

PsbP-induced protein conformational changes around Cl⁻ ions in the water oxidizing center of photosystem II

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Abstract

PsbP is an extrinsic protein of PSII having a function of Ca²⁺ and Cl⁻ retention in the water-oxidizing center (WOC). In order to understand the mechanism how PsbP regulates the Cl⁻ binding in WOC, we examined the effect of PsbP depletion on the protein structures around the Cl⁻ sites using Fourier transform infrared (FTIR) spectroscopy. Light-induced FTIR difference spectra upon the S₁→S₂ transition were obtained using Cl⁻-bound and NO₃⁻-substituted PSII membranes in the presence and absence of PsbP. A clear difference in the amide I band changes by PsbP depletion was observed between Cl⁻-bound and NO₃⁻-substituted PSII samples, indicating that PsbP binding perturbed the protein conformations around the Cl⁻ ion(s) in WOC. It is suggested that PsbP stabilizes the Cl⁻ binding by regulating the dissociation constant of Cl⁻ and/or an energy barrier of Cl⁻ dissociation through protein conformational changes around the Cl⁻ ion(s).

Additional key words: Mn₄CaO₅ cluster; oxygen evolution; photosynthesis.

Introduction

Photosystem II is a multiprotein complex that has a function of oxygen evolution by water oxidation in oxygenic photosynthesis performed by plants and cyanobacteria. The catalytic site of water oxidation is the water-oxidizing center (WOC) that consists of the Mn₄CaO₅ cluster, two Cl⁻ ions, and surrounding amino acid residues (Umena *et al.* 2011, Suga *et al.* 2015). In the WOC, two water molecules are oxidized into one molecular oxygen and four protons by abstraction of four electrons using light energy (Grundmeier and Dau 2012, Messinger *et al.* 2012, Vinyard *et al.* 2013, Shen 2015). The electrons from water are transferred to the primary quinone electron acceptor Q_A and then to the secondary quinone acceptor Q_B, which becomes a quinol upon double reduction and is released into thylakoid membranes (Petrouleas and Crofts 2005). Such abstracted electrons are finally used to reduce CO₂ to synthesize sugars. Reactions in WOC are performed by a cycle of five intermediates called S_i states (*i* = 0–4) (Joliot *et al.* 1969, Kok *et al.* 1970). The reaction starts from the dark-stable S₁ state, and it advances to the S₂ state upon one-electron oxidation. Likewise, S₂ advances to the S₃ state and then to the S₄ state, which is a transient intermediate and immediately relaxes to the S₀

state by releasing a molecular oxygen. The S₀ state is oxidized to the S₁ state to complete a reaction cycle.

Amino acid residues, which ligate or directly interact with the Mn₄CaO₅ cluster and two Cl⁻ ions (designated Cl-1 and Cl-2), are all provided from the D1, D2, and CP43 subunits, which are membrane-spanning intrinsic proteins (Umena *et al.* 2011, Suga *et al.* 2015). However, some extrinsic proteins attached to the lumenal side of PSII are necessary for stabilizing the WOC structure and optimizing water oxidation activity (Seidler, 1996, Enami *et al.* 2008, Ifuku, *et al.* 2008, 2011, Fagerlund and Eaton-Rye 2011, Bricker *et al.* 2012, 2013, Ifuku and Noguchi 2016, Roose *et al.* 2016). PsbO, PsbP, and PsbQ are the extrinsic proteins involved in PSII of higher plants, while cyanobacteria have PsbV and PsbU in addition to PsbO as major extrinsic proteins. The localizations of these extrinsic proteins in PSII complexes have been revealed by X-ray crystallography (Umena *et al.* 2011, Suga *et al.* 2015) or cryo-electron microscopy (Wei *et al.* 2016). Among the extrinsic proteins in higher plants, the role of PsbP has been extensively studied (reviewed in Ifuku *et al.* 2008, 2011, Bricker *et al.* 2013, Ifuku and Noguchi 2016, Roose *et al.* 2016), and its major function was shown

Received 20 April 2017, accepted 31 May 2017, published as online-first 15 July 2017.

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Abbreviations: DCMU – 3-(3,4-dichlorophenyl)-1,1-dimethylurea; FTIR – Fourier transform infrared; Mes – 2-(*N*-morpholino) ethanesulfonic acid; PMS – phenazine methosulfate; WOC – water-oxidizing center.

