁵ T ^d + go, ³⁵⁶ M ^d + aa or go, ³⁵⁷ R ^d + ca. or go, ³⁶⁷ E + gad, ³⁶⁸ P + go, ³⁶⁹ L + ca, ³⁷⁵ L + ca, ³⁷⁶ D + gad, ³⁷⁹ M + go, ⁴⁶⁰ D + go, ⁴⁶¹ R + ca, ⁴⁶² D + gad, ⁴⁶⁹ M + go
CP47	⁵⁸ Q + go, ⁶⁰ M + go, ³²⁷ A + go, ³²⁹ S + go, ³³⁰ M + go or aa, ³⁵⁸ R + go, ³⁵⁹ M + go or aa or ca, ³⁶⁰ P + go or ca, ³⁶¹ T + go, ³⁶³ F ^d + go, ³⁶⁴ E ^d + gad, ³⁶⁵ T ^d + go, ³⁷⁴ D + gad, ³⁷⁶ I + go, ³⁷⁷ V + ca, ³⁷⁸ R + go, ³⁹³ E + gad, ³⁹⁴ Q + ca, ³⁹⁸ T + go, ⁴²³ R + ca, ⁴²⁴ A ^d + go, ⁴²⁵ Q + ca, ⁴²⁸ E + gad, ⁴⁸⁴ P + ca, ⁴⁸⁵ D + gad, ⁴⁸⁶ L + go or ca, ⁴⁸⁷ D + gad, ⁴⁸⁸ V + go, ⁴⁸⁹ Q + ca

^aIndividual residues are listed along with the modifications observed. In some instances, different modifications were observed for the same residue on different peptides. For a complete list of oxidative modification types, the amino acids targeted, and mass modifications searched for in this study, see ref 29. Key: aa, aldehyde addition (−32 amu); ca, carbonyl addition (+14 amu) cysh, cysteine hydroxyl (−16 amu); gad, Glu/Asp decarboxylation (−30 amu); go, general oxidation (+16 amu); stcb, serine/threonine carbonyl (−2 amu). ^bNot resolved in the *Thermosynechococcus vulcanus* structure. ¹³ Buried residues not adjacent to apparent cavities/channels. ^dBuried residues adjacent to cavities/channels partially occupied by structural waters.

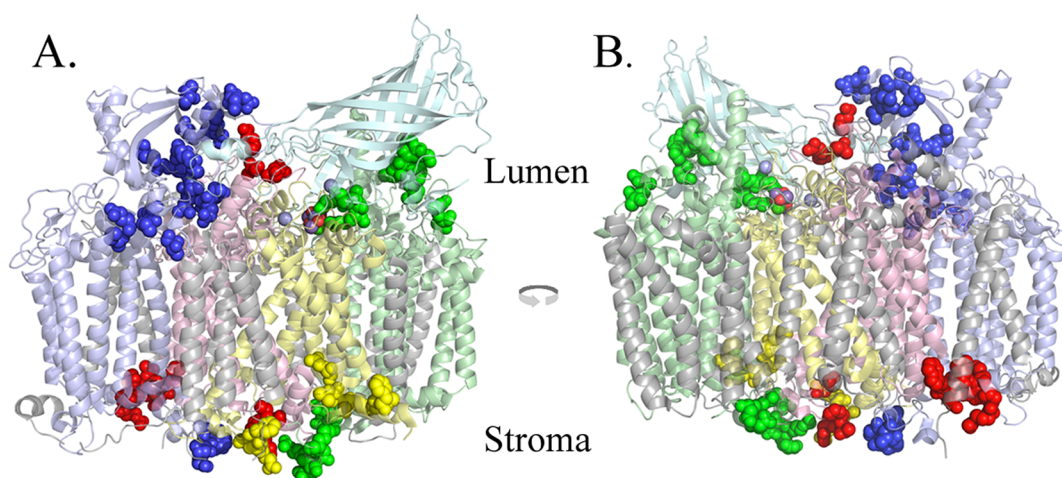


Figure 2. Overview of the oxidized residues identified in the D1, D2, CP43, and CP47 proteins. Oxidized residues are shown as spheres superimposed on monomer I of the *Thermosynechococcus vulcanus* structure. Since the PsbU and PsbV subunits are not present in higher plant PS II, these chains are not shown. (A) The view from outside monomer I, looking toward the dimeric complex within the plane of the membrane. (B) The view from monomer II looking toward its interface with monomer I within the plane of the membrane. Color key: CP47, pale blue; CP43, pale green; D1, pale yellow; D2, pale red; PsbO, pale cyan; all other chains, gray. The modified residues are shown in darker shades of these colors. Chlorides are shown as light blue spheres, and the manganese and oxygen of the Mn₄O₅Ca cluster are shown as purple and red spheres, respectively. It should be noted that many surface residues are modified on the PsbO subunit, but these are not shown in this illustration. Figures 2 and 3 were produced using PYMOL.³¹

