Targeting, Import, and Dimerization of a Mammalian Mitochondrial ATP Binding Cassette (ABC) Transporter, ABCB10 (ABC-me)*

Received for publication, May 6, 2004, and in revised form, June 16, 2004
Published, JBC Papers in Press, June 23, 2004, DOI 10.1074/jbc.M405040200

Soledom A. Graff, Sarah E. Haigh, Erica D. Corson, and Orian S. Shirihai

From the BioCurrents Research Center, Marine Biological Laboratory, Woods Hole, Massachusetts 02543 and Department of Pharmacology, Tufts University, Boston, Massachusetts 02111

ATP binding cassette (ABC) transporters are a diverse superfamily of energy-dependent membrane translocases. Although responsible for the majority of transmembrane transport in bacteria, they are relatively uncommon in eukaryotic mitochondria. Organellar trafficking and import, in addition to quaternary structure assembly, of mitochondrial ABC transporters is poorly understood and may offer explanations for the paucity of their occurrence. Here we examine these processes in ABCB10 (ABC-me), a mitochondrial inner membrane erythroid transporter involved in heme biosynthesis. We report that ABCB10 possesses an unusually long 105-amino acid mitochondrial targeting presequence (mTP). The central subdomain of the mTP (amino acids (aa) 36–70) is sufficient for mitochondrial import of enhanced green fluorescent protein. The N-terminal subdomain (aa 1–35) of the mTP, although not necessary for the trafficking of ABCB10 to mitochondria, participates in the proper import of the molecule into the inner membrane. We performed a series of amino acid mutations aimed at changing specific properties of the mTP. The mTP requires neither arginine residues nor predictable α-helices for efficient mitochondrial targeting. Disruption of its hydrophobic character by the mutation L46Q/H47Q, however, greatly diminishes its efficacy. This mutation can be rescued by cryptic downstream (aa 106–715) mitochondrial targeting signals, highlighting the redundancy of this protein’s targeting qualities. Mass spectrometry analysis of chemically cross-linked, immunoprecipitated ABCB10 indicates that ABCB10 embeds in the mitochondrial inner membrane homodimerizes and homo-oligomerizes. A deletion mutant of ABCB10 that lacks its mTP efficiently targets to the endoplasmic reticulum. Quaternary structure assembly of ABCB10 in the ER appears to be similar to that in the mitochondria.

ABC transporters comprise a large and diverse family of membrane translocases (1). Their function ranges from peptide transport to phospholipid flipping to anion channel formation. Members of the ABC transporter superfamily have been implicated in numerous human diseases (including cystic fibrosis (CFTR/ABCC7), adrenoleukodystrophy (ALDP/ABCD1), Zellweger’s syndrome (PMP70/ABCD3), progressive familial intrahepatic cholestasis (SPGP/ABCB11), and Stargardt macular dystrophy (ABCR/ABCA4)) (2). The basic structure of ABC transporters is relatively well conserved and includes a hydrophilic ATP binding cassette and a hydrophobic membrane-spanning domain (3). Whereas the large members of the family contain two of each domain, the “half-transporters” contain one of each and are predicted to dimerize (3–6). Mammalian ABC transporters are found predominantly in the plasma membrane but are also known to play essential roles in a number of organelles, including the endoplasmic reticulum, peroxisome, and the mitochondrion (2). Mammalian mitochondrial ABC transporters have recently gained attention for their role in heme biosynthesis and iron sulfur cluster synthesis (7–12). Accordingly, their involvement in the pathophysiology of acquired and inherited forms of sideroblastic anemias has been suggested and, in the case of ABCB7, already demonstrated. Only eight mitochondrial ABC transporters have been described to date (7). This is remarkable, given that in bacteria, ABC proteins are the most diverse family of transporters, numbering well over 50 in Escherichia coli. Since mitochondrial ABC transporters are synthesized in the cytosol, they require proper trafficking, import, and assembly to achieve functional activity in their target organelle.

Two dominant mitochondrial import pathways have been described, each involving specialized import machinery and distinct signals (13). One pathway recognizes a cleavable, N-terminal mitochondrial targeting presequence (mTP) and is generally utilized by hydrophilic proteins. The second pathway, employed by the membrane metabolite carrier class of proteins, is used by more hydrophobic proteins bearing internal targeting signals. mTPs, although they lack a consensus primary sequence, share a number of common characteristics. They are rich in arginine residues, feature a net positive charge, and possess amphiphilic α-helices (14–17). Additionally, some mTPs include a short hydrophobic motif that is recognized by Tom20, a component of the mitochondrial import machinery on the outer membrane (18, 19). mTPs lead their preproteins into the matrix in a linear configuration, where they are cleaved off by the mitochondrial processing peptidase, allowing for final assembly of the mature protein in one of the mitochondrion’s compartments (20). Proteins of the metabolite carrier class do not possess a cleavable presequence (21, 22). Rather, they bear three internal targeting modules that cooperatively bind to Tom70. These proteins are transported in a loop configuration directly into the inner membrane, where they assume their functional structure (23).

