A Neuroprotective Function of NSF1 Sustains Autophagy and Lysosomal Trafficking in Drosophila

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ABSTRACT A common feature of many neurodegenerative diseases is the accumulation of toxic proteins that disrupt vital cellular functions. Degradative pathways such as autophagy play an important protective role in breaking down misfolded and long-lived proteins. Neurons are particularly vulnerable to defects in these pathways, but many of the details regarding the link between autophagy and neurodegeneration remain unclear. We previously found that temperature-sensitive paralytic mutants in Drosophila are enriched for those exhibiting age-dependent neurodegeneration. Here we show that one of these mutants, comatose (comt), in addition to locomotor defects, displays shortened lifespan and progressive neurodegeneration, including loss of dopaminergic (DA) neurons. comt encodes N-ethyl-maleimide sensitive fusion protein (NSF1), which has a well-documented role in synaptic transmission. However, the neurodegenerative phenotypes we observe in comt mutants do not appear to depend on defects in synaptic transmission, but rather from their inability to sustain autophagy under stress, due at least in part to a defect in trafficking of lysosomal proteases such as cathepsin-L. Conversely, overexpression of NSF1 rescues α-synuclein-induced toxicity of DA neurons in a model of Parkinson’s disease. Our results demonstrate a neuroprotective role for NSF1 that involves mediation of fusion events crucial for degradative pathways such as autophagy, providing greater understanding of cellular dysfunctions common to several neurodegenerative diseases.

KEYWORDS comatose; neurodegeneration; autophagy

Recent studies have demonstrated the utility of Drosophila models of neurodegeneration in aiding our understanding of a number of neurodegenerative diseases. Specifically, expression of human proteins implicated in diseases including Alzheimer’s disease (Ye and Fortini 1999; Wittmann et al. 2001), Parkinson’s disease (Feany and Bender 2000; Auluck et al. 2002), and polyglutamine expansion disorders (Jackson et al. 1998; Warrick et al. 1998), among others has proven to be a powerful approach for uncovering the mechanisms underlying disease progression.

An alternative strategy to understanding neurodegeneration involves a forward genetics approach, by screening for mutations that result in progressive neurodegeneration to uncover neuroprotective mechanisms that may have further implications for human neurodegenerative diseases. We previously reported a collection of temperature-sensitive (ts) paralytic mutants that is enriched for mutants showing progressive, age-dependent neurodegeneration (Palladino et al. 2002). We have found that other locomotor mutants in our collection also exhibit neurodegeneration. Analysis of these mutants has uncovered several novel neuroprotective functions, ranging from regulation of metabolic processes to regulation of the innate immune response (Palladino et al. 2003; Gnerer et al. 2006; Miller et al. 2012; Cao et al. 2013).

Here we show a neuroprotective role for NSF1 using ts comatose (comt) mutants, which were originally identified in a screen for ts-paralytic mutants (Siddiqi and Benzer 1976). comt encodes N-ethyl-maleimide sensitive fusion protein (NSF1), a AAA-ATPase with a well-characterized role in synaptic transmission (Pallanck et al. 1995; Chen and Scheller 2001). Specifically, NSF1 is involved in the disassembly of SNARE complexes following synaptic vesicle fusion, a process crucial to maintaining the readily releasable pool of synaptic vesicles for subsequent rounds of fusion (Littleton et al. 1998, 2001; Tolar and Pallanck 1998; Sanyal et al. 2001; Hanson and Whiteheart 2005).

Despite its role in synaptic transmission, we find that the neuroprotective function of NSF1 appears to operate via a...
distinct mechanism. Specifically, we find that NSF1 is required to maintain autophagy during periods of stress. Autophagy is a degradative process whereby misfolded proteins, aggregates, organelles, and other cellular materials are engulfed by autophagosomes, which then fuse with lysosomes where the contents are degraded and recycled (Klionsky 2007).

