Backbone chemical shift assignments of mouse HOXA13 DNA binding domain bound to duplex DNA

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Abstract The homeobox gene (Hoxa13) codes for a transcription factor protein that binds to AT-rich DNA sequences and controls expression of many important proteins during embryonic morphogenesis. We report complete backbone NMR chemical shift assignments of mouse Hoxa13 DNA binding domain bound to an 11-residue DNA duplex (BMRB no. 16577).

Keywords HOXA13 · Homeodomain · DNA binding domain · NMR · Duplex DNA

Biological context

Homeobox (Hox) genes encode a conserved family of transcription factor proteins that are critically important in vertebrate development (Krumlauf 1994). In humans, the Hox genes are distributed into four linkage groups (HOXA, B, C, D) comprising 39 genes located on chromosomes 7, 17, 12, and 2. Recently, mutations in HOXA13 have been associated with Hand-Foot-Genital (HFGS) and Guttman syndromes (GS), autosomal dominant disorders that profoundly affect limb and genitourinary development causing defects in the digits, carpal/tarsal bones, uterus, bladder, Mullerian ducts, and the external genitalia (Innis et al. 2002; Jorgensen et al. 2010; Mortlock and Innis 1997). Analysis of the DNA sequences bound by HOXA13 revealed high affinity for a unique duplex containing a consensus sequence of: 5-AAATAAAA-3’ (Knosp et al. 2007). Characterization of this high affinity sequence in mouse models confirmed that HOXA13 binds to this sequence in vivo and regulates the tissue-specific expression of factors such as bone morphogenetic proteins 2 and 7, to facilitate limb skeletal development (Knosp et al. 2004, 2007).

While it is well-established that mutations affecting the DNA binding domain of HOXA13 cause HFGS and GS, less is known about the amino acids that facilitate HOXA13 DNA binding and/or sequence specificity (Innis et al. 2002; Mortlock and Innis 1997). To gain insight into the molecular basis for the phenotypes present in HFGS and GS, an atomic-resolution structure of HOXA13 bound to duplex DNA is needed to define which amino acids facilitate HOXA13 DNA binding and DNA sequence specificity. Here we report the NMR assignments of the mouse HOXA13 DNA binding domain bound to an 11-residue duplex, forming a complex called, A13DBD-DNA. The NMR assignments of A13DBD-DNA are an important first step toward elucidating the amino acids necessary for sequence specific DNA binding and provide a functional explanation for the loss of HOXA13 function in HFGS and GS.

Methods and experiments

Preparation of HOXA13 DNA binding domain bound to duplex DNA (A13DBD-DNA)

Uniformly $^{15}$N-labeled and $^{13}$C, $^{15}$N-labeled A13DBD were prepared as described previously (Zhang et al. 2009).
Two complementary single strands of DNA (1: 5'-CAA ATAAATC-3' and 2: 5'-GATTTTATTTG-3') purchased from Bioneer Inc. were synthesized and HPLC purified. For duplex DNA preparation, the two complementary DNA single strands were dissolved in 1x TE buffer (10 mM TrisHCl, 1 mM EDTA, pH 7.5), mixed in 1:1 M ratio, and annealed at 95°C for 10 min, and then cooled down to room temperature. An aliquot of duplex DNA was added to a stock solution of A13DBD in a 1:1 M ratio and incubated at 15°C for 1 h and the complex was further purified by gel-filtration size-exclusion chromatography (Superdex-75).

NMR spectroscopy

Samples of A13DBD-DNA for NMR analysis were prepared as described above, and exchanged into a buffer containing 20 mM sodium phosphate (pH 6.0) with 5 mM Magnesium chloride and 95% H2O/5% D2O, and finally concentrated to 0.3 ml giving a final protein concentration of 0.5 mM. All NMR experiments were performed at 310 K on a Bruker Avance 800 MHz spectrometer equipped with a four channel interface and triple resonance cryogenic (TCI) probe. The 15N-1H HSQC spectrum (Fig. 1) was recorded with 256 × 2048 complex points for 15N(F1) and 1H(F2). Assignment of backbone resonances was obtained by analyzing the following spectra: HNCA, HNCACB, CBCA(CO)NH, HNCO. The NMR data were processed using NMRPipe and analyzed using Sparky.

Assignments and data deposition

Figure 1 presents the 15N-1H HSQC spectrum of A13DBD-DNA at pH 6.0 to illustrate representative backbone resonance assignments. NMR assignments were based on 3D heteronuclear NMR experiments performed on 13C/15N-labeled A13DBD (residues 1–73). The protein sample in this study consists of 73 native residues without affinity tags or cloning modification. Nearly all non-proline residues exhibited strong backbone amide resonances with uniform intensities in the HSQC spectrum, indicative of α helices.
well-defined three-dimensional protein structure. All backbone resonances (*^1^H^N*, *^1^H^N*, *^1^H^C^α*, *^1^H^C^β*, and *^1^H^C^O*) were assigned except for the first two residues (G1 and S2) that have very weak NMR intensities. The chemical shift index of each amino acid residue was calculated as described previously (Wishart and Sykes 1994) and revealed three *α*-helices (*α*1: T15–T28; *α*2: K34–N45; *α*3: E48–K68) and a short *β*-strand (F31–T33). The protein secondary structure closely resembles the canonical secondary structure and topology seen in other homeobox proteins (Piper et al. 1999). The chemical shift assignments (*^1^H*, *^1^H^N*, *^1^H^C^) of the A13DBD-DNA complex have been deposited in the BioMagResBank (http://www.bmrb.wisc.edu) under accession number 16577.

Figure 2 presents the chemical shift perturbation caused by DNA binding to A13DBD. The weighted average chemical shift difference (A13DBD-DNA vs. free protein) was calculated for backbone amide *^1^H* and *^1^H^N* resonances of each residue. N-terminal residues (K9, R11 and Y14) and residues in the C-terminal helix show the largest chemical shift differences, suggesting that these residues may interact with duplex DNA. Residues I53, Q56, N57 and V60 are all on the same solvent exposed surface of the C-terminal helix and are predicted to make direct sequence-specific contacts inside the major groove analogous to DNA contacts observed in the HoxB1-DNA complex (Piper et al. 1999).

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