Neuronal XBP-1 Activates Intestinal Lysosomes to Improve Proteostasis in C. elegans

Highlights

- Xbp-1s expressed in the neurons or intestine of C. elegans improves proteostasis
- Neuronal xbp-1s drives expression of lysosomal genes in the intestine
- Intestinal lysosomes show enhanced acidity and activity upon xbp-1s expression
- Lysosome function is required for xbp-1s to increase proteostasis and longevity

Graphical Abstract

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In Brief
Expressing the UPRER transcription factor xbp-1s in the neurons or intestine of C. elegans extends lifespan. Imanikia et al. show that neuronal xbp-1s upregulates lysosomal genes and activates lysosomes in the intestine and that intestinal lysosome function is crucial for xbp-1s-mediated increases in proteostasis and longevity.
Neuronal XBP-1 Activates Intestinal Lysosomes to Improve Proteostasis in C. elegans

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SUMMARY
The unfolded protein response of the endoplasmic reticulum (UPRER) is a crucial mediator of secretory pathway homeostasis. Expression of the spliced and active form of the UPRER transcription factor XBP-1, XBP-1s, in the nervous system triggers activation of the UPRER in the intestine of Caenorhabditis elegans (C. elegans) through release of a secreted signal, leading to increased longevity. We find that expression of XBP-1s in the neurons or intestine of the worm strikingly improves proteostasis in multiple tissues, through increased clearance of toxic proteins. To identify the mechanisms behind this enhanced proteostasis, we conducted intestine-specific RNA-seq analysis to identify genes upregulated in the intestine when XBP-1s is expressed in neurons. This revealed that neuronal XBP-1s increases the expression of genes involved in lysosome function. Lysosomes in the intestine of animals expressing neuronal XBP-1s are more acidic, and lysosomal protease activity is higher. Moreover, intestinal lysosome function is necessary for enhanced lifespan and proteostasis. These findings suggest that activation of the UPRER in the intestine through neuronal signaling can increase the activity of lysosomes, leading to extended longevity and improved proteostasis across tissues.

INTRODUCTION
Aging is a process that occurs in a coordinated fashion throughout the body, leading to a susceptibility to deterioration and disease across tissues. The choreography of the aging process is achieved through inter-tissue signaling, often involving the nervous system. In C. elegans, mutations in genes that affect nervous system function can significantly increase longevity [1]. This effect is also observed in other species—in Drosophila melanogaster, sensory perception of food can alter lifespan in a similar way to consumption of that food, while, in mice, loss of the neuronal pain receptor TRPV1 creates animals that live substantially longer, with improved metabolic profiles [2, 3]. The ability of neurons to modulate aging is therefore a conserved phenomenon. However, it is not clear whether these neuronal signals converge upon the same core mechanisms in distal tissues. Understanding these mechanisms could allow them to be engaged directly in order to improve aging phenotypes and increase cellular health.

Recently, the transmission of cellular stress response activation has been identified as a mechanism utilized by the nervous system to increase longevity [4]. Stress responses are organelle-specific pathways that connect the sensing of molecular damage and imbalances in homeostasis to mechanisms that restore equilibrium. They can be activated in distal tissues by signals secreted from neurons, typically leading to marked lifespan extension. One of these stress responses, the endoplasmic reticulum unfolded protein response (UPRER), is an adaptive signaling pathway that restores equilibrium within the secretory pathway. Its three branches are defined by their proximal sensor molecules, IRE1, PERK, and ATF6, which regulate downstream mechanisms that include both transcriptional and translational targets [5]. Activation of IRE1 leads to the regulated splicing and translation of the transcription factor XBP1. In C. elegans, expression of the spliced and active form of XBP-1, XBP-1s, in the nervous system triggers the release of an unidentified signal from neurons that activates this branch of the UPRER in the animal’s intestine [6]. This signaling extends lifespan and increases stress resistance, with signal release dependent upon UNC-13, a neuron-specific regulator of synaptic secretion. The effects of UPRER activation are tissue specific—activation in the intestine, either directly or through neuronal signaling, is beneficial, while expression of XBP-1s in body wall muscle cells is detrimental to longevity. While the downstream targets and mechanisms invoked in distal tissues by XBP-1s to achieve changes in longevity are unknown, it is possible that tissue-specific differences in these XBP-1s targets lead to differing effects on lifespan.

One hallmark of aging is a loss of cellular homeostasis, particularly protein homeostasis (proteostasis) [7–9]. Loss of proteostasis underlies the onset of many diseases of aging, including neurodegenerative diseases, which are often caused by age-associated accumulation of toxic, aggregated proteins. Improving cellular proteostasis extends lifespan, and neurons are able to enhance proteostasis in distal tissues [10–12]. As stress responses can trigger mechanisms that restore proteostasis, one function of the distal activation of these pathways by neurons might be to improve proteostasis in other tissues, counteracting the loss of equilibrium associated with aging and leading to increased longevity.
Figure 1. Expression of xbp-1s in Neurons or the Intestine Restores Proteostasis in Models of Proteotoxicity
(A) Proteotoxicity and UPRIP3 models used in this study. TS, temperature sensitive. *dyn-1(ky51) and let-60(ga89) are expressed in multiple tissues, but their functions in motor neurons and intestine, respectively, are assayed here.
(B) Chemotaxis ability in animals expressing Aβ1–42 in neurons, with and without neuronal, intestinal, or muscle xbp-1s. Graphs represent mean chemotaxis index ± SD. N = 80–150 animals per assay; each assay was independently replicated 3 times. Significance was assessed by one-way ANOVA with Tukey’s multiple comparisons test, *p < 0.05, ****p < 0.0001.
(C) Chemotaxis ability in animals expressing polyQ40::YFP in neurons, with and without neuronal, intestinal, or muscle xbp-1s. Graphs represent mean chemotaxis index ± SD. N = 80–150 animals per assay; each assay was independently replicated 3 times. Significance was assessed by one-way ANOVA with Tukey’s multiple comparisons test, ****p < 0.0001.
(D) Paralysis in dyn-1(ky51) animals at 20°C, with and without neuronal xbp-1s. Animals exhibiting body paralysis were counted daily and the paralyzed fraction of the population plotted against time. N = 100 worms per assay; each assay was repeated 3 times.
(E) Prevalence of osmoregulatory defects in let-60(ga89) and wild-type animals, with and without rab-3p::xbp-1s. Animals were placed in distilled water at day 2 of adulthood, and a swollen body after 5 min was scored as osmoregulation defective (Osm). Data represent mean ± SD of 20–25 animals per strain in 3 independent replicates. Significance assessed by two-way ANOVA with Tukey’s multiple comparison, ***p < 0.001, ****p < 0.0001.
Here, we have explored the mechanisms by which neuronal XBP-1s extends lifespan in C. elegans. Speculating that improved proteostasis might underlie the beneficial effects of XBP-1s on longevity and stress resistance, we have established that neuronal and intestinal XBP-1s improve proteostasis in multiple tissues of the organism by increasing clearance of toxic proteins. To identify transcriptional targets that underlie these improvements in longevity and proteostasis, we conducted tissue-specific RNA sequencing (RNA-seq) analysis in the intestines of animals expressing neuronal XBP-1s and found a prominent upregulation of lysosomal genes. Intestinal lysosomes play a key role in the extension of longevity and improvements in proteostasis downstream of neuronal UPR

RESULTS

To investigate the mechanisms by which XBP-1s might extend longevity, we started by asking whether its expression could improve proteostasis, either cell-autonomously or cell non-autonomously. To do this, we drove expression of spliced xbp-1s in different tissues of the worm (neurons, intestine, and body wall muscle cells, in animals expressing endogenous xbp-1) with models of proteotoxicity, in which aggregation-prone proteins are also expressed in different tissues of the animal (Figure 1A). Proteotoxicity was then assessed using physiological readouts of tissue function.

Neuronal and Intestinal xbp-1s Protects against Proteotoxicity in Neuronal and Intestinal Cells

\( \text{A}_\text{b1–42} \) is a proteotoxic peptide associated with the development of Alzheimer’s disease. To determine whether xbp-1s affects phenotypes associated with proteotoxicity, pan-neuronal expression of \( \text{A}_\text{b1–42} \), with a signal peptide directing it to the secretory pathway [13], was combined with tissue-specific expression of xbp-1s in neuronal, intestinal, and body wall muscle cells, and the function of neurons measured by chemotaxis—movement of worms toward an attractive volatile odorant. Strikingly, expression of xbp-1s in either neurons or the intestine fully rescued the loss of chemotaxis caused by \( \text{A}_\text{b1–42} \) expression, while the effect of muscle cell xbp-1s expression was limited (Figure 1B).

To then ask whether neuronal or intestinal xbp-1s could also exert these protective effects against another toxic protein, tissue-specific xbp-1s was combined with pan-neuronal expression of polyglutamine (polyQ) expansions, associated with disorders including Huntington’s disease and shown to be good sensors for protein folding homeostasis in the cytosol [8, 14]. Again, expression of xbp-1s in neuronal or intestinal, but not muscle cells, fully rescued animals from loss of neuronal function, demonstrating that this transcription factor can mitigate the effects of multiple proteotoxic species, including those directed to the secretory pathway as well as those expressed cytosolically (Figure 1C).

