The intermembrane space protein Mix23 is a novel stress-induced mitochondrial import factor

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The biogenesis of mitochondria requires the import of hundreds of precursor proteins. These proteins are transported post-translationally with the help of chaperones, meaning that the overproduction of mitochondrial proteins or the limited availability of chaperones can lead to the accumulation of cytosolic precursor proteins. This imposes a severe challenge to cytosolic proteostasis and triggers a specific transcription program called the mitoprotein-induced stress response, which activates the proteasome system. This coincides with the repression of mitochondrial proteins, including many proteins of the intermembrane space. In contrast, herein we report that the so-far-uncharacterized intermembrane space protein Mix23 is considerably up-regulated when mitochondrial import is perturbed. Mix23 is evolutionarily conserved and a homolog of the human protein CCDC58. We found that, like the subunits of the proteasome, Mix23 is under control of the transcription factor Rpn4. It is imported into mitochondria by the mitochondrial disulfide relay. Mix23 is critical for the efficient import of proteins into the mitochondrial matrix, particularly if the function of the translocase of the inner membrane 23 is compromised such as in temperature-sensitive mutants of Tim17. Our observations identify Mix23 as a novel regulator or stabilizer of the mitochondrial protein import machinery that is specifically up-regulated upon mitoprotein-induced stress conditions.

Mitochondrial biogenesis relies on the import of hundreds of protein precursors from the cytosol. Proteins of the mitochondrial matrix are synthesized with N-terminal presequences or matrix targeting signals (MTSs) which bind to surface receptors on the mitochondrial outer membrane (1, 2). These receptors are associated with the translocase of the outer mitochondrial membrane (TOM) complex, which serves as a general entry gate for protein import (3–6). After translocation through the protein-conducting channel in the TOM complex, MTSs engage with the TIM23 complex of the inner membrane, which facilitates their further transport into the matrix via a reaction driven by the inner membrane potential and by Hsp70-mediated ATP hydrolysis (7, 8). The interactions between the TOM and TIM23 complex are dynamically regulated by the incoming preproteins and depend on intermembrane space (IMS)-exposed protein domains of several proteins, including Tom22, Tim50, Tim21, and Tim23 (9–13).

Most IMS proteins lack MTSs and employ a unique import mechanism independent of ATP and the inner membrane potential (14–16). Many of these IMS proteins contain characteristic cysteine residues that are part of their targeting signal (17, 18). These mitochondrial IMS sorting signal or IMS targeting signal sequences direct proteins through the protein-translocation pore of the TOM complex into the IMS, where they associate with the hydrophobic substrate-binding region of Mia40 (19, 20). Mitochondrial intermembrane space import and assembly protein 40 (Mia40; coiled-coil-helix-coiled-helix domain containing 4 (CHCHD4) in humans) contributes at least two functions: 1) it traps incoming IMS precursors via a hydrophobic interaction with their mitochondrial IMS sorting signal/IMS targeting signal and 2) it introduces disulfide bonds into these substrates (21–23).

Most Mia40 substrates share a simple helix-loop-helix structure in which both helices are stabilized by two parallel disulfide bonds (24–27). Because of the specific spacing of the cysteine residues in their primary sequence, these proteins are referred to as twin Cx3C or twin Cx4C proteins. However, some Mia40 substrates show different numbers and arrangements of their cysteine residues: whereas Mia40 introduces five disul- fides into the IMS protease Atp23, it is critical for the formation of a single disulfide bond in the inner membrane proteins Tim17 and Tim22 (28–31).

The protein import into mitochondria is under surveillance of the proteasome. Arrested mitochondrial import intermediates induce severe stress, presumably because of the accumulation of non-imported mitochondrial precursor proteins in the cytosol (32–37). Conditions that perturb mitochondrial import trigger a characteristic stress response that leads to an induction in expression of many chaperones and the constituents of the proteasome (Fig. 1A) (33, 34, 38). Here we identify the poorly characterized protein Mix23 as one of few IMS proteins that are induced under these stress conditions. Mix23 is critical for efficient protein import into mitochondria, and deletion of Mix23 in temperature-sensitive tim17 mutants leads to synthetic defects in growth and the import of proteins into the mitochondrial matrix. Our results suggest that Mix23 is a stress-regulated factor that is relevant for efficient protein translocation into mitochondria.
Results