Acknowledgments: This study was supported by the Grants-in-Aid for Scientific Research from the Ministry of Education, Culture, Sports, Science and Technology (17H03662 to T.N.).

to be the retention of Ca^{2+} and Cl^- ions in WOC. In order to understand the mechanism of this PsbP function, the effect of PsbP binding on the structure of WOC has been investigated using light-induced Fourier transform infrared (FTIR) difference spectroscopy (Tomita *et al.* 2009, Ido *et al.* 2012, Kakiuchi *et al.* 2012, Nishimura *et al.* 2014, 2016). This spectroscopy is a powerful method to detect the structural changes of active sites upon reactions in photosensitive proteins, and has been extensively used to study the structure and reactions of WOC (Noguchi and Berthomieu 2005, Chu 2013, Debus 2015, Noguchi 2015). It was shown that PsbP depletion from spinach PSII induced protein conformational changes in WOC (Tomita *et al.* 2009). Similar conformational changes were also detected by depletion of some extrinsic proteins in PSII complexes from a red algae (Uno *et al.* 2013) and a cyanobacterium (Nagao *et al.* 2015). From these observations, it was proposed that PsbP in higher plants and some extrinsic proteins in other phyla play a role in retaining the protein conformations of WOC to regulate its

Materials and methods

Oxygen-evolving PSII membranes of spinach were prepared as reported previously (Ono and Inoue 1986) and suspended in a pH 6.5 buffer (Cl^- -buffer: 40 mM Mes-NaOH, 400 mM sucrose, 40 mM NaCl, 5 mM $\text{Ca}(\text{OH})_2$, pH 6.5). For treatment of NO_3^- , the PSII membranes were suspended [$0.5 \text{ mg}(\text{Chl}) \text{ mL}^{-1}$] in a buffer involving 40 mM NO_3^- [NO_3^- -buffer: 40 mM Mes-NaOH, 400 mM sucrose, 40 mM NaNO_3 , 5 mM $\text{Ca}(\text{OH})_2$, pH 6.5], and washed twice with the same buffer by centrifugation. A buffer involving $\text{Na}^{15}\text{NO}_3$ (Shoko Co. Ltd., 99.1 atom %) instead of natural abundance NaNO_3 ($\text{Na}^{14}\text{NO}_3$) was used for $^{15}\text{NO}_3^-$ treatment. For depletion of the PsbP and PsbQ proteins, the PSII membranes were suspended [$0.5 \text{ mg}(\text{Chl}) \text{ mL}^{-1}$] in the Cl^- -buffer additionally involving 2 M NaCl and incubated for 20 min on ice under dark (the PAGE image of the PsbP, Q-depleted PSII prepared by this treatment was presented in Fig. S1 of Tomita *et al.* 2009). The sample was then washed twice with the Cl^- -buffer or three times with the NO_3^- -buffer by centrifugation.

For $\text{S}_2\text{Q}_\text{A}^-/\text{S}_1\text{Q}_\text{A}$ FTIR measurements, 1 mL of the sample suspension [$0.5 \text{ mg}(\text{Chl}) \text{ mL}^{-1}$] in the Cl^- - or NO_3^- -buffer in the presence of 0.1 mM 3-(3,4-dichlorophenyl)-1,1-dimethylurea (DCMU) and 0.5 mM phenazine methosulfate (PMS) was centrifuged at $170,000 \times g$ for 35 min to obtain a pellet. In the case of $\text{Q}_\text{A}^-/\text{Q}_\text{A}$ FTIR measurement,

Results and discussion

Light-induced FTIR difference spectra upon the formation of an $\text{S}_2\text{Q}_\text{A}^-$ charge separated state ($\text{S}_2\text{Q}_\text{A}^-/\text{S}_1\text{Q}_\text{A}$ difference) and only Q_A reduction ($\text{Q}_\text{A}^-/\text{Q}_\text{A}$ difference) were measured using intact and Mn-depleted PSII membranes of spinach, respectively (Fig. 1, trace a). An

reaction (Tomita *et al.* 2009, Nagao *et al.* 2015, Ifuku and Noguchi 2016). However, the molecular mechanism how PsbP specifically regulates the binding properties of Ca^{2+} and Cl^- in WOC remains to be clarified.

