Regulation of age-related structural integrity in neurons by protein with tau-like repeats (PTL-1) is cell autonomous

Yee Lian Chew1, Xiaochen Fan1, Jürgen Götz2 & Hannah R. Nicholas1

1School of Molecular Bioscience, University of Sydney, Australia, 2Clem Jones Centre for Ageing Dementia Research (CJCADR) at the Queensland Brain Institute (QBI), University of Queensland, Australia.

PTL-1 is the sole homolog of the MAP2/MAP4/tau family in Caenorhabditis elegans. Accumulation of tau is a pathological hallmark of neurodegenerative diseases such as Alzheimer’s disease. Therefore, reducing tau levels has been suggested as a therapeutic strategy. We previously showed that PTL-1 maintains age-related structural integrity in neurons, implying that excessive reduction in the levels of a tau-like protein is detrimental. Here, we demonstrate that the regulation of neuronal ageing by PTL-1 occurs via a cell-autonomous mechanism. We re-expressed PTL-1 in a null mutant background using a pan-neuronal promoter to show that PTL-1 functions in neurons to maintain structural integrity. We next expressed PTL-1 only in touch neurons and showed rescue of the neuronal ageing phenotype of ptl-1 mutant animals in these neurons but not in another neuronal subset, the ventral nerve cord GABAergic neurons. Knockdown of PTL-1 in touch neurons also resulted in premature neuronal ageing in these neurons but not in GABAergic neurons. Additionally, expression of PTL-1 in touch neurons alone was unable to rescue the shortened lifespan observed in ptl-1 mutants, but pan-neuronal re-expression restored wild-type longevity, indicating that, at least for a specific group of mechanosensory neurons, premature neuronal ageing and organismal ageing can be decoupled.

The Caenorhabditis elegans model system has been the focus of many investigations into the processes that contribute towards ageing and longevity (reviewed in1). Surprisingly, initial studies did not detect major age-related changes in neuronal viability or gross axon integrity in the nervous system of C. elegans2, but close examination of neuronal morphology has revealed ageing phenotypes such as abnormal branching and blebbing along neuron processes3–5. In addition to structural changes, it was recently shown using patch-clamp recordings that C. elegans motor neurons demonstrate age-associated declines in activity6. Using the nematode model, we can dissect the relationship between neuronal and whole organism ageing, for example by examining whether accelerated neuronal ageing negatively affects the entire organism.

It was recently demonstrated that C. elegans undergoes progressive, age-dependent changes in the structural integrity of neurons4–5. Several factors that contribute to the maintenance of neuronal structural integrity have been identified, including HSF-11 and [NK-1], which have been demonstrated to act in a cell-autonomous manner. DAF-16, the FOXO transcription factor involved in insulin-like signalling6, also plays a role in neuronal aging. In the case of DAF-16, some data suggest that it functions cell autonomously7, while other evidence suggests a non-cell autonomous role in this ageing process. We have previously shown that the neuronal microtubule-associated protein (MAP) protein with tau-like repeats (PTL-1) is involved in the regulation of both organismal and nervous system ageing in C. elegans8. PTL-1 is the sole C. elegans homolog of members of the mammalian MAPT(tau)/MAP2/MAP4 family9,10. Mutations in the MAPT locus cause neuronal disorders including Frontotemporal dementia (reviewed in11). Furthermore, tau is the main component of neurofibrillary tangles found in the brains of individuals suffering from Alzheimer’s disease or several other neurodegenerative conditions collectively known as tauopathies12–15. Due to this association with age-associated neurodegenerative disease in humans, it was striking to find that PTL-1 is involved in the maintenance of structural integrity in the C. elegans nervous system.

