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The Tim54p–Tim22p Complex Mediates Insertion of Proteins into the Mitochondrial Inner Membrane

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Abstract. We have identified a new protein, Tim54p, located in the yeast mitochondrial inner membrane. Tim54p is an essential import component, required for the insertion of at least two polytopic proteins into the inner membrane, but not for the translocation of precursors into the matrix. Several observations suggest that Tim54p and Tim22p are part of a protein complex in the inner membrane distinct from the previously characterized Tim23p-Tim17p complex. First, multiple copies of the TIM22 gene, but not TIM23 or TIM17, suppress the growth defect of a tim54-1 temperature-sensitive mutant. Second, Tim22p can be coprecipitated with Tim54p from detergent-solubilized mitochondria, but Tim54p and Tim22p do not interact with either Tim23p or Tim17p. Finally, the tim54-1 mutation destabilizes the Tim22 protein, but not Tim23p or Tim17p. Our results support the idea that the mitochondrial inner membrane carries two independent import complexes: one required for the translocation of proteins across the inner membrane (Tim23p–Tim17p), and the other required for the insertion of proteins into the inner membrane (Tim54p–Tim22p).

Mitochondrial function depends on the import of hundreds of different proteins synthesized in the cytosol. Protein import is a multistep pathway which includes the binding of precursor proteins to surface receptors, translocation of the precursor across one or both mitochondrial membranes, and folding and assembly of the imported protein inside the mitochondrion (for review see Ryan and Jensen, 1995; Schatz and Dobberstein, 1996; Jensen and Kinnally, 1997; Pfanner and Meijer, 1997). Most precursor proteins carry amino-terminal targeting signals, called presequences (Hurt et al., 1984, a, b, 1985; Horwich et al., 1985; van Loon et al., 1986), and are imported into mitochondria via import complexes located in both the outer and the inner membrane (IM)1. The mitochondrial outer membrane of Saccharomyces cerevisiae contains at least four proteins, Tom70p, Tom37p, Tom22p, and Tom20p (Hase et al., 1984; Hines et al., 1990; Steger et al., 1990; Moczko et al., 1993; Ramage et al., 1993; Gratzer et al., 1995; Hönlinger et al., 1995), which have been proposed to recognize the targeting signal carried on imported mitochondrial proteins. These four receptors interact with at least six other polypeptides, and together comprise the TOM complex, which facilitate the movement of proteins across the outer membrane (Kiebler et al., 1990; Söllner et al., 1992).

In the IM, a TIM complex mediates the translocation of proteins into the matrix. The TIM complex consists of at least two integral IM proteins, Tim23p and Tim17p, which have been proposed to form part of a protein-translocating channel (Dekker et al., 1993; Emtage and Jensen, 1993; Maarse et al., 1994, 1996; Blom et al., 1995). Tim23p, a 23-kD protein, was first identified as a temperature-sensitive mutant defective in the import of several different matrix-localized precursor proteins (Emtage and Jensen, 1993). TIM17 was identified as a multi-copy suppressor of the tim23 mutant, and was shown to encode a 17-kD protein homologous to the carboxyl-terminal domain of Tim23p (Ryan et al., 1994). The essential Tim23p and Tim17 proteins cooperate with Tim44p, located on the inside of the IM (Scherer et al., 1992; Rassow et al., 1994), and mt-Hsp70, a matrix-localized member of the 70-kD heat shock family (Kang et al., 1990; Scherer et al., 1992; Rassow et al., 1994). Tim44p and mt-Hsp70 are proposed to pull the imported precursor protein through the IM translocation channel into the matrix (Kromdou et al., 1994; Rassow et al., 1994; Schneider et al., 1994; Unger mann et al., 1994, 1996; Blom et al., 1995).

Tim23p, Tim17p, Tim44p, and mt-Hsp70 appear to form functional complexes in the IM. Tim23p and Tim17p cofractionate after detergent solubilization of the mitochondria (Berthold et al., 1995; Blom et al., 1995; Ryan et al., 1994, 1996).
1998). Both proteins can also be chemically cross-linked to each other in intact mitochondria (Berthold et al., 1995; Blom et al., 1995; Ryan et al., 1998). When a precursor destined for the matrix was arrested in transit across the IM, a large complex containing Tim44p, Tim23p, Tim17p, and mt-Hsp70 (and presumably other proteins) communicated with the precursor (Berthold et al., 1995). It has recently been shown that Tim23p may exist in two subcomplexes, one complex consisting of Tim23p, Tim44p, and mt-Hsp70, and another complex with Tim23p, Tim17p, and mt-Hsp70 (Bömer et al., 1997). The function of the two subcomplexes in import is not known.

Recently, an essential IM protein, Tim22p, homologous to both Tim23p and Tim17p, has been identified (Sirrenberg et al., 1996). Tim22p is not required for the import of matrix-localized proteins, but Tim22p appears to be essential for the correct insertion of two membrane proteins, Aac1p, the ATP/ADP carrier, and PiC, the phosphate carrier, into the IM. Whether Tim22p is required for the import of other membrane proteins besides carrier proteins is not known. Tim22p is proposed to be in a separate complex from Tim23p and Tim17p in the IM, but other members of this new complex have not been identified. We have identified a new protein, Tim54p, which is essential for the IM insertion of the Aac1 carrier and Tim23 import proteins. Tim54p is not required for the translocation of precursors across the IM into the matrix. We find that Tim54p physically interacts with Tim22p, and forms an import complex in the IM separate from the Tim23p–Tim17p import machinery. Our results indicate that there are two import pathways in the IM. One pathway uses Tim23p and Tim17p in the translocation of precursors across the IM, and the other requires Tim54p and Tim22p for the insertion of proteins into the IM. Because Tim54p and Tim22p are both essential for yeast cell viability, we suggest that the Tim54p–Tim22p complex plays a key role in the insertion of many, if not all, proteins into the IM.

Materials and Methods

Strains and Relevant Genotypes

MATa gal4 cyh2 trpl-901 leu2-3,112 URA3::GAL1 lacZ LYS2:: GAL1::HIS3 strain Y190 (Bai and Elledge, 1996), MATa/MATa leu2-Δ1 leu2-Δ1 strain YPH501 (Sikorski and Hieter, 1989), MATa/MATa ura3-52 ade2-1 ura3-52 trpl-1::trpl-Δ1 strain YPH857/858 and MATa ade2-1 trpl-Δ1 strain YPH857 (Spencer et al., 1993), MATa/MATa trpl-1::Δtrpl-Δ3 his3-Δ200his3-Δ200 strain FY833/834 and MATa trpl-1::Δtrpl-Δ3 his3-Δ200 strain FY833 (Winston et al., 1995), MATa tim23-1 ura3-52 strain 574 (Ryan et al., 1994), MATa TIM23 tim23 strain 55 and MATa tim23 strain 201 (Emtage and Jensen, 1993), MATa tim23::URA3 leu2 strain 474 (Emtage and Jensen, 1993), and MATa tim17::TRPI leu2 strain 463 (Ryan et al., 1994) have been described. MATa/MATa ade2ade2 ade2ade3 trpl1 ura3ura3 leu2leu2 strain 534 was constructed by crossing strains 4795-202 and 4795-408 (gifts from D. Koshland, Carnegie Institute of Washington, Baltimore, MD). Yeast transformations were performed as described (Schiestl and Geitz, 1989). Standard yeast media and genetic techniques (Rose et al., 1988) were used.

