

Drosophila Trap1 protects against mitochondrial dysfunction in a PINK1/parkin model of Parkinson's disease

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Mitochondrial dysfunction caused by protein aggregation has been shown to have an important role in neurological diseases, such as Parkinson's disease (PD). Mitochondria have evolved at least two levels of defence mechanisms that ensure their integrity and the viability of their host cell. First, molecular quality control, through the upregulation of mitochondrial chaperones and proteases, guarantees the clearance of damaged proteins. Second, organellar quality control ensures the clearance of defective mitochondria through their selective autophagy. Studies in *Drosophila* have highlighted mitochondrial dysfunction linked with the loss of the PTEN-induced putative kinase 1 (PINK1) as a mechanism of PD pathogenesis. The mitochondrial chaperone TNF receptor-associated protein 1 (TRAP1) was recently reported to be a cellular substrate for the PINK1 kinase. Here, we characterise *Drosophila* Trap1 null mutants and describe the genetic analysis of Trap1 function with Pink1 and parkin. We show that loss of Trap1 results in a decrease in mitochondrial function and increased sensitivity to stress, and that its upregulation in neurons of Pink1 mutant rescues mitochondrial impairment. Additionally, the expression of Trap1 was able to partially rescue mitochondrial impairment in parkin mutant flies; and conversely, expression of parkin rescued mitochondrial impairment in Trap1 mutants. We conclude that Trap1 works downstream of Pink1 and in parallel with parkin in *Drosophila*, and that enhancing its function may ameliorate mitochondrial dysfunction and rescue neurodegeneration in PD.

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Accumulating evidence suggests that the disruption of mitochondrial function and dynamics contributes to age-related neurodegenerative diseases, such as Parkinson's disease (PD).¹ Accordingly, cells have developed complex quality control mechanisms to cope with the different challenges constantly imposed on the integrity of mitochondria.^{2,3}

Molecular quality control, the first level of the mitochondrial defence mechanism, involves the upregulation of nuclear genes that encode mitochondrial chaperones and proteases, ensuring the removal of misfolded and non-assembled polypeptides. This stress response is known as the mitochondrial unfolded-protein response (UPR^{mt}).⁴

Furthermore, the highly dynamic nature of mitochondria facilitates quality control through fusion and fission.^{5,6} Thus, damaged mitochondria can fuse with healthy organelles to replenish stores of deficient components. When the damage is excessive, such processes are overwhelmed; mitochondria become depolarised, and their components are targeted for recycling through a specific form of autophagy, termed mitophagy.^{7,8}

Recent findings have demonstrated that PTEN-induced putative kinase 1 (PINK1) is capable of directly modulating the activity of chaperones.^{9,10} The TNF receptor-associated protein 1 (TRAP1) is a mitochondrial chaperone suggested

to act as a cellular substrate of the PINK1 kinase. In cultured cells, PINK1 expression shows a protective effect against oxidative stress-induced apoptosis. Upon the induction of oxidative stress, TRAP1 is phosphorylated in a PINK1-dependent manner, and this phosphorylation is necessary to confer resistance to apoptosis.¹⁰

TRAP1 is a member of the HSP90 family, which has been shown to protect cells from oxidative stress and apoptosis caused by different agents,^{11–14} and its upregulation in tumours has been associated with chemotherapy resistance.^{15,16}

The fruit fly, *Drosophila melanogaster*, has recently emerged as a powerful model system for studying the links between mitochondrial dysfunction and parkinsonian neurodegeneration.¹⁷ In *Drosophila*, the loss of Pink1 leads to mitochondrial dysfunction associated with muscle degeneration and the loss of dopaminergic neurons.^{18,19} In addition, a decreased expression of Trap1 was found to enhance the loss of dopaminergic neurons in flies expressing a mutant form of human α -synuclein that causes PD.²⁰

Here, we describe the consequences of Trap1 loss-of-function in *Drosophila* and its gain-of-function on mitochondrial integrity. We also genetically address the role of Trap1 in mitochondrial quality control through the Pink1/parkin pathway.

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Results

Loss of *Trap1* increases sensitivity to stress and causes a decline in mitochondrial function. To investigate the *in vivo* role of *Trap1* in *Drosophila*, we first determined the consequences of its loss. We obtained *Trap1*⁴ mutant flies, generated by an imprecise excision of the P-element insertion line P(EPgy2)*Trap1*^{EY10238}. The excision deleted the majority of the *Trap1* gene (Figure 1a). We next performed quantitative real-time RT-PCR and confirmed that *Trap1*⁴ mutants show a total loss of *Trap1* mRNA, and that its neighbouring genes were not affected by the imprecise excision of the P-element (Figure 1b). The *Trap1* mutant flies were viable and developed to adulthood; however, they

presented a significantly shorter lifespan compared with the controls (Figure 1c). Given the proposed role of TRAP1 as a mitochondrial chaperone, we decided to investigate the consequences of exposing *Trap1*⁴ mutants to increased levels of stress. We observed a decreased viability of *Trap1*⁴ mutants subjected to heat stress (Figure 1d). The *Trap1*⁴ mutants were also more sensitive to paraquat, a pesticide linked to PD by epidemiological studies²¹ (Figure 1e), and to mitochondrial poisons such as rotenone (Figure 1f) and antimycin (Figure 1g). The *Trap1*⁴ mutants showed an age-dependent impaired climbing ability, suggesting a locomotor deficit (Figure 1h). In *Pink1* mutant flies, locomotor defects result from mitochondrial impairment in the *Drosophila* skeletal muscles, including the indirect flight muscle.^{18,19}

