

Influence of oral supplementation with sesamin on longevity of *Caenorhabditis elegans* and the host defense

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1 **Abstract**

2 *Purpose* Nutritional control has been proposed as a potential therapy for slowing the senescence of immune
3 function and decreasing mortality. This study investigated whether sesamin could modify host defense
4 systems and extend the lifespan of the nematode *Caenorhabditis elegans*.

5 *Methods* Nematodes were fed standard food (the bacterium *Escherichia coli* strain OP50) supplemented
6 with various doses of sesamin/ γ -cyclodextrin inclusion compounds starting from young adulthood. The
7 mean lifespan, muscle function, lipofuscin accumulation, protein carbonyl content, and stress resistance of
8 the worms were examined. Then, *C. elegans* mutants harboring loss-of-function lesions in longevity- and
9 host defense-related signaling pathways were supplemented with sesamin to identify the genes involved in
10 the longevity effects.

11 *Results* Worms supplemented with sesamin displayed higher locomotion and prolongevity, and produced
12 offspring at levels similar to unsupplemented control animals. The growth curves of nematodes were
13 similar to those of controls, suggesting that sesamin did not induce prolongevity effects through dietary
14 restriction. Notably, sesamin made the worms more resistant to infection by *Legionella pneumophila* and
15 more resistant to oxidative stressors such as paraquat and hydrogen peroxide, and prolonged the lifespan of
16 a *mev-1* mutant that produces abundant superoxide anions. However, the accumulation of protein carbonyls
17 and lipofuscin was similar in sesamin-exposed and control worms, suggesting that sesamin is unlikely to
18 work simply as an antioxidant. Sesamin supplementation failed to extend the lifespan of loss-of-function
19 mutants of *daf-2*, *daf-16*, *pmk-1*, and *skn-1*.

20 *Conclusions* Sesamin enhances the host defense of *C. elegans* and increases the average lifespan via
21 activation of both *skn-1* (encoding a component of the p38 MAPK pathway) and *daf-16* (encoding a
22 component of the IGF-1 pathway).

23

24 **Keywords:** Longevity, Nematodes, Sesamin, Aging, Host defense

25

1 **Introduction**

2

3 Senescence of the immune system leads to increased infections, malignancy, and autoimmunity [1-3].
4 Indeed, the mortality of elderly people suffering from viral and bacterial infections is higher than that of
5 younger people [4]. If the senescence of host defense could be slowed through immunonutritional
6 intervention, then the mortality due to infectious diseases might be decreased. However, the difficulty of
7 establishing a suitable animal model has limited studies on this topic. Although an increase of biomarkers
8 related to immunological function has been observed in the elderly with improved nutrition [5], few reports
9 have shown beneficial influences of nutrition on host defense, response to experimental stress, or lifespan
10 [6-8].

11 Lignans constitute the major class of polyphenols in sesame seeds, and sesamin, sesamol, and
12 sesaminol are the most abundant of the lignans. Sesamin, with the highest abundance among the lignans, is
13 particularly noteworthy since the compound has been associated with a variety of health effects such as
14 anti-allergenic [9], anti-carcinogenic [10], anti-hypertensive [11], and hypocholesterolemic [12] effects, as
15 well as decreases in fatty acid synthesis [13]. Although sesamin itself does not exhibit antioxidative activity
16 *in vitro*, the beneficial effects of the compound are likely to be due to the antioxidative effects of sesamin
17 derivatives generated during *in vivo* metabolism [14].

18 *Caenorhabditis elegans* is a small, free-living, bacteriophagous soil nematode. This organism has been
19 extensively used as an experimental system for biological studies because of its morphological simplicity,
20 transparent body, ease of cultivation, and amenability to genetic analysis. In addition, the short and
21 reproducible lifespan of *C. elegans* is particularly advantageous for aging studies [15]. We previously
22 showed that the nematode was a convenient model for investigating the senescence of host defense and the
23 influence of food and nutrition on the system: an opportunistic human pathogen, *Legionella pneumophila*,
24 was pathogenic only in old nematodes but not in young worms, and the presence of probiotic bacteria or
25 orally supplemented antioxidant agents prolonged *C. elegans* lifespan and enhanced the host defense
26 [16-19].

27 Here, we report that sesamin caused prolongevity in *C. elegans*. Although a positive effect of sesamin
28 on longevity has been reported in *Drosophila* [20], the compound's prolongation of lifespan is more
29 apparent in the nematode than in fruit flies. Caloric restriction is known to extend the lifespan of a wide
30 range of organisms from yeasts to mammals [21], although the effect of caloric restriction on lifespan
31 remains controversial in primates [22]. Based on the evaluation of worm biomarkers, including body and
32 brood sizes [23], we assessed whether sesamin extended the lifespan via caloric restriction. While radical

1 oxygen species (ROS) are recognized as key factors causing senescence because of the properties as
2 oxidants [24], ROS are still considered to contribute to anti-senescence by the hormesis effect [25]. Hence,
3 we performed assays for muscle function, lipofuscin accumulation (gut autofluorescence), protein carbonyl
4 content, and stress resistance to determine how sesamin induced the numerous beneficial effects on
5 physiological function and longevity. Additionally, loss-of-function *C. elegans* mutants were grown on
6 medium supplemented with sesamin to identify the genes involved in the longevity-promoting effects of the
7 compound in *C. elegans*.

10 **Materials and methods**

12 Nematode strains

14 *C. elegans* Bristol strain N2 and its derivative mutant strains were kindly provided by the Caenorhabditis
15 Genetics Center, University of Minnesota. The mutants used in this study were KU25 *pmk-1(km25)*, EU1
16 *skn-1(zu67)*, CF1038 *daf-16(mu86)*, CB1370 *daf-2(e1370)*, TK22 *mev-1(kn1)*, and TJ356, a strain
17 harboring a *daf-16::GFP* transgene. Nematodes were maintained and propagated on nematode growth
18 medium (NGM) according to standard techniques [26].

20 Bacterial strains

22 *E. coli* OP50 (OP) was used as the standard feed for nematode cultivation and was grown using tryptone
23 soya agar (Nissui Pharmaceutical, Tokyo, Japan). Cultured bacteria (100 mg wet weight) were suspended in
24 0.5 ml M9 buffer, and 50 μ l of the resulting bacterial suspension then was spread on peptone-free modified
25 NGM (mNGM) in 5.0-cm diameter plates to feed worms. *Salmonella enterica* subsp. *enterica* serovar
26 Enteritidis strain SE1, originally isolated from a diarrheal specimen, and *Legionella pneumophila*
27 serogroup 1 strain JR32 were used as pathogens and were cultured using tryptone soya agar (Nissui) and
28 BCYE agar plates (Oxoid, Basingstoke, UK), respectively as described previously [16, 18].

30 Formulation of sesamin

32 Inclusion compound formulations of γ -cyclodextrin (γ CD) and sesamin were prepared as reported

1 previously [17]. Briefly, a sterile γ CD solution was prepared by filtering a nearly saturated γ CD (230
2 mg/ml) solution. Sterile sesamin ethanol solution (2.5 mg/ml) was prepared by filtration through an organic
3 solvent-resistant filter (SLLG013SL, Millipore, Carrigtwohill, Ireland). One milliliter γ CD solution was
4 mixed with 0.1 ml sesamin solution and stirred with a rotary mixer for 12-24 h at ambient temperature. The
5 solid complex (inclusion compounds) was collected by centrifugation and weighed. Finally, the inclusion
6 compound was suspended in M9 buffer and vortexed so that the nematodes could ingest the compounds.
7 Approximately 1 mg wet weight of the compound, containing 6.3 μ g sesamin, was spread onto each NGM
8 plate.

9

10 Determination of *C. elegans* life span

11

12 Eggs were recovered from adult *C. elegans* worms after exposure to a sodium hypochlorite/sodium
13 hydroxide solution, as previously described [27]. The egg suspension was incubated overnight at 25 °C to
14 allow hatching, and the resulting suspension of L1 stage worms was centrifuged at $156 \times g$ for 1 min. After
15 removing the supernatant by aspiration, the remaining larvae were transferred onto fresh mNGM plates
16 covered with OP and then incubated at 25 °C, with the exception of the *daf-2* mutant, which was
17 maintained at 20 °C. To synchronize pubescence, worms were allowed to feed on OP for 2 days until
18 maturation, as it is well known that the reproductive system regulates aging in *C. elegans* [28].
19 Nematocidal assays were performed by adding 35 three-day-old adult worms to each mNGM plate covered
20 with OP and the sesamin inclusion compound. The plates were incubated at 25 °C, and the numbers of live
21 and dead worms were scored every 24 h. At 25 °C, worms produce progeny that develop into adults in 3
22 days and it is therefore difficult to identify the original worms. To avoid over-estimating the number of
23 living worms, the original worms were transferred daily to fresh mNGM plates for 4 days until completion
24 of their egg-laying phase at 7 days of age. Thereafter the worms were transferred to fresh mNGM plates
25 every second day. A worm was considered dead when it failed to respond to a gentle touch with a worm
26 picker. Worms that crawled off the plate or died from internal hatching were considered lost and not
27 included in the analysis [29]. Nematocidal assays are generally performed using NGM agar plates
28 containing peptone, which allows the overlaid bacteria to proliferate. However, the composition of NGM
29 has been reported to influence the virulence of bacteria, and the *in-situ* production of metabolites by
30 bacteria growing on the medium also may be nematocidal [30]. Thus, to exclude the possibility of
31 bacteria-induced nematocidal effects from nutrients in the medium, the nematocidal assays were performed
32 on mNGM plates lacking peptone. Each assay was carried out in duplicate and repeated twice unless

1 otherwise stated.

2 Mean lifespan was estimated using the formula [31]:

$$3 \quad \text{MLS} = \frac{1}{N} \sum_j \frac{x_j + x_{j+1}}{2} d_j$$

4 where d_j is the number of worms that died in the age interval (x_j to x_{j+1}), and N is the total number of
5 worms. The standard error of the estimated mean lifespan was calculated using the equation:

$$6 \quad \text{SE} = \sqrt{\frac{1}{N(N-1)} \sum_j \left(\frac{x_j + x_{j+1}}{2} - \text{MLS} \right)^2 d_j.}$$

7 Maximum lifespan was calculated as the mean lifespan of the longest-living 15% of worms in each group.

8

9 Effect of sesamin on infections

10

11 After hatching, nematodes were grown on OP for 3 days. The adult worms then were divided into two
12 groups that were given γ CD only or were supplemented with a γ CD inclusion compound containing
13 sesamin. Seven- or 8-day-old worms were fed *Salmonella* or *Legionella*, respectively, instead of OP, as
14 reported previously [16, 18]. Nematode survival was measured as described above.

15

16 Locomotory scoring of aging nematodes

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18 The motility of worms at different ages was examined using a scoring method described in previous reports
19 [32, 33]. Briefly, worms were classified according to a four-point scale: class “A” worms showed
20 spontaneous movement or vigorous locomotion in responding to prodding; class “B” worms did not move
21 unless prodded or appeared to have uncoordinated movement; class “C” worms moved only their head
22 and/or tail in response to prodding; and class “D” worms were dead animals. A minimum of 60 worms
23 provided with sesamin were scored.

24

25 Lipofuscin

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27 To obtain indices of senescence, the autofluorescence of intestinal lipofuscin was analyzed from days 10 to
28 18 of adulthood. Briefly, randomly selected worms were washed five times in M9 buffer and then placed
29 onto fresh mNGM plates coated with 50 mM sodium azide to induce anesthesia. Lipofuscin

1 autofluorescence was measured using a M165 FC fluorescence stereomicroscope (Leica Microsystems,
2 Tokyo, Japan) equipped with a DsRED filter set (excitation, 510-560 nm; emission, 590-650 nm) and a
3 Leica DFC425 C digital microscope camera. The captured data was analyzed using Leica Application Suite
4 imaging software (Version 3.7.0).