Isolation of TIM54 and TIM22

Tim54p was identified as a potential interactor with the Mmm1 protein (Burgess et al., 1994) using the two-hybrid screen (Fields and Song, 1989; Bai and Elledge, 1996). pOK52, a TRPI plasmid which expresses amino acids 123–426 of Mmm1p fused to the Gal4 DNA-binding domain, was constructed by inserting a 2.6-kbp NheI–NcoI fragment from pSB75 into the NheI–CylH site of pAS1-CYLH2 (Bai and Elledge, 1996). pOK52 was transformed into yeast strain Y190 along with a library of yeast cDNAs fused to the Gal4 activation domain carried on the LEU2 plasmid pACT (Bai and Elledge, 1996). Y190 carries both the Escherichia coli lacZ and the yeast HIS3 genes under the control of the GalI promoter region (Bai and Elledge, 1996). 104 transformants were plated onto medium containing 50mM 3-amino-triazeole (Sigma Chemical Co., St. Louis, MO) to select for HIS3 expression, and ~10,000 colonies were subsequently screened for β-galactosidase activity. Five ‘‘gal’’ colonies were identified and the plasmid from one transformant (pACT-15) was shown to specifically interact with pOK52. For example, pACT-15 did not interact with other proteins fused to the Gal4 DNA-binding domain, such as p53, yeast Snf1, and rat lamin. pACT-15 contained a cDNA of 750 bp, which was excised with BglIII and subcloned into the BamHI site of Bluescript SK II + (Stratagene, La Jolla, CA) to form pOK49.

The yeast DNA insert in pOK49 was sequenced completely and shown to encode the carboxyl-terminal region of a novel protein. Database analysis using the BLAST program (Altschul et al., 1990) showed the open reading frame was located upstream of the PEP8 gene (Bachhawat et al., 1994). Plasmid p7171, which carries a 7-kbp SpeI fragment in pRS316 (a gift from E. Jones, Carnegie Mellon University, Pittsburgh, PA), was used to isolate a 3.1-kbp ClaI fragment which was blunt-end ligated into the EcoRV site of Bluescript SK II +, forming pOK21A. Complete DNA sequence of the insert in pOK21A (and subclones from the insert) identified an open reading frame encoding a 54-kD protein.

We identified the TIM22 gene as an open reading frame (YDL217C) on chromosome IV (Jacq et al., 1997) encoding a protein homologous to Tim22p and Tim17p. TIM22 was isolated by PCR-amplifying a 1.4-kbp fragment from yeast genomic DNA using oligonucleotide No. 178 (5’-GATCGAGCTCGTATACAGAGAGATGTC-3’) and oligonucleotide No. 180 (5’-CATGCTCTAAGCCCTGC-3’). The PCR fragment was digested with EcoRI and inserted into the EcoRI site of either the Cen6/URA3 plasmid, pRS316 (Sikorski and Hieter, 1989), or the 2μ-URA3 plasmid, pRS426 (Sikorski and Hieter, 1989), forming pHJ201 and pHJ202, respectively.

Disruption of the TIM54 and TIM22 Genes

Two different disruptions of the TIM54 open reading frame were constructed. First, tim54::URA3 was produced by digesting pOK52 with StyI, which removes nucleotides 1156–1409 from the TIM54 open reading frame, and the DNA ends were filled in with DNA polymerase. A 1.1-kbp Smal fragment containing the URA3 gene was isolated from plasmid pRS304 (gift from S. Mizzen, Johns Hopkins School of Medicine, Baltimore, MD) and blunt-end ligated into the StyI-cut pOK49 vector to form pOK3. For genes disruption in yeast, a 1.5-kbp Xhol fragment containing tim54::URA3 was isolated from pOK3 and transformed into ura3ura3 diploid strain YPH857/858. Southern analysis of this diploid confirmed that one of the two copies of TIM54 had been replaced by tim54::URA3.

A second TIM54 disruption was constructed by inserting a 3.1-kbp XhoI fragment containing LEU2 into the NcoI site located at position 230 in the TIM54 open reading frame. pOK21A, which carries a TIM54 on a 3.1-kbp ClaI fragment blunt-end ligated into the EcoRV site of Bluescript SK II + (Stratagene), was digested with EcoRI and inserted into the EcoRI site of either the Cen6/URA3 plasmid, pRS316 (Sikorski and Hieter, 1989), or the 2μ-URA3 plasmid, pRS426 (Sikorski and Hieter, 1989), forming pHJ201 and pHJ202, respectively.

A complete disruption of the TIM22 open reading frame was constructed as described (Lorenz et al., 1995). A DNA fragment was amplified from pRS304 (Sikorski and Hieter, 1989) using oligonucleotides No. 181 (5’-TAAGATCAAGAAATGAAGTTAATAACAGTTTATACG-3’) and No. 182 (5’-AAATAAT-AAAAACACTTCAGTCTTGGTCAAATGCTATCCAGATGTTA-CTGAGTGATCCAC-3’) using the PCR (Saiki et al., 1985). This DNA fragment contained 40 bp of TIM22 5’ untranslated sequences joined to one end of the TRPI gene and 40 bp of TIM22 3’ untranslated sequences joined to the other end of TRPI. Homologous recombination between the PCR product and the yeast chromosomal DNA produces a precise replacement of the TIM22 ORF by TRPI. The PCR product was transformed into trpl trpl diploid strain FY833/834, and Trp+ transformants were selected. In one transformant (strain 777), one of the two copies of the TIM22 open reading frame was replaced by TRPI. Meiotic analysis of transformed diploids yielded only two viable spores in each tetrad. Similar to the TIM54 disruptions, tim22::TRPI spores germinated and underwent

