The Conserved His-144 in the PsbP Protein Is Important for the Interaction between the PsbP N-terminus and the Cyt $b_{559}$ Subunit of Photosystem II

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**Background:** PsbP is an extrinsic subunit of photosystem II in green plants.

**Results:** H144A mutation in PsbP alters the chloride requirement and affects the interaction between its N terminus and the PsbE component of photosystem II.

**Conclusion:** The N- and C-terminal domains of PsbP cooperate to support PSII activity.

**Significance:** This provides important information about the binding characteristics of PsbP in green plant PSII.

The PsbP protein regulates the binding properties of Ca$^{2+}$ and Cl$^{-}$, and stabilizes the Mn cluster of photosystem II (PSII); however, the binding site and topology in PSII have yet to be clarified. Here we report that the structure around His-144 and Asp-165 in PsbP, which is suggested to be a metal binding site, has a crucial role for the functional interaction between PsbP and PSII. The mutated PsbP-H144A protein exhibits reduced ability to retain Cl$^{-}$ anions in PSII, whereas the D165V mutation does not affect PsbP function. Interestingly, H144A/D165V double mutation suppresses the effect of H144A mutation, suggesting that these residues have a role other than metal binding. FTIR difference spectroscopy suggests that H144A/D165V restores proper interaction with PSII and induces the conformational change around the Mn cluster during the $S_1/S_2$ transition. Cross-linking experiments show that the H144A mutation affects the direct interaction between PsbP and the Cyt $b_{559}$ subunit of PSII (the PsbE protein). However, this interaction is restored in the H144A/D165V mutant. In the PsbP structure, His-144 and Asp-165 form a salt bridge. H144A mutation is likely to disrupt this bridge and liberate Asp-165, inhibiting the proper PsbP-PSII interaction. Finally, mass spectrometric analysis has identified the cross-linked sites of PsbP and PsbE as Ala-1 and Glu-57, respectively. Therefore His-144, in the C-terminal domain of PsbP, plays a crucial role in maintaining proper N terminus interaction. These data provide important information about the binding characteristics of PsbP in green plant PSII.

Photosystem II (PSII)\textsuperscript{2} consists of both membrane-intrinsic and membrane-extrinsic subunits, and functions as a water/plastoquinone oxidoreductase (for reviews, Refs. 1–4). On the thylakoid lumenal side of PSII, a metal cluster of four Mn ions, one Ca$^{2+}$ ion, and five oxo ligands (the Mn cluster) catalyzes the oxygen-evolving reaction. Additionally, two Cl$^{-}$ ions are bound near to the Mn cluster (5). The membrane-intrinsic subunits of PSII are involved in pigment and/or cofactor binding for photochemical reactions, while the membrane-extrinsic subunits surround the catalytic Mn cluster and play crucial roles in stabilizing the Mn cluster and retaining the PSII cofactor ions (6 – 8).

X-ray structural analysis of the cyanobacterial PSII complex at atomic resolution has revealed the location of subunits, pigments, and cofactors, including the exact organization of the subunits within PSII (5, 9 – 11). However, the crystallographic information gained from the study of cyanobacterial PSII cannot necessarily be applied in the context of other eukaryotes because of the differences in PSII subunit composition. In particular, the composition of the PSII extrinsic subunits has undergone significant evolutionary changes; green plants, such as higher plants, green algae, and Euglena have a set of 3 extrinsic proteins, namely PsbO (33 kDa), PsbP (23 kDa), and PsbQ (17 kDa). Alternatively, cyanobacteria have PsbV and PsbU instead of PsbP and PsbQ (12, 13). In cyanobacteria, the PsbP and PsbQ homologs (Cyanop and CyanQ, respectively) are present (14, 15), but they are not included in the current structural models. The PsbP and PsbQ proteins in green plants seem to have evolved from their cyanobacterial homologs, although considerable genetic and functional modifications seem to have occurred to generate the eukaryote-type PsbP (16, 17). The locations and binding topologies of PsbP and PsbQ in the green plant PSII complex have been proposed (17, 18), but this model has not yet been clarified.

PsbP and PsbQ are responsible for the retention of Ca$^{2+}$ and Cl$^{-}$ within PSII (for a review, see Ref. 8). Recent FTIR difference spectroscopy suggests that the protein conformational changes around the Mn cluster, which are induced by PsbP binding,
modulate the binding properties of Ca$^{2+}$ and Cl$^-$ in PSII (19, 20). Truncation of the PsbP N terminus by 15 residues resulted in a loss of the protein Ca$^{2+}$ and Cl$^-$ retention ability (21). In addition, we have recently reported that His-144, a highly conserved residue in PsbP and Cyanop, was important for the ion retention ability (20). However, the functional role of His-144 has not been elucidated.

