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# MrpL35, a mitospecific component of mitoribosomes, plays a key role in cytochrome c oxidase assembly

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**ABSTRACT** Mitoribosomes perform the synthesis of the core components of the oxidative phosphorylation (OXPHOS) system encoded by the mitochondrial genome. We provide evidence that MrpL35 (mL38), a mitospecific component of the yeast mitoribosomal central protuberance, assembles into a subcomplex with MrpL7 (uL5), Mrp7 (bL27), and MrpL36 (bL31) and mitospecific proteins MrpL17 (mL46) and MrpL28 (mL40). We isolated respiratory defective *mrpL35* mutant yeast strains, which do not display an overall inhibition in mitochondrial protein synthesis but rather have a problem in cytochrome c oxidase complex (COX) assembly. Our findings indicate that MrpL35, with its partner Mrp7, play a key role in coordinating the synthesis of the Cox1 subunit with its assembly into the COX enzyme and in a manner that involves the Cox14 and Coa3 proteins. We propose that MrpL35 and Mrp7 are regulatory subunits of the mitoribosome acting to coordinate protein synthesis and OXPHOS assembly events and thus the bioenergetic capacity of the mitochondria.

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## INTRODUCTION

Mitochondrial ribosomes (mitoribosomes) perform the synthesis of a small, but important, subset of the mitochondrial proteome, which is encoded by the mitochondrial genome and represents key components of the oxidative phosphorylation (OXPHOS) system (Borst and Grivell, 1978; O'Brien, 2003; Smits *et al.*, 2007; Kehrein *et al.*, 2015; Greber and Ban, 2016; Ott *et al.*, 2016; Mai *et al.*, 2017). Thus the regulation and activity of mitoribosomes have a major potential impact on an organism's capacity to generate its energy through aerobic respiration. Due to the endosymbiotic origin of mitochondria,

the mitoribosomes are evolutionarily related to bacterial ribosomes and thus share a number of overlapping features, such as antibiotic sensitivities and similarities in protein subunit composition (O'Brien, 2002, 2003; Gruschke *et al.*, 2010; De Silva *et al.*, 2015; Greber and Ban, 2016; Ott *et al.*, 2016). Despite these similarities, mitoribosomes display important compositional and functional differences from their prokaryotic ancestors. Mitoribosomes contain a higher protein:rRNA ratio than bacterial ribosomes, and this extra protein content manifests itself in two ways. First, several of the evolutionarily conserved mitoribosomal proteins are larger than their bacterial counterparts, having acquired additional, extension sequences (often C-terminal) and being termed mitochondrial-specific or "mitospecific" sequences. Second, mitoribosomes contain a number of additional novel proteins, that is, mitospecific ribosomal proteins, which are often conserved throughout eukaryotes but do not have counterparts in the prokaryotic ribosomes (Graack *et al.*, 1988; Koc *et al.*, 2000; O'Brien, 2002; van der Sluis *et al.*, 2015). The significant advances in the structural analysis of mitoribosomes, including from the yeast *Saccharomyces cerevisiae*, have greatly enhanced our knowledge of the location of these mitospecific elements within the mitoribosome structures (Amunts *et al.*, 2014; Brown *et al.*, 2014; Greber *et al.*, 2014a,b, 2015; Kaushal *et al.*, 2014; Amunts *et al.*, 2015; Kaushal *et al.*, 2015; Desai *et al.*, 2017). However, little is known about the function of these mitospecific protein features and how they may contribute to, or regulate, the translational process of the mitoribosome.

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Author contributions: J.B., J.K., and R.A.S. conceived and coordinated the study, and R.A.S. wrote the paper. J.B. designed, performed, and together with R.A.S. analyzed all the experiments shown. The data in Figure 5A represent the initial findings of J.K. J.B. prepared all figures. All authors reviewed the results and approved the final version of the manuscript.

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Abbreviations used: BN-PAGE, blue native-polyacrylamide electrophoresis; COX, cytochrome c oxidase; PEBP, phosphatidylethanolamine-binding protein domain; OXPHOS, oxidative phosphorylation.

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Compositional differences between the mitochondrial and bacterial ribosomes are thought to reflect a divergence in function of the mitochondrial translational apparatus from the bacterial. The mitochondrial proteome, in contrast to the bacterial, is mosaic in origin. The proteins synthesized by mitoribosomes are destined to become components of oligomeric complexes coassembled with these imported nuclear encoded partners, a process facilitated by OXPHOS complex-specific chaperone proteins. Thus there is a need for a close coordination between the activity of the mitochondrial translational apparatus and the components involved in the import and assembly of their cytosol-synthesized partners (Gruschke and Ott, 2010; De Silva *et al.*, 2015; Kehrein *et al.*, 2015; Ott *et al.*, 2016).

The cytochrome *c* oxidase complex (COX), the terminal electron acceptor in the mitochondrial OXPHOS machinery, is an oligomeric enzyme complex composed of mitochondrial translation products (Cox1, Cox2, and Cox3) partnered with multiple nuclear encoded ones, imported from the cytosol. The assembly of this complex is a multi-step, chaperone-facilitated process that involves the uniting of protein subunits from two origins and in a strict stoichiometric manner. Cox1, with its metal centers (two hemes [a+a<sub>3</sub>] and one copper, Cug), forms a central aspect of the catalytic core of the COX complex. The synthesis and assembly of Cox1 into the COX complex is tightly regulated, with the proposed purpose of minimizing accumulation of potentially reactive Cox1-assembly subcomplexes (Khalimonchuk *et al.*, 2010; Mick *et al.*, 2011; McStay *et al.*, 2013; Dennerlein and Rehling, 2015; Mayorga *et al.*, 2016; Richter-Dennerlein *et al.*, 2016; Soto and Barrientos, 2016). In yeast, Cox1 synthesis by the mitoribosomes is under the strict feedback regulation from the downstream COX assembly process and directly involves the Mss51 protein (Figure 1). Mss51, a heme sensing protein, promotes the translation of Cox1 by binding to the 5'-UTR of the COX1 mRNA (Perez-Martinez *et al.*, 2003; Barrientos *et al.*, 2004; Pierrel *et al.*, 2007; Perez-Martinez *et al.*, 2009; Fontanesi *et al.*, 2010; Mick *et al.*, 2010; Fontanesi *et al.*, 2011; Soto *et al.*, 2012; Mayorga *et al.*, 2016). The Mss51 protein can also interact with the newly synthesized Cox1 protein (via the C-terminal end of Cox1), as it is cotranslationally inserted into the membrane, where it forms an Mss51-containing Cox1-assembly intermediate (COA) with the Cox14 and

Coa3(Cox25) proteins (Mick *et al.*, 2010; Fontanesi *et al.*, 2011; Clemente *et al.*, 2013; McStay *et al.*, 2013; Garcia-Villegas *et al.*, 2017). Limiting in levels, the Mss51 remains bound to the Cox1 protein-Coa3 intermediate and thus unavailable to bind the COX1 mRNA to promote further rounds of Cox1 synthesis, until the assembly of Cox1 together with the nuclear encoded COX subunits proceeds. Further progression of Cox1 in its assembly pathway causes the release and recycling of Mss51, making it available to initiate the next round of Cox1 protein synthesis. The presence of Cox14 and Coa3 are required to trap Mss51 in the Cox1-assembly intermediate and they thus act as negative Cox1-translational regulators (Mick *et al.*, 2010, 2011; Fontanesi *et al.*, 2011). It is currently unclear what molecular mechanism drives the subsequent recycling of Mss51 from this Cox1-intermediate and whether, and how, the Cox14 or Coa3 proteins may be directly involved.