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Construction of HA Epitope-Tagged Versions of Tim54p and Tim22p

pOK100, which contains a unique NotI site immediately preceding the termination codon of TIM54, was constructed as follows. Using pOK49, oligonucleotide No. 123 (5'-CTTGGGCGGACATATGTTGCGGATCC-3'), and oligonucleotide No. 98 (5'-AACGCTATGACCATACAAA-3'), we isolated a 2.1-kbp DNA fragment using PCR, and digested the fragment with Xhol and NotI. A 954-bp NotI-BamHI fragment containing a unique NotI site preceding the TIM54 termination codon and the 3' untranslated region of TIM54 was isolated from pJE7 (Emtage and Jensen, 1993). Both fragments were inserted into BamHI-Sall digested pUC18 (Yanisch-Perron et al., 1985) to form pOK101, which encodes Tim54p with the hemagglutinin (HA) epitope at its carboxyl terminal, was constructed by inserting a 114-bp NotI fragment containing three tandem copies of the HA epitope (Field et al., 1988; Tyers et al., 1992; a gift from B. Fuchter, Cold Spring Harbor Laboratory, Cold Spring Harbor, NY) into pOK100. Because pOK101 lacks complete TIM54 sequences, we reconstructed TIM54 by integration into the yeast chromosome. A 3.6-kbp PvuI fragment from pOK101, carrying the TIM54-HA fusion, was isolated into the URA3 integrating vector pRS06 (Sikorski and Hieter, 1989) to form pOK20. pOK102 was transformed into ura3::ura3 diploid YPH57/88, and we targeted the integration to the chromosomal TIM54 gene by digesting pOK102 with HpaI. Stable Ura" transformants were isolated and shown to express Tim54-HA. Integration of the TIM54-HA construct at the chromosomal TIM54 gene was confirmed by Southern analysis. Isolation of mito-specific products from spoulted diploid transformants yielded haploid cells containing the integrated TIM54-HA, which grew on both glucose and glycerol-containing medium at 23°, 30°, and 37°C. Because integration of the pOK102 plasmid inactivates the chromosomal copy of the essential TIM54 gene, our results indicate that the Tim54-HA fusion protein is fully functional.

pH301, a LEU2 plasmid that carries the full-length TIM54-HA gene, was constructed as follows. First, a 1.5-kbp EcoRI fragment carrying the carboxyl-terminal region of Tim54p fused to HA was isolated from pOK102 and inserted into EcoRI-digested pOK21B. pOK21B carries the full-length TIM54 gene on a 3.1-kbp ClaI fragment blunt-end ligated into the EcoRV site of Bluescript SK II (Stratagene). Replacement of the 1.66-kbp pOK102 plasmid that carries the full-length TIM54-HA gene by digesting pOK102 with HpaI. Stable Ura" transformants were isolated and shown to express Tim54-HA. Integration of the TIM54-HA construct at the chromosomal TIM54 gene was confirmed by Southern analysis. Isolation of mito-specific products from spoulted diploid transformants yielded haploid cells containing the integrated TIM54-HA, which grew on both glucose and glycerol-containing medium at 23°, 30°, and 37°C. Because integration of the pOK102 plasmid inactivates the chromosomal copy of the essential TIM54 gene, our results indicate that the Tim54-HA fusion protein is fully functional.

Isolation of the Temperature-sensitive TIM54-1 Mutant

tms5-1 was produced by mutagenesis of the cloned TIM54 gene, pOK22, which carries TIM54 gene on a 3.1-kbp ClaI fragment blunt-end ligated into the EcoRV site of the TRP1-CY2-containing plasmid, pKS1 was passed through bacterial mutator strain XL1-RED (Stratagene) according to manufacturer’s instructions. Mutagenized pOK22 was transformed into ade2 ade3 ade4 ade6 trp1 ura3 leu2 tms5-1 strain 551, which also carries the TIM54::URA3 plasmid, pOK32. tms5-1 mutants of pOK22 were identified by a plasmid-shuffle scheme (Sikorski and Boeke, 1991). Specifically, if the TIM54 gene carried on pOK22 was defective, then transformants were unable to lose the TIM54::ADE3-URA3 plasmid (and the ade2 ade3 cells accumulated a red pigment). If TIM54 on pOK22 is functional, then transformants could lose the pOK32 plasmid (and the ade2 ade3 cells remained white). From 37,400 total transformants, 175 completely red colonies were identified as unable to lose the pOK32 plasmid at 34°C. One transformant was found to be temperature sensitive for TIM54 activity: pOK32 could be lost at 24°C, but not at 34°C. The majority of the other transformants were defective in TIM54 at both temperatures. Plasmid DNA (called pOK24) was isolated from this colony and shown to confer temperature-sensitive growth when reintroduced into strain 551. Subcloning experiments showed that the mutation was located in the TIM54 gene. MATa trp1 tms5-1 strain 809, which carries a chromosomal version of tms5-1, was constructed as follows. trp1 tms54::URA3 strain 835, carrying plasmid pOK24, was patched onto medium containing 5-fluoro-orotic acid (Boeke et al., 1984) to select for cells in which the chromosomal tim54::URA3 gene was replaced with the plasmidborne tim54-1 sequence by homologous recombination.

pOK32 was constructed by inserting TIM54 carried on a 3.5-kbp PvuI fragment isolated from pOK21A into the 2µ-URA3 plasmid pRS426 (Sikorski and Hieter, 1989), forming pOK30. Subsequently, the ADE3 gene was isolated as a 4.8-kbp BamHI-Sall fragment from pDK255 (a gift from D. Koshland) and inserted into BamHI-Sall-cut pOK30 to form pOK32. Yeast strain 551 was constructed as follows. First, a disruption of

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Kerscher et al. Mitochondrial IM Protein Insertion Complex
**Imports into Isolated Mitochondria**

Mitochondria were isolated from yeast strains as described (Daum et al., 1982), except that SEH buffer (250 mM sucrose, 1 mM EDTA, 20 mM Hepes-KOH, pH 7.4) was sometimes used in place of breaking buffer, and temperature-sensitive strains were grown at 24°C. For import reactions, mitochondria were suspended in import buffer (Scherer et al., 1995). Briefly, we added 0.5 ml of sample buffer containing 40 μM indicated reactions were treated with proteinase K (Sigma Chemical Co.) on ice for 30 min, followed by the addition of 1 mM PMSF (Sigma Chemical Co.). The outer membrane of mitochondria were disrupted to form mitoplasts by resuspending mitochondrial pellets in 20 mM Hepes-KOH, pH 7.4, followed by incubation at 0°C for 20 min. After all manipulations, mitochondria and mitoplasts were pelleted by centrifugation at 12,500 g for 10 min, resuspended in 1× sample buffer containing 4% β-mercaptoethanol, and analyzed by SDS-PAGE. Radiolabeled proteins were detected by fluorography (Bonner and Laskey, 1974) or by using a phosphorimager (Molecular Dynamics, Inc.). To quantitate data, we used the Molecular Dynamics ImageQuant Software.

