Letters to the Editor

¹H, ¹³C and ¹⁵N NMR assignments for AlgH, a putative transcriptional regulator from *Pseudomonas aeruginosa* DOI 10.1007/s10858-005-1271-9

AlgH (20.2 kDa, 189 residues) from *Pseudomonas aeruginosa* is known to regulate the biosynthesis of the exopolysaccharide alginate, as well as siderophore, protease, rhamnolipid and nucleoside diphosphate kinase, putatively at the transcriptional level, and has been suggested to be a 'global regulator of several functions' (Schlictman et al., 1995). The mechanism by which AlgH acts has not been defined, and no functional or structural information is currently available for any members of the AlgH family (Pfam PF02622). We have demonstrated previously the production, labeling and purification of this stable, folded, globular protein (Bieber Urbauer et al., 2005). Heteronuclear 2D and 3D methods were used to obtain nearly complete main chain ¹H, ¹³C, and ¹⁵N (>95%) and aliphatic side chain ¹H, and ¹³C assignments (>93%). Unassigned resonances include the 14 Pro ¹⁵N resonances and many ¹³C and ¹H resonances of aromatic groups. The BMRB deposit has Accession No. 6644.

References: Schlictman et al. (1995) J. Bacteriol., 177, 2469–2474: Bieber Urbauer et al. (2005) Prot. Expr. Purif. 43, 57–64.

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Supplementary material is available in electronic format at http://dx.doi.org/10.1007/s10858-005-1271-9

¹H, ¹³C and ¹⁵N resonance assignments of the AT-Rich Interaction Domain (ARID) of Jumonji DOI 10.1007/s10858-005-1282-6

Jumonji (Jmj) is a ubiquitous, transcriptional repressor protein which plays important roles in development, cell growth and gene expression (Takeuchi et al., 1995). Jmj is a member of Jmj transcription factor family, which contains an AT-rich interaction domain (ARID) (Wilsker et al., 2002) and Jumonji-like domains (JmjN/JmjC). Jmj ARID is a putative DNA-binding domain, which shares about 30% sequence identity to the ARID family members. In this study, we prepared non-labeled, uniformly ¹⁵N-labeled and ¹³C/¹⁵N-labeled Jmj ARID (116 residues, L615 to K730) to determine the solution structure of the protein and to examine the structure–function relationships of the protein by NMR experiments. NMR spectra were acquired on a Bruker AVANCE 500 spectrometer equipped with ¹H/¹³C/¹⁵N cryogenic probe at 15 and 25°C. Most of backbone (99% of ¹⁵N and HN, 98% of ¹³CO, 98% of ¹³Ca, 99% of H α , 96% of ¹³C β , 95% of H β) and side-chain resonances (approximately 86% of ¹H and ¹³C resonances) were assigned. BMRB deposit with accession number 6607.

References: Takeuchi et al. (1995) *Genes Dev.*, **9**, 1211–1222; Wilsker et al. (2002) *Cell Growth Differ.*, **13**, 95–106.

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