

## Letter to the Editor

### NMR assignment of peptidyl-tRNA hydrolase from *Mycobacterium tuberculosis* H37Rv

DOI 10.1007/s10858-006-9031-z

Peptidyl-tRNA hydrolase (PTH, EC 3.1.1.29) is an enzyme essential for the viability of bacteria and is responsible for alleviating the toxic effects of accumulation of prematurely dissociated peptidyl-tRNA in the cytosol. We have cloned and over-expressed the protein annotated as a putative PTH (ORF Rv1014c) from the genome of *Mycobacterium tuberculosis* H37Rv. In order to understand the dynamic structure of this protein and the protein and peptidyl-tRNA interactions in solution, we have initiated determination of its 3D solution structure. *M. tuberculosis* PTH (mtPTH) consists of 191 residues and has a molecular weight of 20,550.3 Da. Using 2D and 3D heteronuclear NMR experiments on uniformly  $^{13}\text{C}$ ,  $^{15}\text{N}$ -labeled mtPTH, we have assigned 171 out of a total of 179 non-proline amide residues. The amide residues that could not be assigned are Met1, Ala2, Glu3, Ala15, Asn16, Lys44, Gly111, Asn116, and Arg143. In addition, assignment of 97% of  $\text{C}^\alpha$ ,  $\text{C}^\beta$ , and CO, 95% of non-aromatic and non-carbonyl side-chain carbons, 89% of  $\text{H}^\alpha$ , and 86% of  $\text{H}^\beta$  is complete. Based on the chemical shifts obtained, 7  $\beta$ -strands and 10  $\alpha$ -helices were identified. The overall fold of mtPTH is similar to that of *E. coli* PTH, with which it has 52% sequence homology (Schmitt et al., 1997). BMRB deposits with accession number 7055.

Reference: Schmitt et al. (1997) *EMBO J.*, **16**, 4760–4769.

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**Supplementary material** to this paper is available in electronic format at <http://dx.doi.org/10.1007/s10858-006-9031-z>.