X-ray crystal structural analyses suggest that the C-terminal domain of the PsbP protein from N. tabacum (PDB: 1V2B) and the Cyanop protein from Thermosynechococcus elongatus (PDB: 2X3B) have a similar α/β/α structure, which is characteristic of the PsbP superfamily (22, 23). The crystal structure of spinach PsbP (PDB: 2VU4) has been recently reported (24). Interestingly, the structure of spinach PsbP has one Zn$^{2+}$ ion that is ligated to the Asp-165 and His-144 residues in the C-terminal domain. This Zn$^{2+}$-binding domain is conserved in the structure of Cyanop. PsbP has been reported to bind Mn$^{2+}$ ion, and the metal-binding site has been suggested to be around the His-144, Asp-165, and Glu-177 residues (25, 26). The structural conservation may indicate the importance of metal binding for the function of PsbP and Cyanop. However, it is also possible that Zn$^{2+}$ binding to the PsbP protein is an artifact, due to the conditions during the crystallization process. Therefore, further research is required to determine the importance of metal binding for the function of PsbP in PSII.

In this study we characterized PsbP mutants to analyze how His-144 is involved in the functions of PsbP. We produced three PsbP mutant proteins: PsbP-H144A (H144A), wherein the conserved His-144 residue was substituted with Ala, PsbP-D165V (D165V), in which the Asp-165 residue was substituted with Val, and a mutant containing both substitutions, PsbP-H144A/D165V. The functional properties of the mutated PsbP proteins were studied by conventional reconstitution experiments and FTIR analyses. In addition, interaction between PsbP and PSII was investigated using a chemical cross-linker, a water-soluble carbodiimide, 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide (EDC). The results obtained from these investigations suggest that metal-binding is not required for PsbP function, and that PsbP structure around the His-144 and Asp-165 residues plays a crucial role in maintaining the protein functional interaction with PSII. Finally, cross-linking experiments show that PsbP directly interacts with the cytochrome (Cyt) b$_{599}$ a subunit (the PsbE protein) via its N terminus, and that this interaction is affected by H144A mutation.

**EXPERIMENTAL PROCEDURES**

Plasmid Construction, Recombinant Protein Expression, and Purification—The expression plasmids for the mutated PsbP proteins H144A, D165V, and H144A/D165V were constructed using a site-directed mutagenesis kit (Stratagene) with the designed primers 5’-GGTTAGGAGGTTGAAAGCCCAAGTATTCAGGACGACTG-3’, 5’-GCTCAAGCTGGAGCT-AAGAGATGGTTCC-3’, and their respective complementary primers. The expression plasmid for A186C, in which Ala-186 is substituted with Cys, was constructed with the designed primer pair 5’-GGGAATTCCATATGGCCTATGGAG-3’ and 5’-GGTTAACGAGCTTATAACCCACTG-3’. The recombinant WT and mutated PsbP proteins were expressed in the Escherichia coli strain BL21 (DE3) and purified, as previously reported (27). A186C was labeled with maleimide-PEG2-biotin (Pierce) according to the manufacturer’s protocol. The presence of the desired mutation in each recombinant PsbP and maleimide-PEG2-biotin labeling was confirmed by MALDI-TOF mass spectrometry (Autoflex III, Bruker).

Reconstitution of the PsbP Protein to NaCl-washed PSII, and Measurement of Oxygen-evolving Activity—Oxygen-evolving PSI membranes of spinach were prepared as reported previously (28). The chlorophyll (Chl) concentration was calculated from the equations described in Ref. 29. The activity of isolated PSI membranes was ~480 μmol O$_2$/mg Chl/h. The reconstitution of PsbP to NaCl-washed PSI membranes and the measurement of O$_2$ evolution were performed according to a procedure reported elsewhere (27, 30) with a slight modification: To stabilize the interaction of the extrinsic proteins and the NaCl-washed PSI membranes, 0.4 м sucrose was replaced with 2 м betaine in the reconstitution buffer (25 м Mes-NaOH, pH 6.5, 2 м betaine, 20 м CaCl$_2$) and in the activity measurement buffer (25 м Mes-NaOH, pH 6.5, 2 м betaine). PsbP was reconstituted with PSI in molecular ratio of 2:1 (PsbP:PSII), unless otherwise noted. To measure the oxygen-evolving activity in the presence of Ca$^{2+}$ or Cl$^-$, Ca(OH)$_2$, or NaCl was added to the buffer used for activity measurements, respectively. The SDS-PAGE gels were stained using Flamingo (Bio-Rad) and visualized using a fluoro image analyzer FLA-3000 (FUJIFILM). The amount of protein bound to PSII was determined by measuring the fluorescence intensity with the software Multi Gauge Ver 3.0 (FUJIFILM).

