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Authors

Bhowmick, Asmit

Hussein, Rana

Bogacz, Isabel

et al.

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Asmit Bhowmick^{1,14}, Rana Hussein^{2,14}, Isabel Bogacz^{1,14}, Philipp S. Simon^{1,14}, Mohamed Ibrahim^{2,13}, Ruchira Chatterjee¹, Margaret D. Doyle¹, Mun Hon Cheah³, Thomas Fransson⁴, Petko Chernev³, In-Sik Kim¹, Hiroki Makita¹, Medhanjali Dasgupta¹, Corey J. Kaminsky¹, Miao Zhang¹, Julia Gätcke², Stephanie Haupt², Isabela I. Nangca¹, Stephen M. Keable¹, A. Orkun Aydin³, Kensuke Tono^{5,6}, Shigeki Owada^{5,6}, Leland B. Gee⁷, Franklin D. Fuller⁷, Alexander Batyuk⁷, Roberto Alonso-Mori⁷, James M. Holton^{1,8,9}, Daniel W. Paley¹, Nigel W. Moriarty¹, Fikret Mamedov³, Paul D. Adams^{1,10}, Aaron S. Brewster¹, Holger Dobbek², Nicholas K. Sauter¹, Uwe Bergmann¹, Athina Zouni^{2,12}, Johannes Messinger^{3,12}, Jan Kern¹, Junko Yano^{1,13} & Vittal K. Yachandra^{1,13}

In natural photosynthesis, the light-driven splitting of water into electrons, protons and molecular oxygen forms the first step of the solar-to-chemical energy conversion process. The reaction takes place in photosystem II, where the Mn₄CaO₅ cluster first stores four oxidizing equivalents, the S₀ to S₄ intermediate states in the Kok cycle, sequentially generated by photochemical charge separations in the reaction center and then catalyzes the O–O bond formation chemistry^{1–3}. Here, we report room temperature snapshots by serial femtosecond X-ray crystallography to provide structural insights into the final reaction step of Kok's photosynthetic water oxidation cycle, the S₃→[S₄]→S₀ transition where O₂ is formed and Kok's water oxidation clock is reset. Our data reveal a complex sequence of events, which occur over micro- to milliseconds, comprising changes at the Mn₄CaO₅ cluster, its ligands and water pathways as well as controlled proton release through the hydrogen-bonding network of the ClI channel. Importantly, the extra O atom O_x, which was introduced as a bridging ligand between Ca and Mn1 during the S₂→S₃ transition^{4–6}, disappears or relocates in parallel with Y₂ reduction starting at approximately 700 μs after the third flash. The onset of O₂ evolution, as indicated by the shortening of the Mn1–Mn4 distance, occurs at around 1,200 μs, signifying the presence of a reduced intermediate, possibly a bound peroxide.

Serial femtosecond X-ray crystallography at X-ray free electron lasers (XFELs)⁷ enabled us to collect crystallography data of photosystem II (PS II) in real time as the reaction progresses at physiological temperature. The four photon-induced water-oxidation reaction in PS II (Fig. 1a,b) was initiated with multiple visible laser flashes. Using this capability, intermediate S-state structures (S₀, S₁, S₂ and S₃) have been studied^{4,5,8–10} that revealed the structural changes of the oxygen evolving complex (OEC) of PS II, which is a functional unit composed of the Mn₄CaO₅ cluster and its water–ligand environment (Fig. 1c,d)^{2,11}. Recently, we collected snapshot data at several time points during the S₂→S₃ transition, the step in which one substrate water is introduced into the cluster. The study⁸ suggested the sequence of Mn oxidation, incorporation of an extra oxygen bridge (O_x or O₆ in Suga et al.¹⁰) between the open coordination site at Mn1 and Ca (forming Mn₄CaO₅–O_x in S₃),

the potential entry path for substrate water and the proton release with its gating mechanism^{6,12}. The XFEL studies also clearly established that the electronic and geometric structure of the OEC obtained by these measurements is unaffected by X-ray photoelectrons under the conditions used^{6,8,13}.

In the current study, we investigate the oxygen evolving step of Kok's water oxidation cycle (Fig. 1b), the S₃→[S₄]→S₀ transition (Fig. 1c,d). In dark-adapted PS II samples, this transition is initiated by the third visible laser flash. The OEC is oxidized in this step from the all-Mn(IV) S₃ state to the proposed highly reactive S₄ state with formal oxidation states of Mn(IV)₄O* or Mn(IV)₃(V). This initiates O–O bond formation and O₂ release, and the now vacant binding site is filled by a new water substrate forming the lowest oxidation state of the cluster (S₀). This multistep process, which also involves

¹Molecular Biophysics and Integrated Bioimaging Division, Lawrence Berkeley National Laboratory, Berkeley, CA, USA. ²Department of Biology, Humboldt Universität zu Berlin, Berlin, Germany.

³Molecular Biomimetics, Department of Chemistry – Ångström, Uppsala University, Uppsala, Sweden. ⁴Department of Theoretical Chemistry and Biology, KTH Royal Institute of Technology, Stockholm, Sweden. ⁵Japan Synchrotron Radiation Research Institute, Hyogo, Japan. ⁶RIKEN SPring-8 Center, Hyogo, Japan. ⁷Linac Coherent Light Source, SLAC National Accelerator Laboratory, Menlo Park, CA, USA. ⁸Department of Biochemistry and Biophysics, University of California, San Francisco, CA, USA. ⁹SSRL, SLAC National Accelerator Laboratory, Menlo Park, CA, USA. ¹⁰Department of Bioengineering, University of California, Berkeley, CA, USA. ¹¹Department of Physics, University of Wisconsin–Madison, Madison, WI, USA. ¹²Department of Chemistry, Umeå University, Umeå, Sweden. ¹³Present address: Institute of Molecular Medicine, University of Lübeck, Lübeck, Germany. ¹⁴These authors contributed equally: Asmit Bhowmick, Rana Hussein, Isabel Bogacz, Philipp S. Simon. ✉e-mail: athina.zouni@hu-berlin.de; johannes.messinger@kemi.uu.se; jyano@lbl.gov; vkyachandra@lbl.gov

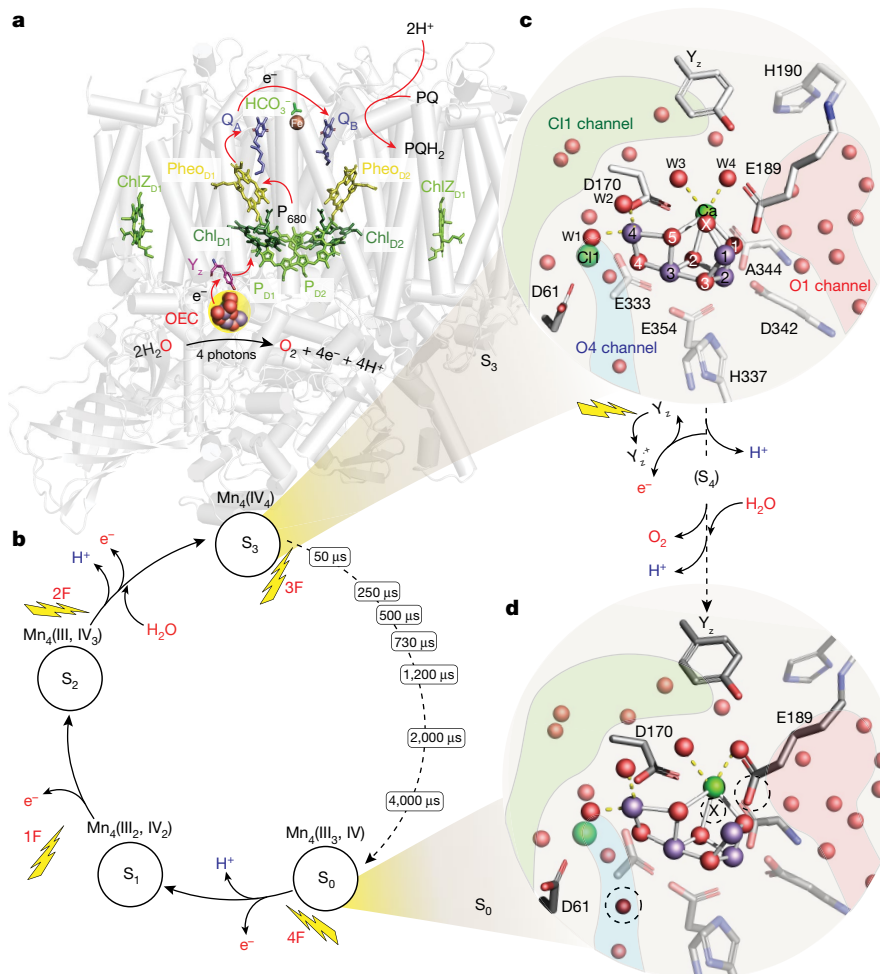


Fig. 1 | An overview of PS II and the electron donor site where water oxidation takes place. a, The structure of PS II with the membrane-embedded helices and the membrane extrinsic regions on the luminal side of PS II shown in gray. The main electron transfer components are shown in colour, which include the reaction center chlorophylls (P680), pheophytins, acceptor quinones Q_A and Q_B, redox-active tyrosine Y_Z and the catalytic Mn₄CaO₅ cluster. The Y_Z and Mn₄CaO₅ cluster are the cofactors of the electron donor site. **b,** Kok cycle of the water oxidation reaction taking place at the donor site that is sequentially driven by charge separations in the reaction center P680 induced by the absorption of

photons (nanosecond light flashes, 1F–4F) in the antenna system of PS II. Room temperature X-ray crystallography data were collected at the time points indicated during the S₃→S₀ transition. **c,d,** The structure of the OEC in the S₃ (**c**) and S₀ (**d**) states and the sequence of events occurring between them. Mn, purple; Ca²⁺, green; O, red. W1, -2, -3 and -4 are water ligands of Mn4 and Ca. The relevant channels for water and proton transfer (O1, O4 and Cl1) are indicated as red, blue and green shaded areas, respectively. The dotted circles mark structural differences between the S₃ and S₀ states.

the release of two protons, has the longest time constant among the S-state transitions, and its kinetics depend on the species and sample preparation^{14–19}.

To provide structural insight into this complex reaction step and specifically, the important interplay between the Mn₄CaO₅ cluster and its protein–water environment (Fig. 1c,d), we collected room temperature crystallography data of PS II at seven different time points during the S₃→S₀ transition, ranging from 50 μs to 4 ms after initiating this transition (Fig. 1b). All datasets have resolutions between 2.00 and 2.16 Å (Extended Data Tables 1 and 2).

The S₃ state was populated by illumination of dark-adapted PS II microcrystals with two in situ visible nanosecond laser flashes (2F in Fig. 1b)⁴, in which the interval between flashes was 200 ms to account for acceptor quinone Q_A and Q_B kinetics and efficiently drive S-state transitions. The time points between the S₃ and S₀ states were generated by giving the third visible pump laser flash at various delay times (Δt) before the crystals were exposed to the femtosecond XFEL pulse (the time labels in Fig. 1b). While our illumination protocol achieves the highest possible populations of particular S states, there is a higher mixing

of S-state populations with increasing flash number due to intrinsic PS II-specific inefficiencies (‘misses’) (Methods)²⁰. We model this distribution in a multicomponent model during structural refinement, with the ‘primary’ component being the centers that advance from the S₃ to S₀ state. The ‘secondary’ and ‘tertiary’ components are the known starting and end points: for example, the decreasing S₃ population and at longer delay times, the increasing S₀ population from centers that have completed the transition (Methods and Extended Data Table 3 have details). We note that the primary component at each time point may consist of a mixture of multiple structures, which are intermediates between S₃ and S₀. All results discussed below correspond to these refined primary components from monomer I (chains annotated as uppercase in the deposited structures).