**Immune Precipitations from Detergent-solubilized Mitochondria**

Mitochondria were isolated from strain 494 that expresses Tim54-HA from the integrated pOK102 construct, TIM22-TRP1 strain 800 carrying TIM22-HA plasmid pH102, or from wild-type strain D273-10b (Sherman, 1964) as described above. For immune precipitations, mitochondria were solubilized as described (Berthold et al., 1995). Briefly, we added 0.5 ml solubilization buffer (0.5% digitonin, 50 mM NaCl, 30 mM Hepes-KOH, pH 7.4, 1 mM PMSF), 1 μg/ml aprotinin (CalBiochem Corp., La Jolla, CA), 1 μg/ml leupeptin (CalBiochem Corp.), 1 μg/ml chymostatin (Sigma Chemical Co.), 1 μg/ml antipain (Sigma Chemical Co.), 1 μg/ml pepstatin A (Sigma Chemical Co.) to 500 μg mitochondria, and incubated the suspension at 4°C with gentle agitation for 10 min. After centrifugation at 12,500 g for 10 min, we added 100 μl of a 1:1 slurry of anti-HA-protein A–Sepharose beads in solubilization buffer to a 1-ml aliquot of the supernatant. Samples were incubated at 4°C with gentle agitation at least 5 h. HA-protein A–Sepharose was prepared using IgG from 12CA5 cells (Niman et al., 1983) and an ImmunoPure™ IgG orientation kit (Pierce) according to manufacturer’s instructions. Alternatively, 20 μl of antisera against Tim23p (Emtage and Jensen, 1993) was added to the supernatant, followed by 150 μl of a 1:1 slurry of protein A–Sepharose beads (Sigma Chemical Co.). Beads were collected by centrifugation. We added 160 μl of 1× sample buffer to each supernatant and heated the supernatants at 65°C for 5 min. Pellets containing the protein A–Sepharose were washed four times with 1.0 ml solubilization buffer, and the bound proteins were eluted by two sequential extractions with 80 μl of 1× sample buffer. 500 μl of solubilization buffer was added to the pellet samples followed by heating at 65°C for 5 min. After separation by SDS-PAGE, proteins were immune blotted with antibodies to the HA epitope, or with antisera to Aac1p (a gift from M. Douglas, mt-Hsp70) (a gift from M. Cumsy, University of California, Irvine, CA), Tim44p (a gift from G. Schatz, Bio-center, Basel, Switzerland), Tim34p, Tim23p, or Tim17p (Ryan et al., 1998).

**Results**

**TIM54 Encodes a Novel 54-kD Protein that Potentially Interacts with Mmm1p**

Mmm1p is a mitochondrial outer membrane protein required for the maintenance of mitochondrial morphology (Burgess et al., 1994). When Mmm1p function is lost, such as in the temperature-sensitive mmm1 mutant, the normal elongated mitochondria collapse into large, spherical organelles. We found that the Mmm1p protein resides in the mitochondrial outer membrane, but its role in mediating mitochondrial shape is unknown. To further investigate Mmm1p function, we identified potential Mmm1p-interacting proteins using the yeast two-hybrid screen (Fields and Song, 1989; Bai and Elledge, 1996). In particular, we fused the bulk of the Mmm1 protein, residues 123–426, to the DNA-binding domain of the yeast Gal4p transcriptional activator. This plasmid construct was cotransformed into Y190 yeast cells along with a cDNA library containing random fusions of yeast genes to the Gal4 activation domain (Bai and Elledge, 1996). Y190 cells contain both the yeast HIS3 gene and the E.coli lacZ gene under the control of Gal4 (Bai and Elledge, 1996). cDNAs that encode Mmm1p-interacting proteins (and therefore bring the activation and DNA-binding domains of Gal4 together) were identified as His+ lacZ+ transfectants. From 10⁶ transformants, we isolated 10 potential Mmm1p-interactors. To determine if the interaction with Mmm1p was specific in any of the cotransformants, we tested each of the 10 plasmids carrying cDNA–Gal4 activation domain fusions for their interaction with other proteins besides Mmm1p (p53, lamin, yeast Snf1) fused to the Gal4-DNA binding domain. One plasmid, pACT-15, contained a yeast cDNA-activation domain fusion that interacted specifically with Mmm1p. DNA sequencing showed that pACT-15 contained a cDNA encoding the carboxyl-terminal 141 amino acids of a novel protein. Complete sequencing of genomic DNA revealed an open reading frame of 1,437 bp, which encodes a protein of 54.2 kD. Subsequent to our work, the yeast genome project revealed that the TIM54 gene is lo-
Tim54p Is an Essential Gene

To investigate the function of the Tim54 protein, we constructed two disruptions of the TIM54 gene (Fig. 1 A). Briefly, we first inserted the yeast LEU2 gene into the NcoI site, which interrupts the TIM54 coding sequence after amino acid 77. We also inserted URA3 into the StyI sites of TIM54 (residues 386 and 470). We then transformed the tim54::LEU2 or tim54::URA3 constructs into diploid strains 410, 411, or 534, and stable Leu− or Ura− transformants were isolated. We sporulated these diploid cells and allowed the haploid progeny to grow at 24°, 30°, or 37°C. In more than 30 total tetrads, none gave rise to more than two viable spores, even after prolonged incubation (Fig. 1 B). All viable spores were shown to carry the wild-type TIM54 gene. TIM54 is thus an essential gene. Spores inferred to be tim54::LEU2 or tim54::URA3 germinated and underwent approximately eight cell divisions, and then arrested in their growth as unbudded cells. We consistently observed that tim54 spores formed microcolonies, never containing more than 250 cells. This “delayed death” phenotype was similar to that seen in disruptions of the IM import protein, Tim22p (Holder, J., unpublished observations).

Tim54p Is Located in the Mitochondrial IM, with its Carboxyl Terminus Facing the Intermembrane Space

To elucidate the function of Tim54p, we first identified its location within the yeast cell. We constructed an epitope-tagged version of Tim54p by inserting a segment from the influenza HA protein at the carboxyl terminus of Tim54p, and then integrated this construct into the yeast genome. The HA epitope is recognized by the monoclonal antibody 12CA5 (Niman et al., 1983). Cells expressing Tim54–HA contained a protein of 58 kD that reacted with the 12CA5 antibodies. The size of this protein was consistent with the addition of the 4-kD HA epitope to the 54-kD Tim54 protein. We also found that the Tim54–HA fusion protein was functional. Cells which only express Tim54–HA, and not wild-type Tim54p, grew on all media tested at 24°, 30°, or 37°C.