To determine whether xbp-1s can also improve phenotypes associated with the misfolding of endogenous proteins, we used C. elegans strains containing temperature-sensitive missense mutations. These metastable proteins are nonfunctional at 25°C but at lower temperatures show an accelerated age-associated loss of folding and function that correlates with age-related decline in endogenous stress responses [7, 15]. A metastable allele of dyn-1, encoding a dynamin GTPase, was examined, as well as a temperature-sensitive allele of let-60, encoding a Ras GTPase homolog. The effect of dyn-1(\text{ky51}) on motor neurons was measured by age-associated motor paralysis; in animals expressing xbp-1s in neurons, paralysis of dyn-1(\text{ky51}) animals was substantially delayed (Figure 1D). To determine the effect of let-60(\text{ga89}) on intestinal function, osmoregulatory capacity was assessed. Xbp-1s expression in neurons significantly decreased the percentage of animals showing defects in osmoregulation (Figure 1E). Neuronal xbp-1s therefore protects against dysfunction associated with endogenous metastable, as well as human disease-associated proteins, in neuronal and intestinal cells.

Neuronal and Intestinal xbp-1s Protects against Proteotoxicity in Muscle Cells

To determine the effects of xbp-1s on proteotoxicity in a tissue other than neurons or the intestine, \( \text{A}_\text{b1–42} \) (targeted to the secretory pathway) was expressed in body wall muscle cells [16-18] in combination with tissue-specific xbp-1s. Muscle function was then assayed by measuring body wall muscle paralysis with age. Surprisingly, although \( \text{A}_\text{b1–42} \) was expressed within muscle cells, xbp-1s delayed toxicity and paralysis when expressed in neurons and intestine, but not when expressed in muscle cells themselves (Figure 1F). Tissue-specific xbp-1s was then combined with cytosolic polyQ35 expression, and muscle cell function measured by swimming rate at day 1 and day 4 of adulthood. Expression of polyQ35 in muscles caused a significant decline in motility, which was rescued by xbp-1s expression in neurons or the intestine, but not in muscle cells (Figure 1G).

Finally, we examined the effect of neuronal xbp-1s expression on a temperature-sensitive metastable allele of the muscle-specific gene unc-15, encoding paramyosin. Upon xbp-1s...
Figure 2. Xbp-1s Expression Reduces Levels of PolyQ

(A) Native western blot analysis of (i) neuronal polyQ<sub>40</sub>::YFP, (ii) intestinal polyQ<sub>40</sub>::YFP, and (iii) muscle polyQ<sub>35</sub>::YFP with and without tissue-specific xbp-1<sup>s</sup>, at day 1 and day 4 of adulthood. Lysates were resolved under native conditions and blotted with an anti-polyQ antibody; the predicted MW of monomeric polyQ<sub>35–40</sub>::YFP is ~32 kDa. Tubulin levels were probed with α-tubulin as a loading control. Blots were quantified using ImageJ (iv–vi). Bar graphs represent mean band intensity relative to day 1 polyQ ± SD and statistical significance was calculated between A and B/C at each time point using two-way ANOVA with Tukey’s multiple comparisons test, *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001. Data are representative of at least 3 independent experiments.

(B) (i) Epifluorescence visualization of neuronal polyQ<sub>40</sub>::YFP, with and without neuronal and intestinal xbp-1<sup>s</sup>, at day 1 and day 4 of adulthood. Enlarged panels are from day 4 animals. Scale bar, 250 μm.

(C) (i) Fluorescence intensity of neuronal polyQ<sub>40</sub>::YFP, quantified from 3 independent experiments using ImageJ and expressed as mean fluorescence intensity relative to day 1 polyQ40 ± SEM. N = 25–30 animals. Significance was assessed between A and B/C at each time point using two-way ANOVA with Tukey’s multiple comparisons test, *p < 0.05, ****p < 0.0001.

(i) Fluorescence intensity of neuronal polyQ<sub>40</sub>::YFP quantified from 3 independent experiments using ImageJ and expressed as mean fluorescence intensity relative to day 1 polyQ<sub>40</sub> ± SEM. N = 25–30 animals. Significance was assessed between A and B/C at each time point using two-way ANOVA with Tukey’s multiple comparisons test, *p < 0.05, ****p < 0.0001.
expression, age-associated paralysis caused by unc-15(1402) was substantially delayed (Figure 1H). In addition, muscle fiber disorganization in unc-15(1402) animals at day 5 of adulthood was largely rescued by neuronal xbp-1s (Figure 1I). Together, these results suggest that xbp-1s in neurons or the intestine exerts a protective effect against proteotoxicity, even when the toxic species is expressed in an entirely different tissue.

Expression of xbp-1s Reduces the Accumulation of Misfolded Proteins

To determine whether xbp-1s exerts its beneficial effects on proteotoxicity through changes in levels of toxic proteins, we co-expressed neuronal or intestinal xbp-1s with Aβ1–42 in neurons or muscle cells. When Aβ1–42 was expressed in neurons, xbp-1s in neuronal or intestinal cells marginally reduced levels of Aβ1–42 at day 4 of adulthood (Figures S1Aii and S1Av). Upon expression of Aβ1–42 in muscle cells, however, neuronal and intestinal xbp-1s strikingly reduced Aβ1–42 levels at both day 1 and day 4 (Figures S1Aii and S1Aiv). Notably, multiple species likely to correspond to soluble oligomers of Aβ1–42, and associated with Aβ1–42 toxicity, were substantially reduced by xbp-1s expression.

We then used native PAGE of YFP-tagged polyQ expansions co-expressed with tissue-specific xbp-1s to assess polyQ::YFP levels. This revealed that polyQ::YFP expressed in neuronal or intestinal [19] cells was substantially reduced by expression of xbp-1s in neurons or the intestine (Figures 2Ai, 2Aii, 2Aiv, 2Av, S1Bii, S1Bii, S1Biv, and S1Bv). Muscle-specific polyQ35::YFP levels were reduced by expression of neuronal xbp-1s, and to a lesser extent by xbp-1s in muscle cells (Figures 2Aii, 2Aiv, S1Bii, and S1Bv). We also analyzed polyQ::YFP under denaturing conditions and found that total polyQ levels were reduced by xbp-1s, although less strikingly than when assessed by native PAGE (and in muscle-specific polyQ35, not at all at day 4) (Figure S1C). Denaturing conditions can resolve both soluble and other forms of polyQ, including insoluble aggregated species, whereas native analysis only resolves soluble proteins; one possibility is that xbp-1s preferentially reduces levels of soluble, lower-molecular-weight polyQ, rather than aggregated forms.

Tissue-specific polyQ::YFP was then visualized by microscopy at days 1 and 4 of adulthood, with and without tissue-specific xbp-1s. Neuronal polyQ40::YFP fluorescence intensity increased between day 1 and day 4 but was reduced by xbp-1s, while muscle-specific polyQ35::YFP aggregates were also reduced by neuronal or intestinal xbp-1s (Figure 2Bii). Intestinal polyQ35::YFP fluorescence also increased with age, although aggregation was not observed, and was very substantially decreased by neuronal or intestinal xbp-1s expression at both day 1 and day 4 (Figures 2Ci and 2Cii). Muscle-specific polyQ35::YFP was co-expressed with neuronal and muscle-specific xbp-1s, at both day 1 and day 4, expression of xbp-1s decreased polyQ35::YFP fluorescence (Figures 2Di and 2Dii). However, polyQ35::YFP aggregate accumulation in muscles was unaffected by xbp-1s (Figure 2Dii). Loss of polyQ35::YFP fluorescence, but not aggregation, suggests that XBP-1s may preferentially reduce levels of more soluble, lower-molecular-weight forms of polyQ35. As animals are healthier (Figure 1G) despite having similar numbers of visible aggregates, we suggest that XBP-1s may reduce proteotoxicity primarily due to reduction in soluble proteotoxic species.

Neuronal and Intestinal xbp-1s Does Not Change Aβ1–42 or PolyQ Transcript Levels

Importantly, no reduction in polyQ::YFP or Aβ1–42 transcript levels were observed upon xbp-1s expression, confirming that changes in protein abundance are unlikely to arise from changes to transcription of these transgenes (Figures S2A and S2B). Although we cannot rule out changes to rates of translation, we hypothesize that reduced levels of proteotoxic species upon neuronal and intestinal xbp-1s expression may potentially be the result of increased misfolded protein clearance driven by xbp-1s.

Intestinal Genes Regulated by Neuronal xbp-1s Include Genes Involved in Lysosome Function

Given the profound effects of xbp-1s expression on proteotoxicity, we decided to explore the mechanisms by which xbp-1s might improve proteostasis. Neuronal and intestinal xbp-1s both protected against proteotoxic species in multiple tissues, while muscle-specific xbp-1s had only minor protective effects. Interestingly, xbp-1s expression in neurons and the intestine, but not muscle cells, extends longevity in C. elegans [6]. Neuronal xbp-1s expression leads to the communication of UPR[ER] activation to intestinal cells, through the release of a signal secreted by neurons. We therefore hypothesized that the intestine may be a key tissue through which xbp-1s activity coordinates organism-wide improvements in lifespan and proteostasis.
To understand how communication of UPRER activation to the intestine leads to these downstream effects, we decided to determine the transcriptional changes occurring in intestinal cells when \( xbp-1s \) is expressed in neurons. To do this, we adapted an existing \( C. \) elegans cellular dissociation protocol and a tissue-specific RNA-seq method previously used to determine transcriptional changes in neurons [20, 21]. GFP-labeled intestinal cells from wild-type and \( \text{rab-3p::xbp-1s} \) animals expressing \( \text{ges-1p::GFP} \), an intestine-specific GFP marker [22], were isolated from dissociated worm cells by fluorescence-activated cell sorting (FACS), and RNA-seq was used to identify transcripts altered by neuronal \( xbp-1s \) (Figure S2C). To validate our approach, we examined the levels of a range of transcripts in our intestinal samples. Confirming previous observations that neuronal \( xbp-1s \)-expressing animals upregulate \( xbp-1s \) activity in the intestine, we observed higher levels of \( xbp-1s \) transcripts in isolated intestinal cells from rab-

\[3p::xbp-1s \] animals compared to wild type (Figure S2D) [6]. Additionally, to confirm enrichment of intestine-specific relative to muscle- and neuron-specific transcripts, expression levels of tissue-specific genes from the Princeton “Tissue-specific expression predictions for \( C. \) elegans” database were examined; this demonstrated enrichment of intestinal transcripts (Figure S2E). Finally, qPCR was used to examine expression levels of intestinal, neuronal, hypodermal, and muscle-specific genes in our RNA samples, identifying significant enrichment of intestinal, and underrepresentation of other tissue-specific transcripts (Figure S2F).

