Sonic hedgehog promotes autophagy in hippocampal neurons

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Summary
The Sonic hedgehog (Shh) signaling pathway is well known in patterning of the neural tube during embryonic development, but its emerging role in differentiated neurons is less understood. Here we report that Shh enhances autophagy in cultured hippocampal neurons. Microarray analysis reveals the upregulation of multiple autophagy-related genes in neurons in response to Shh application. Through analysis of the autophagy-marker LC3 by immunoblot analysis and immunocytochemistry, we confirm activation of the autophagy pathway in Shh-exposed neurons. Using electron microscopy, we find autophagosomes and associated structures with a wide range of morphologies in synaptic terminals of Shh-exposed neurons. Moreover, we show that Shh-triggered autophagy depends on class III Phosphatidylinositol 3-kinase complexes (PtdIns3K). These results identify a link between Shh and autophagy pathways and, importantly, provide a lead for further understanding the physiology of Shh signaling activity in neurons.

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Introduction
The Sonic hedgehog (Shh) signaling pathway is an evolutionarily conserved system that contributes to multiple phases of neural development. At early embryonic phases, Shh signaling plays an essential role in pattern formation of the neural tube (Ingham and McMahon, 2001; Varjosalo and Taipale, 2008). At later embryonic stages, Shh is involved in maintenance of interneuron identity in the ventral telencephalon (Xu et al., 2010). Shh signaling has also been appreciated as a growth factor for postnatal cerebellar granule cell precursors (Dahmane and Ruiz i Altaba, 1999; Wallace, 1999; Wechsler-Reya and Scott, 1999) and adult neural progenitors located in several specific stem cell niches (Lai et al., 2003; Palma et al., 2005; Breunig et al., 2008; Han et al., 2008).

While many studies have focused on functions of the Shh signaling pathway on neural progenitors, its function in differentiated neurons is less well understood. However, recent studies have revealed Shh’s roles in neurons that no longer have stem-cell-like properties. For example, Shh synergizes with NGF to provide trophic support to cholinergic neurons in the basal forebrain (Reilly et al., 2002), and protects mesencephalic dopaminergic neurons against neurotoxins (Tsuboi and Shults, 2002; Dass et al., 2005). Also, Shh is actively involved in adult nigrostriatal circuitry (Gonzalez-Reyes et al., 2012). Shh, which originates from mesencephalic dopaminergic neurons, is necessary for supporting dopaminergic neurons as well as cholinergic and fast spiking GABAergic neurons of the striatum (Gonzalez-Reyes et al., 2012).

Shh and its receptors, Patched (Ptc) and Smoothed (Smo), are also expressed in hippocampal neurons (Traiffort et al., 1999; Sasaki et al., 2010; Petralia et al., 2011a; Petralia et al., 2011b). In a previous study, we found that Shh induces structural and functional changes in presynaptic terminals of cultured hippocampal neurons (Mitchell et al., 2012). However, the molecular mechanisms underlying Shh signaling in hippocampal neurons are unknown. We conducted a pilot study to survey the global gene expression pattern of cultured hippocampal neurons after adding Shh. We noticed an increased expression of a number of genes related to the autophagy pathway, including the well-established autophagy marker LC3 (supplementary material Table S1). This observation was surprising, given no direct evidence to date linking the Shh signaling pathway to autophagy in neurons. For this reason, we focused this study on addressing the question of whether Shh activates the autophagy pathway in hippocampal neurons. Using immunoblotting, immunocytochemistry and electron microscopy, we found that Shh elicits autophagic activity that requires class III Phosphatidylinositol 3-kinase complexes (PtdIns3K).

