Letter to the Editor: Chemical shift and secondary structure conservation of the PNT/SAM domains from the Ets family of transcription factors

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Biological context

The DNA-binding ETS domain defines the Ets family of transcription factors. A subset of this family, including Ets-1, Ets-2, GABPα, Erg, Fli-1 and Tel, also share a conserved Pointed (PNT) or SAM domain that may function in oligomerization, MAP kinase docking or association with other regulatory proteins. The solution structure of the PNT/SAM domain from Ets-1 (PDB: 1qbv; Slupsky et al., 1998a) has an architecture of a core four α-helical bundle interfaced with an additional N-terminal helix. In contrast, the crystal structure of the homologous domain from Tel (1ji7; Kim et al., 2001) shows only the four core helices, as also seen in the structures of SAM domains from non-Ets family members such as p73 (1cok), polyhomeotic (1kw4) and the Ephrin receptors (1b4f, 1sgg, 1b0x). The additional N-terminal helix present in Ets-1 is an integral part of the structured PNT/SAM domain as evident by 15N relaxation measurements and the requirement of these residues for proper protein expression and folding (Slupsky et al., 1998a). However, the extent to which this helix exists in PNT/SAM domains from other Ets proteins remains to be established.

Methods and experiments

The genes encoding the PNT/SAM domains of murine GABPα (residues 168–254) and human Erg (residues 108–201 of isoform p55Erg) were cloned into pET22b and pET28a, respectively, using PCR-generated restriction enzyme sites. Proteins were expressed in BL21(λDE3) grown in Luria Broth or M9 minimal media supplemented with 15NH4Cl (1 g l⁻¹) or 13C6-glucose (3 g l⁻¹). Purification of GABPα(168–254) was achieved with Q-sepharose anion-exchange chromatography, followed by dialysis into 20 mM sodium phosphate (pH 7.2), 20 mM NaCl and 10 mM DTT. His6-tagged Erg(108–201) was purified using Ni2⁺-affinity chromatography, with final dialysis into 20 mM potassium phosphate (pH 7.0), 50 mM NaCl and 2 mM DTT. Following thrombin removal of the His6-tag, there remained three N-terminal residues (Gly-Ser-His) preceding Met108 in Erg(108–201). Samples contained 0.5 to 2.5 mM protein with 10% or 99% D₂O added for the lock. All spectra were recorded at 30 °C using a 500 MHz Varian Unity or a 600 MHz Varian Inova NMR spectrometer, both equipped with a triple resonance gradient probe. The spectra were analyzed using Felix 95.0/2000 (Accelrys, Inc.) and Sparky 3 (T.D. Goddard and D.G. Kneller, University of California, San Francisco). Backbone and aliphatic sidechain assignment were obtained from 13C- and 15N-HSQC, HNCACB, CBCA(CO)NH, HNCO, HACAN, (H)CC(CO)NH- and HCCH-TOCSY spectra (Sattler et al., 1999; Kanelis et al., 2000). Stereospecific assignments were achieved using the method of Senn et al. (1989) for Val and Leu methyl groups and McIntosh et al. (1997) for the sidechain amides of Gln and Asn. Resonances of aromatic sidechains were assigned as described previously (Slupsky et al., 1998b).

We have determined the 1H, 13C and 15N resonance assignments for the PNT/SAM domains of GABPα and Erg. With the exception of the 12 N-terminal residues of Erg(108–201), a striking level of
similarity exists between the $^{13}$C$\alpha$ secondary chemical shifts ($\delta_{\text{obs}} - \delta_{\text{random coil}}$) observed for corresponding residues from the sequence alignment of Ets-1, GABP$\alpha$ and Erg (Figure 1). The secondary chemical shifts of mainchain $^1$H$\alpha$, $^1$H$\beta$, $^1$H$\gamma$, and $^1$H$\delta$ exhibit similar conservation (data not shown). These results confirm the potential of programs that exploit reference chemical shifts from one protein to predict the chemical shifts of a homologous protein (e.g., SHIFTY; Wishart et al., 1997). This observed conservation in secondary shifts should allow for facile NMR spectral assignment of additional PNT/SAM domains.

Deduction of secondary structure from chemical shifts and $^3$J$_{HN-H\alpha}$ values (not shown) also demonstrates that GABP$\alpha$, but not Erg, contains the additional N-terminal helix found in Ets-1. This result confirms a surprising diversity in the architecture of the Ets family PNT/SAM domains that may extend to variation in function.

**Extent of assignments and data deposition**

Complete backbone $^{15}$N, $^1$H$\alpha$, $^{13}$C$'$, and $^{13}$C$\alpha$ assignments of GABP$\alpha$(168–254) and Erg(108–201) have been determined with the exception of proline $^{15}$N chemical shifts in GABP$\alpha$(168–254). $^{13}$C$'$ chemical shifts in Erg(108–201) for residues preceding a proline, and the $^{13}$C$\alpha$ resonance for the C-terminal Pro201. Assignment of resonances from sidechain carbon, nitrogen and non-labile hydrogen nuclei are nearly complete. Methyl groups of Leu and Val have been stereospecifically assigned, as have the sidechain amides of Asn and Gln. The $^1$H, $^{13}$C and $^{15}$N chemical shifts for GABP$\alpha$(168–254) and Erg(108–201) have been deposited in the BioMagResBank database (http://www.bmrb.wisc.edu) under BMRB accession numbers 5401 and 5399, respectively.

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**References**


