The results of genome sequencing projects have shown that up to 30% of human proteins occur in cell membranes. Membrane proteins play crucial roles in many biological functions, including the capture of energy from sunlight by plants, the use of energy in cells, and the movement of molecules across cell membranes. It is essential to study their four-dimensional structures, including dynamics to elucidate the function of these molecules.

At the Japanese XFEL facility, SACLA, we are currently developing a data collection system focusing on dynamic crystallography putting a particular effort on membrane proteins. An experimental system for pump-probe experiments based on serial femtosecond crystallography using a viscous material injector has been developed[1]. After a success to visualize a protein motion in bacteriorhodopsin[2], we have used the system to study the light-induced structural changes and the site of O=O bond formation in PSII[3].

Photosystem II (PSII) is a huge membrane-protein complex consisting of 20 different subunits with a total molecular mass of 350 kDa for a monomer. It catalyses light-driven water oxidation at its catalytic centre, the oxygen-evolving complex, OEC. Previously, the structure of PSII has been analysed at 1.9 Å resolution by synchrotron radiation X-rays, which revealed that the OEC is a Mn4CaO5 cluster organized in an asymmetric, ‘distorted-chair’ form. The mechanism of O=O bond formation, however, remains obscure owing to the lack of intermediate-state structures. We studied the structural changes in PSII induced by two-flash illumination at room temperature at a resolution of 2.35 Å using time-resolved serial femtosecond crystallography. A difference Fourier map between the two-flash and dark-adapted states revealed two areas of apparent changes: around the QB/non-haem iron and the Mn4CaO5 cluster. The changes around the QB/non-haem iron region reflected the electron and proton transfers induced by the two-flash illumination. In the region around the OEC, a water molecule located 3.5 Å from the Mn4CaO5 cluster disappeared from the map after two-flash illumination. This reduced the distance between another water molecule and the oxygen atom O4, suggesting that proton transfer also occurred. Importantly, the two-flash-minus-dark difference Fourier map showed an apparent positive peak around O5, a unique μ4-oxo-bridge between Mn1 and Mn4 ions. This indicates the insertion of a new oxygen atom (O6) close to O5, providing an O=O distance of 1.5 Å between these two atoms. This mechanism for the O=O bond formation is consistent with that proposed previously.


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