Results and Discussion
We investigated autophagy activity in cultured hippocampal neurons that had been exposed to ShhN for 48 hr (9–10 DIV+48 hr) (supplementary material Fig. S1A). Bioactivity and efficacy of the ShhN have been verified by Shh-light2 assay (Mitchell et al., 2012). We focused on the 48 hr-ShhN-exposure because this experimental condition increased expression of...
autophagy-related genes (supplementary material Table S1), and also levels of Ptch (Mitchell et al., 2012) and Gli1 (supplementary material Fig. S1B) – indicators of an activated Shh signaling pathway (Hillman et al., 2011; Dom et al., 2012).

Using immunoblot analysis, we examined the conversion of nonlipidated LC3-I to lipidated LC3-II, a widely used autophagy indicator (Klionsky et al., 2012; Menzies et al., 2012). LC3-II level was low in untreated control neurons. In ShhN-treated neurons, LC3-II level was increased approximately 2-fold as compared to control (Fig. 1A,B; supplementary material Fig. S1C). Likewise, the Shh agonist SAG (Shh agonist) (Chen et al., 2002) also increased LC3-II level (Fig. 1A,B; supplementary material Fig. S1C). LC3-II induction was blocked by co-application of the Shh inhibitor robotnikinin (Stanton et al., 2009) (Fig. 1C), suggesting that ShhN accounts for the autophagy induction. We also tested the effects of 10-NCP, a small compound shown to effectively induce neuronal autophagy (Tsvetkov et al., 2010). 10-NCP (1 μM) in the absence of ShhN – increased LC3-II level in neurons (Fig. 1D), in agreement with results of a previous study (Tsvetkov et al., 2010). When 10-NCP was added to ShhN-treated neurons, it elicited a further LC3-II increase (Fig. 1D, lane 1 and 2; supplementary material Fig. S1A,D). A lower dose of 10-NCP (1 μM) produced a smaller but similar trend of increase in the LC3-II level (Fig. 1D, lane 3 and 4). To determine whether elevated LC3-II level reflects an increase in autophagosome synthesis or a reduction in autophagosome degradation, we tested the effect of ShhN when the last step of autophagy-mediated degradation was inhibited by bafilomycin A (BFA) (Yamamoto et al., 1998). On exposure to BFA, LC3-II level elevated further in ShhN-treated neurons (Fig. 1D, lane 5 and 6; supplementary material Fig. S1A,C), indicating that ShhN induced autophagy.

We then examined LC3-associated autophagic structures by immunocytochemistry. Neurons were identified based on their distinctive morphology and confirmed by expression of the neuronal marker MAP2. Untreated control neurons displayed little LC3 immunoreactivity (Fig. 1E,F). In contrast, addition of ShhN produced a robust increase in the number of LC3-immunopositive puncta (Fig. 1E,F; see supplementary material Fig. S2A–C for additional examples at high- and low-magnification). Robotnikinin blocked the formation of LC3-positive structures (Fig. 1E,F).

In addition to the soma of the neurons, LC3-positive puncta were seen dotted along neurites of the ShhN-treated neurons (Fig. 1G,H; also supplementary material Fig. S3A). These data suggest that ShhN induces autophagy in hippocampal neurons.
discrete puncta likely correspond to synaptic terminals of cultured hippocampal neurons (supplementary material Fig. S3B) (Dean and Scheiffele, 2009; Fletcher et al., 1994; Petralia et al., 2013). Indeed, and as discussed below, autophagosomes are found in the synaptic terminals.

We examined another autophagic/lysosomal pathway marker, the late endosome-specific lipid lysobisphosphatidic acid (LBPA) (Matsuo et al., 2004; Pérez-Sala et al., 2009). LBPA-immunolabeled structures were rarely seen in control neurons (Fig. 1I). By contrast, application of ShhN substantially increased the number of LBPA-labeled structures, and this increased even further upon 10-NCP induction (Fig. 1I,J; see supplementary material Fig. S4 for low-magnification images).

