

The [PSI+] prion modulates cytochrome *c* oxidase deficiency caused by deletion of COX12

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ABSTRACT:

Cytochrome *c* oxidase is a pivotal enzyme of the mitochondrial respiratory chain, which sustains bioenergetics of eukaryotic cells. Cox12, a peripheral subunit of cytochrome *c* oxidase, is required for full activity of the enzyme, but its exact function is unknown. Here, experimental evolution of a *Saccharomyces cerevisiae* Δ cox12 strain for ~300 generations allowed to restore the activity of cytochrome *c* oxidase. In one population, the enhanced bioenergetics was caused by a A375V mutation in the AAA+ disaggregase Hsp104. Deletion or overexpression of *HSP104* also increased respiration of the Δ cox12 ancestor strain. This beneficial effect of Hsp104 was related to the loss of the $[PSI^+]$ prion, which forms cytosolic amyloid aggregates of the Sup35 protein. Overall, our data demonstrate that cytosolic aggregation of a prion impairs the mitochondrial metabolism of cells defective for Cox12. These findings identify a new functional connection between cytosolic proteostasis and biogenesis of the mitochondrial respiratory chain.

INTRODUCTION:

Mitochondrial energy conversion is pivotal for cellular metabolism and bioenergetics. Energy conversion relies on a series of multi-subunit enzymes, which together form the respiratory chain. Electrons are extracted from reduced metabolic intermediates, originating mostly from the TCA cycle, and transported with the help of mobile electron carriers to the terminal enzyme, cytochrome c oxidase (CcO), which reacts these electrons with molecular oxygen. The chemical energy from the reduced metabolites is thereby converted into proton motive force, which drives ATP formation in mitochondria. In the yeast *Saccharomyces cerevisiae*, the metabolism of so-called respiratory carbon sources (like glycerol, acetate, lactate, ethanol) relies on mitochondrial respiration, whereas glucose can be processed independently via glycolysis.

The biogenesis of the respiratory chain necessitates expression of nuclear and mitochondrial genes. Nuclear-encoded proteins are synthesized in the cytoplasm, and are both post- and co-translationally imported into the organelle with the help of sophisticated import complexes (Pfanner *et al.*, 2019). Inside mitochondria, the different complexes are formed by the step-wise addition of subunits, which in turn need to be equipped with redox co-factors aiding in electron transfer. This assembly process and the fact that nuclear gene expression needs to be synchronized with mitochondrial translation (Couvillion *et al.*, 2016; Ott *et al.*, 2016), makes biogenesis of respiratory chain complexes intricate.

The mitochondrial CcO is the terminal enzyme of the respiratory chain and contains 14 protein subunits, of which three, Cox1, Cox2 and Cox3, are typically encoded in the mitochondrial DNA. These three subunits are conserved from the alpha proteobacterial ancestor of mitochondria. Interestingly, only two of these subunits, Cox1 and Cox2, contain redox co-factors (a and $a3$ hemes, Cu_A and Cu_B) necessary for electron transfer (Carr and Winge, 2003), while Cox3 supports the enzyme by establishing a pathway for oxygen diffusion to the active site. Why the mitochondrial enzyme contains 11 additional subunits is currently not fully understood. A few of the subunits have been implicated in regulation of CcO, while others might confer stability to the catalytic core that becomes destabilized because of the comparably rapid rates by which mutations occur in mitochondrial genomes (van der Sluis *et al.*, 2015). Work in *S. cerevisiae* has shown that absence of distinct subunits leads to different consequences, ranging from complete absence of respiration to very small changes in enzyme activity.

In particular, the deletion of Cox12, a peripheral subunit of CcO that resides in the intermembrane space (Rathore *et al.*, 2019), drastically decreased the activity of CcO despite hemes $a + a3$ being detected at ~50% of wild-type (WT) levels (LaMarche *et al.*, 1992). Interestingly, Cox12 is probably not essential for CcO activity *per se* since detergent-purified CcO remained active despite the loss of Cox12 during the purification (LaMarche *et al.*, 1992). Yeast Cox12p is the ortholog of human COX6B and two missense mutations in the COX6B1 isoform caused severe clinical symptoms (Brischigliaro and Zeviani, 2021), supporting the functional importance of the protein. The precise function of Cox12 is currently unknown but Ghosh *et al.* suggested a link with the copper delivery pathway to Cox2 (Ghosh *et al.*, 2016).

Here we used experimental evolution to alleviate the CcO deficiency of Δcox12 cells and we identified the Hsp104 disaggregase as a modulator of the growth defect on respiratory carbon sources. We further demonstrated an unexpected connection between the cytosolic [PSI⁺] prion and the

functionality of CcO in Δcox12 cells. Together, our data reveal a new link in the intricate network that connects the cytosol and mitochondria.

RESULTS:

Experimental evolution yields Δcox12 cells able to grow on respiratory medium:

Δcox12 cells are deficient for CcO activity, and as such they are unable to grow on respiratory media but can propagate on fermentable medium containing glucose (LaMarche *et al.*, 1992). We used experimental evolution to select mutations that would rescue the respiratory growth defect of Δcox12 cells. To this end, we propagated Δcox12 cells in a medium that contained a limiting amount of maltose (as a fermentable carbon source) and an excess of lactate, glycerol and acetate (LGA, as respiratory carbon sources). In those cells, we deleted the *MSH2* gene in order to increase the nuclear mutation rate (Stone and Petes, 2006; Gammie *et al.*, 2007), and thus accelerate the emergence of evolved phenotypes. The cells were cultured in two replicates (A and B) and were serially transferred 43 times, representing approximately 300 generations (Fig. 1A). After transfer number 8, maltose was omitted from the growth medium as the cultures were able to metabolize LGA. The density of the cultures increased over passages (Table S1), showing that cells improved their respiratory capacities. At the end of the evolution experiment, populations A and B showed robust growth on LGA medium, unlike the Δcox12 ancestor (Fig. 1B). We isolated three clones from each population and assayed their growth on media with various carbon sources (Fig. 1C). All evolved clones grew robustly on LGA medium and also on media that contained glycerol-acetate, ethanol or lactate as sole carbon sources (Fig. 1C), supporting that their mitochondria harbored a functional CcO. As the phenotypes of the three tested clones were comparable (Fig. 1C), we selected a single clone per population for further study (Ac and Ba). Interestingly, clone Ba was able to grow on glycerol medium at 37°C unlike clone Ac (Fig. 1D), suggesting that rescue mechanisms might differ in evolved populations A and B.

Since the evolution experiment was conducted with a *msh2::CaURA3* hypermutator strain, the cells accumulated many mutations over 300 generations. Among these mutations, only a few, called here causal mutations, are likely to mediate recovery of respiratory growth in the evolved strains. To characterize the genetic properties of the causal mutation(s), we crossed clones Ac and Ba with the non-hypermutator $\Delta\text{cox12-HM}$ strain that carries wild-type *MSH2*. The Ba diploid did not grow on LGA medium whereas the Ac diploid did (Fig. S1A), indicating that clones Ba and Ac carried recessive and dominant causal mutations, respectively. The Ba diploid was unable to sporulate, consistent with the requirement of a functional mitochondrial respiratory chain for sporulation (Codon *et al.*, 1995). Thus, to evaluate the number of causal mutation(s) in Ac and Ba clones, we crossed them to strain CEN-HM carrying the wild-type *COX12* gene and we analyzed the meiotic progeny by tetrad dissection. As expected, the Δcox12 deletion segregated 2:2 and 50% of the Δcox12 spores were able to grow on LGA (Table S2), supporting the existence of a single causal mutation in the parental clones Ac and Ba. We analyzed the phenotype of two LGA-positive meiotic clones (P1, P2) and found that it was comparable to that of their respective parents, Ac and Ba (Fig. S1B). Clones Ac-P1 and Ba-P2 were selected for further study as they had inherited the wild-type *MSH2* gene after backcross and thus had a normal mutation rate.

