The synaptic vesicle SNARE neuronal Synaptobrevin promotes endolysosomal degradation and prevents neurodegeneration

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Introduction

Regulated SNARE-mediated membrane fusion underlies a plethora of intracellular trafficking, sorting, and exocytosis and has been particularly well studied in neurotransmitter release (Südhof and Rothman, 2009). The synaptic v-SNARE n-syb (neuronal Synaptobrevin)/VAMP2 is a key protein required for synaptic vesicle exocytosis. In this paper, we report that loss of n-syb caused slow neurodegeneration independent of its role in neurotransmitter release in adult Drosophila melanogaster photoreceptor neurons. In addition to synaptic vesicles, n-Syb localized to endosomal vesicles. Loss of n-syb lead to endosomal accumulations, transmembrane protein degradation defects, and a secondary increase in autophagy. Our evidence suggests a primary defect of impaired delivery of vesicles that contain degradation proteins, including the acidification-activated Cathepsin proteases and the neuron-specific proton pump and V0 adenosine triphosphatase component V100. Overexpressing V100 partially rescued n-syb-dependent degeneration through an acidification-independent endosomal sorting mechanism. Collectively, these findings reveal a role for n-Syb in a neuron-specific sort-and-degrade mechanism that protects neurons from degeneration. Our findings further shed light on which intraneuronal compartments exhibit increased or decreased neurotoxicity.

In addition to the synaptic vesicle cycle, neurons use specialized membrane trafficking (Chan et al., 2011) and a neuronal intracellular degradation mechanism during development and adult maintenance (Williamson et al., 2010a,b). Loss of endolysosomal or autophagosomal degradation is sufficient to cause adult-onset degeneration in Drosophila photoreceptors (Lee and Montell, 2004; Xu et al., 2004; Akbar et al., 2009; Williamson et al., 2010a) and neurons in general (Hara et al., 2006; Komatsu et al., 2006; Nixon et al., 2008). Autophagy and the canonical endolysosomal system are considered ubiquitous cellular membrane trafficking machinery that may serve specialized or increased functions in neurons. In addition to the ubiquitous degradation machinery, we recently reported that a neuronal vesicle ATPase (v-ATPase) component provides a neuron-specific intracellular degradation mechanism; loss of v100, the neuron-specific v-ATPase subunit a1, leads to defects in brain...
wiring (Williamson et al., 2010a) and adult-onset degeneration (Williamson et al., 2010b) in Drosophila. To our knowledge, no other neuron-specific membrane trafficking protein required for intracellular degradation has so far been characterized.

We have previously reported that loss of n-syb in the Drosophila visual system leads to fine structural synaptic defects that have an onset before synapse formation (Hiesinger et al., 1999). Recent work has shown that v100 behaves similarly to n-syb. Specifically, loss of either protein leads to neurotransmitter release defects, and both proteins directly interact with the t-SNAREs Syntaxin and SNAP-25 at the synapse (Hiesinger et al., 2005). These observations give rise to the idea that both proteins might exert related functions in endolysosomal trafficking in addition to their roles in neurotransmitter release. Here, we show that loss of n-syb causes intracellular degradation defects that lead to neurodegeneration in adult photoreceptor neurons. n-Syb functions in concert with V100 in a neuronal sort-and-degrade mechanism that is required for neuronal maintenance independent of their roles in neurotransmitter release.

**Results**

**Loss of n-syb causes slow degeneration in adult photoreceptor neurons**

We previously reported remarkable similarities between the v-SNARE n-Syb and the v-ATPase component V100 during synaptic development (Hiesinger et al., 1999; Williamson et al., 2010b) and in synaptic function (Hiesinger et al., 2005). In addition, we have shown that v100 is required to maintain neurons and that this role is independent of its function on synaptic vesicles (Williamson et al., 2010a). To investigate a possible similar function of n-Syb in maintaining neurons, we investigated the morphology and function of n-syb mutant photoreceptor neurons over time compared with photoreceptors mutant for v100. We used the ey3.5FLP method to selectively generate n-syb mutant photoreceptors in otherwise heterozygous animals (Chotard et al., 2005; Mehta et al., 2005). To assay degeneration morphologically, we assessed the loss of rhabdomeres, actin-rich extensions of the photoreceptor apical membrane that contain the light-sensitive Rhodopsin (Harris et al., 1976). To assay degeneration functionally, we recorded electroretinograms (ERGs), extracellular recordings that measure the response of photoreceptors to a light stimulus (Heisenberg, 1971). As shown in Fig. 1, adult n-syb mutant photoreceptors exhibited slow neurodegeneration as indicated by a significant loss in the number of intact rhabdomeres and ERG depolarization in 1- and 5-wk-old flies compared with controls (Fig. 1, A, C, F, and H). The progressive reduction in the ERG response amplitude is accelerated by exposure to constant light and rescued by photoreceptor-specific expression of wild-type n-syb (Fig. 1, D and I). The degeneration phenotype is similar to, but milder than, the phenotype observed for v100 (Fig. 1, B and G). We further compared loss of n-syb or v100 to loss of syt (synaptotagmin), which plays critical roles in both synaptic vesicle exocytosis and endocytosis in Drosophila photoreceptors (Littleton et al., 1993; Reist et al., 1998). In contrast to n-syb and v100, photoreceptors lacking syt exhibited no signs of functional or morphological degeneration, indicating that defective synaptic vesicle cycling does not reduce photoreceptor viability (Fig. 1, E and J–L). Similarly, loss of neurotransmission in a mutant for histamine (hdc<sup>j910</sup>; Burg et al., 1993), the neurotransmitter of Drosophila photoreceptors, caused neither structural nor functional degeneration (Fig. S1). Collectively, our findings reveal that loss of n-syb results in slow degeneration in a manner similar to v100. Comparison with syt and a mutant lacking a neurotransmitter further suggests that degeneration is caused by a mechanism that is independent of neurotransmitter release.