eter. The experiments used the standard electrospray source operating with a source voltage of 5 kV and a capillary temperature of 275 °C. Sheath and auxiliary gas flows were 18 and 5, respectively. A typical scan sequence involved a positive ion FT-ICR scan at 100 K resolution (100 K at *m/z* 400). During the FT-ICR acquisition, six MS² scans were acquired by the linear ion trap determining the parent ions from the six largest ions observed from a preview of the FT-ICR scan. The CID scans are acquired with an isolation width of 2 and a normalized collision energy of 35 (both Thermo Scientific machine settings). After acquiring the tandem mass spectra twice, the ion is placed into an exclusion list for 30 s. Charge state screening is enabled with monoisotopic precursor selection.

Two biological replicate experiments were performed. Identification and analysis of peptides containing oxidative modifications were performed using the MassMatrix Program ver. 1.3.1^{26,27} which had been programmed to include the possible oxidative modifications.^{28,29} A FASTA library containing the spinach PS II protein sequences was searched as was a decoy library that contained these same proteins but with reversed amino acid sequences. No hits to the decoy library

were observed. For the determination of the quality of the peptide calls within MassMatrix, max(*pp*₁, *pp*₂) was ≥8.5 and *pp*_{tag} ≥ 4.0.^{26,27} These parameters yield a *p* value of ≤0.0001; only oxidized peptides which exhibited these extremely low *p* value were considered. Given the high quality of the data used in this study, the union of the replicate data sets was examined. The identified oxidatively modified residues were mapped onto the *T. vulcanus* PS II structure¹³ of the D1, D2, CP43, and CP47 proteins using PYMOL.³¹

RESULTS

Figure 1 illustrates the quality of the mass spectrometry data used for the identification of oxidized amino acid residues in the D1, D2 CP47, and CP43 proteins. Using a *p* value ≤0.0001 assured extremely high quality peptide identifications. In this context it should be noted that in MASCOT searches,³⁰ peptide identifications are typically performed with *p* values ≤0.05 (www.matrixscience.com). The direct consequence of our using such low *p* values is that in all cases nearly complete y- and b-ion series for the oxidatively modified peptides were observed (Figure 1 and Figures S2–S5).

A total of 73 oxidatively modified amino acid residues were identified in the D1, D2, CP43, and CP47 proteins of spinach PS II. This represents $\approx 4\%$ of the residues found in these four proteins. These are summarized in Table 1 along with the type of oxidative modifications observed and the residue location (surface, buried but in contact with cavity/channel, or buried and not in contact with cavity or channel) within the *T. vulcanus* crystal structure. In some instances more than one type of oxidative modification was observed at a particular amino acid residue position. It is important to note that these oxidized residues are not uniformly modified on all PS II complexes. In most instances the oxidation of a particular residue or groups of residues is expected to be quite rare. The actual degree of modification of any particular residue is quite difficult to quantitate and lies well beyond the scope of this communication. Indeed, such quantification is problematic since oxidative modification of many amino acid residues yields derivatives that are more polar than the parent residue. Consequently, peptides containing such modifications are more polar and more readily resolved by reversed-phase HPLC, making quantitative comparison to the unmodified, more hydrophobic, parent peptide difficult. It should also be noted that, in general, mass spectrometry coverage of intrinsic membrane proteins is challenging. The overall sequence coverage observed in this study for the proteins examined was: D1, 24%; D2, 27%; CP47, 41%; and CP43, 26%, which is quite comparable to that observed for these proteins by other investigators (for instance Nakamura et al.³² observed; D1, 21%; D2, 22%; CP47, 23%; and CP43, 32%). However, the coverage of the residues located in the extrinsic loops of these proteins was significantly higher: D1, 35%; D2, 43%; CP47, 55%; and CP43, 43%. These are the domains of principal interest in this study, as the oxygen-evolving complex and proposed oxygen channels are all embedded in the lumenally exposed extrinsic loop regions of these proteins. Within this context it should be noted that any oxidative modifications to amino acid residues lying in the transmembrane helices of these proteins would be very difficult to identify and would probably escape detection due to their poor resolution during reversed-phase chromatography.

