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RESEARCH COMMUNICATION

Against the oxidative damage theory of aging: superoxide dismutases protect against oxidative stress but have little or no effect on life span in *Caenorhabditis elegans*

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The superoxide radical (O_2^-) has long been considered a major cause of aging. O_2^- in cytosolic, extracellular, and mitochondrial pools is detoxified by dedicated superoxide dismutase (SOD) isoforms. We tested the impact of each SOD isoform in *Caenorhabditis elegans* by manipulating its five *sod* genes and saw no major effects on life span. *sod* genes are not required for *daf-2* insulin/IGF-1 receptor mutant longevity. However, loss of the extracellular Cu/ZnSOD *sod-4* enhances *daf-2* longevity and constitutive diapause, suggesting a signaling role for *sod-4*. Overall, these findings imply that O_2^- is not a major determinant of aging in *C. elegans*.

Many forms of pathology lead to elevated levels of damage to biological macromolecules (Halliwell and Gutteridge 2007). This is also true of aging, the poorly understood biological process that leads to progressive deterioration and death. One strategy to discover the underlying mechanisms of aging has been to seek the causes of its associated molecular damage. An important early theory, proposed by Harman (1956), postulates that the cause might be oxygen free radicals. Harman later developed the theory, proposing a central role for the superoxide (O_2^-) radical, issuing from the mitochondrial electron transport chain (Harman 1972). During the last few decades, much effort has been invested in tests of this nexus of theories (for review, see Muller et al. 2007). Despite this, the importance of O_2^- as a cause of aging remains uncertain.

In this study, we take a genetic approach to critically test the role of O_2^- in aging in a short-lived animal, the nematode *Caenorhabditis elegans*, by manipulating expression of genes encoding the antioxidant enzyme superoxide dismutase (SOD). This enzyme catalyzes the dismutation reaction

$$2O_2^- + 2H^+ \rightarrow H_2O_2 + O_2.$$
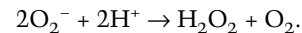
H_2O_2 (hydrogen peroxide) may then be broken down into H_2O and O_2 by catalase or glutathione peroxidase. O_2^- does not readily cross cellular membranes and, consequently, there are distinct extracellular, cytosolic, and mitochondrial O_2^- pools (Missirlis et al. 2003; Muller et al. 2004). In principle, one or more of these three O_2^- pools may play a role in aging.

Instrumental in lowering levels of O_2^- in each pool is a dedicated, compartment-specific SOD isoform. Cytosolic and extracellular O_2^- is consumed by distinct Cu/ZnSOD isoforms, while O_2^- in the mitochondrial matrix is consumed by MnSOD (Weisiger and Fridovich 1973). Most eukaryotes have a single SOD isoform for each compartment, but *C. elegans,* unusually, possesses two isoforms for each compartment. The two cytosolic Cu/ZnSOD isoforms are encoded by *sod-1* and *sod-5* (Larsen 1993; Giglio et al. 1994; Jensen and Culotta 2005). The *sod-4* gene encodes two predicted extracellular Cu/ZnSOD isoforms, products of alternative splicing of mRNA (Fujii et al. 1998). Two mitochondrial MnSOD isoforms are encoded by *sod-2* and *sod-3* (Giglio et al. 1994; Suzuki et al. 1996; Hunter et al. 1997).

This superabundance of SOD isoforms has been a technical hurdle to investigations of the role of SOD and O_2^- in aging in *C. elegans*, and some of the *sod* genes have been barely studied. In situ gel SOD activity assays of a *sod-1* deletion mutant imply that this gene encodes the major cytosolic Cu/ZnSOD (Jensen and Culotta 2005), leaving the function of *sod-5* unclear. *sod-3* mRNA levels are elevated in the dauer larva (Honda and Honda 1999), suggesting that this gene may play a special role in antioxidant defense in this long-lived, stress-resistant diapausal stage, but the role of *sod-2* has remained obscure. In this study, we describe in detail the function of each of the five *sod* genes, characterizing their expression, and the phenotypic effects of manipulating their expression. This has allowed us to assess the effect on life history, especially aging, of each of the three major O_2^- pools, thereby critically testing the role of SOD and, by inference, O_2^-, in longevity assurance and aging.

O_2^- can affect living organisms in a variety of ways. It can cause molecular damage that might contribute to aging; thus, one expectation of our study was that lowering SOD activity and increasing O_2^- levels might accelerate aging, and vice versa. H_2O_2 derived from O_2^- can also act a secondary messenger—for example, in receptor tyrosine kinase signaling pathways (Finkel 1998)—and as an activator of heat-shock factor. O_2^- can also be deployed as a chemical weapon in immune defense against bacterial pathogens in higher animals, and probably in *C.*

elegans as well (Chavez et al. 2007). There is even evidence that in *C. elegans* O_2^- can increase life span, perhaps by activating stress defense processes (Cypser and Johnson 2002; Schulz et al. 2007).

