Three genes for mitochondrial proteins suppress null-mutations in both Afg3 and Rca1 when over-expressed

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Abstract  The AFG3 gene of *Saccharomyces cerevisiae* encodes a mitochondrial inner membrane protein with ATP-dependent protease activity. To gain more insight into the function of this protein, multi-copy suppressors of an afg3-null mutation were isolated. Three genes were found that restored partial growth on non-fermentable carbon sources, all of which affect the biogenesis of respiratory competent mitochondria: *PIM1* (LON) encodes a matrix-localized ATP-dependent protease involved in the turnover of matrix proteins; *OXA1* (PET1402) encodes a putative mitochondrial inner membrane protein involved in the biogenesis of the respiratory chain; and *MBA1* encodes a mitochondrial protein required for optimal respiratory growth. All three genes also suppressed a null mutation in a related gene, *RCA1*, as well as in the combination of afg3- and rca1-null.

Key words  Multi-copy suppression · Mitochondrial membrane · Metalloprotease · *S. cerevisiae*

Introduction

Mitochondrial biogenesis in the yeast *S. cerevisiae* is an intricate process requiring the co-ordinated expression of mitochondrial and nuclear genetic systems (Grivell 1989, 1995). The latter system is quantitatively more important. For instance, with one exception (a mitoribosomal protein), all proteins necessary for mitochondrial gene expression are encoded in the nucleus and imported into the organelle. Nevertheless, mitochondrial genes encode essential components of enzyme complexes in the mitochondrial inner membrane involved in oxidative phosphorylation, making the mitochondrial genetic system essential for respiratory growth.

In yeast, several hundred nuclear genes are required for the maintenance of functional mitochondria, the so-called PET genes (Tzagoloff and Dieckmann 1991). One such gene, AFG3, was isolated previously by complementation of a mutant with a conditional pet phenotype (Guélin et al. 1994). Its translation product, Afg3p, is a protein with several interesting features. It contains the approximately 200 amino-acid domain characteristic of the AAA family of ATPases (Kunau et al. 1993; Confalonieri and Duguet 1995), together with a C-terminal domain containing the consensus sequence motif of a family of zinc metalloproteases (Campbell et al. 1994). Furthermore, hydrophobic regions in the amino-terminal domain mediate membrane association (Pajic et al. 1994). These features are shared with two other mitochondrial membrane proteins, Yme1p and Rca1p (Thorsness et al. 1993; Tzagoloff et al. 1994). Together with prokaryotic homologues, these proteins form a subfamily within the AAA family (Guélin et al. 1994). FtsH, the *E. coli* member of this subfamily, has been implicated in both the assembly/topogenesis of membrane proteins and the translocation of proteins across the plasma membrane (Tomoyasu et al. 1993; Akiyama et al. 1994a, b), as well as in protein degradation (Herman et al. 1993, 1995; Kihara et al. 1995; Tomoyasu et al. 1995). Similarly, Afg3p(Yta10p) is needed for rapid, ATP-dependent degradation of both prematurely terminated and complete mitochondrial translation products (Pajic et al. 1994; Guélin et al. 1996) and together with Rca1p, is required for the formation of mitochondrial inner membrane enzyme complexes (Paul and Tzagoloff 1995). The finding that inactivation of the proteolytic function of Afg3p by site-directed mutagenesis does not affect respiratory growth (Guélin et al. 1996) suggests that Afg3p may have a function besides the proteolysis of mitochondrial translation products.

One way to obtain more insight into Afg3p function is to identify other mitochondrial proteins that can (partially) bypass the need for Afg3p in mitochondrial biogenesis.
Three genes encoding such bypass proteins were identified by selecting for multi-copy suppressors of an afg3-null mutation. Interestingly, these genes also suppress an rca1-null mutation. The implications of these findings for the elucidation of Afg3p and Rca1p function are discussed.

Materials and methods

Strains and media. The S. cerevisiae strains used in this study are listed in Table 1. WDA1 and WDA1R were derived from wild-type strains W303/1A (Muroff and Tzagoloff 1990) and W303ΔRCA1, respectively, by disruption of AFG3 (see below). The following media were used for the propagation of yeast: YPD (2% glucose, 1% peptone, 1% yeast extract); YPGal (2% galactose, 2% peptone, 1% yeast extract); YPGly (2% glycerol, 2% peptone, 1% yeast extract); Lactate (1.5% lactic acid, 2% sodium lactate, 8 mM MgSO4, 45 mM (NH4)2HPO4, 0.5% yeast extract, pH 4.5); and WO [2% glucose, 0.87% yeast extract].

