Cells employ a vast network of regulatory pathways to manage intracellular levels of reactive oxygen species (ROS). An effective means used by cells to control these regulatory systems are sulfur-based redox switches, which consist of protein cysteine or methionine residues that become transiently oxidized when intracellular ROS levels increase. Here, we describe a methionine-based oxidation event involving the yeast cytoplasmic Hsp70 co-chaperone Fes1. We show that Fes1 undergoes reversible methionine oxidation during excessively-oxidizing cellular conditions, and we map the site of this oxidation to a cluster of three methionine residues in the Fes1 core domain. Making use of recombinant proteins and a variety of in vitro assays, we establish that oxidation inhibits Fes1 activity and, correspondingly, alters Hsp70 activity. Moreover, we demonstrate in vitro and in cells that Fes1 oxidation is irreversible and is regulated by the cytosolic methionine sulfoxide reductase Mxr1 (MsrA) and a previously unidentified cytosolic pool of the reductase Mxr2 (MsrB). We speculate that inactivation of Fes1 activity during excessively-oxidizing conditions may help maintain protein-folding homeostasis in a suboptimal cellular folding environment. The characterization of Fes1 oxidation during cellular stress provides a new perspective as to how the activities of the cytosolic Hsp70 chaperones may be attuned by fluctuations in cellular ROS levels and provides further insight into how cells use methionine-based redox switches to sense and respond to oxidative stress.

Oxidative stress is associated with an intracellular accumulation of high levels of reactive oxygen species (ROS). ROS are widely appreciated as damage-inducing agents, based on their potential to irreversibly oxidize biological macromolecules, including lipids, nucleic acids, and proteins. Macromolecule oxidation as a consequence of excessive levels of cellular ROS can lead to a loss of vital cell functions and a deterioration of essential cellular processes. Historically, it has been generally accepted that decreasing or detoxifying ROS is key to limiting the cellular damage associated with oxidative stress. However, more recently, ROS have emerged also as signaling molecules, which can help cells initiate scavenging, protection, and repair pathways during oxidative stress (1). This conceptual shift suggests that preserving cell function during oxidative stress likely depends both on limiting ROS-induced damage and maintaining beneficial signaling events mediated by ROS.

Many of the identified ROS-based signaling pathways employ the use of labile, reversible redox-active post-translational protein modifications (PTMs) (2–7). Akin to the more established PTM phosphorylation, protein oxidation can reversibly alter the conformation and/or activity of a given protein. Altered protein activity can allow for the initiation of distinct downstream signaling events to benefit cells during oxidative stress, including the activation of pathways that serve to scavenge ROS before they exert detrimental effects as well as systems that act to manage and/or repair ROS incurred damage (2–7).

Reversible oxidation of methionine side chains is one example of a redox-based PTM that can elicit beneficial cellular outcomes during oxidative stress (8, 9). Methionine modification by ROS occurs upon oxidation of the sulfur atom in the methionine side chain; addition of an oxygen molecule (from the ROS) to the sulfur generates a structure termed methionine sulfoxide (MetO). In eukaryotic cells, MetO adducts are reduced (reversed) by methionine sulfoxide reductase (Msr) enzymes (10). Advances in mass spectrometry (MS) and chemical biology tools have led to the identification of many proteins with methionine residues that are susceptible to oxidation (11, 12). Yet relative to other post-translational modifications, we are just beginning to uncover the impact of methionine oxidation on the function of individual proteins. Determining how oxidation influences protein function is an essential first step to reveal how modification may affect cell function during oxidative stress conditions.

Here, we report the susceptibility of, and consequences for, oxidation of three methionine residues in the yeast Hsp70 nucleotide-exchange factor (NEF) Fes1. Members of the heat-shock protein 70 (Hsp70) class of molecular chaperones play critical roles in maintaining protein homeostasis during peri-
Fes1 is post-translationally modified in cells by ROS

We observed a striking change in the electrophoretic migration of Fes1 when cell lysates were prepared from yeast exposed to ROS. The most prominent change in Fes1 electrophoretic mobility was seen upon treatment of cells with sodium hypochlorite (NaOCl), a strong oxidant, classified as both a ROS and reactive chloride species (Fig. 1A). The extent of the mobility change in Fes1 intensified as the concentration of NaOCl in the medium was increased. Treatment with 0.5 mM NaOCl for 30 min resulted in several slower migrating Fes1 species (compared with untreated cells), and a single slow-migrating band was recovered in the presence of 1 mM NaOCl (Fig. 1A). Treatment of cells with a range of peroxides also resulted in changes to Fes1 mobility. Treatment with the organic peroxide cumene hydroperoxide (CHP) for 30 min. Fes1 was detected by Western blotting with anti-FLAG antibody. Western images are representative of the data obtained from three independent experiments.

Results

Figure 1. Oxidation of Fes1 methionines occurs in cells exposed to exogenous ROS. Lysates were prepared from WT yeast containing a plasmid encoding Fes1–FLAG after a 30-min treatment with peroxide (5 or 10 mM) or NaOCl (0.5 or 1 mM) (A) or after exposure to elevated heat (37 °C, 1 h), cycloheximide (CHX) (10 or 50 μg/ml, 30 min), DTT (5 or 10 mM, 30 min), or diamide (5 or 10 mM, 30 min) (B). Lysates were prepared by SDS-PAGE, and Fes1 was detected by Western blotting with anti-FLAG antibody. Mobility-shifted Fes1 corresponds to oxidized protein (see text). C, Fes1 methionine mutants complement temperature-sensitive growth of a fes1Δ cells containing CEN plasmids encoding WT or methionine mutant Fes1–FLAG alleles and an HSE-lacZ reporter were cultured at 30 °C to log phase in minimal medium, and cultures were either maintained at 30 °C or shifted to 37 °C for 1 h. Three independent transformants of each strain were grown and assayed in duplicate. Data points represent the averaged values for the three independent transformants. Bars show the averaged values for the three transformants ± S.E. ***, p ≤ 0.0001; **, p ≤ 0.001; *, p ≤ 0.05 is defined as not significant (ns) by one-way analysis of variance, relative to the WT Fes1 expressing strain. E, lysates were prepared from a WT yeast strain containing the indicated Fes1–FLAG-expressing plasmids after treatment (or mock treatment) with 5 mM cumene hydroperoxide (CHP) for 30 min. Fes1 was detected by Western blotting with anti-FLAG antibody. Western images are representative of the data obtained from three independent experiments.
butyl-hydroperoxide (tBHP) did not result in a detectable change in Fes1 migration (Fig. 1A), although even higher concentrations of tBHP were found to induce mobility changes (data not shown). The relative sensitivity of Fes1 to the distinct peroxides is in keeping with the relative yeast toxicity reported for these peroxides; a 10% survival has been established for 4 mM CHP, 6 mM hydrogen peroxide, and 15 mM tBHP, respectively (27, 28). Although exposure to several types of ROS resulted in a clear migration change in Fes1, exposure to other common stressors did not alter Fes1 mobility on SDS-PAGE. Yeast that underwent heat shock, treatment with cycloheximide (CHX), exposure to the reductant (DTT), or incubation with the thiol-oxidant diamide yielded no change in Fes1 mobility (Fig. 1B).

The mobility change we observed for Fes1 was suggestive of a post-translational protein modification triggered by ROS exposure. Prior studies have demonstrated that oxidation of protein methionine residues can cause a pronounced change in protein migration through a reducing SDS-polyacrylamide gel (29–31). We reasoned that the shift in Fes1 mobility could be a consequence of methionine oxidation induced by ROS exposure. To monitor whether Fes1 methionine residues are oxidized in cells exposed to exogenous ROS, we used MS to analyze peptides from Fes1 immunopurified from untreated yeast cells or from cells exposed to 10 mM CHP for 30 min. It is established that methionine oxidation can be induced at several steps during sample preparation for MS analysis (32–34), and care was taken to minimize methionine oxidation post-cell lysis (see “Experimental procedures”). Fes1 contains seven methionines distributed throughout the protein; matched peptides for five of the seven methionines were recovered from both the untreated and treated samples. Methionines in peptides from stressed and unstressed cells were identified in the reduced state (Met; red) and also oxidized to methionine sulfoxide (diox), and oxidized to methionine sulfone (diox). ND, peptide not detected.

| Fes1 Met # | Peptide Fragment | Relative Abundance – CHP + CHP | Fold Enrichment |
|-----------|------------------|-------------------------------|-----------------|
| Met19     | LLQWSIANSQDGKEAM(ox)AR | 1.92 3.28 | 1.71 |
| Met19     | LLQWSIANSQDGKEAM(ox)AR | 1.92 3.28 | 1.71 |
| Met45     | LLQLFGGGGPGDPDTLM(ox)K | 5.18 6.11 | 1.18 |
| Met45     | LLQLFGGGGPGDPDTLM(ox)K | 5.18 6.11 | 1.18 |
| Met49 / Met53 | ESM(red)AVIM(red)NPEVDLETK | 1.00 1.00 | 1.00 |
| Met49 / Met53 | ESM(red)AVIM(red)NPEVDLETK | 1.00 1.00 | 1.00 |
| Met79     | Incomplete coverage | - | - |
| Met126    | AAALSIIGTAVQNNLDSQNNF(ox)K | 7.00 7.00 | 1.00 |
| Meta126   | AAALSIIGTAVQNNLDSQNNF(ox)K | 7.00 7.00 | 1.00 |
| Meta194   | AALSIIGTAVQNNLDSQNNF(ox)K | 0.04 0.05 | 1.25 |

Table 1

**Fes1 methionine-containing peptides identified by mass spectrometry**

Fes1 protein was immunopurified from yeast cells grown in the absence or presence of peroxide, and Fes1 peptides were analyzed by mass spectrometry. Matched peptides containing five of the Fes1 methionines were recovered from both the untreated and treated samples. Methionines were identified in the reduced state (red), oxidized to methionine sulfoxide (diox), and oxidized to methionine sulfone (diox). ND, peptide not detected.
methionine Met-126 (Table 1). We were further struck by the
increased relative abundance of peptides containing oxidized
Met-45, Met-49, and Met-53 under basal conditions (5.2–5.8-
fold relative to reduced peptides), which was further increased
upon peroxide treatment (Table 1). We suggest that increased
oxidation of a given methionine during sample preparation
may also indicate a methionine residue particularly prone to
modification, especially when recovery of the same oxidized
methionine is further noted in peroxide-treated samples. The
three Fes1 methionines that attracted our attention (Met-45,
Met-49, and Met-53) cluster within the area of sequence pre-
dicted to localize to the first α-helix (α1) of the Fes1 core
domain (Fig. S1).