Although previous studies have revealed a link between perturbations in autophagy and neurodegeneration in both mice and humans (Hara et al. 2006; Komatsu et al. 2006; McCray and Taylor 2008), NSF has not previously been implicated in any neurodegenerative disorder. However, recent evidence demonstrates that SNARE proteins such as neuronal synaptobrevin (nSyb) have important neuroprotective functions in addition to mediating synaptic transmission (Haberman et al. 2012). Nonetheless, the question of whether recycling of SNARE complexes following fusion and its dependence on NSF is as important for these neuroprotective functions as for rapid transmitter release in synaptic transmission has not previously been addressed. By investigating the neuroprotective role of NSF in Drosophila and determining which particular mechanisms are particularly sensitive to proper NSF function, we have further elucidated key steps required for maintenance of neuronal integrity that may ultimately fail in some neurodegenerative diseases.

Materials and Methods

Fly stocks

Canton-S was used as a wild-type control. The temperature-sensitive comt alleles comtTP7 (provided by K. S. Krishnan, Tata Institute, Bangalore, India), comtST17, and comtST53 (Siddiqi and Benzer 1976) were previously described. UAS-lamp1-GFP (Pulipparacharuvil et al. 2005) was provided by Helmut Kramer (University of Texas Southwestern Medical Center, Dallas) and UAS-mCherry-GFP-Atg8a (Filimonenko et al. 2007) was provided by Thomas Neufeld (University of Minnesota, Minneapolis). snap25 (Babcock et al. 2004) and UAS-NFS1 (Golby et al. 2001) were provided by Leo Pallanck (University of Washington, Seattle). setTS2 (Jackson et al. 1985) was previously described. The following fly lines were obtained from the Bloomington Stock Center: elav-Gal4, Cg-Gal4, TH-Gal4, UAS-αSyn, UAS-Syx1RNAi, UAS-ubisnapRNAi, UAS-VAMP7RNAi, and UAS-Atg5RNAi. UAS-comtRNAi (no. 105552) was obtained from the Vienna Drosophila RNAi Center.

Histology

Newly eclosed flies were shifted to 29°C for up to 48 hr. Heads were severed and placed in fresh Carnoy’s fixative overnight at 4°C. Samples were then washed in 70% ethanol and embedded in paraaffin using standard histological procedures. The 5-μm frontal sections were cut, stained in hematoxylin and eosin, and imaged under a light microscope. Vacuoles of at least 2 μm in diameter were recorded for each sample. To avoid potential artifacts, vacuoles were only recorded if they appeared in at least two consecutive sections.

Lifespan analysis

Flies were collected shortly after eclosion and cohorts of 10 flies were placed in single vials. A total of 10 vials (100 flies) were used for each genotype. Flies shifted to 29°C were transferred to fresh food every day and the percentage of surviving flies was measured. Flies remaining at room temperature (22°C) were transferred and measured every 2 days.

Climbing assay

Newly emerged flies were collected and placed into cohorts of 10 flies per vial. A total of 10 cohorts of flies were assessed for each genotype. Climbing ability was assessed by transferring a cohort of flies into an empty climbing cylinder (diameter, 2.5 cm; height, 20 cm). For each trial, flies were tapped to the bottom of the vial, and the percentage of flies passing an 8-cm mark within 20 sec was recorded. This percentage of flies was recorded as the climbing index. Each cohort of flies was assessed across five trials, with a 1-min recovery between trials.

Lysotracker staining

Lysotracker staining was performed as previously described (Wong et al. 2012). Briefly, larval fat bodies were dissected in cold PBS and immediately transferred to a solution of Lysotracker Red DND-99 (Invitrogen) (1:500) for 45 min at room temperature. Samples were washed four times in PBS and fixed in 4% formaldehyde in PBS for 20 min at room temperature. Samples were washed four times in PBS and mounted in Vectashield (Vector Laboratories) along with DAPI (1:1000). The size of LysoTracker-positive puncta was measured using ImageJ software (National Institutes of Health, NIH).