**n-Syb localizes to synaptic endosomal compartments as well as synaptic vesicles**

If Syb has a role in maintaining neurons independent of its function in neurotransmitter release, this role is likely to be independent of synaptic vesicles. At synapses, n-Syb colocalized with all synaptic vesicle markers, including Syt and V100 (Fig. 2, A–C). In addition, n-Syb colocalized to a large extent with the early endosomal markers Syx7 (Syntaxin 7)/Avalanche and Rab5 but less extensively with markers of two different endosome types, Hrs and the recycling endosomal marker Rab11, or with several late endosomal and lysosomal markers (Fig. 2, D). We further verified the immunohistochemical colocalization results using YFP-tagged overexpression constructs for Rab5, Rab7, and Rab11. n-Syb colocalized at synapses most extensively with the early endosomal marker Rab5, significantly less with the late endosomal marker Rab7, and only little with the recycling endosomal marker Rab11 (Fig. 2, D–F; quantification is shown in Fig. 2, B, green bars). The colocalization with Rab5 was most prominent at the synaptic terminals (Fig. 2, D, arrow), whereas in the cell bodies, n-Syb colocalized only partly with the endolysosomal markers analyzed here (Fig. 2, D, arrowhead). We corroborated these findings using high-resolution 3D deconvolution that allows to discern individual compartments at the resolution limit of light (Hiesinger et al., 2001). In the synaptic regions of the optic lobe, n-Syb largely colocalized with Syx7- and V100-positive compartments (Fig. 2, G, arrow). In cell bodies, clearly discernible endosomal compartments are visible after deconvolution, which are positive for different combinations of n-Syb, Syx7, V100, and Rab5 (Fig. 2, G [arrowhead] and H). These findings indicate that in addition to synaptic vesicles, n-Syb localizes to a subset of early endosomal compartments, predominantly at synapses.

**Loss of n-Syb results in accumulations of several types of undegraded intracellular membrane compartments**

We have previously reported that loss of functional n-Syb leads to intracellular accumulations of transmembrane receptors and late developmental defects in the optic lobe (Hiesinger et al., 1999). In early experiments, we used tetanus toxin light chain (TNT) expression, which specifically cleaves n-Syb but not n-Syb’s closest homologue Cellubrevin (Sweeney et al., 1995). Photoreceptor-specific expression of TNT mimics all n-syb loss-of-function phenotypes, including impaired synaptic development (Hiesinger et al., 1999) and neurotransmitter release (Sweeney et al., 1995), as well as degeneration of photoreceptors
These data show that loss of n-syb, but not syt, causes a pronounced increase in vesicle content and a several-fold increase in the profile size of synaptic terminals.

Next, we assessed the appearance of late degradative compartments, i.e., multilamellar lysosomal structures (Dermaut et al., 2005) and autophagosomal compartments as recognized by double membranes (Williamson et al., 2010a). For both TNT expression and n-syb–deficient photoreceptor terminals, we found no increase in the appearance of multilamellar lysosomal structures, which occur when degradation in lysosomes is defective (Dermaut et al., 2005; Schultz et al., 2011). However, we did find multivesicular and autophagosomal compartments that are not observed in wild type or syt (Fig. 3, B and D, arrows). These phenotypes were also similar to v100 (Williamson et al., 2010a). We further assessed the emergence of these ultrastructural defects after photoreceptor-specific TNT expression,
Figure 2. **n-Syb localizes to synaptic endosomal compartments in addition to synaptic vesicles.** (A) Intracellular compartments in a wild-type synaptic terminal. Cysteine string protein (CSP), Synaptotagmin (Syt), and the t-SNARE Syntaxin 1A (Syx) function in synaptic vesicle (SV) exocytosis. Early endosomes (EE) and endosomal vesicles (EV) are marked by Rab5 and Avalanche/Syntaxin 7 (Syx7). Early endosomes undergo degradation via multivesicular body (MVB) formation, marked by Rab7 and Hrs, and lysosomes (Ly) marked by carnation (Car), deep orange (Dor), and sunglasses (Sun). Recycling endosomes (RE) are marked by Rab11. (B) Colocalization of 16 markers with n-Syb in developing photoreceptor terminals at ~P+30%. Shown are the percentiles of individually distinguishable compartments for immunolabeling with 13 antibodies (black bars) that are positive for n-Syb coimmunolabeling. Green bars show similar quantification of n-Syb colocalization for compartments marked by YFP-tagged Rab5, Rab7, and Rab11. (C–H) Immunolabeling of photoreceptor terminals at ~P+30%. (C) Immunolabeling for n-Syb, Syx7, and Hrs (green). Single channels are shown in C–C‴. (D–F) Colabeling of n-syb-Gal4>YFP-Rab5 (D),
and wild-type terminals precisely, we generated 50% mutant n-lys mosaics (Fig. 4 A). The early endosome marker Syx7 in particular revealed a striking image of the photoreceptor terminal field with many substantially enlarged terminals and little to no recognizable cartridge structure (Fig. 4, B and F). The often considerably enlarged terminals are consistent with our ultrastructural observations for both TNT expression and loss of n-syb and were not observed for v100. All markers of the endolysosomal system analyzed here were significantly up-regulated, including Syx7, Rab5, Rab7, and Hrs (Fig. 4, C, D, and G; and Fig. 5, A and E). The transmembrane receptor Fas2, which we have previously shown is strongly up-regulated in n-syb photoreceptors (Hiesinger et al., 1999), colocalizes evenly to Syx7 in the enlarged n-syb terminals, suggesting that Fas2 accumulates on Syx7-positive endosomal vesicles (Fig. 4 B). The up-regulation of endosomal markers was further verified with immunoblot analysis of protein extract from complete eyes for Syx7 and Hrs (Fig. 4 H). In contrast from EM observations of terminals at 50 and 75% of pupal development (P+50% and P+75%). An increased overall vesicle density caused the appearance of darkened photoreceptor terminals as early as P+50% (Fig. S3). In addition, terminals at P+75% exhibited numerous larger vesicular structures that have the appearance of multivesicular bodies but lack an obvious double-membrane envelope. Together, these data indicate that intracellular vesicle accumulations at n-lys mutant terminals start during development, with an increase of small vesicle density before the emergence of multivesicular bodies before eclosion and autophagosomes in the adult.