5

6 Protein carbonyl formation

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8 Protein carbonyl groups, which served as biomarkers of oxidative stress, were measured as reported
9 previously [34].

10

11 Measurement of body size

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13 Three-day-old adult worms were placed on mNGM plates covered with OP. The body sizes of live worms
14 were measured every 24 h until the animals reached 7 days of age. Images of adult nematodes were taken
15 with a VCT-VBIT digital microscope (Shimadzu, Kyoto, Japan) and analyzed using ImageJ software. In
16 this system, the area of a worm's projection was estimated automatically and used as an index of body size.

17

18 Brood size

19

20 Eggs isolated with a sodium hypochlorite/sodium hydroxide solution were allowed to develop to 3 days of
21 age on mNGM plates coated with OP at 25 °C. Three hermaphrodites were selected and transferred to an
22 mNGM plate covered with a lawn of BI. The parental animals were transferred every 24 h to fresh mNGM
23 plates until the end of the reproductive period. The resulting progeny were left to develop for 3 days and the
24 progeny number then was determined. Each assay was performed with five plates and repeated twice.

25

26 Stress resistance assays

27

28 Worms were grown from 3 to 7 days of age on mNGM plates with or without sesamin and then subjected to
29 oxidative stress and heat shock assays. To conduct an oxidative stress assay, worms were transferred onto
30 mNGM containing 1.0 or 2.5 mM paraquat, or into M9 buffer with 0.1% cholesterol (5 mg ml⁻¹ in ethanol)
31 that contained either 0.8 mM hydrogen peroxide, 2.0 mM hydrogen peroxide, or 7.0 mM cupric chloride.
32 Plates or suspensions of worms were incubated at 25 °C, and viability then was scored. To assess thermal

1 tolerance, 7-day-old worms were placed in M9 buffer or onto mNGM at 35 °C and then scored for viability.
2 Similarly the 7-day-old worms were exposed to UV irradiation at 250 J / m² or 500 J / m². The survival of
3 worms was determined by touch-provoked movement. Worms were scored as dead when the animals failed
4 to respond to mechanical stimuli with a worm picker. The assays were performed at least twice.

5 6 Statistical analysis

7
8 Nematode survival was calculated by the Kaplan-Meier method, and survival differences were tested for
9 significance using the log-rank test. Differences in protein carbonyl levels and brood size were determined
10 using the Student *t*-test. Differences in lipofuscin levels were analyzed using the Mann-Whitney U test.
11 Differences in body sizes were analyzed using the Kruskal-Wallis test. The level of significance was set at P
12 < 0.05.

13 14 15 **Results**

16 17 Prolongevity due to sesamin

18
19 The lifespan of nematodes that ingested sesamin-containing γ CD was 13.6% longer than that of control
20 worms maintained on mNGM containing γ CD alone (Fig. 1; the data for Fig. 1 is provided in Online
21 Resource Table 1). The effect was dose-dependent with the strongest effect seen at 6.3 μ g / plate;
22 supplementation at higher or lower concentrations of sesamin resulted in weaker or no prolongevity effects.

23 Muscle function and accumulation of lipofuscin, as determined by gut autofluorescence, are known to
24 correlate with age, but vary between age-matched individuals in *C. elegans* [23]. In the present study,
25 locomotory ability was assayed as an indicator of muscle function, as locomotory class is predictive for the
26 remaining lifespan of *C. elegans* worms after 8 days of age [32]. The ratio of worms displaying coordinated
27 sinusoidal locomotion (class A) was apparently higher in the group supplemented with sesamin (Fig. 2).

28 Autofluorescence of lipofuscin, a lipid peroxidation product, in worms supplemented with sesamin was
29 not significantly different from that in control worms (Online Resource Fig. 1). The abundance of protein
30 carbonyls, which is a general biomarker of protein oxidation, in the extracts from worms given sesamin
31 also was similar to that in control worms, and the values were not significantly different (Online Resource
32 Fig. 2a, b).

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Body and brood sizes

To examine whether the prolongevity effects of sesamin were a result of caloric reduction, the body and brood sizes of nematodes supplemented with sesamin were compared with those of control worms fed OP. The supplementation with sesamin did not alter the growth curve of worms (Fig. 3). Although caloric restriction reportedly extends the lifespan by reducing brood size [35], supplementation with sesamin did not decrease the brood size of *C. elegans* (Fig. 4).

Stress resistance assays

The effects of sesamin on resistance to chemical, physical, and biological stresses were examined. Sesamin supplementation (at 6.3 µg / plate) yielded a highly significant (P <0.001 compared to unsupplemented animals) increase in survival against 1.0 mM paraquat, a known source of ROS (Fig. 5a, Online Resource Table 2a). Similarly, when 9-day-old or 15-day-old worms were maintained in M9 buffer containing 2.0 mM hydrogen peroxide, a known source of active oxygen, the sesamin (at 6.3 µg / plate) -supplemented worms exhibited a highly significant (P <0.001 compared to unsupplemented animals) increase in survival time (Figs. 5b and 5c, Online Resource Tables 2b and 2c). However, the worms remained vulnerable to 7.0 mM CuCl₂ solution irrespective of the age and sesamin supplementation status (Online Resource Figs. 3a and 3b, Online Resource Tables 2d and 2e). In the case of physical stresses (Online Resource Figs. 4a to 4d, Online Resource Tables 3a and 3b), sesamin did not prolong the survival time of worms subjected to heat stress (8-day-old worms at 32 °C or 7-day-old worms at 35 °C) or to UV irradiation (9-day-old worms exposed to 250 or 500 J / m²)

Previously we found that the opportunistic pathogen *Legionella* can be virulent in worms older than 8 days old, while young worms fed on the pathogen grow as well as those fed on the standard food bacteria OP [18]. Thus, the opportunistic infection of *C. elegans* by *Legionella* can serve as a unique model to investigate immunosenescence. To study the effect of sesamin on response to biological stress, 7- or 8-day-old worms were fed *Salmonella* or *Legionella*, respectively, instead of OP. Oral supplementation with sesamin failed to enhance the host defense to *Salmonella*, a bacterial strain that exhibits pathogenicity irrespective of age of the nematodes (Online Resource Fig. 5, Online Resource Table 4a). However, when nematodes were exposed to the opportunistic pathogen *Legionella*, sesamin protected against death from the infection (Fig. 6, Online Resource Table 4b).

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Prolongevity effects in mutants

The PMK-1 (p38 mitogen-activated protein kinase; p38 MAPK) and DAF-2/DAF-16 insulin/IGF-1 signaling (IIS) pathways contribute to homeostasis, including host defense against pathogens in *C. elegans*. To investigate whether these two pathways are involved in the prolongevity effects of sesamin, the lifespan of *C. elegans* loss-of-function mutants fed sesamin was compared with that of control worms fed OP. Sesamin failed to prolong the lifespan of strains harboring individual mutations in the *pmk-1* (encoding an ortholog of mammalian p38 MAPK), *skn-1* (encoding an ortholog of mammalian Nrf2), *daf-2* (encoding an ortholog of Insulin/IGF-1-like receptor), or *daf-16* (encoding an ortholog of mammalian FoxO) genes (Online Resource Figs. 6a to 6d, Online Resource Tables 5a to 5d). In contrast, sesamin prolonged the mean lifespan of the *mev-1* mutant by 13% (Online Resource Fig. 6e, Online Resource Table 5e), an increase that was statistically significant ($P < 0.05$). These results suggested the importance of both p38 MAPK and IIS pathways for mediating the prolongevity effect of sesamin. When a strain harboring a *daf-16::GFP* transgene was grown in the presence of sesamin, the DAF-16::GFP transcription factor exhibited translocation into the nuclei of foregut cells (Fig. 7a); although the translocation was not as intense as that observed in worms of the same strain exposed to heat stress in the absence of sesamin supplementation (Fig. 7b).

Discussion

Sesamin, a hydrophobic substance and precursor of the antioxidative sesaminol, clearly extended the lifespan of the nematodes in this study. The worms given sesamin were more locomotive than the controls during old age, and were more resistant to *L. pneumophila*, a species that is selectively pathogenic for aged worms. Further, sesamin protected worms from oxidative stresses produced by selected chemicals (paraquat and peroxide), though not for that associated with cupric chloride. Sesamin supplementation was not protective against physical stresses such as UV irradiation and heat stress, nor against pathogenic *Salmonella*. In the course of developing methods to let *C. elegans* ingest hydrophilic or hydrophobic chemicals, we previously had observed that several antioxidants (including N-acetyl-cysteine, glutathione, tocotrienol, and astaxanthin) prolonged the lifespan of nematodes [17, 19, 36]. In *mev-1* mutants, shortened lifespan is considered to be the result of an increased production of ROS [37]. Our observation, in the

1 present work, of increased longevity in sesamin-treated *mev-1* animals is consistent with the proposed role
2 of ROS in this mutant strain.

3 It is an attractive hypothesis that sesamin works as an antioxidant precursor, slowing senescence
4 compared to control animals, and thereby resulting in prolongevity and increased resistance to the
5 opportunistic pathogen *Legionella*. Oxidative stress is typically associated with protein modifications
6 (including carbonylation of amino acid side chains), and the accumulation of lipofuscin serves as a marker
7 of aging. However, there were no significant differences in the levels of lipofuscin or carbonylated protein
8 between sesamin-supplemented worms and age-matched controls. Thus, sesamin anti-oxidant activity does
9 not appear to affect oxidants produced endogenously in the worms. Sesamin may reduce harm caused by
10 exogenous oxidants; although it remains unclear why the supplementation failed to yield protective effects
11 against copper chloride, a compound that is expected to induce exogenous ROS production [38, 39]. If
12 sesamin does not play a direct role as an antioxidant, sesamin is expected to render worms indirectly
13 resistant to the oxidative chemicals used in this study. Among these indirect roles, antioxidants can work as
14 pro-oxidant hormetins [40], a mechanism that is not expected to decrease the levels of carbonylated protein
15 and lipofuscin. Instead, the resultant hormesis would enhance the host defense via signal induction
16 pathways, in the same way that low levels of ROS can cause so-called redox signaling [41, 42]. This
17 hormetic effect would explain why higher doses of sesamin were not as effective in extending worm
18 lifespan.

19 Caloric restriction is well recognized as a mechanism for the extension of longevity, and such
20 restriction has been shown to slow senescence not only in numerous non-mammalian taxa, but also in
21 mammals [21]. However, sesamin does not appear to cause dietary restriction in nematodes, as evidenced
22 by the fact that growth curves were similar regardless of sesamin exposure. Furthermore, the
23 sesamin-supplemented nematodes produced offspring in numbers similar to those seen in the control
24 worms. This observation confirms our hypothesis that sesamin's effects are not mediated by effects on
25 dietary restriction, given that Bishop and Guarente [5] demonstrated that the prolongevity effect of dietary
26 restriction reflects reductions in brood sizes. Indeed, if sesamin were causing caloric restriction, the
27 compound should extend the lifespan of a *C. elegans daf-2* mutant [35], which we did not observe in the
28 present work.