To learn more about AFG3 function we set out to identify genes which, when over-expressed, can compensate for the absence of functional AFG3. For this purpose, WDA1 was transformed with a yeast genomic DNA library on a multi-copy plasmid and the transformants were screened for growth on glycerol. From about 40 000 transformants, 27 fast-growing and 21 slow-growing clones were identified. Plasmids isolated from the fast-growing clones all exhibited a restriction pattern typical of the AFG3 locus (data not shown) and were not examined further. Upon re-transformation of the plasmids isolated from the slow-growing clones into WDA1, 14 were able to induce a slow-growth phenotype on glycerol, thus confirming the presence of suppressor genes on these plasmids. By restriction mapping, the inserts in these plasmids were divided into three classes. Class A consisted of only one insert (Fig. 1), class B of five indistinguishable inserts (Fig. 2), and class C of eight inserts grouped into three overlapping sets (Fig. 3). For the first two inserts, the sequences obtained were present in the EMBL database, and the restriction maps obtained corresponded to those predicted for the region between each set.

Isolation of multi-copy suppressors of an afg3-null mutant

The chromosomal regions from which the three classes of inserts were derived were identified by sequencing into the borders of the inserts in YEp13/34 (class A), YEp13/36 (class B), and pJN-C1, a plasmid containing the region of overlap between the inserts of class C (Fig. 3). For the first two inserts, the sequences obtained were present in the EMBL database, and the restriction maps obtained corresponded to those predicted for the region between each set.

Table 1 Genotypes and sources of the S. cerevisiae strains used

<table>
<thead>
<tr>
<th>Strain</th>
<th>Nuclear genotype</th>
<th>Mitochondrial genotype¹</th>
<th>Source</th>
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<tbody>
<tr>
<td>WDA1</td>
<td>ade2-1, his3-11, 15, leu2-3, 112, ura3-1, trp1-1, can1-100, afg3::URA3</td>
<td>(W303) This study</td>
<td></td>
</tr>
<tr>
<td>W303ΔRCA1</td>
<td>ade2-1, his3-11, 15, leu2-3, 112, ura3-1, trp1-1, can1-100, rca1::URA3</td>
<td>(W303) Tzagoloff et al. (1994)</td>
<td></td>
</tr>
<tr>
<td>WDA1R</td>
<td>ade2-1, his3-11, 15, leu2-3, 112, ura3-1, trp1-1, rca1::URA3, afg3::TRP1</td>
<td>(W303) This study</td>
<td></td>
</tr>
<tr>
<td>DAY6</td>
<td>his3-11, 15, leu2-3, 112, ura3-1, afg3::URA3</td>
<td>(167) Guélin et al. (1996)</td>
<td></td>
</tr>
</tbody>
</table>

¹ (167) Mitochondrial DNA is from KAR(167) and contains no introns (Séraphin et al. 1987)
of sequences. Subsequent deletion analysis localized the suppressor activities to single open reading frames (ORFs) on each DNA-fragment (Figs. 1 and 2). Both ORFs were characterized earlier and encode a mitochondrial ATP-dependent protease, Pim1p or Lon (Suzuki et al. 1994; Van Dyck et al. 1994), and a putative mitochondrial membrane protein, Oxa1p or Pet1402p (Bauer et al. 1994; Bonnefoy et al. 1994), respectively. The third insert was hybridized to a chromosome blot, and found to be homologous to sequences on chromosome II (data not shown). The sequence of the insert of plasmid pJN-C1 was kindly provided by M. Jacquet and was later published as a contribution to the chromosome-II sequencing project (Demolis et al. 1994).

As shown in Fig. 3, deletion analysis identified YBR1307 as the ORF responsible for suppression. The new gene, termed MBA1, for (Multi-copy By-pass of AFG3), encodes a putative 278 amino-acid protein whose N-terminus displays features characteristic of mitochondrial targeting sequences (Hartl et al. 1989). We have indeed found that MBA1 encodes a mitochondrial protein that is necessary for optimal mitochondrial function (Rep and Grivell 1996).