The D1, D2, CP47, and CP43 proteins are highly homologous between higher plants and cyanobacteria (>85% similarity). Consequently, it is possible to map the modified residues present on these intrinsic proteins from spinach directly to the homologous residues in the *T. vulcanus* crystal structure. Indeed, 90% of the observed oxidatively modified residues in these spinach proteins are conserved (or conservatively replaced) in the corresponding *T. vulcanus* sequences. These results are shown in Figure 2. As expected, the vast majority (81%) of the observed modifications are located on the surface of the PS II complex, which is exposed to the oxygenated bulk solvent. Additionally, four oxidized residues at the N-terminus of the D1 (²T, ³A, ⁵L, and ⁶E) are also almost certainly surface-exposed; however, this could not be verified by direct comparison to the *T. vulcanus* structure, as these residues are not resolved. Consequently, they were not included in the above calculations and are not shown in Figure 2. Eight residues were observed to be modified which, while not located on the surface of the complex, are adjacent to cavities/channels within the complex, while three residues are buried and are not adjacent to such cavities/channels (Table 1).

Figure 3 illustrates all of the modified residues identified which are located within 15 Å of the metal cluster. The residues

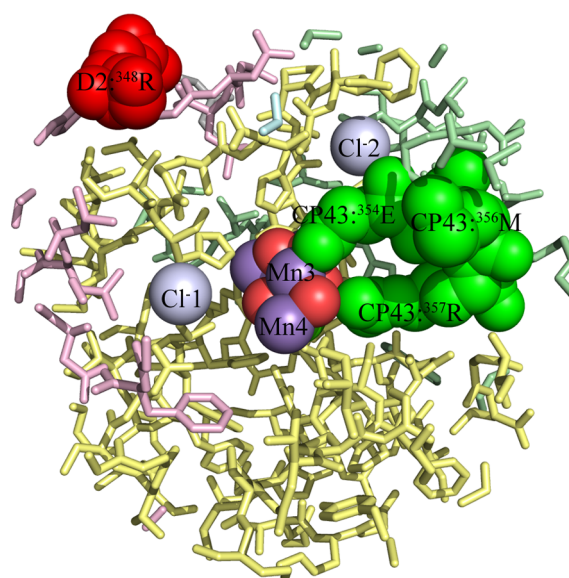


Figure 3. Oxidized residues in the vicinity of the $\text{Mn}_4\text{O}_5\text{Ca}$ cluster active site of PS II. A 15 Å sphere centered on Mn3 of the $\text{Mn}_4\text{O}_5\text{Ca}$ cluster is shown. The CP43 residues ³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R are all buried residues which are in contact with cavities/channels on the interior of the complex. Please note that the D2 residue ³⁴⁹R of spinach is the equivalent of ³⁴⁸R of *T. vulcanus* and is a surface residue within the cyanobacterial structure. In this view ³⁵⁵T is not visible. Color key: CP47, pale blue; CP43, pale green; D1, pale yellow; D2, pale red; PsbO, pale cyan; all other chains, gray. The modified residues are shown in darker shades of these colors. The oxidized residues are labeled, as are Cl⁻1 and Cl⁻2 and Mn3 and Mn4. These are shown as spheres. Some foreground residues, principally of the D1 chain, have been removed for clarity.

³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R of CP43 and residue ³⁴⁹R of the D2 protein are oxidatively modified (Figure 3). This latter residue is exposed on the surface of the complex. The FT-ICR spectra identifying the oxidative modification of ³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R of CP43 residues and the analogous unmodified peptides from both biological replicates are shown in Figure 1 and Figures S2–S5. In all cases, extremely high quality data were obtained with excellent b- and y- ion coverage. It should be noted that ³⁵⁴E is an inner-sphere bidentate ligand to the $\text{Mn}_4\text{O}_5\text{Ca}$ cluster, bridging Mn2 and Mn3. ³⁵⁷R is a second sphere ligand to the metal cluster and is hydrogen bonded to O2 and O4.¹³ All four of these modified residues are in contact with cavities/channels within the photosystem. ³⁵⁴E is in contact with a channel partially occupied with structurally resolved water molecules (W538, W539, W397, W393, etc.) and is also in contact with a cavity containing Cl⁻2 and W498, W506, W547, W548, and W943. ³⁵⁵T is also in contact with this cavity. ³⁵⁶M is in contact with a channel partially occupied by the structurally resolved water molecules (W369, W546, W402, etc.). ³⁵⁷R is in contact with the same channel, with the closest structurally resolved water molecules being W1015, W1016, and W550. This channel leads to the surface of the complex.

