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Compartment-specific perturbation of protein handling activates genes encoding mitochondrial chaperones

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Summary

Protein folding in the mitochondria is assisted by nuclear-encoded compartment-specific chaperones but regulation of the expression of their encoding genes is poorly understood. We found that the mitochondrial matrix HSP70 and HSP60 chaperones, encoded by the *Caenorhabditis elegans* *hsp-6* and *hsp-60* genes, were selectively activated by perturbations that impair assembly of multi-subunit mitochondrial complexes or by RNAi of genes encoding mitochondrial chaperones or proteases, which lead to defective protein folding and processing in the organelle. *hsp-6* and *hsp-60* induction was specific to perturbed mitochondrial protein handling, as neither heat-

shock nor endoplasmic reticulum stress nor manipulations that impair mitochondrial steps in intermediary metabolism or ATP synthesis activated the mitochondrial chaperone genes. These observations support the existence of a mitochondrial unfolded protein response that couples mitochondrial chaperone gene expression to changes in the protein handling environment in the organelle.

Supplemental data available online

Key words: Protein folding, Chaperones, Organelle, Signaling, Genetics

Introduction

Chaperones are a diverse group of proteins with important roles in nascent polypeptide folding, assembly of multimeric protein complexes, protein translocation across membrane barriers, integration into membranes and protein degradation (Bukau and Horwich, 1998; Horwich et al., 1999; Hartl and Hayer-Hartl, 2002). In eukaryotes, chaperones are segregated to specific organelles. For example, the cytoplasm, endoplasmic reticulum (ER) lumen and mitochondrial matrix each contain unique DnaK/Hsp70-type and DnaJ/Hsp40-type chaperones encoded by specific nuclear genes (Martin, 1997; Fewell et al., 2001; Hartl and Hayer-Hartl, 2002), whereas distinct GroE/Hsp10/60-type chaperonins are found in the cytoplasm and mitochondrial matrix but not in the ER lumen (Horwich and Willison, 1993).

Chaperone engagement depends on the balance between chaperone levels and client protein load. The latter is set by the rate of protein synthesis and translocation into specific organelles and is influenced by the intrinsic ability of client proteins to fold, enter into productive complexes with other proteins and retain their properly folded state. Increased chaperone engagement by perturbations that impede proper protein folding, encourage protein complex disassembly or promote misfolding, activates chaperone-encoding genes. For this reason, many chaperones were first identified as heat-

shock proteins, induced at elevated temperature (Lindquist and Craig, 1988).

Studies of chaperone gene expression have revealed a conserved paradigm in which signaling pathways activating chaperone-encoding genes are repressed by free chaperones that are not engaged by client proteins. In *E. coli* the transcription factor activating the heat shock operon, σ_{32} , is repressed by free DnaK (Bukau, 1993; Biaszczak et al., 1999). In eukaryotes signal transduction pathways that regulate expression of ER chaperones are repressed by binding of the ER chaperone BiP to the luminal domains of the proximal signal transducers IRE1, PERK and ATF6 (Bertolotti et al., 2000; Okamura et al., 2000; Liu et al., 2002; Shen et al., 2002). By contrast, cytosolic chaperones specifically repress the heat-shock sensitive transcription factors (HSFs) that regulate their expression (Morimoto, 1998). These observations suggest that cells monitor the folding environment in specific compartments by determining the sufficiency of their chaperone reserves and defend this reserve selectively in each compartment. Thus, segregation of chaperones to specific organelles implies that chaperone-encoding genes respond selectively to the needs of specific organelles.

Compartment specificity in signaling to chaperone-encoding genes has been demonstrated for the ER and cytoplasm. This was established by studies that relied on biochemical tools to

perturb protein folding in a compartment-selective manner. For example, treatment of cells with arsenite, which preferentially perturbs protein folding in the cytoplasm, selectively activates cytoplasmic chaperone gene expression (Mosser et al., 1993; Liu et al., 1994; Harding et al., 1999; Calfon et al., 2002), whereas tunicamycin, an agent that blocks the ER-specific modification of N-linked glycosylation preferentially activates pathways promoting the expression of ER-localized chaperones (Gething and Sambrook, 1992). The molecular basis for this specificity is partially understood, as some of the proximal transducers that selectively activate genes encoding cytoplasmic or ER chaperones have been identified (Morimoto, 1998; Kaufman, 1999; Patil and Walter, 2001; Harding et al., 2002).

Less well understood are the mechanisms that regulate genes encoding chaperones localized to mitochondria. It has recently been shown that over-expression of a folding-incompetent mitochondrial matrix protein, mutant ornithine transcarbamylase, activates mitochondrial chaperonin-encoding genes in mammalian cells. The stress-induced transcription factor CHOP was shown to play an important role in this signaling pathway (Zhao et al., 2002). However, the pathways linking CHOP activation to mutant OTC expression are not understood. Furthermore, given that CHOP is a vertebrate-specific transcription factor, whereas mitochondria are present in all eukaryotes, it seemed likely that other pathways for activating mitochondrial chaperones existed. We report on the establishment of genetic and pharmacological tools for perturbing protein handling in the mitochondria of the nematode *C. elegans* and used these to reveal a mitochondrial unfolded protein response that depends on transmitting information from the organelle to the nucleus.

Materials and Methods

Transgenic *C. elegans*

The strain containing the ER stress reporter *hsp-4::gfp(zcls4)* *V* has been previously described (Calfon et al., 2002). The mtHSP70 reporter, *hsp-6::gfp* was constructed by ligating a 1.7 kb *HindIII*-*Bam*HI PCR fragment derived by amplification of *C. elegans* genomic DNA with the primers: C37H5.5.2AS (TCGAGTCCATA-CAAGACTC) and C37H5.8.2AS (GGGGGATCCGAAGACAA-GAATGATCGTGC) into the GFP reporter plasmid pPD95.75 (a gift of Andy Fire, Baltimore MD, USA). *hsp-6::gfp* contains the 5' flanking region and encodes the predicted first 10 amino acids of HSP-6 fused to GFP. The *hsp-60::gfp* reporter was constructed by ligating a 2.3 kb *Sal*I-*Bam*HI PCR fragment derived by amplification of *C. elegans* genomic DNA with the primers: CeHSP10.6AS (AA-GAGTTCGACTCGCGGAAGATTGAGTATTCC) and CeHSP60.2AS (CTGAGGATCCCTTCTGGCGAGGGGAAGCATC) into pPD95.75. *hsp-60::gfp* contains the 5' flanking region and encodes the predicted first 7 amino acids of HSP-60 fused to GFP. The *myo-3::mgfp* expressing a mitochondrially localized GFP with a cleavable mitochondrial import signal peptide (Labrousse et al., 1999) was gift from Alexander van der Bliek (University of California Los Angeles, USA) and the strain expressing *sod-3::gfp* (muIs84, CF1553) (Libina et al., 2003) was a gift from Malene Hansen and Cynthia Kenyon (University of California San Francisco, USA). The *hsp-6::gfp* plasmid (25 µg/ml) and the *hsp-60::gfp* plasmid (5 µg/ml) were co-injected with a *lin-15* rescuing plasmid, pSK1 (25 µg/ml) into *lin-15(n765ts)* strain. The extrachromosomal arrays were integrated into the chromosome with ultraviolet/trimethylpsoralen treatment. Several independent lines were produced for each reporter, with identical GFP expression

patterns and one of each, *hsp-6::gfp(zcls13)* *V*, and *hsp-60::gfp(zcls9)* *V* were backcrossed four times into wild-type N2, eliminating the *lin-15* mutation. The *cts-1::gfp* reporter was constructed by ligating a 0.8 kb *Sal*I-*Bam*HI PCR fragment derived by amplification of *C. elegans* genomic DNA with the primers: T20G5Sal.1S (TTTGGTTCGACATGGTTCATCGGCGGAAGAG) and T20G5Bam.2AS (GCGAGGATCCATCCAGAGAGCGACATTTCTG) into pPD95.75 and the *aco-2::gfp* was constructed by ligating a 0.9 kb *Sal*I-*Bgl*II PCR fragment derived by amplification of *C. elegans* genomic DNA with the primers: F54H12Sal.1S (GCGGGTTCGACGCTCCTCAACGATCTTTGC) and F54H12Bgl.2AS (AGGTAGATCTAAGTGGCTCAACCGAAGAAGAG) into pPD95.75. Three independently produced extrachromosomal arrays of each line were maintained and two, *cts-1::gfp(zcEX12)* and *aco-2::gfp(zcEX13)* were used in this study. The cytoplasmic HSP70 (C12C8.1) reporter, *hsp-70::gfp(rmls8)* was a gift from Jim Morley and Richard Morimoto from Northwestern University (Chicago, IL, USA).

Pharmacological treatment

Gravid animals were bleached to obtain eggs that hatched to produce a synchronous population of young adults that were used in the treatments described below.

Ethidium bromide (Sigma) was dissolved in water at 10 mg/ml and added to agar plates to a final concentration of 5-125 µg/ml. 2,4 dinitrophenol (Sigma) was dissolved in ethanol at 1 M and added to agar plates to a final concentration of 0.1-2.5 mM. Bacteria were seeded on these plates and the animals were allowed to feed and produce a brood. The brood was harvested for study on day 3 after egg laying. Animals were heat shocked at 30°C or treated with 1 µg/ml tunicamycin, for 6 hours.

Tetramethylrhodamine ethyl ester (TMRE; Molecular Probes), a lipophilic cation whose mitochondrial uptake depends on the $\Delta\Psi$ -potential (Farkas et al., 1989; Loew et al., 1993) was dissolved in DMSO at 50 µM and applied to the agar plate at final concentration of 0.1 µM. Untreated and DNP-treated animals were placed on the TMRE plates for 12 hours, removed, washed with M9 medium, embedded in agarose and examined using rhodamine filters in a Zeiss Axiophot microscope. Fluorescence and transmission photomicrographs were obtained using a digital CCD camera and processed in Adobe Photoshop.

