Coordinate Regulation of Mature Dopaminergic Axon Morphology by Macroautophagy and the PTEN Signaling Pathway

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Abstract

Macroautophagy is a conserved mechanism for the bulk degradation of proteins and organelles. Pathological studies have implicated defective macroautophagy in neurodegeneration, but physiological functions of macroautophagy in adult neurons remain unclear. Here we show that Atg7, an essential macroautophagy component, regulates dopaminergic axon terminal morphology. Mature Atg7-deficient midbrain dopamine (DA) neurons harbored selectively enlarged axonal terminals. This contrasted with the phenotype of DA neurons deficient in Pten – a key negative regulator of the mTOR kinase signaling pathway and neuron size – that displayed enlarged soma but unaltered axon terminals. Surprisingly, concomitant deficiency of both Atg7 and Pten led to a dramatic enhancement of axon terminal enlargement relative to Atg7 deletion alone. Similar genetic interactions between Atg7 and Pten were observed in the context of DA turnover and DA-dependent locomotor behaviors. These data suggest a model for morphological regulation of mature dopaminergic axon terminals whereby the impact of mTOR pathway is suppressed by macroautophagy.

Citation: Inoue K, Rispoli J, Yang L, MacLeod D, Beal MF, et al. (2013) Coordinate Regulation of Mature Dopaminergic Axon Morphology by Macroautophagy and the PTEN Signaling Pathway. PLoS Genet 9(10): e1003845. doi:10.1371/journal.pgen.1003845

Editor: Gregory S. Barsh, Stanford University School of Medicine, United States of America

Received February 23, 2013; Accepted August 14, 2013; Published October 3, 2013

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Funding: This work was supported by NIH grants NS064433-01, NS060876, and NS082068-01 to AA; Michael J. Fox Foundation to AA; and the Kanae Foundation for the Promotion of Medical Science and Research Foundation ITSUU Laboratory to KI. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

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Introduction

Macroautophagy is an intracellular protein degradation mechanism that engulfs cytoplasmic constituents and entire organelles within double-membrane vesicles and delivers these to lysosomes [1,2]. Genetic deletion of the essential macroautophagy components Atg5 or Atg7, during mouse central nervous system (CNS) development, leads to neuronal loss and inclusion formation [3,4]. Furthermore, Atg7 deficiency confined to cerebellar Purkinje cells leads to dystrophic axons and subsequent cell death within several weeks [5,6]. In addition to pathological roles, protein degradation pathways may also play important physiological functions in neurons. Several studies have underscored the role of cytoplasmic protein degradation through the ubiquitin-proteosome system (UPS) in the regulation of neuronal morphology and function [7]. However, the role of macroautophagy in this context is unclear.

A key regulator of neuronal morphology and size is PTEN (phosphatase and tensin homolog) [8,9], an intracellular lipid phosphatase that opposes phosphatidylinositol 3-kinase (PI3K) activity. Deletion of Pten in mouse hippocampus neurons disinhibits the mTOR (mammalian target of rapamycin) signaling pathway, leading to morphological enlargement as well as altered synaptic plasticity and plasticity-related behaviors [10–13]. mTOR is a major activator of protein translation as well as a key inhibitor of macroautophagy, but the relative contribution of these different downstream mechanisms on neuronal size remain unclear. Thus, we hypothesized that altered macroautophagy may play a role in the regulation of neuronal morphology and function in the mammalian CNS, either downstream of or in conjunction with the PTEN/PI3K/mTOR pathway.

Cell size regulation in the mammalian CNS appears to be highly dependent on developmental stage, cell type, and subcellular region. The deletion of Pten at embryonic stages leads to a profound enlargement of neurons and glia [10,11], whereas the deletion at later developmental stages – such as postnatally or in young adult animals – appears to have a lesser impact [12,13]. Postnatal deletion of Pten in certain neuronal subtypes fails to alter the size of neurite processes [13–15], suggesting selective subcellular and developmental regulatory mechanisms. For instance, deletion of Pten in post-mitotic midbrain dopamine (DA) neurons leads to soma hypertrophy, but axonal terminal morphology appears unaltered [14,15].

Here we investigated the role of Atg7 and macroautophagy in the regulation of mature midbrain DA neuron morphology, and contrasted this with the impact of the PI3K/mTOR pathway. Atg7 deficiency in mature DA neurons led to enlargement of axon terminals, whereas DA neuron soma size was only modestly altered. This phenotype was distinct from that of Pten-deficient...
mature midbrain DA neurons, which showed robust soma hypertrophy but no significant alteration at axon terminals. Mature midbrain DA neurons deficient in both Pten and Atg7 showed a dramatic enhancement of the axon terminal enlargement phenotype seen with Atg7 deficiency alone. A similar synergistic genetic interaction was similarly observed between Pten and Atg7 in the context of DA metabolism (turnover) in the striatum, and with respect to DA-associated locomotor behaviors. Taken together, these data support a model whereby macroautophagy activity normally limits the impact of the PTEN/PI3K/mTOR pathway, such that mature dopaminergic axon terminals are unaffected by Pten loss. However in the absence of macroautophagy activity, the impact of the PI3K/mTOR pathway is unmasked and leads to a profound further enlargement.