29 The IIS pathway has a conserved role in modulation of lifespan, and the pathway includes DAF-16, a
30 forkhead family transcription factor that regulates genes that promote stress resistance and extend the
31 lifespan of nematodes [43]. Since sesamin failed to prolong the lifespan of the *daf-2* or *daf-16* mutants, this
32 pathway is presumably involved in the prolongevity effects of sesamin. Similarly, sesamin failed to prolong

1 the lifespan of the *C. elegans pmk-1* and *skn-1* mutants, suggesting that these genes, components of the p38
2 MAPK pathway, mediate the prolongevity effect of the compound. Thus, sesamin appears to promote the
3 induction of phase-2 detoxification enzymes in *C. elegans* through activation of SKN-1, an ortholog of
4 mammalian Nrf transcription factors [44]. As SKN-1 appears to be involved in damage control and stress
5 resistance [45], endogenous antioxidant systems upregulated via activated SKN-1 may account for the
6 enhanced resistance to stressors and the extension of lifespan. Both the p38 MAPK and IIS pathways are
7 evidently required for the effects of sesamin (Figure 8).

8 Inhibition of the TOR (target of rapamycin) is another well-known intervention method for
9 prolongevity: dietary restriction / caloric restriction is likely to extend lifespan by inhibiting the TOR
10 kinase. This protein, which is essential for growth but associated with aging, forms two distinct complexes,
11 designated TORC1 and TORC2. Recently, Robida-Stubbs *et al.* [38] showed that TORC1 inhibition
12 resulted in extension of lifespan in *C. elegans*, a process that depended on both the p38 MAPK and IIS
13 pathways. In contrast, while TORC2 inhibition also increased life span, this process required SKN-1 but
14 not DAF-16 [46]. We previously observed that the presence of bifidobacteria extended lifespan in *C.*
15 *elegans* and suppressed the age-associated increase of the sensitivity to *Salmonella* or *Legionella* [16, 18].
16 Unlike sesamin, however, bifidobacteria elicit their effects mainly via the p38 MAPK pathway, and not via
17 the IIS pathway. The lower lipofuscin and protein carbonyl levels in the worms fed bifidobacteria were
18 taken as evidence of enhanced antioxidant systems, although the bifidobacteria-eating worms were as
19 vulnerable to the chemical and physical stresses used in that study as the control worms. The distinction
20 between the effects of sesamin and those of bifidobacteria might reflect differential effects on the TOR
21 pathway. In this model, sesamin would be a mimetic of caloric restriction, which is known to decrease the
22 activity of TORC1.

23 In conclusion, the present study suggests that sesamin extends the lifespan of *C. elegans* via
24 modulation of both the p38 MAPK and IIS pathways, possibly by suppression of TORC1, although this
25 effect is not mediated by caloric restriction *per se*. We note that sesamin seems to function as a hormetin
26 rather than as an antioxidant, and that the compound enhances the host defense of nematodes against a
27 subset of chemical oxidative stressors and opportunistic pathogens. Agents that promote particular SKN-1
28 or DAF-16 activities likely cause beneficial effects of TOR inhibition without interfering with critical TOR
29 functions and inducing side effects, in contrast to rapamycin, which is known to impair immune function
30 and reduce insulin sensitivity. The role of sesamin as a possible suppressor of TORC1 will be the subject of
31 further studies. We additionally propose that *C. elegans* will serve as a useful model for the identification
32 and characterization of other TOR inhibitors.

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Figure legends

Fig. 1. Survival curves of *C. elegans* fed OP supplemented with sesamin at different doses as compared with the lifespan of control worms fed only OP. Each plate contained 10 mg (wet weight) of bacteria. Worms were 3 days old on the graph's nominal Day 0. *, **, and *** indicate P values of <0.05, <0.01, and <0.001, respectively, for statistically significant differences between control worms fed only OP and worms fed OP supplemented with sesamin. Mean and maximum lifespan are presented in Online Resource Table 1.

Fig. 2. Locomotory activity of *C. elegans* fed OP supplemented with sesamin. Young adult worms fed OP for 2 days after hatching were transferred to plates containing 10 mg of OP with or without sesamin at 6.3 µg / plate. Worms were 3 days old on the nominal Day 0. Animals were classified into four classes based on their locomotion: class A, robust, coordinated sinusoidal locomotion (white bars); class B, uncoordinated and/or sluggish movement (light gray bars); class C, no forward or backward movement, but head movements or shuddering in response to prodding (dark gray bars); and class D, dead animals (black bars). The rates of each class at the indicated time point are indicated.

Fig. 3. Growth curve of worms fed OP supplemented with sesamin (6.3 µg / plate). Images of adult nematodes were recorded using a digital microscope; the area of the worm's projection was measured using image processing software and used to generate an index of body size. The body sizes of the worms fed BI at all ratios were similar to those of the control worms fed only OP.

Fig. 4. Brood size of worms fed OP supplemented with sesamin (6.3 µg / plate). Total brood size was determined from 16 animals and the mean per two worms was calculated.

Fig. 5. Influence of sesamin-supplementation on susceptibility of worms to oxidative stresses. Worms were provided with sesamin at 6.3 µg / plate from 3 days of age and then incubated with (a) 1.0 mM paraquat from 7 days of age, (b) 2 mM hydrogen peroxide from 9 days of age, or (c) 2 mM hydrogen peroxide from 15 days of age. ***, Statistically significant difference compared to control worms fed OP only (P <0.001). Mean and maximum lifespan are presented in Online Resource Tables 2a to 2c.

Fig. 6. Survival of *C. elegans* infected with pathogenic bacteria. Worms were fed OP with or without sesamin (6.3 µg / plate) from 3 days of age. The nematodes were transferred to agar plates covered with *L. pneumophila* virulent strain JR32 at 7 or 8 days of age, respectively. The survival curves were compared with that of worms not supplemented with sesamin. *** Statistical significance at P < 0.001. Mean and maximum lifespan are presented in Online Resource Table 4a.

Fig. 7. Effects of sesamin on translocation of the DAF-16::GFP transcription factor. When the *daf-16::GFP*

mutant was supplemented with sesamin (6.3 μg / plate), translocation of the DAF-16::GFP transcription factor into the nucleus was observed at 8 days of age (panel a). Note that translocation into the nucleus was limited to cells around the foregut and was of lower intensity than that seen in animals subjected to heat stress (37 °C for 1 h; panel b).

Fig. 8. Pathways predicted to be involved in the longevity effect of sesamin. Red lines indicate signals that upregulate or activate genes, and black lines indicate signals involved in suppression. Established pathways are shown by solid lines. Dotted lines suggest hypothetical pathways that remain to be elucidated.

Online Resource Figure legends

Online Resource Fig. 1. Lipofuscin accumulation in the intestine of nematodes. Intestinal autofluorescence from lipofuscin in age-synchronized worms on days 17, 19, and 21 of adulthood was quantified using ImageJ software to determine the lipofuscin levels. Each bar represents the mean value for an n of ten worms, except for the bar corresponding to day 18 (n=6 OP-fed worms). The bar graph depicts the percent difference of the mean value in arbitrary units relative to that of control (OP-fed) worms on day 17. There was no statistically significant difference from control worms fed OP without sesamin (6.3 μg / plate) at a P value of < 0.05 . Error bars represent the SE.

Online Resource Fig. 2. Influence of sesamin on protein oxidation evaluated using an OxyBlot kit. Non-derivatized proteins were used as negative controls for each sample. (a) Worms were grown in the absence or presence of sesamin (6.3 μg / plate); proteins recovered from lysed animals were labeled with DNP solution (Oxyblot) to detect the presence of protein carbonyls and separated on SDS-PAGE gels (lanes labeled OP and sesamin, respectively). After electrophoresis, the proteins were transferred to a PVDF membrane and visualized using a chemiluminescence detection assay. The membrane was re-probed with anti-actin antibody to correct for loading variation. (b) Band densities were quantified to determine the protein carbonyl levels using ImageJ software. The bar graph depicts the percent difference of the mean value of five independent experiments in arbitrary units relative to that of control worms fed only OP; error bars represent the SE. Actin or intensity of Coomassie Brilliant Blue R-stained bands was used to compensate for loading variation.

Online Resource Fig. 3. Influence of sesamin-supplementation on susceptibility of worms to oxidative stresses. Worms were provided with sesamin at 6.3 μg / plate from 3 days of age and then incubated with (a) 7.0 mM CuCl_2 solution from 9 days of age, or (b) 7.0 mM CuCl_2 solution from the 15 days of age. ***, Statistically significant difference compared to control worms fed OP only ($P < 0.001$). Mean and maximum lifespan are presented in Online Resource Tables 2d to 2e.

Online Resource Fig. 4. Influence of sesamin-supplementation on susceptibility of worms to physical stresses. Worms were provided with sesamin (6.3 μg / plate) from 3 days of age and then (a) incubated at 32 $^{\circ}\text{C}$ for 24 h at 8 days of age, (b) at 35 $^{\circ}\text{C}$ for 8 to 10 hours at 7 days of age, (c) exposed to UV irradiation at 250 J / m^2 , or (d) at 500 J / m^2 at 7 days of age. NS, not significant ($P > 0.05$). Mean and maximum lifespan are presented in Online Resource Tables 3a and 3b.

Online Resource Fig. 5. Survival of *C. elegans* infected with pathogenic bacteria. Worms were fed OP

with or without sesamin (6.3 µg / plate) from 3 days of age. The nematodes were transferred to agar plates covered with *Salmonella* Enteritidis at 7 days of age, respectively. The survival curves were compared with that of worms not supplemented with sesamin. Mean and maximum lifespan are presented in Online Resource Table 4b.

Online Resource Fig. 6. Effects of sesamin on the lifespan of *C. elegans* mutants. Survival curves of (a) *pmk-1(km25)*, (b) *skn-1(zu67)*, (c) *daf-2(e1370)*, (d) *daf-16(mu86)*, and (e) *mev-1(kn1)* mutant hermaphrodites fed OP with or without sesamin (6.3 µg / plate) from 3 days of age (nominal Day 0 in the figure). *Statistical significance at $P < 0.05$. Mean and maximum lifespan are presented in Online Resource Tables 5a to 5e.