Immunohistochemistry

Brains and fat body samples were dissected and fixed in 4% formaldehyde in PBS for 20 min at room temperature. Samples were then placed in blocking buffer (PBS with 0.1% Triton X-100 and 0.1% normal goat serum) for 1 hr. Samples were then incubated in primary antibodies overnight at 4°C. Next, samples were washed five times in PBS, then incubated in secondary antibodies for 2 hr at room temperature. Finally, samples were washed five times in PBS and mounted in Vectashield. Primary antibodies used were: rabbit anti-tyrosine hydroxylase (1:100) (Millipore), mouse anti-GFP (1:500) (Invitrogen), rabbit anti-DsRed (1:500) (Clontech Laboratories), and mouse anti-bruchpilot (nc82) (1:50) (Developmental Studies Hybridoma Bank, University of Iowa). Secondary antibodies used were: goat anti-rabbit Alexa-568, goat anti-mouse Alexa-488, and goat anti-rabbit Alexa-488 (1:200) (Invitrogen).

Imaging and quantification

Images were obtained on a Leica LSM 500 confocal microscope. Serial 1-μm z-stacks were obtained for each image using a ×20 numerical aperture (N.A.) 10.8 or ×63 N.A. 1.4 oil objective. Brightness and contrast were adjusted using ImageJ software (NIH) and Adobe Photoshop CS5 (Adobe).
Dopaminergic neurons were counted as previously described (Whitworth et al. 2005; Barone and Bohmann 2013). Briefly, dissected brains were mounted between two glass coverslips to aid in visualization of PPL1 neurons. The number of neurons in each cluster was assessed with the scorer blind to the genotype and condition of each sample.

**Starvation**

Flies were collected shortly after eclosion and placed in groups of 10 in individual vials. Ten vials (100 flies) were used per genotype. Flies were raised on standard fly media for 2 days, and then transferred to empty vials lined with water-soaked filter paper to prevent desiccation. Flies were transferred to fresh vials every 8 hr, and the percentage of surviving flies was recorded. For the larval starvation, early third-instar larvae were collected and plated in a 20% sucrose solution for 3 or 12 hr.

**Western blotting**

Two adult flies of each genotype were homogenized using 20 μl of homogenization buffer (100 mM Tris-HCl, pH 6.8, 2% SDS). The homogenate was then boiled for 10 min, and centrifuged at 21,000 × g for 10 min to remove debris. A total of 5 μl of each sample was run on a 10% acrylamide gel. Western blots were probed with mouse anti-cathepsin-L at 1:5000 (R&D Biosystems) and goat anti-actin at 1:1000 (Santa Cruz Biotechnology). Protein levels were quantified using an Odyssey Imaging System (Li-Cor) by normalizing band intensities to the level of actin. Experiments were repeated at least three times.

**Statistical analysis**

Significance for lifespan and starvation survival curves was analyzed using the log-rank test. Climbing behavior and number of dopaminergic neurons were analyzed using the Student’s t-test, with Bonferroni corrections for multiple comparisons when applicable. Statistical analysis was computed using SPSS software (IBM Corporation).

**Results**

**comt** mutants exhibit shortened lifespan, locomotor defects, and neurodegeneration

**comt** is a recessive X-linked ts-paralytic mutant with a well-documented defect in synaptic transmission at temperatures above 38° (Littleton et al. 1998, 2001; Tolar and Pallanck 1998). Although **comt** flies are not paralyzed below 38°, we found that lifespan was greatly reduced compared with wild-type controls for flies shifted to 29° after eclosion. Three different hypomorphic alleles of **comt** had severely reduced lifespans, with median lifespans ranging from 2 to 4 days, compared with >30 days for controls (Figure 1A). When raised at room temperature (22°), **comt** mutants also displayed shortened lifespan compared with controls, but the phenotype was much milder (Figure 1B). These results are consistent with the idea that **comt** mutants have an underlying unconditional impairment other than a synaptic transmission defect, with the consequences of this impairment becoming more severe under the stress of elevated temperature. In parallel with the shortened lifespan, we also observed progressive neurodegeneration in **comt** mutant brains. This neurodegeneration was assessed in histological sections as vacuolar pathology in the neuropil after 48-hr exposure to 29° (Figure 1, C–F).

Along with neurodegeneration, **comt** mutants also displayed a marked decrease in climbing ability (Figure 2A). All **comt** alleles tested showed significantly reduced climbing as early as 12 hr after shifting to 29°, with a complete lack of climbing ability by 24 hr, even though the restrictive temperature for paralysis is about 38°. The most severe allele, **comt**T7P7, had a moderate climbing defect even before shifting to 29°, again indicating an underlying mutant defect that is independent of temperature.

