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Regulation of ligand-independent notch signal through intracellular trafficking

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Abbreviations: ESCRT, endosomal sorting complex required for transport; MVBs, multivesicular bodies

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Notch signaling is an evolutionarily conserved mechanism that defines a key cell fate control mechanism in metazoans. Notch signaling relies on the surface interaction between the notch receptor and membrane bound ligands in an apposing cell. In our recent study,22 we uncover a non-canonical receptor activation path that relies on a ligand-independent, intracellular activation of the receptor as it travels through the endosomal compartments. We found that notch receptor, targeted for degradation lysosomal degradation through multivesicular bodies (MVBs) is “diverted” toward activation upon mono-ubiquitination through a synergy between the ubiquitin ligase Deltex, the non-visual β-arrestin Kurtz and the ESCRT-III component Shrub. This activation path is not universal but appears to depend on the cellular context.

Regulation of Notch Trafficking

The notch pathway acts throughout development to link fate choices of a cell to those of its immediate cellular neighbors, ultimately affecting proliferation, apoptosis and differentiation.1-3 Notch malfunction has been associated with aberrant development in all metazoans and with various human diseases.4 The dosage of the notch signal defines an extraordinarily sensitive parameter that regulates the developmental outcome of signaling. Hence, cellular mechanisms controlling the dosage of activated notch receptors are of importance for the biology and pathobiology of the pathway.

Numerous studies, mostly in flies, implicated intracellular trafficking of notch receptor in both ligand-dependent and ligand-independent notch signaling.5-13 Endocytosis of notch may result in either down- or upregulation of signaling depending on how the receptor is sorted within the cell.14,15 Although these studies indicated an important role for intracellular trafficking in notch signaling, the compartments and molecular mechanisms underlying notch activation—one the receptor enters the endocytic path—remain unclear. Available evidence suggests that there may be several mechanisms that can lead to intracellular notch activation, either in a ligand-dependent or a ligand-independent manner. The process capable of modulating such intracellular notch signaling remains unknown, but ligand-independent activation of the receptor has been recently shown to be essential for the normal development of Drosophila blood cells.16

An important element of the endosomal sorting machinery is defined by the ESCRT (endosomal sorting complex required for transport) system.17 ESCRT is crucial in mediating the various steps leading to the sorting of membrane proteins into multivesicular bodies (MVBs) on their way to lysosomal degradation. It is also required for the morphogenesis of intraluminal vesicles and for the sorting of ubiquitinated cargo into these vesicles.17 As elaborated below, we identified an ESCRT member as a notch signal modifier. The ESCRT system consists of ESCRT-0, -I, -II, -III, and Vps4.17 ESCRT-0, -I, -II contain ubiquitin-binding domains, and are primarily
involved in cargo sorting and in the recruitment and activation of ESCRT-III. The ESCRT-III subunits Vps20, Vps32 (Shrub), Vps24, Vps4 are assembled in this order, and contribute to the budding and scission of intraluminal vesicles into the vesicles. Vps4 is the ATPase required for ESCRT-III disassembly. Loss of ESCRT function modulates notch trafficking and leads to ectopic activation of notch signaling. The mechanisms underlying these events are complex, as different members of the ESCRT system exhibit distinct, non-overlapping phenotypic characteristics. For instance, the Vps22, Vps25, Vps36 members of ESCRT-II display non-identical mutant phenotypes involving notch activity, proliferation, or apoptotic resistance. Moreover, there may be tissue specific functions of each component of ESCRTs.

Shrub is a Key Modulator of the Notch Signal Mediated by Deltex/Kurtz

We previously reported interactions between the non-visual β-arrestin Kurtz and the ubiquitin ligase Deltex leading to the regulation of notch trafficking, its degradation and consequential loss of signaling. We identified Shrub, a member of the ESCRT-III complex, as a major modulator of Deltex/Kurtz on notch signaling. Our genetic and molecular studies indicate that absence of Shrub suppresses the classic loss-of-notch wing-nicking phenotype associated with notch degradation due to the co-expression of Deltex and Kurtz. Use of Cut, a direct downstream target of notch signals, as a readout of notch activation revealed that its activation by Deltex can be suppressed by coexpression of Shrub. Expression of Shrub also amplifies the inhibiting effect of Kurtz on notch. But when Shrub activity is inhibited in a Deltex overexpressing background, we found a striking concomitant notch activation. This activation does not depend on the presence of the ligands for Delta or Serrate. In addition, biochemical analyses demonstrated that this activation was coupled with an accumulation of mono-ubiquitinated notch. ESCRT-III, through Shrub, seems to play a distinct role in these phenomena, as ESCRT-I and -II components are not required for this Deltex-dependent notch activation. Consistent with this, mutants of ESCRT-I or ESCRT-II did not suppress the wing phenotype caused by the degradation of notch associated with the overexpression of Deltex and Kurtz. Therefore, our data suggest a check-point by ESCRT-III and not ESCRT-I or -II in the ligand-independent and Deltex/Kurtz-dependent regulation of notch signaling.

Context Specific Regulation of the Notch Signal

A characteristic of the unconventional non-canonical activation of notch signaling we uncovered, is that it is context specific.
dependent. In the wing disc for example, the notch signal induced by Deltex overexpression is seen mostly in the ventral region.2,22 Similarly, we found that the notch signal regulated by deltex-shrub is only manifested in the ventral region of the wing disc, indicating that this ligand-independent, intracellular activation of notch, depends on the cellular context.22 This context specificity is not associated with the glycosyltransferase Fringe (unpublished data), previously shown to regulate the differential activity of the notch receptor in the ventral vs. dorsal region of the wing disc.23 However, treatment with the lysosomal inhibitor chloroquine abolishes this context specificity.22 The mechanism underlying the involvement of lysosomal degradation in this context dependent response is unclear but certainly noteworthy and the subject of future inquiries.

A Model for the Intracellular Activation of Notch

On the basis of these observations, we propose a model for Shrub-Deltex-Kurtz dependent modulation of notch activity (Fig. 1). Shrub, Deltex, and Kurtz regulate the trafficking of notch, and modulate the degradation of notch in the late-endosome/MVBs compartment. In the presence of Shrub, notch is sorted to the degradation pathway, resulting in downregulating the notch signal. In the absence of Shrub and/or Kurtz, Deltex promotes mono-ubiquitination, leading to notch activation. Previous studies have indicated that there may be more than one mechanism that can potentially activate notch during its endosomal sorting.5-13 Our contribution is based on linking notch activation to its ubiquitination state, modulated by the synergy of Deltex, Kurtz and Shrub. What remains unclear is the contribution of such a ligand-independent, context-dependent, intracellular activation in the normal biology and in the pathobiology of the notch signaling pathway.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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