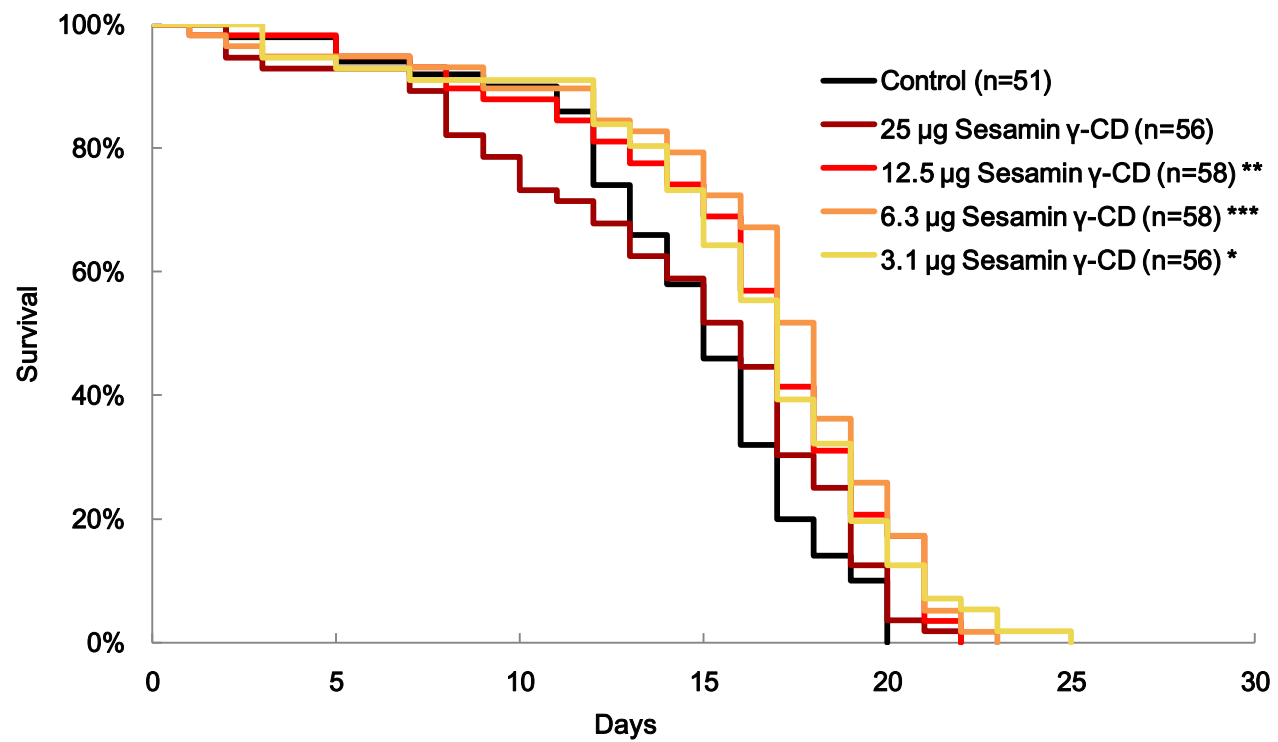


Fig. 1

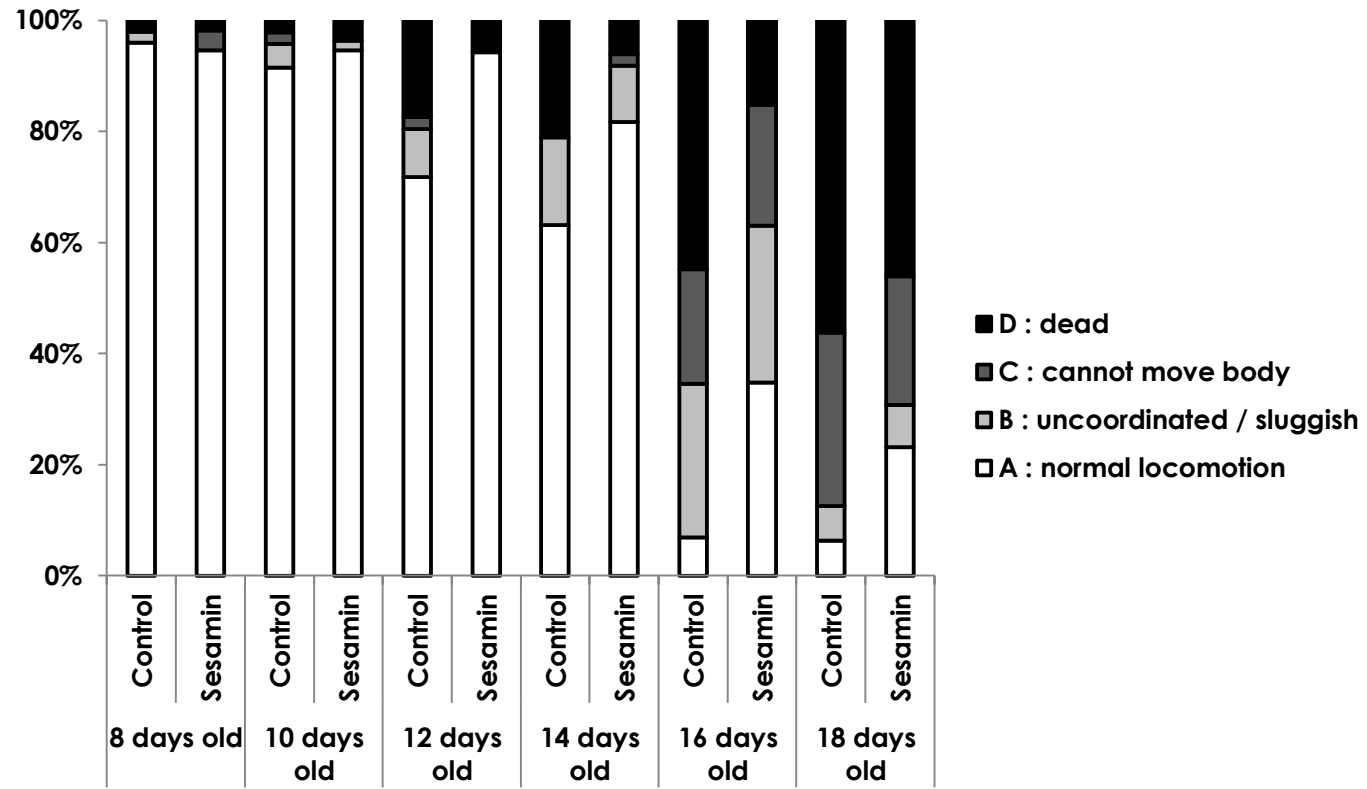


Fig. 2

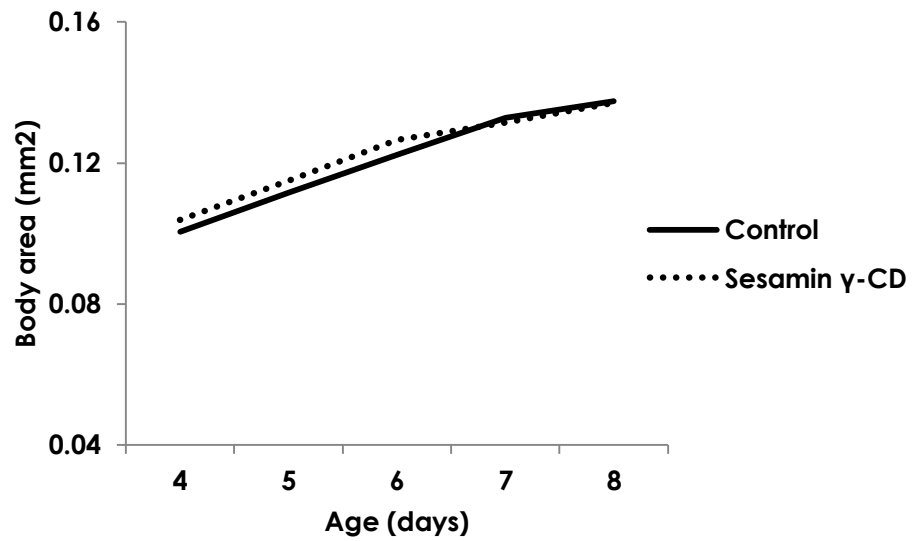


Fig. 3

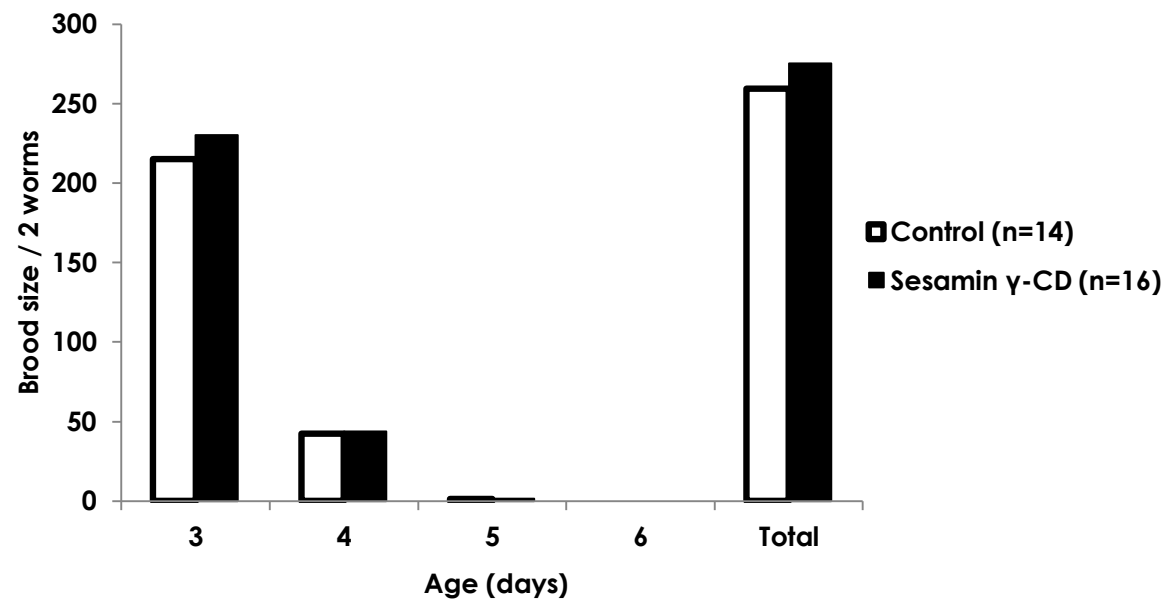


Fig. 4

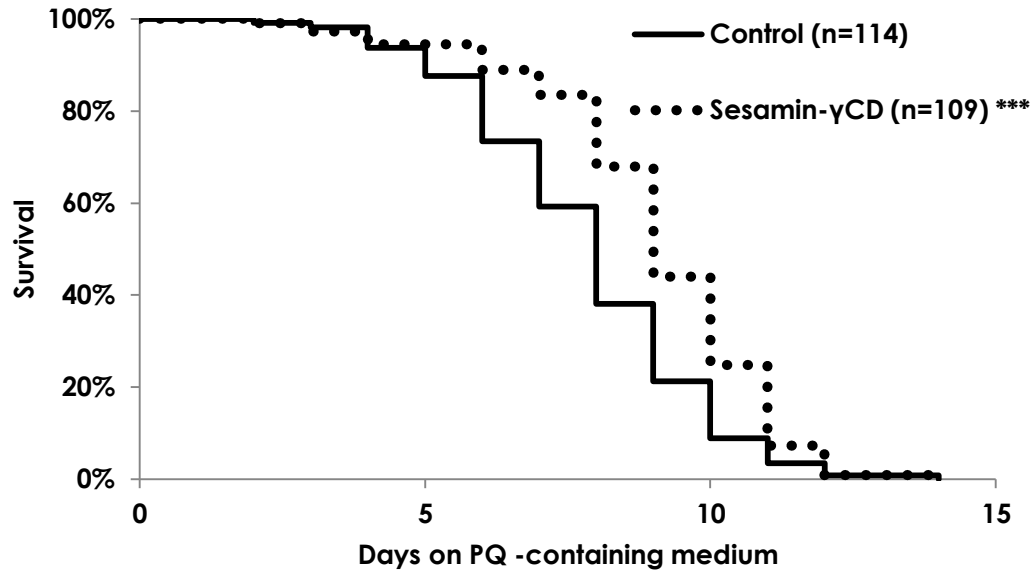


Fig. 5a

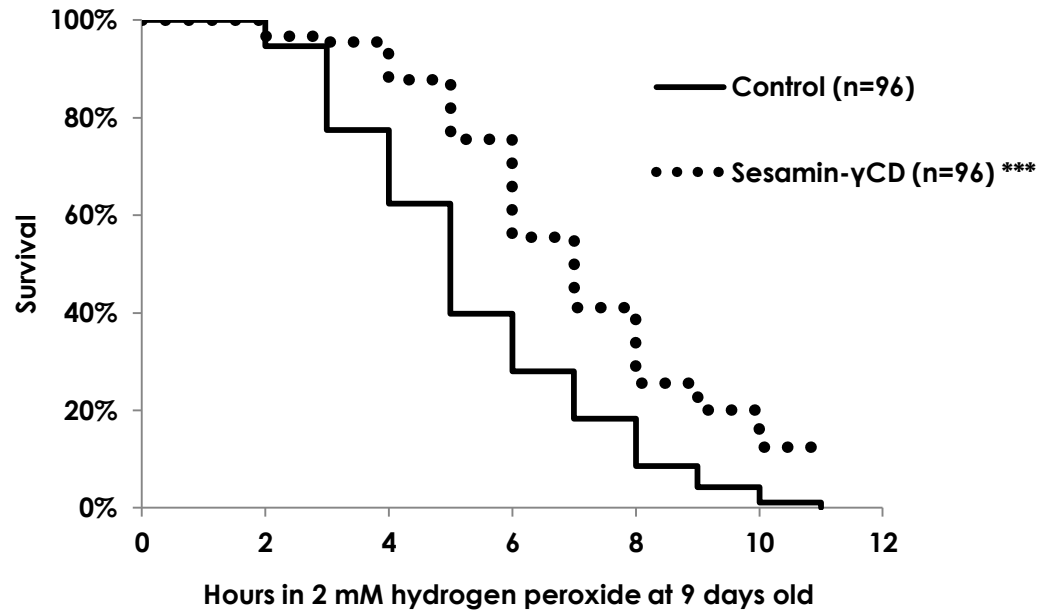


Fig. 5b

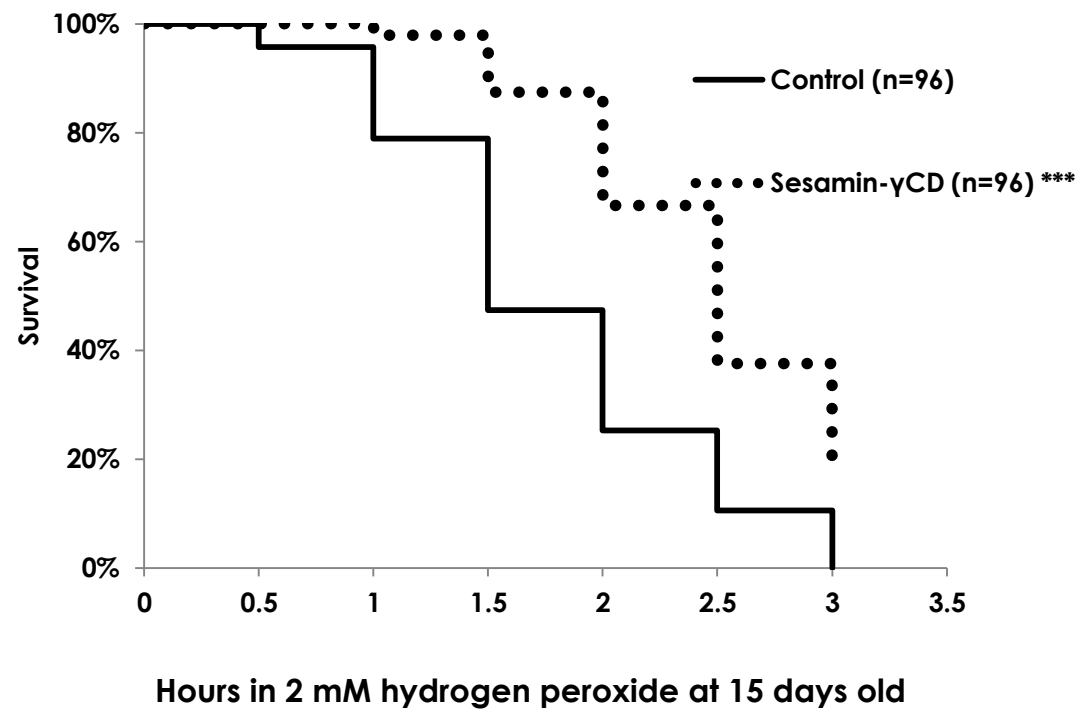


Fig. 5c

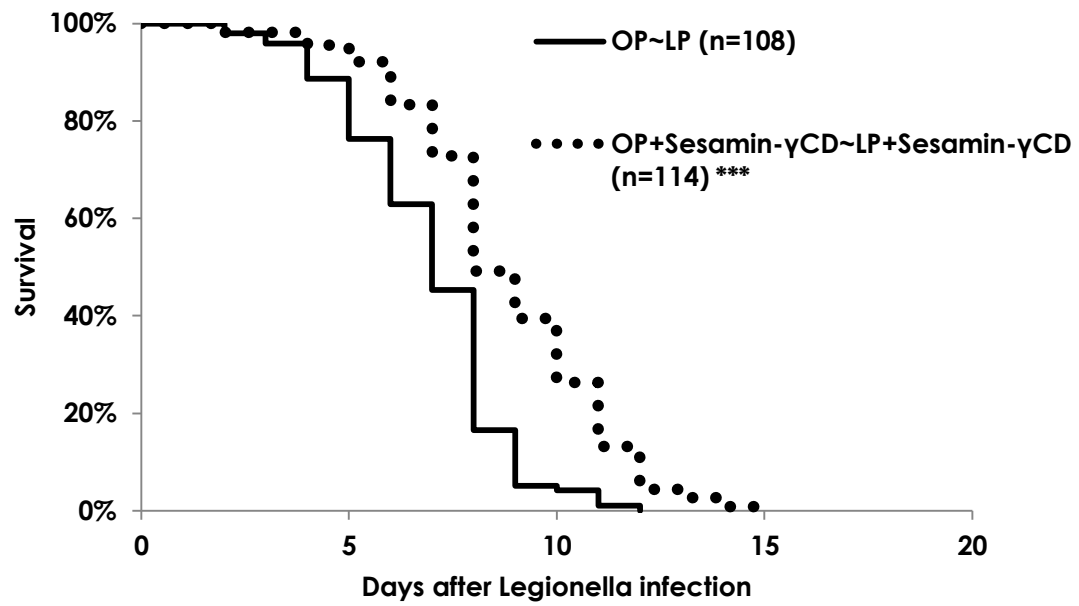


Fig. 6



Fig. 7a

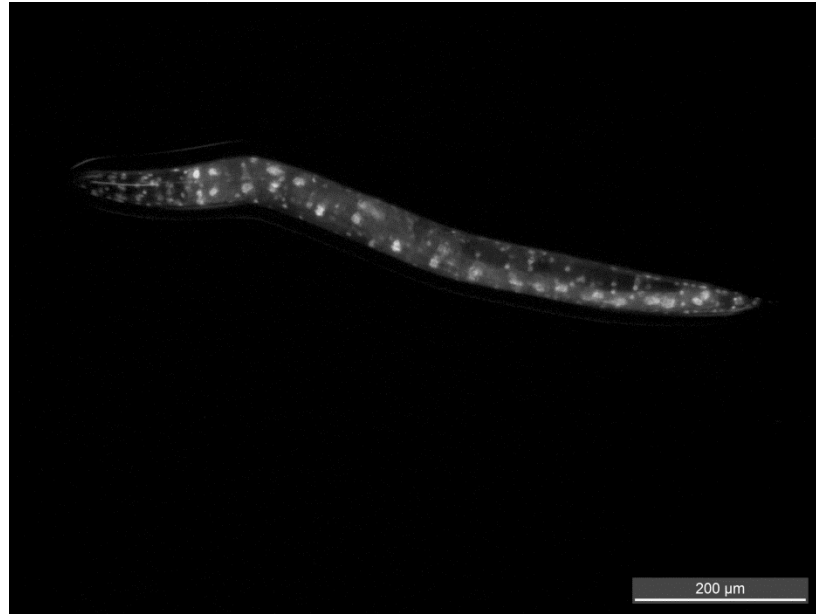


Fig. 7b

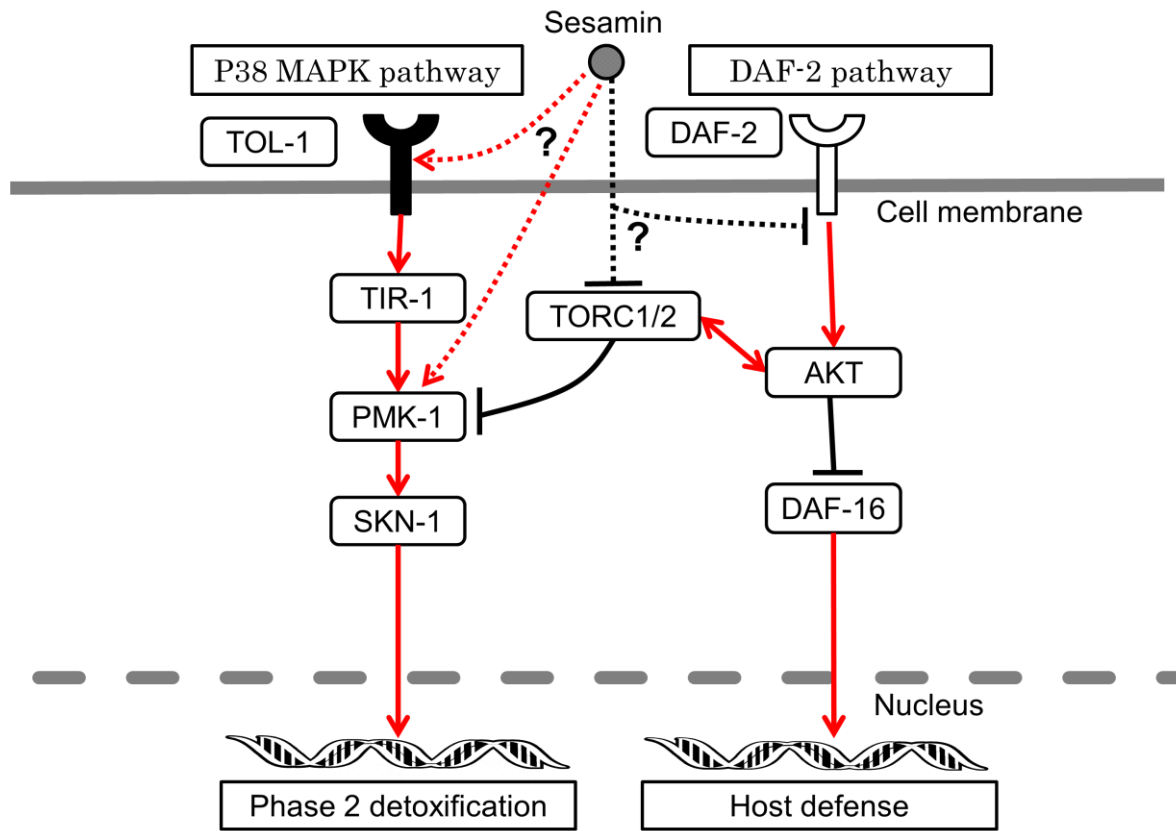


Fig. 8

Online Resource Table 1 for Fig. 1.

Dose of sesamin	No. of experiments ^a	Mean life span ± SE (days old)	Maximum life span ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control		17.56 ± 0.66	23.00 ± 0.27	51 (9)	
25 µg		17.70 ± 0.70	23.75 ± 0.31	56 (4)	0.45307
12.5 µg	1	19.48 ± 0.61	24.75 ± 0.16	58 (2)	0.00804
6.3 µg		20.03 ± 0.63	25.00 ± 0.20	58 (2)	0.00089
3.1 µg		19.41 ± 0.62	24.86 ± 0.39	56 (4)	0.01636

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 2a for Fig. 5a.