A powerful approach to investigate mechanisms of aging is the mutational analysis of genes with effects on life span. In *C. elegans*, mutations affecting the insulin/IGF-1 signaling (IIS) pathway can strikingly increase adult life span. For example, mutation of *daf-2*, which encodes an insulin/IGF-1 receptor, can increase adult life span by more than twofold (Kenyon 2005). Severe *daf-2* loss of function can also cause constitutive formation of dauer larvae, which are developmentally arrested, long-lived, diapausal third-stage larvae (Riddle and Albert 1997).

One possibility is that increased SOD levels and reduced damage from O_2^- contribute to the longevity of IIS mutants. *daf-2* mutants do show increased SOD and catalase activity levels, and resistance to oxidative stress (Vanfleteren 1993; Honda and Honda 1999). *sod-3* mRNA and protein levels are elevated in *daf-2* mutants (Honda and Honda 1999; Yanase et al. 2002; Dong et al. 2007), suggesting a possible role for mitochondrial MnSOD in longevity assurance. However, a more critical test is to examine the effects of alteration of SOD activity on life span.

In this study, we first examine the biology of the five *C. elegans sod* genes and show that *sod-1* and *sod-2* encode the major Cu/ZnSOD and MnSOD isoforms in reproductive development, while *sod-5* and *sod-3* encode minor, auxiliary isoforms mainly expressed in dauer larvae. We then critically test the importance of SOD and, by extension, O_2^- in *C. elegans* aging and in the *daf-2* Age phenotype by means of *sod* gene deletion and overexpression.

Results and Discussion

To understand the respective roles of the five *sod* genes, we characterized their expression using several techniques, including RT-PCR, Western blot analysis, SOD activity assays, and analysis of expression of *sod::gfp* transgenes. Expression was studied mainly in wild-type third-stage (L3) larvae and dauer larvae, *daf-2(m577)* mutants at the L3 stage, and mutants with deletions in each of the five *sod* genes (Supplemental Fig. S1A; for a detailed account of *sod* gene expression, see the Supplemental Material).

We report that *sod-1* and *sod-2* encode the major cytosolic Cu/ZnSOD and MnSOD isoforms, respectively. *sod-1* contributes ~80% of total SOD activity and is ubiquitously expressed (Supplemental Fig. S4C), and SOD-1 protein is localized to the cytosol and mitochondrial intermembrane space (Fig. 3A, below). By contrast, *sod-5* and *sod-3* are, respectively, minor cytosolic Cu/ZnSOD and MnSOD isoforms whose expression is upregulated in dauer larvae (Supplemental Figs. S1B–E, S4D,F). In wild-type L3s, *sod-5::gfp* expression is largely restricted to the ASI, ASK, and ASG amphid neurons (Supplemental Fig. S3B,C), which influence longevity and dauer larva formation (Bargmann and Horvitz 1991; Alcedo and Kenyon 2004).

It has been shown previously that in *daf-2* mutants,

there is an elevation in levels of SOD activity (Vanfleteren 1993) and large fold increase in *sod-3* mRNA (Honda and Honda 1999; Yanase et al. 2002). We confirmed this but also saw increases in expression of *sod-1* and *sod-5* (Supplemental Figs. S1B, S4B–D; Supplemental Material). Although *sod-3* is the most highly up-regulated of the *sod* genes in terms of fold change in expression, its relative contribution to SOD levels remains very small, both in terms of overall SOD activity and MnSOD protein levels (Supplemental Fig. S1B–E).

We examined the organismal effects of deletion alleles of *sod-2* and *sod-3* MnSOD genes on *C. elegans*. We find that *sod-2(0)* but not *sod-3(0)* results in delayed development (data not shown), delayed and reduced reproduction, and a slowed defecation cycle (Supplemental Fig. S5). However, *sod-3(0)* further reduces fertility of *sod-2(0)* animals (Supplemental Fig. S5B), implying functional redundancy between *sod-2* and *sod-3*.

As expected, loss of MnSOD increases sensitivity to oxidative stress. *sod-2(0)* causes a moderate reduction in resistance to the O_2^- generator paraquat, while *sod-3(0)* does not, either alone or when added to *sod-2(0)* (Fig. 1A). In combination, *sod-2(0)* and *sod-3(0)* cause severe hypersensitivity to hyperoxia (60% oxygen), while each mutation alone has no effect (Fig. 1B). Thus, *sod-2* and *sod-3* are functionally redundant when defending against mild but not severe oxidative stress.