Our previous data indicated that the neurons of ptl-1(ok621) null mutants prematurely develop markers of neuronal ageing in touch receptor neurons (TRNs) and ventral nerve cord GABAergic neurons7. In our current investigation, we re-expressed PTL-1 in the ptl-1(ok621) null mutant specifically in all neurons to generate a pan-neuronal transgenic line, or in TRNs alone to generate a TRN-specific transgenic line. The TRNs consist of six...
mechanosensory neurons with specialised microtubule structures important for the response to gentle touch16,17, and incidentally are also the neurons in which PTL-1 is most highly expressed10.19. We examined both lifespan and neuronal ageing in these transgenic lines and found that (i) PTL-1 functions through neurons to regulate ageing, (ii) PTL-1 regulates neuronal ageing in a cell-autonomous manner, (iii) knockdown of ptl-1 by RNAi specifically in the TRNs affected TRN ageing but not ageing of another neuronal subset, and (iv) the processes that regulate organismal ageing and tissue-specific ageing are separable.

Results
The short-lived phenotype of ptl-1 null mutants is rescued by re-expression in all neurons but not in touch neurons alone. Previously, we demonstrated that re-expressing PTL-1 under the regulation of its endogenous promoter is sufficient to rescue both neuronal ageing and lifespan phenotypes observed in ptl-1 mutant animals1. PTL-1 expression is not restricted to neurons10.18; it is also expressed in non-neuronal tissues including vulval cells and stomatotelial muscle16. By using a transgenic line expressing a PTL-1::GFP translational fusion protein, we confirmed this expression pattern (Figure 1a). Because we had already shown that PTL-1 expressed under the control of the ptl-1 promoter can rescue both neuronal and whole organism ageing, we aimed to test if re-expression specifically in neurons, or in the subset of TRNs (where PTL-1 is most highly expressed) would also rescue these phenotypes.

We generated transgenic lines expressing ptl-1::V5 cDNA under the regulation of either the aex-3 pan-neuronal promoter or the mec-7 TRN-specific promoter. For clarity, we describe transgenic lines as either “Pan-neuronal Tg” or “TRN Tg” depending on the transgene, followed by “ptl-1(ok621)” if it is in the ptl-1 null mutant background. We immunostained for the V5 epitope tag present at the C-terminus of PTL-1 and confirmed that PTL-1 displayed broad neuronal expression (Figure 1bi) or expression in TRNs only (Figure 1ci) in these transgenic lines, respectively. We then performed lifespan assays to determine if either of these transgenes would rescue the shortened lifespan of ptl-1(ok621) null mutants. We found that expressing PTL-1 in all neurons in “Pan-neuronal Tg” and “Pan-neuronal Tg: ptl-1(ok621)” populations resulted in a wild-type lifespan (Figure 1bii; Supplementary Figure S1a). In contrast, expressing PTL-1 in TRNs alone in the “TRN Tg; ptl-1(ok621)” strain did not rescue the lifespan phenotype of the ptl-1(ok621) null mutant, as the survival curve of this transgenic line was not significantly different from that of the ptl-1(ok621) control (log-rank test p-value = 0.25) (Figure 1ci; Supplementary Figure S1b). We also observed a small detrimental effect when PTL-1 was over-expressed in TRNs because the “TRN Tg” line without the ptl-1(ok621) mutation had significantly shorter lifespan compared with wild-type controls according to the log-rank statistical test (p-value = 0.01) (Figure 1cii). This difference was not significant using the Wilcoxon test (p-value = 0.15), which places more weight on earlier deaths when comparing survival curves (GraphPad Prism 6).