Δ cox12 evolved clones compensate reduction of CcO protein levels and activity:

Next, we assessed if the restoration of respiratory growth in the evolved clones was related to altered proteins levels of CcO subunits. For that purpose, we investigated yeast cells grown either in fermentable glucose or in galactose to avoid any glucose repression impacting mitochondria. As expected, we observed a severe reduction of all analyzed CcO proteins in the Δ cox12 deletion strain. Interestingly, this effect was largely reverted in clones Ac-P1 and Ba-P2 (Fig. 2A), which showed increased levels of Cox1, Cox2 and Cox13. To evaluate CcO activity, we first determined the amount of CcO molecules in mitochondria isolated from each strain. For this purpose, heme spectra were recorded (Fig. 2B-D) and CcO molecules were quantified (applying the molar extinction coefficient $\epsilon = 26 \text{ mM}^{-1} \text{ cm}^{-1}$). Subsequently, oxygen consumption was evaluated from isolated mitochondria of indicated strains and normalized to the determined amount of CcO molecules to provide a readout of CcO activity (Fig. 2E). The massive reduction of CcO activity in the Δ cox12 strain (approx. 25% of WT) was alleviated in the evolved Ac-P1 and Ba-P2 clones (approx. 60% of WT, Fig. 2E). From these results, we concluded that the evolved clones Ac-P1 and Ba-P2 recovered respiratory growth because of a causal mutation that increases CcO activity despite the absence of Cox12.

HSP104 A375V is the causal mutation in evolved Δ cox12 clones isolated from population B:

In order to identify the causal mutations in clones Δ cox12 Ac and Δ cox12 Ba, we used an approach that consists in sequencing the genome of clones selected phenotypically within a meiotic progeny, as originally described by Murray and colleagues (Koschwanetz *et al.*, 2013). In our case, the causal mutations of interest would confer growth on LGA to evolved Δ cox12 clones. Thus, we reasoned that in the meiotic progeny of evolved Δ cox12 clones Ba and Ac crossed to the CEN-HM strain, all Δ cox12 clones able to grow on LGA (P clones = Positive for growth) should contain the causal mutation, whereas those unable to grow on LGA (N clones = Negative for growth) should contain the wild-type allele. In contrast, non-causal mutations should distribute evenly in pools of P clones or N clones that we constituted from 10 to 15 individual Δ cox12 clones selected in the meiotic progeny. We sequenced these pools at $\sim 100 \times$ coverage, together with the ancestor Δ cox12 strain, the wild-type (WT) strain, and individual clones from the meiotic progeny able to grow on LGA (Ac-P1, Ba-P2, see Fig. S1B). In order to identify the causal mutation, we filtered the sequencing results for mutations present in the Ba-P1 clone and Ba-P pool, but absent in the WT, in the Δ cox12 ancestor and in the Ba-N pool. This analysis yielded three candidate mutations (Table S3). Two of them were positioned in intergenic regions, the third (XII:89746 C \rightarrow T substitution) was present in the coding sequence of the *HSP104* gene, and this *hsp104*-C1124T caused a A375V missense variant in the Hsp104 protein. A similar analysis did not yield any candidate causal mutation for clone Ac, which was therefore not further considered in this study.

Interestingly, Sanger sequencing revealed that the *hsp104*-C1124T mutation was also present in Δ cox12 Bb and Bc (Fig. S2A), two other clones originally isolated from population B (Fig. 1C). Hsp104 is a heat shock protein that is induced under various stresses. It exhibits disaggregase activity in cooperation with Ydj1p (Hsp40) and Ssa1p (Hsp70) to refold and reactivate denatured, aggregated proteins (Glover and Lindquist, 1998). *S. cerevisiae* Hsp104 counts 908 residues and the A375V mutation lies in the nucleotide binding domain 1 (Fig. S2B), which provides most of the ATPase activity necessary to drive protein disaggregation (Hattendorf and Lindquist, 2002). This region shows a high level of conservation

and A375 is strictly conserved in sequences from distant species (Fig. S2C). To test if the A375V mutation affected the function of Hsp104, we evaluated its capacity to promote induced thermotolerance, as described previously (Sanchez and Lindquist, 1990; Hung and Masison, 2006). As seen in figure 3A, the $\Delta hsp104$ strain showed a decreased survival at 50°C compared to the WT strain, in agreement with previous reports (Sanchez and Lindquist, 1990; Hung and Masison, 2006). Chromosomal integration of the *hsp104*-C1124T gene encoding Hsp104-A375V restored thermotolerance similar to WT (Fig. 3A), showing that the A375V allele maintained at least some level of disaggregase activity.

Deletion and overexpression of HSP104 rescue the respiratory growth defect of $\Delta cox12$:

Next, we wanted to verify whether the *hsp104*-C1124T mutation is sufficient to restore LGA growth of the $\Delta cox12$ strain. For this, we crossed a $\Delta cox12$ strain to the $\Delta hsp104$ strain with integrated *hsp104*-C1124T and isolated double mutants from the meiotic progeny after tetrad dissection (data not shown). The $\Delta cox12 \Delta hsp104$ + Hsp104-A375V strain showed robust growth on respiratory medium, but surprisingly, so did the $\Delta cox12 \Delta hsp104$ strain (Fig. 3B). As such, the phenotype of the $\Delta cox12 \Delta hsp104$ + Hsp104-A375V strain might therefore result exclusively from the *hsp104* deletion. We then evaluated the effect of plasmids encoding Hsp104 or Hsp104-A375V on the respiratory phenotype of the $\Delta cox12$ ancestor. Low-copy plasmids allowed intermediate growth and high copy plasmids provided strong complementation, whether they carried WT Hsp104 or the A375V allele (Fig. 3C). Together, our results show that either elimination or overexpression of Hsp104 influence positively the respiratory capacities of the $\Delta cox12$ strain.

Guanidium hydrochloride improves the growth of $\Delta cox12$ cells on respiratory medium:

Besides its role in thermotolerance, Hsp104 has also been connected with the maintenance of prions, particularly [*PSI*⁺] (Chernoff *et al.*, 1995; Jung and Masison, 2001; Kryndushkin *et al.*, 2003; Liebman and Chernoff, 2012; Dulle and True, 2013). [*PSI*⁺] is a naturally occurring amyloid prion consisting of highly ordered fibrous aggregates of the Sup35 protein (Kelly and Wickner, 2013), which normally acts as a translation termination factor (Stansfield *et al.*, 1995; Zhouravleva *et al.*, 1995). The Hsp104 disaggregase activity is required for prion replication, likely by fragmenting the fibers, thus creating new seeds for prion propagation (Kushnirov and Ter-Avanesyan, 1998; Kryndushkin *et al.*, 2003; Liebman and Chernoff, 2012). The overexpression of Hsp104 has also been shown to efficiently prevent the transmission of [*PSI*⁺] to the mitotic progeny (Chernoff *et al.*, 1995; Derkatch *et al.*, 1996). Thus, the inactivation of Hsp104 or its overexpression can both cure the [*PSI*⁺] prion from *S. cerevisiae* (Liebman and Chernoff, 2012; Greene *et al.*, 2020). Given that deletion of *hsp104* and overexpression of *HSP104* restored respiratory growth of the $\Delta cox12$ strain (Fig. 3B, 3C), we investigated whether this phenotype was linked to [*PSI*⁺].

Guanidine hydrochloride (Gu) is known to inactivate Hsp104 and consequently blocks the replication of prions, particularly [*PSI*⁺] (Jung and Masison, 2001). Classically, Gu is added at 5 mM in culture medium (Chacinska *et al.*, 2000; Jung and Masison, 2001). However, a high proportion of WT cells treated with 5 mM Gu lost their respiratory capacity since they formed mostly small colonies on *pet* selection medium containing a limiting amount of glucose and an excess of glycerol (Fig. S3A). After two

subsequent cultures with 5 mM Gu, only very few cells were still able to respire (Fig. S3A), showing that 5 mM Gu interfered with respiration in WT cells. 3 mM Gu seemed better suited as a much higher proportion of cells formed large colonies on *pet* selection medium after two cultures in the presence of Gu (Fig. S3B). Therefore, we used 3 mM Gu in subsequent experiments and found that the Δcox12 ancestor strain was able to form large colonies on *pet* selection medium only after culture in the presence of 3 mM Gu (Fig. 4A). We randomly selected three large colonies (Gu1-Gu3) originating from Gu-supplemented cultures and monitored $[\text{PSI}^+]$ and $[\text{PIN}^+]$ prions by analyzing cellular extracts with semi-denaturing detergent agarose gel electrophoresis (SDD-AGE) followed by immunodetection for Sup35 and Rnq1, respectively. Sup35 and Rnq1 aggregates were seen at high molecular weight in extracts of the Δcox12 ancestor strain but not in Gu1-Gu3 clones (Fig. 4B and Fig. S4A), demonstrating that Gu had efficiently cured $[\text{PSI}^+]$ and $[\text{PIN}^+]$. The growth of the Gu1-Gu3 clones was improved on respiratory media compared to the Δcox12 ancestor strain (Fig. 4C). Consistently, we found significantly increased CcO activity and higher Cox2 protein levels in Gu1 and Gu2 clones compared to the Δcox12 ancestor strain (Fig. 4D-E). Together, these results demonstrate that prions negatively impact Cox2 levels, CcO activity and respiratory metabolism in Δcox12 cells.