OEC and the Y_Z region

Figure 2 shows the omit map density of selected atoms at the OEC and its surroundings of the refined population at the time points (Δt = 250, 500, 730, 1,200 and 2,000 μs) after the third flash (we use

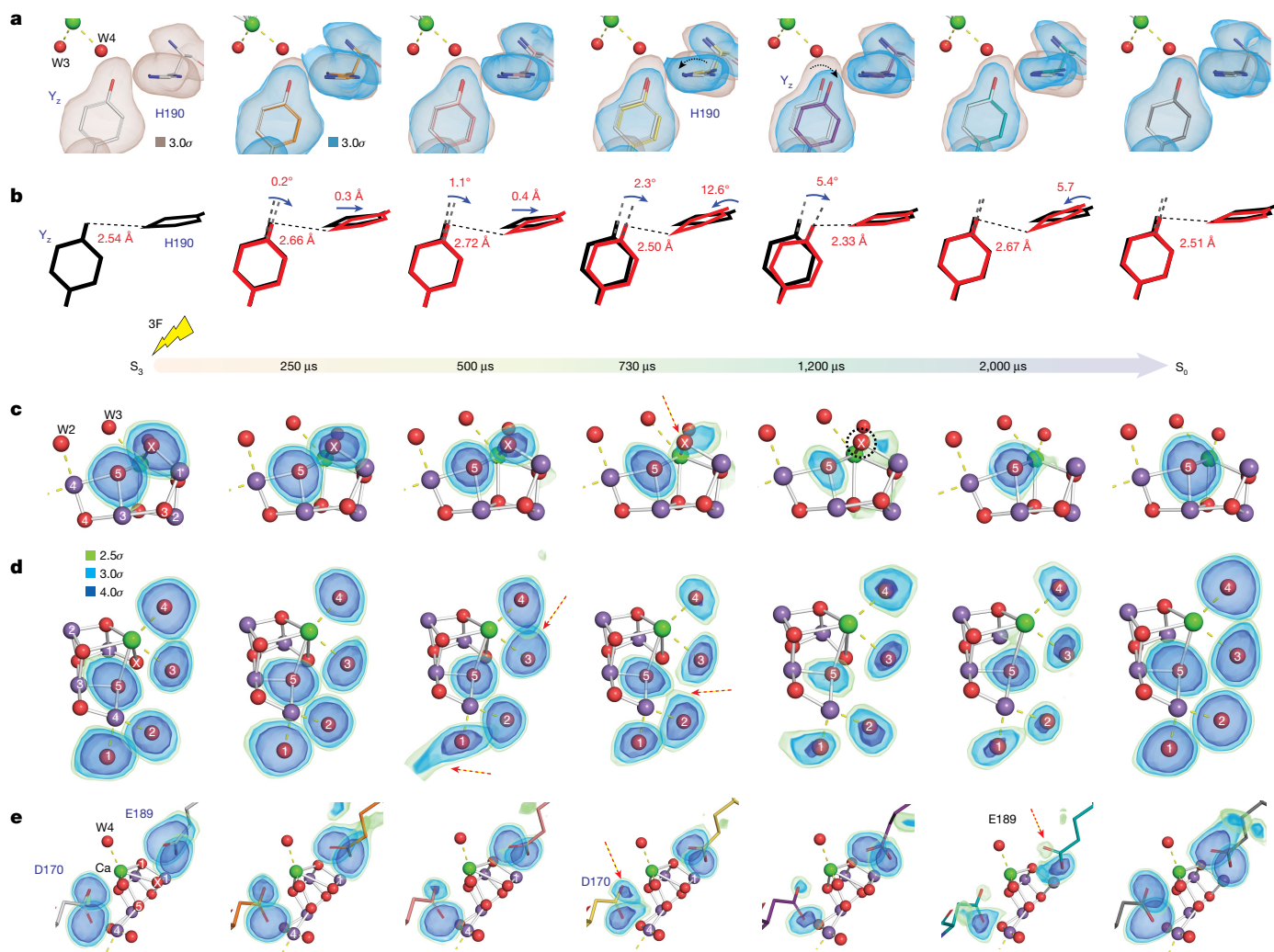


Fig. 2 | $mF_{\text{obs}} - DF_{\text{calc}}$ electron density omit map of key components of the redox active donor site of PS II at five time points along the $S_3 \rightarrow S_0$ transition as well as the S_3 and S_0 states. **a**, Residues D1-Y161 (Y_z) and D1-H190. The omit map from the S_3 state reference is shown in light brown for comparison with the time point data (blue). **b**, A simplified representation of the structural changes observed at the Y_z region. **c**, Omit density of atoms O_5 and O_x of the OEC. **d**, Omit density of atoms O_5 and the terminal water ligands W_1 , W_2 , W_3 and W_4

of the OEC. **e**, Omit density of carboxylate oxygen atoms of D1-E189 and D1-D170. All omit maps shown in **a** and **c–e** were generated by omitting the atom or residue of interest individually, and only the primary component (that is, the state that is advancing to S_0) was used. Notable features are highlighted with red arrows and black dashed circle. All omit maps shown are contoured at 2.5σ , 3σ and 4σ using the colour scheme annotated in **d** for easier visualization. See also Supplementary Information Video 1.

the nomenclature of $3F(\Delta t \mu\text{s})$). To visualize the sequence of events during the $S_3 \rightarrow [S_4] \rightarrow S_0$ transition, we follow the changes in three areas: Y_z and D1-H190 (Fig. 2a,b) and O_x and O_5 (Fig. 2c), as well as the water ligands W_1 – W_4 (Fig. 2d) and two carboxylate ligands of the $\text{Mn}_4\text{CaO}_5/\text{Mn}_4\text{CaO}_5\text{-O}_x$ cluster that bridge between Mn and Ca (Fig. 2e). Additionally, selected atomic distance changes are shown in Fig. 3, including the earlier (50 μs) and later (4,000 μs) time point data.

The distance between Y_z and D1-H190 has been established previously to be an indicator of the oxidation state of Y_z ⁶. In the reduced state, a strong hydrogen bond between Y_z and D1-H190 leads to a short distance of about 2.6 Å. Upon oxidation of Y_z by P680^+ , the phenolic proton of Y_z is transferred to D1-H190, the distance increases to 2.8 Å and a movement of the His ring plane is observed (Fig. 2b). Our data show that the distance already increases between the S_3 data and the first time point (50 μs), indicating that Y_z is fully oxidized by this time (Fig. 3). Thereafter, this distance remains constant until 500 μs and returns to the base level between 730 and 1,200 μs . Thus, the data demonstrate that Y_z^{ox} reduction by the $\text{Mn}_4\text{CaO}_5\text{-O}_x$ cluster starts only after around 500 μs (Extended Data Table 4)^{16,21,22} and appears to be complete by the

1,200- μs time point. Additional distance changes between Y_z and His190 are observed at 2,000 and 4,000 μs , which may be due to the rearrangement of the hydrogen bonding network related to the last proton release but are not well understood currently. Interestingly, a shift in the position of a Tyr residue next to the special pair Chl upon light excitation was previously noted in time-resolved crystallography data of the purple bacterial reaction center and interpreted as originating from a change in the hydrogen-bonding interactions of this Tyr upon deprotonation²³.

The extra oxygen O_x , present in the S_3 state, is lost upon S_0 formation, indicating that O_x may participate in the O–O bond formation, and changes in its density probably inform on the onset of this process. For tracing the O_x population, O_x was eliminated from the OEC model, and its omit map density is shown in Fig. 2c. O_5 was separately omitted, allowing comparison of their densities (see also Extended Data Fig. 1). The O_x omit density becomes asymmetric starting at 250 μs , with a clear reduction in intensity after 500 μs and dropping to the noise level between 1,200 and 2,000 μs , and O_x can only be modeled with a population of less than 20% in the 2,000 μs time point. The O_x intensity

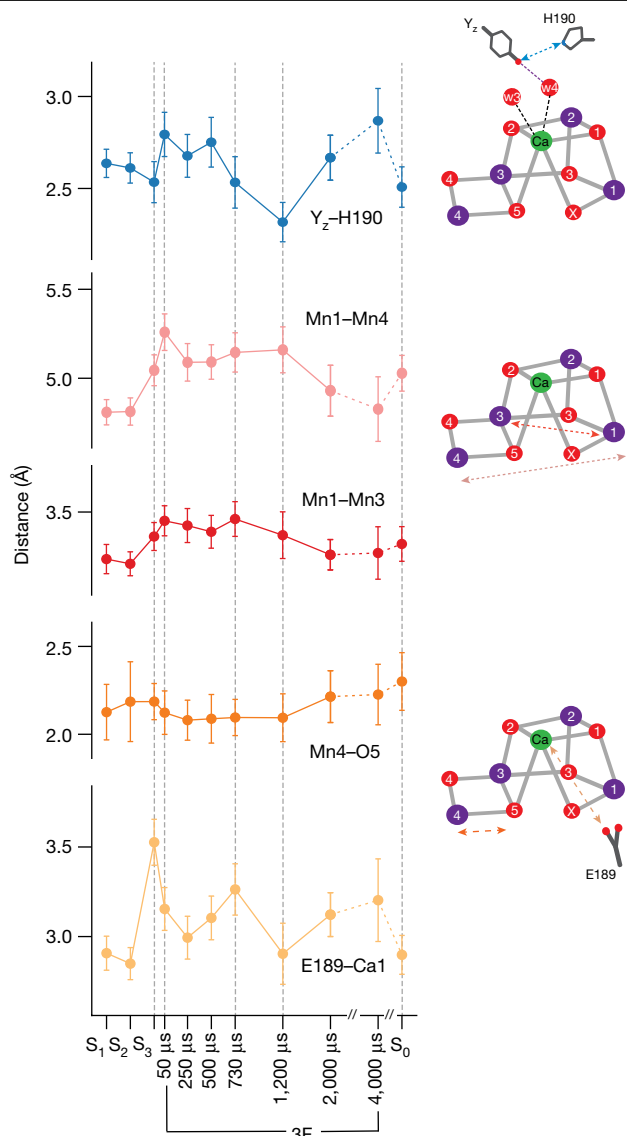


Fig. 3 | Distance changes between selected atoms/residues in the OEC during the $S_3 \rightarrow S_0$ transition. All distances are taken from the refined component of each time point (that is, the state that is advancing to the S_0 state). Error bars are calculated from the end/rapid approach described in Methods and are an upper limit. Data here are shown as mean values \pm standard deviation. The error bars for each time point were obtained from $n = 100$ independent END/RAPID refinements. More details about the END/RAPID procedure can be found in Methods. Dashed arrows in the schematics of the OEC on the right indicate the location of the individual distances. Mn is shown as purple spheres, and O is shown as red spheres.

changes occur concomitantly with Y_z^{Ox} reduction, indicating that the O–O bond formation occurs between 500 and 1,200 μ s. During the entire $S_3 \rightarrow [S_4] \rightarrow S_0$ transition, the O5 density remains approximately constant, except for a decrease of its electron density at 1,200 μ s.

Additional markers for the presence of O_x in the cluster are Mn–Mn distances. As reported previously, the Mn1–Mn4 distance increases during the $S_2 \rightarrow S_3$ transition due to the insertion of O_x (ref. 6). Figure 3 shows that the Mn1–Mn4 distance remains elongated in the 3F structures until 1,200 μ s (Methods has a more detailed analysis) and then declines over the next 3 ms to attain the same value as seen in the S_1 and S_2 states. A similar trend is also seen in the Mn1–Mn3 distance. Thus, there is a delay between the onset of O–O bond formation (500–730 μ s based on the changes observed for Y_z and O_x) and the time when the

Mn–Mn distances start to decrease (1,200 μ s). The finding indicates that during this period, a water oxidation intermediate likely exists before the release of O_2 .

Changes in shape and intensities are observed for the omit map density of the water ligands W1–W4, which are displayed in Fig. 2d. Specifically, a slight elongation of the O5 density toward W2 and a slight elongation of the W4 density toward W3 are observed at 250 μ s. This latter trend continues, and at 500 μ s, an overlap of the W3 and W4 densities is seen, with an $mF_{obs} - DF_{calc}$ peak (F_{obs} and F_{calc} are the experimental and model structure factors respectively, while m and D are weighting factors) at 2.5σ between W3 and W4 (Extended Data Fig. 2). At the same time point, the W1 density becomes extended toward D1–D61 and W19, indicating a higher mobility of W1. We speculate that all these motions are related to the deprotonation of the OEC and proton transfer toward the ClI channel (see the next section and Fig. 4).