RNAi

Interference with gene function by RNAi followed an established protocol (Timmons et al., 2001). Briefly, double stranded RNA was produced in the HT115 strain of *E. coli* transformed with pPD129 plasmids containing cDNA fragments of genes being studied: a 675 bp *Eco*RI-*Xho*I of the *hsp-60* EST yk515d7 (a gift from Yoji Kohara), a 735 bp *Kpn*I-*Bam*HI fragment of the Paraplegin homologue (Y47G6A.10, *spg-7*)-derived EST yk282e3, a 595 base-pair *Sac*I-*Bam*HI fragment of *hsp-6* (C37H5.8)-derived EST yk313b1, a 423 nucleotide *Cl*aI-*Xho*I fragment from *ero-1* (Y105E8B.8) a 794 bp PCR fragment of the PDI homologue (H06O01.1, *pdi-3*) derived by the primers TTCCAGCCCAAGACTACGA and TTTCCA-ACGGCAACCTTAAC. Inactivation of *aco-2* (F54h12.1a), *mdh-1* (F20H11.3), *fum-1* (H14A12.2a) and *cts-1* (T20G5.2) was carried out by the primer pairs as described in Wormbase (<http://www.wormbase.org/>). RNAi of genes on chromosome I was carried out by sequential feeding of an arrayed library of validated genomic fragments obtained from the United Kingdom Human Genome Mapping Project Resource Centre (Cambridge, UK) as described previously (Fraser et al., 2000). Isopropyl-β-D-thiogalactopyranoside (IPTG, 1 mM) was added to the bacterial growth medium to induce transcription of the double stranded RNA and L4 staged animals of defined genotype were added to plates

individually and produced their brood. RNAi phenotypes were evaluated in their F1 progeny 3 days later.

Northern blots, Southern blots and immunoblots

Total RNA was prepared using the guanidine thiocyanate-acid-phenol extraction method and 15 µg were applied to each lane of a 1.5% agarose gel. The hybridization probes used to detect *hsp-6*, *spg-7*, *cts-1*, *aco-2*, F1F0 ATP synthase α -chain (H28O16.1) and cytochrome *c* oxidase subunit IV (W09C5.8) were derived from the fragments used in RNAi (see above). Mitochondrially encoded *cox-1* mRNA was detected using a 1075 bp PCR fragment derived by the primers TTACATGCAGCAGGGTTAAG and TCAGACTGATAACTGTG-ACC and *hsp-4* was detected as previously described (Calfon et al., 2002).

Total cellular DNA (mitochondrial and nuclear), 10 µg, was digested with *Eco*RI resolved on a 1% agarose gel and hybridized sequentially with the radiolabeled *cox-1* probe described above or with a 1065 bp fragment of the nuclear gene *K08H10.2*, derived by the PCR primers TGCTAAAGACAAGACTTCGG and ACGA-TACGGAGTCAGCAACG.

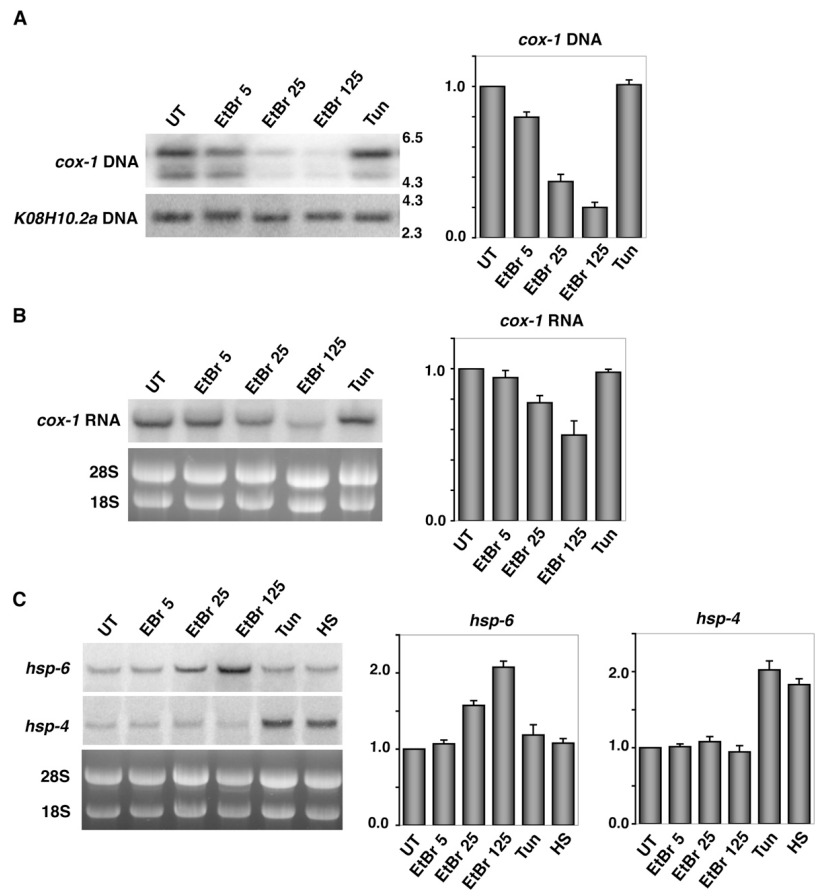
Immunoblot analysis of protein levels were performed on whole animal extracts prepared by washing animals in M9 medium to remove adherent bacteria, crushing on ice with a mechanical Teflon homogenizer in 3 volumes of lysis buffer (20 mM Hepes, pH 7.5, 150 mM NaCl, 1% Triton X-100, 10% glycerol, 1 mM EDTA 10 mM tetrasodium pyrophosphate, 100 mM NaF, 17.5 mM β -glycerophosphate, 1 mM phenylmethylsulfonyl fluoride, 4 µg/ml aprotinin, and 2 µg/ml pepstatin A). The lysate was clarified by centrifugation for 15 minutes at 13,000 *g* at 4°C. Aliquots containing 50 µg of protein were loaded in each lane of a 15% SDS-PAGE for electrophoresis and the immunoblot developed with a rabbit polyclonal serum to GFP at 1:10,000 dilution (gift from Pam Silver, Boston MA, USA) or goat polyclonal antiserum to UNC-32 (Santa Cruz Biotechnology sc-15649, lot #A042). To measure the solubility of GFP extracted by Triton X-100, the crude lysates were clarified by centrifuging at 14,000 *g* for 10 minutes in a table-top centrifuge and the supernatant was then centrifuged at 100,000 *g* for 1 hour to obtain an insoluble pellet and soluble supernatant which was then solubilized in SDS. Equal fractions of the total lysates, 100,000 *g* supernatant and pellet were resolved by SDS-PAGE and subjected to immunoblot as described above.

Fig. 1. Attenuated function of the mitochondrial genome induces mtHSP70 (*hsp-6*) mRNA. (A) Autoradiogram of a Southern blot of *Eco*RI-digested total DNA from animals raised in the indicated concentration of ethidium bromide (EtBr; µg/ml), or tunicamycin (Tun; 1 µg/ml). The blot was hybridized sequentially to a labeled DNA fragment of the mitochondrially encoded *cox-1* gene and the nuclear encoded *K08H10.2a* gene. (B) Autoradiogram of the northern blot of total RNA from animals cultured on agar plates containing the indicated concentration of ethidium bromide or tunicamycin. (Upper panel) The blot was hybridized to a labeled *cox-1* DNA fragment. (Lower panel) Ethidium bromide staining of the ribosomal bands of the RNA. (C) Autoradiogram of a northern blot of total RNA from animals treated with ethidium bromide, tunicamycin or heat-shock (HS). The blot was hybridized sequentially to radiolabeled fragment from the mtHSP70 gene (*hsp-6*) and the *C. elegans* BiP homologue, *hsp-4*. Bar charts on the right of each panel show the relative intensity of the hybridization signal.

Results

Proteins encoded by nuclear genes are imported into mitochondria. During import, they associate with mitochondrial chaperones and are released upon proper folding, often only after incorporation into multi-subunit complexes (Neupert and Pfanner, 1993; Neupert, 1997). Many such complexes also contain mitochondrially encoded proteins, which are required for complex stability (Stuart and Neupert, 1996). Thus, the fixed stoichiometry of the nuclear and mitochondrially encoded components of these multimeric complexes predicts that disrupting the supply of mitochondrially encoded proteins will interfere with complex assembly and increase the association of the mitochondrial chaperones with unassembled components. Ethidium bromide preferentially impairs replication and transcription of the mitochondrial genome in cultured cells, reducing the synthesis of mitochondrially encoded polypeptides (King and Attardi, 1989; Hayashi et al., 1990). We tested whether *C. elegans* raised on medium containing ethidium bromide would also have reduced mitochondrial DNA and reduced expression of mitochondrially encoded genes.

Hybridization analysis revealed a dose-dependent decrease in mitochondrial *cox-1* DNA and mRNA in animals raised on ethidium bromide (Fig. 1A,B). Remarkably, animals survived ethidium bromide treatment, but generation time was prolonged and brood size reduced. We studied the effect of ethidium bromide on activation of the mitochondrial matrix HSP70 (mtHSP70) gene, *hsp-6*. Northern blots showed a dose-dependent increase in *hsp-6* mRNA in animals raised on ethidium bromide (Fig. 1C). By contrast, ethidium bromide did



not activate *hsp-4*, which encodes a *C. elegans* homologue of the ER-localized chaperone, BiP. As previously noted, *hsp-4* was activated by treatment with tunicamycin or by culture at elevated temperature (Fig. 1C). However, these stressful treatments did not increase *hsp-6* mRNA levels.