Immunofluorescence studies indicate that Tim54p is a mitochondrial protein (Fig. 2 A). Yeast cells containing plasmid pOK27, which expresses the Tim54–HA fusion protein, were fixed, permeabilized, and then incubated with antibodies to the HA-epitope and the mitochondrial ATPase β subunit (F1β) protein (Takeda et al., 1985). When immune complexes were visualized using fluorescence microscopy, we found that the Tim54–HA protein colocalized with the mitochondrial F1β protein. We also found in cell fractionation experiments that Tim54p was a mitochondrial protein (Fig. 2 B). Cells expressing the Tim54–HA fusion protein were homogenized, and separated into a mitochondrial fraction and a postmitochondrial supernatant. We found that Tim54p cofractionated with F1β, whereas little or no Tim54p was found in the supernatant along with the cytosolic hexokinase protein.

Tim54p could not be extracted from mitochondrial membranes following treatment with 1.5 M sodium chloride, 0.1 M sodium carbonate (Fig. 2 C), or 7 M urea (not shown), and is therefore an integral membrane protein. Hydrophathy analysis (Kyte and Doolittle, 1982) suggests that Tim54p may contain one or two potential membrane-spanning segments (residues 37–54 and 358–386 in the Tim54 protein). To determine in which of the two mitochondrial membranes Tim54p resides, we isolated mitochondria from cells expressing the Tim54–HA protein and prepared membrane vesicles by sonicatation. As shown in Fig. 2 D, when the mitochondrial outer membrane vesicles were separated from IM vesicles on sucrose gradients, Tim54p cofractionated with the mitochondrial F1β protein, and not with the outer membrane OM45 protein.
The bulk of the Tim54 protein faces the intermembrane space. Tim54p was imported into mitochondria requires an inner membrane potential. Mitochondria from strain 494 were isolated from wild-type strain D273-10B and incubated with 50 μM carbonyl cyanide m-chlorophenylhydrazone and 1 μM valinomycin prior to the import reaction. After 15 min at 30°C, import was stopped by incubation at 0°C. Mitochondria were treated with 50-μg/ml protease K for 20 min at 0°C, isolated by centrifugation, and solubilized in SDS-sample buffer. Proteins were separated on SDS-polyacrylamide gels, and the radiolabeled Tim54 protein was identified by fluorography. (G) The bulk of the Tim54 protein faces the intermembrane space. Tim54p was imported into wild-type mitochondria in the presence (+ Δϕ) or absence (− Δϕ) of inner membrane potential as described above. After import, the mitochondrial outer membrane was disrupted by osmotic shock, and the resulting mitoplasts were digested with 50 μg/ml proteinase K for 20 min. Mitoplasts were recovered by centrifugation and subjected to SDS-PAGE and fluorography. The full-length Tim54 protein (Tim54p) and a fragment of Tim54p (Tim54p*) protected from protease digestion are indicated.
in intact mitochondria. When the mitochondrial outer membrane was disrupted by osmotic shock forming mitoplasts, both the Tim23 and Tim54 proteins were removed, while the matrix-localized F1β protein was not digested. Our Tim23p antibodies recognize the amino-terminal domain of the Tim23 protein (Ryan et al., 1998), which has been shown to face the intermembrane space (Bauer et al., 1996; Bömer et al., 1997; Dekker et al., 1997; Hauke and Schatz, 1997; Emtage, J., O. Kerscher, and R.E. Jensen, manuscript submitted for publication). Since our antiserum to Tim54p was raised to the carboxy-terminal third of the Tim54 protein, we conclude that the carboxy terminus of Tim54p faces the intermembrane space. Supporting this view, we found that the bulk of the radiolabeled Tim54 protein that was imported into isolated mitochondria was located in the intermembrane space (Fig. 2 C). After import into energized mitochondria, trypsin digestion of mitoplasts yielded only a 5–6-kD fragment of Tim54p (Tim54p*). This protected fragment was not seen if the mitochondrial IM potential (Δψ) was dissipated prior to the import reaction.

Tim54p Is Required for the Insertion of Proteins into the IM, but Not for the Translocation of Proteins into the Matrix

All essential mitochondrial IM proteins to date have been shown to play a crucial role in protein import (Blom et al., 1993; Dekker et al., 1993; Emtage and Jensen, 1993; Maarse et al., 1994; Ryan et al., 1994). Because we found that Tim54p is an essential IM protein, we examined its potential role in import. To facilitate these studies, we first isolated a temperature-sensitive tim54 mutant by mutagenizing the cloned TIM54 gene. As described in Materials and Methods, we passed a plasmid carrying TIM54 through a bacterial strain defective in DNA repair (Greener and Callahan, 1994), and identified a temperature-sensitive tim54 mutant, tim54-1, using a plasmid-shuffle scheme (Sikorski and Boeke, 1991). Cells that contain the tim54-1 mutation as the sole source of TIM54 grew at normal rates at 24°C, but failed to grow at 35 or 37°C. We subsequently found that mitochondria isolated from the tim54-1 mutant grown at 24°C contained reduced levels of the altered Tim54 protein (see Fig. 6).

tim54-1 mutants are defective in the insertion of proteins into the mitochondrial inner membrane, but not in the translocation of proteins across the IM into the matrix. We isolated mitochondria from wild-type cells and the tim54-1 mutant, and examined their ability to import 35S-labeled precursor proteins in vitro. As shown in Fig. 3 A, we first examined the import of two proteins: Su9-DHFR, a fusion protein consisting of the amino-terminal mitochondrial targeting sequence of the matrix-localized N. crassa ATPase subunit 9 protein fused to mouse dihydrofolate reductase (Pfanner et al., 1987), and Aac1p, the yeast ATP/ADP carrier protein which resides in the mitochondrial IM (Lawson and Douglas, 1988). We found that Su9-DHFR was efficiently imported and processed to the mature form in mitochondria isolated from both wild-type and tim54-1 cells. This import of Su9-DHFR into tim54-1 mitochondria appeared to be complete. Sonication of wild-type and tim54-1 mitochondria after Su9-DHFR import liberated similar amounts of the mature form of Su9-DHFR into a supernatant fraction (see Fig. 4 B). In contrast to the matrix-localized Su9-DHFR protein, the import of IM-destined Aac1p protein into tim54-1 mitochondria was reduced at least fivefold relative to wild-type mitochondria. As shown in Fig. 3 C, we also found that tim54-1 mitochondria were defective in the import of another inner membrane protein, Tim23p. The import of Tim23p into the IM can be assessed by protease digestion: a 14-kD fragment of Tim23p (Tim23p*), which is resistant to protease digestion when the outer membrane is disrupted, is diagnostic of its correct insertion into the IM (Bömer et al., 1997; Dekker et al., 1997; Hauke and Schatz, 1997; Emtage et al., manuscript submitted for publication). We found that the rate of insertion of Tim23p into the IM, as measured by the appearance of Tim23p* was at least 40% reduced in tim54-1 mitochondria as compared to wild-type. The import of Tim23p*, was dependent upon inner membrane potential (Δψ): the 14-kD Tim23p fragment protected from protease digestion was not seen in either wild-type or tim54-1 imports when the inner membrane potential was dissipated by valinomycin.