Results

Generation of Atg7 deficient mice specifically within mature midbrain DA neurons

Mice deficient in Atg7 specifically within mature midbrain DA neurons (DatCre/+;Atg7flox/flox [Atg7 cKO]) were generated [16] by interbreeding mice that express Cre recombinase (CRE) under the dopamine transporter promoter (DatCre/+; Figure 1B) [17] with mice that harbor Atg7 allele flanked by loxP sites (Atg7flox/flox, Figure 1A) [18]. Mutant animals appeared grossly normal and survival was not significantly altered (data not shown). To confirm alteration in macroautophagy activity, we initially quantified the alteration in macroautophagy activity, we initially quantified the lipidated conversion of LC3 (LC3-II), a marker for autophagy, and found that the lipidated conversion of LC3 was significantly reduced in 2-month-old midbrain tissues including substantia nigra (Figure 1C). LC3 conversion was significantly reduced in 2-month-old midbrain cKO mouse midbrains (Figure S1A). Reduction in LC3-II was only partial, likely due to ATG7 activity in non-dopaminergic cKO mouse midbrains (Figure S1A). Reduction in LC3-II was only partial, likely due to ATG7 activity in non-dopaminergic neurons (however, declined progressively from 2 month of age, and approximately 50% of TH-positive cells were lost by 1 year of age [16]). Furthermore, ubiquitin (Ub)- and p62-positive inclusions were apparent in Atg7 cKO DA neuron cell bodies and dendrites from 1-month of age (Figure 1D) [16].

Midbrain DA neurons lacking Atg7 display enlarged axon terminals

Grossly enlarged TH-positive dopaminergic axon terminal structures were observed at their striatal target in Atg7 cKO mice (Figure 1E). The axon terminal enlargement was apparent from 2 weeks of age, thus preceding other phenotypes observed in Atg7 cKO mice, and did not progress with age (Figure 1E). With aging, striatal fiber density of dopaminergic axon terminals declined slowly in Atg7 cKO mice (Figure 1E), which correlated with the slowly progressive loss of midbrain DA neuron as we have previously reported [16]. The enlarged axon terminals within the striatum were stained positively with an antibody to dopaminergic presynaptic component vesicular monoamine transporter 2 (VMAT2) (Figure 2A). In contrast to the soma inclusions (Figure 1D), the enlarged axon terminals were not stained with antibodies to Ub and p62 (Figure 2B, C).

Biochemical analysis of striatal synaptosomal preparations from 2-month-old Atg7 cKO mice and their littersmates revealed predominantly unchanged levels of pre- and post-synaptic proteins including Synapsin 1, Synaptophysin, Synaptotagmin, Synaptic vesicle protein 2A (SV2A), z-Synuclein, Synaptosomal-associated protein 25 kDa (SNAP25), Syntaxin 1A, Growth associated protein 43 (GAP43), Postsynaptic density protein 95 (PSD95), and gephrin (Figure 3C). Levels of early endosomal compartment markers present at presynaptic terminals, including early endosome antigen-1 (EEA1) and Rab5, were reduced, whereas late endosomal/lysosomal markers including Rab7 and Cathepsin B appeared unchanged (Figure 3B). An additional presynaptic marker protein, Synaptobrevin II, appeared increased in accumulation in Atg7 cKO synaptosomal preparations (Figure 3C). Thus, macroautophagy deficiency in midbrain DA neurons leads to axonal terminal enlargement associated with modest alterations in the accumulation of presynaptic regulatory proteins.

Consistent with these findings, immunoelectron microscopy for DA neuron marker, TH, showed significant enlargement of Atg7 deficient dopaminergic axon terminals but otherwise normal appearing morphology, including presynaptic terminals, synaptic vesicles, and mitochondria (Figure 2D). Furthermore, no inclusions or membrane swirls were apparent, in contrast to those described within dystrophic neuronal terminals in Atg7 deficient Purkinje neurons [5]. Thus, the morphology phenotype does not appear to be a consequence of the accumulation of unfolded protein in the context of aberrant degradation, nor represent dystrophic changes as described in other neuronal populations deficient in Atg7 [5].

We sought to further address whether the axon terminal morphological change in the context of Atg7 deficiency is a consequence of altered development or altered maintenance of mature axon terminals. Thus, adult 2-month-old Atg7flox/flox mice, which remain intact for Atg7 expression, were stereotactically injected with adenovo-associated virus-2 (AAV2) that harbors Cre/ green fluorescence protein (GFP) or GFP control into the ventral midbrain unilaterally (Figure 4A), effectively transducing a large fraction of TH-positive DA neurons (Figure 4B). Analysis of these mice 8 weeks after viral transduction revealed the dramatic enlargement of dopaminergic axon terminals within the striatum only of the AAV2-Cre/GFP virus transduced Atg7flox/flox mice (Figure 4C), consistent with the phenotype in Atg7 cKO mice. Thus, these studies confirm a role for Atg7 in the morphological plasticity of mature dopaminergic axon terminals.
In addition to the axon terminal enlargement, at time points as early as 1-month of age, morphometric analysis of TH-positive cell soma in Atg7 cKO mice (or Atg7 cWT mice) revealed a significant albeit minor increase in soma size (15% increase, Figure 5A, B). This phenotype appeared similar in older animals (and thus not age-dependent; Figure 5A), and thus was not correlated with the neurodegenerative phenotype seen with aging.