ABC half-transporters include a ~200-amino acid hydro-
Mitochondrial Targeting, Import, and Dimerization of ABCB10

Experimental Procedures

Cell Culture and Transfection—HEK 293T and COS cells were cultured in low glucose DMEM without phenol red (Invitrogen) supplemented with 10% standard fetal bovine serum (HyClone, Logan, UT), 2 mM L-glutamine (Invitrogen), and antibiotics. Transient transfections of HEK 293T cells and COS cells were performed using FuGene 6 (Roche Applied Science) during quenching. Immunoprecipitation was performed using Protein G-agarose beads (Roche Applied Science) and anti-V5 (1:500 dilution) (Invitrogen) and anti-e-Myc (1:500 dilution) (Invitrogen) antibodies as described (25).

One-dimensional SDS-PAGE, Silver Staining, and Western Blotting—Mitochondrial samples were taken up in sample buffer (62.5 mM Tris, 10% glycerol, 2% SDS, 5% β-mercaptoethanol, bromphenol blue), loaded on a 4–15% gradient Tris-HCl gel (Bio-Rad), and electrophoresed at 160 V in running buffer using a Bio-Rad MiniProtein 3 cell. Proteins to be identified by MS/MS analysis were revealed by silver staining using the Silver Stain Plus kit (Bio-Rad).

For detection by Western blotting, proteins were transferred onto nitrocellulose or polyvinylidene membranes. Transblot SD rapid transfer cell in Towbin buffer at 140 mA for 1 h at room temperature. Primary antibodies were anti-EGFP (1:2500 dilution) (Clontech) and anti-V5-horseradish peroxidase (5.0000 dilution) (Invitrogen). Protein was revealed using a Super Signal chemiluminescence horseradish peroxidase detection system as directed by the manufacturer (Pierce) and exposed to Kodak BioMax MR film (Eastman Kodak Co.).

Mass Spectrometry and N-terminal Sequencing—Protein identification of SDS-PAGE silver-stained bands was done by MS/MS analysis of tryptic peptides using the MS-fit program as previously described (26). N-terminal protein sequencing was performed by automated Edman degradation on an Applied Biosystems model 491 Procisor sequencer. Samples separated by SDS-PAGE were transferred onto polyvinylidene difluoride membrane and stained with Coomassie Blue. Model 610A version 2.1 software was employed for data acquisition and processing.

Proteasine K Digestion of Extramitochondrial Proteins—Extramitochondrial protein digestion was carried out to verify complete mitochondrial insertion of each mTP and its passenger protein. It was performed on all constructs that targeted partially or entirely to mitochondria. Those constructs that targeted to mitochondria but were not fully imported are mentioned under “Results.” To accomplish extramitochondrial protein digestion, gradient-purified mitochondria were incubated with 100 μg/ml proteasine K (Sigma) on ice for 30 min. Proteasine K was added, and the mitochondria were pelleted by centrifugation at 17,500 × g for 10 min at 4 °C. Mitochondria were resuspended in buffer C (0.67% sorbitol (Fisher), 20 mM HEPES, pH 7.4) supplemented with 2 Complete protease inhibitor mixture (Roche Applied Science), 2 mM phenylmethylsulfonyl fluoride, and 5.0000 (v/v) trichloroacetic acid (Sigma), heated to 70 °C for 5 min, and then incubated on ice for an additional 5 min. Mitochondria were resolated by centrifugation at 12,000 × g for 10 min and immediately resuspended in sample buffer for separation by one-dimensional SDS-PAGE. 2× Tris, pH 9, was added dropwise to neutralize the solution.

White Cell Protein Analysis—Whole cell protein analysis was performed to control for post-translational stability of transfected constructs. The procedure was carried out on all constructs that did not target to mitochondria. Those constructs that were cleaved prior to mitochondrial targeting, thus reducing targeting efficiency, are men-

phospho ATP binding cassette and a ~400-amino acid hydrophobic transmembrane domain. Therefore, these proteins might present a challenge to mitochondrial import machinery. This study marks the first comprehensive analysis of the mitochondrial targeting, import, and assembly properties of an ABC transporter. We have previously reported on the murine mitochondrial transporter, ABCB10 (ABC-m). ABCB10 is induced during erythroid differentiation, and this is involved in heme biosynthesis (8). We report here that the ABCB10 preprotein includes an unusually long mTP of 105 residues that provides mitochondrial targeting redundancy and flexibility. Point mutation experiments reveal that ABCB10 additionally possesses downstream (aa 106–715) targeting signals capable of assisting a damaged mTP. To differentiate between protein trafficking and protein import, we combined confocal microscopy of living cells and biochemical analysis of isolated organelles. We show that removing part of the mTP of ABCB10 results in efficient trafficking but compromised import into the mitochondria. Quaternary structure analysis of ABCB10 indicates that it homo-oligomerizes in the mitochondrial inner membrane and does not appear to assemble with other proteins. ABCB10 that lacks its mTP is targeted not to mitochondria but to the ER, where it also oligomerizes.

