Mitochondrial contact site and cristae organizing system (MICOS) machinery supports heme biosynthesis by enabling optimal performance of ferrochelatase

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ABSTRACT

Heme is an essential cofactor required for a plethora of cellular processes in eukaryotes. In metazoans the heme biosynthetic pathway is typically partitioned between the cytosol and mitochondria, with the first and final steps taking place in the mitochondrion. The pathway has been extensively studied and its biosynthetic enzymes structurally characterized to varying extents. Nevertheless, understanding of the regulation of heme synthesis and factors that influence this process in metazoans remains incomplete. Therefore, we investigated the molecular organization as well as the physical and genetic interactions of the terminal pathway enzyme, ferrochelatase (Hem15), in the yeast Saccharomyces cerevisiae. Biochemical and genetic analyses revealed dynamic association of Hem15 with Mic60, a core component of the mitochondrial contact site and cristae organizing system (MICOS). Loss of MICOS negatively impacts Hem15 activity, affects the size of the Hem15 high-mass complex, and results in accumulation of reactive and potentially toxic tetrapyrole precursors that may cause oxidative damage. Restoring intermembrane connectivity in MICOS-deficient cells mitigates these cytotoxic effects. These data provide new insights into how heme biosynthetic machinery is organized and regulated, linking mitochondrial architecture-organizing factors to heme homeostasis.

1. Introduction

Heme is an essential cofactor and signaling molecule required for diverse physiological processes, from the mitochondrial electron transport chain to oxygen transportation through the bloodstream [1–4]. Most metazoans synthesize heme through a highly conserved and well-characterized eight-step pathway [5]. In the first step, the mitochondrial enzyme aminolevulinic acid synthase (Alas; ALAS in humans; Hem1 in yeast) catalyzes the condensation of glycine and succinyl-CoA to form aminolevulinic acid (ALA), which is then transported out of the mitochondrion by an as-yet-uncharacterized transporter. Once in the cytosol, two ALA molecules are condensed into a monopyrrole,
porphobilinogen, by the enzyme porphobilinogen synthase (PbgS); subsequently, four porphobilinogen molecules are joined together by hydroxymethylbilane synthase (HmbS) to form the linear tetrapyrrole hydroxymethylbilane (HMB). HMB then undergoes cyclization to form uroporphyrinogen III (catalyzed by uroporphyrinogen synthase, Uros) and decarboxylation of its pyrrole acetate side chains (catalyzed by uroporphyrinogen decarboxylase, Urod) to yield coproporphyrinogen III. In Saccharomyces cerevisiae, coproporphyrinogen III is converted to protoporphyrinogen IX by coproporphyrinogen oxidase (Cpox) in the cytosol [6,7], while in humans this reaction occurs in the mitochondrial intermembrane space (IMS) [8,9].

Protoporphyrinogen IX is transported into the mitochondrial matrix and converted to protoporphyrin IX (PPIX) by the matrix-localized enzyme protoporphyrinogen oxidase (Ppox) [10]. How protoporphyrinogen IX is transported across the inner mitochondrial membrane (IM) is currently unclear though evidence suggests that the protoporphyrinogen transporter TMEM14C facilitates this process during erythroid cell maturation [11]. The final step of heme synthesis is catalyzed by ferrochelatase (Fech; FECH in humans; Hem15 in yeast), which resides on the matrix side of the IM [12] and mediates the insertion of ferrous iron into PPIX, thereby yielding heme (also known as protoporphyrin IX or heme b).

Even though the proteins involved in heme synthesis have been structurally characterized and in some cases examined mechanistically [3,13], the processes by which the relative rates of heme synthesis are regulated remain obscure. Understanding of this regulation is mainly limited to the transcriptional level in multicellular organisms with erythrocytes. In these organisms, most heme production occurs in the developing erythrocyte, and specific transcription factors are known to play a role in regulating expression of all the biosynthetic enzymes [14]. The mechanisms of regulation of heme synthesis for many other cells in these organisms and in unicellular eukaryotes like S. cerevisiae, all of which require less heme, remain uncharacterized.

In terms of enzyme activity, the rate-limiting step of the pathway in mammals is the first step (catalyzed by ALAS), with the last step (catalyzed by FECH) being the second regulatory point of the mammalian pathway [14]. In S. cerevisiae, two other pathway enzymes, PbgS and HmbS, have been proposed to catalyze the rate-limiting steps [7,15]. Recent data suggest additional levels of regulation at the post-translational level [16,17].

Proteomic studies in mammalian cells show that the mitochondrial heme biosynthetic machinery exists as a large supercomplex, termed the heme metabolon [17–19]. FECH was identified as a component of a multimeric assembly that includes ALAS and PPOX, the porphyrinogen transporter TMEM14C, and succinyl-CoA synthase [17,18]. The mitochondrial iron import mediator [20] and two ATP-binding cassette proteins (ABCB7 and ABCB10) [20–22] have also been shown to bind to FECH. Additional candidate interaction partners of FECH include factors that mediate IM dynamics and ultrastructure [17,19], most notably components of the conserved mitochondrial cristae and cristae organizing system (MICOS). As suggested by its name, MICOS maintains components of the IM and the outer mitochondrial membrane (OM) [23–25] and has been postulated to facilitate bidirectional transport of hydrophobic molecules such as phospholipidic acid [26] and coenzyme Q biosynthetic intermediates [27]. However, the physiological significance of its connection to FECH remains unaddressed.

