Deletion of the Huntingtin Polyglutamine Stretch Enhances Neuronal Autophagy and Longevity in Mice

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Abstract

Expansion of a stretch of polyglutamine in huntingtin (htt), the protein product of the IT15 gene, causes Huntington’s disease (HD). Previous investigations into the role of the polyglutamine stretch (polyQ) in htt function have suggested that its length may modulate a normal htt function involved in regulating energy homeostasis. Here we show that expression of full-length htt lacking its polyglutamine stretch (ΔQ-htt) in a knockin mouse model for HD (Hdh140Q/140Q), reduces significantly neuropil mutant htt aggregates, ameliorates motor/behavioral deficits, and extends lifespan in comparison to the HD model mice (Hdh140Q/140Q). The rescue of HD model phenotypes is accompanied by the normalization of lipofuscin levels in the brain and an increase in the steady-state levels of the mammalian autophagy marker microtubule-associate protein 1 light chain 3-II (LC3-II). We also find that ΔQ-htt expression in vitro increases autophagosome synthesis and stimulates the Atg5-dependent clearance of truncated N-terminal htt aggregates. ΔQ-htt’s effect on autophagy most likely represents a gain-of-function, as overexpression of full-length wild-type htt in vitro does not increase autophagosome synthesis. Moreover, Hdh140Q/140Q mice live significantly longer than wild-type mice, suggesting that autophagy upregulation may be beneficial both in diseases caused by toxic intracellular aggregate-prone proteins and also as a lifespan extender in normal mammals.

Introduction

In vertebrates, the polyQ stretch within htt is located close to the protein’s N-terminus, and separates a highly conserved 17 amino acid N-terminal domain (N1–17) that can act as a membrane association signal [1], from a proline-rich region that is implicated in protein-protein interactions [2–4]. Expansion of htt’s polyQ stretch (>37Q) causes Huntington’s disease (HD), a neurodegenerative disorder characterized by the appearance of cytoplasmic (neuropil) and nuclear aggregates of mutant htt, and selective cell death in the striatum and cortex [5–9]. Although the mechanism of pathogenesis is still unclear, HD is recognized as a toxic gain-of-function disease, where the expansion of the polyQ stretch within htt confers new deleterious functions on the protein. The extent to which the polyQ expansion affects normal htt function is also unclear, although there is accumulating evidence that loss of normal htt function likely contributes to HD pathogenesis [10]. The polyQ stretch is conserved in vertebrate htt, and its non-pathogenic size varies from 4Q in fish, to 37Q in humans [11–13]. However, the polyQ stretch is absent in Ciona and Drosophila htt, and present as only a short hydrophilic NHQQ stretch in sea urchin htt, suggesting that addition of a htt polyQ stretch may be a late evolutionary feature acquired sometime after protostome-deuterostome divergence [14].

In lymphoblastoid cell lines derived from HD patients, polyQ length (in both the normal and mutant htt alleles) affects energy status, with a longer polyQ stretch correlating with a reduced cellular ATP/ADP ratio [15]. Deletion of the normal short polyQ stretch (7Q) in mouse htt (ΔQ-htt) also results in elevated ATP levels in fibroblasts derived from embryonic and adult Hdh140Q/140Q mice [16]. In addition, adult Hdh12Q/12Q mice exhibit subtly enhanced performance on the rotarod, and altered behavior in the Barnes maze learning and memory test.

To assess ΔQ-htt function in the presence of expanded polyQ htt expression, we generated mice expressing both ΔQ-htt and 140Q-htt (Hdh140Q/140Q). We found that ΔQ-htt expression in the HD mouse model rescued behavioral/motor deficits, reduced the number of neuropil htt aggregates, normalized brain lipofuscin levels, and enhanced lifespan relative to the HD mouse model. Clearance of htt aggregates and the accumulation of lipofuscin are mediated by autophagy, a catabolic pathway that encompasses several distinct processes in mammalian cells [17]. Macroautophagy generally involves the non-selective turnover of bulk cytoplasmic contents, including organelles and aggregated protein, and is an essential pathway for the survival of organisms during nutrient deprivation [18]. Upregulation of autophagy reduces truncated mutant htt aggregation and toxicity in both in vitro and in vivo models [19–22], and recently, the acetylation of soluble...
**Author Summary**

Expansion of a stretch of glutamines near the amino-terminus of huntingtin (htt), the protein product of the IT15 gene, is a deleterious mutation that causes Huntington's disease (HD). Here we show, in contrast, that deletion of htt's normal polyglutamine stretch (ΔQ-htt) is a potentially beneficial mutation that can ameliorate HD mouse model phenotypes when ΔQ-htt is expressed together with a version of htt with the HD mutation. In addition, ΔQ-htt expression can enhance longevity when expressed in either an HD mouse model or in non–HD mice. ΔQ-htt's effects on both lifespan and HD model phenotypes are likely due to an increase in autophagy, a major recycling pathway in cells that is involved in the turnover of cellular components, and aggregated protein. Based on our results, we suggest that development of therapeutic agents that can stimulate autophagy may help both in treating neurodegenerative disorders like HD and also in increasing longevity.

full-length htt has also been reported to assist its recognition by the autophagic apparatus [23]. In Htt14Q/+ and Htt14Q/ΔQ mice, we observed enhanced microtubule-associated protein 1 light chain 3 (LC3, [24]) immunostaining, and increased levels of the LC3-II autophagic marker. Expression of ΔQ-htt, but not wild-type htt, induced the formation of autophagosomes in SK-N-SH neuroblastoma cells, and enhanced the clearance of truncated 74Q-htt aggregates in an autophagy-dependent process. Based on our observations, we hypothesize that deletion of the polyQ stretch within huntingtin enhances neuronal macroautophagy resulting in the more efficient clearance of neuropil mutant htt and phenotypic rescue in Htt14Q/ΔQ mice. Moreover, we have observed that mice homozygous for ΔQ-htt expression live significantly longer than wild-type mice, an observation that is compatible with the view that enhancing constitutive autophagy may also be beneficial in normal ageing.

**Results**

**Rescue of Htt14Q/+ motor and behavioral deficits in Htt14Q/ΔQ mice**

To evaluate the impact of expressing a version of wild-type htt lacking its short polyQ stretch on the motor and behavioral phenotypes exhibited by a mouse model for HD, Htt14Q/+ mice were crossed with the CAG140 knock-in mouse expressing full-length htt with a chimeric human/mouse htt exon 1 containing an expanded stretch of 140 glutamines [25], (for a diagram of the knockin alleles used in this study, see Figure 1A). Htt14Q/ΔQ, Htt14Q/+ and wild-type control littermates were assessed using the accelerating rotarod, the Barnes maze, and an activity cage. Mice were tested on an accelerating rotating rod at 1, 5, and 19 months of age (Figure 1B). At one month of age, there were no significant differences between the wild-type controls, Htt14Q/+ mice, and the Htt14Q/ΔQ mice (n = 6 for each genotype at 1 and 5 months, n = 4 of each genotype at 19 months). A two-way repeated measures ANOVA revealed that enhancing constitutive autophagy may also be beneficial in treating neurodegenerative disorders like HD and also in increasing longevity.

At five months of age, the Htt14Q/ΔQ/+ mice performed poorly in comparison to both the wild type control group and the Htt14Q/+ group (genotype effect; F(2,6) = 5.4; P<0.03). Interestingly, at five months, the Htt14Q/ΔQ/+ mice were indistinguishable from either other (genotype effect; F(2,6) = 6.3; P<0.04), although all mice were performing more poorly at 19 months relative to their performance at 5 months of age.

