The Mitochondrial Permeability Transition Pore Activates a Maladaptive Mitochondrial Unfolded Protein Response

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SUMMARY
Mitochondrial activity determines aging rate and the onset of chronic diseases. The mitochondrial permeability transition pore (mPTP) is a pathological pore in the inner mitochondrial membrane thought to be composed of the F-ATP synthase (complex V). Oligomycin sensitivity-conferring protein (OSCP), a subunit of F-ATP synthase, helps protect against mPTP formation. How the destabilization of OSCP may contribute to aging, however, is unclear. We have found that loss OSCP in the nematode Caenorhabditis elegans initiates the mPTP and shortens lifespan specifically during adulthood, in part via initiation of the mitochondrial unfolded protein response (UPRmt). Genetic or pharmacological inhibition of the mPTP inhibits the UPRmt and restores normal lifespan. The mitochondria of long-lived mutants are buffered from the maladaptive UPRmt, partially via the transcription factor FOXO3a/daf-16. Our findings reveal how the mPTP/UPRmt nexus may contribute to aging and age-related diseases and how inhibition of the UPRmt may be protective under certain conditions.

INTRODUCTION
As mitochondrial function declines with age, the frequency of the mPTP increases (Rottenberg and Hoek, 2017). The mPTP is central to early-stage pathologies associated with several age-related diseases, including Alzheimer’s and Parkinson’s disease (AD, PD) and late-stage pathologies of ischemia-reperfusion injuries including heart attack and stroke (Ong et al., 2015; Panel et al., 2018). The mPTP is a pathological channel that forms in the inner mitochondrial membrane in response to excessive cytosolic Ca²⁺ or high ROS conditions. Sustained opening of the mPTP leads to outer mitochondrial membrane rupture, release of Ca²⁺ into the cytosol, and cell death (Bernardi and Di Lisa, 2015). Cyclosporine A (CysA), a well-characterized mPTP inhibitor, inhibits the mPTP by binding and sequestering cyclophilin D, a mitochondrially-localized peptidyl prolyl isomerase that helps catalyze pore formation (Basso et al., 2005; Nakagawa et al., 2005). Genetic inhibition of cyclophilin D protects against mPTP formation (Baines et al., 2005). The mPTP is also modulated by adenine nucleotide translocases (ANTs), which exchange ADP and ATP across the IMM. Genetic inhibition of ANT5 also help prevent pore formation (Karch et al., 2019; Kokoszka et al., 2004). Both CysA and loss of ANTs have been shown to extend lifespan in C. elegans (Farina et al., 2008; Ye et al., 2014). Thus, the mPTP appears to be an important modulator of healthspan and lifespan.
Though the identification of the proteins that make up the mPTP is controversial, recent studies have pointed to F-ATP synthase (complex V) as the most probable candidate due to its ability to bind cyclophilin D and form Ca\(^{2+}\) currents (Bernardi and Di Lisa, 2015; Mnatsakanyan and Jonas, 2020). Some models posit that dimeric forms of F-ATP synthase open to form a pore while other models have suggested that the pore occurs via the membrane-bound proton rotor (Alavian et al., 2014; Azarashvili et al., 2014; Bonora et al., 2013; Bonora et al., 2017; Giorgio et al., 2013; Mnatsakanyan et al., 2019; Neginiskaya et al., 2019; Urbani et al., 2019). ATP5O, also known as oligomycin sensitivity-conferring protein (OSCP), a subunit of the F-ATP synthase that regulates ATPase rotational activity to provide efficient ATP production (Murphy et al., 2019), has emerged as an important regulator of the mPTP. OSCP levels decrease with normal aging and loss of OSCP increases propensity for pore formation in vitro and in mouse models of AD (Beck et al., 2016b; Gauba et al., 2017; Giorgio et al., 2013). Conversely, OSCP confers protection against the pore under low pH conditions and OSCP overexpression protects from mPTP initiation in AD and cardiac dysfunction models (Antoniel et al., 2018; Beck et al., 2016b; Guo et al., 2020). Thus, OSCP appears to be an important regulator of aging and disease progression, possibly via its ability to modulate mPTP formation.

Under stress, the mitochondria attempt to repair the damage, recycle damaged mitochondria, or, under deleterious circumstances, initiate cell death. Similar to the endoplasmic reticulum unfolded protein response (UPR\textsuperscript{ER}) and the cytoplasmic heat shock response (HSR), the mitochondrial unfolded protein response (UPR\textsuperscript{mt}) is capable of initiating a broad-range transcriptional response that, among other functions, aids in the refolding of mitochondrial matrix proteins (Naresh and Haynes, 2019). Recent studies also show that a loss of mitochondrial membrane potential correlates with activation of the UPR\textsuperscript{mt}, and disruption of mitochondrial processes other than protein misfolding, such as those involved in TCA cycle and lipid catabolism, also induce the UPR\textsuperscript{mt} (Rolland et al., 2019). UPR\textsuperscript{mt} activation is associated with longevity and improvement in neurodegenerative models (Durieux et al., 2011; Houtkooper et al., 2013; Kim et al., 2016; Merkwirth et al., 2016; Sorrentino et al., 2017; Tian et al., 2016), but it has also conversely been shown to increase neurodegeneration, propagate mtDNA mutations, and exacerbate ischemic conditions (Lin et al., 2016; Martinez et al., 2017; Yung et al., 2019), underscoring its complexity. If left unmitigated, UPRs can initiate cell death (Iurlaro and Munoz-Pinedo, 2016; Munch and Harper, 2016). Thus, the context or cellular environment are important determinants of whether UPR\textsuperscript{mt} induction results in beneficial or detrimental effects.

In \textit{C. elegans}, mild mitochondrial perturbations early in life can extend lifespan. Loss of OSCP/atp-3 has previously been shown to extend lifespan when initiated during larval development (Dillin et al., 2002; Rea et al., 2007). In contrast, here, we have determined that loss of OSCP/atp-3 during adulthood leads to initiation of the mPTP, the UPR\textsuperscript{mt}, and a shortened lifespan. Surprisingly, \textit{atfs-1}, the UPR\textsuperscript{mt} master transcription factor (Haynes et al., 2010; Nargund et al., 2012), helps drive the reduction of lifespan, suggesting that the UPR\textsuperscript{mt} program promotes aging during adulthood. The adult UPR\textsuperscript{mt} is responsive to mPTP regulators, including the immunosuppressive drug, CysA, as well as a mitochondrially-localized cyclophilin and ANT, pointing to a previously undiscovered coordination between the UPR\textsuperscript{mt} and the mPTP. We find that proton rotor subunits as well as subunits important for dimerization of the F-ATP synthase are essential for transducing the adult UPR\textsuperscript{mt}. Loss of these subunits as well as pharmacological CysA treatment restores lifespan due to loss of OSCP/atp-3. These results are
consistent with current models that posit that the F-ATP synthase forms the mPTP (Bernardi and Di Lisa, 2015). Remarkably, we find that long-lived mutants which have previously been shown to be protected from the mPTP (Zhou et al., 2019) expressing high FOXO3a/\textit{daf-16} activity do not initiate a UPR\textsuperscript{mt} during adulthood. Overall, our findings point to a model in which loss of OSCP/\textit{atp-3} in adults induces mPTP formation and subsequent detrimental activation of the UPR\textsuperscript{mt}, from which long-lived mutants are buttressed. Understanding the relationship between these two mitochondrial processes will undoubtedly further our understanding of aging as well as disparate age-related disorders, including neurodegenerative diseases, cancer, heart-attack, and stroke.

**RESULTS**

**Loss of OSCP/\textit{atp-3} Induces Mitochondrial Permeability Transition Pore (mPTP) Characteristics**

The opening of the mPTP is characterized by a loss of membrane potential as well as an increase in cytosolic Ca\textsuperscript{2+}. We observed that a reduction in the abundance of OSCP/\textit{atp-3} by RNA interference (RNAi) caused a loss of membrane potential as measured by the mitochondrial dye, TMRM, in young adult worms (Figure 1A, 1B). Strikingly, RNAi of other OXPHOS subunits from complex I, IV, and V had no effect on membrane potential. RNAi of OSCP/\textit{atp-3} during adulthood also caused an increase in cytosolic Ca\textsuperscript{2+} as measured by the intestinal FRET-based Ca\textsuperscript{2+} reporter, KWN190, while RNAi of other F-ATP synthase subunits, \textit{F6/\textit{atp-4}} and \textit{d/\textit{atp-5}}, did not (Figure 1C). Loss of OSCP/\textit{atp-3} also induced mitochondrial swelling and fragmentation (Figure 1D, 1E). In contrast, we observed no swelling or fragmentation following the reduction in expression of the F-ATP synthase subunit \textit{d/\textit{atp-5}} (Figure 1F). These results suggest that loss of OSCP/\textit{atp-3} during adulthood uniquely induces the mPTP in \textit{C. elegans}.

