NMR structure of the death domain of the p75 neurotrophin receptor

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The intracellular domain of the p75 neurotrophin receptor (p75ICD) lacks catalytic activity but contains a motif similar to death domains found in the cytoplasmic regions of members of the tumor necrosis factor receptor family and their downstream targets. Although some aspects of the signaling pathways downstream of p75 have been elucidated recently, mechanisms of receptor activation and proximal signaling events are unknown. Here we report the nuclear magnetic resonance (NMR) structure of the 145 residue long p75ICD. The death domain of p75ICD consists of two perpendicular sets of three helices packed into a globular structure. The polypeptide segment connecting the transmembrane and death domains as well as the serine/threonine-rich C-terminal end are highly flexible in p75ICD. Unlike the death domains involved in signaling by the TNF receptor and Fas, p75ICD does not self-associate in solution. A surface area devoid of charged residues in the p75ICD death domain may indicate a potential site of interaction with downstream targets.

Keywords: death domain/NMR/p75 neurotrophin receptor

Introduction

The low-affinity p75 neurotrophin receptor (p75) was identified 10 years ago as a membrane protein with the ability to bind nerve growth factor (NGF) (Chao et al., 1986; Radeke et al., 1987). Later works established that p75 was capable of binding all members of the NGF family (Rodriguez-Tébar et al., 1990, 1992; Rydén et al., 1995), also known as the neurotrophins, a group of structurally and functionally related polypeptides that regulate neuronal survival, differentiation and repair (Ibañez et al., 1995; Lewin and Barde, 1996). Because the short intracellular domain of p75 (p75ICD) lacked features found in other receptors with known catalytic activity, the subsequent discovery of the Trk family of receptor tyrosine kinases as ligand-specific signaling receptors for the neurotrophins cast doubts on the intrinsic signaling capabilities of p75 (Kaplan et al., 1991; Klein et al., 1991). Various models involving the ligand-binding domain of p75 have been presented to account for the documented ability of p75 to aid in the formation of high affinity binding sites (Hempstead et al., 1991; Mahadeo et al., 1994) and in Trk signaling (Barker and Shooter, 1994; Hantzopoulos et al., 1994; Verdi et al., 1994), including models of ligand presentation, ligand concentration and changes in receptor conformation.

The extracellular domain of p75 has structural similarities to the tumor necrosis factor p55TNFR receptor and to the Fas antigen (Meakin and Shooter, 1992), which, upon ligand binding, signal through a cascade of protein–protein interactions triggered in part by a 90 residue long motif in the cytoplasmic tail of these receptors. Because of the cytotoxic activities of p55 TNFR and Fas, this motif has been known as the death domain (Tartaglia et al., 1993). Several downstream targets of the p55TNFR and Fas death domains have been identified and these also contain death domain sequences (Boldin et al., 1995; Chinnaiyan et al., 1995; Hsu et al., 1995; Stanger et al., 1995; Duan and Dixit, 1997) indicating a signaling mechanism triggered by the association of death domain-containing proteins (Varfolomeev et al., 1996). The similarity of the ligand-binding domain of p75 to those of p55TNFR and Fas has been reinforced by the discovery of a somewhat divergent death domain in p75ICD (Chapman, 1995; Feinstein et al., 1995). This second subtype of death domain has also been found in several other proteins, including p100 NF-κB, p105 NF-κB, DAP kinase and myD88 (Feinstein et al., 1995). The functional role of the death domain in any of these proteins is, however, unknown. Recently, evidence has begun to accumulate supporting the intrinsic signaling capacities of p75. It has been shown that neurotrophin binding to p75, in a cell devoid of cognate Trk receptors, stimulates sphingomyelin hydrolysis and ceramide production (Dobrowsky et al., 1994, 1995). In addition, only NGF among the neurotrophins has been shown to trigger the nuclear translocation of NF-κB in a p75-dependent fashion in Schwann cells and p75-expressing fibroblasts (Carter et al., 1996). Finally, using the yeast two-hybrid system, at least one novel protein has been isolated that is capable of interacting with p75ICD (Carter et al., 1996a). A link between p75 and cell death has also been established, although there are discrepancies as to whether the death induced by p75 is ligand-dependent or ligand-independent (Rabizadeh et al., 1993; Barrett and Bartlett, 1994; Casaccia-Bonnefil et al., 1996; Frade et al., 1996). The importance of the death domain as a signaling module, and of p75 in the control of cell death and neurotrophin signaling, prompted us to elucidate the three-dimensional structure of p75ICD, comprising the p75 death domain and most of the juxtamembrane region. Recently, the nuclear magnetic resonance (NMR) structure of the Fas death domain was reported (Huang et al., 1996). We present here the NMR structure of the p75 death domain which, unexpectedly, shows significant differences to the death domain of Fas. Our structural data
on the full-length p75ICD suggest novel mechanisms of signaling for proteins containing a subtype 2 death domain as in p75.

