Enhancing autophagy by redox regulation extends lifespan in *Drosophila*

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Redox signalling is an important modulator of diverse biological pathways and processes, and operates through specific post-translational modification of redox-sensitive thiols on cysteine residues¹⁻⁴. Critically, redox signalling is distinct from irreversible oxidative damage and functions as a reversible ‘redox switch’ to regulate target proteins. H₂O₂ acts as the major effector of redox signalling, both directly and through intracellular thiol redox relays^{5,6}. Dysregulation of redox homeostasis has long been implicated in the pathophysiology of many age-related diseases, as well as in the ageing process itself, however the underlying mechanisms remain largely unclear^{7,8}. To study redox signalling by H₂O₂ *in vivo* and explore its involvement in metabolic health and longevity, we used the fruit fly *Drosophila* as a model organism, with its tractable lifespan and strong evolutionary conservation with mammals⁹. Here we report that inducing an endogenous redox-shift, by manipulating levels of the H₂O₂-degrading enzyme catalase, improves health and robustly extends lifespan in flies, independently of oxidative stress resistance and dietary restriction. We find that the catalase redox-shifted flies are acutely sensitive to starvation stress, which relies on autophagy as a vital survival mechanism. Importantly, we show that autophagy is essential for the lifespan extension of the catalase flies. Furthermore, using redox-inactive knock-in mutants of Atg4a, a major effector of autophagy, we show that the lifespan extension in response to catalase requires a key redox-regulatory cysteine residue, Cys102 in Atg4a. These findings demonstrate that redox regulation of autophagy can extend lifespan, confirming the importance of redox signalling in ageing and as a potential pro-longevity target.

To explore the role of endogenous redox signalling *in vivo*, we used the UAS/GAL4 expression system to up-regulate catalase in wild-type (WT) flies. Global up-regulation of catalase under control of the *daughterless* promoter (da-GAL4>UAS-cat) extended the median and maximum lifespan of female flies (typically by ~10-15%; Fig. 1a). Importantly, for these experiments we used the *white Dahomey* (*w^{Dah}*) background, which is a long-lived and outbred WT, hence we are extending healthy lifespan and not rescuing a short-lived defect. Catalase was over-expressed ~5-fold at the mRNA level in whole flies (Fig. S1a). Lifespan was not extended in males (Fig. 1a), despite similar catalase over-expression (Fig. S1a), however interventions modulating nutrient-sensing in *Drosophila* often show gender-specific effects on survival in females, that are absent or marginal in males¹⁰.

The da-GAL4>UAS-cat flies were mildly delayed in eclosing, without affecting the proportion of larvae surviving to adulthood (Fig. S1b). To exclude developmental effects, we showed that the lifespan extension could be fully recapitulated using the inducible GeneSwitch system, with over-expression from d2 of adulthood onwards (da-GS>UAS-cat ±RU; Figs. 1b, S1a).

Varying the dose of the inducer drug RU (50-400 μ M) still did not extend lifespan in males and had only marginal effects on lifespan extension in females (Fig. S1c), suggesting that the catalase benefits are at maximum potential. Over-expressing catalase under the control of an alternative ubiquitous driver (actin5c-GAL4>UAS-cat) also extended lifespan (Fig. S1d). Furthermore, catalase-mediated lifespan extension was independent of *Wolbachia* status (Figs. S1e,f), which can influence fly longevity and physiology^{11,12}. In addition to lifespan extension, the catalase females also exhibited increased healthspan¹³, as inferred from their enhanced climbing ability with age (Fig. 1c). To explore effects on age-specific mortality, trajectories derived from the survival curves revealed a shift in the intercept, but not the slope (Fig. 1d), indicating that catalase over-expression decreased the overall risk of death, rather than slowing its rate of increase with age¹⁴. Therefore, the catalase over-expressing flies were healthier for longer.

