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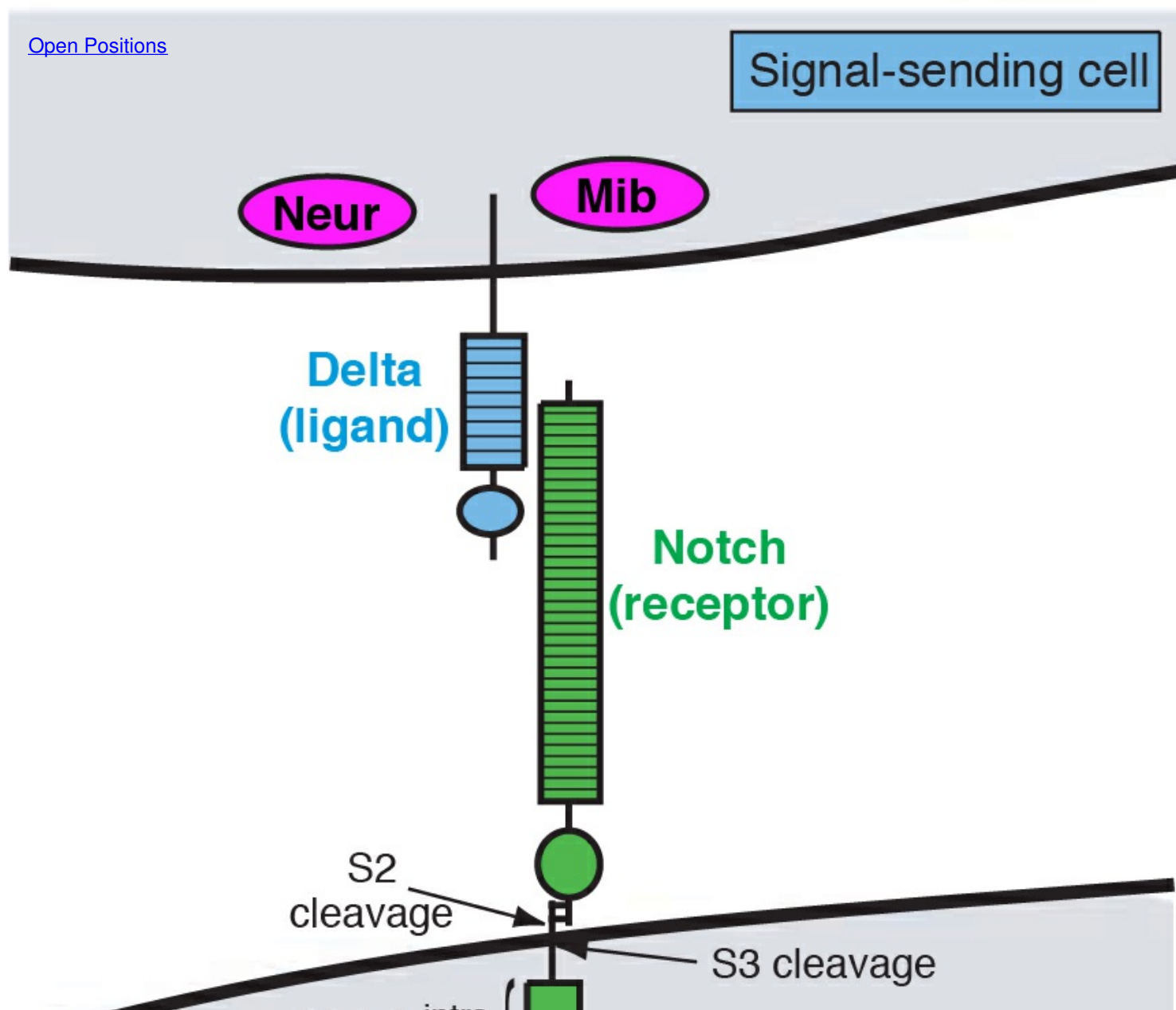
Notch Signaling and Neural Transcription Factors