To determine whether oxidation of methionines 45, 49, and
53 accounts for the slower mobility of Fes1 observed upon per-
oxide treatment (Fig. 1A), we generated alleles of Fes1 with
Met–to–Ile, Met–to–Phe, or Met–to–Gln triple mutant did not
show a slower electrophoretic mobility (similar to that
observed in cells, which is likely a consequence of the negative
charge associated with the glutamine side chain. Similarities
between protein methionine sulfoxide and glutamine have
been previously described (37, 38), and we speculate that the
Met–to–Gln allele could be mimicking (in part) the oxidized
form (see under “Discussion”).

**Oxidative modification of Fes1 is reversed by R- and S-type
methionine sulfoxide reductases**

Protein methionines oxidized to methionine sulfoxide
(MetO) can be reversed by methionine sulfoxide reductase
(Msr) enzymes (39). To determine whether the activity of the
yeast Msrs could reverse oxidation of Fes1, we established a
cell-free assay to monitor Fes1 oxidation. Recombinant Fes1
was purified from bacteria and treated with NaOCl. Similar to
what was observed in cells, we saw a clear change in Fes1 mobility
when protein was incubated with increasing concentrations
of oxidant (Fig. 2D). We suggest that this mutant may be partially com-
promised for Fes1 activity. Importantly, consistent with the MS
data, we found that the electrophoretic mobility of a Fes1 Met–
to–Ile, Met–to–Phe, or Met–to–Gln triple mutant did not
change when cells were exposed to CHP (Fig. 1E), supporting
the conclusion that the Fes1 size shift induced by oxidant ex-
sposure is a consequence of methionine oxidation at residues Met-
45, Met-49, and Met-53. A Fes1 Met–to–Gln triple mutant
showed a slower electrophoretic mobility (similar to that
observed for oxidized WT Fes1) in both untreated and oxidant-
treated cells, which is likely a consequence of the negative
charge associated with the glutamine side chain. Similarities
between protein methionine sulfoxide and glutamine have
been previously described (37, 38), and we speculate that the
Met–to–Gln allele could be mimicking (in part) the oxidized
form (see under “Discussion”).
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higher) molar excess of NaOCl, there was very little, or a complete absence, of any Fes1 species co-migrating with untreated (reduced) Fes1 (Fig. 2A, lanes 5 and 6). Recombinant Fes1 Met–to–Ile and Met–to–Phe triple mutants did not undergo a mobility change in the presence of oxidant, establishing that the same Fes1 methionines oxidized in cells are susceptible to modification in vitro (Fig. 2B).

Yeast encode two Msr enzymes (Mxr1 and Mxr2) that reduce protein-imporporated MetO modifications in a stereospecific manner. Yeast Mxr1 (MsrA) and Mxr2 (MsrB) have been linked to the reduction of protein S-MetO and R-MetO, respectively (Fig. 2C) (40). We purified recombinant versions of Mxr1 and Mxr2 and set up an in vitro system to monitor activity toward oxidant-treated Fes1. Recombinant Fes1 protein was oxidized with NaOCl and then incubated with substoichiometric amounts (a 1–to–10 ratio of Mxr–to–Fes1) of Mxr1 and Mxr2, either alone (Fig. 2D, lanes 5 and 6) or in combination (Fig. 2D, lanes 7–10). To catalyze multiple reaction turnovers, DTT or a reconstituted yeast thioredoxin system (consisting of thioredoxin (Trx1), thioredoxin reductase (Trt1), and NADPH) was provided to reduce (reduce) the Msrs. Note, the presence of DTT alone is insufficient to reduce protein–MetO adducts. We observed a clear loss of the slowest migrating, oxidant-induced Fes1 band in the presence of both Msr enzymes and a recycling system (Fig. 2D, lanes 7 and 9). Both the thioredoxin and DTT recycling systems yielded a similar reduction in the slower-migrating Fes1 species (Fig. 2D). Incubation of Fes1 in the presence of a single Msr led to a very modest decrease in the more slowly migrating oxidized Fes1 species (Fig. 2D, lanes 5 and 6 versus 4), and we hypothesize that the synergistic effect of Mxr1 and Mxr2 results from their combined ability to reduce the anticipated mixtures of R- and S-MetO Fes1 species. Of note, a complete restoration of a band that co-migrates with reduced Fes1 was not obtained, and we speculate that some portion of the in vitro-modified Fes1 contains methionines oxidized to methionine sulfone (MetO_S), which is resistant to reduction by Mrss. Such an oxidation of protein methionines in the cell-free system is not unexpected, given the 10-fold excess of oxidant used to oxidize Fes1 in the absence of any cellular system that may serve to limit overoxidation. In addition, we cannot eliminate the possibility that Fes1 is oxidized also at nonmethionine residues, which would not be reversed by Mrss activity; however, the oxidant-treated Fes1 methionine mutants did not show a shift in size upon oxidant treatment (Fig. 2B), suggesting that the remaining size-shifted species in the Mrs-treated oxidized Fes1 samples (Fig. 2D) reflect methionine modification. Altogether, these data confirm that the migration change for Fes1 induced in the presence of oxidant is a by-product of methionine oxidation, which can be reversed (in part) through the activity of the yeast Mrss enzymes Mxr1 and Mxr2.

Given the ability of recombinant Mxr1 and Mxr2 to reverse Fes1 oxidation in vitro, we next sought to establish the ability of either or both enzyme(s) to modulate Fes1 oxidation in vivo. We reasoned that if the Mrss enzymes mediate reduction of Fes1 in cells, yeast strains lacking Mrss activity would accumulate more oxidized Fes1 protein. In keeping with our expectation, we observed yeast deleted for either Mrs enzyme (mxr1Δ or mxr2Δ) and treated with peroxide showed an increased level of Fes1 oxidation, compared with WT cells treated with the same concentration of oxidant (Fig. 2E). In contrast to our in vitro experiments, we did not see a clear combinatorial effect for a loss of both Mxr enzymes on Fes1 oxidation; an mxr1Δ mxr2Δ double deletion strain showed no further increase in oxidized Fes1 levels compared with the single mutant strains (Fig. 2E). Although we expected a loss of both enzymes would yield a further increase in the accumulation of oxidized Fes1, due to their distinct activities toward MetO isomers, we speculate that the observed yeast phenotype could reflect compensatory yeast systems that allow for the growth and viability of the double mutant strain.

Cytoplasmic pool of Mxr2 influences Fes1 oxidation

Previous studies have localized yeast Mxr1 to the cytoplasm and Mxr2 within the mitochondria (40, 41), and thus we were initially surprised to observe an impact for loss of Mxr2 activity on the oxidation of cytoplasmic Fes1. Although the impact for loss of Mxr2 on cellular Fes1 oxidation could be indirect, we were intrigued by the alternative possibility that an uncharacterized cytoplasmic pool of Mxr2 may exist that could directly reduce Fes1, as observed in our in vitro system. Mining the compiled ribosome-profiling and mRNA-sequencing database (42) revealed the potential for prominent use of an internal methionine start codon in Mxr2 (Met-34), numbered relative to the annotated start site (Met-1) (Fig. S2). We hypothesized that translation initiated from Met-34 would generate a protein with an intact enzymatic core that would lack the amphipathic mitochondrial targeting sequence encoded by residues 1–30 (43). We speculated that an Mxr2 protein translated from Met-34 would be localized to the cytoplasmic space, where it could facilitate reduction of oxidized proteins. A similar binary start-site mechanism has been described for other proteins dual-localized to the cytosol and mitochondria (44).