As axon terminal enlargement appeared non-progressive and was not associated with intracellular inclusions or other apparent pathological changes, this was unlikely to be secondary to the late-onset progressive degeneration seen in midbrain DA neurons of Atg7 cKO mice [19,20]. We note that in vitro studies using primary midbrain cultures prepared from Atg7 cKO or littermate cWT embryos further support this interpretation. TH-positive DA neurons...
neurons in Atg7 cKO cultures at day 5 in vitro (5th DIV) displayed increased total neurite length (Figure S1B). Importantly, TH-positive DA neuron number and appearance was otherwise not altered in the Atg7 cKO primary cultures (Figure S1B), and thus this early phenotype is not likely to reflect degeneration.

mTOR pathway modification in Atg7 deficient dopaminergic axon terminals

The mTOR kinase signaling pathway is a key regulator of mammalian cell size [8,9,21–23]. As mTOR is also a major negative regulator of macroautophagy [24], we hypothesized that Atg7 may function in a common pathway with mTOR, or parallel to mTOR, in the context of dopaminergic axon terminal size regulation. To this end, we evaluated mTOR pathway activation within dopaminergic axon terminal projections in the striatum of Atg7 cKO or cWT mice. Crude striatal synaptosomal protein fractions were prepared and analyzed by Western blotting. As expected, Atg7 cKO synaptosomes displayed evidence of reduced macroautophagy, as the lipidation of LC3 (as well as of the related autophagosome marker protein GABARAPL1) was reduced (Figure 3A). However, the pattern of mTOR pathway component modification was not consistent with canonical activation of the

Figure 2. Characterization of enlarged TH-positive axon terminals of Atg7 cKO mice. (A) Enlarged axon terminals in the striatum of 2-month-old Atg7 cKO mice were positive for the axon terminal proteins of midbrain DA neurons. Enlarged axon terminals (arrows in green) in Atg7 cKO mice were stained with VMAT2 (arrows in red). Bars, 10 μm. (B, C) Enlarged axon terminals in 2-month-old Atg7 cKO mice are Ub- and p62-negative. Enlarged axon terminals (arrows in green) in the striatum of Atg7 cKO mice were not stained with the markers for protein inclusions such as Ub (red in 'B') and p62 (red in 'C'), suggesting that they are distinct from the inclusions seen in the cell somas of Atg7 cKO mice (Figure 1D). Bars, 10 μm. (D) Ultrastructural analysis of dopaminergic axon terminals in the striatum of 3-month-old Atg7 cWT or Atg7 cKO mice by immunoelectron microscopy with an antibody to TH. Red circles indicate the gold particle-conjugated anti-TH antibody. Bars, 200 nm. (right) Quantification of the size distribution of axon terminals in striatal sections. Each dot (○) represents approximately 1.6% of the total axon terminal number. n = 611 terminals for cWT and 592 terminals for cKO sections.

doi:10.1371/journal.pgen.1003845.g002

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PI3K/mTOR pathway. Phosphorylation of eukaryotic translation initiation factor 4E binding protein 1 (4EBP1), a downstream target and effector of mTOR signaling [25], was significantly increased at Ser 65 and Thr 70 in Atg7 cKO mice (Figure 3D), but phosphorylation of other typical downstream targets of mTOR – ribosomal protein S6 kinase (S6K) (Thr 389) and S6 (Ser 235/236) – appeared unchanged (Figure 3D). Furthermore, canonical PI3K pathway-associated modifications of mTOR kinase – in terms of the accumulation of phosho-mTOR (Ser 2448 and Ser 2481; Figure 3D, 6C) – or of the upstream PI3K pathway component AKT kinase (Ser 473), were not evident in Atg7 cKO striatum (Figure 3D). Taken together, these findings argue against a simple model whereby Atg7 deficiency may modify axonal process morphology through the modification of PI3K/mTOR downstream pathway (Figure S3A). A caveat to the interpretation of mTOR pathway modification using striatal synaptosome extracts.
is that non-dopaminergic axon terminals are also present; however, Atg7 deletion was restricted to midbrain DA neurons.