Here, a yeast genetic model was employed to investigate the putative molecular interaction of Fech (Hem15 in yeast) with MICOS. Data from biochemical and genetic analyses are consistent with a dynamic association of Hem15 with the core MICOS component Mic60. Loss of MICOS negatively impacts Hem15 activity and results in accumulation of cytotoxic pathway intermediates. These data provide insights into how the heme biosynthetic machinery is organized and supported, linking mitochondrial architecture to heme homeostasis.

2. Results

2.1. Functional Hem15 forms a high-mass complex

Hem15 is known to exist as a homodimer [28,29], but its protein–protein interaction has not been systematically studied. To that end, an antibody was developed against Hem15, which specifically recognizes S. cerevisiae Hem15 (Fig. 1A). Using this antibody and purified Hem15 as a reference, mitochondria from galactose-cultured wild type (WT) cells in the mid-log growth phase were found to contain approximately 1.0 ng of endogenous Hem15 per milligram of mitochondrial protein (Supplementary Fig. S1A). This concentration is comparable to other mitochondrial heme synthesis enzymes as well as other IM proteins [30]. To assess the oligomerization state of Hem15, mitochondria from WT cells were lysed with digitonin and the lysate proteins subjected to sucrose gradient ultracentrifugation followed by SDS-PAGE immunoblotting of the gradient fractions. Hem15 was detected in the ~250–440 kDa molecular weight range (Fig. 1B), suggesting that endogenous yeast Hem15 exists as a high-mass complex, which likely includes additional associated components, as seen for the mammalian enzyme [17].

To determine whether the formation of this complex depends on the functional state of Hem15, the same analysis was performed using a catalytically impaired Hem15 variant, H235C. The yeast Hem15 residue H235 is equivalent to the human FECH residue H263, which is an essential catalytic histidine involved in proton abstraction prior to clelation [31]. Conversion of this residue to a cysteine yielded a variant that is stably expressed yet unable to support heme-independent growth or exhibit in vitro Fech activity (Supplementary Figs. S1B and S1C). While WT Hem15 predominantly copurifies with bound heme, the H235C variant protein copurifies with PPIX (Supplementary Fig. S1D), further demonstrating this variant of Hem15 is impaired in catalyzing formation of heme from PPIX substrate. This H235C variant was found in fractions of lower molecular weight when mitochondrial lysates were subjected to sucrose gradient ultracentrifugation (Fig. 1B), suggesting the formation of the full complex is impaired when Hem15 is catalytically inactive. The Hem15 complex was also found to shift to lower molecular weight fractions in gradient ultracentrifugation of mitochondrial lysates derived from heme synthesis-deficient cells lacking Als (yeast Hem1; Fig. 1B). These findings suggest that Hem15 oligomerization is at least partially dependent on the functional state of both Hem15 and an earlier step of the heme biosynthetic pathway.

2.2. Human ferrochelatase complements the yeast deletion strain

Human FECH was discovered to be fully functional in yeast, complementing the yeast hem15Δ deletion mutant (Fig. 2A), which cannot survive in the absence of exogenous heme supplementation. This finding reveals that essential interactions between Fech and other proteins are conserved in S. cerevisiae. This result is also interesting because there are known differences between the yeast and human ferrochelatase enzymes. The human enzyme, but not the yeast enzyme, possesses a [2Fe–2S] cluster, which is required for enzyme activity [32–34]. Complementation of the hem15Δ strain with wild-type human FECH, but not the human FECH variant C406S lacking an essential [32,35] cluster ligand residue (Fig. 2A), suggests that the FECH iron-sulfur cluster is assembled properly in yeast.

2.3. A genetic model was developed to assess Hem15 functional roles

While re-expression of either yeast or human ferrochelatase (under the control of the native yeast ferrochelatase promoter or a constitutive, high-efficiency MET25 promoter, both on a Yep vector) in the yeast hem15Δ strain allows cell growth in the presence of glucose, it does not always permit growth of this strain on respiratory media (Fig. 2B). This phenomenon has been observed previously [36] and has thus far posed
This inability of plasmid borne *HEM15* to complement the *hem15Δ* deletion mutant was hypothesized to be due to the *hem15Δ* strain becoming petite, i.e., mitochondrial DNA is lost upon deletion of the *HEM15* gene. Consistent with this interpretation, employing a known strategy to stabilize the mitochondrial genome resolved this obstacle. Overexpression of the dNTP checkpoint enzyme Rnr1, which encodes the large subunit of ribonucleotide reductase, has previously been very effective to stabilize mitochondrial DNA in various petite mutants via an unknown mechanism [37–39]. Employing this strategy in *hem15Δ* cells following Hem15 re-expression using a YEp vector under the control of its native promoter (HEM15↑) or a constitutive, high-efficiency *MET25* promoter (HEM15↑↑) restored respiratory growth (Fig. 2B), returned both respiratory complexes and their subunit proteins to WT levels, and permitted complementation with human FECH (Fig. 2C; Supplementary Fig. S2).