To verify the stability of the mitochondrial membrane potential after RNAi of other OXPHOS subunits, we checked for RNAi efficiency via qPCR; we observed efficient mRNA reduction from subunit \textit{cco-1} from complex IV as well as from several of the complex V subunits, including \textit{atp-1}, \textit{atp-2}, \textit{atp-4}, and \textit{atp-5} (Figure S1A-S1E). We also examined protein levels for subunits in which antibodies were available and observed that RNAi of the complex I subunit NUO-2 and complex V subunits ATP-1 and ATP-2 resulted in significant knockdown of protein levels (Figure S1F-S1H). These findings show that knockdown of OXPHOS subunits other than OSCP/\textit{atp-3} do not lead to a loss of membrane potential.

**Loss of OSCP/\textit{atp-3} Induces a Unique UPR\textsuperscript{mt} During Adulthood**

A recent study showed that a loss of membrane potential is associated with inducing the UPR\textsuperscript{mt} (Rolland et al., 2019). To determine if loss of OSCP/\textit{atp-3} selectively induces the UPR\textsuperscript{mt} during adulthood due to the observed drop in membrane potential, we utilized a GFP reporter under the promoter of the mitochondrial UPR chaperone, \textit{hsp-6} (\textit{phsp-6::GFP}) (Yoneda et al., 2004) and compared it to select representative OXPHOS genes encoding complex I, III, IV, and V subunits. RNAi of OXPHOS subunits induced little to no UPR\textsuperscript{mt} if initiated after the last larval stage (L4), which we termed the post-developmental UPR\textsuperscript{mt} (pdvUPR\textsuperscript{mt}). Only RNAi of OSCP/\textit{atp-3} induced a robust UPR\textsuperscript{mt} in young adults (Figure 2A, 2B, Table S1), the timing of which corresponded with the drop in membrane potential (Figure 1A, 1B). In contrast, RNAi of all same genes induced a robust UPR\textsuperscript{mt} if initiated during the early larval stages (L1, L2, and L3) of
Figure 1: Loss of OSCP/atp-3 Selectively Recapitulates mPTP-like Characteristics
(A) RNAi of various OXPHOS subunits in N2 wild-type worms did not lead to changes in the membrane potential as measured by the TMRM dye, except for RNAi of OSCP/atp-3. RNAi was administered for 48 hours beginning at young adulthood. TMRM was administered via the NGM plates.
(B) Quantification of TMRM intensity from (A). Data are the mean ± SEM of ≤ 15 animals combined from three biological experiments. **p ≤ 0.01 by Student’s t-test. I, IV, and V correspond to OXPHOS complexes.
(C) RNAi of OSCP/atp-3 leads to an increase in cytosolic calcium as measured by the FRET-based calcium indicator protein D3cpv/cameleon. RNAi was administered for 48 hours beginning at young adulthood. Data are the mean ± SEM of ≤ 15 animals combined from three biological experiments. **p ≤ 0.01 by Student’s t-test.
(D-F) Confocal micrographs of intestinal mitochondria labeled with GFP (pges-1::GFPmt) in young adults. RNAi and was administered for 48 hours beginning at young adulthood, then worms were removed from the RNAi and aged until Day 7 of adulthood followed by collection for microscopy. Top panels, fluorescent channel; bottom panels, rendering of individual mitochondria. See Materials and Methods for details on rendering.
development (dvUPR\textsuperscript{mt}) (Figure 2A, 2B). These results are consistent with previous reports demonstrating that the UPR\textsuperscript{mt} is robustly induced during development but poorly induced during adulthood in \textit{C. elegans} (Durieux et al., 2011; Labbadia and Morimoto, 2015). Loss of membrane potential \textit{per se} via FCCP, a potent mitochondrial uncoupler, however, did not induce a UPR\textsuperscript{mt} in young adults, suggesting that uncoupling and its associated loss in membrane potential is not sufficient to induce a pdvUPR\textsuperscript{mt} (Figure S1I). Post-developmental loss of OSCP/atp-3 increased endogenous transcript levels of \textit{hsp-6} as well as endogenous HSP-6/mtHSP70 protein levels (Figure S2A, S2B). Post-developmental loss of OSCP/atp-3 mildly induced the mitochondrial chaperone reporter \textit{phsp-60::GFP} (Figure S2C, S2D). Neither the UPR\textsuperscript{ER} nor the HSR were induced by post-developmental loss of OSCP/atp-3 (Figure S2C).

RNAi of other mitochondrial genes that are known to induce a dvUPR\textsuperscript{mt}, \textit{clk-1} (coenzyme Q hydroxylase), \textit{mrps-5} (mitochondrial ribosome), and \textit{tomm-22} (translocase of outer mitochondrial membrane) (Baker et al., 2012; Bennett et al., 2014; Houtkooper et al., 2013), did not induce the pdvUPR\textsuperscript{mt} (Table S1). Importantly, we found that the pdvUPR\textsuperscript{mt} was dependent on the master UPR\textsuperscript{mt} transcription factor, \textit{atfs-1}, as well as its previously described co-factors, the peptide exporter \textit{haf-1} and the ubiquitin-like protein \textit{ubl-5} (Figure 2C, 2D) (Haynes et al., 2010), demonstrating that the pdvUPR\textsuperscript{mt} is regulated similarly to the previously described dvUPR\textsuperscript{mt}. Thus, loss of OSCP/atp-3 induces a robust and specific pdvUPR\textsuperscript{mt} which is dependent on the conserved transcription factor ATFS-1.

The Post-Developmental UPR\textsuperscript{mt} is Temporally Confined and Reversible

Given that the UPR\textsuperscript{mt} has been studied during development in \textit{C. elegans}, we sought to determine the window of the UPR\textsuperscript{mt} during adulthood. We initiated RNAi of OSCP/atp-3 beginning at the last larval stage (L4 stage) and every few hours thereafter into adulthood. GFP expression was examined 48 hours after RNAi initiation (Figure S2E). We observed that the pdvUPR\textsuperscript{mt} is initiated up to 6 hours after the L4 stage, after which RNAi of OSCP/atp-3 no longer induced the UPR\textsuperscript{mt}. In contrast, RNAi of COX5B/cco-1 of complex IV had no effects on the UPR\textsuperscript{mt} at any of these stages (Figure S2E). For all subsequent post-development experiments, RNAi was therefore administered at the young adult stage corresponding to 4-hours after the L4 stage. Thus, the pdvUPR\textsuperscript{mt} is confined to pre-gravid stages of adulthood, corresponding with previous reports showing a global decline in stress responses at the onset of egg-laying (Labbadia and Morimoto, 2015).

Previous studies have shown that developmental RNAi of \textit{cco-1} RNAi leads to persistent activation of the UPR\textsuperscript{mt} into adulthood, even after removal from RNAi (Durieux et al., 2011). Similarly, we observed that developmental RNAi of OSCP/atp-3 initiated a UPR\textsuperscript{mt} that persisted into adulthood, even after removal from \textit{atp-3} RNAi (Figure S2F). In contrast, removal from post-developmental \textit{atp-3} RNAi treatment led to a steady decline in the GFP signal (Figure S2F), suggesting the activation of the pdvUPR\textsuperscript{mt} is reversible.

Loss of ATP Synthase ATPases Induce a Post-Developmental UPR\textsuperscript{mt}

F-ATP synthase is composed of a membrane-bound proton rotor (F\textsubscript{o}), a catalytic ATPase that converts ADP to ATP (F\textsubscript{i}), and peripheral stalk and supernumerary subunits that help bridge these two portions together (Figure 2F). OSCP/atp-3 sits atop the ATPase and helps tether it to the peripheral stalk subunits. We systematically tested via RNAi whether loss of F-ATP synthase subunits other than OSCP/atp-3 could induce a pdvUPR\textsuperscript{mt}. During adulthood, loss of rotor
Figure 2: Loss of OSCP/atp-3 and ATPases Initiate a Post-developmental UPRmt

(A) Developmental RNAi of OXPHOS subunits induced the phsp-6::GFP reporter while only OSCP/atp-3 RNAi induced the reporter when initiated during post-development. For developmental treatment, worms were exposed to RNAi beginning from eggs for 72 hours. For post-developmental treatment, worms were exposed to RNAi beginning from young adulthood for 48 hours. CV, control vector RNAi; dev, development; post-dev, post-development.

(B) Quantification of GFP intensity from (A). Data are the mean ± SEM of ≤ 15 animals combined from three biological experiments. *p ≤ 0.05, **p ≤ 0.01, and ***p ≤ .0001 by Student’s t-test. dv, development; pdv, post-development.

(C) Post-developmental RNAi of OSCP/atp-3 induced the phsp-6::GFP reporter, which is abrogated via co-treatment with atfs-1, haf-1, or ubl-5 RNAi. Worms were exposed to RNAi beginning from young adulthood for 48 hours. Left panels, fluorescent channel; right panels, bright-field channel. CV, control vector RNAi; post-dev, post-development.