Using the inducible GeneSwitch system showed that induction of catalase from middle-age (d28 and d42) or old-age (d56) was sufficient to extend lifespan (Figs. 1e, S1g), although not to the full extent from induction at d2. By d56, the -RU control flies had already started dying, yet switching to +RU treatment even at this late stage still enhanced survival. This implies that for full benefits the redox shift needs to occur early in life, yet late-onset still offers protection. The level of catalase over-expression induced by RU was equivalent at all ages, as were the levels of endogenous catalase in the controls, eliminating any contribution from changes in RU consumption or endogenous catalase expression with age (Fig. S1h). We conclude that ubiquitous up-regulation of catalase improves healthspan and extends lifespan in female WT flies. Interestingly, tissue-specific catalase up-regulation using a range of drivers (e.g. tubule, gut, fat body, neuronal; Figs. S1i-m), did not recapitulate the strong lifespan extension obtained by the ubiquitous drivers, suggesting that catalase is either acting in an untested tissue (or combination of tissues), or alternatively is required at a more global organismal level.

The catalase over-expressors were exceptionally resistant to multiple modes of oxidative stress - by exogenous H₂O₂ (Fig. 1f), the redox cyler paraquat both upon feeding (Fig. 1g) and injection (Fig. 1h), as well as hyperoxia (Figs. 1i, S1n). However, this enhanced oxidative stress resistance is unlikely to explain the lifespan extension in females, because catalase over-expression protected males to a similar extent against oxidative stress without increasing longevity.

To explore the mechanism underlying the catalase-mediated lifespan extension, we examined its relationship to dietary restriction (DR), which is a robust and evolutionary conserved nutritional intervention known to have health and longevity benefits¹⁵. We measured the lifespan response of catalase over-expressor females to DR by varying the yeast content (i.e. protein source) in the food, while maintaining the sugar content constant¹⁶. This generated a typical tent-shaped response (Fig. 2a), with lifespan decreased at very low yeast levels (0.1x), highest under restricted

conditions (0.5x), then gradually shortened towards more fully-fed conditions (1.5x). The lifespan of the catalase flies was enhanced relative to controls at all yeast levels (Fig. 2a), while fecundity increased with yeast content throughout the 0.1-1.5x range for both the control and catalase females (Fig. S2a). Therefore, the catalase over-expressor females exhibit a normal DR response, and the lifespan extension upon catalase up-regulation is not mediated by the activation of DR pathways.

While the DR experiment revealed improved survival compared to control at a range of yeast concentrations (Fig. 2a), including extremely poor nutritional conditions (0.1x-yeast; Fig. 2b), we unexpectedly observed that the catalase over-expressing females, but not males, were acutely sensitive to complete starvation (Fig. 2c). There was no difference in triacylglyceride (TAG) levels at d7 (t=0 in Fig. 2d) and during a starvation time course (Fig. 2d). Similarly, the levels of glycogen storage and mobilisation were the same in control and over-expressor females (Fig. 2e). Therefore, the starvation sensitivity of the catalase flies was not due to differences in metabolic energy reserves or their mobilisation.

Autophagy is a known longevity assurance process, involved in the response to nutritional challenges such as starvation¹⁷⁻²². Furthermore, there is evidence for redox-regulation of autophagy²³. Therefore, we next explored the involvement of autophagy in the differential starvation response and longevity of the catalase flies. To monitor autophagy status *in vivo* directly, we quantified the levels of Atg8 (LC3 in mammals), a major autophagosome marker. The levels of both the de-lipidated (Atg8-I) and lipidated (Atg8-II) forms were strongly elevated in the catalase over-expressors (Fig. 2f), indicating that autophagy is induced. Indeed, levels of autophagy are physiologically fine-tuned, with both autophagy inhibition and excessive activation shown to induce starvation sensitivity *in vivo*^{24,25}. Furthermore, similarly to the starvation stress, the catalase flies were also sensitive to treatment with the autophagy inhibitor chloroquine (Fig. 2g), confirming that autophagy is affected.

To test the involvement of autophagy in the longevity of the catalase flies, we down-regulated autophagy by RNAi of Atg5 (Figs. S2b). Atg5 knock-down enhanced sensitivity to starvation (Fig. S2c)²⁴, but did not affect the lifespan of control females under fed conditions (da-GS>UAS-Atg5RNAi ±RU; Fig. 2h). Importantly, the lifespan extension by catalase over-expression was abolished in an Atg5-RNAi background (da-GS>UAS-Atg5RNAi+UAS-cat ±RU; Fig. 2i). Therefore, autophagy is required for the enhanced longevity by catalase up-regulation.