Loss of n-syb results in accumulations of endosomal vesicles

To assess the molecular nature of the intracellular accumulations observed with EM, we performed wild-type colocalization analyses. To assess even subtle level changes between mutant and wild-type terminals precisely, we generated 50% mutant n-syb mosaics (Fig. 4 A). The early endosome marker Syx7 in particular revealed a striking image of the photoreceptor terminal field with many substantially enlarged terminals and little to no recognizable cartridge structure (Fig. 4, B and F). The often considerably enlarged terminals are consistent with our ultrastructural observations for both TNT expression and loss of n-syb and were not observed for v100. All markers of the endolysosomal system analyzed here were significantly up-regulated, including Syx7, Rab5, Rab7, and Hrs (Fig. 4, C, D, and G; and Fig. 5, A and E). The transmembrane receptor Fas2, which we have previously shown is strongly up-regulated in n-syb photoreceptors (Hiesinger et al., 1999), colocalizes evenly to Syx7 in the enlarged n-syb terminals, suggesting that Fas2 accumulates on Syx7-positive endosomal vesicles (Fig. 4 B). The up-regulation of endosomal markers was further verified with immunoblot analysis of protein extract from complete eyes for Syx7 and Hrs (Fig. 4 H). In contrast
to endolysosomal proteins, synaptic vesicle markers were unaltered or slightly down-regulated, similar to observations previously made for v100 mutant photoreceptor terminals (Fig. 4 G; Williamson et al., 2010a). Interestingly, V100 itself is mostly lost at n-syb mutant synapses (Fig. 4 C). These findings suggest that the accumulated vesicles observed in n-syb mutant terminals with EM do not represent either synaptic vesicles or vesicles lacking the normal complement of synaptic vesicle proteins.
Collectively with the wild-type localization data, these observations suggest that n-Syb has a hitherto unrecognized function in endolysosomal progression.

Next, we directly compared the endosomal defects in n-syb and v100 mutant synaptic terminals. Immunolabeling of whole laminae reveals that synaptic accumulations of the early endosomal proteins Rab5, Syx7, and the transmembrane receptor Chaoptin (Fig. 5, A, C, and E) are substantially increased in n-syb compared with v100. In contrast, the recycling endosomal marker Rab11 is not noticeably altered in either mutant. Interestingly, degeneration is more severe in the v100 mutants (Fig. 1), yet the endosomal accumulations and swelling of photoreceptor terminals are considerably more severe in the n-syb mutant. These findings suggest that the severity of degeneration depends on the type of endolysosomal defect and is not simply a function of the amount of endosomal vesicle accumulation.

Many enlarged n-syb mutant terminals have a diameter of 5 µm or more, large enough to permit an analysis of the subcellular distribution of intracellular compartment markers. As shown in Fig. 4 (E and F) with high-resolution sections from 3D deconvolved confocal datasets, such enlarged terminals appear evenly filled with Syx7 and the photoreceptor-specific transmembrane receptor Chaoptin (Krantz and Zipursky, 1990), which is not observed in wild type. Hence, membrane receptors, including Fas2 (Fig. 4 B) and Chaoptin (Fig. 4 F), exhibit a distribution and colocalization to early endosomal markers that suggest accumulation of large numbers of endosomal vesicles that evenly fill n-syb terminals. Because the loss of n-syb caused no obvious defects in eye development, cell type specification, or axon pathfinding, we infer that spatiotemporally regulated exo- and endocytosis of developmentally required receptors do not require n-syb. In contrast, loss of syx7 causes early developmental and tumor growth phenotypes not observed in the n-syb mutant (Lu and Bilder, 2005), and n-Syb does not directly interact with Syx7 (Fig. S4 A; Antonin et al., 2000) but may reside on the same endocytic vesicles, as suggested by the wild-type colocalization. Collectively with the wild-type localization data, these observations suggest that n-Syb has a hitherto unrecognized function in endolysosomal progression.

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Loss of n-syb causes a primary defect in delivering degradation machinery vesicles to endosomal compartments

What intracellular membrane fusion reaction is disrupted by loss of n-syb? Because loss of n-syb leads to a failure to degrade heterogeneous endosomal, multivesicular, and autophagosomal compartments in a manner reminiscent of loss of v100, a possible explanation is a failure to deliver proteins required for degradation. Such Golgi-derived degradation machinery vesicles might contain the v-ATPase, including V100, as well as lysosomal proteases.

