Rilmenidine mimics caloric restriction via the nischarin II-imidazoline receptor to extend lifespan in *C. elegans*

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**Keywords:** aging, dietary restriction, drug repurposing, longevity, nischarin receptor, autophagy, mTOR
Abstract

Caloric restriction increases lifespan across species and has health benefits in humans. Because complying with a low-calorie diet is challenging, here we investigated pharmacological interventions mimicking the benefits of caloric restriction. Searching for compounds that elicit a similar gene expression signature to caloric restriction, we identified rilmenidine, an I1-imidazoline receptor agonist and prescription medication for the treatment of hypertension. We then show that treating *C. elegans* with rilmenidine at young and older ages increases lifespan. We also demonstrate that the stress-resilience, healthspan, and lifespan benefits upon rilmenidine treatment in worms are mediated by the I1-imidazoline receptor *nish-1*, implicating this receptor as a potential longevity target. Furthermore, we show that rilmenidine treatment increased ERK phosphorylation via NISH-1. Consistent with the shared caloric-restriction-mimicking gene signature, supplementing rilmenidine to caloric restricted *C. elegans*, genetic reduction of TORC1 function, or rapamycin treatment did not further increase lifespan. The rilmenidine-induced longevity required the transcription factors FOXO/DAF-16 and NRF1,2,3/SKN-1, both important for caloric restriction-mediated longevity. Furthermore, we find that autophagy, but not AMPK signaling, was needed for rilmenidine-induced longevity. Lastly, we find that treating mice with rilmenidine showed transcriptional changes in liver and kidney similar to caloric restriction. Overall, our findings reveal rilmenidine as a caloric restriction mimetic and as a novel geroprotective compound.
Introduction

Individuals over 65 are now the fastest-growing demographic group worldwide, a fact that emblematizes the global aging population (Jarzebski et al. 2021). Unfortunately, at present, with age comes age-related chronic disease and death (Fontana et al. 2014), and as such, the estimated benefits of delaying aging, even if the effect is rather small, are immense (Farrelly 2010; Goldman et al. 2013; de Magalhães et al. 2012). A large body of evidence has demonstrated that the aging rate can be markedly slowed in model organisms. So far, caloric restriction (CR) is the most robust anti-aging intervention (Liang et al. 2018), and CR promotes longevity across species (Fontana et al. 2010). However, studies of CR in humans have had mixed results, low compliance, and many side effects (Most et al. 2017), meaning finding medications that can mimic the effect of caloric restriction is the most reasonable anti-aging target (Ingram & Roth 2015). However, only a few compounds have been identified to mimic the beneficial effects of CR.

Previously, we detailed a drug repositioning method in which we identified potential caloric restriction mimetics (CRMs), yielding biologically relevant results (Calvert et al. 2016). We did this by comparing drug-gene signatures to the in-vitro transcriptome of caloric restriction, looking for drugs with an overlapping profile. One drug identified during this work was the imidazoline receptor agonist allantoin, which was further confirmed to extend lifespan in C.elegans (Admasu et al. 2018). This has highlighted the potential of imidazoline agonists as caloric restriction mimetics. Allantoin, however, does not display oral bioavailability, obviating its use in humans (Kahn & Nolan 2000). We hypothesized, therefore, that a more potent and specific agonist of the imidazoline receptor, the widely-prescribed, oral antihypertensive rilmenidine (Dardonville & Rozas 2004), would have a similar, if not better, longevity effect in C.elegans but with more translatability to humans in the future.
Indeed, rilmenidine elicits a similar transcriptional profile to CR (Calvert et al. 2016). In another study, we also identified rilmenidine to reprogram human cell transcription profiles to a more youthful state (Statzer et al. 2021). Additionally, we identified rilmenidine by searching for gene expression signatures of longevity in mammals and subsequently have shown that the gene expression profile of liver from mice subjected to a diet containing rilmenidine for 1 month displays a strong positive association with the CR transcriptome profiles (Tyshkovskiy et al. 2019). Yet, whether rilmenidine acts as a CR mimetic to promote healthy aging is unknown.

In this work, we first validated the geroprotective properties of rilmenidine in *C.elegans* employing lifespan, healthspan, and stress assays. Rilmenidine extended the lifespan of *C.elegans* when commenced in early adulthood and also in aged animals. It did this via the nematode ortholog of the I1-imidazoline receptor (I1R/nischarin/IRAS), *fl3e9.1/nish-1*, and was dependent upon increased autophagy and manipulation of three CR nexuses (DAF-16/FOXO, SKN-1/NRF, and mTOR) (Blackwell et al. 2015; Blackwell et al. 2019). Lifespan extension was not possible in genetic models of caloric restriction (*eat-2*), suggesting CR mimicry. Lastly, rilmenidine improved thermotolerance and attenuated PolyQ aggregation. This data demands further clinical investigation into the geroprotective effect of rilmenidine.
Results

Rilmenidine improves survival in Caenorhabditis elegans

Previous computational studies predicted rilmenidine as a longevity drug and CR mimetic (Calvert et al. 2016; Tyshkovskiy et al. 2019). To test if rilmenidine is involved in lifespan regulation, we treated wild-type (WT) C. elegans with different concentrations of rilmenidine at the larval stage L4 and performed lifespan assays. We found that rilmenidine extended the lifespan of WT animals by ~19% compared to DMSO-treated WT control, at an optimal concentration of 200 μM (Figure 1A, B, Supplementary Table 1). Thus, we show that rilmenidine is indeed a geroprotective drug promoting longevity.

One goal is to administer geroprotective drugs at geriatric ages. However, not all longevity-promoting drugs work during old age. For instance, the geroprotective drug metformin, when administered at day 10 of C. elegans’ adulthood or 20-months in mice, instead shortened lifespan and accelerated age-related pathologies (Espada et al. 2020; Zhu et al. 2021). Interestingly, we found that in worms rilmenidine prolonged the lifespan to the same extent (~33%) whether exposure to the drug was initiated during youth, day 1 of adulthood, or when old, day 12 of adulthood (Figure 1C, D, Supplementary Table 1). However, this pro-longevity effect was not additive to lifespan extension exerted by early-life treatment, suggesting a possible cellular mechanism that upon activation lasts through adulthood and aging (Figure 1C, D, Supplementary Table 1). These results indicate that rilmenidine might be employed as a pharmacological intervention during old ages to extend lifespan.

Identification of the conserved imidazoline receptor NISH-1 in C. elegans

Rilmenidine has been identified as a classical imidazoline type 1 receptor (I1R/IRAS/Nischarin) agonist in mammals (Zhang & Abdel-Rahman 2006). The
amino-terminal sequence of Nischarin displays strong homology to a C. elegans protein encoded by the gene f13e9.1 (Figure 2A) (Alahari et al. 2000). Our comparison of the C. elegans f13e9.1 full-length protein sequence to the human orthologue IRAS protein sequence revealed high similarity in the following functional domains; PHOX (PX) domain, Leucine-rich repeats (LRRS), and coiled-coil (CC) domain (Figure 2A). The PHOX domain entails phosphoinositide-3-phosphate-binding capacity, likely enabling Nischarin to be anchored to the intracellular surface of the plasma membrane (Piletz 2003). The coiled-coil domain is the predicted imidazoline binding site (Sun et al. 2007) (Figure 2A). Thus, we renamed the f13e9.1 gene to nish-1 and will refer to it as nish-1 throughout this study.