At five months of age, the mice were also tested on the Barnes maze, a measure of spatial learning and memory [26]. Wild-type mice produced better scores on the Barnes maze distance test than Htt14Q/ΔQ/+ mice, but did not differ significantly from the Htt14Q/+ mice (n = 5 of each genotype) (Figure 1C). The distance score measures how effectively the mice are using spatial cues to locate the escape tunnel. A two way repeated measures ANOVA revealed a significant effect of genotype (F(2,5) = 5.96; P<0.02) and a significant effect of trial day (F(5,25) = 2.2; P<0.04). In addition, wild-type and Htt14Q/ΔQ/+ mice made fewer errors than Htt14Q/+ mice before finding the Barnes maze target (Figure 1C). A two-way repeated measures ANOVA revealed a significant effect of genotype (F(2,5) = 25.28; P<0.001), and a significant effect of trial day (F(5,25) = 3.33; P<0.003).

At 6 and 20 months of age, the mice were also tested in an activity cage (n = 5 of each genotype) (Figure 1D). Previous analyses of Htt14Q/+ mice revealed that they exhibit a period of hyperactivity, followed by hypoactivity when tested at night in an activity cage [25]. Based on total horizontal activity, the Htt14Q/ΔQ/+ mice were more hypoactive at night than the wild-type mice at 6 months, but the exploratory activity of the Htt14Q/ΔQ/+ mice did not differ significantly from wild-type controls (one-way ANOVA F(2,12) = 6.63; P<0.02; post-hoc analysis wild-type versus Htt14Q/ΔQ/+ or Htt14Q/+ mice did not differ significantly from wild-type controls (F(2,12) = 0.87; P>0.05), although there was a significant effect of trial day (F(8,4) = 2.2; P<0.04). Bonferroni post-hoc analysis showed the Htt14Q/ΔQ/+ mice to be significantly hypoaactive when compared to the Htt14Q/+ mice (P<0.01).

Htt14Q/ΔQ/+ mice also exhibited a significant increase in their lifespan (median age of 31+/−0.8 months) in comparison to either Htt14Q/+ or Htt14Q/ΔQ mice (median ages of 24+/−2.3 and 27+/−1.7 months, respectively, Htt14Q/+ versus Htt14Q/ΔQ mice). log-rank test, χ² = 11.7, P<0.002; Htt14Q/+ versus Htt14Q/ΔQ mice). log-rank test, χ² = 9.9, P<0.003 for n = 8 females of each genotype) (Figure 2). However, we could not detect any significant difference in the lifespan of the Htt14Q/ΔQ/+ and Htt14Q/ΔQ/+ mice (log-rank test, χ² = 0.03, P = 0.953).

**Reduction of neuropil htt aggregates in Htt14Q/ΔQ/+ mice**

To determine if the rescue of behavioral phenotypes in the Htt14Q/ΔQ/+ mice correlated with a change in the number and distribution of htt aggregates, we examined Htt14Q/+ and Htt14Q/ΔQ/+ (control) brains (n = 4 of each genotype) using an antibody recognizing aggregated mutant htt in inclusions (MW8 [27]) (Figure 3A). At 4 months of age, we were unable to detect htt aggregates in either Htt14Q/+ or Htt14Q/+ mouse brain (Figure 3B). At 1 year and 2 years of age, the aggregate load increases dramatically in the Htt14Q/+ brain, with the number of striatal neuropil aggregates growing more quickly with age than the number of nuclear aggregates (Figure 3B). The significant reduction in the number of striatal neuropil aggregates that was observed at 6 months of age in the Htt14Q/ΔQ/+ brain in comparison to the Htt14Q/+ brain, P<0.01 (Figure 3B). At 1 year and 2 years of age, the aggregate load increases dramatically in the Htt14Q/+ brain, with the number of striatal neuropil aggregates growing more quickly with age than the number of nuclear aggregates (Figure 3B). The significant reduction in the number of striatal neuropil aggregates that was observed at 6 months of age in the Htt14Q/ΔQ/+ brain in comparison to the Htt14Q/+ brain, P<0.01 (Figure 3B).
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Figure 1. ΔQ-htt expression ameliorates motor and behavioral deficits in Hdh^{140Q/ΔQ} mice. (A) Diagram of the Hdh^{140Q}, wild-type (Hdh^{7Q}), and Hdh^{140Q} exon 1, and a schematic of the breeding scheme used to generate the mice employed in this study. The expansion of the polyQ stretch in the Hdh^{140Q} allele also includes human exon 1 sequences (purple) between a conserved XmnI restriction site within exon 1 and a KpnI restriction site located within intron 1, while the deletion of the polyQ stretch in the Hdh^{ΔQ} allele also includes the insertion of a FLAG epitope tag (green) following the initiation codon. Endogenous mouse sequence is shown in light blue. (B) Accelerated rotarod testing of wild-type (+/+) and Hdh^{140Q/+} (140Q/+) mice (n = 6 for each genotype). Hdh^{140Q/+} versus +/+ at 5 months, ANOVA for genotype; P<0.03 and at 19 months; P<0.04. The performance of the Hdh^{140Q/ΔQ} mice did not differ significantly from the wild-type controls. (C) Barnes maze testing of 5 month old wild-type (+/+) and Hdh^{140Q/+} and Hdh^{140Q/ΔQ} mice (n = 5 of each genotype). Hdh^{140Q/+} mice performed poorly on their distance scores compared to wild-type and Hdh^{140Q/ΔQ} mice (ANOVA for genotype; F(2,4) = 5.96, P<0.02). Hdh^{140Q/+} and wild type mice also made significantly fewer errors than Hdh^{140Q/ΔQ} mice before finding the target (ANOVA for genotype; F(2,4) = 25.28; P<0.001). *Significant differences on individual trial days between Hdh^{140Q/ΔQ} and Hdh^{140Q/+} mice (Holm-Sidak post hoc analysis; P<0.001 to 0.05). (D) Horizontal activity in a novel environment was assessed in wild-type, Hdh^{140Q/+}, and Hdh^{140Q/ΔQ} mice (n = 5 of each genotype) at 6 and 20 months of age. At both ages, the Hdh^{140Q/ΔQ} mice were significantly more hypoactive than the Hdh^{140Q/+} and wild-type mice (one-way ANOVA F(2,12) = 6.63, P<0.02 and Bonferroni post-hoc analysis, P<0.02 at 6 months; one-way ANOVA (F(2,14) = 6.78, P<0.02 and Bonferroni post-hoc analysis Hdh^{140Q/ΔQ} versus Hdh^{140Q/+} at 20 months, P<0.01).

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respectively. In the cortex, a similar marked decrease in Hdh140Q/DQ neuropil aggregates was observed at 6 months, 1 year, and 2 years of age ($P_{0.001}$ to $P_{0.005}$) (Figure S1). In both striatum and cortex, nuclear aggregates were also reduced significantly at 1 year of age, but the magnitude of the decrease was less than that observed for the neuropil aggregates.

Lipofuscin deposits are reduced in Hdh140Q/DQ mice

Increased lipofuscin has been observed in the HD brain and in the R6/2 transgenic mouse model for HD [28–30]. Accumulating in the lysosomes of neurons and other post-mitotic cells, lipofuscin is a yellowish-brown autofluorescent aging pigment that is composed of oxidized lipid and aldehyde cross-linked protein [31]. Lipofuscin is believed to be the byproduct of the incomplete autophagic catabolism of cellular organelles, such as mitochondria that are rich in iron. Iron and peroxide-catalyzed oxidation of incompletely digested lipid and protein results in the slow accumulation of lipofuscin in autolysosomes at a rate that correlates with metabolic activity and age of the organism [32]. In HD, oxidative stress may enhance the formation of lipofuscin, resulting in the appearance of large perinuclear lipofuscin deposits in neurons. In aged cells with high levels of lipofuscin, autophagy is diminished [33,34], and in C. elegans, lower levels of lipofuscin in age-matched worms correlated with greater motility, suggesting that lipofuscin accumulation reflects biological versus chronological age [35].