Stalled import intermediates selectively induce Mix23 expression

Cloggers are slowly imported mitochondrial precursor proteins that competitively inhibit mitochondrial translocation in vivo and activate the mitoprotein-induced stress response. Because of their slow import, inner membrane proteins with N-terminal stop-anchor sequences (also called bipartite precursors) have particularly strong competitive effects (32, 33). These proteins include model substrates such as cytochrome b$_{2}$-DHFR and endogenous inner membrane proteins such as cytochrome c oxidase 11 (Cox11) or Cox5a. To assess the effect of clogger expression on proteins of the mitochondrial IMS, we analyzed the data from our previous study (33) in which the cellular mRNA levels were followed after induction of the model protein cytochrome b$_{2}$-DHFR for 90 min compared with control cells, which did not express the clogger protein. Whereas most chaperones and proteasome subunit (red) genes were induced by clogger expression, C, most IMS proteins (red) were not significantly increased upon mitoprotein-induced stress. However, clogger expression leads to an up-regulation of Mix23. D, shown are the relative changes in clogger versus control expression of genes coding for IMS proteins (39). Dually localized proteins that are predominantly located in the cytosol were omitted. E and F, RNA levels of MIX23 and PRE2 (coding for a proteasome subunit) are shown for different times after clogger induction. See (33) for details.

Figure 1. The expression of clogger induces MIX23 expression. A, schematic representation of the mitochondrial import machinery and the cytochrome b$_{2}$-DHFR construct (Clogger) (33). B, cells expressing the clogger construct or an empty vector for control were shifted for 90 min from lactate medium to lactate + 0.5% galactose medium. Cellular mRNA was isolated from three biological replicates and sequenced (33). Many chaperone (blue) and proteasome subunit (red) genes were induced by clogger expression. C, most IMS proteins (red) were not significantly increased upon mitoprotein-induced stress. However, clogger expression leads to an up-regulation of Mix23. D, shown are the relative changes in clogger versus control expression of genes coding for IMS proteins (39). Dually localized proteins that are predominantly located in the cytosol were omitted. E and F, RNA levels of MIX23 and PRE2 (coding for a proteasome subunit) are shown for different times after clogger induction. See (33) for details.

Mix23 stabilizes mitochondrial import
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Figure 2. Mitoprotein-induced stress conditions specifically increase Mix23 expression. A, WT cells with empty vector (e.v.) or GAL1-driven cytochrome b2-DHFR plasmid (Clogger) were shifted to galactose-containing lactate medium for 4 h. RNA was isolated and Mix23 mRNA levels (and controls) were quantified by qPCR. The values shown were normalized as described under "Experimental procedures." Mean values of six (three biological and three technical) replicates are shown and were used for a Student’s t test. *p < 0.05; **p < 0.01; ***p < 0.005. B, the experiment was repeated as in (A) but GAL-COXL1 was used instead of the clogger plasmid. C and D, Mix23 expression was analyzed in WT, tim17-5, and mia40-3 cells at permissive (25 °C) growth conditions and after a 24-h shift to semi-restrictive (33 °C) conditions. E, WT cells were grown in galactose medium and shifted for 2 h to the indicated temperatures. F, WT cells were transformed with plasmids for expression of Htt25-GFP, Htt97-GFP, or α-synuclein (SNCA), or with an empty vector for control. After the shift to galactose for 4 or 6 h, MIX23 RNA levels were quantified. G, MIX23 levels were measured in Δrpn4 cells as described in (A). Values were normalized to WT levels. H, Rp4 was overexpressed from a multicopy plasmid under control of the constitutive CYC1-promoter (49). I, mitoprotein-induced stress leads to the simultaneous induction of MIX23 and the proteasome in an Rp4-dependent manner.

Mix23 is a Mia40 substrate of the IMS

Mix23 has been previously identified in a proteomic screen for components that are released from isolated mitochondria by a BCL2-associated X (Bax)-mediated opening of the outer membrane (39). We confirmed the localization of Mix23 in the IMS of mitochondria because it became accessible to protease after rupturing the outer membrane by hypotonic swelling (Fig. 3A). Mix23 homologs are found throughout fungi and animals, all containing four conserved cysteine residues (Fig. 3B and Fig. S1). Mix23 lacks a presequence (Fig. 3B and Fig. S2), and its cysteine pattern is distinct from that of the twin Cx3C or twin Cx9C proteins that represent well-studied substrates of the Mia40 import pathway (20).