At 730 μ s, the O5, W2 and W3 densities become anisotropic, all pointing toward a region between these three oxygen atoms (Fig. 2d), which indicates increased mobility of these ligands. At the same time, the O_x density is also highly anisotropic. This movement of all four oxygen atoms is likely related to the formation of the water oxidation intermediate. We note that accurate modeling of the O_x position will require higher-resolution data. At 1,200 μ s, the densities for all the terminal water ligands (W1–W4) and the bridging O5 become weakest. A comparison of these omit map densities within the OEC with that of the O2 atom, which is believed to not play a prominent role in the $S_3 \rightarrow S_0$ step, shows that the reduction of the density is specific to these five atoms (W1–W4 and O5) and O_x (Extended Data Fig. 1). The overlap of the O5, W2 and W3 densities is no longer observed at 1,200 μ s.

At 2,000 μ s after the third flash, the O5 omit map density is restored considerably compared with the S_3 and S_0 states, but omit map densities of the waters W1–W4 have not yet reached a similar level and remain elongated. We note a similar overlap of W1/W2 as observed at 500 μ s. This could indicate the onset of the second proton release, known to occur in the $S_3 \rightarrow S_0$ transition, after the binding of a water that refills the vacant site formed by O_2 release^{21,24,25}. Interestingly, the elongated shape of the W3 density persists even in the S_0 state, which we modeled previously with two possible positions of W3 (ref. 4).

Figure 2e shows that the D1–D170 and D1–E189 ligands, which both bridge between an Mn and Ca, change their conformation during O_2 formation and release. Consistent with the high mobility observed for the W1 and W2 water ligands, the connection between D1–D170 and Ca appears to be weakened between 730 and 2,000 μ s and is only fully restored at the S_0 state (3F(200 ms)).

From 1,200 to 4,000 μ s, several structural changes occur, which are reversed upon formation of the stable S_0 state (3F(200 ms)). These include the increase of the Y_z –D1–H190, Ca–D1–E189 and Mn4–O5 distances, as well as a decrease of the Mn1–Mn4 and Mn1–Mn3 distances. Most of these changes are indicative of O_2 release and/or water insertion via the Ca ion^{26,27}, possibly from the O1 channel (see below). This indicates that O_2 release and refilling of the cluster by bulk water and resetting of the catalytic center occur over an extended timescale.

Water and proton channels

PS II has several hydrophilic channels that extend from the OEC to the luminal side of the thylakoid membrane^{11,28–31}, and some of these are proposed to play a critical role in transporting protons and substrate waters during the catalytic cycle (O1, O4 and ClI channels are shown in Fig. 1)^{24,32,33}. In our recent study of the $S_2 \rightarrow S_3$ transition¹², we assigned the O1 channel, which extends from the O1 and Ca of the OEC to the bulk, to be a substrate water channel and the ClI channel, which extends from W1 and W2 of the OEC to the bulk, to be a proton release channel during the $S_2 \rightarrow S_3$ transition⁶. The $S_3 \rightarrow S_0$ transition also involves the insertion of one substrate water into the OEC and the release of two

protons to the bulk; it has been suggested that one proton is released before the O–O bond formation and the other after the rebinding of a water molecule to the OEC^{24,25,34}.

Figure 4 shows the time point data for the O1 and the ClI/O4 channels near the OEC during the $S_3 \rightarrow S_0$ transition. The electron density of water molecules in the O1 channel in the vicinity of the OEC changes substantially as shown in the $2mF_{obs} - DF_{calc}$ maps (Fig. 4a), similar to what was observed during the $S_2 \rightarrow S_3$ transition⁶. Waters W27, W28 and W32 (Supplementary Table 1 has water numbering) have low electron density and high B factors (about 50 \AA^2) (Extended Data Fig. 3) relative to the more stable waters, such as W29 (B factor of about 37 \AA^2), in particular at 1,200 μs . We interpret this as an indication of the high mobility of these waters and hypothesize that this region could serve as the inlet for the substrate water that refills the OEC after the release of molecular oxygen¹². We, therefore, propose that PS II uses the O1 channel for the substrate intake in both the $S_2 \rightarrow S_3$ and $S_3 \rightarrow S_0$ transitions. Among the group of five waters (W26–30) ('water wheel' in ref. 6), W26 shows high electron density throughout the transition, with a substantially elevated density at 500 μs . As W26 is within hydrogen-bonding distance to O1 of the OEC, this interaction may be important for balancing the charge on the cluster, when the OEC advances through the last oxidation step (that is, S_4 state formation) and the subsequent four-electron reduction to form the S_0 state.

Changes are also observed in the ClI channel. At 250 μs , the D1-E65 residue rotates by 19° toward W40, resulting in shortening the distance between D1-E65 and W40 by 0.3 \AA (Fig. 4b). This is the time point when the O_x density starts to become asymmetric, which is even more pronounced at 500 μs (Fig. 2c). At 500 μs , the W1 omit map density also becomes elongated toward the region of D1-D61 and W19 (Fig. 4b). This coincides with a decrease of the D1-D61 carboxylate oxygen density that is within hydrogen bond distance of W40 and a shortening of the distance between W40 and D1-E65 by 0.5 \AA due to rotation (25°) of the side chain. Consequently, a continuous hydrogen bond network is formed that connects the OEC to the D1-E65/D2-E312 region. We speculate that the changes are related to the first proton transfer from the OEC toward the ClI channel. These changes are reversed by the 730- μs time point (Fig. 4b). An early deprotonation event has also been suggested by other studies using different methods, with time constants that range from 50 to 300 μs (that is, before the last oxidation event; the transient S_4 state formation)^{15–17,21,24,34,35}.

At 1,200 μs , the D1-E65 residue rotates toward W40 for the second time during this transition, accompanied by a shortening of the W42–D1-E65 distance by 0.4 \AA . Thus, a hydrogen bond network from the OEC to the D1-E65/D2-E312 region, similar to what we observed at 500 μs , is reformed at this time point. The D1-E65/D2-E312 distance elongates from roughly 2.6 to roughly 3.2 \AA , which points to a substantial weakening of the interaction, likely forming a configuration that can accept the next proton. This change coincides with the time at which the O_x electron density decreases below the detection level in the omit map (Fig. 2), and the W28, W27 and W32 densities decrease in the O1 channel (Fig. 4a). We interpret this series of changes to be related to the onset of the recovery process of the Mn_4CaO_5 cluster (that is, initiated by the insertion of water into the OEC along with a deprotonation). At 2,000 and 4,000 μs , the side chain of D1-E65 is rotated almost 40° from its position in the S_3 structure toward W119 (Fig. 4b). The distance D1-E65–W119 is around 2.5 \AA , suggesting a shared proton or very tight interaction between these two groups. D1-R334 also moves by 20° , forming a hydrogen-bonding interaction with W41.

The changes around the D1-E65/D2-E312 region are indicative of the proton release to the bulk since they are reminiscent of what we observed for proton release during the $S_2 \rightarrow S_3$ transition¹². In the $S_3 \rightarrow S_0$ transition, we hypothesize that the earlier changes of this region (250–730 μs) are related to the first proton transfer and that the later changes (1,200–4,000 μs) are related to the second proton transfer

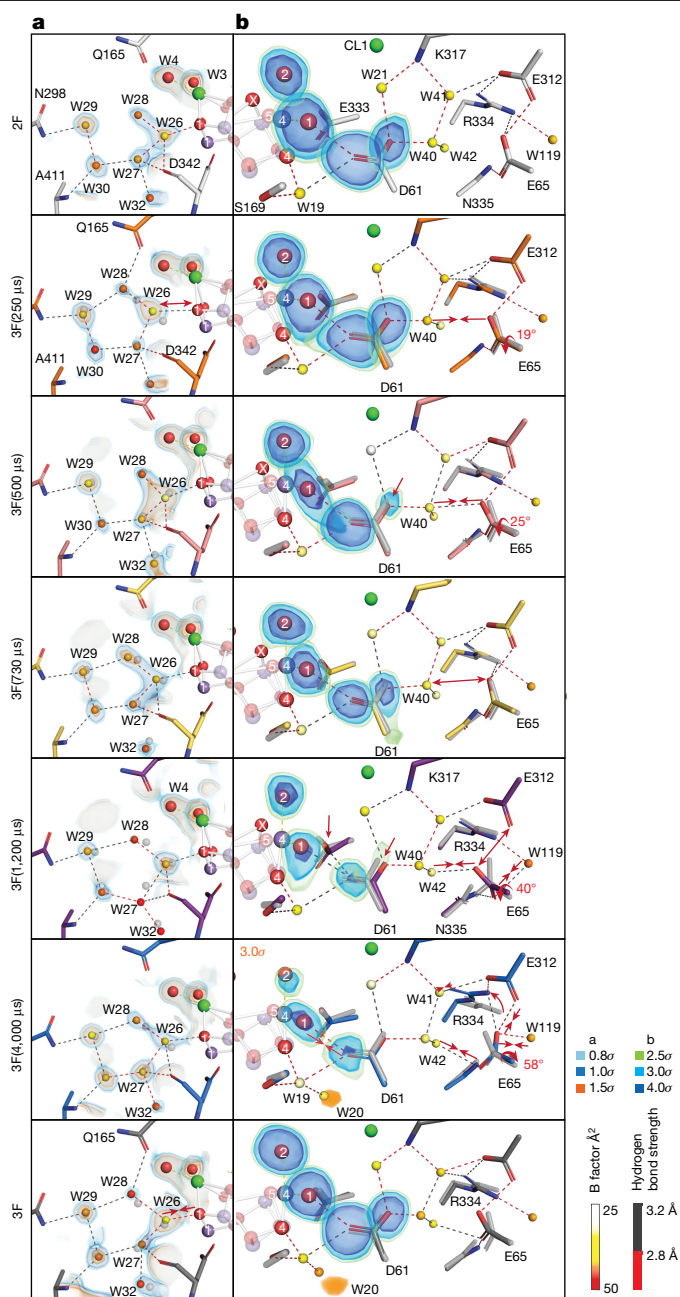


Fig. 4 | Structural changes in select regions of the water and proton channels of PS II during the $S_3 \rightarrow S_0$ transition. a, The terminus of the O1 channel near the OEC that includes the group of five waters (W26–W30) in this region. Overlaid is the $2mF_{obs} - DF_{calc}$ electron density map contoured at 0.8σ , 1.0σ and 1.5σ .

b, The O4 and ClI channels (branch A) that include the D1-D61 and D1-E65/D2-E312 region that is suggested to function as a proton gate. Overlaid are $mF_{obs} - DF_{calc}$ omit maps for W1, W2 and D61 shown at 2.5σ , 3.0σ and 4.0σ . Also shown is the $F_{obs}(\text{time point}) - F_{obs}(2F)$ difference density map within a 1.5-\AA radius of W19/W20/W48 in the O4 channel at 3σ (orange map). The observed rotation angle at the side chain of E65 at particular time points is calculated with respect to the corresponding side chain position at the 2F state. Major changes are highlighted with an arrow or dashed circle. All waters are coloured by their B factors according to the diverging colour scheme shown in the figure. Important hydrogen-bond interactions are shown with a binary colour scheme to indicate strength (distance $< 2.8 \text{ \AA}$ is red and $2.8\text{--}3.2 \text{ \AA}$ is gray).

from the OEC to the bulk, through D1-D61^{36,37} via the rotation of D1-E65. Thus, the current result suggests that the D1-E65/D2-E312 region functions as a gate for proton release twice during the $S_3 \rightarrow S_0$ transition.