Next, we tested the effect of *sod-2(0)* and *sod-3(0)* on life span. Mean life span is unaffected by either mutation alone (Fig. 1C; Supplemental Table S1) or both mutations combined (Fig. 1D; Supplemental Table S1). Thus, surprisingly, complete absence of MnSOD has no effect on life span. We also find that activity of cytosolic and
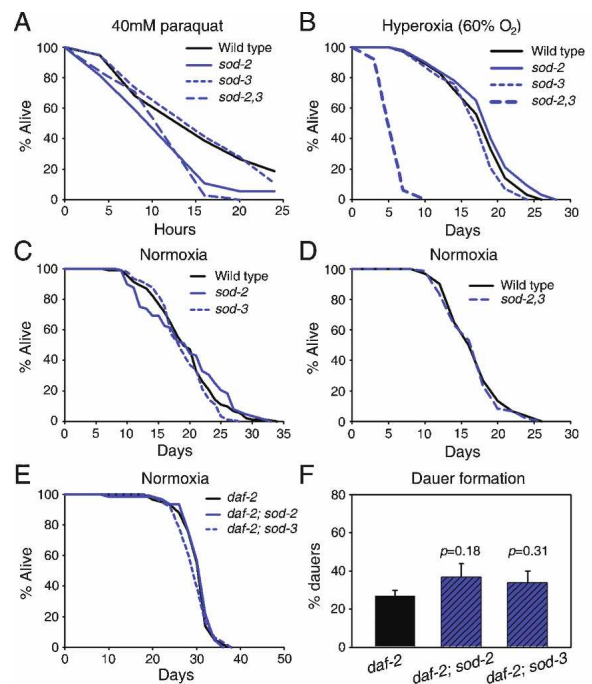

Figure 1. Oxidative stress resistance and longevity of MnSOD mutants. (*A*) Survival in 40 mM paraquat. (*B*) Survival under hyperoxia (60% O_2). (*C,D*) Life span of MnSOD single (*C*) and double (*D*) mutants under normoxia (20°C). (*E*) Effect of *sod-2(0)* or *sod-3(0)* on *daf-2(m577)* longevity (25°C). (*F*) Effect of *sod-2(0)* or *sod-3(0)* on *daf-2(m577)* Daf-c.

mitochondrial aconitase (an oxidation-sensitive iron–sulfur protein) is not detectably reduced in *sod-2; sod-3* animals (data not shown), suggesting no major increase in oxidative damage to protein. To test for a role in *daf-2* mutant longevity, we examined the impact of *sod-2(0)* or *sod-3(0)* on *daf-2* longevity but there is none (Fig. 1E; Supplemental Table S2). Altogether, these results strongly imply, against expectation, that mitochondrial matrix O_2^- has no effect on aging and that MnSOD does not contribute to longevity assurance in *C. elegans*.

Next, we describe the effects of deletion of *sod-1* and *sod-5* cytosolic Cu/ZnSOD genes. *sod-1(0)* increases sensitivity to paraquat, but *sod-5(0)* does not, either alone or when added to *sod-1(0)* (Fig. 2A). By contrast, neither *sod-1(0)* nor *sod-5(0)* has a marked effect on sensitivity to mild hyperoxia (Fig. 2B). *sod-1(0)* also decreases mean life span, by 15%–31%, while *sod-5(0)* has no effect, either alone or when added to *sod-1(0)* (Fig. 2C,D; Supplemental Table S1). Moreover, addition of *sod-2(0)* does not further reduce the life span of *sod-1* mutants (data not shown), supporting the view that O_2^- does not move between mitochondrial and cytosolic pools. Potentially, *sod-1(0)* shortens life span by accelerating the age increase in molecular damage. We therefore examined the effect of *sod-1(0)* on the age increase in damage to protein and lipid but could not detect any acceleration (Supplemental Material), perhaps because the impact of *sod-1(0)* is relatively subtle and difficult to detect.

If the shorter life span of *sod-1* mutants is due to accelerated aging, then overexpression of *sod-1* should increase life span. To test this, we first examined the effect of expression of the *sod-1::gfp* transgene on life span but

saw no effect (data not shown). However, we subsequently discovered that fusion of GFP to SOD-1 reduces Cu/ZnSOD-specific activity (Supplemental Fig. S7; Supplemental Material). Next, we generated transgenic lines with multiple copies of the *sod-1* gene, focusing initially on two lines bearing integrated transgene arrays, *wuIs152* and *wuIs154*. Overall SOD activity is increased approximately twofold by *wuIs152* (Fig. 3B) and *wuIs154* (data not shown), and both lines show increased Cu/ZnSOD immunoreactivity (Fig. 3A; data not shown).

Against expectation, *sod-1* overexpression increases sensitivity to paraquat (Fig. 3E). Potentially, this could result from elevated levels of H_2O_2 due to faster conversion of paraquat-generated O_2^- into H_2O_2. To test this, we generated an integrated transgene array, *wuIs151*, with multiple copies of the entire *ctl-1 ctl-2 ctl-3* gene cluster, which produces a 10-fold increase in catalase activity (Fig. 3C). Catalase overexpression suppresses the increased paraquat sensitivity resulting from *sod-1* overexpression (Fig. 3E), implying that this hypersensitivity is indeed due to elevated H_2O_2 levels.

wuIs152 caused a statistically significant ($P < 0.05$) increase in life span in five out of 11 trials, and in no instance did it decrease life span (Supplemental Table S3). Combined data for these 11 trials defines a 21.5% increase in mean life span (Fig. 3F, $P < 0.0001$). Although *wuIs154* did not increase life span, increased life span was seen in three further lines with extrachromosomal arrays (*wuEx125*, *wuEx122*, and *wuEx123*) (Supplemental Table S3). From this we conclude that elevated SOD-1 can slightly increase life span and that the absence of an effect in *wuIs154* likely reflects a life-shortening mutation associated with chromosomal insertion of the transgene array.