To determine the subcellular distribution of Mxr1 and Mxr2 in yeast, we fractionated lysates prepared from cells expressing GFP-tagged Mxr1 or Mxr2 proteins. Lysates were separated into mitochondrial and cytosolic fractions, and proteins in each fraction were probed for Mxr1/2 (using anti-GFP serum) or for established mitochondrial (porin; Por1/2) and cytosolic (phosphoglycerate kinase; Pgk1) proteins, to confirm the separation of the mitochondrial and cytoplasmic protein pools (Fig. 3A). Consistent with prior reports, Mxr1 fractionated with the cytosolic marker Pgk1 (Fig. 3A) (40). Some Mxr1 and Pgk1 signals were detected in the mitochondrial fraction (Fig. 3A); given the established localization of Pgk1 in the cytoplasm, we speculate that the mitochondrial signal for these proteins is a consequence of carryover from the cytosolic fractions during the experimental protocol. Strikingly, the Mxr2 protein also appeared predominantly in the cytosolic fraction (Fig. 3A); no signal for Mxr2 was detected in the mitochondrial fraction. We cannot rule out that a lower level of mitochondrial Mxr2 protein exists in addition to the cytosolic pool and that this mitochondrial Mxr2 protein pool is below the limit of detection of our assay. Yet, overall our data suggest a predominant cytosolic localization for both Mxr1 and Mxr2. To link the presence of a cytosolic Mxr2 pool and the potential use of an internal (Met-34) start codon, we next mutated the two potential start sites in
Mxr2, generating Met→Leu substitution alleles at Mxr2 Met-1 and Met-34. An Mxr2 lacking the annotated Met-1 start codon (Mxr2–M1L) resulted in a cytosolic localization for Mxr2, which appeared unchanged from the expression level and localization of WT Mxr2 (Fig. 3A). These data suggest the cytosolic pool is generated by initiation at a methionine other than the annotated Met-1 start site. In contrast, signal from an Mxr2–M34L protein was not detected in any cellular fractions (Fig. 3A). Although it is possible that the lack of detectable Mxr2–M34L expression arises from instability (and subsequent turnover) of a mutant protein (initiated at Met-1), we suggest this result indicates Met-34 is the predominant start codon for Mxr2 translation in cells. Indeed, use of Met-34 as a primary start site is supported by data from the compiled ribosome-profiling database, which reveals minimal footprints for both scanning and elongating ribosomes at the Met-1 site compared with Met-34 (Fig. S2).

To determine whether the cytosolic Mxr2 modulates Fes1 oxidation, we took advantage of the increase in Fes1 oxidation observed in an mxr2Δ strain treated with peroxide (Figs. 2E and 3B, lane 5). We confirmed first that addition of a plasmid encoding WT Mxr2 could rescue the increased Fes1 oxidation phenotype of the mxr2Δ strain; we established that the presence of a MXR2 plasmid resulted in less Fes1 oxidation upon peroxide treatment (Fig. 3B, lane 2) similar to what was observed in a WT yeast strain (Fig. 3B, lane 1). Addition of a plasmid encoding Mxr2–M1L protein resulted in a similar outcome as observed for a plasmid encoding WT Mxr2 (Fig. 3B, lane 3), suggesting that a cytoplasmic pool of Mxr2 (initiated at a start site distinct from Met-1) accounts for the relative decrease in accumulated Fes1 oxidation in a WT strain versus an mxr2Δ. A strain containing a plasmid coding for Mxr2–M34L accumulated higher levels of oxidized Fes1 (Fig. 3B, lane 4), analogous to the levels in an mxr2Δ strain. The absence of detectable, tagged Mxr2–M34L protein (Fig. 3A) leads us to suggest that the absence of Fes1 MetO-reduction activity in this strain is a consequence of no (or low levels of) functional Mxr2 protein.

Our results indicate an underappreciated cytoplasmic pool of Mxr2 that can influence the level of oxidized Fes1; these results contrast with prior reports of a sole mitochondrial localization for Mxr2 (40, 41). A distinction between our experi-
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ments and these previous studies is our use of constructs wherein Mxr2 expression is controlled by the endogenous MXR2 promoter; prior localization of Mxr2 to the mitochondria was established using constructs with stronger, nonnative promoters to drive Mxr2 expression: GAL (40) or TEF (41). Notably, in each case, this strong promoter was fused proximal to the Met-1 codon sequence; correspondingly, the Met-34 codon would be more distant in sequence (~100 bp) from the nonnative promoter sequence. To follow localization of Mxr2 under similar conditions as those previously reported, we generated C-terminally–tagged mNeonGreen Mxr constructs under control of the TEF2 promoter. We assessed the localization of these proteins using fluorescence confocal microscopy (Fig. 3C). Under these conditions, Mxr1 showed a diffuse fluorescence distribution throughout the cell (Fig. 3C); the cytoplasmic localization of Mxr1 was consistent both with previous studies (40) and also the localization we observed for Mxr1 expressed from its endogenous promoter (Fig. 3A). Importantly, we observed that the TEF-promoted Mxr2 co-localized with the mitochondria, visualized by a mitochondrial-targeted RFP (Fig. 3C). Substitution of Met-34 had no impact on localization of the TEF-promoted Mxr2; the Mxr2–M34L proteins localized to the mitochondria similar to WT Mxr2 (Fig. 3C). In contrast, an Mxr2–M1L protein resulted in a cytoplasmic localization for Mxr2 (Fig. 3C), suggesting that use of an internal start methionine allows for generation of a cytoplasmic form of Mxr2 (in the absence of Met-1). Altogether, we conclude that a mitochondria-localized Mxr2 protein is translated when the annotated open reading frame (ORF) is juxtaposed with a strong promoter. Yet, our data suggest that when translation is mediated by endogenous sequences surrounding the annotated ORF, translation preferentially starts at an internal methionine. We suggest that this internal methionine is Met-34; in support of this conclusion, mutation of Met-34 to Leu, in a construct containing the native 5’ UTR, results in no detectable Mxr2 protein (Fig. 3A).

To determine whether the subcellular localization of Mxr2 influences its ability to modulate Fes1 oxidation, we took advantage of the distinct intracellular distribution of the TEF-promoted WT and mutant Mxr2 proteins (Fig. 3C). We observed that the predominantly cytoplasmic Mxr2 (Mxr2–M1L) was most efficient at limiting the accumulation of oxidized Fes1 in peroxide-treated cells (Fig. 3D, lane 3). In contrast, more oxidized Fes1 accumulated in peroxide-treated cells containing the mitochondria-localized Mxr2 and Mxr2–M34L proteins (Fig. 3D, lanes 2 and 4). Of interest, the strain expressing TEF-promoted Mxr2 showed higher levels of oxidized Fes1 than a WT strain (Fig. 3D, lane 2 versus 1), suggesting that overexpression of Mxr2 from a nonnative promoter results in less reduction of Fes1 than observed in a strain containing a lower level of Mxr2 expression mediated by its endogenous promoter. We speculate that endogenous Mxr2, present in the cytoplasm of a WT strain, is able to reduce cytoplasmic Fes1, whereas a mitochondrial pool of Mxr2 (driven by the nonnative promoter) is not sufficient to fully rescue the observed elevated levels of Fes1 oxidation in an mxr2Δ strain.

Our data suggest the potential for both cytoplasmic and mitochondrial Mxr2 forms that modulate MetO oxidation of protein substrates within these respective compartments. Such a cross-organelle distribution would be consistent with the reported multiple isoforms of the mammalian methionine sulf oxide reductase MrsB, which are localized to multiple intracellular compartments, including the cytosol, mitochondria, nucleus, and endoplasmic reticulum (45).

Fes1 oxidation alters its ability to modulate Hsp70 activities

Having established the susceptibility of Fes1 to methionine oxidation, we next sought to determine how oxidation impacts the established abilities of Fes1 to facilitate Hsp70 nucleotide exchange and substrate release (Fig. 4A) (22, 24, 26, 35).

To monitor NEF activity, we employed stopped-flow spectroscopy and the fluorescent ADP analog MABA-ADP. When bound to Hsp70, MABA-ADP shows enhanced fluorescence; nucleotide release from the Hsp70 manifests as a decrease in fluorescence (46, 47). Yeast contain two classes of cytoplasmic Hsp70s, grouped based on sequence homology: the Ssa and Ssb proteins (48). Prior reports have demonstrated that Fes1 shows a similar affinity toward Ssa1 and Ssb1 (22) and that Fes1 is effective as an Ssa1 and Ssb1 NEF (22, 24). A majority of published reports focus on Ssa1 as the model cytoplasmic Hsp70, and we continue this trend for most of our biochemical assays. However, we found that the Ssa1 protein purified with bound nucleotide, which we were unable to effectively remove (see “Experimental procedures”). Thus, for the NEF assays we made use of recombinant Ssb1 protein, which in its apo-state could be efficiently loaded with MABA-ADP. Complexes were formed between recombinant Ssb1 and MABA-ADP. Release of MABA-ADP from Ssb1 (in the presence of an excess of unlabelled ADP) was found to be relatively slow in the absence of a NEF (Fig. 4, B and C); an average MABA-ADP off-rate of 0.189 ± 0.009 s⁻¹ was measured (Fig. 4C), which is similar to the previously reported rates of 0.205 ± 0.001 s⁻¹ (22) and 0.27 s⁻¹ (24). Addition of Fes1 accelerated the rate of MABA-ADP release from Ssb1 ~9-fold, increasing the off-rate to 1.634 ± 0.098 s⁻¹ (Fig. 4C). These data are consistent with the previously reported ability of Fes1 to stimulate MABA-ADP release from Ssb1 (22, 24).