We compared the extent of lipofuscin accumulation in the striatum and cortex of wild-type, Hdh140Q+/+, Hdh140Q/140Q, and Hdh140Q/140Q mice at 4 months, 6 months, 1 year, and 2 years of age (n = 8 mice of each genotype) (Figure 4A and 4B). Consistent with prior observations in the R6/2 HD transgenic mouse model and in postmortem HD brain tissue, we observed a significant increase in lipofuscin (measured as the pixel area of deposits in confocal images) in the striatum and cortex of Hdh140Q/140Q mice as they aged in comparison to wild-type mice, $P_{<0.05}$ to $P_{<0.001}$ (Figure 4B). Lipofuscin accumulation was greater in the striatum,

Figure 2. Hdh<sup>140Q</sup> expression enhances longevity in a knockin mouse model for HD. Kaplan-Meier survival curves are shown for Hdh<sup>140Q</sup>/+, Hdh<sup>140Q/140Q</sup>, and Hdh<sup>140Q/140Q</sup> mice (n = 8 mice of each genotype). The Hdh<sup>140Q/140Q</sup> mice lived significantly longer than either the Hdh<sup>140Q/140Q</sup> mice or the Hdh<sup>140Q/+</sup> mice (log-rank test; $\chi^2 = 9.9$, $P_{<0.003}$ and $\chi^2 = 11.7$, $P_{<0.002}$, respectively). The lifespans of the Hdh<sup>140Q/140Q</sup> and Hdh<sup>140Q/140Q</sup> mice did not differ significantly ($\chi^2 = 0.03$, $P_{= 0.958}$).

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Figure 3. Reduced htt neuropil aggregates in the Hdh<sup>140Q/140Q</sup> striatum.

(A) Representative confocal images of the striatum from Hdh<sup>140Q</sup>/+, Hdh<sup>140Q/140Q</sup>, and Hdh<sup>140Q/140Q</sup> mice at 4 months, 6 months, 1 year, and 2 years of age that were immunostained with an antibody recognizing htt aggregates (MW8, red signal). Nuclei were stained with To-Pro-3 (blue). Enlarged images of the areas enclosed by dashed white boxes are shown in the bottom panels. Open and solid white arrowheads indicate neuropil and nuclear aggregates, respectively. Scale bars = 25 µm (top panels), 10 µm (bottom three panels). (B) Total; T, nuclear; N, and neuropil; C, htt aggregate numbers from the Hdh<sup>140Q/140Q</sup> striatum (n = 4 mice of each genotype). The aggregate numbers (mean ± s.e.m.) represent counts/field from 8 images of the ventral and lateral striatum from each mouse. *$P_{<0.05}$, **$P_{<0.001}$.

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relative to the cortex in the Hdh\textsuperscript{140Q/+} brain. In both the Hdh\textsuperscript{140Q/+} cortex and striatum, however, neuronal lipofuscin accumulation was similar to that observed in wild type controls at all ages examined.

Altered autophagy in Hdh\textsuperscript{DQ/+} and Hdh\textsuperscript{140Q/DQ} mice

To determine if clearance of the neuropil htt aggregates and the reduction in lipofuscin in the Hdh\textsuperscript{140Q/DQ} brain may be related to altered autophagy, we performed immunohistochemical analyses and western blot analyses of cellular fractions obtained from wild-type, Hdh\textsuperscript{DQ/+}, Hdh\textsuperscript{140Q/+}, and Hdh\textsuperscript{140Q/DQ} whole brains and dissected brain regions, respectively, using an antibody to LC3. LC3 is encoded by the mammalian homolog of the yeast Atg8 gene, and is widely used as a marker for autophagy in mammalian cells because it associates tightly with autophagic membranes beginning at vesicle nucleation, and ending with its turnover in autolysosomes [24]. Western blotting with antibodies recognizing the N-terminus of LC3 detects two species with apparent molecular weights of 18 kD (LC3-I) and 16 kD (LC3-II). LC3 is processed proteolytically at its C terminus to form cytosolic LC3-I, which is conjugated to phosphatidylethanolamine on autophagosome membranes to form LC3-II. LC3-II associates specifically with autophagosome and autolysosome membranes, and LC3 vesicle numbers or levels of LC3-II correlate with autophagosome numbers [24,36].

LC3 immunostaining was enhanced in the striatum of Hdh\textsuperscript{140Q/DQ} mice beginning at 6 months of age in comparison to age-matched wild-type, Hdh\textsuperscript{DQ/+}, and Hdh\textsuperscript{140Q/+} mice (n = 4 of each genotype, Figure S2). At 1 year of age, the Hdh\textsuperscript{140Q/DQ} striatum continued to exhibit enhanced LC3 immunostaining, and at 2 years of age, elevated LC3 immunostaining was now detected in both the Hdh\textsuperscript{DQ/+} and Hdh\textsuperscript{140Q/DQ} striatum (Figure 5A). In contrast, LC3 immunostaining in the Hdh\textsuperscript{140Q/DQ} striatum at 1 year and 2 years of age was not increased substantially in comparison to age-matched wild-type controls. Moreover, co-localization of LC3 immunostaining with neuropil htt aggregates was observed in the Hdh\textsuperscript{140Q/DQ}

Figure 4. ΔQ-htt expression in Hdh\textsuperscript{140Q/+} mice normalizes the increased lipofuscin accumulation that is observed in the Hdh\textsuperscript{140Q/+} brain. (A) Confocal images of the parietal cortex and striatum from one year old Hdh\textsuperscript{10Q/+}, Hdh\textsuperscript{140Q/+}, and Hdh\textsuperscript{140Q/DQ} mice (n = 4 of each genotype). Lipofuscin deposits are yellow (white arrowheads), and nuclei are stained with To-Pro-3 (blue). Scale bar = 25 μm. (B) Lipofuscin deposits at 4 months, 6 months, 1 year, and 2 years of age in wild-type (+/+), Hdh\textsuperscript{DQ/+}, Hdh\textsuperscript{140Q/+}, and Hdh\textsuperscript{140Q/DQ} parietal cortex and striatum were quantified by measuring their pixel area/field (mean ± s.e.m.). *P<0.05, **P<0.001. doi:10.1371/journal.pgen.1000838.g004

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striatum at 1 and 2 years of age, but was difficult to detect in the
Hdh\(^{140Q/+}\) striatum (Figure 5A).

To confirm that the enhanced LC3 immunostaining observed in the
Hdh\(^{140Q/+}\) and Hdh\(^{140Q/14Q}\) striatum was due to an increase in
LC3-II levels, dissected striata from 2 year old wild-type (+/+), Hdh\(^{140Q/+}\),
Hdh\(^{140Q/14Q}\), and Hdh\(^{140Q/14Q}\) mice (n = 4 of each genotype) were
homogenized and separated into supernatant (NP40-soluble) and
pellet (NP40-insoluble) fractions, and then analyzed by western
blotting with an antibody that recognizes both LC3-I and LC3-II
(Figure 5B and 5C). In the soluble protein fractions, an increase in
LC3-II was observed in the Hdh\(^{140Q/14Q}\) striatum. Interestingly,
LC3-II was also enriched in the striatal pellet fractions from both
Hdh\(^{140Q/+}\), Hdh\(^{140Q/14Q}\), and Hdh\(^{140Q/14Q}\) mice (n = 4 of each genotype).
Blots were stripped and re-probed with an antibody against \(\beta\)-actin to control for
protein loading. The positions of protein standards (in kD) are indicated on the left. (C) Quantification of LC3-II levels in the supernatant and pellet
fractions relative to actin. *P<0.004 versus +/+.