To investigate neurodegeneration in **comt** mutants in more detail, we asked whether specific populations of neurons were preferentially lost. Because dopaminergic (DA) neuron loss is strongly correlated with locomotor dysfunction, including impaired climbing ability in *Drosophila* (Feany and Bender 2000), we first tested whether DA neurons are among those lost in **comt** mutants. DA neurons are found in distinct clusters throughout the central brain of *Drosophila*. Several of these clusters, particularly on the posterior side of the brain, contain relatively small numbers of neurons, making them amenable to counting (Figure 2B). We found that several DA neuron populations in the protocerebral posterior lateral 1 (PPL1) cluster were lost in **comt** mutant brains after 48 hr at 29° (Figure 2, C–G). Thus, we conclude that NSF1 function is required for neuronal viability, particularly under stressful conditions such as elevated temperature.

**The neuroprotective role of NSF1 is distinct from its role in synaptic transmission**

As mentioned previously, NSF1 has a well-characterized role in synaptic transmission. Although synaptic transmission is blocked in photoreceptors at 38° in **comt** mutants (Littleton et al. 1998, 2001), without direct electrophysiological recordings from the synaptic partners of DA neurons, it is not possible to determine whether synaptic transmission is completely normal in these cells at the 29° incubation temperature used in the experiments above. Thus, to test whether a defect in synaptic transmission can cause loss of DA neurons, we hyperpolarized DA neurons by expressing the inward-rectifying potassium channel, Kir2.1 (Baines et al. 2001) using the TH-Gal4 driver, which should block synaptic transmission in these cells. Because TH-Gal4 also drives expression in non-neuronal cells required for embryonic development (Friggi-Grelin et al. 2003), we repressed Gal4 expression until the adult stage by using *tubulin-Gal80* driver (McGuire et al. 2003) and raising the flies at 18° until eclosion, after which flies were shifted to 29° to inactive Gal80. We found that expression of UAS-Kir2.1 had no effect on the
viability of DA neurons (Figure 3A), although it did disrupt climbing ability (Figure 3B). This result suggests that a defect in synaptic transmission alone in DA neurons is capable of disrupting signaling of these neurons, but is not sufficient to cause their loss.

Additionally, if neurodegeneration and shortened lifespan in comt were due to impaired recycling of SNARE complexes involved in synaptic transmission, we would expect the phenotype to be exacerbated in combination with hyperexcitability mutants, as the pool of fusion-competent vesicles would be depleted more rapidly. However, we found that comt; seir^{2} double mutants (Jackson et al. 1985; Elkins and Ganetzky 1990; Titus et al. 1997; Wang et al. 1997) were no more severe than comt mutants alone when measuring lifespan, locomotor defects, or loss of DA neurons (Figure 3, C–E).

Although we cannot completely rule out the possibility of some synaptic defect in comt mutants at 29°C that contributes to neurodegeneration, our results are consistent with previous data showing that selectively blocking dopamine synthesis in DA neurons, which would effectively eliminate all synaptic transmission in these cells, causes no apparent deficit in their viability (Riemensperger et al. 2011). These results are also consistent with a recent study demonstrating that loss of nSyb in Drosophila photoreceptors causes degeneration that is independent of synaptic transmission defects (Haberman et al. 2012). Together, these data suggest that the neurodegeneration observed in comt mutants is likely to involve a mechanism other than synaptic transmission.

We also tested whether known binding partners of NSF1 modify the shortened lifespan and neurodegeneration phenotypes we observed. One such protein is α-SNAP (Clary and
Rothman 1990), which serves as an adaptor between NSF1 and various SNARES. We found that the shortened lifespan of comt mutants is dominantly enhanced by SNAP55 (Babcock et al. 2004), which significantly lowered the median lifespan (Figure 3F). This observation suggests that the neuroprotective role of NSF1 is still likely to involve SNARE-mediated processes.