When mitochondrial proteins were denatured with SDS and treated with the alkylating agent mmPEG24 (methyl-PEG24-maleimide), Mix23 only shifted in size after the disulfide bonds were reduced with TCEP (tris(2-carboxyethyl)phosphine), consistent with the cysteines in endogenous Mix23 protein being oxidized (Fig. 3C). The formation of disulfide bonds in IMS proteins are catalyzed by the import component Mia40. Mia40 is also crucial for the import of Mix23, because mitochondria from mia40-3 cells did not import radiolabeled Mix23 in an in vitro import experiment whereas mitochondria of WT cells imported Mix23 efficiently (Fig. 3D). The cysteine residues of Mix23 are crucial for its Mia40-dependent import because a mutant in which the six cysteine residues were replaced by serines was not imported (Fig. 3E). Thus, the import of Mix23 shows a similar Mia40-dependence as that of the twin Cx9C protein Tim9 (Fig. S3A). Further, the import of presequence-containing proteins such as Oxa1 does not require Mia40 (Fig. S3B). From this we conclude that Mix23 is an IMS

Mutant (Fig. 2D) in which the import of proteins into the IMS but not into the matrix is compromised (45).

We next tested whether Mix23 expression is also induced upon other stress conditions. However, increased temperature did not result in Mix23 induction, whereas the target of the heat shock response HSP82 was induced (Fig. 2E). The presence of neither aggregate in the cytosol led to Mix23 induction, because neither the overexpression of a polyQ model protein (huntingtin 97 (Htt97)-GFP) nor of α-synuclein significantly changed the transcript levels of Mix23 (Fig. 2F).

Because we observed that Mix23 is coexpressed with genes of proteasome subunits, we tested whether clogger-mediated Mix23 induction depends on regulatory particle non-ATPase 4 (Rpn4), which serves as a master transcription factor for components of the proteasome machinery (46–48). Indeed, cytochrome b2-DHFR expression in Δrpn4 cells did not induce Mix23 (Fig. 2G, cf. Fig. 2A). However, overexpression of Rpn4 in yeast cells (49) induced Mix23 expression as it does of the established Rpn4 target putative proteasome subunit 2 (PLP2) (Fig. 2H). Under the conditions here, the induction is mild (about 1.4-fold) presumably because of the fact that the levels of the Rpn4 protein are tightly regulated by a feedback mechanism (46). From this we conclude that Mix23 and proteasome subunits are coregulated via one common transcription factor (Fig. 2I).

Mix23 is up-regulated upon mitoprotein-induced stress but not upon heat stress

Next, we performed real-time qPCR measurements to study the expression of Mix23 in more detail. This confirmed a strong effect of cytochrome b2-DHFR on the transcript levels of Mix23, which were up-regulated more than 4-fold (Fig. 2A). Mix23 transcript levels were also significantly increased upon overexpression of Cox11, an endogenous protein that is synthesized with a bipartite presequence (41), albeit to a lower degree (Fig. 2B). Mitoprotein-induced stress was also detected in mutants of the mitochondrial TIM23 machinery (34, 42–44), and we observed increased MIX23 levels in temperature-sensitive tim17-5 mutants at permissive (25 °C) and semi-permissive (33 °C) temperatures (Fig. 2C). In contrast, MIX23 levels were more moderately affected in a temperature-sensitive mia40-3 mutant (Fig. 2D) in which the import of proteins into the IMS but not into the matrix is compromised (45).
protein with a conserved pattern of cysteine residues that is imported into mitochondria via the Mia40 disulfide relay (Fig. 3F).

**High Mix23 levels lead to a severe growth defect**

As reported above, mitoprotein-induced stress leads to a considerable up-regulation of Mix23. We therefore tested the effects of increased levels of Mix23 by expression of the protein from a regulatable galactose metabolism 1 (GAL1) promoter (Mix23\(^{\Delta}\)). In the presence of galactose, where Mix23 levels are strongly increased, we observed a considerable reduction in growth of Mix23\(^{\Delta}\) cells irrespective of temperature (Fig. 4A and Fig. S3D). Obviously, increased amounts of Mix23 are not well tolerated by yeast cells, potentially because the unphysiologically high levels of Mix23 clog the import machinery. A hallmark of the mitoprotein-induced stress response triggered by cloggers is the induction of the Rpn4 response (33). Indeed, we observed that Mix23\(^{\Delta}\) strongly induced the expression of a YFP reporter from a reporter that contains the proteasome-associated control element (PACE) to which Rpn4 binds (Fig. 4B).

Because yeast cells can grow by fermentation on galactose plates, the slow growth is not due to defects in respiration but rather in a function of more general relevance.

We therefore tested whether increased levels of Mix23 interfere with the import of proteins into isolated mitochondria. Interestingly, we found that the import of the matrix protein Atp1 and the inner membrane protein Oxa1 into Mix23\(^{\Delta}\) mitochondria was strongly diminished (Fig. 4, C–F). Similarly, the import of the IMS protein Cmc1 into Mix23\(^{\Delta}\) mitochondria was reduced (Fig. 4, G and H).