Comparison of transcripts in neuronal \( xbp-1s \) and control worms revealed broad transcriptional remodeling in the intestine, with 1,388 genes upregulated and 830 downregulated in neuronal \( xbp-1s \) animals (Figures 3A and 3B; Data S1). The gene ontology categories most enriched among these differentially regulated transcripts included the endoplasmic reticulum unfolded protein response, endoplasmic reticulum lumen, and protein processing in the endoplasmic reticulum, as expected (Figure 3C; Data S1). Upregulated genes included known UPRER components such as \( xbp-1 \) and \( hsp-4 \), confirming that neuronal XBP-1s can trigger activation of the UPRER in the intestine cell non-autonomously [6].

However, several other gene classes were also differentially regulated, suggesting that the transcriptional landscape of the cell non-autonomous UPRER extends beyond canonical UPRER targets. Of these, a prominent upregulation of lysosomal genes
A

\( \text{Imp-1 RNAi} \)

Survival

Days

\( \text{N2 = control RNAI} \)
\( \text{N2 + Imp-1 RNAI} \)
\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + Imp-1 RNAI} \)

\( \text{cup-5 RNAI} \)

Survival

Days

\( \text{N2 = control RNAI} \)
\( \text{N2 + cup-5 RNAI} \)
\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + cup-5 RNAI} \)

\( \text{vha-18 RNAI} \)

Survival

Days

\( \text{N2 = control RNAI} \)
\( \text{N2 + vha-18 RNAI} \)
\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + vha-18 RNAI} \)

\( \text{asp-3 RNAI} \)

Survival

Days

\( \text{N2 = control RNAI} \)
\( \text{N2 + asp-3 RNAI} \)
\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + asp-3 RNAI} \)

B

\( \text{bec-1 RNAI} \)

Survival

Days

\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + vps-3d RNAI} \)
\( \text{rab-3p::xbp-1s + vps-3d RNAI + vps-3d RNAI} \)

\( \text{vps-3d RNAI} \)

Survival

Days

\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + vps-3d RNAI} \)
\( \text{rab-3p::xbp-1s + vps-3d RNAI + vps-3d RNAI} \)

\( \text{atf-13 RNAI} \)

Survival

Days

\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + atf-13 RNAI} \)
\( \text{rab-3p::xbp-1s + atf-13 RNAI + atf-13 RNAI} \)

\( \text{atf-18 RNAI} \)

Survival

Days

\( \text{rab-3p::xbp-1s + control RNAI} \)
\( \text{rab-3p::xbp-1s + atf-18 RNAI} \)
\( \text{rab-3p::xbp-1s + atf-18 RNAI + atf-18 RNAI} \)

C

\( \text{Control RNAI} \)

Chemotaxis Index

Neuronal A8

A

B

C

D

\( \text{Imp-1 RNAI} \)

Chemotaxis Index

Neuronal A8

A

B

C

D

\( \text{cup-5 RNAI} \)

Chemotaxis Index

Neuronal A8

A

B

C

D

\( \text{vha-15 RNAI} \)

Chemotaxis Index

Neuronal A8

A

B

C

D

\( \text{asp-3 RNAI} \)

Chemotaxis Index

Neuronal A8

A

B

C

D

D

\( \text{Control RNAI} \)

Chemotaxis Index

Neuronal polyQ

A

B

C

D

\( \text{Imp-1 RNAI} \)

Chemotaxis Index

Neuronal polyQ

A

B

C

D

\( \text{cup-5 RNAI} \)

Chemotaxis Index

Neuronal polyQ

A

B

C

D

\( \text{vha-15 RNAI} \)

Chemotaxis Index

Neuronal polyQ

A

B

C

D

\( \text{asp-3 RNAI} \)

Chemotaxis Index

Neuronal polyQ

A

B

C

D

(legend on next page)
stood out as a candidate mechanism for XBP-1s-mediated reductions in toxic protein levels, as lysosomes and autophagy have been implicated in clearance of protein aggregates (Figures 3C and 3D) [23]. qPCR analysis of a selection of these genes confirmed their upregulation in neuronal xbp-1s-expressing animals (Figure 3E). We also found that a majority of these lysosomal genes had putative XBP-1s binding sites in their promoters, suggesting that they may represent direct transcriptional targets (Figure S3A). Lysosomal genes upregulated by XBP-1s are involved in a range of lysosomal functions and include proteases, suggesting that increased lysosomal proteolysis might underlie the effects of XBP-1s on proteotoxicity (Figure 3D). Interestingly, when we examined transcript levels of several of these lysosomal genes in short-lived animals expressing xbp-1s in muscle cells, we found that they were not increased—in fact, transcription of these lysosomal genes was reduced (Figure S3B), suggesting a correlation between lysosomal gene transcription and longevity. We therefore decided to explore the role of lysosomes in XBP-1s-mediated improvements to lifespan and proteostasis.

**Lysosome Function Is Required for Extended Longevity and Protection against Proteotoxicity by Neuronal xbp-1s**

To determine whether lysosome function is required for the long lifespan of rab-3p::xbp-1s animals, we used RNAi to knock down LMP-1, a homolog of mammalian LAMP1, which localizes to the membranes of lysosomes (and some endosomal vesicles), thus inhibiting lysosome function. Lmp-1 knockdown suppressed the extended lifespan of rab-3p::xbp-1s worms, while longevity of N2 animals was unaffected (Figure 4Ai; Table S1). We also extended lifespan of (B) Lifespan analysis of (iii) N2, control (black), median lifespan 17 days; N2, independently replicated 3 times. Significance between neuronal A

Figure 4. Lysosome Function Is Required for the Effects of xbp-1s on Lifespan and Proteotoxicity

(A) Lifespan analysis of N2 and rab-3p::xbp-1s animals grown on control (empty vector) or lysosomal gene RNAi; (i) imp-1, (ii) cup-5, (iii) vha-18, (iv) asp-3. Graphs were plotted as Kaplan-Meier survival curves, and p values were calculated by Mantel-Cox log-rank test; N = 80–120 animals per lifespan.

(i) N2, control (black), median lifespan 15 days; N2, imp-1 (gray), median lifespan 21 days, p = 0.8860; rab-3p::xbp-1s, control (green), median lifespan 23 days; rab-3p::xbp-1s, imp-1 (blue), median lifespan 21 days, p < 0.0001.

(ii) N2, control (black), median lifespan 21 days; N2, cup-5 (gray), median lifespan 21 days, p = 0.0950; rab-3p::xbp-1s, control (green), median lifespan 25 days; rab-3p::xbp-1s, cup-5 (blue), median lifespan 21 days, p = 0.01.

(iii) N2, control (black), median lifespan 15 days; N2, vha-18 (gray), median lifespan 17 days, p = 0.8118; rab-3p::xbp-1s, control (green), median lifespan 25 days; rab-3p::xbp-1s, vha-18 (blue), median lifespan 23 days, p < 0.0001.

(iv) N2, control (black), median lifespan 21 days; N2, asp-3 (gray), median lifespan 21 days, p = 0.9913; rab-3p::xbp-1s, control (green), median lifespan 25 days; rab-3p::xbp-1s, asp-3 (blue), median lifespan 23 days, p < 0.001.

(B) Lifespan analysis of rab-3p::xbp-1s animals grown on control (empty vector) and xbp-1 RNAi; as well as RNAi against autophagy genes: (i) bec-1, (ii) vps-34, (iii) atg-13, (iv) atg-18. Graphs were plotted as Kaplan-Meier survival curves, and p values were calculated by Mantel-Cox log-rank test.

(i) rab-3p::xbp-1s, control (black), median lifespan 25 days; rab-3p::xbp-1s, xbp-1 (green), median lifespan 20 days, p < 0.0001; rab-3p::xbp-1s, bec-1 (blue), median lifespan 26 days, p < 0.0001.

(ii) rab-3p::xbp-1s, control (black), median lifespan 25 days; rab-3p::xbp-1s, xbp-1 (green), median lifespan 20 days, p < 0.0001; rab-3p::xbp-1s, vps-34 (blue), median lifespan 25 days, p = 0.0976.

(iii) rab-3p::xbp-1s, control (black), median lifespan 25 days; rab-3p::xbp-1s, xbp-1 (green), median lifespan 21 days, p < 0.0001; rab-3p::xbp-1s, atg-13 (blue), median lifespan 27 days, p < 0.0001.

(iv) rab-3p::xbp-1s, control (black), median lifespan 25 days; rab-3p::xbp-1s, xbp-1 (green), median lifespan 21 days, p < 0.0001; rab-3p::xbp-1s, atg-18 (blue), median lifespan 25 days, p = 0.1527.

(C) Chemotaxis ability in animals expressing A\textsubscript{1},\textsubscript{42} in neurons in combination with neuronal and intestinal xbp-1s, grown on (i) control (empty vector), (ii) imp-1, (iii) cup-5, (iv) vha-18 (from L4), (v) vha-18, or (vi) asp-3 RNAi. Graphs represent mean chemotaxis index ± SD. N = 85–140 animals per assay; each assay was independently replicated 3 times. Significance between neuronal A\textsubscript{1},\textsubscript{42} (B) and xbp-1s-expressing (C/D) strains was assessed by two-way ANOVA with Dunnett’s multiple comparisons test. ***p < 0.0001.