EXPERIMENTAL PROCEDURES

Cell Culture and Transfection—HEK 293T and COS cells were cultured in low glucose DMEM without phenol red (Invitrogen) supplemented with 10% standard fetal bovine serum (HyClone, Logan, UT), 2 mM L-glutamine (Invitrogen), and antibiotics. Transient transfections of HEK 293T cells and COS cells were performed using FuGene 6 (Roche Applied Science) during quenching. Immunoprecipitation was performed using Protein G-agarose beads (Roche Applied Science) and anti-V5 (1:500 dilution) (Invitrogen) and anti-e-Myc (1:500 dilution) (Invitrogen) antibodies as described (25).

One-dimensional SDS-PAGE, Silver Staining, and Western Blotting—Mitochondrial samples were taken up in sample buffer (62.5 mM Tris, 10% glycerol, 2% SDS, 5% β-mercaptoethanol, bromphenol blue), loaded on a 4–15% gradient Tris-HCl gel (Bio-Rad), and electrophoresed at 160 V in running buffer using a Bio-Rad MiniProtein 3 cell. Proteins to be identified by MS/MS analysis were revealed by silver staining using the Silver Stain Plus kit (Bio-Rad).

For detection by Western blotting, proteins were transferred onto nitrocellulose or polyvinylidene membranes. Transblot SD rapid transfer cell in Towbin buffer at 140 mA for 1 h at room temperature. Primary antibodies were anti-EGFP (1:2500 dilution) (Clontech) and anti-V5-horseradish peroxidase (5.0000 dilution) (Invitrogen). Protein was revealed using a Super Signal chemiluminescence horseradish peroxidase detection system as directed by the manufacturer (Pierce) and exposed to Kodak BioMax MR film (Eastman Kodak Co.).

Mass Spectrometry and N-terminal Sequencing—Protein identification of SDS-PAGE silver-stained bands was done by MS/MS analysis of tryptic peptides using the MS-fit program as previously described (26). N-terminal protein sequencing was performed by automated Edman degradation on an Applied Biosystems model 491 Procisor sequencer. Samples separated by SDS-PAGE were transferred onto polyvinylidene difluoride membrane and stained with Coomassie Blue. Model 610A version 2.1 software was employed for data acquisition and processing.

Proteasine K Digestion of Extramitochondrial Proteins—Extramitochondrial protein digestion was carried out to verify complete mitochondrial insertion of each mTP and its passenger protein. It was performed on all constructs that targeted partially or entirely to mitochondria. Those constructs that targeted to mitochondria but were not fully imported are mentioned under “Results.” To accomplish extramitochondrial protein digestion, gradient-purified mitochondria were incubated with 100 μg/ml proteasine K (Sigma) on ice for 30 min. Proteasine K was added, and the mitochondria were pelleted by centrifugation at 17,500 × g for 10 min at 4 °C. Mitochondria were resuspended in buffer C (0.67% sorbitol (Fisher), 20 mM HEPES, pH 7.4) supplemented with 2 Complete protease inhibitor mixture (Roche Applied Science), 2 mM phenylmethylsulfonyl fluoride, and 5.0000 (v/v) trichloroacetic acid (Sigma), heated to 70 °C for 5 min, and then incubated on ice for an additional 5 min. Mitochondria were resolated by centrifugation at 12,000 × g for 10 min and immediately resuspended in sample buffer for separation by one-dimensional SDS-PAGE. 2× Tris, pH 9, was added dropwise to neutralize the solution.

White Cell Protein Analysis—Whole cell protein analysis was performed to control for post-translational stability of transfected constructs. The procedure was carried out on all constructs that did not target to mitochondria. Those constructs that were cleaved prior to mitochondrial targeting, thus reducing targeting efficiency, are men-
Mitochondrial Targeting, Import, and Dimerization of ABCB10

Identification of ABCB10 Presequence Cleavage Site—ABCB10 fused to V5 at its C-terminal was expressed in HEK 293T cells, immunoprecipitated, and separated by SDS-PAGE. Silver staining revealed a band at \( \sim 65 \text{ kDa} \) (Fig. 1A). This band was subjected to tryptic fragment analysis by tandem mass spectrometry and was identified as ABCB10, with the fragments obtained indicating that a larger than expected portion of the N terminus could be missing (data not shown). N-terminal sequencing performed on the band identified the first 17 amino acids as LGNDSQRPAATGRSEV, which correspond to the N terminus of ABCB10, suggesting that an alternative subcellular distribution of ABCB10 or isoforms of ABCB10 may exist. Subcellular localization of \( \Delta 105-\text{ABCB10} \) was additionally examined in COS cells, yielding identical ER preference (data not shown).

