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**Direct in vivo imaging of ferrous iron dyshomeostasis in ageing Caenorhabditis elegans**

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Iron is essential for eukaryotic biochemistry. Systematic trafficking and storage is required to maintain supply of iron while preventing it from catalysing unwanted reactions, particularly the generation of oxidising reactive species. Iron dyshomeostasis has been implicated in major age-associated diseases including cancers, neurodegeneration and heart disease. Here, we employ population-level X-ray fluorescence imaging and native-metalloproteomic analysis to determine that altered iron coordination and distribution is a pathological imperative of ageing in the nematode, *Caenorhabditis elegans*. Our approach provides a method to simultaneously study iron metabolism across different scales of biological organisation, from populations to cells. Here we report how and where iron homeostasis is lost during *C. elegans* ageing, and its relationship to the age-related elevation of damaging reactive oxygen species. We find that wild types utilise ferritin to sustain longevity, buffering against exogenous iron and showing rapid ageing if ferritin is ablated. After reproduction, escape of iron from safe-storage in ferritin raised cellular Fe^{2+} load in the ageing *C. elegans*, and increased generation of reactive species. These findings support the hypothesis that iron-mediated processes that drive senescence. We propose that loss of iron homeostasis may be a fundamental and inescapable consequence of ageing that may represent a critical target for therapeutic strategies to improve health outcomes in ageing.

**Introduction**

The near-universal requirement of iron in life is due to its ability to mediate single-electron chemistry under physiological conditions, from ATP production to oxygen transport. The unique chemical properties of iron facilitate innumerable processes, suggesting that iron dyshomeostasis may be an upstream event preceding biological senescence.

Age is the greatest risk factor for a number of diseases including cancer, heart disease and neurodegeneration. The associated impairments to iron metabolism and increased iron-mediated oxidative stress have been implicated in disease pathogenesis. The precise mechanism through which iron imparts cytotoxicity depends on the chemical state in which the metal exists within the cell. Biological iron is predominantly either ferrous (Fe^{2+}) or ferric (Fe^{3+}), with the majority of bioavailable iron tightly bound by dedicated enzymes and storage proteins, and a small proportion likely existing in dynamic equilibrium as low molecular weight complexes. The normally desirable propensity of iron to participate in single-electron redox chemistry is non-specific; unchecked interaction of Fe^{2+} with biological oxygen species can generate powerful oxidants and thus become cytotoxic. The reactivity of iron is determined by local bonding environment. Biological ligands typically prevent iron from engaging in side reactions particularly the indiscriminate reduction of endogenous cellular peroxides (i.e. the Fenton reaction) to generate the damaging OH• radical.

Iron accumulation appears to be a common feature of ageing organisms. In the human brain, for instance, both iron and proteins associated with iron metabolism are elevated in aged individuals, particularly in regions most susceptible to damage in diseases including Alzheimer’s and Parkinson’s. Stable isotope tracing in rats has identified an imbalance favouring iron influx over efflux, leading to an approximately 30% increase in brain iron as the animals aged. In concert with this gradual yet significant elevation in iron levels, a changing redox environment favouring unrestrained oxidative activity of Fe^{2+} is an intriguing candidate that might explain both age-related disease and general senescence. For example,
ceruloplasmin-deficient mice that lack the capacity to oxidise Fe^{2+} to Fe^{3+} display marked age-dependent neurodegeneration.\(^8\)

Direct appraisal of iron redox chemistry in vivo is an analytical challenge. Post mortem artefacts arising from tissue handling, fixation and cryoprotection are all potential sources of experimental error, including metal redistribution and altered cellular redox state.\(^9\) Achieving mechanistic insight into the specific redox chemistry associated with iron in an ageing organism requires the capacity to obtain snapshots of true biophysical conditions with minimal disturbance to normal Fe^{2+}/Fe^{3+} balance. When applied to this problem the results of any analytical technology are only as relevant as the degree to which the native redox state of the cell is maintained. For instance, fluorescent probes offer one method to assess native redox state.\(^10\) Hard X-ray microprobes and mass spectrometry also offer an opportunity to probe biological microstructures at high resolution,\(^11\) and are compatible with hydrated specimens.\(^12\)

Synchrotron-based X-ray fluorescence microspectroscopy allows direct quantitative appraisal of the abundance of inorganic species, as well as a means to assess coordination environment within intact specimens at micrometer length scales. Combined with the ability to image live, hydrated biological samples, this unique approach is a step closer to permitting accurate measurement of the true in vivo chemical environment in which iron exists in the ageing cell.