Redox regulation of autophagy has been described for Atg4 in the context of starvation-induced ROS production *in vitro*²⁶. Atg4 is the only cysteine peptidase amongst the autophagy components, and is essential for autophagosome biogenesis²³. Atg4 regulates autophagy by processing Atg8 at two critical stages: 1) the initial cleavage of Atg8, mediated by the redox-insensitive catalytic cysteine of Atg4, therefore this first step promoting Atg8 lipidation is redox-

independent; and 2) the subsequent redox-dependent de-lipidation of Atg8, which is selectively inactivated upon oxidation of an adjacent redox-regulatory cysteine in Atg4. Therefore, under oxidising conditions, lipidated Atg8 accumulates due to the redox-mediated suppression of de-conjugation by Atg4, thereby enhancing autophagosome biogenesis and promoting Atg4-mediated autophagy (Fig. 3a).

We recently showed that fasting for 24 h is associated with a strong oxidising shift of bulk cysteine residues in *Drosophila in vivo*²⁷. We therefore hypothesised that the starvation sensitivity of the da-GAL4>UAS-cat females may be attributed to such thiol redox changes. To explore the effects of catalase up-regulation on global thiol redox state, we applied the same redox proteomic technique, OxICAT²⁷, to the catalase over-expressing females. In OxICAT, samples undergo differential labelling of cysteine residues according to redox status, followed by trypsin proteolysis and enrichment for cysteine-containing peptides, and finally detection by tandem mass spectrometry. This allows both the identification of redox-responsive cysteine residues, as well as determination of their redox state. The bulk redox state of cysteines in control flies does not change with age, with the majority remaining at ~10-15% oxidised²⁷. In contrast, the catalase over-expressors displayed an oxidising shift in cysteine redox state relative to controls with increasing age (Figs. 3b-d, S3a-d, Table S2). This finding is counter-intuitive, since we are up-regulating an antioxidant enzyme. Therefore, we tested the hypothesis that by quenching H₂O₂, catalase over-expression blocks H₂O₂ redox signals that up-regulate other antioxidant systems and redox couples. The Keap1/Nrf2 signalling pathway is an appealing candidate for this process, as it is an oxidative stress response pathway that enhances the expression of a range of redox processes and is known to be redox-regulated in *Drosophila*²⁸. To assess this pathway, we used a transgenic reporter for Keap1/Nrf2 activity (gstD-GFP). This pathway was up-regulated with age in controls, but not in the long-lived catalase flies (Fig. S3e), suggesting that catalase over-expression prevented the induction of Keap1/Nrf2 signalling with age and thus the induction of a range of redox processes. Altogether, we have shown that catalase flies undergo an unexpected global oxidising thiol redox shift with age. This oxidation is consistent with the enhancement of autophagy via redox-regulation of Atg4.

The protein sequence of Atg4 is evolutionarily conserved, with both the catalytic cysteine (Cys98 in *Drosophila* Atg4a) and the adjacent redox-regulatory cysteine (Cys102 in *Drosophila* Atg4a) present in flies and mammals (Figs. 4a, S4a). To dissect the physiological role of Atg4 redox regulation *in vivo*, we generated a transgenic knock-in fly line by CRISPR, where the regulatory cysteine in endogenous Atg4a was replaced by a redox-inactive serine residue (C102S mutant). Basal levels of autophagy were not affected under control conditions (Fig. 4b, UAS-cat/+, Atg4a-C102S), whereas autophagy induction by catalase over-expression was fully abolished (Fig. 4b, da-

GAL4>UAS-cat, Atg4a-C102S). Therefore, this redox-regulatory cysteine in Atg4 is required for autophagy induction by redox signalling *in vivo*, as previously reported *in vitro* ²⁶.

