Astaxanthin extends lifespan via altered biogenesis of the mitochondrial respiratory chain complex III

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Running title: “Astaxanthin affects mitochondria and life-span”

Abstract

Astaxanthin is a *keto*-carotenoid produced in some bacteria and algae, which has very important industrial applications (i.e., in cosmetics, coloring additive in aquaculture and as a dietary supplement for human). Here, we analyzed the molecular basis of Astaxanthin-mediated prolongevity in the model organism, *Caenorhabditis elegans*. The increased lifespan effects of Astaxanthin are restricted in *C. elegans* to the adult phase and are uninfluenced by various other carotenoids tested. Genetic analyses indicated that the Astaxanthin-mediated life-extension relies on mitochondria activity, via the Rieske iron-sulfur polypeptide-1 (ISP-1), but is not influenced by the functions of other known longevity-related gene-loci, including *CLK-1*, *DAF-2*, *DAT-16*, *EAT-2*, *GAS-1* *GLP-1* or *MEV-1*. Biochemical analyses of native respiratory complexes showed that Astaxanthin affects the biogenesis of holo-complex III (and likely supercomplex I+III, as well). Effects on holo-CIII assembly and activity were also indicated by *in-vitro* assays, with mitochondria isolated from worms, rodents, human and plants, which were treated with Astaxanthin. These data indicated a cross-species effect on the oxidative phosphorylation (OXPHOS) machinery by the carotenoid, and provide with further insights into the molecular mechanism of animals longevity extension by Astaxanthin.

Keywords: *Caenorhabditis elegans*, Carotenoids, Astaxanthin, median lifespan; mitochondria, OXPHOS, Complex III, Rieske iron-sulfur protein,

Significance Statement

Astaxanthin is a widely consumed pigment by animals and human. In this study we find that Astaxanthin, but not other tested carotenoids, significantly extends the lifespan of animals by affecting respiratory complex III (CIII) biogenesis of the mitochondria, in plants, *C. elegans*, rodents and human. We further propose a model to try explaining this effect of astaxanthin on animals' longevity.

Introduction

The nematode *Caenorhabditis elegans* is a leading model organism for studying the biology of aging, and for identifying new pharmacological targets for extended longevity and the treatment of aging-related diseases in animals (Chen *et al.* 2014). Genetic studies indicated to three major aging-related mechanisms in *C. elegans*. These include the insulin pathway, the dietary restriction system, and the mitochondrial respiration system, which were all shown to affect the lifespan of worms and other animals (Amrit *et al.* 2014, Collins *et al.* 2008). Noteworthy, the majority (>80%) of the *C. elegans* proteome was shown to share homology with known human genes (Henricson *et al.* 2004, Lai *et al.* 2000), thus suggesting common aging-related mechanisms between *C. elegans* and human.

Among the key factors that affect longevity in worms is oxidative damage, generated by reactive oxygen species (ROS). Induced ROS level was shown to induce early aging in various animals and plants (Paital *et al.* 2016, Singh *et al.* 2016). Consequently, strong antioxidants as resveratrol, coenzyme Q10 (ubiquinol or 2,3-dimethoxy-5-methyl-6-decaprenyl benzoquinone) and carotenoids as Astaxanthin can extend the lifespan of *C. elegans*, presumably by scavenging ROS and lowering the levels of oxidative stress (Chen, *et al.* 2014). Carotenoids comprise a large class of natural antioxidant pigments that are synthesized and accumulate in photosynthetic organisms, various bacteria and some fungi (Cazzonelli 2011, Rodriguez-Concepcion *et al.* 2018). These isoprenoid compounds are also essential components of the photosynthetic machinery, and are associated with anti-oxidative reactions in bacteria, fungi, algae and plants. In animals, carotenoids that are consumed from the food play important roles in the antioxidant defense mechanism (Fiedor and Burda 2014).

Astaxanthin (i.e., 3,3-dihydroxy- β,β -carotene-4,4-dione) is an orange-red pigment containing 13 conjugated double bonds (Supplementary Fig. S1). As other carotenoids, Astaxanthin is a lipid-soluble pigment, which has self-limited absorption orally, with no known toxic syndromes (Barros *et al.* 2014). Astaxanthin is naturally produced by some marine bacteria, algae and plants. Many red-colored marine animals (e.g., salmon, shrimps) and a few bird species (e.g., flamingo) obtain the Astaxanthin orally, through the food chain (Ambati *et al.* 2014). As a strong antioxidant, Astaxanthin was utilized as a common antioxidant agent and as a dietary supplement, intended for human, animal and aquaculture consumption. Astaxanthin has been implicated in numerous health benefits in humans, including reduction in cardiovascular diseases, enhancement of the immune response and reduction in the occurrence of various cancers. (Ames 2018, Zhang and Wang 2015, Zhang *et al.* 2016). Astaxanthin is

obtained industrially by either chemical synthesis or by extraction from algae or bacteria, both of which are approved by the U.S. Food and Drug Administration as food, feed or coloring additives (Raposo *et al.* 2015, Zhang and Wang 2015). Plant-derived Astaxanthin is generally recognized as safe by the FDA, meaning it can be sold as a dietary supplement (Zhang and Wang 2015), although in the United States is restricted to be used only as a colorant agent to animal feed. Currently, there is great interest around Astaxanthin due to its biochemical characteristics, mainly as a potent antioxidant, which is approximately 10 times more effective than β -carotene or lutein and about 100 times than α -tocopherol (Higuera-Ciapara *et al.* 2006, Rao *et al.* 2015). These studies further suggest that Astaxanthin affects cellular metabolism and protects against fatty acid oxidation.

Astaxanthin was previously shown to extend the lifespan of both flies and worms. *Drosophila melanogaster* mutants in CuZn-superoxide dismutase (SOD1) and Mn-superoxide dismutase (SOD2), as well as mutant-lines with reduced levels of SOD1, SOD2 and catalase, show significant lifespan extension and amelioration of age-related decline in motility when are fed with *Haematococcus pluvialis* (a microalga which accumulates high-levels of Astaxanthin) (Huangfu *et al.* 2013). Likewise, feeding *C. elegans* with Astaxanthin also extends the mean lifespans of both wild-type and long-lived mutant *age-1* animals (Kashima *et al.* 2012, Yazaki *et al.* 2011). The molecular basis of astaxanthin-mediated longevity effects seen in both *D. melanogaster* and *C. elegans* remained unknown, but it was anticipated to be mediated through the scavenging of reactive oxygen species (ROS) produced in these animals by the carotenoid compound.

In this study we show that the lifespan of *C. elegans* fed with Astaxanthin increases by about 20%, which is in accordance with previously published data showing the effects of Astaxanthin on the mean lifespans of the worms (Liu *et al.* 2016). Our data further indicate that the effects of Astaxanthin are independent of the insulin/IGF or dietary restriction of germline pathways, and provide with evidence that the extended longevity effects on *C. elegans* is mediated by the disassembly or altered biogenesis of the native respiratory complexes III₂ and presumably the supercomplex I+III₂. The effects of Astaxanthin on the mitochondrial OXPHOS system seems universal, as was also apparent by ‘*in-organello*’ assays with mitochondria preparations from plant, *C. elegans*, mice rat and human. These results are of great importance as they offer with novel insights into the molecular basis of longevity extension by Astaxanthin, and may also lead to developing new generation of drugs based on carotenoids to prolong lifespan in animals and human.

Materials and Methods

C. elegans strains

C. elegans strains maintenance was performed under standard conditions, essentially as described (Brenner 1974). N2 (Wild-type), *daf-2*(e1370), *daf-16*(mu86), *glp-1*(or178), *eat-2*(ad1116), *clk-1*(e2519), *mev-1*(kn1), *gas-1*(fc21) and *isp-1*(qm150) strains were used in this study. All strains were obtained from the *C. elegans* Genome Center (CGC) and were out-crossed with wild-type (N2) animals.

Lifespan assays

Lifespan assays were performed essentially as described previously (Bar *et al.* 2016). Synchronized *C. elegans* animals were grown on NGM (K₂HPO₄, KH₂PO₄, NaCl, Bacto peptone, Agar, Agarose, MgSO₄, CaCl₂, and Cholesterol) plates containing one of the following bacterial strains: Wild type *Paracoccus marcusii*, which are accumulating high-levels of Astaxanthin, *P. marcusii* ΔcrtB mutant which lacks carotenoids, *Escherichia coli* OP50 strain (Taxonomy ID 637912), recombinant *E. coli* cells carrying the pASTA plasmid for the expression of Astaxanthin (Cunningham and Gantt 2007). The carotenoid composition in different bacterial strains used in this work is indicated in Supplementary Table S1. In addition, the worms were also grown in the presence of *E. coli* cells in agar-plates supplemented with Astaxanthin (extracted from the algae *Haematococcus pluvialis*) solubilized in 5% DMSO (w/v), and agar-plates containing 5% DMSO with *E. coli* (OP50) cells, as control. Experiments were performed at 20°C, with the exception of *glp-1* and *daf-2* mutant-lines which were conducted at 25°C. In the *daf-2* experiments, the worms were grown at 16°C until the L4 stage, and then transferred to 25°C. For each experiment, 80 adult animals were placed on four separate NGM-plates. Animals were considered dead when they no longer responded to gentle prodding with a platinum wire. Scoring was performed every day. For all lifespan experiments, assays were repeated at least twice.