To examine the role of nish-1 as a functional ortholog of Nischarin that could mediate rilmenidine cell signaling, we generated a loss-of-function mutant nish-1(syb767) by CRISPR/Cas9 genome editing. A deletion of 873bp was initiated at the 3rd exon and a stop codon was introduced to generate a potential null allele (see Materials and Methods for details). This resulted in truncated nish-1 protein, containing only the PHOX domain but abolishing LRRs and the coiled-coil domain (Figure 2B). We validated the deletion in the nish-1 mutant by genotyping (Figure S 1A).

Murine nischarin mutants are smaller than their wild-type counterparts (Crompton et al. 2017; Dong et al. 2017; Zhang et al. 2013). Therefore, for the functional characterization of the nish-1 mutant, we first measured their body size. Day-1 adult C. elegans nish-1 mutants were 5.44% smaller than WT (p-value <0.05; Figure 2C). To exclude slower development as the reason for size defects seen in nish-1 mutants, we assessed the development rate of animals via the categorical staging of vulval development (Ludewig et al. 2017). Both WT and nish-1 mutant developed at a similar rate and had reached full vulval development within 48h (Figure S 1B). Also, nish-1 mutants did not exhibit any reproductive developmental abnormalities.
Rilmenidene functions to activate ERK via nish-1 in *C. elegans*

Previously, it has been shown that interaction of rilmenidene with the I1-imidazoline receptor leads to activation of phosphatidylcholine-specific phospholipase C (PC-PLC) and subsequent accumulation of the second messenger diacylglycerol (DAG) from phosphatidylcholine, with the release of phosphocholine. This leads to downstream activation of MAPK/ERK1/2, with phosphorylation of ERK1/2 being a standard read-out for the pathway (Zhang & Abdel-Rahman 2005) (Figure 2D). We utilized this well-characterized pathway read-out for testing if rilmenidene could activate the only orthologue of ERK, MPK-1 in *C. elegans*. The activation of MPK-1 was measured, by way of pMPK-1 to α-tubulin immunoreactivity, in response to 24 hours treatment with rilmenidene at varying concentrations (200 µM, 300 µM, and 400 µM) to establish a working concentration (Figure S 1C). Rilmenidene increased MPK-1 phosphorylation the strongest at the concentration of 200 µM. Since starvation also leads to phosphorylation of MPK-1, we used it as a positive control (You et al. 2006).

To establish whether this increase in activity was mediated via rilmenidene binding to an I1-imidazoline receptor, WT were co-incubated with rilmenidene (200 µM) and efaroxan (1 mM), an established selective I1-imidazoline receptor antagonist. 24 hours of efaroxan treatment alone did not significantly affect baseline MPK-1 activity compared to 1% DMSO vehicle-treated WT. However, efaroxan abolished MPK-1 activation by rilmenidene. This suggests rilmenidene treatment most likely interacts with an endogenous imidazoline binding site in *C. elegans* to elicit increases in MPK-1 activity (Figure 2E).

To further investigate whether the effect of rilmenidene on MPK-1 stimulation in *C. elegans* was mediated by the putative imidazoline ortholog NISH-1, we also measured MPK-1 phosphorylation in the nish-1 mutant strain (Figure 2E). Rilmenidene failed to increase MPK-1 activity in nish-1 mutants. (Figure 2E). In addition, in the transgenic strains
wherein *nish-1* is rescued, the intensity of phosphorylation was higher in the presence of rilmenidine than in control, implying the necessity of *nish-1* for rilmenidine induced MPK-1 activation (Figure 2E).

**Rilmenidine requires *nish-1* for extending lifespan and improving healthspan**

Next, we asked if rilmenidine extends lifespan through the *nish-1* receptor. We found that the lifespan-extending effects of rilmenidine were abolished when *nish-1* was deleted. Rilmenidine-treated *nish-1* mutants lived as long as WT and *nish-1* mutants without rilmenidine (Figure 3A). On the contrary, rescuing the *nish-1* receptor reinstated the increase in lifespan upon treatment with rilmenidine (Figure 3B). This demonstrates that *nish-1* plays a crucial role in relaying the signals downstream to increase lifespan.

Further, we sought to explore if lifespan extension by rilmenidine is accompanied by better healthspan as well. Older animals reduce their movement with time, representing a marker of frailty (Hosono et al. 1980). Therefore, to investigate the healthspan of rilmenidine-treated WT animals, we measured swimming or thrashing assessed as body bends per minute. Rilmenidine showed higher thrashing rates with age in WT but not in *nish-1* mutants (Figure 3C). By contrast, *nish-1* mutants showed comparable numbers of body bends with or without exposure to rilmenidine, indicating beneficial effects of rilmenidine are conferred through the *nish-1* receptor (Figure 3D). The transgenic rescue of *nish-1* abrogated the age-related decline in the motility of *nish-1* mutants by increasing the number of body bends (Figure 3E). Thus, rilmenidine acts via the imidazoline receptor to improve healthspan as well.

**Rilmenidine requires *nish-1* to improve thermotolerance**

Longevity is often associated with better stress survival (Zhou et al. 2011). Here, we determined the thermotolerance of animals exposed to rilmenidine. Rilmenidine treatment
improved the thermotolerance at 37°C of WT animals but not nish-1 mutants (Figure 3F). Transgenic rescuing the NISH-1 receptor in nish-1 mutants restored rilmenidine’s resistance to heat stress (Figure 3F). This indicates that nish-1 is central for rilmenidine to stimulate a conserved protective mechanism that may contribute to lifespan extension.

**Autophagy is required for extension in lifespan by rilmenidine**

Rilmenidine has been shown to induce autophagy in SOD1 or TDP43 mutant mice models of amyotrophic lateral sclerosis (Perera et al. 2017; Perera et al. 2021). This warranted the examination of autophagy induction in *C. elegans* by rilmenidine. We used the mCherry::LGG-1 reporter strain to quantify autophagosome puncta in the posterior intestinal cells (Gosai et al. 2010; Li et al. 2014; Zhang et al. 2015). We found a significant increase in puncta formation by rilmenidine in a dose-dependent manner (Figure 4A). To determine whether the increase in autophagy is merely an association or has a causal effect on the longevity of rilmenidine, we performed lifespan assays and knocked down essential autophagy genes, *lgg-1* and *bec-1*, in WT exposed to rilmenidine or DMSO control. The lifespan extension conferred by rilmenidine was completely abrogated by impairment of autophagy (Figure 4B, C, Figure S2, Supplementary Table 1). This confirms that autophagy induction is a prerequisite for the increase in longevity by rilmenidine, similar to CR (Madeo et al. 2015).

**Rilmenidine attenuates poly-glutamine aggregates**

One important hallmark of neurodegenerative diseases is the accumulation of protein aggregates. These should typically be degraded, perhaps via autophagy. Rilmenidine employs the process of autophagy as a therapeutic strategy reducing the soluble Huntington protein in the Huntington disease (HD) model in mice (Rose et al. 2010). Consistent with this finding in mice, we found that rilmenidine treatments significantly delayed the accumulation of
polyQ40::YFP fusion protein aggregates compared to untreated WT (Figure 4D). Thus, rilmenidine may be considered for delaying the onset of neurodegenerative diseases like HD. Besides, this observation attests to our speculation of rilmenidine’s function as CRM, since DR too, mediates the reduction in the aggregation of polyQ in C. elegans (Matai et al. 2019).