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Figure 5. \(\Delta Q\)-htt expression in mice enhances LC3 immunostaining and LC3-II steady-state levels. (A) Confocal images of LC3 (green)
and htt aggregate (MW8, red) immunostaining in the striatum from 1 year and 2 year old wild-type (+/+), Hdh\(^{140Q/+}\), Hdh\(^{140Q/14Q}\), and Hdh\(^{140Q/14Q}\) mice
(n = 4 of each genotype). Nuclei were stained with To-Pro-3 (blue). Enlarged images of the areas enclosed by dashed white boxes are shown in the
bottom panels. White arrowheads indicate examples of htt aggregates co-immunostaining with LC3 (yellow signal). Scale bars = 25 \(\mu\)m (top panels),
10 \(\mu\)m (bottom panels). (B) Western analyses for LC3-I and LC3-II in striatal supernatant (Sup) and pellet (Pel) fractions from 2 year old wild-type (+/+),
Hdh\(^{140Q/+}\), Hdh\(^{140Q/14Q}\), and Hdh\(^{140Q/14Q}\) mice (n = 4 of each genotype). Blots were stripped and re-probed with an antibody against \(\beta\)-actin to control for
protein loading. The positions of protein standards (in kD) are indicated on the left. (C) Quantification of LC3-II levels in the supernatant and pellet
fractions relative to actin. *P<0.004 versus +/+.

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an increase in LC3-II in both the Hdh4Q/4 and Hdh140Q/140 samples (Figure S3B). We note that we also observed an enrichment of both the autophagy protein beclin 1 and lysosome-associated membrane protein type 1 (Lamp1) levels in the 800×g low-speed P1 fractions from Hdh4Q/4 and Hdh140Q/140 striatal extracts prepared by lysis in the absence of detergent (Figure S3A). Overall levels of beclin 1 and Lamp1 in total brain extract, however, were similar in all genotypes (Figure S3B). Lamp1 is a marker for late endosomes, amphisomosomes (formed after autophagosome-late-endosome fusion), dense autolysosomes and lysosomes that are enriched in the 800×g P1 fraction [37], and these observations, together with our findings related to the alterations in beclin 1 and LC3-II fractionation, suggest that the subcellular distribution of several components of the autophagy pathway are altered by ΔQ-htt expression.

It was proposed recently, that htt’s association with the ER via its N1–17 domain allows it function as a sensor of ER stress, and to potentially regulate autophagy [1,38]. In previous work, we found no obvious difference in the nuclear/cytoplasmic localization of ΔQ-htt in comparison to wild-type htt in early passage wild-type and Hdh4Q/4Q primary mouse embryonic fibroblasts (PMEFs) [16]. To analyze further the subcellular localization of wild-type- and ΔQ-htt together with markers for the ER (calnexin), and to assess a marker for autophagy (LC3), we performed immunocytochemistry on passage 5 (P5) cultures of wild-type and Hdh4Q/4Q primary mouse embryonic fibroblasts (PMEFs) (Figure S4). P5 cultures of wild-type fibroblasts are actively dividing, while P5 cultures of Hdh4Q/4Q fibroblasts are, in contrast, undergoing replicative senescence [16]. Wild-type- and ΔQ-htt were detected in both the cytoplasm and nucleus, and perinuclear localization of wild-type- and ΔQ-htt with the ER marker, calnexin, was also detected in both Hdh4Q/4Q and Hdh4Q/4Q+PMEFs (Figure S4A, S4B). However, nuclear localization of htt appeared to be increased in those cells with a more senescent morphology (i.e. more flattened/spread appearance). Perinuclear LC3 immunoreactivity was also enhanced in the Hdh4Q/4Q PMEFS with a senescent morphology (Figure S4C), suggesting the possibility for increased autophagy in those Hdh4Q/4Q PMEFS undergoing replicative senescence.

Expression of ΔQ-htt enhances autophagosome synthesis in vitro

An alteration to autophagy resulting in the increased steady-state levels of LC3-II can be attributed to either enhanced autophagic flux, or to a block in a later step within the pathway that would interfere with the turnover of LC3-II in the autolysosome [39]. To determine if ΔQ-htt can enhance autophagosome synthesis, we transfected SK-N-SH neuroblastoma cells with full-length wild-type (7Q-htt) or ΔQ-htt cDNA expression constructs (diagrams in Figure S5), and monitored the levels of LC3-II 24 h post-transfection by western blotting (Figure 6A). The levels of LC3-II were increased significantly in the ΔQ-htt transfected cells in comparison to either control vector or 7Q-htt transfected cells. To monitor autophagy by an alternative method, we also transfected HeLa cells with an EGFP-LC3 expression construct, together with pCDNA5.1 (vector control), 7Q-htt or ΔQ-htt in a 1:3 ratio (Figure S6). The proportion of EGFP-positive cells with >10 EGFP-LC3 vesicles was assessed and expressed as an odds ratio with 95% confidence limits. ΔQ-htt transfection, but not 7Q-htt transfection, increased the proportion of cells with EGFP-LC3 vesicles. To measure autophagosome synthesis, the cDNA constructs were also transfected in the presence or absence of the antibiotic bafilomycin A1, a vacuolar H+ ATPase inhibitor that suppresses turnover of LC3-II in autolysosomes [40–42]. Thus, measuring the levels of LC3-II in the presence of bafilomycin A1 measures LC3-II formation, as the antibiotic blocks LC3-II degradation. The levels of LC3-II were increased significantly in the bafilomycin A1-treated and ΔQ-htt-transfected cells in comparison to the bafilomycin A1-treated cells alone or the bafilomycin A1-treated and 7Q-htt-transfected cells, suggesting that ΔQ-htt but not 7Q-htt expression results in increased autophagosome synthesis (Figure 6B). To confirm that an increase in LC3-II formation resulting from ΔQ-htt expression is enhancing autophagic activity that can remove another autophagy substrate, 7Q- or ΔQ-htt constructs were transfected into Agt5+/+ (autophagy-competent) and Agt5−/− (autophagy-deficient) mouse embryonic fibroblasts [43], together with an EGFP-tagged 7Q-htt exon 1 construct (EGFP-HDQ74) expressing an N-terminal fragment of mutant htt that forms aggregates readily in vitro [44]. Aggregate formation in EGFP-positive cells 48 h post-transfection was assessed by calculating odds ratios with 95% confidence limits [44–46] (Figure 6C). The proportion of cells with EGFP-HDQ74 aggregates was significantly reduced in Agt5+/+ cells transfected with ΔQ-htt, but not in Agt5−/− cells. Interestingly, 7Q-htt overexpression also reduced aggregate load in both Agt5+/+ and in Agt5−/− cells. These data suggest that while ΔQ-htt can induce autophagic clearance of mutant htt aggregates, 7Q-htt overexpression may induce a reduction in aggregate numbers or formation via an autophagy-independent mechanism in our in vitro system. Taken altogether, these data support the hypothesis that ΔQ-htt expression can stimulate autophagosome formation and the Agt5-dependent clearance of htt aggregates.

Importantly, we saw no difference in autophagy in 7Q-htt-overexpressing cells versus empty vector transfected cells (Figure 6A and 6B), or when comparing huntingtin knockout (Hdh140Q/140/Hdh4Q/4Q) mouse embryonic stem cells (Hdh/−/−) which were either transfected with empty vector of with wild-type full-length 17Q-Htt (Figure 6D), suggesting that the ability of htt to induce autophagy is a specific consequence of the loss of its polyQ tract.