Measurements of carotenoids concentration

Wild-type *P. marcusii* or *E. coli* producing Astaxanthin were plated on NGM media, harvested from the plates and resuspended in 2 ml of DDW. The cells were then washed once in 2 ml DDW. The bacterial suspension (1.8 ml) was incubated at 105°C for 24 h, after which the weight of the bacteria was measured. For carotenoid extraction, after washing in DDW, the cells were resuspended in 1 ml of acetone and incubated at 65°C for 10 min in the dark. The

samples were centrifuged again at 13,000 xg for 10 min and the acetone supernatant, containing the pigments, was placed in a clean tube. The pigment extract was dried with N₂ and then stored in the freezer at -20°C. Acetone-dissolved pigment extracts were analyzed by high performance liquid chromatography (HPLC), essentially as previously described (Neuman et al. 2014, Ronen et al. 1999), using a Waters 996 photodiode array detector. Carotenoids were identified based on their characteristic absorption spectra and typical retention times, which corresponded to standard compounds (Isaacson *et al.* 2002). For Astaxanthin extraction, wild-type algae *Haematococcus pluvialis* (kindly provided by Prof. Sammy Boussiba, BGU) were grinded with a mortar and pestle in 5 ml of acetone. The cells were incubated at 65°C for 10 min in the dark, and centrifuged at 13,000 xg for 10 min. The supernatant, containing acetone-soluble pigments, was placed in a clean tube, dried with a stream of N₂ and then subjected to HPLC analysis.

Crude mitochondrial preparations from *C. elegans* cells

Mitochondria were isolated from *C. elegans* according to the method described in Grad *et al.* (2007). Bleached embryos were grown at 20°C, on NGM plates seeded with either *P. marcusii* expressing Astaxanthin or *E. coli* (OP50). Before extraction, worms of the specified stages were washed off plates with sterile M9 solution (42 mM Na₂HPO₄, 22 mM KH₂PO₄, 86 mM NaCl, and 1 mM MgSO₄·7H₂O). The worms were pelleted by centrifuging at 3,000g for 5 min, and then resuspended in 10 ml of ice-cold isolation buffer (IB) (210 mM mannitol, 70 mM sucrose, 0.1 mM EDTA, 5 mM Tris-HCl pH 7.4) in the presence of protease inhibitors. The worms were homogenized using a glass homogenizer for 15 strokes. The homogenate was collected in a 50 ml tube and 10 ml of IB was added. Then tubes were centrifuged at 750 g for 10 min at 4°C. The clear supernatant was transferred to a new tube, and centrifuged at 12,000 g for 10 min at 4°C to pellet the mitochondria. The organellar fraction was washed once in 10 ml ice-cold IB (12,000g for 10 min at 4°C), and the mitochondrial pellet was aliquoted and stored at -80°C.

Mitochondrial extraction from cauliflower

Mitochondria were isolated from cauliflower (*Brassica oleracea* var. botrytis) inflorescences, which allows the purification of large quantities of highly enriched organellar preparations (Keren *et al.* 2009, Neuwirt *et al.* 2005, Sultan *et al.* 2016). About 1.0 kg fresh weight inflorescences were cut off from the stems and kept overnight (about 16 h) in cold water (4°C). In the following morning, the inflorescences were ground with ice-cold extraction buffer (0.9 M mannitol, 90 mM Na-pyrophosphate, 6 mM EDTA, 2.4% PVP25 (w/v), 0.9% BSA (w/v), 9 mM cysteine, 15 mM glycine, and 6 mM β-mercaptoethanol; pH 7.5). Mitochondria were

recovered from the extract by differential centrifugations and purified on Percoll gradients. Following fractionation, the mitochondria were pelleted, resuspended in a small volume of wash buffer (0.3 M mannitol, 10 mM K-phosphate, 1 mM EDTA, pH 7.5), aliquoted and stored frozen at -80°C. For the isolation of mitochondria from *Arabidopsis*, we used 2-week-old seedlings grown on MS-plates. The enriched organelle preparation was performed essentially as described in (Keren *et al.* 2012). For analysis of Astaxanthin effects, the mitochondria were incubated with Astaxanthin dissolved in DMSO or DMSO alone for 5 min on ice.

Mitochondrial enriched preparation from mammals

Mitochondrial enriched fraction, was prepared from human (fibroblast) and mouse (heart) by teflon-glass homogenization and differential centrifugation in sucrose buffer (250 mM Sucrose, 10 mM Tris pH 7.4, 50 µg/ml heparin) as described (Saada *et al.* 2003). For enzymatic activities, mitochondria were pre-incubated with Astaxanthin or DMSO for 5 min on ice prior to assay.

Blue native electrophoresis for isolation of native organellar complexes

Blue native (BN)-PAGE of mitochondrial fractions was performed according to the method described by (Eubel *et al.* 2003). Mitochondrial pellets were solubilized with n-dodecyl-β-maltoside (1.5% [w/v]) in ACA buffer (750 mM amino-caproic acid, 0.5 mM EDTA, and 50 mM Tris-HCl, pH 7.0), and then incubated on ice for 30 min. The samples were centrifuged 8 min at 20,000g, and Serva Blue G (0.2% [v/v]) was added to the supernatant. The samples were then loaded onto a native 4% to 16% gradient gel. For non-denaturing-PAGE-Western blotting, the gel was transferred to a PVDF membrane (Bio-Rad) in Cathode buffer (50 mM Tricine and 15 mM Bis-Tris-HCl, pH 7.0) for 16 h at 4°C (const. 40 mA). The membrane was incubated with various antibodies, as indicated for each blot, and detection was carried out by chemiluminescence assay after incubation with an appropriate horseradish peroxidase (HRP)-conjugated secondary antibody.

SDS-PAGE gel electrophoresis

Organellar protein concentration was determined by the Bradford method (BioRad, Catalog no. 5000201) according to the manufacturer's protocol, with bovine serum albumin (BSA) used as a calibrator. For immunoassays, an aliquot equivalent to 600 µg *C. elegans* mitochondrial proteins was suspended in sample loading buffer (Laemmli 1970) and subjected to SDS-PAGE (at a constant 100 V). Following electrophoresis, the proteins were transferred to a PVDF

membrane (BioRad, Catalog no. 1620177) and incubated overnight at 4°C with various primary antibodies (Table S2). Detection was carried out by chemiluminescence assay after incubation with an appropriate horseradish peroxidase (HRP)-conjugated secondary antibody.

Assessment of mitochondrial respiratory chain enzymatic activities

In-gel activity stains of mitochondrial respiratory complexes I, II and IV was carried out according to (Eubel *et al.* 2005). Following BN-electrophoresis, the gels were washed several times with DDW. For Complex I activity staining, the BN-gels were incubated in CI-activity assay solution (100 mM Tris-HCl, pH 7.4, 0.14 mM NADH, 1 mg/ml NBT solution), for 10 to 30 min until the purple staining of CI bands became visible in the gel. Complex II activity staining was carried out with 50 mM KH₂PO₄ buffer (pH 7.4), 84 mM succinate, 0.2 mM PMS and 2 mg/ml NBT solution. For complex IV activity staining, the BN-gels were incubated in 10 mM KH₂PO₄ buffer (pH 7.4), 1 mg/ml DAB solution and 0.2 mg cytochrome c. Reactions were stopped in fixing solution containing 15% (v/v) ethanol and 10% (v/v) acetic acid. All steps were carried out at room temperature. Respiratory chain activities and citrate synthase activity assayed by spectrophotometric methods at 37°C (Kirby *et al.* 2007, Shufaro *et al.* 2012). Briefly, complex I was measured as rotenone sensitive NADH-CoQ reductase at 340 nm in the presence of coenzyme Q₁. Complex II was measured as succinate dehydrogenase (SDH) based on the succinate-mediated phenazine methosulfate reduction of DCPIP (dichloroindophenol indophenol) at 600 nm. Complex II + III was measured as thenoyltrifluoroacetone sensitive succinate cytochrome c reductase at 550 nm. Complex III was measured as antimycin sensitive ubiquinol cytochrome c reductase at 550 nm. Complex IV. (cytochrome c oxidase) was measured by following the oxidation of reduced cytochrome c at 550 nm. Citrate synthase (CS), an ubiquitous Krebs cycle enzyme, serving as a control, was measured in the presence of acetyl-CoA and oxaloacetate by monitoring the liberation of CoASH coupled to 5',5-dithiobis (2-nitrobenzoic) acid (i.e., Absorbance at 412 nm).