To test the effect of Fes1 oxidation on its activity as a NEF, oxidized Fes1 was prepared by treating recombinant protein with a 10-fold molar excess of NaOCl. To prevent oxidation of Ssb1 protein, all the oxidant-treated Fes1 samples (and the oxidant-treated buffer control) were run through a desalting column prior to injection into the stopped-flow apparatus. Treatment with oxidant was observed to lessen Fes1 NEF activity (Fig. 4, B and C). The rate of Ssb1 nucleotide release stimulated by oxidized Fes1 was diminished ~4-fold relative to reduced Fes1, exhibiting an average measured off-rate for MABA-ADP of 0.404 ± 0.083 s⁻¹. The inhibitory effect of oxidant treatment was partially reversible. Treatment of the desalted oxidized Fes1 protein with a mixture of Mxr1 and Mxr2, prior to mixing with Ssb1–MABA-ADP, resulted in a Fes1 protein that stimulates nucleotide release ~5-fold, relative to the rate observed in the absence of Fes1 (Fig. 4C). The partial restoration of Fes1 activity was consistent with the partial restoration in Fes1 mobility seen upon incubation with a combination of Mxr1 and Mxr2 (Fig. 2D). It is worth noting that we were unable to mon-
The association of ATP with apo-Hsp70 induces peptide release. The interaction of ATP with apo-Hsp70 stimulates Hsp70 ATPase activity. To monitor ATPase activity, we followed ATP hydrolysis by Ssa1 protein purified from yeast. Similar to the NEF activity assays, we observed that WT Fes1 stimulated ATP turnover by Ssa1 and that the stimulatory activity of Fes1 was decreased upon treatment of Fes1 with oxidant (Fig. 4D). The damped activity for oxidized Fes1 was reversed by treatment with Mrx1 and Mrx2 (Fig. 4D). Given the location of the oxidation-sensitive methionines in proximity to the N-terminal domain (Fig. S1), we speculated that the lack of activity observed for oxidized Fes1 might be mediated by changes to the N-terminal region. However, we observed that the presence of the N terminus is not necessary to mediate oxidant-induced changes to Fes1 activity (Fig. S4).

A consequence of stimulated Hsp70 nucleotide release by a cytoplasmic NEF is the association of Hsp70 with ATP, which facilitates peptide release (Fig. 4A) (22, 25, 50). The NR peptide (sequence: NRLLLTG) and a derivative of the APPY peptide (Ala-P5: ALLLSAPRR) have been shown to be substrates for the cytoplasmic Hsp70 Ssa1 (25, 50–52). Hence, to monitor the influence of Fes1 on peptide binding, we made use of the Ssa1 protein and an NR peptide labeled with the environmentally-sensitive fluorophore NBD, which shows enhanced fluorescence when complex with an Hsp70. We initially planned to also monitor Ssb1–peptide interactions, but we were unable to detect an association of labeled peptide with Ssb1 (data not shown); a similar lack of detectable interaction between Ssb1 and APPY–peptide has been noted previously (51). Complexes of Ssa1 were formed with the NBD–NR peptide. Using stopped-flow spectroscopy, we monitored the ability of Fes1 to stimulate a decrease in NBD–NR fluorescence upon mixing of the Ssa1–NBD–NR complex with an excess of ATP and unlabeled peptide. In the absence of Fes1, we observed a peptide release off-rate of 0.083 ± 0.002 s⁻¹, which was accelerated ~8-fold by Fes1 (k_{obs} of 0.66 ± 0.017 s⁻¹).
Oxidant-treated Fes1 showed a decreased ability to stimulate peptide release from Ssa1; relative to the off-rate observed in the presence of untreated Fes1, oxidant-treated Fes1 peptide release was slowed by ~5-fold ($k_{\text{obs}}$ of 0.136 ± 0.023 s⁻¹) (Fig. 4, E and F). The activity of oxidant-treated Fes1 was restored when incubated with a combination of Mxr1 and Mxr2, resulting in an average off-rate for the Ssa1–peptide complex of 0.503 ± 0.17 s⁻¹ (Fig. 4F). Interestingly, peptide release kinetics appeared to be more resistant to the effects of Fes1 oxidation, as Fes1 inhibition required more oxidant (20-fold molar excess NaOCl) compared with other Hsp70 activity assays, where Fes1 inhibition was observed upon oxidation with 10-fold molar excess NaOCl (Fig. 4, B–D). It is unclear whether the increased oxidation requirement we observed relates to technical differences for the distinct assays or whether the necessity for additional oxidation to impact peptide release has biological implications.

Most recently, a new activity for the N-terminal region of Fes1 as a pseudo-peptide has been reported, wherein the association of the Fes1 N terminus with the substrate-binding domain prevents the re-association of released peptide with the Hsp70, to further facilitate Hsp70 peptide clearance (Fig. 4A) (26). Fes1 was found to limit association of a dansyl-labeled NR peptide with Ssa1, and this activity of Fes1 depended on the presence of an intact Fes1 N terminus (26). Using anisotropy, we observed a similar decrease in the association of Ssa1 with a 6-carboxyfluorescein (FAM)-labeled NR peptide in the presence of Fes1 (Fig. 4G). We found that oxidant-treated Fes1 was less effective in limiting peptide association (Fig. 4G). Altogether, these data suggest that oxidation diminishes all known Hsp70-associated Fes1 activities; oxidation decreases the ability of Fes1 to facilitate Hsp70 nucleotide exchange, mediate peptide release, and limit peptide association.

**Interaction between Fes1 and Hsp70 is diminished upon Fes1 oxidation**

Structural data for the Fes1 ortholog HspBP1 shows that the core domain adopts a largely $\alpha$-helical structure with a concave face that embraces lobe II of the Hsp70 NBD (25). Four helical repeats (formed by helices 3–14) adopt a structure similar to Armadillo repeats; these repeats are capped at each end by longer helices (a1 and a15–17) (25). Homology modeling based on the HspBP1 core structure predicts that the oxidant-sensitive methionines for Fes1 map along a1 (Fig. S1) (22). It has been suggested that a steric clash between a1 and the Hsp70 NBD is what propels the change in NBD conformation that facilitates nucleotide exchange (25). We reasoned that oxidized Fes1 (when associated with Hsp70) may be unable to facilitate a change in NBD conformation (and nucleotide release). Alternatively, we speculated that conformational changes with a1 induced by oxidation may serve to more globally disrupt the core domain structure and alter the ability for major contacts to form between the Hsp70 lobe II and the Armadillo repeats, precluding the association of Hsp70 and Fes1.

To test the latter possibility, we examined whether the association of Fes1 and Hsp70 was altered upon oxidation. Purified recombinant His$_{6}$-tagged Fes1 was immobilized on Ni-NTA resin after treatment with oxidant (or mock treatment). Immobilized Fes1 was then incubated with an equimolar amount of untagged recombinant Ssb1. Yeast-purified Ssa1 preparations could not be used in this assay, as Ssa1 was tagged with a His$_{6}$-tag. Consistent with prior studies, we observed a stable association of Fes1 with Ssb1 (Fig. 5A) (22). We found that the interaction between Fes1 and Hsp70 was diminished upon Fes1 oxidation (Fig. 5A). The decreased interaction correlated with the degree of Fes1 oxidation; treatment of Fes1 with increasing amounts of NaOCl corresponded with the appearance of increasingly-slower migrating Fes1 species and a decrease in Hsp70 association (Fig. 5A). A modest background binding of Ssb1 with the Ni-NTA beads was observed even in the absence of Fes1 protein (Fig. 5A, lane 7); however, the majority of recovered Ssb1 signal in the pulldowns required the presence of Fes1. The decreased association between oxidized Fes1 and Ssb1 could be reversed by Mxr1/2 treatment; the recovery of Hsp70 bound to immobilized Fes1 was increased in samples where oxidant-treated Fes1 was incubated with Mxr1/2 prior to immobilization (Fig. 5B, lanes 6 versus 5). The presence of Mxr1/2 and the thioredoxin system alone did not alter the association of unmodified Fes1 and Ssb1 (Fig. 5B, lanes 3 and 4).

Similar results were obtained also for experiments wherein a more complex mixture of yeast cell lysate was used in lieu of purified Hsp70 protein. Cytoplasmic Hsp70s of both the Ssa and Ssb class were isolated from yeast lysates by immobilized FLAG-tagged Fes1 (Fig. 5C), similar to what has been reported previously (21, 22). Treatment of Fes1–FLAG with oxidant prior to immobilization decreased the recovery of Ssa–Fes1 and Ssb–Fes1 complexes (Fig. 5C). A striking protein band of ~70 kDa was isolated in complex with untreated (reduced) Fes1, and recovery of this band decreased upon Fes1 oxidation (Fig. 5C). The ~70-kDa band was confirmed to include both Ssa and Ssb proteins by Western blotting (Fig. 5C). We noted above that similar to oxidized Fes1, a Met–to–Gln mutant shows altered NEF activity (Fig. S3). We observed that recovery of both Ssa and Ssb proteins was decreased also for a FLAG-tagged Fes1 Met–to–Gln mutant (Fig. 5D), relative to untreated (reduced) WT Fes1–FLAG. Altogether, these data suggest that Fes1 oxidation disrupts the interaction with Hsp70. A decreased association between Hsp70 and Fes1 upon oxidation is consistent with the overall disruption of Fes1 activities, including the ability to facilitate nucleotide exchange, peptide binding, and peptide clearance (Fig. 4).