(D) Chemotaxis ability in animals expressing polyQ\textsubscript{40} in neurons in combination with neuronal and intestinal xbp-1s, grown on (i) control (empty vector), (ii) imp-1, (iii) cup-5, (iv) vha-18 (from L4), (v) vha-18, or (vi) asp-3 RNAi. Graphs represent mean chemotaxis index ± SD. N = 80–170 animals per assay; each assay was independently replicated 3 times. Significance between neuronal polyQ\textsubscript{40} (B) and xbp-1s-expressing (C/D) strains was assessed by two-way ANOVA with Dunnett’s multiple comparisons test. **p < 0.01, ***p < 0.001, ****p < 0.0001. See also Figure S4 and Table S1.
Figure 5. Intestinal Knockdown of Lysosomal Genes Abolishes Lifespan Extension and Improved Proteostasis in xbp-1s-Expressing Animals

(A) Lifespan analysis of rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1 animals grown on control (empty vector) or (i) xbp-1, (ii) lmp-1, (iii) vha-18, or (iv) asp-3 RNAi. Graphs were plotted as Kaplan-Meier survival curves, and p values were calculated by Mantel-Cox log-rank test; N = 80–120 animals per lifespan. (i) rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, control (black), median lifespan 22 days; rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, xbp-1 (green), median lifespan 16 days, p < 0.0001. (ii) rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, control (black), median lifespan 22 days; rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, lmp-1 (green), median lifespan 16 days, p < 0.0001. (iii) rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, control (black), median lifespan 22 days; rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, vha-18 (green), median lifespan 16 days, p < 0.0001. (iv) rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, control (black), median lifespan 22 days; rab-3p::xbp-1s; rde(n219); nhx-2p::rde-1, asp-3 (green), median lifespan 16 days, p < 0.0001.

(B) Chemotaxis ability in rde(n219); mex-5p::rde-1 animals expressing Aβ1-42 in neurons in combination with neuronal or intestinal xbp-1s, grown on (i) control (empty vector), (ii) xbp-1, or (iii) lmp-1 RNAi. Graphs represent mean chemotaxis index ± SD. N = 80–130 animals per assay; each assay was independently replicated 3 times. Significance between neuronal Aβ1-42 (B) and xbp-1s-expressing (C and D) strains was assessed by two-way ANOVA with Dunnett’s multiple comparisons test, *p < 0.05, **p < 0.01, ****p < 0.0001.

(C) Chemotaxis ability in rde(n219); mex-5p::rde-1 animals expressing polyQ40 in neurons in combination with neuronal or intestinal xbp-1s, grown on (i) control (empty vector), (ii) xbp-1, or (iii) lmp-1 RNAi. Graphs represent mean chemotaxis index ± SD. N = 80–130 animals per assay; each assay was independently replicated 3 times. Significance between neuronal polyQ40 (C) and xbp-1s-expressing (D and E) strains was assessed by two-way ANOVA with Dunnett’s multiple comparisons test, *p < 0.05, **p < 0.01, ****p < 0.0001.
to induce autophagosome formation, combined with RNAi against autophagy genes, we found that autophagosome formation was significantly suppressed in animals grown on RNAi against bec-1, vps-34, atg-13, or atg-18, confirming that these treatments effectively inhibit autophagy (Figure S4C).

**Intestinal Knockdown of Lysosomal Genes Suppresses Lifespan Extension and Protection against Proteotoxicity by xbp-1s**

Neuronal xbp-1s expression activates the UPRER in the intestine, leading to upregulation of lysosomal genes in this tissue (Figure 3) [6]. In addition, expression of xbp-1s only in intestinal cells is sufficient to protect against proteotoxicity in multiple tissues. We therefore wondered whether intestine-specific knockdown of lysosomal genes would affect the ability of neuronal xbp-1s to extend lifespan and protect against proteotoxicity. To address this, we used an rde-1(n219) mutation that prevents effective RNAi, coupled with replacement of rde-1 only in intestinal cells [24, 25]. Intestine-specific rde-1 expression utilized the nhx-2p promoter; however, because these animals were somewhat short lived, we also used animals in which rde-1 was replaced in both the intestine and germline (but no additional somatic tissues) using the mex-5p promoter [26]. First, we established that rab-3p::xbp-1s animals were still long lived in an rde-1(n219) mutant background and when rde-1 was expressed under either intestinal promoter (Figures S5Aii–S5Aiii; Table S1). We then confirmed that xbp-1 RNAi did not shorten the lifespan of rab-3p::xbp-1s; rde-1(n219) animals (Figure S5Aiv; Table S1). However, when rde-1 expression was rescued only in the intestine, xbp-1 RNAi was able to significantly reduce longevity (Figure 5Aii). Likewise, intestinal knockdown of the lysosomal genes imp-1, vha-18, and asp-3 in rab-3p::xbp-1s; rde-1(n219); nhx-2p::rde-1 animals also significantly reduced lifespan, demonstrating that intestinal lysosome function is required for extended longevity upon neuronal xbp-1s expression (Figures 5Aiii–5Aiv; Table S1). The same effects were seen in animals expressing rde-1 under the mex-5 promoter (Figure S5B; Table S1).

Next, we asked whether intestinal knockdown was sufficient to abrogate neuronal xbp-1s-mediated protection against proteotoxicity in animals expressing neuronal Aβ1–42 or polyQ40. We found that intestinal xbp-1 or imp-1 knockdown prevented rescue of chemotaxis by neuronal and intestinal xbp-1s expression in both models (Figures 5B and 5C), suggesting that the activity of lysosomes in the intestine is important to the protection against proteotoxicity mediated by neuronal or intestinal xbp-1s. Finally, to determine whether xbp-1s in the intestine is required for upregulation of lysosomal gene targets in this tissue, we used qRT-PCR in rab-3p::xbp-1s; rde-1(n219); nhx-2p::rde-1 animals to ask whether intestine-specific knockdown of xbp-1 altered expression of lysosomal genes. Indeed, intestine-specific xbp-1 RNAi was sufficient to prevent upregulation of several lysosomal transcripts in these animals (Figure S5C).

**HLH-30 Is Necessary for Lifespan Extension and Protection against Proteotoxicity by xbp-1s**

The *C. elegans* homolog of the TFEB transcription factor, HLH-30, is required for lysosomal biogenesis and activation [27, 28]. We found that several of the lysosomal genes upregulated in xbp-1s-expressing animals had predicted HLH-30-binding CLEAR domains in their promoters (Figure S3A) and therefore asked whether HLH-30 was required for neuronal xbp-1s to extend lifespan. We found that hlh-30 knockdown substantially reduced longevity in rab-3p::xbp-1s animals (Figure 5D; Table S2). Indeed, intestine-specific RNAi against hlh-30 was sufficient to abolish lifespan extension, and adult-only intestinal knockdown of either hlh-30 or xbp-1, by transfer of day 1 adults to RNAi, was again sufficient to reduce longevity (Figures S6A–S6C; Table S2).

We also asked whether RNAi against hlh-30 could abolish xbp-1s-mediated protection against proteotoxicity. Again, rescue of chemotaxis by xbp-1s was substantially reduced by hlh-30 knockdown in both Aβ1–42 and polyQ40 models, confirming that functional lysosomes are required for improved proteostasis (Figure 5E). Intestine-specific knockdown of hlh-30 prevented induction of lysosomal genes in rab-3p::xbp-1s animals (Figure S5C). However, unlike starvation, a known HLH-30-activating stimulus, neuronal xbp-1s did not induce nuclear localization of HLH-30::GFP, suggesting that xbp-1s may not directly activate HLH-30 but may instead rely on its basal roles in lysosome biogenesis and activity (Figure S6D).

**Lysosomes Are Required for xbp-1s-Induced Reductions in PolyQ Levels**

Expression of xbp-1s in the intestine or nervous system of *C. elegans* reduces levels of toxic proteins expressed in multiple tissues. To determine whether lysosomes are required for this reduction, imp-1 was knocked down in worms expressing neuronal and intestinal polyQ40 concurrently with neuronal or intestinal xbp-1s, and polyQ40::YFP levels determined by native PAGE. Upon xbp-1s expression, polyQ40 levels were reduced; however, upon imp-1 knockdown, this effect was diminished (Figure 6A), suggesting that lysosomes may be involved in XBP-1s-mediated polyQ40 degradation.

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replicated 3 times. Significance between neuronal polyQ40 (B) and xbp-1s-expressing (C and D) strains was assessed by two-way ANOVA with Dunnett’s multiple comparisons test, *p < 0.05, ***p < 0.0001.

(D) Lifespan analysis of rab-3p::xbp-1s animals grown on control (empty vector) or hlh-30 RNAi, rab-3p::xbp-1s, control (black), median lifespan 25 days; rab-3p::xbp-1s, hlh-30 (green), median lifespan 19 days, p < 0.0001. Graphs were plotted as Kaplan-Meier survival curves, and p values were calculated by Mantel-Cox log-rank test; N = 80–120 animals per lifespan.

(E) (i) Chemotaxis ability in animals expressing Aβ1–42 in neurons in combination with neuronal and intestinal xbp-1s, grown on control (empty vector) or hlh-30 RNAi. Bar graphs represent mean chemotaxis index ± SD. N = 70–130 animals per assay; each assay was independently replicated 3 times. Significance was assessed by two-way ANOVA with Dunnett’s multiple comparisons test.