### 2.4. Hem15 is physically associated with MICOS machinery

Protein interaction partners of Hem15 were determined by immunoprecipitation (IP) of a FLAG-tagged Hem15 construct. Since attachment of a C-terminal tag to FECH results in an inactive enzyme [40], Hem15 was tagged immediately after its N-terminal mitochondrial targeting sequence, so that the natural proteolytic removal of the targeting sequence upon mitochondrial import results in a protein bearing the FLAG tag on the structurally disordered amino terminus (Supplementary Fig. S3A). This Hem15-iFLAG construct is stably expressed and functional (Supplementary Figs. S3B and S3C) and was therefore used for IP experiments.

Clarified mitochondrial lysates from *hem15Δ* cells stabilized with Rnr1 and expressing Hem15-iFLAG were incubated with anti-FLAG affinity resin, and affinity-purified proteins were analyzed by LC-MS/MS to identify co-purifying proteins. In addition to Hem15 and some of its previously known binding partners such as PpoX (Hem14) [17], the IM
GTpase Mgm1 and five of the eight known subunits of the MICOS complex (Mic60, Mic19, Mic26, Mic27, and Mic12) were all identified as physically interacting with Hem15 (Fig. 3A). Consistent with these findings, targeted co-IP experiments showed the core MICOS subunit Mic60 co-purifies with Hem15-iFLAG (Fig. 3B), further suggesting physical association between Hem15 and MICOS. These findings are consistent with interactions observed in IP experiments with yeast MICOS [41] and human FECH [17,19].

To examine the functional significance of this Hem15-MICOS association, the impact of MIC60 deletion on Hem15 was experimentally determined. Steady-state levels of Hem15 were not affected in the mic60Δ deletion mutant (Supplementary Fig. S3D). However, the sucrose gradient migration profile of the Hem15 high-mass complex derived from mic60Δ mutant mitochondria was altered, indicating a shift toward smaller complex size (Fig. 3C). This shift in Hem15 oligomerization pattern is similar to that observed for both the nonfunctional H235C Hem15 variant and WT Hem15 in the hem1Δ mutant (Fig. 1B). Collectively, these results suggest that Hem15 associates with the MICOS complex and Hem15 oligomerization is impaired in the absence of MICOS.

2.5. MICOS facilitates proper heme biosynthetic activity of Hem15

Respiratory growth in the absence of Mic60 was observed to be slightly impaired, and this growth defect enhanced markedly when Hem15 levels were elevated by overexpression from a plasmid (Fig. 4A). This result was observed for mic60Δ strains in both the W303 and BY4741 genetic backgrounds and with plasmids expressing Hem15 from either its native promoter or a strong heterologous promoter. Hem15 protein levels remained similar in the presence versus the absence of Mic60 (Fig. 4B), ruling out reduced levels of Hem15 as a possible cause for the growth defect. Therefore, the activity of that Hem15 protein was examined.

WT and mic60Δ cell lysates do not have significantly different levels of Fech activity, and overexpression of Hem15 in each cell type results in significantly increased PPIX metabolism by Hem15 (Fig. 4C). Strikingly, when Hem15 is overexpressed, mic60Δ cells have dramatically lower Fech activity than WT cells, suggesting an inability to maximize heme production in the absence of Mic60 despite abundant Hem15 protein (Fig. 4B and C). This apparent defect in Hem15 activity in the absence of Mic60 raises questions about the role of MICOS in the function of Hem15, which were pursued next.
2.6. MICOS is required for optimal porphyrin substrate delivery to Hem15

MICOS could potentially impact Hem15 activity either directly or indirectly. MICOS could be directly responsible for substrate import into or export out of mitochondria due to its key role in maintenance of IM-OM contact sites [24,25], which are postulated to facilitate bidirectional transport of hydrophobic molecules (e.g., phosphatidic acid and coenzyme Q biosynthetic intermediates) [26,27]. Alternatively, the disorganization of the IM stemming from loss of MICOS could negatively impact mitochondrial carrier proteins required for import of substrates or export of heme, thus indirectly impacting Hem15.

In either the direct or indirect involvement scenario, 1) impaired heme export due to the absence of MICOS might lead to mitochondrial heme build-up and inhibition of Hem15 by heme [42], since heme dissociation from FecH is the rate-limiting step in catalysis [43], or 2) impaired import of substrates and intermediates required for heme synthesis into mitochondria might limit substrate availability to FecH. To investigate these two potential mechanisms for how the MICOS machinery contributes to FecH activity, experiments were carried out to assess rates of heme transport across the IM, cellular heme levels, mitochondrial iron levels, and cellular levels of porphyrin biosynthetic precursors to heme, including PPIX.