The $[\text{PSI}^+]$ prion impairs mitochondrial respiration of Δcox12 cells:

We wanted to verify whether the Δcox12 ancestor strain had acquired the prions during the genetic manipulations required for its construction or whether $[\text{PSI}^+]$ and $[\text{PIN}^+]$ were already present in the CEN.PK2-1C/D parental strains obtained from the Euroscarf repository. Thus, we analysed by SDD-AGE the extract of these strains together with those from the evolved clones Δcox12 Ac-P1 and Δcox12 Ba-P2. Sup35 and Rnq1 aggregates were clearly detected in the CEN.PK2-1C/D parental strains and in the Δcox12 ancestor (Fig. S4B), demonstrating that $[\text{PSI}^+]$ and $[\text{PIN}^+]$ were present originally. Interestingly, both evolved clones had cleared $[\text{PSI}^+]$ but Δcox12 Ac-P1 still contained $[\text{PIN}^+]$ (Fig. S4B). This suggested that clearing of $[\text{PSI}^+]$ was responsible for the equally increased CcO activity measured in both evolved strains compared to the Δcox12 ancestor (Fig. 2E). Since we showed that elimination or overproduction of Hsp104 restored respiratory growth of Δcox12 (Fig. 3B-C), we evaluated how prions were impacted in those strains. Consistent with previous reports (Chernoff *et al.*, 1995; Derkatch *et al.*, 1997; Liebman and Chernoff, 2012), deletion of *hsp104* had cured $[\text{PSI}^+]$ and $[\text{PIN}^+]$ whereas Hsp104 overexpression had only cured $[\text{PSI}^+]$ (Fig. S4C). This result again supported that $[\text{PSI}^+]$ and not $[\text{PIN}^+]$ interferes with respiratory growth of Δcox12 .

To further substantiate this, we deleted the *COX12* gene in two isogenic $[\text{pin}^-]$ backgrounds (1414 and 1415) that differ only in the presence or absence of the PSI prion (Jung and Masison, 2001). 1414 $[\text{PSI}^+]$ Δcox12 cells were unable to grow on YPLG and YPLGA plates, whereas 1415 $[\text{psi}^-]$ Δcox12 cells showed appreciable growth (Fig. 5A). Overall, these data demonstrate that $[\text{PSI}^+]$ negatively impacts the respiratory capacity of Δcox12 cells.

Hsp104-A375V cures $[\text{PSI}^+]$ efficiently:

Finally, we wanted to understand the reason for the positive selection of the *hsp104*-C1124T mutation during the evolution experiment. We thus transformed low copy vectors encoding Hsp104 or

Hsp104 A375V in 1414 and 1414 Δcox12 cells and monitored the curing of $[\text{PSI}^+]$ by evaluating the color of colonies on selective plates. Indeed, cells carrying the *ade2-1* nonsense allele appear red on low adenine medium when $[\text{psi}^-]$ but they appear white when $[\text{PSI}^+]$ because the aggregation of the Sup35 protein reduces termination efficiency and causes read-through of the *ade2-1* nonsense codon (Wickner *et al.*, 1995; Liebman and Chernoff, 2012; Yeast Genetic Structures and Functions, 2012). Whereas cells containing the control vector formed white colonies as expected, most clones transformed with the Hsp104 A375V construct showed a red color (Fig. 5B). The cells encoding the WT Hsp104 protein displayed an intermediate color (Fig. 5B), suggesting that the elimination of $[\text{PSI}^+]$ was more efficient in cells expressing the A375V allele. Overall, we propose that the *hsp104*-C1124T mutation was selected in Δcox12 population B during the evolution experiment because it allowed an efficient clearing of $[\text{PSI}^+]$, which in turn improved CcO activity and allowed the use of the respiratory substrates available in the evolution medium.

DISCUSSION

Classical genetic approaches, like high copy suppressor screens or the characterization of spontaneous mutants, have been instrumental to pinpoint the role of several proteins required for the proper function of the mitochondrial respiratory chain (Tzagoloff and Dieckmann, 1990; Lasserre *et al.*, 2015; Rutter and Hughes, 2015). However, such approaches typically yield single modifiers of a phenotype, whereas experimental evolution has the potential to reveal complex epistatic interactions involving several genes (Kawecki *et al.*, 2012; Koschwanez *et al.*, 2013; Plucaín *et al.*, 2014). In this study, we tested whether it is possible to compensate the defect in CcO activity caused by the absence of the peripheral subunit Cox12. Thus, we conducted an evolution experiment over several hundreds of generations to select Δcox12 clones with improved respiration. The two populations showed a rapid improvement of respiratory metabolism (Table S1) and genetic analysis of two representative Δcox12 clones showed that a single causal mutation was involved in each case. Given the high number of mutations accumulated during evolution experiments, the identification of the mutation(s) causative of the phenotype is often challenging (Koschwanez *et al.*, 2013; Fisher and Lang, 2016). Here, we identified *hsp104*-C1124T as the causative mutation in population B via next generation sequencing of the meiotic progeny of evolved clones selected for respiratory growth on LGA medium. The reasons for our incapacity to identify the causal mutation in population A remain unclear but this mutation is interesting as it restored CcO activity and respiratory growth to levels comparable to those obtained in population B (Fig. 1-2). Our analyses showed that the clones from both populations lost the $[\text{PSI}^+]$ prion present in the ancestral strain (Fig S4B), suggesting a possible common rescue mechanism of CcO activity. A potential explanation is that one cell in population A spontaneously lost the $[\text{PSI}^+]$ prion, which would have conferred a fitness advantage to this $[\text{psi}^-]$ lineage in the evolution medium, allowing it to become dominant in population A.

We showed that cells containing the *hsp104*-C1124T allele, which encodes the Hsp104-A375V variant, lost the $[\text{PSI}^+]$ prion more efficiently compared to cells with wild-type Hsp104 (Fig. 5B). The importance of the AAA+ disaggregase Hsp104 for the propagation of $[\text{PSI}^+]$ has been recognized early on with studies showing that $[\text{PSI}^+]$ was cured not only by loss of function of Hsp104, but also by its overproduction (Chernoff *et al.*, 1995; Wegrzyn *et al.*, 2001). The sensitivity to Hsp104 overproduction is a unique feature of $[\text{PSI}^+]$ among yeast prions (Liebman and Chernoff, 2012). The same basic function of

Hsp104, namely the disentangling of misfolded/aggregated proteins by extrusion through the central pore of the Hsp104 hexamer, is needed for its role in prion propagation and induced thermotolerance (Greene *et al.*, 2020). As expected, many point mutations in Hsp104 that decrease thermotolerance also decrease prion propagation (Kurahashi and Nakamura, 2007). Nevertheless, both functions can be dissociated since some mutations were described in Hsp104 to cause defective prion propagation despite maintaining thermotolerance (Jung *et al.*, 2002; Kurahashi and Nakamura, 2007). The A375V mutation that we identified in the NBD1 domain also falls into this class since it maintained thermotolerance and cured *[PSI+]* efficiently.

Several prion proteins have been identified in *S. cerevisiae*, the most studied being *[PSI+]* and *[PIN+]* (Kelly and Wickner, 2013). They correspond to infectious and self-replicating amyloid-like aggregates of Sup35 and Rnq1, respectively. The Sup35 protein functions normally as a translation termination factor, thus aggregation of Sup35 in *[PSI+]* cells increases the rate of ribosomal read-through for nonsense codons to ~1%, as compared to ~0.3% in *[psi-]* cells (Firoozan *et al.*, 1991). As such, *[PSI+]* would promote evolvability and is thus thought to be beneficial in harsh environments (Halfmann *et al.*, 2012), but its rare frequency in nature – presence in ~1% of 700 wild yeast isolates – suggests that it may be detrimental in most cases (Halfmann *et al.*, 2012). Rates of spontaneous appearance of *[PSI+]* in laboratory conditions range from $6 \cdot 10^{-7}$ to $1 \cdot 10^{-5}$ per generation (Lancaster *et al.*, 2010; Speldewinde and Grant, 2017) and increase up to $4 \cdot 10^{-4}$ during chronological aging (Speldewinde and Grant, 2017). Even though the effect of *[PSI+]* on cellular fitness in laboratory conditions is still debated (Kelly *et al.*, 2012; Kelly and Wickner, 2013; Speldewinde and Grant, 2017), the self-propagating properties of prions may result in a rather high frequency of *[PSI+]* in laboratory strains. Importantly, the two CEN.PK2 strains we obtained from the Euroscarf repository carried the *[PSI+]* and *[PIN+]* prions, suggesting that prions might be quite common in strains widely used by the scientific community.

