The Glutathione Reductase GSR-1 Determines Stress Tolerance and Longevity in *Caenorhabditis elegans*

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Abstract

Glutathione (GSH) and GSH-dependent enzymes play a key role in cellular detoxification processes that enable organisms to cope with various internal and environmental stressors. However, it is often not clear, which components of the complex GSH-metabolism are required for tolerance towards a certain stressor. To address this question, a small scale RNAi-screen was carried out in *Caenorhabditis elegans* where GSH-related genes were systematically knocked down and worms were subsequently analysed for their survival rate under sub-lethal concentrations of arsenite and the redox cycler juglone. While the knockdown of γ-glutamylcysteine synthetase led to a diminished survival rate under arsenite stress conditions, GSR-1 (glutathione reductase) was shown to be essential for survival under juglone stress conditions. gsr-1 is the sole GSR encoding gene found in *C. elegans*. Knockdown of GSR-1 hardly affected total glutathione levels nor reduced glutathione/glutathione disulphide (GSH/GSSG) ratio under normal laboratory conditions. Nevertheless, when GSSG recycling was impaired by gsr-1(RNAi), GSH synthesis was induced, but not vice versa. Moreover, the impact of GSSG recycling was potentiated under oxidative stress conditions, explaining the enormous effect gsr-1(RNAi) knockdown had on juglone tolerance. Accordingly, overexpression of GSR-1 was capable of increasing stress tolerance. Furthermore, expression levels of SKN-1-regulated GSR-1 also affected life span of *C. elegans*, emphasising the crucial role the GSH redox state plays in both processes.

Introduction

Aerobic life uses the oxidizing power of O2 for various fundamental biochemical processes including energy generation in form of ATP synthesis realised by a controlled stepwise reduction of molecular oxygen. However, oxygen consumption has severe side effects, since, inevitably, reactive oxygen species (ROS) are produced as by-products. If not countered or buffered, these ROS attack and disturb biological molecules and cellular structures leading to a state of impaired physiology also known as oxidative stress. Hence, acrob organisms have evolved a variety of mechanisms such as antioxidants, detoxification enzymes and repair systems to counteract the deleterious effects of ROS [1,2]. ROS production and/or intake are further enhanced under pathological conditions with inflammatory implications (e.g. rheumatoid arthritis, diabetes, cancer or neurodegenerative diseases) or by environmental stressors [3,4]. Moreover, according to the free radical theory of aging, oxidative stress-mediated accumulation of damaged biological molecules favours the aging process and shortens the life span of organisms [5,6]. In particular, endogenous superoxide radicals derived from mitochondrial respiration have been implicated to be a major cause for aging. In good agreement with that, resistance to oxidative stress was often found to correlate with longevity in different metazoan organisms [7,8], although some concerns over this linkage have recently been raised [9,10].

In animal cells, the tripeptide glutathione (γ-glutamylcysteinyl-glycine, GSH) represents the predominant low molecular weight thiol [11]. Under normal physiological conditions most of the redox-active GSH molecules are reduced and only a minor fraction of the tripeptide is present as glutathione disulphide (GSSG). Accordingly, the GSH/GSSG couple represents a major cellular redox buffer that significantly contributes to the maintenance of the reduced intracellular milieu and, hence, to the antioxidative capacity of cells. GSSG, formed when GSH serves as a biological reductant, has to be recycled by the NADPH-dependent GSSG reductase (GSR, EC 1.6.4.2). In addition, intracellular GSH homeostasis is regulated by a synthesis pathway consisting of a two-step reaction catalysed by γ-glutamylcysteine synthetase (GGS) and GSH synthetase (GSS). Furthermore, exogenous GSH was reported to represent an important source to replenish the intracellular GSH pool, however, only after extracellular breakdown and intracellular re-synthesis of the tripeptide via the γ-glutamyl cycle, where the γ-glutamyl transferase (GGT) catalyses the first and rate limiting step. Being a coenzyme or a substrate for diverse enzymes such as glutathione peroxidases (GPX), glutathione S-transferases (GST) and glutaredoxins (GLRX), GSH...
functions as a central player in redox regulation, ROS defence and phase II detoxification [11,12].

The nematode Caenorhabditis elegans is an established model organism in research on stress defence and aging offering distinct advantages [9,13,14,15]. The worm can be easily cultured on agar plates, reproduces with a rapid life cycle of approximately 3.5 days and has a maximum life span of only about 30 days. C. elegans is genetically tractable by RNA interference or germ-line transformation via microinjection allowing the assessment of gene function and related phenotypes at the organismic level. Moreover, in silico analysis of the approximately 19,000 genes revealed that central pathways related to stress defence and aging including putative homologues of the GSH metabolism genes that are well conserved among metazoa are also found in the worm. Consistent with that it has been demonstrated in many studies that the transcription factors C. elegans DAF-16 (homologous to the mammalian forkhead box family FOXO) and C. elegans SKN-1 (homologous to the mammalian NF-E2 related factor Nrf-2) have a central position in stress resistance and life span determination in metazoan organisms from C. elegans up to mammals, being controlled among others by the insulin-like/IGF and p38 MAP-kinase pathways ([14,16,17], DAF-16 and SKN-1 are responsible for the induction of numerous stress response genes including superoxide dismutases, catalases and GSH-related genes such as gse and gst.

The highly reactive metalloid arsenite causes oxidative stress by binding directly to thiols such as GSH or by the formation of intracellular ROS [18]. It is well known that the susceptibility of cells towards arsenite correlates with their intracellular GSH level and that treatment with the GCS-inhibitor buthionine sulfoximine diminishes resistance [19]. Biotransformation of inorganic arsenic can either involve repetitive reduction and oxidative methylation, the direct methylation of arsenic-glutathione complexes or direct complexation of trivalent arsenic with GSH. These arsenic-GSH complexes are possibly involved in the efflux of arsenicals form cells by ATP-binding cassette membrane transporters. Furthermore, arsenite and the complexes with arsenite are inhibitors of the GSR, creating a negative feedback loop involving reduction and complexation of trivalent arsenic by GSH and modulation of GSR activity [20,21].

Black walnut toxicity is due to the allelopathic naphthoquinone Juglone, that is being used in natural dyeing, as herbicide, dietary supplement and in various folk medicines to treat bacterial, fungal or viral infections of the skin. Its toxicity, and probably also its antibiotic, antiviral, and antifungal properties are due to the intracellular ROS production. Juglone is able to cross cell membranes and generate superoxide anion from molecular oxygen during metabolism [22,23,24]. Juglone response in C. elegans has frequently been analysed by investigating alterations in transcript levels and reporter gene expression. Here a number of genes/pathways that are affected under juglone exposure were revealed, including genes of the GSH metabolism [23,25,26,27,28,29]. However, the induced or decreased expression of a particular gene or even a protein is not the ultimate evidence of its essential role in stress tolerance of an organism.