Given that SOD converts O_2^- into H_2O_2, it is possible that increased levels of SOD-1 lowers cytosolic O_2^- but increases cytosolic H_2O_2, replacing damage from O_2^- with damage from H_2O_2. To test this, we compared the effect on life span of elevated SOD-1, catalase, or both. Overexpression of catalase alone results in a high level of mortality due to internal hatching of larvae (bagging) (Fig. 3D), perhaps reflecting H_2O_2 deficiency. This bagging is suppressed by overexpression of *sod-1* (Fig. 3D), perhaps due to restoration of H_2O_2. In the absence of bagging (prevented by the inhibitor of DNA replication fluorodeoxyuridine, FUdR) overexpression of catalase slightly shortens life span, either alone or in addition to overexpression of *sod-1* (Fig. 3G; Supplemental Table S3). Taken together, these results imply that cytosolic Cu/ZnSOD and, by implication, cytosolic O_2^-, are weak determinants of longevity and aging, respectively. By contrast, H_2O_2 does not seem to contribute to aging.

We next investigated whether *sod-1* and *sod-5* might contribute to the *daf-2(m577)* longevity (Age) phenotype. *daf-2(m577)* is a weak, temperature-sensitive (ts) allele resulting in a moderate increase in life span at 20°C and a large increase at 25°C (Gems et al. 1998; Patel et al. 2008). *m577* was selected because it is a class 1 allele showing fewer pleiotropic effects than class 2 alleles such as *e1370*, making epistasis results easier to interpret. *sod-1(0)* slightly shortens *daf-2* life span at 25°C, perhaps reflecting a minor contribution of *sod-1* to *daf-2* Age, but not 20°C (Fig. 2E; Supplemental Table S2). *sod-5(0)* had little consistent effect (Supplemental Table S2).
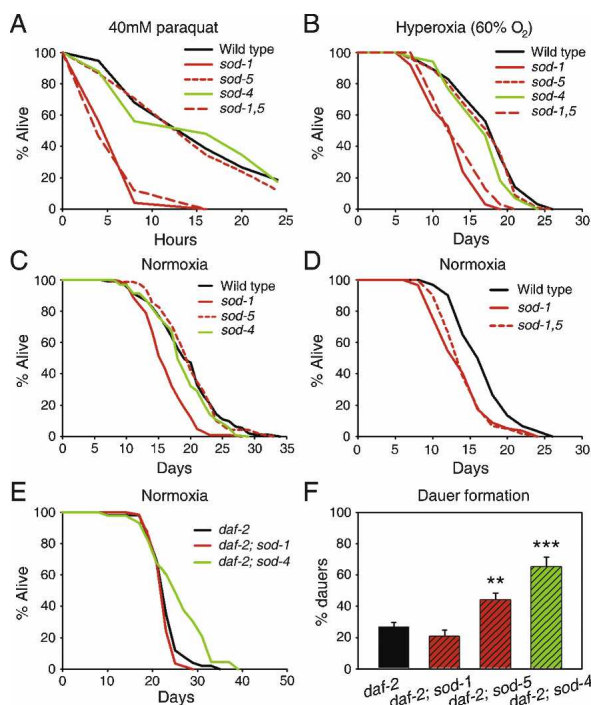

Figure 2. Oxidative stress resistance and longevity of Cu/ZnSOD mutants. (*A*) Survival in 40 mM paraquat. (*B*) Survival under hyperoxia (60% O_2). (*C,D*) Life span of Cu/ZnSOD single (*C*) and double (*D*) mutants under normoxia (20°C). (*E*) Effect of loss of Cu/ZnSOD on *daf-2(m577)* longevity (20°C). (*F*) Effect of loss of Cu/ZnSOD on *daf-2(m577)* Daf-c. (**) $P < 0.01$; (***) $P < 0.0001$, Student's *t*-tests.
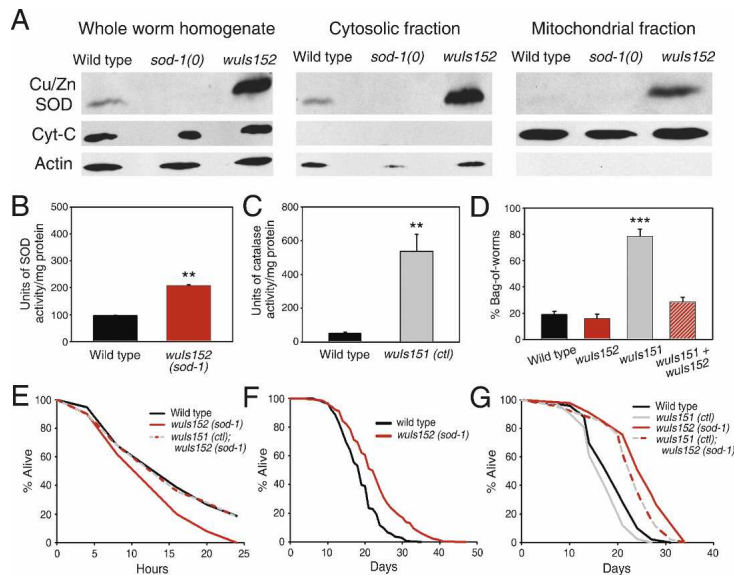
Figure 3. Overexpression of *sod-1* cytosolic Cu/ZnSOD increases life span. (*A*) Western blots of protein extracts of wild-type, *sod-1* mutant, and *sod-1* overexpresser lines using anti-Cu/ZnSOD antibodies. Cytochrome C (Cyt-C) and actin were used as mitochondrion- and cytosol-specific markers, respectively. Increased SOD-1 protein was also detected in *wuIs154* transgenics (data not shown). (*B*) Total SOD activity in protein extracts from lines overexpressing *sod-1*. (**) $P < 0.01$, Student's *t*-test. (*C*) Total catalase activity in line overexpressing catalase. (**) $P < 0.01$, Student's *t*-test. (*D*) Catalase overexpression causes high levels of mortality from internal hatching of larvae (bagging), and this is suppressed by overexpression of *sod-1*. (***) $P < 0.0001$, Student's *t*-test. (*E*) *sod-1* overexpression increases sensitivity to oxidative stress (40 mM paraquat), and this is suppressed by overexpression of catalase. (*F*) Overexpression of *sod-1* increases life span. (*G*) Elevated catalase does not further extend longevity of the *sod-1* overexpresser.