Cathepsins are lysosomal proteases that are synthesized in the endoplasmic reticulum as an inactive, high molecular weight proform that is activated by proteolytic cleavage only in strongly acidified compartments (Lee et al., 2010). The acidification-dependent proteolytic cleavage of pro-Cathepsin generates mature Cathepsins of low molecular weight that can be readily distinguished from the larger proform on Western blots. Hence, analysis of the ratio between the proform and mature Cathepsin provides a means to assay whether vesicles containing degradative enzymes can fuse with lysosomal and autophagosomal compartments. As shown in Fig. 6 A, an antibody against insect Cathepsin L (CatL), which was robustly present in wild type, also exhibited a significant increase that is most pronounced in n-syb or two other degradation mutants, the lysosomal tetraspanin sunglasses or the lysosomal sugar transporter benchwarmer/spinster (Fig. S4 B; Sweeney and Davis, 2002; Xu et al., 2004; Dermaut et al., 2005). These findings are consistent with the idea of a delivery defect of pro-Cathepsin–containing vesicles in the n-syb mutant.

Is the delivery of pro-Cathepsin vesicle a primary defect in the n-syb mutant? In addition to the increase in pro-Cathepsin, we found that mature Cathepsin L, which was robustly present in wild type, also exhibited a significant increase that is most pronounced in n-syb. This is surprising because it implies that more Cathepsin accumulated in both nonacidified (pro-Cathepsin) as well as acidified (mature Cathepsin) compartments. We therefore examined which defect occurs first, the
failed delivery of pro-Cathepsin vesicles or the accumulation of mature Cathepsin in degradative compartments. Interestingly, n-syb mutant eyes from late pupae (P+90%) already exhibited a mild increase in pro-Cathepsin but a temporary decrease of mature Cathepsin, consistent with an initial failure to deliver Cathepsin vesicles to degradative compartments (Fig. 6 A, blue box). These phenotypes differ from V100, which also showed an increase of pro-Cathepsin (albeit less than in n-syb) at 1 d after eclosion but not at P+90%, consistent with our previous analyses (Fig. 6 A; Williamson et al., 2010a). We conclude that the accumulation of pro-Cathepsin precedes the increase of mature Cathepsin in n-syb, supporting the hypothesis that delivery of pro-Cathepsin–containing vesicles is a primary defect. This interpretation is further supported by the EM of developing photoreceptor terminals lacking functional n-Syb in which the accumulation of small vesicles precedes the appearance of larger compartments (Fig. S3).

Because both autophagosomes and the increase in mature Cathepsin only appear in the adult mutant (Fig. 6 A and S3), we set out to test the possibility that autophagy is a secondary response to an earlier trafficking defect caused by loss of n-syb. Autophagy is critically important for neuronal maintenance (Hara et al., 2006; Komatsu et al., 2006), but autophagy is not known to require any neuron-specific proteins. A recently developed probe to measure autophagosome acidification is based on the autophagosome marker Atg8/LC3; double tagging with acidification-sensitive GFP and acidification-insensitive mCherry leads to selective quenching of GFP of this reporter only in acidified autophagosomes (Filimonenko et al., 2007). As shown in Fig. 6 (B and C), autophagosomes in wild-type and n-syb mutant photoreceptor cell bodies exhibited selective quenching of GFP, indicating that they were indeed acidified. Furthermore, if these autophagosomes compartments contained active proteases, we should observe morphologically distinct, late-stage autophagosomes with degraded contents. We therefore counted early stage autophagosome profiles (immature autophagosomal vacuoles [AVIs]; containing large, morphologically recognizable cellular structures; Fig. S5, arrowheads) and late autophagosome profiles (degradative autophagosomal vacuoloes [AVDs]; containing only small, electron-dense debris; Fig. S5, arrows). These counts revealed that almost 90% of all autophagosomes in the n-syb mutant are late AVDs, whereas AVI and AVD profiles appeared in a 50:50 ratio in v100 mutants, which are predicted to partially lack acidification required for degradation (Fig. S5). Finally, cooverexpression of tagged versions of the autophagosome markers LC3/Atg8 and n-Syb revealed that autophagosomes are primarily found in cell bodies, where they only partially overlap with n-Syb. In contrast, n-Syb is transported to the synapses and in the cell bodies marked mostly compartments that are not positive for LC3/Atg8 (Fig. 6 E). This nonoverlapping pattern of expression further supports an autophagy-independent function of n-Syb. We conclude that by all the criteria we have investigated here, autophagy seems to be functional until late stages of autophagosome maturation independent of n-syb and that the pronounced increases in endosomal markers and vesicles are distinct from the increase in the number of autophagosomes. The presence of autophagosomal accumulations may reflect a dramatic increase of functional autophagy, or loss of n-syb causes a defect in removing late autophagosomal compartments. In both scenarios, increased autophagy is likely to be a secondary response to a primary vesicle trafficking defect downstream of endocytosis.

**Early cell death through induction of autophagy or tau overexpression does not depend on n-syb, but loss of n-syb sensitzizes adult photoreceptors to degeneration**

Unchecked induction of autophagy through overexpression of atg1 causes cell death (Berry and Baehrecke, 2007; Scott et al., 2007). Correspondingly, high levels of atg1 overexpression during photoreceptor development cause eye developmental defects that result in a reduced and rough eye (Fig. 7, A and E). atg1-induced cell death thus provides a direct test as to whether n-syb is required for autophagy in photoreceptors. As shown in Fig. 7 (E and F), loss of n-syb did not reduce the ability of atg1-induced autophagy to disrupt eye development. Interestingly, loss of n-syb did not noticeably increase the developmental defect as indicated by indistinguishable reductions in eye size. However, ERG recordings from these eyes in 1-d-old flies revealed a reduced response amplitude in atg1-overexpressing photoreceptors that was further reduced in an n-syb mutant background. These findings indicate that n-syb is not required for atg1-induced autophagy but has an additive negative effect on adult photoreceptor function. The genetics therefore do not indicate a requirement of n-syb for autophagy but rather suggest that loss of n-syb sensitizes adult photoreceptors to neurotoxic insults through an independent mechanism. To test this idea further, we challenged wild-type and n-syb mutant photoreceptors with overexpression of human tau protein, similar to an experiment recently performed for v100 (Wittmann et al., 2001; Williamson and Hiesinger, 2010a). Overexpression of tau caused a defective eye morphology and reduction of the ERG response amplitude similar in strength to atg1 overexpression (Fig. 7, I–L). As in the case of atg1 overexpression, loss of n-syb did not noticeably affect the developmental defect but lead to a further reduction of the response amplitude in the adult (Fig. 7, K and L). We conclude that impaired intracellular degradation through loss of n-syb sensitizes adult photoreceptor neurons to neurotoxic insults and that n-syb is not required for atg1-induced autophagy.