**Rilmenidine acts as a CR mimetic**

Rilmenidine shares the transcriptome profile with CR, as indicated by our meta-analysis studies (Calvert et al. 2016; Tyshkovskiy et al. 2019). Therefore, to discern the signaling pathway through which rilmenidine extends lifespan, we chose a genetic (eat-2 mutant) model of CR in C. elegans (Lakowski & Hekimi 1998). Rilmenidine did not further increase the longevity of the eat-2 mutants, suggesting that rilmenidine may function through the same genetic pathway (Figure 5A, Supplementary Table 1). CR is thought to work through the nutrient sensor, mTOR complex 1, mTORC1 to extend lifespan (Hansen et al. 2007; Statzer et al. 2020). Also, mTOR inhibition leads to autophagy induction, like rilmenidine (Hansen et al. 2007). To investigate the requirement of mTOR in lifespan increase by rilmenidine, we performed lifespan assays using a heterozygous mutant of daf-15, the raptor adaptor protein in mTORC1. We found that rilmenidine could not increase the longevity additively of the daf-15(m81/+ ) mutants (Figure 5B, Figure S3, Supplementary Table 1). Complementary, we inhibited mTOR pharmacologically by rapamycin. Rapamycin significantly increased the lifespan of WT animals. Co-treatment with rilmenidine and rapamycin failed to further increase the mean lifespan extension of WT animals treated either with rilmenidine or rapamycin individually (Figure 5C, Figure S3, Supplementary Table 1). This result strengthens the earlier observation that rilmenidine indeed acts through mTOR1 signaling for increasing the lifespan.

In C. elegans, mTOR inhibition by genetic or pharmacological intervention, like in mammals, leads to activation of key longevity transcription factors such as SKN-1/NRF and
DAF-16/FOXO (Robida-Stubbs et al. 2012). Moreover, on solid media, daf-16 is indispensable for DR-induced longevity (Greer et al. 2007). To explore whether rilmenidine induced longevity effects are DAF-16 and SKN-1 driven, we employed skn-1(tm3411) and daf-16(mgDf50) null mutants to perform lifespan assays. We found that rilmenidine did not significantly extend the lifespan of either daf-16 or skn-1 mutants, revealing a requirement of daf-16 and skn-1 for longevity benefits by rilmenidine (Figure 5D, Supplementary Table 1).

AMPK is another critical metabolic switch, which works antagonistically with the mTOR pathway and is activated by CRM drugs like metformin (Zhang et al. 2019; Onken & Driscoll 2010). Thus, we assessed the necessity of AMPK in the mediation of rilmenidine’s pro-longevity effects. We found that rilmenidine additively extended lifespan in aak-2(gt33) mutants compared to WT, suggesting AMPK is not required for lifespan regulation by rilmenidine (Figure 5D, Supplementary Table 1).

All the above results show that rilmenidine functions through the imidazoline receptor, nish-1, which activates genetic pathways mediated by the established CR nexuses to extend the lifespan.

Rilmenidine produces longevity-associated gene expression effects in mouse tissues

To test if the observed beneficial effects of rilmenidine are recapitulated in mammals, we performed RNA-seq analysis of liver and kidney samples from young UM-HET3 male mice subjected to rilmenidine for one month, together with the corresponding controls. We identified transcriptomic effects of the drug independently for each tissue and compared them with biomarkers of lifespan-extending interventions and aging, using a gene set enrichment analysis (GSEA) based approach (Figure 6A) (Tyshkovskiy et al. 2019). Gene expression signatures of lifespan extension reflect a liver response to individual interventions, such as caloric restriction (CR), rapamycin and mutations leading to growth hormone (GH)
deficiency; gene expression changes shared by different longevity interventions; and genes, which expression is quantitatively associated with the effect on maximum and median lifespan (Tyshkovskiy et al. 2019). Aging signatures correspond to murine age-related gene expression changes in the liver and kidney, according to the Tabula Muris Consortium (Schaum et al. 2020).

Interestingly, changes induced by rilmenidine in both liver and kidney were positively associated with the majority of longevity signatures, including biomarkers of median and maximum lifespan extension (adjusted p-value < 2 \times 10^{-3} for both liver and kidney), CR (adjusted p-value = 0.066 for liver), rapamycin (adjusted p-value = 6.7 \times 10^{-4} for kidney) and common signatures of longevity interventions (adjusted p-value < 0.013 for both liver and kidney). In other words, genes up- and downregulated in long-lived mice were also changed in the same direction after rilmenidine administration (Figure 6B-E). Remarkably, we also observed a significant positive association of rilmenidine effect with biomarkers of aging (median adjusted p-value = 0.03), which may be partly due to the age-related downregulation of insulin and IGF-1 signaling (López-Otín et al. 2013).

To understand if rilmenidine in mice regulates cellular functions similar to those identified in *C. elegans*, we also performed functional GSEA. This method allowed us to test if genes involved in certain pathways, based on REACTOME, KEGG and GO BP annotation, were enriched among those affected by the compound. We observed significant enrichment of genes involved in mTOR signaling and proteolysis among genes upregulated by rilmenidine in murine liver and kidney, respectively (adjusted p-value < 7 \times 10^{-3}) (Figure 6A). These results suggest that longevity-associated molecular mechanisms induced by rilmenidine in *C. elegans* are reproduced in mammals, pointing to a conserved geroprotective effect and downstream signaling (Figure 6F).
Discussion

Challenges with humans’ long-term caloric restriction (CR) routines raise an unmet need to explore pharmacological interventions like calorie restriction mimetics (CRM). These may provide an alternative to CR, conferring the same longevity benefits without the challenges or side effects of low-calorie diets. Studies detailing the therapeutic efficacy of CRMs when administered at geriatric stages are rarely performed (Ingram & Roth 2015). Identifying drugs that can mirror the benefits of long-term calorie restriction even when the administration is initiated at late stages of life are of great interest.

Our study illustrates that rilmenidine acts as a CRM in *C. elegans*, improving survival, even when administered at older ages (commencing day-12 of adulthood). Rilmenidine targets conventional signaling nodes like mTOR, DAF-16/FOXO, SKN-1/NRF in *C. elegans* and activates stress response pathways, enhancing autophagy, attenuating polyglutamine aggregation, and improving thermotolerance. Rilmenidine acts via a novel mechanism wherein the newly identified imidazoline receptor, *nish-1* (previously *f13e9.1*) in *C. elegans*, mediates signaling to the genetic pathways relating to CR to extend lifespan. Although the imidazoline receptor signaling had been implicated to interact with alpha/beta-integrin or insulin-dependent PI3 kinase/Akt pro-survival pathways (Dontenwill et al. 2003), the connection of the imidazoline receptor to CR or longevity had not been studied. We showed that rilmenidine-induced lifespan and healthspan extension are dependent upon the imidazoline receptor, *nish-1*.