We generated an *hsp-6::gfp* transcriptional reporter and introduced this fusion transgene into animals. GFP was expressed broadly in multiple transgenic lines and expression was noted from L1 larva to adults. ethidium bromide caused time- and dose-dependent increase in *hsp-6::gfp* expression

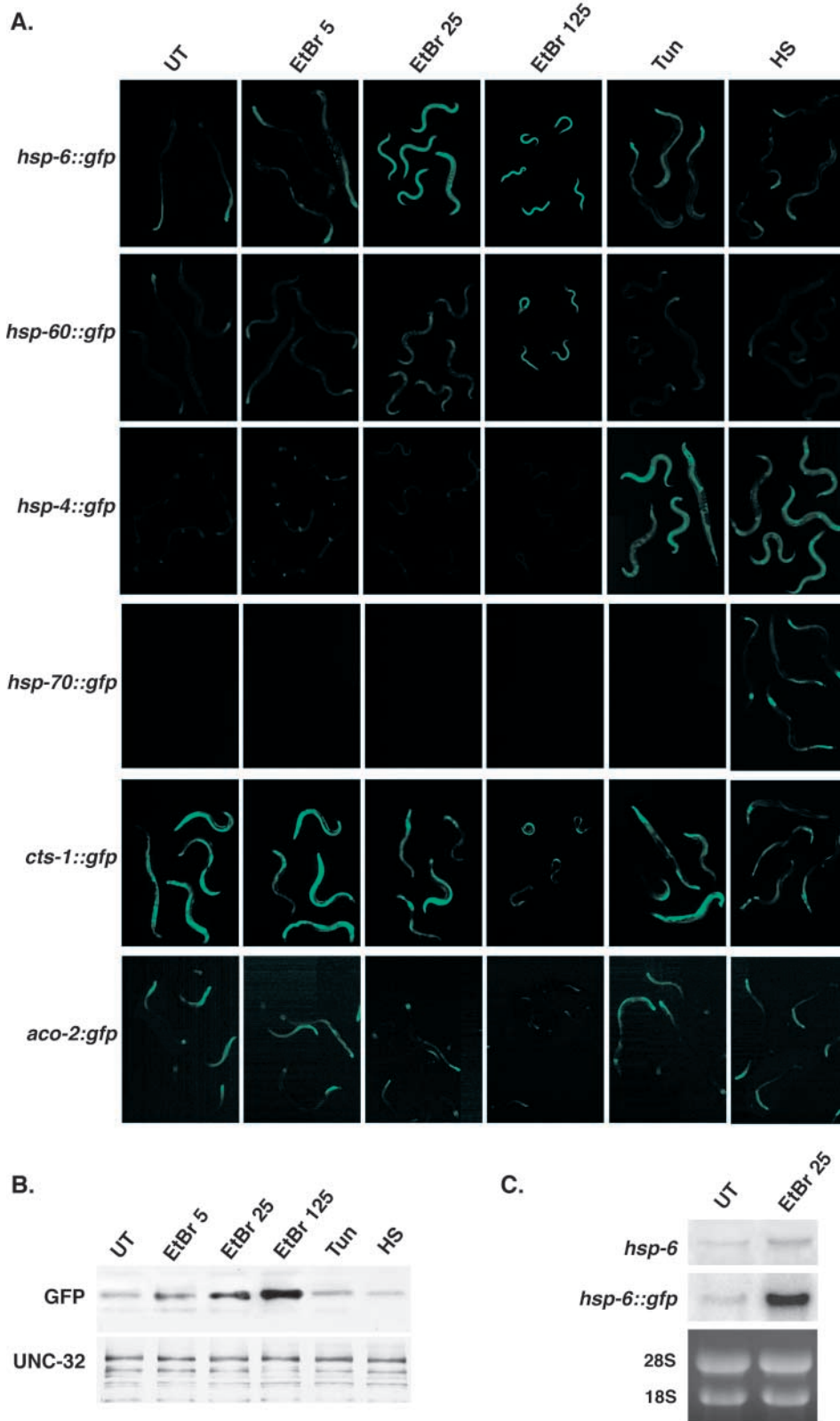


Fig. 2. Reporter genes activated by mitochondrial perturbation. (A) Fluorescence photomicrographs of untreated (UT), ethidium bromide (EtBr; $\mu\text{g/ml}$) or tunicamycin-treated (Tun; $1 \mu\text{g/ml}$) or heat-shocked (HS) animals with reporter transgenes for mitochondrial chaperones (*hsp-6::gfp*, *hsp-60::gfp*), an endoplasmic reticulum chaperone (*hsp-4::gfp*), a cytoplasmic chaperone (*hsp-70::gfp*), and the mitochondrial tri-carboxylic acid cycle enzymes citrate synthetase (*cts-1::gfp*) and aconitase (*aco-2::gfp*). (B) Immunoblot of soluble proteins extracted from the *hsp-6::gfp* animals described in A. The blot was reacted with anti-GFP serum (upper panel) or antiserum to the broadly expressed UNC-32 protein (lower panel). (C) Northern blot of untreated and ethidium bromide-treated *hsp-6::gfp* animals. The blot was hybridized sequentially with a *hsp-6*-coding region probe that detects the endogenous gene and a GFP probe that detects the transgene.

(Fig. 2A,B). A similar pattern of ethidium bromide inducibility was observed in animals transgenic for a mitochondrial chaperonin-encoding gene reporter, *hsp-60::gfp* (Fig. 2A, Fig. 5B). The transgenic reporters were much more sensitive to ethidium bromide than the endogenous genes (compare the levels of endogenous *hsp-6* mRNA to that of *hsp-6::gfp* mRNA in ethidium bromide-treated *hsp-6::gfp* transgenic animals, Fig. 2C). However, the reporter's specificity for mitochondrial perturbations was retained, as reflected in their indifference to tunicamycin or heat shock. By contrast, the *hsp-4::gfp* reporter, like the endogenous gene, was unaltered by ethidium bromide but was markedly induced by either tunicamycin or heat shock and the cytoplasmic *hsp-70::GFP* reporter was inducible only by heat shock (Fig. 1C, Fig. 2A). Furthermore, the induction of *hsp-6* and *hsp-60* was not shared by nuclear genes that encode enzymes active in the mitochondria, as neither the *cts-1::gfp*, which reports on the activity of *C. elegans* citrate synthase, nor the *aco-2::gfp*, which reports on the activity of mitochondrial aconitase, were activated by ethidium bromide (Fig. 2A). These last observations suggest that activation of the mitochondrial chaperones can occur independently of a general upregulation of nuclear genes encoding mitochondrial proteins.

We exploited the heightened sensitivity of the transgenic reporters to explore other perturbations predicted to selectively impede protein folding in the mitochondria. *C. elegans* is highly susceptible to disruption of gene expression by RNA-mediated interference (RNAi). Therefore we examined the impact of *hsp-6(RNAi)* and *hsp-60(RNAi)* on the activity of the *hsp-6::gfp* and *hsp-60::gfp* reporters. *hsp-6(RNAi)* and *hsp-60(RNAi)* animals had severe growth delay and many were arrested before hatching or at early larval stages. Nonetheless, these growth-arrested animals expressed high levels of the GFP reporter. By contrast, *hsp-6(RNAi)* and *hsp-60(RNAi)* did not induce *hsp-4::gfp*, cytoplasmic *hsp-70::gfp*, *cts-1::gfp* or *aco-2::gfp*. RNAi of the *C. elegans* homologues of the ER oxidase, *ero-1* or the ER protein disulfide isomerase, *pdi-3*, markedly activated *hsp-4::gfp* but did not affect *hsp-6::gfp* and *hsp-60::gfp* (Fig. 3).

We sought to perturb mitochondrial protein handling by inactivating a gene that might not exert as profound an impact on animal health and development as *hsp-60(RNAi)* or *hsp-6(RNAi)*. Paraplegin (or SPG7) is a nuclear encoded, mitochondrially localized, membrane-associated protease, mutations in which cause human spinal cord dysfunction (spastic paraplegia) (Casari et al., 1998). As the clinical syndrome associated with loss of function mutations in *SPG7* affects individuals that have otherwise developed normally, we hoped that inactivating a *C. elegans* homologue of *SPG7*, *spg-7*, might likewise have only a modest impact on animal development. Yeast homologues of *SPG7* are involved in proteolytic degradation of mitochondrial membrane proteins and play an important role in assembly of mitochondrial multi-protein complexes (Langer et al., 2001). Therefore, inactivation of the gene might promote accumulation of abnormal mitochondrial proteins, as interfering with protein degradation in other compartments increases the load of malformed proteins in a compartment-specific manner (Bence et al., 2001; Nishitoh et al., 2002).

To determine if *spg-7(RNAi)* affected protein processing in the mitochondria, we used a transgenic strain expressing GFP linked to a mitochondrial import signal in the body wall muscle

(*myo-3::mtGFP*) (Labrousse et al., 1999). Following import into the mitochondria mtGFP folds to a soluble protein that is identical in size to conventional GFP. In *E. coli* GFP folding is chaperone dependent (Wang et al., 2002), we therefore expected that the ability of the processed mtGFP to fold to its soluble form might also be chaperone dependent and report on the capacity of the mitochondria to chaperone matrix proteins. Immunoblot analysis of soluble and Triton insoluble fractions from untreated mtGFP and GFP transgenic animals showed that most of the mtGFP signal is in the soluble fraction and that this signal co-migrates on SDS-PAGE with conventional (cytoplasmic) GFP, which is mostly soluble too (Fig. 4A). The modest amount of insoluble mtGFP in extracts from untreated animals has slightly reduced mobility on the SDS-PAGE (lane 9) suggesting that it represents a precursor form. *spg-7(RNAi)* did not alter the solubility of conventional cytoplasmic GFP, however, it markedly increased the mtGFP signal in the insoluble fraction. These observations are consistent with an adverse affect of *spg-7(RNAi)* on the folding environment in mitochondria, which impairs the ability of the organelle to handle a model nuclear-encoded matrix protein. A similar increase in insoluble processed mtGFP was also observed in animals raised on medium containing ethidium bromide (Fig. 4A, lower panel). This last observation is consistent with the idea that impaired mitochondrial gene expression and an unbalanced load of unassembled components of multi-subunit complexes had taxed the organelle's capacity to process an imported model polypeptide such as GFP.