To investigate the possibility that loss of MICOS impairs heme export, both rates of heme transport across the IM and cellular heme levels were determined. A high-affinity, genetically encoded heme sensor (HS1) [44,45] was employed in an assay to compare rates of heme synthesis from FecH in WT and mic60Δ cells expressing vector control or Hem15 under control of its native promoter (HEM15 †) or the heterologous MET25 promoter (HEM15 ‡). Cells were spotted onto SC media and cultured for 2 days (glucose), 3 days (galactose), or 5 days (glycerol/lactate) at 28°C. (B) SDS-PAGE immunoblot of indicated proteins in mitochondria from cells described in panel A. (C) Ferrochelatase specific activity in mitochondrial lysates from indicated cells. Bars indicate the average and S.D. (error bars) of 3 biological replicates. Asterisks indicate a statistically significant difference by t-test (**p < 0.001).

To test the availability of the other substrate for FecH, PPIX, and intermediates necessary for its synthesis, cellular porphyrin content was measured. The iron content measured by ICP-MS was comparable in WT and mic60Δ cells with or without Hem15 overexpression (Supplementary Fig. S4A), ruling out iron deficiency as the reason for reduced FecH activity in the Hem15-overexpressing mic60Δ mutant. In agreement with this result, the steady-state levels of the mitochondrial iron transporter Mr3 were also unchanged by the absence of Mic60 (Supplementary Fig. S4B). Likewise, loss of Mic60 had no appreciable effect on the activity of the iron-sulfur cluster-containing enzyme succinate dehydrogenase, indicating that iron-sulfur cluster biogenesis is not affected (Supplementary Fig. S4C).
altered in these cells (Supplementary Fig. S5). To rule out the possibility that metabolite transport is universally defective in mic60Δ cells, mitochondrial lipoylation was measured in WT and mic60Δ cells. Levels of the lipoylated mitochondrial pyruvate dehydrogenase subunit dihydrolipoamide acetyltransferase and dihydrolipoyl succinyltransferase were found to be normal in the absence of Mic60, suggesting IM carrier proteins remain generally functional and the elevated porphyrins observed in the absence of Mic60 are the result of a specific effect (Supplementary Fig. S4E).

Interestingly, PPIX was decreased in mic60Δ cells overexpressing Hem15 compared to WT cells overexpressing Hem15 (Fig. 5D), suggesting substrate availability may be a factor that limits Fech activity in these cells. Much like the subtle growth defect and statistically insignificant decrease in Fech activity seen in mic60Δ cells compared to WT cells with endogenous Hem15 levels (Fig. 4), levels of PPIX are not significantly different in the presence versus the absence of Mic60 when Hem15 is present at endogenous levels (Fig. 5D). However, when Hem15 is overexpressed, the growth defect becomes marked, the decrease in Fech activity becomes dramatic, and the decrease in levels of PPIX becomes significant. These three experiments suggest a role for MICOS in Hem15 activity that is more apparent under conditions of high Hem15 levels, when PPIX may become limiting for Hem15 activity.

2.7. MICOS-deficient cells expressing Hem15 have oxidative stress, which can be partially mitigated by synthetic restoration of IM/OM contact sites

Since heme synthesis pathway intermediates like those elevated in the absence of Mic60 are known to be toxic, in part due to their inherent redox reactivity [5,47–49], signs of oxidative damage and stress were evaluated in the absence of Mic60. The metabolic enzyme aconitase was assessed first because of its sensitivity to damage by superoxide. While steady-state levels of aconitase were the same in WT and mic60Δ cells overexpressing Hem15 (Fig. 5E), the cells lacking Mic60 exhibited a significant decrease in aconitase specific activity, consistent with oxidative damage (Fig. 6A). Furthermore, mic60Δ cells overexpressing Hem15 were significantly less tolerant to acute oxidative insults than WT cells overexpressing Hem15 (Fig. 6B). Consistent with these observations of oxidative stress, supplementation with the antioxidant N-acetylcysteine partially rescued a growth defect of Hem15-overexpressing mic60Δ cells on galactose medium (Fig. 6C).

To test whether restoring intermembrane connectivity in MICOS-deficient cells could mitigate some of these cytotoxic effects, an artificial IM-OM tether, mitoT, was generated [50] (Fig. 6D and E). This chimeric protein comprises the N-terminal portion of the IM protein Sco2 (residues 1–112) encompassing the mitochondrial targeting sequence and transmembrane domain, followed by a short (12 amino acid residues) unstructured linker derived from the E. coli LacI protein, a transmembrane helix of the OM import receptor protein Tom20 (residues 1–20), and the GFP moiety. The optimal length of the linker region was deduced from the previously reported analogous construct [26].