Many known CRMs, like metformin, rapamycin, or resveratrol, had been shown to act via AMPK activation, mTOR inhibition, and sirtuin activation, respectively (Shintani et al. 2018). Rilmenidine too can be viewed as a CRM as it regulates the intracellular signaling systems like the mTOR pathway in *C. elegans*. Indeed, rilmenidine induces the recycling process of autophagy, which is considered one of the characteristic features of CRMs and
mTORC1 signaling (Chung & Chung 2019; Mariño et al. 2014). In mouse models, rilmenidine induced similar changes in mTOR regulation and autophagy activation in liver tissue (Figure 6). Since mTOR signaling and AMPK are two key players in regulating autophagy, and AMPK is not involved in lifespan extension by rilmenidine, rilmenidine probably induced autophagy by inhibition of inhibition mTOR signaling. The exact signal transduction pathway from the interaction of rilmenidine with \textit{nish-l} to the inhibition of mTOR might be worth addressing in future studies. Rilmenidine has been reported to enhance the phosphorylation of ERK/MAPK via the phosphatidylcholine-specific phospholipase C pathway (Zhang & Abdel-Rahman 2005). We speculate, therefore, that ERK activation may inhibit the mTOR complex. In fact, mTOR inhibition upregulates phospho-ERK signaling through p70S6K in a feedback loop (Sunayama et al. 2010; Alonso et al. 2016). This might explain the upregulation of phospho-ERK upon rilmenidine binding to nischarin-1/IRAS. In any case, questions remain as to what signal transduction components are implicated by rilmenidine to induce autophagy.

By modulating autophagic flux, CRMs protect against neurotoxic protein aggregates (Yang & Zhang 2020; Sanchis et al. 2019). Previous studies reported autophagy to be a major checkpoint for the protein aggregates in many neurodegenerative polyglutamine diseases such as spinocerebellar ataxias, Huntington disease, etc, among model organisms (Ravikumar et al. 2002; Ravikumar et al. 2004). For instance, pharmacologically inducing autophagy using the CRM rapamycin encouraged the degradation of ataxin-3 aggregates (Ravikumar et al. 2004; Berger et al. 2006). Similarly, rilmenidine was proven effective for removing polyQ aggregates in mice models of Huntington disease and amyotrophic lateral sclerosis by upregulating autophagy (Rose et al. 2010; Perera et al. 2018). We further showed rilmenidine to be beneficial against the polyQ aggregates in \textit{C. elegans}. Additionally, we observed
upregulation of proteolysis in the rilmenidine treated kidney tissue in mice, contributing to increased clearance of the aggregated proteins.

Cytoprotective autophagy is also activated by heat shock response (Kumsta et al. 2017). Furthermore, the heat shock response helps to curb proteotoxicity due to aggregation (Hsu et al. 2003). We found that rilmenidine improved thermotolerance in C. elegans. These observations indicate that autophagy induction by rilmenidine is an important stress evasion mechanism, which supports proteostasis, possibly through regulating the proteasome pathway and heat stress response and thus, extending lifespan.

Alongside being a clinically-approved anti-hypertensive drug, rilmenidine improves plasma lipid and blood glucose in patients with hypertension and metabolic syndrome (Luca et al. 2000; Reid 2001). Given its contribution towards lowering blood glucose and increasing insulin sensitivity, rilmenidine may have an anti-diabetic function. Since I1R and I3R do not share similar ligand binding sites and I3R is a regulator of insulin secretion, the potential interaction of rilmenidine with the I3type imidazoline receptor might explain its effects on improved insulin resistance and, thus, glucose levels (Bousquet et al. 2020). As rilmenidine improves glucose tolerance, it can likely be put in the insulin sensitizer class of CRMs (Shintani et al. 2018, Ingram et al. 2006).

Since it is already in use, it becomes feasible for rilmenidine to be repositioned against insulin resistance, metabolic syndrome, and polyglutamine diseases. Moreover, conservation of findings between C. elegans, mice, human cell culture, such as induction of autophagy, points towards the more considerable potential for translating the longevity benefits to humans. In conclusion, rilmenidine is a new addition to the list of potential CRMs that could be viable therapeutic interventions administered late in life, and thus, warrants further examinations.
Figures and legends

Figure 1. Improved survival of *C. elegans* treated with rilmenidine.

A) Pooled survival curve showing significant lifespan extension in WT animals treated with rilmenidine at all concentrations, except 400 µM, compared to 1% DMSO vehicle control. B) Bar graph showing quantified lifespan data in terms of mean lifespan (days) for each rilmenidine concentration (100-400 µM) treated WT, compared to DMSO vehicle control; 200 µM concentration of rilmenidine provides maximum lifespan increase (19%) in WT. C) Pooled survival curve showing late-life treatment with rilmenidine (200 µM) at day 12 of adulthood increased lifespan in WT compared to DMSO vehicle control. D) Quantified data for lifespan assay showing percentage increase in mean lifespan of adult WT fed 200 µM rilmenidine at different times (day-1 or day-12 adulthood) compared to DMSO vehicle control. Error bars represent SEM; adjusted p-value was derived from log-rank test and Bonferroni correction. Kaplan Meir survival analysis was performed on pooled data from at
least three independent trials. Quantitative data and statistical analyses for the representative experiments are included in Supplementary Table 1.

Figure 2. Characterization of NISH-1.

A) BLASTp alignment of *C. elegans* NISH-1 protein sequence (CELE_F13E9.1 – UniProtKB-K8F807) against truncated human NISCHARIN isoform 3 (HUMAN_NISCH –
UniProtKB-Q9Y2I1-3) revealing 38% sequence similarity, with conserved secondary structure elements like alpha helices shown in red and beta-strand in blue. Functional protein motifs like PHOX (Px) domains, LRRs, and coiled-coil domains are underlined. B) Schematic diagram illustrating exons of nish-1 gene, alongside translated protein motifs and the deleted regions in the homozygous nish-1 mutant. C) nish-1 mutants exhibit reductions in body size compared to WT at day 1 of adulthood as shown by body area (defined as width x length). A two-tailed $t$-test was used for analysis; *p<0.05. Representative pictures of WT and nish-1 mutant captured in brightfield at 10X objective on a Zeiss Axio Observer following paralysis in 20 mM tetramisole. Animal heads are left. Scale bar: 200 μm. D) Model illustrating the phosphorylation of ERK following interaction of rilmenidine to the NISH-1/IRAS receptor. E) Western blots showing MPK-1 phosphorylation in WT, nish-1 mutants, or nish-1 rescue transgenic animals, in response to 24h pharmacological intervention. α-tubulin was used as the loading control. Rilmenidine significantly increased MPK-1 phosphorylation *(%CV vs. FC = >1.5), however neither efaxoran alone nor rilmenidine and efaxoran in combination significantly increased MPK-1 phosphorylation n/s (%CV vs. FC = <1.5). Rilmenidine failed to reproducibly increase phosphorylation of MPK-1 in nish-1 mutants. n/s (%CV vs. FC = <1.5). Transgenic rescue strains significantly increased MPK-1 phosphorylation upon rilmenidine treatment. **(%CV vs. FC = >2 in PHX945) and (%CV vs. FC =>1.5 in PHX946). Data is quantified as log-fold densitometric ratio change relative to α-tubulin. Data is then expressed as log fold change compared to DMSO vehicle control. Coefficient of variation (%CV), defined as the percent standard deviation: mean ratio; the significant difference in groups if % fold change is x1.5* greater or x2** than % CV.
Figure 3. *nish-1* is required for an extended lifespan and better healthspan.