In the present study we have employed a small scale RNAi screen in C. elegans to address the question which components of the GSH-metabolism are required for the tolerance towards sub-lethal intracellular oxidative stress generated by arsenite and the redox cycler juglone. Here, it is important to note that knockdown by RNAi can be variable and can lead to false negative results. Since only the induction of a detectable phenotype is a reliable indicator of a positive RNAi result, only these enzymes will be considered. Furthermore, our experimental setup cannot exclude combined effects of several enzymes.

Whereas tolerance towards arsenite was shown to be GSH synthesis-dependent, the gene C46F11.2 encoding GSR-1, a protein orthologous to human mitochondrial glutathione reductase, was identified to be absolutely essential for survival under sub-lethal juglone stress. Our studies revealed that expression levels of SKN-1-regulated GSR-1 determine not only the stress tolerance (primarily against endogenous oxidative stress) but also the life span of C. elegans, emphasising the crucial role the GSH redox state plays in both processes.

Materials and Methods

Oligonucleotides

Oligonucleotide sequences are listed in the Table 1.

C. elegans Culturing and Strains

C. elegans were cultured at 20°C under standard conditions on nematode growth medium (NGM) seeded with Escherichia coli OP50 (Caenorhabditis Genetics Center) as a food source [30] unless otherwise noted. Worm populations were synchronized by alkaline hypochlorite lysis [40% sodium hypochlorite, 5% 10 N NaOH] [31]. The following strains were obtained from the NIH National Center for Research Resources: Wild-type N2 Bristol, GE23 pha-1[e2123III], TJ356 zls356 IV [DAF-16::GFP], CL2166 dvl19[pAF15(gst-6::GFP::NLS)III], zcs14[myo-3::GFP[mit]]. The strains N2 Ex003[gor-1::GFP] and N2 Is007[SKN-1::GFP] [32,33] were kindly provided by Keith Blackwell (Harvard Medical School, Boston). The balanced mutant gsr-1 strain VB2729 (gor-1[m3574]/hT2[bl-4(e937) let-7b(qf182) qIs48]) was kindly provided by Simon Tuck (Umeå Center for Molecular Medicine, Umeå University).

RNAi Assays

RNAi was induced by feeding worms with E. coli HT115 strains producing dsRNA following standard protocols [34]. The bacterial RNAi clones were grown in Luria-Bertani (LB) medium containing 50 μg ml⁻¹ ampicillin and 12.5 μg ml⁻¹ tetracyclin for 16 h at 37°C, before being spread on fresh NGM agar plates supplemented with 50 μg ml⁻¹ ampicillin and 2.5 mM isopropyl β-D-thiogalactoside (IPTG). Worms were transferred to RNAi plates either at L4 stage and F1 L4/young adults were analysed or as eggs/L1 and the adult worms of the same generation were inspected. Greatest care was applied to avoid contaminating bacteria and/or fungi during feeding of worms. Furthermore, to assess the efficiency of the feeding RNAi method, we used dpy-5 and bli-1 as target genes. RNAi resulted in either a dumpy or blistered phenotype.

Identification of GSH Metabolism Genes and RNAi Screen for Juglone Tolerance

Genes that are potentially involved in the GSH-metabolism of C. elegans were identified by combining database mining of annotated genes and Blast searches of WormBase using known genes of GSH-metabolism from other organisms. In total, 67 genes were identified that encode proteins of GSH-synthesis (gor-1, gsr-1) and GSH-redox cycle (gpx-1 to gpx-6, gsr-1), GST (ggt-1 to ggt-44), GGT (gtt-1 to gtt-6) and GLRX (glrx-5, glrx-10, glrx-21). Most of the E. coli HT115 strains that produce the corresponding dsRNA were found in the RNAi library (Geneservice, Cambridge, UK). For genes that were not present in the library, open reading frames were amplified from C. elegans CDNA by polymerase chain reaction (PCR) using the oligonucleotides listed in Table 1. PCR products were cloned into pL4400 using standard methods. Positive clones
validated by DNA-sequencing, were transformed into *E. coli* HT115 cells. Stress tolerance assays with RNAi treated animals were performed as follows: In pre-tests we determined a 90% survival rate of wildtype worms at a juglone concentration of 0.15 mM for an incubation time of 18 h. Using a NaAsO₂ concentration of 3 mM ensured 80% survival of wildtype worms after 18 h incubation. RNAi assays were carried out as described in [25]. F₁ young adults were transferred to RNAi-plates supplemented with 0.15 mM juglone. Animals were cultured for additional 18 h at 20°C. The percentage of survivors was determined.

**Nucleic Acid Preparation and Northern Blot Analysis**

Synchronised *C. elegans* N2 wild-type worms were cultured on NGM agar under standard conditions until they reached the L4/young adult stage. Animals were allocated and cultured on standard control plates or in the presence of 10 mM juglone for 3 h before being harvested. Total RNA was prepared using TRIzol extraction according to the manufacturer’s instructions (Invitrogen, Carlsbad, CA, USA). For Northern blotting, total RNA was separated on an agarose formaldehyde gel and transferred to a positively charged nylon membrane (Millipore Co., Bedford, MA, USA). The membrane was hybridized with a radio-labelled probe of the *gsr-1* coding region (amplified from *C. elegans* cDNA by using the primer pair CeGR-5306-S and GGR-5307-AS, Table 1) in 50% formamide, 5x standard saline citrate (SSC), 5x Denhardt’s solution, and yeast tRNA at 55°C overnight, followed by washing with 2x SSC and 0.1% sodium dodecyl sulfate (SDS) at 60°C.

**Table 1. Oligonucleotide sequences.**

| Primer name | Sequence 5’ to 3’ | Restriction sites |
|-------------|-------------------|------------------|
| Primer for GSR cloning: | | |
| CeGR-EX-S | CATATGCTGGCGTCAAGGAGTTTCG | NdeI |
| CeGR-EX-AS | CTACGGTTATCCGCTGCTACACCTCC | XhoI |
| CeGR-3165-S | CTGGCGCCATCTTGGGACTGAGCTTCCG | PstI |
| CeGR-3166-AS | GAGTCCTTATGCTGGGACTGAGCTTCCG | BamHI |
| CeGR-5306-S | ATGTCCTGAGTTATTCCGGCTGACCTTGGACAC | – |
| CeGR-5307-AS | TTATCCGGCTACACCTCCGCA | – |
| CeGR-5326-AS | CTCGGGCTCTGGGCTCACCTCC | SmaI |