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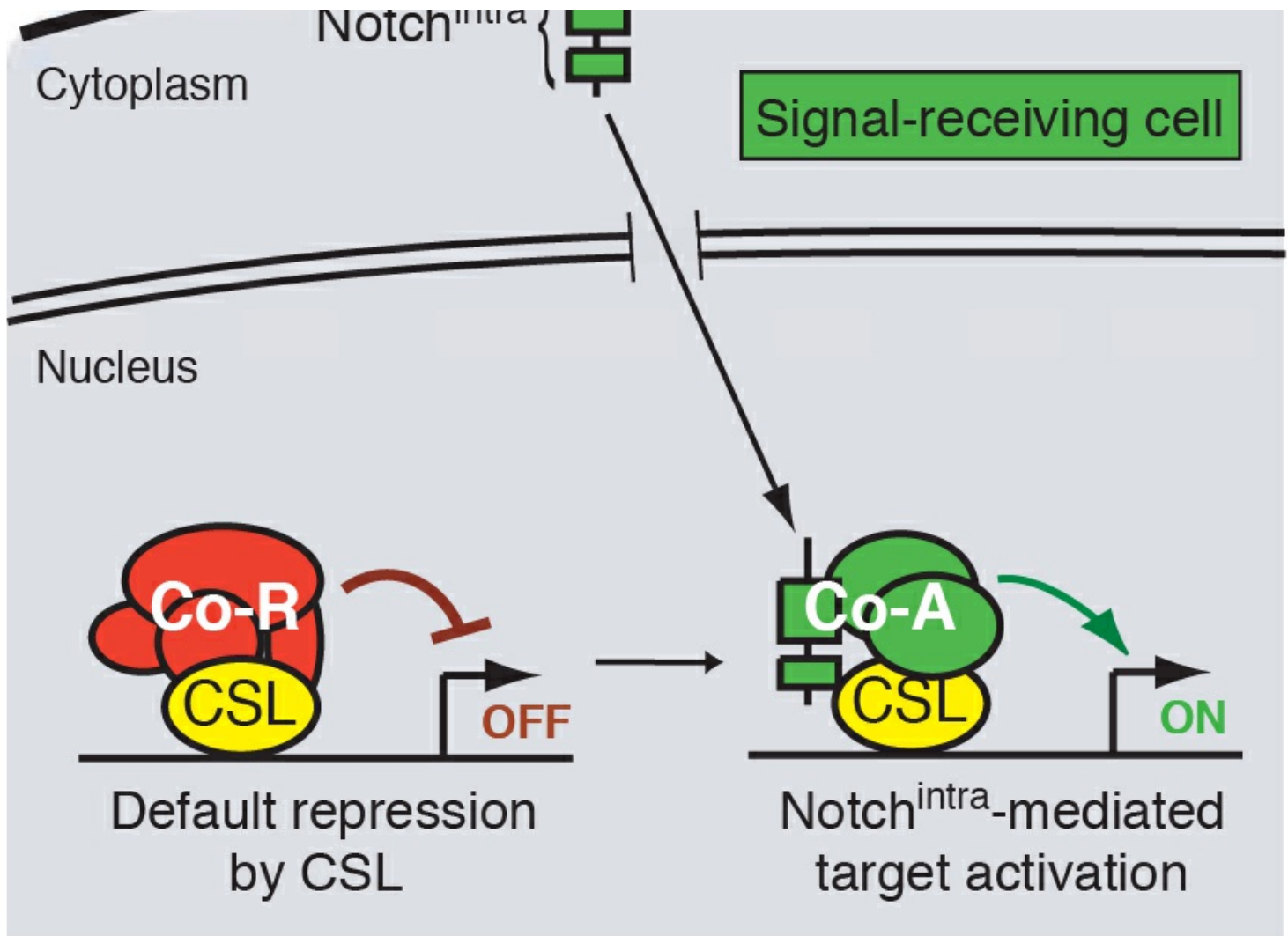


Figure 1 — A highly simplified model of Notch signaling. Neur and Mib are E3 ubiquitin ligases that activate DSL ligands. Interaction between the extracellular domains of DSL and Notch triggers cleavage and release of the Notch intracellular domain (Nintra).

Notch signal transduction

Coordinated development requires that cells communicate with each other. This is made possible by molecular mechanisms of cell-cell signaling, by which cells can influence each other's fate and behavior. A cell signaling mechanism of fundamental importance to animal development is the Notch pathway (Figure 1). It may be safely said that Notch signaling is required, in some reasonably direct fashion, for the development of most tissues in all animal species. As such, misregulation of Notch signaling underlies a variety of human diseases and cancers.

Many ways to use a pathway

We focus on a few developmental settings that are paradigms of Notch-regulated events. One broad category is "inhibitory" signaling, whereby Notch prevents a cell from adopting a particular cell fate, usually through the repression of celltype-specific determinants. For example, Notch signaling singles out individual neural precursors from clusters of cells with neural potential. Loss of Notch signaling therefore causes excess neural differentiation. A second broad category is "inductive" signaling, whereby activation of Notch induces the expression of molecules with tissue-organizing activity. For instance, Notch signaling specifies the wing margin, a line of cells that organizes the patterning and outgrowth of the wing proper. Loss of Notch signaling here results in loss of wing tissue. We have utilized both inhibitory and inductive signaling events to study the roles of different Notch pathway factors, and how they regulate the activity of resident tissue and cell identity genes.