The mTP of ABCB10 is Necessary for Mitochondrial Import of ABCB10 and Sufficient for Import of EGFP—To explore the mitochondrial targeting and import properties of ABCB10 and its mTP, EGFP was fused at their C termini, and the proteins were expressed in HEK 293T cells. Their colocalization with mitochondria stained with TMRE was analyzed by confocal microscopy (Fig. 2). Metamorph image analysis software generated per-pixel scatter plots of red and green signal intensities. The MMCC is a measure of the scatter plot’s tendency for positive linear correlation. Thus, a value of +1 represents perfect mitochondrial colocalization, 0 represents no mitochondrial preference, and −1 represents perfect mitochondrial exclusion. Since HEK 293T fibroblasts grow in adherent colonies, cells were selected from at least 12 colonies to minimize the impact of clonal variation. EGFP has been employed previously for similar subcellular localization studies (18, 29). EGFP alone is cytosolically localized and shows no organellar preference (Fig. 2E). EGFP carries consecutive negative charges at residues 6E and 7E of the EGFP molecule. Concentrated negative charge is highly unfavorable to mitochondrial targeting (16). Therefore, it is believed that C-terminal EGFP fusion does not enhance mitochondrial targeting of a molecule.

As expected, ABCB10-EGFP co-localizes with mitochondria, giving an MMCC of 0.81 ± 0.02 (Fig. 2A). The mTP of ABCB10 alone also targets EGFP to the mitochondria, demonstrating its sufficiency as an mTP. The MMCC for aa 1–105-EGFP is 0.81 ± 0.02 (Fig. 2B). To ensure that these proteins were fully imported into mitochondria, in addition to being trafficked to the mitochondrial surface, biochemical analysis of their position was performed. Density gradient-purified mitochondria were isolated from HEK 293T expressing the protein of interest. Mitochondria were then subjected to proteinase K digestion of extramitochondrial proteins, and the results were analyzed by Western blotting. Western blots of aa 1–105-EGFP pre- and postdigestion were identical, indicating that complete import was accomplished (data not shown). This validation procedure was carried out on all subsequently examined constructs that targeted to mitochondria (see “Experimental Procedures”).

Removal of the mTP from ABCB10-EGFP drastically reduces mitochondrial targeting, resulting in an MMCC of \( \Delta 105-\text{ABCB10}-\text{EGFP} \) of 0.33 ± 0.03 (Fig. 2C). For experimental control, mitochondrial targeting was additionally calculated for EGFP C-terminally fused to a standard, commercially available mTP, subunit VIII of human cytochrome c oxidase (Clonech), and EGFP alone (Fig. 2, D and E). The negative MMCC for EGFP alone indicates that it avoids mitochondria. To ensure that the failure of \( \Delta 105-\text{ABCB10}-\text{EGFP} \) to target to mitochondria was not due to post-translational cleavage, cell lysate was prepared and analyzed by Western blotting with anti-EGFP. All \( \Delta 105-\text{ABCB10}-\text{EGFP} \) that was detected appeared at an apparent molecular weight consistent with the predicted value (data not shown). This procedure was carried out on all subsequently examined constructs that failed to efficiently target to mitochondria (see “Experimental Procedures”).

Removal of the mTP Results in the Mis-targeting of ABCB10 to the ER—\( \Delta 105-\text{ABCB10}-\text{EGFP} \) does not appear ubiquitously in the cytoplasm but rather is identified as a meshlike structure characteristic of the ER membrane (Fig. 2C). Cells expressing \( \Delta 105-\text{ABCB10}-\text{EGFP} \) were co-stained with the ER-specific dye ER-Tracker blue-white DPX (E-12353) and subjected to ER colocalization quantification according to the same technique as mitochondrial colocalization (Fig. 2F). \( \Delta 105-\text{ABCB10}-\text{EGFP} \) is targeted into the ER with high efficiency, suggesting that an alternative subcellular distribution of ABCB10 or isoforms of ABCB10 may exist. Subcellular localization of \( \Delta 105-\text{ABCB10} \) was additionally examined in COS cells, yielding identical ER preference (data not shown).