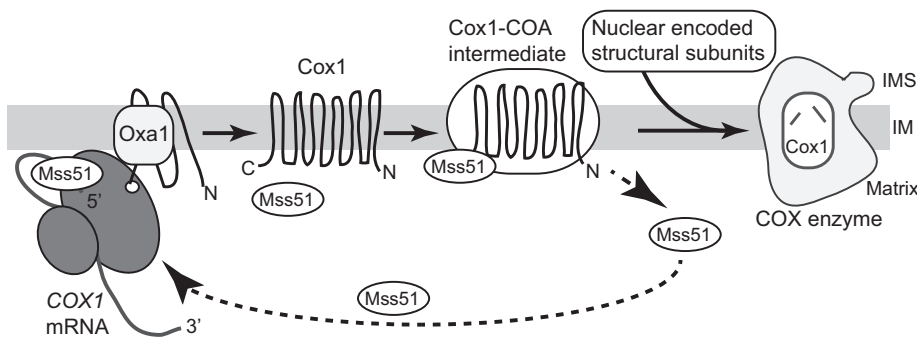
In this present study, we have used the yeast model *S. cerevisiae* to investigate the functional relevance of the mitospecific ribosomal protein, MrpL35 (also known as mL38 using the standardized mitoribosomal protein nomenclature). Conserved throughout eukaryotes (where in higher eukaryotes is referred to as MrpL38), MrpL35 (mL38) is a component of the central protuberance region of the large (54S) mitoribosomal subunit, where its functional relevance is unknown. Our evidence indicates that MrpL35 (mL38) protein is coassembled into a subcomplex with two other mitochondrial-specific ribosomal proteins, MrpL17 (mL46) and MrpL28 (mL40), and together with proteins that are homologues of bacterial ribosomal proteins, MrpL7 (uL5), Mrp7 (bL27), and MrpL36 (bL31), respectively. MrpL35/mL38 proteins contain a C-terminal domain that displays homology to members of the conserved phosphatidylethanolamine-binding protein domain (PEBP) family. The PEBP family comprises a diverse group of proteins present in many organisms and whose functions have been linked to the regulation of key cell signaling pathways (Banfield *et al.*, 1998; Bruun *et al.*, 1998; Serre *et al.*, 1998; Yousuf *et al.*, 2014). Through mutational analysis of the PEBP-like domain, we demonstrate here that MrpL35 plays a key role in coordinating the synthesis and assembly of the COX complex. Our findings demonstrate a connection among MrpL35, Cox1 synthesis, and the COX assembly components, in particular Cox14 and Coa3, and

thus illustrate a link between the mitochondrial translation apparatus and the downstream OXPHOS assembly chaperones.

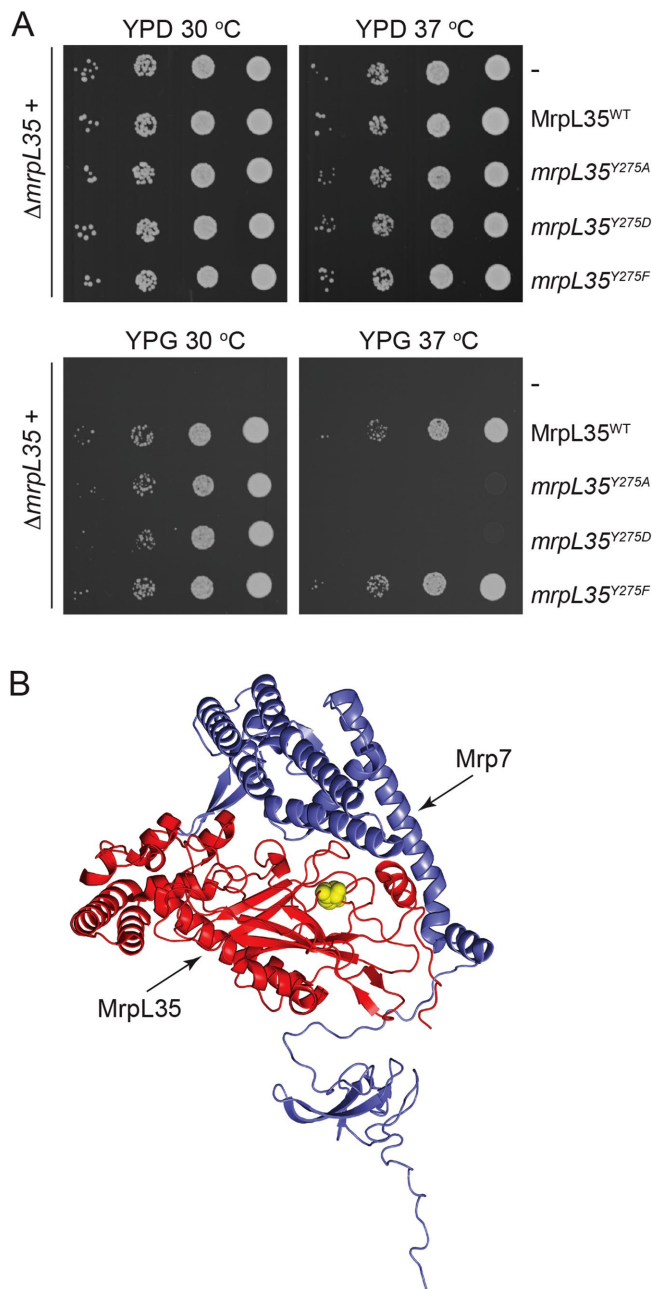
## RESULTS

### Mutation of residue Y275 of MrpL35 results in a temperature-sensitive phenotype

The *S. cerevisiae* yeast *mrpL35* null mutant strain ( $\Delta mrpL35$ ) expressing the wild-type MrpL35 protein (from a plasmid-borne gene) exhibited the ability to grow on the nonfermentable carbon (glycerol) source, whereas the strain lacking MrpL35 (i.e.,  $\Delta mrpL35$ +empty plasmid) was completely unable to do so (Figure 2A). This result illustrates the essential nature of MrpL35 for respiratory/OXPHOS-based growth. Through mL38 protein sequence alignments, we sought to identify highly conserved residues present in the PEBP-like homology domain, located at the C-terminal end of the MrpL35 (mL38) proteins. One of these residues was a conserved



**FIGURE 1:** Overview of the Cox1 synthesis and assembly pathway. Cox1 is synthesized on mitoribosomes tethered to the inner membrane at the site of the cotranslation insertion machinery containing the Oxa1 protein. Synthesis of Cox1 is promoted on activation by the translational activators Mss51 (and Pet309, not depicted) by their binding to the 5'-UTR of the COX1 mRNA. Following insertion into the membrane, the newly synthesized Cox1 protein forms an assembly intermediate with COX-specific assembly factors, including Cox14 and Coa3, which support the recruitment of Mss51 and its binding to the C-terminal end of the Cox1 protein. Further maturation of Cox1 (e.g., incorporation of heme) and assembly with nuclear encoded COX partner subunits, facilitated by specific chaperones, leads to the assembly of Cox1 into the active COX enzyme and promotes the release of Mss51, which in turn activates the next round of Cox1 synthesis. Abbreviations: IM, inner mitochondrial membrane; IMS, intermembrane space; Matrix, mitochondrial matrix space.



**FIGURE 2:** Mutation of residue Y275 of MrpL35 can result in a respiratory deficient phenotype. (A) Serial 10-fold dilutions of  $\Delta mrpL35$  strain harboring a pRS413 plasmid containing gene inserts encoding either the wild-type MrpL35 protein (*MrpL35*<sup>WT</sup>), no insert (-) or genes encoding the mutated *mrpL35* derivatives (*mrpL35*<sup>Y275A</sup>, *mrpL35*<sup>Y275D</sup>, or *mrpL35*<sup>Y275F</sup>), as indicated, were spotted on YP plates containing glucose (YPD) (2 d) or glycerol (YPG) (3 d) and grown at 30° and 37°C. (B) The images of MrpL35 (in red) and Mrp7 (in slate blue) structures were generated using PDB ID:5MRC. The position of the Y275 residue in MrpL35 targeted for mutagenesis is indicated in yellow.

Tyr(Y) residue, corresponding to Tyr(Y)275 of the MrpL35 protein from *S. cerevisiae*, present in all mL38 proteins analyzed, and located on the periphery of the PEBP-like domain of MrpL35 and in a region of the protein that is adjacent to its neighboring protein Mrp7 (bL27) (Figure 2B). We targeted the Y275 residue (indicated in yellow in Figure 2B) for site-directed mutagenesis, initially changing it either to Ala(A) or Asp(D). Expression of the resulting

derivatives, *mrpL35*<sup>Y275A</sup> and *mrpL35*<sup>Y275D</sup>, respectively, supported respiratory-based growth of the  $\Delta mrpL35$  mutant at 30°C but not at the elevated temperature of 37°C (Figure 2A). In contrast, the relatively conservative exchange of Y275 for another aromatic residue Phe(F) did not appear to impact MrpL35's ability to support aerobic growth at both temperatures tested (Figure 2A). Thus we conclude that the highly conserved Y275 residue must play a critical structural/functional role in the MrpL35 protein that is compromised by mutation to a smaller, or negatively charged, residue.