Results and discussion

Solution structure of p75ICD

Different linewidths were observed for the $^1$H NMR signals of p75ICD, indicating substantially increased mobility on a subnanosecond timescale for much of the polypeptide chain. A quantitative measurement of the transverse relaxation of the amide protons showed that residues 336–418 were structurally rigid, whereas the first 55 and the last six amino acid residues of p75ICD were highly flexible (Figure 1). The flexible N- and C-terminal residues of p75ICD retained their mobility also in the presence of high osmolyte concentration (2 M glycine) which, in other examples, protects protein conformations against denaturation (Matthews and Leatherbarrow, 1993). The conformationally structured segment comprising residues 336–416 represents the death domain of p75ICD (Feinstein et al., 1995).

The NMR structure of the death domain of p75ICD (Table I) contains six helices (Figure 2). Their approximate lengths are from residues 343 to 351, 356 to 363, 368 to 375, 380 to 390, 396 to 406 and 409 to 416. The helices are arranged in two perpendicular sets of three helices each, with approximately parallel helix axes within each set. The N- and C-terminal polypeptide segments of the death domain are close together. A search of the Brookhaven protein data bank (Bernstein et al., 1977) with the program DALI (Holm and Sander, 1993) did not reveal any other protein or protein domain of similar three-dimensional structure. Thus, the only known structure of similar topology is represented by the recently described death domain of the Fas antigen (see below), the coordinates of which are not yet accessible in the database. The death domain of p75 is a small globular structure of ~25–30 Å diameter in all dimensions. The side chains of the charged residues are more or less uniformly distributed over most of its surface, with the exception of a surface area facing the viewer in the orientation shown in Figure 2. This part of the domain is largely devoid of charged residues and therefore is the most plausible contact area for putative hydrophobic interactions with as yet unknown downstream signaling proteins.

The conformation of the death domain of p75ICD was found to be not very stable. Only a few amide protons of helix 5 were significantly protected from exchange with the solvent in experiments in which lyophilized p75ICD was dissolved in D$_2$O and the $^1$H NMR signals of the amide protons monitored (data not shown). The limited stability of the death domain is also indicated by the onset of denaturation at pH 5 and below.

Structural comparison with other death domains

The NMR structure of the death domain of p75 is the first example of a subtype 2 death domain (Feinstein et al., 1995). Compared with the NMR structure recently described for the subtype 1 death domain of the Fas antigen (Huang et al., 1996), the death domain of p75 shows a number of important similarities and differences which allow the delineation of the conserved structural features of death domains.

The total number of helices (six) and the overall fold of the death domain is similar in p75 and in Fas. Yet, while the relative orientations of helices 2–6 are conserved, helix 1 in p75ICD is almost perpendicular to the orientation of helix 1 in the Fas death domain. This helix lies about parallel to helices 2, 3 and 4 in Fas, but is more closely aligned to helices 5 and 6 in p75ICD (Figure 2). The extent of this structural rearrangement is unusually large for a conserved protein domain. On the other hand, there is little sequence similarity in helix 1 between p75 and Fas (Figure 3) (Feinstein et al., 1995), and the death domain of Reaper appears to lack helix 1 altogether (Golstein et al., 1995). The amino acid sequence of helix 1 is better conserved among subtype 2 than subtype 1 death domains (Feinstein et al., 1995), suggesting that the structures of subtype 2 death domains will in general be close to the death domain structure of p75ICD.