In this study, we investigated the effect of PsbP on the protein structures of the Cl^- binding sites of WOC using light-induced FTIR difference spectroscopy. For this purpose, we examined the effect of replacement of Cl^- with NO_3^- on the PsbP-induced structural changes in WOC, which were detected by FTIR difference measurement upon the $\text{S}_1 \rightarrow \text{S}_2$ transition. It has been shown that NO_3^- substitution for Cl^- retains O_2 evolution activity but with a decreased efficiency in the $\text{S}_3 \rightarrow \text{S}_0$ transition (Sinclair 1984, Wincencjusz *et al.* 1999, Hasegawa *et al.* 2004, Suzuki *et al.* 2012). It is predicted that NO_3^- substitution induces perturbations in the protein moieties around Cl^- ions. The obtained results clearly showed that PsbP binding affects the protein conformations around the Cl^- ions, which should cause the alteration in the binding properties of the Cl^- ions in WOC.

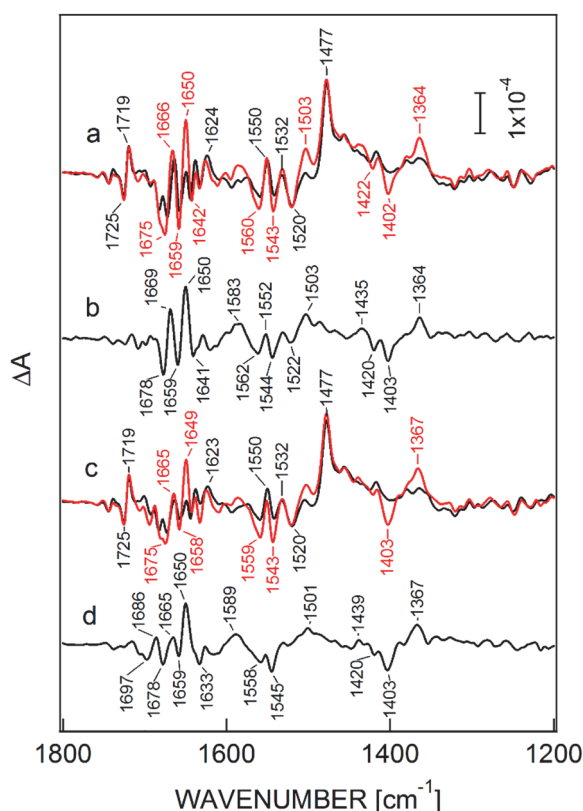
the last buffer for centrifugation additionally involved 10 mM NH_2OH , which depletes the Mn_4CaO_5 cluster and functions as an exogenous electron donor during measurement. The pellet obtained by centrifugation was sandwiched between two CaF_2 plates (25 mm diameter). One of the CaF_2 plates had a circular groove (10 mm inner diameter, 1 mm width), and the sample cell was sealed with silicone grease laid on the outer part of the groove (Noguchi and Sugiura 2001). The sample temperature was adjusted to 283 K by circulating cold water in a copper holder.

Light-induced $\text{S}_2\text{Q}_\text{A}^-/\text{S}_1\text{Q}_\text{A}$ and $\text{Q}_\text{A}^-/\text{Q}_\text{A}$ FTIR difference spectra were recorded using a Bruker VERTEX 80 spectrophotometer equipped with an MCT detector (D313-L, InfraRed Associates, Inc., USA). Flash illumination was performed using a Q-switched Nd:YAG laser (Quanta-Ray INDI-40-10, Spectra-Physics, USA; 532 nm; ~ 7 ns fwhm; $\sim 7 \text{ mJ pulse}^{-1} \text{ cm}^{-2}$). Single-beam spectra (10-s scans) were recorded before and after a single flash followed by dark relaxation for 5 min. This cycle was repeated 40–80 times and averaged spectra were used to calculate a light-induced difference spectrum. Spectra measured using a couple of samples were averaged to improve signal-to-noise ratios.