**A)** Survival curve showing lifespan extension by rilmenidine in WT, but not in *nish-1* mutant (Statistics and raw data are in Supplementary Table 1). **B)** Rescue of *nish-1* enabled extension in lifespan by 200 μM rilmenidine in *nish-1* mutant. DB06 (=PHX893) is a *nish-1* knockout mutant and DB03 (=PHX945) and DB04 (=PHX946) are two rescue lines expressing wild-type *nish-1* gene copies. Quantitative data and statistical analyses for the representative experiments are included in Supplementary Table 1. **C)** Quantified data of body bends representing motility deterioration with age in WT (C), *nish-1* mutants (D), and two *nish-1* rescue mutants (PHX945 (=DB03) and PHX946 (=DB04)) (E) in the presence or absence of rilmenidine. Rilmenidine at 200 μM significantly reduced age-related motility deterioration in WT and *nish-1* rescue strains, but not in *nish-1* mutants. Data is represented as a pooled mean of 10 animals per time-point and condition/genotype repeated over three independent trials overlaid on a box plot representing quartiles. Error bars, upper: Q3 + 1.5*IQR; minimum: Q1 -1.5*IQR (adj. p<0.05; 2-way ANOVA and Tukey post hoc). **F)** Thermotolerance:
Rilmenidine increases the percentage survival of WT exposed to 37°C for 3 hours and ~20h recovery period, dependent on *nish-1*. Bar graph represents mean % survival ± S.E.M from three independent trials of at least 150 animals per strain and/or condition. *p<0.05* (one-way ANOVA with Tukey post hoc comparisons).

**Figure 4. Induced autophagy by rilmenidine perturbed polyQ aggregation.**

**A)** Representative images of day 2 adult transgenic animals, expressing the intestinal specific autophagy reporter gene *Pnhx-2::mCherry::lgg-1* showing increased autophagy, when exposed to varying concentrations of rilmenidine for 24h compared to 1% DMSO vehicle.
Arrows indicate autophagosome puncta formation. Scale bar = 20 μm. Graph shows the interquartile distribution of the mean number of mCherry::LGG-1 puncta in the posterior intestine of the animals in each condition. Error bars, upper: Q3 + 1.5*IQR; minimum: Q1 -1.5*IQR. **P<0.01, *P<0.05; one-way ANOVA followed by a Tukey’s post hoc test. B,D) Inhibition of autophagy abrogates the pro-longevity effect of rilmenidine as shown by survival curves of WT animals fed either RNAi bacteria expressing an empty vector (L4440), or bec-1 (B) or lgg-1 (D) dsRNA from day 1 adulthood in the presence or absence of 200 μM rilmenidine. Kaplan Meir survival analysis was performed on pooled data from at least three independent trials. Groups tested by Log-rank with Bonferroni correction; p<0.05. Quantitative data and statistical analyses for the representative experiments are included in Supplementary Table 1. C,E) Representative fluorescent micrographs of the transgenic strain expressing polyQ40::YFP in body-wall muscles showed a reduction of polyQ40 aggregation by differing rilmenidine concentrations. Scale bars = 100 μm. The graph depicts quantified data of Q40::YFP aggregates per entire nematode after treatment with differing rilmenidine concentrations for the indicated times from L1. Data represented as the pooled mean no. aggregates per animal. Error bars are ± S.E.M. Significance derived from 2-way repeated-measures ANOVA p<0.05.
Figure 5. Effects of rilmenidine treatment on survival of CR-associated mutants.

Survival curves showing the inability of rilmenidine to extend life span in (A) DR412 daf-15(m81/+), (B) rapamycin-treated WT, (C) DR1116 eat-2(ad116), and (D) GR1307 daf-16(mgDf50), LD1057 skn-1(tm3411), but TG38 aak-2(gt33) mutants. Raw data, quantitative data, additional trials, and statistical analyses for the representative experiments are included in Supplementary Table 1.
Figure 6. Rilmenidine treatment in mice induces gene expression changes associated with a lifespan-extending effect.

A) Association of the liver (top) and kidney (bottom) responses to rilmenidine with signatures of aging (left), lifespan-extension (middle), and longevity-related intracellular processes (right). The significance score was calculated as \( \log_{10}(\text{adjusted p-value}) \) corrected by a sign of 26.
regulation. Dotted lines represent the FDR threshold of 0.1. B-E) GSEA enrichment plots with running normalized enrichment scores (top) and distributions (bottom) of selected up-(red) and downregulated (blue) gene signatures of lifespan-extending interventions among genes perturbed by rilmenidine in the liver (B-D) and kidney (E). Perturbed genes were sorted based on the log_{10}(p-value) of their differential expression between control and rilmenidine-treated samples corrected by a sign of regulation. The resulting index was divided by the number of genes in the dataset. Dotted lines represent NES calculated for up- (red) and downregulated (blue) gene signatures. F) Working model representing possible pro-longevity signaling by rilmenidine to extend life span in C. elegans. NES: Normalized enrichment score; CR: Caloric restriction; GH: Growth hormone; mTOR: mammalian target of rapamycin.
Figure S1. A) Endpoint single worm PCR genotyping of WT and nish-1 deletion mutant following x4 outercrossing. Using oligonucleotides flanking the deletion site (external primer...
pairs) generated either a 1968 bp or a 1936 bp amplicon depending on oligonucleotide pair in WT or a correspondingly 873 bp reduced amplicon size in the genomic DNA of homozygous nish-1 mutants. Likewise, an oligonucleotide pair targeted to sequences within the CRISPR deletion site produced a 1558 bp amplicon in WT but no amplicon in homozygous nish-1 mutants. Control amplifications were conducted on the lgc-37 gene coded on chromosome III. Primer sequences are in Supplementary Table 2. B) Effect of different concentrations on rilmenidine-induced MPK-1 phosphorylation as measured by Western Blotting. Phosphorylation of MPK-1 was assessed in WT in response to 24h pharmacological intervention. Data is then expressed as log fold change compared to 1% DMSO vehicle control. Rilmenidine at 200μM and 400μM significantly increased MPK-1 phosphorylation Coefficient of variation (%CV), defined as the percent standard deviation: mean ratio; significant difference in groups if % fold change is x1.5* greater or x2** than % CV. **(%CV vs. FC = >2). C) nish-1 mutants and rilmenidine-fed WT do not exhibit delayed vulval development from the L3 stage. In the chart, developmental stages are listed on the Y-axis, and the number of hours after a 24h exposure to food from L1 is plotted on the x-axis. The areas of the circles in the chart reflect the percentage of the population at each stage of development; n≥30 from 3 independent pooled trials for each time point. Statistical difference established by way of 2-way ANOVA wherein a mean development score could be ascertained through pre-L3 corresponding to a score of 0, Early L3 is 1 and Mid L3 a 3 etc. No groups displayed any significant difference in development rate.
**Figure S2:** Related to figure 4B and 4C. Quantified data for mean lifespan (days) and % change in survival of WT upon knockdown of autophagy genes, *bec-1* and *lgg-1* alongside rilmenidine treatment. Rilmenidine significantly increased lifespan by 8.29% in WT animals fed HT115 *E.coli* expressing an empty vector (*p*<0.001), however, failed to significantly affect lifespan in animals fed HT115 *E.coli* expressing *bec-1* or *lgg-1* dsRNA; n/s = *p*>0.05. See Supplementary Table 1 for details.