Co-expression of mitoT in Hem15-overexpressing mic60Δ cells improved growth on galactose and glycerol-lactate media, reflecting a partial rescue effect (Fig. 6F). Interestingly, mitoT expression had the opposite effect in Hem15-overexpressing WT cells, suggesting that excess intermembrane tethering has a negative effect on the normal function of mitochondria. MitoT expression also resulted in increased steady-state levels of Hem15 protein in both WT and mic60Δ cells (Fig. 6E) although the reason for this effect is unclear at present. The rescue appears to be due to a remarkable increase in oxidative stress tolerance in the Hem15-overexpressing mic60Δ cells co-expressing mitoT (Fig. 6G).

3. Discussion

Towards the goal of better understanding the molecular and functional organization of yeast ferrochelatase (Hem15) and, more broadly, the regulation of the heme biosynthetic pathway, this study has revealed several new findings.

The functional state of Hem15 was assessed and the yeast enzyme compared to human ferrochelatase. Hem15 assembles into a high-mass oligomer, which likely represents the functional enzyme, as it is compromised both in hem1Δ cells blocked at a very early stage of heme biosynthesis and in cells expressing the catalytically inactive H235C Hem15 mutant. This study confirmed that the yeast H235 residue is critical for Ferch activity in yeast, much like the homologous H263 residue in humans [31], and is similarly hypothesized to be critical for proton abstraction from PPIX for metalation and heme formation.

To determine how the active-site structure of Hem15 is impacted by the H235C mutation, the H235C variant was crystallized and its structure solved at 2.4 Å resolution. This structure reveals that the location and orientation of active-site residues in the WT protein and H235C Hem15 variant are similar, as are the active-site hydrogen-bonding...
networks (Supplementary Fig. S1E). This intact network differs from the human H263C variant structure where the network is disrupted [31]. It is unclear why this difference between the yeast and human catalytic mutants exists, though both variants are inactive due to the absence of the base (H235 in yeast and H263 in humans) necessary for proton abstraction.

Human FECH is able to efficiently replace the yeast enzyme, despite the human enzyme requiring a \([2Fe–2S]\) cluster for enzyme activity [32–34] that is absent in the yeast enzyme. This iron-sulfur cluster is believed to serve as a sensor for the redox state and/or pH of the mitochondrial matrix [32,51]. The finding that WT human FECH, but not a mutant bearing the C406S substitution for one of the cluster-binding ligands, complements the yeast deletion mutant suggests that the cluster is successfully formed in the context of yeast mitochondria.

Hem15-deficient cells exhibit a petite-inducing phenotype that can be rescued genetically through re-expression of Hem15 in conjunction with overexpression of ribonucleotide reductase subunit Rnr1, a genetic manipulation known to stabilize mitochondrial DNA in petite mutants [39,52]. This finding explains previously described difficulties in establishing a robust genetic complementation of the \(hem15\)Δ mutant with plasmid-borne Hem15 [36]. While the exact molecular underpinnings of the petite-inducing phenotype of hem15Δ are currently unclear, they might arise from impaired hemylation of mitochondrial
matrix hemoproteins with connections to the mitochondrial genome, such as the yeast flavohemoglobin Yhb1 or translational activator Mss51 [53–56]. Further studies are warranted to explore the connection between loss of Hem15 and instability of mitochondrial DNA.

The high-mass Hem15 complex was found by proteomic analysis to contain components of the MICOS machinery. A portion of the core MICOS subunit Mic60 is exposed to the matrix side of the IM [24,25] and is thus likely to mediate this Hem15–MICOS association. Loss of Mic60 affects the high-mass oligomeric species of Hem15 akin to the destabilizing effect of the catalytic H235C Hem15 variant and the Alas (Hem1) deletion mutant, consistent with the idea that MICOS contributes to the molecular organization of Fech. These results agree with earlier reports suggesting that mammalian FECH is a component of a large complex termed the mitochondrial heme metabolon [17–19]. Studies in mammalian cells have shown the heme metabolon contains components of MICOS, underscoring the conservation and importance of this connection.

The yeast MICOS presents as a series of large oligomers [41] and, since Mic60 is about 1.5-fold more abundant than Hem15 [30,57], it is unlikely that its subunits form stable stoichiometric complexes with Hem15. Instead, there may be a dynamic association between Hem15 and MICOS that is not essential for Hem15 activity but is nevertheless important for its ideal performance. In agreement with this idea is the observation that basal Hem15 activity is not significantly affected in the mic60Δ mutant, whereas the added strain caused by overexpression of Hem15 results in a profound respiratory growth defect, reduces levels of PPIX, attenuates Hem15 activity, elevates levels of intermediate porphyrins, and results in oxidative damage and stress, compared to similarly treated WT cells. Porphyrin profiling under these conditions revealed that the levels of some porphyrins like coproporphyrins and PPIX are decreased in the absence of Mic60, while others like hexacarboxylporphyrins and heptacarboxylporphyrins are increased. Low PPIX levels could be attributed to reduced Ppox activity, a downstream effect of low Cpxox activity since Cpxox is inhibited by heme [58], or a more complex result of this apparent disruption of porphyrin transport and homeostasis. The elevated intermediate porphyrins are likely causing the oxidative damage and stress observed when Hem15 is overexpressed.