Pten deficient midbrain DA neurons display enlarged cell soma but unaltered axon terminals

We next sought to more directly compare the roles of PI3K/mTOR pathway modification and macroautophagy in the context of dopaminergic axon terminal size. To this end, we generated mice deficient in Pten specifically within midbrain DA neurons (DatCre+/+Ptenfloxflox [Pten cKO]). As expected, Pten cKO midbrain sections displayed canonical activation of PI3K/mTOR pathway, quantified in terms of accumulation of phospho-AKT at Ser 473, phospho-mTOR at Ser 2448 and phospho-S6 at Ser 235/236 (Figure 6B) [14,15]. In contrast with Atg7 cKO mice, Pten cKO mice displayed no significant alteration in axon terminal size in the striatum (Figure 7C) [14]. Nonetheless, Pten cKO mice displayed robustly increased soma size of DA neurons (30% increase, Figure 7D), consistent with two prior studies of Pten deficiency [14,15], and which was much more profound than the modest soma alteration in the context of Atg7 deficiency (Figure 7D). Thus, although both Pten deficiency and Atg7 deficiency modify cell morphology, the phenotypes are distinct, and thus the effect of Atg7 deficiency cannot simply reflect altered PI3K/mTOR pathway activation alone.

Synergistic enlargement of axon terminal size in Atg7 and Pten double deficient DA neurons

To further consider the genetic relationship of Atg7 and Pten in the context of axon morphology, we generated double mutant mice lacking both Atg7 and Pten specifically in midbrain DA neurons (Atg7/Pten double cKO), and compared these to single cKO mice (either Atg7 cKO or Pten cKO alone) as well as to control WT mice (Figure 7A). PI3K/mTOR pathway activation was apparent in the Atg7/Pten double cKO mice, as expected, with accumulation of phospho-AKT at Ser 473, phospho-mTOR at Ser 2448, and phospho-S6 at Ser 235/236 in midbrain DA neurons (Figure 6D) comparable to that observed in the Pten single cKO mice (Figure 6B). Surprisingly, dopaminergic axon terminals were dramatically larger in the double cKO mice than those in Atg7 cKO mice or control animals (Figure 7C). GIANT axon terminals in Atg7/Pten double cKO mice were positive to VMAT2, but negative to Ub and p62 (data not shown). Atg7/Pten double cKO mice also showed mildly potentiated soma enlargement (79% increase, Figure 7D) relative
to either single cKO mice. Thus, PI3K/mTOR pathway activation alone is not sufficient to modify dopaminergic axon terminal size, but its impact is unmasked in the context of Atg7 deficiency (Figure 8E). A further consequence of Pten deficiency in the context of Atg7 loss is that subsequent progressive midbrain DA neuron degeneration, as seen in 2-month-old Atg7 cKO mice and thereafter, is suppressed (Figure 7B). As such ‘rescue’ of neurodegeneration in Atg7/Pten double cKO mice failed to prevent the enlarged axon terminal phenotype (but instead actually enhanced the enlargement; Figure 7C), this further validates the notion that axon terminal enlargement in Atg7 cKO mice is not a consequence of neurodegeneration.

Synergistic impact of Atg7 and Pten deficiency on DA metabolism and DA-associated locomotion

The studies above detail a synergistic role for Atg7 and Pten in the regulation of DA neuron morphology. We sought to expand the synergistic effects to functional changes in Atg7 cKO mice. Striatal DA accumulation and its metabolites, 3,4-dihydroxypheynlactic acid (DOPAC) and homovanillic acid (HVA), were quantified in 3-month-old single and double mutant mice (Figure S2A–C). DA turnover (DOPAC/DA and HVA/DA) in the striatum, which is a reflection of dopaminergic axon terminal activity, was increased in Atg7 cKO mice (relative to control cWT mice; Figure 8A, B). This phenotype was further enhanced in...
**Discussion**

The PTEN/PI3K/mTOR signaling pathway plays a central role in the regulation of neuronal morphology and size in developing vertebrate and invertebrate species [8,9,21,22]. However, a number of studies have provided evidence that in the context of the mature mammalian CNS, as well as within certain subcellular compartments of neurons such as axonal processes, the impact of the PTEN/PI3K/mTOR pathway on size can be highly regulated. For instance, although deletion of PTEN in post-mitotic DA neurons leads to enlarged soma (Figure 7D) [14,15], dopaminergic axon terminals are not altered (Figure 7C) [14,15]. In contrast to these observations, PTEN deletion at earlier developmental stages or in other neuronal types, such as in the progenitors of dentate gyrus granule neurons [10,11], leads to

**Figure 7.** Atg7 and Pten double deficiency synergistically increases axon terminal size in midbrain DA neurons. (A) Schema of mouse mating to obtain Atg7 and Pten double deficient mice. DatCre/+ background animals were used for the analyses. Animals lacking Cre (Ptenflox/*Atg7flox/*)) were not used in this study. Asterisk indicates ‘+’ or ‘flox’. (B) Neurodegeneration in Atg7 cKO mice was rescued by secondary deletion of Pten. Secondary Pten deletion (Atg7/Pten double cKO) suppressed the loss of TH-positive DA neurons in the substantia nigra of 2-month-old Atg7 cKO mice. Representative TH-stained midbrain sections are presented. Bar, 250 μm. (right) Quantification of TH-positive DA neuron number in the substantia nigra of Atg7/Pten double cKO mice. n = 4 per genotype. **, p<0.01. (C) The enlarged axon terminal phenotype of Atg7 cKO mice was greatly enhanced in Atg7/Pten double cKO mice, whereas Pten deficiency alone (Pten cKO) did not significantly change the axon terminal size. (left) Giant (arrowheads, >9.8 μm²) and moderately enlarged (arrows, 4.4–9.8 μm²) axon terminals were seen in the striatum of Atg7/Pten double cKO mice, whereas only moderately enlarged axon terminals (arrows) were seen in Atg7 cKO mice and no enlarged axon terminals were seen in Pten cKO mice. Bars, 20 μm. (right) Quantification of enlarged axon terminal distribution. Black bar, giant terminals (>9.8 μm²); white bar, moderately enlarged terminals (4.4–9.8 μm²). n = 6 per genotype. **, p<0.01. (D) The soma of TH-positive DA neurons in Atg7/Pten double cKO mice were dramatically enhanced (79% increase versus control cWT) relative to Atg7 cKO mice (15% increase versus control cWT) and Pten cKO mice (32% increase versus control cWT). (left) Representative sections stained with anti-TH antibody. Bars, 10 μm. (right) Quantification of the average cell size, presented as a fraction of DA neuron soma size in control cWT mice. n = 290–417 TH-positive DA neurons per genotype. **, p<0.01.