ΔQ-htt expression stimulates autophagy via an mTOR–independent pathway

A central regulator of metabolism and autophagy in both invertebrates and vertebrates is TOR (Target of Rapamycin) kinase, and inhibition of TOR kinase activity by rapamycin and its analogs has been used successfully to stimulate autophagic clearance of mutant htt aggregates in both Drosophila and mouse models for HD [48]. To determine if the activity of mammalian TOR (mTOR) is inhibited by ΔQ-htt expression, we examined the phosphorylation status of mTOR in the striatum of two year old wild-type, Hdh4Q/4Q, Hdh140Q/140, Hdh140Q/140, and Hdh140Q/4Q mice, and also the phosphorylation status of downstream targets of mTOR in our in vitro system (Figure 7). Phospho-mTOR (p-mTOR) levels correlate positively with mTOR kinase activity and inversely with mTOR inhibition and the activation of macroautophagy [48], although autophagy can also be regulated by mTOR-independent pathways. We observed no difference in p-mTOR levels in the supernatant fractions of all genotypes examined (Figure 7A). However, we did detect an enrichment of p-mTOR in the striatal pellet fractions from the Hdh140Q/140 and Hdh140Q/4Q brains. This association of p-mTOR with the pellet fraction likely represents p-mTOR association with htt aggregates, as was observed previously both in vitro, and in a transgenic HD mouse model [48].

To confirm our in vivo analyses, SK-N-SH cells were transfected with either 7Q- or ΔQ-htt expression constructs and the phosphorylation status of two targets of mTOR kinase activity
Figure 6. ΔQ-htt expression increases autophagosome synthesis and Atg5-dependent reduction of EGFP-74Q truncated htt aggregates in vitro. (A) SK-N-SH cells were transfected with control vector (pCDNA3.1), 7Q-htt, or ΔQ-htt constructs (transfections performed in triplicate at least twice), and cell lysates were analyzed for LC3-II levels 24 hr post-transfection by western blotting with an anti-LC3 antibody (for comparison, both longer and shorter exposures are shown), and densitometric analysis relative to actin levels. ΔQ-htt versus control; (***P<0.001), ΔQ-htt versus 7Q-htt; (P<0.002). (B) SK-N-SH cells were transfected with control vector, 7Q-htt, or ΔQ-htt constructs and treated with either DMSO (pCDNA3.1 transfected cells) or with 400 nM bafilomycin A1 (Baf) in DMSO (pCDNA3.1, 7Q-htt, or ΔQ-htt transfected cells) for 4 hours at 20 hr post-transfection. Autophagosome synthesis was analyzed by western blotting using an anti-LC3 antibody and densitometric analysis relative to actin. ***P<0.0001, **P<0.004. (C) Atg5+/+ (autophagy-competent) and Atg5−/− (autophagy-deficient) mouse embryonic fibroblasts were transfected with an EGFP-HDQ74 construct together with vector control, 7Q-htt, or ΔQ-htt constructs, and then assessed for the proportion of EGFP-positive cells with EGFP-74Q-htt aggregates by calculating the odds ratio 48 h post-transfection. ***P<0.0001 for both ΔQ-htt versus control, and for 7Q-htt versus control in Atg5+/+ cells, P<0.0001 for 7Q-htt versus control in Atg5−/− cells. ΔQ-htt's ability to reduce aggregates is abolished in Atg5−/− cells (P=0.36 in comparison with control vector transfected cells). (D) Hdh+/−/− knock-out (Hdh−/−) mouse embryonic stem (ES) cells were transfected with EGFP along with either pCI (empty vector) or full-length wild-type huntingtin (1:5 ratio) for 4 h, and then treated with DMSO (− Baf) or 400 nM bafilomycin A1 in DMSO (+ Baf) for the last 2 h of the 48 h post-transfection period. Cells were then FACS-sorted for the EGFP-positive cells, in which LC3-II levels were assessed by western blotting and densitometric analysis relative to actin. Htt levels were assessed by western blotting using MAB2166. There were no significant changes in LC3-II levels in the huntingtin transfected Hdh−/− ES cells compared to empty vector transfected Hdh−/− ES cells in the absence (P=0.7103) or presence (P=0.4063) of bafilomycin A1.

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were assessed 24 h post-transfection (Figure 7B). The levels of phospho-S6 kinase and phospho-S6 ribosomal protein were not significantly different in the cells transfected with 7Q- or ΔQ-htt, supporting the hypothesis that ΔQ-htt’s upregulation of autophagy is not mediated by a reduction in mTOR kinase activity.

ΔQ-htt expression extends lifespan in the mouse

Our observations suggest that expression of a version of htt lacking its normal stretch of polyQ can enhance autophagic clearance of neuropil mutant htt inclusions. During normal aging, misfolded and aggregated proteins accumulate due to an apparent decline in the function of lysosomal degradation pathways [49]. In Caenohabditis elegans, autophagy is an essential component in the mechanism that extends lifespan upon dietary restriction. Knockdown of essential autophagy genes, for example, shortens lifespan in C. elegans, and suppresses lifespan extension induced by dietary restriction, reduced mitochondrial function, and alterations in insulin/IGF-1 or TOR signaling [50]. Moreover, enhancing basal levels of autophagy in the nervous system of Drosophila by Atg8a overexpression increases both longevity and resistance to oxidative stress [51]. However, the ability of autophagy upregulation to extend mammalian lifespan has not previously been tested.

To determine if ΔQ-htt expression has an effect on longevity in the absence of 140Q-htt expression, we assessed the lifespan of HdhΔQ/ΔQ mice in comparison to wild-type mice (n = 15 mice of each genotype). While the wild-type controls lived to 28 ± 1.2 months (median age ± s.e.m.), the HdhΔQ/ΔQ mice lived to a median age of 33 ± 1.1 months, representing an 18% extension of lifespan (log-rank test, c² = 9.6, P < 0.005) (Figure 8). The oldest HdhΔQ/ΔQ mouse survived for approximately 3.5 years in our colony, compared to 3 years for the oldest wild-type mouse.

Discussion

We provide evidence that expression of a version of mouse htt that lacks its short 7Q polyglutamine domain can stimulate the formation of autophagosomes in vitro and enhance the clearance of htt neuropil aggregates, ameliorate behavioral/motor pheno-
types, and extend lifespan in a mouse model for HD. When expressed in homozygosity, ΔQ-htt can also significantly extend lifespan in the mouse. Recently, it was proposed that htt may participate directly in autophagy because of its structural similarity with mTOR, and also because it co-localizes partially with autophagosomes [38]. These results, and our data, do not address directly whether or not htt has a normal function involved in autophagy. However, our results do suggest that deletion of htt’s short 7Q stretch can enhance basal autophagy, and are compatible with a hypothesis suggesting that htt’s polyQ stretch may modulate a normal function for htt in this process. Expansion of htt’s polyQ stretch beyond the pathogenic threshold may, in contrast, suppress such a function, and account for our observation that we did not observe autophagy induction in the Hdh140Q/+ brain. In this scenario, ΔQ-htt’s gain-of-function in autophagy would be dominant to a potential loss-of-function in autophagy caused by the expansion of the polyQ stretch.