**Overexpression of V100 partially rescues n-syb-dependent neurodegeneration**

Loss of either n-syb or v100 causes synaptic dysfunction, neurodegeneration, and accumulation of undegraded endosomal compartments. Furthermore, loss of n-syb leads to reduced V100 localization at synapses. To test genetic interactions between the two mutants, we analyzed both RNAi-mediated loss of v100 (Williamson and Hiesinger, 2010a) and v100 overexpression in n-syb mutant photoreceptors. v100 RNAi does not rescue n-syb–dependent degeneration (unpublished data). In contrast, overexpressing V100 partially rescues both
Our results indicate that the acidification-independent function of V100 is sufficient to significantly protect n-syb mutant photoreceptor neurons from degeneration independent of neurotransmitter release.

V100 overexpression does not reduce intracellular accumulations but increases undegraded cargo colocalization with early endosomes.

The rescue with acidification-defective v100R755A is not consistent with increased degradation as a rescuing parameter. To the contrary, we found even more widespread and evenly morphological and functional degeneration (Fig. 8). We undertook ERG and immunohistochemical analyses in flies that had been exposed to constant light for 7 d. Under these conditions, wild-type photoreceptors exhibited both a normal ERG and morphology (Fig. 8, A and E). In contrast, n-syb mutant photoreceptors exhibited a significant reduction in the number of rhabdomere profiles (Fig. 8 B) and an ~50% reduction of depolarization (Fig. 8, F and J). Photoreceptor-specific expression of v100 in the n-syb mutant terminals partially but significantly rescued the defect in rhabdomere number (Fig. 8, C and K) as well as ERG depolarization (Fig. 8, G and J). Next, we tested whether the acidification function of V100 is required for this rescue. We have previously generated the v100R755A mutant that selectively disrupts proton translocation (Williamson et al., 2010a). Surprisingly, v100R755A expression in photoreceptors rescued the degeneration defect to an extent indistinguishable from wild-type v100 overexpression. Note that the loss of neurotransmitter release (as measured by the ERG “on” transient) in n-syb mutant neurons was not rescued by either wild-type or acidification-defective v100 (Fig. 8 I).

Figure 7. Early cell death through induction of autophagy or tau overexpression does not depend on n-syb, but loss of n-syb sensitizes adult photoreceptors to degeneration. (A–L) Eye pictures and sample ERG traces for control (A and C), n-syb (B and D), autophagy induction through atg1 expression in wild type (E and G), and n-syb mutant (F and H) photoreceptors. Expression of human tau in wild-type (I and K) and mutant (J and L) photoreceptors. (M) Quantification of ERG response amplitudes. Note that loss of n-syb does not obviously alter the developmental eye defects but leads to further reduced response amplitudes in adult photoreceptors. All error bars are SEM. ctrl, control.
We next asked how Cathepsin sorting and processing is affected by \( v100 \) overexpression in \( n-syb \) mutant photoreceptors. As shown in Fig. 9 K, overexpression of \( v100 \) in \( n-syb \) mutant photoreceptors caused a further increase of both pro-Cathepsin and mature Cathepsin, whereas overexpression of \( v100 \) in wild-type photoreceptors caused only very mild changes. This result suggests that some Cathepsin was normally degraded independent of \( n-syb \) but now rerouted by V100. Based on the \( v100 \) overexpression results, we hypothesized that V100 is a component of Cathepsin-containing vesicles and that it plays a role during vesicle sorting (Fig. 10). As shown in Fig. 9 K, expression of \( v100 \)-R755A in the \( n-syb \) mutant lead to a substantial increase in pro-Cathepsin at the expense of mature Cathepsin. This observation strongly supports the idea that the overexpressed V100 indeed resides on vesicles that contain Cathepsin and that the vesicles require V100’s acidification function to mature. Note that expressing the acidification-defective \( v100 \)-R755A only had a very mild effect in wild-type neurons, whereas its expression in \( n-syb \) mutant, alone among all genotypes analyzed here, led to a ratio reversal to more pro- than mature Cathepsin (Fig. 9 K). Furthermore, \( v100 \)-R755A expression in the \( n-syb \) mutant decreased mature Cathepsin when compared with the \( n-syb \) mutant alone. Because expression of V100-R755A rescues degeneration to the same extent as wild-type \( v100 \), these findings are not consistent with increased degradative capacity as a rescuing parameter. Instead, the \( v100 \)-R755A experiment suggests that V100 reroutes Cathepsin vesicles to compartments for whose acidification V100 itself is required. Finally, we observe that the further increase of cargo colocalization with early endosomal compartments in these rescue experiments coincides with reduced neurodegeneration compared with the \( n-syb \) mutant.
Discussion

We have characterized an unexpected function of the neuronal v-SNARE n-Syb in endolysosomal trafficking that is independent of its known function in neurotransmitter release. Loss of...
this function leads to neurodegeneration in adult fly photoreceptor neurons. These findings explain our earlier observations of late developmental defects during brain wiring that do not occur when neuronal activity or neurotransmitter release is impaired (Hiesinger et al., 1999, 2006). Genetic and cell biological evidence indicates that n-Syb functions in concert with the neuronal v-ATPase component V100 in a neuronal sort and degrade pathway. Specifically, our evidence supports a primary defect in the fusion of vesicles that contain degradation machinery (including degradation proteases and V100-containing ATPases required to activate them) with endosomal compartments.