FTIR difference spectrum upon the $\text{S}_1 \rightarrow \text{S}_2$ transition (S_2/S_1 difference) was obtained by subtraction of the $\text{Q}_\text{A}^-/\text{Q}_\text{A}$ spectrum from the $\text{S}_2\text{Q}_\text{A}^-/\text{S}_1\text{Q}_\text{A}$ spectrum so as to cancel the strong band at $1,477 \text{ cm}^{-1}$ arising from the CO/CC stretching vibration of the Q_A^- semiquinone anion

(Berthomieu *et al.* 1990, Ashizawa and Noguchi 2014) (Fig. 1, trace *b*). This S_2/S_1 spectrum is virtually identical to the spectrum of spinach PSII membranes reported previously (Noguchi *et al.* 1995, Tomita *et al.* 2009). Prominent bands in the 1,700–1,600 cm^{-1} and 1,450–1,350 cm^{-1} have been assigned to the amide I vibrations (CO stretches) of backbone amides and the symmetric COO^- stretching vibrations of carboxylate groups around the Mn_4CaO_5 cluster, respectively (Noguchi and Sugiura 2003, Nakamura and Noguchi 2016). In the 1,600–1,500 cm^{-1} region, bands of the amide II vibrations (NH bends + CN stretches) of backbone amides and those of the asymmetric COO^- vibrations of carboxylate groups overlap (Noguchi *et al.* 1995). The presence of strong amide I and II bands reflects the perturbation of protein conformations around the Mn_4CaO_5 cluster coupled with the $S_1 \rightarrow S_2$ transition.

The S_2/S_1 difference spectrum of PSII membranes, in which Cl^- ions in WOC were replaced with NO_3^- ions, was obtained in an analogous way (Fig. 1, trace *d*) as a double difference spectrum between the $\text{Q}_\text{A}^-/\text{Q}_\text{A}$ and $\text{S}_2\text{Q}_\text{A}^-/\text{S}_1\text{Q}_\text{A}$ difference spectra (Fig. 1, trace *c*). Note that at the present stage it is uncertain whether both of the two Cl^- ions (Cl-1 Cl-2) were replaced with NO_3^- ions or only one of them



traces *b*, *d*). The intensities of the peaks at 1,678/1,669 cm^{-1} significantly decreased and minor peaks appeared at 1,697/1,686 cm^{-1} . These changes indicate that protein conformations around the Cl^- ion(s) are perturbed by NO_3^- substitution.

Comparison of the S_2/S_1 difference spectrum of PsbP-depleted PSII membranes (Cl^- -bound PSII) with that of intact PSII showed some spectral changes (Fig. 3*A*, trace *a*). These are clearly shown in their double difference spectrum (Fig. 3*A*, trace *b*); prominent bands were observed at 1,683, 1,666, 1,657, 1,649, 1,635, and 1,325 cm^{-1} in the amide I region together with some minor bands at 1,560 and 1,539 cm^{-1} in the amide II region, although no appreciable bands were found in the symmetric COO^- region (1,450–1,300 cm^{-1}). This spectral feature is identical to that of a corresponding spectrum in our previous study (Tomita *et al.* 2009), in which the large amide I changes were attributed to the perturbation of the protein conformations of WOC induced by the removal of PsbP. Note that although PsbQ was also removed together with PsbP by high-concentration NaCl treatment, only PsbP is effective to the protein conformation of WOC, because rebinding of PsbP recovered the spectral features (Tomita *et al.* 2009).

Corresponding S_2/S_1 spectra of NO_3^- -substituted PSII membranes and a double difference spectrum (Fig. 3*A*, traces *c* and *d*, respectively) showed a similar tendency to those of Cl^- -bound PSII; the spectral feature in the amide I region (1,700–1,600 cm^{-1}) was significantly changed, whereas that in the symmetric COO^- region (1,450–1,300 cm^{-1}) was little changed. The double difference spectra of NO_3^- -substituted PSII and Cl^- -bound PSII were compared in an expanded view of the amide I region (Fig. 3*B*). Upon NO_3^- substitution, the prominent positive peak at 1,666 cm^{-1} almost disappeared, and the negative peak at 1,683 cm^{-1} was replaced with more complex peaks at 1,694/1,686/1,680 cm^{-1} . In addition, the intensities of the positive peaks at 1,649 and 1,625 cm^{-1} slightly increased. These changes indicate that PsbP-induced protein conformational changes in WOC were affected by NO_3^- substitution for Cl^- . In other words, we detected specific perturbations of the protein conformations around the Cl^- ions by the depletion of PsbP. It is likely that the largely changed amide I bands at 1,683/1,666 cm^{-1} are attributed to the polypeptide chains surrounding the Cl^- ions in WOC.