Premature ageing of touch neurons in ptl-1(ok621) mutants can be rescued by pan-neuronal and touch neuron-specific re-expression of PTL-1. We visualised cell body branching and axon blebbing in anterior TRNs on days 1, 5, 7 and 9 of adulthood using the Punc-47::gfp(oxIs12) reporter. As noted previously10.47, GABAergic neurons in young animals do not show branching in the dorsally-projecting commissures, which are often branched in older adults (Figure 3a). We found that re-expressing PTL-1 in all neurons in the ptl-1 null mutant background completely rescued premature neuronal ageing observed in ptl-1(ok621) mutant animals (Figure 3b). For example, at day 5 of adulthood, 10% of wild-type animals displayed branching along the commissures of GABAergic neurons, compared with 36% of ptl-1(ok621) animals and 15% of “Pan-neuronal Tg; ptl-1(ok621)” animals (Figure 3b). In contrast, we did not observe a substantial difference in the incidence of abnormal structures between “TRN Tg; ptl-1(ok621)” animals (53%) at day 7 of adulthood compared with ptl-1(ok621) mutants at the same time point (55%), which are both significantly higher than wild-type (24%) (p-value = 0.0003 or 0.0002) (Figure 3c). This demonstrates that regulation of neuronal ageing by PTL-1 occurs cell-autonomously within the neuron. Additionally, over-expression of PTL-1 in TRNs alone does not negatively affect ageing in the GABAergic neurons, as the incidence of branching in “TRN Tg” animals at this time point was 25%, and therefore not significantly different from wild-type (p-value = 0.85). As for the neuronal ageing experiments in TRNs, we observed the same phenotypes at 25°C as at 20°C (Supplementary Figure S2a; bii).

Premature ageing of GABAergic neurons in ptl-1(ok621) mutants can be rescued by pan-neuronal but not touch neuron-specific re-expression of PTL-1. To complement the above experiment, we investigated if PTL-1 expression specifically in one neuronal subset, the TRNs, is able to regulate neuronal ageing in other neurons, such as the GABAergic neurons. We visualised GABAergic neuron branching on days 1, 5, 7 and 9 of adulthood using the Punc-47::gfp(oxIs12) reporter. As noted previously10.47, the GABAergic neurons in young animals do not show branching in the dorsally-projecting commissures, which are often branched in older adults (Figure 3a). We found that re-expressing PTL-1 in all neurons in the ptl-1 null mutant background completely rescued premature neuronal ageing observed in ptl-1(ok621) mutant animals (Figure 3b). For example, at day 5 of adulthood, 10% of wild-type animals displayed branching along the commissures of GABAergic neurons, compared with 36% of ptl-1(ok621) animals and 15% of “Pan-neuronal Tg; ptl-1(ok621)” animals (Figure 3b). In contrast, we did not observe a substantial difference in the incidence of abnormal structures between “TRN Tg; ptl-1(ok621)” animals (53%) at day 7 of adulthood compared with ptl-1(ok621) mutants at the same time point (55%), which are both significantly higher than wild-type (24%) (p-value = 0.0003 or 0.0002) (Figure 3c). This demonstrates that regulation of neuronal ageing by PTL-1 occurs cell-autonomously within the neuron. Additionally, over-expression of PTL-1 in TRNs alone does not negatively affect ageing in the GABAergic neurons, as the incidence of branching in “TRN Tg” animals at this time point was 25%, and therefore not significantly different from wild-type (p-value = 0.85). As for the neuronal ageing experiments in TRNs, we observed the same phenotypes at 25°C as at 20°C (Supplementary Figure S2a; bii).