**A role in neuronal vesicle fusion independent of synaptic vesicle exocytosis**

The ultrastructure of n-syb mutant terminals reveals accumulations of both small vesicles and large degradative and autophagosomal compartments. n-Syb is a small (<200 amino acids) protein that is very well characterized as a synaptic v-SNARE required for vesicle fusion (Broadie et al., 1995; Galli et al., 1995; Deitcher et al., 1998; Südhof and Rothman, 2009). In addition, synaptic vesicles are sorted through synaptic endosomes (Uytterhoeven et al., 2011). Hence, the most straightforward explanation for endolysosomal trafficking defects in n-syb mutant neurons would be a defect in vesicle fusion.

Several observations indicate that the endolysosomal defect is not directly linked to n-Syb’s function in synaptic vesicle exocytosis. First, loss of neuronal activity or neurotransmitter release does not cause endolysosomal, developmental or degenerative phenotypes in *Drosophila* photoreceptors as shown here and previously (Hiesinger et al., 2006). Second, although vesicles accumulate, synaptic vesicle markers do not accumulate correspondingly. Instead, we find that several markers of the endolysosomal system are up-regulated. In particular, the early endosomal t-SNARE Syx7, together with transmembrane receptors, including Chaoptin and Fasciclin 2, exhibits a strong up-regulation and evenly fills the profiles of enlarged synaptic terminals. Together with the EM data, we conclude that synaptic terminals are filled with large numbers of endosomal vesicles rather than with synaptic vesicles.

Several lines of evidence indicate that n-syb is not required for autophagosome maturation, at least up to a late stage. First, n-Syb colocalizes only little with autophagosomes but marks mostly endosomal compartments and synaptic vesicles. Second, autophagosomes are acidified and contain active proteases (Cathepsins) and degraded material based on EM, Atg8-mCherry-GFP measurements, and mature Cathepsin accumulations. Third, induction of autophagy by *atg1* expression does not depend on n-syb. Finally, the accumulation of undegraded Chaoptin and Fasciclin 2 receptors that evenly fill terminals is not consistent with their accumulation in autophagosomes. Because autophagy typically initiates in response to nutrient deprivation or other neurotoxic insults, the appearance of autophagosomes may be a secondary response (Tooze and Schiavo, 2008; McPhee and Baehrecke, 2009).
ponent V100 defines a neuronal sort-and-degrade mechanism (Fig. 10). The inverse correlation between endosomal accumu-
v100 yet a significantly slower progression of degeneration than in
v100 netic and cell biological interactions indicate that
localization, while further ameliorating the degeneration. The ge-
compartments, with which fusion of small endosomal vesicles
vesicles with neuronal endolysosomal compartments before the
delivery to autophagosomes is apparently unimpaired. Further-
mechanism) vesicles with to-be-degraded compartments
mechanism suggests that the synaptic vesicle cycle may be
interlinked with v100- and n-syb-mediated endosomal trafficking.
Furthermore, these observations suggest the resulting
maintenance function predominantly operates at synapses. We
therefore propose that v100 and n-syb mutants reveal a neuron-
specific synaptic extension of the endolysosomal system that
serves the specialized and extensive demands of neurons, and
of synapses in particular, on intracellular membrane trafficking.

Materials and methods

Genetics and fly culture
Eye clones were generated using the ey3.5FLP system [Chotard et al., 2005; Mehta et al., 2005] to generate photoreceptor clones in otherwise heterozy-
gous animals. The alleles n-syb^{F33B} (Deitcher et al., 1998), v100^{100P} (Hiesinger et al., 2005), and syx7^{M2} (DiAntonio et al., 1993) were used for all ex-
periments. The transgenes upstream activation sequence (UAS)-v100^{47} (Hiesinger et al., 2005), UAS-v100^{P546} (Williamson et al., 2010a), UAS-
n-syb [Bhattacharya et al., 2002], UAS-atg1 (Scott et al., 2007), UAS-
GFp-mCherry-atg8a (Filimonenko et al., 2007), UAS-YFP-rab5, UAS-YFP-rab7, and UAS-YFP-rab 11 (Zhang et al., 2007) have all been previously described.

For degeneration assays, flies were raised at 22°C. Flies were either
exposed to standard room light for ~12 h followed by darkness for 12 h,
or they were constantly exposed to light stimulation by four light-emitting
diode bulbs (2026LED-65K; 1.5 W, 30 mA) in a 30 x 30 x 40-cm aluminum foil-lined box.