The nematode Caenorhabditis elegans provides an excellent model to explore the bioinorganic chemistry of ageing in an entire multicellular organism.\(^13,14\) Like higher-order counterparts, C. elegans deposit fat, accumulate lipofuscin, develop sarcopenia and suffer neurodegeneration with age. Genetic modulation of ageing has been well characterised in this animal,\(^15\) however, the biochemical processes and underlying molecular mechanisms driving these events are uncertain.
We have previously identified that *C. elegans* express high concentrations of iron in intestinal cells,\textsuperscript{16,17} which house the majority of metabolic processes in this animal. To expand our understanding of the chemistry of ageing we here imaged iron coordination chemistry within live *C. elegans*, quantitated age-related iron accumulation, and assessed biochemical iron alterations in this genetic model of longevity. A complementary native metalloproteomic analysis identified senescent changes in endogenous iron ligands. We propose that systematic failure of iron metabolism is a major contributor to the ageing process, which has implications for understanding the risk for several age-related diseases.

**Results**

**Iron accumulation and redistribution during ageing**

We have previously shown that *C. elegans* adults exhibit a highly compartmentalised bioinorganic architecture,\textsuperscript{16,17} including high iron concentrations in the metabolically active intestinal cells. Using synchrotron-based X-ray fluorescence microscopy (XFM; \(\sim 2\) µm resolution) we quantified the spatial distribution of iron in young and aged intact wild-type adults (Fig. 1a). This revealed a 77 % increase in mean total iron from young (4 days post egg lay, 62.5 pg iron per individual) to post-reproductive adults (12-day old, 110.4 pg iron per individual; two-tailed unpaired Student’s *t*-test \(p < 0.001\), Fig. 1b), with marked intracellular accumulation in the intestinal cells (Fig. 1a). Measurement of iron using inductively coupled plasma-mass spectrometry (ICP-MS) in aged *C. elegans* cohorts (\(n = 100\) adults per aliquot) closely corresponded and confirmed a significant increase in total iron ([Supplementary Fig. 1a](#)). For instance, post-reproductive senescent 10-day old adults had a 66 % increase in mean iron compared to 6-day old adults (two-tailed unpaired Student’s *t*-test, \(p < 0.001\)). Iron also increased with age when normalised against dry weight, indicating that the iron elevation was not due to increased body mass ([Supplementary Fig. 1b](#)).

[Fig. 1](#) Elevated iron in *C. elegans* promotes oxidative stress in *vivo*. (a) Live imaging of exchangeable iron using calcein-AM fluorescence in (top) young (4-day) and (bottom) aged (12-day) *C. elegans*. Fluorescence is quenched by chemically accessible iron in the aged intestine. Bright field image (above) and fluorescence image (below) with an outline in white. (b) In vivo oxidative activity was detected by dichlorofluorescein (DCF) fluorescence and increased in the intestine of old (bottom) compared to young (top) *C. elegans*. (c) Quantification of *in vivo* reactive species by DCF fluorescence. In the ageing worm, fluorescence significantly increased with age relative to 4-day old adults (mean \(\pm\) SEM, \(n = 9, 7\) and 5 individuals worms respectively; two-tailed unpaired Student’s *t*-test, \(* * *\) \(p < 0.001\)). (d) Ex vivo ROS generation is a product of iron accumulation. Long-lived daf-2 mutants lacked the increase in levels of reactive species with age (measured by DCF fluorescence; relative to 4-day old adults) compared to wild type and daf-16:daf-2 mutants (mean \(\pm\) SEM, \(n = 4\); two-tailed unpaired Student’s *t*-test, \(* * *\) \(p < 0.001\)).

To examine the presence of potentially harmful non-heme iron, we used histological Perls’ staining, visually enhanced with 3,3’-diaminobenzidine (DAB).\textsuperscript{22} Consistent with the XFM data, we found an age-dependent accumulation of non-heme iron in intestinal cells, progressing from discrete vesicular to dispersed distribution ([Fig. 1c](#)). Aged individuals also exhibited iron staining within intestinal cell nuclei, and dense inclusions in the head. Adult *C. elegans* are post-mitotic and lose intestinal cell nuclei during ageing by an unknown process,\textsuperscript{23} which our histological findings indicate might be caused by toxic accumulation of nuclear non-heme iron. In addition, we observed conspicuous iron accumulation in the germ nuclei of post-reproductive adults ([Supplementary Fig. 4](#)).

We hypothesised that age-dependent elevation of iron is a significant source of harmful redox activity if the accumulation is not coordinated in a redox-silenced manner. Both XFM and Perls staining revealed significant changes in iron concentration, cellular distribution and chemical state. Consequently, we assessed the contribution of elevated iron to age-related oxidative stress.