In order to identify autophagosomes more definitively and to visualize synaptic terminals directly, we next examined these neurons at higher resolution using electron microscopy. We used older neurons (18–19 DIV+48 hr ShhN) as their synapses are more mature. In synaptic terminals of control neurons, it was uncommon to find clearly identifiable autophagic structures, irrespective of pre- or post-synaptic distinction (data not shown). In contrast, synaptic terminals of ShhN-treated neurons contained a wide range of endocytic structures and autophagic structures (Fig. 2) (Boland et al., 2008; Eskelinen et al., 2011; Fader and Colombo, 2009; Liou et al., 1997; Ylä-Anttila et al., 2009). Endocytic structures included late sorting endosomes (Fig. 2A,E,G), as well as multivesicular bodies, which are characterized by the presence of distinctive intraluminal vesicles (mv in Fig. 2A–D, Fig. 2G). Autophagic structures or vacuoles displayed different morphological characteristics, representing autophagosomes at various stages (Boland et al., 2008; Eskelinen et al., 2011; Fader and Colombo, 2009; Liou et al., 1997; Ylä-Anttila et al., 2009). Fig. 2E shows an example of initial autophagic vacuoles (avi) based on the presence of cytoplasmic constituents surrounded by multiple layers of membranes. Fig. 2C,E show examples of degradative autophagic vacuoles (avd), which are filled with degrading or degraded components. Fig. 2G is a striking example of a ShhN-treated presynaptic terminal that houses a continuous sequence of (from left to right) a late sorting endosome (se), a multivesicular body (mv), and an autophagosome with a portion in the early stages of degradation on the left (avi) and a portion in later stages on the right (avd). Furthermore, Fig. 2F shows an example of a presynaptic terminal containing phagophores (ph) – a fully formed one on the left and a partially formed (U-shaped) one on the right.

Fig. 2. Electron micrographs showing various stages in the formation of autophagosomes and associated structures at synapses of ShhN-treated hippocampal neurons. (A) A presynaptic terminal (p) contains a late sorting endosome (se) and a multivesicular body (mv). A postsynaptic spine (s) apposing to the presynaptic terminal has a clathrin-coated pit (c). (B) Another example of a presynaptic terminal containing a multivesicular body. The postsynaptic spine appears to be degenerating. (C) A presynaptic terminal contains a multivesicular body – but this example has additional irregular membranes, possibly representing a combination of a multivesicular body with an autophagosome to form an amphisome. Notice that a process to the lower right is filled with a late or degradative autophagosome (avd) complex with the characteristic osmiophilic structures. (D) This synapse has a multivesicular body in the presynaptic terminal and an autophagosome complex filling much of the spine. (E) A presynaptic terminal expansion of an axon containing a late sorting endosome (se) on the left, an early stage autophagosome (avi) near the center, and a late stage autophagosome (avd) on the right. (F) Another example of presynaptic terminal expansion but it contains two putative phagophores (ph), and a clathrin-coated pit (c) nearby. (G) A presynaptic terminal expansion contains a continuous sequence (from left to right) of a late sorting endosome, a multivesicular body, an early autophagosome, and a late stage autophagosome filled with osmiophilic contents. Scale bar: 100 nm.

Fig. 3. Electron micrographs showing presynaptic autophagosomes in hippocampal neurons treated with ShhN. (A–D,F) Examples of early autophagosomes (avi). (D) A synapse with a presynaptic terminal (p) and a postsynaptic spine (s). Above this typical synapse, a large process is almost completely occupied by autophagosomes resulting in a few barely visible obscured vesicles. (E) An example of a late autophagosome (avd) based on the characteristic osmiophilic contents. Scale bar: 100 nm. (G) Quantification of presynaptic terminals that contain a multivesicular body or autophagosome. n=229 presynaptic terminals for control; n=229 presynaptic terminals for ShhN-treated.
We focused our analysis on the presynaptic terminals of the ShhN-treated neurons (Fig. 3). Many presynaptic compartments were crowded with, or nearly filled by autophagic structures, resulting in the resident synaptic vesicles being pushed against the presynaptic membrane or almost completely obscured (Fig. 3C–F). We surveyed clearly traceable presynaptic compartments (229 for control and ShhN-treated) and counted those containing multivesicular bodies or autophagosomes. ShhN treatment induced a marked increase in the occurrence of these structures in presynaptic terminals (Fig. 3G).