It was shown before that IMS proteins that fail to be imported into mitochondria are degraded by the proteasome (50–52). The Mia40 substrate Cox12 was identified as a protein that is particularly unstable in the cytosol but stable in the IMS so that its stability can be used to monitor its intracellular distribution (53, 54). We therefore tagged Cox12 by chromosomal integration of an HA epitope in WT or Mix23\(^{\Delta}\) cells. When synthesis of new proteins was inhibited by addition of cycloheximide, Cox12 remained stable in the WT cells but was rapidly degraded in the Mix23\(^{\Delta}\) mutant (Fig. 4I). The matrix protein

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**Figure 3. Mix23 IMS import is Mia40-dependent.** A, isolated mitochondria were subfractionated by hypotonic swelling and incubation with proteinase K (protease). The matrix protein Ilv5 remained inaccessible to protease at these conditions, whereas the IMS protein Mia40 and Mix23 were digestible once the outer membrane (OM) was ruptured by swelling. B, schematic representation of Mix23 homolog sequences from different organisms (see Fig. S1 for details and nonabbreviated species names). Positions of conserved cysteine residues are indicated in yellow. Mix23 was isolated from yeast mitochondria and analyzed by MS. Identified peptides are shown in green. See Fig. S2 for details. C, cell extracts were treated with the reductant TCEP and the alkylating agent mmPEG\(_{24}\). Mix23 was detected by Western blotting. In the presence of TCEP, but not in its absence, Mix23 was alkylated by mmPEG\(_{24}\), indicating that cysteine residues are oxidized in Mix23 in vivo. D, Mix23 was synthesized in reticulocyte lysate in the presence of \(^{[35S]}\)methionine and incubated for the indicated times with mitochondria isolated from WT or mia40-3 cells that were both grown at 33 °C. Mitochondria were isolated and treated with proteinase K where indicated. Proteins were resolved by SDS-PAGE and visualized by autoradiography. E, radiolabeled Mix23CS protein in which all six cysteine residues had been replaced by serine residues was incubated with WT mitochondria and further treated as described in (D). F, schematic representation of the Mia40-mediated import of Mix23 into the IMS, also showing the mixed disulfide form as a reaction intermediate.
mitochondrial ribosomal protein L40 (Mrpl40) served as a control in this experiment and was not degraded by Mix23 overexpression. Thus, in summary, we found that the synthesis of high amounts of Mix23 interferes with the translocation of proteins into mitochondria.

Mix23 is required for efficient preprotein import into tim17-5 mitochondria

Next, we tested the relevance of Mix23 for mitochondrial functionality. Mutants lacking Mix23 did not show growth defects on fermentative and nonfermentative carbon sources, and the import or oxidation of matrix and IMS proteins was not perturbed in the Δmix23 single mutant (Fig. 5 and Fig. S4, A–E).

However, when MIX23 was deleted in the background of a temperature-sensitive tim17-5 mutant, a synthetic growth defect was observed even on fermentative glucose medium at permissive temperatures (25 and 30°C). Under these conditions, the tim17-5 mutant grew indistinguishable from WT cells. The tim17-5 Δmix23 mutant still grew at these conditions, albeit with reduced colony size (Fig. 5B). No synthetic defect was observed when Mix23 was deleted in a temperature-sensitive mia40-3 mutant (Fig. S3C), suggesting that Mix23 is particularly relevant for matrix targeting but not for the biogenesis of IMS proteins using the mitochondrial disulfide relay.

This synthetic genetic interaction of MIX23 and TIM17 was confirmed by in vitro import experiments. The import of Oxa1 and Atp1 into tim17-5 Δmix23 double mutant mitochondria was considerably reduced, particularly at elevated temperatures (Fig. 5, C–F and Fig. S5, A–D), whereas the single mutants were either not affected or less affected. The import of the Mia40 substrate Cx9C motif containing protein 1 (Cmc1) was not diminished in the tim17-5 Δmix23 mitochondria, confirming a specific relevance of Mix23 for matrix-destined proteins (Fig. 5, G and H and Fig. S5, E and F). It should be noted that tim17-5 Δmix23 mitochondria were isolated from cells that were grown at permissive temperature (25°C) at which the tim17-5 mutant shows no obvious difference from WT cells (44). Apparently, efficient Oxa1 import into the matrix of tim17-5 mitochondria requires Mix23.
This suggests that Mix23 has a stabilizing or stimulating activity on mitochondrial protein import. This is also supported by the observation that Cox12 has a much lower stability in Δmix23 cells, in line with a less efficient protein import in this in vivo model (Fig. 5). From these results we conclude that Mix23 is not essential for mitochondrial protein import per se but stimulates or stabilizes protein translocation (Fig. 5).