With the exception of helices 2, 5 and 6, the helical segments are of similar lengths in the death domains of p75 and Fas (Figure 3). Helix 5 in p75ICD starts with a classical N-cap residue (Thr395) (Richardson and

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**Table I. Parameters characterizing the NMR structure of p75ICD**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
</tr>
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<tbody>
<tr>
<td>Number of distance constraints (residues 334–418)</td>
<td>914</td>
</tr>
<tr>
<td>Number of dihedral angle constraints (residues 334–418)</td>
<td>210</td>
</tr>
<tr>
<td>Dist. constraint violations $&gt;$0.1 Å (per conformer)</td>
<td>$0.55 \pm 0.67$</td>
</tr>
<tr>
<td>Dihedral angle constraint violations $&gt;$2.5° (per conformer)</td>
<td>$0.40 \pm 0.58$</td>
</tr>
<tr>
<td>Intra-protein AMBER energy (kcal/mol)</td>
<td>$-3910 \pm 59$</td>
</tr>
<tr>
<td>R.m.s.d. to the mean for N, C$\alpha$ and C' of residues 336–416</td>
<td>$0.6 \pm 0.1$</td>
</tr>
<tr>
<td>R.m.s.d. to the mean for all heavy atoms of residues 336–416</td>
<td>$1.0 \pm 0.1$</td>
</tr>
</tbody>
</table>

The NMR structure of p75ICD is represented by 20 conformers calculated with the program DIANA (Güntert et al., 1991) starting from random structures. The structures were calculated closely following the protocol of Liepinsh et al. (1994). The conformers were energy minimized with the program OPAL (Luginbühl et al., 1996).
Fig. 2. Solution structure of the death domain of rat p75ICD. (A) Ribbon representation of residues 336–416. The positions of the N-terminal residues of the helices are labeled. (B) Stereo view showing a superposition of the backbone atoms of residues 336–416 in the 20 conformers representing the NMR structure (Table I). Same orientation and labeling as in (A). (C) Stereo view of the conformer closest to the mean structure of the 20 conformers shown in (B). Residues 336–416 are shown in a heavy atom representation in the same orientation as in (A) and (B). The polypeptide backbone is drawn in purple. The following colors were used for the side chains: blue, Arg and Lys; red, Glu and Asp; yellow, Ala, Cys, Ile, Leu, Phe, Pro, Trp, Val; white, Asn, Gln, Ser, Thr, Tyr, His. Tyr366 at the site of the lpr<sup>+</sup> mutation is labeled.
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While most of the subtype 1 death domains associate with subtype 1 death domains (Huang et al., 1995) for optimum alignment of the helical secondary structure. Observed for this N-terminal segment of our p75ICD. Furthermore, rat p75 has a deletion of three amino acid residues sequence alignment used for helix 1 differs from that of Feinstein juxtamembrane domain. Interestingly, high flexibility was wide bars identify, respectively, little and significant reduction of self- mechanisms, perhaps via phosphorylation of its serine/ threonine counterparts. Binding could then involve the dissociation of these residues with the p75ICD does not seem to contain a death domain (Carter et al., 1996a). The recruitment of p75 death domains in signaling therefore appears to involve different mechanisms, perhaps via phosphorylation of its serine/threonine-rich C-terminal segment, or involvement of the juxtamembrane domain. Interestingly, high flexibility was observed for this N-terminal segment of our p75ICD structure which comprised all but the first seven amino acid residues of the polypeptide linker between the transmembrane and death domains. Fully extended, this linker would span >200 Å. Flexibility in this long linker region in vivo would make it difficult to explain the apparent capacity of p75 to be activated selectively by different ligands. It has been noted previously that p75 can signal elevated cytoplasmic levels of ceramide in cultured oligodendrocytes as well as activation of NF-κB in cultured Schwann cells only after binding to NGF, but not to other neurotrophins such as BDNF or NT-3 (Carter et al., 1996b; Casaccia-Bonnefil et al., 1996), despite the ability of all these ligands to bind p75 with comparable affinity. Clearly, the flexible juxtamembrane linker of p75ICD would effectively prevent the propagation to the death domain of any conformational differences in the extracellular domain which might be expected from the binding of different ligands. The association of intracellular factors with the flexible N-terminal domain of p75ICD could conceivably stabilize the juxtamembrane linker and allow the recruitment of the p75 death domains for downstream signaling in a ligand-specific manner. For example, the p75 death domain might be constitutively associated with intracellular factors; a first step in receptor activation after ligand binding could then involve the dissociation of these proteins. At this point, no structural information for this linker region is available for any of the receptors containing subtype 1 death domains (Huang et al., 1996). It has been noted previously, however, that deletion of most of the linker between the transmembrane and death domains did not affect signaling in p55TNFR (Tartaglia et al., 1993), indicating that this region is not critical for activation of receptors with death domains capable of self-association.