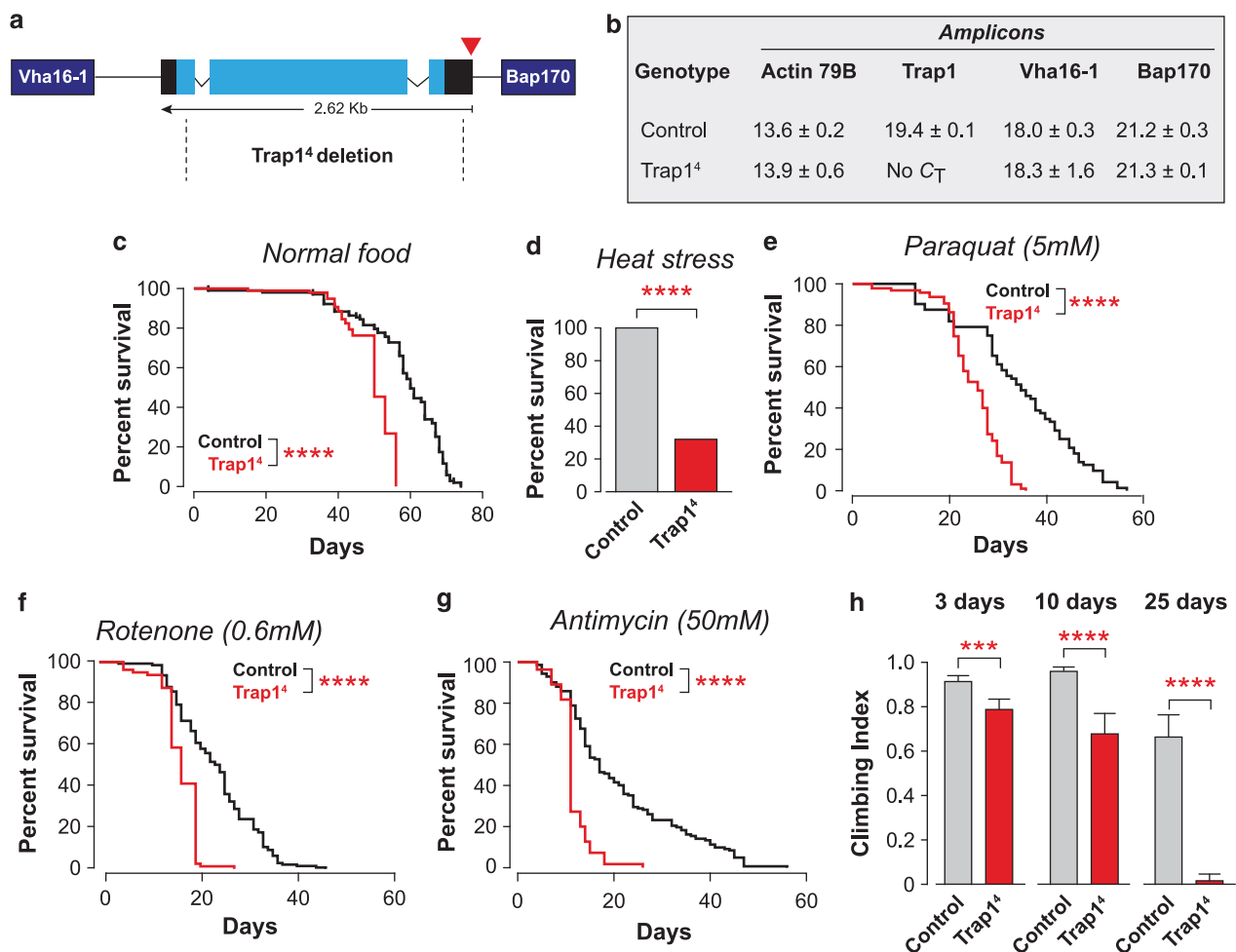


Figure 1 The loss of *Trap1* in *Drosophila* causes motor impairment and an increased sensitivity to stress. (a) Genomic map of *Trap1* (cytological location 42B2). Black, untranslated regions; light blue, exons. The P-element insertion (EY10238) is indicated by the red triangle. The neighbouring genes (*Vha16-1* and *Bap170*) are indicated in dark blue. *Trap1*⁴ deletion, delimited by the dashed lines, removes most of the *Trap1* gene. (b) Analysis of the expression levels of *Trap1* and its neighbouring genes. Expression levels were measured by real-time PCR in 3-day-old flies with the indicated genotypes (mean Ct ± S.D., *n* = 4 per genotype). Expression of actin was used as a control. No Ct for the *Trap1* transcript was detected in *Trap1*⁴ mutants. (c) *Trap1*⁴ mutants (red) have a reduced lifespan compared with the controls (black). Fly viability was scored over a period of 75 days, using a minimum of 100 flies per genotype. The statistical significance is indicated (log-rank, Mantel-Cox test). (d–g) *Trap1* mutant flies show enhanced sensitivity to stress. (d) Flies were subjected to heat stress, and viability was assessed after 24 h. A total of 100 males (4 days old) were assayed for each genotype. The asterisks indicate significant values (Fisher's exact test, two-sided, $\alpha < 0.05$). (e–g) Flies were maintained on food supplemented with the indicated drugs, and the viability was scored over a period of 60 days, using a minimum of 100 flies per genotype. The statistical significance is indicated by the asterisks (log-rank, Mantel-Cox test). (h) *Trap1*⁴ mutants show a decrease in motor performance. Flies with the indicated genotypes and ages were tested using a standard climbing assay (mean ± S.D., *n* ≥ 80 flies for each genotype). The asterisks indicate significant values (two-way ANOVA with the Bonferroni multiple comparison test)

