Crystal Structures of pmbA and CsrA: Both Reveal New Folds <u>C. Rife</u>^{a,b}, H. Axelrod^{a,b}, M. Miller^{a,b}, R. Schwarzenbacher^{a,c}, Q. Xu^{a,b}, *^aJoint Center for Structural Genomics.* ^bSSRL, Stanford University, Menlo Park, CA, ^cUniversity of California, San Diego, La Jolla, CA. E-mail: crife@slac.stanford.edu

The crystal structure of pmbA reveals a new fold. PmbA, which is encoded by the TM0727 gene of *Thermatoga maritima*, functions in the production of the antibiotic peptide microcin B17[1]. Additionally, pmbA is a putative modulator of DNA gyrase that may function with carbon storage regulator A (CsrA)[2]. The structure was determined using MAD phasing, and two monomers were refined to 1.95Å. The pmbA monomer is composed of two domains, with the N-terminal domain forming a long anti-parallel six-stranded β -sheet, and the Cterminal domain containing three anti-parallel β -sheets, five α -helices and regions of extended coil.

The crystal structure of the carbon storage regulator A (CsrA) gene of *Pseudomonas putida* also reveals a new fold. The structure of dimeric CsrA was determined with MAD phasing and refined to 2.05Å. Each monomer is composed of five consecutive anti-parallel β -strands and one α -helix, with the dimer formed by the intertwining of a pair of β -strands. *E. coli* CsrA is an RNA binding protein which, in conjunction with CsrB-RNA, negatively regulates glycogen biosynthesis, glyconeogenisis and glycogen metabolism, while having a positive regulatory effect on glycolysis[3].

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