These results suggest the FECH–MICOS association has functional significance. We propose a model wherein IM–OM contact sites formed by MICOS facilitate the transfer of intermediate porphyrinogen precursors across the mitochondrial membranes, thereby ensuring optimal substrate delivery to Fech as well as PPOX (Fig. 7). Data in the current study do not exclude the possibility that MICOS is directly involved in this transport of heme biosynthetic intermediates, analogous to its proposed role in transporting phospholipids such as phosphatidic acid [26] and coenzyme Q biosynthetic intermediates [27]. However, it may be more likely that MICOS plays an indirect role through one or more plausible mechanisms. For instance, the complex may aid the heme intermediate transport process through bridging the OM and IM to create a proximity conduit between these membranes, as suggested by the rescue of oxidative stress tolerance seen in Hem15-overexpressing mic60Δ cells co-expressing a synthetic mitoT [50] tether. In mammalian cells, MICOS might be required for spatial organization of the currently elusive coproporphyrinogen III and protoporphyrinogen IX transporters in the OM and IM, respectively, thus accounting for the accumulation of tetrapyrrole intermediates arising from cellular accumulation of the heme synthesis intermediates during uroporphyrinogen III decarboxylation in MICOS-deficient cells (Fig. 5C and Supplementary Fig. S5). It is unclear under which conditions MICOS and Fech physically interact, but this interaction is expected to be dynamic and transient, as discussed above.

Intriguingly, Mic60 is highly conserved with homologs found in α-proteobacteria, wherein the gene clusters with heme biosynthetic genes [59,60], further supporting a potential functional link between MICOS and the heme biosynthetic pathway. The additional identification of the conserved dynamin-like GTPase Mgm1 (OPA1 in humans) as a potential interacting partner of Fech supports earlier proteomic studies in mammalian mitochondria [17,19]. Our recent study established that the mgm1Δ mutant exhibits a marked defect in nuclear heme levels [46], further supporting a model in which IM ultrastructure-related factors like Mgm1 and MICOS may cooperate with Fech to facilitate delivery of its substrate and/or distribution of its product.

4. Materials and Methods

4.1. Yeast strains, plasmids, and growth conditions

Yeast strains used in this work were of the W303–1B genetic background (MATa ade2-1 can1-100 his3-11,15 leu2-3 trp1-1 ura3-1), unless specified otherwise. The ΔMICOS strain included in a Supplementary figure lacks the genes MIC10, MIC60, MIC19, MIC27, MIC26, and MIC12. Plasmids for wild-type and variant Hem15 and human ferrochelatase expression in S. cerevisiae were produced as previously described [61,62]. The HEM15 ORF with its 350-bp native promoter region was subcloned into pRS424 and pRS426 vectors from the pRS316-Hem15 plasmid described before [61]. This plasmid also served as a template to generate a pRS426-Hem15 vector in which the HEM15 ORF is under the control of the heterologous MET25 promoter.

![Fig. 7. Model for involvement of MICOS in heme biosynthesis. IM–OM contacts formed by MICOS may facilitate the transfer of intermediate porphyrin precursors (purple crosses) across the mitochondrial membranes into the matrix, thus ensuring optimal delivery of substrate to Fech for synthesis of heme (red crosses). Loss of Mic60/MICOS results in accumulation of porphyrin intermediates and subsequent oxidative damage. See Discussion for additional details. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)](image-url)
To generate a plasmid expressing Hem15-iFLAG, the mitochondrial targeting sequence (MTS) of Hem15 (amino acid residues 1–31) with a flanking region containing the FLAG epitope tag at the 3’-end was PCR-amplified from the pRS424-Hem15 plasmid using primers 5’-CGGGATCCGTAGCTTCCAGAACAATCGTACAAGGTTCTCCTCTAA GAAGATCAACACTGACATT-3’ and 3’-CTCTTGCATGCAGGCCTTGT GAGTTCGATGTGAAATCGAGCTGTAATTGCGATTTG GATCTTCTT-5’. In a separate reaction, Hem15 excluding the MTS (amino acid residues 31–393) containing the FLAG epitope tag at the 5’-end was PCR amplified from the same plasmid using primers 5’-GAAT CAAAGAGTACGACAAATGAATTCAACAGCCGAAGATTACCC AACAGGAT TTGGTTATGGAACATGGGTGGC-3’ and 3’-GGGGCTAGATGA TATGCTGATGCAATACCGTATTTACCAGGTCATTGAGCTTAC GCTT-3’. Gel-purified products of the reactions described above were fused together by overlap extension PCR using 5’-CGGGATCC CATGCTTCCCGAAGATCGTACAAGGTTCTCCTCTTACAGGA AGATTTGAACATGGGTGGC-3’ and 3’-GGGGCTAGATGTA ATTGCTGATGCAATACCGTATTTACCAGGTCATTGAGCTTAC GCTT-3’ including 5’-BamH I and 3’-Xho I restriction sites, respectively. The obtained product was cloned into pRS423 vector under the control of the MET25 promoter and CYC1 terminator.