BEN proteins are novel CSL co-repressors in the Notch pathway

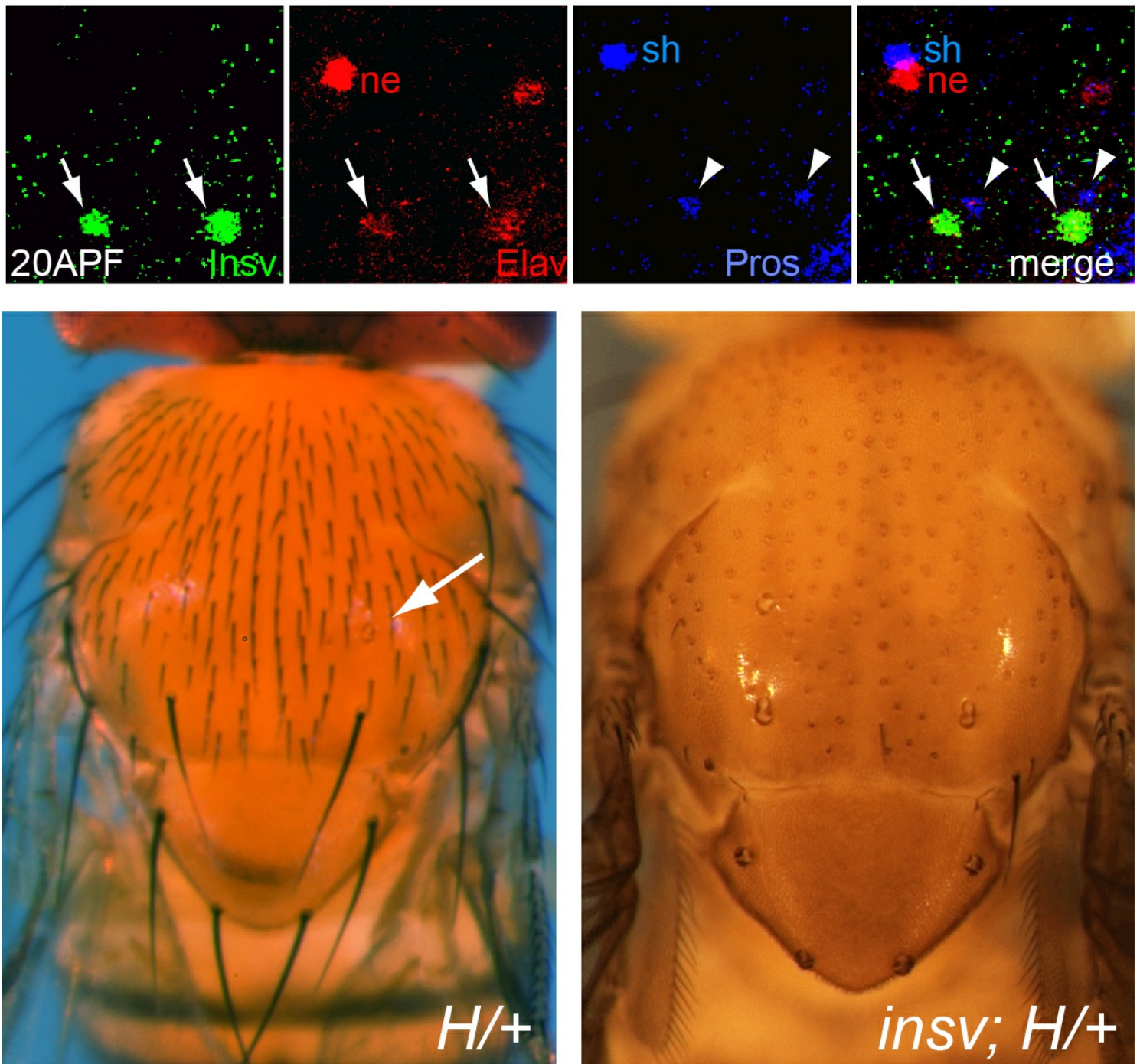


Figure 2 — Insensitive (*Insv*) is a novel Notch antagonist. (Top) Immunostaining of pupal notum sensory organs shows *Insv* (green) is a nuclear factor that is an early neuronal antigen, as seen by coexpression with neural marker *Elav* (red) an exclusion with sheath marker *Prospero* (blue). As neurons mature and acquire full *Elav* levels, *Insv* disappears. (Bottom) *Insv* antagonizes Notch-dependent cell fate. *Hairless* (*H*) is a classical Notch repressor, and heterozygotes exhibit a mild double socketting phenotype (arrow) indicating shaft-to-socket cell fate conversion. Mutation of *insv* in this background results in complete double socketting of all sensory bristle organs.