**Accumulated iron contributes to oxidative stress**

The relationship between reactive iron elevation and oxidative damage within post-mitotic intestinal cells during ageing was examined in living *C. elegans*. The calcium sensor calcein exhibits decreased fluorescence in the presence of chemically available iron. Consistent with an accumulation of calcein-accessible iron, *in vivo* fluorescence was suppressed in aged adults ([Fig. 2a](#)). This suppression could not be explained by a
Iron storage capacity of ferritin is compromised during ageing. (a) Native soluble iron-binding species separated and detected by size exclusion chromatography-inductively coupled plasma-mass spectrometry (SEC-ICP-MS). In ageing C. elegans, greater levels of iron are associated with high molecular weight (Peak #1) and low molecular weight (Peak #3) species, whilst ferritin iron is decreased (Peak #2). (b) Integration of these three major chromatographic peaks across age (mean ± SEM, n = 3, one-way ANOVA with Dunnett’s post hoc test, **p < 0.01, *** p < 0.001). Linear regression analysis showed correlation between iron levels in each peak and age. (c) Loss of ftn-2 (null) ablated soluble ferritin-associated iron in young adults. In contrast, ftn-2 nulls did not differ from wild type, retaining the same iron-binding protein profile. (d) Representative X-ray fluorescence micrographs of wild type (i), ftn-1 null (ii) and ftn-2 null (iii) young adults confirmed loss of ftn-2 and resulted in a specific decrease in total iron with unaltered calcium levels. (e) Quantification of iron per adult revealed ftn-2 (ferritin) accounts for approximately 40% total iron in young C. elegans (mean ± SEM, n = 3, two-tailed unpaired Student’s t-test, **p < 0.01).

Increased accessible iron appeared concurrently with elevated formation of reactive species in vivo, as measured by oxidation of 2',7'-dichlorodihydrofluorescein (DCF) to dichlorofluorescein (DCF; Fig 2b). DCF fluorescence was markedly increased in the intestine compared to young adults (two-tailed unpaired Student’s t-test, p < 0.001; Fig. 2c), consistent with a pro-oxidant chemical environment within the aged C. elegans intestine.

To confirm whether age-dependent iron elevation is a catalytic source of the in vivo oxidative stress, we measured DCFH oxidation in the soluble fraction of lysed C. elegans. This ex vivo oxidative activity from wild type C. elegans increased throughout lifespan (two-tailed unpaired Student’s t-test, p < 0.001; Fig. 2d), matching the observed steady-state increase in oxidation in vivo (Fig. 2b). The increased rate of ex vivo DCF fluorescence is consistent with an increased pool of catalytically reactive iron because it was abolished by the redox-silencing chelator diethylenetriamine penta-acetic acid (DTPA) (Supplementary Fig. 6a). Paralleling iron levels, pro-oxidant chemistry was significantly attenuated in the long-lived daf-2 mutants at each of the ages studied, remaining below the reference baseline rate (4-day old wild type), even by 16 days of age (Fig. 2d, Supplementary Fig. 6b). As with its impact on iron levels with ageing, loss of daf-16 function increased oxidative activity in the abbreviated lifespan of the daf-16;daf-2 double mutants.

Normally, careful chaperoning of the metal by protein ligands prevents unregulated iron activity. Elevated oxidative stress arising from an increased pool of accessible iron, typical of many age-related disease states, implicates a failure of containment as a feature of the ageing process.
Ageing impairs iron buffering

The cellular reservoir of exchangeable iron is normally stored by ferritin, which oxidises ferrous (Fe\(^{2+}\)) iron to the more chemically stable hydrous ferric (Fe\(^{3+}\)) form, protecting the cell from unwanted incidental redox reactions. Ferritin proteins are highly conserved and typically organise as a 24-mer capable of storing up to ~4,500 atoms of iron, although are rarely saturated.

To determine whether the age-related tandem elevation of iron and oxidative activity were caused by loss of ferritin sequestration of iron we used native size-exclusion chromatography with on-line inductively coupled plasma-mass spectrometry (SEC-ICP-MS) to examine the soluble iron-binding protein profile of wild type C. elegans homogenates. Soluble protein-bound iron resolved into three major peaks (Fig. 3a). Peak #2 represented most (~84 %) of the total soluble iron in 4-day-old adults. Protein purification and analysis by mass spectrometry identified Peak #2 as ferritin, FTN-2, (Supplementary Fig. 7). Soluble iron redistributed from ferritin towards high molecular weight (HMW, > 1 MDa; Peak #1), and low molecular weight (LMW)-iron species (< 30 kDa; Peak #3; Fig. 3b). In total, by 13 days of age, the amount of iron associated with ferritin was reduced by approximately 50 %, despite a near doubling of total iron levels (Fig. 1b, c).

Integration of all peaks indicated that total soluble iron did not change with age, rather, the increased burden of iron we observed was driven by elevations in the insoluble fraction (two-tailed unequal Student’s t-test, p < 0.001, Supplementary Fig. 8).

The C. elegans genome encodes two heavy-chain ferritin orthologs, ftn-1 and ftn-2.\(^{14}\) FTN-1 (predicted MW = 19524.8 Da) expression is induced in the intestine by high iron exposure, while FTN-2 (experimentally observed MW = 19502.2 Da; Supplementary Fig. 7f) has constitutive as well as iron-responsive expression in intestinal cells.\(^{25,26}\) Mass spectrometry detected no peptides of FTN-1 within Peak #2. Therefore, ageing is associated with the escape of now-chemically available iron from previously redox-protected storage in FTN-2 (Fig. 3b) to become species that can facilitate oxidation (Fig. 2).