(ii) Chemotaxis ability in animals expressing polyQ40 in neurons in combination with neuronal and intestinal xbp-1s, grown on control (empty vector) or hlh-30 RNAi. Bar graphs represent mean chemotaxis index ± SD. N = 70–130 animals per assay; each assay was independently replicated 3 times. Significance was assessed by two-way ANOVA with Dunnett’s multiple comparisons test, **p < 0.01. See also Figures S5 and S6 and Table S1.
Figure 6. Lysosomes Are Required for xbp-1s-Mediated Reduction in PolyQ Levels

(A) Western blot analysis of (i) neuronal or (ii) intestinal polyQ40::YFP at day 1 and day 4 of adulthood, expressed in combination with neuronal and intestinal xbp-1s and grown on control (empty vector) or lmp-1 RNAi. Lysates containing total proteins were resolved under native conditions and blotted with an anti-polyQ antibody. Tubulin levels were probed with α-α-tubulin as a loading control. Data are representative of at least 3 independent experiments. Quantification of (iii) neuronal or (iv) intestinal polyQ40::YFP was conducted using ImageJ. Graphs represent mean band intensity relative to day 1 polyQ ± SD. Statistical significance was assessed between control and lmp-1 for each genotype and time point, using two-way ANOVA with Tukey’s multiple comparisons, *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001.

(B) (i) Representative confocal images of the intestine of animals expressing LMP-1::GFP, with and without rab-3p::xbp-1s. Animals were grown on OP50 and imaged at days 2 and 5 of adulthood. Imaging was performed at X63 magnification. Scale bar, 10 μm.

(legend continued on next page)
**Neuronal xbp-1s Increases Lysosomal LMP-1 Labeling**

As xbp-1s expression increases transcription of lysosomal genes, and lysosome activity is required for clearance of toxic species, we asked whether the number of lysosomes in intestinal cells was increased by neuronal xbp-1s expression. To do this, we used animals expressing GFP-tagged LMP-1 to label lysosomes [29]. When LMP-1::GFP was combined with rab-3p::xbp-1s, a small increase in lysosome number was observed (Figures 6Bi and 6Bii), while when we examined animals expressing LGG-1 tagged with GFP, we found that there was no increase in number of GFP::LGG-1-labeled intestinal autophagosomes in animals expressing neuronal xbp-1s (Figure S6E) [30]. Furthermore, when we measured the GFP fluorescence intensity of individual LMP-1::GFP-labeled lysosomes, we found a significant increase in LMP-1 labeling in rab-3p::xbp-1s nematodes (Figures 6Bi and 6Bii). This might suggest an increase in LMP-1 recruitment to lysosomes and led us to speculate that their activity may be altered by neuronal xbp-1s.

**Neuronal xbp-1s Increases Lysosomal Acidity**

Measures of autophagic flux in *C. elegans* suggest a decline in lysosomal activity with age [31]. The activity of lysosomes depends on their acidity, which is necessary for the activity of lysosomal hydrolases. We therefore asked whether neuronal xbp-1s could increase or preserve lysosomal acidity with age.

To explore this, we utilized a technique to sensitively measure lysosomal acidity in the proximal intestine of the worm [32]. Using carboxy-2',7'-dichlorofluorescein diacetate (cDCFDA), which is hydrolyzed to fluorescent cDCF in acidic conditions, we found that neuronal xbp-1s significantly increases the acidity of intestinal lysosomes, particularly as animals age; at day 2, and especially at day 5 of adulthood, lysosomes of rab-3p::xbp-1s animals were significantly more acidic, and larger, than those in wild-type worms (Figure 7A). While tissue-specific promoters may lose some specificity from day 5 [33], we were reassured by the observation that increased acidity was already present at day 2. Greater activity of lysosomal protein hydrolases in rab-3p::xbp-1s animals was also observed—while activity declined between day 2 and 5 in wild-type animals, it increased significantly in rab-3p::xbp-1s nematodes (Figure 7B). To ensure that cDCFDA was truly detecting changes to lysosomal acidity, we confirmed that RNAi against *daf-2* increased acidity-dependant staining of lysosomes, as expected, and that knockdown of the downstream transcription factor *daf-16* reduced acidity (Figure S7A). In addition, staining was reduced by knockdown of two subunits of the acidifying vacuolar ATPase, *vha-2* and *vha-8* (Figure S7B). We then asked whether RNAi against the vacuolar ATPase subunit upregulated by xbp-1s, *vha-18* also suppressed lysosomal acidity. We found that, indeed, *vha-18* knockdown reduced lysosomal acidification in rab-3p::xbp-1s animals, suggesting that elevated levels of this subunit might be a mechanism by which xbp-1s orchestrates higher lysosomal acidity and activity (Figure 7C).

This difference in lysosome acidity was also apparent in animals expressing neuronal polyQ<sub>40</sub>. Background intestinal fluorescence was substantially increased in these animals, possibly reflecting an increase in overall cellular acidity; this increase, as well as the acidity of lysosomes, was lost by day 5 of adulthood but was significantly increased in animals expressing neuronal xbp-1s, suggesting a mechanism for improved polyQ clearance (Figure S7C).

To confirm these findings, we stained intestinal lysosomes with LysoTracker, another pH-dependent dye. Lysosomal staining was again increased in animals expressing neuronal or intestinal xbp-1s at both day 2 and particularly at day 5 of adulthood; neuronal polyQ<sub>40</sub> was also associated with age-dependent collapse in lysosomal acidity and number, rescued by co-expression of neuronal or intestinal xbp-1s (Figures S7D and S7E). Similarly, muscle-specific polyQ<sub>35</sub> decreased number and staining of intestinal lysosomes by day 5, rescued by expression of neuronal but not muscle-driven xbp-1s (Figure S7F). Together, these results confirm that xbp-1s expression in neurons or the intestine increases the acidity of intestinal lysosomes, rescuing the decline seen with age and in animals expressing misfolded proteins.

**Neuronal xbp-1s Promotes Maturation of Autophagosomes to Autolysosomes**

Finally, we asked whether xbp-1s expression was able to promote the maturation of autophagosomes to autolysosomes. To address this, we used a tandem tag reporter, with GFP and mCherry conjugated to LGG-1 [31]. Autophagosomes are labeled with both GFP and mCherry, while the acidification associated with maturation of autophagosomes into autolysosomes quenches GFP, so that autolysosomes are only labeled with mCherry. Using this system, we observed a striking enrichment in autolysosomes at day 2 and especially at day 5 in neuronal xbp-1s animals, while autophagosome number decreased in older animals (Figure 7D). This suggests that autolysosomes are more prevalent, while autophagosome turnover may be increased, in xbp-1s-expressing animals, which would be consistent with increased function of the lysosomal compartment. Together, these results suggest that xbp-1s transcriptionally activates lysosomes in the intestine of *C. elegans*, leading to improved clearance of proteotoxic species and increased longevity (Figure 7E).

**DISCUSSION**

In this work, lysosomes are identified as a key organelle mediating the downstream effects of cell non-autonomous UPR<sub>ER</sub> activation. Expression of xbp-1s in neurons or the intestine improves proteostasis across tissues, suppressing the toxicity associated with aggregation-prone proteins by reducing levels of toxic species. This is accomplished by increased transcription of lysosomal genes in the intestine and increases in lysosomal

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(ii) Quantification of lysosome number. LMP-1::GFP-positive punctae were counted in 10–15 worms per genotype at day 2 and day 5 of adulthood using ImageJ, from 3 independent biological replicates. Statistical analysis was carried out using one-way ANOVA with Tukey’s multiple comparisons test, *p < 0.05.

(iii) Quantification of lysosomal LMP-1::GFP labeling. Fluorescence intensity of LMP-1::GFP per lysosome was quantified using ImageJ in 10–15 worms per genotype, at day 2 and day 5 of adulthood, in 3 biological replicates. Statistical significance was calculated using one-way ANOVA with Tukey’s multiple comparisons test, **p < 0.01, ***p < 0.0001. See also Figure S6.
Figure 7. Neuronal xbp-1s Increases Lysosomal Acidity

(A) (i) Confocal imaging of lysosomal acidity in N2 and rab-3p::xbp-1s intestines. Animals were grown on OP50 and transferred to plates containing cDCFDA 16 h prior to imaging. Imaging was conducted in the anterior intestine at 63× magnification in day 2 and day 5 adults. Yellow arrowheads indicate representative lysosomes. Scale bar, 10 μm.

(ii) Quantification of lysosomal acidity in wild-type and rab-3p::xbp-1s intestines. Animals were grown on OP50 and transferred to plates containing cDCFDA 16 h prior to imaging; imaging was conducted as described above, and fluorescence quantified from 3 experiments using ImageJ. Plots represent 5–10 animals per replicate. Significance was assessed between N2 and rab-3p::xbp-1s at each time point by one-way ANOVA with Tukey’s multiple comparisons, *p < 0.05; ***p < 0.001.

(B) Protease activity in rab-3p::xbp-1s and N2 animals. Following incubation with a fluorescent substrate, relative fluorescence units (RFU) at 400/505 nm were measured. Bar graphs represent mean ± SD from 3 independent biological replicates. Significance was calculated using one-way ANOVA with Tukey’s multiple comparisons test; ns, not significant, **p < 0.01.

(C) (i) Confocal imaging of lysosomal acidity in N2 and rab-3p::xbp-1s animals grown on control (empty vector) or vha-18 RNAi and transferred to plates containing cDCFDA 16 h prior to imaging. Imaging was conducted as described above. Scale bar, 10 μm.

(ii) Quantification of lysosomal acidity in N2 and rab-3p::xbp-1s animals grown on control (empty vector) or vha-18 RNAi. Animals were imaged as above, and quantification of fluorescence from 3 experiments were carried out using ImageJ. Plots represent 8–10 animals per replicate. Significance was assessed between N2 and rab-3p::xbp-1s at each time point by one-way ANOVA with Tukey’s multiple comparisons. ***p < 0.0001.