Interestingly, *[PSI+]* has been previously connected to mitochondrial function in yeast. Indeed, *[PSI+]* cells showed a highly fragmented mitochondrial network and a decreased mitochondrial abundance of the prohibitins Phb1 and Phb2, probably as a result of being trapped in cytosolic *[PSI+]* aggregates (Sikora *et al.*, 2009). Since prohibitins stabilize newly synthesized mitochondrial proteins (Nijtmans *et al.*, 2002), Cox2 was proposed to be indirectly destabilized by *[PSI+]* via the decrease of Phb1 and Phb2 (Sikora *et al.*, 2009). Decreased levels of Cox2 in *[PSI+]* cells were also observed in the *nam9-1* strain, which carries a point mutation in the Nam9 mitoribosome subunit (Chacinska *et al.*, 2000). Although we cannot entirely rule out the possibility that *[PIN+]* might also contribute to the phenotype of the Δcox12 cells, the following arguments are in favor of *[PSI+]* playing the major role in modulating CcO activity: i) evolved clones Ac-P1 *[psi-][PIN+]* and Ba-P2 *[psi-][pin-]* displayed similar levels of CcO activity (Fig. 2E), ii) the overproduction of Hsp104 improved respiratory growth and cured PSI but not *[PIN+]* (Fig. 3C and S4C), iii) the respiratory growth of isogenic *[pin-]* Δcox12 strains was decreased in *[PSI+]* cells compared to *[psi-]* cells (Fig. 5A).

In *S. cerevisiae*, CcO is composed of 11 nuclear-encoded subunits and 3 mitochondrial-encoded subunits. Overall, our data show that the function of CcO *in vivo* is not strictly dependent on Cox12 since the Δcox12 ancestor displayed a ~20% residual CcO activity and the evolved Δcox12 clones Ba-P2 and Ac-P1 recovered ~50% activity of WT cells (Fig. 2E). These data are consistent with previous work showing that purified CcO was active despite the loss of the Cox12 subunit during the purification (LaMarche *et al.*, 1992). Since several studies reported that CcO subunits are degraded when the enzyme is not properly assembled (Zee and Glerum, 2006; Barrientos *et al.*, 2009; Soto *et al.*, 2012), it is tempting to

speculate that the low levels of Cox1,2,13 proteins observed in Δcox12 cells reflect an assembly defect of CcO, which is partially corrected in the evolved clones (Fig. 2A). Interestingly, Cox12 is rapidly ubiquitinated and degraded by the proteasome when its mitochondrial import is defective (Kowalski *et al.*, 2018). Thus, the addition of Cox12 into CcO, which corresponds to one of the last step in the assembly process (Timón-Gómez *et al.*, 2018), appears to be tightly controlled by cytosolic proteostasis.

The assembly of functional CcO is a complex process that requires multiple accessory factors to coordinate the synthesis, maturation and assembly of the 14 subunits (Soto *et al.*, 2012; Timón-Gómez *et al.*, 2018; Watson and McStay, 2020). In this study, we demonstrated in two genetic backgrounds (CEN.PK2 and 1414/5) that [PSI+] is detrimental to the respiratory metabolism of Δcox12 cells and that [PSI+] negatively impacts the activity of CcO. Several mechanisms might underlie these phenotypes. First, the assembly of CcO might be impaired by lower levels of particular CcO subunits or assembly factors, as a result of cytosolic sequestration within Sup35 aggregates or via destabilization due to decreased abundance of stabilizing proteins like prohibitins (Sikora *et al.*, 2009). Even though a recent proteomic study demonstrated that the proteomes of [psi-] and [PSI+] cells are very similar (Chan *et al.*, 2017), variations in subcellular levels, notably of mitochondrial proteins, are possible. Second, mitochondrial biogenesis and cytosolic protein aggregation are tightly linked. Most mitochondrial proteins are synthesized in the cytoplasm and are post-translationally imported into the organelle. Cytosolic chaperones maintain these proteins in an import-competent form, and occupation of chaperones by other cytosolic proteins like prions can reduce import efficiency (Dimogkioka *et al.*, 2021). This might be particularly relevant for small proteins of the mitochondrial intermembrane space that are imported via the MIA pathway. Interestingly, this relationship between import and aggregation appears to be mutual (Nowicka *et al.*, 2021; Schlagowski *et al.*, 2021). Third, [PSI+] is known to decrease the efficiency of translation termination, which could perturb the synthesis of critical nuclear-encoded CcO subunits or assembly factors and thus impair the concerted assembly of CcO. However, the rather modest increase in read-through rate - from 0.3% in [psi-] cells to 1% in [PSI+] cells (Firoozan *et al.*, 1991)- suggests that its impact is likely limited. Whatever the mechanism at play, we clearly demonstrated in this study that [PSI+] impairs mitochondrial respiration in Δcox12 cells. Our work reveals that [PSI+] is an important modulator of the complex interplay between the cytosol and mitochondria. Whether [PSI+] affects other respiratory complexes besides CcO and whether other prions may also impact mitochondrial bioenergetics are interesting questions that remain to be addressed in future studies.

MATERIAL AND METHODS

Yeast culture conditions:

Yeast strains were typically grown at 30°C and 180 rpm shaking in either YP-based rich medium (1% [w/v] yeast extract, 2% [w/v] peptone), or in YNB-based synthetic medium (YNB wo AS, US Biological) supplemented with ammonium sulfate (5g/L) and nutrients to cover the strains' auxotrophies in quantities as described in (Sherman, 2002). Carbon sources from 20% stocks sterilized by filtration were added at the following final concentration, unless indicated otherwise: glucose (2% [w/v]), lactate (2% [w/v]), ethanol (2% [v/v]), acetate (2% [w/v]), glycerol (2% [v/v]), LGA was 1% lactate-0.1% glycerol-0.1% acetate. When G418 was added to synthetic medium (0.2mg/mL), ammonium sulfate was omitted and

monosodium glutamate (MSG, 1g/L) was used as sole nitrogen source. ½ YPD (0.5% yeast extract, 2% peptone, 2% glucose) was used to monitor the PSI status of *ade2-1* strains. Bacto Agar (Euromedex) was added at 1.6% (w/v) for solid media. Guanidium hydrochloride (Gu, Sigma) was added at 5 mM or 3 mM from a 1 M aqueous solution sterilized by filtration over 0.2 µm. *pet* selection medium consisted of YP 2% glycerol, 0.1% glucose (Tzagoloff and Dieckmann, 1990).

Construction of strains and plasmids:

Saccharomyces cerevisiae strains used in this study are listed in Table S4. Transformations were performed using the lithium acetate method (Burke *et al.*, 2000) and genomic DNA was prepared according to (Dymond, 2013). The Δ *cox12* strain used in the evolution experiment was constructed from the CEN.PK2-1D strain in several steps including insertion of *Saccharomyces kluyveri* *HIS3* gene at the *MET17* locus, restoration of *LEU2* at the *leu2-3_112* locus, deletion of *cox12* by the KanMX cassette and deletion of the DNA mismatch repair gene *msh2* by *Candida albicans* *URA3*. First, the *HIS3* gene from *S. kluyveri* was amplified from the pFA6a:His3MX6 plasmid using the oligonucleotides 5SkHis3Met17 and 3SkHis3Met17 (Table S5) and the PCR product was transformed in CEN.PK2-1D. Recombinant clones were selected on medium lacking histidine and were checked for methionine auxotrophy. Correct replacement of *MET17* ORF by the SkHIS3 gene was verified by PCR with the 5verifMet17 and 3verifMet17 primers and one clone, called CEN-H, was selected for subsequent modification of the *leu2-3_112* locus. A PCR product corresponding to the *LEU2* gene was amplified from the pRS415 plasmid with oligonucleotides 5verifLeu2 and 3verifLeu2 and transformed into CEN-H. Recombinant clones were selected on medium lacking leucine and correction of the *LEU2* locus in strain CEN-HL was verified by DNA sequencing. Alternatively, the *leu2-3_112* locus of CEN-H was replaced with the *MET17* gene by homologous recombination with a PCR fragment obtained with the primers 5Met17Leu2 and 3Met17Leu2. Recombinants were selected on medium lacking methionine and correct insertion of the *MET17* gene at the *leu2* locus was verified by PCR with 5verifLeu2 and 3verifLeu2 primers on the genomic DNA. One correct clone was selected and named CEN-HM. Strain CEN-HL was crossed to CEN.PK2-1C, the diploid was selected on medium lacking leucine and methionine, and after sporulation according to a published procedure (Sherman, 2002), tetrads were dissected on a Nikon 50i microscope equipped with a micromanipulator. Strains CEN-HLa and CEN-Ha were isolated after selecting the appropriate markers in the meiotic progeny and verifying the loci by PCR. *COX12* was deleted in strains CEN-HLa and CEN-HM by homologous recombination of a fragment obtained by PCR amplification with the primers 5cox12 and 3cox12 on the genomic DNA of the BY4741 Δ *cox12::KanMX4* strain. Recombinants were selected on YPD medium containing G418, positive clones were then plated on YPLG to verify respiratory growth and strains Δ *cox12-HLa* and Δ *cox12-HM* were confirmed by PCR amplification with the primers 5cox12 and 3cox12. Finally, the *MSH2* gene was deleted in the Δ *cox12-HLa* strain by homologous recombination with a fragment obtained by PCR using the primers 5KOMsh2 and 3KOMsh2 on genomic DNA of *Candida albicans* SC5314. Transformants were selected on medium lacking uracil and correct insertion of CaURA3 at the *msh2* locus was verified by PCR with primers 5verifMsh2 and 3verifMsh2 on the genomic DNA of candidates. The resulting strain was named Δ *cox12-msHLa* and was used as the ancestral strain (Δ *cox12* anc.) to initiate the evolution experiment.

Deletion of *cox12* in the 1414 and 1415 strains (derived from the 779-6A strain (Jung and Masison, 2001)) was conducted as described above with recombination of the *kanMX4* cassette at the *cox12* locus. Hsp104 was deleted in the CEN.PK2-1C strain by replacing the *hsp104* locus with a URA3 cassette, amplified with primer Hsp104_fwd and Hsp104_rev from a pUG72 plasmid.

All plasmids used in this study are listed in Table S6. pRS424 plasmids expressing wild type Hsp104 or the mutant Hsp104-A375V were obtained by cloning a PCR fragment amplified from the genomic DNA of Δ *cox12* anc. or Δ *cox12* Ba-P2 with primers cHsp104_fwd and cHsp104_rev. The PCR fragment contained the *HSP104* ORF with its endogenous promoter and terminator and was cloned into pRS424 using *XhoI* and *NotI*. After verification by sequencing, inserts were subcloned into the pRS414 plasmid. Hsp104-A375V was also subcloned into a pRS305 plasmid using *XhoI* and *NotI* and subsequently integrated into Δ *hsp104* to yield Δ *hsp104*+Hsp104-A375V. Strains Δ *cox12* Δ *hsp104* and Δ *cox12* Δ *hsp104*+Hsp104-A375V were isolated from the F1 meiotic progeny of a diploid obtained by mating Δ *cox12*-HM with Δ *hsp104*+Hsp104-A375V. Diploids were selected on –leucine – histidine medium, sporulated and tetrads were dissected. Germinated spores were screened for LEU2, URA3 and *kanMX4* markers, loci were verified by PCR and Δ *cox12* Δ *hsp104* (*URA3*, *kanMX4*, *leu2*) and Δ *cox12* Δ *hsp104*+Hsp104-A375V (*URA3*, *kanMX4*, *LEU2*) were obtained.

Experimental evolution:

The medium was composed of 1.67g/L YNB (without pABA and folate, MP Biomedicals) supplemented with 5g/L ammonium sulfate, 1g/L monosodium glutamate, 0,008% yeast extracts, 0,15% maltose, 1% Lactate / 0,2% Glycerol / 0,2% Acetate (LGA), tryptophan and methionine to cover the auxotrophies (Sherman, 2002). The pH of the medium was adjusted to 4.4 with 5 M NaOH and the carbon sources (sterilized by filtration) were added after autoclaving. The Δ *cox12*-*msHLa* strain was inoculated in duplicate in 250 mL Erlenmeyer flasks containing 20 mL medium and incubated at 30°C with 180 rpm shaking. Both cultures were diluted and transferred daily into fresh medium for 28 days, (see Table S1 for dilution factors), then from day 29, the cultures were transferred only every two days. Before transfer, growth was evaluated by measuring the optical density at 600nm (Table S1) in a Tecan microplate reader (NanoQuant Infinite M200PRO). From transfer 8, maltose was omitted from the culture medium. Every 3 to 4 transfers, glycerol stocks were constituted by sampling 0.8 mL of culture, adding 0.2 mL glycerol, and freezing in liquid nitrogen before storing at -80°C.

Genetic analysis of clones from experimental evolution:

Among the three clones Aa-Ac and Ba-Bc, which we isolated from each population at the end of the evolution experiment (after growth of cells from transfer 43), Ac and Ba were selected for further analysis. To differentiate between dominant and recessive causal mutations, Ac and Ba were crossed with the Δ *cox12*-HM strain and diploids were selected on medium lacking leucine and methionine.

To evaluate the number of causal mutation(s) in clones Ac and Ba, they were crossed to strain CEN-HM and diploids were selected on medium lacking leucine and methionine. We sporulated the diploids and performed tetrad dissection on glucose synthetic medium. After 3 days at 30°C, we replicated about 15 complete tetrads for each strain on synthetic glucose media lacking leucine or

methionine, synthetic LGA medium containing or not G418, on YPD + G418. The plates were incubated for 2 days (glucose) or 4 days (LGA) at 30°C. Clones that grew on the YPD + G418 carried the $\Delta\text{cox12}::\text{kanMX4}$ marker and were subdivided into P clones and N clones depending on their growth phenotype on the LGA plate (P = Positive for growth, N = Negative for growth). 10-15 individual P and N clones were grouped to constitute the P and N pools used for whole genome sequencing. Ac-P1/2 and Ba-P1/2 clones were characterized phenotypically and Ac-P1, Ba-P2 were characterized biochemically (Fig. 2).

Whole genome sequencing:

Selected clones were grown in 10 mL cultures in synthetic glucose medium for 30 hours at 30°C, 180 rpm shaking. For cultures of P and N clones, 2 units OD600 were mixed together to obtain Ac-P, Ac-N, Ba-P and Ba-N pools. Cells were collected by centrifugation 3200g, 4°C, 5 min and genomic DNA was prepared from cell pellets according to (Dymond, 2013). Illumina sequencing was performed by the Max Planck-Genome-centre Cologne, Germany (<https://mpgc.mpipz.mpg.de/home/>). Genomic DNA was sheared by Covaris Adaptive Focused Acoustics technology (COVARIS, Inc.) to average fragment size of 250bp with settings: intensity 5, duty cycle 10%, 200 cycles per burst and 180s treatment time. Library preparation was done with NEBNext Ultra DNA Library preparation kit, and then sequenced as a 2x150bp paired end reads on a HiSeq 3000 to approximately 3 million reads per sample. The sequencing data were deposited in BioProject at NCBI under the identification PRJNA768319.

Analysis of whole genome sequencing data:

Reads were analyzed with fastQC (<https://www.bioinformatics.babraham.ac.uk/projects/fastqc/>) and trimmed with cutadapt based on the fastQC results (Martin, 2011). Trimmed reads were aligned on the S288C R64-1-1 reference genome using bwa-mem (arXiv:1303.3997v2). The resulting BAM files were sorted and indexed with samtools (Li *et al.*, 2009). GATK HaplotypeCaller 3.8 in BP_RESOLUTION mode was used to produce GVCF files (McKenna *et al.*, 2010). The resulting GVCF files were merged with mergeGVCFs.pl (a component of <https://github.com/ntm/grexome-TIMC-Primary>) to obtain a single GVCF file. Since some samples were clonal while others were pools of 10 to 15 haploid strains, the GATK genotype calls (GT) were often irrelevant; they were therefore discarded, and the ALT alleles and allele frequencies (AF) produced by GATK were used to make our own genotype calls, using a simple yet reliable algorithm. Briefly, a minimum depth DP of 10 was imposed to make a call, and any allele supported by at least 15% of the reads was called for that sample.

