Backbone and side chain NMR assignments for the intrinsically disordered cytoplasmic domain of human neuroligin-3

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Abstract Neuroligins act as heterophilic adhesion molecules at neuronal synapses. Their cytoplasmic domains interact with synaptic scaffolding proteins, and have been shown to be intrinsically disordered. Here we report the backbone and side chain $^1$H, $^{13}$C and $^{15}$N resonance assignments for the cytoplasmic domain of human neuroligin 3.

Keywords Intrinsically disordered protein · Neural cell adhesion · PDZ-binding domain · Cholinesterase-like adhesion molecule

Biological context

Synaptic adhesion proteins bridge pre- and postsynaptic specializations at neuronal synapses. As well as providing mechanical bridging, they participate in formation, function, and plasticity of such synapses (Dalva et al. 2007). Cholinesterase-like (ChE-like) adhesion molecules (CLAMs) are a class of neuronal cell adhesion molecules that are characterised by an extracellular N-terminal domain displaying sequence homology to acetylcholinesterase (although lacking catalytic activity), a single transmembrane segment, and a small intracellular C-terminal domain (Gilbert and Auld 2005).

Neuroligins (NLs) are a family of CLAMs, of which four isoforms have been identified (Baudouin and Scheiffele 2010), that are embedded in the postsynaptic membrane. Their ChE-like extracellular domains interact with their presynaptic partners, the neurexins.

The intracellular domain of human NL3 (hNL3-cyt) is disordered in the absence of binding partners (Paz et al. 2008), and has been shown to interact with several scaffolding proteins, including PSD-95 (Irie et al. 1997), via a C-terminal PDZ-recognition motif.

Mutations in NL3 and NL4 have been shown to be associated with autism (Jamain et al. 2003), and the neurexin-NL-PSD95 interaction has been implicated in ischemia (Li et al. 2007). Conformational characterization of the disordered cytoplasmic domains of the NLs, and of their interactions with partner proteins, is important for understanding structural and functional features of synaptic junctions, and may also lead to possible targets for the treatment of neurological disorders.

Here we report the backbone and side-chain NMR resonance assignments of hNL3-cyt.
Methods and experiments

Expression and purification of uniformly labelled hNL3-cyt were performed as described previously (Paz et al. 2008). The NMR sample contained 1 mM of uniformly 13C, 15N-enriched protein in a solution of 5 mM dithiothreitol (DTT)/20 mM phosphate, pH 6.0, containing 0.1 mM DSS (2,2-Dimethyl-2-silapentane-5-sulfonic acid). Sodium azide was added to 0.02% (w/v), and D2O to 7% (v/v).

All experiments were performed at 25°C on a Varian Unity Inova 600 MHz spectrometer, equipped with a triple-resonance room temperature probehead and single-axis pulsed field gradient capabilities.

2D [1H–15N]-HSQC, 3D HNCA/HNCACB/CBCA(CO)NH, 15N-edited NOESY-HSQC (Sattler et al. 1999), 3D HN(CA)NO (Panchal et al. 2001), and 3D (H)CC(CO)NH-TOCSY (Logan et al. 1993) experiments were used to obtain the sequential backbone resonance assignment. Side chain assignments were achieved with 3D CO-CA (Dijkstra et al. 1994), 3D 15N-edited TOCSY-HSQC (Marion et al. 1989), 3D HC(C)H-TOCSY (Bax et al.1990), and 3D (H)CC(CO)NH-TOCSY (Logan et al. 1993) experiments.

Data were processed with the NMRPipe/NMRDraw software package (Delaglio et al. 1995), and analyzed using the program Sparky (Goddard and Kneller 2004). All chemical shifts are referenced to DSS according to IUPAC recommendation (Markley et al. 1998).