(D) (i) lgg-1p::mCherry::GFP::lgg-1 and rab-3p::xbp-1s; lgg-1p::mCherry::GFP::lgg-1 animals were grown on OP50 and imaged at days 2 and 5 of adulthood at X63 magnification. Arrowheads in lgg-1p::mCherry::GFP::lgg-1 indicate representative autophagosomes; arrowheads in rab-3p::xbp-1s; lgg-1p::mCherry::GFP::lgg-1 indicate representative autolysosomes. Scale bar, 10 μm.

(ii) Quantification of autolysosomes. mCherry::LGG-1-positive (magenta) punctae were counted in 10–15 worms per genotype at day 2 and day 5 of adulthood using ImageJ. Data are derived from 3 independent biological replicates. Statistical analysis was carried out using one-way ANOVA with Tukey’s multiple comparisons test, ***p < 0.0001.

(iii) Quantification of autophagosomes. mCherry::LGG-1::GFP-positive (white) punctae were counted in 10–15 worms per genotype at day 2 and day 5 of adulthood using ImageJ. Data are derived from 3 independent biological replicates. Statistical analysis was carried out using one-way ANOVA with Tukey’s multiple comparisons test, **p < 0.01.

(E) Schematic of the regulation of lifespan and proteostasis by neuronal and intestinal xbp-1s through increased lysosome activity. See also Figure S7.
variety of physiological processes [48–50]. They also play roles in lipid signaling pathways that modulate aging—lysosomal lipases can release lipid-derived molecules able to mediate nuclear responses that promote longevity [51]. Future work to measure the number and pH status of lysosomes in tissues other than the intestine would confirm directly whether lysosome activation is achieved throughout the organism following intestinal UPRER activation. The identification of such a lysosome-activating signaling pathway would then be of great interest.

Lysosome activity is critical for extension of longevity by neuronal xbp-1s. Fusion with lysosomes is the end point of autophagy, and a wealth of data links autophagy to aging and neurodegeneration [52, 53]. However, we do not observe a requirement for autophagy genes in xbp-1s-mediated lifespan extension. Some evidence already suggests that, as here, lysosomes may play autophagy-independent roles in aging and proteostasis. In the C. elegans germline, for example, rejuvenation of the proteome and clearance of aggregates depends upon the lysosome, without a requirement for autophagy. Similarly, prevention of protein aggregate seeding and spreading in the worm depends on lysosomes, but not autophagosomes [39, 54].

Lysosomes have been shown to decline in function with age; measures of autophagic flux in C. elegans suggest age-dependent deterioration in lysosome activity, and, in yeast, age-associated decline in the acidity of the lysosome-like vacuole limits lifespan; preventing this decline extends longevity [31, 55]. Vacuolar acidity is increased by caloric restriction, leading to lifespan extension, and acidic vacuoles are restored in daughtier cells upon budding, renewing lifespan [55]. In C. elegans oocytes, sperm trigger the activation of lysosomes through the activation of the vacuolar H⁺-ATPase, clearing protein aggregates and renewing the proteostasis state of the cell [54].

This accumulating evidence suggests that vacuolar or lysosomal acidification could act as a molecular “switch” to rejuvenate the proteome of cells and restore youthfulness. Our results indicate that xbp-1s may be able to activate this lysosomal switch through physiological inter-tissue signaling. Understanding the nature of these signals, both those that communicate UPRER activity between neurons and the intestine, and those that may communicate lysosomal activation between the intestine and other tissues, is now an important goal for research into aging and neurodegeneration.

**STAR+METHODS**

Detailed methods are provided in the online version of this paper and include the following:

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**QUANTIFICATION AND STATISTICAL ANALYSIS**

**DATA AND CODE AVAILABILITY**

Supplemental Information can be found online at https://doi.org/10.1016/j.cub.2019.06.031.

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**AUTHOR CONTRIBUTIONS**

S.I. and R.C.T. conceived, performed, and analyzed experiments and wrote the manuscript. N.P.O. developed methodology, performed and analyzed experiments, and curated data. C.K. provided expertise and analyzed data. M.O.C. provided expertise and resources and reviewed the manuscript.

**DECLARATION OF INTERESTS**

The authors declare no competing interests.

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## STAR METHODS

### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| Antibodies          |        |            |
| α-Aβ                | BioLegend | 6E10; Cat#803001; RRID: AB_2564653 |
| α-polyQ             | Sigma Aldrich | 3B5H10; Cat#MABN821; RRID: AB_532270 |
| α-GFP/YFP           | ChromoTek | 3H9; Cat#3h9-100; RRID: AB_10773374 |
| α-α-tubulin         | Sigma Aldrich | B512; Cat#T5168; RRID: AB_477579 |
| Bacterial and Virus Strains |        |            |
| E. coli OP50        | CGC     | WB OP50; RRID: WB-STRAIN:OP50 |
| E. coli HT115       | CGC     | WB HT115; RRID: WB-STRAIN:HT115 |
| L4440 RNAi          | Addgene | Cat#1654 |
| bec-1 RNAi          | Vidal ORF-RNAi library, Source Bioscience | Cat#3320_Cel_ORF_RNAi |
|yps-34 RNAi          | Vidal ORF-RNAi library, Source Bioscience | Cat#3320_Cel_ORF_RNAi |
| xbp-1 RNAi          | Vidal ORF-RNAi library, Source Bioscience | Cat#3320_Cel_ORF_RNAi |
|asp-3 RNAi           | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|atg-13 RNAi          | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|atg-18 RNAi          | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|cup-5 RNAi           | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|hnh-30 RNAi          | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|lmp-1 RNAi           | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|vha-15 RNAi          | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|vha-18 RNAi          | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|vha-2 RNAi           | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|vha-8 RNAi           | Ahringer library, Source Bioscience | Cat#3318_Cel_RNAi_complete |
|daf-2 RNAi           | Dillin lab, UC Berkeley | N/A |
|daf-16 RNAi          | Dillin lab, UC Berkeley | N/A |
| Chemicals, Peptides, and Recombinant Proteins |        |            |
| Carbenicillin       | Formedium | Cat#CAR0025 |
| IPTG                | Generon | Cat#orb340528 |
| Sodium azide        | Sigma Aldrich | Cat#S2002 |
| Levamisole          | Sigma Aldrich | Cat#L0380000 |
| Tetramisole         | Sigma Aldrich | Cat#T1512-5G |
| Benzaldehyde        | Sigma Aldrich | Cat#B1334-250ML |
| AlexaFluor 546-Phalloidin | Life Technologies Ltd. | Cat#A22283 |
| Papain              | AppliChem GmbH | Cat#A3824 |
| Fetal bovine serum  | GIBCO, Life Technologies | Cat#10270106 |
| Trizol LS           | Ambion, Life Technologies | Cat#10296010 |
| SYBRGreen Master Mix | Applied Biosystems | Cat#4472897 |
| cDCFDA              | ThermoFisher | Cat#C369 |
| LysoTracker DeepRed | ThermoFisher | Cat#L12492 |
| Critical Commercial Assays |        |            |
| Pierce BCA protein assay kit | Thermo Scientific | Cat#23227 |
| Direct-zol kit      | Zymo Research | Cat#R2060 |
| NuGEN Ovation RNaseq v2 kit | NuGEN | Cat#7102 |
| Agilent High Sensitivity DNA kit | Agilent | Cat#5067-4626 |
| Ovation Ultralow System v2 kit | NuGEN | Cat#0344 |

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Continued

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| KAPA Library Quantification Kit | Illumina Platforms | Cat#07960140001 |
| Agilent RNA 6000 Pico Kit | Agilent | Cat#5067-1513 |
| QuantiTect reverse transcription kit | QIAGEN | Cat#205310 |
| Fluorometric assay kit | Abcam | Cat#ab65300 |

**Deposited Data**

| NCBI | BioProject | Identifier |
|------|------------|------------|
|      | PRJNA482604 |            |