Respiration activity

Oxygen consumption (O₂-uptake) measurements were performed with a Clarke-type oxygen electrode (Oxytherm System, Hansatech Instruments, Norfolk, UK), and the data feed was collected by Oxygraph-Plus software, essentially as described previously (see e.g., (Shevtsov *et al.* 2018). The electrode was calibrated with O₂-saturated water, and by depletion of the oxygen in the electrode chamber with the addition of excess sodium dithionite. Total respiration was measured at 25°C in the dark, following the addition of 250 µL *C.elegans* suspension from

different treatments ($\pm$ Astaxanthin) and mutant-lines to a 2.25 ml of M9 solution in the respiration chamber. The *C.elegans* suspension was prepared by 1/16 dilution of dense packed worms from 5 plates in M9 buffer. Oxygen consumption was also measured in the presence of various inhibitors, including Rotenone (Rot), Antimycin A (Anti A) and potassium cyanide (KCN).

Statistical Analyses

Lifespan curves were analyzed by plotting Kaplan-Meier survival curves (Goel *et al.* 2010), and by conducting Log-rank tests (Mantel 1966). Mean lifespan data was compared using Log-rank test with multiple comparisons test, OASIS-2 (Han *et al.* 2016). Differences with P-value smaller than 0.05 were considered as statistically significant.

Results

Feeding adult *C. elegans* with Astaxanthin cause significant extension in their lifespan

Synchronized *C. elegans* animals were grown for 5 days at 20°C on NGM plates containing *P. marcusii* cells (expressing Astaxanthin) or in the presence of *P. marcusii* Δ crtB mutant-line (lacks carotenoids expression). Following 5 days, the intestine of the worms was cleaned by transferring the animals to new NGM plates, containing *E. coli* (OP50) cells, for additional 24 hours (Images of *C. elegans* before and after intestine cleaning procedure are provided in Supplementary Figure S2). Under these conditions, the average lifespan of wild-type *C. elegans* (N2), grown on agar plates and fed with *Paracoccus marcusii* (containing on average 716 μ g Astaxanthin per gram of dry material), was about 20% longer (i.e., 24 Days vs. 20 Days) than *C. elegans* fed with *P. marcusii* Δ crtB mutant that lacks carotenoids (P-value = 0.0038) and about 45% longer (24 Days vs. 17 Days) compared with those of *C. elegans* fed with *E. coli* OP50 cells (P-value = 0.00003) (Fig. 1). Likewise, feeding *C. elegans* with a recombinant *E. coli* cell-line which produces Astaxanthin (pASTA, 146 μ g Astaxanthin per gram of dry material), was also sufficient to significantly increase the lifespan of the worms by about 20% (P-value = 0.0114, Fig. 1B). Feeding *C. elegans* with Astaxanthin extracted from the algae *Haemayococcus pluvialis* and placed on *E. coli* OP50 strain increased the average lifespan by ~20% (P-value = 0.0173, Fig. 1C). No effect on the lifespan was observed in *C. elegans* fed with *P. marcusii* strains expressing the carotenoids Zeaxanthin (P-value = 0; *P. marcusii* Δ crtW, Fig. S2 and Table S1), while Lycopene (*P. marcusii* Δ crtY, Table S1) was found to be toxic to the animals (animals arrested at the L2 stage).

Next, we determined the developmental period where Astaxanthin has the most notable effect on the lifespan of *C. elegans* (see Fig. S3). For this purpose, we used four different feeding conditions. These include animals that were fed from the larval L1 until adulthood with Astaxanthin (i.e., in the presence *P. marcusii*), and later transformed to new plates lacking Astaxanthin (i.e., plates containing *P. marcusii* Δ crtB) (Fig. S3, L1-Mature +Asta, Adult -Asta). The second condition involved animals from the larva L1 stage until adulthood that were fed with *P. marcusii* Δ crtB (lacks Astaxanthin) and later transferred to new plates containing wild-type *P. marcusii* (produces Astaxanthin) (Fig. S3, L1-Mature -Asta, Adult +Asta). In addition, as controls we also used worms that were fed during their entire lifespan on either *P. marcusii* wild-type (Fig. S3, +Asta) or *P. marcusii* Δ crtB (Fig. S3, -Asta).

As indicated Figure S3, feeding animals that are only at their adult stage (Fig. S3, L1-Mature -Asta, Adult +Asta) with Astaxanthin resulted with an increase in lifespan which was similar to that of animals fed with Astaxanthin throughout their entire life (P-values <0.05). However, no significant change in longevity was apparent when the animals were fed with Astaxanthin from the larval L1 stage until young adulthood, but were avoided from being exposed to the pigment, during their adult stage (Fig. S3, L1-Mature +Asta, Adult -Asta). Similarly, no increase in lifespan was seen when the animals are grown on *P. marcusii* Δ crtB cells lacking any carotenoids.

We also noted that *C. elegans* animals fed with *P. marcusii* showed a strong red-orange color of the gut (Fig. S4, left panel). Feeding *C. elegans* with *P. marcusii* cells expressing Astaxanthin followed by 24 hours feeding with *E. coli* (OP50) resulted with a strong decolorization of the guts, but some residual red color was still observed in the bod cells (Fig. S4, middle panel). We therefore hypothesize that feeding *C. elegans* at the adult stage with Astaxanthin resulted with the accumulation of Astaxanthin in the cells, an effect that likely correlates with the observed extension in the lifespan.

The effects of Astaxanthin on *C. elegans* longevity are tightly associated to the mitochondria-aging pathway

There are several known pathways that affect aging in *C. elegans* (i.e., the insulin pathway, the dietary restriction system, and the mitochondrial respiration system) (Amrit, *et al.* 2014, Collins, *et al.* 2008). Here, we performed genetic analyses to determine whether and which of the three known aging pathways might be associated with the prolongevity effect of Astaxanthin. We reasoned that animals with mutations in pathways affected by Astaxanthin would show reduced

effects in the presence of the carotenoid on the lifespan extension. Among of the longevity-associated pathway in *C. elegans* is the DAF-16/DAF-2 insulin pathway (Lin *et al.* 2001). We noticed that similarly to the-wild-type animals, Astaxanthin caused an (additional) increase in the lifespan of the *daf-2 (e1370)* (P-value = 0.001) and *daf-16 (mu86)* (P-value = 0.000046) mutant animals (Fig. 2). Similarly, feeding *C. elegans glp-1 (or178)* mutants (Libina *et al.* 2003, TeKippe and Aballay 2010) with Astaxanthin resulted in additional extension of the lifespan (P-value = 0.000024, Fig. 2). We therefore concluded that Astaxanthin effect on *C. elegans* lifespan are independent of the insulin/IGF signaling pathway. Another well-characterized pathway of lifespan extension involves dietary restriction (Lakowski and Hekimi 1998). Here too, feeding *eat-2 (ad1116)* mutant animals with Astaxanthin resulted with an extended lifespan (P-value = 0.011, Fig. 2), indicating that Astaxanthin effect on *C. elegans* lifespan are also independent of the dietary restriction mechanism.

In addition to the insulin/IGF signaling and dietary restriction pathways, life-extending in animals also involves the mitochondrial respiration system (i.e., mitochondrial dysfunctions resulting with extended lifespan) (Murakami and Murakami 2005). It is postulated that mutations in genes encoding components of the mitochondrial electron transport chain (ETC) lower the levels of ROS and thereby may increase the lifespan of the animals (Murakami and Murakami 2005). Electron leakage from complex III is suggested to be a major contributor to the generation of ROS within the mitochondria (Brand 2010, Chen *et al.* 2003, Quinlan *et al.* 2013). To test whether reduced respiration activities may influence the prolongevity-mediated effects by Astaxanthin, we analyzed the lifespan of *C. elegans* mutants in genes encoding respiratory-related proteins. Mutants in the Rieske iron-sulfur protein (RISK) of complex III (i.e., *isp-1* mutants, also denoted as *qm150*), showed reduced (shortened) lifespan when they were treated (fed) with Astaxanthin (P-value = 0.000011, Fig. 2). These results strongly suggest that the Astaxanthin-mediated longevity effects involve complex III. The effects of Astaxanthin on the lifespan was further analyzed in *C. elegans* mutants affected in other mitochondrial proteins. *CLK-1* encodes a demethoxyubiquinone hydroxylase enzyme required for ubiquinone biosynthesis. *C. elegans clk-1* mutants show a pleiotropic phenotype that includes slowed development and aging (Branicky *et al.* 2006). However, the data showed that Astaxanthin-mediated longevity was not affected in the *clk-1 (e2519)* animals (P-value = 1.1×10^{-7} , Fig. 2). Feeding *C. elegans gas-1 (fc21)* (affected in complex I) or *mev-1 (kn1)* (affected in complex II) mutants with Astaxanthin, resulted with mild extensions in longevity (P-values 0.0024 and 0.2831, respectively). Taken together, these data indicate that the Astaxanthin-extended

lifespan effects we see in *C. elegans* rely on the level or activity of respiratory complex III, and maybe to some extent on other respiratory-mediated functions.