Fragmentation of the mTP of ABCB10 and Identification of a Critical Targeting Region—The primary amino acid sequence

\[
\text{NH}_2-\text{MRAFRAALLLFRPRPAVRAWAFVSEKVLAS}^{105}\text{COO}^{-}
\]

FIG. 1. A, silver stain of immunoprecipitated ABCB10 isolated from mitochondria of HEK 293T cells transfected with ABCB10-V5. Lane 1, negative control of immunoprecipitation (IP) using anti-c-Myc antibody. Lane 2, immunoprecipitation using anti-V5 antibody. A band (verified as ABCB10-V5 by mass spectrometry) appears at \( \sim 65 \text{ kDa} \) (indicated by an arrow). An artifact of immunoprecipitation appears in both lanes at \( \sim 55 \text{ kDa} \). ABCB10-V5 was excised from the gel and subjected to N-terminal sequencing. B, the first 123 amino acids of ABCB10 and the position of the protein in the mitochondria. N-terminal sequencing of mature protein identified the first 17 amino acids as LGNDSQRPAATGRSEV, corresponding to Leu\(^{106}\)-Val\(^{123}\), indicating that the mTP is composed of aa 1–105. Basic residues are in boldface type, and acidic residues are in gray. Predicted α-helices are underlined. Note that the hydrophobic transmembrane domain (TMD) and hydrophilic nucleotide binding domain (NBD) each represent large fractions of the imported protein.
The distribution of both positive charge and hydrophobic moment (a measure of amphipathicity) along the mTP of ABCB10 follows a pattern that delineates three distinct subdomains (Fig. 3, A and B). Similarly, the three putative α-helices are...
Targeting of the subdomains of the mTP of ABCB10 and role of aa 1–35 in mitochondrial import of ABCB10. A, plot of charge distribution along the mTP of ABCB10. A sliding window of 12 aa was used to calculate average charge. B, hydrophobic moment plot of the mTP calculated according to Eisenberg’s algorithm using a β-angle of 100° and a consensus hydrophobicity scale, a measure of amphiphilicity. Note the presence of three regions of the mTP, each containing a maximum in positive charge, a peak in hydrophobic moment, and a single predicted α-helix. C–E, the central subdomain is as effective at targeting EGFP to mitochondria as the entire mTP. aa 1–35 and 71–105 demonstrate reduced mitochondrial targeting compared with the entire mTP. An asterisk indicates significant difference. F and G, removal of aa 1–35 from the mTP and...
dispersed one per subdomain (Fig. 1B). To investigate underlying independent functionality of the three subdomains, fragmentation and mutation experiments were performed. The mTP was divided into three 35-aa fragments: aa 1–35, 36–70, and 71–105. The two negatively charged residues of the mTP of ABCB10 are located in positions 35 and 105, the C-terminal residues on the first and third fragments, respectively. Their proline to EGFP minimizes their effect on the targeting properties of the mTP. Since the average length of all confirmed mTPs is 34 aa, the targeting efficiency of the fragments is not compromised by mTP length alone (28). Thus, examination of the targeting of the mTP fragments offers insight into the inherent properties in each region.

The central 35 aa of the mTP of ABCB10 is more effective at targeting EGFP to mitochondria than the flanking subdomains (Fig. 3). aa 1–35 and 71–105 only partially target EGFP to the mitochondria (MMCC of 0.50 ± 0.04 and 0.50 ± 0.05, respectively). aa 36–70 targets EGFP to the mitochondria as efficiently as the entire mTP of ABCB10 (Figs. 2B and 3D). aa 1–70, which is expected to target to mitochondria well, has a low MMCC of 0.27 ± 0.05 (data not shown). However, Western blot analysis of cell lysate found that the protein is unstable (Fig. 3H). Ambiguous cleavage events free EGFP from the rest of the protein, thereby causing EGFP to localize to the cytoplasm. A similar process occurred when aa 36–70 alone was N-terminally fused to Δ105-ABC10-EGFP to determine the capacity of the isolated central subdomain of the mTP to target ABCB10 to the mitochondria. The resulting protein was unstable in both HEK 293T cells and COS cells (data not shown).

Functional Analysis of aa 1–35: Evidence for a Role in the Translocation of ABCB10 across Mitochondrial Membranes—The N-terminal subdomain of the mTP, aa 1–35, is not essential for the trafficking of ABCB10 to mitochondria. Both aa 36–105-EGFP and Δ35-ABC10-EGFP are brought to the mitochondrial surface (Fig. 3, F and G). Western blot analysis suggested, however, that Δ35-ABC10-V5 is not fully inserted into mitochondria despite reaching their outer membranes. Western blot analysis of density gradient-purified mitochondria was carried out on cells expressing either ABCB10-V5 or Δ35-ABC10-V5. Wild-type ABCB10-V5 appears as a major band corresponding to the molecular weight of the mature protein from which the mTP has been cleaved (Fig. 3J). When SDS-PAGE is performed within 24 h post-transfection, a faint, higher molecular weight band corresponding to the approximate weight of the immature, uncleaved protein is present (as seen in Fig. 1A). Δ35-ABC10-V5 appears as two bands of nearly equal strength, indicating that a substantial fraction of the protein is left uncleaved. Since mitochondrial presequences are known to be cleaved by enzymes located in the mitochondrial matrix, significant amounts of Δ35-ABC10-V5 probably fail to translate across both mitochondrial membranes. Extramitochondrial protein digestion by protease K performed on isolated mitochondria results in the complete disappearance of the high molecular weight Δ35-ABC10-V5 band and leaves the low molecular weight Δ35-ABC10-V5 band unchanged, supporting the role of aa 1–35 in the mitochondrial import of ABCB10 into the inner membrane. Unlike Δ35-ABC10-V5, aa 36–105-EGFP does not significantly degrade in the presence of protease K. This suggests that full mitochondrial insertion and proper prescission cleavage takes place. Therefore, the downstream domain of ABCB10 (aa 106–715) appears to present a challenge to mitochondrial import machinery not found in EGFP alone. Although not essential for efficient mitochondrial targeting, aa 1–35 is required for proper mitochondrial insertion.