**C. elegans strains**

| Strain | Description | Source |
|--------|-------------|--------|
| N2     | Wild type, Bristol | CGC [56] |
| AGD927 | uhsIs270[rab-3p::xbp-1s, myo-2p::tdTomato] | CGC [6] |
| CL2355 | dvlIs50[snb-1::Aβ1-42, mtl-2::GFP] | CGC [13] |
| CX51   | dyn-1(ky51) | CGC [7] |
| SD551  | let-60(ga89) | CGC [7] |
| AM140  | rmlIs132[unc-54p::Q35::YFP] | CGC [8] |
| CL2006 | dvlIs2[unc-54p::Aβ1-42, rol-6(su1006)] | CGC [17] |
| CB1402 | unc-15(e1402) | CGC [7] |
| WM27   | rde-1(ne219) | CGC [24] |
| VP303  | rde-1(ne219); kils7 [php-2::rde-1 + rol-6(su1006)] | CGC [29] |
| AMJ345 | rde-1(ne219); jamS12 [mex-5::rde-1] | CGC [28] |
| RT258  | pwlIs50 [imp-1::imp-1::GFP, Cbr-unc-119(+)]] | CGC [29] |
| DA2123 | adls2122[lgg-1p::GFP::lgg-1, rol-6(su1006)] | CGC [30] |
| MAH215 | sqIs11[lgg-1p::mCherry::GFP::lgg-1 + rol-6] | CGC [31] |
| MAH240 | [sqIs17 [hhl-30::hhl-30::GFP + rol-6] | CGC [27] |
| AGD197 | rmlIs110[gef-1p::Q40::YFP] | Dillin lab (outcrossed from [14]) |
| AGD195 | uhsIs393[vha-6p::Q40::YFP, rol-6(su1006)] | Dillin lab (integrated from [19]) |
| AGD179 | uhsIs388[glv-19p::xbp-1s, myo-2p::tdTomato] | This paper (integrated from [8]) |
| AGD191 | uhsIs390[unc-54p::xbp-1s, myo-2p::tdTomato] | This paper (integrated from [8]) |
| AGD1412 | qcls18(ges-1p::GFP) | This paper (outcrossed from [22]) |
| RCT51  | uhsIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; dvlIs2[unc-54p::Aβ1-42, rol-6(su1006)] | This paper |
| RCT2   | uhsIs388[glv-19p::xbp-1s, myo-2p::tdTomato]; dvlIs2[unc-54p::Aβ1-42, rol-6(su1006)] | This paper |
| RCT3   | uhsIs390[unc-54p::xbp-1s, myo-2p::tdTomato]; dvlIs2[unc-54p::Aβ1-42, rol-6(su1006)] | This paper |
| RCT23  | uhsIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; dvlIs50[snb-1::Aβ1-42, mtl-2::GFP] | This paper |
| RCT25  | uhsIs388[glv-19p::xbp-1s, myo-2p::tdTomato]; dvlIs50[snb-1::Aβ1-42, mtl-2::GFP] | This paper |
| RCT1   | uhsIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; rmlIs132[unc-54p::Q35::YFP] | This paper |
| RCT146 | rmsEx15[glv-19p::xbp-1s, myo-2p::tdTomato]; rmlIs132[unc-54p::Q35::YFP] | This paper |
| RCT4   | uhsIs390[unc-54p::xbp-1s, myo-2p::tdTomato]; rmlIs132[unc-54p::Q35::YFP] | This paper |
| AGD1399 | uhsIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; rmlIs110[gef-1p::Q40::YFP] | This paper |
| RCT31  | uhsIs388[glv-19p::xbp-1s, myo-2p::tdTomato]; rmlIs110[gef-1p::Q40::YFP] | This paper |
| RCT32  | uhsIs390[unc-54p::xbp-1s, myo-2p::tdTomato]; rmlIs110[gef-1p::Q40::YFP] | This paper |
| RCT33  | uhsIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; uhsIs393[vha-6p::Q40::YFP, rol-6(su1006)] | This paper |

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**LEAD CONTACT AND MATERIALS AVAILABILITY**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Rebecca Taylor (rtaylor@mrc-lmb.cam.ac.uk).

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| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| uthIs388[gly-19p::xbp-1s, myo-2p::tdTomato]; uthIs393[ha-6p::Q40::YFP, rol-6[su1006]] | This paper | RCT26 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; dyn-1[ky51] | This paper | RCT59 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; unc-15[e1402] | This paper | RCT60 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; let-60[ga89] | This paper | RCT112 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; zcls18[ges-1p::GFP] | This paper | RCT52 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; rde-1(ne219); kbls7 [ntx-2p::rde-1 + rol-6[su1006]] | This paper | RCT145 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT105 |
| dvls50[snb-1::A1-42, mtl-2::GFP]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT109 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; dvls50[snb-1::A1-42, mtl-2::GFP]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT110 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; dvls50[snb-1::A1-42, mtl-2::GFP]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT111 |
| rmls110[rgef-1p::Q40::YFP]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT106 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; rmls110[rgef-1p::Q40::YFP]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT107 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; rmls110[rgef-1p::Q40::YFP]; rde-1(ne219); jamSi2 [mex-5p::rde-1] | This paper | RCT108 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; sqs17 [h1h-30p::h1h-30::GFP + rol-6] | This paper | RCT94 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; pwls50 [imp-1p::imp-1::GFP, Cbr-unc-119+] | This paper | RCT103 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; adls2122[ggg-1p::GFP::lgg-1, rol-6[su1006]] | This paper | RCT104 |
| uthIs270[rab-3p::xbp-1s, myo-2p::tdTomato]; sqs111[ggg-1p::mCherry::GFP::lgg-1+ rol-6] | This paper | RCT98 |

Oligonucleotides

See Table S2

Software and Algorithms

| SeqMonk | Babraham Institute | https://www.bioinformatics.babraham.ac.uk/projects/seqmonk/ |
| g:Profiler | ELIXIR | https://biit.cs.ut.ee/gprofiler/index.cgi |
| Promoterome Database | Harvard University | http://woldb.dfci.harvard.edu/promoteromedb/ |
EXPERIMENTAL MODEL AND SUBJECT DETAILS

**C. elegans maintenance and RNAi knockdown**

*C. elegans* strains were maintained at 20°C on nematode growth medium (NGM) plates seeded with OP50 bacteria unless otherwise stated [56]. For feeding RNAi experiments either L4440 empty vector or the designated RNAi bacteria was used [57]. Plates for RNAi analysis were prepared by supplementation of agar with 100 µg/mL carbenicillin and 1mM IPTG after autoclaving. 24 hours prior to each assay plates were spotted with 100 µL of overnight bacterial culture. Where dual RNAi knockdown was performed, equal volumes of two cultures normalized to equivalent OD600 were mixed before plates were seeded.

METHOD DETAILS

**Chemotaxis assays**

Animals were synchronized by timed egg lay on 55 mm NGM plates. Chemotaxis assays were conducted at day 1 of adulthood as described (section 4.4 of [58]). Briefly, animals were raised on OP50 or designated RNAi bacteria until day 1 of adulthood, when they were collected and washed three times using M9 buffer, with worms settled between washes by gravity. Assay plates were prepared using a 1 µL spot of ethanol (solvent spot) and a 1 µL spot of chemo-attractant, 1:100 benzaldehyde (target spot) on opposite sides. 1 µL of sodium azide (50mM) was added to solvent and target spots once dried. Worms were dispensed at the center of the plate and animals kept in the dark for 60 min at room temperature before worms at each spot were counted. Chemotaxis indices were then calculated as (# worms at target spot - # worms at vehicle spot)/total number of worms on the plate. Indices were reported as ~1 to 1, where 1 indicates that 100% of animals have arrived at the target spot. All assays were repeated at least 3 times, and significance was assessed by one-way ANOVA with Tukey’s multiple comparisons test.

**Paralysis assays**

Eggs were isolated by bleaching and allowed to develop on a bacterial lawn. At the L4 larval stage, individual nematodes were picked and transferred to 55mm NGM plates seeded with OP50. Paralysis was evaluated daily from day 1 to day 12 of adulthood and scored by touching the animal’s nose with a platinum wire. Worms able to move the head but not the rest of the body were scored as paralyzed [16]. Dead animals or those with other phenotypes (e.g., vivipary) were censored from the analysis. All assays were performed with at least 100 animals and repeated at least 3 times.

**Osmoregulatory assays**

Age-matched animals were grown at 20°C. At day 1 of adulthood, worms were placed into a drop of distilled water for 5 minutes, after which animals with a swollen, fluid-filled shape were scored as osmoregulation defective (Osm), as described [15]. All assays were replicated 3 times and significance was assessed by two-way ANOVA with Tukey’s multiple comparison.

**Motility assays**

Animals were synchronized by timed egg lay and grown to adulthood on OP50 or RNAi bacteria. 10-15 day 1 or day 4 adult worms of each genotype were transferred to a 24-well dish containing 700 µL of M9 buffer and allowed to settle for 1 minute, after which each well was filmed for 30 s at a frame rate of 15 fps. Movies were analyzed manually to assess thrashing rate. Each assay was repeated 3 times and significance assessed by one-way ANOVA with Tukey’s multiple comparisons test.

**F-actin staining**

Synchronized worms were freeze cracked to permeabilize the cuticle at day 5 of adulthood. Staining was performed using AlexaFluor 546-Phalloidin to visualize F-actin in body wall muscles, as described (section 2.6 of [59]; section 2.3 of [60]) in 3 independent biological replicates.

**Fluorescence microscopy**

Micrographs of worms were acquired using a Leica M205 FA microscope and LAS X software. 5-10 animals were anesthetized using levamisole (25 mM) or sodium azide (50 mM) prior to imaging. All fluorescence microscopy analysis was independently replicated at least 3 times.

**Confocal microscopy**

For live microscopy nematodes were mounted on a 2% agarose pad and anesthetized using levamisole (25 mM). For HLH-30::GFP animals, sodium azide (50 mM) was used to avoid the effects of levamisole on nuclear localization of HLH-30. Fixed animals were prepared as described for F-actin staining. Images were acquired using a Zeiss LSM 710 confocal microscope using the 20x air, or 40x and 63x oil immersion objectives. A single section was acquired for all imaging and the pinhole used was 1 AU for optimal section thickness using the smart setup function. Image analysis utilized ImageJ or Fiji. For tandem-tagged nematodes, confocal microscopy was performed as described [31], where the Z-position was selected so that nuclei were clearly in focus in...
order to visualize autolysosomes and autophagosomes. All confocal microscopy analysis was independently replicated at least 3 times.

**Protein extraction and Western Blotting**

For native extracts, nematodes were synchronized by bleaching and allowed to grow on designated bacterial strains until day 1 or 4 of adulthood. Worms were then washed three times with M9 buffer and pelleted before addition of 75-80 µL of native lysis buffer (as described in [61]). Samples were flash frozen in liquid nitrogen prior to use. Nematodes were thawed on ice and mechanically disrupted using a Precellys (Bertin Instruments) programmed for 3x15 s pulses, with 30 s between each pulse. Samples were then centrifuged for 5 minutes (8000 g) and supernatant containing total proteins was transferred to fresh tubes to resolve using NativePAGE 4%–16% Bis-Tris (Invitrogen). Gels were transferred using the iBlot 7-Minute Blotting System (Thermo Fisher Scientific) and imaged using ChemiDoc (BioRad). ImageLab software (BioRad) was further used to analyze the bands. For denaturing conditions, total cell lysate was prepared in RIPA buffer (with 2% SDS). Samples were centrifuged for 15min at high speed and pellets were resuspended in urea (7M) and treated with DTT. Samples were then boiled at 95°C for 5 minutes prior to loading on NuPAGE 4%–12% Bis Tris gels (Invitrogen). Gels were transferred and imaged as described above. For both native and denaturing conditions total protein concentrations were assayed using a Pierce BCA protein assay kit. All western blotting experiments were repeated at least 3 times.