Astaxanthin affects the biogenesis of the native respiratory complex III, *in vitro*

The Astaxanthin-mediated effects we see suggest that Astaxanthin increases the lifespan of *C. elegans* by influencing the biogenesis or function of the respiratory system via CIII and maybe CI and CII as well (Fig. 2). To analyze whether Astaxanthin directly affect the biogenesis or activities of these respiratory complexes, we used *in vitro* analyses with crude mitochondria preparations (i.e., ‘*in-organello*’ assays), obtained from both animals and plants. These include *C. elegans*, rodents, human and cauliflower mitochondria (*Brassica oleracea* var. botrytis) (see Materials and Methods). For this purpose, Astaxanthin was extracted from the algae *Haematococcus pluvialis*, solubilized in 5% DMSO and then incubated with the mitochondria for 30 min on ice. The integrity of native respiratory complexes in *C. elegans*, cauliflower, rodents and human, was investigated by BN-PAGE analyses (Fig. 3 and Fig. S5).

Figure 3A shows the effects of increased Astaxanthin concentrations (i.e., 0, 9 and 14 μ M) on native respiratory complexes in animal mitochondria. A decreased level of a high-molecular mass band (about 500 kDa), which we expected to corresponds to a native complex III dimer (i.e., holo-CIII₂), was observed in *C. elegans* mitochondria treated with Astaxanthin. Notably, this Astaxanthin-mediated reduction in the ~500 kDa band, was followed by the appearance of a ~350 kDa band, which may corresponds to a partially disassembled CIII particle (i.e., CIII*). BN-PAGEs of crude mitochondria preparations obtained from rodents and human treated Astaxanthin showed the same patterns (i.e., reduced intensities of the ~500 kDa complex and an accumulation of ~350 kDa particles) (Fig. 3A). To analyze whether similar effects may occurs *in vivo*, we analyzed the integrity of respiratory complexes in *C. elegans* fed with Astaxanthin. For this purpose, young adult worms were grown for 7 days in the presence of wild-type *P. marcusii*, washed several times to remove bacterial contaminants and crude mitochondria preparations were obtained, essentially as described previously (Grad, *et al.* 2007). Organellar proteins from *E.coli* (-Asta) and *P. marcusii* (+Asta) fed *C. elegans* were separated by under native conditions. The BN-PAGE assays indicated that the absorption or uptake of Astaxanthin (Fig. S4) also resulted with a reduction in the 500 kDa band and the accumulation of the 350 kDa particles (Fig. S5).

We next analyze the effects of Astaxanthin on mitochondria isolated from cauliflower inflorescences, as this tissue allows the purification of large quantities of highly-pure mitochondria preparations (Keren, *et al.* 2009, Neuwirt, *et al.* 2005, Sultan, *et al.* 2016).

Mitochondria respiratory complexes were then separated by BN-PAGE analysis (Fig. 3B and Supplementary Fig. S6). Arrows indicate to native respiratory complexes CI (~1,000 kDa), a CIII dimer, (CIII₂, about 500 kDa), CIV (~220 kDa) and CV (~600 kDa) in cauliflower mitochondria. In plant's mitochondria as well, the addition of Astaxanthin led to a notable reduction in the levels of the 500 kDa corresponding to CIII₂, followed by the accumulation of a lower (about 300 kDa band) corresponding to sub-CIII particle (CIII*) (Fig. 3B). These were correlated with the levels of Astaxanthin added to the isolated mitochondria (Fig. 3C). BN-PAGEs followed by immunoblot analyses with antibodies raised to RISP subunit (iron-sulfur protein of complex III), indicated that both the ~350 kDa and 500 kDa particles contain RISP (Fig. 3 and Fig. S6), strongly supporting that these particles correspond to holo-CIII₂ and sub-CIII*. The Astaxanthin-mediated effects (i.e., decreased CIII₂ levels and increased amounts of the 350 kDa band) seem specific to Astaxanthin, as mitochondria treated with DMSO alone had no obvious effects on the levels or integrity of respiratory complexes I, III and V in cauliflower mitochondria (Fig. 3B), while the addition of β-Carotene or Zeaxanthin had no obvious effects on the integrity of the organellar complexes (Fig. S5B). Mass-spectrometry (LC-MS/MS) analyses of gel slices corresponding to holo-CIII₂ (upper band) and CIII* particles (lower band) obtained from human (Table S3) and plant (Table S4) mitochondria, indicated the presence of numerous subunits of CIII (see Table 2). The LC-MS/MS data also indicated the presence of many other peptides that are corresponding to CI (mainly), CIII, CIV, CV and various other known mitochondrial proteins. These may be associated with the CIII particles. However, it is anticipated that these proteins rather correspond to various sub-particles of respiratory complexes that are migrating with similar masses to those of the holo-CIII₂ and CIII*, as only the immunoblots with RISP analyses indicated a correlation with the reduction in the CIII dimer and to increased levels of the lower (350 kDa) band (Figs. 3C and S6). As indicated by immunoblots, the presence of RISP protein in the holo-CIII₂ and CIII* of both human and cauliflower mitochondria, was also supported by the LC-MS/MS data (Table 2 and Supplementary Tables S3 and S4).

In addition to CIII, Astaxanthin may also affects the accumulation of CI (using anti-CA2 antibodies) and the ATP-synthase (CV, anti-ATPA antibodies) in cauliflower mitochondria (Fig. 3C and Fig. S6). BN-PAGE analyses followed by immunoblots corresponding to CI (with anti-CA2 antibodies) and ATP-synthase (CV, anti-ATPA antibodies) indicated to reduced holo-CI (~1,000 kDa) and a ~1,500 kDa particle, corresponding to a 'supercomplex' containing CI and CIII₂ (i.e., CI+CIII₂), which was also apparent in the

immunoblots with anti-RISP antibodies (Figs. 3C and S6). Yet, the pigment has no obvious effect on the levels and biogenesis of the native complex V (Fig. 3C and Fig. S6).

The steady-state levels of various mitochondria proteins in Astaxanthin-fed *C. elegans* was assayed by immunoblot analyses, with antibodies generated to various respiratory complexes, including CI (C25H3.9), CIII (ISP-1), CIV (CTC-1) and CV (ATP-5) and cytochrome C (CYC-1) (see Table S2). In each assay, the steady-state levels of different proteins in the Astaxanthin-fed animals were compared to those of untreated wild-type animals of the same age at different total protein loadings (i.e., 0.15 up to 1.5 times) (Fig. 4). The levels of various mitochondrial proteins, including C25H3.9 (CI), CTC-1 (CIV) and ATP-5 (CV) did not changed significantly between the control and Astaxanthin-fed animals or the *C. elegans isp-1* mutants (Fig. 4B). A reduction in the signal of cytochrome C (CYC-1) was observed in *C. elegans isp-1* mutants, but not in animals fed with Astaxanthin. We also noticed a notable reduction in RISP signals in both *isp-1* animals and wild-type *C. elegans* fed with Astaxanthin. While a reduction in RISP is expected in the case of the *isp-1* mutant-line (Fig. 4, ISP-1 panel), it remains unclear why a notable reduction in RISP is seen in animals fed with Astaxanthin, whereas the protein remains intact in the ‘*in organello*’ assays. A reasonable explanation is that when the worms are fed with Astaxanthin they undergo continues disassembly of CIII (or the suppercomplex of CI-CIII). Under these conditions, various complex III (and possibly CI) subunits, including the RISP protein, may become more susceptible to proteolysis by organellar proteases and peptidases, while much lower degradation rates are expected under the in vitro conditions (i.e., short incubations on ice).