doi:10.1371/journal.pgen.1003845.g007

Atg7/Pten double cKO mice, whereas Pten cKO alone appeared normal (Figure 8A, B), mirroring the morphological findings. The absolute level of DA accumulation, in contrast to DA turnover, was significantly reduced in Atg7 cKO mice (Figure S2A), which may reflect cell loss at this age. However, this reduction in DA levels was not ‘rescued’ by PTEN loss (Figure S2A). Given the altered DA accumulation in Atg7 cKO mice, as well as prior studies demonstrating physiological changes in these animals [26], we sought to identify possible behavioral correlates. To this end, basal locomotor activity was quantified in an open field chamber that was novel to the animals over a 30-min period. Ambulatory distance travelled was increased in Atg7 cKO mice (relative to control cWT mice; Figure 8C, D). This phenotype was further enhanced in Atg7/Pten double cKO mice (Figure 8C, D), whereas Pten single cKO mice behavior did not appear significantly altered (Figure 8C, D) [14]. Other activity parameters, including jump counts and vertical activity, appeared similarly altered in the Atg7 cKO mice and Atg7/Pten double cKO mice (Figure S2E, F). In contrast, velocity of ambulation was not altered (Figure S2D). Thus, Atg7 single cKO and Atg7/Pten double cKO mice displayed alterations in motor behavior that correlated with their cellular changes and alterations in DA turnover.
prominent enlargement of axon terminals as well as soma. Thus, mechanisms that regulate the impact of the PI3K/mTOR pathway on neuronal morphology are of particular interest. Here we show that deletion of the essential macroautophagy component Atg7 unmasks the impact of Pten deletion on dopaminergic axon terminal size. These data implicate macroautophagy as a negative regulator of PTEN/PI3K/mTOR pathway regulation of neuronal morphology (Figure 8E).

Although the precise mechanism by which Pten deletion or PI3K pathway activation leads ultimately to cellular hypertrophy remains unclear, multiple studies in Drosophila [27] and mice [23,28,29] have implicated activation of the ribosomal protein S6 by a family of related kinases that include S6K1 and S6K2, leading to increased protein translation [30]. As neither S6 phosphorylation nor activation of S6K appeared modified in the context of Atg7 deficiency (Figure 3D), the axonal process enlargement seen with macroautophagy deficiency is likely to be through a distinct mechanism. Our genetic studies strongly validate this notion, as PI3K/mTOR pathway activation (by means of PTEN deficiency) and Atg7 deficiency act cooperatively and synergistically in modifying axon terminal morphology. We favor an interpretation whereby at the axonal terminus, macroautophagy-mediated protein degradation is typically able to overcome the increased protein production in the context of Figure 8E.

Figure 8. Atg7 and Pten double deficiency synergistically increases DA turnover and DA-associated behaviors. (A, B) DA turnover, quantified as the ratio of the DA metabolites DOPAC or HVA to DA (DOPAC/DA [A] or HVA/DA [B]) was synergistically increased in Atg7/Pten double cKO mice. The concentrations of DA, DOPAC, and HVA are shown in Figures S2A–C. n = 10–12 per genotype. **, p < 0.01. (C, D) Basal locomotor activity was synergistically increased in Atg7/Pten double cKO mice. Traces (D) display total distance traveled over a 30-min period in an open field environment. Walking velocity was unchanged (Figure S2D). n = 10–12 per genotype. *, p < 0.05; **, p < 0.01. (E) In midbrain DA neurons at baseline, Atg7-mediated macroautophagy inhibits the enlargement of dopaminergic axon terminals, in part by masking the impact of the PTEN/PI3K/mTOR pathway. Also see Figure S3.

doi:10.1371/journal.pgen.1003845.g008
mTOR pathway activation, and thus macroautophagy typically suppresses the impact of Pten deletion on dopaminergic axon terminal size. However, in the context of defective macroautophagy with Atg7 deficiency, the impact of Pten deletion is unmasked (Figure 8E). In addition to suppressing PTEN/PI3K/mTOR pathway-mediated regulation, macroautophagy likely plays additional roles in determining dopaminergic axon terminal size and function, as the impact of Atg7 deficiency was observed regardless of Pten deficiency (Figure 7C).