The pRS424-Rnr1 plasmid was generated by subcloning a 2.8-kbp fragment from the YEp181-B1 Rnr1 vector [39]. All plasmids were validated by DNA sequencing.

Depending on experiment, cells were cultured in yeast extract-peptone (YP) or synthetic complete (SC) media lacking nutrients (amino acids or nucleotides) necessary to maintain plasmid selective pressure [64] containing either 2% glucose, 2% galactose, or 2% glycerol/2% lactic acid as the carbon source. To culture heme synthesis-deficient strains, cells were grown in the presence of either hemin or Tween-80/ergosterol/methionine mix as described previously [65,66]. Growth tests to assess respiratory capacity and quantify hydrogen peroxide sensitivity were carried out as before [67–69], except cells lacking functional Hem15 were cultured in 20 μM hemin. Cultures used for growth tests were grown overnight (or two days in the case of cells lacking functional Hem15) in SC media lacking relevant nutrients to maintain plasmid selection then normalized to OD600 of 1 and spotted onto SC plates with or without 20 μM hemin.

4.2. Heme and porphyrin analysis –

Total cellular porphyrin content was measured using a previously described porphyrin fluorescence assay [44]. Briefly, 1 × 10^8 log-phase cells were harvested, washed in sterile ultrapure water, resuspended in 500 μL of 20 mM oxalic acid and stored at 4°C overnight (16–18 h) in the dark. The next day, 500 μL of 2 M oxalic acid was added to the cell suspensions. Half the cell suspension was transferred to a heat block set at 95°C and heated for 30 min to deplete the heme iron and yield fluorescent protoporphyrin IX. The other half of the cell suspension was kept at room temperature for 30 min. All suspensions were centrifuged at 4000 rpm for 5 min. Cells were then washed in 5 mL ice cold water and re-centrifuged. Pellets were then resuspended on ice in 4 mL lysis buffer (50 mM Tris-MOPS, pH 8.0, 100 mM KCl, 1% sodium cholate) with 40 μL of a protease inhibitor cocktail (Sigma P-8215) and transferred to 50-mL conical tubes. Glass beads (4 mL, 0.5 mm) were then added, and cells were vortexed for 1 min at maximum speed followed by 1 min on ice, repeated for a total of 5 vortex cycles. Tubes containing lysate and beads were then centrifuged at ~9000 × g for 5 min. Supernatant was removed using a 1-mL pipette, taking care not to disturb the pellet. The supernatant was then transferred to 2-mL micro-centrifuge tubes and further centrifuged at 15000 × g for 10 min. Supernatant was transferred to fresh micro-centrifuge tubes and stored on ice for up to 2 h during measurements. Total protein concentration of yeast lysate was measured via a NanoDrop spectrophotometer with 1 A = 1 mg/mL setting. Heme and porphyrin analysis were carried out at the University of Utah Center for Iron and Heme Disorders core facility as previously described [11,17].

4.3. Hem15 activity assays –

Lysate volumes ranging from 0 to 200 μL were brought to a total volume of 850 μL with lysis buffer in 3-mL glass tubes. Master mix (150 μL) consisting of 20 mM ferrous sulfate, ~1 mM protoporphyrin IX, and 33.3 mM β-mercaptoethanol was added to start the assay. Samples were incubated in the dark at 37°C for 15 min. Heme produced in lysate activity assays was quantified by pyridine hemochromogen assay. To stop the assay reactions and produce the pyridine hemochrome derivative, 1 mL 50% (v/v) pyridine:0.2 N NaOH solution was added to 1-mL assay samples. Heme content was measured via differential spectra of oxidized and reduced samples as described previously [69–71].

4.4. Variant construction, protein expression, purification, and X-ray crystallography –

The Hem15 variant H235C was constructed using QuickChange site-directed mutagenesis (Agilent) and verified by sequencing. His-tagged mature wild-type and variant Hem15 were produced as previously described [40,72]. In vivo ferrochelatase activity of the variant was assessed by complementation and rescue of a strain of Escherichia coli lacking functional ferrochelatase, pppCΔ [73,74]. The H235C variant protein was concentrated to ~400 μM and crystals were grown by hanging drop with mother liquor composed of 0.1 M Bis-Tris pH 6.5, 25% PEG 3350 within 48 h.

All data sets were collected at the Advanced Photon Source and SER-CAT on beamline 22-ID. Phases were obtained by using a monomer of wild-type Hem15 (PDB ID 1LBQ) as a molecular replacement search model. Molecular replacement was performed using the program CNS [75]. Iterative rounds of model building and refinement were performed with the programs COOT [76] and CNS, respectively. Data collection, refinement statistic, and PDB ID for the H235C structure are listed in Table S1. Structural representations were created using PyMol [77].

4.5. Mass spectrometry and data analysis –

Affinity purification and mass spectrometry were carried out as described for human ferrochelatase [17], except 1% digitonin was used to solubilize purified mitochondria. Subsequent analysis of candidate hits against the CRAPome database (www.crapome.org) eliminated known contaminants and non-specific interactors.