(D) Quantification of GFP intensity from (C). Data are the mean ± SEM of ≤ 15 animals combined from three biological experiments. **p ≤ 0.01 by Student’s t-test. pdv, post-development.

(E) Post-developmental RNAi of ATPase and c-ring subunits but not peripheral stalk or supernumerary subunits of the F-ATP synthase induced the phsp-6::GFP reporter. Worms were exposed to RNAi beginning from young adulthood for 48 hours. post-dev, post-development.

(F) Schematic of monomeric F-ATP synthase. White subunits, ATPase; black subunits, H+-rotor/c-ring; grey subunits, peripheral stalk and supernumerary subunits; red subunit, oligomycin sensitivity-conferring protein (OSCP/atp-3).

(G) Quantification of GFP intensity from (E). Data are the mean ± SEM of ≤ 15 animals combined from three biological experiments. *p ≤ 0.05, **p ≤ 0.01, and ***p ≤ .0001 by Student’s t-test. pdv, post-development.
subunits (c-ring/Y82E9BR.3), peripheral stalk or supernumerary subunits (F6/atp-4, d/atp-5, b/asb-2, e/R04F11.2, f/R53.4), induced little to no UPR\textsuperscript{mt} (Figure 2E, 2G, Table S2). Loss of the ATPase subunits (α/atp-1; β/atp-2; δ/F58F12.1; ε/hpo-18) induced a mild to moderate UPR\textsuperscript{mt}, though none as robustly as loss of OSCP/atp-3. Subunits α and ε induced a moderate UPR\textsuperscript{mt}, while loss of the β and δ subunits induced a limited, though significant, UPR\textsuperscript{mt} (Figure 2E, 2G, Table S2). In contrast, during development, loss of rotor subunits, ATPase subunits, or peripheral stalk and supernumerary subunits all induced a robust UPR\textsuperscript{mt} (Figure S2G, S2H). Thus, in addition to the robust induction of the pdvUPR\textsuperscript{mt} caused by loss of OSCP/atp-3, loss of F-ATP synthase ATPases moderately to mildly induce the pdvUPR\textsuperscript{mt}.

The Post-Developmental UPR\textsuperscript{mt} is dependent on Mitochondrial Permeability Transition Pore (mPTP) Factors

To determine if the pdvUPR\textsuperscript{mt} is initiated in response to the mPTP, we tested pharmacological and genetic modulators of the mPTP on induction of the pdvUPR\textsuperscript{mt}. The mPTP is known to be inhibited by the immunosuppressive drug, cyclosporin A (CysA). CysA binds cyclophilins, which in the cytoplasm regulates calcineurin signaling (Liu et al., 1991; Takahashi et al., 1989), while in the mitochondria inhibits the mPTP (Nicoli et al., 1996). To parse out the mitochondrial versus cytoplasmic functions of CysA, we also tested the cytoplasmic-only immunosuppressive drug, FK506, which acts similarly to CysA in that it modulates calcineurin signaling in the cytoplasm (Liu et al., 1991). We observed that CysA strongly inhibited the pdvUPR\textsuperscript{mt} but not dvUPR\textsuperscript{mt} in a dose-dependent manner (Figure 3A, 3B, Figure S3A). In contrast, we found that FK506 had no effect on the pdvUPR\textsuperscript{mt} (Figure 3A, 3B, Figure S3B), demonstrating that CysA acts in the mitochondria to suppress the pdvUPR\textsuperscript{mt}. The mPTP has been shown to be regulated by adenine nucleotide translocases (ANTs) of the inner mitochondrial membrane and loss of ANT\textsubscript{s} helps prevent the mPTP (Karch et al., 2019). We tested \textit{ant-1.1}, which is ubiquitously expressed in \textit{C. elegans}, and \textit{ant-1.2}, which is expressed predominantly in the pharynx and intestines (Farina et al., 2008). RNAi of \textit{ant-1.1} moderately suppressed the pdvUPR\textsuperscript{mt} while RNAi of \textit{ant-1.2} strongly suppressed the pdvUPR\textsuperscript{mt}, but not the dvUPR\textsuperscript{mt} (Figure 3C, 3D). In mammals, CysA acts in the mitochondria to inhibit the mPTP by binding and sequestering cyclophilin D, a peptidyl prolyl isomerase (Nicoli et al., 1996). \textit{C. elegans} contains 17 poorly defined cyclophilins, of which 2 are predicted to be mitochondrially localized, \textit{cyn-1} and \textit{cyn-17} (Table S3). RNAi of \textit{cyn-1} did not inhibit the pdvUPR\textsuperscript{mt} while \textit{cyn-17} did (Figure 3D), suggesting that \textit{cyn-17} may act similarly to cyclophilin D in mediating a conformation change that leads to the mPTP. RNAi of \textit{cyn-17} did not affect the dvUPR\textsuperscript{mt} (Figure 3C). Together, these results show that the pdvUPR\textsuperscript{mt} is regulated by canonical pharmacological and genetic mPTP factors.

Loss of Essential F-ATP Synthase Subunits Important for the mPTP Suppress the Post-Developmental UPR\textsuperscript{mt}

Current models posit that the F-ATP synthase forms a pore that is capable of releasing Ca\textsuperscript{2+} under conditions of high oxidative stress, leading to rupturing of the mitochondria and initiation of cell death cascades. Some models suggest that ATP synthase dimers are required for the mPTP (Figure 3G) and that peripheral and supernumerary subunits are essential for pore formation (Carraro et al., 2014; Guo et al., 2019). Other models demonstrate that ATP synthase monomers (Figure 2F) are sufficient for the mPTP and specify the c-ring proton rotor as the
Figure 3: The Post-developmental UPR\textsuperscript{mt} is Regulated by Pharmacological and Genetic Modulators of the mPTP

(A, B) 15 µM of the immunosuppressants cyclosporin A (CysA) and FK506 had no effect on the dvUPR (A) but selectively suppressed the pdvUPR\textsuperscript{mt} in the phsp-6::GFP reporter strain (B). Data are the mean ± SEM of \( \leq 15 \) animals combined from three biological experiments. ***\( p \leq .0001 \) by Student’s \( t \)-test.

(C, D) Loss of cyclophilin cyn-17 and adenine nucleotide translocase ant-1.2 via RNAi had no effect on the dvUPR\textsuperscript{mt} (C) while their loss suppressed the pdvUPR\textsuperscript{mt} in the phsp-6::GFP reporter strain (D). ant-1.1 partially suppressed the pdvUPR\textsuperscript{mt} (D). Data are the mean ± SEM of \( \leq 15 \) animals combined from three biological experiments. *\( p \leq .05 \), ***\( p \leq .0001 \) by Student’s \( t \)-test.

(E) Concomitant post-developmental RNAi of OSCP/atp-3 and individual F-ATP synthase subunits inhibited the post-developmental UPR\textsuperscript{mt} to varying degrees.

(F) Quantification of GFP intensity from (E). Data are the mean ± SEM of \( \leq 15 \) animals combined from three biological experiments. **\( p \leq .001 \), ***\( p \leq .0001 \) compared to CV + atp-3 condition by Student’s \( t \)-test. pdv, post-development.

(G) Schematic of ATP synthase dimers. White subunits, ATPase; black subunits, H\(^+\) rotor/c-ring; grey subunits, peripheral stalk and supernumerary subunits; red subunit, oligomycin sensitivity-conferring protein (OSCP/atp-3).
actual pore-forming component (Alavian et al., 2014; Azarashvili et al., 2014; Bonora et al., 2013; Bonora et al., 2017; Giorgio et al., 2013; Mnatsakanyan et al., 2019; Neginskaya et al., 2019; Urbani et al., 2019). To determine whether the structural integrity of F-ATP synthase subunits were required for the pdvUPR<sup>mt</sup>, we systematically knocked down OSCP/atp-3 as well as one additional F-ATP synthase subunit via RNAi. When we knocked down the c-ring subunits (color coded black) as well as peripheral and supernumerary subunits (color coded grey) via RNAi in adults, we observed nearly complete inhibition of the OSCP/atp-3 RNAi-mediated pdvUPR<sup>mt</sup> ((Figure 3E, 3F). When we knocked down the ATPase subunits in adults, we observed that loss of the β/atp-2 subunit robustly suppressed the OSCP/atp-3 RNAi-mediated pdvUPR<sup>mt</sup>, possibly due to its role in modulating Ca<sup>2+</sup> in the mPTP (Giorgio et al., 2017), while loss of α/atp-1 moderately inhibited the pdvUPR<sup>mt</sup> (Figure 3E, 3F, color coded white). Loss of the ATPase subunits δ/F58F12.1 or ε/hpo-18 in adults did not affect the pdvUPR<sup>mt</sup>. In contrast, dual loss of subunits during development all robustly activated the dvUPR<sup>mt</sup> (Figure S3C, S3E). Dual loss of subunits from other OXPHOS complexes (NDUFS3/muo-2, complex I; COX5B/cco-1, complex IV) had no effect or slightly increased the dvUPR<sup>mt</sup> and the pdvUPR<sup>mt</sup> (Fig S3D, S3F). Thus, we find subunits critical for dimerization (peripheral and supernumerary subunits) and proton translocation (c-ring rotor) are required to transduce the OSCP/atp-3 RNAi-mediated pdvUPR<sup>mt</sup>. We also find that the β/atp-2 subunit, previously found to play an important role in Ca<sup>2+</sup> mediated mPTP, is required to transduce the OSCP/atp-3 RNAi-mediated pdvUPR<sup>mt</sup>. Taken together, these findings support a model in which inhibition of the mPTP via deletion of critical F-ATP synthase subunits inhibits the pdvUPR<sup>mt</sup>.