Knockdown of PTL-1 in touch neurons only has a cell-autonomous effect on neuronal ageing. In C. elegans, SID-1 is a transmembrane protein that allows the passive uptake of dsRNA and is required for systemic RNAi knockdown. Wild-type neurons are refractory to RNAi as they do not express SID-119.20; however, Calixto et al. showed that neuronal re-expression of SID-1 in a sid-1 null mutant, as such in TRNs, allowed RNAi knockdown to occur in these neurons21. The authors demonstrated that feeding mec-4 RNAi bacteria to these animals resulted in an expected loss of touch sensitivity, indicating that SID-1 expression in TRNs efficiently sensitised these neurons to knockdown by RNAi treatment22. We therefore complemented the assays involving tissue-specific re-expression of PTL-1 by reducing PTL-1 levels in one set of neurons. We investigated the effect of PTL-1 knockdown in the neuronal subset of TRNs on the age-associated loss of structural integrity. To this end, we first tested whether PTL-1 in TRNs could be effectively knocked down using this system. We fed empty vector (EV) and ptl-1 RNAi bacteria to animals that express both SID-1 only in the TRNs and a PTL-1::GFP fusion protein. These animals showed
Figure 1 | Pan-neuronal but not TRN-specific re-expression of PTL-1 rescues the short-lived phenotype of the ptl-1(ok621) null mutant. The presence of the ptl-1(ok621) mutation in the genetic background of each transgenic line is indicated by the addition of "ptl-1(ok621)" in the strain name. (a) PTL-1 expression as shown by a translational GFP fusion is enriched in neurons and is also present in non-neuronal tissues. i–iii) Micrographs from a single transgenic animal indicating particular anatomical features where PTL-1::GFP is expressed including the nerve ring in the head, cells in the vulva and seam cells. iv–vii) Micrographs from other transgenic animals indicating PTL-1::GFP expression in muscle, seam cells, vulval cells and neurons. Anatomical features are highlighted by white arrows. (b) i. Staining for PTL-1::V5 in pan-neuronal re-expression line. The left panel shows the head, with the white arrow indicating neurons of the nerve ring and the pink outline indicates the approximate position of the pharynx. The right panel shows the tail, with the white arrowheads indicating tail neurons. ii. Lifespan assay conducted for pan-neuronal transgenic worms. (c) i. Staining for PTL-1::V5 in TRN-specific re-expression lines. The left panel indicates the tail, with PLM neurons indicated by white arrowheads. The right panel indicates the mid-body of the worm, showing an ALM neuron. The white arrowhead here indicates the cell body. ii. Lifespan assay for TRN-specific transgenic worms. Survival curves for control wild-type and ptl-1(ok621) animals in both graphs were obtained in the same experiment. n = 120 at day 0. Results of statistical analysis are indicated by p-values underneath each graph. Details of p-values are shown in Supplementary Table S1. Lifespan experiments were conducted twice independently, and the representative data shown are from one experiment. Survival curves obtained from the second independent lifespan experiment are shown in Supplementary Figure S1.
Figure 2 | Pan-neuronal and TRN-specific re-expression of PTL-1 rescues the neuronal ageing phenotype in TRNs that is observed in the \textit{ptl-1(ok621)} null mutant. The TRNs were visualised using the \textit{Pmec-4:gfp} (zdIs5) reporter. The presence of the \textit{ptl-1(ok621)} mutation in the genetic background of each transgenic line is indicated by the addition of “\textit{ptl-1(ok621)}” in the strain name. (a) Representative images of a young TRN, and TRNs showing cell body branching (arrow) or axon blebbing. (b) Neuron imaging assay conducted for pan-neuronal transgenic worms (“Pan-neuronal Tg”) showing (i) cell body branching and (ii) axon blebbing. (c) Neuron imaging assay conducted for TRN-specific transgenic worms (“TRN Tg”). Data for wild-type and \textit{ptl-1(ok621)} animals in both graphs were obtained in the same experiment. \( n \) for each time-point is indicated under the graphs. The chi-squared statistical test was used to determine statistical significance. P-value is indicated by ns = not significant, * \(< 0.05\). Details of p-values are shown in Supplementary Table S1. Experiments were conducted twice independently, and the representative data shown are from one experiment.
a loss of GFP signal in the TRNs in the 
ptl-1 RNAi treated cohort but not in those fed EV control RNAi bacteria (Supplementary Figure S3), demonstrating the effectiveness of the 
ptl-1 RNAi treatment.