The next question is how the PsbP binding/depletion affects the protein conformations at the Cl^- binding sites. The recent spinach PSII structure at a 3.2 Å resolution obtained by cryo-electron microscopy showed the interactions of extrinsic proteins with the intrinsic proteins forming WOC (Fig. 4) (Wei *et al.* 2016). Also, the high-resolution (1.9–1.95 Å) X-ray crystallographic structures of cyanobacterial PSII revealed the detailed structure of WOC involving the Cl^- -binding sites (Umena *et al.* 2011, Suga *et al.* 2015). Cl^- 1 is anchored by N181 and E333 of the D1 protein and K317 of the D2 protein, while Cl^- 2 is anchored by N338 and F339 of the D1 protein and D354

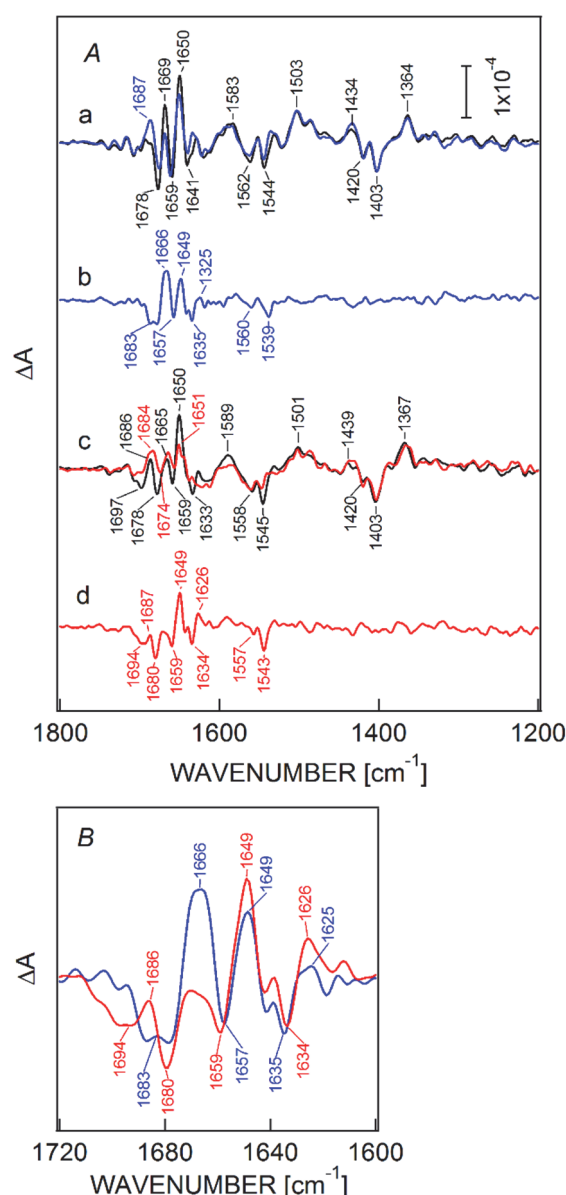


Fig. 3. (A) S_2/S_1 FTIR difference spectra of PsbP-depleted PSII membranes (*a* – blue line; *c* – red lines) in comparison with those of control (PsbP-intact) PSII membranes (*a*, *c* – black lines), and control-minus-PsbP-depleted difference spectra (*b*, *d*). (*a*, *b*) Cl^- -bound PSII; (*c*, *d*) NO_3^- -substituted PSII. (B) Expanded view of the amide I region of the control-minus-PsbP-depleted difference spectra of Cl^- -bound (blue line) and NO_3^- -substituted (red line) PSII membranes.

of the CP43 protein (Umena *et al.* 2011, Suga *et al.* 2015). Here, we assume van der Waals interactions that are determined only by a close contact between subunits, because the higher resolution structures of side chains are necessary to argue electrostatic and hydrogen bonding interactions. PsbP interacts with the D1 protein mainly in the two regions (Fig. 4*A*): the region formed by E140, G141, and K166 in PsbP interacting with P340 and L341 in the C-terminus loop region of D1, and the region formed