Genetic ablation of FTN-2 abolished Peak #2 (ferritin-bound iron) and increased iron associated with Peak #1, whereas loss of FTN-1 had only minor effects compared to wild type (Fig. 3c), likely due to its low constitutive expression.\(^{28}\) Populations of whole wild type and null mutants for ftn-1 and ftn-2 were imaged using XFM (Fig. 3d), which revealed that ftn-2 nulls had ~46 % less total iron compared to wild type (Fig. 3e). Total iron in ftn-1 nulls was not altered. Young ftn-1;ftn-2 null animals had no detectable histological non-heme iron (Supplementary Fig. 9), demonstrating that ferritin accounts for the bulk of Perls staining in young animals (Fig. 1i).
Fig. 5  
Ageing disturbs iron redox balance  (a) Integrated intensity of 71 X-ray fluorescence micrographs spanning the iron K-edge (7100 to 7170 eV; dashed grey lines) for the anterior portion of hydrated (i; n = 4) 5-day old wild type, (ii; n = 2) 5-day old ftn-2;ftn-1 nulls; and (iii; n = 2) 12-day old wild type adults. (b) Spatial distribution of ROIs defined by grouping pixels with similar spectra using k-means clustering. ROI-1 corresponds predominately to non-intestinal tissues, whilst ROI-2 through 5 are localized to the intestine. Scale bar = 100 um. (c) Derivative iron K-edge spectra (AXRF/ΔEnergy) for each ROI. Derivative spectra were modeled as the sum of two Gaussians between 7114 and 7134 eV (solid black lines) and the peak area associated with the 1s→4s (light shaded curves) and 1s→4p (dark shaded curves) transitions used to assess the Fe$^{2+}$/Fe$^{3+}$ balance of each ROI. Impaired ferritin function in aged wild type and ftn-2;ftn-1 null adults animals resulted in a proportional elevation of Fe$^{3+}$.  

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ferritin function and/or cellular iron homeostasis, with resultant increases in pro-oxidant reactive iron (Figs. 2&3). Therefore, we sought to investigate how the aggregate coordination environment responsible for maintaining the cellular Fe\(^{2+}/Fe^{3+}\) balance changes within age.

**Corrupted buffering shifts iron oxidation state in vivo**

X-ray absorption near-edge spectroscopy (XANES) reports on the electronic structure and organisation of nearest neighbors around the absorbing atom. Ideally, XANES assessment of metal coordination in *vivo* requires procedures that avoid inducing chemical disruption, (e.g. artefacts that arise from fixation). To visualise the coordination chemistry of iron in *vivo*, we stepped beyond single point projection-based XANES and performed fluorescence-XANES imaging on intact hydrated *C. elegans*. Serial scanning of young and aged wild type, and young *fin*-2 null animals (2–4 individuals per group) across an incident energy range spanning the Fe K-edge (7112 eV) produced a hyperspectral dataset (incident energy, x position, y position, each with an associated fluorescence intensity); alternately, each pixel possesses an iron XANES spectrum.

As the relevant ligand set is difficult to define *a priori* we did not seek to localise individual iron species, rather we gained an overview of iron metabolism by identifying aggregate modes of iron coordination associated with physiological structures (Fig. 5 and Supplementary Fig. 11a-c). For biological systems the aggregate iron coordination environment is composed of numerous iron complexes (including ferritin, Fe-S and heme proteins, etc) with distinct spatial distributions and present at a wide range of concentrations. We used the approach of Lerotic *et al.*\(^{28}\) to identify the spatial distribution of unique coordination states. Principal component analysis was used to decompose the XANES spectra using k-means clustering (Supplementary Fig. 10) into distinct aggregate iron coordination states, which we used to segment the XANES image into 5-6 regions of interest (ROI).

Consistent with our earlier results, we observed decreased total iron in *fin*-2 nulls and a marked increase in aged wild types (Fig. 5a) with distinct ROIs corresponding to areas of similar iron metabolism. In all animals ROI1 (grey) was localised to tissues outside the intestine, while all other ROIs defined intestinal structures (Fig. 5b,c).

Both the 1s→3d pre-edge peak and edge position (defined as the peak in the first derivative spectra) shift to higher energy with oxidation state, independent of coordination geometry\(^{29}\) and can be used to assess the relative proportion of Fe\(^{2+}\) and Fe\(^{3+}\). Both of these features revealed changes in the dominant iron oxidation state in young *fin*-2 nulls and aged wild type animals. Compared to young wild types all *fin*-2 null ROI K-edges were shifted to lower energies (wild type having a K-edge at approximately 7127 eV). This is consistent with a biochemical shift towards increased Fe\(^{2+}\) content within these animals that are unable to safely store iron in ferritin as Fe\(^{3+}\). Aged wild types also displayed a similar shift of ROI K-edge indicative of elevated Fe\(^{2+}\) content. These data are supported by a concomitant shift in the energy of iron pre-edge feature (Supplementary Fig. 11). A unique mode of Fe coordination was identified in the aged animals (ROI6 red; Fig. 5b,c), representing distinct foci of aggregate iron coordination localised to the intestine.