| Primer for RNAi clones: | | |
| GST-11-S | CTGCAGAGCCGGCAAGGTGTTTCTTCC | PstI |
| GST-11-AS | CTACGGTTATCCGCTGCTACACCTCC | XhoI |
| GST-17-S | CTGGCGCCATCTTGGGACTGAGCTTCCG | PstI |
| GST-17-AS | CTGGCGCCATCTTGGGACTGAGCTTCCG | PstI |
| GST-35-S | CTACGGTTATCCGCTGCTACACCTCC | XhoI |
| GST-35-AS | CTACGGTTATCCGCTGCTACACCTCC | XhoI |
| GST-43-S | CTGCAGTACACTGTAAGCTGCTAGTG | PstI |
| GST-43-AS | CTGCAGTACACTGTAAGCTGCTAGTG | XhoI |
| GSTD-1-S | CCAAGCTTATGGTATTAAAAGCAGTGATAC | HindIII |
| GSTD-1-AS | CCAAGCTTATGGTATTAAAAGCAGTGATAC | XhoI |
| GLRX-3-S | TGCGCCGAGCTCTGAGGACCTGACAAAAGCAGTGCTAGG | XbaI |
| GLRX-3-AS | TGCGCCGAGCTCTGAGGACCTGACAAAAGCAGTGCTAGG | XbaI |
| GLRX-22-S | GCCGCGCGCCGAGCTCTGAGGACCTGACAAAAGCAGTGCTAGG | XbaI |
| GLRX-22-AS | GCCGCGCGCCGAGCTCTGAGGACCTGACAAAAGCAGTGCTAGG | XbaI |

Underlined nucleotides correspond to restriction sites.

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**Plasmid Constructs and Transgenic *C. elegans***

**Pgsr-1::GFP.** A 2.4 kb fragment including 1996 bp of the 5’ upstream region, the first exon, the first intron and part of exon II of *gsr-1* was amplified from the *C. elegans* genomic DNA using Expand Long Template PCR system (Roche, Mannheim, Germany) and the primers CeGR-5165-S and CeGR-5166-AS (Table 1). The PCR product was cloned into pPD95.77 provided by A. Fire (Carnegie Institute, Baltimore, USA).

**Pgsr-1::gsr-1::GFP.** A 3864 kb genomic fragment that includes 1996 bp of the 5’ upstream region and the complete open reading frame of *C. elegans* *gsr-1* was amplified by PCR using the primers CeGR-5165-S and CeGR-5326-AS (Table 1). The PCR product was cloned into pPD95.77.

Germline transformation with the constructs *Pgsr-1::GFP* and *Pgsr-1::gsr-1::GFP* were performed by co-injecting the vector constructs (20–50 μg ml⁻¹) with the pRF4 plasmid (80 μg ml⁻¹) encoding the dominant marker gene rol-6 [35]. The *rol-6* construct was used for selection of transgenic animals. Only transformed progeny survive and can be easily maintained by cultivation at 25°C. To investigate the cell-specific, developmentally regulated transcription of *gsr-1*, GFP expression patterns were analysed by fluorescence microscopy.
Analyses of gsr-1 Promoter Activity under Stress Conditions

To analyze alterations in gsr-1 promoter activity, worms carrying the P_{gsr-1::GFP} reporter construct were allowed to grow from L1 to the L4/young adult stage in the presence of skn-1(2)(RNAi), daf-16(RNAi) or control RNAi bacteria before being transferred to corresponding (i) standard RNAi plates, (ii) RNAi plates supplemented with 150 μM juglone or (iii) starvation plates without bacteria. After 16 h incubation worms were inspected for GFP expression intensities.

GFP Reporter Evaluation

Expression of the intestinal promoter reporters P_{gsr-1::GFP} and P_{gsr-4::GFP} as well as of the intestinal nuclear translocation reporters DAF-16::GFP and SKN-1::GFP was analysed and scored following the method described in Tullet et al. [16]. P values were determined from a Fisher test.

Stress Tolerance Assays

For stress tolerance assays synchronised N2 wildtype L1 larvae were fed with gsr-1(RNAi), gsr-1(RNAi) or pL4440 control bacteria under standard RNAi conditions until reaching L4. Worms were then transferred to RNAi plates supplemented with 3 mM cumene hydroperoxide, 0.15 mM juglone, 50 mM paraquat, 7.5 mM tert-butylhydroperoxide or 3 mM NaAsO_2. Similarly, L4 worms overexpressing GSR-1 in the pha-1(e2123) background and promoter::GFP control worms were transferred to N2 NGM plates containing 0.25 mM juglone. Animals were cultured for another 16 h, before their survival rate was evaluated. A worm was scored as dead when it did not respond to a mechanical stimulus. Each experiment was performed at least three times, and the data were analysed using GraphPad Prism software.

Determination of Life Span and Brood Size

Phenotype analyses of gsr-1(RNAi) and pL4440(RNAi) control worms were performed at 20°C. Transgenic worms that carry the thermo-selectable pBX in the pha-1(e2123) genetic background were maintained at 25°C. To determine brood sizes, worms were cultured individually on NGM plates. Adults were transferred daily to new plates until egg production ceased. Plates with progeny were retained and counted when larvae reached L3/L4. For life span assays, synchronised worms were incubated at 20°C until they reached adulthood. 100 fluorescent P_{gsr-1::GFP} and P_{gsr-4::GFP} worms were transferred on 10 NGM plates and were scored as dead, when they no longer responded to touch-provoked movement. Animals that crawled away from plates or died from internal itching of progeny were replaced by worms from a parallel substitutes’ plate. Survival plots were compared using the log-rank test (GraphPad Prism software).

Cloning, Expression, Purification and Enzymatic Activity Assay of the Recombinant GSR-1

The coding region of GSR-1 was amplified from C. elegans cDNA by PCR using the oligonucleotides GGR-EXS and GGR-EXAS (Table 1). The PCR product was cloned into the prokaryotic expression plasmid pJG40 [36] and transformed into the E. coli strain BL21[DE3] (Novagen, Madison, WI, USA). After expression, recombinant GSR-1 was purified from the supernatant by chelating chromatography on Ni^{2+}-nitrilotriacetate (Ni-NTA) agarose according to the manufacturer’s instructions (Qiagen, Hilden, Germany). Subsequently, GSR-1 was subjected to fast protein liquid chromatography on a Superdex 75 column (Amersham-Pharmacia Biotech, Piscataway, NJ, USA). The protein concentration was determined by the method of Bradford [37] and the homogeneity of the enzyme preparation was analysed with 12.5% SDS-PAGE. Proteins were revealed by Coomassie Blue staining or Western blot analysis.

The activity of the GSR-1 was determined spectrophotometrically at 25°C (UVikon-932 spectrophotometer, Goebel, Germany) as previously described by Muller et al. [38]. The decrease of absorbance at 340 nm due to the oxidation of NADPH was recorded. Background rates were determined in the absence of GSSG. The activity that catalyzes the consumption of 1 μmol of substrate per minute was defined as 1 U. To determine the steady-state kinetic parameters, the enzyme assay was carried by varying the concentration of one substrate (10–500 μM for NADPH and 10–200 μM for GSSG) in the presence of saturating concentration of the other.