FTIR Analysis—Preparation of membrane samples and FTIR measurements were performed as reported previously (19), except that $S_2$Q$_A$-minus-S$_1$Q$_A$ difference spectra (hereafter termed $S_2$Q$_A$/S$_1$Q$_A$ difference spectra) were analyzed, instead of the $S_2$/S$_1$ difference spectra, as in the previous study. This modification was introduced to avoid the effect of potassium ferricyanide, which can act as an exogenous electron acceptor, on PsbP and PsbQ binding. The PSI samples in the presence of 0.1 м DCMU were centrifuged, and the resulting pellet was sandwiched between the CaF$_2$ plates. Light-induced S$_2$Q$_A$/S$_1$Q$_A$ FTIR difference spectra were recorded using a Bruker IFS-66/S spectrophotometer equipped with an MCT detector (Infrared D316/8) at a resolution of 4 cm$^{-1}$ (31). Illumination was performed using a Q-switched Nd:YAG laser (Quanta-Ray GCR-130, 532 nm, ~7 ns full width at half-maximum, ~7 mJ pulse$^{-1}$ cm$^{-2}$ at the sample surface). Single-beam spectra (150 s scan) were recorded before and after single-flash illumination, providing a difference spectrum via subtraction of the initial spectrum from the spectrum obtained after illumination. The measurements were repeated three to five times for each sample. The samples were warmed to 285 K to relax the S$_2$ state, and then cooled again to 250 K. One to two samples were used for the measurements, and the obtained spectra were averaged.

Cross-linking Experiments—NaCl-washed PSI membranes, at a concentration of 0.5 м Chl/ml were cross-linked with the recombinant PsbP proteins in the cross-linking buffer (25 мm
Hepes-NaOH, pH 7.2, 25 mM CaCl₂) by using 6.25 mM EDC and 5 mM N-hydroxysulfosuccinimide (sulfo-NHS). The solution was incubated for 2 h in darkness, and the reaction was terminated by adding ammonium acetate to a final concentration of 0.2 M. The solution was centrifuged for 5 min at 20,400 × g at 4 °C, and the pellet was subjected to SDS-PAGE. Proteins separated by SDS-PAGE were transferred to PVDF membranes and analyzed by immunoblotting using specific antibodies. To cross-link the carboxyl groups on the NaCl-washed PSII and the primary amines on the PsbP, NaCl-washed PSII was treated with 6.25 mM EDC and 5 mM sulfo-NHS in activation buffer (25 mM Mes-NaOH pH 6.0, 500 mM NaCl) for 15 min in darkness, and then centrifuged for 5 min at 20,400 × g at 4 °C. The pellet was washed once with the cross-linking buffer and the activated PSII was subsequently incubated with PsbP in the cross-linking buffer without EDC or sulfo-NHS for 2 h in darkness.

Rabbit antibodies against PsbP and D1 were prepared by the authors. Rabbit antibody against PsbO was provided by the late Dr. A. Watanabe of Tokyo University. Anti-CP47 rabbit antibody was provided by Dr. A. Tanaka of Hokkaido University, while rabbit antibody against CP43 was a gift from Dr. Y. Kashino of Hyogo Prefectural University. Rabbit antibodies against D2, PsbR and PsbE were purchased from Agrisera.

Purification of Cross-linked Products—PSII membrane cross-linked with A186C labeled with maleimide-PEG2-biotin was solubilized in 100 mM Hepes-NaOH pH 7.2, 150 mM NaCl, 2% n-octyl-β-D-glucoside and incubated with Strep-Tactin Sepharose (IBA) at 4 °C for 1 h. The resin was washed three times with 100 mM Hepes-NaOH pH 7.2, 150 mM NaCl and 1% SDS at room temperature for 10 min. The proteins bound to the resin were eluted by boiling the resin in SDS-PAGE sample buffer (62.5 mM Tris-HCl pH 6.8, 10% (v/v) glycerol, 2.5% (w/v) SDS, 2.5% (v/v) 2-mercaptoethanol) at 90 °C for 10 min, and used for subsequent analysis.