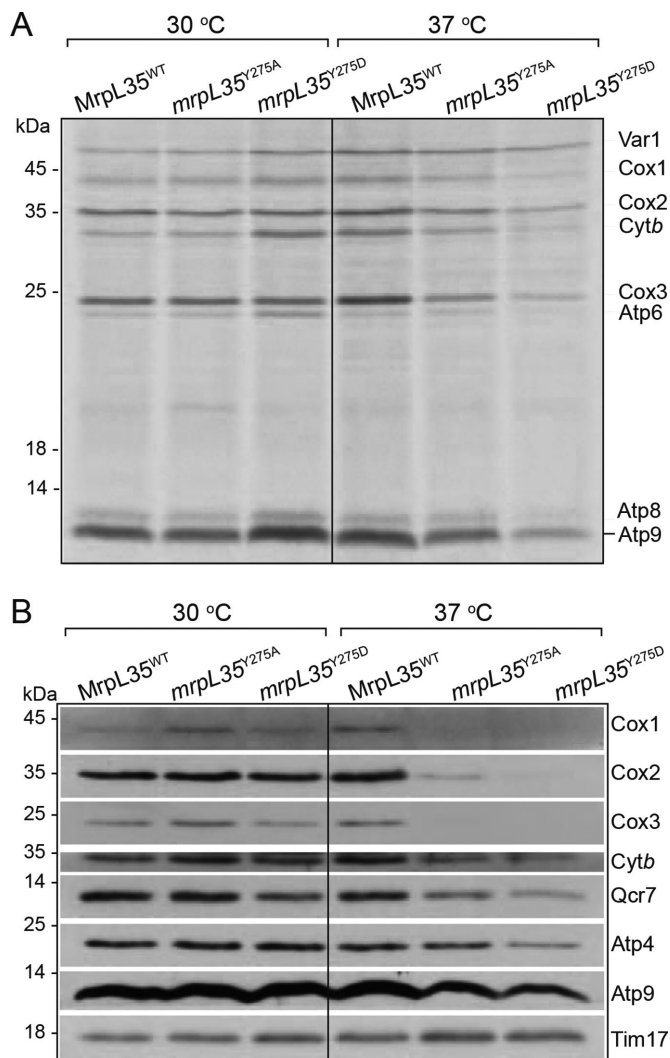
### Mutation of Y275 residue does not prevent mitoribosome translation

Using an in vivo radiolabeling approach in the presence of cycloheximide (to block cytosolic protein synthesis), we initially investigated if the translational capacity of the mitoribosome was inhibited in *mrpL35*<sup>Y275A</sup> and *mrpL35*<sup>Y275D</sup> mutants grown either at the permissive and nonpermissive temperature. In *S. cerevisiae*, the mitoribosomes synthesize eight proteins, Var1 (a small ribosomal subunit component) and OXPHOS subunits, Cox1, Cox2, and Cox3 of the COX (core subunits of the COX enzyme), and cytochrome *b* (of the cytochrome *bc*<sub>1</sub> enzyme) and F<sub>1</sub>F<sub>0</sub>-ATP synthase subunits, Atp6, Atp8, and Atp9. Both mutant strains displayed normal capacity for mitochondrial protein synthesis at the permissive temperature, and they retained the ability to synthesize also at 37°C (Figure 3A). Thus we conclude that the lack of OXPHOS-based growth of the *mrpL35* mutants at 37°C is not due to an absence of mitoribosomal translation at the elevated temperature. An overall reduction in total translation levels, in particular for the *mrpL35*<sup>Y275D</sup> mutant, was observed at 37°C, however, and may indicate a role for MrpL35 in mediating efficient mitochondrial translation or simply that the mitochondria of these cells were bioenergetically compromised after the prolonged growth period at the nonpermissive temperature, thus indirectly affecting their capacity for the energy-driven process of translation. When analyzed in isolated mitochondria, where exogenously added energy (ATP, GTP, and OXPHOS substrate NADH) were supplemented, the *mrpL35*<sup>Y275D</sup> mitochondria displayed normal level of protein synthesis (Figure 4A).

Western blotting of the protein extracts from the cells grown at the different temperatures indicated that the COX subunits, Cox1, Cox2, and Cox3, were strongly reduced in the *mrpL35*<sup>Y275A</sup> and more so in the *mrpL35*<sup>Y275D</sup> mutant strains grown at the nonpermissive temperature of 37°C (Figure 3B). Although the *mrpL35* mutants display the capacity to synthesize these COX subunits at 37°C, their strongly reduced steady-state protein amounts are indicative of a possible COX assembly defect at elevated temperatures, which would promote turnover of the newly synthesized Cox1, Cox2, and Cox3 proteins. Control experiments demonstrated that the preexisting COX complex assembled in the mutant at the permissive temperature did not display an enhanced proteolytic instability (i.e. relative to the wild-type control assayed in parallel) when shifted to 37°C (Supplemental Figure S1). These results indicate that if assembled at the permissive temperature, the COX complex in the mutant cells displays a similar resilience as that of the wild-type control when shifted to 37°C. We conclude therefore that the reduced levels of COX complex in the *mrpL35* mutants when grown at 37°C is indicative of a compromised ability to initially assemble the COX enzyme when the mitoribosomal protein synthesis and assembly processes occur at the nonpermissive temperature.

Although somewhat reduced, the levels of cytochrome *b* and Atp9 (and of Qcr7 and Atp4, two proteins dependent on the presence of cytochrome *b*- and Atp9-containing complexes, respectively,





**FIGURE 3:** The *mrpL35* mutants maintain the ability to translate even at nonpermissive temperature. (A) MrpL35 wild-type *mrpL35*<sup>Y275A</sup> and *mrpL35*<sup>Y275D</sup> mutant strains were grown on minimal media supplemented with galactose grown at either 30°C or 37°C for 24 h, as indicated. Mitochondrial translation capacity was monitored in vivo for 10 min with [<sup>35</sup>S]methionine and in the presence of cycloheximide (200 µg/ml). Equivalent amounts of cells were reisolated and proteins extracted and analyzed by SDS-PAGE, Western blotting, and autoradiography. (B) The indicated strains were grown as described in A, total cellular proteins were extracted, and steady-state levels of indicated mitochondrial proteins were measured following SDS-PAGE, Western blotting, and immunodecorated with antibodies indicated. Tim17 served as loading control.

for their stability), were clearly less affected in the *mrpL35* mutants than the COX subunits were (Figure 3B).

### The *mrpL35* Y275 mutants display a cytochrome c oxidase defect

To analyze the OXPHOS defect in the *mrpL35*<sup>Y275A</sup> and *mrpL35*<sup>Y275D</sup> mutant strains in more depth, mitochondria were isolated from cultures grown in galactose/lactate media at the permissive temperature of 30°C. First, in organello translation in the presence of [<sup>35</sup>S]methionine was performed, where we observed that the overall translational capacity of the mitochondrial ribosomes was not adversely affected in the *mrpL35* mutants (Figure 4A). The *mrpL35*<sup>Y275D</sup> mutant

mitochondria, however, displayed an altered translation profile with the levels of Cox1, cytochrome b, and Atp6 elevated, relative to the wild-type control, and the levels of Cox2 were somewhat reduced (Figure 4A, left panel). The enhanced levels of newly synthesized Cox1 protein in the *mrpL35* mutant mitochondria, in particular in the *mrpL35*<sup>Y275D</sup> mutant, indicated that the tightly regulated Cox1 synthesis pathway may be misregulated (Figure 4A, right panels).

Western blotting of the isolated *mrpL35* mutant mitochondrial preparations showed that the steady-state amounts of COX subunits Cox1, Cox2, and Cox3 were reduced, in particular in the *mrpL35*<sup>Y275D</sup> mitochondria, indicating that although synthesized at relatively normal (Cox2 and Cox3) or elevated (Cox1) levels in organello, the COX subunits fail to efficiently assemble in the *mrpL35* mutant (Figure 4B). Blue native gel electrophoresis (BN-PAGE) analysis of the individual OXPHOS complexes confirmed that the levels of assembled COX complex were strongly reduced in the *mrpL35* mutant mitochondria relative to the other OXPHOS complexes analyzed (Figure 4C, upper and lower panels). A parallel reduction in the level of COX enzyme activity was also measured in the *mrpL35* mutants (Figure 4D). In contrast, the levels of the cytochrome *bc*<sub>1</sub> and F<sub>1</sub>F<sub>o</sub>-ATP synthase complexes, as indicated by steady-state analysis of individual subunits, BN-PAGE analysis of the complexes and their levels and assembly states, and the measurement of the cytochrome *bc*<sub>1</sub> enzyme activity were not affected to the same extent as the COX complex was in the *mrpL35*<sup>Y275A</sup> and *mrpL35*<sup>Y275D</sup> mitochondria (Figure 4, B–D).

