C-136 CRYSTALLOGRAPHY OF BIOLOGICAL MACROMOLECULES

PS04.02.33 CRYSTALLIZATION OF SUPEROXIDE DISMUTASE FROM A HALOPHILIC ORGANISM. Terence P. Lo, Ilona L. Canestrelli, John A. Tainer and Elizabeth D. Getzoff. Department of Molecular Biology, The Scripps Research Institute, 10666 North Torrey Pines Road, La Jolla, California, 92037, USA

Superoxide dismutase from the phototrophic purple sulfur bacterium Ectothiorhodospira halophila has been isolated, purified and crystallized. The importance of this enzyme lies in its ability to scavenge toxic superoxide free radicals. Although these free radicals are a normal byproduct of metabolism, they are responsible for the breakdown of biological macromolecules and thus contribute to such conditions as aging and cancer. In humans, Cu,Zn superoxide dismutase has been implicated in the neurodegenerative disease Familial Amyotrophic Lateral Sclerosis, and so has been the subject of intense scrutiny.

Ectothiorhodospira halophila is a halophilic bacterium found in sunny, hypersaline lakes, and is able to survive in solutions having a salt concentration approaching 4 M. Ectothiorhodospira halophila is also slightly thermophilic, having optimum growth at 42 degrees Celsius. The enzyme was purified directly from the bacterium by assaying for superoxide dismutase activity. While superoxide dismutases from a variety of sources have been found to be highly thermostable, the enzyme from Ectothiorhodospira halophila is extremely thermostable, retaining activity even after boiling. In addition, this superoxide dismutase is highly resistant to cleavage by proteases.

Superoxide dismutase from Ectothiorhodospira halophila has been crystallized in the orthorhombic space group P212121 with a=218.21, b=368.43, c=237.86, a=b=g=90.0. C2221 with a=218.21, b=368.43, c=237.86, a=b=g=90.0. A hexamer per asymmetric unit, it gives a Vm=2.6A³/Da. NSF MCB 9305250

PS04.02.35 DMSO REDUCTASE FROM RHODOBACTER CAPSULATUS. A. S. McAlpine, S. Bailey, CCLRC Daresbury Laboratories, Daresbury, Warrington, WA4 4AD, Cheshire, England

DMSO reductase from the photosynthetic bacteria Rhodobacter capsulatus has been crystallised in two forms which are suitable for X-ray structure determination. DMSO reductase from Rhodobacter is the simplest molybdenum oxotransferase known and as a result makes it an ideal model to study the structure and function of this class of enzymes.

Both crystal forms belong to the same space group, P412121 and have cell dimensions a=b=80.81 Å c=229.75 Å (type I) and a=b=89.30 Å c=230.05 Å (type II). Data have been collected from crystal type I at both room temperature and at 100K.

Several heavy atom derivatives have been collected leading to poor MIR phases. Density modification has improved the electron density maps and interpretation of these maps is underway.

PS04.02.36 STRUCTURAL ROLES OF CRITICAL RESIDUES IN COOPERATIVITY OF SCAPHARCA DIMERIC HEMOGLOBIN. A. Pardanani, M. Bonham, W. E. Royer, Jr., Program in Molecular Medicine, University of Massachusetts Medical Center, Worcester, MA 01605

High resolution atomic structures of the cooperative dimeric hemoglobin found in Scapharca inaequalis have implicated several residues in the cooperative mechanism. Of these, residues Phe 97, Thr 72 and His 101 have been mutated and subjected to functional and structural analysis in order to elucidate their effects on cooperative oxygen binding. Phe 97, whose side-chain undergoes the largest ligand-linked movement, has been hypothesized to be the key regulator of oxygen affinity. Mutation to leucine increases oxygen affinity by an order of magnitude with cooperativity diminished but not eliminated. Crystallographic analysis reveals that many aspects of ligand-linked transitions observed with wild-type are still present in F97L mutant. An important aspect of the ligand-linked structural transitions of Scapharca dimeric hemoglobin is a dramatic change in a cluster of interface water molecules upon ligation. To probe these interface water molecules, we have mutated Thr72 to Val which removes a single hydrogen bond between each subunit and the water network of the deoxygenated species. This isosteric mutation increases oxygen affinity to an even greater extent than seen with the F97L mutant. Interestingly, cooperativity is completely preserved, and may even be marginally enhanced. Structural studies have defined the only change to be the loss of the single hydrogen bonded water molecule, demonstrating the importance of these water molecules in the cooperative mechanism. This proximal histidine, residue 101, provides the only direct coordination of the heme group to the polypeptide chain. Following work of Barrick on sperm whale myoglobin, we have mutated this residue to glycine, and are using exogenous imidizole to functionally replace the lost histidine side chain. This mutation will directly test the importance of the direct link of the iron with the polypeptide chain in the cooperative mechanism. These mutations are providing novel insights into the manner by which two subunits can communicate.