Finally, we examined the phenotypic effects of loss of the *sod-4* extracellular Cu/ZnSOD. *sod-4(0)* does not affect sensitivity to oxidative stress (Fig. 2A,B) or life span in otherwise wild-type animals (Fig. 2C). Surprisingly, *sod-4(0)* enhances *daf-2* Age at both 20°C and 25°C (Fig. 2E; Supplemental Table S2). One possibility is that this reflects an effect on signaling. SOD-4, like mammalian Cu/ZnSOD, might generate H_2O_2, which then crosses into the cell and promotes insulin signaling by inhibiting redox-sensitive, signal-quenching phosphatases (Goldstein et al. 2005). In *C. elegans*, treatment with H_2O_2 increases PIP_3 levels and promotes cytosolic retention of DAF-16 (Weinkove et al. 2006). If *sod-4(0)* does reduce IIS, then it should enhance other *daf-2* mutant traits, including constitutive dauer larva formation (Daf-c). We therefore tested the effect of loss of each *sod* gene on *daf-2(m577)* Daf-c and report that *sod-4(0)* significantly enhances Daf-c (Figs. 1F, 2F). Thus, *sod-4* may contribute to IIS.

O_2^- has long been viewed as a possible major determinant of aging (Harman 1956, 1972). In this study, we explored the importance of the three major O_2^- pools on aging in *C. elegans*. Overall, our results imply that O_2^- is not a major determinant of aging in this model organism, either in wild-type animals or long-lived *daf-2(m577)* mutants; however, it remains possible that SOD contributes substantially to longevity in other contexts (e.g., under dietary restriction). Our findings point to the novel conclusion that each O_2^- pool is different in terms of its effect on aging. In wild-type *C. elegans*, the cyto-

solic O_2^- pool contributes weakly to aging while mitochondrial O_2^-, long considered a likely determinant of aging, and extracellular O_2^- have no detectable effect on normal aging. In *daf-2* mutants, extracellular O_2^- appears to promote longevity; however, it is more likely that SOD-4 converts O_2^- into H_2O_2, which then weakly activates IIS, thereby shortening life span. The effects of SOD and O_2^- on aging therefore vary according to cell compartment and to genotype (summarized in Fig. 4).

Our findings paint a clearer picture of the role of the various SOD isoforms in *C. elegans*. *sod-1* and *sod-2* encode the major Cu/ZnSOD and MnSOD isoforms, corresponding to the equivalent isoforms in other eukaryotes. By contrast, *sod-5* and *sod-3* encode inducible, auxiliary Cu/ZnSOD and MnSOD isoforms. The presence of these supernumerary isoforms, like that of the stress-resistant dauer larva stage in which they are up-regulated, may reflect the hostile soil environment in which *C. elegans* has evolved.

We examined the effect on life span of loss of SOD in each of the three major cellular compartments. The oxidative damage theory of aging predicts that loss of SOD should cause accelerated aging, particularly cytosolic Cu/ZnSOD and MnSOD, both of which contribute to scavenging of mitochondrially generated O_2^- (the former in the mitochondrial intermembrane space). In fact, only loss of *sod-1* shortened life span, and then only modestly. However, overexpression of *sod-1* did increase life span slightly (Fig. 3F; Supplemental Table S3), implying a small contributory role of O_2^- to aging.