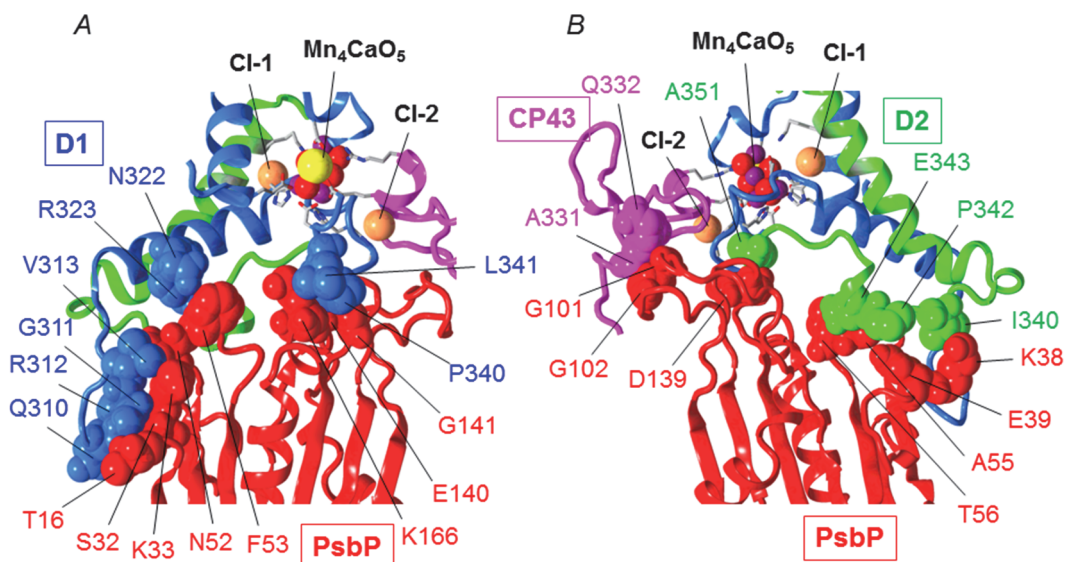


Fig. 4. Interactions of PsbP with the D1 subunit (A) and the D2 and CP43 subunits (B). The protein structures were produced from the PDB data (entry code: 3JCU) of spinach PSII obtained by cryo-electron microscopy (Wei *et al.* 2016). Different colors are used to express individual subunits: PsbP (red), D1 (blue), D2 (green), and CP43 (magenta). The Mn₄CaO₅ cluster, Cl⁻ ions, and amino acid residues interacting between PsbP and other subunits are shown in balls with van der Waals radii.

by T16, S32, K33, N52, and F53 in PsbP interacting with Q310–V313 in the loop region and N322 and R323 in the helical region of D1. The first region is very close to the Cl-2 binding site and may directly affect it, while the latter interaction may be effective to the Cl-1 site. The interaction with the D2 protein exists in the region formed by K38, E39, A55, and T56 in PsbP interacting with I340–E343 in D2 (Fig. 4B). This interaction could affect the Cl-1 site though the C-terminal helix of D2. In addition, D139 in PsbP interacts with A351 in D2 (Fig. 4B), which interacts with the loop region of D1 near Cl-1 and hence can affect the Cl-1 site. Furthermore, G101 and G102 in PsbP interact with A331 and Q332 in CP43 near Cl-2 (Fig. 4B). These interactions of PsbP with D1, D2, and CP43 may contribute to the protein conformational changes around the Cl⁻ ions upon PsbP removal. It is noteworthy that similar sites in the D1 and D2 proteins (K310–V313, R323, P340, and L341 of the D1 protein and E343 of the D2 protein) interact with the PsbV extrinsic protein in the cyanobacterium *Thermosynechococcus vulcanus* (Suga *et*

al. 2015), suggesting a similar effect on the WOC structure between PsbP in higher plants and PsbV in cyanobacteria. In contrast to PsbP, PsbQ interacts with neither the D1 nor the D2 protein, but interacts with the CP43 protein at sites rather far from the Cl⁻ ions, which is consistent with the previous FTIR observation that PsbQ binding did not affect the FTIR spectrum (Tomita *et al.* 2009).

It has been known that depletion of PsbP enhances the Cl⁻ requirement in O₂ evolution, and decreases the activity at a low concentration of Cl⁻ (Miyao *et al.* 1988, Seidler 1996, Ifuku *et al.* 2008). The results of the present study showed that PsbP binding induces the protein conformational changes around the Cl⁻ ion(s). It is highly likely that such perturbations of protein conformations change the dissociation constant of the Cl⁻ ions and/or the energy barrier of Cl⁻ dissociation from the binding sites. We thus conclude that the PsbP extrinsic protein regulates the Cl⁻ binding and hence the reaction of WOC by perturbing the protein conformations around the Cl⁻ binding sites.

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