Glutathione Determination

Synchronised three-days-old young adult worms were cultured under RNAi conditions on plates containing (i) control HT115 carrying empty pL4440 vector and (ii) plus 150 μM juglone or (iii) gsr-1(RNAi) feeding bacteria and (iv) plus 150 μM juglone. After 4 h incubation, worms were harvested and the GSH redox states and γ-glutamylcysteine level were determined by high-performance liquid chromatography (Spectrasystem P2000 pumps, Thermo Fisher) and fluorometric detection (Jasco 821-FP, Jasco) according to Neuschwander-Tetri & Roll [39]. Briefly, worms were homogenized with a glass-embedded Potter-Elvehjem tissue grinder in 100 mM Tris-HCl, pH 8.5 containing 10 mM EDTA. For the determination of GSSG, worms were homogenized in the presence of 2 mM N-ethylmaleimide to eliminate free thioldiols, before the addition of DTT. Proteins were precipitated by 2 N perchloric acid. Thios were derivatized for two minutes with orthophthalaldehyde (OPA-reagent) in 0.4 M potassium borate, pH 9.9 before samples were neutralized by the addition of 100 mM sodium phosphate, pH 7.0. Separation of OPA-labeled thios was performed isocratically on a Nucleosil 120-5 C18 column (250×3 mm; Machery-Nagel) at a flow rate of 0.55 ml min^{-1} with 7.5% methanol, 92.5% 150 mM sodium acetate buffer, pH 7.0. Peaks were detected at 420 nm after excitation at 340 nm.

Results

GSR-1 is Required for Tolerance Towards the Pro-oxidant Juglone

A survey based on homology searches and already confirmed gene functions revealed 67 genes that are directly involved in the GSH metabolism of C. elegans (Table 2). Microarray analyses have demonstrated that the transcription level of numerous of these genes are altered under ROS and other stress conditions [40]. However, changes in mRNA levels do not indicate whether a gene is required for organismic tolerance towards a stressor. To address this question, we carried out a small scale RNAi screen where C. elegans was exposed to sub-lethal concentrations of the pro-oxidant juglone. Among the 67 GSH-metabolism genes tested only C46F11.2 was found to be absolutely essential for juglone tolerance (juglone survival rate 3.3±2.3, p<0.001, Table 2; a more moderate effect was observed with the previously characterised omega-class gst, GSTO1 (juglone survival rate 16.4±1.7, p<0.01) [25].
### Table 2. Small scale RNAi screen testing juglone and arsenite tolerance.

| Gene      | Annotation                    | RNAi clone | Juglone survival | Arsenite survival |
|-----------|-------------------------------|------------|------------------|-------------------|
| Control   | –                             | L4440      | 90.8±7.4         | 90.0±5.8          |
| GST       | Glutathione synthetase (GSS)  | F37B12.2   | 67.8±1.9         | 0***              |
| Glutathione reductase (GSR) |                     | M176.2     | 85.6±13.5        | 76.7±14.1         |
| Glutathione S-transferases (GST) |                     | C46F11.2*  | 3.3±7.3***       | 87.8±8.4          |
| R107.7    | gsr-1                         | II-4J12    | 94.7±5.1         | 76.7±4.7          |
| K08F4.11  | gsr-2                         | IV-3I02    | 93.3±2.7         | 74.5±7.8          |
| K08F4.10  | gsr-3                         | IV-5I10    | 91.7±1.9         | 81.5±2.1          |
| K08F4.4   | gsr-4                         | IV-5I04    | 94.2±5.7         | 78.3±11.8         |
| R03D7.6*  | gsr-5                         | II-7H21    | 94.2±5.7         | 63.3±26.5         |
| F11G11.2  | gsr-6                         | II-4E11    | 82.5±11.0        | 76.7±14.5         |
| F11G11.1  | gsr-7                         | II-4E09    | 88.3±8.4         | 58.9±15.8         |
| R05F9.5   | gsr-8                         | II-4I01    | 83.3±7.2         | 64.4±16.8         |
| Y45G12.2  | gsr-9                         | V-2E10     | 89.2±12.6        | 93.3±9.4          |
| R11G13.1  | gsr-10                        | cloned     | 88.1±10.3        | n.d.              |
| F37B1.2   | gsr-11                        | II-8F10    | 85.0±10.3        | 56.2±22.9         |
| T26C5.1   | gsr-12                        | II-6J13    | 89.2±9.6         | 91.7±7.1          |
| F37B1.3   | gsr-13                        | II-8F12    | 83.3±8.2         | 67.8±10.2         |
| F37B1.4   | gsr-14                        | II-8F14    | 81.7±10.0        | 90.0±1.5          |
| F37B1.5   | gsr-15                        | II-8F16    | 90.0±7.2         | 78.7±7.1          |
| F37B1.6   | gsr-16                        | cloned     | 96.4±3.6         | n.d.              |
| F37B1.7   | gsr-17                        | II-8F18    | 89.2±9.8         | 75.0±7.1          |
| F37B1.8   | gsr-18                        | II-8F20    | 85.8±13.2        | 78.3±21.2         |
| ZK697.6*  | gsr-19                        | V-1J14     | 77.5±14.8        | 66.7±17.6         |
| F21H7.1   | gsr-20                        | V-10H12    | 83.3±12.5        | 85.0±11.8         |
| T28A11.11 | gsr-21                        | V-2N04     | 92.5±6.9         | 83.3±4.7          |
| F37F2.3   | gsr-22                        | I-1O23     | 86.7±4.7         | 96.7±0.0          |
| Y53F4.29  | gsr-23                        | II-9O02    | 91.1±6.9         | 73.3±23.6         |
| Y53F4.30  | gsr-24                        | II-9O04    | 87.8±1.9         | 90.0±9.4          |
| Y53F4.31  | gsr-25                        | II-9O06    | 92.2±7.7         | 91.7±2.4          |
| Y53F4.32  | gsr-26                        | II-9O08    | 91.1±5.1         | 47.0±26.7         |
| Y53F4.35  | gsr-27                        | II-9O14    | 91.1±6.9         | 71.7±16.5         |
| Y53F4.37  | gsr-28                        | II-9O18    | 87.8±6.9         | 24.0±13.2         |
| Y1H11.1   | gsr-29                        | V-10H12    | 85.6±8.4         | 63.3±2.3          |
| Y1H11.2   | gsr-30                        | cloned     | 94.6±2.1         | n.d.              |
| R07B1.4   | gsr-31                        | X-4J08     | 87.8±5.1         | 77.3±10.4         |
| Y32G9A.1  | gsr-32                        | V-13E20    | 92.2±3.8         | 73.3±1.0          |
| F35E8.8   | gsr-33                        | V-10F05    | 91.1±6.9         | 40.0±17.6         |
| Y53F4.33  | gsr-34                        | II-9O10    | 91.9±8.8         | 88.3±7.1          |
| F56B3.10  | gsr-35                        | IV-1E20    | 88.6±11.7        | 63.3±23.6         |
| R13D7.7   | gsr-36                        | V-5I06     | 89.8±3.7         | 78.3±2.4          |
| D1053.1   | gsr-37                        | X-5J07     | 91.1±9.6         | 60.8±12.4         |
| Y71F9A.5  | gsr-38                        | cloned     | 95.5±1.1         | n.d.              |
| F13A7.10  | gsr-39                        | V-10F14    | 90.0±3.3         | 15.0±6.4***       |
| C29E4.7   | gsr-40                        | cloned     | 16.4±1.7**       | 86.5±9.0          |
| C02D5.3   | gsr-41                        | III-4F21   | 80.0±3.3         | 64.1±27.4         |
C46F11.2 encodes a member of the pyridine nucleotide-disulphide oxidoreductase family orthologous to human mitochondrial GSR (www.wormbase.org) and shares 65% and 58% amino acid sequence identity with the GSR from the filarial nematode *O. volvulus* and humans (data not shown). *gsr-1* is the sole GSR encoding gene found in *C. elegans*. The gene contains 5 distinct gt-ag introns and transcription produces four alternatively spliced mRNAs (C46F11.2a, C46F11.2b1-3). Whereas the long transcript C46F11.2a, starting from exon 1 has a coding region of 1422 bp (459 aa), the shorter transcripts C46F11.2b1-3, encompassing exons 2–5, have a coding region of 1380 bp (459 aa). As with other anti-oxidant factors, the *gsr-1* is clearly expected to reside in the cytosol and mitochondria. Since both a mitochondrial and a cytosolic isoform of the *GSR-1* could be generated from a single gene through alternative splicing, analysis of the long isoform was performed, using PSORT [41]. This program allows the prediction of protein localization sites and a mitochondrial localization with a probability of 43.5% was calculated. This is in agreement with the MITOPROT analysis [42], however, a mitochondrial matrix peptidase cleavage site could not be predicted. In worms carrying the *Pgsr-1::gsr-1::GFP* reporter construct (1996 bp of the 5′ upstream region and the complete open reading frame of *gsr-1* tagged with GFP), only strong cytosolic GFP expression was observed, possibly obscuring lesser granular mitochondrial GFP signals.