Because comt mutants affect viability of DA neurons, which also undergo degeneration in Parkinson’s disease models, we tested whether the neuroprotective role of NSF1 intersects with one of these models. Expression of human α-synuclein in DA neurons results in progressive neurodegeneration (Auluck et al. 2002). This neurodegeneration is rescued by coexpression of heat-shock protein 70 (HSP70), which belongs to the AAA family of ATPases and is involved in protein quality control mechanisms. Since NSF1 is a member of the same family of ATPases, we tested whether NSF1 could also rescue α-synuclein-mediated loss of DA neurons. Consistent with previous studies, we found that expression of α-synuclein in DA neurons resulted in neurodegeneration by 20 days (Figure 4, A–C). This phenotype was rescued when UAS-NSF1 (Golby et al. 2001) was coexpressed in the same population of neurons (Figure 4D). To rule out the possibility that this suppression is due to the presence of multiple UAS-linked transgenes that titrate Gal4 levels and limit α-synuclein expression, we also coexpressed α-synuclein with UAS-GFP. In this case, the loss of DA neurons was identical to expression of α-synuclein alone (Figure 4E), supporting the hypothesis that it is the expression of UAS-NSF1 that alleviates α-synuclein-induced toxicity.

**comt mutants cannot sustain autophagy**

Although the neuroprotective function of NSF1 seems to be independent of its role in synaptic transmission, there are many other significant cellular functions requiring the recycling of SNARE complexes. For example, previous studies in yeast found that the NSF homolog Sec18p was required for fusion of autophagosomes to the vacuole (Ishihara et al. 2001), a critical step in autophagy. Because autophagy has been associated with neurodegeneration, we investigated the possibility that this process is impaired in comt mutants.

If comt mutants were defective in autophagy, we hypothesized that they would exhibit many of the same phenotypes as mutants known to play key roles in the autophagy pathway. Indeed, shortened lifespan and neurodegeneration have been observed in autophagy-defective (Atg) mutants in *Drosophila* (Juhasz et al. 2007; Simonsen et al. 2008). Another common phenotype among autophagy mutants is an increased sensitivity to starvation (Scott et al. 2004; Juhasz et al. 2007), due to the inability to break down stored nutrients. To examine if comt mutants are more sensitive to starvation, 3-day-old comt mutant flies were placed in empty vials with only water-soaked filter paper to prevent desiccation. Whereas control flies had a median lifespan of ~80 hr under starvation conditions, comt mutants died much sooner, with median lifespans ranging from 40 to 48 hr (Figure 5A). These data are consistent with a defect in autophagy in comt mutants.

To assay autophagy in comt mutants more directly, we examined starvation-induced autophagy in the larval fat body (Scott et al. 2004) using a tandem-tagged Atg8a reporter (UAS-mCherry-GFP-Atg8a) containing a pH-sensitive GFP and a pH-insensitive RFP to measure autophagic flux (Filimonenko et al. 2007). Because Atg8a localizes to the autophagosome, fusion with a lysosome can be seen by the loss of GFP fluorescence, leaving RFP+ puncta. We expressed this reporter in the larval fat body in wild type and comt mutants using the CG-Gal4 driver, and found that
fed larvae displayed minimal autophagic flux (Figure 5, B and C). After 3 hr of starvation, the induced autophagy response is near its peak, with several fusion events visible per cell (Figure 5, D and E, arrowheads). It is noteworthy that the initial response in comt mutants is indistinguishable from that of controls, suggesting that comt mutants are capable of normal induction of autophagy. We also monitored autophagic flux after longer periods of starvation. After 12 hr, autophagic flux in wild-type larvae is still strong. However, autophagic flux in comt mutants at this point has greatly diminished (Figure 5, F and G). In fact, we found a complete absence of autophagosomes at this point, suggesting that membrane recycling required for the formation of new autophagic compartments could be defective in comt. The fact that autophagic flux is only impaired in comt mutants following prolonged stress suggests that the defect is not with the induction of autophagy, but rather with sustaining it under periods of stress. This explanation is consistent with the role of NSF1 in synaptic transmission, where neurotransmission is normal until prolonged activity caused...
a depletion of recycled SNARE complexes for subsequent rounds of fusion (Littleton et al. 2001; Sanyal et al. 2001).