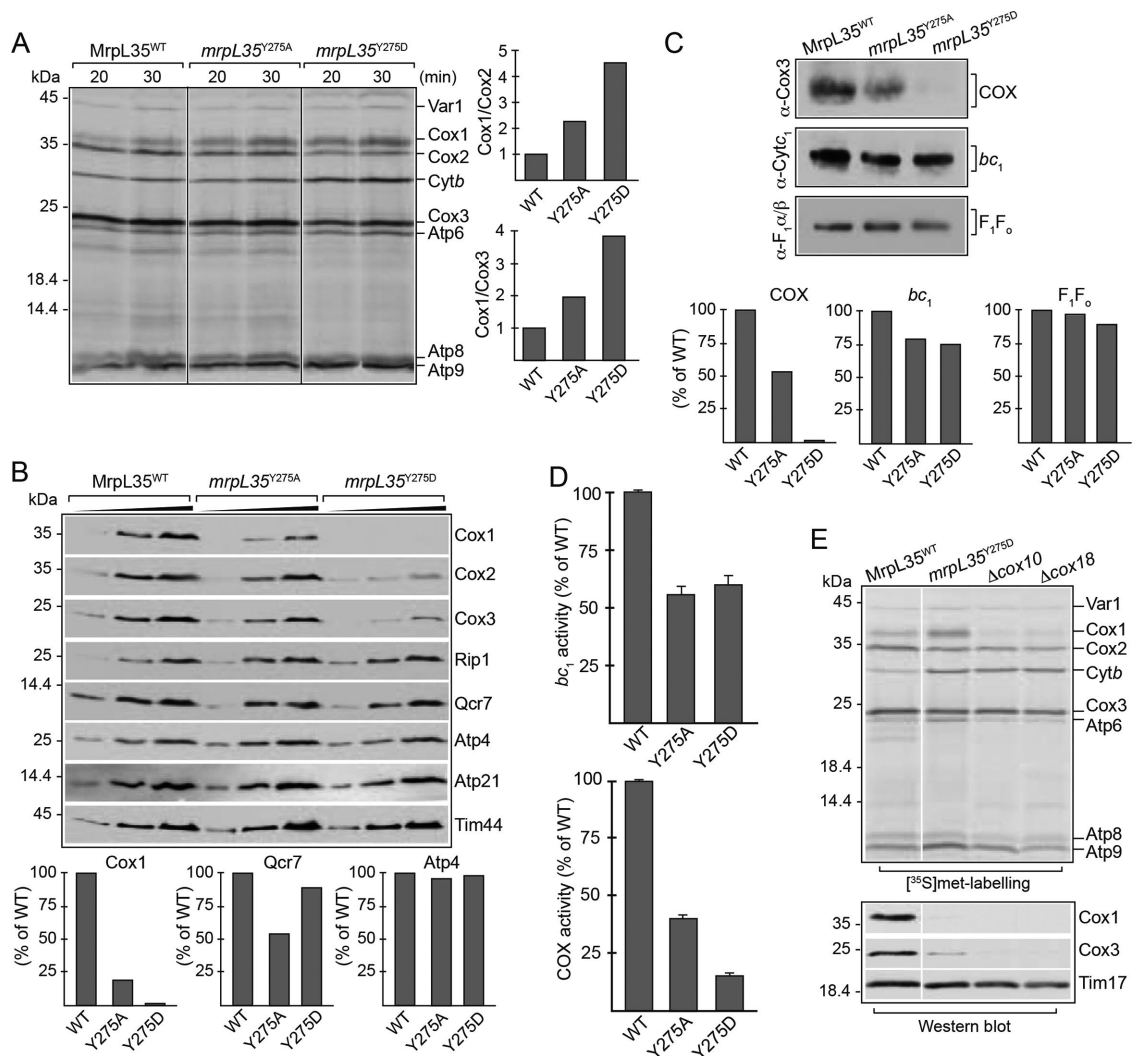
The ability of the *mrpL35*<sup>Y275</sup> mutants to continue to synthesize Cox1, and moreover at elevated levels relative to the wild-type MrpL35 control, was not anticipated given the COX complex assembly defect observed in these mitochondria. As outlined earlier, synthesis of Cox1 by the mitoribosomes is under strict feedback regulation with its assembly into the COX complex (Figure 1). Hence, as previously reported, Cox1 synthesis is characteristically reduced in COX assembly mutants, for example,  $\Delta$ cox10 and  $\Delta$ cox18 mitochondria, where the steady-state levels of COX subunits are strongly reduced (Figure 4E). In contrast, the *mrpL35*<sup>Y275</sup> mutant, despite having strongly reduced COX subunit levels, not only retains the ability but also displays enhanced capacity to synthesize the Cox1 protein (Figure 4E).

Taken together, these data demonstrate that mutation of the MrpL35 protein at residue Y275 does not cause an overall inhibition in mitochondrial protein translation but rather results in an altered translational profile and a notable misregulated Cox1 synthesis behavior and a COX assembly defect.

### Mutation in the Y275 residue of MrpL35 reveals a mitoribosomal connection to the COX assembly chaperones, Coa3 and Cox14

The observed elevated levels of Cox1 synthesis in the absence of assembly of the COX complex suggests that the Mss51 feedback regulatory system may be perturbed in the *mrpL35* mutant mitochondria. Analysis of the steady-state levels of Mss51, Cox14, Shy1, and Coa3 assembly factors indicated them to be present in the *mrpL35* mutant mitochondria at amounts similar to the wild-type MrpL35 control (Figure 5A). Thus the aberrant Cox1 synthesis and/or lack of efficient COX enzyme assembly observed in the *mrpL35* mutant mitochondria was not due to altered levels of these key COX regulatory/assembly factors.

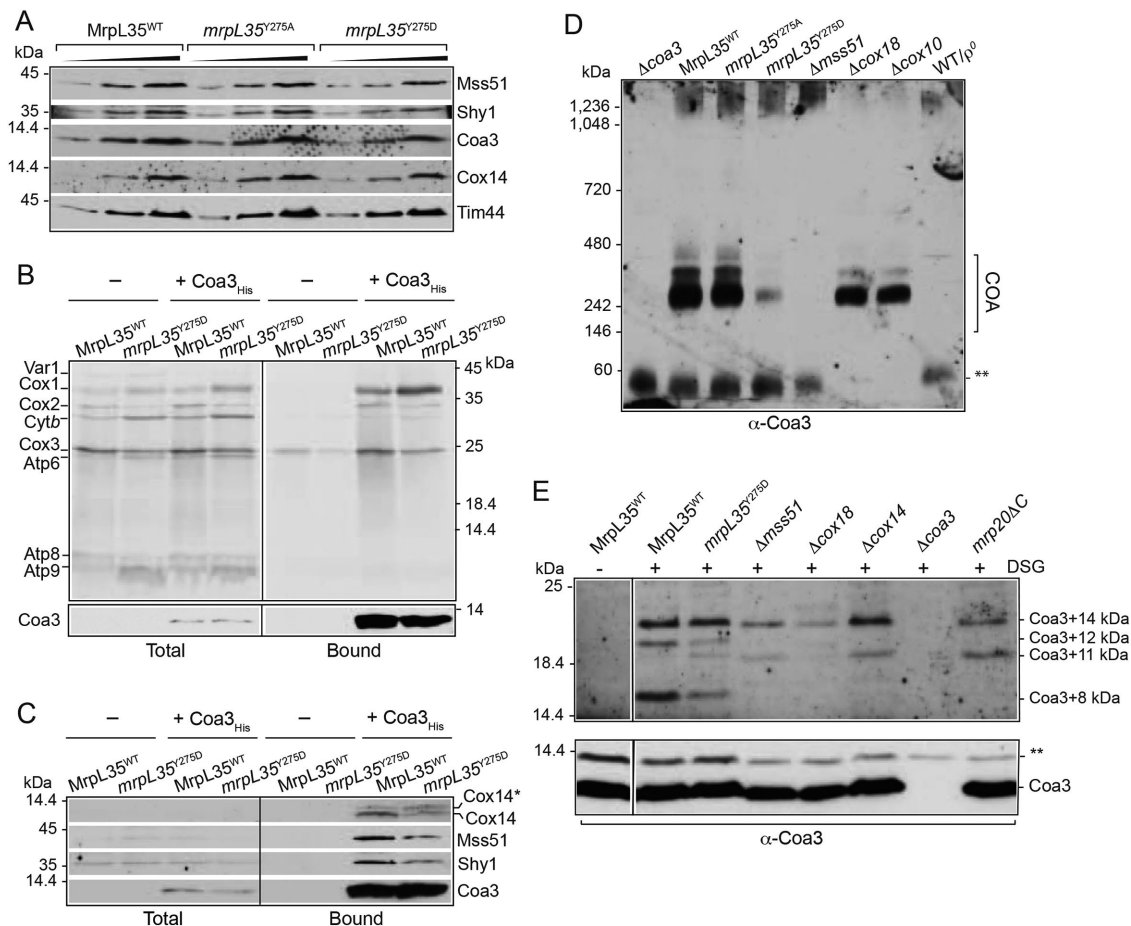
Following insertion into the membrane, newly synthesized Cox1 forms a COA assembly intermediate with Coa3 and other factors (including Cox14, Mss51, Coa1, and Coa2), which facilitates its subsequent assembly into the COX enzyme. We therefore tested