Since C46F11.2 had not been characterised before, we cloned the corresponding open reading frame of 1380 bp for recombinant expression in *E. coli*. Consistent with the theoretical molecular mass, the purified His-tag fusion protein that forms an enzymatically active homodimer had a molecular mass of approximately 52 kDa (Figure S1 A). The protein catalysed the reduction of GSSG by using NADPH as an electron donor. The Km values for GSSG and NADPH were found to be 34.1 ± 8.8 and 8.8 ± 1.1, respectively (Figure S1 B).

Initially we wanted to use the balanced *gsr-1* knockout strain (allele tm3574, mutation site: 9752/9753-10135/10136), however, even under normal laboratory conditions, the impairment of homozygous sterile mutants was so severe, making stress experiments futile. Therefore, to further evaluate the role of *C. elegans GSR-1* in stress tolerance, *gsr-1(RNAi)* worms were exposed to environmental stressors. To demonstrate the effect of *gsr-

### Table 2. Cont.

| Gene     | Annotation | RNAi clone | Juglone survival | Arsenite survival |
|----------|------------|------------|------------------|-------------------|
| C25H3.7  | gsr-like   | II-4H13    | 90.0 ± 8.8       | 78.3 ± 11.8       |
| R11A8.5  | gsr-like   | IV-5B15    | 81.1 ± 9.6       | 85.3 ± 3.5        |

* `-Glutamyltransferase ( `-GT`)

| T03D8.6* | -          | V-13A06    | 83.3 ± 23.1      | 70.0 ± 4.7        |
| C53D5.5  | -          | I-1C15     | 90.0 ± 5.8       | 70.0 ± 4.7        |
| Y7A9A.1  | -          | IV-7H18    | 84.4 ± 5.1       | 46.7 ± 23.3       |
| H14N184* | -          | V-6E10     | 77.8 ± 24.1      | 58.9 ± 22.2       |
| Y97E10A.2* | -        | V-13B05    | 92.2 ± 7.7       | 75.0 ± 11.8       |
| T19H12.6 | -          | V-3L18     | 86.7 ± 8.8       | 58.9 ± 17.1       |

Glutathione peroxidase (GPX)

| F26E4.12 | gpx-1      | I - 4L14   | 96.7 ± 3.3       | 60.0 ± 18.9       |
| C11E4.2  | gpx-3      | X - 4D12   | 90.0 ± 11.5      | 83.3 ± 4.7        |
| C11E4.1  | gpx-5      | X - 4D10   | 80.0 ± 14.5      | 13.3 ± 9.4***     |
| T09A122* | gpx-6      | IV - 4M01  | 84.4 ± 11.7      | 71.7 ± 2.4        |
| R03G5.5* | gpx-7      | X - 4E07   | 91.1 ± 3.8       | 61.7 ± 21.2       |
| F55A5.5  | gpx-8      | I - 5O10   | 92.2 ± 3.8       | 66.7 ± 23.6       |

Glutaredoxin (GLRX)

| D2063.3* | glrx-3     | cloned     | 89.3 ± 0.5       | 96.6 ± 1.0        |
| Y49E10.2 | glrx-5     | III - 6C14 | 88.9 ± 10.7      | 53.7 ± 27.8       |
| Y34D9A.6 | glrx-10    | I - 7D15   | 90.0 ± 8.8       | 83.3 ± 14.1       |
| ZK1211*  | glrx-21    | III - 2F14 | 94.4 ± 3.8       | 79.8 ± 9.2        |
| C07G1.8  | glrx-22    | cloned     | 94.8 ± 9.8       | 81.8 ± 2.1        |

Phytochelatin synthase (PCS)

| F54D5.1* | pcs-1      | II - 7J22  | 78.1 ± 8.5       | 89.1 ± 1.4        |

Glyoxalase (GLOD)

| C16C10.10 | glod-4     | III - 2G01 | 87.9 ± 11.2      | 82.7 ± 2.0        |

Genes involved in the *C. elegans* GSH-metabolism were classified according to their reported or predicted functions. Each of the 67 listed genes was targeted by RNAi starting in L4 animals. F1 L4/young adults of the treated worms were transferred to 0.15 mM juglone and 3 mM arsenite stress plates, respectively, and the survival rates (± SD) were determined after 18 h at 20°C. Significance levels were determined by student’s t-test (n ≥ 3, at least 90 animals, *P* ≤ 0.05, **P** ≤ 0.01, and ***P** ≤ 0.001). RNAi clones of the RNAi library (Geneservices, positions are shown) were verified by PCR using T7-primer followed by analytical restriction. Where indicated, RNAi constructs for genes that are not included in the library were generated by standard cloning of PCR products into the pL4440 vector, n.d. not determined.* genes possess different isoforms, however, all isoforms are affected by the RNAi constructs used.