**Suppression of rca1-null**

Af3p belongs to the FtsH subfamily of the AAA family which includes both mitochondrial and prokaryotic members (Confalonieri and Duguet 1995). Yeast mitochondria contain two additional members of this subfamily, Yme1p(Yta11p) and Rca1p(Yta12p). Of these, Rca1p is the most similar to Af3p in sequence and structure. Disruption of RCA1 and AFG3 leads to similar phenotypes (see Discussion), and both proteins probably share the same topology in the mitochondrial inner membrane (Guélin et al. 1994; Schnall et al. 1994). The list of these similarities is here extended by the observation that the three multi-copy suppressors of af3-null suppress an rca1-null mutation to a similar extent (Fig. 4 A). However, Af3p and Rca1p cannot replace each other, as witnessed by the pet phenotype of the respective disruption strains. Even over-expression of either gene cannot overcome the res-

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**Fig. 1** Restriction map and subcloning of the class-A insert. This insert contains two complete open reading frames (ORFs): HAP3 (Hahn et al. 1988) and PIM1/LON (see text). PIM1 was identified as the gene responsible for the suppression of af3-null by showing that (1) pJN-A1, which contains only HAP3, does not suppress af3-null, (2) pJN-A3, which contains an internal (DraIII-Bsu36I) deletion in HAP3 (causing a frameshift after codon 57), still suppresses af3-null, and (3) pJN-A4, which contains PIM1 but has most of its promoter deleted, does not show suppression. Boxes indicate ORFs, arrows the direction of transcription. Thin lines indicate yeast genomic DNA sequences, bold lines indicate YEp13-sequences. Restriction sites used for mapping and/or subcloning are shown (the second BamHI site is not present in the sequence deposited with the EMBL database; it presumably reflects strain polymorphism). Suppression (growth on glycerol of a transformed af3-null strain) is indicated by +, no suppression by –.

**Fig. 2** Restriction map and subcloning of the class-B insert. This insert contains three complete and two partial ORFs. Subclones that allowed identification of OXA1 as the gene responsible for the suppression of af3-null are shown. See legend of Fig. 1 for explanation of symbols.
piratory defect caused by the absence of the other (data not shown). One interpretation of these observations is that both proteins are required for a single activity. If this is true, the three multi-copy suppressors should also suppress an \textit{afg3}/\textit{rca1} double disruption. This is indeed the case (Fig. 4B).

Discussion

A search for multi-copy suppressors of an \textit{afg3-null} mutation has been employed to obtain clues to the primary defect in mitochondrial function caused by deletion of \textit{AFG3} and its homologue \textit{RCA1}. This defect manifests itself in the strong reduction in the amount and activity of respiratory chain complexes (Tauer et al. 1994; Tzagoloff et al. 1994) and a deficiency in the assembly of ATP synthase (Paul and Tzagoloff 1995). Clearly, the assembly of the mitochondrial inner membrane complexes does not pro-
ceed normally in \textit{afg3} and \textit{rcal} mutants. This fits well with the proposed function of the \textit{E. coli} homologue of these genes, \textit{FtsH}, in the assembly and/or topology of plasma membrane proteins (Tomoyasu et al. 1993; Akiyama et al. 1994 a, b).

On the other hand, mutations in \textit{FtsH} have been found to stabilize different cellular proteins (Herman et al. 1993, 1995; Kihara et al. 1995), and disruption of \textit{AFG3} (\textit{YTA10}) leads to stabilization of otherwise rapidly degraded incomplete (Pajic et al. 1994) and complete (Guélin et al. 1996) mitochondrial translation products. Although involvement of \textit{FtsH} in protein degradation has been suggested to be indirect, by presenting target proteins to a protease (Herman et al. 1993; Kihara et al. 1995), a protease activity of the members of the \textit{FtsH}-subfamily is supported by their conserved metalloprotease active-site motif (Campbell et al. 1994), combined with the dependence of turnover of mitochondrial translation products on divalent metal-ions (Nakai et al. 1994; Yasuhara et al. 1994). Moreover, degradation of the heat-shock transcription factor sigma32 by purified \textit{FtsH} has recently been demonstrated in vitro (Tomoyasu et al. 1995). These observations have led some investigators to conclude that all defects in \textit{ftsH} cells will eventually be shown to result from the defective turnover of different cellular proteins (Herman et al. 1995).