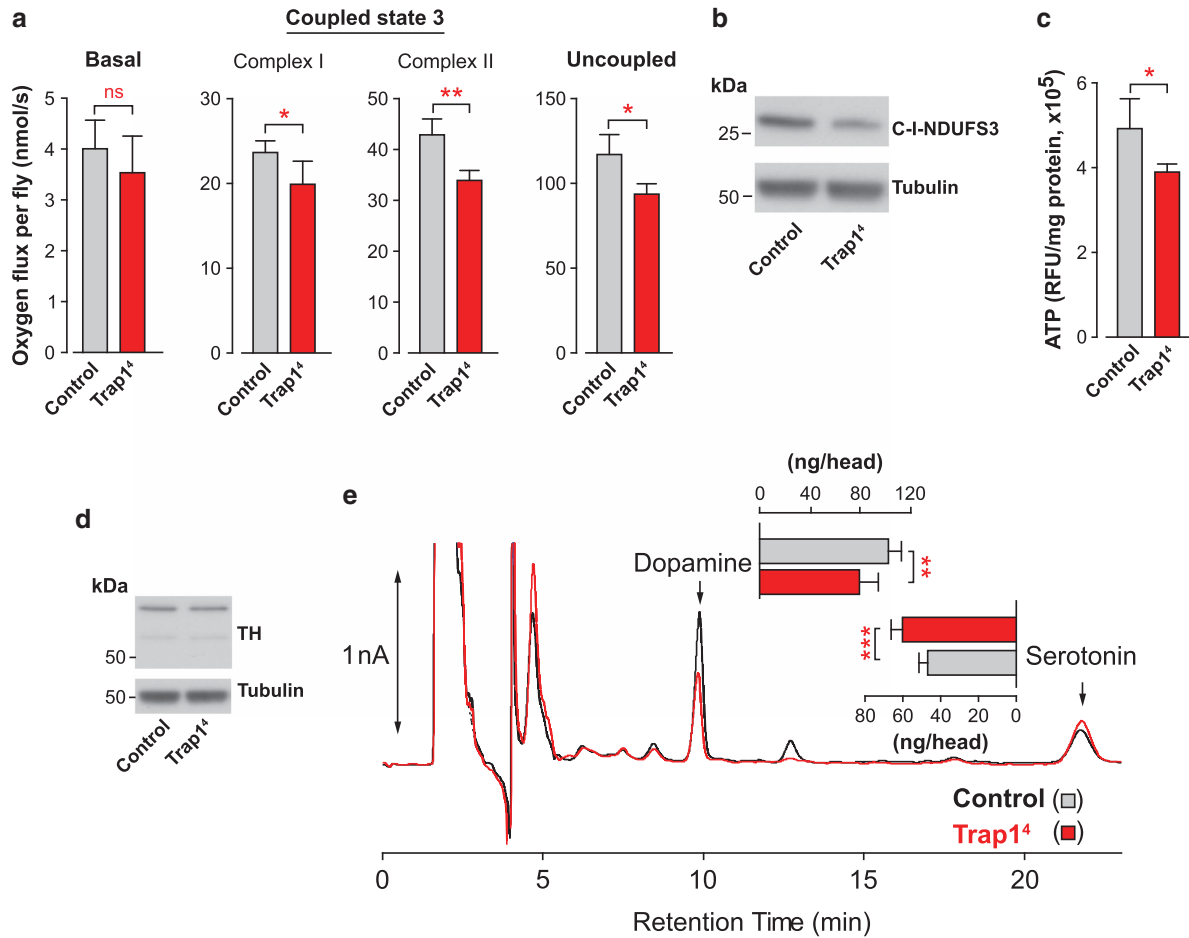


Figure 2 The loss of *Trap1* in *Drosophila* results in a reduction of mitochondrial function and brain dopamine levels. (a) Decreased respiration in *Trap1*⁴ mutant flies. Activity was measured by high-resolution respirometry in 20-day-old flies. Data are shown as the means \pm S.D. ($n \geq 4$ in each genotype). Significant values relative to the control are indicated by asterisks (two-tailed unpaired *t* test). (b) Decreased complex I protein levels in *Trap1*⁴ mutant flies. Whole-fly (20 days old) protein lysates were subjected to western blot analysis with the indicated antibodies. (c) *Trap1*⁴ mutants have lower levels of ATP compared with the controls. The ATP levels were measured using a bioluminescent assay. Data are shown as the means \pm S.D. ($n = 4$ for each genotype). The statistical significance is indicated by asterisks (two-tailed unpaired *t*-test). (d) Analysis of TH levels in *Trap1* mutant flies. Fly-head protein lysates from 20-day-old flies were used for western blotting analysis with the indicated antibodies. (e) *Trap1* mutant flies have decreased dopamine and increased serotonin levels. Neurotransmitter levels were assessed by HPLC with electrochemical detection. Data are shown as the means \pm S.D. ($n \geq 5$ for each genotype). The statistical significance is indicated by asterisks (two-tailed unpaired *t*-test)

To determine if the loss of *Trap1* lead to mitochondrial impairment, we compared the respiration rates in the controls and mutants. This analysis revealed a significant decrease in the respiratory function of the *Trap1* mutants (Figure 2a). *Trap1*⁴ mutants also showed a decrease in the levels of the mitochondrial complex I (Figure 2b). Mitochondria are responsible for the production of the majority of cellular energy in the form of ATP. The measurement of ATP levels in *Trap1*⁴ mutants revealed a significant decrease when compared with controls (Figure 2c). Taken together, these results suggest that the loss of the mitochondrial chaperone *Trap1* results in a decrease in mitochondrial function, which is associated with a loss of ATP in adult flies. The loss of dopaminergic neurons can be indirectly assessed through the analysis of the expression levels of tyrosine hydroxylase (TH), an enzyme expressed in dopaminergic neurons.²² We failed to detect any differences in the TH levels of *Trap1*⁴ mutants (Figure 2d); however measuring neurotransmitter levels in the heads of *Trap1*⁴

mutants revealed a significant decrease in the dopamine content, compared to the controls (Figure 2e).

Trap1 expression enhances motor performance and protects against oxidative stress.

Recently, TRAP1 was suggested to work downstream of PINK1 in the prevention of mitochondrial dysfunction associated with PD pathogenesis.¹⁰ We therefore decided to test the epistatic relationship between *Trap1* and *Pink1* in flies. We generated transgenic flies expressing full-length *Trap1*, and using UAS-GAL4 system²³ we confirmed the expression of the transgene by western blot analysis (Figure 3a). To investigate its intracellular localisation, we crossed UAS-*Trap1* flies with a transgenic line expressing mitochondria-targeted green-fluorescent protein (mitoGFP). This analysis revealed that *Trap1* localises to the mitochondria (Figure 3b). To further investigate the phenotypic consequences of *Trap1* expression, we compared the locomotor performance of *Trap1* flies with that of controls using a climbing