**Figure S3:** Related to figure 5B and 5C. Quantified data for mean lifespan (days) and % change in lifespan upon rilmenidine administration in *daf-15(m81/+)* mutants and rapamycin-treated WT. Error bars represent ± SEM; adjusted p-value was derived from log-rank test and Bonferroni correction. See Supplementary Table 1 for details.
Supplementary Table 1: Lifespan summary and details.

Supplementary Table 2: Primer sequences.
Materials and Methods

C. elegans Strains

All strains were maintained on OP50 E.coli seeded NGM agar supplemented with 50 IU/mL penicillin (Sigma #P4333), 50 µg/ml streptomycin (Sigma #P4333), and 50 IU/mL Nystatin (Sigma #1638) to prevent contamination (Stiernagle 2006). The strains used in the study are: WT (N2 Bristol), PHX893 (=DB06) nish-1(syb767), PHX117 sybIs62 [f13e9.1::EGFP::3xFLAG, unc-119(+)] + Pmyo-2::mCherry], PHX116 sybIs62 [f13e9.1::EGFP::3xFLAG, unc-119(+)] + Pmyo-2::mCherry], PHX945 (=DB03) and PHX946 (=DB04) are two independent integrated lines of f13e9.1 (syb767) IV; sybIs62 [f13e9.1::EGFP::3xFLAG, unc-119(+)] + Pmyo-2::mCherry], VK1093 vkEx1093 [Pnhx-2::mCherry::lgg-1], AM141 rmls133 [Punc-54::Q40::YFP], GR1307 daf-16(mgDf50) I; DA1116 eat-2(ad1116) II; LD1057 skn-1(tm3411) (Ewald et al. 2015) IV, DR412 daf-15(m81)/unc-24(e138) IV; and TG38 aak-2(gt33) X.

The transgenic strains PHX893 (=DB06) nish-1(syb767), PHX117 sybIs62 [f13e9.1::EGFP::3xFLAG, unc-119(+)] + Pmyo-2::mCherry], and PHX116 sybIs62 [f13e9.1::EGFP::3xFLAG, unc-119(+)] + Pmyo-2::mCherry] were generated by SunyBiotech. In PHX893 (=DB06) nish-1(syb767), the 3rd exon, 4th exon, and part of 5th exon, was deleted and replaced by a splice site and stop codon that terminates transcription and resulted in 873 bp homozygous deletion in the nish-1 gene. This strain was subsequently outcrossed four times against WT background using PCR screening at each stage, and the final outcrossed strain was confirmed by sequencing. The details of the primers are documented in Supplementary Table 2. PHX117 and PHX116 contain integrated full-length rescue of the nish-1 locus within the fosmid generated in the WT background. Specifically, the integrated rescue contained the
WRM0616C_F02(pRedFlp-Hgr)(F13E9.1[33366]::S0001_pR6K_Amp_2xTY1ce_EGFP_FR T_rpsl_neo_FRT_3xFlag)dFRT::unc-119-Nat) fosmid construct derived from the copy-number inducible vector pCC1Fos grown up in *E.coli* strain EPI300. Fosmid DNA was extracted and added to a 20 ng/µL DNA mixture containing both the purified fosmid and pCFJ90 *Pmyo-2::mcherry*. The solution was microinjected into 10-30 adult WT gonads to generate a line of extrachromosomal transgenic arrays. Stable lines were then generated by irradiating with a γ dose of 40 GRAY to produce two lines of the same genotype, PH117, and PH116. The previously produced, PHX893 (F13E9.1(syb767)IV) was then crossed 4 times with PX116 to generate the two rescue lines: PHX945 (=DB03) and PHX946 (=DB04) *f13e9.1 (syb767) IV; sybIs62 [f13e9.1::EGFP::3xFLAG, unc-119(+)] + Pmyo-2::mCherry*.

**Generation of UV-Killed OP50 *E.coli***

32 mL of unconcentrated live OP50 *E.coli* cultures were aliquoted to each 50 mL falcon tubes, mixed with absolute ethanol (8 mL) to yield a 20% ethanol solution as described in Calvert et al., 2016 and pipetted into T160 flasks to a volume of 150ml. Flasks were transferred to a UV-Linker machine (CL-1000 Ultraviolet crosslinker UVP) and irradiated for 120 minutes at 999,900 microjoules/cm². 50ml aliquots of UV-killed *E.coli*-ethanol solution was pelleted at 3000 rpm for 10 minutes and resuspended to a 10X concentrate in 5 mL Lennox LB broth.

**Drug Treatment**

All drug treatments, unless otherwise stated, were administered to *C.elegans* via the addition of the respective compound to NGM (Zheng et al. 2013). Aliquots of stock solutions of the drugs (rilmenidine and rapamycin) were prepared by dissolving them into DMSO solvent,
such that final volume in NGM agar did not exceed 1%. NGM agar was cooled to below 65°C after same-day autoclaving, and respective drugs were concomitantly added. Plates were gently swirled to homogenize the solution and left to dry for 1 hour in a hood. 60 µl of 10X concentrated UV-killed OP50 *E.coli* was then spotted to the centre of the plate and again the plates were left to dry for 30 minutes in a hood. Unless otherwise stated, plates were prepared one day before use.

**Lifespan Assay**

Well-fed gravid hermaphrodites were L1 synchronized overnight at 20°C, after hypochlorite treatment. The next day, L1s were transferred to NGM agar plates seeded with 10X OP50 *E.coli* and allowed to develop for 52 hours at 20°C until late L4/adult. At L4/adult molt, animals were transferred to 6-well lifespan assay plates, containing 3 ml of NGM per well with test compounds and 100 µg/ml 5-Fluoro-2’deoxyuridine (FUdR) (Calvert et al. 2016). Plates were maintained at 20°C in the dark, wrapped in parafilm to moderate humidity. Animals were counted every 2 to 3 days, and dead ones (failure to respond to no more than 3 "prods" with a platinum pick) were removed at each inspection. 60 µL of UV-killed OP50 was added three times a week to all plates until day 15 of the assay by which point low-food consumption negated the need for additional *E.coli* (Calvert et al. 2016).

For late administration lifespan assays, at day 12, animals were washed of lifespan assay plates with M9 buffer and carefully pipetted onto new drug plates prepared on day 11, seeded with UV-killed *E.coli*, and contained freshly prepared test compounds and 100 µg/ml FUdR.

For RNAi lifespan analyses, L1 synchronized WT *C. elegans* were grown on live OP50 *E.coli* until late L4-stage. At L4 stage, animals were then transferred to NGM plates with 100 µM IPTG, 400 µM FUdR, and 50 µg/ml carbenicillin with either 200 µM rilmenidine or vehicle
seeded with live *E.coli* (HT115) expressing an empty vector, *bec-1* dsRNA or *lgg-1* dsRNA. Plates were incubated for 1–2 days at room temperature prior to inducing dsRNA expression. Animals were maintained on the same drug plates for the entirety of their lifespan and scored for survival every 2-3 days.