**Discussion**

We identified the yeast cytoplasmic NEF Fes1 as prone to methionine oxidation under excessively-oxidizing conditions. Our data show that when a cluster of three methionines (Met-45, Met-49, and Met-53) in the Fes1 core domain are oxidized, the physical association of the cytoplasmic Hsp70s and Fes1 is diminished (Fig. 5), which correlates with an observed decrease in the ability of oxidized Fes1 to stimulate Hsp70 nucleotide exchange, peptide release, and/or peptide association (Fig. 4). Although the impact for Fes1 oxidation on yeast physiology remains a focus for future research efforts in the lab, we speculate that altered activity of Fes1 upon oxidation may benefit cells during oxidative stress by helping to maintain cytoplasmic proteostasis (53).
A variety of proteins post-translationally modified by MetO have been identified from a range of species, including many proteins identified as oxidized in proteome-wide studies of human cells treated with various oxidants (9, 12, 54, 55). Recently, a database (MetOSite) was constructed, which catalogs MetO sites from 3562 different proteins identified across 23 species (56). Notably few S. cerevisiae methionine oxidation–susceptible proteins are annotated in the MetOSite, and our identification of Fes1 further extends the known targets of MetO oxidation in this widely-studied model organism. MetOSite lists only two yeast proteins identified as susceptible to methionine oxidation: Mge1 (41) and enolase (shown to be oxidized in vitro) (12).

Interestingly, one of the two reported yeast proteins susceptible to oxidation (Mge1) is a NEF for the mitochondrial Hsp70s, and it is attractive to consider that changes to Hsp70 chaperone function through modulation of NEF activity could be a common theme in the response to oxidative stress in the mitochondria and cytoplasm of yeast. The human orthologs of Fes1 have also been identified with MetO adducts in proteomic studies, and we further speculate that modulation of Hsp70 activity through NEF oxidation may be conserved across species. Similar to what we report for Fes1, directed studies of yeast Mge1 have demonstrated that methionine oxidation lessens the interaction between Mge1 and Hsp70 and decreases Mge1 NEF activity (41, 58, 59). 

In vitro oxidation of GrpEL similarly was shown to lessen the ability of GrpEL to stimulate Hsp70 ATPase activity (58).

We propose that a decrease in Fes1 activity upon oxidation may be of benefit to cells during oxidative stress. A global inactivation of cellular NEF activity during oxidative stress could result in a prolonged association of ADP-bound Hsp70 with peptide, which could serve to limit protein aggregation during overly-oxidizing conditions. However, cells contain multiple NEFs from distinct structural families (GrpE, Hsp110, Fes1, and BAG family proteins) (13), and not all cellular NEF proteins have been identified as prone to oxidation. In addition, for the subset of NEFs identified as susceptible to MetO oxidation in prior proteomic studies (mammalian HspBP1, GrpEL, and BAG proteins), only GrpEL has been shown to become inactivated upon methionine oxidation (58). Thus, we find it interesting to consider specifically how inactivation of Fes1 activity might benefit cells.

It has been previously shown that a strain disrupted for Fes1 activity (a<sup>fes1Δ</sup>) displays a decreased translation rate (35) and a constitutively up-regulated heat-shock response (21). Given that oxidation inactivates Fes1 activity (as would be true for a <sup>fes1Δ</sup>), we speculate that Fes1 MetO modification could acti-
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In vertebrates, beneﬁcial stress responses such as translation attenuation or the HSR, which would help cells cope with excess ROS during oxidative stress. Ssb1/2 have an established role in protein translation (60), and consistent with a change in translation upon Fes1 oxidation, we found Fes1 oxidation lessened the association of Fes1 with Ssb1/2 (Fig. 5). It has been suggested that Fes1 may help to unload (release) the nascent chain from Ssb1 (22), and a loss of Fes1 activity could slow translation by stalling the nascent chain release. A decrease in protein load during unfavorable folding conditions could beneﬁt cells by decreasing the folding burden on an already stressed cytoplasmic system. We also observed an elevated HSR for cells containing a Fes1 Met→Gln mutant (Fig. 1). Similar functional outcomes have been reported for proteins with methionine(s) modiﬁed with methionine sulfoxide or mutagenized to glutamine (37, 38). Consistent with the behavior of the Fes1 Met→Gln substitutions as oxidomimetics, we observed oxidized Fes1 and a Fes1 Met→Gln mutant both showed a decrease in NEF activity (Fig. 4 and Fig. S3) and a lessened association with Hsp70 (Fig. 5). We suggest that the elevated HSR seen for the Fes1 Met→Gln allele would be consistent with HSR up-regulation as a consequence of Fes1 methionine oxidation. We speculate that an increased production of HSR targets (including additional chaperones) could serve to buffer against an unfavorable folding environment during excessively oxidizing conditions.

Moreover, the recently reported role for the Fes1 N-terminal region in peptide clearance suggests a unique activity for Fes1, relative to the other classes of cytoplasmic NEFs (Hsp110 and BAG proteins) that may be modulated by oxidation (26). It was reported that the unstructured Fes1 N-terminal domain associates with the Hsp70 SBD to limit Hsp70 re-association of aggregation-prone substrates; aggregation-prone substrates are proposed to include substrates with multiple exposed hydrophobic patches that possess low apparent Hsp70 off-rates (26). We suggest that during oxidative stress, the chance for productive folding is low. We hypothesize that Fes1 inactivation during these suboptimal folding conditions would prolong the interaction of substrates with the cytoplasmic Hsp70s, due to the continued re-association of released substrates in the absence of the action of the Fes1 N terminus. We propose this extended Hsp70 binding may serve to limit the release and subsequent aggregation of aggregation-prone substrates. It is inferred that substrates prevented from re-associating with Hsp70 (due to normal Fes1 activity) are delivered to the proteasome to be degraded (26), and it is interesting to consider why cells under oxidative stress might prioritize prolonged Hsp70–substrate interaction over substrate degradation. We propose that an extended Hsp70 association of aggregation-prone substrates during oxidative stress may be advantageous due to the compromised activity of the 26S proteasome and ubiquitination-activating/conjugation machinery during oxidative stress (61).

Alternatively (but not mutually exclusively), a decreased association of oxidized Fes1 with the cellular Hsp70s may promote the interaction of Hsp70 chaperones with other binding partners that in turn act to beneﬁcially modulate Hsp70 chaperone activity. In mammals, HspBP1–Hsp70 complex formation precludes the association of Hsp70 with the ubiquitin ligase CHIP (62), and HspBP1–Hsp70 association has been shown to interfere with CHIP-induced degradation of misfolded Hsp70 substrates, including the cystic ﬁbrosis transmembrane regulator (CFTR) and sequestered antigens (62–64). Although yeast do not contain a CHIP homolog, it is tempting to speculate that additional Hsp70 interactors exist that may complex with Hsp70 in the absence of Fes1 to alter Hsp70 chaperone function during oxidative stress. In mammals, we speculate that oxidation of HspBP1 could promote the association of Hsp70 with CHIP to facilitate protein turnover.

How oxidation of Met-45, Met-49, and Met-53 disrupts Hsp70 association (and decreases Fes1 activity) remains to be determined. For the structurally distinct NEF Mge1 (65), oxidation was demonstrated to disrupt dimer formation (59), and it was speculated that the dissociation of Mge1 dimers accounts for the concomitant decrease in Hsp70 association (59). Given the monomeric structure of HspBP1 (25), we anticipate that Fes1 oxidation disrupts Hsp70 association through a different means. Based on homology modeling, we suggest that all three Fes1 methionines localize to one face of the helix (α1) in the Fes1 core domain (Fig. S1). Physical contact between the Hsp70 NBD and the analogous α1 helix of human HspBP1 or the yeast ER ortholog Sii1 was not observed in the published Hsp70–NEF complex structures (25, 66). Thus, we do not expect Fes1 oxidation to disrupt an Hsp70–NEF interface. Yet, both structures were solved for only a portion of the NEF (the core domain), and it remains possible that the α1 helix forms direct Hsp70 contacts that remain to be observed and that may be altered upon NEF oxidation. We can envision also that oxidation of methionines in α1 may perturb the structure of helix 1, which could limit Hsp70 association as a consequence of concomitant changes to the positioning of the ﬂanking N-terminal region and/or to the conformation of the Fes1 core. Helix destabilization would be consistent with other reported examples of the outcome for methionine oxidation; for example, a loss of the secondary helical structure as a consequence of methionine sulfoxide formation was shown to disrupt interactions normally formed by the β-amyloid peptide and small heat-shock proteins (67, 68). Similar to what we observed for Fes1, methionine residues within and around the α1 helix of human HspBP1 (Met-93 and -104) have been identiﬁed as susceptible to oxidation (9, 54, 55). Despite an overall poor primary sequence homology between yeast Fes1 and human HspBP1 (Fig. S1), we speculate that oxidation may occur at structurally analogous positions and serve to similarly impact NEF activity. Although our understanding of the exact nature of MetO-induced structural perturbations in Fes1 remains imprecise, it is evident that altering the oxidant-susceptible methionine residues in the α1 region perturbs Fes1 function; all of the site-directed mutations we generated at these residues resulted in proteins with compromised activity (Fig. S3).

Our characterization of Fes1 oxidation also revealed a cytoplasmic pool of the methionine sulfoxide reductase enzyme Mxr2 (Fig. 3), which was characterized previously as a mitochondrial enzyme (40). Our data establish Fes1 as the ﬁrst substrate of cytoplasmic Mxr2 activity (Fig. 3); Mge1 has been characterized as a mitochondrial Mxr2 substrate (41). To the
best of our knowledge, our data characterize Fes1 also as the first substrate of yeast Mxr1.