**Loss of OSCP/atp-3 During Adulthood Promotes Aging Partially via the UPR<sup>mt</sup>**

Previous reports have shown that loss of OSCP/atp-3 initiated during development robustly increases lifespan in *C. elegans* (Dillin et al., 2002), but how loss of OXPHOS subunits during adulthood affects lifespan has not been well studied. We initiated RNAi during both development and post-development. As previously reported, we found that developmental RNAi treatment led to lifespan extension (Figure 4A, Table S4). Worms continuously exposed to OSCP/atp-3 RNAi during post-development experienced a high incidence of matricide (data not shown). To circumvent this outcome, we administered OSCP/atp-3 RNAi to young adults for 48 hours of adulthood and observed approximately a 38% decrease in lifespan independent of matricide (Figure 4A, 4B, Table S4). Surprisingly, when atfs-1 mutants were exposed to OSCP/atp-3 RNAi during adulthood for 48 hours, we observed only about a 19% decrease in lifespan, suggesting that the initiation of the pdvUPR<sup>mt</sup> via atfs-1 contributes to reduced lifespan (Figure 4B, Table S4). Loss of other OXPHOS subunits had little or no effect on lifespan when administered during adulthood for 48 hours (Figure 4C, Table S4). Thus, we have determined that loss of OSCP/atp-3 induces an age-accelerating, transcriptionally-dependent pdvUPR<sup>mt</sup> not previously described in *C. elegans*.

**Pharmacological or Genetic Inhibition of the mPTP Restores Lifespan**

To determine if inhibition of the mPTP could also inhibit its associated toxicity, we tested the mPTP inhibitor, CysA, in lifespans. We found that CysA treatment was sufficient to rescue the shortened lifespan caused by RNAi of OSCP/atp-3 in adults (Figure 4D). Based on our observations that loss of peripheral stalk subunits are capable of suppressing the pdvUPR<sup>mt</sup>, we tested if their loss would also suppress the shortened lifespan. RNAi with either F6/atp-4 or
Figure 4: Loss of OSCP/atp-3 Shortens Lifespan, which is Restored by Pharmacological or Genetic Inhibition of the mPTP

(A) Survival curves of wild-type N2 animals on CV or OSCP/atp-3 RNAi initiated during development or post-development. Representative curves selected from three biological experiments. **p ≤ .0001 by Log Rank (Mantel Cox). CV, control vector RNAi.

(B) Survival curves of atfs-1(tm4525) when OSCP/atp-3 RNAi is initiated during post-development. Representative curves selected from three biological experiments. **p ≤ 0.01 by Log Rank (Mantel Cox). Bar graph is a quantification of percent change in lifespan of N2 or atfs-1(tm4525) mutant. **p ≤ 0.01 by Student's t-test is the mean of the percent change in lifespan ± SEM from three biological experiments. CV, control vector RNAi.

(C) Survival curves of wild-type N2 animals on indicated RNAi initiated during post-development. Representative curves selected from three biological experiments. **p ≤ 0.01 by Log Rank (Mantel Cox).

(D) Survival curves of wild-type N2 animals on CV or OSCP/atp-3 RNAi initiated during post-development and treated with 15µM CysA. Lifespan curves from two pooled biological replicates, ***p ≤ .0001 by Log Rank (Mantel Cox) between CV and OSCP/atp-3 RNAi. CV, control vector RNAi.

(E) Survival curves of wild-type N2 animals on CV or OSCP/atp-3 RNAi with the additional loss of either F6 subunit/atp-4 or d subunit/atp-3 subunits via RNAi initiated during post-development. Lifespan curves from two pooled biological replicates, ***p ≤ .0001 by Log Rank (Mantel Cox). CV, control vector RNAi.

(F) Quantification of membrane potential as measured by TMRM after post-developmental RNAi of OSCP/atp-3 with either F6 subunit/atp-4 or d subunit/atp-3 RNAi. Data are the mean ± SEM of ≤ 15 animals combined from three experiments. *p ≤ 0.05 by Student’s t-test. CV, control vector RNAi.

(G, H) 63X confocal micrographs of intestinal mitochondrial GFP worms (pges-1::GFPpwe) treated with post-developmental dual-RNAi of d subunit/atp-5 with CV or OSCP/atp-3 RNAi. Top panel, fluorescent channel, bottom panels, rendering of individual mitochondria. See Materials and Methods for details on rendering.
d/atp-5 rescued the shortened lifespan caused by RNAi of OSCP/atp-3 in adults as well as restored the loss in membrane potential (Figure 4E, 4F). Loss of d/atp-5 subunit also suppressed the mitochondrial swelling and fragmentation observed after OSCP/atp-3 RNAi (Figure 4G, 4H). Thus, inhibition of key subunits of F-ATP synthase as well as pharmacological inhibition of the mPTP by CysA rescues the mPTP/pdvUPR\textsuperscript{mt} toxicity and rescues its effects on aging.

To verify that the use of dual-RNAi did not interfere with knockdown of OSCP/atp-3, we assessed an endogenously-expressing patp-3::ATP-3::GFP translational reporter generated via CRISPR-Cas-9 (Figure S4A). We observed efficient knockdown of OSCP/atp-3 via RNAi in the presence of either b/atp-2 or d/atp-5 RNAi, two subunits that suppress the pdvUPR\textsuperscript{mt}, demonstrate that the inhibition of the pdvUPR\textsuperscript{mt} is not due to ineffective RNAi of OSCP/atp-3 but rather the functional consequences of removal of additional ATP synthase subunits. To examine the effects of dual-RNAi another way, we examined how loss of ATP synthase subunits impacted the pdvUPR\textsuperscript{mt} after a/atp-1 RNAi, which induced the second most robust pdvUPR\textsuperscript{mt} (Figure 2E, 2G). Remarkably, we see the same pattern of pdvUPR\textsuperscript{mt} activation and inhibition as with loss of OSCP/atp-3: loss of the ATPase subunit b/atp-2 and peripheral stalk subunit d/atp-5 suppressed the a/atp-1 RNAi-mediated pdvUPR\textsuperscript{mt} while loss of NDUFS3/nuo-2 and COX5B/cco-1 had no effect (Figure S4B). Importantly, immunoblots against a/atp-1 showed similar protein knockdown under all conditions (Figure S4B), confirming that dual-RNAi is an effective method to assess the structural components of the F-ATP synthase.

**Longevity Mutants Display Exceptional Mitochondria Partially via FOXO3a/daf-16**

Given that the mPTP can accelerate aging, we tested whether long-lived worms, glp-1(e2141) (Notch-1 germline signaling), daf-2(e1370) (insulin/IGF-1-like signaling), and eat-2(ad465) (caloric restricting), were protected from the mPTP/pdvUPR\textsuperscript{mt}. We also included fem-1(hc17) (sperm deficient) mutants, which we identified as long-lived (Figure S5A, S5B). RNAi of OSCP/atp-3 induced little to no pdvUPR\textsuperscript{mt} in glp-1(e2141), fem-1(hc17), and daf-2(e1370) backgrounds, while a moderate pdvUPR\textsuperscript{mt} was induced in eat-2(ad465) (Figure 5A, 5B). For simplicity, we next focus on the germline mutants, fem-1(hc17) and glp-1(e2141). We found that fem-1(hc17) and glp-1(e2141) were able to induce a robust UPR\textsuperscript{mt} similar to control when RNAi was initiated during development (Figure S5C). The mutants also initiated a normal UPR\textsuperscript{ER} and heat shock response (HSR) during adulthood (Figures S5D-F). Thus, mutations that confer longevity specifically repress the pdvUPR\textsuperscript{mt}.