Variants were then filtered, imposing a homozygous variant call in samples Ba-P2 and Ba-P pool and a homozygous reference call in all other samples. This resulted in three candidate variants (Table S3). Annotation using VEP (McLaren *et al.*, 2016) showed that a single variant had more than “MODIFIER” impact, and directly affected a coding sequence: the XII:89746 C→T substitution, which results in a A375V missense mutation in HSP104. Similar analyses with the Ac pool and Ac-P1 clone did not yield any promising candidates.

All scripts developed for these analyses are available (https://github.com/ntm/cox12_evolutionExperiment).

Growth assay by serial dilution:

Overnight cultures were adjusted in sterile water to $OD_{600} = 1$ and ten-fold serial dilutions were performed in sterile water in 96 well plates. 5 μ L of each dilution was spotted on various plates using multi-channel pipette and plates were typically imaged after incubation at 30 °C for three days (fermentable carbon source) or 5 to 10 days (respiratory carbon sources).

Induced thermotolerance assay:

Cells from overnight cultures were used to inoculate 4 mL YPD cultures at $OD_{600nm} = 0.2$. The cultures in glass tubes were incubated at 30°C to mid log phase ($OD_{600nm} \sim 1$) at which point they were switched to 37°C for one hour with 180 rpm shaking. Then, cultures were transferred at 50°C in a water bath with occasional shaking and 200 μ L aliquots were taken at different time intervals (3, 7, 10, 15, 20, 30 and 45 min), diluted in series (5 dilutions of 10 fold each) in 96 well plates. 5 μ L of each dilution were spotted on a YPD agar plate and the plate was incubated for 2 days at 30°C.

Immunoblotting:

To determine steady state protein levels, strains were grown either on YPD or YPGal to mid-log phase and cells equivalent to an OD_{600} of 3 were harvested. Pellets were resuspended in 200 μ L of 0.1 M NaOH and incubated at room temperature (RT) for 5 min with 1400 rpm shaking. Samples were centrifuged (1500 rcf, 5 min, RT) and pellets were resuspended in 50 μ L of reducing Laemmli buffer (50 mM Tris-HCl, 2% SDS, 10% glycerol, 0.1% bromophenol blue, 100 mM DTT; adjusted to pH 6.8). Subsequently, samples were incubated for 5 min at RT, 1400 rpm shaking and heated for 10 min to 65°C prior to SDS-PAGE. 10 μ L of the sample were applied for standard SDS-PAGE and immunoblotting followed standard protocols. Blots were decorated with antibodies against Tom70, Cor1, Cor2, Cox1, Cox2, Cox4, Cox12, Cox13 and Rcf2 as described previously (Berndtsson *et al.*, 2020).

SDD-AGE:

Strains were grown to mid-log phase either in YPD or in YNB glucose. Protein extraction was performed according to the published protocol (Kushnirov *et al.*, 2006) but with a modified lysis buffer (100 mM Tris-HCl pH 7.5, 50 mM NaCl, 10 mM 2-mercaptoethanol, 2% (v/v) protease inhibitor cocktail (Sigma, USA), 2 mM phenylmethylsulfonyl fluoride (PMSF)). Cells were homogenized with a FastPrep-24 benchtop homogenizer (MP Biomedicals, USA) at 6.0 M/S for 30 s and then incubated on ice for at least 30 s. The procedure was repeated 7 times. The SDD-AGE was performed according to the published protocol (Kryndushkin *et al.*, 2003; Drozdova *et al.*, 2020). Gels with 1.5% (w/v) agarose were run at 30 V for 200–240 min. Proteins were transferred onto a polyvinylidene fluoride (PVDF) membrane according to the published protocol (Halfmann and Lindquist, 2008). The rabbit polyclonal anti-Sup35 (SE4290, (Chabelskaya *et al.*, 2004)) and anti-Rnq1 (unpublished data) antibody were used to detect Sup35 or Rnq1, respectively. The HRP conjugated secondary anti-rabbit antibody (GE Healthcare) was used for detection. Signals were recorded with the GeneGnome device (SynGene).

Mitochondrial isolation:

Mitochondria were isolated according to (Meisinger *et al.*, 2006). In brief, yeast cells were grown in YPGly media to mid-log phase and harvested by centrifugation (3000 rcf, 5 min, RT). Cells were washed once in distilled water and resuspended in 2 mL/g cell wet weight MP1 buffer (0.1 M Tris, 10 mM dithiothreitol, pH 9.4). After incubation for 10 min at 30°C, 170 rpm shaking, cells were washed once in 1.2 M sorbitol and subsequently resuspended in 6.7 mL/ cell wet weight MP2 buffer (20 mM potassium phosphate, 0.6 M sorbitol, pH 7.4, containing 3 mg/g of cell wet weight zymolyase 20T). Spheroplasts were created via incubation for 1 h at 30°C and harvested by centrifugation (3000 rcf, 5 min, 4°C). After careful resuspension in 13.4 mL/g of cell wet weight in ice-cold homogenization buffer (10 mM Tris, 0.6 M sorbitol, 1 mM ethylenediaminetetraacetic acid, 1 mM phenylmethylsulfonyl fluoride, pH 7.4), lysates were created by mechanical disruption via ten strokes with a Teflon plunger. Homogenates were centrifuged for 5 min at 3000 rcf, 4°C and the resulting supernatants were subsequently centrifuged at 17000 rcf for 12 min, 4°C. Pelleted mitochondria were resuspended in isotonic buffer (20 mM HEPES, 0.6 M sorbitol, pH 7.4) to a concentration of 10 mg/mL and stored at -80°C.

UV-VIS spectroscopy and CcO activity:

A Cary4000 UV-VIS spectrophotometer (Agilent Technologies) was used to record optical spectra (400-650 nm) from isolated mitochondria. 200 µg of mitochondria were lysed in 20 µL of lysis buffer (50 mM KPi pH 7.4, 150 mM KCl, 1× Complete Protease Inhibitor cocktail (Roche), 1 mM PMSF, 1% n-Dodecyl β-D-maltoside) for 20 min at 4°C and subsequently mixed with 130 µL of dilution buffer (50 mM KPi pH 7.4, 150 mM KCl) in a microcuvette and applied for measurement. To obtain reduced spectra, small amounts of sodium dithionite were admixed and measurements were repeated. Heme concentrations were determined from the differential spectrum (reduced-minus-oxidized) from 2 replicates by applying calculations as described recently (Berndtsson *et al.*, 2020). Concentrations of CcO were determined by applying an extinction coefficient of $\epsilon = 26 \text{ mM}^{-1} \text{ cm}^{-1}$ and volumes of isolated mitochondria corresponding to 10 nmol CcO were used for CcO activity measurements. Thereby, mitochondria were lysed in lysis buffer (50 mM Tris pH 7.4, 100 mM KCl, 1 mM EDTA, 1× Complete Protease Inhibitor cocktail (Roche), 1mM PMSF, 2% Digitonin) for 10 min, 4°C and clarified by centrifugation (25000 rcf, 10 min, 4°C). Lysates were diluted in measurement buffer (50 mM Tris pH 7.4, 100 mM KCl, 1 mM EDTA) and transferred into a Clark-type oxygen electrode. The measurement was started by adding 20 mM Na-Ascorbate, 50 µM Yeast Cyt. c, 40 µM N,N,N',N'-tetramethyl-p-phenylenediamine (TMPD) and recorded for 1-2 min. The slope of each analysis as a measure for oxygen consumption (µmol O₂/ml/min) was evaluated from 3 replicates and normalized to the average of the slope from wild type mitochondrial lysates to describe CcO activity as fold change.

Statistical analysis:

Outliers were defined as data points outside 1.5-fold interquartile range and of note, no outliers were detected with this method. Normal distribution of the data was confirmed by a Shapiro-Wilk's test (R studio, shapiro_test) and homogeneity of variances by a Leven's test (R studio, leveneTest). The means were compared by a One-Way ANOVA followed by a Tukey post hoc test (R studio, anova_test, tukey_hsd). Significances are indicated with asterisk (***P < 0.001, **P < 0.01, *P < 0.05, n.s. P > 0.05) and graphs were created via R studio using the ggplot2 library.