Uncovering a pool of cytoplasmic Mxr2 begins to explain how reversible methionine oxidation is achieved for cytoplasmic yeast proteins. It is anticipated that oxidation of proteins results in a mixture of S-MetO and R-MetO adducts. A MetO epimerase enzyme (capable of interconverting S- and R-MetO species) has not been identified in yeast (or any other organism), and until now the cytoplasm was expected to contain a single enzyme with select activity toward S-MetO (40). A cytosolic pool of both Mxr1 and Mxr2 explains how yeast can reduce cytoplasmic proteins modified by both S-MetO and R-MetO. The efficient reduction of MetO adducts allows cells to regulate signaling events brought about by MetO protein modifications and also limits the overoxidation of methionine sulfoxide residues to irreversible methionine sulfoxide species, which may lead to protein aggregation and damage. Although identification of two cytoplasmic Msrs begins to clarify how reversible methionine oxidation is achieved in the yeast cytosol, we speculate that other yeast Msr proteins likely remain to be identified, including a mitochondrial enzyme with activity toward S-MetO as well as enzymes localized to the nucleus or endoplasmic reticulum. Furthermore, we anticipate that yet to be uncovered mechanisms may exist to regulate the distribution of Mxr2 between the cytoplasm and mitochondria depending on the redox conditions within these organelles.

In conclusion, our data establish a role for reversible methionine oxidation of Fes1 in regulating the interaction between Fes1 and Hsp70 and in modulating Hsp70 activities. Our results provide a new perspective on how activities of the well-studied Hsp70 family can be attuned by ROS and further define how methionine modification is used as a means for cells to sense and respond to oxidative stress. We anticipate that Fes1 MetO modification may be conserved in other species, and we speculate that inactivation of Fes1 via MetO modification may be an important means to regulate the critical activities of Hsp70 chaperones during oxidative stress.

**Experimental procedures**

**Plasmid construction**

A complete list of plasmids can be found in Table S1. Yeast expression plasmids derive from the pRS vector series (69). Gene sequences were amplified from genomic DNA prepared from S288C strains CSY5 or BY4741 (70). For expression from the endogenous promoter elements, sequences coding for the ORF plus ~1 kb and ~500 bp of the 5′ and 3′ UTR, respectively, were amplified. Epitope-tagged versions of Fes1 were constructed by insertion of sequence coding for a 3×FLAG epitope or GFP immediately prior to the stop codon. Plasmids pEN254–257 were constructed by insertion of amplified sequences plus sequence coding for a C-terminal mNeonGreen were generated by insertion of amplified ORF plus the endogenous promoter elements, sequences coding for the yeast Mxr1, Mxr2 (beginning at residue 30), Trx1, and Trr1 complete Fes1 coding sequence or sequence coding for Fes1 residues 38–290 into a pET21b vector (EMD Millipore) to generate an in-frame C-terminal His6-tag. The coding sequences for yeast Mxr1, Mxr2 (beginning at residue 30), Trx1, and Trr1 and were similarly cloned into pET28a (EMD Millipore) to create an in-frame N-terminal His6-tag and to generate plasmids pEN55, pBM1, pEN208, and pEN108. Complete Ssb1 and Fes1 coding sequences were inserted into a modified version of the pET28a vector containing a His6–SUMO tag to construct pEN130 and pEN253. For pEN253, a C-terminal 3×FLAG was inserted immediately prior to the stop codon. To make pEN21, DNA sequence coding for the Ydj1 J-domain (residues 1–69) was inserted into pGEX-5X-3 (GE Healthcare). pCS980 was constructed by inserting the upstream sequence from SSA3 (nucleotides −468 to +6) prior to the bacterial lacZ sequence in a pRS315 vector. QuikChange site-directed mutagenesis (Agilent Technologies) was performed to generate amino acid substitutions. All plasmids and mutations generated were confirmed by sequencing.

**Yeast strains and growth conditions**

*S. cerevisiae* strains were grown and genetically manipulated using standard techniques (71). Cultures were grown in rich medium (1% Bacto-yeast extract and 2% Bacto-peptone) containing 2% dextrose (YPD) or 2% galactose (YPGal) or minimal medium (0.67% nitrogen base without amino acids supplemented with 14 amino acids, uracil, and adenine) containing 2% dextrose (SMM) or 2% galactose (SGal). Uracil or leucine supplements were removed from minimal media to select for plasmids as needed.

A complete list of yeast strains used in this study can be found in Table S2. Strain CSY751 was made by one-step gene replacement of the coding region of *FES1* with KanMX in BY4741 (72). Strains CSY974–976 were generated from standard crosses initiated from the *mxr1Δ* and *mxr2Δ* BY4741 and BY4742 deletion collection strains (Research Genetics). Strain CSY825 was a kind gift from Dr. Kevin Morano (University of Texas Health Science Center) and is described in Ref. 21. Strain AJMY28 was generously provided by Dr. Jeffrey Brodsky (University of Pittsburgh) and is described in Ref. 73.

**Cell-based assays to monitor levels of oxidized Fes1 in vivo**

Yeast cells expressing Fes1–FLAG constructs were grown to mid-log phase in minimal medium, and cultures were treated for the final 30 min prior to harvesting at 30 °C with the following stressors: 5–10 mM hydrogen peroxide (EMD), 5–10 mM tert-butyl hydroperoxide (Alfa Aesar), 5–10 mM cumene hydroperoxide (CHP) (Alfa Aesar), 0.5–1 mM sodium hypochlorite (Sigma), 10–50 µg/ml cycloheximide (Amresco), 5–10 mM dithiothreitol (DTT) (GoldBio), or 5–10 mM diamide (TCI). Heat-shocked samples were shifted to a 37 °C water bath for 1 h prior to harvest. Cells (5 OD600) were harvested and washed in sterile water, and lysates were prepared by alkaline lysis (74). Proteins were suspended in 100 µl of SDS sample buffer (60 mM Tris-HCl, pH 6.8, 5% glycerol, 2% SDS, 0.0025% bromphenol blue) with 5% BME, and samples were boiled at 100 °C for 5 min. Insoluble material was removed with a 1-min spin at 21,130 × g. Soluble proteins were separated by reducing SDS-PAGE (12% gel) and transferred to nitrocellulose. Proteins were detected using an anti-FLAG mouse mAb (Agilent Technologies; catalog no. 200472) or anti-FLAG rat mAb (Thermo Fisher Scientific; clone L5, catalog no. MA1-142), and an Alexa
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488 – conjugated anti-mouse (Invitrogen; catalog no. A11029) or an Alexa 488- or Alexa 546 – conjugated anti-rat (Invitrogen; catalog nos. A11006 and A11081) secondary antibody. Fluorescent signals were detected with a ChemiDoc MP system (BioRad). Images shown are representative of at least three independent experiments.

Heat-shock activation assay

Yeast (CSY825) transformed with an HSE-lacZ reporter plasmid (pCS980) and the indicated CEN FES1 plasmids (pEN139 and pEN141–143) were grown in SMM medium at 30 °C until mid-log phase (OD600 between 0.5 and 0.7). Heat-shocked samples were shifted to a 37 °C water bath for the last 1 h prior to harvest. Cells were permeabilized using Y-PER (Thermo Fisher Scientific), and β-gal activity was quantified as described (75). For each experiment, three yeast transformants of each strain were assayed in duplicate. Error bars denote standard deviation of the average from three independent experiments.

Mass spectrometry

WT cells expressing Fes1–FLAG from a 2-μm plasmid were grown in SMM medium at 30 °C until mid-log phase and were treated with 10 mM CHP for 30 min at 30 °C prior to harvest. Cells (200 OD600) were collected, washed in sterile water, and lysed by alkaline lysis (74). Proteins were suspended in 100 μl of SDS sample buffer with 5% BME, and samples were boiled at 100 °C for 5 min. Lysate proteins were incubated for 30 min at room temperature in 1 ml of IP buffer (50 mM Tris-HCl, pH 7.4, 150 mM NaCl, 1% Triton X-100). Unbroken cells and insoluble material were removed by centrifugation for 5 min at 21,130 × g. The remaining supernatant (1 ml) was added to 250 μl of anti-FLAG bead resin slurry (Sigma) and incubated overnight at 4 °C. Beads were washed five times with 1 ml of IP buffer, and bound proteins were eluted using 250 μl of SDS sample buffer with 5% BME and boiled at 100 °C for 5 min. To concentrate proteins, 1.5 ml of cold acetone was added and incubated at −20 °C overnight. Protein precipitate was collected after a 21,130 × g spin at 4 °C for 15 min, dried, and resuspended in 40 μl of SDS sample buffer with 5% BME. Proteins were separated by reducing SDS-PAGE (10% gel) and visualized with SYPRO Ruby (Invitrogen). Gel bands corresponding to Fes1–FLAG were extracted and analyzed by MS at the Cornell University Proteomics and Mass Spectrometry Core facility. To limit spontaneous methionine oxidation, all sample preparation steps at the facility were conducted in the presence of 20 mM methionine, and trypsin digestion was limited to 3 h at room temperature, and the HPLC trapping column was pre-washed with 20 mM EDTA three times prior to nanoLC-MS/MS analysis.