We next crossed animals carrying a TRN GFP reporter with those that express SID-1 only in the TRNs. These animals that are henceforth referred to as the “TRN SID-1” strain carry a sid-1(qt2) mutation and a TRN-specific SID-1 rescue transgene (parent strain TU340321). TRN SID-1 animals were fed empty vector (EV) and 
ptl-1 RNAi feeding clones for two generations and were imaged on days 1, 3, 5, and 7 of adulthood to observe neuronal ageing. As an additional control, we carried out an experiment using animals containing only the TRN GFP reporter that are wild-type at the sid-1 locus and do not have the TRN SID-1 transgene. Here, we would expect that no RNAi knockdown would occur within neurons in these animals as the neurons do not express SID-1. As expected, these animals did not show a substantial difference in the incidence of abnormal neuronal structures between RNAi treatments (Figures 4a, 4b). In contrast, when observing the anterior TRNs in TRN SID-1 transgenic animals, we saw that worms fed with 
ptl-1 RNAi bacteria had a higher incidence of cell body branching and axon blebbing compared with those fed EV bacteria. Specifically, 70% of TRN SID-1 animals fed 
ptl-1 RNAi bacteria displayed cell body branching at day 7 compared with 47% in the EV RNAi control, and 53% of 
ptl-1 RNAi-fed animals displayed axon blebbing compared with 30% of the control population (Figure 4ai). In addition, we observed that TRN SID-1 animals displayed a relatively high

![Image](https://example.com/image.png)
Figure 4 | Knockdown of PTL-1 in touch neurons has a cell autonomous effect on neuronal ageing. Strains labelled as zdIs5 or oxIs12 indicate the allele name of the gfp reporter, are wild-type at the SID-1 locus and do not contain a SID-1 transgene. SID-1 transgenic worms are labelled as “sid-1(qt2);TRN SID-1” to indicate the presence of the sid-1 mutation and Pmec-18::sid-1 transgene. The RNAi treatment is either empty vector (EV) or ptl-1 and is indicated after the strain name. (a) i. TRN imaging assay for animals carrying the zdIs5 reporter, indicating data for cell body branching, axon blebbing and axon branching. (a) ii. Micrograph shows an example of the axon branching phenotype. (b) GABAergic imaging assay for animals carrying the oxIs12 reporter. n for each sample is indicated in the table. Statistical analysis: chi-squared test, p-value is indicated by ns (not significant), *, p<0.05. Details of p-values are shown in Supplementary Table S1. Experiments were conducted twice independently, and the representative data shown are from one experiment.
frequency of axon branching (Figure 4aii), which is normally only present in late stage (day 10) adult animals. TRN SID-1 animals fed with both control and ptl-1 RNAi treatments showed considerably higher levels of axon branching in the anterior TRNs compared with wild-type controls, although transgenic animals fed ptl-1 bacteria displayed a higher incidence of this phenotype compared with EV treatments (Figure 4ai). This could be due to the expression of multiple TRN-specific transgenes (both Pmec-18::sid-1 and Pmec-4::gfp) that may alter the development or ageing process of the neuron.

We further tested the effect of the loss of PT1-1 in TRNs by examining neuronal ageing in a different set of neurons. For this, we generated a strain expressing a GABAergic neuron GFP reporter in a TRN SID-1 background. As SID-1 is only expressed in the TRNs, we would not expect PT1-1 levels in the GABAergic neurons to be affected by feeding RNAi bacteria in this strain. Therefore, although PT1-1 levels in the TRNs of TRN SID-1 animals should decrease with ptl-1 feeding RNAi, PT1-1 expression in the GABAergic neurons of these animals should remain at endogenous levels. As above, we also performed a control RNAi experiment using animals containing only the GABAergic neuron GFP reporter and not the TRN SID-1 transgene. When GABAergic neuron ageing in TRN SID-1 transgenic worms was monitored, we observed no difference between ptl-1 RNAi treatment and EV treatment (Figure 4b). This demonstrates that loss of PT1-1 in the TRNs appears not to affect neuronal ageing in the GABAergic neurons.