We previously showed that Tim23p is essential for the import of proteins into the matrix (Emtage and Jensen, 1993). To determine if Tim23p also played a role in IM protein import, we isolated mitochondria from the temperature-sensitive tim23-1 mutant, and compared their ability to import different precursor proteins to mitochondria isolated from wild-type and tim54-1 strains. Consistent with our earlier results, we found that the matrix-localized Cox4p precursor was efficiently imported and processed to the mature form in wild-type mitochondria, but was imported at greatly reduced rates into tim23-1 mitochondria (Fig. 3 D). The tim23-1 mutation had no effect on the import of the Aac1 protein: Aac1p was imported into wild-type and tim23-1 mitochondria at virtually identical rates. We found that the imported Aac1p was correctly inserted into the inner membrane. Import of Aac1p was dependent upon the inner membrane potential, and protease treatment of mitoplasts after import yielding characteristic protected fragments of Aac1p (Aac1p* and Aac1p**; Sirrenberg et al., 1996; Dekker et al., 1997). In contrast to the inner membrane Aac1 protein, tim54-1 mitochondria imported Cox4p into the matrix at a rate similar to wild-type mitochondria (Fig. 3 D). We found that Cox4p was completely translocated across the IM into the matrix of tim54-1 mitochondria. Similar amounts of the mature form of Cox4p were protected from protease digestion after disruption of the mitochondrial outer membrane (Fig. 3 D). Our results thus suggest that there are two separate import pathways in the mitochondrial inner membrane. Tim23p mediates the translocation of precursors into the matrix, whereas Tim54p is required for the insertion of at least some proteins into the IM, including at least one inner membrane component (Tim23p) of the alternative pathway.

TIM54 and TIM22 Genetically Interact with Each Other, but Not with TIM23 or TIM17

To further explore the role of Tim54p in the import of inner membrane proteins, we looked for genetic interactions between TIM54 and other IM import components. In par-
Figure 3. Mitochondria isolated from the tim54-1 mutant are defective in the insertion of proteins into the inner membrane, but not in the import of matrix proteins. (A) Su9-DHFR and Aac1p imports: Mitochondria were isolated from tim54-1 strain 809 and wild-type strain YPH857, and incubated with the S-labeled Aac1p or Su9-DHFR proteins. After 20 min at 30°C, import was stopped by the addition of 40 μM valinomycin and incubation at 0°C. An aliquot of the mitochondria was treated with 200 μg/ml proteinase K for 20 min at 0°C (+ PK). Samples were isolated by centrifugation, and pellets were solubilized in SDS-sample buffer. Proteins were separated on SDS-polyacrylamide gels, and the radiolabeled proteins were identified by fluorography. Precursor (p) and mature (m) forms of the Su9-DHFR are indicated. 20% of the precursor added to each import reaction is also shown. (B) Su9-DHFR is imported into the matrix in tim54-1 mitochondria. Mitochondria were isolated from tim54-1 strain 809 and wild-type strain YPH857, incubated with the S-labeled Su9-DHFR protein, and treated with proteinase K as described above. Mitochondria were sonicated and then centrifuged at 200,000 g for 45 min. Equal aliquots of the pellet (p) and supernatant (s) fractions were subjected to SDS-PAGE and fluorography. The mature form of Su9-DHFR is shown. (C) Tim23p imports: S-labeled Tim23 protein was imported into either wild-type or tim54-1 mitochondrial for 3, 9, or 18 min at 30°C. Import was stopped by the addition of valinomycin, and mitochondria were treated with 200 μg/ml trypsin for 20 min at 0°C. After the addition of 1 mg/ml soybean trypsin inhibitor and centrifugation, the outer membrane of the mitochondria was disrupted by resuspending the mitochondrial pellet in 20 mM Hepes-KOH pH 7.4, and incubation for 20 min at 0°C. Proteins were digested by treatment with 100 μg/ml proteinase K for 20 min at 0°C. Samples were then isolated by centrifugation, and analyzed by SDS-PAGE and fluorography. Tim23+ indicates the 14-kD protease-protected fragment of Tim23p indicative of correct insertion into the inner membrane. In some reactions, the inner membrane potential (ΔΨ) was dissipated prior to import by the addition of 40 μM valinomycin. (D) Import of Aac1p and Cox4p into wild-type, tim23-1, and tim54-1 mitochondria: Mitochondria were isolated from wild-type strain 55, tim23-1 strain 201, and tim54-1 strain 809 and incubated with the S-labeled Aac1 or Cox4 proteins for the indicated times at 30°C. After import, mitochondria were reisolated by centrifugation, converted to mitoplasts, and treated with protease as described above.

Tim54p and Tim22p Are Part of a Protein Complex Separate from the Tim23p–Tim17p Complex

To test the possibility that the Tim54 and Tim22 proteins interact with each other, but are not part of the Tim23p–Tim17p complex, we asked whether Tim54p and Tim22p could be co-precipitated from detergent-solubilized mitochondria. We isolated mitochondria from cells expressing the Tim54–HA protein as the sole source of Tim54p, and solubilized the mitochondria in buffer con-
taining 0.5% digitonin. We then immune precipitated Tim54–HA using antibodies to the HA epitope and analyzed the pellet and supernatant fractions by immune blotting. As shown in Fig. 5 A, all of the Tim54–HA protein was found in the pellet fraction, and little or no Tim54–HA was seen in the supernatant. When we examined our fractions with antibodies to Tim22p, we found that all of the Tim22 protein coprecipitated along with Tim54–HA, indicating that all of the Tim22p in the mitochondria is associated with the Tim54 protein. Other IM proteins not involved in import, such as the ATP/ADP carrier protein Aac1p (Fig. 5 A), or the F1β and Cox4 proteins (not shown), did not precipitate with Tim54–HA. Thus the interaction of Tim54p and Tim22p was specific. Furthermore, proteins previously shown to be part of an IM complex mediating the translocation of proteins into the mitochondrial matrix did not coprecipitate with Tim54–HA: Tim23p, Tim17p, Tim44p, and mt-Hsp70 all remained in the supernatant fraction after Tim54–HA precipitation.