We next sought to determine the mechanism of suppression of the pdvUPR\textsuperscript{mt} by mutations conferring longevity. The longevity of glp-1 and daf-2 has previously been shown to be dependent on the transcription factor FOXO3A/daf-16 (Hsin and Kenyon, 1999; Kenyon et al., 1993) and we also found that the longevity of fem-1(hc17) to be dependent on FOXO3a/daf-16 (Figure S5A, S5B). On the other hand, eat-2 has previously been found to be dependent on FOXA/pha-4 (Panowski et al., 2007). Thus, we tested whether FOXO3a/daf-16 was required for the suppression of the pdvUPR\textsuperscript{mt} and found that introducing the mutation, daf-16(mu86), partially restored the pdvUPR\textsuperscript{mt} in glp-1(e2141) mutants and completely restored it in fem-1(hc17) mutants (Figure 5C). Previous studies have suggested that an increase in mitochondrial biogenesis may protect against opening of the mPTP (Goncalves et al., 2016). To determine if fem-1(hc17) and glp-1(e2141) exhibited enhanced mitochondrial content via FOXO3a/daf-16, we examined the mitochondria in the intestines where the UPR\textsuperscript{mt} is most robustly activated. We
Figure 5: Long-lived Mutants Suppress the Post-Developmental UPR\textsuperscript{mt} Via FOXO3a/\textit{daf-16}

(A, B) Long-lived germline mutants with a \textit{phsp-6::GFP} reporter strain background do not initiate a robust pdvUPR\textsuperscript{mt} due to loss of OSCP/\textit{atp-3} via RNAi compared to \textit{phsp-6::GFP} controls. RNAi was administered for 48 hours beginning at young adulthood. Temperature sensitive mutants \textit{glp-1(e2141)} and \textit{fem-1(hc17)} and controls were developed at 25°C to initiate sterility and shifted to 20°C and RNAi plates as young adults. Data are the mean ± SEM of ≤15 animals combined from three biological experiments. ** p ≤ .001, ***p ≤ .0001 by Student’s t-test. CV, control vector RNAi.

(C) The pdvUPR\textsuperscript{mt} is fully restored in \textit{fem-1(hc17); daf-16(mu86)} mutants and partially restored in \textit{glp-1(e2141); daf-16(mu86)} mutants. Temperature sensitive mutants and controls were developed at 25°C to initiate sterility and shifted to 20°C and RNAi as young adults. Data are the mean ± SEM of ≤15 animals combined from three biological experiments. *p ≤ .01 by Student’s t-test. CV, control vector RNAi.

(D) Germline mutants \textit{fem-1(hc17)} and \textit{glp-1(e2141)} with a \textit{pges-1::GFP\textsuperscript{mt}} reporter display enhanced intestinal mitochondrial content. Germline mutants with a \textit{daf-16(mu86)} mutation no longer display enhanced intestinal mitochondria. Temperature sensitive mutants and controls were developed at 25°C to initiate sterility and shifted to 20°C and RNAi as young adults. Data are the mean ± SEM of ≤15 animals combined from three biological experiments. ** p ≤ .001 by Student’s t-test.

(E) Model: Loss of OSCP/\textit{atp-3} initiates the mPTP, leading to activation of a maladaptive UPR\textsuperscript{mt}, contributing to aging and age-related diseases. FOXO3a/\textit{daf-16} protects the mitochondria and suppresses mitochondrial dysfunction.
observed significantly higher mitochondrial mass in the intestines of glp-1(e2141) and fem-1(hc17), which was dependent on FOXO3a/daft-16. To determine if the increased mitochondrial content may be due to a block in mitophagy or autophagy, we examined mitochondrial content after RNAi of pdr-1 (parkin homology) and hhl-30 (TFEB homolog). RNAi of pdr-1 or hhl-30 significantly increased the mitochondrial content in glp-1(e2141) and fem-1(hc17) mutants, suggesting that a block in mitophagy/autophagy was not the responsible mechanism. Overall, these findings suggest that glp-1(e2141) and fem-1(hc17) mutants are protected from initiating a maladaptive mPTP/pdvUPR, possibly due to enhanced mitochondrial biogenesis via FOXO3a/daft-16.

DISCUSSION

While loss of the ATP synthase subunit OSCP/atp-3 during development leads to lasting activation of the UPR and is associated with longevity (Dillin et al., 2002; Rea et al., 2007), we have discovered that loss of this subunit during adulthood induces the mPTP and activates a reversible and atfs-1-dependent UPR (pdvUPR). Furthermore, we observed that activation of the UPR during adulthood helps drive aging. Suppression of the mPTP/UPR via genetic or pharmacological interventions is protective. Similarly, long-lived mutants from various longevity paradigms dependent on FOXO3a/daft-16 are protected from the mPTP/UPR during adulthood. The fact that loss of other OXPHOS subunits or loss of membrane potential via FCCP are not capable of inducing the UPR during adulthood suggests a specific activation of the UPR due to the mPTP. In contrast, it does not appear that the UPR is activated in response to the mPTP during development. Thus, loss of OXPHOS subunits can have drastically different effects depending on the life stage or cellular milieu of the organism and suppression of the UPR can be beneficial.

While the mPTP detrimental role in health is well established, considerable evidence suggests that the UPR contributes to health and longevity (Houtkooper et al., 2013; Merkwirth et al., 2016; Mouchiroud et al., 2013; Sorrentino et al., 2017). Activating the UPR in neurons can activate protective cell non-autonomous signals and also epigenetically rewire C. elegans to live longer (Durieux et al., 2011; Merkwirth et al., 2016; Tian et al., 2016; Zhang et al., 2018). NAD+ boosters appear to activate the UPR and contribute to longevity and ameliorate AD (Mouchiroud et al., 2013; Sorrentino et al., 2017). However, an unbiased screen that identified activators of the UPR found no correlation between UPR activation and longevity (Bennett et al., 2014) and constitutive activation of the UPR in dopaminergic neurons lead to increased neurodegeneration (Martinez et al., 2017). Activation of the UPR in our setting is distinct in that it is specifically linked to activation of the mPTP. Thus, it is possible that preemptively boosting the UPR may ward of aging and disease while activation in a diseased setting may exacerbate conditions, akin to instances of inflammation in disease.

We propose a model in which loss of OSCP/atp-3 induces a conformational change in F-ATP synthase that leads to pore formation and activation of the UPR during adulthood but not during development (Figure 5D). Previous reports have shown that loss of OSCP increases susceptibility to Ca2+-induced mPTP formation and that key residues within the OSCP are required to suppress the mPTP during conditions of low pH (Antoniel et al., 2018; Giorgio et al., 2013), suggesting that a functional and intact OSCP protects against pore formation. OSCP
levels have also been shown to decrease with age while concomitantly increasing its binding to amyloid β, suggesting that loss of OSCP destabilizes the remaining ATP synthase to increase pore formation (Beck et al., 2016b). However, it has also been shown that OSCP provides the binding site for cyclophilin D and thus it has been proposed to be critical in the formation of the mPTP (Giorgio et al., 2017). However, these findings have been contested as knock out studies of OSCP show that cyclophilin D still binds to ATP synthase and EM studies have identified a cyclophilin D structure positioned on peripheral stalk subunits lateral to the OSCP (Daum et al., 2013; He et al., 2017a). Thus, our findings support a model in which loss of OSCP/atp-3 induces a conformation change that is favorable for cyclophilin binding to the remaining peripheral stalk proteins or to sites independent of the F-ATP synthase, leading to destabilization of ATP synthase, pore formation with a loss of membrane potential, and subsequent activation of the pdvUPRmt.

The loss of OSCP/atp-3 may also be more impactful than other subunits. Unlike other ATP synthase subunits, OSCP/atp-3 is prominently accessible in the mitochondrial matrix and has a diverse set of binding partners, such as estradiol and p53, which can modulate ATP production (Bergeaud et al., 2013; Zheng and Ramirez, 1999). Its loss may induce a strong protein misfolding cascade, reminiscent of the He and Lemasters mPTP model first proposed in 2002 (He and Lemasters, 2002). In this model, it was proposed that exposure to activators of the mPTP, such as oxidants, causes protein misfolding of integral membrane proteins, thereby recruiting chaperones such as cyclophilin D to repair. If the protein misfolding is unable to be repaired, then Ca²⁺, along with cyclophilin D, could catalyze pore formation in a CysA-dependent manner. Indeed, both paraquat and manganese have been shown to induce both the UPRmt and the mPTP in separate studies (Angeli et al., 2014; Costantini et al., 1995; Nargund et al., 2012; Rao and Norenberg, 2004). However, more research is needed to clarify the relationship between protein misfolding and the mPTP.

Despite mounting evidence implicating F-ATP synthase as the pore-forming component of the mPTP, this remains controversial. The Walker lab has systematically deleted nearly every subunit of ATP synthase in a cell model and showed that a pore still forms (Carroll et al., 2019; He et al., 2017a; He et al., 2017b), leading some groups to suggest that in the absence of an intact F-ATP synthase, smaller low conductance CysA-dependent pores distinct from the mPTP still form (Neginskaya et al., 2019). Other groups have proposed that the mPTP can be mediated by ANT, as well as a cyclophilin D binding structure, which supports a “multi-pore model”, which would explain why deletion of putative pore components may still yield pore formation (Carraro et al., 2020; Karch et al., 2019).