Ethidium bromide and *spg-7(RNAi)* specifically activated *hsp-6::gfp* and *hsp-60::gfp* but not *hsp-4::gfp* or cytoplasmic *hsp-70::gfp* (Fig. 3). Northern blot analysis showed that *spg-7(RNAi)* also activated the endogenous *hsp-6* gene but had no effect on the endogenous *aco-2* and *cit-1* genes that encode enzymes of the tri-carboxylic acid cycle or on the α -chain of the F1F0 ATP synthase or cytochrome c oxidase subunit IV-encoding genes, involved in oxidative phosphorylation (Fig. 4B). These observations suggest that the mitochondrial chaperones are not activated as part of a general increase in organelle biogenesis attendant upon the perturbation in protein handling, as biogenesis would be expected to increase the expression of genes involved in key mitochondrial metabolic activities and oxidative phosphorylation. As the *spg-7(RNAi)* animals were readily recovered in large quantities, it was also possible to confirm by northern blotting the expected decreased expression of *spg-7* mRNA (Fig. 4B).

The experiments described above revealed a link between perturbed mitochondrial protein handling and activation of genes encoding mitochondrial chaperones. However, it remained to be determined if any perturbation of mitochondrial function would activate the chaperone-encoding genes or if there were specificity in their response to perturbations affecting protein handling in the organelle. 2,4 dinitrophenol (DNP) uncouples oxidative phosphorylation in mammalian and *C. elegans* mitochondria (Murfitt et al., 1976). It does so by promoting a proton leak across the inner mitochondrial membrane, thus dissipating the electromotive chemical gradient ($\Delta\Psi^-$) required for ATP synthesis and aspects of mitochondrial import. Therefore, to determine if DNP treatment impacts on mitochondrial function in *C. elegans*, we exposed live untreated and DNP-treated animals to tetramethylrhodamine ethyl ester (TMRE), a lipophilic

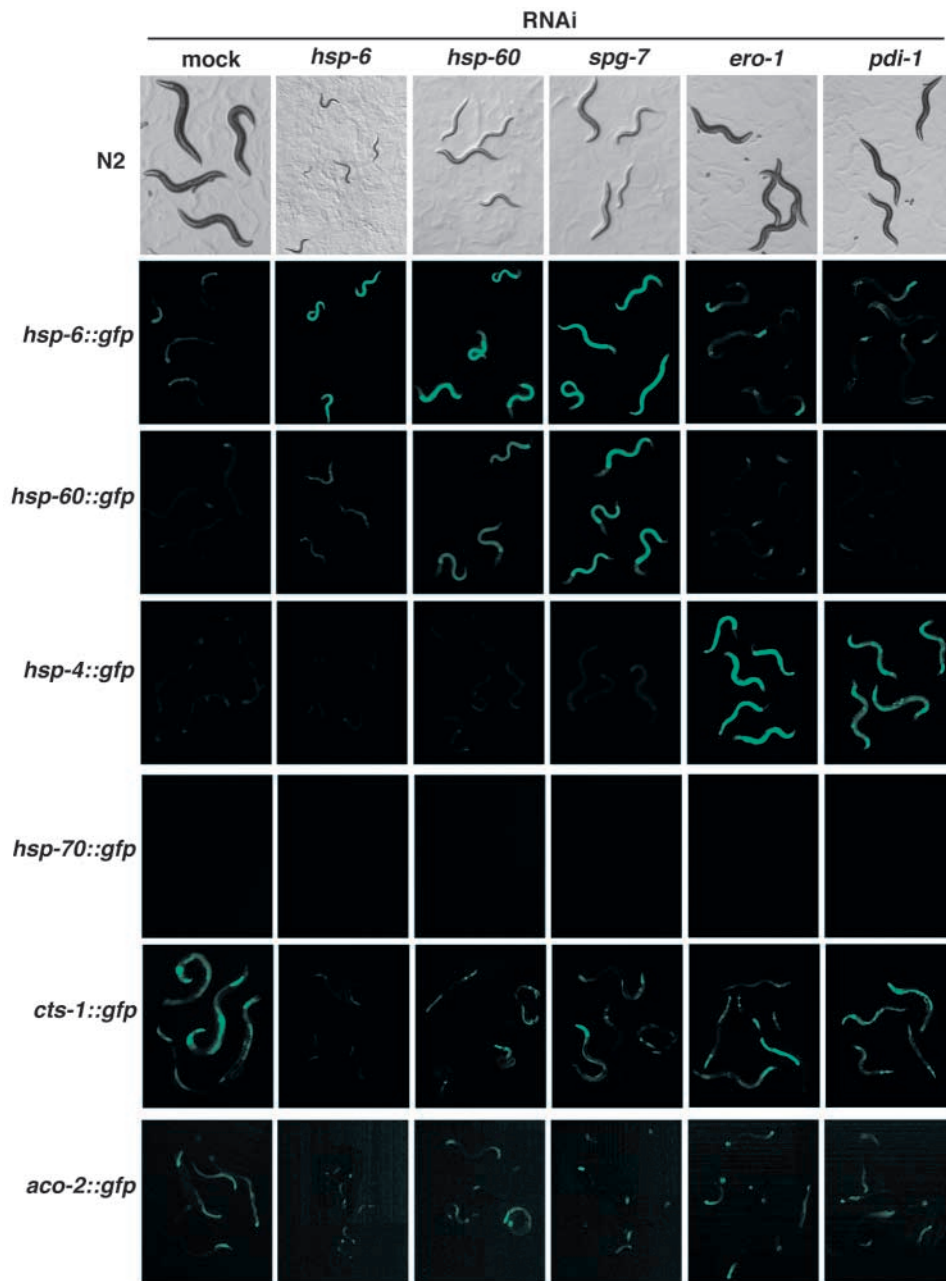


Fig. 3. Interference with genes that process mitochondrial proteins selectively activates mitochondrial chaperone genes. The top row of panels shows photomicrographs of wild-type (N2) animals exposed to the indicated RNAi; the panels below are photomicrographs of GFP fluorescence of *hsp-6::gfp*, *hsp-60::gfp*, *hsp-4::gfp*, *hsp-70::gfp*, *cts-1::gfp* and *aco-2::gfp* animals exposed to mock RNAi or RNAi of the indicated genes.

fluorescent cation whose mitochondrial uptake depends on $\Delta\Psi^-$ (Farkas et al., 1989; Loew et al., 1993).

As shown in Fig. 5A, *C. elegans* takes up TMRE from the culture medium and incorporates it broadly into its tissues. Animals exposed to DNP exhibited a dose-dependent decrease in TMRE fluorescence, consistent with dissipation of their mitochondrial proton gradient, $\Delta\Psi^-$. DNP-mediated decrease in TMRE fluorescence accrued slowly, over several days, indicating that direct interference with TMRE fluorescence by DNP is unlikely. Furthermore, DNP treatment also delayed development of the animals and reduced their fertility, indicating that exposure to this mitochondrial uncoupler has significant effects in vivo. As TMRE can also be taken up by other organelles, a more general ATP shortage affecting other electrochemical gradients may have contributed to reduced

TMRE fluorescence in DNP-treated animals. Despite evidence that DNP strongly impairs mitochondrial function in vivo, it did not activate mitochondrial chaperone-encoding genes (Fig. 5B,C).

Mitochondria are a major source of reactive oxygen species and perturbed mitochondrial function might further promote their accumulation by interfering with the normal flow of electrons to molecular oxygen (Shigenaga et al., 1994). To examine the potential role of reactive oxygen species in signaling the activation of the mitochondrial chaperones we compared the activation of the *hsp-60::gfp* reporter to that of *sod-3::gfp*, reporting on the manganese superoxide dismutase gene that is known to respond to accumulation of reactive oxygen species (Honda and Honda, 1999; Libina et al., 2003). RNAi of the mitochondrial chaperone *hsp-60*, the