MS Analysis—The purified proteins were separated by SDS-PAGE and subjected to in-gel digestion with Mass Spec Grade Modified Trypsin (Promega). The peptides were extracted and loaded on a column (100 μm internal diameter, 15 cm length; L-Column, CERI) using a Paradigm MS4 HPLC pump (Michrom Bioresources) and an HTC-PAL autosampler (CTC Analytics). Buffers were (A) 0.1% (v/v) acetic acid and 5% (v/v) acetonitrile in water and 0.1% (v/v) acetic acid, and (B) 90% (v/v) acetonitrile in water. A linear gradient of buffer B from 5% to 45% was employed, and peptides eluted for 26 min from the column were introduced directly into an LTQ-Orbitrap XL mass spectrometer (Thermo Fisher Scientific) with a flow rate of 200 nl min⁻¹ and a spray voltage of 2.0 kV. The range of the mass spectrometric scan was a mass-to-charge ratio of 450 to 1500, and the top three peaks were subjected to tandem mass spectrometry analysis.

Search for Cross-linked Sites—Thermo RAW files were converted to mzXML files with MM File Conversion Tools (www.massmatrix.net). Cross-linked peptides and cross-linked sites were identified with the MassMatrix Data base Search Engine (32). The settings were as follows: enzyme, trypsin, no P rule; decoy database, reversed; missed cleavages, 4; variable modifications, oxidation of M, amidation of DE, carbamidomethyl of C; precursor ion tolerance, 10 ppm; product ion tolerance, 0.8 Da; maximum number of PTM per peptide, 2; mass type, monoisotopic; minimum peptide length, 5; maximum peptide length, 40; cross link, EDC; cross link mode, exploratory; cross link site cleavability, non-cleavable by enzyme or not applicable; maximum number of cross links per peptide, 2.

RESULTS

Introduction of the D165V Mutation to the H144A Mutant PsbP Protein Restored Cl⁻ Retention Ability—Fig. 1 shows the structural models of spinach PsbP (A, PDB ID: 2VU4) and of tobacco PsbP (B, PDB ID: 1V2B). α-Helices and β-strands are shown in blue and orange, respectively. The side chains of the His-144, Asp-165, and Glu-177 residues are shown as stick models. A Zn²⁺ ion is shown as a magenta sphere.

FIGURE 1. Comparison of the PsbP structures of spinach (PDB ID: 2VU4; A) and tobacco (PDB ID: 1V2B; B). α-Helices and β-strands are shown in blue and orange, respectively. The side chains of the His-144, Asp-165, and Glu-177 residues are shown as stick models. A Zn²⁺ ion is shown as a magenta sphere.
The Functional Role of the His-144 of PsbP in Photosystem II

The oxygen-evolving activity of PSII reconstituted with WT, H144A, D165V, and H144A/D165V respectively, and the binding ability of these PsbP proteins to PSII. A, oxygen-evolving activity was measured in the absence of Ca\(^{2+}\) and Cl\(^{-}\) ions. WT-reconstituted PSII activity (216 μmol O\(_{2}/\)mg Chl/h) was set at 100%; the quantities of the PSII-bound PsbP following reconstitution were quantified from the intensity of the fluorescence in the SDS-PAGE gel. The band intensity in the mock lane was subtracted from the intensity of each test lane. The intensity of the WT was set at 100%; n = 3, error bars = S.D. B, The oxygen-evolving activity was measured in the absence of Ca\(^{2+}\) and Cl\(^{-}\) ions. WT-reconstituted PSII measured in 20 mM Cl\(^{-}\) (white bars), 10 mM CaCl\(_{2}\) (black bars), and 10 mM NaCl (light gray bars). WT-reconstituted PSII activity measured in the absence of Ca\(^{2+}\) and Cl\(^{-}\) (289 μmol O\(_{2}/\)mg Chl/h) was set at 100%; n = 3, error bars = S.D. C, binding profiles of WT (squares, black line) and H144A (triangles, gray line) to NaCl-washed PSII. PSII was reconstituted with WT or H144A in various molecular PsbP:PSII ratios, and the amount of PSII-bound PsbP was quantified as in B. The intensity of WT:PSII = 4:1 was set at 100%; n = 4, error bars = S.D.

The ability of PsbP to bind to PSII (Fig. 2B). We also analyzed the importance of Glu-177 for PsbP function. Both the oxygen-evolving activity and the amount of PsbP bound to the PSII of the E177V-reconstituted PSII were similar to those of the WT-reconstituted PSII, indicating that Glu-177 is not essential for the ion retention and binding ability of PsbP (supplemental Fig. S1). These results suggest that the metal binding around His-144 and Asp-165 is not essential for PsbP function.