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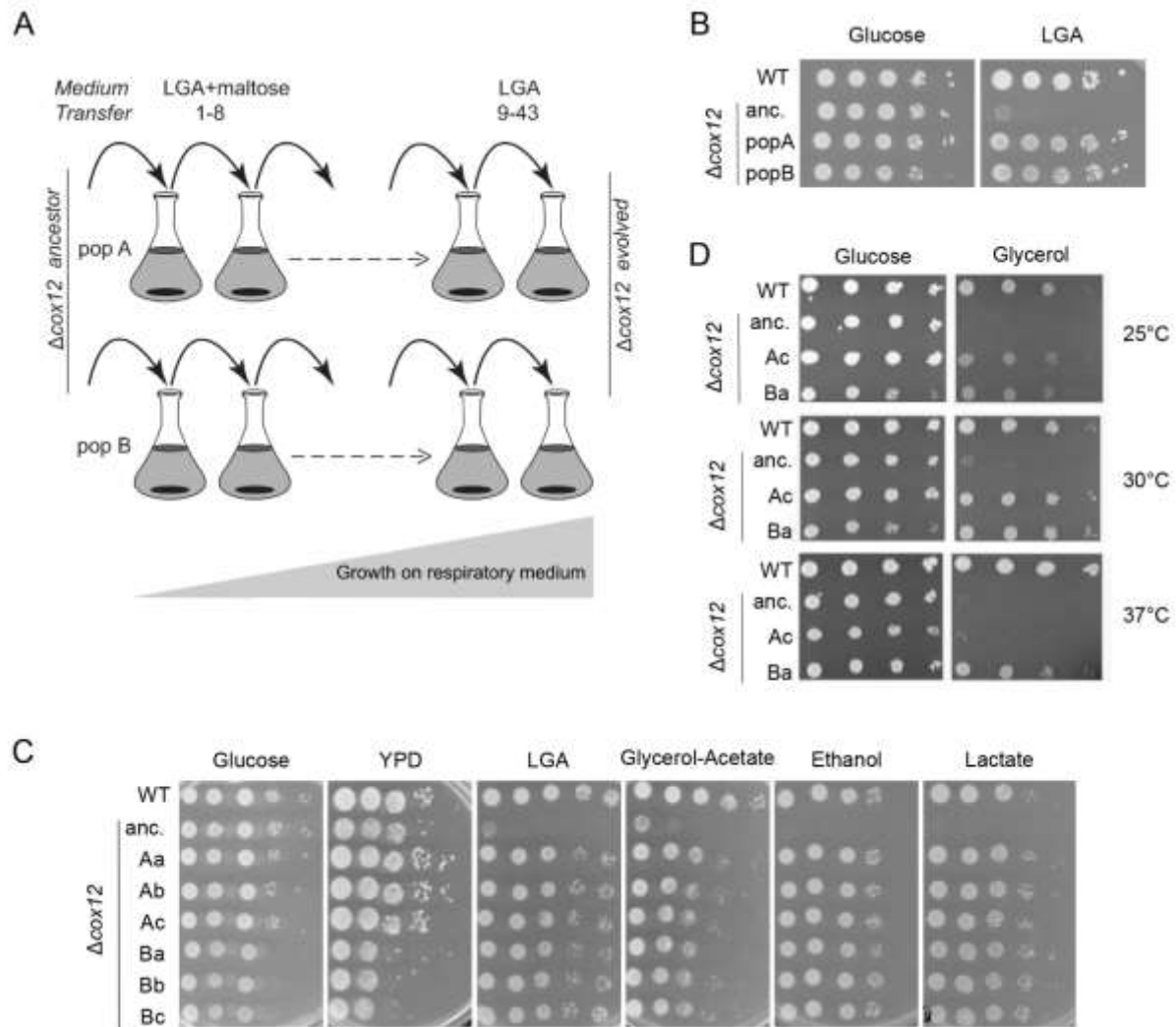


FIGURE LEGENDS

Figure 1: Phenotypic analysis of evolved Δcox12 cells. A) Schematic representation of the evolution experiment. B) 10-fold serial dilutions of control strains (WT and Δcox12 ancestor) and of populations A and B after transfer 43. The synthetic media contained either glucose, or lactate-glycerol-acetate (LGA) as in the evolution experiment. C) Serial dilution of control strains (WT and Δcox12 ancestor) and of clones a-c isolated from populations A and B. The plates contained either YPD medium or synthetic medium supplemented with the indicated carbon sources. The plates were imaged after 2 days (YPD and Glucose) or 4 days at 30°C. D) The indicated strains were tested for growth at various temperatures on synthetic medium containing glucose or glycerol. The plates are representative of 2 independent experiments (B-D).

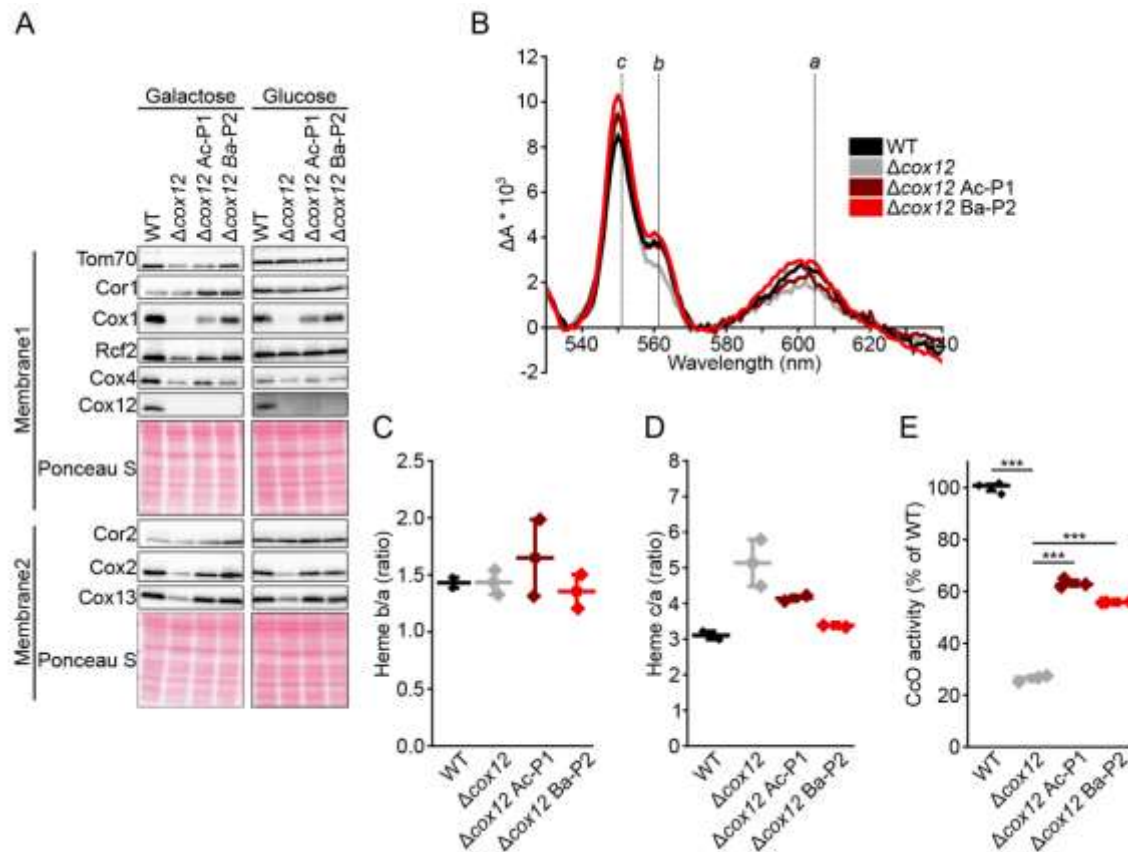


Figure 2: CcO protein levels and activity. A) Steady state levels of indicated proteins were analyzed in depicted strains during exponential growth in YP medium containing either galactose or glucose as carbon source. Ponceau S staining of respective membranes was performed as loading control. B-D) Heme spectra and heme ratios recorded from mitochondria isolated from indicated strains (n=2). E) CcO activity was measured as oxygen consumption in mitochondrial lysates as described in material and method section (n=3). 10 mmol CcO were used for each measurement, as quantified via heme content (B-D). Data was subsequently normalized to CcO activity from wild type mitochondrial extracts. Analyses in C-E are depicted as mean (square) \pm s.e.m., median (center line) and single data points (diamonds). Data in E was statistically analyzed by a One-Way ANOVA followed by a Tukey Post Hoc test. Single main effects are depicted as *** $P < 0.001$.