**Quantification of protein band intensity**

Protein bands or lanes were quantified using ImageJ. The designated band or lane was selected and the band intensity calculated. Intensity was then normalized to the appropriate control and significance assessed by two-way ANOVA with Tukey’s multiple comparisons test [62].

**C. elegans cellular dissociation**

Using animals with an intestinal GFP marker (RCT51 and RCT52), intestinal cells were isolated following the method described in [21]. Briefly, animals were harvested at day 1 of adulthood and washed several times to remove bacteria from their gut and surroundings. Cuticles were then permeabilized using a Triton X-100-SDS-DTT solution. For enzymatic digestion, papain (10 mg/mL) was used and animals were also mechanically disrupted using an electronic hand homogenizer (IKA T10 basic, ULTRA-TURRAX). Following this step, cells were recovered in fetal bovine serum (FBS) solution and kept on ice for sorting.

**C. elegans cellular isolation by FACS**

Intestinal cell suspensions (PBS/2% FBS) were passed through 35 and 40 mm nylon cell strainers (FALCON). Filtered cells were then diluted in the same media as above and sorted by a Sony iCyt Synergy Dual Channel, high speed cell sorter (488 nm excitation). Gates were used to eliminate cells with tdTomato and autofluorescence. GFP positive fluorescent events were collected in 1.5 mL eppendorf tubes containing 10-20 µL PBS/2% FBS. Cells were kept on ice for the RNA extraction step. Collected positive fluorescent events varied between 20,000 and 60,000 events for intestinal samples.

**RNA extraction and amplification**

FACS-sorted intestinal cells were centrifuged and cell volumes normalized to 100 µL. Immediately 400 µL of Trizol LS was added to the tubes which were then snap frozen in liquid nitrogen. RNA was extracted, DNase digested and cleaned using a microprep Direct-zol kit (Zymo Research). Initial RNA quantities for downstream analysis were normalized to 1.5 ng. Agilent Bioanalyzer RNA Pico chips were used to assess the quantity and quality of RNA. Amplified cDNA was generated using the NuGEN Ovation RNaseq v2 kit. DNA quality and quantity were then evaluated using an Agilent High Sensitivity DNA kit and Nanodrop (Nanodrop 2000c, Thermo Scientifics). Thereafter, cDNA (1 µg) was sheared using a Covaris M220 sonicator (Covaris) to an average size of 150-200bp.

**Library preparation, RNA sequencing and analysis**

Library preparation was performed using the NuGEN Ovation Ultralow System v2 kit. Input cDNA was normalized to 90 ng. After purification, libraries were assessed using a Bioanalyzer High Sensitivity DNA chip. Fragment size was around 300 bp. Samples (10,000x and 20,000x dilutions) were quantified using the KAPA Library Quantification Kit. Samples were then pooled and quantified as before and further diluted to 15 nM for sequencing on an Illumina HiSeq4000 (CRUK CI, Cambridge). Reads were trimmed using Trim Galore (v0.4.5, cutadapt 1.15) and mapped to C. elegans WBcel235 using Hisat2 (v2.1.0). rRNA reads were removed and analysis performed using SeqMonk (https://www.bioinformatics.babraham.ac.uk/projects/seqmonk/). Differentially expressed genes were identified using an intensity difference filter with a cut-off point of p < 0.05. Gene ontology was analyzed using g:Profiler (https://biit.cs.ut.ee/gprofiler/index.cgi), with a custom-created background gene list based on the genes expressed in intestinal cells of rab-3p::xbp-1s worms, with log2 RPMK values between 0-100. Intestine-specific, muscle-specific and neuron-specific gene lists were obtained using the Princeton database of tissue-specific expression predictions for C. elegans (http://worm-tissue.princeton.edu/search/download), where only the genes with a score of 1 for the relevant tissue and a score of 0 for all the other tissues were chosen.
Quantitative RT-PCR
Total RNA was extracted as described above and 2 ng of purified RNA was used for cDNA synthesis using a QuantiTect reverse transcription kit. SYBRGreen quantitative RT-PCR was performed using either the Corbett system and following the Rotor-Gene 6000 Series Software manual, or a Vii7 Real-Time PCR (ThermoFisher Scientific). Data from 3 biological repeats were analyzed using the comparative 2ΔΔCt method. Significance was assessed by one-way ANOVA with Dunnett’s multiple comparisons test.

Lifespan analysis
Lifespan analyses were performed at 20°C and were repeated at least twice. Typically, a minimum of 100 animals was used per condition, and worms were scored for viability every second day, from day 1 of adulthood (treating the pre-fertile day preceding adulthood as t = 0). Lifespans were performed on E. coli OP50, unless otherwise indicated, and animals were treated with 100 μg/mL FUDR at t = 0 and again at day 5 of adulthood, to avoid issues associated with early death through vivipary (“bagging”) in rab-3p::xbp-1s animals. Prism 7 software was used for statistical analysis, and significance calculated using the log-rank (Mantel–Cox) method.

Quantification of lysosomes
RT258 and RCT103 worms expressing LMP-1::GFP were used to quantify the number of fluorescently-labeled lysosomes in the intestine of 10-15 animals per genotype from 3 independent biological replicates. ImageJ was used to analyze the data by counting numbers of LMP-1::GFP positive punctae. Fluorescence intensity of LMP-1::GFP punctae was also quantified at each time point and xbp-1s-expressing animals normalized to LMP-1::GFP alone. Quantification was performed using the oval-shaped selection tool to select the region of interest (ROI). GFP intensity was then quantified using the Measure function and background fluorescence intensity subtracted prior to normalization. Statistical analysis was carried out using a one-way ANOVA with Tukey’s multiple comparisons test.

Lysosome acidity assays
To measure lysosomal acidity, the fluorescent pH-sensitive dye cDCFDA (5(6)-carboxy-2’,7’-dichlorofluorescein diacetate), which converts to cDCF (5-(6)-carboxy-2’,7’-dichlorofluorescein) through hydrolysis, was fed to live worms [32]. Animals were age-matched and grown on OP50 until day 1 or 4 of adulthood, then transferred to assay plates prepared by the addition of 100 μL of freshly prepared cDCFDA (10 mM) to a fresh lawn of OP50 bacteria in a 2cmx2cm patch. Plates were then kept in the dark to absorb dye into the agar/bacterial patch. Single worms picked onto assay plates were allowed to feed on the bacteria/cDCFDA patch for at least 16 hours prior to imaging the anterior section of the intestine, in 5-10 animals per genotype from 3 replicates. Data was analyzed using ImageJ, and statistical analysis carried out using a one-way ANOVA with Tukey’s multiple comparisons test, similarly to the intensity quantification of LMP-1::GFP.

LysoTracker staining
C. elegans were grown from hatch to day 2 and 5 of adulthood on OP50- or RNAi bacteria-seeded plates containing 25 μM LysoTracker DeepRed or an equivalent volume of DMSO. Worms were then imaged by confocal microscopy, using 8-10 animals per genotype and 3 independent biological replicates; quantification was carried out using ImageJ, and statistical analysis conducted by one-way ANOVA with Tukey’s multiple comparisons test.

Protease activity assays
Enzymatic activity of lysosomal proteases was determined using an Abcam fluorometric assay kit. Age-matched worms were grown on OP50 and lysed at day 2 and 5 of adulthood. Total proteins were extracted as described previously and protein concentration quantified and normalized. After adjusting volumes to 50 μL/well, 50 μL of reaction buffer was added per well followed by 2 μL of 10mM substrate (Ac-RR-AFC). Samples were incubated at 37°C for 1-2 hours and protease activity assessed using a fluorescent microplate reader (TECAN) at Ex/Em = 400/505 nm. All assays were prepared in triplicate and repeated 3 times independently, and statistical analysis carried out using a one-way ANOVA with Tukey’s multiple comparisons test.

Tandem quantification of autophagosomes and autolysosomes
MAH215 worms expressing mCherry::GFP::LGG-1 were used to quantify the number of fluorescently-labeled autophagosomes and autolysosomes in the intestine of 10-15 animals per genotype from 3 independent biological replicates. ImageJ was used to analyze the data by counting numbers of mCherry::GFP positive autophagosomes, and mCherry positive autolysosomes. Statistical analysis was carried out using a one-way ANOVA with Tukey’s multiple comparisons test.
QUANTIFICATION AND STATISTICAL ANALYSIS

Tests used to determine statistical significance were one-way ANOVA with Dunnett’s multiple comparisons test (qRT-PCR); one-way ANOVA with Tukey’s multiple comparison test (chemotaxis, motility and protease activity assays; quantification of lysosomes, autophagosomes, autolysosomes, lysosomal acidity, and Lysotracker staining); two-way ANOVA with Tukey’s multiple comparisons test (osmoregulation assays and quantification of protein bands); and Mantel-Cox log-rank test (lifespan assays). Statistical information for each experiment can be found in the corresponding figure legend.

DATA AND CODE AVAILABILITY

The accession number for the RNA-seq data reported in this paper is NCBI BioProject number PRJNA482604.