Our data argue against an alternative model whereby mTOR pathway activation dictates dopaminergic axon terminal morphology through the downstream inhibition of macroautophagy (rather than through downstream effects on the translation machinery) (Figure S3A), as has been suggested based on the recent in vitro analyses of acutely prepared striatal slice preparations treated with the mTOR inhibitor rapamycin [26]. The impact of the PTEN/PI3K/mTOR pathway on mature dopaminergic axon terminal morphology in our present study was apparent in the complete absence of macroautophagy (when comparing Atg7 single mutant mice with Atg7/Pten double mutant mice), and thus the mechanism of action cannot be explained simply by alterations in macroautophagy activity. It remains possible that alterations in macroautophagy activity play some role downstream of mTOR pathway activation in the context of more acute physiological changes at axon terminals [26]. Nonetheless, our findings support a distinct model whereby macroautophagy plays a key role in suppressing the impact of mTOR pathway activation in the context of mature dopaminergic axon terminals (Figure 8E).

Previous studies have reported that midbrain DA neuronespecific Atg7 loss [19], as well as loss of Atg7 in other neuronal classes, leads to enlarged but dystrophic axons [5,6]. These findings have generally been interpreted as secondary effects of the accumulation of pathological inclusions (‘engorgement’). However, in our analyses of Atg7 deficient midbrain DA neurons, dopaminergic axon terminal enlargement preceded degeneration, appeared non-progressive (Figure 1E), and was not associated with protein aggregates (Figure 2B, C). Furthermore, this phenotype was enhanced – rather than suppressed – in the context of additional Pten deficiency (Atg7/Pten double deficient animals) (Figure 7C), although such additional Pten deficiency effectively suppressed the late-onset degeneration phenotype of the Atg7 deficient midbrain DA neurons (Figure 7E) [16].

Prior studies in invertebrate species have suggested a role for macroautophagy in axon terminal morphology. For instance, at the Drosophila neuromuscular junction (NMJ), macroautophagy has been reported to promote synapse development through selective degradation of the Highwire ubiquitin ligase [31]. Mutations in C. elegans unc-51 – a macroautophagy regulator – lead to developmental axonal defects [32]. There is also precedent of a role for macroautophagy in cell size homeostasis: induction of macroautophagy by Atg7 leads to the reduced Drosophila fat body cell size in TOR signaling-dependent manner [33]. In mammalian models, loss of macroautophagy-associated proteins other than Atg7 has similarly been implicated in axonal morphology, although the mechanism has remained unclear. Mammalian Ulk1/2 orthologues of Unc51, have been reported to regulate axonal outgrowth [34–36]. Taken together, we speculate that regulation of axon terminal size by macroautophagy may play an important role in structural plasticity at mature adult axon terminals. Extrinsic cues such as glial derived neurotrophic factor (GDNF) impact dopaminergic axon terminal structures and modify signaling through the PI3K pathway [37,30]; it will be of interest to pursue the role of macroautophagy in such changes, which have been implicated clinically in pathological movements associated with experimental therapeutics for Parkinson’s disease [39–42].

Future studies will seek to identify specific molecular components that may mediate dopaminergic axon terminal enlargement in the context of defective macroautophagy. Our initial screen of known axon terminal proteins revealed the increased accumulation of Synaptobrevin II in the context of Atg7 deficient dopaminergic axon terminal preparations (Figure 3C). In addition to proteins, macroautophagy also plays a role in cell membrane regulation [43], and this may also impact axon terminal morphology. Finally, it is interesting to note that axon terminal morphology, DA neurons, and the PI3K pathway [44–47] have all been implicated in the etiology of autism spectrum disorders (ASD), which are characterized by cognitive difficulties and can be associated with hyperactivity [48,49]. Furthermore, recent studies have suggested a role for alterations in the protein degradation machinery in ASD [50–52]. We thus speculate a role for macroautophagy regulation of axon terminal morphology in the context of brain disorders.

Materials and Methods

Animal

DatCre/+ mice, Atg7+/flx/+ mice, and Pten+/flx/+ mice were generated previously [17,18,53]. All animals were maintained in the animal facility of Columbia University Medical Center. All of the experimental protocols were approved by the Institutional Animal Care and Use Committees. All mice used were DatCre/+ background, as Dat heterozygous KO mice show some defects in their behavior and physiology [54].

Histology

Mice were perfused in 4% paraformaldehyde and 50 μm coronal sections were made by a vibratome. The antibodies used here were listed in Text S1.

Electron microscopy

Electron microscopic analysis was according to the previous paper [55] and Immunogold incubation protocol for general application (Electron Microscopy Sciences, Hatfield, PA).

Cell size determination

After the TH staining of midbrain sections by DAB, pictures were taken at 400× magnification. The size of TH-neuron was measured manually by Image-J (NIH) [56]. More than 200 TH-neurons from 4 mice were analyzed per group.

Western blotting

Preparation of the striatal synaptosomal fractions was according to the previous paper [57]. The antibodies used here were listed in Text S1.