With death domain structures of subtypes 1 and 2 in hand, the effects of mutations in death domains can be rationalized. A systematic mutational analysis has been performed with the TRADD death domain (Park and Baichwal, 1996). Sequence alignment of p75, Fas and TRADD death domains (Figure 3) shows insertions in

Richardson, 1988) which is highly conserved in most other death domains except that of Fas (Feinstein et al., 1995). The N-terminal extension of helix 5 may thus be specific to the Fas receptor. Similarly, helix 6 extends much further towards the C-terminus in Fas than in p75ICD, which has instead a flexible C-terminal segment rich in serine and threonine (Figures 1 and 3).

Despite the structural variability observed between death domains of different subtypes, the hydrophobic core appears to be built by homologous residues. Using the sequence alignment of Feinstein et al., 1995), the 19 core residues conserved in different death domains are buried in the interior of the death domain of p75ICD with side chain solvent accessibilities below 15% (Figure 4). Of the two conserved residues with the highest solvent accessibility, one is charged (Asp372 in p75ICD) and one is a classical N-cap residue for helix 5 (Thr395 in p75ICD) in almost all other death domains. About 85% of the residues with <15% side chain solvent accessibility in p75ICD are hydrophobic. Most of these residues are also hydrophobic in the other death domains (Feinstein et al., 1995).

**Functional implications**

While most of the subtype 1 death domains associate with each other and with other death domains of the same subtype, none of these death domains have been found to associate with p75ICD (Varfolomeev et al., 1996). Furthermore, we found no sign of self-association for p75ICD. Thus, the mechanism of activation of p75 appears to be different from that of Fas and p55TNFR, in which the direct association between intracellular death domains is crucial for signaling after ligand binding (Boldin et al., 1996; Muzio et al., 1996). Interestingly, the only protein isolated so far that is capable of specifically interacting with the p75ICD does not seem to contain a death domain

![Fig. 4. Solvent accessibility of the amino acid side chains in the death domain of p75. The solvent accessibilities are expressed as a percentage of their accessibilities calculated for the situation where the respective amino acid residues are located in a hypothetical poly-Gly α-helix with a fully extended side chain (Sevilla-Sierra et al., 1994). The values were averaged over the 20 NMR conformers. Hydrophobic and hydrophilic core conserved residues (Feinstein et al., 1995) are identified by large filled and open circles, respectively. The helix locations in p75ICD are indicated at the top.](image-url)
TRADD, compared with p75, in the loops between helices 3 and 4, and helices 4 and 5. The sequence of TRADD appears to be compatible with the p75 fold, in that the pattern of buried hydrophobic and solvent-accessible charged residues is conserved. The only charged residue predicted to be buried in TRADD is located in helix 1 (Asp218 in TRADD), illustrating the ambiguities involved in aligning sequences corresponding to this helix. The charged amino acid side chains of TRADD appear to be relatively uniformly distributed over the surface of the molecule, and there is no evidence for the presence of an extended hydrophobic surface area as in p75ICD (Figure 5). The same applies to the charged residues in the death domains of p55TNFR and MORT1/FADD when mapped onto the structure of p75ICD (data not shown). The mutational analysis of TRADD involved the systematic exchange of consecutive stretches of three to four residues into alanine (Park and Baichwal, 1996). Most mutations affected buried residues, which in p75ICD correspond to side chains with <15% solvent accessibility, and may therefore have disrupted the structure of the molecule. However, some of the mutations that affected the self-association of TRADD occurred in residues of presumably high solvent accessibility, in particular the loop regions between helices 1 and 2, and on either side of helix 4 (Figures 3 and 5). Because these sites are too far apart to be buried simultaneously in the interface of a homodimer, diastereotopic methylene protons and methyl groups. Distance constraints established from the pattern of buried hydrophobic and solvent-accessible and cells were harvested 5 h after induction. The cells were sonicated and...and two-dimensional (NOESY, TOCSY, DQF-COSY) NMR spectra (Ernst et al., 1991) spectra, respectively. Stereospecific assignments were obtained for 63 pairs of...The inactivating lprΔ mutation in the Fas receptor...In conclusion, the NMR structure determination of p75ICD revealed a globular death domain flanked by flexible regions and allowed the delineation of the conserved structural features of different types of death domains. In addition, our data point to possible mechanisms of p75 activation which appear to be distinct from those previously described in receptors containing subtype 1 death domains such as p55TNFR and Fas. The p75ICD structure will now facilitate the design of decisive mutational studies to unravel mechanisms of p75 activation and proximal signaling events.