Group	No. of experiments ^a	Mean survival time ± SE (days old)	Maximum survival time ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control	2	15.30 ± 0.20	18.57 ± 0.28	114 (3)	0.0009
Sesamin		16.54 ± 0.20	19.17 ± 0.21	109 (3)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 2b for Fig. 5b.

Groups	No. of experiments ^a	Mean survival time ± SE (hours)	Max. survival time ± SE (hours)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control	2	5.79 ± 0.23	8.37 ± 0.13	96 (0)	0.00001
Sesamin		ND	ND	96 (0)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 2c for Fig. 5c.

Groups	No. of experiments ^a	Mean survival time ± SE (hours)	Max. survival time ± SE (hours)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control	2	2.53 ± 0.07	3.5 ± 0.11	96 (0)	0.00000
Sesamin		ND	ND	96 (0)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 2d for Supplementary Fig. 3a.

Groups	No. of experiments ^a	Mean survival	Max. survival	No. of	Log-rank test
		time	time	nematodes	p value
		± SE (hours)	± SE (hours)	(No. of lost worms) ^b	
Control	1	3.27 ± 0.19	4.93± 0.14	48 (0)	0.15891
Sesamin		4.31± 0.25	6.50 ± 0.21	48 (0)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 2e for Supplementary Fig. 3b.

Groups	No. of experiments ^a	Mean survival	Max. survival	No. of	Log-rank test
		time	time	nematodes	
		± SE (hours)	± SE (hours)	(No. of lost worms) ^b	p value
Control	1	1.69 ± 0.07	2.14± 0.19	48 (0)	0.35136
Sesamin		1.92± 0.09	2.79 ± 0.13	48 (0)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 3a for Supplementary Fig. 4c.

Groups	No. of experiments ^a	Mean survival	Maximum	No. of	Log-rank test
		time	survival time	nematodes	
		± SE (days old)	± SE (days old)	(No. of lost worms) ^b	p value
Control	1	15.76 ± 0.22	17.88± 0.26	54 (1)	0.30640
Sesamin		16.33± 0.18	18.75 ± 0.37	60 (0)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 3b for Supplementary Fig. 4d.

Groups	No. of experiments ^a	Mean survival time ± SE (days old)	Maximum survival time ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control	1	14.82 ± 0.26	18.13± 0.50	61 (2)	0.05050
Sesamin		15.66± 0.22	18.38 ± 0.30	61 (1)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 4a for Supplementary Fig. 5.

Groups	No. of experiments ^a	Mean survival time ± SE (days old)	Maximum survival time ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control	2	13.52 ± 0.16	16.90± 0.36	115 (1)	0.05837
Sesamin		14.12± 0.17	17.50 ± 0.38	117 (3)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 4b for Fig. 6.

Groups	No. of experiments ^a	Mean survival time ± SE (days old)	Maximum survival time ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
Control	2	15.38 ± 0.20	18.17± 0.27	108 (3)	0.00000
Sesamin		17.26± 0.23	21.10 ± 0.25	114 (2)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 5a for Supplementary Fig. 6a.

Groups	No. of experiments ^a	Mean life span ± SE (days old)	Max. life span ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
<i>Δ pmk-1</i> Control	1	14 ± 0.55	20 ± 0.26	67 (6)	0.84015
<i>Δ pmk-1</i> Sesamin		14 ± 0.58	21 ± 0.39	61 (5)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 5b for Supplementary Fig. 6b.

Groups	No. of experiments ^a	Mean life span ± SE (days old)	Max. life span ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
<i>Δ skn-1</i> Control	1	16 ± 0.54	22 ± 0.16	67 (1)	0.29481
<i>Δ skn-1</i> Sesamin		17 ± 0.50	23 ± 0.26	62 (5)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 5c for Supplementary Fig. 6c.

Groups	No. of experiments ^a	Mean life span ± SE (days old)	Max. life span ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
<i>Δ daf-2</i> Control	1	28 ± 2.90	38 ± 0.56	64 (1)	0.18890
<i>Δ daf-2</i> Sesamin		29 ± 3.12	39 ± 0.12	68 (3)	

^a Number of independent trials. Each trial was performed using two parallel plates.

^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 5d for Supplementary Fig. 6d.

Groups	No. of experiments ^a	Mean life span ± SE (days old)	Max. life span ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
<i>Δ daf-16</i> Control	1	20 ± 0.43	24 ± 0.35	53 (6)	0.41851
<i>Δ daf-16</i> Sesamin		19 ± 0.52	24 ± 0.25	57 (7)	

^a Number of independent trials. Each trial was performed using two parallel plates.

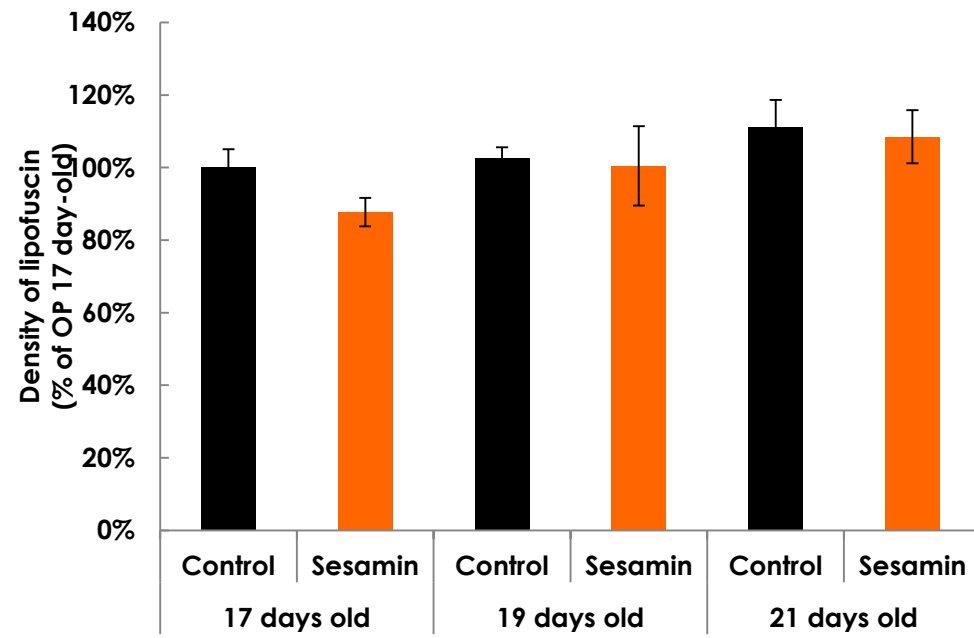
^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.

Online Resource Table 5e for Supplementary Fig. 6e.

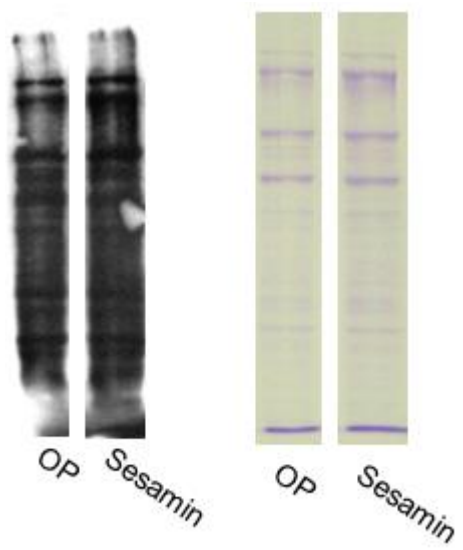
Groups	No. of experiments ^a	Mean life span ± SE (days old)	Max. life span ± SE (days old)	No. of nematodes (No. of lost worms) ^b	Log-rank test p value
<i>Δ mev-1</i> Control	2	14 ± 0.52	23 ± 0.30	139 (10)	0.0381
<i>Δ mev-1</i> Sesamin		16 ± 0.53	24 ± 0.34	118 (23)	

^a Number of independent trials. Each trial was performed using two parallel plates.

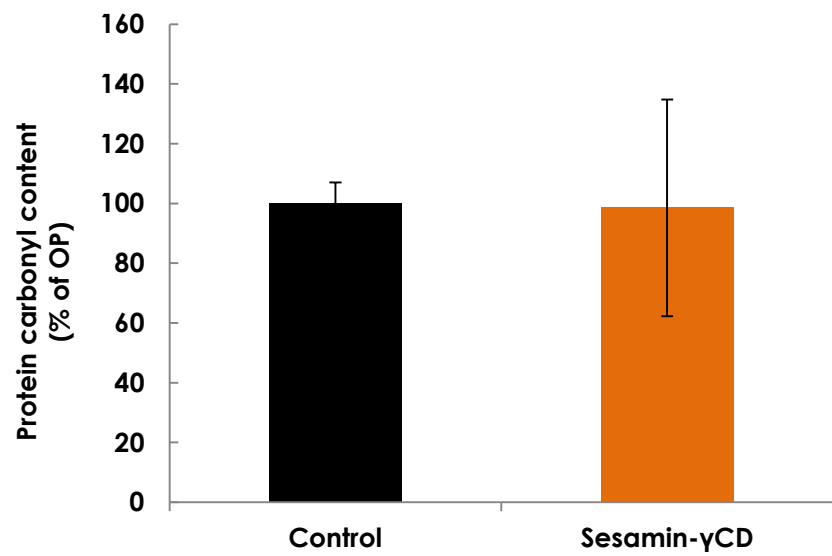
^b Worms that crawled off the plate or died from internal hatching were considered lost and not included in the analysis.