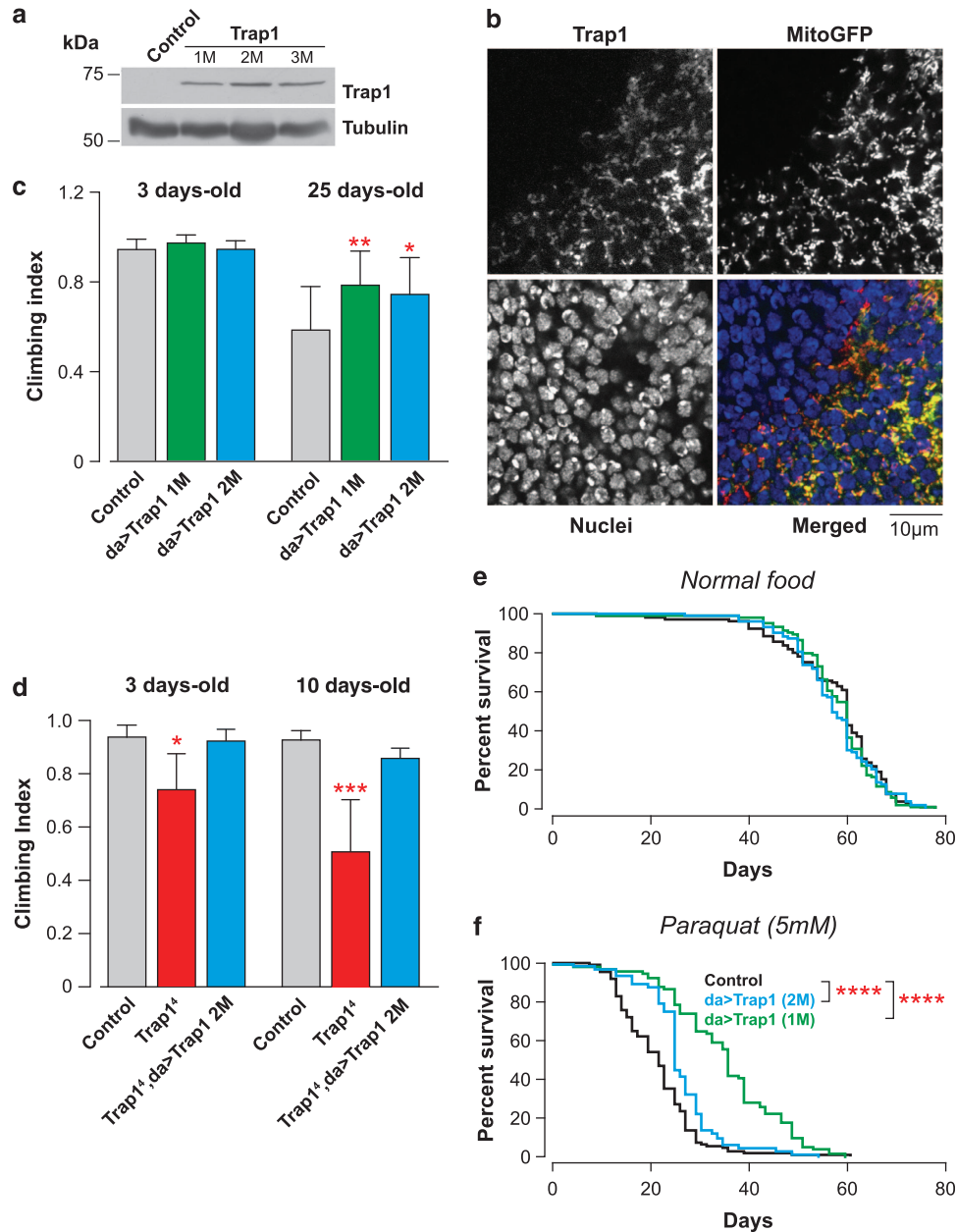


Figure 3 Expression of *Drosophila* Trap1 confers resistance to stress. (a) Analysis of the Trap1 expression levels in transgenic flies. Whole-fly lysates were analysed by western blotting with the indicated antibodies. (b) Trap1 localises to the mitochondria. Confocal analysis of the posterior compartment of larval wing discs coexpressing Trap1 and a mitoGFP under the control of the *enGAL4* driver. In the merged colour image, red corresponds to Trap1, blue to nuclei and green to mitoGFP. (c) The expression of Trap1 results in increased motor performance. Flies with the indicated genotypes and ages were tested using a standard climbing assay (mean \pm S.D., $n \geq 120$ flies for each genotype). The asterisks indicate significant values relative to the control (two-way ANOVA with the Bonferroni multiple comparison test). (d) The expression of Trap1 reversed the decrease in motor performance in *Trap1⁴* mutants. Flies with the indicated genotypes and ages were tested using a standard climbing assay (mean \pm S.D., $n \geq 80$ flies for each genotype). The asterisks indicate significant values relative to the control (two-way ANOVA with the Bonferroni multiple comparison test). (e) Trap1 expression does not affect the total lifespan. A total of 100 flies per genotype were maintained on normal food for a period of 80 days. (f) Trap1-expressing flies (green and blue) show enhanced resistance to paraquat toxicity compared with the controls (black). Fly viability was scored over a period of 60 days using a minimum of 100 flies per genotype. The statistical significance is indicated by the asterisks (log-rank, Mantel–Cox test)

assay. This analysis revealed that Trap1 expression leads to a significantly enhanced climbing performance in aged flies (Figure 3c) and suppressed the climbing defects of *Trap1⁴* mutants (Figure 3d). We previously showed that treating flies with paraquat leads to the upregulation of mitochondrial chaperones as a means to protect mitochondria from the oxidative stress triggered by this pesticide.²⁴ Additionally,

TRAP1 phosphorylation was reported to be enhanced in response to oxidative stress.¹⁰ Flies expressing Trap1 show a normal lifespan compared with the controls (Figure 3e). However, Trap1 expression significantly increased the lifespan of paraquat-treated flies (Figure 3f), indicating that the expression of this mitochondrial chaperone protects against oxidative stress *in vivo*.

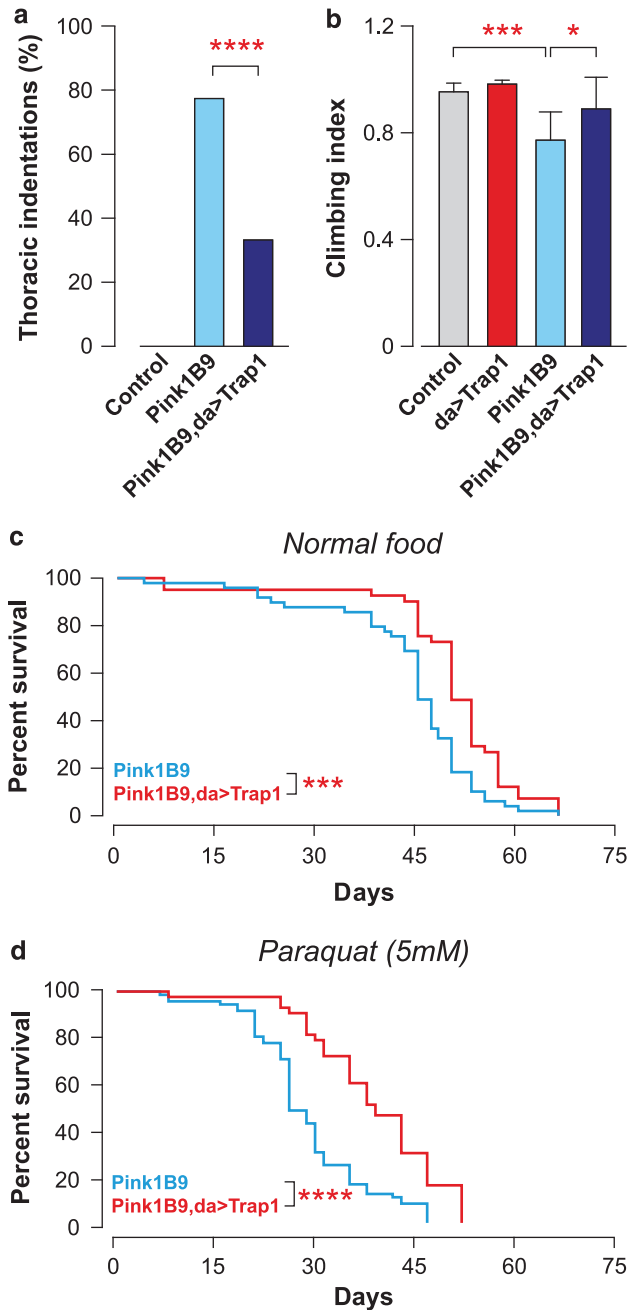


Figure 4 *Trap1* gain-of-function rescues *Pink1* mutant flies. (a) Expression of *Trap1* rescues the thoracic defects of *Pink1*^{B9} mutants. Thoracic indentations were counted up to 24 h after eclosion ($n > 900$ for each genotype). The asterisks indicate significance (χ^2 , two-sided, $\alpha < 0.05$). (b) *Trap1* expression suppresses motor impairment in the *Pink1*^{B9} mutants. 10-day-old flies with the indicated genotypes were tested using a standard climbing assay (mean \pm S.D., $n = 120$ flies per genotype). The statistical significance relative to *Pink1*^{B9} is indicated by asterisks (one-way ANOVA with Dunnett's multiple comparison test). (c and d) *Trap1* expression enhances the lifespan of *Pink1*^{B9} mutants. A minimum of 80 flies per genotype were maintained on either normal (c) or paraquat-containing food (d). Fly viability was scored over a period of 70 days. The statistical significance is indicated by the asterisks (log-rank, Mantel-Cox test)

Expression of *Trap1* suppresses mitochondrial dysfunction in *Pink1* mutants. The expression of TRAP1 in cultured cells was shown to be required for PINK1-dependen-

dent inhibition of oxidative stress-induced cytochrome *c* release and cell death.¹⁰ The mitochondrial dysfunction in *Pink1* mutants is associated with muscle pathology, affecting the indirect flight muscles, leading to the appearance of thoracic indentations and impaired motor performance.^{18,19} The expression of *Trap1* partially rescued the degree of thoracic indentations (Figure 4a) and the impaired motor performance in the *Pink1*^{B9} mutants (Figure 4b). Mitochondrial defects present in the *Pink1* mutants are also associated with a decreased lifespan.¹⁸ We noted that the expression of *Trap1* increased the lifespan of the *Pink1* mutants (Figure 4c) and decreased their sensitivity to paraquat (Figure 4d). Taken together, these observations suggest that the ubiquitous expression of *Trap1* protects against the phenotypic consequences of *Pink1* mutations, and that *Trap1* acts genetically downstream of *Pink1*. The mitochondrial dysfunction in the *Pink1* mutant flies is reflected by a loss of mitochondrial proteins and decreased respiration. By expressing *Trap1* in *Pink1*^{B9} mutants, we observed a recovery of the mitochondrial protein content, indirectly assessed through the analysis of the levels of a complex I subunit (Figure 5a); a partial recovery of the respiration rates (Figure 5b) and ATP levels (Figure 5c). Taken together, these observations indicate that *Trap1* expression results in a recovery of the mitochondrial function in *Pink1* mutants.