In addition to their presence in the presynaptic terminals (Figs 2, 3), various autophagosomes were also found in other parts of the ShhN-treated neurons (supplementary material Fig. S5). Supplementary material Fig. S5A shows a U-shaped phagophore (ph) in a presynaptic terminal, and a process adjacent to this presynaptic terminal is also filled with multiple autophagosomes. Supplementary material Fig. S5B shows an array of degenerative phagophores with whirls of multilayered membranes residing in a cell soma. Taken together, our observations suggest that Shh induces autophagy pathway activation in hippocampal neurons.

We went on to test whether the observed Shh-induced autophagy was mediated through phosphatidylinositol 3-kinase 3-kinase complexes (PtdIns3K) that are known to be essential during autophagy (Chen and Klionsky, 2011). Different classes of PtdIns3Ks exert distinctly different effects: class I PtdIns3K inhibits autophagy, whereas class III PtdIns3K stimulates autophagy (Petiot et al., 2000). The PtdIns3K signaling has been also linked to the Shh signaling pathway (Kenney et al., 2004; Riobo et al., 2006). We first asked whether class I PtdIns3K was involved in ShhN-induced autophagy. Upon ShhN- or SAG-treatment, while increased LC3-II was once again detected, the level of Akt phosphorylation was not noticeably different between treated neurons and untreated control neurons (Fig. 4A). These results indicate that class I PtdIns3K is not directly involved or does not play a prominent role in Shh-induced autophagy in the hippocampal neurons.

We then examined class III PtdIns3K by adding 3-methyladenine (3-MA) to ShhN-treated neurons. 3-MA blocks the class III PtdInsK activity necessary for autophagy induction (Seglen and Gordon, 1982). Addition of 3-MA effectively prevented ShhN-elicited LC3 elevation (Fig. 4B; supplementary material Fig. S1D), indicating that Shh-induced autophagy in hippocampal neurons is possibly mediated through class III PtdIns3K. We also used a fluorescent reporter, FYVE-DsRed. The FYVE domain is a zinc finger protein domain that binds PtdIns(3)P specifically (Kutateladze et al., 1999). It has been used as a reporter for levels of intracellular PtdIns(3)P, and thus is a sensor for activity of class III PtdIns3K (Zhang et al., 2007). We transiently transfected neurons (9–10 DIV) with FYVE-DsRed and treated these neurons with ShhN (10%) for 48 hr. Confocal microscope images showed that, without addition of ShhN, transfected neurons have diffuse fluorescence signals with occasional punctate structures (Fig. 4C; supplementary material Fig. S6A). In contrast, with addition of ShhN, transfected neurons displayed large punctate FYVE-DsRed structures and quantification showed an approximately 2-fold increase in FYVE fluorescence intensity (a.u., arbitrary unit) in the soma of neurons (n=45 neurons from 3 experiments). **P<0.001 by Student’s t test.

score (Fig. 4C,D; additional examples in supplementary material Fig. S6B). Moreover, the increased FYVE was completely blocked by 3-MA (Fig. 4C,D; also supplementary material Fig. S6B). Thus, our data suggest that Shh-triggered autophagy requires class III PtdIns3K activity.

Our results present evidence that hippocampal neurons can respond to Shh signaling by upregulating autophagy activity. In a previous study while addressing the subcellular location of the
Ssh’s receptor Ptch, we repeatedly – and puzzlingly – noticed that Ptch immunolabeling was often located in close vicinity to autophagosomes that were found in normal hippocampal neurons of adult brain (Petralia et al., 2011a). This intimate physical association of Ptch and autophagosomes is not, in and of itself, sufficient to determine whether Ptch affects autophagosome formation or functions, or whether Ptch is being processed or degraded by autophagosomes. The new data described here imply that the association may be more than merely physical: that a functional association between Ptch – or the Shh signaling pathway – and autophagosomes probably exists. While our findings do not directly address this question, they provide the impetus to design functional experiments in future studies.