The relative intensity of the 1s→4s (7114 – 7123 eV) and 1s→4p (7124 – 7134 eV) transitions provides a means for quantifying the Fe\(^{2+}/Fe^{3+}\) ratio.\(^{20,30}\) Fitting the contribution of these electronic transitions to the structure of the first derivative we determined the relative abundance of Fe\(^{2+}\) within each ROI. This analysis showed both the aged wild type and *fin* null had significantly more Fe\(^{2+}\) content than the young wild type (Fig. 5 & Supplementary Fig. 11).

**Discussion**

Accumulation of iron with age may be a universal feature of multicellular organisms, from invertebrates to humans.\(^5,5,31\) Although iron is an essential redox-active element for eukaryotes, it can induce uncontrolled oxidative chemistry if it is not appropriately regulated. The human brain appears particularly vulnerable to improper iron handling, probably because its exceptionally high metabolic output calls for constitutively high iron concentrations.

We have recapitulated the biochemical features of the ageing brain in the well-characterised animal model *C. elegans*. Using several advanced analytical approaches to study bioinorganic chemistry we identified iron dyshomeostasis as an imperative feature of *C. elegans* senescence, and we have characterised a possible mechanism for this. Our data indicate that the *C. elegans* intestine is particularly susceptible to senescent iron changes. The *C. elegans* intestine is the primary site responsible for digestion, macromolecule storage, endocrine function, detoxification of exogenous toxins, and host-pathogen interactions. The intestine comprises approximately one third of the somatic mass of *C. elegans*, has the greatest relative concentrations (>10 \(\mu\)M) of iron,\(^17\) and is the predominant site of ferritin expression and function. Collapse of iron homeostasis is marked in the intestine of aged *C. elegans*. The ILS pathway acts on intestinal DAF-16 transcription factor activity to modulate ageing,\(^32\) and therefore loss of normal intestinal cell function could mirror the accelerated ageing effects of *daf-16* mutation. Other lifespan-affecting transcription factors such as SKN-1 (a nuclear factor E2-related factor family member), also act in the intestinal cells to alter ageing rates.\(^33\) Intestinal cell nuclei are progressively lost as *C. elegans* age,\(^23\) and the resultant cumulative loss of intestinal function is likely to further compound disruption of iron homeostasis. We observed nuclear iron accumulation (Fig. 1f) within the intestine that is likely to contribute to demise of these nuclei via elevated iron-mediated free radical generation. Assays designed to directly appraise *H*\(_2*O*\)2 have also suggested nuclear accumulation of this damaging oxidant.\(^34\) Intra-nuclear aggregates of iron-rich ferritin have also been observed in mammalian models of toxic iron overload.\(^35,36\)
The age-dependent rise in iron is markedly delayed in long-lived daf-2 mutants. However, at their median lifespan, daf-2 mutants still show a significant increase in total iron (Supplementary Fig. 3), demonstrating that iron elevation invariably heralds death even where the rate of ageing is slowed. The suppression of iron accumulation accounts for the comparatively low levels of reactive species generated by the daf-2 mutants versus age-matched wild type (Fig. 2d). Loss of daf-16 by mutation reverses the longevity effects of the daf-2 mutant and not only restores but also exaggerates the age-dependent rise in iron (Fig. 1b). DAF-16 need be expressed only in the intestine to slow ageing in C. elegans. This is consistent with uncontrolled iron elevation and altered coordination in the metabolically critical intestinal cells being a primary contributor to oxidative load in aged C. elegans.

The formation of non-ferritin iron collections in HMW soluble- and insoluble forms (Fig. 3; Supplementary Fig. 8) precedes death, and is reminiscent of iron-filled intracellular inclusions, lipofuscin, neuromelanin and hemosiderin, which all feature in the pathologies of ageing and age-related disease, but have questionable or unclear roles in cytotoxicity. We hypothesise that the underlying mechanism of ageing in wild type C. elegans is the escape of iron from safe storage in ferritin (FTN-2), where it emerges as redox-active Fe³⁺ species that redistribute primarily to soluble HMW and LMW-iron, as well as insoluble precipitates, where it catalyses toxic free radical generation. HMW-iron may represent misfolded, mis-metallated proteinaceous material transitioning towards an insoluble aggregate typical of ageing pathology. The withdrawal of soluble iron from the cytoplasm into these aggregates could explain why the rise in total somatic iron is not sensed by the transcriptional mechanisms that should upregulate the expression of ferritin and other iron-regulatory elements.

The C. elegans genome encodes two ferritin genes; ftm-1 and ftm-2, however we determined that only ftm-2 contributes to iron storage under basal conditions. Although understanding the specific function of ftm-1 will require further investigation, our data are consistent with previous reports showing the impact of ftm-1 is negligible with respect to lifespan under normal conditions.