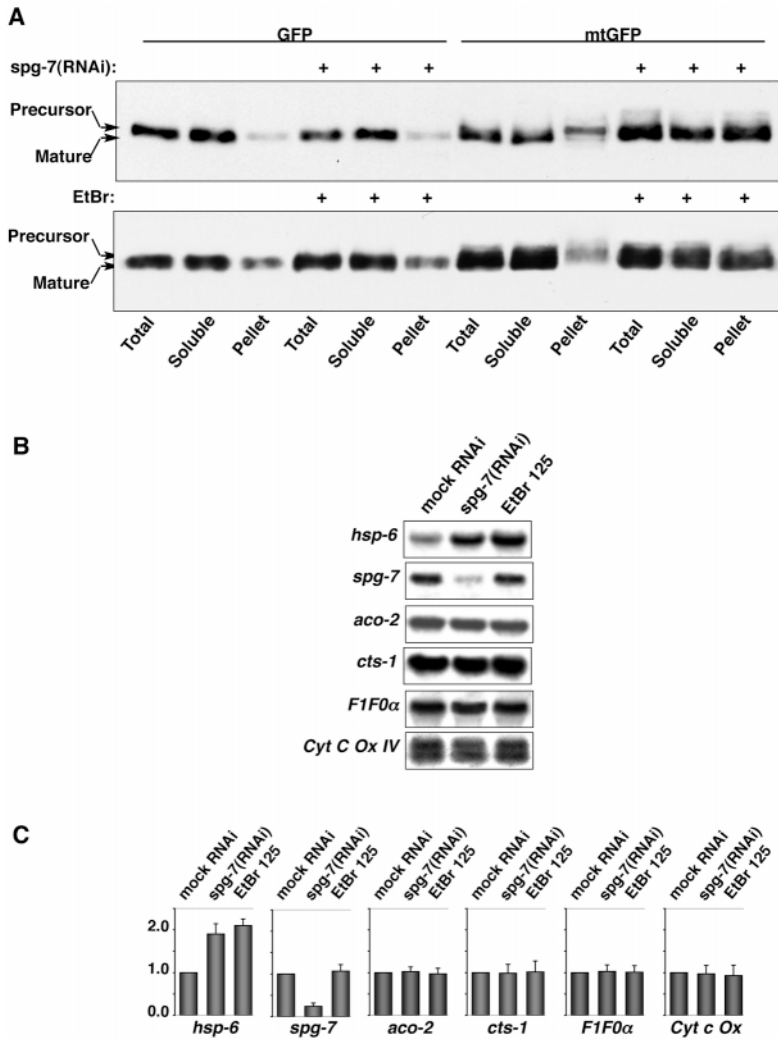


Fig. 4. Interference with *spg-7* (predicted to encode a mitochondrial protease) or ethidium bromide treatment perturbs protein processing in the mitochondria and activate the endogenous *hsp-6* gene. (A) Immunoblot of GFP in detergent extracts from transgenic animals expressing conventional GFP (a cytoplasmic protein) or GFP with an N-terminal mitochondrial import sequence (mtGFP), both in the body wall muscle. The animals were exposed to mock RNAi or *spg-1(RNAi)* or cultured on 125 µg/ml ethidium bromide (EtBr) as indicated. Equal fractions of the total extract (Total), 100,000 g soluble (Soluble) and 100,000 g pellet (Pellet) of the whole animal detergent extract were loaded on the gel as indicated.

(B) Autoradiogram of a northern blot of total RNA from mock-RNAi, *spg-7(RNAi)* or ethidium bromide (EtBr) treated animals. The blot was hybridized sequentially to radiolabeled fragments from the *hsp-6*, *spg-7*, *aco-2*, *cts-1*, F1F0 ATP synthase α -chain (F1F0 α) and cytochrome c oxidase subunit IV (Cyt C Ox IV) genes. (C) Quantification of the radiolabeled signal on the blots shown in B. The hybridization signal for each gene at the untreated point was set as 1.

mitochondrial protease *spg-7* or *phb-2* (the latter encodes a component of the prohibitin complex that is implicated in assembly of components of the respiratory chain (Artal-Sanz et al., 2003) all strongly activated the *hsp-60::gfp* reporter, as predicted. However, no activation of *sod-3::gfp* was noted (Fig. 5D). Exposure of animals to Paraquat, an agent that promotes accumulation of reactive oxygen species, potentially activated *sod-3::gfp*, serving as a positive control. Interestingly, Paraquat exposure also activated *hsp-60::gfp*, consistent with the damaging effect of reactive oxygen species on mitochondrial proteins. Conversely, exposure to ethidium bromide activated both *hsp-6::gfp* and *sod-3::gfp* (not shown). These experiments show that while conditions that promote expression of mitochondrial chaperones may lead to accumulation of reactive oxygen species, and accumulation of reactive oxygen species may promote the putative mitochondrial unfolded protein response, the two can be clearly uncoupled.

The observation cited above suggested that perturbations affecting the metabolic activity of mitochondria do not necessarily activate mitochondrial chaperone gene expression. To further examine the link between mitochondrial chaperone expression and perturbations affecting the folding environment in the organelle, we performed an unbiased screen for genes

whose loss of function activated mitochondrial chaperones in *C. elegans*. We sequentially exposed *hsp-6::gfp* and *hsp-60::gfp* animals to RNAi directed to all the genes on *C. elegans* chromosome I that were predicted to encode mitochondrial proteins and to all the 1160 predicted genes in a randomly selected interval on *C. elegans* chromosome I, between Y58G1C and *unc-13*. The latter component of the collection was included in an effort to uncover a potential class of genes whose RNAi might affect the expression of *hsp-6::gfp* and *hsp-60::gfp* but who are not annotated as encoding mitochondrial proteins.

RNAi of 32 predicted genes in this collection reproducibly activated the *hsp-6* and *hsp-60* reporter (Table 1). Activation was apparent from the earliest larval stages and was also observed in the embryo in some cases. Ten of these 32 genes encode proteins predicted to play a role in expression of the mitochondrial genome. These include seven mitochondrial ribosomal proteins, a mitochondrial tRNA synthetase, a mitochondrial translation elongation factor and a mitochondrial DNA helicase. The activation of the mitochondrial chaperones by RNAi of genes in this class is consistent with the observations made on animals raised on ethidium bromide, which also impairs mitochondrial genome

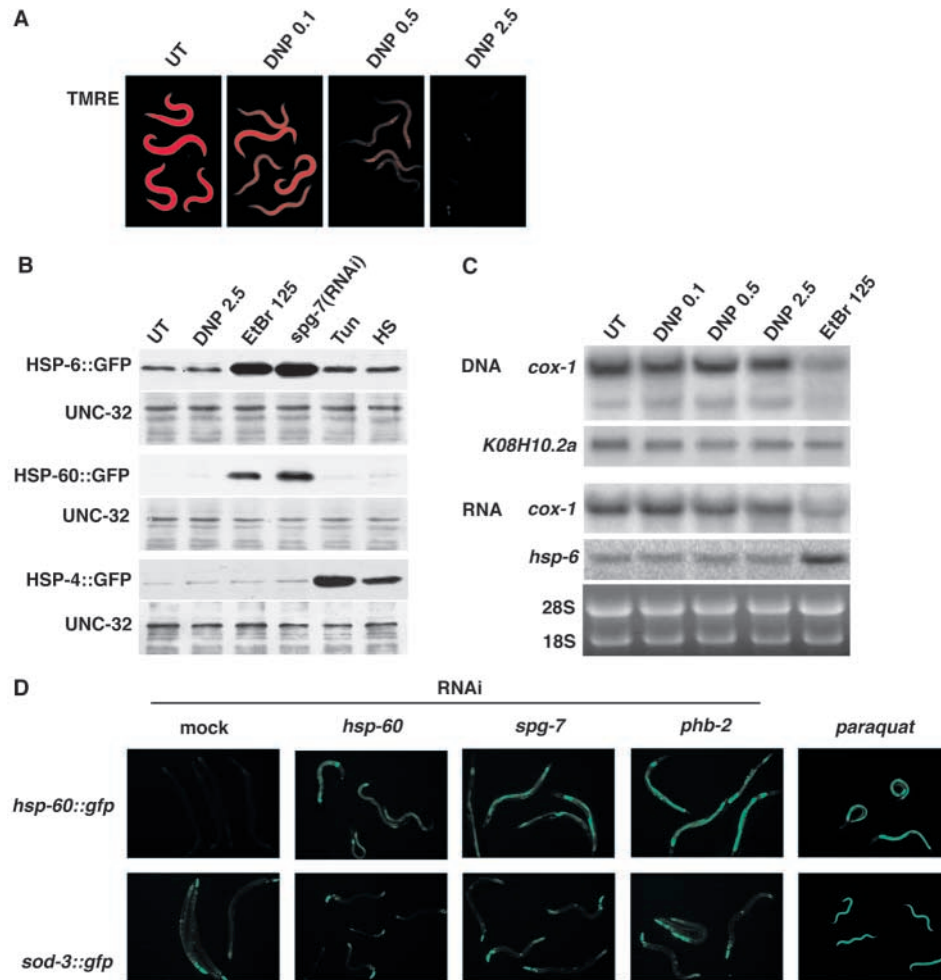


Fig. 5. Activation of mitochondrial chaperone genes can be uncoupled from depolarization of the mitochondrial membrane potential and the accumulation of reactive oxygen species. (A) Fluorescence photomicrographs of animals cultured on plates containing the indicated concentration of dinitrophenol (DNP, mM) and subsequently exposed to the fluorescent dye tetramethylrhodamine ethyl ester (TMRE). (B) Immunoblot of GFP (upper panel) or UNC-32 (lower panel) in lysates from untreated (UT), dinitrophenol (DNP; mM), ethidium bromide (EtBr, 125 μ g/ml), *spg-7(RNAi)*, tunicamycin (Tun; 1 μ g/ml) treated or heat shocked (HS) animals with *hsp-6::gfp*, *hsp-60::gfp* and *hsp-4::gfp* reporters. (C) Autoradiogram of a northern blot or Southern blot of total RNA or DNA from untreated animals (UT), dinitrophenol (DNP; mM) or ethidium bromide (EtBr, 125 μ g/ml) treatment. The blots were hybridized to a radiolabeled fragment from the *C. elegans* mitochondrially encoded *cox-1* gene or nuclear *hsp-6* or *K08H10.2a* genes. (D) Fluorescent photomicrographs of *hsp-60::gfp* and *sod-3::gfp* transgenic animals following RNAi of genes involved in mitochondrial protein processing or exposure to the toxin Paraquat (2 mM).

expression (Fig. 2). Two genes in this pool of activators are predicted to be involved in processing of mitochondrial proteins, one, a protease, SPG-7, has been described above and the second, a *C. elegans* homologue of yeast Oxal1p, is predicted to play a role in membrane insertion of mitochondrially encoded components of the respiratory apparatus. Sixteen activators encode mitochondrial proteins with other roles (oxidative phosphorylation, intermediary metabolism, protein import). Remarkably, 15 of these are subunits of multi-protein complexes, whose assembly requires precise stoichiometry of nuclear and/or mitochondrially encoded components. Only one of these 16 genes, that encoding cytochrome_c1, is not part of a heteromeric complex.