***C. elegans* treated with Astaxanthin display altered respiratory functions**

Analyses of native mitochondrial complexes, using denaturing PAGE profiles, suggested to alter CIII and maybe CI biogenesis in *C. elegans* fed with Astaxanthin. To determine whether the respiratory activity was affected in animals fed with Astaxanthin, we monitored the oxygen (O₂) uptake rates of non-treated versus Astaxanthin-fed wild-type *C. elegans* and *isp-1* mutants, using a Clark-type electrode (Hansatech Instruments, Norfolk, UK). For this purpose, synchronized late L4 worms were collected and washed several times with M9 buffer. Oxygen consumption was monitored in the worms in a sealed chamber (see e.g., (Cohen *et al.* 2014, Keren, *et al.* 2012, Shevtsov, *et al.* 2018, Zmudjak *et al.* 2013). Each respiration chamber contained 250 µl of packed *C. elegans* suspension diluted 10 times in M9 buffer. These measurements indicated that *C. elegans* fed on Astaxanthin and the *isp-1* mutants have reduced respiration activities (i.e., 15.03 ± 3.03 and 14.33 ± 2.88 nmol O₂·min⁻¹·mg⁻¹ total protein),

respectively, compare to those of the non-treated wild-type animals (i.e., 28.83 ± 3.07 nmol $O_2 \cdot \text{min}^{-1} \cdot \text{mg}^{-1}$ total protein) (Fig. 4).

Here, we also measured the O_2 -uptake rates in the presence of various inhibitors, including Rotenone (Rot), which inhibits complex I electron transport, the complex III inhibitor Antimycin A (Anti A) and Potassium cyanide (KCN) that blocks electron transport from complex IV to oxygen. As shown in Figure 4, both Rot and Anti-A have noticeable effects on the respiratory activities of the non-treated wild-type, compared to those of *isp-1* mutant or *C. elegans* fed with Astaxanthin ($55\% \pm 2.95$ and $66\% \pm 0.70$ compared to $34\% \pm 0.99$ $O_2 \cdot \text{min}^{-1} \cdot \text{mg}^{-1}$ total protein, respectively). In contrast, KCN had similar effects on the O_2 -uptake rates of wild-type worms, worms fed on Astaxanthin and the *isp-1* mutants ($85\% \pm 5.18$ and $90\% \pm 2.35$, and $95\% \pm 0.19$, $O_2 \cdot \text{min}^{-1} \cdot \text{mg}^{-1}$ total protein, respectively) (Fig. 4). These results further indicate to altered respiratory CI- and CIII-mediated activities.

Astaxanthin inhibits enzymatic activities of mammalian complexes I and III.

The BN-PAGE (Fig. 3 and Suppl. Figs. S4-S6), LC-MS/MS (Table S4) and the respiration activity data (Fig. 4) indicate that Astaxanthin has a strong effect on the biogenesis and functions of CIII, and presumably affects CI and the supercomplex I+III, as well. These data were further supported by analyses of the enzymatic activities of native respiratory complexes in the presence or absence of Astaxanthin. Table 1 shows the effects on the respiratory functions of mitochondrial complexes isolated from mouse heart that were pre-incubated with DMSO alone (Mock) or with purified Astaxanthin dissolved in DMSO (see Materials and Methods). The biochemical assays indicated to significant decreases in the activities of both complexes I and III. The inhibitory effect of Astaxanthin was even more pronounced in enzymatic activity assays of CI+CIII, which approached the respiratory assay sensitivity limits. These analyses also indicated that complexes II, IV and citrates synthase enzyme were not, or only slightly, affected by the addition of Astaxanthin to animals mitochondria.

Discussion

The nematode *Caenorhabditis elegans* is a well-established animal system to study organismal aging, and for the characterizing and identifying new drugs that can extend lifespan and improve the quality of old age in animals. Here, we used *C. elegans* to study the molecular basis of the anti-aging effects of Astaxanthin consumed by various organisms. Our results show that Astaxanthin extends *C. elegans* lifespan by 20%-40%. The effects on the life span seem specific

to Astaxanthin, as other carotenoids tested here, including Zeaxanthin and Lycopene had no lifespan extension effects. In addition, the anti-aging role of Astaxanthin had no obvious effects on the reproduction, age pigmentation and locomotion of the animals. The life-extension phenotype, induced by Astaxanthin, relies mainly on *isp-1* mutants that are affected in mitochondria respiratory complex III, but not on *gas-1* (complex I) and *clk-1* (ubiquinol biosynthesis), nor on other aging-related pathways (i.e., the insulin/IGF-1 and dietary restriction pathways), as indicated by lifespan measurements of *daf-2/daf-16*, *eat-2* or *glp-1* mutants fed with Astaxanthin.

Remarkably, Astaxanthin added to isolated mitochondria causes a reduction in the levels of holo-complex-III₂ and supercomplex I+III, and to the appearance of lower molecular-mass particles that correspond to sub-CIII particles. Accordingly, BN-PAGE and LC-MS/MS analyses indicated the presence of various CIII subunits in both the high and lower mass particles. The presence of various complex I subunits in these gel-bands, may relate to the strong association of complexes I and III, which together with complex IV also form a stable supercomplex known as the Respirasome (I₁III₂IV₁) (Schafer *et al.* 2006). Alternatively, it remains possible that sub-complex I, IV and V (natively formed or disassemble during mitochondria preparations or gel-run) may migrate with similar masses under the experimental conditions. Together, these analyses indicate that Astaxanthin affects the assembly or biogenesis of complex III and the CI-CIII₂ supercomplex, whose functional roles and assembly are under investigation (Signes and Fernandez-Vizarra 2018). As CIII biogenesis / disassembly are observed in both animal and plant mitochondria (Figs. 3, S5 and S6), the data imply to a universal effect caused by Astaxanthin to CIII, in all eukaryotes.

Reduced accumulation of CIII is expected to affect the respiratory functions of *C. elegans* cells fed with Astaxanthin. The O₂-uptake rates of worms fed on Astaxanthin or *isp-1* mutants were reduced by about 2x folds (Fig. 5), compared to those measured in the non-treated (control) *C. elegans*. Similarly, the oxygen consumption of Astaxanthin-treated mitochondria, isolated from mouse cells was lowered by more than 50%, and to reduced activities of the respiratory complexes I and III. Currently, it remains unclear how Astaxanthin affects the assembly or biogenesis of complex III in plants and animals. It is possible that Astaxanthin interacts with complex III and thereby interfere with the assembly process or destabilizes the integrity of the native complex. Due to its skeletal properties (long polyene chain as well as two oxygenated β-ionone-type rings), Astaxanthin seems to share some chemical equivalents with ubiquinol (contains a single benzylquinone head-group ring). Astaxanthin may therefore interfere with the association of ubiquinol with CIII, affecting RISP stability, the assembly state

of the holo-CIII or the integrity of the CI-CIII₂ supercomplex. Another possible effect of Astaxanthin may involve cardiolipin, a unique phospholipid that is an important component of the inner-mitochondrial membrane. Crystal structures show that cardiolipin (CL) is tightly associated with CIII in vivo (Lange *et al.* 2001) and affects the assembly of various respiratory complexes (Paradies *et al.* 2014). The absorption of Astaxanthin (in vivo or in vitro) may interfere with the association of CL with CIII and thereby affects its biogenesis and function. However, such speculations require further investigation.

At this point, it is also not clear to us why RISP was reduced in *C. elegans* fed with Astaxanthin, but seem to remain intact in the ‘*in organello*’ assays of mitochondria treated with Astaxanthin (Figs. 3 and S4). The data indicate that Astaxanthin affects the assembly of CIII, as evident by the appearance of lower molecular mass particles corresponding to sub-CIII particles in the mitochondria. Disassembly of CIII is expected to affect the stability of various CIII subunits, including RISP, leading to their degradation over time, whereas under *in organello* assays (i.e., short incubations on ice; see Materials and Methods), RISP (or various other CIII subunits) remain intact and are less prone for degradation by the organellar proteases.

Mitochondria are major sites for energy metabolism and ATP production in eukaryotic cells, through the oxidative phosphorylation pathway. In light of the expected significance of mitochondria functions to animal and plant physiology, why does the reduced CIII activity and level in Astaxanthin-fed animals or the *isp-1* mutants result with an extended life span? Currently, we cannot provide a definitive explanation, but we speculate that these phenotypes relate to reduced mtROS production. ROS are known to induce oxidative damages of fatty acyl chains of membrane lipids, by ‘lipid-peroxidation’ (Tsuchiya *et al.* 1994). Altered membrane integrity has many pathological consequences in human, such as neurological diseases, inflammations, diabetes, cancer, and aging. Astaxanthin is a strong antioxidant that was recently suggested to reduce mtROS production in *C. elegans* and other animals (Kuraji *et al.* 2016, Liu, *et al.* 2016, Wu *et al.* 2015). Our data indicate to an additional mechanism by which Astaxanthin affects the life span of animals. The association of Astaxanthin with CIII destabilizes the native complex, resulting with lower molecular mass CIII intermediates and reduced respiration. Lowered respiration activities are likely associated with reduced rates of ‘toxic’ electron leakage from complex III, a major contributor to the generation of mtROS (Brand 2010, Chen *et al.* 2003, Quinlan *et al.* 2013), and thereby can protect the cells (and organelles) from oxidative damages. Accordingly, the aging related mechanism through mitochondria activities have been previously related to ROS production (Murakami and Murakami 2005, Quinlan, *et al.* 2013).