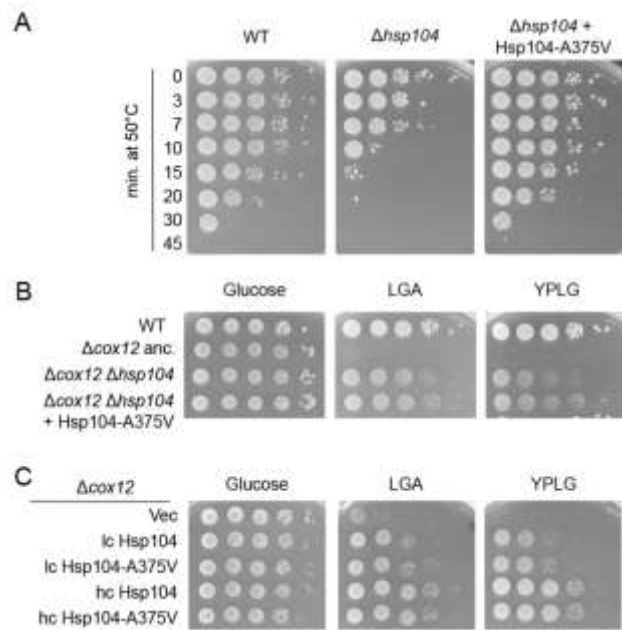


Figure 3: Deletion or overexpression of HSP104 rescue the respiratory deficiency of $\Delta cox12$. A) 10-fold serial dilutions of cultures of various strains exposed to a temperature of 50°C for the indicated time (0, 5, 10, 15, 20, 45 min) were deposited on YPD plates and grown for 2 days at 30°C. The data are representative of 2 independent experiments. B) 10-fold serial dilutions of control strains (WT and $\Delta cox12$ ancestor) and of $\Delta cox12 \Delta hsp104$ containing or not an integrated version of the Hsp104-A375V allele. C) 10-fold serial dilutions of the $\Delta cox12$ ancestor containing an empty plasmid (vec), low copy (lc) or high copy (hc) plasmids encoding Hsp104 or Hsp104-A375V.

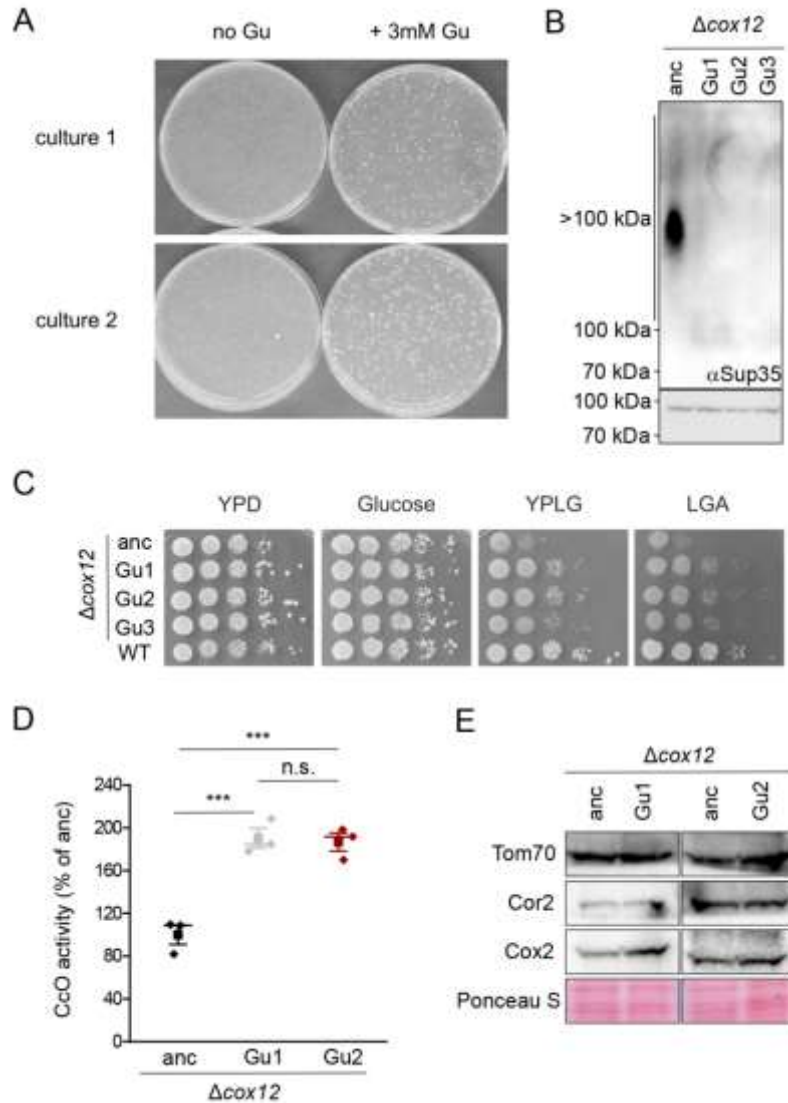


Figure 4: Gu treatment cures *[PSI⁺]* and improves respiratory growth and CcO activity of Δcox12 . A) Δcox12 cells were cultured in liquid YPD medium in the absence (no Gu) or presence of 3 mM Gu for 24 hours (culture 1) after which a second culture (culture 2) was inoculated in the same medium with a 100 fold dilution. After 24 hours, the cultures were diluted 5000 fold and an aliquot was spread on *pet* selection medium (YP 0.1% glucose, 2% glycerol). The plates were imaged after 4 days at 30°C and the data are representative of 2 independent experiments. B) SDD-AGE (upper panel) and SDS-PAGE (lower panel) results for protein lysates of analyzed strains. SDS-PAGE was used to check the presence of the Sup35 protein. Anti-Sup35 polyclonal antibodies were used for western blot hybridization. C) 10-fold serial dilutions of the indicated strains on YP-glucose (YPD), YP- 1% Lactate 1% Glycerol (YPLG) or synthetic medium supplemented with 2% glucose or 1% Lactate 0.2% Glycerol 0.2% Acetate (LGA). D) CcO activity measured via oxygen consumption in mitochondrial lysates (n=3) from the Δcox12 ancestor strain (anc) strain, as well as two independent clones of Gu treated cultures (Gu1 and Gu2). Data was normalized to the CcO activity from anc mitochondrial extracts and is depicted as mean (square) \pm s.e.m., median (center line) and single data points (diamonds). A One-Way ANOVA followed by a Tukey Post Hoc test was performed for statistical analysis and p values are depicted as ***P<0.001 and n.s. (not significant)>0.05. E) Steady state levels of indicated proteins analyzed in strains described in (D) during exponential growth. Ponceau S staining of respective membranes was performed as loading control.

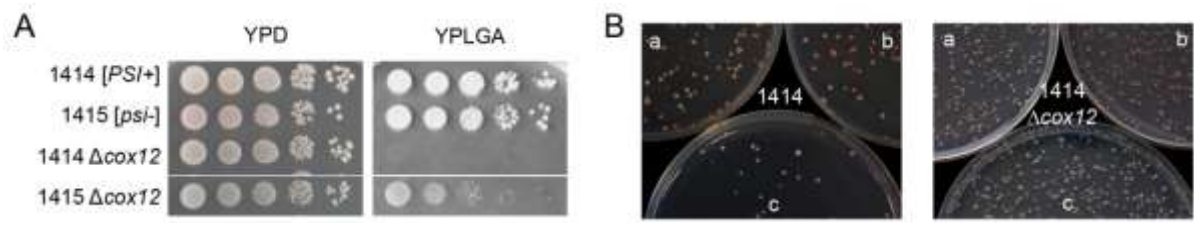


Figure 5: *[PSI+]* impairs the respiratory growth of Δcox12 and is cured efficiently by Hsp104-A375V. A) 10-fold serial dilutions of the indicated strains on YP-glucose (YPD) and YP supplemented with 10X adenine and 1% Lactate 0.2% Glycerol 0.2% Acetate (YPLGA). The plates were imaged after 3 days (YPD) or 8 days (YPLGA). B) Images of colonies from 1414 or 1414 Δcox12 strains transformed with a low copy vector encoding Hsp104 (a), Hsp104-A375V (b), or the empty vector (c) and grown for 4 days on YPD $\frac{1}{2}$ plates. Results are representative of 2 independent experiments (A-B).