Crystal Structure of the â-subunit of Acetyl-CoA

Carboxylase in C. Glutamicum. <u>Ryo Natsume^a</u>, Minoru Yamada^{a,b}, Miki Senda^a, Tsuyoshi Nakamatsu^b, Sueharu Horinouchi^c, Hisashi Kawasaki^b, Toshiya Senda^d, ^aJBIRC, JBIC. ^bMaterials Sci. Eng., Grad. Sch. Eng., Tokyo Denki Univ. ^cDept. of Biotech., Grad. Sch. of Agriculture and Life Sciences, the Univ. of Tokyo. ^dBIRC, AIST. E-mail: rnatsume@jbirc.aist.go.jp

Acetyl-CoA carboxylases (ACCs) catalyze the first committed step of fatty acid biosynthesis. Although ACC is an essential enzyme (complex) in every organism, the structure-function relationship of ACC remains to be unclear. As the first step for elucidating the structurefunction relationship of ACC, we started the crystallographic analysis of DtsR1. DtsR1 is the β-subunit of ACC multisubunit complex in Corynebacterium glutamicum, which catalyzes the transcarboxylation between biotin and acetyl-CoA.

DtsR1 was over-expressed in E. coli, purified, and crystallized by the sitting-drop vapor diffusion method using PEG 6000 as a precipitant. The approximate dimensions of the obtained crystals were 0.07x0.07x0.03mm³. Diffraction data of the crystals were collected at NW12 of the Photon Factory (Tsukuba), revealing that the crystals belong to the space group R32. The crystal structure of DtsR1 was solved at 3.2Å resolution by the molecular replacement method using single-subunit coordinates of the 12S transcarboxylase (PDB ID: 10N3) as a search model. The obtained structure suggests that the biological unit of DtsR1 is a ring-shaped hexamer with the 32-point group symmetry. Crystallographic refinement of DtsR1 is in progress at 2.7Å

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