Neuronal expression of *Trap1* is sufficient to revert the phenotype of *Pink1* mutants. The genetic removal of the function of *Drosophila Pink1* results in apoptotic muscle degeneration and a decrease in the brain dopamine levels as a consequence of the degeneration of dopaminergic neurons.¹⁹

The loss of mitochondria at neuronal synapses causes defects in neurotransmission of the *Drosophila* neuromuscular junction (NMJ).²⁵ Given that one of the most important features of *Pink1* mutant flies is the degeneration of both the indirect flight muscle and dopaminergic neurons, we next tested whether the degeneration of the indirect flight muscle involves a presynaptic component, and is therefore preceded by neuronal loss.

We detected a decrease in the TH levels in *Pink1*^{B9} mutants that was reversed upon the expression of *Trap1* using a pan-neuronal *elav*-GAL4 driver (Figure 6a). This result indicates that the neuronal expression of *Trap1* can suppress neurodegeneration in *Pink1* mutant flies. Next, we evaluated the consequence of the neuronal expression of *Trap1* on phenotypes associated with muscle degeneration. The neuronal expression of *Trap1* was sufficient to suppress muscle degeneration in *Pink1*^{B9} mutants, which was reflected in a decrease in the degree of thoracic indentations (Figure 6b) and an improved climbing performance (Figure 6c). Finally, the neuronal expression of *Trap1* significantly reversed the respiration deficit present in *Pink1* mutant flies (Figure 6d). Taken together, these results indicate that the neuronal rescue of mitochondrial function in *Pink1* mutants by *Trap1* is sufficient to suppress muscle degeneration and motor impairment.

Genetic epistasis analysis of *parkin* and *Trap1*. Upon mitochondrial damage, PINK1 has been shown to recruit

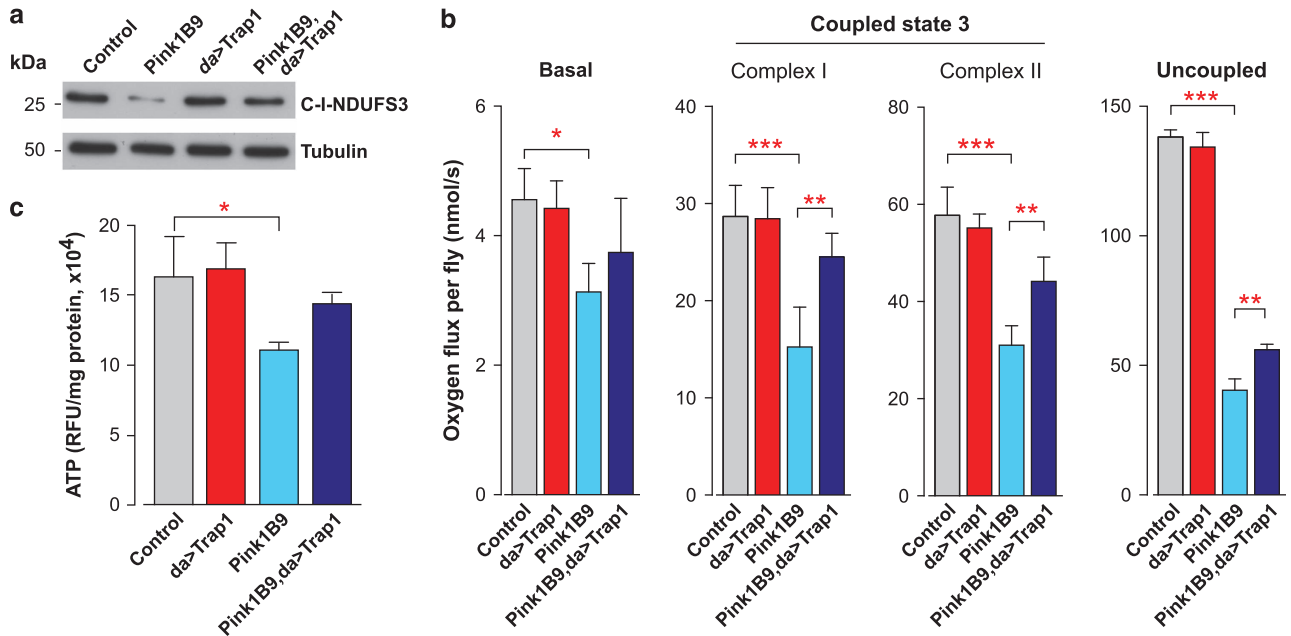


Figure 5 *Drosophila* Trap1 reverses mitochondrial dysfunction in *Pink1* mutants. (a) Trap1 expression restores the mitochondrial complex I protein levels in the *Pink1^{B9}* mutant flies. Whole-fly protein lysates were used for western blotting analysis with the indicated antibodies. (b) Trap1 expression partially rescues the mitochondrial respiratory defects in the *Pink1^{B9}* mutants. Measurements of state three respiration of complex I and complex II, and complex II uncoupled respiration by high-resolution respirometry. The data are shown as the means \pm S.D. ($n = 6$ for each genotype). The statistical significance is indicated by the asterisks (one-way ANOVA with Dunnett's multiple comparison test). (c) Expression of Trap1 increased the ATP levels in the *Pink1^{B9}* mutants. ATP levels were measured using a bioluminescent assay. Data are shown as the means \pm S.D. ($n = 4$ for each genotype). The statistical significance is indicated by the asterisks (one-way ANOVA with Dunnett's multiple comparison test relative to the control)