To interrogate the role of Atg4a Cys102 in mediating the longevity of the catalase flies, we performed survival assays with the Atg4a-WT CRISPR control line, and reproduced the catalase lifespan extension in this background (Fig. 4c). The Atg4a-C102S point mutation did not affect survival of control flies, confirming that this knock-in alone is not deleterious (Fig. 4d). Critically, in contrast to the Atg4a-WT control, the lifespan extension upon catalase up-regulation was fully abolished in the Atg4a-C102S mutant background (Fig. 4d). Therefore, redox-regulation of autophagy via Atg4a-Cys102 mediates the longevity upon catalase over-expression. Enhancing autophagy is an evolutionarily conserved intervention associated with health and survival benefits, and here we demonstrate that selective redox-mediated up-regulation of autophagy can extend lifespan.

Many attempts have been made to extend lifespan in model organisms by enhancing their antioxidant capacity, notably through the over-expression of antioxidant enzymes, including catalase ²⁹⁻³². These trials have been largely unsuccessful, casting doubt on the causative role of ROS and oxidative damage in ageing ³³. Therefore, our finding that catalase over-expression extends lifespan was at first surprising. The original study over-expressing catalase in *Drosophila* found no effect on lifespan and only modest resistance to oxidative stress by H₂O₂ ²⁹. However, the study used an extra chromosomal copy under its endogenous promoter, resulting in far lower over-expression of catalase (~1.75-fold at the mRNA level and ~1.5-fold increased enzyme activity). The degree of catalase over-expression is therefore likely to be important for the lifespan extension. Furthermore, the earlier study used only males, while our findings show a robust effect specifically in females. The results of the two studies are therefore not discordant with one another.

Our redox proteomic analysis has revealed that the catalase flies undergo a thiol oxidising shift in bulk cysteine redox state. Interestingly, this pattern is similar to our earlier observations in WT flies under starvation stress ²⁷. Nutrient deprivation can directly affect redox homeostasis by depleting the provision of important reducing equivalents, such as NADPH and glutathione, mediating an intracellular oxidising shift ³⁴. Therefore, we suggest that the catalase flies undergo a thiol oxidising shift that is perceived as an internal state of starvation, which triggers the induction of autophagy as a protective response (Fig. 4e). Autophagy plays a fundamental role in healthy physiology, such as cellular differentiation, tissue remodelling, and mitochondrial homeostasis, as well as in the response to stress and the clearance of cellular damage ²². Consequently, enhanced autophagy is a common denominator of many evolutionary conserved interventions that extend lifespan ^{21,22}, both genetically, such as down-regulation of insulin signalling ³⁵, and pharmacologically, for instance rapamycin treatment ³⁶. Furthermore, direct up-regulation of

autophagy has been shown to exert health benefits and extend lifespan in a range of model organisms including worms, flies and mice³⁷⁻³⁹. Several components of the autophagy pathway are known to be redox regulated, including Atg3 and Atg7⁴⁰, as well as the focus of our study Atg4²⁶. Overall, we have shown that shifting the *in vivo* redox state of *Drosophila* through over-expression of catalase extends lifespan and healthspan in females through redox regulation of autophagy via a key redox-regulatory cysteine in Atg4a. Our findings further emphasise the importance of fine-tuning autophagy in health and disease, and demonstrate how manipulation of redox signalling *in vivo* can ameliorate the effects of ageing. Furthermore, our data are consistent with a growing view in the ageing field that many effects of ROS on longevity are likely to be through alterations in redox signalling rather than through lessening of oxidative damage^{7,8}.

Methods

Fly strains and husbandry. The *white Dahomey* (w^{Dah}) strain of *Drosophila melanogaster* was used as the WT background. The *Dahomey* stock was collected in 1970 in Dahomey (presently the Republic of Benin), and maintained since then as large population cages, ensuring outbreeding and overlapping generations. The w^{Dah} stock was derived by incorporation of the w^{1118} mutation into the outbred *Dahomey* background by back-crossing. Flies were either negative (w^{Dah}) or positive (w^{Dah+}) for the bacterial cytoplasmic endosymbiont *Wolbachia*, with infection status confirmed by PCR using published primers against *wsp*¹¹. The w^{Dah} stock was originally achieved by tetracycline treatment of w^{Dah+} ¹¹. All transgenic lines were back-crossed into the appropriate w^{Dah} background for at least 6-10 generations. See the Supplementary Information for details of all fly strains. Experimental flies were incubated at 25°C on a 12 h light:12 h dark cycle with 65% humidity.