In summary, our data provide with novel insights into the molecular mechanism of longevity extension mediated by Astaxanthin in *C. elegans*. Biochemical and genetic studies indicate that Astaxanthin affects complex III biogenesis and activity, and may increase longevity by protecting the animal cells against ROS and oxidative stresses, due to complex III-associated ‘electron leakage’ during respiration. These effects may be universal among different eukaryotes, as similar effects on CIII biogenesis or assembly were noted in plants, worms, rats and human. Our data may also become useful for the development of multi-target drugs designed to CIII in order to prolong the lifespan of animals.

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Table 1. The effects of Astaxanthin on the enzymatic activities of mouse heart mitochondrial respiratory complexes.

Assay	Activity ¹		
	Vehicle (DMSO ²)	Astaxanthin ³	residual activity
NADH-CoQ reductase (CI)	354	142	40%
NADH-cytochrome c reductase (CI+III)	1464	78	5.3%
Succinate dehydrogenase (CII)	498	513	103%
Ubiquinone- cytochrome c reductase (CIII)	4160	1781	43%
Cytochrome c oxidase (CIV)	3638	2828	77%
Citrate synthase	1376	1402	101%

¹ - Enzyme activity (nmol O₂ · min⁻¹ · mg⁻¹)

² - 5% (v/v) DMSO

³ - 10 μM Astaxanthin

Table 2. Summary of mass-spectrometry (MS) of the ~500 kDa (higher) and ~350 kDa (lower) particles in animals (human) and plant (cauliflower) mitochondria, treated with Astaxanthin.

	Human		Cauliflower	
Associated respiratory complex	No. of subunits identified in the 500 kDa band	No. of subunits identified in the 350 kDa band	No. of subunits identified in the 500 kDa band	No. of subunits identified in the 350 kDa band
CI (NADH-dehydrogenase)	40	20	11	9
CII (succinate dehydrogenase)	2	2	2	2
CIII (Cytochrome c reductase)	13	12	11	10
Rieske iron-sulfur protein	+	+	+	+
CIV (Cytochrome c oxidase)	15	11	1	1
CV (ATP-synthase)	13	11	8	7

Figure legends

Figure 1. The effects of Astaxanthin on the lifespan of *C. elegans*.

Survival curves of *C. elegans* fed with Astaxanthin. Between 70-80 animals were used in each experiment. **(A)**. Animals were fed with wild-type *Paracoccus marcusii* (produce Astaxanthin, green), *P. marcusii* Δ crtB mutant (lacks carotenoids, orange) or *E. coli* (OP50) cells (blue). The results are the average of 4 independent experiments. **(B)**. *C. elegans* fed with *E. coli*-pASTA (expressing Astaxanthin, green) and *E. coli* cells containing an empty vector (*E. coli* pACYC184, orange). The mean lifespan (in days) was 19.97 ± 0.74 , compared to 17.34 ± 0.69 in the control cells. The results are the average of 5 independent experiments. **(C)**. Animals fed with purified Astaxanthin solubilized with DMSO (green), *E. coli* (OP50) cells (orange) and *E. coli* (OP50) cells + DMSO (blue). The results are the average of 5 independent experiments. **(D)**. The half-life's of worms fed with *P. marcusii* (produce Astaxanthin, green) and *P. marcusii* Δ crtB mutant (lacks carotenoids, orange). The results are the average of 7 independent experiments. The values are means of five biological replicates (error bars indicate one standard deviation). Asterisk in panel D indicates a significant difference from control (Student's T-test, P 0.05). Lifespan curves were analyzed by plotting Kaplan-Meier survival curves (Goel, *et al.* 2010). Mean lifespan data was compared using Log-rank test (Mantel 1966) with appropriate correction for multiple comparisons OASIS-2.

Figure 2. The effects of Astaxanthin on the longevity of *C. elegans* mutated in different life-extending genes

Survival curves of nematodes mutated in genes that regulate lifespan through different pathways supplemented with Astaxanthin. Worms mutated in the *daf-2(e1370)*, *daf-16(mu86)*, *eat-2(ad1116)*, *glp-1(or178)*, *isp-1(qm150)*, *mev-1(kn1)*, *gas-1(fc21)* or *clk-1(e2519)* were fed with *P. marcusii* (produce Astaxanthin, red), *P. marcusii* Δ crtB mutant (lacks carotenoids, blue). Nematode survival was calculated with the Kaplan-Meier method (Goel, *et al.* 2010), and differences in survival rates were tested for significance using the log-rank test (Mantel 1966), with appropriate correction for multiple comparisons OASIS-2.

Figure 3. The Effect of Astaxanthin on isolated plant mitochondria.

(A). Astaxanthin was added at different concentrations (i.e., 0, 9 and 14 μ M) to crude mitochondria preparations, obtained from *C. elegans*, human and mice. Native organellar complexes were separated by BN-PAGE, followed by Coomassie Blue staining. The position

of complex III, which partially disassembled in the presence of Astaxanthin, is indicated by arrows. **(B)**. The effects of Astaxanthin on various respiratory complexes in cauliflower mitochondria was analyzed by BN-PAGE assays. **(C)** Increasing amounts of Astaxanthin (as indicated in the gel) were added to isolated mitochondria of cauliflower. Arrows indicate to native complexes I, III and V. The effects of Astaxanthin on CIII was analyzed by western-blot analysis with antibodies raised against plant-specific RISP (Rieske iron sulfur) protein. The blots indicate that Astaxanthin cause a reduction in the levels of the holo-complex III and supercomplex I+III, and to the appearance of protein-bands of lower molecular-masses.

Figure 4. Changes in the mitochondria RISC complex of *C. elegans* grown on bacteria expressing Astaxanthin. **(A)**. Western-blot analysis of crude mitochondria obtained from wild-type *C. elegans* at different concentrations, *C. elegans* fed with Astaxanthin (+Asta), and *C. elegans isp-1* mutant-line. Antibodies against NUF5 (a subunit of the mitochondrial NADH dehydrogenase (ubiquinol) of the mitochondrial complex I (ortholog in *C. elegans* is termed as C25H3.9), Rieske iron sulphur protein (ISP-1) subunit of the mitochondrial complex III, COX1 (cytochrome c oxidase subunit 1) of the mitochondrial complex IV (the ortholog in *C. elegans* is termed as CTC-1), ATP5A (ATP synthase subunit alpha subunit) of mitochondrial complex V (the orthologue in *C. elegans* is ATP-5), and Cyt C (cytochrome c reductase; the ortholog in *C. elegans* is CYC-1), were used in immunoblotting assays. **(B)**. SDS-PAGE followed by Coomassie staining of isolated mitochondria from *C. elegans* fed with Astaxanthin (+Asta). Controls included crude mitochondria isolated from wild-type *C. elegans* that were not fed with Astaxanthin, and mitochondria isolated from the *C. elegans isp-1* mutant-line.

Figure 5. Respiration activities in *C. elegans* wild-type and *isp-1* mutants fed with Astaxanthin. O₂-uptake rates of wild-type, Astaxanthin-fed animals and *isp-1* mutants were analyzed with a Clark-type electrode as described previously (Cohen, *et al.* 2014). For each assay, equal amounts of dense packed worms from 5 different plates were submerged in 2.25 mL M9 buffer and applied to the electrode in a sealed glass chamber in the dark. O₂-uptake rates were measured in the absence (Control) or in presence of rotenone (+ROT, 50 µM), Antimycin A (Anti A, 50 µM) and KCN (1 mM) which inhibit complexes I, III and IV activities (respectively). The values are means of four biological replicates. Error bars indicate one standard deviation.

Supplementary data

Table S1. Carotenoid composition in bacterial strains used in this work.

Table S2. List of antibodies used for the analysis of astaxanthin effects on *C. elegans* lifespan.

Table S3. Mass-spectrometry (MS) of the ~500 kDa (higher) and ~350 kDa (lower) particles in human mitochondria treated with Astaxanthin. Peptides highlighted with orange color indicate to CI subunits, in blue CII subunits, in red CIII subunits, in green CIV subunits, in yellow CV (i.e., ATP-synthase).

Table S4. Mass-spectrometry (MS) of the ~500 kDa (higher) and ~350 kDa (lower) particles in cauliflower mitochondria treated with Astaxanthin. Peptides highlighted with orange color indicate to CI subunits, in blue CII subunits, in red CIII subunits, in green CIV subunits, in yellow CV (i.e., ATP-synthase).