Materials and methods

Preparation of p75ICD

A DNA fragment corresponding to the intracellular domain of the neurotrophin low affinity receptor p75 was amplified by PCR from a plasmid containing the rat p75 cDNA, using primers that had Sall restriction sites to aid subcloning, and an N-terminal methionine for cytochrome c (Cmbr) cleavage. This fragment was inserted into a pQE31 vector (Quagen) which added on a histidine tail (His-tag) to facilitate purification. This construct produced a fusion protein containing 20 extra amino acids due to the linker sequence and the His-tag. The fusion protein was expressed in the Escherichia coli strain M15(pREP4) (Quagen) grown in 20 l of modified M9 minimal medium (containing 10% NH4Cl as the sole nitrogen source) at 37°C. Expression was induced with 1 mM isopropyl-β-D-thiogalactopyranoside (IPTG) at OD600 = 0.8, and cells were harvested 5 h after induction. The cells were sonicated...and...The inactivating lprΔ mutation in the Fas receptor (Val238Asn, corresponding to Tyr366 in p75) has also been shown to interfere with homo- and hetero-association of the death domains of p55TNFR and MORT1/FADD (Varfolomeev et al., 1996). This residue defines the spacing between helices 2/3 and 4/5 in the death domain (Figure 2C) (Huang et al., 1996). Since different death domains have different residues at this position, this spacing is likely to be different even between death domains of the same subtype. Interestingly, the signaling capacity of TRADD was hardly affected when this residue was mutated to alanine (Park and Baichwal, 1996).

In conclusion, the NMR structure determination of p75ICD revealed a globular death domain flanked by flexible regions and allowed the delineation of the conserved structural features of different types of death domains. In addition, our data point to possible mechanisms of p75 activation which appear to be distinct from those previously described in receptors containing subtype 1 death domains such as p55TNFR and Fas. The p75ICD structure will now facilitate the design of decisive mutational studies to unravel mechanisms of p75 activation and proximal signaling events.

Fig. 5. Mapping of the amino acid sequence of TRADD onto the death domain of p75. The death domain of p75 is drawn as a ribbon highlighting the Cα positions of those residues which are positively (blue) or negatively (red) charged in TRADD. Loop regions in p75ICD corresponding to sites in TRADD where mutations to alanine suppressed the self-association of the TRADD death domain (Park and Baichwal, 1996) are shown in magenta (see also Figure 3).
References


Liepinsh,E., Berndt,K.D., Sillard,R., Mutt,V. and Otting,G. (1994) Involvement of MACH, a novel MORT1/FADD-interacting protease, and Nicolas Gilles for advice on 15N-labeling. NMR characterization of p75ICD during the initial stages of the project. We thank Drs William Griffiths and Leodevico Llag for molecular weight determinations using mass spectroscopy and gel filtration, respectively, Dr Judith Murray-Rust for comments on the manuscript, Dr Moses Chao for a p75ICD plasmid construct and antisem used in preliminary work, Ella Cedrelund for N-terminal sequencing, Ann-Sofie Nilsson for technical assistance, and Nicolaus Gilles for advice on 15N-labeling. NMR spectra at 800 MHz were recorded at the European large scale facility for biomolecular NMR at the University of Frankfurt. This work was supported by grants from the Swedish Natural Science Research Council, the Swedish Medical Research Council and the Biotechnology Programme of the European Commission.

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