Two recent studies reported that Shh signaling indeed regulates autophagy (Li et al., 2012; Jimenez-Sanchez et al., 2012). One discrepancy, however, between these two studies was how the autophagy pathway responded to Shh signaling activity. While the study by Li et al., found enhanced autophagy by Shh that is consistent with our findings, the study by Jimenez-Sanchez et al., reported the opposite – suppressed autophagosome synthesis by Shh signaling. This contradiction could result from differences in experimental conditions and designs such as the extent and duration of Shh pathway activation, the basal autophagy level, the method used in inducing autophagy, and perhaps most physiologically relevant, the type of cells studied.

The biological significance of the interaction between the Shh signaling pathway and autophagy in neurons is not yet clear. In neural stem cells, Shh serves as a mitogen (Lai et al., 2003; Palma et al., 2005; Breunig et al., 2008; Han et al., 2008), although it is unknown if it is mediated through autophagy. In differentiated neurons, Shh-enhanced autophagy could be responsible for some of Shh’s functions, including the newly described role in presynaptic differentiation of hippocampal neurons (Mitchell et al., 2012), and in circuit formation and maintenance of other types of neurons in the adult brain (Gonzalez-Reyes et al., 2012; Harwell et al., 2012). The chief function of autophagy is to remove damaged proteins and organelles (Mizushima, 2007). In addition, autophagy participates in normal turnover of various intracellular components (Mizushima, 2007), a critical process that takes place in developing synaptic terminals, and is used by growing neurons. Indeed, autophagy has been reported to promote synapse formation in Drosophila (Shen and Gansetzy, 2009), although whether Shh activity is involved has not been tested. Given that autophagy also serves as a cellular adaptive response to stress, it is not surprising that autophagy has been shown to protect neurons against degeneration in animal models of Alzheimer’s disease (Yang et al., 2011) and ischemic stroke (Wang et al., 2012). Finally, one might speculate that Shh-induced autophagy could also control or modify Shh signaling activity, a feedback regulatory mechanism analogous to the mechanism used by the Wnt signaling pathway (Gao et al., 2010). In total, this study demonstrates that the interaction between Shh signaling and autophagy may be involved not only in autophagy regulation, but also in regulation of Shh signaling activity in neurons.

Materials and Methods

Hippocampal neuron culture and reagents

All animal procedures were approved by the NIA Animal Care and Use Committee, and complied with the NIH Guide for Care and Use of Laboratory Animals. Cultures of hippocampal neurons were prepared from embryonic day 18 rat brains, and dissociated neurons were grown in Neurobasal medium/B27 as described (Matthson et al., 1989; Kaeche and Banker, 2006; Bushlin et al., 2008).

Shh-N-conditioned medium from HEK 293 cells was prepared as described (Chen et al., 2002). Control medium was prepared from non-transfected HEK 293 cells. Throughout this study, we use ShhN to refer to Shh-N-conditioned medium. ShhN- or control-medium was used at 10%. The efficacy of ShhN was validated using the Shh-light2 assay (Taipale et al., 2000; Mitchell et al., 2012). SAG (Shh agonist) was purchased from Axcora (ALX-279-426) and used at 100 nM (Chen et al., 2002). Robotnikinin was from BioVision and used at 50 μM (Stanton et al., 2009). 10-NCP was from Calbiochem, 124020. BFA and 3-MA were from Sigma. LC3 antibody was kindly provided by Dr Jay Debnath (UCSF) and also purchased from Cell Signaling. LBPA antibody was from Exchelon Biosciences, Inc. (Z-SLBPA). MAP2 antibody was from Sigma. FYVE-DsRed was a kind gift of Dr Junying Yuan (Harvard Medical School).
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