Conclusion

Extreme levels of iron can be toxic, however, our results indicate that the age-related accumulation of iron observed in C. elegans is, by itself, insufficient to cause senescence. Rather than being determined by simple accumulation, we find that normal ageing is characterised by a loss of ferritin function and/or cellular iron homeostasis, with resultant increases in reactive iron and oxidative stress. With ferrous iron a catalyst for radical generation, the clear shift from Fe²⁺ to Fe³⁺ in both aged and ferritin null worms in vivo provides a mechanism to drive senescence. These observations have broader implications for understanding how cellular iron dyshomeostasis may contribute to major age-related human diseases. Strategies derived to delay or mitigate the risks associated with ageing may consider the inevitability of contributions of late-life iron.

Methods and materials

Strains

N2 (wild-type), TJ1060: spe-9(hc88); fer-15(b26), CB1370: daf-2(e1370), and DR1309: daf-16(m26); daf-2(e1370), RB2603; ftm-1(ok3625) and RB668; ftm-2(ok404) were obtained from the Caenorhabditis Genetics Center. As the ftm-1(ok3625) deletion allele has not been mapped, we sequenced the genomic DNA across the putative deletion site. A 495 bp deletion was identified that removes all of exon 2 and most of exon 3 (Supplementary Fig. 12), resulting in a premature stop codon (UGA), so that the likely product is truncated to 20 amino acids (out of the predicted 170). The ftm-2(ok404) is a previously-characterised null allele. RB2603; ftm-1(ok3625) and RB668; ftm-2(ok404) were each backcrossed four times to wild type prior to further analysis. New strains reported in this study are listed in Supplementary Table 1. All strains were maintained at 20 °C on standard nematode growth media (NGM) and aged at 25 °C as required, with the exception of the fertility mutant TJ1060 which was maintained at 16 °C.

DNA primers 5'-ATGTTGTCAGATTTCGCC-3' and 5'-GAACCTTTTGTTGCAATA-3' were used to track the ftm-1(ok3625) deletion, and 5'-CCAGGCCAAACTACATCA-3' and 5'-AAAGAAGATGCAGACGACA-3' for the ftm-2(ok404) deletion respectively during back crossing to wild type and crossing to generate the double null strains GMC005 and GMC007.

X-ray fluorescence microscopy (XFM)

Aged C. elegans were prepared as previously described on Si₃N₄ windows (Silson) for analysis at the XFM beamline at the Australian Synchrotron and the experimental setup has been described previously. Briefly, the distribution of metals was mapped using a beam of 12.7 keV X-rays focused to 2 µm (full-width at half-maximum) using a Kirkpatrick–Baez mirror pair. The X-ray energy was chosen to induce K-shell ionisation of elements with atomic numbers below 34, while also separating the elastic and inelastic scatter from fluorescence lines of lighter elements. Specimens were continuously scanned through X-ray focus and the real-time event-mode processing approach used in the Maia detector system, installed at the Australian Synchrotron XFM beamline, allowed flexible selection of effective per pixel dwell times, though 8 ms was typical. Resulting elemental maps ranged up to 50,000 pixels in size and the total acquisition time varied around 7 minutes per specimen. Two single-element foils of known areal density, Mn and Pt (Micromatter, Canada), were scanned during the experiment as references for quantification of elements with atomic numbers below 34, while蕉 continuous scanning through X-ray fluorescence spectra was performed using a GeoPIXE v7.0f (CSIRO, Australia) and calibrated using the GeoPIXE v7.0f (CSIRO, Australia).
metal foil measurements, with corrections made for self-absorption in the sample, absorption in air, and the efficiency response of the detector.

Comparison of total iron quantitated from whole worms was examined via one-way ANOVAs with a Dunnett’s post-hoc test. Within each genotype aged cohorts were compared to young (4 days of age) adults.

Histology

C. elegans were washed in S-basal, fixed overnight in 10 % (v/v) neutral buffered formalin (NBF) at 4 °C, embedded in 2 % (w/v) agar in phosphate buffered saline (PBS) blocks and then fixed again in 10 % NBF overnight. Following processing of the agar blocks into paraffin, 5 µm sections were prepared, dewaxed and stained with DAB-enhanced modified Perl’s Prussian blue following a standard protocol. Samples were counter stained with Harris haematoxylin solution (Amber Scientific).

Live imaging of iron

C. elegans cultures were aged as indicated, washed in S-basal, then co-cultured in S-basal containing 1 x 10^6 cells OP50 (E. coli) and 0.05 µg mL^-1 calcein-AM (Invitrogen) for 1 h and then in S-basal with 1 x 10^6 cells OP50 for 1 h. Samples were then mounted for epi-fluorescence microscopy using standard techniques. Calcein fluoresces in the presence of calcium ions in solution, but this fluorescence is quenched by ionic iron. Calcein has a slight selectivity for Fe^{2+} over Fe^{3+}. Other divalent metal ions, Cu^{2+}, Ni^{2+} and Co^{2+} can also quench calcein fluorescence, however, these were found to be present at ≥ 2 orders of magnitude lower than Fe in the intestine and therefore not considered to be able to interfere with the signal.