To compare the binding ability of the WT and H144A PsbP proteins more precisely, reconstitution experiments were performed with various PsbP:PSII ratios. As shown in Fig. 2C, the WT and H144A PsbP proteins bound to PSII in a very similar manner when the reconstitution was performed with PsbP:PSII ratios from 0.25:1 to 2:1, indicating that the specific manner with which PsbP binds to PSII and the affinity of this binding is not affected by H144A mutation. Alternatively, when PsbP was reconstituted with a PsbP:PSII ratio of 4:1, the binding affinity of the H144A protein appeared greater than that of the WT. This may suggest that H144A binds to PSII in a nonspecific manner when an excess amount of H144A is present.

To evaluate the effect of the H144A mutation on the Ca\(^{2+}\) and Cl\(^{-}\) retention ability of PsbP separately, the oxygen-evolving activity of WT- and H144A-reconstituted PSII was measured under four conditions: in the presence of 10 mM Ca\(^{2+}\), in the presence of 20 mM Cl\(^{-}\), in the presence of 10 mM CaCl\(_{2}\), and in Ca\(^{2+}\) - and Cl\(^{-}\)-free conditions (Fig. 3A). Addition of 10 mM Ca\(^{2+}\) did not restore the oxygen-evolving activity of H144A-reconstituted PSII when compared with the activity in the absence of Ca\(^{2+}\) and Cl\(^{-}\). In contrast, addition of 20 mM Cl\(^{-}\) purified to metal-free forms. The absence of any divalent cations was confirmed by inductively coupled plasma mass spectrometry (data not shown). We then performed conventional release-reconstitution experiments to determine whether these PsbP variants could restore the oxygen-evolving activity of the NaCl-washed PSII and bind to PSII. The oxygen-evolving activity was measured in the absence of Ca\(^{2+}\) and Cl\(^{-}\) ions. WT-reconstituted PSII activity (216 μmol O\(_{2}/\)mg Chl/h) was set at 100%; the quantities of the PSII-bound PsbP following reconstitution were quantified from the intensity of the fluorescence in the SDS-PAGE gel. The band intensity in the mock lane was subtracted from the intensity of each test lane. The intensity of the WT was set at 100%; n = 3, error bars = S.D. C, binding profiles of WT (squares, black line) and H144A (triangles, gray line) to NaCl-washed PSII. PSII was reconstituted with WT or H144A in various molecular PsbP:PSII ratios, and the amount of PSII-bound PsbP was quantified as in B. The intensity of WT:PSII = 4:1 was set at 100%; n = 4, error bars = S.D.
significantly increased the activity of H144A-reconstituted PSII to about 80% of that of the WT. The activities of WT- and H144A-reconstituted PSII measured in the presence of 10 mM CaCl₂ was comparable to the activity measured in the presence of 20 mM Cl⁻. We then examined the dependence of the oxygen-evolving activity of WT- and H144A-reconstituted PSII on Cl⁻ concentration (Fig. 3B). The maximum activity of WT-reconstituted PSII occurred at 1.25 mM Cl⁻. Higher concentrations of Cl⁻ gradually decreased the activity. In contrast, the maximum activity of H144A-reconstituted PSII occurred at 10 mM Cl⁻. These data suggest that the His-144 in PsbP is required to suppress the Cl⁻ requirement of PSII for oxygen-evolving activity.

D165V Mutation of the H144A Mutant PsbP Protein Restored the Ability of the Protein to Induce Conformational Change around the Mn Cluster—FTIR spectroscopy suggests that the reconstitution of PsbP restored the conformational change around the Mn cluster, and that this conformational recovery is related to the ability of PSII to retain Ca²⁺ and/or Cl⁻ (19). We have recently reported that H144A cannot restore this conformational change (20). Because H144A showed a defect in Cl⁻ retention in PSII, the conformational change induced by PsbP would be relevant to Cl⁻ binding in PSII. We then examined whether D165V and H144A/D165V could also restore the conformational change around the Mn cluster.

Fig. 4 shows the S₂Qₓₐ/SₓQₓ FTIR difference spectra of the untreated, NaCl-washed, and PsbP WT- or mutant-reconstituted PSII membranes. Only the bands in the 1700–1600 cm⁻¹ region that arise from the amide I vibrations (C=O stretches of backbone amides) of the polypeptide main chain are shown, as the changes in the other spectral regions were marginal as reported previously (19). When PsbP and PsbQ were released from PSII by washing with NaCl, the shape of the amide I bands changed from that of untreated PSII, indicating that the dissociation of PsbP and PsbQ triggered the conformational change around the Mn cluster (Fig. 4A, curve d). These changes are expressed more clearly in an untreated-minus-NaCl-washed double-difference spectrum (Fig. 4B, curve a). Reconstitution of H144A did not restore the spectral changes, and the spectrum obtained was very similar to that obtained from NaCl-washed PSII (Fig. 4, A and B, curve c), indicating that H144A could not properly restore the conformational change. Alternatively, reconstitution of WT, D165V, and H144A/D165V to NaCl-washed membranes restored the spectra, producing spectra similar to that of untreated PSII (Fig. 4, A and B, curves b, d, and e), indicating that WT, D165V, and H144A/D165V can induce proper conformational recovery during the S₁→S₂ transition of the Mn cluster. This suggests that H144A/D165V restores functional interaction between PsbP and PSII, which is required for Cl⁻ retention in PSII.