Such an interpretation seems less likely for explaining the \textit{pet} phenotype of yeast cells upon disruption of \textit{AFG3}, since specific mutational inactivation of the protease activity of \textit{Afg3p} does not affect respiratory growth (Guélin et al. 1996). Rather, Afg3p may be directly involved in complex assembly. The identification of \textit{OXA1} as one of the suppressors supports this notion, as it also has a role in the assembly of integral membrane complexes. Disruption of \textit{OXA1} results in a complete absence of cytochrome \textit{aa3} and reduction of cytochrome \textit{b}, while mitochondrial translation products are present (Bonnefoy et al. 1994). Other researchers have isolated the same gene as \textit{PET1402} by a screen for mutants defective in the processing of \textit{CoxlI} (Bauer et al. 1994). In addition to the \textit{CoxlI}-processing defect, a general reduction of mitochondrial protein synthesis was observed in \textit{pet1402} mutants. Finally, \textit{FIF0-ATPase} assembly is also affected in \textit{oax1} cells (Altamura et al. 1996). Whatever the exact function of \textit{Oxa1p}, it is not specific for the assembly of a mitochondrial-type respiratory chain since \textit{Oxa1} homologues have also been found in bacteria (Bonnefoy et al. 1994). It will be interesting to see if these homologues (or \textit{Oxa1} itself) can suppress mutations in \textit{FtsH}.

Preliminary data indicate that the second suppressor, \textit{MBA1}, also affects the assembly of the respiratory chain, although to a lesser extent than \textit{AFG3}, \textit{RCA1} or \textit{OXA1} (data not shown). However, the finding that \textit{PIM1} (\textit{LON}) is one of the suppressors apparently disturbs the notion of assembly being the primary defect in \textit{afg3-null} cells. \textit{PIM1} encodes an ATP-dependent protease of the mitochondrial matrix which until now has only been implicated in the maintenance of mitochondrial DNA and the turnover of proteins localized to the matrix (Suzuki et al. 1994; Van Dyck et al. 1994; Wagner et al. 1994). It is therefore tentative to assume that \textit{Pim1p} can take over a degradation function of Afg3p when present in high amounts. However, preliminary data indicate that suppression by \textit{PIM1} may be independent of its protease activity (Rep et al. 1996).

Our observation that the three multi-copy suppressors act on both \textit{afg3-null} and \textit{rcal-null} mutations, as well as on a combination of these, even though \textit{AFG3} and \textit{RCA1} have non-overlapping functions, suggests that Afg3p and Rcalp are both needed for a single activity which is essential for the biogenesis of respiratory competent mitochondria. This is also supported by the similar assembly defects seen in the single- and double-disruption mutants (Paul and Tzagoloff 1995). One obvious possibility is that they are part of a heteromultimeric complex. Complex formation has been suggested before for \textit{Yme1p} (Thorsness and Fox 1993; Thorsness et al. 1993; Nakai et al. 1995) and recently demonstrated for \textit{FtsH} (Akiyama et al. 1995). Moreover, formation of (ring-shaped) multimers appears to be common among members of the \textit{AAA} family (Peters et al. 1990, 1993; Whiteheart et al. 1995; Frohlich et al. 1995).

In conclusion, this study shows that the roles of \textit{OXA1} and \textit{AFG3/RCA1} in the assembly of mitochondrial inner membrane complexes are partially overlapping, and identifies two new actors, \textit{PIM1} and \textit{MBA1}, in the same process. The main question that we are now addressing is how these proteins affect the assembly process at the molecular level.

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\textbf{References}


Demolis N, Mallet L, Jacquet M (1994) A 12.5-kb fragment of the yeast chromosome X contains two adjacent genes encoding ribosomal proteins and six putative new genes, one of which encodes a putative transcriptional factor. Yeast 10:1151–1152


Muroff I, Tzagoloff A (1990) CBP7 codes for a co-factor required in conjunction with a mitochondrial maturase for splicing of its cognate intervening sequence. EMBO J 9:2765–2773


Nakai T, Yasuhara T, Fujiki Y, Ohashi A (1995) Multiple genes, including a member of the AAA family, are essential for degradation of unassembled subunit 2 of cytochrome c oxidase in yeast mitochondria. Mol Cell Biol 15:4441–4452


Thorsness PE, Fox TD (1993) Nuclear mutations in Saccharomyces cerevisiae that affect the escape of DNA from mitochondria to the nucleus. Genetics 134:21–28