Loss of MnSOD isoforms had no effect on life span, either in a *daf-2(+)* or *daf-2(m577)* background (Fig. 1C–E; Supplemental Tables S1, S2). This strongly implies that O_2^- within the mitochondrial matrix is not a significant cause of aging in *C. elegans*. An alternative possibility is that other mechanisms protect *C. elegans* mitochondria against O_2^-; however, the oxygen hypersensitivity of *sod-2; sod-3* mutants argues against this.
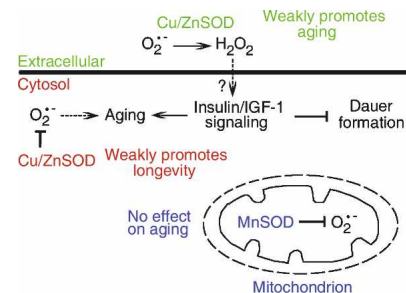

Figure 4. Influence of SOD and O_2^- on aging. This scheme shows a synthesis of conclusions drawn from the present study. Different SOD isoforms (and by deduction, the corresponding O_2^- pools) have different effects on aging. Extracellular Cu/ZnSOD weakly inhibits dauer formation and promotes aging, potentially by generating H_2O_2, which crosses into the cell and stimulates insulin/IGF-1 signaling by inhibiting redox-sensitive phosphatases. Cytosolic Cu/ZnSOD weakly promotes longevity, perhaps by protecting against molecular damage (*sod-1* does not influence *daf-2* Daf-c). Mitochondrial MnSOD has no detectable effect on aging. Arrow with dotted line implies a weak effect.

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Very recently, another study also reported, using different mutant alleles, that *sod-2(0); sod-3(0)* does not affect life span in an otherwise wild-type background (Honda et al. 2008), confirming our findings. These investigators also observed that in a *daf-2(e1370)* mutant, *sod-2(0)* slightly shortened life span and lessened Daf-c while *sod-3(0)* had the opposite effect. *daf-2* alleles fall into two phenotypic classes: Class 2 alleles are more pleiotropic than class 1 alleles and show more complex epistatic interactions with other mutations (Gems et al. 1998; Patel et al. 2008). The difference in effects of loss of MnSOD on *daf-2(m577)* (this study) and *daf-2(e1370)* (Honda et al. 2008) is interesting since *m577* is a class 1 allele and *e1370* a class 2 allele and suggests that MnSOD exerts a selective influence on class 2-specific defects.

Loss of function of many genes involved in mitochondrial function result in a Clk or Mit phenotype, which includes a delayed reproductive schedule, lowered fertility, slowed defecation cycle, and increased life span (Wong et al. 1995; Rea 2005). In this study, we found that *sod-2(0)* results in all these traits (Supplemental Fig. S5) except the increase in life span (Fig. 1C,D). RNAi of *sod-2* enhances *clk-1(qm30)* mutant longevity (Yang et al. 2007), suggesting a possible cryptic effect of *sod-2* on aging via a Clk/Mit-type mechanism.

Loss of *sod-4* enhances the *daf-2* Age and Daf-c phenotypes (Fig. 2E,F), suggesting that *sod-4* may play a role in IIS regulation of dauer formation and life span. We postulate that SOD-4-generated H_2O_2 promotes IIS by inhibiting IIS antagonistic phosphatase enzymes, as occurs in mammals (Goldstein et al. 2005).

While our findings imply that SOD and O_2^- have, at most, minor effects on aging in *C. elegans*, the question remains: What is the relevance of these findings to higher animals? The effects on life span of altering SOD gene expression vary between *C. elegans*, *Drosophila*, and the mouse (for review, see Muller et al. 2007). Loss of MnSOD causes early lethality in the fly and the mouse but not the worm, while loss of cytosolic Cu/ZnSOD causes only small decreases life span in the worm and the mouse but a large (~80%) decrease in life span in the fly. Overexpressing cytosolic Cu/ZnSOD slightly increases life span in *C. elegans* (this study), and in *Drosophila* overexpression of cytosolic Cu/ZnSOD and MnSOD seem to increase life span, although findings vary (Muller et al. 2007); however, overexpression of cytosolic Cu/ZnSOD does not increase life span in the mouse (Huang et al. 2000). By contrast, in the filamentous fungus *Podospora anserina*, reduction of mitochondrial ROS production results in a dramatic retardation of aging: Instead of aging rapidly after a brief period of growth, hyphae grow continuously and, apparently, indefinitely (Dufour et al. 2000).

It has been suggested that damage from reactive oxygen species might represent a public (i.e., evolutionarily conserved) rather than a private (i.e., lineage-specific) mechanism of aging (Martin et al. 1996). Our findings imply that the role of O_2^- in aging is to some extent public and to some extent private: Cytosolic O_2^- appears to contribute to aging in *C. elegans* and *Drosophila* but not mice. By contrast, loss of MnSOD is lethal to the fly and the mouse but has no effect on aging in the worm. Thus, O_2^- in some cellular compartments seems to contribute to some degree to aging in some species, but in other contexts (e.g., *C. elegans*) it appears unimportant to aging.

Materials and methods

Nematode culture

Nematodes were cultured on NGM agar seeded with *E. coli* OP50 as described previously (Sulston and Hodgkin 1988). Strains were maintained at 20°C unless otherwise noted. Mutant alleles and transgenic arrays used or generated in this study are listed in the Supplemental Material.