The rescue of motor and behavioral phenotypes in the Hdh140Q/AC mice starting at 5–6 months of age correlates with a reduction in neuropil htt aggregates, and a normalization of lipofuscin levels. Neuritopi aggregates are an early phenotypic feature of HD, and our results are compatible with a recent study demonstrating that clearance of cytoplasmic htt aggregates by the expression of an intrabody specific for aggregated htt can rescue motor and behavioral deficits in a transgenic mouse model for HD [52]. The reduced number of nuclear inclusions observed in the Hdh140Q/AC brain at one year of age may be due to htt’s ability to shuttle between the nucleus and cytoplasm [1]. The stimulation of autophagy in other HD mouse models, for example, can also reduce htt nuclear aggregate number [48]. The normalization of lipofuscin deposits in the Hdh140Q/AC brain may be the consequence of reduced oxidative stress that is secondary to the reduction in mutant htt aggregate load. Although we had previously detected increased lipofuscin in the HdhQ/AC brain [16], we observed similar levels of lipofuscin in the HdhQ/+ and wild-type brain. As lipofuscin is an end-product of autophagy, the increased lipofuscin accumulating in the HdhQ/AC brain may represent the increased turnover of mitochondria due to the expression of two HdhQ alleles. In this case, increased lipofuscin accumulation could represent increased autophagic activity instead of increased oxidative stress. This causes a positive feedback loop resulting in further damage. There is accumulating evidence that enhanced autophagy correlates with increased longevity in C. elegans and Drosophila. In C. elegans, autophagy is an essential component in the mechanism that extends lifespan upon dietary restriction [50]. Knockdown of essential autophagy genes, for example, shortens lifespan in C. elegans, and suppresses lifespan extension induced by dietary restriction, reduced mitochondrial function, and alterations in insulin/IGF-1 or TOR signaling. In a complementary experiment, enhancing basal levels of autophagy in the nervous system of Drosophila by Agl8 overexpression increases both longevity and resistance to oxidative stress [51].

The 18% increase in HdhQ/AC lifespan relative to wild-type controls is comparable to that observed with mouse mutations in insulin signaling pathways that result in increased longevity. Mice heterozygous for a knock-out of the insulin-like growth factor 1 receptor gene (Igf1r) exhibit a 26% increase in mean lifespan [54], while mice expressing a mutant insulin receptor gene in adipose tissue exhibit an 18% increase in lifespan [55]. Mice lacking expression of the insulin receptor substrate 1 gene (Irs1-/-) also live 18% longer than wild-type mice [36], and brain-specific knock-out of one Irs2 allele in the mouse results in an 18% extension of lifespan [57]. Our findings do not permit us to determine if increased lifespan in the HdhQ/AC mice is the consequence of neuronal or global expression of ΔQ-htt. Neuronal-specific overexpression of Agl8 in Drosophila is sufficient to extend lifespan [51], but analogous experiments in mice have not yet been performed.

The effect of ΔQ-htt expression on HdhQ/AC lifespan appears, at first, to be incompatible with the premature replicative senescence phenotype that we observed in HdhQ/AC PMEFs cultured in vitro [16]. However, the detection of both increased senescence-associated (SA)-β-galactosidase staining [16,58], and LC3 immunoreactivity (Figure S4C) in senescent PMEFs supports the data of Narita and colleagues suggesting that upregulating autophagy may facilitate the mitotic senescence transition in vitro [59]. In this scenario, ΔQ-htt expression may have opposite effects on cellular senescence and mammalian lifespan. A similar, apparently contradictory, response to mammalian SIRT1 expression (a homolog of the yeast Sir2 factor involved in extending replicative lifespan) has been described in PMEFs where absence of SIRT1 expression increases replicative lifespan [60]. Thus, in these examples, replicative lifespan in vitro may not always correlate positively with organismal lifespan.

Upregulation of autophagy has the potential to be a therapeutic strategy for Huntington’s disease and related disorders. Although rapamycin and its analogs have proven to be very useful in stimulating increased clearance of N-terminal truncated mutant htt aggregates in various animal models for HD via a “pulsitile” upregulation of autophagy, our data suggest that tonic long-term autophagy upregulation via ΔQ-htt expression is not associated with overt side-effects. This genetic method for autophagy upregulation is apparently mTOR-independent based on our inability to detect a significant decrease in soluble p-mTOR levels in vivo, and alterations in the phosphorylation status of downstream mTOR kinase targets in vitro. In this regard, we have identified a series of novel compounds that influence autophagy in an mTOR-independent fashion [45,61]. Although further work is required to elucidate the pathway responsible for ΔQ-htt’s affect on autophagy, our findings support the view that the development of both genetic and small molecule-based therapeutic strategies aimed at stimulating the autophagic clearance of aggregated protein may be of use in both the treatment of neurodegenerative disease, and in lifespan extension.

**Methods**

**Generation of mice**

HdhQ/+-, HdhQ/Q-, HdhQ/Q+, and HdhQ/Q/Q+ mice were obtained from heterozygous intercrosses between HdhQ/Q+ and HdhQ/Q+ mice that were maintained in a mixed 129/Sv and C57BL6 background. HdhQ/Q and HdhQ/Q/Q+ mice were obtained from HdhQ/Q+ and HdhQ/Q+ intercrosses, respectively. All protocols for animal use were approved by the Institutional Animal Care and Use Committee of the University of Virginia, and were in accordance with NIH guidelines. For routine genotyping, PCR was used to confirm the presence of the different Hdh alleles: ΔQ allele; ΔQ-for = 5'-GACGGGCCCAAGATGGAGCGACTTGAGCGACTCGAAAG-3' and ΔQ-rev = 5'-GGCGGTGCGCCAGACTTT-3' amplify a 226 bp product from only the ΔQ allele, while Epi-for = 5'-GAGGTATGGTGGCAGTAGTGCGAGCCGCTCAAAG-3' and Epi-rev = 5'-CGGACTTTGGAAGAGAAGAAG-3' flank the site of the
FLAG epitope in the ΔQ allele and amplify either a 112 bp product from the wild-type allele or a 136 bp product from the ΔQ allele. 140Q allele; 140-for = 5’-CTGCACGGACGGT-GAGGTTCC-3’ and 140-rev = 5’-GAAGGCCAGTTCAATGCTGA-C-3’ flank a small intron-1 deletion created during the generation of the 140Q allele. A wild-type allele (or ΔQ allele) will generate a 235 bp product, while the 140Q allele will generate a 150 bp product. To verify that the mean CAG repeat length in the Hdi140Q allele was similar in the Hdi140Q/+ and Hdi140Q/ΔQ mice that were used for our analyses, the CAG repeat was amplified using CAG-1 = 5’-CTTCTGAGCTCCTGAGTCCTTCC-3’ and CAG-2 = 5’-GGTTGGCGGCGCTGTCTCTTG-3’ (data not shown). These oligonucleotides are specific for the human sequence surrounding the CAG repeat in the Hdi140Q allele and will generate a ~450 bp product using the Expand High Fidelity PCR system (Roche Molecular Diagnostics).

Motor and behavioral analyses
The accelerating rotarod test was performed at 1, 3, and 19 months of age (n = 6 mice of each genotype at 1 and 5 months of age, and n = 4 mice of each genotype at 19 months of age; all mice in the same cohort) as described [16]. At each time-point, there were 3 separate testing sessions of 5 days (3 trials per day) to control for environmental factors. The Barnes maze testing was performed at 5 months of age according to methods described previously (n = 5 mice of each genotype) [16]. There were 3 separate testing sessions of 9 days to control for environmental factors. Activity testing was performed at 6 and 20 months of age (separate cohorts) according to methods described previously, except that tests were performed between the hours of 7 pm - 6 am, as open field activity is dependent upon the resting state of the mouse, with more activity anticipated during nocturnal hours [25]. There were 3 separate testing sessions with a mix of genotypes in each session.