**FIGURE 4:** The *mrpL35* mutants are COX assembly defective and display an altered Cox1 synthesis profile. (A) Mitochondria were isolated from indicated strains, which had been grown at 30°C. Following a preincubation for 10 min at 37°C, in organello labeling in the presence of [<sup>35</sup>S]methionine was performed for 20 min and 30 min, as indicated, followed by a chase for 5 min in the presence of puromycin and cold methionine. Mitochondria were reisolated and samples analyzed by SDS–PAGE, Western blotting, and autoradiography (left panel). The amounts of newly synthesized Cox1 were quantified by subsequent phosphorimaging and expressed as a ratio of the levels of Cox2 and Cox3 synthesized in the same time period (30 min) (right panels). (B) Steady-state levels of the OXPHOS subunits in mitochondria (25, 50, 100 µg) isolated from indicated strains detailed in A were determined following SDS–PAGE, Western blotting, and immunodecoration. Tim44 was used as a loading control (upper panel). The amounts of Cox1, Qcr7, and Atp4 were quantified, normalized to Tim44 levels, and expressed as a percentage of the wild-type control (lower panels). (C) Mitochondria isolated from indicated strains were solubilized in DDM (0.6%) and subjected to BN–PAGE analysis, Western blotting, and immunodecoration with antibodies against the cytochrome *bc*<sub>1</sub> (*bc*<sub>1</sub>) subunit, cytochrome *c*<sub>1</sub> ( $\alpha$ -Cyt<sub>c1</sub>), Cox3 (COX), and F<sub>1</sub>F<sub>0</sub>-ATP synthase (F<sub>1</sub>F<sub>0</sub>) subunits ( $\alpha$ -F<sub>1</sub> $\alpha$ / $\beta$ ) (upper panels). The amounts of these complexes were quantified in the mutant mitochondria and expressed as a percentage of the control wild type (lower panels). (D) The specific activities of the cytochrome *bc*<sub>1</sub> (top panel) and COX (bottom panel) enzymes were measured in the *mrpL35* mutant and wild-type MrpL35<sup>WT</sup> control (WT) mitochondria, as described under *Materials and Methods* and expressed as a percentage of the wild-type control. Measurements were performed in triplicate and the statistical analysis and values were analyzed for statistical significance by Student's *t* test. Data are presented as means  $\pm$  SD of percentage of control. (E) In organello translation in the presence of [<sup>35</sup>S]methionine was performed for 30 min in the indicated mitochondria (upper panel) and further analyzed as described in A. The steady-state levels of indicated OXPHOS subunits were analyzed (lower panel), as described in B. Tim17 is used as a loading control.

whether radiolabeled Cox1 in the *mrpL35* mutants could be recovered in a complex with Coa3, and to do so we employed MrpL35 wild-type and *mrpL35*<sup>Y275D</sup> mutant mitochondria harboring a His-tagged Coa3 derivative (Coa3<sub>His</sub>). Following in organello synthesis, radiolabeled Cox1 protein was recovered in a specific manner with

the affinity-purified Coa3<sub>His</sub> species from both the control and *mrpL35* mutant mitochondria (Figure 5B). The observed association of Cox1 with Coa3<sub>His</sub> in the *mrpL35*<sup>Y275D</sup> mitochondria suggests that the COX assembly defect in this mutant is downstream of the Cox1 synthesis and Coa3-Cox1 assembly intermediate formation events.



**FIGURE 5:** *mrpL35* mutant exhibits alterations in Cox14 and Coa3 behavior. (A) Steady-state levels of proteins in isolated *mrpL35* mutant mitochondria were determined as described in Figure 4B. (B) In organello translation in the presence [<sup>35</sup>S]methionine was performed for 30 min in isolated MrpL35<sup>WT</sup> or *mrpL35*<sup>Y275D</sup> mitochondria harboring Coa3<sub>His</sub>, or not, as indicated. Following translation, mitochondria were solubilized with 0.6% DDM and affinity purification of Coa3<sub>His</sub> on Ni-NTA beads was performed, followed by SDS-PAGE, Western blotting, and autoradiography (upper panel). The recovery of Coa3<sub>His</sub> on the Ni-NTA beads was demonstrated by immunodecoration of the resulting blot with anti-His antibody (lower panel). Total, 5% of solubilized material; Bound, 100% of affinity purified material recovered from the Ni-NTA beads. (C) Same as B, except the in organello step was omitted, mitochondria were solubilized in 1% digitonin, and immunodecoration with indicated antibodies was performed following Western blotting. The Cox14 and novel, slower migrating Cox14\* species enriched with the affinity purified Coa3<sub>His</sub> complex is indicated. (D) BN-PAGE analysis of Coa3-containing complexes following solubilization of indicated isolated mitochondria with 1% digitonin. Western blotting and immunodecoration with Coa3 antisera was performed and protein complex that nonspecifically reacts with the Coa3 antiserum is indicated by \*\*. (E) Chemical cross-linking using DSG (+) or the mock treatment of solvent DMSO alone (-) was performed on mitochondria isolated from the indicated yeast strains. Following SDS-PAGE and Western blotting, decoration with Coa3- antisera was performed. Note the area encompassing the 14.4- to 25-kDa range of the gel is only shown (upper panel), as no Coa3-specific cross-link adducts were observed in the higher molecular mass range of the gel (not shown). The minus cross-linking (DMSO) control is only shown for the MrpL35<sup>WT</sup> sample, as this reflected the results for all other mitochondrial types analyzed, that is, the complete absence of any Coa3-immunoreactive material in this molecular weight range. The positions of the Coa3 monomer (lower panel and lighter exposure) and the four DSG-Coa3 adducts observed (upper panel) are indicated. A protein (14 kDa) that nonspecifically reacts with the Coa3 antiserum and observed in the lower panel, is indicated by \*\*.

Affinity-purified Coa3<sub>His</sub> complexes were also analyzed to ascertain whether the association of Coa3 with Mss51, Shy1, and Cox14 was perturbed in the *mrpL35* mutant (Figure 5C). Both Shy1 and Mss51 copurified with Coa3<sub>His</sub>; however, the levels of both proteins associated with the Coa3<sub>His</sub> complex from the *mrpL35*<sup>Y275D</sup> mitochondria were reduced, relative to those from the parallel wild-type control. Cox14 also detected in the affinity purified Coa3<sub>His</sub> complex from the *mrpL35*<sup>Y275D</sup> mitochondria; however, a significant

fraction of the protein recovered was present as a novel, slower migrating species, termed here Cox14\* (Figure 5C). A smaller proportion of this novel Cox14\* species (relative to mature Cox14 protein) was also observed in the Coa3<sub>His</sub> complex purified from wild-type MrpL35 control mitochondria. The enrichment of this novel Cox14\* species with the Coa3-complex from the *mrpL35*<sup>Y275D</sup> mitochondria would indicate that the behavior of the Cox14 protein is altered when the function of the MrpL35 protein is compromised.

BN-PAGE analysis indicated that the assembly of Coa3-containing complexes was significantly affected in the *mrpL35<sup>Y275D</sup>* mutant mitochondria (Figure 5D). In the wild-type MrpL35 control mitochondria, Coa3 was recovered in COA complexes of ~240–480 kDa in mass, which have been proposed to also include the Cox1–Coa3 assembly intermediates of the COX complex (McStay *et al.*, 2013; Mayorga *et al.*, 2016). The level of these Coa3–COA complexes was strongly reduced in the *mrpL35<sup>Y275D</sup>* mutant mitochondria, indicating that despite the fact that newly synthesized Cox1 could associate with Coa3 (Figure 5B), the assembly state of the predominant Coa3–COA-containing complexes was strongly perturbed in this mutant. As the steady-state levels of the Coa3 protein were not reduced in the *mrpL35<sup>Y275D</sup>* mitochondria (Figure 5A), we conclude that the majority of the Coa3 protein in the mutant must be present in complexes smaller than 60 kDa under these conditions and thus not retained in the BN-PAGE gel. In contrast to the *mrpL35<sup>Y275D</sup>* mutant, the assembly state of the Coa3 complex did not appear to be affected in the *mrpL35<sup>Y275A</sup>* mutant mitochondria. As these mitochondria had both been isolated from cells grown at the permissive temperature, it appears that the phenotype of the *mrpL35<sup>Y275D</sup>* mutant is more severe and penetrated into cells grown at 30°C, in contrast to the *mrpL35<sup>Y275A</sup>* mutant. Consistently reduced COX levels were observed in these isolated *mrpL35<sup>Y275D</sup>* mitochondria and to a lesser extent in the *mrpL35<sup>Y275A</sup>* mitochondria (Figure 4, B–D). The aberrant assembly state of Coa3 observed in the *mrpL35<sup>Y275D</sup>* mutant was not simply due to the absence of an assembled COX enzyme, as Coa3-containing COA complexes were detected in mitochondria from the  $\Delta\text{cox10}$  and  $\Delta\text{cox18}$  COX assembly mutants, analyzed in parallel. The perturbed assembly of the Coa3–COA complexes in the *mrpL35<sup>Y275D</sup>* mutant mitochondria mirrored, however, that observed in the  $\Delta\text{mss51}$  and wild-type/ $\rho^0$  mitochondria (Figure 5D). In contrast to these mitochondria where Cox1 synthesis is absent (Barrientos *et al.*, 2004), the *mrpL35<sup>Y275D</sup>* mitochondria retain the capacity to synthesize Cox1 and form Cox1–Coa3 intermediates. We conclude therefore that in the ribosomal mutant *mrpL35<sup>Y275D</sup>*, the further (or stable) assembly of Coa3–Cox1 into larger Coa3–COA complexes was perturbed.