Therefore, 27 of the 32 genes whose loss of function activated the mitochondrial chaperones are predicted to function directly in protein metabolism or to have the potential to tax chaperone reserves by causing accumulation of unassembled subunits of mitochondrial complexes. Of the remaining five genes whose RNAi activated *hsp-6::gfp* and *hsp-60::gfp*, one, *rls-1*, encodes a protein related to transcriptional repressors of the polycomb group and whose RNAi non-specifically up-regulated other reporter transgenes (data not shown). A second gene encodes a DnaJ-like chaperone whose subcellular localization is not known. Two genes encode a pair of related predicted cytoplasmic ribosomal proteins whose RNAi may cross-react with mitochondrial

ribosomal protein genes, and the fifth gene encodes the aforementioned cytochrome_c1.

The genes in this collection whose RNAi did not activate the chaperones were also informative as 30 of them encode predicted mitochondrial proteins (Table 1). Remarkably, 24 of these 30 encode monomeric proteins or proteins that form homo-oligomeric complexes. Their inactivation is therefore not predicted to result in accumulation of unassembled subunits of mitochondrial complexes. Four of the remaining six genes do encode components of multimeric complexes, however it is not known if the RNAi procedure sufficiently diminished their expression. Similar considerations apply to a fifth gene, which encodes a predicted homologue of a mitochondrial Lon-like protease, whereas the oligomeric status of the sixth gene, encoding a homologue of Metataxin, is not known.

To further explore the role of impaired mitochondrial metabolism in activation of the chaperones, we inactivated, by RNAi, four genes that encode key monomeric or homo-oligomeric enzymes of the tri-carboxylic acid cycle. RNAi of all four genes resulted in significant embryonic lethality and arrested growth (as previously described) (Kamath et al., 2003). However the *hsp-6::gfp* or *hsp-60::gfp* reporter genes were either inactive [in the case of *aco-2(RNAi)* and *mdh-1(RNAi)*] or were activated very late in development of the RNAi animals [in the case of *fum-1(RNAi)* and *cts-1(RNAi)*] (Table 1). The associated RNAi phenotypes are unlikely to

Table 1. Activators and non-activators of the mitochondrial UPR

| Activators, Chr I | | | Non-activators, Chr I | | | | |
|-------------------|-----------------------------|--|--|-----|-----------|--|--------------------------|
| No. | Gene name | Description | Class | No. | Gene name | Description | Class |
| 1 | Y18H1A.6, <i>pf1-1</i> | Mitochondrial DNA helicase | Mitochondrial DNA maintenance | 1 | Y71F9B.9 | Similar to mitochondrial hydroxyacyl-coA dehydrogenase | Soluble, homodimer |
| 2 | Y47G6A.247, <i>f. sfg-7</i> | Mitochondrial protease | Membrane, proteolysis | 2 | C32E8.9 | Enoyl-coA hydratase | Soluble, homohexamer |
| 3 | R12E2.12 | Mitochondrial ribosomal protein S6 | Mitochondrial protein synthesis | 3 | T09B4.8 | Mitochondrial alanine-glyoxylate aminotransferase | Soluble, monomeric |
| 4 | C43E11.4 | Mitochondrial elongation factor EF-TU | Mitochondrial protein synthesis | 4 | T08B2.7 | Enoyl-coA hydratase | Soluble, homohexamer |
| 5 | ZK973.10 | NADH-coQ oxidoreductase subunit | Mitochondrial membrane, multimeric | 5 | K06A5.6 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 6 | D1007.6 | Cytosolic ribosomal protein (rps-10) | Cytoplasmic protein synthesis | 6 | C55B7.4 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 7 | D1007.12 | Cytosolic ribosomal protein (rpl-24) | Mitochondrial protein synthesis | 7 | T10E9.9 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 8 | B0261.4 | Mitochondrial ribosomal protein L47 | Mitochondrial protein synthesis | 8 | W02D3.6 | ATP-ADP transporter | Membrane, monomeric |
| 9 | T19B4.4 | Dnj-21, DNA homologue | Chaperone, unknown localization | 9 | W02D3.2 | Mitochondrial dihydroorotate dehydrogenase | Membrane, monomeric |
| 10 | T09B4.9 | Tim44 homologue | Mitochondrial membrane, multimeric | 10 | C37A2.3 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 11 | T08B2.8 | Mitochondrial ribosomal protein L23 | Mitochondrial protein synthesis | 11 | C30F12.7 | Isocitrate dehydrogenase subunit | Soluble, multimeric |
| 12 | T10E9.7 | NADH-coQ oxidoreductase subunit | Mitochondrial membrane, multimeric | 12 | F13G3.7 | Mitochondrial carrier | Membrane, monomeric |
| 13 | F22D6.4 | NADH-coQ oxidoreductase subunit | Mitochondrial membrane, multimeric | 13 | T25G3.4 | Mitochondrial glycerol 3-phosphate dehydrogenase homologue | Soluble, homodimer |
| 14 | W01A8.4 | NADH-coQ oxidoreductase subunit | Mitochondrial membrane, multimeric | 14 | F27D4.5 | Mitochondrial 2-oxoisovalerate dehydrogenase beta subunit | Soluble, multimeric |
| 15 | R06C7.7, <i>Rls-1</i> | Polycomb group | Transcription repressor | 15 | F30F8.2 | Mitochondrial glutaminase | Soluble, monomeric |
| 16 | F52A8.5 | Mitochondrial glycine cleavage system protein H | Mitochondrial, multimeric | 16 | F10D11.1 | Sod-2 | Soluble, homoletramers |
| 17 | C25A1.7, <i>Irs-2</i> | Mitochondrial isoleucyl-tRNA transferase | Mitochondrial protein synthesis | 17 | F43G9.3 | Mitochondrial carrier | Membrane, monomeric |
| 18 | D2030.4 | NADH-coQ oxidoreductase subunit | Mitochondrial membrane, multimeric | 18 | T05F1.8 | Mitochondrial carrier | Membrane, monomeric |
| 19 | F27D4.1 | Electron transfer flavoprotein α -subunit | Mitochondrial matrix, multimeric | 19 | C34B2.6 | Similar to ion proteases | Membrane, proteolysis |
| 20 | C54G4.8 | Cytochrome c1 | Mitochondrial membrane, multimeric | 20 | C34B2.7 | Succinate dehydrogenase complex subunit | Membrane, multimeric |
| 21 | F43G9.1 | Isocitrate dehydrogenase subunit | Mitochondrial, multimeric | 21 | T26E3.7 | ATP synthase alpha chain | Membrane, multimeric |
| 22 | K07A12.3 | ATP synthase subunit gamma | Mitochondrial membrane, multimeric | 22 | F39B2.11 | Metaxin homologue | Membrane, protein import |
| 23 | F25H5.6 | Mitochondrial ribosomal subunit L54 | Mitochondrial protein synthesis | 23 | Y71F9B.9 | Similar to mitochondrial hydroxyacyl-coA dehydrogenase | Soluble, homodimer |
| 24 | F26E4.6 | Cytochrome c oxidase subunit | Mitochondrial membrane, multimeric | 24 | C32E8.9 | Enoyl-coA hydratase | Soluble, homohexamer |
| 25 | F26E4.9 | Cytochrome c oxidase subunit | Mitochondrial membrane, multimeric | 25 | T09B4.8 | Mitochondrial alanine-glyoxylate aminotransferase | Soluble, monomeric |
| 26 | F59C6.5 | NADH-coQ oxidoreductase subunit | Mitochondrial membrane, multimeric | 26 | T08B2.7 | Enoyl-coA hydratase | Soluble, homohexamer |
| 27 | B0511.8 | Mitochondrial ribosomal protein S30 | Mitochondrial protein synthesis | 27 | K06A5.6 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 28 | H28Q16.1 | ATP synthase α -chain (F1F0 α) | Mitochondrial membrane, multimeric | 28 | C55B7.4 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 29 | C01A2.3 | Cytochrome oxidase biogenesis:oxal p homologue | Mitochondrial membrane, protein processing | 29 | T10E9.9 | Acyl-coA dehydrogenase homologue | Soluble, homoletramers |
| 30 | W09C5.8 | Cytochrome c oxidase subunit IV | Mitochondrial membrane, multimeric | 30 | W02D3.6 | ATP-ADP transporter | Membrane, monomeric |
| 31 | Y92H12BR.8 | Mitochondrial ribosomal protein L15 | Mitochondrial protein synthesis | | | | |
| 32 | K11B4.1 | Mitochondrial ribosomal protein S27 | Mitochondrial protein synthesis | | | | |

| Chr III | | | |
|---------|-------------------------|------------------------------------|------------------------------|
| No. | Gene name | Description | Class |
| 1 | F54h12.1a, <i>aco-2</i> | Mitochondrial acornitase | Soluble, monomeric |
| 2 | F20H11.3, <i>mdh-1</i> | Mitochondrial malate dehydrogenase | Soluble, homodimer/letramers |
| 3 | H14A12.2a, <i>fum-1</i> | Mitochondrial fumarase | Soluble, homoletramers |
| 4 | T20G5.2, <i>cit-1</i> | Mitochondrial citrate synthase | Soluble, monomeric |

Activators: Genes on chromosome I in the interval between Y48G1C and *umc-13* whose RNAi activated *hsp-6::gfp* and *hsp-60::gfp*.