In addition to the tandem-tagged reporter, we also monitored autophagy using LysoTracker, a pH-sensitive dye often used to label lysosomes or autolysosomes (Berry and Baehrecke 2007; Scott et al. 2007). In fed larvae, there was minimal staining with LysoTracker both in cont mutants and in wild-type controls (Figure 5, H and I). After a 12-hr starvation period, however, robust staining could be seen in both cont mutants and control samples but with some important differences between the two genotypes (Figure 5, J and K). Notably, there appeared to be fewer, although much larger lysosomes in cont mutants (Figure 5, J–L, arrowheads). This enlargement of lysosomes is similar to that observed in cathepsin-D mutants (Khurana et al. 2010), as well as in a variety of lysosomal storage diseases (Fukuda et al. 2006). The enlarged lysosomes could result from a lack of lysosomal turnover. A defect in recycling membranes out of lysosomes is consistent with the absence of autophagosomes under similar conditions (Figure 5G). Alternatively, lysosomal enlargement could be a form of compensation that attempts to increase lysosomal capacity in response to the decrease in autophagy. These data provide further support for the idea that cont mutants are defective in the critical events necessary to sustain autophagy.

Loss of DA neurons in cont mutants is due to inability to sustain autophagy

Neurons are often particularly vulnerable to defects in the ability to degrade aberrant proteins and organelles. It was thus of interest to examine the role of NSF1 and autophagy in maintaining the viability of DA neurons. We knocked down expression of NSF1 in DA neurons by expressing UAS-contRNAi with TH-Gal4, along with a UAS-dicer2 transgene to enhance RNAi knockdown. Knockdown of NSF1 in DA neurons caused a loss of neurons after 48-hr exposure to 29°C comparable to that observed in cont mutants (Figure 6A). This observation indicates that the role of NSF1 in the loss of DA neurons is cell autonomous. We found a similar loss of DA neurons when expressing UAS-Atg5RNAi, showing that a reduction in autophagy is sufficient to cause degeneration of DA neurons. We also tested knockdown of three SNARE proteins recently discovered to be required for the fusion of autophagosomes to late endosomes and lysosomes (Takats et al. 2013) for their effects on DA neuron survival. We found that knocking down syntaxin17, ubisnap, or VAMP7 using TH-Gal4 resulted in loss of DA neurons (Figure 6A), demonstrating that activity of these SNARE proteins required for autophagy is necessary to maintain viability of DA neurons.

Taken together, the results presented so far support the idea that degeneration of DA neurons in cont mutants results from a defect in some step in autophagy. To test this hypothesis, we monitored autophagic flux directly in DA neurons by expressing the UAS-mCherry-GFP-Atg8a reporter using the TH-Gal4 driver. Because DA loss in cont mutants is evident by 48 hr at 29°C, we examined whether there was a decrease in autophagic flux at earlier time points. When flies were maintained at room temperature, we noted autophagic flux in nearly all neurons both in cont mutants and controls (Figure 6, B, C, and F). We saw similar autophagic activity when control flies were placed at 29°C for 36 hr (Figure 6D). In cont mutants, however, we found that ~30% of DA neurons showed little to no autophagic activity (Figure 6E, arrow). Importantly, the percentage of neurons lacking autophagic activity closely matches the percentage of those ultimately lost in cont mutants (Figure 6F and Figure 2G). From these observations, we conclude that a decrease in autophagic flux precedes neurodegeneration of DA neurons in cont mutants.

Lysosomal trafficking is disrupted in cont mutants

As described above, LysoTracker staining indicated that acidification of lysosomes was normal (Figure 4K), but this technique does not measure the degradative capacity of the

Figure 4 NSF1 is protective against α-synuclein-mediated toxicity of dopaminergic neurons. (A) Average number of dopaminergic neurons in each PPL1 cluster for 20-day-old flies raised at 25°C expressing α-synuclein in dopaminergic neurons alone, or coexpressed with NSF1 or GFP using TH-Gal4. (B–E) Representative images of PPL1 clusters for each genotype. Dopaminergic neurons are stained with anti-tyrosine hydroxylase. Bar in E, 10 μm. (**P < 0.01, Student’s t-test with Bonferroni correction for multiple comparisons.)