Figure S1. Chemical structures of the carotenoids Zeaxanthin and Astaxanthin compared with ubiquinone.

Molecular structures of Zeaxanthin and Astaxanthin as well as ubiquinone.

Figure S2. Lifespan of *C. elegans* fed with Zeaxanthin.

A survival curve of nematodes supplemented with zeaxanthin in different stages of the development. About 75 animals were fed with *P. marcusii* bacteria that produce zeaxanthin (red), control *P. marcusii* bacteria (blue) or control *E. coli* OP50 bacteria (green). Animal survival was calculated with the Kaplan-Meier method, and survival differences were tested for significance using the log-rank test compared with the control. The mean lifespans (in days) of the different groups were 18.32 ± 0.53 , 23.43 ± 0.57 and 16.29 ± 0.73 , respectively. Lifespan curves were analyzed by plotting Kaplan-Meier survival curves (Goel, *et al.* 2010). Mean lifespan data was compared using Log-rank test (Mantel 1966) with appropriate correction for multiple comparisons OASIS-2.

Figure S3. Astaxanthin effect on the median lifespan at the adult stage.

A survival curve of nematodes supplemented with Astaxanthin in different stages of the development. About 75 animals were used in the experiment. Worms were fed with *Paracoccus marcusii* with no carotenoids expression from the embryo stage until L4 stage, and from L4 stage until their death with *P. marcusii* which produce Astaxanthin (blue), another group of

worms were fed from the embryo stage until L4 stage with *P. marcusii* which produce Astaxanthin, and from L4 stage until their death with P.M. with no carotenoids expression (red). The two control groups included (i) worms which were fed from the embryo stage until their death, with *P. marcusii* with Astaxanthin (green); (ii) worms which were fed from the embryo stage until their death with *P. marcusii* with no carotenoids expression (purple). *C. elegans* survival was calculated with the Kaplan-Meier method, and survival differences were tested for significance using the log rank test compared to the control. The mean lifespans (in days) of the groups given *P. marcusii* with no carotenoids expression during development and *P. marcusii* with Astaxanthin during adulthood, *P. marcusii* with Astaxanthin during development and *P. marcusii* with no carotenoids expression during adulthood, *P. marcusii* with Astaxanthin during all their lives and *P. marcusii* with no carotenoids expression during all their lives were 19.87 ± 0.48 , 17.95 ± 0.57 , 20.18 ± 0.55 and 18.22 ± 0.49 , respectively. Lifespan curves were analyzed by plotting Kaplan-Meier survival curves (Goel, *et al.* 2010). Mean lifespan data was compared using Log-rank test (Mantel 1966) with appropriate correction for multiple comparisons OASIS-2.

Figure S4. Astaxanthin from *P. marcusii* is absorbed by *C. elegans* cells.

(A). Stereo microscope photograph of Astaxanthin-fed animals. *C. elegans* fed with *P. marcusii* expressing Astaxanthin from the embryo stage for 5 days (left panel). After 5 days the worms were fed with *E. coli* bacteria for 24 hours, in order to wash their intestines (middle panel). The control worms were fed just with *E. coli* bacteria (right panel). (B). Quantification of Astaxanthin in *P. marcusii* and in *E. coli* cells by HPLC. The amount of Astaxanthin is shown as microgram per gram dry weight ($\mu\text{g/gDW}$). Stereomicroscope images of Astaxanthin-fed animals. *C. elegans* were fed with *P. marcusii* containing Astaxanthin (left and middle panels) or control *P. marcusii* (left panel) from the embryo stage until adulthood. Then one group of worms were fed with *E. coli* for 24 hours, which cleans their intestine (middle panel).

Figure S5. Accumulation of respiratory complexes in *C. elegans* mitochondria treated with astaxanthin.

(A). Crude mitochondria were obtained from *C. elegans* grown in the absence (-) and presence of *P. marcusii*. Mitochondrial complexes were separated by BN-PAGE. (B). Crude mitochondria obtained from *C. elegans* were treated in absence (-) or presence (+) of Astaxanthin (10 μM), Lycopene (5 and 10 μM), and Zeaxanthin (5 and 10 μM), and then

separated by BN-PAGE. The positions of the holo-complex III dimer ($CIII_2$) and sub- $CIII$ ($CIII^*$) particles are indicated by arrows.

Figure S6. Accumulation of respiratory complexes in cauliflower mitochondria treated with different concentrations of Astaxanthin.

Astaxanthin, at various concentrations (as indicated in the blots), was added to cauliflower mitochondria. The mitochondrial complexes were then separated by BN-PAGE, and the gel was stained by Coomassie brilliant blue. Antibodies raised against CA2 (Carbonic anhydrase subunit 2) and ATPA (ATP synthase subunit alpha) were used to assay the levels and integrity of the organellar complexes. The positions of the native CI (~1,000 kDa), CV (about 600 kDa) and supercomplex I+III are indicated.

Figure 1

Figure 2

Figure 3

A

B

Figure 4

	Respiration	% of inhibition		
	Mock	ROT (50 μM)	Anti-A	KCN (1 mM)
WT	28.83 ± 3.07	55.17 ± 2.95	66.17 ± 0.70	85.71 ± 5.18
<i>isp-1</i>	14.33 ± 2.88	34.17 ± 1.33	34.71 ± 0.99	95.22 ± 0.19
<i>astaxanthin</i>	15.03 ± 3.03	39.09 ± 1.44	40.20 ± 0.84	90.05 ± 2.35

Figure 5

Zeaxanthin

Astaxanthin

Ubiquinone

Supplementary Figure S1. Chemical structures of the carotenoids Zeaxanthin and Astaxanthin compared with ubiquinone.

Supplementary Figure S2. Lifespan of *C. elegans* fed with Zeaxanthin.

Supplementary Figure S3. Astaxanthin effect on the median lifespan at the adult stage.

A

C. elegans fed with *P. marcusii*

C. elegans after intestine cleaning

C. elegans (control)

B

HPLC quantitative analysis of astaxanthin in bacteria

Sample	Mass of sample (mg)	Amount of astaxanthin (μg/gDW)
<i>E.coli</i> + Asta	0.311	146.3066
<i>P. marcusii</i> + Asta	0.422	715.9167

Supplementary Figure S4. *Astaxanthin from P. marcusii is absorbed by C. elegans cells.*

A

B

Supplemental Figure S5. Accumulation of respiratory complexes in *C. elegans* mitochondria treated with Astaxanthin, Lycopene and Zeaxanthin.

Supplemental Figure S6. Accumulation of respiratory complexes in cauliflower mitochondria treated with different concentrations of astaxanthin.

Supplementary Table S1. Carotenoid composition in bacterial strains used in this work

Name	Carotenoid composition
<i>Escherichia coli</i> WT	None
<i>E. coli</i> pASTA	Astaxanthin
<i>E. coli</i> pACYC184	None
<i>Paracoccus marcusii</i> (<i>P. marcusii</i>) wild-type	Largely Astaxanthin Minor: Phoenicoxanthin Adonixanthin, 3'-Hydroxyechinenone
<i>P. marcusii</i> ΔcrtB	None
<i>P. marcusii</i> ΔcrtW	Zeaxanthin
<i>P. marcusii</i> ΔcrtY	Lycopene

Supplementary Table S2. List of antibodies used for the analysis of astaxanthin effects on *C. elegans* lifespan.

Antibody	Protein I.D.	origin	serum	dilution	Reference / source
Atp-A	Mitochondrial ATP-synthase subunit α	<i>Zea mays</i>	Mouse (monoclonal)	1/5,000	(Michael <i>et al.</i> 1993)
ATP-5	ATP synthase subunit, mitochondrial	<i>Caenorhabditis elegans</i>	Rabbit (polyclonal)	1/1,000	Thermo Fisher Scientific
C25H3	Uncharacterized protein, mitochondrial respiratory complex I	<i>Caenorhabditis elegans</i>	Rabbit (polyclonal)	1/1,000	Sigma-Aldrich Co. LLC
CA2	γ -carbonic anhydrase-like subunit 2	<i>Arabidopsis thaliana</i>	Rabbit (polyclonal)	1/1,000	(Perales <i>et al.</i> 2005, Sunderhaus <i>et al.</i> 2006)
CTC-1	Cytochrome c oxidase subunit 1, complex IV, mitochondrial	<i>Caenorhabditis elegans</i>	Mouse (monoclonal)	1/1,000	Thermo Fisher Scientific
CYC-1	Cytochrome C, mitochondrial	<i>Caenorhabditis elegans</i>	Rabbit (polyclonal)	1/1,000	Schagger H.H. et al. (2002)
ISP-1	Cytochrome b-c1 complex subunit Rieske, mitochondrial	<i>Caenorhabditis elegans</i>	Mouse (monoclonal)	1/1,000	Thermo Fisher Scientific
RISP	Rieske iron-sulfur protein	<i>Arabidopsis thaliana</i>	Rabbit (polyclonal)	1/5,000	Gift of Prof. Ian Small, UWA