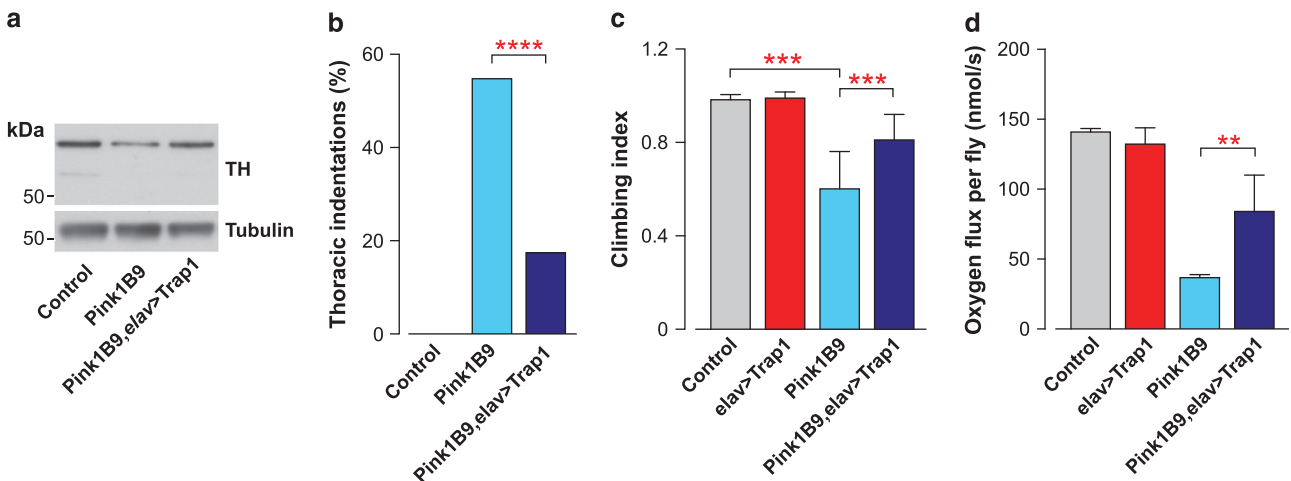


Figure 6 Targeted neuronal expression of Trap1 rescues mitochondrial dysfunction in *Pink1* mutants. (a) Decreased TH levels observed in the *Pink1^{B9}* mutants can be partially restored by the expression of Trap1 in neurons. Fly-head protein lysates from 20-day-old flies were used for western blotting analysis with the indicated antibodies. (b) Neuronal expression of Trap1 is capable of diminishing the degree of thoracic indentations in the *Pink1^{B9}* mutants. Thoracic indentations were counted up to 24 h after eclosion ($n > 300$ for each genotype). The asterisks indicate the significance relative to *Pink1^{B9}* (χ^2 , two-sided, $\alpha < 0.05$). (c) Targeted neuronal expression of Trap1 improves the climbing performance of *Pink1^{B9}* mutants. 10-day-old flies with the indicated genotypes were tested using a standard climbing assay (mean \pm S.D., $n = 120$ flies per genotype). The statistical significance relative to *Pink1^{B9}* is indicated by the asterisks (one-way ANOVA with Dunnett's multiple comparison test). (d) Expression of Trap1 specifically in neurons is sufficient to improve mitochondrial respiration in the *Pink1^{B9}* mutants. Complex II-uncoupled respiration was measured by high-resolution respirometry. Data are shown as the means \pm S.D. ($n = 6$ for each genotype). The statistical significance is indicated by the asterisks (one-way ANOVA with Dunnett's multiple comparison test relative to the *Pink1^{B9}* mutant)

Parkin, a cytosolic E3 ubiquitin ligase, to damaged mitochondria.^{26,27} Parkin recruitment to damaged mitochondria has been proposed to cause the ubiquitination of the profusion

factor Mfn to label terminally damaged mitochondria for autophagic degradation,²⁸ a process that has been defined as organellar quality control.^{3,29} The importance of Parkin as

a downstream effector of PINK1 was confirmed in a series of genetic studies in *Drosophila*, which showed that the expression of *parkin* could rescue the phenotype of *Pink1* mutant flies.^{18,19} To address the epistatic relationship between *parkin* and *Trap1*, we first tested whether *Trap1* expression could also rescue the *parkin* mutant phenotype. The expression of *Trap1* partially rescued the degree of thoracic indentations (Figure 7a) but not the impaired motor performance in the *parkin* mutants (Figure 7b). We also observed that the expression of *Trap1* had a modest effect on the suppression of the ATP loss present in *parkin* mutants (Figure 7c) and caused a recovery in the mitochondrial protein content, indirectly assessed through the analysis of the levels of a complex I subunit (Figure 7d). Moreover, *Trap1* expression in *parkin* mutants led to an increased level of survival in flies maintained on normal food (Figure 7e) and on paraquat-containing food (Figure 7f). We next tested whether *parkin* expression could also rescue the *Trap1* mutant phenotype. When we expressed *parkin* in *Trap1*⁴ mutant flies, we observed a significant suppression of their climbing defects (Figure 7g), as well as a marked recovery of the levels of mitochondrial complex I (Figure 7h). Taken together, these results show that the enhanced expression of *Trap1* can partially compensate for the loss of *parkin*, and reciprocally, the enhanced expression of *parkin* can also ameliorate some of the *Trap1* mutant-associated phenotypes. This suggests that both *Trap1* and *parkin* act in a parallel pathway that modulates mitochondrial competence.

Discussion

We have characterised the *Drosophila* orthologue of the TRAP1 chaperone and found that, similar to its mammalian counterpart, it localises to the mitochondria. Using a loss-of-function approach, through the generation of *Trap1* mutant flies that lack *Trap1* expression, we determined that its loss compromises mitochondrial function. Mutations in either *Pink1* or *parkin* influence mitochondrial dynamics and function, causing an impairment of fission, which affects male fertility by compromising the normal development of mitochondrial derivatives (also known as nebenkern).^{18,19,30,31} During spermatogenesis, *Drosophila* mitochondria undergo significant morphological changes that require fusion to form a large spherical nebenkern and subsequent fission when the nebenkern unfurls to yield two mitochondrial derivatives.³² The *Trap1* mutants developed normally to adulthood and were both viable and fertile, suggesting that their mitochondrial defects do not compromise the mitochondrial fission and fusion processes required for normal spermatogenesis.

Although *Trap1* mutants show no defects in development, they had a decreased lifespan, progressive locomotor defects and a decrease in the brain dopamine levels. Taken together, these results suggest that this mitochondrial chaperone is important to minimise the age-dependent decline of mitochondrial function in post-mitotic cells, such as muscle cells and neurons, which rely heavily on mitochondrial function. The milder phenotype of *Trap1* mutants compared with the *Pink1* mutants suggest that other effectors of molecular quality control acting downstream of *Pink1* can compensate for the potential defects caused by the loss of *Trap1*. In this

respect, the serine protease HtrA2, a downstream target of PINK1^{9,33} involved in mitochondrial molecular quality control,³⁴ could potentially partially compensate for any disturbances in protein folding in the intermembrane space of mitochondria.

The expression of *Trap1* was sufficient to enhance the climbing performance of aged flies and promote resistance to oxidative stress (Figure 3). Mitochondria in skeletal muscles are highly specialised towards energy production for contractile activity,³⁵ and therefore, the expression of this chaperone might contribute to the maintenance of mitochondrial performance in tissues with high-energy demand, such as skeletal muscles, thereby enhancing the motor performance.

Our results indicate that the neuronal expression of *Trap1* in *Pink1* mutants is sufficient to suppress muscle degeneration. Mitochondria have been shown to be concentrated in both the presynaptic nerve terminal and the postsynaptic endplates in muscle,³⁶ indicating that a decline in mitochondrial bioenergetic function may affect synaptic transmission and result in a loss of NMJ innervation. Our data suggest that mitochondrial dysfunction at the presynaptic terminal of the NMJ is a major determinant of the pathologies of the skeletal muscle present in the *Pink1* mutants, and that the compromise of the motor neuron function in the *Pink1* mutants causes muscle dysfunction; therefore, is likely to precede muscle degeneration.

In *Drosophila*, the expression of *parkin* can rescue the phenotype of *Pink1* mutants, presumably by enhancing the organellar quality control through autophagic disposal of defective mitochondria.²⁹ Our data suggest that *Trap1* expression can partially compensate for the mitochondrial defects present in *parkin* mutant flies. Given that *Trap1* is likely to act as a molecular chaperone in the clearance of misfolded proteins, its expression may lead to enhanced molecular quality control in mitochondria, and therefore decrease the demand for organellar quality control as a clearance mechanism for mitochondria overwhelmed by excessive protein misfolding. Enhanced expression of *Trap1* may in turn decrease the demand on organellar quality control in *parkin* mutants, leading to a decrease in the degree of dysfunctional mitochondria in these insects. We also observed that the expression of *parkin* was capable of suppressing some of the phenotypic consequences associated with *Trap1* mutations. This suggests that both the molecular and organellar pathways for mitochondrial quality control work in parallel, and the enhancement of either one of these can relieve pressure on the other and contribute to a better mitochondrial health.