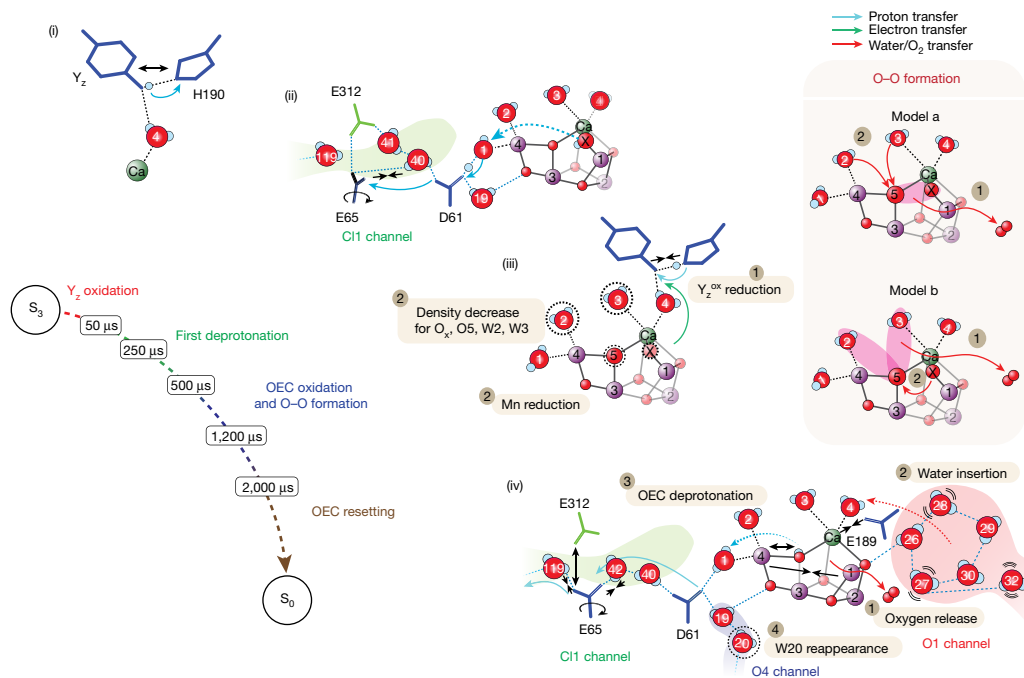


Fig. 5 | Schematic of the $S_3 \rightarrow S_0$ transition and proposed mechanism for O–O formation. The sequence of events (i–iv) leading to the first deprotonation event, the molecular oxygen release, the water insertion and the second deprotonation event. The OEC atoms are shown in purple (Mn), green (Ca) and red (O). The O1 channel is shown in red, the O4 channel is in blue and the Cl1 channel in green. The ligands of the OEC and the residues forming the water–proton channels are coloured based on the subunit they belong to

W20, which forms a tight hydrogen bonding network with O4 via W19 in the O4 channel, disappears during the $S_1 \rightarrow S_2$ transition and reappears in the S_0 state⁴. Its return during the $S_3 \rightarrow S_0$ transition is therefore an indicator for the full recovery of the S_0 state. The first clear indication of the return of W20 is found in the $F_{\text{obs}}(4,000 \mu\text{s}) - F_{\text{obs}}(2\text{f})$ difference map (orange density in Fig. 4b), and it was modeled at 40% occupancy in the 4,000- μs refined component. This implies that the W20 restoration happens in the later stage of the OEC recovery.

Sequence of events during $S_3 \rightarrow S_0$

The snapshots of the structures of PS II during the $S_3 \rightarrow S_0$ transition show the sequence and the progression of each of the events at multiple locations with different time constants. The structural changes can be broadly grouped into four sections with different onset times and kinetics, as shown in Fig. 5; they are the redox state changes of Y_z , first deprotonation, OEC oxidation and O₂ formation and the complete recovery and resetting of the Kok clock in the S_0 state.

In the S_3 state, all four Mn are formally in the (+IV) oxidation state. We note that oxidation of ligand instead of Mn (formation of oxyl or an oxo–oxyl bond) during the $S_2 \rightarrow S_3$ transition has been suggested in the literature⁵, but this is not in line with the room temperature X-ray emission spectroscopy (XES) data, which show the oxidation of Mn^{6,13}. Upon the third flash, the oxidation of Y_z occurs by donation of an electron to P680⁺ after the charge separation at the reaction center chlorophylls. A distance increase between Y_z and D1-H190 is observed in comparison with the S_3 state in 3F (50 μs) (Figs. 3 and 5(i)). This change is assigned to the Y_z^{ox} formation, which is known to occur within 30 μs after photoexcitation³⁸, and related proton translocation between Y_z^{ox} and D1-H190.

The Y_z^{ox} formation triggers the first deprotonation event likely during the time period of 200–500 μs ^{15–17,24,34,39}. We observed the start of

the rotation of D1-E65, which is proposed to be part of the proton gate, and the formation of the hydrogen bond pathway from the OEC to this region at 250 μs . Along with these changes, the electron densities at W1 and D61 become more prominent at 500 μs (Figs. 2d and 4b), potentially related to a proton release from the OEC to the proton gate residues D1-E65 and D2-E312 (Fig. 5(ii)).

In the early stage of the 500- to 1,200- μs period, the last oxidation event (transient S_4 state formation, with Mn(IV)₄O^{*} or Mn(IV)₃(V)) occurs, and subsequently, the reduction of Mn takes place (Fig. 5(iii)). The O–O bond formation should be triggered by this final oxidation event of the OEC to the potentially short-lived S_4 state. The change in distance we observe for Y_z –D1-H190 between 500 and 730 μs suggests that the reduction of Y_z^{ox} takes place during this time, through the electron transfer from the OEC to Y_z . After the transient formation of S_4 , the four-electron reduction may proceed in one step with the O–O bond formation and immediate release of O₂ or in two steps with the presence of an intermediate before the release of molecular oxygen from the OEC. In the latter case, a peroxo species formed by an initial two-electron reduction appears most likely as an intermediate.

Our data show that there is a delay between the onset of O–O bond formation (500–730 μs) as indicated by the Y_z –D1-H190 distance/rotation and the decrease of the O_x electron density and the onset of the O₂ release supported by the Mn1–Mn4 distance contraction (1,200 μs). This onset time for O₂ release is also in line with studies of O₂ evolution^{22,40}. The delay indicates that there is an intermediate state, possibly a peroxide-like species, pointing toward the two-step electron reduction mechanism.

Several O–O bond formation sites have been proposed in the literature based on theoretical studies (Fig. 5)^{3,41–49}. Among these, O5–O_x best account for our data because of their proximity and the reduced occupancy of O5 around 1,200 μs (Fig. 5, model a). However, two other possibilities in which O5 reacts with either W2 or W3 and O_x replaces O5

cannot be excluded at this time (Fig. 5, model b). While other mechanisms that do not involve O5 cannot be ruled out, there is no clear evidence to support those in the current data.

At 1,200 μs , the O_x omit map density is below the 2.5σ threshold, indicating that a predominant fraction of O_x has shifted from its original positioning in the cluster. The contraction of the Mn1–Mn4 and Mn1–Mn3 distances starting at this time point suggests that the onset of O_2 release happens around this time. Once O_2 is released, refilling of the cluster with a new substrate water seems to occur immediately. This is based on the observation that there is no missing oxygen density besides O_x , although the omit map densities of all the terminal waters (W1–W4) and bridging O5 are weakened at 1,200 μs . The data support that the O_2 release and refilling of the site are highly coordinated and occur likely via a terminal water already ligated to the OEC.

At both 2,000 and 4,000 μs (Fig. 5(iv)), O_x density is within the noise level, which implies that Mn1 becomes predominantly five coordinate. Mn4 is six coordinate, although the Mn4–O5 interaction is weak (2.2–2.3 Å), suggesting that O5 may be a hydroxide. The Ca–D1–E189 distance is still more elongated at these time points than in the S_0 state. Other slow recoveries are observed in the Y_z region (Fig. 2b), the ‘water wheel’ (Fig. 4a) and the proton gate regions (D1–E65/D2–E312) (Fig. 4b). We hypothesize that a water from the ‘water wheel’ region in the O1 channel, similar to the $\text{S}_2 \rightarrow \text{S}_3$ transition^{6,12}, replaces the terminal water ligand of the OEC. The changes in the proton gate region might indicate the deprotonation of the newly inserted water³⁶. Concomitantly, the amino acid coordination environment, the hydrogen bonding network around the OEC and the waters in the channels reset to the S_0 state. This includes recovery of W20 in the O4 channel, which is proposed to be involved in proton release during the $\text{S}_0 \rightarrow \text{S}_1$ transition^{4,50}.

In the current study, room temperature snapshots of PS II structures through the final step of Kok’s clock ($\text{S}_3 \rightarrow [\text{S}_4] \rightarrow \text{S}_0$) reveal details of the molecular processes for photosynthetic water oxidation. Until now, these processes were interpreted largely based on kinetic studies. Importantly, the results reported here provide experimental support for a two-step reduction mechanism of the $\text{Mn}_4\text{CaO}_5\text{-O}_x$ cluster upon the O–O bond formation and O_2 release with a transient intermediate, most likely a bound peroxide. This is a major step forward toward understanding the chemistry of the water oxidation reaction. The results also show how biological catalysts, such as the OEC in PS II, enable multielectron/multiproton reactions through the interplay between the metal center, the protein environment and the water network. The active role of the microenvironment in natural enzymes provides inspiration for how to control such reactions in artificial photosynthetic systems that can be made from earth-abundant elements.

Online content

Any methods, additional references, Nature Portfolio reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at <https://doi.org/10.1038/s41586-023-06038-z>.