Identification of Elements of the mTP Essential for Mitochondrial Targeting—A series of specific amino acid mutation experiments were performed to elucidate the features of aa 36–70 that are necessary for its mitochondrial targeting. Mutations directed at removing arginine residues, disrupting the predicted α-helix, and replacing hydrophobic residues were carried out. aa 36–70 contains two arginine residues at positions 41 and 55, and these were mutated to alanine individually and in tandem. Alanine possesses similar helix formation properties to arginine, thereby limiting the effect of arginine replacement to removing its positive charge. All three mutants, R41A, R55A, and R41A/R55A, targeted EGFP to mitochondria nearly as efficiently as wild-type: MMCC values of 0.71 ± 0.04, 0.76 ± 0.03, and 0.74 ± 0.03, respectively (Table I). This was surprising in light of the presumed importance of arginine in mTPs. aa 36–70 effectively targets EGFP to the mitochondria with no arginine residues and a single positively charged residue, Lys46.

To test the importance of the region’s α-helix, the secondary structure of aa 36–70 was disrupted. The predicted α-helix of aa 36–70 falls at aa 37–47. Mutating either Ala42 to Pro or Arg41 to Pro is expected to destroy this conformation. Both mutations were carried out, and neither significantly reduced the mitochondrial targeting efficiency (Table I). The mTP of ABCB10 therefore does not require a predictable α-helix for normal mitochondrial targeting.

The aa 36–70 fragment possesses considerable hydrophobic character. We removed one hydrophobic residue pair with the double mutation I46Q/I47Q. Glutamine residues were substituted for leucine and isoleucine to maintain the helix formation and neutral charge properties of wild type. This mutation resulted in drastic reduction in mitochondrial targeting of aa 36–70: MMCC = 0.30 ± 0.06 (Fig. 4A). The mitochondrial targeting efficiency of aa 36–70 was reduced significantly further with the addition of the R41A mutation (MMCC = 0.15 ± 0.02), indicating that a threshold was breached that prevents the ability of mTP to overcome the mutation of Arg41 (Fig. 4B).

The Addition of aa 1–35 and 71–105 to Hydrophobically Mutated aa 36–70 Fails to Fully Rescue Mitochondrial Targeting

<table>
<thead>
<tr>
<th>Mutation</th>
<th>MMCC</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>0.77 ± 0.02</td>
<td>12</td>
</tr>
<tr>
<td>aa 36–70-R41A</td>
<td>0.71 ± 0.04</td>
<td>12</td>
</tr>
<tr>
<td>aa 36–70-R55A</td>
<td>0.76 ± 0.03</td>
<td>12</td>
</tr>
<tr>
<td>aa 36–70-R41A-R55A</td>
<td>0.74 ± 0.03</td>
<td>28</td>
</tr>
<tr>
<td>aa 36–70-R41P</td>
<td>0.73 ± 0.02</td>
<td>19</td>
</tr>
<tr>
<td>aa 36–70-A2P</td>
<td>0.75 ± 0.03</td>
<td>16</td>
</tr>
</tbody>
</table>
ing—The flanking thirds of the mTP, aa 1–35 and 71–105, only partially rescue the effect of the L46Q/I47Q mutations on aa 36–70, demonstrating that the Leu46-Ile47 hydrophobic pair of the mTP of ABCB10 is essential for its mitochondrial targeting (Fig. 4C). Since additional mutation of Arg41 to Ala in the hydrophobic mutant results in no further decrease in mitochondrial targeting, the importance of the R41A mutation observed in the aa 36–70 hydrophobic mutant is negated by the presence of the flanking regions (Fig. 4, B and D).

**Downstream (aa 106–715) Elements of ABCB10 Rescue Impaired Targeting of Mutated mTP**—The internal domain of ABCB10 (aa 106–715) completely restores mitochondrial targeting of the mutated mTP. The R41A/L46Q/I47Q mutations were carried out in full-length ABCB10 (715 aa) fused to EGFP (Fig. 4E). The protein targeted to mitochondria as efficiently as wild-type and at a significantly greater level than the mutated 105-aa mTP alone. Thus, internal regions of ABCB10, in addition to the mTP, possess mitochondrial targeting information.

