

## NMR resonance assignments of the human high mobility group protein HMGA1

Garry W. Buchko · Shuisong Ni ·  
Natacha M. Lourette · Raymond Reeves ·  
Michael A. Kennedy

Received: 6 October 2006 / Accepted: 27 October 2006 / Published online: 6 January 2007  
© Springer Science+Business Media B.V. 2007

Human high mobility group protein HMGA1 is a 107-residue, non-histone chromatin protein with a wide sphere of influence including embryogenesis, apoptosis, differentiation, cell proliferation, and cancer development (Reeves 2001). Due to the repetitive nature of the three DNA-binding domains, C-terminal strings of glutamic acid residues, and unstructured nature in the absence of A-T rich regions of DNA and/or other proteins, resonance assignments were challenging. Especially useful was the HNN experiment (Panchal et al. 2001), a set of truncated HMGA1 constructs, and some high resolution NMR data ( $^1\text{H}$  900 MHz). Except for absolute assignment of R60 and R86, all 82 amides were assigned to cross peaks in the  $^1\text{H}$ - $^{15}\text{N}$  HSQC spectrum and many of the side chain  $^{13}\text{C}$  and  $^1\text{H}$  resonances were assigned (BMRB—7279). The intensity of the amide cross peaks for residues

E3–S9 and S64–K67 were much weaker than the other amide cross peaks in the  $^1\text{H}$ - $^{15}\text{N}$  HSQC spectrum suggesting that even in this unstructured protein there are regions experiencing motion different from the molecule as a whole.

**Acknowledgements** This work was supported by an LDRD project at EMSL, a national scientific user facility sponsored by a US. DOE BER program located at PNNL.

### References

- Reeves R (2001) Molecular biology of HMGA proteins: hubs of nuclear function. *Gene* 277:63–81  
Panchal SC et al. (2001) Improved 3D triplet resonance experiments, HNN and HN(C)N, for  $\text{H}^N$  and  $^{15}\text{N}$  sequential correlations in labeled proteins: application to unfold proteins. *J Biomol NMR* 20:135–147

**Electronic supplementary material** The online version of this article (doi: 10.1007/s10858-006-9116-8) contains supplementary material, which is available to authorized users.

G. W. Buchko (✉) · S. Ni · N. M. Lourette ·  
M. A. Kennedy  
Pacific Northwest National Laboratory, Richland, WA  
99352, USA  
e-mail: garry.buchko@pnl.gov

R. Reeves  
Department of Biochemistry and Biophysics, Washington  
State University, Pullman, WA 99164, USA