- Kok, B., Forbush, B. & McGloin, M. Cooperation of charges in photosynthetic O_2 evolution. 1. A linear four-step mechanism. *Photochem. Photobiol.* **11**, 457–475 (1970).
- Yano, J. & Yachandra, V. K. Mn_4Ca cluster in photosynthesis: where and how water is oxidized to dioxygen. *Chem. Rev.* **114**, 4175–4205 (2014).
- Cox, N., Pantazis, D. A., Neese, F. & Lubitz, W. Biological water oxidation. *Acc. Chem. Res.* **46**, 1588–1596 (2013).
- Kern, J. et al. Structures of the intermediates of Kok’s photosynthetic water oxidation clock. *Nature* **563**, 421–425 (2018).
- Suga, M. et al. An oxyl/oxo mechanism for oxygen-oxygen coupling in PSII revealed by an x-ray free-electron laser. *Science* **366**, 334–338 (2019).
- Ibrahim, M. et al. Untangling the sequence of events during the $\text{S}_2 \rightarrow \text{S}_3$ transition in photosystem II and implications for the water oxidation mechanism. *Proc. Natl Acad. Sci. USA* **117**, 12624–12635 (2020).
- Brändén, G. & Neutze, R. Advances and challenges in time-resolved macromolecular crystallography. *Science* **373**, eaba0954 (2021).
- Kern, J. et al. Simultaneous femtosecond X-ray spectroscopy and diffraction of photosystem II at room temperature. *Science* **340**, 491–495 (2013).
- Young, I. D. et al. Structure of photosystem II and substrate binding at room temperature. *Nature* **540**, 453–457 (2016).
- Suga, M. et al. Light-induced structural changes and the site of O=O bond formation in PSII caught by XFEL. *Nature* **543**, 131–135 (2017).
- Umeha, Y., Kawakami, K., Shen, J.-R. & Kamiya, N. Crystal structure of oxygen-evolving photosystem II at a resolution of 1.9 Å. *Nature* **473**, 55–60 (2011).
- Hussein, R. et al. Structural dynamics in the water and proton channels of photosystem II during the S_2 to S_3 transition. *Nat. Commun.* **12**, 6531 (2021).
- Fransson, T. et al. Effects of x-ray free-electron laser pulse intensity on the Mn $\text{K}\beta_{1,3}$ x-ray emission spectrum in photosystem II—a case study for metalloprotein crystals and solutions. *Struct. Dyn.* **8**, 064302 (2021).
- Babcock, G. T., Blankenship, R. E. & Sauer, K. Reaction-kinetics for positive charge accumulation on water side of chloroplast photosystem II. *FEBS Lett.* **61**, 286–289 (1976).
- Rappaport, F., Blanchard-Desce, M. & Lavergne, J. Kinetics of electron transfer and electrochromic change during the redox transitions of the photosynthetic oxygen-evolving complex. *Biochim. Biophys. Acta Bioenerg.* **1184**, 178–192 (1994).
- Razeghifard, M. R. & Pace, R. J. EPR kinetic studies of oxygen release in thylakoids and PSII membranes: a kinetic intermediate in the S_3 to S_0 transition. *Biochemistry* **38**, 1252–1257 (1999).
- Gerencsér, L. & Dau, H. Water oxidation by photosystem II: $\text{H}_2\text{O-D}_2\text{O}$ exchange and the influence of pH support formation of an intermediate by removal of a proton before dioxygen creation. *Biochemistry* **49**, 10098–10106 (2010).
- Dilbeck, P. L. et al. The D1-D61N mutation in *Synechocystis* sp. PCC 6803 allows the observation of pH-sensitive intermediates in the formation and release of O_2 from photosystem II. *Biochemistry* **51**, 1079–1091 (2012).
- Noguchi, T. Fourier transform infrared difference and time-resolved infrared detection of the electron and proton transfer dynamics in photosynthetic water oxidation. *Biochim. Biophys. Acta Bioenerg.* **1847**, 35–45 (2015).
- Han, G., Chernev, P., Styring, S., Messinger, J. & Mamedov, F. Molecular basis for turnover inefficiencies (misses) during water oxidation in photosystem II. *Chem. Sci.* **13**, 8667–8678 (2022).
- Haumann, M. et al. Photosynthetic O_2 formation tracked by time-resolved x-ray experiments. *Science* **310**, 1019–1021 (2005).
- Bao, H. & Burnap, R. L. Structural rearrangements preceding dioxygen formation by the water oxidation complex of photosystem II. *Proc. Natl Acad. Sci. USA* **112**, E6139–E6147 (2015).
- Wöhri, A. B. et al. Light-induced structural changes in a photosynthetic reaction center caught by Laue diffraction. *Science* **328**, 630–633 (2010).
- Noguchi, T., Suzuki, H., Tsuno, M., Sugiura, M. & Kato, C. Time-resolved infrared detection of the proton and protein dynamics during photosynthetic oxygen evolution. *Biochemistry* **51**, 3205–3214 (2012).
- Suzuki, H., Sugiura, M. & Noguchi, T. Monitoring proton release during photosynthetic water oxidation in photosystem II by means of isotope-edited infrared spectroscopy. *J. Am. Chem. Soc.* **131**, 7849–7857 (2009).
- Tso, J., Sivaraja, M. & Dismukes, G. C. Calcium limits substrate accessibility or reactivity at the manganese cluster in photosynthetic water oxidation. *Biochemistry* **30**, 4734–4739 (1991).
- Kim, C. J. & Debus, R. J. One of the substrate waters for O_2 formation in photosystem II is provided by the water-splitting Mn_4CaO_5 cluster’s Ca^{2+} ion. *Biochemistry* **58**, 3185–3192 (2019).
- Murray, J. W. & Barber, J. Structural characteristics of channels and pathways in photosystem II including the identification of an oxygen channel. *J. Struct. Biol.* **159**, 228–237 (2007).
- Ho, F. M. & Styring, S. Access channels and methanol binding site to the CaMn_4 cluster in Photosystem II based on solvent accessibility simulations, with implications for substrate water access. *Biochim. Biophys. Acta* **1777**, 140–153 (2008).
- Gabdulkhakov, A. et al. Probing the accessibility of the Mn_4Ca cluster in photosystem II: channels calculation, noble gas derivatization, and cocrystallization with DMSO. *Structure* **17**, 1223–1234 (2009).
- Vassiliev, S., Zaraiskaya, T. & Bruce, D. Exploring the energetics of water permeation in photosystem II by multiple steered molecular dynamics simulations. *Biochim. Biophys. Acta Bioenerg.* **1817**, 1671–1678 (2012).
- Ishikita, H., Saenger, W., Loll, B., Biesiadka, J. & Knapp, E. W. Energetics of a possible proton exit pathway for water oxidation in photosystem II. *Biochemistry* **45**, 2063–2071 (2006).
- Service, R. J., Hillier, W. & Debus, R. J. Evidence from FTIR difference spectroscopy of an extensive network of hydrogen bonds near the oxygen-evolving Mn_4CaO_5 cluster of photosystem II involving D1-Glu65, D2-Glu312, and D1-Glu329. *Biochemistry* **49**, 6655–6669 (2010).
- Klauss, A., Haumann, M. & Dau, H. Alternating electron and proton transfer steps in photosynthetic water oxidation. *Proc. Natl Acad. Sci. USA* **109**, 16035–16040 (2012).
- Allgöwer, F., Gamiz-Hernandez, A. P., Rutherford, A. W. & Kaila, V. R. I. Molecular principles of redox-coupled protonation dynamics in photosystem II. *J. Am. Chem. Soc.* **144**, 7171–7180 (2022).
- Capone, M., Narzi, D. & Guidoni, L. Mechanism of oxygen evolution and Mn_4CaO_5 cluster restoration in the natural water-oxidizing catalyst. *Biochemistry* **60**, 2341–2348 (2021).
- Debus, R. J. Evidence from FTIR difference spectroscopy that D1-Asp61 influences the water reactions of the oxygen-evolving Mn_4CaO_5 cluster of photosystem II. *Biochemistry* **53**, 2941–2955 (2014).
- Renger, G. Mechanism of light induced water splitting in photosystem II of oxygen evolving photosynthetic organisms. *Biochim. Biophys. Acta Bioenerg.* **1817**, 1164–1176 (2012).

39. Ishida, N. et al. Biosynthetic exchange of bromide for chloride and strontium for calcium in the photosystem II oxygen-evolving enzymes. *J. Biol. Chem.* **283**, 13330–13340 (2008).
40. Assuncao, R., Zaharieva, I. & Dau, H. Ammonia as a substrate-water analogue in photosynthetic water oxidation: influence on activation barrier of the O₂-formation step. *Biochim. Biophys. Acta Bioenerg.* **1860**, 533–540 (2019).
41. Siegbahn, P. E. Structures and energetics for O₂ formation in photosystem II. *Acc. Chem. Res.* **42**, 1871–1880 (2009).
42. Sproviero, E. M., Gascon, J. A., McEvoy, J. P., Brudvig, G. W. & Batista, V. S. Quantum mechanics/molecular mechanics study of the catalytic cycle of water splitting in photosystem II. *J. Am. Chem. Soc.* **130**, 3428–3442 (2008).
43. Rummel, F. & O'Malley, P. J. How nature makes O₂: an electronic level mechanism for water oxidation in photosynthesis. *J. Phys. Chem. B* **126**, 8214–8221 (2022).
44. Guo, Y., Zhang, B., Kloo, L. & Sun, L. Necessity of structural rearrangements for O-O bond formation between O5 and W2 in photosystem II. *J. Energy Chem.* **57**, 436–442 (2021).
45. Messinger, J. Evaluation of different mechanistic proposals for water oxidation in photosynthesis on the basis of Mn₄O₄Ca structures for the catalytic site and spectroscopic data. *Phys. Chem. Chem. Phys.* **6**, 4764–4771 (2004).
46. Nilsson, H., Cournac, L., Rappaport, F., Messinger, J. & Lavergne, J. Estimation of the driving force for dioxygen formation in photosynthesis. *Biochim. Biophys. Acta* **1857**, 23–33 (2016).
47. Rapatskiy, L. et al. Detection of the water-binding sites of the oxygen-evolving complex of Photosystem II using W-band ¹⁷O electron-electron double resonance-detected NMR spectroscopy. *J. Am. Chem. Soc.* **134**, 16619–16634 (2012).
48. Davis, K. M. et al. Rapid evolution of the photosystem II electronic structure during water splitting. *Phys. Rev. X* **8**, 041014 (2018).
49. Yamaguchi, K. et al. Geometric, electronic and spin structures of the CaMn₄O₅ catalyst for water oxidation in oxygen-evolving photosystem II. Interplay between experiments and theoretical computations. *Coord. Chem. Rev.* **471**, 214742 (2022).
50. Sakashita, N., Ishikita, H. & Saito, K. Rigidly hydrogen-bonded water molecules facilitate proton transfer in photosystem II. *Phys. Chem. Chem. Phys.* **22**, 15831–15841 (2020).

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Methods

Sample preparation

The X-ray diffraction measurements of 20- to 60- μm crystals prepared from PS II dimers of *Thermosynechococcus vestitus* (previously named *Thermosynechococcus elongatus*) were performed in 100 mM 2-(*N*-morpholino)ethanesulfonic acid, pH 6.5, 100 mM ammonium chloride and 35% (wt/vol) PEG 5000 (refs. 51,52). PS II crystal suspension, at about 0.5–1.2 mM chlorophyll (Chl) concentration, was loaded into a syringe (Hamilton gastight syringe, 1 ml) and dark adapted for 1 h before data collection. Membrane inlet mass spectroscopy (MIMS) was used to determine the O_2 evolution, turnover parameters and S-state populations^{4,6}. The PS II crystals showed no Mn (II) contamination based on XES and electron paramagnetic resonance measurements⁵³ and exhibited an activity of $2500 \pm 100 \mu\text{mol O}_2 (\text{mg(Chl)} \times \text{h})^{-1}$.

Sample injection and illumination

Acoustic droplet ejection⁵⁴ was used in combination with the Drop-on-Tape sample delivery method⁵⁵. For capturing the stable intermediates S_2 , S_3 and S_0 , each droplet of the crystal suspension was illuminated by 120-ns laser pulses at 527 nm using an Nd:YLF (yttrium lithium fluoride) laser (Evolution, Coherent) at Linac Coherent Light Source (LCLS) or by 8-ns laser pulses at 532 nm using a combination of two Nd:YAG (yttrium aluminium garnet) lasers (Minilite, Continuum) at Spring-8 Angstrom Compact free electron Laser (SACLA) via three fiber-coupled outputs with a delay time of 200 ms between each illumination and of 200 ms between the last illumination and the X-ray probe, similar to what was used previously to accommodate the acceptor quinone Q_A and Q_B kinetics and efficiently drive S-state transitions^{4,6,55}. We implemented a feedback control system of the belt speed and deposition delay, and the flashing delay and droplet phase were adjusted accordingly⁵⁵. To achieve time delays shorter than 200 ms between illumination and the X-ray probe, a fourth ‘free space’ laser was utilized. This was either an Opolette 355 LD laser (Opotek, 530-nm wavelength, 7-ns pulse width) at the macromolecular femtosecond crystallography/LCLS instrument or an NT230 OPO laser system (530-nm wavelength, 5-ns pulse width, EKSPILA Co.) at SACLA. This free space laser was triggered to be synchronized with the X-ray pulse with an adjustable delay that was set between 50 and 4,000 μs for this study. The laser was guided with optics to the X-ray interaction spot, and its position was fine-tuned for each delay time to ensure that the laser spot position coincides with the position of the sample droplet at the selected delay timing. At the XFELs, a light intensity of $120 \pm 10 \text{ mJ per cm}^2$ was applied as O_2 evolution was found to be saturated at 70 mJ per cm^2 for the dimensions and concentrations of samples used in our experiments⁴. A light intensity of 120 mJ per cm^2 corresponds to about 140 photons absorbed per PS II monomer in the front 5- μm layer of the crystal and approximately 9 photons per PS II monomer when assuming a 60- μm thickness of the crystal (which is the upper size limit of the crystals used in this study) for the back 5- μm layer. This photon density ensures saturation over the entire crystal volume, even in the case of two crystals stacked on top of each other in the laser beam. Given a minimum pulse length of 5 ns and 35 Chl per PS II monomer, the light intensity used averages to 0.8 photons per (Chl and nanosecond) for the front and 0.05 photons per (Chl and nanosecond) for the back part of the crystal. If a PS II center is undergoing charge separation, additional photons absorbed by the internal antenna Chl are rapidly dissipated in the form of fluorescence with an average fluorescence lifetime of around 0.5–1 ns, hence preventing any overexcitation of the reaction center or causing any heating artefacts.

X-ray data collection

The crystallography data were collected at various facilities, and details are listed in Supplementary Table 2. The experimental beam conditions and detector configurations used to collect each dataset are also

tabulated. The sample was delivered into the X-ray interaction region using the previously described Drop-on-Tape setup⁵⁵. Illumination conditions for populating different S states are detailed in ref. 4.