ROS detection

2',7'-dichlorodihydrofluorescein diacetate (DCFH-DA) at 10 µM was used as a fluorescent probe for reactive oxygen species (ROS) detection in vivo using standard protocols. DCFH-DA enters and accumulates in the intestinal cells, where it is oxidised by several ROS (including hydroxyl radicals) to form the fluorophore DCF. Analysis of fluorophore fluorescence rate increase was performed on samples sonicated in 1x TBS, pH 7.4 and recovered as the supernatant from a 100,000 g centrifugation at 4 °C. Total protein concentration was determined by a Nanodrop spectrophotometer (Thermo Scientific). Ex vivo ROS detection were made by bringing lysate supernatants (50 µg total protein) to a reaction volume of 200 µl with 200 mM ammonium acetate pH 7.5, 100 µM DCFH-DA (Sigma-Aldrich, made from a 10 mM stock in acetonitrile), and 400 µM ascorbate, in black 96-well micro titer plates. Fluorescence in 8 replicate wells was quantitated (E_s: 485 nm, E_m: 535 nm using a 495 nm cut-off) by a FlexStation ( Molecular Devices) plate reader, using 30 reads every minute for 1 h at medium PMT setting. Values from lysate-free (negative) controls were subtracted and the fluorescence data was then baseline corrected. A linear regression fitted from 20-40 minutes and the slopes were normalised against 4-day old wild type and plotted (Prism v5.0d, Graphpad Software).

Fluorescence microscopy

Animals were mounted on a glass slide with 2 % (w/v) agar pad containing 2 % NaCl under a glass cover slip and examined using an Olympus BX40 epifluorescence microscope equipped with SPOT RTKE cooled color CCD camera (Diagnostic Instruments, MI). The GFP fusion protein was visualised by using an Olympus U-MWG filter set (E_l: bandpass 530 ± 20 nm, E_m: longpass 590 nm), and the images imported into SPOT software (Diagnostic Instruments). ImageJ v1.45s (NIH, USA) was used for image preparation and overlays.

Bulk iron quantitation

Total iron was measured using a 7700 series (Agilent) inductively coupled plasma mass spectrometry (ICP-MS) as previously reported. Samples consisted of 100 adults per replicate for different aged cohorts as indicated.

Purification of C. elegans ferritin

10 g of mixed stage wild type cultured on 8P media at 20 °C were frozen as pellets in liq-N_{2}. Pellets were crushed in a liq-N_{2} chilled mortar and pestle, then added to 20 ml PBS pH 7.4 with EDTA-free proteinase inhibitors (Roche). The lysate was further disrupted with 20 strokes of an ice-cooled Dounce homogeniser (in 50 mL). The extract was centrifuged at 3300 g at 4 °C and the supernatant (~30 mL) dialyzed overnight in 18 MΩ pure H_{2}O (Millipore) at 4 °C using pleated dialysis tubing with a 10 kDa molecular weight cut-off (Thermo Scientific). 20 ml of dialysate was then diluted to 60 ml with pH 3-10 ampholytes and iso-electrically focused via a Rotofor II (Bio-Rad), as per manufacturer’s protocols to 2500 Vh. Fractions were collected and iron content measured by graphite furnace atomic absorption spectroscopy (AAS). A standard method was used with ashing and atomisation temperatures of 700 °C and 2300 °C respectively, and with linear absorbance to a concentration of 100 µg L^{-1}. The three highest contiguous iron-containing fractions were then pooled and re-focused via the Rotofor II as above. Fractions containing iron were identified by AAS. To the six contiguous iron containing fractions NaCl was added to a final concentration of 150 mM and then size-excluded via FPLC (Bio-Rad) using a Superdex 200 10/300 GL column and PBS buffer at 0.6 mL min^{-1}. Fractions were collected and iron was measured by AAS. Fractions 9 and 10 (F9/10) were identified, pooled and concentrated to 300 µl via vacuum centrifugation (SpeedVac, Savant). Aliquots were then frozen at -80 °C until required for further analysis.

Electrophoretic analysis

Samples were suspended in 1x Laemml sample buffer (with 10 mM TCEP, 6 M urea, and 2 % SDS), boiled for 10 min, and analyzed by SDS-PAGE (NuPage 4-12 % Bis-Tris, Invitrogen). Samples were prepared in parallel were either stained with Oriole (Bio-Rad) in preparation for mass spectrometry or immunoblotted using a 1:1000 dilution of polyclonal anti-horse...
spleen ferritin antibody produced in rabbit (Sigma-Aldrich), and imaged via standard chemiluminescence.

**Mass spectrometry**

Matrix-assisted laser desorption/ionisation- mass spectrometry (MALDI-MS) using an UltraflexXtreme (Bruker Daltronics), a two-layer sample preparation method and α-Cyano-4-hydroxycinnamic acid as the MALDI matrix, was used to determine the protein parent mass of the purified *C. elegans* F9/10 as ferritin. For further analysis the purified *C. elegans* ferritin was digested with LysC (Roche) in 4 M urea in a ratio of 50:1 protein to LysC overnight at 37 °C. Digested material was then desalted and concentrated through a C4 ZipTip (Millipore) for MALDI-MS/MS analysis (as above) to generate a peptide mass fingerprint. The peptide masses were searched using MASCOT (Matrix Science).

