



Axon-guidance defects in a *Drosophila* MICAL loss-of-function (LOF) mutant. Courtesy of Alex Kolodkin, Johns Hopkins School of Medicine, Baltimore, USA.

## AXON GUIDANCE

## MICAL — a missing link

The semaphorins are important axon-guidance molecules in invertebrates and vertebrates, and they can cause attraction or repulsion. The repulsive response is mediated by the plexin family of receptors, but little is known about how the plexins signal to the cytoskeleton to induce growth cone collapse. Now, a new study implicates a family of oxidoreductases, the MICALs, as a missing link in this pathway.

The identification of the vertebrate MICAL-1 (molecule interacting with CasL) was reported earlier this year, and now Terman *et al.* describe the isolation of a *Drosophila* counterpart, and of vertebrate MICAL-2 and -3. MICALs are large cytosolic proteins that consist of several domains. In the *Drosophila* embryo, MICAL shows a similar expression pattern to Plexin A (PlexA) on central nervous system and motor axons, and the authors showed that it interacts physically with PlexA. MICAL also contains an actin-binding domain and an intermediate-filament-binding region, so it could provide a direct link between the plexins and the cytoskeleton. Vertebrate MICAL-1 interacts with CasL, and as Cas-family proteins interact with signalling proteins that influence neuronal morphology, MICAL could link plexins to cytoskeletal regulators through Cas proteins.

Terman *et al.* looked for evidence that MICAL acts in the same pathway as Semaphorin 1a (Sema-1a) and PlexA. They found that MICAL loss-of-function caused the same axon-guidance and defasciculation defects as were seen in PlexA and Sema-1a mutants, and overexpression of MICAL had the same effect as overexpression of PlexA. Further evidence for a common pathway came from

embryos that were doubly heterozygous for mutations in MICAL and PlexA or Sema-1a, which showed similar axon-guidance defects to Sema-1a;PlexA double heterozygotes.

Which domains in MICAL are important for its function in semaphorin-mediated repulsion? The amino terminus is highly conserved between species, and this region contains a flavoprotein monooxygenase domain, which catalyses an oxidation–reduction (redox) reaction that results in the insertion of an oxygen atom into a substrate. Altering key amino-acid residues within this domain that are conserved within this enzyme class and are required for their activity compromises MICAL function in motor axons *in vivo*. In an *in vitro* rat growth cone assay, Terman *et al.* showed that semaphorin-mediated axonal repulsion can be inhibited by selective flavoprotein monooxygenase inhibitors.

So, these findings not only tell us more about the signalling pathway that underlies semaphorin-mediated repulsion, but in combination with recent data that implicate 12/15-lipoxygenase in semaphorin signalling, they also provide support for the involvement of redox reactions in this pathway. Oxidation of actin is known to promote its depolymerization, so it will be interesting to see whether redox reactions are a common feature of repulsive neuronal guidance.

Heather Wood

### References and links

**ORIGINAL RESEARCH PAPER** Terman, J. R. *et al.* MICALs, a family of conserved flavoprotein oxidoreductases, function in plexin-mediated axonal repulsion. *Cell* **109**, 887–900 (2002)

**FURTHER READING** Finkel, T. Redox-dependent signal transduction. *FEBS Lett.* **476**, 52–54 (2000) | Tamagnone, L. & Comoglio, P. M. Signalling by semaphorin receptors: cell guidance and beyond. *Trends Cell Biol.* **10**, 377–383 (2000)

## PRION PROTEINS

## Protector prions

Two papers published in *EMBO Journal* hint at a neuroprotective function for the cellular prion protein, PrP<sup>C</sup>. Mutations in this protein, or changes in conformation induced by the infective prion protein PrP<sup>Sc</sup>, can cause spongiform encephalopathies such as scrapie, bovine spongiform encephalopathy and Creutzfeldt–Jakob disease.

However, the function of PrP<sup>C</sup> is still a mystery. It is not even clear whether the neurodegeneration that is associated with spongiform encephalopathies is caused by a loss of the normal function of PrP<sup>C</sup> or by a gain of toxic function by PrP<sup>Sc</sup>. The apparent absence of a phenotype in prion-knockout mice argued for the latter, but evidence is accumulating that cellular prions might have a neuroprotective role, the loss of which could cause cell death in prion diseases.

The two new papers both come from the same research group. One, by Zanata *et al.*, shows that the cellular prion binds to a heat-shock-related protein, stress-inducible protein 1 (STI1), and that the interaction between these two proteins at the cell surface can rescue cultured retinal cells from apoptosis induced by treatment with anisomycin. A functional role for this interaction is supported by the finding of both proteins on neuronal cell surfaces in the central nervous system. The second paper, by Chiarini *et al.*, shows that binding of PrP<sup>C</sup> by a peptide that recognizes the STI1-binding site (known as PrR) also protects neurons against apoptosis *in vitro*. It goes further to demonstrate that this protection depends on an increase in the levels of cyclic AMP that activates protein kinase A.

Many questions are outstanding. For example, it is unclear how the binding of PrP<sup>C</sup> at the outer membrane activates adenylyl cyclase, which is normally regulated by G proteins on the inner membrane. And it remains to be shown that the neuroprotective effect of PrP<sup>C</sup>–STI1 binding is physiologically relevant *in vivo*, or that it is disrupted by infection with PrP<sup>Sc</sup>. But a fuller understanding of signalling and neuroprotection by PrP<sup>C</sup> should help us to understand the pathogenesis of prion disorders as well as the underlying cell biology of these proteins.

Rachel Jones

### References and links

**ORIGINAL RESEARCH PAPER** Zanata, S. M. *et al.* Stress-inducible protein 1 is a cell surface ligand for cellular prion that triggers neuroprotection. *EMBO J.* **21**, 3307–3316 (2002) | Chiarini, L. B. *et al.* Cellular prion protein transduces neuroprotective signals. *EMBO J.* **21**, 3317–3326 (2002)