Taken together, these data from tissue-specific re-expression and RNAi knockdown experiments indicate that the effect of ptl-1 on neuronal ageing is cell autonomous.

Discussion

Given the clinical relevance of tau pathology in neurodegenerative disease, we aimed to investigate the function of the tau-like protein PT1-1 in the context of neuronal ageing. The key findings of our study in C. elegans are: (i) the tau/MAP2/MAP4 homologue PT1-1 functions through neurons to regulate organisational ageing, (ii) PT1-1 regulates neuronal ageing in a cell-autonomous manner, and (iii) the processes that regulate organisational ageing and tissue-specific ageing can be separable.

Previous research has highlighted the importance of neurons, and sensory neurons in particular, in the regulation of lifespan. Examples include the role of olfactory neurons and ciliated sensory neurons in inhibiting longevity, and that of thermosensory neurons in regulating lifespan. Examples of processes that regulate organismal ageing and tissue-specific ageing include the role of olfactory neurons and ciliated sensory neurons in particular, in the regulation of lifespan. Examples include the role of olfactory neurons and ciliated sensory neurons in inhibiting longevity, and that of thermosensory neurons in regulating lifespan. Examples of processes that regulate organismal ageing and tissue-specific ageing include the role of olfactory neurons and ciliated sensory neurons in inhibiting longevity, and that of thermosensory neurons in regulating lifespan. Examples of processes that regulate organismal ageing and tissue-specific ageing

Our previous work has demonstrated that ptl-1(ok621) null mutants display both premature organisational and premature neuronal ageing. We have now shown that animals that express PT1-1 in TRNs alone have a shortened lifespan compared with wild-type as have ptl-1(ok621) null mutants, but that their TRNs age at the same rate as wild-type animals (Figure 1Bii, C). This demonstrates that PT1-1-mediated processes that influence lifespan or neuronal ageing can be decoupled from each other. A similar segregation has been observed for components of the insulin-like/IGF-1 signalling pathway. Long-liveddaf-2/insulin receptor mutants show less neuronal branching in early adulthood compared with wild-type, and this effect is ameliorated in daf-2;daf-16/FOXO transcript factor double mutants, which have a wild-type lifespan. However, RNAi-mediated knockdown of daf-16 in non-neuronal tissues in a daf-2 mutant results in an animal with wild-type lifespan, but has no effect on the daf-2-mediated delay of neuronal ageing.

Therefore, although neuronal cell body branching and axon blebbing increase with temporal age, tissue-specific ageing and whole organism ageing appear to be separable. This suggests that the mechanisms that define these ageing processes contain both distinct and overlapping processes. It is of considerable interest in the fields of ageing and neurodegeneration to identify the genetic players involved in these processes.

We have shown that PT1-1 in C. elegans functions in neurons to regulate both whole organism ageing and neuronal ageing. Although several lines of evidence indicate that PT1-1 is expressed mainly in the nervous system until now it remained to be determined whether PT1-1 was indeed acting through neurons to mediate these processes. We have proven this by demonstrating that a transgenic line expressing PT1-1 only in neurons is able to rescue both premature lifespan and neuronal ageing in a ptl-1 mutant. We have now also shown that the regulation of neuronal ageing by PT1-1 is cell autonomous. The role of PT1-1 in regulating these ageing processes as a neuronal MAP is particularly striking due to its homology with mammalian tau, which plays a critical role in neurodegenerative disease. Understanding the physiological roles of a tau-like protein in C. elegans may provide important information on human neurodegeneration and ageing.