Project Overview										Detailed Task List										Resource Allocation										Timeline & Milestones									
Project Name	Manager	Status	Priority	Start Date	End Date	Budget	Actual Cost	Progress %	Risk Level	Task ID	Task Name	Sub-Task	Assignee	Start Date	End Date	Progress %	Resource	Capacity	Availability	Task ID	Task Name	Sub-Task	Assignee	Start Date	End Date	Progress %	Resource	Capacity	Availability	Task ID	Task Name	Sub-Task	Assignee	Start Date	End Date	Progress %	Resource	Capacity	Availability
Project A: New Product Development										Task 1.1: Market Research										Resource 1: Marketing Team										Milestone 1.1: Research Complete									
Project B: System Upgrade										Task 1.2: System Analysis										Resource 2: IT Department										Milestone 1.2: Analysis Complete									
Project C: Customer Service Improvement										Task 1.3: User Feedback Collection										Resource 3: Customer Support Team										Milestone 1.3: Feedback Analysis									
Project D: Marketing Campaign Launch										Task 1.4: Campaign Strategy Development										Resource 4: Marketing Team										Milestone 1.4: Strategy Approved									
Project E: HR Policy Review										Task 1.5: Policy Drafting										Resource 5: HR Department										Milestone 1.5: Draft Review									
Project F: Financial Reporting Automation										Task 1.6: Requirement Gathering										Resource 6: Finance Department										Milestone 1.6: Requirements Document									
Project G: Legal Compliance Audit										Task 1.7: Audit Preparation										Resource 7: Legal Department										Milestone 1.7: Audit Report									
Project H: IT Security Enhancement										Task 1.8: Vulnerability Assessment										Resource 8: IT Security Team										Milestone 1.8: Assessment Report									
Project I: Sales Process Optimization										Task 1.9: Process Mapping										Resource 9: Sales Department										Milestone 1.9: Process Map									

1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	100	101	102	103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118	119	120	121	122	123	124	125	126	127	128	129	130	131	132	133	134	135	136	137	138	139	140	141	142	143	144	145	146	147	148	149	150	151	152	153	154	155	156	157	158	159	160	161	162	163	164	165	166	167	168	169	170	171	172	173	174	175	176	177	178	179	180	181	182	183	184	185	186	187	188	189	190	191	192	193	194	195	196	197	198	199	200	201	202	203	204	205	206	207	208	209	210	211	212	213	214	215	216	217	218	219	220	221	222	223	224	225	226	227	228	229	230	231	232	233	234	235	236	237	238	239	240	241	242	243	244	245	246	247	248	249	250	251	252	253	254	255	256	257	258	259	260	261	262	263	264	265	266	267	268	269	270	271	272	273	274	275	276	277	278	279	280	281	282	283	284	285	286	287	288	289	290	291	292	293	294	295	296	297	298	299	300	301	302	303	304	305	306	307	308	309	310	311	312	313	314	315	316	317	318	319	320	321	322	323	324	325	326	327	328	329	330	331	332	333	334	335	336	337	338	339	340	341	342	343	344	345	346	347	348	349	350	351	352	353	354	355	356	357	358	359	360	361	362	363	364	365	366	367	368	369	370	371	372	373	374	375	376	377	378	379	380	381	382	383	384	385	386	387	388	389	390	391	392	393	394	395	396	397	398	399	400	401	402	403	404	405	406	407	408	409	410	411	412	413	414	415	416	417	418	419	420	421	422	423	424	425	426	427	428	429	430	431	432	433	434	435	436	437	438	439	440	441	442	443	444	445	446	447	448	449	450	451	452	453	454	455	456	457	458	459	460	461	462	463	464	465	466	467	468	469	470	471	472	473	474	475	476	477	478	479	480	481	482	483	484	485	486	487	488	489	490	491	492	493	494	495	496	497	498	499	500	501	502	503	504	505	506	507	508	509	510	511	512	513	514	515	516	517	518	519	520	521	522	523	524	525	526	527	528	529	530	531	532	533	534	535	536	537	538	539	540	541	542	543	544	545	546	547	548	549	550	551	552	553	554	555	556	557	558	559	560	561	562	563	564	565	566	567	568	569	570	571	572	573	574	575	576	577	578	579	580	581	582	583	584	585	586	587	588	589	590	591	592	593	594	595	596	597	598	599	600	601	602	603	604	605	606	607	608	609	610	611	612	613	614	615	616	617	618	619	620	621	622	623	624	625	626	627	628	629	630	631	632	633	634	635	636	637	638	639	640	641	642	643	644	645	646	647	648	649	650	651	652	653	654	655	656	657	658	659	660	661	662	663	664	665	666	667	668	669	670	671	672	673	674	675	676	677	678	679	680	681	682	683	684	685	686	687	688	689	690	691	692	693	694	695	696	697	698	699	700	701	702	703	704	705	706	707	708	709	710	711	712	713	714	715	716	717	718	719	720	721	722	723	724	725	726	727	728	729	730	731	732	733	734	735	736	737	738	739	740	741	742	743	744	745	746	747	748	749	750	751	752	753	754	755	756	757	758	759	760	761	762	763	764	765	766	767	768	769	770	771	772	773	774	775	776	777	778	779	780	781	782	783	784	785	786	787	788	789	790	791	792	793	794	795	796	797	798	799	800	801	802	803	804	805	806	807	808	809	810	811	812	813	814	815	816	817	818	819	820	821	822	823	824	825	826	827	828	829	830	831	832	833	834	835	836	837	838	839	840	841	842	843	844	845	846	847	848	849	850	851	852	853	854	855	856	857	858	859	860	861	862	863	864	865	866	867	868	869	870	871	872	873	874	875	876	877	878	879	880	881	882	883	884	885	886	887	888	889	890	891	892	893	894	895	896	897	898	899	900	901	902	903	904	905	906	907	908	909	910	911	912	913	914	915	916	917	918	919	920	921	922	923	924	925	926	927	928	929	930	931	932	933	934	935	936	937	938	939	940	941	942	943	944	945	946	947	948	949	950	951	952	953	954	955	956	957	958	959	960	961	962	963	964	965	966	967	968	969	970	971	972	973	974	975	976	977	978	979	980	981	982	983	984	985	986	987	988	989	990	991	992	993	994	995	996	997	998	999	1000	1001	1002	1003	1004	1005	1006	1007	1008	1009	1010	1011	1012	1013	1014	1015	1016	1017	1018	1019	1020	1021	1022	1023	1024	1025	1026	1027	1028	1029	1030	1031	1032	1033	1034	1035	1036	1037	1038	1039	1040	1041	1042	1043	1044	1045	1046	1047	1048	1049	1050	1051	1052	1053	1054	1055	1056	1057	1058	1059	1060	1061	1062	1063	1064	1065	1066	1067	1068	1069	1070	1071	1072	1073	1074	1075	1076	1077	1078	1079	1080	1081	1082	1083	1084	1085	1086	1087	1088	1089	1090	1091	1092	1093	1094	1095	1096	1097	1098	1099	1100	1101	1102	1103	1104	1105	1106	1107	1108	1109	1110	1111	1112	1113	1114	1115	1116	1117	1118	1119	1120	1121	1122	1123	1124	1125	1126	1127	1128	1129	1130	1131	1132	1133	1134	1135	1136	1137	1138	1139	1140	1141	1142	1143	1144	1145	1146	1147	1148	1149	1150	1151	1152	1153	1154	1155	1156	1157	1158	1159	1160	1161	1162	1163	1164	1165	1166	1167	1168	1169	1170	1171	1172	1173	1174	1175	1176	1177	1178	1179	1180	1181	1182	1183	1184	1185	1186	1187	1188	1189	1190	1191	1192	1193	1194	1195	1196	1197	1198	1199	1200	1201	1202	1203	1204	1205	1206	1207	1208	1209	1210	1211	1212	1213	1214	1215	1216	1217	1218	1219	1220	1221	12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Genomic coordinates (chr:pos)										chr	pos	ref	alt	qual	filter	info	format
chr1:100000000										chr1	100000000	A	G	100	0	AD=100	0
chr1:100000001										chr1	100000001	A	G	100	0	AD=100	0
chr1:100000002										chr1	100000002	A	G	100	0	AD=100	0
chr1:100000003										chr1	100000003	A	G	100	0	AD=100	0
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chr1:100000070										chr1	100000070	A	G	100	0	AD=100	0
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chr1:100000123										chr1	100000123	A	G	100	0	AD=100	0
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chr1:10000																	