X-ray diffraction data processing

The data collected for the different illumination states were processed using the program `dials.stills_process` with a target unit cell of $a = 117.0 \text{ \AA}$, $b = 221.0 \text{ \AA}$, $c = 309.0 \text{ \AA}$, $\alpha = \beta = \gamma = 90^\circ$ and the space group $\text{P}2_12_12_1$. Bragg spots were integrated to the edge of the detector. A Kapton absorption correction due to the conveyor belt of our sample delivery system was applied to each integrated Bragg spot, taking into account the droplet size, tape thickness, tape angle and the position of the diffraction spots on the detector with respect to the crystal position. Before integration, we also performed ensemble refinement of the crystal and detector parameters using the program `cctbx.xfel.stripe_experiment`, which has been shown to narrow the unit cell distribution and improve the final isomorphous difference maps⁵⁶. Finally, the intensities were merged using the program `cctbx.xfel.merge`, which applies a per-image resolution cutoff and filtering of the lattices using a unit cell threshold of 1% from the reference model. To merge the reflections, we use the best practices described in ref. 57. The unit cells and number of lattices merged for each dataset are tabulated in Extended Data Tables 1 and 2.

Final merged datasets were acquired for the 2F, 3F(50 μs), 3F(250 μs), 3F(500 μs), 3F(730 μs), 3F(1,200 μs), 3F(2,000 μs), 3F(4,000 μs) and 3F(200 ms) states to resolutions between 2.16 and 2.0 \AA , obtained by merging between 6,659 and 39,199 lattices (Extended Data Tables 1 and 2). The final merged datasets before model building were also scaled on a per-resolution bin basis to a reference dataset (in this case, the reference dataset is the PS II dataset published in PDB ID code 7RF1)¹². This allows us to conduct a more accurate comparison of $mF_{\text{obs}} - DF_{\text{calc}}$ omit maps and $2mF_{\text{obs}} - DF_{\text{calc}}$ maps between different datasets.

Model building and map calculation

Each dataset was refined using a high-resolution PS II structure (1.89 \AA) that was published in a previous work (PDB ID code 7RF1)¹² as the starting point using the program `phenix.refine`⁵⁸. The refinement is done in several stages. First, the B factors of the starting model are set to 30, and all waters and the atoms of the OEC are removed. An initial rigid body refinement coupled with refinement of `xyz` coordinates and isotropic B factors was done for 15 cycles to adjust the model into the unit cell. Next, the OEC atoms are added back and refined with custom bonding restraints for several cycles. We also use custom bonding restraints for chlorophyll-*a* (to allow correct placement of the Mg relative to the plane of the porphyrin ring) and unknown lipid-like ligands (steric acid) in the refinement. After initial refinement of the OEC + protein complex, waters were added to the model using the `phenix.refine` water picking protocol as well as manual placement of waters via `coot`⁵⁹ and doing multiple cycles of refinement.

At this stage, we split the model in the vicinity of the OEC and the OEC itself (only protein and OEC atoms) into multiple components (Extended Data Table 3). The split was done only in parts of chains A/a, C/c and D/d. The rationale and population of the components in each time point used are described in the section Estimating population distribution in each time point. In each dataset, the primary conformer (defined as the intermediate that is advancing from S_3 to S_0) is refined using a strategy of reciprocal `xyz` + isotropic B-factor refinement. For the secondary/tertiary components (whose structures are known as they are either in the S_3 or S_0 state), only the group B factors are adjusted (group `adp` strategy in `phenix.refine`) to adjust them to the resolution of the dataset. For the remaining part of the model that is not split, regular reciprocal `xyz` refinement and isotropic B-factor refinement are performed in tandem for multiple cycles. All waters (except for the terminal waters ligated to the OEC; that is, W1–W4) were refined as a single component.

The refinement of the OEC in the primary component of the multi-component model was done using custom restraints that were used

to model the S_3 state. However, for all the time points, we used slightly looser estimated s.d. values for the restraints (0.1 Å for bonds, 10° for angles) to allow the OEC atoms during refinement to move toward where the electron density is optimally modeled and reduce strain in the refinement while at the same time maintaining the overall shape of the cluster. The OEC in the S_0 state was modeled with restraints used for our previously published S_0 state structure. The restraints used to model the OEC atoms in the time points have been provided as text files (schemes 1–3 in Supplementary Data).

Estimating population distribution in each time point

The S-state population distribution in the $S_3 \rightarrow S_0$ transition is a heterogeneous distribution consisting of (1) centers that are advancing from the S_3 to S_0 state, (2) centers that are lagging behind by one transition and hence, are advancing from the S_2 to S_3 state and (3) centers that have transitioned over to the S_0 state. While the majority of centers are in category (1), due to the intrinsic inefficiencies ('misses') of the Kok cycle in PS II²⁰, a certain fraction of centers is in category (2). In addition, after a certain time in the $S_3 \rightarrow S_0$ transition, a substantial number of centers will have formed the stable S_0 state (category (3)).

Given this context, it is important to account for this population heterogeneity in our structural modeling to obtain accurate electron density maps and models. We do this by splitting up our structural model near the active site region (including the OEC) into multiple components. The primary component in each dataset is category (1), which is the intermediate transitioning from S_3 to S_0 . The nature of the secondary and tertiary components depends on the dataset under consideration. In each dataset, the coordinates/isotropic B factors of only the primary component are refined (category (1)). The secondary and tertiary component structures are modeled from known or previously deposited structures and only adjusted for resolution using a group B-factor refinement. The identity of the secondary (and tertiary if used) component depends on which time point is being processed. For example, in the 3F(50 μ s) dataset, a two-component model is constructed with the 2F(50 μ s) model coordinates/B factors being used for the secondary component. In the 3F(1,200 μ s) data, we use a three-component model with the secondary and tertiary components being the S_0 and S_3 states. The populations for each of the components in the various time points are given in Extended Data Table 3. We used numbers available in the literature to perform a kinetic analysis yielding an estimate of the population distribution. Since populations below 10% are in the noise level for structural refinement, we adjusted our populations to avoid any conformer with such low populations.

The population distribution in each of the metastable S states has been previously determined using the MIMS technique. In our work, the starting 2F state, which is generated by illuminating with two visible lasers with a flash interval of 200 ms, consists of approximately 65% S_3 state and 35% S_2 state based on studies conducted on crystals. With the third visible flash, the $S_3 \rightarrow S_0$ transition is initiated. The reader is referred to the extended data in ref. 4 for more details on how the S-state populations for each of the flash states were estimated, accounting for miss parameters calculated from XES and MIMS data, and crossillumination (this was negligible at the speed of the tape and deposition frequency of the acoustic droplet ejection that was used in the present study). All the results described in this paper are from monomer I (chains annotated as uppercase in the published structures). Similar trends are observed for monomer II (chains annotated as lowercase).

Estimating the effect of population on the Mn1–Mn4 distance

The Mn1–Mn4 distance stays elongated until 3F(1,200 μ s) in the intermediate undergoing the $S_3 \rightarrow S_0$ transition, after which a decrease is seen in the next 3 ms. We tested the robustness of the elongated distance at 3F(1,200 μ s) by constructing an alternative hypothesis to explain this observation, postulating that it could be due to two separate populations in the primary component: (1) increased Mn1–Mn4 distance

due to misses that form additional S_3 or (2) decreased Mn1–Mn4 distance (with/without disappearance of O_x) that is a property of the intermediate undergoing the $S_3 \rightarrow S_0$ transition. We modeled this scenario by increasing the S_3 population from 35 to 55% and decreasing the primary component (with/without O_x) from 40 to 20% in the 3F(1,200 μ s) time point. The resulting refinement gave Mn1–Mn4 distances of 5.14 Å (with O_x) and 5.09 Å (without O_x). Both numbers are similar to the distance given in Fig. 3 and within the measurement error. The tests thus show no contraction compared with the S_3 state and allow us to reject the hypothesis. We reiterate that the S_3 population estimate of 35% in the 3F(1,200 μ s) time point is well established using multiple independent experiments as detailed in the previous section and past publications.

Estimated positional precision

To estimate the positional precision of the OEC atoms and the surrounding amino acids for each time point, we used the END/RAPID procedure⁶⁰, similar to what was previously employed⁶. Briefly, in this method, we perturb the structure factors by a random amount in between $\pm(mF_{\text{obs}} - DF_{\text{calc}})$. The atomic coordinates of the final model for that time point are also perturbed by a small amount to allow the model to explore greater phase space (only the primary conformer is perturbed). Subsequently, 100 such synthetic datasets are generated for each time point, and they are then each refined separately. From the ensemble of these refined datasets, we can estimate the error associated with the distance metric of interest. The obtained errors should be considered an upper bound as the introduced perturbations in the structure factors are an overestimate of the true errors in the experiment.

Implementation details can be found at <https://bl831.als.lbl.gov/END/RAPID/end.rapid/Documentation/end.rapid.Manual.htm>.

$mF_{\text{obs}} - DF_{\text{calc}}$ difference omit density

All $mF_{\text{obs}} - DF_{\text{calc}}$ omit maps shown in the manuscript were generated using the phenix.polder program and using the normal omit map coefficients from the output.mtz file (not polder map coefficients)⁶¹. For peak height calculation wherever stated, we used custom python scripts that average the $mF_{\text{obs}} - DF_{\text{calc}}$ omit map value about a 0.5-Å radius of the atom of interest.

Reporting summary

Further information on research design is available in the Nature Portfolio Reporting Summary linked to this article.

Data availability

The atomic coordinates and structure factors have been deposited in the Protein Data Bank, www.pdb.org (PDB codes 8EZ5 for the 2F data; 8F4D for the 3F(50 μ s) data; 8F4E for the 3F(250 μ s) data; 8F4F for the 3F(500 μ s) data; 8F4G for the 3F(730 μ s) data; 8F4H for the 3F(1,200 μ s) data; 8F4I for the 3F(2,000 μ s) data; 8F4J for the 3F(4,000 μ s) data; 8F4K for the 3F(200 ms) data and 8F4C for the 2F-alternate data). The raw X-ray free electron laser data have been deposited in the Coherent X-Ray Imaging Database, www.cxidb.org (ID 215).

Code availability

The open source programs `dials.stills_process`, `cctbx.xfel GUI`, `cctbx.xfel.stripe_experiment`, `cctbx.xfel.merge` and legacy program `cx.merge` are distributed with DIALS packages available at <http://dials.github.io>. Instructions and details for using these programs can be found in refs. 56,57 with further documentation available at <http://cci.lbl.gov/xfel>. Figures shown in the paper were rendered using the PyMOL software v.2.5 (ref. 62). Custom code used for calculating peak height values is publicly available at <https://github.com/asmit3/eden> (ref. 63).

51. Hellmich, J. et al. Native-like Photosystem II superstructure at 2.44 Å resolution through detergent extraction from the protein crystal. *Structure* **22**, 1607–1615 (2014).
52. Ibrahim, M. et al. Improvements in serial femtosecond crystallography of photosystem II by optimizing crystal uniformity using microseeding procedures. *Struct. Dyn.* **2**, 041705 (2015).
53. Fransson, T. et al. X-ray emission spectroscopy as an in situ diagnostic tool for X-ray crystallography of metalloproteins using an X-ray free-electron laser. *Biochemistry* **57**, 4629–4637 (2018).
54. Roessler, C. G. et al. Acoustic injectors for drop-on-demand serial femtosecond crystallography. *Structure* **24**, 631–640 (2016).
55. Fuller, F. D. et al. Drop-on-demand sample delivery for studying biocatalysts in action at X-ray free-electron lasers. *Nat. Methods* **14**, 443–449 (2017).
56. Brewster, A. S. et al. Improving signal strength in serial crystallography with DIALS geometry refinement. *Acta Crystallogr. D Struct. Biol.* **74**, 877–894 (2018).
57. Brewster, A. S., Young, I. D., Lyubimov, A., Bhowmick, A. & Sauter, N. K. Processing serial crystallographic data from XFELs or synchrotrons using the cctbx.xfel GUI. *Comput. Crystallogr. Newsl.* **10**, 22–39 (2019).
58. Liebschner, D. et al. Macromolecular structure determination using X-rays, neutrons and electrons: recent developments in. *Phenix. Acta Crystallogr. D Struct. Biol.* **75**, 861–877 (2019).
59. Emsley, P., Lohkamp, B., Scott, W. G. & Cowtan, K. Features and development of Coot. *Acta Crystallogr. D Struct. Biol.* **66**, 486–501 (2010).
60. Lang, P. T., Holton, J. M., Fraser, J. S. & Alber, T. Protein structural ensembles are revealed by redefining X-ray electron density noise. *Proc. Natl Acad. Sci. USA* **111**, 237–242 (2014).
61. Liebschner, D. et al. Polder maps: improving OMIT maps by excluding bulk solvent. *Acta Crystallogr. D Struct. Biol.* **73**, 148–157 (2017).
62. Schrödinger, LLC. The PyMOL Molecular Graphics System, version 1.8 (2015).
63. Bhowmick A. asmit3/eden: release v.0.1.1. Zenodo <https://doi.org/10.5281/zenodo.773470> (2023).