The PsbP Protein Is Directly Associated with PsbE—Our current data suggest that the C-terminal domain of PsbP, particularly the structure around the His-144 residue, is important for protein functional interaction with PSII. It has been suggested that PsbP associates with PSII via PsbO in higher plants (34). However, it can be deduced that the amide I bands perturbed by PsbP dissociation originate from the intrinsic subunits of PSII, as opposed to PsbO, because the release of PsbO from PSII did not affect any oxygen-evolving center (OEC) structures coupled to the S₁→S₂ transition (19). Therefore, it is probable that PsbP has multiple interacting sites within the PSII complex, and it should have a direct interaction with the intrinsic subunit(s) of PSII.

To address this issue, we cross-linked PsbP and PSII using the chemical cross-linker EDC with sulfo-NHS. EDC, a zero-length cross-linker, cross-links a primary amine and a carboxyl group that are electrostatically associated. Sulfo-NHS stabilizes the intermediate products of cross-linking and increases the EDC cross-linking efficiency. The cross-linked PSII complexes were analyzed by SDS-PAGE, and the cross-linked pattern of the PSII subunits was visualized by immunoblotting using specific antibodies. Fig. 5 shows the results of immunoblotting for PSII membranes cross-linked in the presence or absence of PsbP reconstitution (Fig. 5, A and B, respectively). The cross-linking pattern of major PSII intrinsic subunits such as D1, D2, CP43, CP47, and extrinsic PsbO was not significantly affected by PsbP reconstitution. Contrasting, an additional band appeared in PsbE (Cyt b₅₉₉ α subunit) immunoblotting in the presence of PsbP, and a band with the same molecular mass was also detected in the PsbP immunoblot (Fig. 5A, indicated by the arrow). This band was not observed when the PSII was treated with EDC and sulfo-NHS in the absence of PsbP (Fig. 5B). The molecular weight of this cross-linked product was shown to be about 30 kDa in the SDS-PAGE gel. This size is consistent with the theoretical molecular mass of the PsbP-PsbE cross-linked product, indicating that PsbP and PsbE were cross-linked. This cross-linked product was also observed when the carboxyl groups on NaCl-washed PSII were activated with EDC and sulfo-NHS, and subsequently incubated with PsbP (Fig. 5C). However, the product was not detected when the carboxyl groups on the PsbP protein were activated, and then incubated with non-activated PSII membranes (data not shown). This suggests that the primary amine(s) on PsbP were cross-linked with carboxyl group(s) on PsbE.

It is worth noting that a possible cross-linked product of PsbR and PsbE was also detected in the immunoblotting,
both in the presence and absence of PsbP (Fig. 5, A and B, indicated by asterisks). This suggests that PsbR is located in close proximity to PsbE. Since PsbR is not present in the cyanobacterial PSII complex, its location in PSII has not yet been clarified. Because PsbP seems to interact directly with PsbE (Fig. 5A), the location of PsbR in the vicinity of PsbE is consistent with a previous study showing that PsbR is required for the stable binding of PsbP to PSII (35, 36). However, the cross-linked products between PsbP and PsbR or including PsbP, PsbR, and PsbE were not detected in our analysis. This may indicate that PsbR plays auxiliary functions in facilitating the interaction between PsbP and PsbE, as suggested by the recent observation that PsbP can associate with PSII without PsbR.3

We then analyzed whether H144A and H144A/D165V mutations affected the cross-linking between PsbP and PsbE. The amount of the PsbP-PsbE cross-linked product was significantly decreased in the H144A-reconstituted PSII complex compared with the WT-reconstituted PSII complex, and was restored in the H144A/D165V-reconstituted PSII (Fig. 6A). These results suggest that the H144A mutation affected the PsbP local structure around His-144, which inhibited PsbP-PsbE cross-linking, and that H144A/D165V restored the structural change.