AD and PD both display evidence of the mPTP and in some instances, elevated UPRmt profiles have also been observed (Beck et al., 2016a; Beck et al., 2016b; Ludtmann et al., 2018; Perez et al., 2018; Sorrentino et al., 2017), but the relationship between these two mitochondrial processes have not been fully explored. Ischemic reperfusion directly causes the mPTP but little is known about the UPRmt under these conditions. Establishing a clearer understanding of the relationship between the UPRmt and the mPTP in these disease states could result in the development of new therapeutics for these and related disorders.
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AUTHOR CONTRIBUTIONS
Conceptualization, S.A., J.K.A., and G.J.L.; Methodology, S.A., J.K.A., G.J.L., and A.C.F.; Investigation, S.A., A.C.F., T.H.P., M.C., D.B. and A.A.S., Validation, S.A. and A.C.F.; Resources, A.A.G.; Writing –Original Draft, S.A., J.K.A., and G.J.L.; Visualization, S.A., Supervision, S.A., J.K.A, G.L., Funding Acquisition, J.K.A. and G.J.L.

DECLARATION OF INTEREST
G.J.L. is cofounder of GeroState alpha and declares no financial interest related to this work. A.A.G. is founder of Image Analyst MKII Software and declares a financial interest.

MATERIAL AND METHODS

KEY RESOURCES

**Chemicals**
- Tetramethylrhodamine methyl ester (perchlorate) (TMRM)
  - Cayman Chemical
  - Item # 21437
- Cyclosporin A
  - Cayman Chemical
  - Item # 12088
- FK506 (Tacrolimus)
  - Cayman Chemical
  - Item # 10007965
- FCCP (Trifluoromethoxy carbonylcyanide phenylhydrazone)
  - Cayman Chemical
  - Item # 15218

**Antibodies**
- Mouse monoclonal anti-GRP 75 (D9)
  - Santa Cruz Biotechnology
  - Cat# sc-133137
- Mouse monoclonal anti-β Tubulin Antibody (D-10)
  - Santa Cruz Biotechnology
  - Cat# sc-5274
- Mouse monoclonal anti-β-Actin (8H10D10)
  - Cell Signaling Technology
  - Cat# 3700
- Mouse monoclonal anti- ATP5A1 Monoclonal Antibody (15H4C4)
  - Invitrogen
  - Cat# 43-9800
- Mouse monoclonal anti-ATP Synthase beta Monoclonal Antibody (3D5AB1)
  - Invitrogen
  - Cat# A-21351
- Mouse monoclonal anti-NDUF5 Monoclonal Antibody (17D95)
  - Invitrogen
  - Cat# 43-9200

**Experimental Models: Organisms/Strains**
- *C. elegans*: N2 Bristol
  - Caenorhabditis Genetics Center
  - RRID:WB-STRAIN:N2_(anscestral)
- *C. elegans*: SJ4100 (zcIs13[hsp-6p::GFP])
  - Caenorhabditis Genetics Center
  - RRID:WB-STRAIN:SJ4100
- *C. elegans*: SJ4058 (zcIs9[hsp-60p::GFP + lin-15(+)])
  - Caenorhabditis Genetics Center
  - RRID:WB-STRAIN:SJ4058
CONTACT FOR REAGENT AND RESOURCE SHARING
Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contacts, Gordon Lithgow, glithgow@buckinstitute.org, Julie Andersen, jandersen@buckinstitute.org, and Suzanne Angeli, sangeli@buckinstitute.org

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Nematode Strains
The following strain was obtained from the National BioResource Project: atf-1(tm4525). The following strains were obtained from the Caenorhabditis Genetic Center (CGC): SJ4100 (zcls13[hsp-6p::GFP]), SJ4058 (zcls9[hsp-60p::GFP]), SJ4005 (zcls4[hsp-4p::GFP]), CL2070 (dvl[hsp-16.2p::GFP]), SJ4143 (zcls17 [ges-1p::GFP(mt)]), KWN190 ([pha-1(e2123) III]; [him-5(e1490) V]; rnyEx109), CF1880 [daf-16(mu86)], CL2070 (dvl70[hsp-16.2p::GFP], [rol-6(su1006)], SJ4005 (zcls4[hsp-4p::GFP]), DA1116 [eat-2(ad1116)] and N2 Bristol (wild-type). The following strain, SJ4100 (zcls13[hsp-6p::GFP]) was backcrossed 6 times to wild-type N2 to generate GL347, which was used for all hsp-6p::GFP experiments. BA17 [fem-1(hc17)] and CB4037 [glp-1(e2141)] were back-crossed 3 times with wild-type N2 to create GL355 and GL345 respectively and used for the strains generated in this study. The following crossed strains were created in the Lithgow Lab for this study: GL364 [fem-1(hc17)], (zcls13[hsp-6p::GFP]); GL346 [glp-1(e2141)], (zcls13[hsp-6p::GFP]); GL380 [daf-2(el370)], (zcls13[hsp-6p::GFP]); GL384 [eat-2(ad1116)], (zcls13[hsp-6p::GFP]); GL361 [fem-1(hc17)], (zcls17 [ges-1p::GFP(mt)]); GL362 [glp-1(e2141)], (zcls17 [ges-1p::GFP(mt)]); GL365 [glp-1(e2141)], (zcls13[hsp-6p::GFP], [daf-16(mu86)]); GL370 [fem-1(hc17)], [daf-16(mu86)]; GL396 [glp-1(e2141)], [daf-16(mu86)]; GL361 [fem-1(hc17)], (zcls17 [ges-1p::GFP(mt)]); GL351 [daf-16(mu86)], (zcls17 [ges-1p::GFP(mt)]); GL389 [fem-1(hc17)], (zcls17 [ges-1p::GFP(mt)]); [daf-16(mu86)]); GL358 [glp-1(e2141)] (zcls17 [ges-1p::GFP(mt)]); [daf-16(mu86)]; GL376 [fem-1(hc17)], (zcls4[hsp-4p::GFP]); GL336 [glp-1(e2141)], (zcls4[hsp-4p::GFP]); GL397 [fem-1(hc17)], (dvl70[hsp-16.2p::GFP], [rol-6(su1006)]); GL342 [glp-1(e2141)], (dvl70[hsp-16.2p::GFP], [rol-6(su1006)]).
CRISPR-Cas9 strain
The translational atp-3::ATP-3::GFP was generated by SunyBiotech (http://www.sunybiotech.com/) using CRISPR-Cas9. The strain PHX1826 atp-3(syb1826)/hT2[bli-4(e937) let-7(q782) qIs48] was verified by PCR sequencing. The sgRNA target site used is as follows:
Sg1: CCCTTGCCACCGCCATCTAAatt
Sg2: CCGCATCTAAatttttcccaaa.

Nematode and Bacterial Culture Conditions
Nematodes were maintained on nematode growth medium (NGM) plates. NGM plates were seeded with *E. coli* OP50 obtained from CGC that was grown in LB broth at 37˚C for 18 hours shaking at 225 rpm. Plates with bacteria were dried for 48 hours before use.

For RNAi experiments, *E. coli* HT115 (DE3) bacteria obtained from the Ahringer and Vidal RNAi Library was used (Kamath et al., 2003; Rual et al., 2004). All RNAi clones were verified via sequencing (Eurofins™). RNAi plates were prepared by cooling NGM to 55˚C and supplementing with a final concentration of 50ug/ml carbenicillin and 1mM Isopropyl β-d-1-thiogalactopyranoside (IPTG). RNAi bacteria was inoculated with one colony of RNAi bacteria into LB with 50ug/ml carbenicillin and was grown shaking overnight for 18 hours at 37˚C at 225 rpm.

Post-Developmental Timing
To achieve synchronous nematode populations, day 1 adult nematodes were allowed to lay eggs for 2 hours on seeded NGM plates. For convenience, nematodes were developed at 25˚C on *E. coli* OP50 until worms were visibly past the L4 stage (loss of crescent) but not yet gravid, approximately 45 hours for wild-type (although the time it takes for the worms to reach the young adult stage varies by strain). Nematodes were shifted to 20˚C once they reached adulthood.

METHOD DETAILS

RNA Interference (RNAi) Treatment
For developmental treatments, synchronized eggs were moved onto plates seeded with RNAi bacteria and developed at 20˚C for 72 hours. Nematodes were then either collected for analysis or for lifespans, remained on RNAi bacteria for the remainder of their life for survival analysis. For post-developmental treatments, synchronized eggs were developed on plates with *E. coli* OP50 at 25˚C until the young adult stage and then transferred to RNAi plates at 20˚C. Nematodes were collected after 48 hours for analysis or for lifespans, moved onto *E. coli* OP50 for the remainder of their life.