Oxidative stress resistance assays

For paraquat, young adults were placed overnight on agar plates containing 40 µM fluorodeoxyuridine (FUdR), and then picked into microtiter wells with 100 µl of 40 mM paraquat (Sigma, 856177) in M9 buffer. Viability was assayed over a 20-h period. For hyperoxia, young adults were picked onto plates containing 40 µM FUdR and placed in a sealed chamber under 60% O_2 (22°C). Animals were briefly removed from the O_2 chamber to score viability every 2–3 d.

Life span measurements

Life span assays were performed as described previously (Gems et al. 1998), at either 20°C or 25°C (see Supplemental Tables S1–S3). Survivorship of populations was compared statistically using the log rank test, performed using JMP 7.0.1 (SAS).

Construction of transgenic lines

Reporter constructs for each of the *sod* genes are full-length, translational fusions of GFP to the C terminus (see Supplemental Figure S2). For *sod* and *ctl* (catalase) overexpression, gDNA fragments were generated by PCR and microinjected directly. For primer sequences for reporter and overexpression constructs, see the Supplemental Material. Unless otherwise noted, pRF4 [*rol-6(su1006)*] was used as a marker of transformation. Integrated lines were generated by X-irradiation, and backcrossed to wild type (N2) at least five times before further study.

SOD and catalase activity assays

SOD activity was measured using an assay involving the inhibition of superoxide-induced lucigenin chemiluminescence by SOD, as described previously (Lenaerts et al. 2002). Catalase activity was assayed at 25°C according to a standard method (Aebi 1984), adapted for use in microtiter plate format.

Dauer formation assays

Eggs from strains bearing *daf-2(m577)* were collected by hypochlorite lysis and cultured at 23.5°C. The proportion of dauer larvae was scored 72 h later.

References

Aebi, H. 1984. Catalase in vitro. *Methods Enzymol.* **105:** 121–126.

Alcedo, J. and Kenyon, C. 2004. Regulation of *C. elegans* longevity by specific gustatory and olfactory neurons. *Neuron* **41:** 45–55.

Bargmann, C.I. and Horvitz, H.R. 1991. Control of larval development by chemosensory neurons in *Caenorhabditis elegans. Science* **251:** 1243–1246.

Chavez, V., Mohri-Shiomi, A., Maadani, A., Vega, L.A., and Garsin, D.A. 2007. Oxidative stress enzymes are required for DAF-16-mediated immunity due to generation of reactive oxygen species by *Caenorhabditis elegans. Genetics* **176:** 1567–1577.

Cypser, J.R. and Johnson, T.E. 2002. Multiple stressors in *Caenorhabditis*

elegans induce stress hormesis and extended longevity. *J. Gerontol.* **57:** B109–B114.

Dong, M.Q., Venable, J.D., Au, N., Xu, T., Park, S.K., Cociorva, D., Johnson, J.R., Dillin, A., and Yates III, J.R. 2007. Quantitative mass spectrometry identifies insulin signaling targets in *C. elegans. Science* **317:** 660–663.

Dufour, E., Boulay, J., Rincheval, V., and Sainsard-Chanet, A. 2000. A causal link between respiration and senescence in *Podospora anserina. Proc. Natl. Acad. Sci.* **97:** 4138–4143.

Finkel, T. 1998. Oxygen radicals and signaling. *Curr. Opin. Cell Biol.* **10:** 248–253.

Fujii, M., Ishii, N., Joguchi, A., Yasuda, K., and Ayusawa, D. 1998. Novel superoxide dismutase gene encoding membrane-bound and extracellular isoforms by alternative splicing in *Caenorhabditis elegans. DNA Res.* **5:** 25–30.

Gems, D., Sutton, A.J., Sundermeyer, M.L., Larson, P.L., Albert, P.S., King, K.V., Edgley, M., and Riddle, D.L. 1998. Two pleiotropic classes of *daf-2* mutation affect larval arrest, adult behavior, reproduction and longevity in *Caenorhabditis elegans. Genetics* **150:** 129–155.

Giglio, M.-P., Hunter, T., Bannister, J.V., Bannister, W.H., and Hunter, G.J. 1994. The manganese superoxide dismutase gene of *Caenorhabditis elegans. Biochem. Mol. Biol. Int.* **33:** 37–40.

Goldstein, B.J., Mahadev, K., and Wu, X. 2005. Redox paradox: Insulin action is facilitated by insulin-stimulated reactive oxygen species with multiple potential signaling targets. *Diabetes* **54:** 311–321.

Halliwell, B. and Gutteridge, J.M.C. 2007. *Free radicals in biology and medicine*. Oxford University Press, Oxford, UK.

Harman, D. 1956. Aging: A theory based on free radical and radiation chemistry. *J. Gerontol.* **11:** 298–300.

Harman, D. 1972. The biologic clock: The mitochondria? *J. Am. Geriatr. Soc.* **20:** 145–147.

Honda, Y. and Honda, S. 1999. The *daf-2* gene network for longevity regulates oxidative stress resistance and Mn-superoxide dismutase gene expression in *Caenorhabditis elegans. FASEB J.* **13:** 1385–1393.