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Author contributions R.A.-M., N.K.S., U.B., A.Z., J.M., J.K., J.Y. and V.K.Y. designed the experiment. R.H., M.I., R.C., M.D., M.Z., J.G., S.H., I.I.N., A.Z. and J.K. prepared samples. R.C., M.H.C., T.F., J.G., S.H., A.O.A. and F.M. characterized sample activity. K.T., S.O., L.B.G., F.D.F., A.Batyuk and R.A.-M. prepared and operated the X-ray free electron laser (XFEL) beam lines. I.B., P.S.S., P.C., I.-S.K., H.M., F.D.F. and J.K. developed, tested and ran the sample delivery system. A.Bhowmick, R.H., I.B., P.S.S., M.I., R.C., M.D.D., M.H.C., T.F., P.C., I.-S.K., H.M., M.D., C.J.K., M.Z., J.G., S.H., S.M.K., K.T., S.O., L.B.G., F.D.F., A.Batyuk, R.A.-M., D.W.P., A.S.B., N.K.S., U.B., A.Z., J.M., J.K., J.Y. and V.K.Y. performed the XFEL experiment. A.Bhowmick, J.M.H., D.W.P., N.W.M., P.D.A., A.S.B. and N.K.S. developed new software for data processing. A.Bhowmick, M.D., S.M.K., D.W.P., A.S.B. and N.K.S. processed XFEL data. A.Bhowmick, R.H., I.B., P.S.S., M.I., M.D.D., H.D., N.K.S., A.Z., J.M., J.K., J.Y. and V.K.Y. analysed and interpreted data. A.Bhowmick, R.H., I.B., P.S.S., M.D.D., A.Z., J.M., J.K., V.K.Y. and J.Y. wrote the manuscript with input from all authors.

Competing interests The authors declare no competing interests.

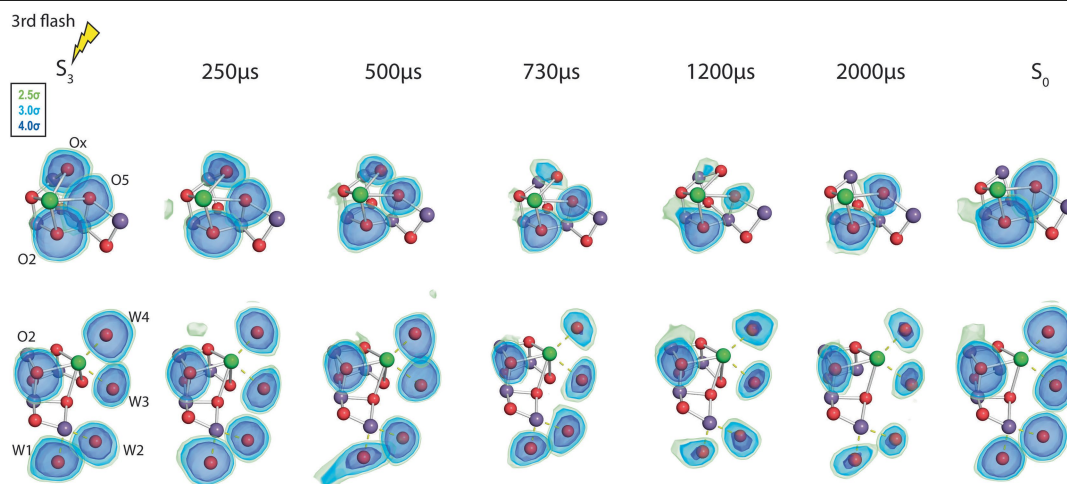
Additional information

Supplementary information The online version contains supplementary material available at <https://doi.org/10.1038/s41586-023-06038-z>.

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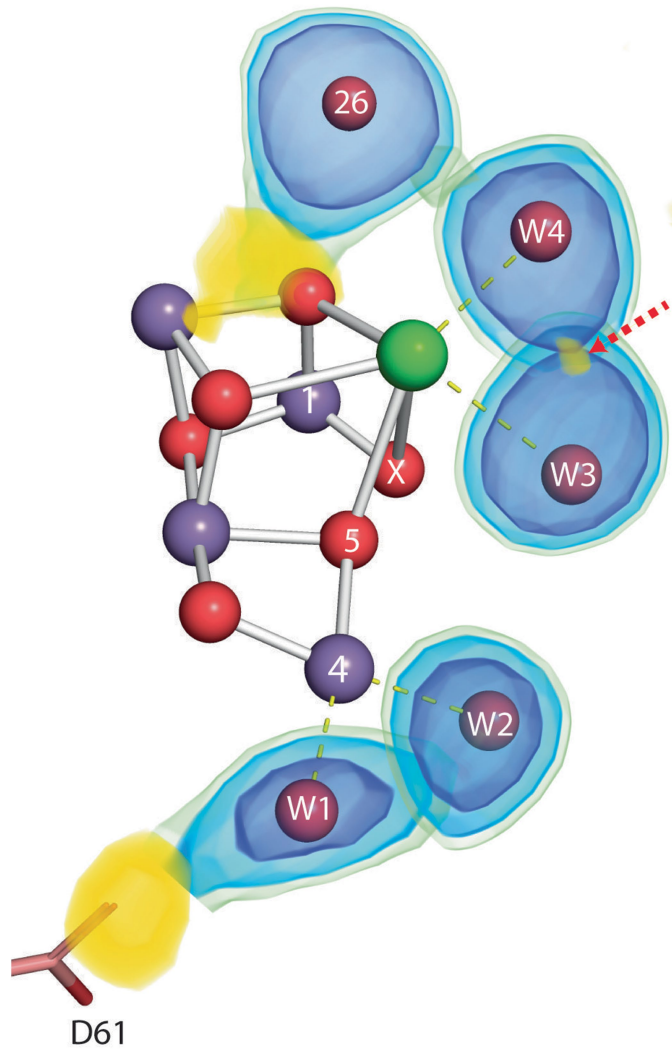
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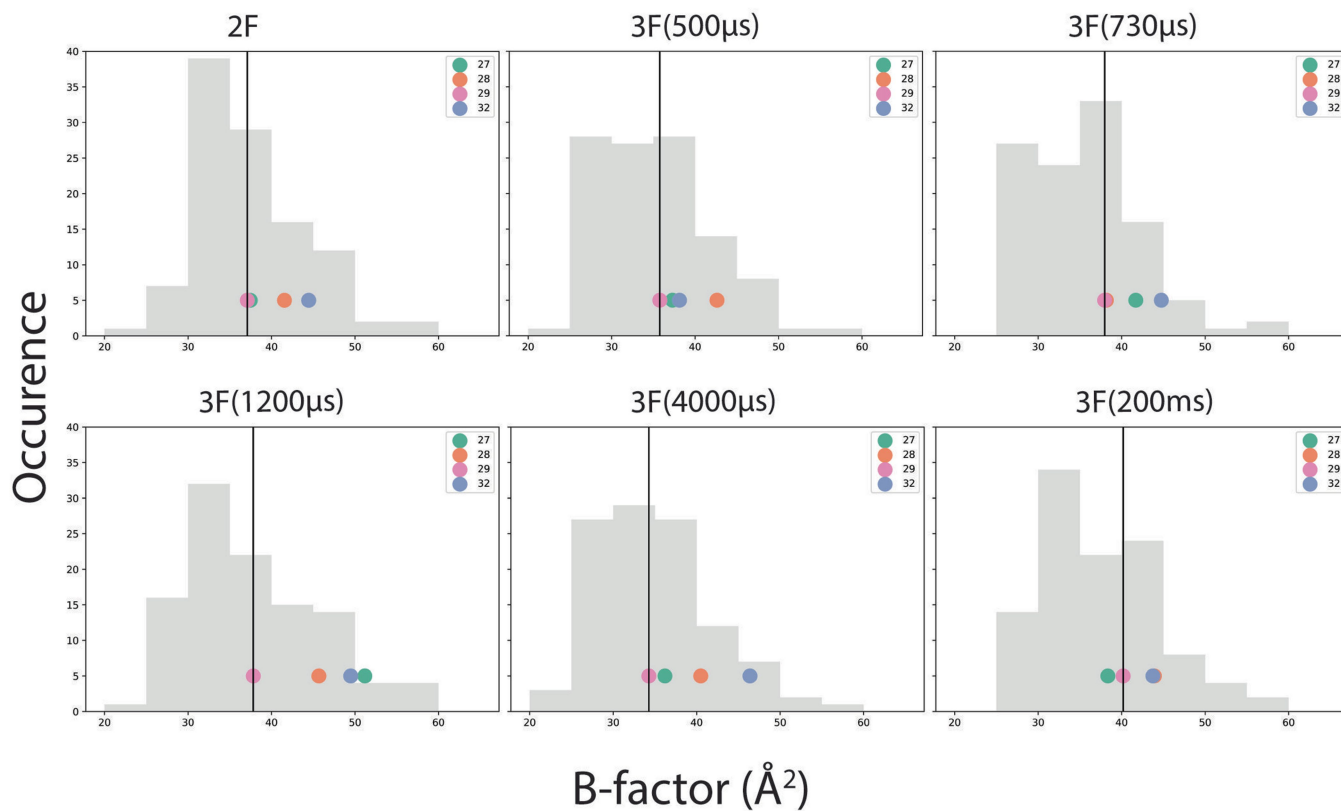


Extended Data Fig. 1 | $mF_{\text{obs}} - DF_{\text{calc}}$ electron density omit map of relevant atoms in the OEC for time points along the $S_3 \rightarrow S_0$ reaction. For comparison of the peak height level, the omit map of the oxygen atom, O2, is also shown. Omit maps for each atom were generated by individually omitting the atom only in

the primary conformer in each dataset. The maps are shown at contour levels of 2.5, 3 and 4 σ . (a) Omit maps of O_x, O5 and O2 (b) Omit maps of W1, W2, W3, W4 and O2. A clear reduction in omit map peak height of O_x, O5, W1, W2, W3, W4 is observed with respect to the reference O2 omit map between 730–2000 μ s.



Extended Data Fig. 2 | $mF_{\text{obs}} - DF_{\text{calc}}$ electron density map near the OEC region in the 3F(500 μs) time point. The map is shown at a contour level of 2.5σ (in yellow). Electron density is observed at this level in-between W3 and W4 (see red arrow), possibly indicating a transient water motion that was also corroborated by the overlap of the omit map densities. For comparison, the individual omit maps of W3 and W4 are also overlaid at 2.5 , 3 and 4σ (color scheme same as in Fig. 2 in main text). Stronger electron density is also observed around W1/D61 possibly related to motion involving a proton transfer as discussed in the main text related to Fig. 2. Electron density around the O1 is possibly related to changes observed in the W26-W30 (referred to as the 'water wheel' region) in this time point (discussed in main text related to Fig. 4).