Immunohistochemical and immunocytochemical analyses
Rapidly frozen mouse brains were sectioned at 14 μm using a cryostat (Bright Instrument Co.). Sections were washed briefly in PBS, fixed for 10 min in 4% paraformaldehyde in 0.1M phosphate buffer pH 7.4 or in 4% paraformaldehyde for 10 min, followed by rinse in PBS and a second fixation step in 100% methanol for 15 min on ice (both conditions yielded identical results, data not shown). Sections were washed in PBS before blocking with 5% donkey serum, 0.1% Triton X-100, in PBS for 1 h at RT, and then incubated o/n at 4°C with primary antibody diluted in 5% donkey serum, 0.1% Triton X-100, and donkey anti-mouse IgG FAB (1:400) for 1 h at RT. The cells were then washed three times in PBS (10 min each wash at RT), and then blocked again in PBS containing 5% donkey serum, 0.1% Triton X-100 for 1.5 h at RT. The cells were incubated with primary antibody diluted in blocking solution for 2 h at RT, and then washed three times for 5 min at RT in PBS. The cells were then incubated with secondary antibody diluted in blocking solution, and then washed again three times for 5 min each at RT in PBS. Slides were then immersed in 70% ethanol for 5 min and then treated with 1 drop of autofluorescence eliminator reagent (Millipore/Chemicon) for an additional 5 min. Slides were coverslipped in aqueous mounting medium and imaged using an Olympus BX51 microscope equipped with an Olympus MagnaFire CCD camera. Primary antibodies used were: goat anti-Calnexin (C-20), Santa Cruz Biotechnology, 1:200; mouse anti-LC3 (5F10), Nanotools, 1:100; mouse anti-FLAG M2 (F3165), Sigma, 1:100; and mouse anti-htt MAB2166, Millipore, 1:100.

Quantification of htt aggregates and EGFP-LC3 vesicles
Tissue sections. Sections were imaged with a 60× objective using a Nikon C1 confocal microscope, and the numbers of protein aggregates were counted manually (blind to genotype) from 14 μm coronal and sagittal sections through the striatum. Neuropil and nuclear inclusions were counted separately. A z-stack was performed using the confocal microscope to confirm the presence of MW8-positive aggregates in both the cytoplasm and the nucleus (data not shown). Htt aggregates in the cortex from the same sections used to obtain the striatal htt aggregates counts were also quantified.

Cell culture analyses. The percentage of EGFP-positive cells with EGFP-HDQ74 aggregates was determined as previously described [44–46]. Analysis and acquisition of images were done with a Nikon Eclipse E600 fluorescence microscope (plan-apo 60×/1.4 oil immersion lens). Quantification of cells with EGFP-LC3 vesicles was performed as described previously [45].

Lipofuscin imaging and quantification
Lipofuscin accumulation analyses were performed using 14 μm fresh frozen brain sections. Sections were fixed for 15 min on ice in 100% methanol, washed in PBS, and then incubated with To-Pro-3 iodide (1:10,000 dilution in PBS) for 1 h at RT. Sections were then washed in PBS and mounted using Vectashield. Confocal images were acquired in the green and red channels (lipofuscin has a broad autofluorescent emission spectrum from 500 nm to 650 nm). Yellow pixel areas corresponding to the lipofuscin deposits were quantified using ImagePro 4.5 (Media Cybernetics) software from 8 images of the ventral striatum or parietal cortex obtained from each brain (n = 4 brains of each genotype for each age analyzed).

Tissue fractionation
Dissected striata from an individual brain were homogenized on ice in 500 μl 50 mM Tris-HCl pH 8.5, 100 mM NaCl, 2 mM MgCl2, 1 mM EDTA, 0.5% NP-40 supplemented with 5 mM NaF, 1 mM NaN3, and a protease inhibitor mixture (Complete -EDTA tablets, Roche). The tissue homogenate (total or unfractionated sample) was then centrifuged for 10 min at 4°C at 16,100 xg to obtain crude cytoplasmic (supernantant)
and nuclear pellet fractions. The pellet was suspended by
dounce homogenization in 100 μl homogenization buffer, and
incubated with 0.2 mg/ml final concentration of DNase I for
60 min on ice. The suspension was then centrifuged for 10 min
at 4°C at 16,100×g to obtain the final pellet fraction. The pellet
was resuspended in 100 μl homogenization buffer, and the
protein concentration in the supernatant and pellet samples
was determined using the BCA assay (Pierce). Typically, there was a
20-fold excess of protein recovered from the supernatant
fraction relative to the pellet fraction. 30 μg of each fraction
was analyzed by western blotting. Although some htt
N-terminal fragments can be solubilized by SDS-PAGE sample buffer.

**Plasmids**

The EGFP-HDQ74 construct expressing a truncated htt exon
1 fragment with 7Q fused to an MAG reporter was described previously [44]. The EGFP-LC3 expression plasmid was a gift from T. Yoshimori, while the full-length 17Q-Htt construct was a gift from M. R. Hayden [62]. The full-length 7Q-htt and ΔQ-htt expression constructs (Figure S5) were assembled from a genomic fragment containing mouse exon 1 with a portion of the flanking intron 1, a portion of the full-
length mouse htt cDNA extending from exon 2 through exon 67 including a synthetic 3′ splice acceptor site, and a poly(A)
addition sequence from the bovine growth hormone gene. The
mouse exon 1 fragments contained either wild-type sequences
encoding the 7Q stretch, or sequences derived from our HdhQ42
targeting construct lacking the polyQ stretch. A 3×FLAG
epitope tag was also inserted at the htt N-terminus between amino acids 1 and 2. A phosphoglycerol kinase (pgk)
gene promoter was used to drive expression of the 7Q- and ΔQ-htt constructs in the transfected cells.

**Cell culture and transfection**

SK-N-SH cells, wild-type Atg5 (Atg5+/+), and Atg5-deficient
(Atg5−/−), HeLa cells, wild-type P5, and HdhQ42/ΔQ P5 mouse
embryonic fibroblasts (MEFs) were maintained in DMEM
(D6546, Sigma) supplemented with 10% FBS, 100 U/ml
penicillin/streptomycin, and 2 mM L-glutamine (Sigma) in a
37°C, 5% CO2 humidified incubator. HdhQ42 or HdhΔQ
knockout (Hdh−/−) mouse ES cells were cultured on 0.1%
gelatin coated tissue culture flasks in DMEM (D6546, Sigma) supple-
mented with 15% FBS, 1 mM L-Glutamine, 1 mM penicillin/streptomycin,
1× essential amino acids, 3.5 μl (per 500 ml media) 2-mercaptoethanol (Sigma) and 1000 U/ml ESGRO
(ESG1107 with LI, Chemicon/Millipore), and incubated in a
37°C, 5% CO2 humidified incubator. Cells were transfected with
DNA constructs for 4 h using Lipofectamine 2000 (Invitrogen) according
to the manufacturer’s protocol, and either processed
for western blotting analysis 24 h post-transfection by harvesting
the cells and lysing the cell pellet on ice for 30 min in SDS-PAGE
sample buffer (62.5 mM Tris-HCl pH 6.8, 2% SDS, 5% β-
mercaptoethanol, 10% glycerol, 0.01% bromophenol blue) or
fixed with 4% paraformaldehyde (Sigma) 48 h post-transfection
and mounted with ProLong Gold antifade reagent containing
4′,6-diamidino-2-phenylindole (DAPI) (Invitrogen) for aggrega-
tion analysis.

**Western blotting**

Samples (30 μg unless otherwise noted) were fractionated on
SDS-PAGE, and then transferred electrophoretically onto
0.45 μm PVDF membranes (Invitrogen). Membranes were
processed for western blotting using standard procedures.

**Statistical analyses**

For behavioral tests, data was analyzed using the SigmaStat
program (Systat Software). One-way ANOVA, two-way repeated
measures ANOVA (with Bonferroni or Holm-Sidak post-hoc
tests), and unpaired Student t-tests were used to analyze data.

**Supporting Information**

Figure S1 Reduced htt neuropil aggregates in the HdhQ42/ΔQ
cortex. (A) Confocal images of the parietal cortex from HdhQ42/ΔQ+,
Hdh<sup>140Q/+</sup> and Hdh<sup>140Q/40Q</sup> mice at 4 months, 6 months, 1 year, and 2 years of age (n = 4 of each genotype) immunostained with the MW8 antibody recognizing htt aggregates (red). Nuclei were stained with To-Pro-3 (blue). Enlarged images of the areas enclosed by the dashed white boxes are shown in the bottom panels. Scale bars = 25 μm (top panels), 10 μm (bottom three panels). (B) Total; T, nuclear; N, and neuropil; C, htt aggregate numbers from the Hdh<sup>140Q/+</sup> and Hdh<sup>140Q/40Q</sup> cortex (n = 4 of each genotype). The aggregate numbers represent counts/field (mean ± s.e.m.) from 8 images of the parietal cortex from each mouse. *P<0.05, **P<0.001.