Methods

Strain information. C. elegans strains were cultured on nematode growth media (NGM) agar plates seeded with the Escherichia coli strain OP50. Hermaphrodite animals were used for all experiments. The wild-type strain used for all experiments is N2 (Bristol), Strains N2, Rb809 ptl-1(ok621), T2U34 c3is4251 [Pmyo-3::gfp(NE)(L):lacZ] [Pmyo-3::gfp] + Pmyo-2::gfp[SA4] + dpy-20 [+] [plus71] (pCF90[Pmyo-2::mCherry] + Pmech-18::sid-1::GFP [pQ2]). C3H175::Pmec-4::gfp + lin-15 [+] + 1 and EG1285 ox1212[Punc-47::gfp + lin-15 [+] X] were obtained from the Caenorhabditis Genetics Centre (CGC), and FX00543 ptl-1(tm543) was obtained from the National Bioresource Project, Japan (Dr S. Mitani). ptl-1 mutant lines Rb809 and FX00543 were both outcrossed six times to wild-type and renamed APD004 and APD015, respectively. For neuron imaging assays, strains involved were crossed with CX10175 to visualise TRNs and EG1285 to visualise GABAergic motor neurons. For tissue-specific RNAi strains, sid-1(q2) mutants were selected by screening for insensitivity (lack of paralysis) to knockdown of unc-22 and the missense mutation confirmed by sequencing.

List of strains generated. APD050: zdIs5; sid-1(q2); uIs71. APD066: oxIs12; sid-1(q2); uIs71. APD070: apdIs10[Pmec-7::PTL1-1::3′UTR; Pmyo-2::mCherry; Pprl-28::PuroR::rpl-16::outron; Neoxy::let-858::3′UTR], outcrossed six times to wild-type. APD074: apdIs9; ox121; sid-1(q2). APD075: apdIs9; zdIs5; ptl-1(ok621). APD076: apdIs9; sid-1(q2); uIs71. APD081: apdIs9; ox121. APD082: apdIs9; zdIs5. APD096: apdIs10[Pac-3::PTL1-1::3′UTR; Pmyo-2::mCherry; Pprl-28::PuroR::rpl-16::outron; Neoxy::let-858::3′UTR], outcrossed six times to wild-type. APD097: apdIs10; zdIs5. APD098: apdIs10; zdIs5; ptl-1(ok621). APD099: apdIs10; ox121. APD100: apdIs10; ox121; sid-1(q2). APD101: apdIs10; ptl-1(ok621). APD130: apdIs10[ptl-1::PTL1-1::GFP::puro-54::3′UTR; rpl-28::PuroR::rpl-16::outron; Neoxy::let-858::3′UTR; sid-1(q2); uIs71.

Generation of transgenic lines. The ptl-1 promoter, PT1-1:GFP tagged at the C-terminus with VS, and the PT1-1 3′UTR was amplified as previously described.
Neuronal imaging experiments. Age-matched animals synchronised by egg-laying were cultured on plates at 20 °C or 25 °C. At the start of the experiment, animals were one day-old adults in all cases. To separate adult worms from their progeny, adult worms were moved to new NGM plates every second day until the assay was completed. Assays were conducted blind to the genotype of the worms. To score the incidence of aberrant neuronal structures in TRNs, worms were scored as positive if at least one of the observed commissures displayed branching. The proportion of animals scored as positive was expressed then as a percentage of the sample size observed at that time point.

Lifespan assays. Age-matched animals synchronised by egg-laying were cultured on plates at 25 °C and the number of surviving animals recorded every day until death. Two independent lifespan experiments were conducted, each with 120 one-day-old adults per strain plated at the start of each assay. Animals that were lost or displayed internal hatching or bursting were censored. Pan-neuronal and TRN-specific Tg lines were assayed together with the same cohort of wild-type and knockdown animals expressing branching or blebbing at the cell body or axon. For scoring, all TRN-BAC neurons, worms were scored as positive if at least one of the observed commissures displayed branching. The proportion of animals scored as positive was expressed then as a percentage of the sample size observed at that time point.

Survival curves were generated using GraphPad Prism 6.