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1 (RNAi) on gsr-1 expression levels, gsr-1 (RNAi) was performed on Pgsr-1::gsr-1::GFP worms. Here, the dimmed GFP fluorescence signal and a reduced amount of GSR-1::GFP clearly indicates the efficiently reduced gsr-1 expression (Figure S2).

As shown in Figure 1A, GSR-1 was also found to be crucial for the tolerance against other pro-oxidants including paraquat and to a lesser extent cumene hydroperoxide and tert-butylhydroperoxide. Knockdown of GSR-1, however, did not affect the sensitivity of C. elegans towards arsenite. Remarkably, protection against this stressor required the presence of the GSH synthesis enzyme gcs-1 (Figure 1B). In contrast to that, RNAi-mediated inhibition of GCS-1 hardly altered the susceptibility to juglone.

GCS and - to a Lesser Extend gst-32, gst-44 and an Extracellular GPX C11E4.11 - are Required for Tolerance Towards Arsenite

Several studies have shown that GSH synthesis plays a critical role in the protection of C. elegans against inorganic arsenite, with GCS-1 being involved in the protection [43,44]. We can confirm that the first rate-limiting enzyme of GSH synthesis, the GCS is essential for protection (0% survival rate of gcs(RNAi)-worms under the given arsenite concentration) (Figure 1b). Furthermore, knockdown of an alpha-class related GST (gst-32), an omega-class GST (gst-44) and an extracellular glutathione peroxidase (C11E4.11) less profoundly affected the survival rate under arsenite stress (arsenite survival rate of 24.0 ± 6.13 (p > 0.01), 15.0 ± 6.14 (p < 0.001) and 13.3 ± 9.4 (p < 0.001), respectively). Knockdown of gsr-1 expression had no effect on the survival rate of worms under arsenite stress conditions (Table 2). However, this negative result

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Figure 1. Analyses of GSR-1 and GCS-1 dependent stress tolerance. Wild-type eggs/L1 were placed on RNAi plates (A) gsr-1(RNAi) or (B) gcs-1(RNAi) and cultured until they have reached the L4/young adult stage, before being transferred to corresponding RNAi plates containing the indicated stressors. As control, plates with empty control vector pL4440 were used. Following 16 h incubation at 20°C, the survival rate was determined. Data represent means of at least three independent triplicate determinations (n≥120 animals, p values from Fisher test).
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must be treated with caution since small amounts of residual GSR-1 might be sufficient for arsenite tolerance.

The Effect of gsr-1(RNAi) and Juglone Stress on the GSH-redox State

The physiological role of GSR is the recycling of oxidized GSSG to GSH, thereby maintaining the intracellular GSH redox state, which is essential for cellular integrity. To test the role of *C. elegans* gsr-1 in the regulation of the redox state, we therefore analysed the GSH/GSSG ratio in *gsr-1(RNAi)* worms (Figure 2). Interestingly, knockdown of GSR-1 hardly affected the total GSH-level of about 40 nmol/mg protein, nor the GSH/GSSG ratio found in control worms (70:1). However, when worms were exposed to oxidative stress using juglone, the effect on GSH-related metabolites, that is enhanced GSSG- and γ-glutamylcysteine (γ-GC) levels and reduced GSH/GSSG ratios, was potentiated (Figure 2). The GSH-precursor γ-GC changed from

Figure 2. Determination of γ-glutamylcysteine, GSH and GSSG levels in lysate of synchronised 3-days-old worms. (A) Worms were cultured on RNAi-plates containing (i) HT115 bacteria carrying empty pL4440 vector or (ii) HT115 bacteria carrying empty pL4440 vector plus 150 μM juglone or (iii) *gsr-1(RNAi)* feeding bacteria or (iv) *gsr-1(RNAi)* feeding bacteria plus 150 μM juglone. (B) Analysis of the respective GSH/GSSG ratios. doi:10.1371/journal.pone.0060731.g002
2.3 to 8.1 nmol/mg protein, indicating that GSH-synthesis is boosted under oxidative stress conditions. Simultaneously, the GSSG-concentration altered from 2.0 to about 6.0 nmol/mg protein in control worms and \textit{gsr-1}(RNAi) worms exposed to juglone stress, leading to GSH/GSSG ratios of about 20:1 and 6:1, respectively. This explains the enormous effect of \textit{gsr-1} knockdown on the tolerance towards juglone stress observed in the above mentioned results from the juglone toxicity RNAi-screening (\textit{Table 2}).

Expression Pattern of Stress-inducible GSR-1

To analyse the temporal and spatial expression pattern of \textit{C. elegans gsr-1}, we generated transgenic  \textit{Pgsr-1::GFP} and  \textit{Pgsr-1::gsr-1::GFP} worms expressing the GFP reporter under the control of the \textit{gsr-1} promoter. In both cases bright fluorescence was seen mainly in the intestine, vulval muscle, the pharynx and some unidentified cells in the tail region (\textbf{Figure 3A}). Although the overall spatial GFP expression pattern was similar throughout all postembryonic life stages, a general increase in the fluorescence signals was observed during the development from L1 to adult worms. As control, \textit{gsr-1}(RNAi) on \textit{Pgsr-1::gsr-1::GFP} worms was
performed and efficiently reduced gsr-1 expression as indicated by a dimmed GFP fluorescence signal and a reduced amount of GSR-1::GFP fusion protein (Figure S2). Notably, the high GFP expression level was further elevated when the reporter strains were exposed to juglone (Figure 3B).

Northern blot analyses of N2 wildtype worms confirmed the enhanced gsr-1 expression in the presence of the naphthoquinone stressor (Figure 4).

Juglone and Starvation Induced gsr-1 Expression is Regulated by SKN-1, but not by DAF-16

In C. elegans the transcription factors DAF-16 and SKN-1 are crucial for the coordination of stress response and both have been shown to respond to juglone [16,28]. To analyse the contribution of these transcription factors to the regulation of the stress-responsive gsr-1 promoter of C. elegans, we used the Pgsr-1::GFP worms. As shown in Figure 5, the knockdown of DAF-16 does not affect GSR-1::GFP expression under standard condition and, moreover, had no effect on the induction of GSR by juglone. Conversely, reduction of SKN-1 expression by RNAi led to a reduced GFP signal under standard conditions and to a significantly reduced induction of reporter gene expression under juglone stress (Figure 6). These data strongly suggest that the gsr-1 promoter is regulated by SKN-1 and not by DAF-16. Furthermore, the obtained data indicate that SKN-1 is required for GSR-1 expression per se and for the drastic response observed under juglone stress. Accordingly, in silico analyses revealed three well conserved potential SKN-1 binding sites [45] at position -443, -617 and -651 of the gsr-1 promoter region, while DAF-16 binding sites are lacking within the 1992 bp of the upstream region (Figure S3).