Quantitative RT-PCR
Approximately 300 adult nematodes were collected; nematode pellets were resuspended in 300uL RNA Lysis Buffer and frozen. Pellets was thawed, vortexed, and snap frozen three times. Zymo Research Quick-RNA MiniPrep kit was used to extract RNA. The primers used for qPCR are as follows:
act-1, Forward: ACGACGAGTCCGGGCCCCATCC
act-1, Reverse: GAAAGCTGGTGTTGACGATGGTT
**atp-1**, Forward: GAAGGACAAATCTCCCCACA
**atp-2**, Reverse: CGCCACATTCTTCTTTTTC
**atp-3**, Forward: GCCTAATTTCTTAATTTTGCAGGT
**atp-3**, Reverse: GAACCTGAATCGGGGTCTTC
**atp-4**, Forward: AATATGTTGCCTCCTTCTGAT
**atp-4**, Reverse: GGAACAAAAACGTTCATTCG
**atp-5**, Forward: TCTTCGACGTGCCGACAA
**atp-5**, Reverse: AAATGGTAGGAGAGCGATAAGG
**nuo-2**, Forward: TGAAGTTGCTGAGCCAACAC
**nuo-2**, Reverse: TCCACACTAACAGAAAATGAGTCT
**cco-1**, Forward: TTTCGGCTATTGTTCGATT
**cco-1**, Reverse: GCCGTCTTAGCAAGTGCATC

**Lifespans**
Day 1 adult nematodes were allowed to lay eggs for 2 hours on seeded NGM plates to obtain a synchronous aging population. 5-fluoro-2-deoxyuridine (FUdR) was omitted from plates due to its potentially confounding effects (Angeli et al., 2013). Worms are transferred to freshly seeded bacterial plates every day for the first 7 days of adulthood and then as needed afterwards. Worms were scored as dead when they failed to respond to gentle prodding with a platinum wire. Worms that experienced matricide or bagging were censored.

**Tetramethylrhodamine Methyl Ester (TMRM) Staining**
Plates were prepared by spotting seeded NGM plates with a TMRM solution diluted in water to a final concentration of 0.1 µM in the plates. Water was used as a solvent control. Plates were allowed to dry for 24 hours before use. For developmental experiments, synchronized eggs were placed on plates for 72 hours and then nematodes were collected for analysis. For post-developmental experiments, young adult nematodes were placed on plates for 48 hours and then collected for analysis.

**Cyclosporine A (CysA) Treatment**
Plates were prepared by spotting seeded NGM plates with a CysA (stock solution in DMSO) solution diluted in 100% ethanol. Comparable amounts of DMSO and ethanol were used as solvent controls. Plates were allowed to dry for 24 hours before use. For developmental experiments, synchronized eggs were placed on plates for 72 hours and then nematodes were collected for analysis. For post-developmental experiments, young adult nematodes were placed on plates for 48 hours and then collected for analysis; for lifespans, worms were moved to regular NGM plates for the remainder of their life after 48 hours on drug-treated RNAi bacteria.

**Microscopy**
Worms were anesthetized with 2mM levamisole and mounted on 2% agarose pads on glass slides. Fluorescence micrographs of GFP and TMRM were taken using a Zeiss Imager A2 at 5x magnification with 600ms exposure using the ZEN software. GFP expression was enhanced using the brightness/contrast tool in Photoshop. The same parameters were used for all images.
Confocal micrographs of mitochondrially targeted GFP and the cytosolic calcium sensor D3cpv (Zhang et al., 2016) were taken using a Zeiss LSM780 laser scanning confocal microscope using a 63× Plan Apochromat NA1.4. To visualize outlines of mitochondria, Image Analyst MKII (Image Analyst Software, Novato, CA) was used. Selected rectangular regions of interest (ROI) from the worm intestine were segmented and converted to outlines by a modification of the “Segment mitochondria” pipeline. Emission ratio images of D3cpv were excited at 440nm and captured at 450-490nm and 520-560nm and analyzed in Image Analyst MKII. Images were Wiener filtered and the ratio of the 540nm over the 470nm channel, indicative of cytosolic calcium concentration was calculated, and showed in pseudo-color coding. Emission ratios were determined in ROIs in the posterior intestine by the Plot Ratio function.

**Western Blot**
Approximately 30 to 50 adult worms were collected in S-basal buffer. Supernatant was removed and nematodes were flash-frozen. Worms pellets were resuspended in 2% SDS sample buffer with 2.5% β–mercaptoethanol and samples were boiled for 10 minutes. Samples were subjected to SDS-PAGE using 4-12% SDS gels (Invitrogen™) and transferred to Immun-Blot PVDF Membrane (BIO-RAD™) using BIO-RAD™ western blot Criterion apparatus. Membranes were blocked with 5% non-fat dry milk blocking solution; concentrations for antibodies were 1:1000 for primary antibodies and 1:2000 for secondary antibodies.

**QUANTIFICATION, STATISTICS, AND PREDICTION**

**Statistics**
Significance between control and experimental groups was determined by using two-tailed Student’s t test. Asterisks denote corresponding statistical significance *p < 0.05; **p < 0.01; ***p < 0.0001. Error bars were generated using the standard error of the mean (SEM), typically from three pooled biological replicates. Graphpad Prism 7™ was used to plot survival curves. Log Rank (Mantel Cox) test in Prism™ was used to determine significance between the control and experimental group.

**GFP Quantification**
GFP intensity of worms was quantified using Image J 1.52A. The ‘integrated density’ of GFP expression and length of worms was measured using Image J tools. Integrated Density value was normalized by number of worms and average length of worms. The final value is in arbitrary units.

**Mitochondrial Targeting Prediction**
Mitochondrial presequences were predicted using the MitoFates tool (Fukasawa et al., 2015).

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SUPPLEMENTAL FIGURES
Figure S1: RNAi Efficiency and FCCP in Young Adult *C. elegans*

(A-E) qRTPCR from N2 worms after RNAi of subunits from complex IV and V. RNAi was administered for 48 hours beginning at young adulthood. Data are the mean ± SD ≤ 150 animals combined from two experiments. **p ≤ 0.01, ***p ≤ .0001 by Student’s *t*-test.

(F-H) Immunoblots from N2 worms after RNAi of subunits from complex I and V. RNAi was administered for 48 hours beginning at young adulthood. Representative immunoblots from two biological experiments. Actin was used as a loading control.

(I) Dose-response treatment with the mitochondrial uncoupler, FCCP, did not significantly induce the *p*hs*6*::GFP reporter. FCCP was administered for 24-48 hours beginning at young adulthood. Data are the mean ± SEM of ≤ 15 animals from three biological experiments. n. s., not significant by Student’s *t*-test.
Figure S2: Loss of the ATP Synthase Subunit, OSCP/atp-3, Induces a Specific, Temporally Confined, and Reversible Post-Developmental UPR\textsubscript{mt}

(A) qRTPCR of hsp-6 and hsp-60 from N2 worms on post-developmental CV or atp-3 RNAi. Data are the mean ± SEM of ≤150 animals combined from four biological experiments. **p ≤ 0.01, n.s. not significant by Student’s t-test. CV, control vector RNAi.

(B) Western blot of mtHSP70/HSP-6 and tubulin (loading control) from worms on post-developmental CV or atp-3 RNAi. CV, control vector RNAi. Representative blot from three biological experiments.

(C) Post-developmental RNAi of atp-3 mildly induced the mitochondrial p\textsubscript{hsp-60}::GFP reporter but not the UPR\textsubscript{ER} p\textsubscript{hsp-4}::GFP or HSR p\textsubscript{hsp-16.2}::GFP reporters. Left panel, fluorescent channel; right panel, bright-field channel. CV, control vector RNAi.

(D) Quantification of p\textsubscript{hsp-6}::GFP and p\textsubscript{hsp-60}::GFP reporters from (A). Data are the mean ± SEM of ≤15 animals combined from three biological experiments. **p ≤ 0.01, *p ≤ 0.05 by Student’s t-test.

(E) Post-developmental RNAi of atp-3 or cco-1 RNAi that was initiated at L4, young adult, and adult stages using the p\textsubscript{hsp-6}::GFP reporter at 25°C. GFP expression was assessed 48 hours after RNAi initiation. Representative pictures from two biological experiments.

(F) The p\textsubscript{hsp-6}::GFP reporter worms were treated with developmental or post-developmental atp-3 RNAi. Worms were removed from HT115 RNAi bacteria and placed onto OP50 bacteria and GFP expression was assessed at indicated time-points. Representative pictures from two biological experiments.

(G) Developmental RNAi of all ATP synthase subunits tested induced the p\textsubscript{hsp-6}::GFP reporter. For developmental treatment, worms were exposed to RNAi beginning from eggs for 72 hours.

(H) Quantification of GFP intensity from (G). Data are the mean ± SEM of ≤15 animals combined from three biological experiments. ***p ≤ 0.0001 by Student’s t-test. dv, development. See Table S3 for additional details.
Figure S3: The Post-developmental but not Developmental UPR\textsuperscript{mt} is Modulated by the mPTP

(A) Dose response of CysA administered during development or post-development on atp-3 RNAi-treated phsp-6::GFP reporter worms.

(B) Dose response of FK506 administered during post-development on atp-3 RNAi-treated phsp-6::GFP reporter worms.

(C) Concomitant developmental RNAi of atp-3 and individual ATP synthase subunits induced the phsp-6::GFP reporter. For developmental treatment, worms were exposed to RNAi beginning from eggs for 72 hours.