Genotypes. For the analysis of mutant photoreceptors the following were used: control [ey3.5FLP;FRT82B;FRT82B], v100 [ey3.5FLP;FRT82B,t.
FRT82Bv100P;Hiesinger et al., 2005], n-syb^{ey3.5FLP;FRT82B;w,cl>FRT808
n-syb^{F33B}; Deitcher et al., 1998); n-syb rescue [ey3.5FLP;GMRA4/UAS-
n-syb,FRT80B;w,cl>FRT80B; syx7^{M2}; syx7^{M2}, ey3.5FLP;FRT40A,w,cl>FRT40A
syx^{M2}; Littleton et al., 1993] TNT inactive [GMRA4=UAS-TNT; Sweeney et al., 1995], TNT active [GMRA4=UAS-TNT-H; Sweeney et al., 1995];
atg1 expression [GMRA4=UAS-atg1; Scott et al., 2007]; atg1 expression in
n-syb photoreceptors [ey3.5FLP;GMRA4/UAS-atg1; FRT80B;w,cl>FRT808
n-syb^{F33B}], tau expression [GMRA4=UAS-etu; Wittmann et al., 2001], tau expression in n-syb photoreceptors [ey3.5FLP;GMRA4/UAS-atg1;
FRT80B;w,cl>FRT80B; syx7^{M2}, v100 expression in n-syb photoreceptors
[ey3.5FLP;GMRA4/UAS-v100; FRT80B;w,cl>FRT80B; syx7^{M2}], and
v100^{R755A} expression in n-syb photoreceptors [ey3.5FLP;GMRA4/UAS-
v100^{R755A}; FRT80B;w,cl>FRT80B; syx7^{M2}]. All genotypes were verified by
unequivocal markers on balancer chromosomes (Cy for CyO; Hu and Tb for
TM6b, Hu; Tb; Sb for TM3, Sb) as well as independent markers (e.g., eye
mosaicism, eye roughness, and antibody staining).

A neuronal sort-and-degrade mechanism and its implications for neurodegeneration

We recently reported that the neuron-specific v-ATPase com-
ponent V100 defines a neuronal sort-and-degrade mechanism that
increases neuronal degradative capacity [Williamson and
Hiesinger, 2010a; Williamson et al., 2010a]. Loss of n-syb and
v100 reveals striking similarities with one remarkable difference:
endosomal accumulations in n-syb are considerably more
severely compromised when amino acids and other molecules are not
recycled and accumulate in degraded compartments; second, the
accumulations themselves may be toxic. The latter hypothesis
is typically discussed for accumulations in late degradative
compartments, as they have a highly toxic content in terms of
pH and proteases (leaky lysosomes). This idea is consistent with
our observation of reduced toxicity through increased
accumulations in early endosomal compartments in n-syb and
further with v100 overexpression in the n-syb mutant. We there-
fore speculate that the appearance of more neurotoxic late degra-
dative compartments is the likely cause of death in n-syb
mutant photoreceptors.

Notably, both V100 and n-Syb are synaptic proteins that
have first been characterized as components of synaptic vesi-
cles. Our finding that both proteins additionally function in neu-nonal maintenance through an endolysosomal sort-and-degrade
mechanism suggests that the synaptic vesicle cycle may be
interlinked with v100- and n-syb-mediated endosomal trafficking.
What then is n-Syb’s other vesicle fusion role? We have
firstly, the complete absence of any early developmental de-
defect implies that n-syb cannot be required for the canonical
secretory pathway that regulates morphogen and receptor
exocytosis. Similarly, spatiotemporally regulated receptor
endocytosis appears unaffected because receptor accumula-
tions distribute evenly in the enlarged n-syb mutant terminals.
We conclude that receptors accumulate in small endocytic vesi-
cles downstream of endocytosis and upstream of further sorting
or degradation. Collectively, these observations pinpoint a role
for n-Syb in the fusion of CathepsinV100-containing (degra-
dation machinery) vesicles with to-be-degraded compartments
(Fig. 10). The latter do not include autophagosomes, as Cathepsin
delivery to autophagosomes is apparently unimpaired. Further-
more, the large numbers of Syx7-positive small endosomal vesi-
cles are not likely a fusion partner because Syx7 does not directly
interact with n-Syb as shown here and in vertebrates (Antoini
et al., 2000). Instead, we can narrow down the function to a
fusion reaction between V100/Cathepsin degradation machinery
vesicles with neuronal endolysosomal compartments before the
formation of lysosomes. The n-Syb-mediated endolysosomal
fusion event could be directly between degradation machinery
vesicles and larger or intermediate endosomal vesicles or sorting
compartments, with which fusion of small endosomal vesicles is
also blocked in the absence of n-syb (Fig. 10). In either case,
the intravesicular endosomal fusion uncovered by loss of n-syb
represents a neuronal specialization and possible addition to
the ubiquitous endolysosomal machinery, as shown in the model
in Fig. 10.

endosomal accumulations are not per se neurotoxic and less toxic
than accumulations in late degradative compartments.

What then kills n-syb mutant neurons? Two main mecha-
nisms are thought to underlie cell death in most neurodegen-
erative conditions. First, the neuron may become metabolically
compromised when amino acids and other molecules are not
recycled and accumulate in degraded compartments; second, the
accumulations themselves may be toxic. The latter hypothesis
is typically discussed for accumulations in late degradative
compartments, as they have a highly toxic content in terms of
pH and proteases (leaky lysosomes). This idea is consistent with
our observation of reduced toxicity through increased
accumulations in early endosomal compartments in n-syb and
further with v100 overexpression in the n-syb mutant. We there-
fore speculate that the appearance of more neurotoxic late degra-
dative compartments is the likely cause of death in n-syb
mutant photoreceptors.
Immunohistochemistry, microscopy, and image processing

Adult brains and eyes, as well as pupal brains and eye-brain complexes, were dissected as previously reported (Williamson and Hiesinger, 2010b). In brief, the tissues were fixed in PBS with 3.7% formaldehyde for 30 min and washed in PBS with 0.4% Triton X-100. High-resolution light microscopy was performed at room temperature using a resonance-scanning confocal microscope (SP5; Leica) using a 63x HCX Plan Apochromat, NA 1.45, glycerin lens. Imaging data were processed and quantified using Amira 3.2 (Indeed), Photoshop (CS2; Adobe), and Imagej (National Institutes of Health). All confocal data were obtained as 3D dataset, and all images shown are either single sections or maximum intensity projections of small subsets that were equally processed for controls and mutants. Blind deconvolution was applied as previously published (Hiesinger et al., 2001).