**ABCB10 Oligomerizes in the Inner Membrane**—The membrane-permeable cross-linker DFDNB was used to cross-link endogenous ABCB10 in mitochondria isolated from differentiating erythroid cells and V5 epitope-tagged ABCB10 in HEK 293T cells. When DFDNB was added, a band at ~130 kDa, the approximate mass of an ABCB10 homodimer, appeared and strengthened in intensity with increasing amounts of DFDNB over the range of 0.1–10 mg/mg of protein (Fig. 5A). The band at ~65 kDa weakened with more DFDNB. The ~130-kDa band similarly appeared when the mitochondrial preparation expressing ABCB10-V5 was run on SDS-PAGE in the absence of β-mercaptoethanol (Fig. 5B). A higher molecular mass band (>200 kDa) also appeared with cross-linking, its intensity similarly related to the concentration of DFDNB.

**ABCB10 Forms Homodimers and Homo-oligomers**—To determine whether the ABCB10 complexes include ABCB10 ho-
Mitochondrial ABC transporters play a key role in iron metabolism and heme biosynthesis. Consisting of hydrophobic as well as hydrophilic domains, ABC transporters may present a challenge to mitochondrial import machinery. Four mammalian mitochondrial transporters besides ABCB10 have been discovered: ABC7, M-ABC1, M-ABC2, and MTABC3. ABC7, M-ABC2, and M-ABC1 have been shown to possess mTPs (12, 30, 31). Computational analysis of MTABC3 predicts that it does not possess an mTP and therefore is likely to enter the mitochondrion via the Tom70 pathway. All mitochondrial ABC transporters described to date are of the half-transporter type and therefore predicted to dimerize; however, their dimerization partners have not yet been identified. We report here the first comprehensive analysis and characterization of the import of a mitochondrial ABC transporter, ABCB10, and its assembly into a homodimer in the inner membrane. We applied combined microscopy and cellular fractionation to study the function of different components of the mTP in trafficking, import, and dimerization of ABCB10. Taking this approach, we show that whereas the middle segment of the mTP contains hydrophobic residues essential for trafficking of the protein to the organelle, the N-terminal subdomain is necessary for proper insertion of the protein into the organelle. Further, we show that ABCB10 homodimerizes in the inner membrane and can similarly assemble into dimers in the ER when expressed without the mTP.

We determined that the mTP of ABCB10 is sufficient for mitochondrial import of EGFP and that the middle fragment (aa 36–70) contains the minimal information necessary to direct these processes. Net positive charge, derived particularly from arginine residues, and amphiphilic α-helices are thought to play important roles in the mitochondrial targeting properties of mTPs (14, 17). Removal of these elements by mutation in aa 36–70-EGFP results in wild-type phenotype. We therefore speculate that certain mTPs possess flexible and redundant mitochondrial targeting properties that allow...
them to bypass the use of conventional targeting mechanisms. Further mutation experiments identified two key hydrophobic residues that are essential for efficient mitochondrial targeting. A hydrophobic pentapeptide motif has been identified as essential for recognition by the TOM complex both in vitro and in vivo (18, 19). This motif occurs three times in aa 36–70, one of which includes Leu\(^{46}\)Ile\(^{47}\). Mutating these residues to glutamine destroyed most of the mitochondrial targeting of aa 36–70 and left the peptide susceptible to additional reduction in targeting efficiency by arginine replacement. This indicates that, like plant \(P_1\)ATP synthase (18), ABCB10 associates with Tom20 via hydrophobic interactions, as described by Abe et al. (32). Fusing the flanking regions of the mTP to aa 36–70 failed to completely rescue the effect rendered by the hydrophobic mutations yet did attenuate the additional insult imparted by the R41A mutation. This suggests that the distribution of positive charge in the mTP of ABCB10 has margin for error, supporting its role as a provider of thermodynamic energy for translocation and not for precise recognition and binding with import machinery (33). On the contrary, the hydrophobic residues responsible for Tom20 interaction must meet stringent accessibility and location requirements, since all nine additional hydrophobic pentapeptide motifs of the aforementioned type introduced by the flanking regions failed to completely rescue the L46Q/I47Q mutation. Fusion of the internal region of ABCB10, aa 106–715, to the hydrophobically mutated 105-aa mTP fully rescued mitochondrial targeting. This implies that ABCB10 contains internal signals that provide targeting information in addition to the mTP. This has been observed in the inner membrane protein cytochrome c1 but is a relatively rare feature of nuclear encoded mitochondrial proteins (34). Although the hydrophobically mutated mTP and the internal domain independently are incapable of targeting EGFP to the mitochondria, their union creates an efficiently targeted protein (without the introduction of a necessary targeting element at their junction). A similar phenomenon was demonstrated by Galanis et al. (35), who by doubling a mitochondrial presequence increased its targeting potency. Considering the challenge to mitochondrial import that ABCB10 presents, the protein is an ideal candidate for possessing redundant import features.