**DNA sequencing**

The *fmn-1* ORF amplicon was amplified using the following nested primers: Outer-forward 5'-ATGTGTCTCAGATTTCGCC, Inner-forward 5'-GGTTGAACCTTTTAGAAGTCGC, Inner-reverse 5'-ACAGTCCCGACACGTAATC and Outer-Reverse 5'-GAACCTTTGGTGCCATA. Sequencing was performed using the inner primer pair at the Applied Genetic Diagnostic facility (Department of Pathology, University of Melbourne) using ABI3130xl capillary genetic analyzers and BDV3.1 terminators. Three independent amplicons, from both wild type and *fmn-1(ok3625)*, were sequenced on both complementary DNA strands.

**Size exclusion chromatography-inductively-coupled plasma mass spectrometry (SEC-ICP-MS)**

Samples were homogenised in TBS (pH 8.0) with added proteinase inhibitors (EDTA-free, Roche), then clarified by a 15 min centrifugation at 175,000 g, 4 °C. The protein concentration of the supernatant was determined via UV absorbance (Nanodrop, ThermoScientific) and equivalent protein amounts were size-excluded using a Bio SEC-5 (4.6 x 300 mm, 5 μm, Agilent) column with 200mM ammonium nitrate (trace analysis grade, Sigma) pH 8.0 buffer the flow rate was 0.4 mL/min at 30 °C. The eluant from the column was directly connected to the ICP-MS for elemental detection as previously described.49,50

**Lifespan analysis**

The effects of genetic ablation of *fmn-2* and *fmn-1* on wild type was measured using established protocols.51,52 For the life span effects of elevated exogenous iron we used ferric ammonium citrate (FAC) supplemented NGM. Due to the semi defined chemical formulae of FAC, xFe₃y(NH₄)₆C₁₂H₁₄O₆, we determined the iron content of our batch of FAC (9.60 +/- 0.61 % by weight; n = 3), NGA (246.24 ng p g of media, n = 10) and the NGA + 5 mg mL⁻¹ FAC high iron media (1.172 mg g⁻¹ of media, n = 10) by ICP-MS. For the media measurements the media was aliquoted into ~500 mg aliquots, weighed and then freeze dried, digested in 50 μL 60 % (w/v) HNO₃ + 50 μL 30 % (w/v) H₂O₂ at 60 °C for 5 min, diluted to 1 mL and analysed by ICP-MS. All life span assays were conducted at 25 °C following adulthood.

**Spatially resolved fluorescence X-ray absorption near edge structure (XANES) spectroscopy.**

Hydrated, anesthetised *C. elegans* samples were prepared for XFM as previously described.16 A series of 100 XFM maps were collected using energies 7282 – 7042 eV (energy steps: 7282 – 7242 in 20 eV increments; 7222 – 7197 in 5 eV increments; 7192 – 7164 in 2 eV increments; 7162 – 7093 in 1 eV increments; 7092 – 7042 in 10 eV increments). To reduce the applied dose and limit radiation damage the beam was defocused to 3 μm and specimens scanned with a 7 μm vertical step. Data was binned every 7 μm in the horizontal to produce 7 x 7 μm pixel with an effective dwell time of ~27.34 ms per pixel.

As above, deconvolution of the Maia data was performed using GeoPIXE v7.1f (CSIRO, Australia). The X-ray photons from each pixel were related to calculated-model fluorescence X-ray yield for an assumed specimen composition and thickness as previously described.16 To minimise the effects of sample movement subsequent analysis was limited to the information rich iron K-edge (7100 – 7170 eV, consisting of 71 micrographs). The distribution of diffusible calcium remained unchanged throughout the data collection and was used to correct small positional discrepancies between micrographs. Cross correlation as implemented in MANTIS v2.06 was used to align the 71 iron micrographs into the final XANES stack. For each group the maximum pixel displacement through all series was < 1 pixel (Supplementary Fig 13). Pixels were grouped by spectral similarity using MANTIS v2.06 into distinct regions of interest. The associated XANES spectra for each region (Supplementary Fig 10) were normalised to the intensity of the white line and the K-edge structure analysed. Edge position was determined by fitting a single Gaussian to the first derivative spectra between 7114 – 7134 eV. The relative intensities of the 1s → 4s and 1s → 4p electronic transitions within the structure of the first derivative was used to indicate the Fe²⁺: Fe³⁺ ratio.29,30

**Statistical tests**

Kaplan–Meier survival curves were generated and compared via non-parametric Log rank tests and t-tests performed using Prism v5.0d (Graphpad Software). Comparisons between the goodness of fit for nested models, i.e. Gaussian (X ~ (μ,σ²)) vs. a sum of Gaussians (Σ^n i=1 Xi) were assessed using the Akaike information criterion and the exact sum of squares F-test. The dependency and co-variance between parameters of candidate models were monitored and models containing redundant parameters were rejected.
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Notes and references

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