Fly food. Flies were raised on standard sugar-yeast-agar medium (SYA) consisting of: 5% w/v sucrose (granulated sugar, Tate & Lyle), 10% w/v yeast (#903312, MP Biomedicals), 1.5% w/v agar (A7002, Sigma), supplemented with nipagin (Sigma H5501; 30 mL/L of 10% w/v nipagin in 95% EtOH) and propionic acid (Sigma P1386; 0.3% v/v) as mould inhibitors, added once the food had cooled down to ~60°C¹⁶. Expression via the inducible GeneSwitch system was achieved by addition of the drug RU (RU486/mifepristone; M8048, Sigma) to standard SYA once cooled down to ~60°C, typically at 200 µM from a 0.1 M stock in EtOH. For dietary restriction (DR) experiments, the yeast content was varied to give 1% (0.1x), 5% (0.5x), 7.5% (0.75x), 10% (1x = SYA) or 15% (1.5 SYA) w/v yeast⁴¹.

Experimental flies. For all experiments, eggs were collected over a defined period (<24 h) to ensure a synchronous population and reared at constant density in 200 mL bottles with SYA⁴¹.

Eclosing adults of a defined age were kept as a mixed population for ~48 h to allow mating, then separated into males and females under mild CO₂ anaesthesia, and maintained as separate sexes from then on.

Lifespan & stress assays. Lifespan assays were set up as above, typically with n~10-15 flies per vial and a total of n~100-250 flies per condition. Flies were transferred to fresh food without gassing every ~2-3 days, with deaths and censors recorded. Stress assays were performed on d7 flies (typically n>100 per condition in groups of ~15-20 flies per vial), with deaths scored regularly following initiation of treatment. See Table S1 for full survival assay information. For H₂O₂ resistance, flies were transferred onto medium containing 5% v/v H₂O₂ (Sigma H1009), 5% w/v sucrose, 1.5% w/v agar. For paraquat stress, flies were either transferred onto standard SYA food supplemented with 20 mM paraquat (Sigma 856177), or injected with 75 nL of 1 mg/mL paraquat in Ringers buffer (3 mM CaCl₂, 182 mM KCl, 46 mM NaCl, 10 mM Tris base, pH 7.2 HCl) and maintained on standard SYA⁴². Starvation stress was assayed by transferring flies to 1.5% w/v agar medium, which lacks nutrients but allows hydration. Chloroquine (10 mM, Sigma C6628) was prepared in 5% w/v sugar, 1.5% w/v agar. Hyperoxia was performed by incubating flies on standard SYA vials in a glove box chamber set at 90% O₂ using a ProOx controller (BioSpherix). The majority of lifespans (Figs. 1a-b, 2b,h-i, 4c-d, S1d-f, S2c) and stress assays (Figs. 1f-g, 2c,g) were repeated at least twice as independent biological experiments, except Figs. 1h-i and S1c,g,i-m, which were performed once.

Development time. Eggs were collected from flies in cages onto grape juice agar plates over a defined time window (~4 h). After ~24 h, the resulting L1 larvae were picked onto SYA food at a density of 50 per vial (n=500 total per genotype), and the time to adult eclosion was monitored.

Climbing assay. Climbing ability (negative geotaxis) was assayed essentially as described⁴³. Briefly, groups of 15 flies were transferred to a sawn-open 25 mL serological pipette (35 cm long, 1.5 cm diameter), with the base sealed by parafilm. The flies were tapped down within the column and observed during 45 s, after which their location was recorded. The column was separated into three sections: top 10 cm, middle, bottom 3 cm. Each cohort was evaluated 3 times, using 5 groups per genotype. The climbing performance index was calculated as: $1/2 (n^{\text{total}} + n^{\text{top}} - n^{\text{bottom}}/n^{\text{total}})$.