When we immune precipitated the Tim22 protein, we found that part, but not all of the Tim54 protein, associated with Tim22p (Fig. 5 B). We isolated mitochondria from cells expressing Tim22–HA, solubilized the mitochondria in digitonin-containing buffer, and precipitated the Tim22–HA protein using anti–HA antibodies. Immune blots showed that while all of the Tim22–HA could be precipitated, only about half of Tim54p was in the pellet and the other half was present in the supernatant. Similar to our precipitations using Tim54–HA, the Tim23, Tim17, Tim44, and mt-Hsp70 proteins were found only in the supernatant fraction after Tim22–HA precipitation.

Thus, neither Tim54p nor Tim22p interact with the Tim23p/Tim17p complex using our assay conditions. As a control, mitochondria isolated from cells expressing Tim22–HA were solubilized and immune precipitated with antiserum to the Tim23 protein (Fig. 5 C). Consistent with previous studies we found a tight association between Tim23p and Tim17p (Berthold et al., 1995; Blom et al., 1995; Ryan et al., 1998): both Tim23p and Tim17p were found in the pellicle fraction after Tim23p immune precipitation. In contrast, all of the Tim54 and Tim22 proteins were found in the supernatant fraction and thus did not associate with Tim23p.

When we precipitated Tim54–HA, all of the Tim22 protein coprecipitated (Fig. 5 A), but precipitation of Tim22–HA, on the other hand, brings down only half of the Tim54 protein (Fig. 5 B). We conclude that not all of the Tim54 protein in the inner membrane is interacting with Tim22p. We suggest that there may be two pools of Tim54p, one interacting with Tim22p, the other playing an unknown function. Supporting this possibility, we found that the Tim54–HA protein was approximately two to three times more abundant than Tim22–HA when both proteins were expressed from identical centromere-containing plasmids (Holder, J., and R. Jensen, unpublished observations).

**Mitochondria Isolated from the tim54-1 Mutant Contain Reduced Amounts of the Tim54 and Tim22 Proteins, but Normal Levels of Tim23p and Tim17p**

Mutations in one subunit of a multi-subunit complex often destabilize the complex and cause the rapid turnover of other subunits (Dowhan et al., 1985; Fang and Green, 1994; Shani et al., 1996). Consequently, if Tim54p and Tim22p are part of the same IM complex then *tim54-1* may have an effect on the steady-state level of the Tim22 protein. To test this possibility, we isolated mitochondria from wild-type cells and the *tim54-1* mutant grown at the per-
Discussion

We have identified a new mitochondrial protein, Tim54p, which is required for the insertion of proteins into the inner membrane, but not for the translocation of proteins across the IM into the matrix. Mitochondria that contained reduced amounts of both Tim54p and Tim22p, depleted in Tim54p and Tim22p, were present in similar amounts in wild-type and tim54-1 mitochondria. In contrast, the levels of the Tim54 and Tim22 proteins were at least 10-fold reduced in Tim54p–Tim22p mitochondria as compared to wild-type. The Tim54-1 mutation thus appears to destabilize the Tim54p–Tim22p complex. Consistent with this idea, we find that both Tim54p and Tim22p mediate the import of essential mitochondrial proteins by immune blotting. As shown in Fig. 6, the α-subunit of the F1-ATPase, Tim17p, and Tim23p were not part of the Tim54p–Tim22p complex. Consistent with this idea, we find that mitochondria isolated from a temperature-sensitive tim54-1 mutant have lower levels of the Tim23 and Tim17 proteins, but normal levels of Tim54p and Tim22p (Fig. 6). Similar amounts of the outer membrane protein, Tom70, and the inner membrane protein, Aac1p, were found in all three mitochondria (not shown).

The role of Tim54p appears similar to that of Tim22p, which has recently been shown to mediate the IM import of the Aac1p and PiC carrier proteins (Sirrenberg et al., 1996). However, in addition to Aac1p, Tim54p–Tim22p complex mediates the import of the Tim23 protein. In preliminary studies, we have also found that the import of the Tim22 protein is defective in tim54-1 mitochondria (Kerscher, O., and R. Jensen, manuscript in preparation). Because Tim23p and Tim22p are not obvious members of the carrier family, our results raise the possibility that Tim54p and Tim22p may be required for the insertion of many, if not all, inner membrane proteins. Supporting this idea, we find that both Tim54p and Tim22p are essential genes. The Aac1p and PiC proteins, on the other hand, are not essential for yeast cell viability, but are only required for growth on nonfermentable medium (Lawson and Douglas, 1988; Zara et al., 1991). It is therefore likely that both Tim54p and Tim22p mediate the import of essential proteins, including the Tim23p and Tim22p import components. To determine the complete repertoire of substrates for Tim54p and Tim22p, we are currently examining the import of additional inner membrane proteins, as well as proteins sorted to other mitochondrial locations.

Several support our results indicating that Tim23p is not required for the insertion of at least some inner membrane proteins. For example, Dekker et al. (1997) recently found that mitochondria isolated from a different tim23 mutant, tim23-2, were not defective in the insertion of Aac1p, Tim23p, or Tim17p into the inner membrane, but were defective in the import of several precursors into the matrix. Dekker et al. (1997) also found that chemical amounts of a matrix-destined precursor blocked the import of additional matrix proteins, but not the import of the inner membrane Tim23 and Aac1 proteins. Similarly, while we showed that Tim23p was required for the import of different matrix proteins (Emtage and Jensen, 1993), we have recently found that the tim23-1 mutation and antibodies to Tim23p do not block the import of additional matrix proteins.
block the insertion of the Aac1, Tim23, or Tim17 proteins into the IM (Emtage, J., O. Kerscher, R.E. Jensen, manuscript submitted for publication). In contrast to the above studies, Volker and Schatz (1997) have reconstituted the protein insertion machinery of the mitochondrial inner membrane, and find that Tim23p is required for the insertion of Aac1p and Tim23p into the IM. Why the reconstituted system apparently differs from studies with intact mitochondria awaits further analyses.

We find that Tim54p and Tim22p physically interact in the inner membrane, but that Tim54p and Tim22p are not part of the previously-characterized Tim23p–Tim17p complex. Tim22p can be coprecipitated from detergent-solubilized mitochondria along with Tim54p, but the Tim23 and Tim17 proteins do not associate with Tim54p. In addition, Tim44p and mt-Hsp70, two members of the Tim23p–Tim17p complex (Berthold et al., 1995; Blom et al., 1995; Bömer et al., 1997; Ryan et al., 1998), fail to interact with either Tim54p or Tim22p. This observation may not be surprising since Aac1p and Tim23p, both imported via the Tim54p-dependent pathway, do not require mt-Hsp70 function for their import (Bömer et al., 1997; Emtage, J., O. Kerscher, R.E. Jensen, manuscript submitted for publication). Our genetic studies support the possibility that Tim54p and Tim22p are part of a complex separate from Tim23p–Tim17p. Multiple copies of the TIM22 gene, but not TIM23 or TIM17, suppress the growth defect of a tim54-I strain. Moreover, TIM22 and TIM54 do not suppress the tim23-I mutant. Our observations are also strengthened by recent findings that TIM22 is one of the genes identified in a library screen for multi-copy suppressors of the tim54-I mutant (Leung, R., and R. Jensen, unpublished observations).