High performance liquid chromatography (HPLC)

The striatal tissues were used for HPLC analysis. Concentrations of DA and its metabolites were measured according to the previous paper [58].

Statistical analysis

All of the comparisons were made with Mann-Whitney U-test (for 2 samples) or non-repeated measures ANOVA (for multiple samples). The values are expressed as the means ± SEM. A p value less than 0.05 is considered significant.
Supporting Information

**Figure S1** Characterization of enlarged axon terminals of Atg7 cKO mice. (A) Decreased macroautophagy activity in midbrain extracts from Atg7 cKO mice. The conversion of LC3-I to LC3-II was reduced in 2-month-old Atg7 cKO mice, n = 5 per genotype, *, p<0.05. (B) Increased neurite length of Atg7 cKO midbrain TH-positive primary neurons. Total neurite length was significantly increased in Atg7 cKO primary neurons (bottom right), whereas the total number of TH-positive neurons per well unchanged. Primary midbrain neuron cultures were prepared from 3 embryos per genotype. **, p<0.01. (TIF)

**Figure S2** Characterization of Atg7/Pten double cKO mice. (A–C) Concentrations of DA, DOPAC, and HVA in the striatum of Atg7/Pten double cKO mice. (A) DA. (B) DOPAC. (C) HVA. n = 10–12 mice per genotype. **, p<0.01. (D) Quantifications of the parameters in open field test. (D) Walking velocity. (E) Jump counts. (F) Vertical behavior counts. n = 10–12 per genotype. *, p<0.05. (TIF)

**Figure S3** Models for the role of macroautophagy in midbrain DA neuron. Two distinct models for the role of macroautophagy in regulating DA axon terminal morphology and function. (A) In the linear model proposed by the prior study [26], the primary action of mTOR on DA axon morphology is directly through the inhibition of macroautophagy. (B) In our sculptural model, macroautophagy plays a key role in suppressing the action of mTOR signaling at DA axon terminal morphology. (TIF)

**Text S1** Supporting materials and methods. (DOC)

Acknowledgments

We would like to thank G. DiPaolo, and O. Hobert for comments on the manuscript, T. Chiha, M. Komatsu and K. Tanaka for providing Atg7 flox/flox mice, J. Dunning for some help in mouse analysis, T. Iwasato for advice on immunostaining, T. Yamashita for valuable discussion, S. Itohara, M. Noda, K. Yamamaka, A. Yamamoto and M. Sakurai for helpful suggestions.

Author Contributions

Conceived and designed the experiments: KI AA. Performed the experiments: KJ JR LY. Analyzed the data: KI AA. Contributed reagents/materials/analysis tools: KI LY DM MFB EK. Wrote the paper: KI AA.