Metabolic and molecular assays. Flies for molecular experiments were rapidly transferred to pre-chilled microtubes via a small plastic funnel and snap frozen in liquid nitrogen, then stored at -80°C until required. Flies were always frozen at approximately the same time of day to minimise any

circadian variation. For some assays, frozen flies were separated into body segments using forceful manual impact ⁴⁴.

Western blotting. Frozen fly samples were homogenised directly into 2X Laemmli loading buffer (Bio-Rad) supplemented with 5% v/v β -mercaptoethanol (Sigma) using a pellet pestle and motor (usually 5/10 females into 100/200 μ L) and separated by standard SDS-PAGE. The following primary antibodies were used at the indicated dilutions: anti-actin (AbCam Ab1801; 1:1,000), anti-Atg8 (a generous gift from K. Köhler ⁴⁵; 1:1,000), anti-catalase (Sigma C0979; 1:10,000), anti-GFP (Cell Signaling #2955; 1:1,000). Blots were developed using standard ECL, followed by analysis with FIJI (ImageJ) software.

Energy storage assays. Whole body triacylglyceride (TAG) and glycogen levels were measured in d7 females (n=5 flies per sample, n=6-8 replicates per genotype) under control (fed) conditions and in response to starvation. For the TAG assay, flies were homogenised in 0.05% v/v Tween-20 and assayed using the Triglyceride Infinity Reagent (Thermo-Scientific TR22421) in a 96-well plate measuring absorbance at 540 nm. For the glycogen assay, flies were homogenised in saturated sodium sulphate, then the subsequent pellet was resuspended in anthrone reagent (Sigma 319899) and assayed in a 96-well plate measuring absorbance at 620 nm ⁴⁶.

OxICAT. To measure the redox state of protein cysteine residues, we performed redox proteomics using OxICAT, with protein isolation, cysteine-residue labelling, peptide preparation and LC-MS/MS analysis performed exactly as described previously ²⁷. See the Supplementary Information for further details.

Statistical analysis. Lifespan and stress assays were plotted as cumulative survival curves, and statistical analysis was performed by Log-Rank test. Other data were analysed by Student's t-test or ANOVA as appropriate in GraphPad Prism 8.

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Author contributions

Conceived the project: HMC and LP. Performed experiments: HMC, IB, FC, JICQ and CL. Provided technical support: AF, MB and JA. Generated the CRISPR mutants: SG. Conducted and analysed the redox proteomics: KEM and AMJ. Directed the work: HMC, MPM and LP. Wrote the manuscript: HMC, MPM and LP, with input from all the authors.

Competing interests

The authors declare no competing interests.