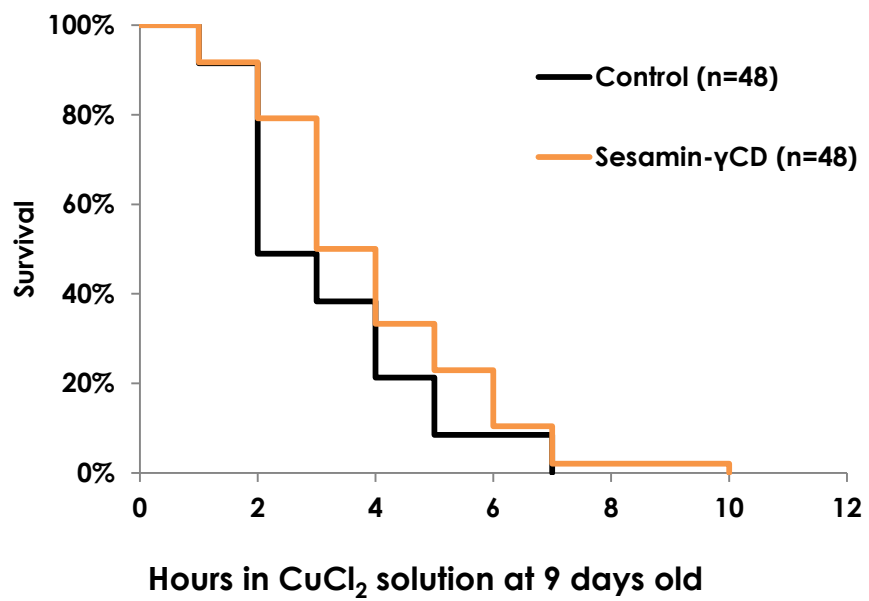
Online Resource Fig. 1



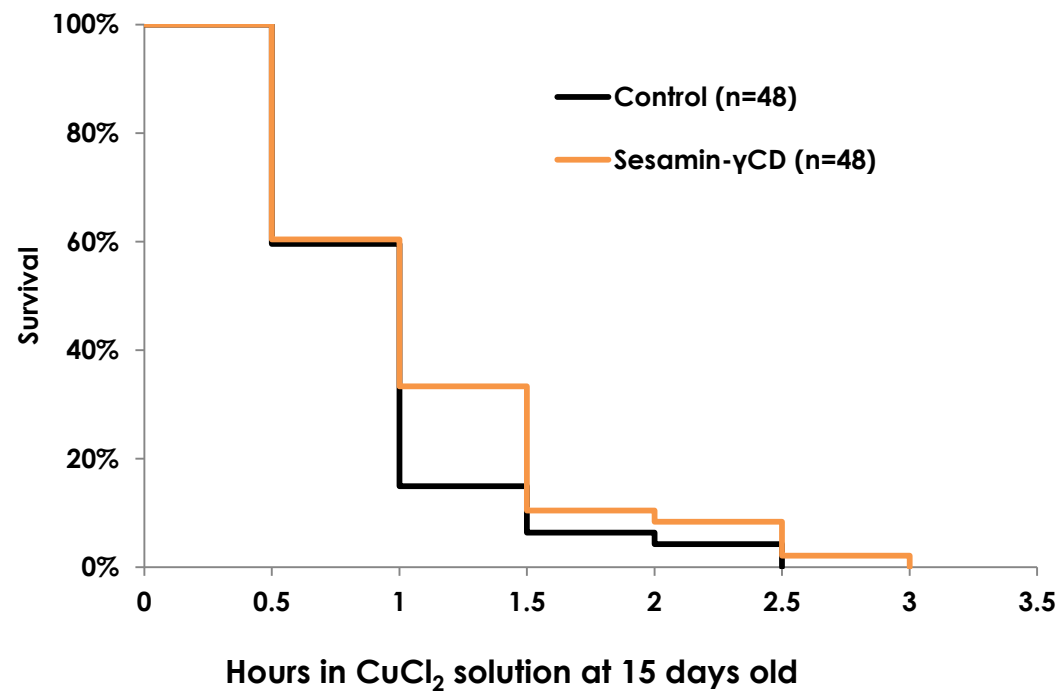
Online Resource Fig. 2a



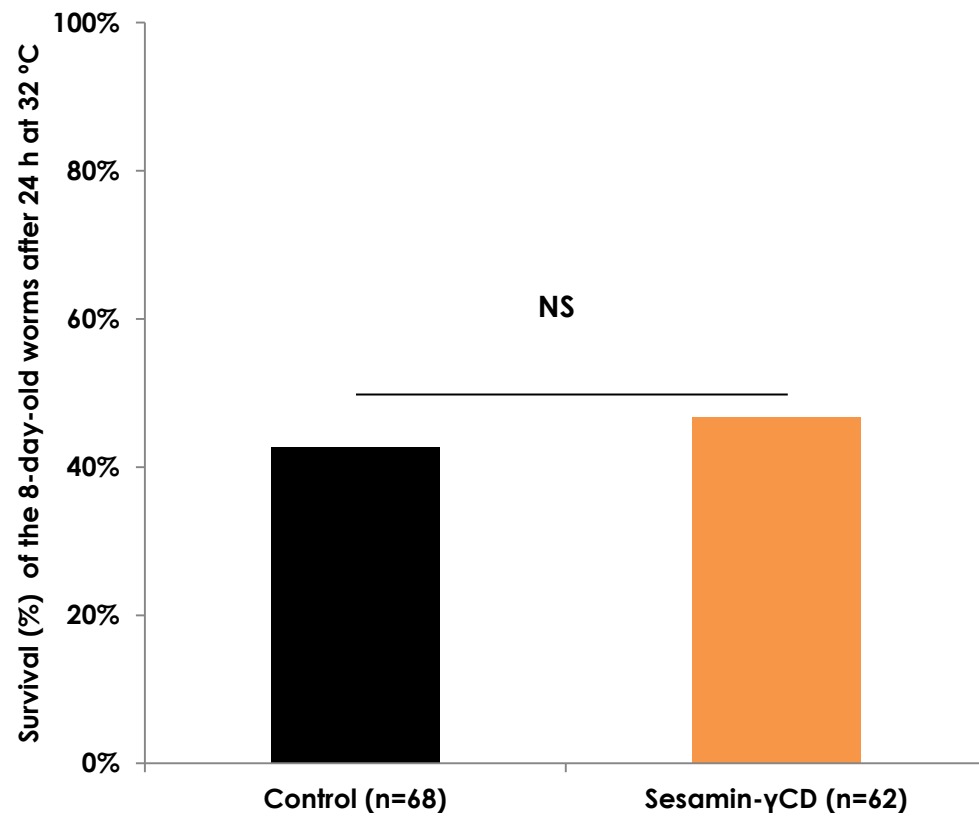
Online Resource Fig. 2b



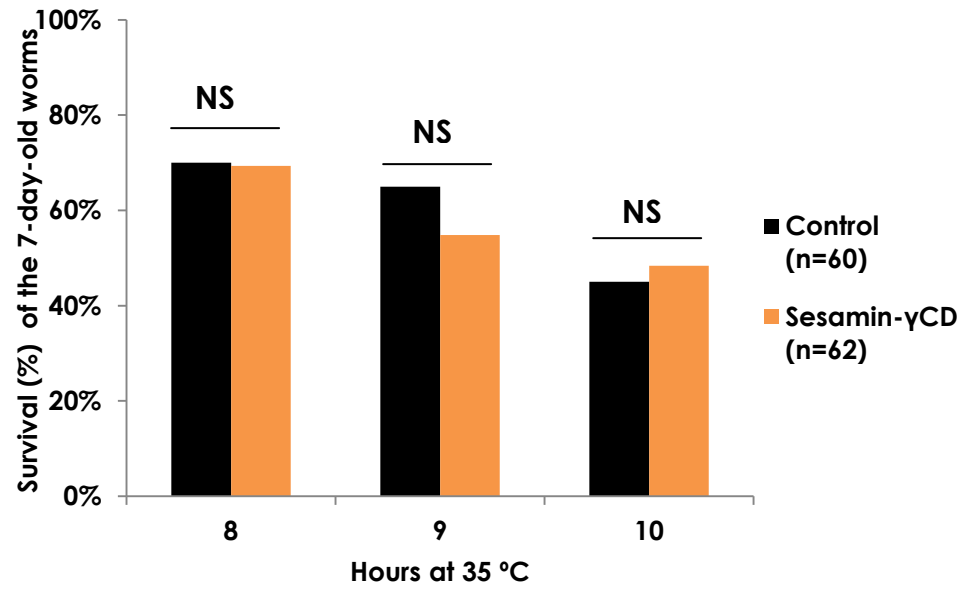
Online Resource Fig. 3a



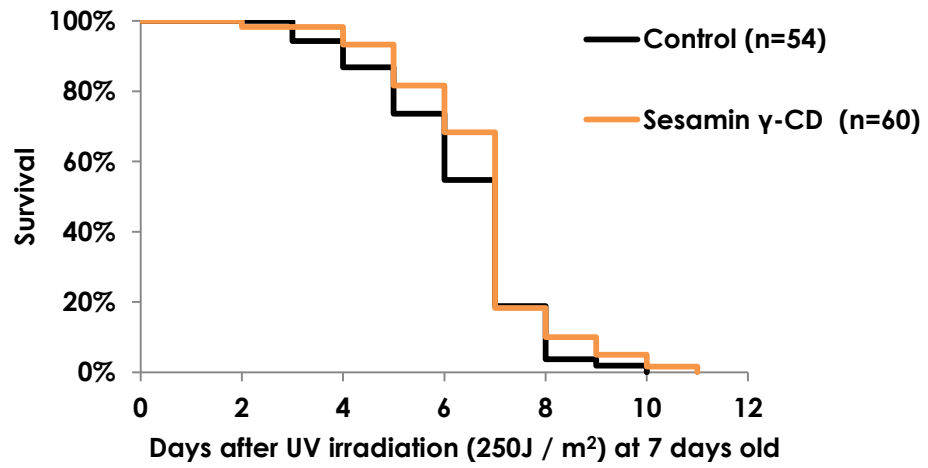
Online Resource Fig. 3b



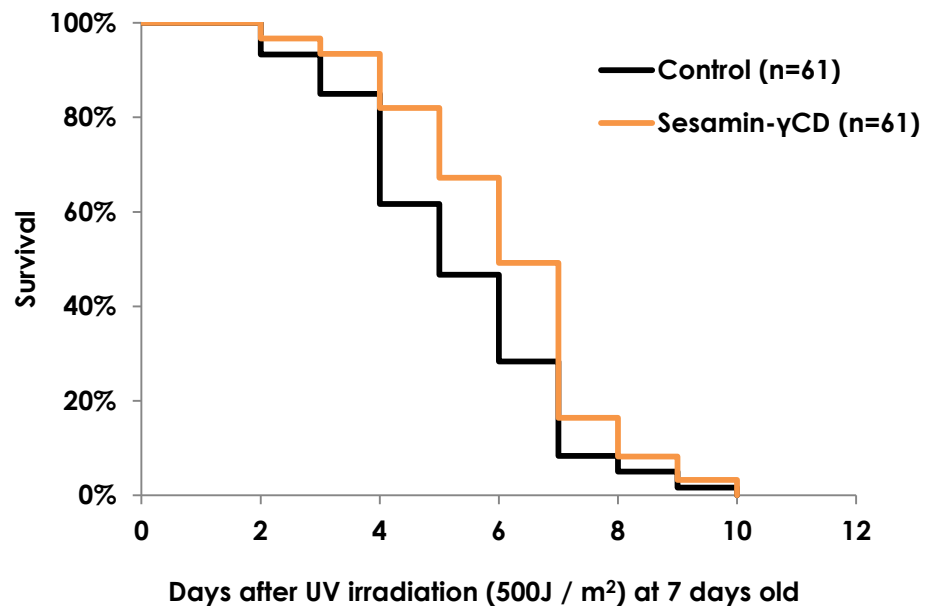
Online Resource Fig. 4a



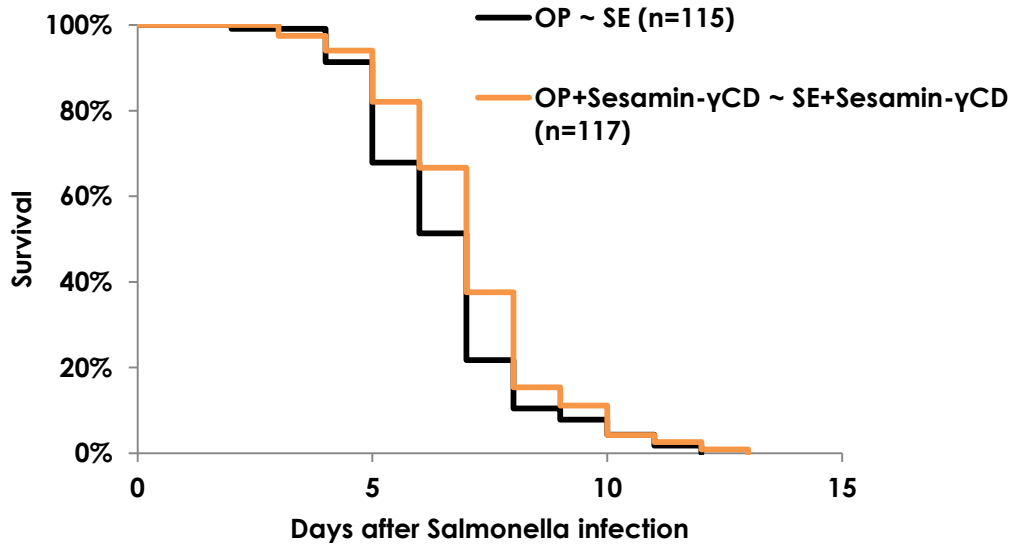