Finally, the molecular environment of Coa3 in intact *mrpL35* mutant mitochondria was independently probed with the chemical cross-linking agent disuccinimidyl glutarate (DSG) (Figure 5E). In the wild-type MrpL35 control mitochondria, endogenous Coa3 (8 kDa) was observed to form three small adducts in the presence of DSG, which corresponded to Coa3 cross-linked to partners of ~14, 12, and 8 kDa, respectively. A reduction in the formation of the smaller two of the Coa3-adducts, and a parallel acquisition of a novel fourth Coa3-adduct, corresponding to Coa3 cross-linked to an 11-kDa protein, was observed instead in the *mrpL35<sup>Y275D</sup>* mitochondria. The Coa3-adduct profile in the *mrpL35<sup>Y275D</sup>* mitochondria bore similarities to that obtained in the  $\Delta\text{mss51}$  and  $\Delta\text{cox14}$  mutant mitochondria, analyzed in parallel, where loss of the Coa3+12 kDa and Coa3+8 kDa adducts, and the gain of the novel Coa3+11 kDa adduct, was observed (Figure 5E). The altered Coa3-adduct profile in these mitochondria may be indicative of their COX-assembly defective phenotype, as it was also observed in  $\Delta\text{cox18}$  mitochondria and also in the ribosomal mutant *mrp20 $\Delta$ C* mitochondria, which are defective in mitochondrial translation and, consequently, COX assembly. Although the identities of the Coa3 cross-linked partners in these adducts are unknown at this time, the observed altered Coa3 cross-linking pattern in the *mrpL35<sup>Y275D</sup>* mitochondria is further evidence of an aberrant Coa3 condition in this mitoribosomal mutant.

In summary, the molecular environment of Coa3, and the nature of the Cox14 species associated with it, is altered in the *mrpL35*

mutant. These perturbations may impact the productive nature of the Cox1-assembly intermediate formed with the Coa3 and thus the further assembly of Cox1 into the COX enzyme.

### Mutation of Y275 residue perturbs the close relationship of MrpL35 with Mrp7 protein

The cryo-EM structures of yeast mitoribosomes indicate MrpL35 (mL38) to be a peripheral component of the central protuberance where it is embraced by mitospecific elements of the Mrp7 (bL27) protein. To gain more insight into MrpL35 and the possible effect of the Y275 mutation on its interactions with partner proteins, we affinity purified His-tagged wild-type and mutant MrpL35 derivatives, which had been expressed in the ribosome assembly yeast mutant, *mrp20 $\Delta$ C* (Kaur and Stuart, 2011). Stable assembly of the 54S particles has been reported to be defective in this *mrp20 $\Delta$ C* mutant, and instead ribosomal proteins can be purified in subassemblies or subcomplexes, enabling their compositional analysis (Kaur and Stuart, 2011). The affinity purified wild-type MrpL35<sub>His</sub> was recovered in a complex with other proteins, which were shown either through specific antibody decorations or mass spectrometry analysis (Supplemental Table S1) to include Mrp7 (bL27), MrpL36 (bL31), and MrpL7 (uL5) and two mitospecific proteins, MrpL17 (mL46) and MrpL28 (mL40) (Figure 6A). Western blotting with available antibodies against Mrp7 (bL27) and MrpL36 (bL31) confirmed their specificity of their association with MrpL35<sub>His</sub>, as they were not detected in the Mrp20 ribosomal exit pore subcomplex, purified parallel via MrpL25<sub>His</sub> (mL59), a known component of the Mrp20 exit pore subcomplex (Kaur and Stuart, 2011) (Figure 6, B and C). Reciprocal affinity purifications of MrpL7<sub>His</sub>, MrpL17<sub>His</sub>, and Mrp7<sub>His</sub> from the *mrp20 $\Delta$ C* mutant strain also confirmed their specific association with Mrp7, MrpL36, and MrpL35 proteins (Figure 6B). The ability of Mrp7 to be copurified with the His-tagged derivatives of *mrpL35<sup>Y275A</sup>* and *mrpL35<sup>Y275D</sup>* proteins was strongly reduced relative to the wild-type MrpL35<sub>His</sub> control (Figure 6C). The recovery of MrpL36 (bL31) in the mutant *mrpL35* subcomplexes was also reduced, albeit not to the same extent as the Mrp7 protein. These results indicate that the interaction of MrpL35 with Mrp7 (and possibly other components of the central protuberance subcomplex) is perturbed through the mutation of the Y275 residue, such that it does not remain associated with the Y275 mutated *mrpL35* protein under these detergent solubilization and purification conditions.

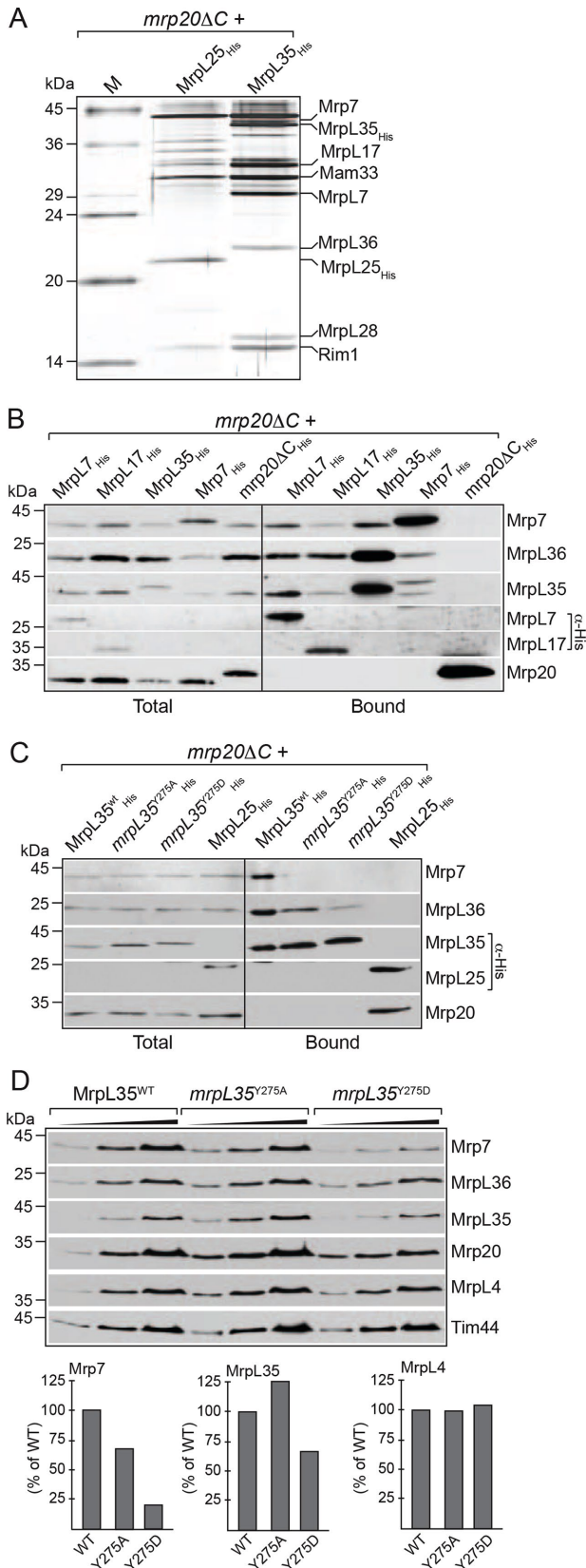
Steady-state analysis of protein levels indicated that a reduction in Mrp7 levels was observed in the *mrpL35<sup>Y275D</sup>* mutant (Figure 6D, upper and lower panels). Mutation of the Y275 to Ala(A) residue did not greatly compromise the stability of the MrpL35 protein, though a decrease in MrpL35 steady-state levels was seen with the *mrpL35<sup>Y275D</sup>* mitochondria. In contrast to Mrp7, the levels of MrpL36, another central protuberance constituent, or other 54S particle proteins, for example, Mrp20 and MrpL4, were relatively unaffected in the *mrpL35* mitochondria (Figure 6D).