Non-activators: Genes encoding predicted mitochondrial protein, localized to the above interval that did not activate *hsp-6::gfp* and *hsp-60::gfp*. Also listed under non-activators are four genes from chromosome III who encode enzymes of the tri-carboxylic acid cycle and whose RNAi presumably results in a severe disruption of mitochondrial metabolism reflected in a high incidence of larval arrest and embryonic lethality (Kamath et al., 2003) but whose effect on *hsp-6::gfp* and *hsp-60::gfp* is confined to late induction in adult RNAi animals (in the case of *aco-2* and *cit-1*).

Description: Predicted protein function.

Class: Hypothesized mechanism by which the RNAi procedure perturbs (or fails to perturb) protein processing in the mitochondria (see text for further discussion).

References for this Table can be found in the supplemental data (see Table S1; <http://jes.biologists.org/supplemental/>).

have obscured reporter activation, as many of the activators listed in Table 1 had similar severe RNAi phenotypes and these did not interfere with detection of reporter activity in early embryonic or larval stages (also see Fig. 2A). Collectively these data indicate that mitochondrial chaperones are activated by manipulations predicted to tax the organelle's protein handling machinery, but not by manipulations that impede its metabolic activity.

Discussion

The lack of tools to specifically manipulate the folding environment in the mitochondria had restricted the ability to study mitochondrial chaperone gene expression. In this study we took advantage of the ease with which gene function can be manipulated in *C. elegans* to show that reduced function of genes encoding chaperones and proteases specifically involved in protein processing in the mitochondria selectively activates genes encoding two major mitochondrial chaperones. Using sensitive reporter genes we found that mitochondrial chaperones are upregulated transcriptionally in response to perturbations that affect protein processing in the mitochondria. The robustness of the responses we observed and their specificity strongly suggests the existence of a mitochondrial unfolded protein response. It is important, however, to emphasize that we have not proved that signaling in the pathway culminating in *hsp-6* and *hsp-60* expression initiates with direct recognition of unfolded proteins or recognition of reduced capacity to handle proteins in the organelle; this remains a working hypothesis to be tested as the components of the pathway are revealed by future work.

In this study, we have cast the problem of regulating mitochondrial chaperone-encoding genes in the context of the need to respond to compartment-specific changes in protein folding capacity and chaperone occupancy; in other words in the context of an unfolded protein response. However, the problem also needs to be considered in the broader context of the communication that must take place between the mitochondrial and the nuclear genome. Inactivation of genes involved directly in protein processing by the mitochondria gives rise to a very early and strong activation of the chaperone reporter genes. However, impaired protein folding in the mitochondria will also affect organelle function. Therefore, it is informative that manipulations primarily predicted to degrade mitochondrial energy production, such as dissipation of the mitochondrial inner membrane proton gradient with DNP or RNAi of genes encoding enzymes of the tri-carboxylic acid cycle (which interrupts the supply of electrons for oxidative phosphorylation), induced the chaperone reporters only minimally, or resulted in markedly delayed activation. Furthermore, manipulations that impair protein handling in the organelle selectively activate chaperone genes as these do not result in measurable upregulation of genes encoding mitochondrial enzymes (Fig. 2A and Fig. 4B). Together these observations suggest that genes encoding mitochondrial chaperones are responsive to a signaling pathway(s) that is selectively activated by perturbed protein folding and processing in the organelle rather than being activated by pathways primarily responsive to the metabolic consequences of mitochondrial dysfunction.

Previous studies have addressed the role of transcriptional

control in cellular adaptation to mitochondrial dysfunction (Liao and Butow, 1993; Amuthan et al., 2001) and have begun to shed light on mechanisms that coordinately upregulate nuclear genes encoding mitochondrial proteins (Wu et al., 1999; Lin et al., 2002). As mitochondrial biogenesis entails expression of many nuclear-encoded genes whose products must be processed and folded by mitochondrial chaperones, one would expect activation of nuclear-encoded mitochondrial chaperones to accompany mitochondrial biogenesis, as has been described in mammalian adipocytes (Wilson-Fritch et al., 2003). It is tempting to speculate that the delayed induction of *hsp-6::gfp* and *hsp-60::gfp* observed in the *cts-1(RNAi)* and *aco-2(RNAi)* (data not shown) reflects such an indirect response to the metabolic consequences of mitochondrial dysfunction. Furthermore, Lemire and colleagues have emphasized the physiological increase in mitochondrial mass during *C. elegans* germline development (Tsang and Lemire, 2002). It is interesting in this regard to note that activation of the *hsp-6::gfp* and *hsp-60::gfp* reporter by *spg-7(RNAi)* or ethidium bromide treatment was most conspicuous in L4 larvae and in young adults, stages of development associated with expansion of the germline. This observation is consistent with crosstalk between the pathways that control mitochondrial biogenesis and the hypothesized mitochondrial UPR. Our study did not further address this issue as we have not identified conditions that selectively manipulate mitochondrial biogenesis in *C. elegans*.

It is also important to emphasize the limitations inherent in the use of pharmacological and genetic criteria to distinguish between signals initiated by altered metabolism and altered protein handling by the mitochondria. The potential for diminished chaperone reserve to secondarily affect metabolic function has been discussed above. But altered metabolism may also affect protein handling by the organelle. For example, dissipation of the inner membrane proton gradient (by DNP) impairs mitochondrial import (Fig. 5A), which is likely to have pleiotropic effects on mitochondrial function. Perhaps the reduced expression of mitochondrial genes in animals exposed to high dose DNP (Fig. 5C) offsets the predicted imbalance between imported and mitochondrially encoded components, explaining why chaperones are not induced in DNP-treated animals.

Another variable addressed by our study is the role of reactive oxygen species in activating mitochondrial chaperone-encoding genes. Interactions between these two processes are revealed by the observations that both Paraquat, which leads to accumulation of massive amount of reactive oxygen species and ethidium bromide, which primarily affects the integrity of the mitochondrial genome, activate *hsp-60::gfp* and the oxidative stress inducible reporter *sod-3::gfp* (Fig. 5D and data not shown). However, the two responses are uncoupled by milder manipulations that primarily affect protein processing in the mitochondria. Thus, our data do not support an essential role for reactive oxygen species in signaling the putative mitochondrial unfolded protein response.

The severity of the *hsp-6(RNAi)* and *hsp-60(RNAi)* phenotype is consistent with the essential role of HSP60 and mtHSP70 in yeast (Cheng et al., 1989; Craig et al., 1989). More surprising is the relative resistance of *C. elegans* to degradation of mitochondrial function by culture in ethidium bromide or DNP. Though brood size was markedly diminished, consistent

with the established effect of mitochondrial dysfunction on germline development (Tsang et al., 2001; Tsang and Lemire, 2002), the treated animals remained viable and surprisingly mobile under conditions in which much of their mitochondrial DNA had been lost (Fig. 1) or in face of evidence for loss of mitochondrial membrane potential (Fig. 5). This tolerance of nematodes for impaired mitochondrial function suggests that it may be possible to recover animals with mutations affecting the hypothesized mitochondrial UPR.

