

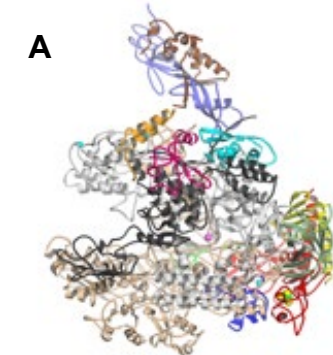


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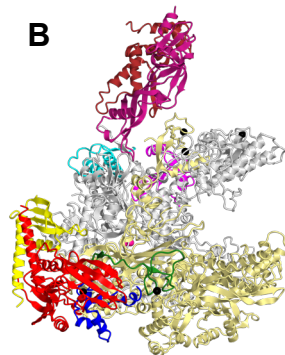
Structural basis for the molecular mechanisms of catalytic reaction and substrate recognition by nucleic acid enzymes.

Associate professor Akira Hirata

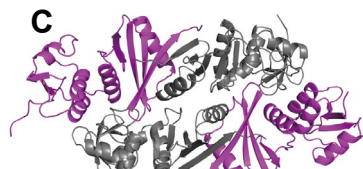
Fig. 1 X-ray structures of nucleic acid enzyme determined.



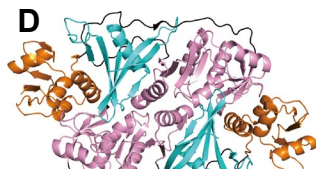
A
Sulfolobus solfataricus
RNA polymerase
Nature 451 (2008)



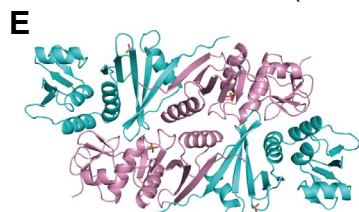
B
Thermococcus kodakarensis
RNA polymerase
Nature commun 5 (2014)



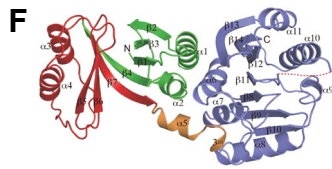
C
Aeropyrum pernix
RNA-splicing endonuclease
Nucleic Acids Res 39 (2011)



D
Candidatus Micrarchaeum acidiphilum
RNA-splicing endonuclease
Nucleic Acids Res 40 (2012)



E
Methanopyrus kandleri
RNA-splicing endonuclease
Nucleic Acids Res 46 (2018)



F
Thermococcus kodakarensis
tRNA methyltransferase Trm11
Nucleic Acids Res 44 (2016)

Content:

Our research goal is to reveal the molecular mechanisms of catalytic reaction and substrate recognition of nucleic acid enzymes regarding transcription and post-transcription events. X-ray crystallography is a powerful tool to determine the structures of proteins and nucleic acids (DNA and RNA). Based on the structural information, we perform biochemical studies of the enzymes. These results obtained often provide a novel insight into the substrate specificity of enzymes.

Our favorite model microorganism is Archaea, which is one of three domains of life. There is strikingly similarity of structure and function of enzymes related to transcription and RNA maturation between archaea and eukaryote because the two domains are evolutionally divided from a common ancestor. So far, we have determined the X-ray structures of archaeal enzymes, two RNA polymerases (Fig. 1A and 1B), three RNA-splicing endonucleases (Fig 1C, 1D and 1E) and tRNA methyltransferase Trm11 (Fig. 1F).

Keywords: gene expression, Archaea, X-ray crystallography

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