4.6. Mitochondrial isolation and assays –

Mitochondria-enriched fractions were isolated using established protocols [78]. For ICP-MS measurements, mitochondrial fractions were further purified using discontinuous 14%/22% Nicodenz gradients as described [79]. Total mitochondrial protein concentrations were
determined using the Coomassie Plus kit (Thermo Scientific). Proteins or protein complexes were separated by SDS-PAGE, blue native (BN)-PAGE, or continuous 12–50% sucrose density gradient ultracentrifugation as previously described [67,80]. Aconitase specific activity was determined as before [81].

4.7. Heme trafficking dynamics assay –

Heme trafficking rates were monitored as previously described [46]. Briefly, in this three-step assay: 1) heme synthesis is first inhibited with succinylacetone (SA) in sensor-expressing cells, 2) the block in heme synthesis is then removed by resuspending cells into media lacking SA, and 3) the time-dependent change in the heme occupancy of HS1 is monitored. The fractional heme occupancy of the sensor can be determined using previously established sensor calibration protocols [44]. The percent of sensor bound to heme (% Bound) is calculated by determining the sensor eGFP/mKATE2 fluorescence ratio (R) under a given test condition relative to the eGFP/mKATE2 fluorescence ratio when the sensor is 100% (% Bound) bound to heme, as described previously [44, 46, 82]. R_min is determined by measuring the HS1-eGFP/mKATE2 ratio in parallel cultures that are conditioned with succinylacetone (SA), which inhibits the second enzyme in the heme biosynthetic pathway, Phgs [83], and R_max can be determined by permeabilizing cells and adding an excess of heme to saturate the sensor [44]. Given HS1 is quantitatively saturated with heme in the cytosol, nucleus, and mitochondria of WT yeast, R_max is typically determined by measuring the HS1-eGFP/mKATE2 ratio in parallel WT cultures grown without SA [44].

Growth for the heme trafficking dynamics assay was accomplished by culturing HS1-expressing cells with or without 500 μM SA (Sigma-Aldrich) in SC media lacking leucine. Triplicate 5-ml cultures were seeded at an initial optical density of OD600nm = 0.1–0.02 (~2–4 x 10^5 cells/ml) and grown for 14–16 h at 30°C and shaking at 220 rpm until cells reached a final density of OD600nm ~1.0 (~2 x 10^8 cells/ml). After culturing, 1 OD (or ~2 x 10^7 cells) were harvested, washed twice with 1 mL of ultrapure water, and resuspended in 1 mL of fresh media. The cells that were pre-cultured without SA provided HS1 R_max values. The SA-conditioned cells were split into two 500-μl fractions. One fraction was treated with 500 μM SA to give HS1 R_min values. The other fraction was not treated with SA so that heme synthesis could be re-initiated to give compartment-specific heme trafficking rates. HS1 fluorescence was monitored on 200 μL of a 1 OD/ml (~2 x 10^7 cells/ml) cell suspension using black Greiner Bio-one flat bottom fluorescence plates and a Synergy Mx multi-modal plate reader. Fluorescence of eGFP (λ_exc. = 488 nm, λ_em. = 510 nm) and mKATE2 (λ_exc. = 588 nm, λ_em. = 620 nm) was recorded every 5 min for 4 h, with the plate being shaken at “medium-strength” for 30 s prior to each read. Background fluorescence of cells not expressing the heme sensors was recorded and subtracted from the eGFP and mKATE2 fluorescence values.

4.8. Immunoblotting –

Separated proteins were transferred to either nitrocellulose or PVDF membranes, blocked in 5% non-fat milk in PBS with 0.1% Tween-20, and incubated with relevant primary antibodies and goat anti-mouse or goat anti-rabbit horseradish peroxidase-coupled secondary antibodies (Santa Cruz Biotechnology). Proteins of interest were visualized by incubation of membranes with chemiluminescence reagents (Thermo Scientific) and exposure to X-ray film. For assessment of Hem15 endogenous levels, proteins were detected using the Odyssey FC imaging system (LI-COR Biosciences) and quantified using built-in Image Studio software. The following primary antibodies were used: mouse anti-porin (459500, Thermo Scientific), mouse anti-Cox1 (ab110270, Abcam), mouse anti-Cox2 (ab110271, Abcam), mouse anti-Cox3 (ab110259, Abcam), and mouse anti-FLAG (sc-66335, Santa Cruz Biotechnology). We also used rabbit sera against S. cerevisiae Hem15 (produced in the Dalay lab), Rip1 (provided by Dr. D. Winge), Mic60 (provided by Dr. N. Pfanner), Aco1 (provided by Dr. R. Lill), and β-subunit of F1 ATP synthase (provided by Dr. A. Tzagoloff). All antibodies were tested for reliability to ensure specificity of detection.

Declaration of competing interest

The authors declare no competing interests.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.redox.2021.102125.

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