Taken together, we conclude that the MrpL35 can assemble into a central protuberance subcomplex with the MrpL36, Mrp7, MrpL7, MrpL17, and MrpL28 proteins and that the association with, and possibly stability of, Mrp7 is perturbed through the mutation of the Y275 residue in MrpL35.

## DISCUSSION

Mitochondrial translation and the assembly of the OXPHOS complexes are processes of critical importance to aerobic metabolism and cell viability. Mutation and impairment of mitoribosome capacity has been shown to be directly associated with a range of





**FIGURE 6:** Mutation of MrpL35 perturbs the association with and stability of Mrp7. (A) Isolated *mrp20ΔC* mitochondria harboring His-tagged derivatives of wild-type MrpL35 or MrpL25, as indicated, were solubilized with 1.0% Triton X-100 and subjected to affinity purification protocol with Ni-NTA beads. The Ni-NTA purified material was further analyzed by SDS-PAGE and silver staining. The positions

pathophysiological neuromuscular conditions (Lightowers *et al.*, 2014; De Silva *et al.*, 2015). Investigating the molecular details of mitoribosome activity, regulation, and coupling of translation to the downstream OXPHOS assembly events is therefore critical for our understanding of normal and dysfunctional cellular metabolism. MrpL35 (mL38) is a mitospecific component of the yeast mitoribosomal 54S particle. Located in the central protuberance region, aspects of MrpL35 and the mitospecific domain of its partner protein Mrp7 (bL27) form an externally exposed element of this region of the mitoribosome (Amunts *et al.*, 2014; Desai *et al.*, 2017). Mutation of residue Tyr(Y)275 of the MrpL35 protein was found to compromise its ability to support aerobic growth and OXPHOS assembly, in particular at elevated temperatures, with the exchange for the negatively charged Asp(D) residue being more deleterious than Ala(A). Surprisingly, the molecular basis for the defective OXPHOS function in the *mrpL35* mutants was not due to a primary inhibition in mitochondrial protein translation, but rather our OXPHOS complex analysis supports a pronounced COX assembly defect in these *mrpL35* mutants. The consequences of the *mrpL35* mutation for the assembly of the cytochrome *bc<sub>1</sub>* or *F<sub>1</sub>F<sub>o</sub>*-ATP synthase complexes, which both also contain mitoribosome synthesized subunits, were not as pronounced as they were for the COX complex assembly. As the *mrpL35* mutants retained the capacity to synthesize the mitochondrially encoded COX subunits, the strong reduction in COX complex steady-state levels indicates that a defect may lie in a posttranslational step required for the assembly of the COX complex. Furthermore, the elevated levels of Cox1 synthesized in the mutants, despite their gross COX assembly defect, points to a breakdown in the Mss51/Cox14/Coa3-dependent feedback loop used to negatively regulate Cox1 synthesis under conditions when further assembly of Cox1 into a COX complex is perturbed.

Our data indicate that newly synthesized Cox1 can form the Cox1–Coa3-containing assembly intermediate in the *mrpL35* mutant, but the further progression of Cox1 assembly from this intermediate into the Coa3–COA intermediate complex, and thus subsequently to the fully assembled COX enzyme, are defective. Both Cox14 and Coa3 act as negative regulators of the Cox1 synthesis regulatory pathway (Mick *et al.*, 2010; Fontanesi *et al.*, 2011). Thus the elevated levels of Cox1 synthesized in the *mrpL35* may also be indicative of a breakdown in the function of the Cox14 and/or Coa3 proteins. We observed the presence of a novel form of

of proteins identified to copurify with the MrpL35<sub>His</sub> and MrpL25<sub>His</sub> proteins are indicated. Note Rim1 and Mam33 were found to copurify with both the MrpL35<sub>His</sub> and MrpL25<sub>His</sub> subcomplexes, which was confirmed by Western blotting (Rim1) and reciprocal Mam33<sub>His</sub> purification studies (not shown). M, molecular mass marker proteins (kDa) used for SDS-PAGE. (B) Affinity purification of His-tagged MrpL7, MrpL17, MrpL35, Mrp7, and *mrp20ΔC* derivatives from *mrp20ΔC* mitochondria was performed as described in A with the exception that following SDS-PAGE, Western blotting and immunodecoration with indicated specific antibodies was performed. Total, 5% of solubilized material; Bound, 100% of affinity purified material recovered from the Ni-NTA beads. (C) Affinity purification of His-tagged MrpL35<sup>WT</sup>, *mrpL35 Y275A*, or *mrpL35 Y275D* derivatives or MrpL25<sub>His</sub> from *mrp20ΔC* mitochondria was performed and analyzed as described in B. (D) The steady-state amounts of mitoribosomal proteins were determined by analyzing indicated mitochondria, as described in Figure 4B (upper panel). The amounts of Mrp7, MrpL35, and MrpL4 were quantified, normalized to Tim44, and expressed as a percentage of the wild-type control (lower panel).

Cox14, the Cox14\* species enriched with the purified Coa3 complex and observed both in the wild type, and at greatly increased proportions in Coa3 complexes from the *mrpL35* mutant. The Cox14\* species has not been reported in the literature to date and may represent a novel, post-translationally modified form of the Cox14 protein. The abundance of Cox14\* with Coa3 complex in the *mrpL35* mutant would indicate its recycling to its regular Cox14 status is perturbed in this mitoribosomal mutant, which in turn may stall the progression of Cox1–Coa3 assembly intermediate. Both BN–PAGE and chemical cross-linking data further illustrate that the stable assembly state of the Coa3–COA complexes is defective in the *mrpL35*<sup>Y275D</sup> mutant. Although the nature of this modification is currently unknown, we speculate here that the formation of Cox14\*, and its possible recycling to Cox14, may influence the productive nature of the Cox1–Coa3 complex as an assembly intermediate and/or its ability to recruit or retain Mss51 and thus Cox14's ability to negatively control the Cox1 synthesis levels.

MrpL35, like the other members of the mL38 family, contains a PEBP-like domain in its C-terminal region. PEBP proteins are a highly diverse group of proteins (i.e., not exclusively mitoribosomal proteins) involved in signaling pathways. PEBP-domains form anion ligand-binding pockets, which can interfere with kinase-based signaling pathways when they form complexes with anionic ligands (Banfield *et al.*, 1998; Banfield and Brady, 2000; Yousuf *et al.*, 2014). The cryoEM structural analyses of mitoribosomes indicates that the PEBP-region of MrpL35/mL38 proteins is exposed to an exterior surface of the CP (Amunts *et al.*, 2014, 2015; Greber *et al.*, 2015; Desai *et al.*, 2017). This aspect of the MrpL35 may represent a binding site for an external regulatory ligand or feature of a nonribosomal protein (e.g., a phosphorylated residue) in the central protuberance of the mitoribosome. Residue Y275 is conserved in all MrpL35 (mL38) proteins and is located in the PEBP-like domain of MrpL35 (mL38). It is unclear whether the PEBP-like domain of MrpL35 binds anionic ligands, but the conserved nature of this domain, and the detrimental effect of the Y275 mutation, may suggest a role for MrpL35 in a signaling pathway involving the mitoribosome. It is important to note that the region of MrpL35 where residue Y275 is located is directly adjacent to the fungal-specific, mitospecific domain of Mrp7 (bL27), which embraces the PEBP-like domain of MrpL35 (Amunts *et al.*, 2014; Desai *et al.*, 2017). We observed that mutation of residue Y275 destabilized the association of MrpL35 with its partner protein Mrp7 and led to a reduction in Mrp7 levels in the mutant mitochondria. Some of the reported phenotypes in the *mrpL35* mutants could therefore be attributed to a disturbance in the relationship of MrpL35 with Mrp7 protein rather than (or in addition to) a defect in MrpL35.