References

1. Levine B, Klionsky DJ (2004) Development by self-digestion: molecular mechanisms and biological functions of autophagy. Dev Cell 6: 463–477.
2. Klionsky DJ (2008) Neurodegeneration: good riddance to bad rubbish. Nature 441: 819–826.
3. Hara T, Nakamura K, Matsu M, Yamamoto A, Nakahara Y, et al. (2006) Suppression of basal autophagy in neural cells causes neurodegenerative disease in mice. Nature 441: 815–819.
4. Komatsu M, Waguri S, Chiba T, Murata S, Isawa J, et al. (2006) Loss of autophagy in the central nervous system causes neurodegeneration in mice. Nature 441: 880–884.
5. Komatsu M, Wang QJ, Helkoste GR, Friedrich VL Jr., Isawa J, et al. (2007) Essential role for autophagy protein Atg7 in the maintenance of axonal homeostasis and the prevention of axonal degeneration. Proc Natl Acad Sci U S A 104: 14489–14494.
6. Nishiyama J, Mizu E, Mizunuma N, Watanabe M, Yuzaki M (2007) Aberrant membranes and double-membrane structures accumulate in the axons of Atg7-null Parkinje cells before neuronal death. Autophagy 3: 591–596.
7. Wan HI, DiAntonio A, Fetter RD, Bergstrom K, Strauss R, et al. (2000) Highwire regulates synaptic growth in Drosophila. Neuron 26: 313–325.
8. Backman S, Stambolie V, Suzuki A, Haight J, Elia A, et al. (2001) Deletion of Pten in mouse brain causes seizures, ataxia and defects in soma size resembling Lhermitte-Duval disease. Nat Genet 29: 396–403.
9. Kwon CH, Zhu X, Zhang J, Baker SJ (2003) mTOR is required for hypertrophy of Pten-deficient neuronal soma in vivo. Proc Natl Acad Sci U S A 100: 12923–12928.
10. Suresh P, Kuralwalla-Martinez C, Guo W, Gregor C, Wang Y, et al. (2011) Selective deletion of Pten in peripheral beta-cell homestasis in vivo. Mol Cell Biol 26: 4511–4518.
11. Spires BL, Kuralwalla-Martinez C, Guo W, Gregor C, Wang Y, et al. (2011) Selective deletion of Pten in peripheral beta-cell homestasis leads to increased islet mass and resistance to STZ-induced diabetes. Mol Cell Biol 26: 2772–2781.
12. Chalhoub N, Kozma SC, Baker SJ (2006) S6k1 is not required for Pten-deficient neuronal hypertrophy. Curr Opin Neurobiol 16: 516–522.
39. Tomac A, Lindqvist E, Lin LF, Ogren SO, Young D, et al. (1995) Protection and repair of the nigrostriatal dopaminergic system by GDNF in vivo. Nature 373: 335–339.
40. Hudson J, Granholm AG, Gerhardt GA, Henry MA, Hoffman A, et al. (1995) Glial cell line-derived neurotrophic factor augments midbrain dopaminergic circuits in vivo. Brain Res Bull 36: 425–432.
41. Rosenblad C, Kirik D, Bjorklund A (2000) Sequential administration of GDNF into the substantia nigra and striatum promotes dopamine neuron survival and axonal sprouting but not striatal reinnervation or functional recovery in the partial 6-OHDA lesion model. Exp Neurol 161: 503–516.
42. Rangasamy SB, Soderstrom K, Bakay RA, Kordower JH (2010) Neurotrophic factor therapy for Parkinson’s disease. Prog Brain Res 36: 425–432.
43. Zhang M, Scheiman R (2013) Cell biology. Unconventional secretion, unconventional solutions. Science 340: 559–561.
44. Bourgeron T (2009) A synaptic trek to autism. Curr Opin Neurobiol 19: 231–234.
45. Delorme R, Ey E, Toro R, Leboyer M, Gillberg C, et al. (2013) Progress toward treatments for synaptic defects in autism. Nat Med 19: 685–694.
46. Nakamura K, Sekine Y, Ouchi Y, Tsuji M, Yoshikawa E, et al. (2010) Brain serotonin and dopamine transporter bindings in adults with high-functioning autism. Arch Gen Psychiatry 67: 59–68.
47. Gadow KD, Devineci CJ, Obert DM, Psearevskaya V, Hatchwell E (2010) Association of DRD4 polymorphism with severity of oppositional defiant disorder, separation anxiety disorder and repetitive behaviors in children with autism spectrum disorder. Eur J Neurosci 32: 1058–1065.
48. Taurines R, Schwenck C, Westervald E, Sacher M, Sinatchkin M, et al. (2012) ADHD and autism: differential diagnosis or overlapping traits? A selective review. Atten Defic Hyperact Disord 4: 113–139.
49. Rommele NN, Franke B, Geurts HM, Hartman CA, Butelaar JK (2010) Shared heritability of attention-deficit/hyperactivity disorder and autism spectrum disorder. Eur Child Adolesc Psychiatry 19: 281–295.
50. Glessner JT, Wang K, Cai G, Kovatska O, Kim CE, et al. (2009) Autism genome-wide copy number variation reveals ubiquitin and neuronal genes. Nature 456: 569–573.
51. Busan M, Abrahams BS, Wang K, Glessner JT, Herman EL, et al. (2009) Genome-wide analyses of exonic copy number variants in a family-based study point to novel autism susceptibility genes. PLoS Genet 5: e1000336.
52. Scheuerle A, Wilson K (2011) PARK2 copy number aberrations in two children presenting with autism spectrum disorder: further support of an association and possible evidence for a new microdeletion/microduplication syndrome. Am J Med Genet B Neuropsychiatr Genet 156B: 413–420.
53. Groszer M, Erickson R, Scripture-Adams DD, Lesche R, Truempy A, et al. (2001) Negative regulation of neural stem/progenitor cell proliferation by the Pten tumor suppressor gene in vivo. Science 294: 2186–2189.
54. Spielewoy C, Biala G, Rouhert C, Hamon M, Betancur C, et al. (2001) Hypolocomotor effects of acute and daily d-amphetamine in mice lacking the dopamine transporter. Psychopharmacology (Berl) 159: 2–9.
55. Mengual E, Pickel VM (2004) Regional and subcellular compartmentation of the dopamine transporter and tyrosine hydroxylase in the rat ventral pallidum. J Comp Neurol 468: 395–409.
56. MacLeod D, Dowman J, Hammond R, Leete T, Inoue K, et al. (2006) The familial Parkinsonism gene LRRK2 regulates neurite process morphology. Neuron 52: 587–593.
57. Teng L, Crooks PA, Somailla PK, Dwoskin LP (1997) Lobeline and nicotine evoke [3H]overflow from rat striatal slices preloaded with [3H]dopamine: differential inhibition of sympotomosomal and vesicular [3H]dopamine uptake. J Pharmacol Exp Ther 280: 1432–1444.
58. Yang L, Calingasan NY, Chen J, Ley JJ, Becker DA, et al. (2005) A novel azulenyl nitrone antioxidant protects against MPTP and 3-nitropropionic acid neurotoxicities. Exp Neurol 191: 86–93.
59. Ishioto T, Nomura R, Ando R, Ikeda T, Tanaka M, et al. (2004) Dorsal telencephalon-specific expression of Cre recombinase in PAC transgenic mice. Genesis 38: 130–138.