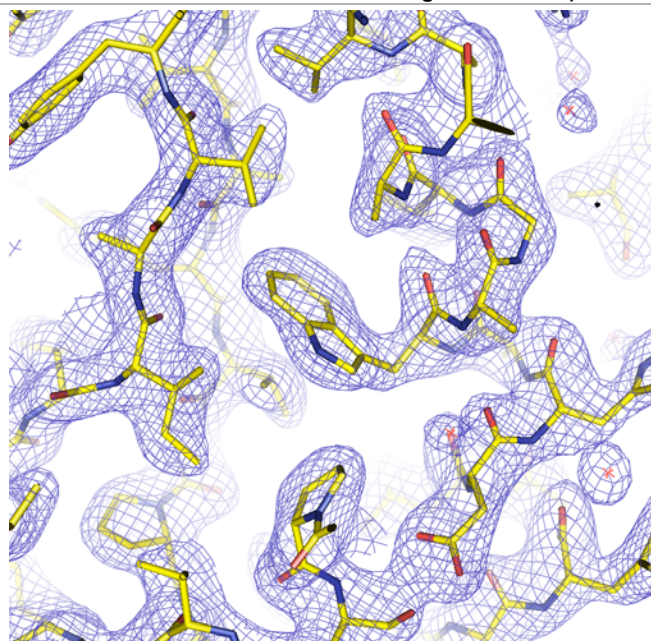
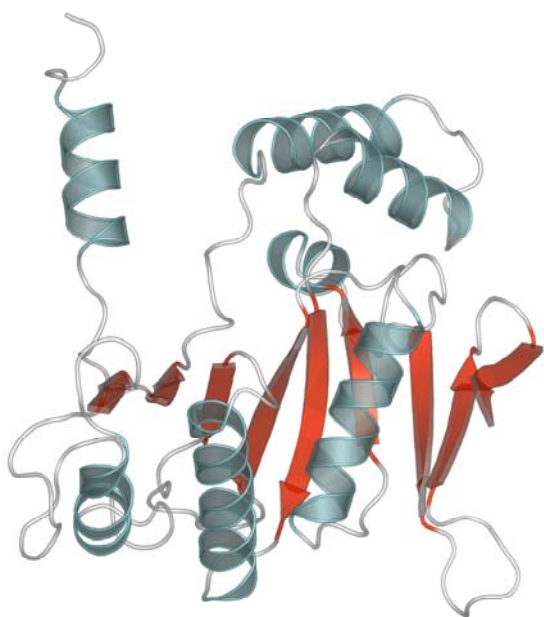


# Center for Eukaryotic Structural Genomics

## Protein Structure Initiative



<b>Target ID</b>	GO.36704	
<b>Source Organism</b>	<i>Homo sapiens</i>	
<b>Target Name</b>	Hs.433573	
<b>PDB Entry</b>	1ZTP	Deposition: 27-May-2005
<b>Function</b>	basophilic leukemia-expressed protein (BLES03, p5326) (FF/Refine: 2Q4K)	
<b>Produced From</b>	<i>E. coli</i> B834, pRARE2, pVP-16	
<b>Structure by X-ray</b>	Resolution: 2.50Å	R-value (R-free): 18.8% (24.5%)
	No. of Residues/ASU: 686 (753)	Complexes/ASU: 3
<b>Data Collected At</b>	Advanced Photon Source NSLS X29A 05-May-2005	
<b>Authors</b>	E. Bitto, C.A. Bingman, H. Robinson, S.T.M. Allard, G.E. Wesenberg, G.N. Phillips, Jr.	



### Structural Features

The crystal structure of BLES03 was solved by SeMet SAD using data collected at the mail-in program at NSLS. BLES03 shows no reliable sequence similarity to any functionally characterized proteins. The structure of BLES03 adopts a fold similar to the eukaryotic transcription initiation factor 4E (eIF4E) a protein that binds the mRNA cap structure. In addition to fold similarity, the general features of the electrostatic surface potential of BLES03 and eIF4E are similar. It is proposed that BLES03 is involved in a biochemical process that requires recognition of nucleic acids.

*References:* (1) Bitto, E., Bingman, C.A., Robinson, H., Allard, S.T.M., Wesenberg, G.E., Phillips, G.N., Jr. (2005) The structure at 2.5Å resolution of human basophilic leukemia-expressed protein BLES03. *Acta Crystallogr Sect F* 61(Pt 9):812-7.

<b>Percent Identity with Nearest PDB Structure at Time Solved</b>	9% (1IPB)
<b>Pfam Cluster</b>	Pfam-B_53833
<b>Sequence Cluster Size : Structures in PDB</b>	18 NR at e<0.1

Center for Eukaryotic Structural Genomics (CESG), University of Wisconsin-Madison Biochemistry Department, 433 Babcock Drive, Madison, WI 53706-1549; phone: 608.263.2183; fax: 608.890.1942; email: [cesginfo@biochem.wisc.edu](mailto:cesginfo@biochem.wisc.edu); website: <http://www.uwstructuralgenomics.org>. This research funded by NIH / NIGMS Protein Structure Initiative grants U54 GM074901 and P50 GM064598.