For all lifespan assays, unless otherwise stated, at least 150 animals per genotype and/or condition were used across at least three independent trials. Animals that crawled into the agar, or experienced matricidal events were removed from the assay and not included in the life span calculations. The Kaplan-Meir nonparametric method was used for estimating survival curves. For each condition and/or genotype, survival results were pooled. Lifespan statistics were calculated using the OASIS2 online tool (Han et al., 2016). P-values relating to survival differences between populations were calculated using a log-rank (Mantel-Cox method) test.

**Automated Lifespan Measurements**

Automated survival analysis was performed as described in (Statzer et al. 2020) employing the lifespan machine setup developed by Stroustrup and colleagues (Stroustrup et al. 2013). Briefly, *C. elegans* were age-synchronized using bleach lysis and kept in L1 culture (52h, 20°C), placed on live OP50 until L4 stage, and subsequently transferred to plates containing the drug, FUdR (50 µg/ml) and dead OP50 from L4 to day 4. Lastly, the animals were moved to tight-fitting Petri dishes (BD Falcon Petri Dishes, 50 x 9 mm) containing the drug, dead bacteria, and FUdR and imaged until the end of life.

Rilmenidine hemifumarate was dissolved in 1% DMSO and supplemented to the agar just before pouring to yield 200 µM, 300 µM, and 400 µM final concentrations. When using dead
bacteria, the agar was additionally supplemented with Nystatin (44 U/ml) and Penicillin-Streptomycin (50 U/mL).

Air-cooled Epson V800 scanners were utilized for all experiments operating at a scanning frequency of one scan per 30 minutes. To limit condensation, the tight-fitting plates were dried without lids in a laminar flow hood for 40 minutes before starting the experiment.

Furthermore, temperature probes (Thermoworks, Utah, U.S.) were used to monitor the temperature on the scanner flatbed and maintain 20.0°C. Animals that left the imaging area during the experiment were censored. Automated lifespan results were validated using manual lifespan measurements as described in (Ewald et al. 2016) by picking L4 from normal culturing plates onto the corresponding assay plates. Lifespans were calculated from the L4 stage (= day 0).

**Body Size Phenotyping**

Synchronized day 1 *C. elegans* hermaphrodites cultured on live OP50 *E. coli* NGM plates were measured for body length and width in brightfield at 10X objective on a Zeiss Axio Observer following paralysis in 20 mM tetramisole. Between 10-20 animals per data point were used. The animal's length was measured from head (most visibly anterior buccal line) to a position where the tail tapered to a 10 ± 0.5 μm diameter (Petzold et al., 2011). Body width measurements were taken from the posterior vulval peak to the corresponding outer edge of the intestinal cuticle (Collins 2007). Measurements were made using the segmented lines function on Image J software normalized to the scale bar.

**Developmental Measurements**
Bleach synchronized animals that had been halted in L1 starvation for 24h were measured for development following exposure to food source (Schindler et al. 2014). ~ 500 L1s per genotype/condition were placed onto NGM plates containing rilmenidine and spotted with live OP50 *E.coli* and allowed to develop at 20°C. After 24 hours, measurement of development was conducted at different time points: 0 hours (24 hours exposure to food source), 3 hours, 6 hours, and then 24 hours (48 hours exposure to food source). Per trial and time point, 10 animals were immobilized and mounted onto glass slides in 20 mM solution of tetramisole hydrochloride and then imaged at a 10X objective for body length and 60X objective for vulval development in brightfield on a Zeiss Axio Observer. Vulval development was scored using visual identification of late larval stage vulval checkpoints detailed by Schindler (2014). Statistical significance of differences in vulval development was determined by Two-way ANOVA.
**Motility Assay**

NGM agar plates were firmly tapped onto the microscope to stimulate movement. In responding animals, not impeded by OP50, body bends were counted for 30 seconds in 10 animals for each condition at each time point (5, 10, and 15 days post-L4 molt) per trial across three independent trials to a total of 30 animals per genotype and condition. Mean deterioration in motility per genotype and/or condition was compared by 2-factor repeated ANOVA and Tukey post hoc correction whilst individual time-point comparisons were tested by student’s $t$-test.

**Protein Extraction**

~500 day-1 adult animals were transferred to either empty NGM plates to induce starvation (+ve control) or to UV-killed OP50 *E.coli* NGM plates containing requisite drugs or vehicles. After 24 hours, animals were collected in an M9 buffer and centrifuged at 1000 rpm for 2 minutes to yield a wet pellet. Animals were then re-suspended on ice in 50 µl of RIPA buffer with added phosphatase inhibitors (Cell Signalling Technologies #9806) and complete EDTA-free protease inhibitor cocktail (Roche # 04693159001) before sonication: 10 sec ON/1 minute OFF, 14% amplitude 3 times until the vast majority of animals were dissolved. Lysates were then spun down twice at 13,000 g/ 20 minutes 4°C and immediately frozen at -20°C for no more than one week before use.

**Western Blot Analysis**

Total protein lysate concentrations were quantified using a bicinchoninic acid (BCA) assay (BioRAD) according to the manufacturer’s protocol.
Protein lysates were heated for 5 minutes at 95°C, cooled on ice for 2 minutes, and pelleted for 3 minutes at 13,000 g. 25 µg of protein was loaded per well into an 18 well 10% TGX™ Precast Gel (BioRad), submerged in running buffer (25 mM Tris, 192 mM glycine, 0.1% SDS, pH 8.3) and separated at 300V for approximately 30 minutes. Gels were transferred via Trans-Blot® Turbo™ Transfer System to nitrocellulose membranes and blocked for 60 minutes in blocking buffer (Licor Odyssey® Blocking Buffer #P/N 927-40100). Blots were incubated overnight with the primary antibody Phospho-ERK (Cell Signalling Technology, Catalogue number: 9101) at 1:1000 in blocking buffer (Licor Odyssey® Blocking Buffer #P/N 927-40100) (Gee et al., 2013) with gentle rocking. The following morning, blots were washed three times in TBST before being incubated for 1 hour with the secondary antibody IRDye® 800CW Goat anti-Rabbit IgG at a concentration of 1:10,000 in blocking buffer w/ 0.1% tween. α-tubulin was used as the loading control [Primary anti-α-tubulin : ab72910 and Secondary antibody: IRDye® 680RD Goat anti-Mouse IgG]. Bound antibody was detected using Odyssey® CLx Imaging System. Changes in ERK phosphorylation are often calculated using phospho-ERK:Total ERK ratios (Kao et al. 2004; Nykamp et al. 2008). However, given the frequent use of ERK as a loading control, we calculated changes in phospho-ERK via the surrogate loading control α-tubulin.