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References

- Amuthan, G., Biswas, G., Zhang, S. Y., Klein-Szanto, A., Vijayarathay, C. and Avadhani, N. G. (2001). Mitochondria-to-nucleus stress signaling induces phenotypic changes, tumor progression and cell invasion. *EMBO J.* **20**, 1910-1920.
- Artal-Sanz, M., Tsang, W. Y., Willems, E. M., Grivell, L. A., Lemire, B. D., van der Spek, H., Nijtmans, L. G. and Sanz, M. A. (2003). The mitochondrial prohibitin complex is essential for embryonic viability and germline function in *Caenorhabditis elegans*. *J. Biol. Chem.* **278**, 32091-32099.
- Bence, N. F., Sampat, R. M. and Kopito, R. R. (2001). Impairment of the ubiquitin-proteasome system by protein aggregation. *Science* **292**, 1552-1555.
- Bertolotti, A., Zhang, Y., Hendershot, L., Harding, H. and Ron, D. (2000). Dynamic interaction of BiP and the ER stress transducers in the unfolded protein response. *Nat. Cell. Biol.* **2**, 326-332.
- Biaszczak, A., Georgopoulos, C. and Liberek, K. (1999). On the mechanism of FtsH-dependent degradation of the sigma 32 transcriptional regulator of *Escherichia coli* and the role of the Dnak chaperone machine. *Mol. Microbiol.* **31**, 157-166.
- Bukau, B. (1993). Regulation of the *Escherichia coli* heat-shock response. *Mol. Microbiol.* **9**, 671-680.
- Bukau, B. and Horwich, A. L. (1998). The Hsp70 and Hsp60 chaperone machines. *Cell* **92**, 351-366.
- Calfon, M., Zeng, H., Urano, F., Till, J. H., Hubbard, S. R., Harding, H. P., Clark, S. G. and Ron, D. (2002). IRE1 couples endoplasmic reticulum load to secretory capacity by processing the *XBP-1* mRNA. *Nature* **415**, 92-96.
- Casari, G., de Fusco, M., Ciarmatori, S., Zeviani, M., Mora, M., Fernandez, P., de Michele, G., Filla, A., Coccozza, S., Marconi, R. et al. (1998). Spastic paraplegia and OXPHOS impairment caused by mutations in paraplegin, a nuclear-encoded mitochondrial metalloprotease. *Cell* **93**, 973-983.
- Cheng, M. Y., Hartl, F. U., Martin, J., Pollock, R. A., Kalousek, F., Neupert, W., Hallberg, E. M., Hallberg, R. L. and Horwich, A. L. (1989). Mitochondrial heat-shock protein hsp60 is essential for assembly of proteins imported into yeast mitochondria. *Nature* **337**, 620-625.
- Craig, E. A., Kramer, J., Shilling, J., Werner-Washburne, M., Holmes, S., Kosic-Smithers, J. and Nicolet, C. M. (1989). SSC1, an essential member of the yeast HSP70 multigene family, encodes a mitochondrial protein. *Mol. Cell. Biol.* **9**, 3000-3008.
- Farkas, D. L., Wei, M. D., Febroriello, P., Carson, J. H. and Loew, L. M. (1989). Simultaneous imaging of cell and mitochondrial membrane potentials. *Biophys. J.* **56**, 1053-1069.
- Fewell, S. W., Travers, K. J., Weissman, J. S. and Brodsky, J. L. (2001). The action of molecular chaperones in the early secretory pathway. *Annu. Rev. Genet.* **35**, 149-191.
- Fraser, A. G., Kamath, R. S., Zipperlen, P., Martinez-Campos, M., Sohrmann, M. and Ahringer, J. (2000). Functional genomic analysis of *C. elegans* chromosome I by systematic RNA interference. *Nature* **408**, 325-330.
- Gething, M. J. and Sambrook, J. (1992). Protein folding in the cell. *Nature* **355**, 33-45.
- Harding, H., Zhang, Y. and Ron, D. (1999). Translation and protein folding are coupled by an endoplasmic reticulum resident kinase. *Nature* **397**, 271-274.
- Harding, H. P., Calfon, M., Urano, F., Novoa, I. and Ron, D. (2002). Transcriptional and translational control in the mammalian unfolded protein response. *Annu. Rev. Cell. Dev. Biol.* **18**, 575-599.
- Hartl, F. U. and Hayer-Hartl, M. (2002). Molecular chaperones in the cytosol: from nascent chain to folded protein. *Science* **295**, 1852-1858.
- Hayashi, J., Tanaka, M., Sato, W., Ozawa, T., Yonekawa, H., Kagawa, Y. and Ohta, S. (1990). Effects of ethidium bromide treatment of mouse cells on expression and assembly of nuclear-coded subunits of complexes involved in the oxidative phosphorylation. *Biochem. Biophys. Res. Commun.* **167**, 216-221.
- 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.
- Horwich, A. L. and Willison, K. R. (1993). Protein folding in the cell: functions of two families of molecular chaperone, hsp 60 and TF55-TCPI. *Philos. Trans. R. Soc. London B. Biol. Sci.* **339**, 313-325.
- Horwich, A. L., Weber-Ban, E. U. and Finley, D. (1999). Chaperone rings in protein folding and degradation. *Proc. Natl. Acad. Sci. USA* **96**, 11033-11040.
- Kamath, R. S., Fraser, A. G., Dong, Y., Poulin, G., Durbin, R., Gotta, M., Kanapin, A., le Bot, N., Moreno, S., Sohrmann, M. et al. (2003). Systematic functional analysis of the *Caenorhabditis elegans* genome using RNAi. *Nature* **421**, 231-237.
- Kaufman, R. J. (1999). Stress signaling from the lumen of the endoplasmic reticulum: coordination of gene transcriptional and translational controls. *Genes Dev.* **13**, 1211-1233.
- King, M. P. and Attardi, G. (1989). Human cells lacking mtDNA: repopulation with exogenous mitochondria by complementation. *Science* **246**, 500-503.
- Labrousse, A. M., Zappaterra, M. D., Rube, D. A. and van der Blik, A. M. (1999). *C. elegans* dynamin-related protein DRP-1 controls severing of the mitochondrial outer membrane. *Mol. Cell* **4**, 815-826.
- Langer, T., Kaser, M., Klanner, C. and Leonhard, K. (2001). AAA proteases of mitochondria: quality control of membrane proteins and regulatory functions during mitochondrial biogenesis. *Biochem. Soc. Trans.* **29**, 431-436.
- Liao, X. and Butow, R. A. (1993). RTG1 and RTG2: two yeast genes required for a novel path of communication from mitochondria to the nucleus. *Cell* **72**, 61-71.
- Libina, N., Berman, J. R. and Kenyon, C. (2003). Tissue-specific activities of *C. elegans* DAF-16 in the regulation of lifespan. *Cell* **115**, 489-502.
- Lin, J., Wu, H., Tarr, P. T., Zhang, C. Y., Wu, Z., Boss, O., Michael, L. F., Puigserver, P., Isotani, E., Olson, E. N. et al. (2002). Transcriptional co-activator PGC-1 alpha drives the formation of slow-twitch muscle fibres. *Nature* **418**, 797-801.
- Lindquist, S. and Craig, E. A. (1988). The heat-shock proteins. *Annu. Rev. Genet.* **22**, 631-677.
- Liu, R., Corry, P. and Lee, Y. (1994). Regulation of chemical stress-induced hsp70 gene expression in murine L929 cells. *J. Cell Sci.* **107**, 2209-2214.
- Liu, C. Y., Wong, H. N., Schauerer, J. A. and Kaufman, R. J. (2002). The protein kinase/endonuclease IRE1a that signals the unfolded protein response has a luminal amino-terminal ligand-independent dimerization domain. *J. Biol. Chem.* **277**, 18346-18356.
- Loew, L. M., Tuft, R. A., Carrington, W. and Fay, F. S. (1993). Imaging in five dimensions: time-dependent membrane potentials in individual mitochondria. *Biophys. J.* **65**, 2396-2407.
- Martin, J. (1997). Molecular chaperones and mitochondrial protein folding. *J. Bioenerg. Biomembr.* **29**, 35-43.
- Morimoto, R. I. (1998). Regulation of the heat shock transcriptional response: cross talk between a family of heat shock factors, molecular chaperones, and negative regulators. *Genes Dev.* **12**, 3788-3796.
- Mosser, D. D., Duchaine, J. and Massie, B. (1993). The DNA-binding activity of the human heat shock transcription factor is regulated in vivo by hsp70. *Mol. Cell. Biol.* **13**, 5427-5438.
- Murfitt, R. R., Vogel, K. and Sanadi, D. R. (1976). Characterization of the mitochondria of the free-living nematode, *Caenorhabditis elegans*. *Comp. Biochem. Physiol. B* **53**, 423-430.
- Neupert, W. (1997). Protein import into mitochondria. *Annu. Rev. Biochem.* **66**, 863-917.

- Neupert, W. and Pfanner, N.** (1993). Roles of molecular chaperones in protein targeting to mitochondria. *Philos. Trans. R. Soc. London B. Biol. Sci.* **339**, 355-361.
- Nishitoh, H., Matsuzawa, A., Tobiume, K., Saegusa, K., Takeda, K., Inoue, K., Hori, S., Ishikawa, K., Mizusawa, H., Kakizuka, A. et al.** (2002). ASK1 is essential for endoplasmic reticulum stress-induced neuronal cell death triggered by expanded polyglutamine repeats. *Genes Dev.* **16**, 1345-1355.
- Okamura, K., Kimata, Y., Higashio, H., Tsuru, A. and Kohno, K.** (2000). Dissociation of Kar2p/BiP from an ER sensory molecule, Ire1p, triggers the unfolded protein response in yeast. *Biochem. Biophys. Res. Commun.* **279**, 445-450.
- Patil, C. and Walter, P.** (2001). Intracellular signaling from the endoplasmic reticulum to the nucleus: the unfolded protein response in yeast and mammals. *Curr. Opin. Cell Biol.* **13**, 349-355.
- Shen, J., Chen, X., Hendershot, L. and Prywes, R.** (2002). ER stress regulation of ATF6 localization by dissociation of BiP/GRP78 and unmasking of Golgi localization signals. *Dev. Cell* **3**, 99-111.
- Shigenaga, M. K., Hagen, T. M. and Ames, B. N.** (1994). Oxidative damage and mitochondrial decay in aging. *Proc. Natl. Acad. Sci. USA* **91**, 10771-10778.
- Stuart, R. A. and Neupert, W.** (1996). Topogenesis of inner membrane proteins of mitochondria. *Trends Biochem. Sci.* **21**, 261-267.
- Timmons, L., Court, D. L. and Fire, A.** (2001). Ingestion of bacterially expressed dsRNAs can produce specific and potent genetic interference in *Caenorhabditis elegans*. *Gene* **263**, 103-112.
- Tsang, W. Y. and Lemire, B. D.** (2002). Mitochondrial genome content is regulated during nematode development. *Biochem. Biophys. Res. Commun.* **291**, 8-16.
- Tsang, W. Y., Sayles, L. C., Grad, L. I., Pilgrim, D. B. and Lemire, B. D.** (2001). Mitochondrial respiratory chain deficiency in *Caenorhabditis elegans* results in developmental arrest and increased life span. *J. Biol. Chem.* **276**, 32240-32246.
- Wang, J. D., Herman, C., Tipton, K. A., Gross, C. A. and Weissman, J. S.** (2002). Directed evolution of substrate-optimized GroEL/S chaperonins. *Cell* **111**, 1027-1039.
- Wilson-Fritch, L., Burkart, A., Bell, G., Mendelson, K., Leszyk, J., Nicoloro, S., Czech, M. and Corvera, S.** (2003). Mitochondrial biogenesis and remodeling during adipogenesis and in response to the insulin sensitizer rosiglitazone. *Mol. Cell. Biol.* **23**, 1085-1094.
- Wu, Z., Puigserver, P., Andersson, U., Zhang, C., Adelmant, G., Mootha, V., Troy, A., Cinti, S., Lowell, B., Scarpulla, R. C. et al.** (1999). Mechanisms controlling mitochondrial biogenesis and respiration through the thermogenic coactivator PGC-1. *Cell* **98**, 115-124.
- Zhao, Q., Wang, J., Levichkin, I. V., Stasinopoulos, S., Ryan, M. T. and Hoogenraad, N. J.** (2002). A mitochondrial specific stress response in mammalian cells. *EMBO J.* **21**, 4411-4419.