生命分子化学セミナーのお知らせ

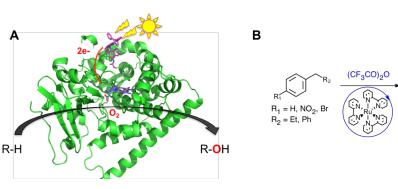
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日時:11月28日(水曜)15:10~16:40

Our laboratory has developed hybrid P450 enzymes containing a Ru(II)-diimine photosensitizer covalently attached to non-native cysteine residues of P450 heme domains. This approach has enabled to harness their synthetic potential upon visible light excitation.¹ High total turnover



numbers and initial reaction rates were obtained in the light-driven hydroxylation of natural long-chain fatty acid substrates.² The crystal structure of the most efficient hybrid enzyme revealed that the photosensitizer is ideally positioned to deliver electrons to the heme active site utilizing the natural electron transfer pathway.³ Our current efforts in optimizing the biocatalyst photocatalytic activity has included a combination of rational and directed evolution approaches while taking advantages of the unique properties of the Ru(II)-diimine complexes.⁴⁻⁶ Selected mutants from a directed evolution screen display several folds enhancement in photocatalytic activity towards various substituted arenes. We also probed the effect of systematically varying the para-substituents on the Ru(II)-diimine photosensitizer on the photocatalytic of the hybrid enzymes and gained insights into the rate limiting step of the photocatalytic process.⁵ Recently, the merging of photoredox catalysis with the hybrid enzyme approach has enabled the selective light-driven chemoenzymatic trifluoromethylation hydroxylation of a wide range of substituted arenes with exquisite selectivity.⁶

連絡先:理学研究科 化学専攻 木村哲就 (ext. 5789) 本セミナーはK-CONNEXのサポートを受けて開催されます

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