

Serial crystallography captures dynamic control of falvoenzyme

Jeeru Sucharita

Laboratories India, Science Park & Road, Science, India

 jeerusuchrita98@hotmail.com

Abstract

Flavin coenzymes are universally found in biological redox reactions. DNA photolyases, with their flavin chromophore (FAD), utilize blue light for DNA repair and photoreduction. The latter process involves two single-electron transfers to FAD with an intermittent protonation step to prime the enzyme active for DNA repair. Here we use time-resolved serial femtosecond X-ray crystallography to describe how light-driven electron transfers trigger subsequent nanosecond-to-microsecond entanglement between FAD and its Asn/Arg-Asp redox sensor triad. We found that this key feature within the photolyase-cryptochrome family regulates FAD re-hybridization and protonation. After first electron transfer, the FAD•⁻ isoalloxazine ring twists strongly when the arginine closes in to stabilize the negative charge. Subsequent breakage of the arginine–aspartate salt bridge allows proton transfer from arginine to FAD•⁻. Our molecular videos demonstrate how the protein environment of redox cofactors organizes multiple electron/proton transfer events in an ordered fashion, which could be applicable to other redox systems such as photosynthesis.

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Biography

I am an energetic, reliable, responsible and self-motivated First Class Marine Engineer with deverse experience developed through past employment with Saudi ARAMCO and British Petroleum (contractor for both Companies). I am excellent at working with others, always willing and able to take an increased responsibility whenever necessary to

Achievements

- More than 19 years in the Offshore industry without any negative remarks to the Chief Engineer or Port Engineer duties.

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achieve specified objective on time and with excellence.