To investigate the reason for the functional recovery of the H144A/D165V mutant protein, we simulated the effects of H144A mutation on spinach PsbP structure by using the Fold X program (37). The Zn$^{2+}$ ion in the structure of spinach PsbP was ignored in this analysis. The predicted free energy change ($\Delta G$) caused by the H144A mutation was $\sim$1.5 kcal/mol, indicating that the H144A mutation has a minor destabilizing effect on PsbP folding. In fact, circular dichroism spectroscopy and analysis of Trp fluorescence did not detect major structural differences between WT and H144A PsbP (data not shown). As indicated in the structure of tobacco PsbP (Fig. 1B), His-144 should form a salt bridge with the carboxyl group of Asp-165 at physiological pH (pH 6.5, Fig. 6B). The H144A mutation is predicted to be able to break this salt bridge and alter the orientation of the Asp-165 side chain (Fig. 6B). Very similar results were obtained when the structure of tobacco PsbP was used in the analysis. It is probable that the liberated Asp-165 in the H144A protein interferes with the normal interaction between PsbP and PSII, which is required for ion retention in PSII. It is likely that the PsbP His-144 residue plays a role in maintaining the orientation of the Asp-165 residue via the formation of a salt

3 K. Ido, F. Sato, and K. Ifuku, unpublished data.

FIGURE 5. Cross-linking of PsbP with PSII membranes using EDC and sulfo-NHS. NaCl-washed PSII membranes were incubated with 6.25 mM EDC and 5 mM sulfo-NHS in either (A) the presence or (B) the absence of WT PsbP. Cross-linked proteins corresponding to 2 $\mu$g of Chl were loaded onto each lane. The proteins were immunodetected with antisera against CP47, CP43, PsbO, D2, D1, PsbP, PsbR, and PsbE, as shown on each lane. Protein size markers are shown on the left. C. NaCl-washed PSII was treated with 6.25 mM EDC and 5 mM sulfo-NHS to activate the carboxyl groups on the PSII. After activation, the PSII was washed and incubated either in the presence (+PsbP) or in the absence (−PsbP) of PsbP. As a positive control, the sample used in A was analyzed simultaneously.
The Functional Role of the His-144 of PsbP in Photosystem II

The results presented clarify the mechanism of how His-144 mutation affects PsbP function: the H144A mutation alters the interaction between the PsbP N terminus and PsbE, resulting in the increased Cl⁻ requirement of PSII. Furthermore, the interaction site between PsbP and PSII has been identified as the Ala-1 on PsbP and the Glu-57 of PsbE. Previous reports have suggested that the truncation of the first nine N-terminal residues specifically impairs the ability of PsbP to retain Cl⁻ (38). This suggests that the C-terminal domain of PsbP, including the local structure around His-144 and Asp-165, has a role of binding its N-terminal sequence to exactly the right position in PSII inducing proper conformation around the Cl⁻ binding site.

We also provide solid evidence that the Zn²⁺-binding site observed in the structure of spinach PsbP is not required for Ca²⁺ or Cl⁻ retention in PSII, because the additional mutation of D165V into the H144A mutant protein restored the ability of H144A to retain Cl⁻ ions and fully activated oxygen evolution. It was reported that a low concentration of Zn²⁺ (~2 mM) inhibited the activity of PSII and caused dissociation of PsbP and PsbQ from the core complex in spinach PSII (39, 40). Presumably, the Zn²⁺ incorporated into PsbP would hinder proper interaction between the His-144 and Asp-165 residues, and interfere with the function of PsbP. In the absence of Zn²⁺, His-144 seems to form a salt bridge with Asp-165 in PsbP. This suggests that His-144 in the PsbP plays a role in maintaining the orientation of Asp-165 such that it will not interfere with the PsbP-PSII interaction (Fig. 6B). Because the charge of the His side chain is sensitive to pH fluctuations within the physiological range, it is possible that the His-144 residue in PsbP may control the functional interaction with PSII in a pH-dependent manner.

It is widely believed that PsbO directly binds to the PSII core, PsbP binds to PsbO, and PsbQ binds to PsbP (34). However, several reports suggest that PsbO is not required for the interaction between PsbP and PsbQ (41–43), indicating direct association of PsbP and PsbQ with the PSII intrinsic subunits. The direct interaction between the PsbP N terminus and PsbE is in accordance with the previous reports, showing that N-terminal residues of PsbP are important for the binding of the protein to PSII (21, 38). The interaction between PsbP and PsbE was also reported in Chlamydomonas, although cross-linked residues have not been determined (44). However, because both Ala1 on PsbP and Glu-57 on PsbE are conserved between Chlamydomonas and higher plants, the manner of interaction...
between PsbP and PsbE is likely to be conserved in the green plant lineage.