In C. elegans, SKN-1 is also involved in the regulation of gene expression under conditions of dietary deprivation [46]. Therefore, we transferred Pggr-1::GFP L4/young adult worms from standard NGM plates containing E. coli OP50 to starvation plates without bacterial food. As shown in Figure 3C, the GFP signal was considerably enhanced under starvation conditions. In good agreement with the results obtained for juglone stress, RNAi assays revealed that the observed induction of the gsr-1 promoter activity was found to be SKN-1 but not DAF-16 dependent (Figure 7).

Knockdown of gsr-1 Induces gcs-1 Expression but not Vice Versa

Since the cellular GSH status is regulated by redox cycling and de novo synthesis, we next asked whether a reduced GSH redox cycling affects GSH synthesis and, vice versa, inhibition of GSH synthesis affects GSR-1 expression. As shown in Figure 8, inhibition of GSH synthesis by gcs-1(RNAi) did not affect gsr-1 promoter activity. In contrast to that, gsr-1(RNAi) resulted in a significantly enhanced GFP signal in the intestinal cells of the Pgcs-1::GFP reporter strain (Figure 8 and Figure S4 A) indicating that GSH synthesis is induced when GSSG recycling is impaired.

Figure 4. Northern blot analyses. C. elegans N2 wild-type worms were cultured on NGM agar under standard conditions (lane 1) or exposed to 150 μM (lane 2) and 300 μM juglone for 4 h, before total RNA was isolated and applied to Northern blot analyses using a radiolabeled gsr-1 probe. Since there is only a 40 bp difference in transcript size (C46F11.2a, C46F11.2b1-3), only one band can be observed.

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Figure 5. Juglone-induced GSR-1 expression is not regulated by DAF-16. Worms carrying the Pggr-1::GFP reporter construct were grown on HT115 bacteria carrying empty pL4440 vector (A, B) or subjected to daf-16(RNAi) (C, D). The respective effects on GFP-expression were monitored under standard culture conditions (A, C) or after induction by juglone (B, D) (quantified by ImageJ). Scale bars = 100 μm.

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Under stress conditions, *gcs-1* promoter activity is stimulated by the transcription factor SKN-1 [33]. Since the expression of another SKN-1 responsive gene, *gst-4*, was similarly induced by *gsr-1(RNAi)* (Figure 8 and Figure S4 B), we next analysed the nuclear translocation of a *SKN-1b/c::GFP* fusion reporter, indicative for the activation of the transcription factor. However, consistent with a recent report [47], the knockdown of *gsr-1* did not increase the nuclear accumulation of SKN-1 (data not shown). Furthermore, *gsr-1(RNAi)* did not affect the nuclear translocation of the stress responsive transcription factor DAF-16 (data not shown).

**GSR-1 Overexpression Protects against Juglone Toxicity**

Since the expression level of GSR-1 was found to be elevated when worms were exposed to juglone, we next examined whether overexpression of GSR-1 is capable of increasing the stress tolerance of *C. elegans*. Transgenic *Pgsr-1::gsr-1::GFP* worms exhibited a significantly higher resistance towards juglone than the control strain (Figure 9). 29% of control and 66% of *gsr-1p::GFP* worms survived when exposed to 0.25 mM juglone for 16 h.

**GSR-1 Expression Level Affects Life Span**

According to the ROS theory of aging, the degree of stress resistance is often related to the life span of an organism [7,8]. Likewise, the knockdown of GSR-1 led to a significantly reduced mean life span of 17.7 days when compared with control animals with a mean value of 20.7 days (Figure 10A), thus confirming the results obtained by [29].

Mean values of the 10% longest living worms were 22.4 days and 28.0 days, respectively. We observed no retardation in postembryonic development. No effects on brood size and number on progeny were observed (control: 252±35, *gsr-1(RNAi)*: 269±45; n = 30 worms).

On the other hand worms that overexpress GSR-1 displayed a prolonged life span when compared to control animals with mean values of 19.4 and 16.4 days, respectively (Figure 10B). Control worms had a maximal life span of 24.5 days, GSR-1 overexpressers of 27.9 days (mean values of the 10% longest living worms). The life span assay was performed at 20°C.

**Discussion**

In this study we conducted a small-scale RNAi screen aiming to identify components of the GSH-metabolism that are essential for tolerance towards the pro-oxidative stressors arsenite and juglone. In *C. elegans*, RNAi screening is a highly efficient and convenient method to determine loss-of-function phenotypes of an individual gene. A systematic analysis of the RNAi library used in this study showed that 90% of genes that were identified by classical genetics to cause embryonic lethality also showed the same phenotype using the RNAi feeding method, showing its effectiveness and quality of the library [48]. However, it is important to note that the effectiveness of RNAi in the suppression of gene expression can be variable and we cannot confirm a complete abrogation of gene function for each RNAi clone used that did not cause a detectable phenotype. Additionally, our RNAi experiments cannot exclude combined effects of several enzymes. However, we can be sure of the RNAi clones that showed a detectable phenotype, like the *gsr-1*, *gst-44*, *gst-32*, *gst0-1*, *gcs-1* and the extracellular GPX. Interestingly, a recent publication clearly demonstrates the combined action of thioredoxin reductase and GSR-1 during molting. Here the regulated reduction of cuticle components is driven by thioredoxin reductase and GSR-1 [49].

Arsenic-mediated toxicity is thought to be due to oxidative stress on the one hand and on the other hand due to biotransformation by methylation and accumulation of the metalloid in the cells. As described before, tolerance towards arsenite is primarily GSH
synthesis-dependent, with p38 MAPK being activated by arsenite treatment, the PMK-1 phosphorylating the transcription factor SKN-1, thereby inducing its nuclear translocation and the activation of gcs-1 gene expression [32,43,44]. Additionally, our RNAi screen indicated the involvement of an antioxidant protein, the extracellular GPX (C11E4.1) and two phase II xenobiotic-metabolizing enzymes, thegst-32 andgst-44. Both, the alpha-class relatedgst-32 and the GPTX are implicated in the oxidative stress response by way of their GSH-dependent reduction of inorganic and organic peroxides or by their ability to eliminate secondary products of lipid peroxidation. However, the role ofgst-44 remains unclear. Since methylation of arsenic involves an omega-class GST that functions as a monomethylarsonate reductase [22], an involvement ofgst-44 in arsenic biotransformation seems feasible. Also, the enzyme is exclusively localizes in the cytoplasm of the H-shaped cell associated with the excretory system. Besides osmoregulation and secretion of molting fluid or homones, one of the proposed functions of the excretory system is the excretion of metabolic waste (Wormatlas database) However, like Drosophila melanogaster, C. elegans does not possess an arsenic methyltransferase homolog, making methylation processes of inorganic arsenic highly unlikely to occur in C. elegans [50]. Another possible function of the omega-class GST could be the reduction of GSH-mixed disulphides once oxidative stress has been removed. Knockdown of the GSR-1 does not seem to affect arsenite sensitivity. However, as mentioned above, negative RNAi results must be treated with caution since small amounts of residual GSR-1 might be sufficient for arsenite tolerance. Literature search also gave ambiguous results. For example, exposure of porcine endothelial cells to trivalent arsenics caused an increase in total glutathione levels and higher enzymatic activity of GPX and GST, with no increase being observed in GSR-activity [51]. Whereas in rice roots, the enzymatic activity of the GSR responded in a dose-dependent manner to arsenic [52], in ferns the GSR activity remained unchanged, with the GR not being inhibited or activated by arsenite in enzymatic assays [53].