Assignments, data deposition and intrinsic exchange rates

Figure 1 illustrates the 2D [1H–15N]-HSQC spectrum and assignments of the amide resonances of hNL3-cyt; its unfolded nature is evident from the small dispersion in the amide proton resonances. The construct contains 139 residues, the first 10 of which are a 6xHis-tag, followed by a short linker sequence (see also Fig. 4). Following a standard sequential assignment procedure, 98% of the observable backbone 1H, 15N and 13C resonances were assigned (excluding the tag). The observable 1H/15N pairs are displayed in Fig. 1, labeled by their one-letter amino acid code and residue number. Considering only aliphatic 1H and 13C atoms, side chain assignments are complete to 92%.

The resonance assignments were deposited in the BioMagResBank (http://www.bmrb.wisc.edu/) under accession number 17290.

To understand whether the absence of some backbone amide proton resonances in our experiments was due to fast exchange with the solvent, intrinsic exchange rates were calculated (Bai et al. 1993) using the program Sphere (http://www.fccc.edu/research/labs/roder/sphere/sphere.html) and these values are plotted in Fig. 2. The data show that in all cases where the amide proton resonance was not assigned (for residues H20, H64, H134, and S135), intrinsic exchange rates faster than 15 s⁻¹ are predicted.

Secondary structure prediction

Considering charge and hydrophobicity in a so-called Uversky plot (Uversky et al. 2000), hNL3-cyt was found to lie very close to the boundary between proteins predicted to be unfolded and folded (Zeev-Ben-Mordehai et al. 2003). A bioinformatics analysis using several predictors had been performed earlier for hNL3-cyt (Paz et al. 2008). It
predicted that the N-terminal section (residues 1–70) is completely disordered, while the C-terminal segment is predicted to be somewhat more ordered, depending on the predictor used. Here we use two bioinformatic methods to predict structural propensity in hNL3-cyt: PONDR\textsuperscript{/C210} (Romero et al. 2001) and Jpred 3 (Cole et al. 2008), which consider sequence information alone. These results were compared to secondary structural propensity (SSP) scores (Marsh et al. 2006), derived from the experimental chemical shift data.

PONDR\textsuperscript{/C210} predicts that 73\% of hNL3-cyt is disordered, consistent with the small dispersion of amide chemical shifts measured by NMR. PONDR scores of the sequence are displayed in Fig. 3, where a score between 0.5 and 1 indicates disorder, and a score between 0 and 0.5 indicates order. The figure shows that the first 75 residues are predicted to be disordered, and that order and disorder alternate in the remainder of the sequence. Specifically, two ordered stretches are predicted, viz., residues 76–92 and 111–128.

Secondary structure prediction for hNL3-cyt was also performed based on sequence similarity using Jpred 3, 88\% of the residues are predicted to be disordered by the Jpred 3 algorithm. As seen in Fig. 4, three short helical stretches are predicted, residues 25–29, 53–60 and 84–86.

A prediction of secondary structural propensity was also performed based on chemical shift analysis using SSP. The analysis used measured chemical shifts of C\textsuperscript{\alpha}, C\textsuperscript{\beta} and H\textsuperscript{\alpha} atoms, and an averaging window of five residues (Fig. 5). A positive SSP score indicates a propensity for \alpha-structure, and a negative score indicates a propensity for \beta- or extended structure. Residues in fully formed \alpha-helices and \beta-strands are given scores of +1 and −1, respectively. As can be seen in Fig. 4, the bulk of hNL3-cyt yields SSP scores within the range of −0.2 to 0.2, indicating that the sequence has little propensity to assume a stable secondary
structure. Only two segments have values >0.2, residues 51–56 and 82–86.

Our experimental data clearly show that hNL3-cyt does not adopt a well-defined tertiary structure, but is intrinsically disordered. The analysis of measured chemical shifts only agrees with the two prediction algorithms used here on one point: the presence of a small stretch with helical propensity around residue 84. It furthermore agrees with the Jpred 3 prediction that a labile helical segment is present around 52–60. The experimental data do not support the prediction by Jpred 3 of a helical segment around residue 27, and indicate that the chain adopts an extended structure in this region.

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