Found at: doi:10.1371/journal.pgen.1000838.s001 (3.67 MB TIF)

Figure S2 ΔQ-htt expression enhances LC3 immunostaining in the 6 month old Hdh<sup>106Q/106Q</sup> striatum. Confocal images of LC3 (green) and htt aggregate (MW8, red) immunostaining in the striatum from 6 month old wild-type (+/+), Hdh<sup>106Q/+</sup>, Hdh<sup>106Q/106Q</sup>, and Hdh<sup>140Q/40Q</sup> mice (n = 4 of each genotype). Nuclei were stained with To-Pro-3 (blue). Enlarged images of the areas enclosed by dashed white boxes are shown in the bottom panels. Scale bars = 25 μm.

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Figure S3 The lysosomal marker, Lamp1, is enriched in the Hdh<sup>106Q/+</sup> and Hdh<sup>140Q/40Q</sup> 800xg P1 fraction. (A) Striata dissected from wild-type (+/+), Hdh<sup>106Q/106Q</sup>, and Hdh<sup>140Q/40Q</sup> mice (n = 2 of each genotype) were homogenized and then centrifuged at 800xg, to generate a low-speed P1 fraction (see Methods). Aliquots of the P1 fraction were analyzed by western blotting using antibodies specific for Lamp1 (marker for lysosomes and autolysosomes) and beclin 1 (an essential autophagy protein involved in autophagosome nucleation). Blots were then stripped and re-probed with a tubulin antibody (loading control). Both lamp1 and beclin 1 are enriched in the P1 fractions from the Hdh<sup>106Q/+</sup> and Hdh<sup>140Q/40Q</sup> striata, but are difficult to detect in the wild type and Hdh<sup>106Q/106Q</sup> fractions. (B) Striata dissected from wild-type, Hdh<sup>106Q/106Q</sup>, and Hdh<sup>140Q/40Q</sup> mice (n = 2 of each genotype) were homogenized and aliquots of the unfractionated extract were analyzed by western blotting using antibodies specific for LC3, beclin 1, and lamp1. Blots were then stripped and re-probed with a B-actin antibody (loading control).

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Figure S4 Htt, calnexin, and LC3 localization in wild-type and Hdh<sup>106Q/106Q</sup> primary mouse embryonic fibroblasts. (A) Images of wild-type P5 (+/+), and Hdh<sup>140Q/40Q</sup> P5 primary mouse embryonic fibroblasts probed with an antibody specific for the ER marker calnexin (green), and an antibody recognizing both wild-type and ΔQ-htt (2166, red). Nuclei were stained with To-Pro-3 (blue). A merged image indicating overlap of the calnexin and htt immunoreactivity (orange to yellow color) is shown on the right. White arrowheads indicate increased nuclear htt immunoreactivity that correlates with a senescent cellular morphology. (B) Cells were probed with a mixture of calnexin (to visualize the ER; green) and FLAG antibodies (to visualize the N-terminal FLAG epitope tag on ΔQ-htt; red). The white arrowhead indicates increased nuclear ΔQ-htt immunoreactivity in an Hdh<sup>140Q/40Q</sup> senescent fibroblast. (C) Cells were probed with calnexin and LC3 antibodies to visualize ER (green), and autophagosomes (bright red punctate staining). Senescent cells exhibited increased perinuclear LC3 immunostaining. Scale bars = 10 μm.

Found at: doi:10.1371/journal.pgen.1000838.s004 (3.90 MB TIF)

Figure S5 Diagram of the 7Q-htt and ΔQ-htt expression constructs. A DNA fragment containing a synthetic 3′splice acceptor site, mouse htt cDNA sequence extending from exon 2 through exon 67, and a bovine growth hormone poly(A) addition site (located between the Sphl and KpnI restriction sites) was cloned into the pGEM 5zf plasmid vector (Promega). Inserted within the NotI restriction site located at the end of the synthetic splice acceptor site is a BamHI to EcoRV fragment containing a phosphoglycerokinase (pgk) gene promoter, an Hdh exon 1 genomic fragment containing either 7Q or ΔQ that was modified to contain a 3′FLAG epitope tag inserted at the htt N-terminus after the Methionine initiation codon, and a portion of the adjacent intron 1. Selected restriction sites are indicated, and the BamHI restriction site within parentheses indicates that it was destroyed during cloning. N = NotI, K = KpnI, X = XhoI, Xb = XbaI, Sp = SphI, Nfd = NdeI, B = BamHI, Ns = NsiI, R1 = EcoRI, X or Xmn = XmnII. The orientation of transcription is indicated with an arrow.

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Figure S6 ΔQ-htt expression in vitro increases the number of EGFP-LC3-positive vesicles. HeLa cells, transfected with EGFP-LC3 and either pCDNA3.1 (vector control), 7Q-htt or ΔQ-htt in a 1:3 ratio for 4 h, were fixed at 24 h post-transfection. The proportion of EGFP-positive cells with >10 EGFP-LC3-positive vesicles was assessed and expressed as an odds ratio with 95% confidence limits. ΔQ-htt expression (***P<0.001), but not 7Q-htt expression (NS, P = 0.737), increased the proportion of cells with EGFP-LC3-positive vesicles compared to empty vector transfected cells. ΔQ-htt expression also increased the proportion of cells with EGFP-LC3 vesicles compared to 7Q-htt transfected cells (**P<0.001).

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Figure S7 N-terminal htt fragments and htt aggregates are present in the Hdh<sup>140Q/40Q</sup> striatal pellet fraction. (A) Western blot analysis of the supernatants obtained following DNAse I digestion of a 16,100 xg pellet fraction from Hdh<sup>140Q/40Q</sup> striatum (DNAse I), and the supernatants obtained following sequential extraction of the pellet (Pel) with buffers containing 0.1% Triton x100 (Triton), CHAPS, and sodium deoxycholate (DOC). The blot in the top panel was probed with an antibody specific for the expanded polyQ stretch (1C2), while the bottom panel was probed with an antibody against p62/SQSTM1, a polyubiquitin-binding protein associated with htt aggregates [63], for comparison. A low level of soluble truncated htt fragments were recovered in the final pellet. (B) The pellet fractions from striata obtained from 2 year old wild-type (+/+), Hdh<sup>106Q/106Q</sup>, Hdh<sup>140Q/40Q</sup> (n = 1), and Hdh<sup>140Q/40Q</sup> (n = 2) mice were resuspended in SDS-PAGE sample buffer, fractionated by AGERA [64] on a 1% agarose gel, and analyzed by western blotting using an antibody recognizing htt aggregates (MW8, left panel), and an antibody recognizing ubiquitin (right panel). The position of monomeric protein, protein oligomers/aggregates, and the gel origin are indicated on the left. Note that htt aggregates are present in the Hdh<sup>140Q/40Q</sup> pellet fractions, but the amount of aggregated htt appears to be reduced compared to the levels in the Hdh<sup>140Q/40Q</sup> pellet fraction.

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Author Contributions

Conceived and designed the experiments: SZ EBDC SS MF DCR SOZ. Performed the experiments: SZ EBDC SS MF DCR SOZ. Analyzed the data: SZ EBDC SS MF DCR SOZ. Contributed reagents/materials/analysis tools: SZ EBDC SS MF DCR SOZ. Wrote the paper: SZ EBDC SS MF DCR SOZ.
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