Extended Data Fig. 3 | Comparing B-factors of key waters in the water wheel region of the O1 channel in the $S_3 \rightarrow S_0$ transition (W27, W28, W32) against a reference water (W29) for time points shown in Fig. 4 in the main text. The B-factor distribution of all the channel waters are overlaid for each

time point for comparison. The y-axis values of the marker points for the 4 waters are just placeholders. There is a stark shift in the relative B-factor values of W27, W28 and W32 in the 3F(1200 µs) time point which coincides with the disappearance of O_x .

Extended Data Table 1 | Data collection, merging and refinement statistics (2F, 2F-alternate, 3F(50 μ s), 3F(250 μ s), 3F(500 μ s))

Data set	2F*	2F-alt*	3F (50 μ s)	3F (250 μ s)	3F (500 μ s)
PDB ID	8EZ5	8F4C	8F4D	8F4E	8F4F
Data Collection					
Resolution range (\AA)	33.65 - 2.09	19.80 - 2.00	33.74 - 2.15	33.58 - 2.09	31.23 - 2.03
Resolution upper bin (\AA)	2.13 - 2.09	2.03 - 2.00	2.19 - 2.15	2.13 - 2.09	2.06 - 2.03
Wavelength (\AA)	1.302	1.241	1.302	1.302	1.305
Space group	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁
Unit cell parameters (\AA)	a=117.0 b=221.6 c=307.8	a=117.3 b=222.8 c=309.1	a=117.1 b=222.2 c=308.4	a=117.0 b=221.8 c=308.0	a=117.5 b=222.8 c=309.5
Lattices merged	10043	35179	7205	8140	17259
Unique reflections	468875	543134	431113	468955	519407
(upper bin)	23274	26980	21309	23217	25669
Completeness	99.97	99.88	99.96	99.97	99.92
(upper bin)	99.98	100.0	99.96	99.97	99.25
CC _{1/2}	99.0	99.0	99.0	99.1	99.1
(upper bin)	5.0	3.8	5.4	6.9	8.9
I/ $\sigma_{B19}(I)$ [†]	3.60	4.17	3.73	3.77	3.83
(upper bin)	0.38	0.41	0.46	0.50	0.38
Wilson B-factor	38.44	35.60	40.10	38.54	36.67
Refinement					
Resolution range (\AA)	33.65 - 2.09	19.80 - 2.00	33.74 - 2.15	33.58 - 2.09	31.23 - 2.03
Resolution upper bin (\AA)	2.13 - 2.09	2.03 - 2.00	2.19 - 2.15	2.13 - 2.09	2.06 - 2.03
R-factor	18.61	17.98	18.83	18.23	19.92
R-free	23.45	22.44	24.33	23.34	24.85
Number of atoms	51845	52239	51657	52072	52011
Number non-hydrogen atoms	51845	52239	51657	52072	52011
Ligands	192	192	192	192	192
Waters	1626	2020	1438	1845	1792
Protein residues	5308	5308	5308	5308	5308
RMS (bonds)	0.009	0.009	0.010	0.009	0.009
RMS (angles)	1.15	1.16	1.18	1.17	1.16
Ramachandran favored	97.32	97.78	97.32	97.57	97.51
Ramachandran outliers	0.19	0.27	0.27	0.17	0.17
Clashscore	6.81	5.23	7.13	6.05	6.25
Average B-factor	44.19	40.61	44.78	44.02	41.85

Note that all reference to the 2F dataset or the S₃ model in the paper uses the 2.09 \AA model. The 2F-alternate model is only provided here for the isomorphous difference maps shown in Fig. 4b.

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Extended Data Table 2 | Data collection, merging and refinement statistics (3F(730 μ s), 3F(1200 μ s), 3F(2000 μ s), 3F(4000 μ s), 3F(200 ms))

Data set	3F (730 μ s)	3F (1200 μ s)	3F (2000 μ s)	3F (4000 μ s)	3F (200 ms)
PDB ID	8F4G	8F4H	8F4I	8F4J	8F4K
Data Collection					
Resolution range refined (\AA)	29.87 - 2.03	29.48 - 2.10	19.73 - 2.00	19.74 - 2.00	33.52 - 2.16
Resolution upper bin (\AA)	2.06 - 2.03	2.14 - 2.10	2.03 - 2.00	2.03 - 2.00	2.19 - 2.16
Wavelength (\AA)	1.302	1.302	1.241	1.241	1.302
Space group	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁	P2 ₁ 2 ₁ 2 ₁
Unit cell parameters (\AA)	a=117.6 b=222.9 c=310.0	a=117.6 b=223.2 c=310.2	a=117.3 b=222.6 c=308.9	a=117.3 b=222.8 c=309.2	a=117.0 b=221.7 c=307.9
Lattices merged	14568	23681	25065	39199	6659
Unique reflections	520277	471242	543130	543130	425199
(upper bin)	25749	23117	26979	26978	21097
Completeness	99.69	99.88	99.88	99.88	99.96
(upper bin)	99.70	98.93	100.0	99.99	99.94
CC _{1/2}	98.8	98.8	99.3	99.0	98.7
(upper bin)	6.3	7.2	9.2	3.3	4.7
$I/\sigma_{B,19}(I)^\dagger$	3.94	3.54	4.72	4.22	3.15
(upper bin)	0.34	0.34	0.51	0.35	0.39
Wilson B-factor	36.65	38.80	35.30	35.70	40.58
Refinement					
Resolution range refined (\AA)	29.87 - 2.03	29.48 - 2.10	19.73 - 2.00	19.74 - 2.00	33.52 - 2.16
Resolution upper bin (\AA)	2.06 - 2.03	2.14 - 2.10	2.03 - 2.00	2.03 - 2.00	2.19 - 2.16
R-factor	18.02	18.77	18.59	18.19	18.51
R-free	22.25	23.24	23.24	22.63	24.10
Number of atoms	52123	51767	52246	52130	51826
Number non-hydrogen atoms	52123	51767	52246	52130	51826
Ligands	192	192	192	192	192
Waters	1896	1540	2021	1905	1609
Protein residues	5308	5308	5308	5308	5308
RMS (bonds)	0.009	0.009	0.009	0.009	0.009
RMS (angles)	1.16	1.17	1.15	1.17	1.14
Ramachandran favored	97.49	97.32	97.47	97.70	97.13
Ramachandran outliers	0.19	0.27	0.23	0.19	0.27
Clashscore	4.83	5.79	5.44	4.96	6.86
Average B-factor	41.16	45.17	39.84	40.71	46.67

Extended Data Table 3 | The estimated population distribution and identity of each of the conformers used in the multi-component modeling of the datasets

Dataset	Primary conformer	Secondary conformer	Tertiary conformer
2F	S ₃ (65%)	S ₂ (35%)	
3F(50μs)	[S _n] (65%)	2F(50μs) (35%)	
3F(250μs)	[S _n] (65%)	2F(250μs) (20%)	S ₂ (15%)
3F(500μs)	[S _n] (65%)	S ₃ (35%)	
3F(730μs)	[S _n] (45%)	S ₃ (40%)	S ₀ (15%)
3F(1200μs)	[S _n] (40%)	S ₃ (35%)	S ₀ (25%)
3F(2000μs)	[S _n] (35%)	S ₃ (35%)	S ₀ (30%)
3F(4000μs)	[S _n] (25%)	S ₃ (35%)	S ₀ (40%)
3F(200ms)	S ₀ (65%)	S ₃ (35%)	

Refer to the discussion *Estimating population distribution in each timepoint* in Methods for more details regarding the conformers.

Article

Extended Data Table 4 | Source data for the plots shown in Fig. 3 of the main text

Dataset	Yz-H190	Mn1-Mn4	Mn1-Mn3	Mn4-O5	E189-Ca1
S ₁	2.64+/-0.08	4.81+/-0.07	3.24+/-0.08	2.13+/-0.16	2.91+/-0.09
S ₂	2.61+/-0.08	4.81+/-0.08	3.22+/-0.07	2.19+/-0.23	2.85+/-0.09
S ₃	2.54+/-0.11	5.05+/-0.09	3.36+/-0.08	2.19+/-0.10	3.55+/-0.13
3F(50μs)	2.78+/-0.12	5.28+/-0.10	3.44+/-0.08	2.13+/-0.12	3.16+/-0.12
3F(250μs)	2.66+/-0.12	5.11+/-0.11	3.43+/-0.09	2.07+/-0.11	2.99+/-0.12
3F(500μs)	2.72+/-0.14	5.14+/-0.10	3.41+/-0.09	2.09+/-0.14	3.09+/-0.12
3F(730μs)	2.50+/-0.14	5.16+/-0.11	3.44+/-0.09	2.09+/-0.10	3.26+/-0.14
3F(1200μs)	2.33+/-0.11	5.18+/-0.13	3.37+/-0.13	2.11+/-0.14	2.90+/-0.17
3F(2000μs)	2.67+/-0.12	4.93+/-0.14	3.27+/-0.08	2.21+/-0.15	3.13+/-0.12
3F(4000μs)	2.89+/-0.18	4.85+/-0.18	3.29+/-0.14	2.22+/-0.17	3.17+/-0.23
S ₀	2.51+/-0.11	5.02+/-0.10	3.33+/-0.09	2.30+/-0.16	2.91+/-0.11

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- | | |
|-----------------|--|
| Data collection | We used the live version of psana (ana-current) provided by SLAC and development versions of open source programs dials.stills_process, the cctbx.xfel GUI and cxi.merge, which are distributed with DIALS packages (3.13.0) publicly available at http://dials.github.io , for data collection. |
| Data analysis | We used development versions of open source programs dials.stills_process, the cctbx.xfel GUI and cxi.merge, which are distributed with DIALS packages (3.13.0) publicly available at http://dials.github.io and END/RAPID structure factor modification software available at http://bl831.als.lbl.gov/END/RAPID/ . We also used Phenix structure refinement software available at https://phenix-online.org (version 1.19.2-4158), Coot structure visualization and modification software version 0.8.8-pre, and PyMol structure visualization and figure rendering software version 2.5. Any custom code used for this publication are publicly available at https://doi.org/10.5281/zenodo.7734707 . |

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Note that full information on the approval of the study protocol must also be provided in the manuscript.

Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

- Life sciences Behavioural & social sciences Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see nature.com/documents/nr-reporting-summary-flat.pdf

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	<input type="text" value="Sample size is stated for all data sets in the Method section and Supplementary Data Table 1. Sample size was chosen to ensure at least 10 independent observations for each structure factor upto the resolution limit reported."/>
Data exclusions	<input type="text" value="During merging of structure factors, lattices that deviated by 1% or more from the reference unit cell were rejected."/>
Replication	<input type="text" value="Diffraction data was collected once for each state. We simulate the experimental errors possible using the END/RAPID approach as described in the Methods section. The END/RAPID approach gives us n=100 different synthetic datasets for which we performed independent refinements, confirming the results and giving an error estimate for atom positions of interest."/>
Randomization	<input type="text" value="Different illumination states were interleaved as much as possible during data collection within the constraints of the XFEL experiment. Data and refinement quality was assessed based on CC1/2 and R-free statistics and the datasets were randomly subsampled to generate these statistics. Other randomizations were not relevant for the XFEL experiment."/>
Blinding	<input type="text" value="Investigators were not blinded to group allocation during data collection. Blinding was not relevant to this study as data for all different states studied was collected interleaved, ensuring similar experimental conditions. Subsequent data treatment was also performed in the same way for all data sets."/>

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

n/a	Included in the study
<input checked="" type="checkbox"/>	<input type="checkbox"/> Antibodies
<input checked="" type="checkbox"/>	<input type="checkbox"/> Eukaryotic cell lines
<input checked="" type="checkbox"/>	<input type="checkbox"/> Palaeontology and archaeology
<input checked="" type="checkbox"/>	<input type="checkbox"/> Animals and other organisms
<input checked="" type="checkbox"/>	<input type="checkbox"/> Clinical data
<input checked="" type="checkbox"/>	<input type="checkbox"/> Dual use research of concern

Methods

n/a	Included in the study
<input checked="" type="checkbox"/>	<input type="checkbox"/> ChIP-seq
<input checked="" type="checkbox"/>	<input type="checkbox"/> Flow cytometry
<input checked="" type="checkbox"/>	<input type="checkbox"/> MRI-based neuroimaging