Our immune precipitation results suggest that there may be two populations of the Tim54 protein. When we precipitate Tim54p, all of the Tim22 protein coprecipitates. Precipitating Tim22p brings down only about half of the Tim54 protein. Furthermore, in preliminary experiments, the Tim54 protein in mitochondria appears to be present in two to threefold greater amounts than the Tim22 protein (Holder, J., and R. Jensen, unpublished observations). We speculate that there may be two Tim54p-containing complexes in the inner membrane: one containing Tim22p and the other containing unknown components. Tim54p may be similar to Tim23p, which is also found in two subcomplexes (Bömer et al., 1997). One Tim23p-containing subcomplex contains Tim17p, Tim44p, and mt-Hsp70; the other contains Tim17p and mt-Hsp70. Alternatively, the population of Tim54p that does not interact with Tim22p may represent an unassembled pool of Tim54p in the inner membrane. Thus, why Tim54p is present in more than one location awaits further studies.

If separate import pathways exist in the inner membrane, specific signals must then direct proteins to a given pathway. Virtually all matrix-localized proteins carry amino-terminal presequences, which target the protein to receptors on the mitochondrial surface. We propose that the presequence also directs the protein to the Tim23p–Tim17p machinery. For example, we found that the addition of the Cox4 presequence to the amino terminus of Tim23p, caused the mislocalization of Tim23p to the matrix (Emtage et al., manuscript submitted for publication). Because the import of Cox4-Tim23p was defective in tim23-1 mitochondria, we concluded that Cox4-Tim23p was imported via the Tim23p–Tim17p pathway, instead of Tim23p’s normal route via the Tim54p–Tim22p pathway. In similar studies, the addition of a presequence to another IM protein, the mammalian uncoupling protein, UCP, caused its mislocalization to the matrix (Liu et al., 1990). Whereas the presequence appears to target proteins to the Tim23p–Tim17p complex, it is not clear what signal directs membrane proteins to the Tim54p–Tim22p pathway. Inner membrane proteins, such as Aac1p and Tim23p, do not contain amino-terminal presequences, and their import signal has not yet been identified.

Interestingly, when Tim23p and UCP were mislocalized to the matrix by the addition of a presequence, neither protein was inserted into the membrane and both could be readily extracted by alkali (Liu et al., 1990; Emtage, J., manuscript submitted for publication). Mislocalization to the Tim23p–Tim17p pathway prevents membrane insertion, even though the Tim23 and UCP proteins contain multiple hydrophobic transmembrane segments. It is possible that the Tim23p–Tim17p machinery does not have the capacity to recognize the membrane-spanning segments in at least some proteins, and instead behaves like a passive channel (Ungermann et al., 1994; Berthold et al., 1995). Only when Tim23p and UCP are imported through their normal route, the Tim54p–Tim22p pathway, are they correctly inserted into the bilayer.

Why was Tim54p, an inner membrane import protein, identified as a potential protein–protein interactor with Mmm1p, an outer membrane protein required to maintain normal mitochondrial shape (Burgess et al., 1994)? Since the inner membrane appears to contain separate machinery for the import of IM proteins, it is possible that the outer membrane similarly has distinct import machinery for different proteins. Mmm1p may function in a specific import pathway for IM proteins (which includes the Tim54 protein), and may transiently interact with these proteins during their import. We, however, consider this possibility unlikely since we have not been able to detect an import defect in mmm1 mutants or mmm1 mitochondria (Srinivasan, M., S. Burgess, and R. Jensen, manuscript in preparation). Furthermore, IM proteins and matrix proteins use many of the same outer membrane machinery for their import (Pfanner and Neupert, 1987; Pfaller et al., 1988; Hines et al., 1990; Kiebler et al., 1990; Dietmeier et al., 1997; Emtage, J., and R. Jensen, unpublished observations). It is also possible that the Mmm1p and Tim54p interaction is indirect. For example, we have preliminary evidence that Mmm1p is located in contact sites between the inner and outer membrane (Srinivasan, M., S. Burgess and R. Jensen, manuscript in preparation). The interaction between Tim54p and Mmm1p may occur since the protein import machinery is present, at least transiently, in these contact sites (Horst et al., 1995). We have found that the carboxyl terminus of Tim54p faces the intermembrane space and is therefore potentially accessible to outer membrane proteins. Further studies, however, are needed to clarify the relationship between Tim54p and Mmm1p.

Our studies clearly show that Tim54p is required for the insertion of IM proteins, but we have not shown that Tim54p plays a direct role in this import pathway.
mitochondria isolated from the tim54-1 mutant have lowered amounts of Tim22p, it is possible that the primary function of Tim54p is to stabilize or localize the Tim22 protein in the mitochondrial inner membrane, and that Tim22p directly mediates IM protein insertion. Our observations that TIM54 is an essential gene and that Tim54p forms a stable complex with Tim22p argue that regardless of whether Tim54p's role in import is direct or indirect, the function of Tim54p is very important. It is also interesting to note that both Tim23p and Tim17p, and both proteins have been shown to play direct roles in import of matrix proteins (Kürlich et al., 1994; Ryan et al., 1994; Berthold et al., 1995; Dekker et al., 1997). Nonetheless, further experiments are needed to pinpoint the roles of both Tim54p and Tim22p in the insertion of IM proteins.

In summary, the mitochondrial inner membrane contains distinct machinery for the translocation of proteins across the bilayer (Tim23p–Tim17p), and for the insertion of proteins into the membrane (Tim54p–Tim22p). It is possible that the mechanism of the Tim23p–Tim17p machinery differs significantly from that of the Tim54p–Tim22p complex. Tim22p, however, is homologous to both Tim23p and Tim17p, with over 50% similar residues shared between the three proteins. We therefore suggest that the membrane insertion machinery and the translocation machinery in the mitochondrial IM share at least some common activities. Supporting this idea, a single translocon in the ER mediates the transport of both soluble and membrane proteins (Kehry et al., 1980; McEwan et al., 1980; Do et al., 1996; Mothes et al., 1997). It is tempting to speculate whether the two import complexes in mitochondria represent a more primitive or advanced evolutionary state compared to the ER machinery.

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References


peptide is sufficient to direct mitochondrial import of a chimeric protein.