Materials and Methods

Antibodies. The following primary antibodies were used in this study: anti-TRAP1 mouse IgG (clone 30) (612342), BD Transduction Laboratories, Franklin Lakes, NJ, USA; anti-NDUFS3 mouse monoclonal (ab14711), Mitosciences, Eugene, OR, USA; anti-TH rabbit polyclonal (AB152), Millipore (Billerica, MA, USA); and anti- α -tubulin IgG1 clone B-5-1-2, Sigma-Aldrich (St Louis, MO, USA).

Genetics and *Drosophila* strains. Transgenic strains with UAS-*Trap1* were generated by P-element-mediated transformation. Transgenic flies were generated at BestGene Inc. (Chino Hills, CA, USA). Three independent transgenic lines were obtained, and all were tested for protein expression. *Pink1*^{B9}, *park*²⁵, UAS *parkin*^{62B} and *da*-GAL4 flies were a gift from A Withworth (MRC, Centre for

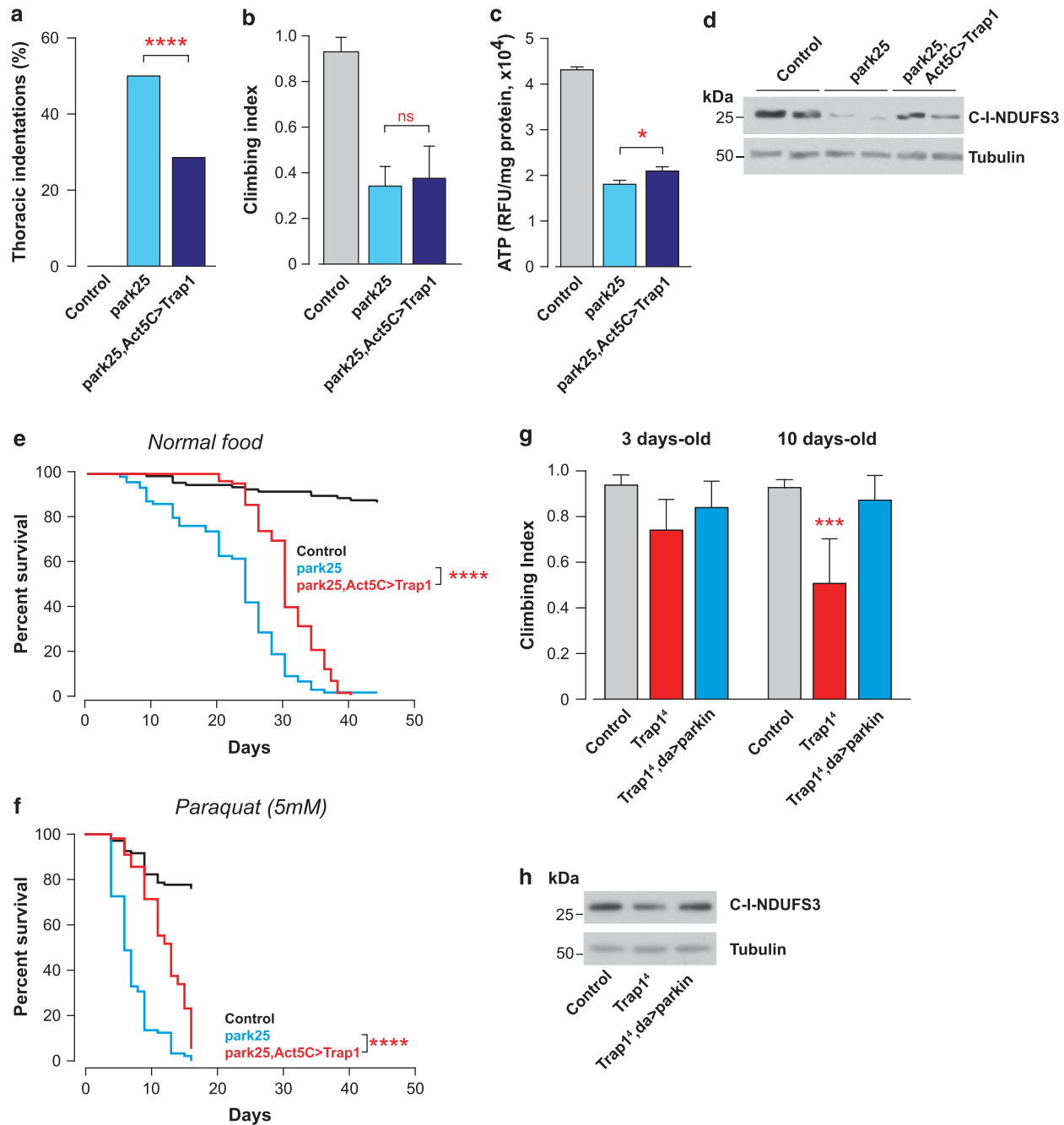


Figure 7 Epistatic relationship between *Trap1* and *parkin*. **(a)** *Trap1* expression decreases the degree of thoracic indentations in *parkin* mutant flies, *park*²⁵. Thoracic indentations were counted up to 24 h after eclosion ($n > 250$ for each genotype). The asterisks indicate the significance (χ^2 , two-sided, $\alpha < 0.05$). **(b)** The expression of *Trap1* fails to rescue the climbing defect of the *park*²⁵ mutants. 3-day-old flies with the indicated genotypes were tested using a standard climbing assay (mean \pm S.D., $n > 120$ flies per genotype). **(c)** *Trap1* expression increases the levels of ATP in the *park*²⁵ mutants. The ATP levels were measured using a bioluminescent assay. Data are shown as the means \pm S.D. ($n = 4$ for each genotype). The statistical significance is indicated by the asterisks (one-way ANOVA with Dunnett's multiple comparison test relative to *park*²⁵). **(d)** *Trap1* expression restores the mitochondrial complex I protein levels in the *park*²⁵ mutant flies. Whole-fly protein lysates were used for western blotting analysis with the indicated antibodies. **(e)** *Trap1* expression enhances the lifespan of the *park*²⁵ mutants. Fly viability was scored over a period of 40 days using a minimum of 100 flies per genotype. The statistical significance is indicated by the asterisks (log-rank, Mantel-Cox test). **(f)** Expression of *Trap1* increases the resistance of *park*²⁵ mutants to paraquat-induced stress. Fly viability was scored over a period of 15 days using a minimum of 80 flies per genotype. The statistical significance is indicated by the asterisks (log-rank, Mantel-Cox test). **(g)** The expression of *parkin* rescues the climbing defects of *Trap1*⁺ mutants. Flies with the indicated genotypes and ages were tested using a standard climbing assay (mean \pm S.D., $n > 60$ flies per genotype). The asterisks indicate significant values (two-way ANOVA with the Bonferroni multiple comparison test). **(h)** *parkin* expression restores the mitochondrial complex I protein levels in *Trap1*⁺ mutant flies. Whole-fly protein lysates from 20-day-old flies were used for western blotting analysis with the indicated antibodies

Developmental and Biomedical Genetics, University of Sheffield, Sheffield, UK). *Trap1^d* line was kindly provided by Dr. JE Treisman (New York University School of Medicine, NY, USA). *w1118*, *Act5C-GAL4*, *enGAL4* and *elav-GAL4* lines were obtained from Bloomington Stock Centre (Indiana University, Bloomington, IN, USA). Fly stocks and crosses were maintained on standard cornmeal agar media at 25 °C. The genotypes of the flies used in each individual figure are listed in Supplementary Table S1.