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Figure 5  Autophagy is defective in comatose mutants. (A) Lifespan analysis of comt mutant flies and controls during prolonged starvation at room temperature (22°C). (B–G) Images of fat body in well-fed larvae (B and C) and after starvation for 3 hr (D and E) or 12 hr (F and G). Bar in G, 10 μm. (H–K) LysoTracker (red) and DAPI (blue) staining of fat body in well-fed larvae (H and I) and after 12-hr starvation (J and K). (L) Average size of LysoTracker-positive puncta for each genotype. Bar in I, 20 μm. J’’’ and K’’’ are enlarged images of boxed areas in J and K, respectively. (***P < 0.01, log-rank test.)
lysosomes. Therefore, to determine more precisely the step at which autophagy could be disrupted, we examined lysosomal trafficking in \textit{comt} mutants using pan-neuronal expression of a LAMP1-GFP fusion protein. LAMP1-GFP is trafficked from the Golgi to late endosomes and lysosomes, where hydrolases degrade the GFP signal (Pulipparacharuvil \textit{et al.} 2005). In wild-type brains, proper lysosomal trafficking prevents the accumulation of LAMP1-GFP (Figure 7A). In \textit{comt} brains, however, we found widespread accumulation of LAMP1-GFP throughout the brain (Figure 7B). The failure to efficiently degrade the GFP signal indicates that lysosomal trafficking is impaired in \textit{comt} mutants.

The ability of lysosomes to carry out their degradative functions depends on the delivery of cargo, including cathepsins (Futerman and van Meer 2004). Cathepsins are lysosomal proteases that are synthesized in a higher molecular weight pro-form, which then undergo proteolytic cleavage into a mature form in acidic compartments such as autophagosomes or lysosomes (Lee \textit{et al.} 2010). The pro- and mature-forms of cathepsins, such as cathepsin-L, can be easily distinguished on a Western blot, providing a measurement of trafficking of cathepsins to the lysosome. We found that mature cathepsin-L was reduced by \(~75\%\) in \textit{comt} mutants (Figure 7, C and D), which is consistent with impaired delivery to the lysosome. If production of mature cathepsin-L is reduced, this might be expected to result in excess accumulation of the uncleaved pro-form of cathepsin-L. We did not observe this increase but instead found a decrease in pro-cathepsin L in \textit{comt} mutants (Figure 7C). However, compared with wild-type controls, the decrease in pro-cathepsin-L levels in \textit{comt} was less than that of the mature-form (Figure 7, D and E) and the ratio of the pro-form to the mature-form is significantly higher in \textit{comt} mutants (Figure 7E). While the reason for lower levels of mature cathepsin-L are unclear, there may be a decrease in synthesis or stability of the pro-form, in addition to the defect in cathepsin-L trafficking in \textit{comt} mutants. Taken together, these data suggest that trafficking of lysosomal proteases is impaired in \textit{comt} mutants, which accounts at least in part for the diminished degradative capacity of lysosomes in these mutants.

**Discussion**

The occurrence of progressive neuronal loss, particularly of DA neurons, in \textit{comt} mutants has enabled us to uncover and characterize a novel neuroprotective role for NSF1. \textit{comt} mutants display stress-induced locomotor defects, severely shortened lifespan, and neurodegeneration. This neurodegeneration is apparently independent of the role of

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**Figure 6** Loss of dopaminergic neurons in \textit{comatose} mutants is due to a defect in autophagy. (A) Average number of dopaminergic neurons in each PPL1 cluster with knockdown of NSF1, ATG5, or SNAREs mediating autophagosome–lysosome fusion after 48 hr at 29°C. \textit{UAS-Dicer2} was used to enhance RNAi knockdown. Controls were Gal4 and UAS transgenes alone. (B–E) Dopaminergic neurons expressing \textit{UAS-mCherry-GFP-Atg8a} in either a wild-type or \textit{comatose} mutant background after 0 hr (B and C) or 36 hr (D and E) at 29°C. Bar in C, 10 μm. (F) Percentage of dopaminergic neurons with at least one autophagic RFP+ punctum. (\(*P < 0.05, **P < 0.01\), Student’s \(t\)-test with Bonferroni correction for multiple comparisons.)
NSF1 in synaptic transmission similar to what has been observed in nSyb mutants (Haberman et al. 2012). Autophagy is disrupted in comt mutants, due at least in part to a defect in trafficking of cathepsin-L to lysosomes. In addition to cathepsin-L, it is likely that other lysosomal proteases such as cathepsin-D are also affected. Flies lacking cathepsin-D exhibit several key features of neuronal ceroid lipofuscinoses, a lysosomal storage disorder, including the accumulation of storage material and neurodegeneration (Myllykangas et al. 2005).