Figure 5. Mix23 facilitates efficient protein import into mitochondria. A, WT and Δmix23 cells were grown on synthetic lactate medium to log phase. Then 10-fold serial dilutions were dropped onto synthetic lactate (2%) medium. B, Mix23 was deleted in the background of the temperature-sensitive tim17-5 mutant. The resulting double mutant (and tim17-5 cells for control) was analyzed by a drop dilution assay on full glucose medium and grown at the temperatures indicated. C–H, mitochondria were isolated from the strains indicated grown at permissive temperature (25 °C). Radiolabeled Oxa1 and Cmc1 proteins were incubated with these mitochondria at the temperatures indicated. I, the cells were grown on glycerol-containing medium. Protein synthesis was inhibited by addition of cycloheximide. Samples were taken 0, 30, and 60 min after cycloheximide addition. The cells were lysed and the levels of Cox12 and the loading control Djp1 were analyzed by Western blotting. J, model of a role of Mix23 as a factor that directly or indirectly influences the efficiency of mitochondrial protein import.
Mix23 stabilizes mitochondrial import

Discussion

The mitochondrial IMS contains a number of small cysteine-containing proteins that are imported by Mia40. In yeast, where IMS proteins are best characterized, Mia40 clients comprise five twin Cx3C proteins, about 15 Cx9C proteins, and a small number of proteins with cysteine residues in other arrangements (25, 26, 39, 55). Here we show that Mix23 is a further Mia40 substrate of the IMS whose cysteine residues do not match the Cx9C pattern. The Mia40-driven import of Mix23 into the IMS depends on its cysteine residues because a cysteine-less mutant was not import-competent.

In yeast, Mix23 has six cysteine residues, four of which are conserved. Mix23 homologs are found ubiquitously in fungi and animals (many hundred homologous sequences in NCBI). In humans, the Mix23-like protein was named coiled-coil domain containing 58, CCDC58. This protein is localized to mitochondria and found to interact with a number of IMS proteins, including ATPase family gene 3 like 2 (AFG3L2), apoptosis inducing factor mitochondria associated 1 (AIFM1), synthesis of SCO1, and cytochrome c in recent high-throughput interaction studies (56–58). Its function is elusive, but increased CCDC58 levels have been identified as an unfavorable prognostic factor in endometrial hyperplasia (59) and in liver and urothelial cancer (60) (RRID:SCR_006710). Consequently, a role for CCDC58 in tumor progression was suggested. Moreover, CCDC58 was identified as one of a few cellular proteins whose absence provides resistance against the intracellular bacterium *Ehrlichia chaffeensis*, the cause of monocytic ehrlichiosis (61).

Despite this pathophysiological relevance and although CCDC58 is expressed in a wide variety of tissues in mice and humans, including skeletal muscle, heart, and brain, (62, 63), mice lacking CCDC58 were healthy and free of obvious symptoms (64). Thus, increased levels of CCDC58 were associated with pathological conditions in cancer or pathogen-infected cells, but its absence is obviously well tolerated under nonstress conditions. This is reminiscent of what we observed in this study. Yeast cells lacking Mix23 did only show mild defects, whereas overexpression of Mix23 strongly affected cellular functionality. However, it should be noted that the overexpression used in this study is certainly higher than that under physiological conditions even in stressed cells.

The expression of Mix23 was strongly and specifically increased by mitoprotein-induced stress, a condition that both up-regulates (33) and activates (34) the cytosolic proteasome system. Moreover, we found that Mix23 expression depends on Rpn4, the master transcription factor of the proteasome system in yeast (46, 49). Thus, Mix23 serves as a stress-induced factor that is co-regulated with the proteasome. The coregulation of MIX23 and proteasome components is also apparent from the Serial Pattern of Expression Levels Locator database, which calculates coexpression scores on the basis of hundreds of gene expression microarray data (65). Here, MIX23 showed a coexpression with genes with a Gene Ontology (GO) annotation “mitochondrial protein complex,” “mitochondrial large ribosomal subunit,” “mitochondrial small ribosomal subunit,” and “proteasome complex,” in line with a role in mitochondrial biogenesis and proteostasis (Fig. S6). However, the fact that Mix23 and the proteasome are under regulation of the same transcription factor Rpn4 alone is not convincing evidence for a function of Mix23 in protein quality control, but just shows that the regulation of their cellular abundance adheres to similar regulatory patterns.