RNA interference experiments. Experiments were conducted at 20 °C. sid-1(q2) animals expressing SID-1 in TRNs only (APD050 and APD066) were synchronised by hyphochlorite bleaching and larval stage 1 worms plated onto carbenicillin plates seeded with RNAi bacteria (HT115 E. coli) that either contained the pL4400 empty vector (EV), or pL4400 containing cloned pti-1 or unc-22 DNA generated by the Ahringer laboratory. Strains that are wild-type at the sid-1 locus were used as negative controls. unc-22 knockdown was used as a control for the experiment; knockdown of unc-22 in muscle results in paralysis, so animals that are wild-type for sid-1 will be paralysed. Bacterial cultures were induced to express dsRNA using 3 mM isopropyl β-D-thiogalactopyranoside (IPTG). After two generations fed on RNAi bacteria, animals were imaged at day 1, 3, 5 and 7 of adulthood to visualise TRNs or GABAergic neurons. To maintain knockdown, animals were transferred to new RNAi plates every second day until the assay was completed. Assays were conducted blind to the genotype of the worms.

Immunofluorescence. One-day-old adults synchronised by egg-laying and cultured at 23 °C were used for all experiments. Animals were permeabilised using a freeze-crack protocol as previously described. Samples were immediately fixed in ice-cold 4% paraformaldehyde at 4 °C for ≥12 hours. The primary antibody used was mouse monoclonal anti-V5 (R900–25, Life Technologies) [1:200]. The secondary antibody used was goat anti-mouse Alexa Fluor-568 (Sigma) [1:500]. All antibody dilutions were made in 30% (v/v) normal goat serum (Life Technologies). Samples were mounted on microscope slides using VectaShield mounting medium (Vector Labs). Samples were imaged using a BX51 Microscope (Olympus). Micrographs were captured using AnalySIS software (Olympus).

Statistical analysis. The t-level is 0.05 for all analyses. Survival curves were analysed in GraphPad Prism 6 (GraphPad Software Inc.) to provide two-tailed p-values. Tests used were the log-rank test, which gives equal weight to deaths at all time points, and the Wilcoxon test, which gives more weight to deaths at early time points. The log-rank test is commonly used for a.c. lifespan data but we also report p-values from the Wilcoxon test to accommodate the possibility that hazard ratios were not consistent through the assay. For neuron imaging experiments, we grouped data into categories of “displays branching/blebbing” or “does not display branching/blebbing” and used Pearson’s chi-squared test in Microsoft Excel (Microsoft) to determine if the incidence of branching/blebbing in a mutant population at a single time point is different from the ‘expected value’ of the wild-type strain at the same time point. This analysis provides one-tailed p-values that were used to determine if the incidence of abnormal neuronal structures in a mutant strain at a time point was different from the wild-type strain.

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23. Ahringer laboratory. Strains that are wild-type at the sid-1 locus were used as negative controls. unc-22 knockdown was used as a control for the experiment; knockdown of unc-22 in muscle results in paralysis, so animals that are wild-type for sid-1 will be paralysed. Bacterial cultures were induced to express dsRNA using 3 mM isopropyl β-D-thiogalactopyranoside (IPTG). After two generations fed on RNAi bacteria, animals were imaged at day 1, 3, 5 and 7 of adulthood to visualise TRNs or GABAergic neurons. To maintain knockdown, animals were transferred to new RNAi plates on days 2, 4 and 6 of the experiment. Scoring was conducted in the same manner as for other neuronal imaging assays. Imaging was conducted blind to the identity of the RNAi clones. To test knockdown of pti-1 in TRNs by feeding RNAi, PTL-1::GFP animals were treated as described above by feeding either EV or pti-1 RNAi bacteria and imaged after two generations of RNAi treatment.

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**Author contributions**
Y.L.C., X.F., J.G. and H.R.N. designed experiments. Y.L.C. and X.F. performed experiments. Y.L.C., X.F., J.G. and H.R.N. analysed data. Y.L.C. wrote the main manuscript text and prepared figures. Y.L.C., X.F., J.G. and H.R.N. reviewed the manuscript.

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