In the course of studying *Drosophila* novel factors that regulate neural development, we elucidated the molecular genetic function of Insensitive (*Insv*), an anonymous protein whose only annotated domain was of unknown function, the BEN domain. We showed that *Insv* is a nuclear factor that inhibits Notch signalling during multiple peripheral nervous system cell fate decisions. A direct role for *Insv* in transcriptional repression was indicated by binding of *Insv* to CSL transcription factor Su(H), and by strong chromatin

immunoprecipitation of endogenous Insv to most E(spl)-C loci. Interestingly, we later found that the mammalian BEN domain factor BEND6 shares the functional attributes of Insv, even though they have only modest sequence similarity. BEND6 binds the mammalian CSL protein CBF1 and antagonizes Notch-dependent target activation. In vivo, BEND6 inhibited Notch-mediated self-renewal of neocortical neural stem cells and promoted neurogenesis. Thus, BEN domain factors represent a conserved type of Notch co-repressor in the nervous system.

The BEN domain reveals a new DNA binding domain

Upon using ChIP-sequencing to examine the extent of cobinding of Insv and Su(H) genomewide, we observed extensive association of Insv to sites bearing a novel palindromic motif. We validated binding of endogenous Insv to genomic regions bearing such sites, whose associated genes are enriched for neural functions and are functionally repressed by Insv. Unexpectedly, we found that the Insv BEN domain binds specifically to this sequence motif and that Insv directly regulates transcription via this motif. Together with Dinshaw Patel's lab (SKI), we determined the crystal structure of the BEN–DNA target complex, revealing homodimeric binding of the BEN domain and extensive nucleotide contacts via a helices and a C-terminal loop. We further demonstrate DNA-binding and repression activities by the mammalian neural BEN-solo protein BEND5, and continued with the molecular and structural elucidation of other *Drosophila* BEN domain factors. Altogether, we define novel DNA-binding activity in a conserved family of transcriptional repressors, opening a molecular window on this extensive gene family.

Insv-BEN domain homodimer bound to its DNA target

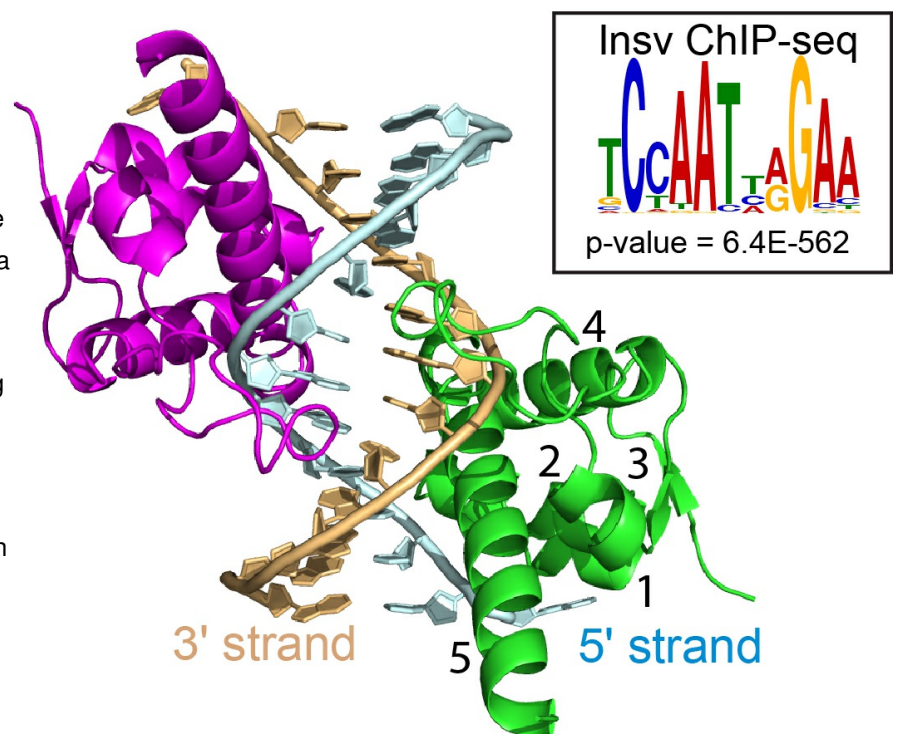


Figure 3 — Insv contains BEN, a prototype of a new type of DNA binding domain.

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