Online Resource Fig. 4b



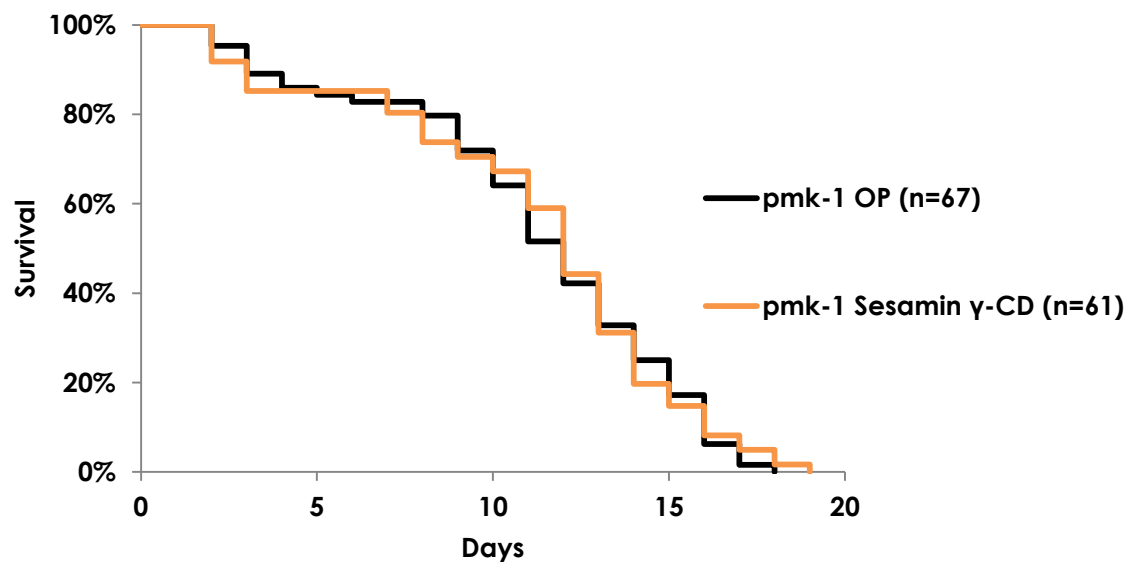
Online Resource Fig. 4c



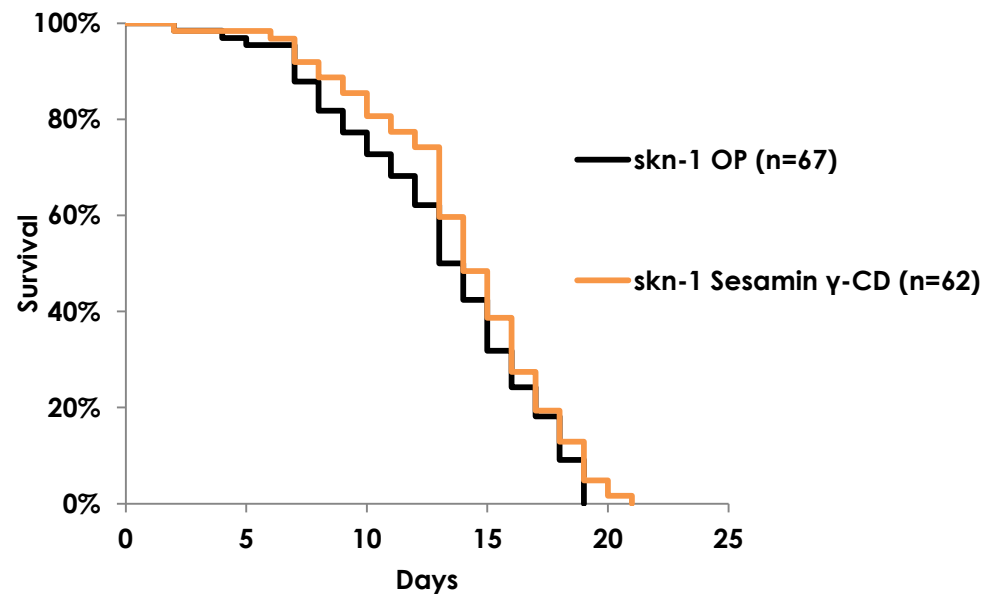
Online Resource Fig. 4d



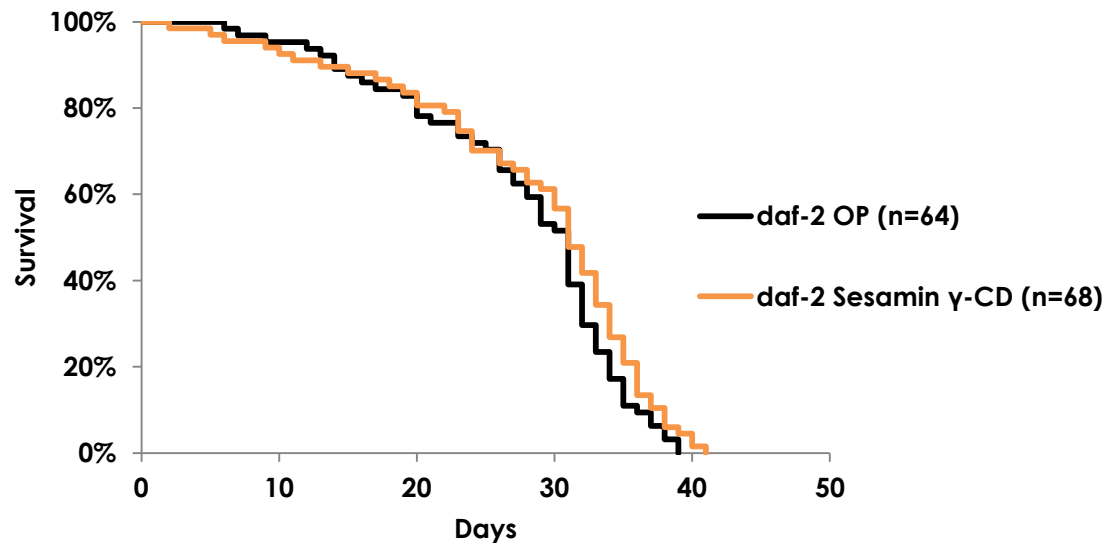
Online Resource Fig. 5



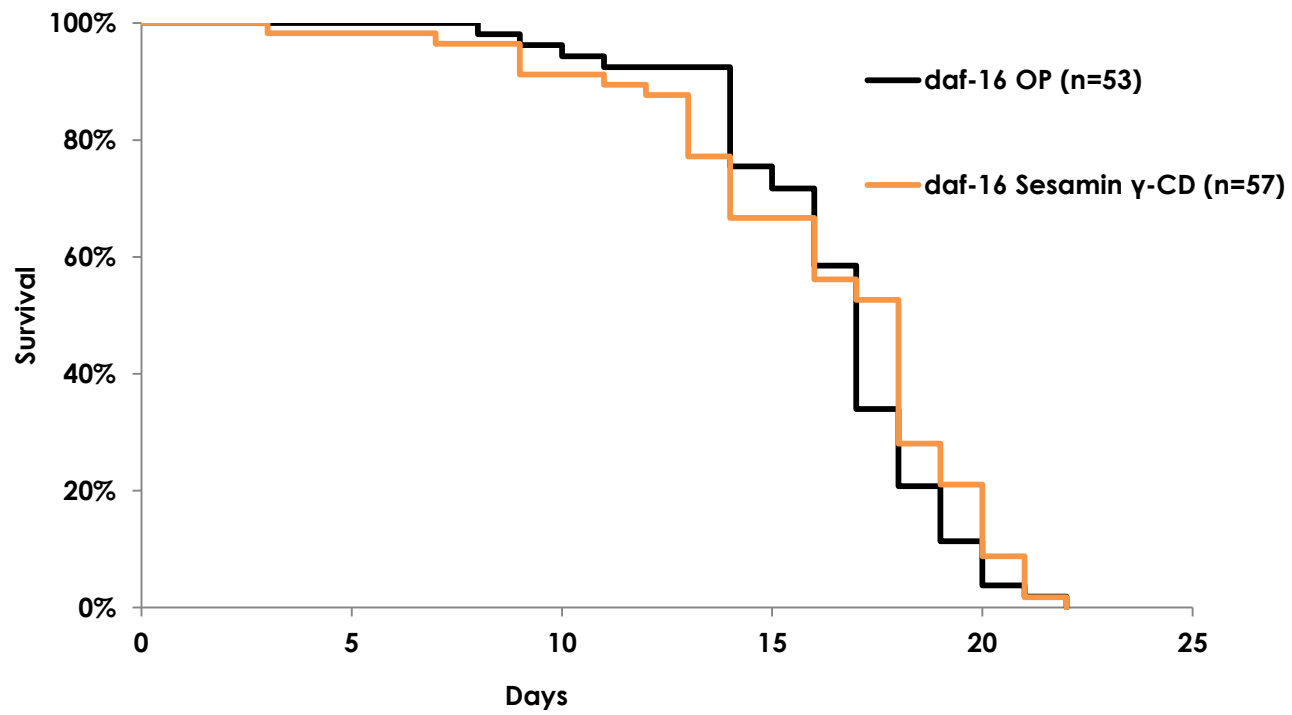
Online Resource Fig. 6a



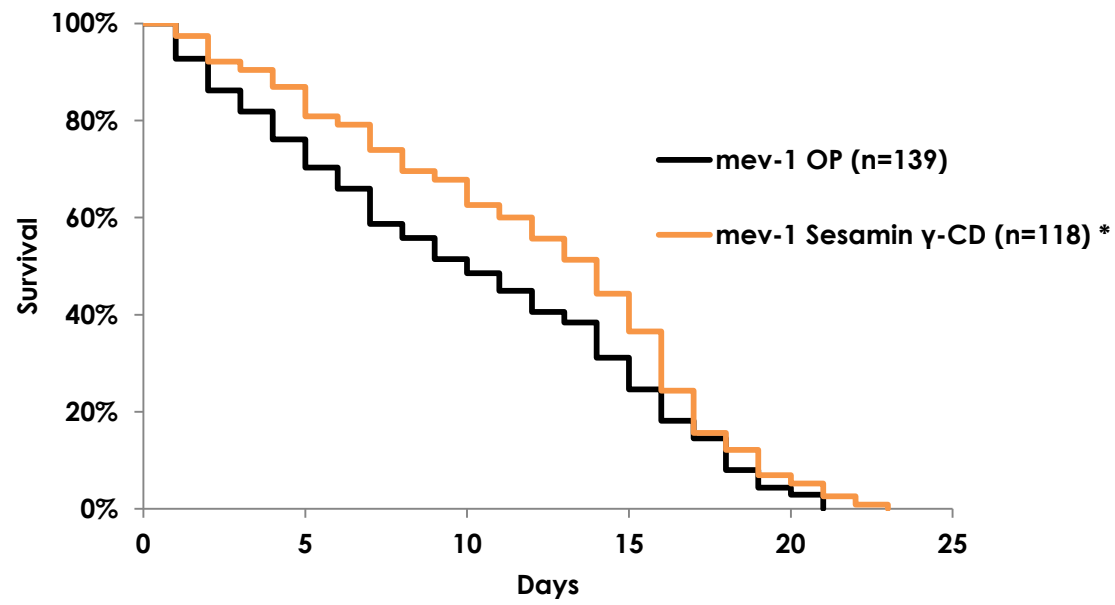
Online Resource Fig. 6b



Online Resource Fig. 6c



Online Resource Fig. 6d



Online Resource Fig. 6e