Figure legends

Fig. 1 | Ubiquitous catalase over-expression extends lifespan in flies, independently of oxidative stress resistance. **a**, Constitutive, ubiquitous catalase over-expression (da-GAL4>UAS-cat) extended the survival of female flies in a *w^{Dah}* (*white Dahomey*) WT background relative to the UAS-cat/+ ($p=5.7 \times 10^{-7}$) and da-GAL4/+ ($p=3.3 \times 10^{-6}$) controls. The control lines were not significantly different from each other ($p=0.7504$). No effect was observed in males ($p>0.05$ for all comparisons). Lifespans were performed with $n=200$ flies per condition. **b**, Inducible catalase over-expression from early adulthood (d2) using the GeneSwitch system extended the lifespan of female flies (da-GS>UAS-cat $\pm$ RU, $p=1.1 \times 10^{-16}$). RU had no effect on the da-GS/+ control line ($p=0.7161$). Lifespans were performed with $n=225-300$ flies per condition. Inset: catalase over-expression assessed by Western blotting in whole d9 females (=d7 of RU induction), with actin as a loading control. **c**, Healthspan, inferred from climbing performance, was improved in catalase over-expressing females. Climbing was assayed on da-GS>UAS-cat females $\pm$ RU to control for effects of eye colour on this behaviour. Data are presented as box-and-whisker plots (min/max error bars) of $n=5$ replicates per condition, each with $n=15$ flies per sample, analysed by unpaired Student's t-test. **d**, Mortality trajectories of the da-GS>UAS-cat $\pm$ RU survival curves from (**b**), fitted with a linear regression trendline (dotted line). **e**, Late onset over-expression of catalase using the inducible GeneSwitch system from either middle-age (d28 and d42) and old-age (d56) extended the lifespan of female flies ($p=7.2 \times 10^{-8}$, $p=1.0 \times 10^{-7}$ and $p=1.4 \times 10^{-3}$, respectively against the -RU control). Lifespans were performed with $n=270$ flies per condition, and were plotted from point of RU induction relative to the remaining -RU control flies at that age (see Fig. S1g for the original survival data). **f**, Catalase over-expressing flies were strongly resistant to exogenous H_2O_2 stress relative to controls (da-GAL4>UAS-cat v. UAS-cat/+; $p=5.0 \times 10^{-61}$ females, $p=3.2 \times 10^{-33}$ males). H_2O_2 treatment (5% v/v in sucrose/agar medium) was initiated at d7, with $n=105$ males ($n=75$ for da-GS>UAS-cat) and $n=120$ females per condition. **g**, Catalase over-expressing flies were resistant to chronic dietary paraquat stress relative to control flies (da-GAL4>UAS-cat v. UAS-cat/+; $p=6.93 \times 10^{-31}$ females, $p=7.99 \times 10^{-16}$ males). Paraquat treatment (20 mM in SYA food) was initiated at d7, with $n=100$ flies per condition. **h**, Catalase over-expressing flies (da-GAL4>UAS-cat) were resistant to acute paraquat stress relative to controls (UAS-cat/+). d7 females were injected with 75 nL of 1 mg/mL paraquat in Ringers buffer (+PQ, $n=150$ flies) or mock injected with buffer alone (-PQ, $n=120$ flies). **i**, Catalase over-expressing flies were resistant to environmental hyperoxia stress relative to controls (da-GAL4>UAS-cat v. UAS-cat/+; $p=1.7 \times 10^{-8}$ females, $p=8.1 \times 10^{-3}$ males). Incubation at 90% O_2 was initiated at d7, with $n=120$ flies per condition (except $n=90$ for UAS-cat/+ females). All survival assays (**a,b,e,f,g,i**) were analysed by Log-Rank test (see Table S1 for full n numbers and p values). n/s, $p>0.05$; *, $p<0.05$; **, $p<0.01$; ***, $p<0.001$.

Fig. 2 | Catalase lifespan extension requires autophagy. **a**, Catalase over-expressor (da-GAL4>UAS-cat) and control (UAS-cat/+) females displayed a normal response to dietary restriction (DR). Median lifespan is plotted against the yeast content of the diet, with 1x corresponding to standard SYA food. Data are the means $\pm$ range of 2 independent lifespan experiments, each set up with n=150-160 flies per genotype, analysed by two-way ANOVA (Fisher's LSD). **b**, Survival curve on 0.1x-fold yeast from (**a**). The catalase over-expressor females (da-GAL4>UAS-cat) were longer-lived than controls (UAS-cat/+) under low yeast nutritional conditions ($p=3.6 \times 10^{-10}$, n=160 flies per genotype). **c**, Catalase over-expressor females (da-GAL4>UAS-cat) were sensitive to starvation stress relative to UAS-cat/+ and da-GAL4/+ controls ($p=5.6 \times 10^{-5}$ and $p=7.6 \times 10^{-7}$, respectively). No difference was observed in males ($p>0.05$ for all comparisons). Assays were performed at d7 with n=120 flies per condition (except n=80 for UAS-cat/+ females). **d-e**, Triacylglyceride (TAG, **d**) and glycogen (**e**) levels in whole females assayed at d7 (t=0) and depletion in response to starvation treatment. Data are presented as box-and-whisker plots (min/max error bars) of n=4-6 replicates per genotype, each with n=5 females per sample, analysed by unpaired Student's t-test ($p>0.05$). **f**, Catalase over-expressor females (da-GAL4>UAS-cat) displayed enhanced autophagy induction compared to UAS-cat/+ controls at d7, assessed by Western blotting against Atg8, normalised to actin. Data are means $\pm$ SD of n=6 biological replicates, each with n=10 abdomens per sample, analysed by paired Student's t-test (*, $p<0.05$). Right, typical bands probed against Atg8 with actin as a loading control (see Fig. S4b for the full blot). **g**, Catalase over-expressor females (da-GAL4>UAS-cat) were sensitive to treatment with the autophagy inhibitor chloroquine (10 mM in sucrose/agar medium) relative to UAS-cat/+ and da-GAL4/+ controls ($p=2.6 \times 10^{-5}$ and $p=3.5 \times 10^{-9}$, respectively). Assays were performed at d7 with n=160 flies per condition. **h**, Global Atg5 knock-down did not decrease lifespan in a WT background under control conditions (da-GS>UAS-Atg5RNAi $\pm$ RU; $p=0.4177$). Lifespan assays were performed on n=225-240 females. **i**, Lifespan extension upon catalase over-expression (da-GS>UAS-cat $\pm$ RU; $p=7.2 \times 10^{-15}$) was abolished in an autophagy-deficient background (da-GS>UAS-cat+UAS-Atg5RNAi $\pm$ RU; $p=0.1701$). Survival assays (**b,c,g,h,i**) were analysed by Log-Rank test (see Table S1 for full n numbers and p values). n/s, $p>0.05$; *, $p<0.05$; ***, $p<0.001$.