Protein expression and purification

Plasmids encoding for recombinant His6-tagged proteins (Fes1, Mxr1, Mxr2, Trx1, and Trr1) were transformed into BL21 (DE3) cells, and fresh transformants were grown overnight at 37 °C in LB medium containing 100 μg/ml ampicillin or 15 μg/ml kanamycin. The following morning, cells were diluted 1:50 into LB containing fresh antibiotic, and cultures were grown at 37 °C until an OD600 of 0.5 was reached. Cells were shifted to 18 °C for 30 min, and protein expression was induced with 0.5 mM isopropyl β-D-thiogalactopyranoside (GoldBio). After 18 h, cells were harvested, and pellets were stored at −80 °C. Cells were thawed and resuspended in lysis buffer (50 mM sodium phosphate, pH 8, 0.5 mM NaCl, 10 mMimidazole, 10% glycerol) containing an EDTA-free protease inhibitor mixture (Pierce), 5 mM BME, 12.5 units/ml benzonase (Pierce), and 2 mg/ml lysozyme (VWR). Resuspended cells were lysed on ice with a microtip sonicator using five 30-s pulses (with a 2-min rest between pulses) at a 60% duty cycle. Insoluble material was removed by centrifugation at 23,700 × g for 20 min at 4 °C. Soluble material was loaded onto a HiTrap chelating column (GE Healthcare) charged with nickel. The column was washed with at least 10 column volumes (cv) of lysis buffer and 10 cv of wash buffer (50 mM sodium phosphate, pH 8, 0.5 mM NaCl, 20 mM imidazole, 10% glycerol) until the A280 signal baseline. Protein was eluted with 5 cv of elution buffer (50 mM sodium phosphate, pH 8, 0.5 mM NaCl, 250 mM imidazole, 10% glycerol). Protein fractions were collected, exchanged into desalting buffer (10 mM Tris-HCl, pH 7.4, 50 mM NaCl, 10% glycerol) using a PD-10 desalting column (GE Healthcare), and concentrated using a centrifugal filter (Millipore). Purified proteins were flash-frozen in liquid nitrogen and stored at −80 °C. Concentrations were determined by BCA protein assay (Thermo Fisher Scientific) using BSA as a standard.

His6–SUMO–Ssb1 (pEN130) and His6–SUMO–Fes1–FLAG proteins (pEN253 and pEN258) were expressed and purified by metal-affinity chromatography as above. Protein elution fractions were collected and concentrated, and the His6–SUMO tag was cleaved off the protein at 4 °C using recombinant yeast His6–Ulp1 protease. Proteins were exchanged into lysis buffer with a PD-10 column (to remove imidazole), and the uncleaved protein, protease (His6-tagged), and the released His6–SUMO tag were removed with a HiTrap chelating column charged with nickel and equilibrated in lysis buffer. Fes1–FLAG was used without additional purification. Untagged Ssb1 was further purified to facilitate nucleotide removal. Ssb1 protein fractions were exchanged into 10 mM Tris-HCl, pH 7.4, 10% glycerol, 10 mM EDTA using a PD-10 desalting column and incubated at 4 °C overnight to allow for nucleotide release. Apoprotein was isolated using a HiTrap Blue HP column (GE Healthcare). Ssb1 was eluted with a salt gradient (0–2 M NaCl), and fractions containing Ssb1 were collected and concentrated, and Ssb1 protein was stored in desalting buffer (10 mM Tris-HCl, pH 7.4, 50 mM NaCl, 10% glycerol) at −80 °C. Purified protein was confirmed to be nucleotide-free by HPLC.

Plasmid coding for GST–Ydj1(1–69) (pEN21) was similarly transformed into BL21 (DE3) cells, and protein was expressed as above. Harvested cells were thawed and resuspended in PBS containing an EDTA-free protease inhibitor mixture, 5 mM BME, 12.5 units/ml benzonase, and 2 mg/ml lysozyme and lysed by sonication as above. After sonication, 0.1% Triton X-100 final was added to the lysate mixture. Insoluble material was removed by centrifugation at 23,700 × g for 20 min at 4 °C, and soluble material was loaded onto a GSTrap column (GE Healthcare). The column was washed with at least 20 cv of wash buffer 1 (PBS with 2 mM EDTA), 20 cv of wash buffer 2 (PBS...
with 2 mM EDTA, 1 mM KCl, and 0.1% Triton X-100), 20 cv of ATPase wash buffer (50 mM Tris-HCl, pH 7.4, 2 mM ATP, 10 mM MgOAc₂, 200 mM KOAc), and 10 cv of PBS until the A₂₈₀ signal of the column flow-through baselined. Protein was eluted with 5 cv of elution buffer (50 mM Tris-HCl, pH 8, 10 mM reduced GSH, 10% glycerol). Protein fractions were collected, desalted, concentrated, flash-frozen, and stored as above.

The yeast strain AJMY28 was generously provided by Dr. Jeffrey Brodsky (University of Pittsburgh), and His₆–Ssa1 was purified from this strain as described previously (73), with minor alterations. Cells freshly streaked from a frozen glycerol stock were grown in SGal-ura medium at 30 °C to saturation. Saturated cultures were diluted 1:20 into YPGal, and cultures were grown overnight (~20 h) at 30 °C until cells reached late log phase (OD₆₀₀ between 2.5 and 3). Cell pellets were harvested and resuspended at a 1:1 volume ratio in Ssa1 lysis buffer (50 mM sodium phosphate, pH 7.8, 300 mM NaCl, 10 mM imidazole, 5 mM BME) containing 1 mM PMSF (VWR) and 1 μM pepstatin A (Alfa Aesar). Suspended cells were frozen by drop-wise addition of the cell slurry into liquid nitrogen, and frozen cells were stored at −80 °C. Cells were lysed with 10 2-min runs/1-min rests in a cryomill (SPEX SamplePrep) set at 12 cps. Insoluble material was removed by centrifugation at 23,700 g for 20 min at 4 °C, and soluble material was incubated with Ni-NTA–agarose resin (Thermo Fisher Scientific) for 2 h at 4 °C. The resin was washed with 10 cv of Ssa1 lysis buffer, and bound protein was eluted with 3 cv of Ssa1 elution buffer (50 mM sodium phosphate, pH 7.8, 300 mM NaCl, 200 mM imidazole, 5 mM BME). Ssa1 protein fractions were subsequently diluted with an equal volume of ATP-agarose buffer (25 mM HEPES-KOH, pH 7.4, 10 mM NaCl, 10 mM KCl, 5 mM MgCl₂, 2.5% glycerol, 5 mM BME) and loaded onto a 5-ml ATP-agarose column (Sigma, catalog no. A2767). The column was washed with 5 cv of ATP-agarose buffer and then with 5 cv of ATP-agarose buffer with 50 mM NaCl. Bound protein was eluted from the column with 3 cv of ATP-agarose buffer containing 2 mM ATP. Final eluted fractions were collected, exchanged into desalting buffer (10 mM HEPES-KOH, pH 7.4, 50 mM NaCl, 10% glycerol), and concentrated using a centrifugal filter (Millipore). Purified proteins were flash-frozen in liquid nitrogen and stored at −80 °C. Concentrations were determined by a BCA protein assay as above.

**In vitro Fes1 oxidation and reduction**

Purified recombinant Fes1–His₆ protein (10 μM) was diluted in desalting buffer (10 mM Tris-HCl, pH 7.4, 50 mM NaCl) with sodium hypochlorite (Sigma) in quantities ranging from equimolar (10 μM NaOCl) to 20-fold molar excess (200 μM NaOCl) for 60 min at room temperature. To monitor the extent of oxidation, proteins were separated by reducing SDS-PAGE (12% gel) and visualized by staining with Coomassie Brilliant Blue.

To prepare oxidized Fes1 for later reduction with methionine sulfoxide reductase enzymes, Fes1–His₆ (50 μM) was incubated with 500 μM NaOCl in desalting buffer for 60 min at room temperature, and excess oxidant was removed with a Bio-Gel P6 Microspin column (Bio-Rad). For all Hsp70 assays making use of oxidant-treated Fes1, buffer with oxidant was similarly desalted for the no Fes1 control. To reduce Fes1 and recycle the Mxr enzymes, oxidized Fes1–His₆ (5 μM) was added to His₆–Mxr1 (0.5 μM), His₆–Mxr2 (Δ1–29) (0.5 μM), and either 0.5 mM DTT or a mixture of 0.25 μM His₆–Trx1, 0.25 μM His₆–Trx1, and 0.5 mM NADPH in a final reaction volume of 50 μl. For Hsp70 assays, oxidized Fes1 was reduced by incubation with equimolar His₆–Mxr1 and His₆–Mxr2 (Δ1–29) in the presence of 10 mM DTT. Samples were incubated with Mxrs for 60 min at room temperature. When analyzed by electrophoresis, samples were quenched with the addition of SDS sample buffer, and proteins were separated by reducing SDS-PAGE (12% gel) and visualized by staining with Coomassie Brilliant Blue.