The toxicity of juglone is due to redox cycling as well as direct electrophilic interaction with glutathione [22]. In our small-scale RNAi-screen, only the GSR-1 was identified to be absolutely essential for survival under sub-lethal juglone stress. This indicates that the maintenance of low GSSG concentrations is vital to the

Figure 7. Starvation and regulation of gsr-1 promoter activity. Worms carrying the Pgsr-1::GFP reporter construct were grown on HT115 bacteria carrying empty pL4440 vector (A and B) or subjected to skn-1(RNAi) (C and D) or daf-16(RNAi) (E and F). The respective effects on GFP expression were monitored under standard culture conditions (A, C, E) or after hunger (B, D, F) (quantified by ImageJ, Scale bars = 100 μm).
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preservation of an adequate reduction potential when exposed to increasing oxidative stress.

At our applied juglone concentration, no changes in the total GSH level of stressed control worms were found, however, enhanced GSSG- and γ-GC levels and a reduced GSH/GSSG ratio were observed. Under oxidative insult, the existing GSH pool is oxidized to GSSG by nonenzymatic or enzymatic routes. Here GPX uses GSH to reduce peroxides, while GRX reduces protein disulphides directly via its active site dithiol, which is converted to a disulphide needing to be reduced again by GSH and thereby producing GSSG. The availability of GSH is ensured by GSSG recycling and/or by de novo synthesis, both of which are up-regulated when oxidative stress occurs [54]. Another protective mechanism during oxidative insult is the rapid and active transportation of excessive GSSG out of the cell, demonstrating the acute need of the cell to prevent its toxic accumulation [55]. Using our current experimental setup of derivatising whole worms, we cannot detect whether active GSSG export is taking place during juglone stress and/or paired with gsr-1(RNAi), but this can be the explanation of why we observe constant total GSH levels. However, the observed changes in γ-GC levels strongly indicate that de novo GSH synthesis is boosted under juglone stress and that it is potentiated by simultaneous gsr-1(RNAi).

During juglone stress, gsr-1(RNAi) aggravates intracellular oxidative challenge, as any GSSG generated in the cell cannot be reduced back to GSH, thereby severely limiting the efficient removal of peroxides by GPX or the reduction of protein disulphides by GRX, which is at the expense of GSH to GSSG. Therefore, these functions can only be performed until the supply of GSH is exhausted. This explains the enormous effect of gsr-1 knockdown on the tolerance towards juglone stress observed in the above mentioned results. Likewise, overexpression of GSR-1 is

Figure 8. Analyses of the interrelation between GSH synthesis and GSH redox cycling. Applying RNAi, GCS-1 expression was suppressed in the C. elegans Pgsr-1::GFP reporter strain (left) or GSR-1 expression in the Pgcs-1::GFP (middle) or Pgst-4::GFP reporter strain (right). RNAi-treated worms were analysed by fluorescence microscopy and GFP-expression was compared with control worms grown on HT115 bacteria carrying empty pL4440 vector. Fluorescence intensities for Pgsr-1::GFP reporter strain were scored as low (A), medium (B) and high (C) according to [16]. Fluorescence intensity scoring for Pgcs-1::GFP and Pgst-4::GFP reporter strains is given in Figure S4. Significant alterations in GFP expression was detected only in Pgsr-1::GFP (p<0.001) and Pgst-4::GFP worms (p<0.001) exposed to GSR-1(RNAi). Results are the means of at least three independent assays (n = 100 worms; p values from Fisher test).

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Figure 9. Enhanced stress tolerance by GSR-1 overexpression. L4/young adult of C. elegans overexpressing GSR-1::GFP fusion protein or of control worms expressing a GFP protein only were exposed to 0.25 mM juglone. Survival rate was determined after 16 h incubation. Owing to the pha-1(e2123) genetic background, assays were performed at 25°C. Results are means of three independent triplicate assays (n=120 animals, Fisher test).

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Glutathione Reductase of C. elegans
The affinity between the GCS and its substrates. In age-related accumulation of homocysteine is causing a loss of experimental studies by Rebrina & Sohal [56] suggest that the overexpression of GSR-1 led to longevity (p < 0.001, results are cumulative from three independent experiments with 50 worms per trial), whereas (B) overexpression of GSR-1 is associated with an enhanced stress tolerance of C. elegans, capable of increasing the stress tolerance of C. elegans overexpression of GSR-1 led to longevity (p < 0.001, results are cumulative from three independent experiments with 100 worms per trial). As control, pha-1 transgenic worms were used that express GFP in a similar expression pattern (see Material and Methods).

Figure 10. GSR-1 dependent life span. (A) gsr-1(RNAi) resulted in a reduced life span (log-rank test p < 0.001, results are cumulative from three independent experiments with 50 worms per trial), whereas (B) overexpression of GSR-1 led to longevity (p < 0.001, results are cumulative from three independent experiments with 100 worms per trial). As control, pha-1 transgenic worms were used that express GFP in a similar expression pattern (see Material and Methods).

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Supporting Information

Figure S1 Recombinant expression and biochemical characterisation of C. elegans GSR-1.
(DOCX)

Figure S2 Gsr-1(RNAi) on gsr-1p::GFP worms.
(DOCX)

Figure S3 GSR-1 promoter region. Putative SNK-1 conserved binding sites were manually determined using the consensus sequence for SKN-1 binding WWT

(DOCX)

Figure S4 Representative fluorescence intensities for Pgsr-1::GFP reporter strain and Pgst-4::GFP reporter strain are shown.

(DOCX)

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Author Contributions
Conceived and designed the experiments: KL EL PH. Performed the experiments: DS MD IA CA JD CW. Analyzed the data: DS KL EL MD CJW. Wrote the paper: KL EL.

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