The molecular function of Mix23 in the IMS and the reasons that this factor is up-regulated during mitoprotein-induced stress remain unclear. An Rpn4-dependent regulation was also described before for Mia40 (33, 66, 67). Still, we found no indication that Mix23 is of relevance for the functionality of the mitochondrial disulfide relay in the IMS, and the deletion of MIX23 did not lead to synthetic defects in *mia40*-3 mutants. However, the observed synthetic defect with the *tim17*-5 mutation suggests that Mix23 facilitates the import of presequence-containing proteins (Fig. 5F). Such a role might be direct, for example by interaction with IMS-exposed regions of the TOM or the TIM23 subunits, which are well-known to be part of a highly complex network of regulatory interactions (9–13). But this role also could be more indirect, for example by an effect of Mix23 on the local distribution of the translocases and their interaction with proteins and lipids of the outer and inner membranes. Several examples for such modulating factors that influence the import machinery were reported in recent years, such as Tim21 or Mgr2 (10, 11, 44, 68, 69).

It will be interesting to unravel the molecular mechanisms in detail by which Mix23 regulates mitochondrial protein translocation and, in particular, its specific function in the context of the cytosolic stress conditions that coregulate it together with the proteasome and other Rpn4 targets.

Experimental procedures

Yeast strains and plasmids

All yeast strains used in this study are based on the WT strains BY4742 (MATα his3 leu2 lys2 ura3), YPH499 (MATα ura3 lys2 ade2 trp1 his3 leu2), or YPH499 Δarg4 (MATα ura3 lys2 ade2 trp1 his3 leu2 Δarg4). The *mia40*-3 and *tim17*-5 mutants were described previously (44, 45). For the detection of Cox12 and Mix23 in Western blotting, a HIS3MX6 cassette was genomically integrated downstream of the COX12 and MIX23 loci, respectively (70). The mutants *tim17*-5 Δmix23 and *mia40*-3 Δmix23 were generated by genomically integrating the NatNT2 cassette into the MIX23 ORF. Positive colonies were verified by PCR.

For generation of the Mix23CS mutant, a variation of the MIX23 DNA sequence was synthesized in which all cysteine-specific codons were replaced by those for serine. This sequence was cloned into pGEM4 using the restriction sites EcoRI and BamHI. The plasmid was verified by sequencing.

To generate a Mix23 overexpression plasmid, the coding region of MIX23 was amplified from genomic DNA and ligated into a pYX225 empty vector plasmid between the GAL1 promoter and an HA tag by using the restriction sites EcoRI and SalI. For the Rpn4 activity reporter, the MIX23 sequence was cloned into a p426 GALI vector, and a pNH603 PACE-YFP reporter plasmid was used as described before (33).

To produce radiolabeled lysate of Mix23, the coding region of MIX23 was amplified from genomic DNA and cloned into a...
pGEM4 plasmid using the restriction sites EcoRI and XmaI. All plasmids were verified by sequencing.

All yeast strains used in this study are listed in Table S1. Strains were grown in yeast complete medium (1% yeast extract and 2% peptone) containing 2% of the carbon sources galactose, glucose, or glycerol as indicated. Temperature-sensitive strains (mia40-3 and tim17-5) were grown at 25 °C before switching them to 33 °C. Strains containing plasmids were grown at 30 °C in minimal synthetic medium containing 0.67% yeast nitrogen base and 2% lactate as carbon source. To induce the expression from the GAL1 promoter, cultures were supplemented with 0.5% galactose.

**Drop dilution assay**

To test growth on plates, drop dilution assays were conducted. Yeast cells were grown in YPGal media or lactate selective media to mid-log phase. After harvesting 0.1 OD600 of cells and washing formed. Yeast cells were grown in lactate selective media to mid-log phase. After harvesting 0.1 OD600 of cells and washing with sterile water, a 1:10 serial dilution was prepared in sterile water. Equal amounts of the dilutions were dropped on agar plates to determine growth differences. Pictures of the plates were taken after 2 to 3 days of incubation.

**Growth curve assay**

To test growth in liquid media, growth curves were performed. Yeast cells were grown in lactate selective media to mid-log phase. After harvesting 0.1 OD600 of cells and washing with sterile water, the cell pellets were resuspended in the experimental media and transferred into a clear 96-well plate. Automated A measurements were performed at 600 nm in the ELx808™ Absorbance Microplate Reader (BioTek®). A's were measured for 72 h every 10 min at 30 °C in technical triplicates.

