

Crystal structure analysis of Hmc

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High molecular mass cytochrome c_3 (Hmc) from *Desulfovibrio vulgaris* Miyazaki F is an electron transfer protein composed of a single polypeptide and 16 heme c groups (total molecular weight is 67 kDa). Primary and tertiary structures of Hmc have not been reported but Hmc may have four cytochrome c_3 domains since the primary structure of Hmc from Hildenborough strain indicates that Hmc is divided into four cytochrome c_3 units.

The function of Hmc *in vivo* remains unclear but the protein must transport electrons because cytochrome c_3 is an electron transfer protein. Redox potentials of heme groups are from positive ($E = 60$ mV) to negative ($E = -260$ mV), which cover the redox potentials of other electron transfer proteins, such as cytochrome c_3 , cytochrome c_{553} , ferredoxins I and II, and rubredoxin. Hmc may receive electrons from hydrogenase.

Data sets for native crystals were collected at BL41XU equipped with the Rigaku R-AXIS IV imaging plate detector system. A wavelength of an incident beam

was 0.708 \AA and a crystal-to-detector distance was 400 mm. The crystals diffracted to 3.0 \AA for the first frame, but we could not complete the data collection because of immediate radiation damage of Hmc crystals.

The image data were processed by using the programs *DENZO* and *SCALEPACK*. The programs indicate that the crystals belong to an orthorhombic space group of $P2_12_12_1$ with unit cell dimensions of $a=60.5 \text{ \AA}$, $b=85.2 \text{ \AA}$, $c=127.6 \text{ \AA}$.

The crystals subjected to the data collections were weak for radiation damage. We have searched new crystallization conditions to improve the quality of the crystals and then we have obtained a new Hmc crystal using a precipitant solution containing 20 % glycerol. It may improve the quality of the crystals and we will collect a complete data set at 100 K using the new crystal. We hope we can complete the data collection at SPring-8 using the new crystals if the occasion arises.