For all immunoblotting, three independent trials for each genotype and/or condition were completed and mean densitometric ratios between loading control and pERK minus local median background were calculated using Image Studio Lite as per software guidelines (Licor Biosciences 2013). Results were considered significant if the magnitude of the %change exceeded the CV by at least x1.5.
**Thermotolerance and Recovery Assay**

Well-fed gravid hermaphrodites were bleach synchronised and resultant embryos were allowed to grow for 52 hours at 20°C on NGM plates seeded with live OP50 *E.coli*. At late-L4 stage, animals were transferred to seeded UV-killed OP50 *E.coli* NGM plates containing 200-400 µM rilmenidine, or vehicle for 24h. After 24h of the respective drug exposure, plates were upshifted to an incubator preset to 37°C to induce heat shock. After 3 hours, plates were moved to 20°C and scored for survival after a ~20h “recovery period” (Kumsta et al. 2017). At least 100 animals per condition across three identically designed independent trials were scored as either dead or alive by their ability to respond to no more than 3 "prods" with a platinum pick. Percentage survival was calculated and populations were compared by one-way ANOVA of variance with Tukey post hoc correction.

**Measurement of Autophagy**

Gravid hermaphrodites were bleach synchronized and resultant embryos were allowed to grow for 52 hours at 20°C on NGM plates seeded with live OP50 *E.coli*. At late-L4 stage, animals were transferred to either empty NGM plates to induce starvation (+ve control) or to UV-killed OP50 *E.coli* NGM plates containing requisite drugs or vehicle (1% DMSO). Animals were then incubated at 20°C for 24 hours to maximize drug absorption and efficacy (Zheng et al., 2013). After 24 hours, approximately 10 day-1 adults per condition were imaged at a 10X objective on a Zeiss Axio Observer using a 150 ms exposure time. In total, at least 25 images across three independent trials were collected and pooled for each condition (Eisenberg et al. 2009; Morselli et al. 2010). Levels of autophagy induction were quantified using Zen software. Three 1 mm² boxes were sequentially assigned to the most fluorescent
areas of the posterior intestine for each animal and with set histogramic parameters (black: 0
gamma:1.0 white: 1000 or 5000 for highly fluorescent animals), the total number of visible
puncta were manually counted in each box, and a mean number of puncta calculated per mm²
of animal intestine. Median number of puncta per mm² posterior intestine for all animals in
each condition was calculated and compared using one-factor (ANOVA) variance analysis
corrected by the post hoc Bonferroni test.

PolyQ Protein Aggregation Assay

Gravid hermaphrodites were bleached, and synchronized L1s were transferred to NGM plates
containing rilmenidine or DMSO. After incubation for the indicated periods of time,
approximately 30 animals per condition across three trials were imaged under a Zeiss Axio
Observer microscope at 10X objective. The number of polyQ40::YFP aggregates in body wall
muscle was counted using ZenBlue software with set histogramic parameters (black: 500
gamma:1.0 white: 5000). Approximately 50 animals across three independent trials were
randomly selected for each treatment group and scored for the number of aggregates. Groups
were compared by two-factor repeated measures ANOVA and Tukey posthoc correction.

RNA-sequencing Data Processing

RNAseq data of gene expression changes induced by rilmenidine in mouse liver after 1 month
of oral administration was obtained from (Tyshkovskiy et al. 2019). In addition, we performed
RNA sequencing of corresponding kidney samples from 8 control mice and 4 mice subjected
to the drug for 1 month. RNA was extracted from tissues with the PureLink RNA Mini Kit as
described in the protocol. Libraries were prepared and sequenced with 100 bp read length option on the Illumina HiSeq 2500.

Quality filtering and adapter removal were performed using Trimmomatic (version 0.32). Processed/cleaned reads were then mapped with STAR (version 2.5.2b) and counted via featureCounts (version 1.5). To filter out genes with low number of reads, we left only genes with at least 6 reads in at least 66.6% of the samples separately for each tissue. Filtered data was then passed to RLE normalization (Anders & Huber 2010). Differentially expressed genes between control and treated mice were identified using edgeR. For each gene, we calculated the p-value of its logFC in response to rilmenidine compared to control independently for every tissue. We then converted them to $\log_{10}(p$-value) corrected by the sign of regulation, calculated as:

$$-\log_{10}(pv) \times sgn(lfc),$$

where $pv$ and $lfc$ are p-value and logFC of a certain gene, respectively, and $sgn$ is signum function (is equal to 1, -1 and 0 if the value is positive, negative and zero, respectively). The resulting pre-ranked list of genes was used to identify significant associations with the biomarkers of aging, lifespan extension and specific intracellular functions.

**In-silico Association Analysis**

To identify functions enriched by genes perturbed by rilmenidine, we performed gene set enrichment analysis (GSEA) (Subramanian et al. 2005) on a pre-ranked list of genes, obtained as described earlier. REACTOME, KEGG, and GO BP from Molecular Signature Database (MSigDB) have been used as gene sets for GSEA. Only functions related to the processes
identified in *C. elegans* (insulin, FOXO, ERK and mTOR signaling, autophagy and proteolysis) were chosen for the analysis.

To identify associations between rilmenidine’s effect in mice and biomarkers of lifespan-extending interventions and aging, we employed a GSEA-based algorithm developed in our previous study (Tyshkovskiy et al. 2019). First, for every trait, we specified significant genes, using the FDR threshold of 0.05. Among the remaining genes, we selected 1000 genes with the highest absolute logFC and divided them into up- and downregulated genes. These lists were considered as gene sets. Signatures of lifespan-extending interventions were taken from (Tyshkovskiy et al. 2019), while signatures of aging in liver and kidney were identified from GSE132040 dataset derived by Tabula Muris Consortium (Schaum et al. 2020). Then we calculated normalized enrichment scores (NES) separately for the up- and downregulated lists of each gene set as described in (Lamb 2006) and defined the final NES as a mean of the two.

To calculate the statistical significance of the NES, we performed a permutation test where we randomly selected gene sets of the same size. p-value of association was calculated as the frequency of observed final NES being bigger by absolute value than random final NES obtained from 5,000 random permutations. The resulting p-values were further adjusted for multiple hypothesis testing using Benjamini-Hochberg (BH) procedure (Benjamini & Hochberg 1995). Association was considered significant if the adjusted p-value was smaller than 0.1. To visualize the results of the association analysis, we converted adjusted p-values into significance scores as:

\[
\text{significance score} = - \log_{10}(\text{adj.pv}) \times \text{sgn}(\text{NES}),
\]

where adj.pv and NES are BH adjusted p-value and final NES, respectively.
**Author contributions**

All authors participated in analyzing and interpreting the data. DFB, JPdM, and CYE designed the experiments. DFB and CYE performed manual lifespan assays, and CS performed automated lifespan machine assays. AT and VNG performed mice experiments and analyses. All other experiments were performed by DFB. AG, CWB, CYE, and JPdM wrote the manuscript in consultation with the other authors.

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**Acknowledgment**

We thank Dominic Alcock and Alan Morgan for assistance and advice with the worm experiments. Further thanks to the de Magalhães and Ewald labs for critical feedback on the manuscript and to Wormbase for curating and updating the *nish-1* gene information. Some strains were provided by the CGC, which is funded by NIH Office of Research Infrastructure Programs (P40 OD010440). Graphic illustrations in Figure 2D and 6F were created with BioRender.com (PQ22YK2PVV and LT22YK2745). Funding from the Swiss National
Science Foundation 163898 and 190072 to CS, AG, and CYE, and from LongeCity and the Biotechnology and Biological Sciences Research Council (BB/R014949/1) to JPM.
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