**Analysis of mRNA levels by quantitative real-time PCR**

To determine relative mRNA expression levels, quantitative real-time PCR was conducted. Yeast strains were treated as indicated and grown until mid-log phase before RNA isolation. The equivalents of two optical densities (600 nm) were used for each RNA sample. RNA isolation was performed using the RNeasy Mini Kit (Qiagen) with a column DNase digestion (Qiagen). 500 ng of each RNA sample were used to synthesize complementary DNA with a qScript™ cDNA Synthesis Kit (Quantabio). To measure relative mRNA levels, the iTaq Universal SYBR Green Supermix (Bio-Rad) was used with 2 μM of a 1 in 10 dilution of cDNA. Measurements were performed in technical triplicates with the CFX96 Touch Real-Time PCR Detection System (Bio-Rad). Relative mRNA expression was calculated by the $2^{-ΔΔCT}$ method (71). For normalization, the housekeeping genes TFC1 or TFA2 were used because of their stability in published gene expression data (72). See Table S3 for primer sequences.

Statistical significance was assessed using a paired one-tailed Student’s t test.

**Cox12 degradation assay**

To investigate the role of Mix23 in the degradation of mitochondrial proteins of the IMS, a Cox12 degradation assay was carried out essentially as previously described (54). Cells expressing Cox12-HA were grown in yeast complete medium containing 2% glycerol (or synthetic lactate medium with 0.5% galactose in case of the Mix23 overexpression strain) before 0.1 mg/ml cycloheximide was added to stop protein translation. Whole cell lysates were taken over time and visualized by Western blotting.

**Isolation of mitochondria**

To isolate crude mitochondria, yeast strains were cultivated either in full or selective media under respiratory conditions. The cells were grown to mid-log phase and harvested by centrifugation (5 min, 3000 × g). After washing the pellets with water and centrifugation (5 min, 3000 × g), the weight of the cell pellet was determined. The pellets were resuspended in 2 ml/g wet weight MP1 (100 mM Tris and 10 mM DTT), incubated for 10 min at 30 °C, and centrifuged again (5 min, 3000 × g). The pellets were washed with 1.2 M sorbitol and centrifuged (5 min, 3000 × g) before resuspending in 6.7 ml/g wet weight MP2 (20 mM KPi buffer, pH 7.4, 1.2 M sorbitol, and 3 mg/g wet weight zymolyase from Seikagaku Biobusiness) and incubated at 30 °C for 60 min. The following steps were conducted on ice. After centrifugation (5 min, 2800 × g), the pellets were resuspended in 13.4 ml/g wet weight homogenization buffer (10 mM Tris, pH 7.4, 1 mM EDTA, 0.2% BSA, 1 mM PMSF, and 0.6 M sorbitol), and a cooled glass potter was used to homogenize the sample with 10 strokes. After homogenization, the extract was centrifuged three times (5 min, 2800 × g) while always keeping the mitochondria-containing supernatant. To pellet mitochondria, the samples were centrifuged for 12 min at 17,500 × g. The pellets were resuspended in SH buffer (0.6 M sorbitol and 20 mM HEPES, pH 7.4). The concentration of the purified mitochondria was adjusted to 10 μg/ml protein. Aliquots were snap-frozen in liquid nitrogen and stored at −80 °C.

**Import of radiolabeled precursor proteins into mitochondria**

To prepare radiolabeled ([35S]methionine) proteins for import experiments, the TNT® Quick Coupled Transcription/Translation Kit from Promega was used. To determine the ability of proteins to be imported into mitochondria, in vitro import assays were conducted. Mitochondria were resuspended in a mixture of import buffer (500 mM sorbitol, 50 mM Heps, pH 7.4, 80 mM KCl, 10 mM Mg(OAc)2), and 2 mM ATP) with 2 mM NADH and 2 mM NADPH to energize them for 10 min at 25 °C. To start the import, the radiolabeled lysate was added to the mitochondria. Import was stopped at different time points by transferring the mitochondria into cold SH buffer (0.6 M sorbitol and 20 mM HEPES, pH 7.4). The remaining precursors outside of the mitochondria were removed by protease treatment for 30 min. 2 mM PMSF was added to stop protein degradation. After centrifugation (10 min, 25,000 × g, 4 °C), the supernatant was removed. The pellets were resuspended in SH buffer containing 500 mM KCl and 2 mM PMSF and centrifuged again (10 min, 25,000 × g, 4 °C). The pellets were then lysed in 1× Laemmli buffer containing 50 mM DTT and heated to 96 °C for 3 min. The samples were run on a 16%
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SDS–gel, blotted onto a nitrocellulose membrane, and visualized with autoradiography.