How does mutation of MrpL35, a mitoribosomal protein of the central protuberance, affect the COX assembly pathway and at the level of the Coa3/Cox14 proteins? One possibility is a physical connection between elements of the central protuberance, involving the PEBP-domain of MrpL35 and/or Mrp7, and the COX assembly chaperones, such as Cox14 and Coa3 proteins in the membrane, which is important for the correct operation of the COX assembly chaperones. The mitoribosomes are physically tethered to the inner membrane; however, to date this association has been shown to involve aspects of the 54S particle in the vicinity of the polypeptide exit channel and thus distanced from the central protuberance region (Jia *et al.*, 2009; Gruschke *et al.*, 2010; Haque *et al.*, 2010; Pfeiffer *et al.*, 2015). It is therefore unlikely that MrpL35 in context of the translating ribosome can have a direct physical connection to the Cox14/Coa3 proteins; however, we cannot rule out that MrpL35

(and/or Mrp7) independent of the assembled ribosome (e.g., alone or in context of a central protuberance subcomplex) could do so. Furthermore, it is also possible that MrpL35 and/or components of the central protuberance may serve to recruit Mss51 to the ribosome and thus modulate its capacity to act as a Cox1 translational activator; however, we have not obtained data to support a Mss51–MrpL35 (or Mrp7) interaction to date. Another possibility is that MrpL35, through its PEBP-like domain, serves to affect COX assembly by recruiting a common anionic factor important for signaling between the ribosome and the Coa3–Cox14–Mss51 proteins and thereby to balance the Cox1 synthesis rate with the downstream chaperoned assembly events. Mutation of the Y275 residue may impair this ligand binding and thus promote more Cox1 synthesis in a manner that is uncoupled from the downstream assembly events. Finally, the mutation of the MrpL35 protein could also cause alterations in the ribosomal structure/organization that affects the accessibility of the newly synthesized Cox1 protein to the Mss51 protein following its insertion into the membrane. Mss51 binds to the extreme C-terminal tail of the newly synthesized Cox1 protein, a step required to recruit the chaperone into the Cox1–Coa3 assembly intermediate (Shingu-Vazquez *et al.*, 2010; Garcia-Villegas *et al.*, 2017). Factors that hinder the access to the C-terminal region of the newly synthesized Cox1 may influence the recruitment of Mss51 to the Cox1–Coa3 complex. Delayed recycling of the ribosome at the termination of translation and/or impaired disengaging of the ribosome from Oxa1, the membrane site of cotranslational insertion of proteins, may both prevent Mss51's timely access to the C-tail of Cox1. Interestingly, yeast mutants deficient in the ribosome-recycling factor (Rrf1), which promotes ribosomal subunit dissociation and recycling, display an OXPHOS assembly defect that was not caused by an inhibition in mitoribosomal translation *per se* (Ostojic *et al.*, 2016). Rather, similarly to the *mrpL35* mutants, the *rrf1* mutant mitochondria displayed an altered profile of synthesis products, including increased levels of newly synthesized Cox1, and reduction in COX complex assembly levels. However, in contrast to the *mrpL35* mutants, the yeast *rrf1* mutants also exhibit a strong impairment in the assembly of the F<sub>1</sub>F<sub>0</sub>–ATP synthesis complex, in addition to the observed COX assembly defect (Ostojic *et al.*, 2016). The molecular defect underpinning the COX assembly deficiency in the *mrpL35* Y275 mutant would thus appear to be different from that *rrf1* mutant and ribosome recycling where a more pleiotropic OXPHOS assembly defect is described. Taking this together with other *mrpL35* results reported here, namely i) the maintained ability to form the Cox1–Coa3 intermediate, ii) the enrichment of Cox14\* with Coa3, and iii) the altered assembly state of the Coa3 complex (as evidenced by chemical cross-linking and BN–PAGE approaches), we favor that the *mrpL35*<sup>Y275</sup> mutation does not cause a general ribosome translation inhibition defect but rather has direct consequences for functional integrity of the Mss51, Coa3, or Cox14 proteins.

In summary, our findings here unexpectedly highlight that components of the mitoribosome, MrpL35, and possibly its closely associated Mrp7 protein, play a role in regulating the COX assembly process, in particular by influencing the molecular environments of the Mss51, Coa3, and Cox14 proteins. Furthermore, this study also represents the first report of a novel Cox14 species, Cox14\* associated with the Cox1–Coa3 complex, suggesting that Cox14 may be regulated by a posttranslational event. Future experimentation will ascertain the nature of the Cox14 modification and the role it plays in Cox14's ability to support Coa3, Mss51, and the productive Cox1-assembly pathway and how this may involve the mitoribosomes and MrpL35.

## MATERIALS AND METHODS

### Yeast strains and growth conditions

All *S. cerevisiae* strains used in this study are in the haploid W303-1A genetic background (W303-1A, *mat a leu2, trp1, ura3, his3, ade2*). They include the  $\Delta mrpL35$  (MRPL35::KAN) mutant and those harboring plasmid borne copies of the wild-type MRPL35 gene or *mrpL35* mutant derivatives, created through a plasmid shuffling approach, as described below. The W303-1A-based strain *mrp20 $\Delta$ C* + GAL10-*mrp20 $\Delta$ C* (termed here *mrp20 $\Delta$ C*) was also employed (Kaur and Stuart, 2011). Yeast strains were cultured using standard protocols on either minimal medium supplemented as appropriate with uracil, tryptophan, leucine, adenine, histidine, or on full YP media, as indicated. Mitochondria were isolated from cultures grown at 30°C in minimal medium and containing 0.5% lactate and 2% galactose as carbon sources.

### Generation of *mrpL35* mutants

The yeast chromosomal MRPL35 open reading frame (ORF) was replaced with the KAN cassette in a haploid W303-1A yeast strain harboring a wild-type MRPL35 (MRPL35<sup>WT</sup>) genomic insert (corresponding to the MRPL35 ORF plus 650 base pairs 5' and 300 base pairs 3') cloned into a pRS316 URA3-based plasmid. The resulting strain then transformed with the centromeric plasmid pRS413 (HIS3) harboring either a similar genomic fragment containing the wild-type or mutant *mrpL35* derivative or no insert (i.e., the empty plasmid). Following plasmid shuffling process, involving growth on 5'FOA to promote the loss of the pRS316(URA3)-MRPL35<sup>WT</sup> plasmid,  $\Delta mrpL35$  transformants harboring only the pRS413(HIS3) plasmids were selected. The resulting strains ( $\Delta mrpL35$ +MRPL35<sup>WT</sup> or  $\Delta mrpL35$ +*mrpL35*<sup>Y275</sup> derivatives) are referred to here as the  $\Delta mrpL35$  null or MrpL35<sup>WT</sup> (wild-type) and the *mrpL35*<sup>Y275A</sup>, *mrpL35*<sup>Y275D</sup>, or *mrpL35*<sup>Y275F</sup> mutant strains. Mutations in the Y275 residue of MRPL35 gene in the pRS413(HIS3)-MRPL35 plasmid were generated using a PCR site-directed mutagenesis strategy.

### Generation of His-tagged MrpL35 and Coa3 derivatives

The Yip351-LEU2 vector containing the open reading frame encoding either the wild-type MrpL35 or *mrpL35*<sup>Y275D</sup> derivatives or the Coa3 protein as C-terminal His<sub>12</sub>-tagged proteins and downstream of the galactose-inducible GAL10 promoter were created. The resulting plasmids were linearized with BstEII and integrated into the *leu2* loci of indicated yeast strains and LEU<sup>+</sup> transformants were selected. The presence of His-tagged derivative proteins in isolated mitochondria was confirmed using Western blot approaches with a His-specific antibody.

### Affinity purification of His-tagged proteins

Isolated mitochondria (200  $\mu$ g protein) harboring the His-tagged protein derivatives were solubilized in lysis buffer (160 mM KCl, 20 mM HEPES-KOH, 10 mM MgCl<sub>2</sub>, 0.5 mM phenylmethylsulfonyl fluoride [PMSF], pH 7.4), with 1.0% Triton X-100, 0.6% *n*-dodecyl  $\beta$ -D-maltoside (DDM), or 1% digitonin, as indicated, for 30 min on ice. Following a clarifying spin, Ni-NTA purification of His-tagged proteins was performed as previously described (Jia *et al.*, 2007).

### BN-PAGE analysis

BN-PAGE analysis of digitonin (1%) or DDM (0.6%) solubilized mitochondrial extracts (30  $\mu$ g protein) was performed using Invitrogen Nu-PAGE gradient (3–12%) gels according to the manufacturer's protocol and followed by Western blotting and antibody decoration, as indicated.

### Miscellaneous

All experiments presented were performed at least three times; results were consistently reproducible, and representative examples of data are given in the figures. Spectral measurements of specific enzyme activities of the cytochrome *bc*<sub>1</sub> and COX complexes were performed as previously described (Tzagoloff *et al.*, 1975). In organello translation with [<sup>35</sup>S]methionine and chemical cross-linking with DSG (7.7 Å, homobifunctional, amine-specific), were performed as previously described (Hell *et al.*, 2001). Mitochondrial isolation, protein determination, and SDS-PAGE were performed as previously described (Dienhart and Stuart, 2008). The Cox1 and Cox3 antibodies used in this study were commercially obtained (Cox1: Molecular Probes, anti-yeast Cox1, mouse monoclonal 11D8-B7, lot# 6251-1; Cox3: Invitrogen/Novex Anti-Cox3 monoclonal, 459300, lot#H3578). All other antibodies used were rabbit polyclonal against the respective purified yeast proteins and generated either in the Stuart lab or received as gifts (see *Acknowledgments*). Antigen-antibody complexes were detected by horseradish peroxidase-coupled secondary antibodies and chemiluminescence detection on x-ray films. Resulting signals were quantified following scanning using Image Studio software.

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