(D) Concomitant developmental or post-developmental RNAi of atp-3 and either cco-1 or nuo-2 induced the phsp-6::GFP reporter.

(E, F) Quantification of GFP intensity from (C) and (D). Data are the mean ± SEM of ≤ 15 animals combined from three biological experiments. (E) No significant differences from CV + atp-3 condition. ** p ≤ .0001, ***p ≤ .0001 by Student’s t-test. dvUPR\textsuperscript{mt}, development; pdv, post-development.
Figure S4: Effects of Loss of OXPHOS Subunits on the dvUPR\textsuperscript{mt} and pdvUPR\textsuperscript{mt}

(A) Dual-RNAi effectively knocks down ATP-3 protein synthesis as measured by the translational patp-3::ATP-3::GFP reporter, i. 10X photomicrographs of homozygous and heterozygous (bright pharyngeal GFP expression) worms, ii. 40X photomicrographs of head region of control homozygous worms, iii. 40X photomicrographs of mid-region of control homozygous worms, iv. 40X photomicrographs of tail region of control homozygous worms.

(B) RNAi of atp-1 induces a post-developmental UPR\textsuperscript{mt} similar to atp-3 in the p\textit{hsp-6}::GFP reporter. Representative results from two biological experiments. Immunoblot of ATP-1 from p\textit{hsp-6}::GFP reporter worms after dual-RNAi treatment. Representative blots from two biological experiments.
Figure S5: Characterization of germline mutant longevity, stress response, and mitophagy/autophagy processes.

(A) Germline mutant fem-1(hc17) is long-lived, which is abolished in daf-16(mu86) mutant background. Worms were developed at 25°C and shifted to 20°C beginning at young adulthood.

(B) Germline mutant fem-1(hc17) is long-lived, which is abolished after treatment with daf-16 RNAi. Worms were developed at 25°C and shifted to 20°C beginning at young adulthood.

(C) Germline mutants fem-1(hc17) and glp-1(e2141) with a p[hsp-6]::GFP reporter strain background induce robust UPRmt when OSCP/atp-3 RNAi is administered during development. Worms were developed at 25°C and exposed to RNAi beginning from the egg stage and collected for microscopy after approximately 48 hours.

(D, E) Germline mutants fem-1(hc17) and glp-1(e2141) with a p[hsp-4]::GFP reporter strain background induce robust UPRER when exposed to 4ug/ml tunicamycin. Worms were developed at 25°C and exposed to tunicamycin as young adults and collected for microscopy after approximately 24 hours. Tunicamycin was spotted onto seeded NGM plates and water was used as the solvent control.

(F) Germline mutants fem-1(hc17) and glp-1(e2141) with a p[hsp-16.2]::GFP reporter strain background induce a robust heat-shock response when to 33°C for 2 hours. Worms were developed at 25°C and exposed to heat as young adults following immediate collection for microscopy.

(G) Germline mutants fem-1(hc17) and glp-1(e2141) with a pges-1::GFPmt reporter strain background exhibit higher intestinal mitochondrial mass after RNAi with pdr-1 and hlh-30. Worms were developed at 25°C and shifted to 20°C and exposed to RNAi as young adults; worms were collected for microscopy after approximately 48 hours.
### Supplemental Table 1

| gene | dvUPR<sub>mt</sub> | pdvUPR<sub>mt</sub> | pdvUPR<sub>mt</sub> fold change | p-value |
|------|---------------------|---------------------|----------------------------------|---------|
| **Complex I** |                      |                     |                                 |         |
| *nuo-1* | yes                 | yes                 | 2.6                              | 0.0002  |
| *nuo-2* | yes                 | yes                 | 2.1                              | 0.0155  |
| **Complex III** |                    |                     |                                 |         |
| *cyc-1* | yes                 | yes                 | 2.1                              | < 0.0001 |
| **Complex IV** |                    |                     |                                 |         |
| *cco-1* | yes                 | no                  | 1.9                              | 0.0511  |
| **Complex V** |                      |                     |                                 |         |
| *atp-2* | yes                 | yes                 | 2.7                              | 0.0029  |
| *atp-3* | yes                 | yes                 | 71.6                             | < 0.0001 |
| **Other** |                      |                     |                                 |         |
| *clk-1* | yes                 | no                  | -1.5                             | 0.0064  |
| *mrps-5* | yes                 | no                  | 1.0                              | 0.9671  |
| *tomm-22* | yes                 | no                  | -1.8                             | 0.0004  |

**Supplemental Table 1**: List of genes tested for whether they induce a developmental UPR<sub>mt</sub> (dvUPR<sub>mt</sub>) or post-developmental UPR<sub>mt</sub> (pdvUPR<sub>mt</sub>).
| Complex V     | gene     | subunit | dvUPR<sup>mt</sup> | pdvUPR<sup>mt</sup> | fold change | p-value |
|--------------|----------|---------|--------------------|---------------------|-------------|---------|
| F1 ATPase    | atp-1    | α       | yes                | yes                 | 19.7        | 0.0004  |
| F1 ATPase    | atp-2    | β       | yes                | yes                 | 2.7         | 0.0029  |
| F1 ATPase    | F58F12.1 | δ       | yes                | yes                 | 1.5         | 0.0410  |
| F1 ATPase    | hpo-18   | ε       | yes                | yes                 | 12.3        | < 0.0001|
| F1 peripheral stalk | atp-3 | OSCP   | yes                | yes                 | 55.4        | < 0.0001|
| F1 peripheral stalk | asb-2 | b       | yes                | no                  | 1.9         | 0.0797  |
| F1 peripheral stalk | atp-5 | d       | yes                | no                  | 1.1         | 0.6388  |
| F1 peripheral stalk | atp-4 | F6      | yes                | no                  | 1.2         | 0.2206  |
| Fo rotor     | Y82E9BR.3 | c     | yes                | yes                 | 3.8         | 0.0003  |
| Fo supernumerary | R04F11.2 | e     | yes                | no                  | 1.0         | 0.9604  |
| Fo supernumerary | R53.4 | f     | yes                | no                  | 1.1         | 0.7152  |

**Supplemental Table 2**: Summary of the effects of RNAi of ATP synthase subunits on the developmental UPR<sup>mt</sup> (dvUPR<sup>mt</sup>) or post-developmental UPR<sup>mt</sup> (pdvUPR<sup>mt</sup>).
| gene   | MTS strength |
|--------|--------------|
| cyn-1  | 0.941        |
| cyn-2  | 0.0          |
| cyn-3  | 0.0          |
| cyn-4  | 0.0          |
| cyn-5  | 0.0          |
| cyn-6  | 0.0          |
| cyn-7  | 0.0          |
| cyn-8  | 0.0          |
| cyn-9  | 0.0          |
| cyn-10 | 0.0          |
| cyn-11 | 0.0          |
| cyn-12 | 0.0          |
| cyn-13 | 0.0          |
| cyn-14 | 0.0          |
| cyn-15 | 0.0          |
| cyn-16 | 0.0          |
| cyn-17 | 0.154        |
| atfs-1 | 0.117        |

Supplemental Table 3: List of *C. elegans* cyclophilins and their predicted mitochondrial localization using the MitoFates mitochondrial targeting sequence (MTS) prediction tool.
|                | Strain          | Median lifespan (days) | Deaths/Events | Censored Animals | Deaths/Censored | p-value |
|----------------|-----------------|------------------------|---------------|------------------|-----------------|---------|
| **Development**| N2; cv          | 15                     | 48            | 20               | 48/20           |         |
|                | N2; atp-3       | 26.5                   | 34            | 30               | 34/30           | *** < 0.0001 |
|                |                 |                        |               |                  |                 |         |
|                | N2; cv          | 15                     | 161           | 38               | 161/38          |         |
| **Post-development** | N2; atp-3     | 10                     | 181           | 20               | 181/20          | *** < 0.0001 |
|                | atfs-1; cv      | 12                     | 117           | 63               | 117/63          |         |
|                | atfs-1; atp-3   | 9                      | 157           | 31               | 157/31          | *** < 0.0001 |
|                | N2; cv          | 17                     | 116           | 27               | 116/27          |         |
|                | N2; cyc-1       | 17                     | 137           | 22               | 137/22          | 0.6097  |
|                | N2; cco-1       | 17                     | 139           | 6                | 139/6           | 0.2826  |
|                | N2; atp-2       | 15                     | 131           | 22               | 131/22          | ** 0.0067 |
|                | N2; nuo-2       | 17                     | 128           | 27               | 128/27          | 0.5893  |
| **Post-development** | N2; cv        | 15                     | 124           | 33               | 124/33          |         |
|                | N2; atp-3       | 13                     | 130           | 16               | 130/16          | *** < 0.0001 |
|                | N2; cv 15uM CysA| 17                     | 85            | 24               | 85/24           |         |
|                | N2; atp-3 15uM CysA | 17             | 112           | 35               | 112/35          | 0.6084  |

Supplemental Table 4: Summary of lifespans.