Cysteine redox state detection by mmPEG24 shifts

To determine the oxidative state of proteins, mmPEG24 shifts were conducted. mmPEG24 binds to reduced thiols, inducing a size shift on SDS gels. For each free thiol, the addition of mmPEG24 increases the size of the protein by 1.2 kDa. Mitochondria were incubated in 80 mM Tris, pH 7, 10% glycerol, 2% SDS, and 0.05% bromocresol blue. Then, mmPEG24 was added to alkylate free thiols. To gain a maximum shift (all thiols reduced), the reducing agent tris(2-carboxyethyl)phosphine was added to a control sample before the mmPEG24 treatment to reduce all disulfide bonds. The samples were analyzed by Western blotting.

Mitochondrial subfractionation

To determine the localization of Mix23 inside mitochondria, a swelling assay was performed. Here, reduced osmolarity of the HEPES buffer causes swelling and subsequent rupture of the outer membrane of mitochondria. Protease treatment subsequently digests proteins of the outer membrane and the IMS. Therefore, mitochondria were incubated on ice in either SH buffer (no swelling) or 20 mM HEPES buffer (swelling and rupture) with and without the presence of 100 µg/ml proteinase K. After 30 min, SH buffer with 20 mM PMSF was added. Mitochondria were then pelleted by centrifugation (10 min, 14,000 rpm, 4 °C). The pellets were resuspended in Laemmli buffer containing 50 mM DTT, heated at 96 °C for 3 min, and visualized by Western blotting.

YFP reporter assay

The cells were grown to mid-log phase (OD600 value of 0.5–0.8) in selective lactate medium. 3 h after addition of galactose, equal amounts of cells (4 OD600 × ml) were collected by centrifugation (20,000 × g, 5 min, room temperature), resuspended in 400 µl synthetic defined medium (100 µl per OD600 and OD600 × ml). 100 µl of this cell suspension were transferred to flat-bottomed black 96-well imaging plates (BD Falcon) in technical replicates. The cells were sedimented by gentle spinning (30 × g, 5 min, room temperature), and fluorescence (excitation 497 nm, emission 540 nm) was measured using a ClarioStar Fluorescence plate reader (BMG Labtech). A corresponding WT strain not expressing YFP was used for background subtraction of autofluorescence. Statistical significance was assessed using a paired one-tailed Student’s t test.

Identification of coexpressed genes and GO enrichment analysis

To identify genes coexpressed with MIX23 across all transcriptomics datasets deposited in the yeast Saccharomyces genome database, the Serial Pattern of Expression Levels Locator database (65) was queried for MIX23 with the online interface provided by the Saccharomyces genome database (https://spell.yeastgenome.org). Genes with an adjusted correlation score of at least 1.4 were considered as coexpressed with MIX23 and subjected to a GO enrichment analysis with GOrilla (RRID: SCR_006848) (73). The 170 coexpressed genes and MIX23 were set as an unranked target list, and the complete Saccharomyces cerevisiae genome was set as the background list. p values for enrichment were calculated according to a hypergeometric distribution and corrected for multiple testing using the Benjamini–Hochberg procedure.

Whole cell lysates

To determine the protein levels of whole cells, whole cell lysates were prepared. To do so, yeast at an A of 1.5 (600 nm) were harvested and washed with water. The pellets were resuspended in Laemmli buffer containing 50 mM DTT. After adding glass beads, the cells were lysed using a cell beater at 4 °C. Then, the cells were heated at 96 °C for 3 min. Lysates were stored at −20 °C until visualization by Western blotting.

Antibodies

Antibodies were produced in rabbits using recombinant purified proteins (see Table S4). Secondary antibodies were ordered from Bio-Rad (Goat Anti-Rabbit IgG (H + L)-HRP Conjugate). The horseradish peroxidase–coupled HA antibody for Western blotting was obtained from Roche (Anti-HA-Peroxidase, High Affinity (3F10), catalog no. 12013819001). The Anti-HA agarose antibody for immunoprecipitation was obtained from Sigma-Aldrich (monoclonal Anti-HA agarose antibody produced in mouse, A2095). Antibodies were diluted in 5% nonfat dry milk-TBS (Roth, T145.2).

Data availability

All relevant data for this study are contained within this article and in the supporting information. Materials and strains are available from the authors.

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Conflict of interest—The authors declare that they have no conflicts of interest with the contents of this article.

Abbreviations—The abbreviations used are: IMS, intermembrane space; TIM, translocase of the inner membrane; MTSS, matrix targeting signals; TOM, translocase of the outer mitochondrial membrane; DHFR, dihydrofolate reductase; mmPEG, methyl-PEG-maleimide; TCEP, tris(2-carboxyethyl)phosphine; PACE, proteasome-associated control element.

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