Fig. 3 | Catalase flies undergo an oxidising shift in global thiol redox state. **a**, Scheme showing the dual function of Atg4 in autophagy: 1) initial redox-independent cleavage of Atg8, to expose a C-terminal glycine residue enabling lipidation by PE (phosphatidyl-ethanolamine) via the E1-like enzyme Atg7 and the E2-like enzyme Atg3. The conjugated Atg8-PE is involved in autophagosome elongation/closure; 2) redox-dependent de-lipidation of Atg8-PE, allowing interaction and fusion

of the autophagosome with the endosomal-lysosomal compartments, and recycling of cleaved Atg8. Oxidation of a redox-regulatory cysteine selectively inactivates the Atg8-PE de-conjugation activity of Atg4, promoting autophagosome biogenesis and therefore enhancing Atg4-mediated autophagy. **b**, Redox proteomic (OxICAT) analysis of d7, d28 and d56 catalase over-expressing females (da-GAL4>UAS-cat) compared to control (UAS-cat/+). Distribution of total cysteine residue oxidation levels, plotted as the proportion of the total number of peptides containing unique cysteine residues in each 5% quantile of percentage oxidation. Data are means $\pm$ SEM of n=5 biological replicates. **c-d**, Oxidation state of cysteine residues present, comparing control versus catalase over-expressor females at d7 (**c**) and d56 (**d**). Data points above the diagonal dotted line (slope=1) indicate cysteine residues more oxidised upon catalase up-regulation, with red symbols designating significance ($p < 0.05$), assessed by unpaired two-tailed Student's t-test.

Fig. 4 | Redox regulation of autophagy via Atg4a Cys102 extends lifespan. **a**, Multiple sequence alignment of the Atg4a protein from *Drosophila* with the mouse and human orthologues, showing the catalytic cysteine residue (Cys98 in *Drosophila*) and the redox-regulatory cysteine (Cys102 in *Drosophila*). See Fig. S4a for the full sequence. **b**, Autophagy levels assessed by Western blotting against Atg8. The induction of autophagy in response to catalase over-expression was maintained in Atg4a-WT flies, but abolished in the Atg4a-C102S mutant background. See Fig. S4b for the full blots. Quantification of Atg8 levels by densitometry, normalised to actin as a loading control. Data are means $\pm$ range of n=2 biological replicates, analysed by two-way ANOVA (Fisher LSD). **c-d**, Catalase over-expression extended lifespan in the Atg4a-WT control (**c**), but not in the redox-insensitive Atg4a-C102S knock-in background (**d**). Survival assays (**c,d**) were analysed by Log-Rank test (see Table S1 for full n numbers and p values). n/s, $p > 0.05$; ***, $p < 0.001$. **e**, Scheme illustrating the mechanism underlying the lifespan extension upon redox regulation of autophagy.

Figure 1

Figure 2

Figure 3

Figure 4