Most of our focus on neurodegeneration here has centered on the loss of DA neurons for several reasons: the stereotypical clusters of these neurons in the brain are amenable to quantitative analysis of neurodegeneration; they are a particularly valuable subset of neurons for studying mechanisms of neurodegeneration because of their vulnerability to insults such as oxidative stress; and they are especially relevant to human neurodegenerative disease because DA neurons are preferentially lost in Parkinson’s disease. However, the widespread accumulation of LAMP1-GFP throughout the brain (Figure 7B), reveals that disruption of lysosomal degradation is not limited to DA neurons. Thus, it will be interesting to examine the consequences of decreased NSF1 function in various other subsets of neurons and to determine whether DA neurons are for some reason more susceptible to impairments in autophagy and lysosomal trafficking. If so, we anticipate that association studies for Parkinson’s disease in humans will eventually uncover hits in genes affecting these pathways. In this context, it is also of interest that overexpression of NSF1 rescues α-synuclein-induced toxicity of DA neurons in a fly model of Parkinson’s disease.

Although our results indicate that neurodegeneration in comt mutants is not a secondary consequence of impaired synaptic transmission, there are similarities in the role of NSF1 between the two processes. For example, NSF1 is critical for the recycling of SNARE complexes involved in synaptic transmission, allowing for subsequent rounds of fusion. Similarly, there are a number of SNARE-dependent fusion events required for efficient endolysosomal signaling. These include the trafficking of lysosomal proteins, as well as the fusion between autophagosomes and lysosomes. Our results suggest that recycling of these SNARE complexes requires NSF1 for disassembly and that disassembly is
required to allow subsequent rounds of fusion. This idea is consistent with our demonstration that α-SNAP as well as the SNARE proteins syntaxin17, ubisnap, or VAMP7, are also required to maintain viability of DA neurons.

We have demonstrated that trafficking of cathepsin-L, a lysosomal protease, is disrupted in cont mutants. Although this disruption is likely to contribute to the observed defects in autophagic degradation in cont mutants, it should be emphasized that cathepsin-L merely serves as a convenient marker to assess lysosomal trafficking. There is no reason to believe that trafficking of this protein is specifically impaired by cont or that this impairment alone is responsible for the observed defects in neuronal maintenance in cont mutants. Indeed, it is likely that trafficking of many, if not most, proteins required for autophagic degradation is disrupted in cont mutants. As such, it is difficult to determine the primary defect in cont mutants because there are numerous fusion events involved, all of which are likely to require NSF1 for recycling. Thus, cont mutants exhibit deficits at multiple stages of the degradative pathway. The earliest problems to arise would likely be fusion events involving SNAREs with the smallest available pool, or those that require the most frequent recycling.

There is still much to be discovered about which SNARE proteins are involved in which specific fusion events in the cell. Recent studies examining specific fusion events have been very successful, however, including the discovery of particular SNARE complexes required for fusion of autophagosomes with lysosomes (Takats et al. 2013). Matching SNARE proteins to a particular fusion could be complex, however. As illustrated by the recent characterization of retinal degeneration in nSyb mutants in flies (Haberman et al. 2012), there is precedent for the same SNARE complex proteins independently mediating various distinct fusion events.

In summary, our results demonstrate a neuroprotective role for NSF1, which is required to maintain autophagy under stress. Our findings highlight the requirement of recycling SNARE complexes to maintain the integrity of the degradative pathway and advance our understanding of the cellular mechanisms that may underlie a number of neurodegenerative disorders.

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