**ATP-hydrolysis measurements**

ATP hydrolysis was measured using a modified version of the EnzChek Phosphate Assay kit (Thermo Fisher Scientific) as described previously (76). In brief, ATP hydrolysis was assayed in ATPase buffer (50 mM Tris-HCl, pH 7.4, 50 mM KCl, 5 mM MgCl₂, 1 mM DTT) containing 200 μM 2-amino-6-mercapto-7-methylpurine riboside, 0.2 units/ml purine nucleoside phosphorylase, and 5 mM ATP. Phosphate generation was monitored by following the change in absorbance at room temperature using a BioTek Synergy 2 microplate reader. Final protein concentrations were as follows: 0.5 μM His₆–Ssa1, 1 μM GST–Ydj1 (1–69), and 1 μM Fes1–His₆ (reduced or oxidized with 10-fold molar excess NaOCl). Fes1–His₆ or GST–Ydj1 proteins were assayed alone to confirm an absence of contaminating ATP hydrolysis activity.

**Stopped-flow nucleotide and peptide release assays**

Experiments were carried out in HKM buffer (25 mM HEPES-KOH, pH 7.4, 150 mM KCl, 5 mM MgCl₂, 1 mM DTT), and data were collected with a Hi-Tech SFA-20 stopped-flow coupled to a Fluoromax-4 spectrofluorometer. To monitor nucleotide release, Ssb1–MABA-ADP complexes were prepared by incubation of 2 μM nucleotide-free Ssb1 and 2 μM MABA-ADP (Jena Bioscience) in a final volume of 2 ml at room temperature for 5 min. Dissociation of the fluorescent nucleotide analog was monitored upon addition of Fes1–His₆ and an excess ADP, for a final reaction of 1 μM Ssb1, 1 μM MABA-ADP, 125 μM ADP, and 1 μM Fes1–His₆ (reduced or oxidized with 10-fold molar excess NaOCl). Data were collected at 0.05-s intervals during a 60-s time course, with an excitation wavelength of 335 nm, an emission wavelength of 440 nm, and a 3-nm slit width. To monitor peptide release, His₆–Ssa1–NBD–NR complexes were prepared by incubation of 5 μM His₆–Ssa1 and 2 μM NBD–NRLLTGT peptide (NBD–NR; Elim Biopharmaceuticals) in a final volume of 2 ml at room temperature for 1 h. Dissociation of the fluorescently-labeled peptide was monitored upon addition of excess unlabeled NR peptide (Elim Biopharmaceuticals), ATP, and 5 μM Fes1–His₆ (reduced or oxidized with 20-fold molar excess NaOCl). The final reaction consisted of 2.5 μM His₆–Ssa1, 1 μM NBD–NR, 250 mM ATP, 125 μM NR peptide, and 2.5 μM Fes1–His₆. Data were collected at 0.05-s intervals during a 60-s time course, with an excitation wavelength of 465 nm, an emission wavelength of 533 nm, and a 5-nm slit width. Data curves for nucleotide and
peptide release assays were fit to a one-phase exponential decay using GraphPad Prism (version 8.0).

**Peptide-association measurements**

Experiments were carried out in HKM buffer, and anisotropy measurements were performed with a Fluoromax-4 spectrofluorometer using a sub-micro quartz fluorometer cell (Starna). Ssa1–Fes1 complex was prepared by incubation of 50 μM Hsp84–Ssa1 with an equimolar concentration of Fes1–His6 (reduced or oxidized with 10-fold molar excess NaOCl) in a final volume of 50 μl at room temperature for 30 min. Fluorescently-labeled peptide association was monitored upon addition of a 10 μM complex to a reaction containing 50 nM fluorescein-labeled NR peptide (FAM–NR; Elim Biopharmaceuticals) in a 150-μl final volume. Fluorescence anisotropy measurements were collected at 12-s intervals over 60 min at an excitation wavelength of 492 nm, an emission wavelength of 516 nm, and a 12-nm slit width.

**Cellular subfractionation**

WT yeast cells expressing GFP-tagged Mxr constructs were grown to mid-log phase in minimal medium. Cells (100 OD600) were harvested with sterile water, washed with a buffer of 50 mM Tris-HCl, pH 7.4, 10 mM sodium azide. Cells were incubated in 100 mM Tris-SO4, pH 9.4, with 50 mM BME for 15 min at room temperature, and cells were harvested and resuspended in 1.2 M sorbitol with 10 mM Tris-HCl, pH 7.4, plus lyticase. After a 45-min incubation at room temperature, spheroplasts were harvested by centrifugation, washed with 10 mM Tris-HCl, pH 7.4, 1.2 M sorbitol, and lysed on ice using a Dounce homogenizer in 1 ml of lysis buffer (0.2M sorbitol, 1 mM EDTA) containing a yeast Protease Arrest protease inhibitor mixture (G Biosciences). Cell debris was cleared by centrifugation at 3000 × g at 4 °C, and mitochondria-containing pellets were collected by centrifugation at 20,000 × g at 4 °C. The mitochondrial-containing pellet was resolubilized and saved for further analysis. A portion (250 μl) of the post-mitochondrial (cytosolic) fraction was re-centrifuged at 70,000 × g at 4 °C to remove high-density membranes and protein aggregates, and the remaining soluble material was saved as a representative of the cytosol. SDS sample buffer and 5% BME final were added to proteins collected in each fraction sample, and proteins were separated by reducing SDS-PAGE (12% gel). Proteins were transferred to nitrocellulose, and Mxr proteins were detected using an anti-GFP mouse mAb (Clontech; catalog no. 632569). Control proteins were detected using mouse monoclonal antiporin (Invitrogen; clone 16G9E6BC4, catalog no. 459500) or anti-Pgk1 (Invitrogen; clone 22C5D8, catalog no. 459250); antibodies. Protein signals were visualized using an Alexa 488–conjugated anti-mouse secondary antibody and a ChemiDoc MP system. A representative image of three independent experiments is shown.

**Fluorescence microscopy**

Images were acquired using a CSU-X spinning-disk confocal microscopy system (Intelligent Imaging Innovations) equipped with a DMI 6000B microscope (Leica) and a ×100 1.47NA field planarity apochromat objective lens and a scientific complementary metal-oxide-semiconductor (scCMOS) camera (ORCA Flashv4.0 V2, Hamamatsu). WT cells transformed with pTB3390 (Mito-RFP) (obtained from Dr. Anthony Bretscher, Cornell University, New York) and the indicated Mxr-mNeonGreen plasmid were grown to log phase at 30 °C in SMM medium, and cells were harvested, attached to a glass-bottomed dish coated with concanavalin A (EY Laboratories, Inc.), and washed with cell medium. Cells were live-imaged at room temperature. SlideBook 6.0 software was used to operate the microscopy system and analyze the images. Multiplane images to reach the top and bottom of each cell were taken at 0.3-μm steps (~20–25 total planes) and used to create z-stack maximum projections.

**Protein pulldown assays**

To assess complex formation between recombinant Fes1 and Ssb1, Fes1–His6 (10 μg) was incubated with 20 μl of HisPur Ni-NTA magnetic bead suspension (Thermo Fisher Scientific) in a total volume of 100 μl of binding buffer (50 mM sodium phosphate, pH 8, 100 mM NaCl, 10 mM imidazole, 0.1% Igepal, 2% glycerol, 1 mM DTT, 1 mM pepstatin A). Samples were rotated for 1 h at 4 °C, and beads were collected by magnetization. Beads were washed three times with 500 μl of binding buffer to remove unbound proteins, and washed beads were suspended in a final volume of 100 μl of binding buffer. Untagged Ssb1 (20 μg) was added to the beads, and samples were rotated at 4 °C for 1 h. Beads were magnetized and washed three times with 500 μl of binding buffer. Bound proteins were solubilized in 25 μl of sample buffer containing 5% BME. Samples were resolved on an SDS-acrylamide gel (12%) and visualized with Coomassie Brilliant Blue. Band intensities were quantified using Image Lab software (version 5.2). A representative image of three independent experiments is shown.

To assess association of recombinant Fes1 with yeast cytosolic proteins, Fes1–FLAG (~40 μg) or an equivalent amount of BSA (Calbiochem) was incubated with 25 μl of anti-FLAG bead slurry (Sigma) in a total volume of 200 μl of desalting buffer. Samples were rotated for 1 h at 4 °C, and beads were collected by centrifugation at 1000 × g for 2 min. Beads were washed three times with 500 μl of binding buffer to remove unbound proteins. Cytosol was prepared from WT yeast (BY4741), which were grown to saturation at 30 °C, harvested by centrifugation, and resuspended in a 1:1 volume ratio of desalting buffer containing 1 μM peptatin A and 1 mM PMSF. Cells were lysed using a cryomill (SPEx SamplePrep) using 10 2-min runs (with 1-min rest between cycles) at 12 cps. Insoluble lyse material was removed by centrifugation at 23,700 × g for 20 min at 4 °C, and the remaining cytosolic fraction (1 ml) was added to the prepared FLAG beads and incubated for 2 h at 4 °C. Beads were collected by centrifugation at 1000 × g for 2 min and washed three times with 500 μl of binding buffer, and proteins were solubilized in 50 μl of sample buffer containing 5% BME. Proteins were resolved on an SDS-acrylamide gel (12%) and visualized with Coomassie Brilliant Blue stain or by Western blotting. Primary antibodies against the yeast cytosolic Hsp70s (anti-Ssa, anti-Ssa3/4, and anti-Ssb) were generously provided by Dr. Elizabeth Craig (University of Wisconsin-Madison). Proteins were visualized using an Alexa 488–conjugated anti-rabbit secondary antibody (Invitrogen; catalog no. A21206).
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