DISCUSSION

The oxidative modification of these four CP43 residues (³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R) appears to indicate that they have been modified by ROS generated within the PS II complex. The close proximity of these residues to the $\text{Mn}_4\text{O}_5\text{Ca}$ cluster

supports the hypothesis that these residues are associated with an O₂/ROS exit pathway leading from the metal cluster to the surface of the complex. In the earlier computational studies ³⁵⁴E, ³⁵⁶M, and ³⁵⁷R were identified as being associated with a number of proposed channels. All three residues were located in the “narrow” channel³³ (also termed the “D, E, F” channel²¹) which was hypothesized to be a proton exit pathway. ³⁵⁶M and ³⁵⁷R were also proposed to be associated with the “back”,³³ “(i)”,¹⁸ or “A1, A2”²¹ channel. Ho and Styring³³ and Gabdulkhakov et al.²¹ hypothesized that this channel was involved in vectoring water toward the Mn₄O₅Ca cluster, while Murray and Barber¹⁸ suggested that this was an O₂ exit pathway. Finally, ³⁵⁴E was also suggested to be located in the “large”,³³ “(ii)”,¹⁸ or “B1, B2”²¹ channel. Ho and Styring³³ and Gabdulkhakov et al.²¹ suggested that this channel was involved in vectoring O₂ away from the Mn₄O₅Ca cluster, while Murray and Barber¹⁸ hypothesized that this was a water/proton exit pathway. ³⁵⁵T was not identified in any of these prior studies as being associated with any channel. It should be noted that several limitations, in large measure, exist with all of these largely computational studies. First, both the CAVER program,¹⁹ used by Murray and Barber¹⁸ and Gabdulkhakov et al.,²¹ and calculations based on solvent accessibility used by Ho and Styring,³³ examine static crystal structures. This is also true of the noble gas^{20,21} and dimethyl sulfoxide²¹ cocrystallization studies which have been reported. Large protein complexes exhibit substantial movement on the nanoseconds time scales.³⁴ Such movement in PS II could substantially alter the overall shape and dimensions of the identified channels within the photosystem. For substrate water, this problem has been recognized and addressed computationally by molecular dynamic simulations of water movement within the photosystem;^{34,35} however, at this time, the movement of O₂ has not been examined by these methods. Second, the majority of these studies have been performed on PS II structures of moderate resolution (≥ 2.9 Å). At these resolutions the accurate modeling of the location of the amino acid side chains is problematic. The accurate positioning of residue side chains could have important consequences with respect to the continuity and radius of proposed channels. At this time, no similar studies based on the high-resolution (1.9 Å) PS II structure of Umena et al.¹³ have been reported with respect to the identification of O₂ channels. As noted above, noble gas derivitization has been used in an attempt to identify oxygen egress channels in PS II. These gases bind to hydrophobic domains hypothesized to be associated with O₂ transport within the photosystem.^{20,21} However, it should be noted that in other systems hydrophilic O₂ binding domains have been identified which do not interact with noble gases but which do interact with chloride and water.^{36,37} Finally, all of these studies have been performed on crystal structures of PS II complexes which, while uniformly reported to be in S₁, are clearly in an unknown, more highly reduced S-state.³⁸ Even if these crystal structures were to accurately represent the protein architecture of the S₁ state, an implicit assumption has been made that no conformational changes occur during the various S-state transitions, particularly the S₃ → [S₄] → S₀ state transition associated with O₂ formation, release, and water binding which could affect the location, shape, and dimensions of possible O₂/ROS exit pathways. Numerous recent studies have indicated that protein conformational changes do occur during S-state cycling,^{39–43} although the specific residues involved and the effects on

possible channels within the photosystem have not been determined.