The location of the PsbP-binding site around Cyt b$_{559}$ is also supported by genetic studies of tobacco and *Arabidopsis* plants. Transplastomic tobacco plants that have a mutation in PsbE showed lower associations between PsbP and PSII in the thylakoid membranes (45). It was reported that PsbJ in the vicinity of Cyt b$_{559}$ is required for the association of PsbP with PSII and the assembly of PsbR in tobacco (35, 46), and that PsbR is required for the stable binding of PsbP and PsbQ in *Arabidopsis* (36). In addition, we observed a putative cross-linking between PsbR and PsbE (Fig. 4), suggesting that PsbR is likely to be localized near Cyt b$_{559}$. Knockdown of PsbP expression is correlated with the decreased accumulation of PsbQ, D2, and CP47 (47), while knockdown of PsbO expression is correlated with decreased accumulation of the D1 and CP43 subunits (48), which are in close contact with PsbO. These observations strongly suggest that PsbP is localized on or near PsbE, PsbJ, and PsbR, where PsbE and PsbJ have a close interaction with the D2 subunit in the cyanobacterial PSII structure.

Although the N terminus of PsbP was determined to be involved in the interaction with PsbE, it is still unclear how the
C-terminal domain of PsbP is associated with PSII. The observation that the His-144 and Asp-165 residues in PsbP affect the functional interaction with PSII suggests that these residues should be located near the interface between PsbP and PSII. By chemical modification with N-succinimidyl propionate, which modifies primary amino groups, Tohri et al. identified the modified Lys residues on PsbP that drastically decreased the binding of PsbP to PSII. Among those residues, Lys11, 13, 33, 38, 143, 166, 170, and 174 are highly conserved in the PsbP of higher plants (49). Interestingly, these Lys residues are all located on one surface of the PsbP protein (50). The His-144 and Asp-165 residues are also located on this surface. Therefore, it is probable that His-144 and Asp-165, together with the surrounding Lys residues, are located on the interface with PSII. Presumably, the negative charge of Asp-165 that is liberated by the H144A mutation would interfere with the interaction between the Lys residues of PsbP and the negative charges of the PSII core. Further study is necessary to elucidate the amino acid residues in the C-terminal domain of PsbP that interact with PSII subunits.

In the recently described crystal structure of cyanobacterial PSII, the two Cl\(^{-}\)-binding sites were visualized near the OEC (5). At the present time, however, we cannot conclude which Cl\(^{-}\) binding site was affected by the PsbP-H144A mutation. The Cl\(^{-}\)\(1\) site is surrounded by the amino group of D2-Lys317 and the backbone nitrogen of D1-Glu333, together with two water molecules. The other Cl\(^{-}\)\(2\) site is found close to the backbone nitrogens of D1-Asn338 and CP43-Glu354, together with two water molecules. Because the side chains of D1-Glu333 and CP43-Glu354 are coordinated with the Mn cluster directly, the two Cl\(^{-}\) anions are suggested to stabilize the coordination environment of the Mn cluster. The direct interaction of PsbP with PsbE may suggest that FTIR spectroscopy of PsbP binding may reflect the changes around the Cl\(^{-}\)\(1\) binding site near the D2 side. A recent study suggests that Cl\(^{-}\)\(1\) has an important role in the transfer of protons to the lumen via hydrogen-bond networks, starting from the Mn cluster and extending toward the luminal bulk solution (51). It is possible that PsbP might be involved in this process; however, further research is required to confirm this hypothesis.

The results of this study are summarized as schematic models in Fig. 8. PsbP maintains the protein conformation around the Mn cluster in order to retain the Ca\(^{2+}\) and Cl\(^{-}\) ions in the OEC (Fig. 8A). H144A can bind to PSII; however, binding of H144A does not retain the OEC conformation for Cl\(^{-}\) retention, because the liberated Asp-165 residue alters the proper association with PSII, which affects the functional interaction between the N terminus of PsbP and PsbE (Fig. 8B). Reconstitution with the H144A/D165V double mutant restores the conformational change and ion retention because of the absence of the liberated Asp-165 that interferes with the interaction between PsbP and PSII, resulting in the recovery of PsbP-PsbE interaction (Fig. 8C).

Recently, in silico docking experiments using PSII and Cyanop structures predicted that Cyanop interacts with PsbE in the cyanobacterial PSII complex (52). If Cyanop directly interacts with PsbE, the binding sites of PsbP and Cyanop in PSII could be conserved across phyla. Alternatively, PsbP may take the place of PsbV in cyanobacterial PSII, because the N terminus of PsbV in cyanobacteria is also directly associated with PsbE. Identification of the exact binding site of Cyanop and PsbP in PSII will provide important clues for understanding how PsbP developed its crucial function in PSII during the process of evolution.

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