For the quantification of up/down-regulation in mutant versus wild-type photoreceptor terminals, 50% mutant photoreceptor mosaics were created and verified using anti-n-Syb immunolabeling or other established markers. 3D tetroxide, all as previously reported. Given that tetrad synapses are distributed relatively evenly along the length of the terminal, we sampled synaptic organelles from different levels in a single section to obtain averages. Each terminal’s profile was captured with a camera (MegaView II; Kodak) and software (Soft Imaging Solutions; Olympus) at a magnification of 4,900–9,700x using a microscope (201C or Tecnai; Philips) operated at 80 kV. Samples of ≥15 terminal profiles per fly were evaluated from at least three flies per genotype. Tests of statistical significance were made between terminals from paired genotypes using a two-tailed t test of the means of the mean counts or measurements.

ERGs

ERGs were recorded as previously described (Fabian-Fine et al., 2003; Williamson et al., 2010a). In brief, flies were reversibly glued on slides using nontoxic school glue. Light stimulus was provided in 1-s pulses by a computer-controlled white light-emitting diode system (MC1500; Schott). Data were recorded using Clampex (version 10.1; Axon Instruments) and measured using Clampfit (version 10.2; Axon Instruments).

Western blots of mutant eye-lamina complexes, coimmunoprecipitations, and pull-downs

Eyes were dissected from 1-d-old flies in HL3 medium (Stewart et al., 1994). Groups of 20 eyes were collected in HL3 and then crushed in 10 μl extraction buffer [20 mM Tris, pH 7.4, 150 mM NaCl, 1 mM PMSF, and protease inhibitor [Complete; Roche]]. Samples were incubated on ice for 15 min and then centrifuged at 16,000 RCF for 10 min at 4°C to precipitate debris. Next, 9 μl of sample was then added to 9 μl of 2× Laemmli buffer to make a final concentration of 1 eye/μl. 10 eyes were run on each lane of a 12.5% PAGE gel.

Communoprecipitations. Flies were frozen in liquid nitrogen, and heads were collected with a sieve. Total protein was extracted in immuno-precipitation buffer containing 20 mM Tris, 150 mM NaCl, 1 mM PMSF, and Complete protease inhibitors, pH 7.4. The fly head extract was mixed well in 1% Triton X-100 [Bio-Rad Laboratories] and incubated for 1 h at 4°C. Samples were centrifuged at 16,000 g for 15 min at 4°C to remove cell debris. The resulting supernatant was incubated with 20 μl anti-Syb7 antibody (a gift from H. Krämer, University of Texas Southwestern Medical Center, Dallas, TX). 8C3 anti-Syntaxin antibody (Hiesinger et al., 2005), and anti-lab antibody (Novus) coupled to protein A/G beads (Santa Cruz Biotechnology, Inc.) for 1 h at 4°C. After removing the supernatant, the beads were washed four times with immunoprecipitation buffer. A nonspecific preimmune serum was used as a control. The immunoprecipitates were eluted by boiling the beads in 50 μl SDS sample buffer and analyzed by Western blot with anti-Syb and anti-Syx7 antibodies.

GST pull-down assays. For in vivo GST pull-down assays, the GST Syx7 fusion protein was bound to glutathione-Sepharose 4B [GE Health-care] and washed with standard buffer [28 μM Tris/150 mM NaCl, 1 mM PMSF, and Complete protease inhibitors, pH 7.4] at 4°C overnight. For in vitro GST pull-down assays, His-tagged n-Syb protein was incubated with the indicated GST fusion proteins in binding buffer [20 μM Tris/150 mM NaCl, pH 7.4, 0.2% Triton X-100] at 4°C overnight. After binding, the beads were washed three times in binding buffer, and protein samples were eluted with SDS sample buffer for Western blot with anti-Syb and anti-V100 antibodies (Hiesinger et al., 2005).

EM

Newly hatched adults or staged pupae were prepared for EM and fixed as previously described (Meinertzhagen, 1996). In brief, for pupae, head capsules were carefully opened under Karnovsky fixative, and brains were dissected and split with forceps within 2 min. Tangential sections of the lamina were cut at 65 nm. Fixation was performed in modified Karnovsky fixative (Meinertzhagen, 1996) followed by veronal-buffered 2% osmium tetroxide, all as previously reported. Given that tetrads are distributed relatively evenly along the length of the terminal, we sampled synaptic organelles from different levels in a single section to obtain averages. Each terminal’s profile was captured with a camera (MegaView II; Kodak) and software (Soft Imaging Solutions; Olympus) at a magnification of 4,900–9,700x using a microscope (201C or Tecnai; Philips) operated at 80 kV. Samples of ≥15 terminal profiles per fly were evaluated from at least three flies per genotype. Tests of statistical significance were made between terminals from paired genotypes using a two-tailed t test of the means of the mean counts or measurements.

Online supplemental material

Fig. S1 shows loss of neurotransmitter release in Drosophila photoreceptors in a mutant for histamine synthesis does not cause degeneration. Fig. S2 shows that lack of synb but not syx7 in photoreceptors disrupts lamina development. Fig. S3 shows developmental analysis of synaptic terminals in the lamina. Fig. S4 shows that n-Syb does not directly interact with the early endosomal Syntaxin Syx7 and loss of n-syb causes more severe pro-Cathepsin accumulations than lysosomal degradation mutants. Fig. S5 shows that n-syb mutant synaptic terminals contain many late autophago- somal compartments. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.201108088/DC1.

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References


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