As noted above, oxidative modification of amino acid residues requires that they be exposed to ROS. PS II, particularly when under stress, can produce a variety of ROS at a variety of sites.¹⁵ Several studies have identified the production of ROS, particularly OH•, by the donor side of PS II. OH• is produced at or near the Mn₄CaO₅ cluster under moderately high temperature^{44,45} conditions. Additionally, it has been documented that the Mn₄O₅Ca cluster can directly absorb near-UV radiation,^{46–48} leading to damage to the manganese cluster. This damage could also lead to the partial oxidation of water, H₂O₂ production, and the generation of OH•. Both of these conditions could be experienced by the field-grown spinach used in these studies. It has also recently been shown that loss of the chloride associated with the active site under these conditions leads to the production of H₂O₂.⁴⁹ The loss of chloride also leads to the loss of manganese associated with the Mn₄O₅Ca cluster.⁵⁰ This manganese has been hypothesized to react with the H₂O₂ in a classical Fenton reaction to produce OH•.¹⁵ It must be stressed, however, that at this time we cannot discriminate between these or other mechanisms that produce the ROS responsible for the oxidative modifications that we observe. Using mass spectrometry, it is also very difficult, and in most cases impossible, to distinguish hydroxide ion radical modifications of amino acid side chains from those produced by singlet oxygen, superoxide, or other oxidative species.^{51,52}

Our results indicate that the residues ³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R of CP43 are oxidatively modified in spinach PS II, and consequently, these residues have been exposed to oxygen/ROS. These residues probably represent only a subset of the total oxidatively modified residues in the photosystem. For instance, oxidatively modified residues in the transmembrane domains of these proteins would generally be difficult to detect given the hydrophobic nature of the tryptic peptides derived from these helical domains. We hypothesize that ³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R of CP43 are adjacent to molecular oxygen/ROS egress channel(s) which vector these products away from the Mn₄O₅Ca cluster toward the surface of the PS II complex. One possibility is that ³⁵⁶M and ³⁵⁷R which are in contact with the so-called “back” channel³³ (also known as the “(i)”¹⁸ or “A1, A2”²¹ channels) that leads to surface of the PS II is an oxygen/ROS exit pathway as was originally suggested by Murray and Barber.¹⁸ ³⁵⁴E and ³⁵⁵T, however, do not appear to be in contact with this “back” channel in these analyses of the static crystal structures.^{18,21,33} Dynamic motions of PS II may modulate the dimensions and/or the precise location of the “back” channel, allowing ³⁵⁴E and ³⁵⁵T to become oxidatively modified. It is additionally possible, however, that the “narrow” channel³³ (also termed the “D, E, F” channel²¹) that is adjacent to ³⁵⁴E, ³⁵⁶M, and ³⁵⁷R is an oxygen/ROS egress pathway. While in static structures this channel appears too narrow to allow the egress of oxygen/ROS, dynamic molecular motions or S-state-dependent conformational changes may facilitate this process. Alternatively, all of the oxidized residues (³⁵⁴E, ³⁵⁵T, ³⁵⁶M, and ³⁵⁷R) which we have identified in the vicinity of the Mn₄O₅Ca cluster may be associated with additional exit pathways for oxygen/ROS which have not been identified computationally. The differentiation between these and other hypotheses is currently underway.

Finally, it has not escaped our attention that another group of oxidized residues which we have identified (the D1 residues ^{130}E , ^{133}L , and ^{135}F) may be involved in an ROS egress pathway operating on the reducing side of the photosystem. Oxidative modifications of amino acid residues in this vicinity have been previously observed, although the identities of the modified residues were not determined.⁵³ These residues are in close proximity to Pheo_{D1} which, under stress conditions, has the potential ($E_m = -610$ mV, pH 7.0) of being a source of reducing-side ROS.¹⁵ This possibility will be examined in a separate communication.

■ ASSOCIATED CONTENT

■ Supporting Information

One figure showing the efficacy of the nonoxidizing gel system in suppressing artifactual electrophoresis-induced protein oxidation; additionally, four figures showing the MS² of the unmodified and modified peptides containing residues ^{354}E , ^{355}T , ^{356}M , and ^{357}R of CP43 and their associated tables of the identified ions in these spectra. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

■ ABBREVIATIONS

amu, atomic mass units; CID, collision-induced dissociation; FT-ICR, Fourier transform-ion cyclotron resonance; LiDS, lithium dodecyl sulfate; PAGE, polyacrylamide gel electrophoresis; PS II, Photosystem II; ROS, reactive oxygen species; S_n , oxidation states ($n = 1, 2, 3, 4$) of the $\text{Mn}_4\text{O}_5\text{Ca}$ cluster.

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