Honda, Y., Tanaka, M., and Honda, S. 2008. Modulation of longevity and diapause by redox regulation mechanisms under the insulin-like signaling control in *Caenorhabditis elegans. Exp. Gerontol.* **43:** 520–529.

Huang, T., Carlson, E., Gillespie, A., Shi, Y., and Epstein, C. 2000. Ubiquitous overexpression of CuZn superoxide dismutase does not extend life span in mice. *J. Gerontol.* **55:** B5–B9.

Hunter, T., Bannister, W.H., and Hunter, G.J. 1997. Cloning, expression, and characterization of two manganese superoxide dismutases from *Caenorhabditis elegans. J. Biol. Chem.* **272:** 28652–28659.

Jensen, L.T. and Culotta, V.C. 2005. Activation of CuZn superoxide dismutases from *Caenorhabditis elegans* does not require the copper chaperone CCS. *J. Biol. Chem.* **280:** 41373–41379.

Kenyon, C. 2005. The plasticity of aging: Insights from long-lived mutants. *Cell* **120:** 449–460.

Larsen, P.L. 1993. Aging and resistance to oxidative stress in *Caenorhabditis elegans. Proc. Natl. Acad. Sci.* **90:** 8905–8909.

Lenaerts, I., Braeckman, B., Matthijssens, F., and Vanfleteren, J. 2002. A high-throughput microtiter plate assay for superoxide dismutase based on lucigenin chemiluminescence. *Anal. Biochem.* **311:** 90–92.

Martin, G.M., Austad, S.N., and Johnson, T.E. 1996. Genetic analysis of ageing: Role of oxidative damage and environmental stresses. *Nat. Genet.* **13:** 25–34.

Missirlis, F., Hu, J., Kirby, K., Hilliker, A., Rouault, T., and Phillips, J. 2003. Compartment-specific protection of iron-sulfur proteins by superoxide dismutase. *J. Biol. Chem.* **278:** 47365–47369.

Muller, F.L., Liu, Y., and Van Remmen, H. 2004. Complex III releases superoxide to both sides of the inner mitochondrial membrane. *J. Biol. Chem.* **279:** 49064–49073.

Muller, F.L., Lustgarten, M.S., Jang, Y., Richardson, A., and Van Remmen, H. 2007. Trends in oxidative aging theories. *Free Radic. Biol. Med.* **43:** 477–503.

Patel, D.S., Garza-Garcia, A., Nanji, M., McElwee, J.J., Ackerman, D., Driscoll, P.C., and Gems, D. 2008. Clustering of genetically defined allele classes in the *Caenorhabditis elegans* DAF-2 insulin/IGF-1 receptor. *Genetics* **178:** 931–946.

Rea, S.L. 2005. Metabolism in the *Caenorhabditis elegans* Mit mutants. *Exp. Gerontol.* **40:** 841–849.

Riddle, D.L. and Albert, P.S., eds. 1997. *Genetic and environmental regulation of dauer larva development*. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.

Schulz, T.J., Zarse, K., Voigt, A., Urban, N., Birringer, M., and Ristow, M. 2007. Glucose restriction extends *Caenorhabditis elegans* life span by inducing mitochondrial respiration and increasing oxidative stress. *Cell Metab.* **6:** 280–293.

Sulston, J. and Hodgkin, J. 1988. Methods. In *The Nematode* Caenorhabditis elegans (ed. W.B. Wood), pp. 587–606. Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY.

Suzuki, N., Inokuma, K., Yasuda, K., and Ishii, N. 1996. Cloning, sequencing and mapping of a manganese superoxide dismutase gene of the nematode *Caenorhabditis elegans. DNA Res.* **3:** 171–174.

Vanfleteren, J.R. 1993. Oxidative stress and ageing in *Caenorhabditis elegans. Biochem. J.* **292:** 605–608.

Weinkove, D., Halstead, J.R., Gems, D., and Divecha, N. 2006. Long-term starvation and ageing induce AGE-1/PI 3-kinase-dependent translocation of DAF-16/FOXO to the cytoplasm. *BMC Biol.* **4:** 1. doi: 10.1186/1741-7007-4-1.

Weisiger, R.A. and Fridovich, I. 1973. Superoxide dismutase. Organelle specificity. *J. Biol. Chem.* **248:** 3582–3592.

Wong, A.E., Boutis, P., and Hekimi, S. 1995. Mutations in the *clk-1* gene of *Caenorhabditis elegans* affect developmental and behavioral timing. *Genetics* **139:** 1247–1259.

Yanase, S., Yasuda, K., and Ishii, N. 2002. Adaptive responses to oxidative damage in three mutants of *Caenorhabditis elegans* (*age-1, mev-1* and *daf-16*) that affect life span. *Mech. Ageing Dev.* **123:** 1579–1587.

Yang, W., Li, J., and Hekimi, S. 2007. A measurable increase in oxidative damage due to reduction in superoxide detoxification fails to shorten the life span of long-lived mitochondrial mutants of *Caenorhabditis elegans. Genetics* **177:** 2063–2074.