Molecular biology. Expression constructs encoding *Drosophila Trap1* sequence were generated by polymerase chain reaction using appropriate adaptor primers. The PCR fragments were cloned into the GAL4-responsive pUAST expression vector for the generation of transgenic fly strains.

HPLC measurement of dopamine and serotonin. For sample preparation, 15 male fly heads (20 days old) were dissected under mild CO₂ anaesthesia, and quickly frozen in dry ice. The heads were then homogenised in 50 µl chilled 0.1M perchloric acid using a motorised, hand-held tissue homogeniser. The chilled homogenates of fly heads were filtered through a low-binding Durapore (0.22 µm) PVDF membrane using Ultrafree-MC centrifugal devices. The dopamine and serotonin measurements were performed immediately after sample preparation. Supernatant fluid (30 µl) was eluted at a flow rate of 50 µl/min through a 150 × 1.0 mm C18 column. The mobile phase contained 50 mM phosphoric acid, 8 mM NaCl, 0.1 mM EDTA, 12.5% (v/v) methanol, 500 mg/l octanesulfonic acid (OSA) (pH 6.0). Analysis was performed using an Alexys LC-EC system equipped with a DECADE II electrochemical detector.

Immunofluorescence and confocal microscopy. Third instar larvae were dissected, fixed in 4% paraformaldehyde, stained with the anti-TRAP1 antibody and incubated with Hoechst 33342. Wing imaginal discs were mounted on glass slides with Vectashield (Vector Laboratories, Burlingame, CA, USA) and imaged on a Zeiss LSM510 confocal microscope.

Statistical analysis. The data are presented as mean values, and the error bars indicate ± S.D. Inferential statistical analysis was performed using the Prism and StatMate software packages (www.graphpad.com). The significance level is indicated as **** for $P < 0.0001$, *** for $P < 0.001$, ** for $P < 0.01$, * for $P < 0.05$ and ns for $P > 0.05$.

ATP assays. Five male flies (10 or 20 days old) were homogenised in 100 µl of 6 M guanidine-HCl in extraction buffer (100 mM Tris and 4 mM EDTA, pH 7.8) to inhibit ATPases.

Homogenised samples were subjected to rapid freezing in liquid nitrogen, followed by boiling for 5 min. Samples were cleared by centrifugation, and supernatant was diluted (1/50) with extraction buffer and mixed with a luminescent solution (CellTiter-Glo Luminescent Cell Viability Assay, Promega, Fitchburg, WI, USA). Luminescence was measured on an Infinite M200Pro multifunction reader (TECAN, Männedorf, Switzerland). The relative ATP levels were calculated by dividing the luminescence by the total protein concentration, which was determined by the Bradford method.

Climbing assays. The climbing assays were performed as previously described,³⁷ using a countercurrent apparatus developed initially for phototaxis experiments. A total of 20 male flies were placed into the first chamber, taped to the bottom and then given 20 s to climb a distance of 10 cm. Those flies that successfully climbed 10 cm or beyond in 20 s were then shifted to a new chamber, and both sets of flies were given another opportunity to climb the 10-cm distance. This procedure was repeated a total of five times. After five trials, the number of flies in each chamber was counted. At least 120 flies were used for each genotype tested.

Respirometry analysis. Mitochondrial respiration was assayed at 37 °C by high-resolution respirometry using an OROBOROS Oxygraph. DatLab software package (OROBOROS, Innsbruck, Austria) was used for the data acquisition (2-s-time intervals) and analysis, including the calculation of the time derivative of the oxygen concentration, signal deconvolution dependent on the response time of the oxygen sensor and correction for instrumental background oxygen flux. Respiration was assayed by homogenising two flies using a pestle in MiRO5

respiration buffer (20 mM HEPES, 10 mM KH₂PO₄, 110 mM sucrose, 20 mM taurine, 60 mM K-lactobionate, 0.5 mM EGTA, 3 mM MgCl₂, 1 g/l fatty acid-free BSA). The basal respiration rates were assayed in the absence of exogenous respiration substrates. For coupled (state 3) assays, complex I activity was assayed in MiRO5 respiration buffer in the presence of 2 mM malate, 10 mM glutamate and 5 mM ADP. Complex II was assayed in respiration buffer supplemented with 1 mM rotenone, 10 mM succinate and 5 mM ADP. ADP was not added for the measurement of respiration in uncoupled mitochondria obtained by one freeze (liquid nitrogen)–thaw cycle of the fly tissue.

Lifespan analysis. Groups of 10 newly eclosed males of each genotype were placed into separate vials with food and maintained at 25 °C. The flies were transferred to vials containing fresh food every 2–3 days, and the number of dead flies was recorded. The data are presented as Kaplan–Meier survival distributions, and the significance was determined by log-rank tests.

Drug treatments. All the drugs used in this study were incorporated into the fly food. The concentrations of each of the drugs in the food were the following: 5 mM for paraquat, 600 µM for rotenone and 50 mM for antimycin A. The newly eclosed males were placed with normal food in groups of 10 for 1 day prior to their transfer to vials containing drug. The flies were transferred to new vials every 2–3 days, and the number of dead flies was recorded. The assay was completed after all the flies died. A minimum of 80 flies per genotype were analysed. The results are presented as Kaplan–Meier survival distributions, and the significance was determined by log-rank tests.

Preparation of *Drosophila* protein lysates and western blot analysis. Single adult flies (3, 10 or 20 days old) were homogenised using a motorised pestle in lysis buffer (0.1% Triton-X100, 10 mM EDTA, 1 mM DTT, 100 mM KCl, 20 mM HEPES (pH 7.5), 5% glycerol, 1 µg/ml chymostatin, 1 µg/ml leupeptin, 1 µg/ml pepstatin A and 1 µg/ml antipain). The lysates were subsequently cleared by centrifugation. The proteins were separated on 12 or 10% SDS-PAGE gels and transferred to Immobilon PVDF membranes (Millipore). The membranes were blocked in 5% milk and then incubated with the indicated primary antibody prior to incubation with the appropriate HRP-conjugated secondary antibody. The antibody complexes were visualised by enhanced chemiluminescence (ECL). The results are representative of at least three biological replicates per genotype.

RNA extraction and quantitative real-time RT-PCR. The isolation of total RNA from three adult flies per sample was performed using the RNeasy Mini Kit (Qiagen, Hilden, Germany). Quantitative real-time RT-PCR was performed on an M × 4000 (Stratagene, La Jolla, CA, USA) real-time cycler using the QuantiTect SYBR Green RT-PCR system (Qiagen). Primers used (QuantiTect Primer Assay, Qiagen) were Cat. No. QT00967393 (*actin 79B*, used as a housekeeping gene), Cat. No. QT00940520 (*Trap1*), Cat. No. QT00501130, (*Vha16-1*) and Cat. No. QT00940527, (*Bap170*). The quantification was performed using the comparative Ct method.³⁸

Heat stress. For heat sensitivity, 4-day-old male flies were divided into groups of 15 and placed in empty vials (no food or water). The flies were maintained at 25 °C for 30 min, after which the thermo stress was applied by placing the vials in a water bath at 37 °C for 1 h. This step was followed by a 30-min recovery at 25 °C and a new heat stress for 1 h at 37 °C. The flies were then transferred to vials containing fresh food and maintained at 25 °C. The number of living flies was recorded 24 h later. A total of 100 flies per genotype were tested for thermo stress.

Conflict of Interest

The authors declare no conflict of interest.

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