To determine whether aa 1–35 improves mitochondrial targeting by providing additive information to the targeting properties of aa 36–70, aa 1–70-EGFP was constructed; however, the resulting protein was unstable. This demonstrates the potential instability of EGFP-tagged proteins and the necessity to verify that unanticipated cleavage does not occur. Instead, we assessed the effect that the absence of aa 1–35 had on targeting efficiency by constructing \(\Delta35\)-ABCB10-EGFP/V5. Fluorescence microscopy found that the absence of aa 1–35 does not significantly reduce mitochondrial targeting of ABCB10-EGFP. However, protease K digestion of extramitochondrial proteins found that translocation of \(\Delta35\)-ABCB10-V5 across the mitochondrial membrane is reduced, indicating that the processes of mitochondrial targeting and import can be separated. aa 1–35, although necessary for proper mitochondrial import of ABCB10-V5, is not required for that of EGFP, a purely hydrophilic protein. Thus, the distinction between mitochondrial trafficking and import may be more easily observed in proteins that challenge the trafficking/import machinery. The mechanism by which aa 1–35 directs the mitochondrial import of ABCB10 is potentially related to a number of characteristics of ABCB10 and the import machinery. Since mitochondrial proteins are predicted to enter the mitochondria either through a linear or loop conformation, depending on the import pathway, aa 1–35 may affect the folding of ABCB10 and thereby influence the pathway selected (13, 36). Since each mitochondrial membrane possesses its unique import machinery, aa 1–35 may be specifically required for import across the inner membrane but not the outer. Mislocalization of imported proteins among the mitochondrial compartments has been demonstrated in cytochrome c oxidase subunit Va (37). Alternatively, aa 1–35 might simply add import elements needed to overcome a threshold, similar to the additive targeting properties observed in ABCB10. Further investigation will help to tease apart mitochondrial trafficking and import mechanisms and explain the obstacles that ABCB10 presents to each.

ABCB10 lacking its 105-aa mTP efficiently localizes to the ER, raising the possibility that an isoform of ABCB10 is present in the ER under physiological conditions or that the start codon is mutated. A number of proteins have been described that employ alternate splicing or post-translational modification to direct ER or mitochondria-specific targeting as required. In such cases, the C-terminal end of ABCB10 is potential instability of EGFP-tagged proteins and the necessity to provide targeting information in addition to the mTP. This has been observed in the inner membrane protein cytochrome c1 but is a relatively rare feature of nuclear encoded mitochondrial proteins (34). Although the hydrophobically mutated mTP and the internal domain independently are incapable of targeting EGFP to the mitochondria, their union creates an efficiently targeted protein (without the introduction of a necessary targeting element at their junction). A similar phenomenon was demonstrated by Galanis et al. (35), who by doubling a mitochondrial presequence increased its targeting potency. Considering the challenge to mitochondrial import that ABCB10 presents, the protein is an ideal candidate for possessing redundant import features.

To determine whether aa 1–35 improves mitochondrial targeting by providing additive information to the targeting properties of aa 36–70, aa 1–70-EGFP was constructed; however, the resulting protein was unstable. This demonstrates the potential instability of EGFP-tagged proteins and the necessity to verify that unanticipated cleavage does not occur. Instead, we assessed the effect that the absence of aa 1–35 had on targeting efficiency by constructing \(\Delta35\)-ABCB10-EGFP/V5. Fluorescence microscopy found that the absence of aa 1–35 does not significantly reduce mitochondrial targeting of ABCB10-EGFP. However, protease K digestion of extramitochondrial proteins found that translocation of \(\Delta35\)-ABCB10-V5 across the mitochondrial membrane is reduced, indicating that the processes of mitochondrial targeting and import can be separated. aa 1–35, although necessary for proper mitochondrial import of ABCB10-V5, is not required for that of EGFP, a purely hydrophilic protein. Thus, the distinction between mitochondrial trafficking and import may be more easily observed in proteins that challenge the trafficking/import machinery. The mechanism by which aa 1–35 directs the mitochondrial import of ABCB10 is potentially related to a number of characteristics of ABCB10 and the import machinery. Since mitochondrial proteins are predicted to enter the mitochondria either through a linear or loop conformation, depending on the import pathway, aa 1–35 may affect the folding of ABCB10 and thereby influence the pathway selected (13, 36). Since each mitochondrial membrane possesses its unique import machinery, aa 1–35 may be specifically required for import across the inner membrane but not the outer. Mislocalization of imported proteins among the mitochondrial compartments has been demonstrated in cytochrome c oxidase subunit Va (37). Alternatively, aa 1–35 might simply add import elements needed to overcome a threshold, similar to the additive targeting properties observed in ABCB10. Further investigation will help to tease apart mitochondrial trafficking and import mechanisms and explain the obstacles that ABCB10 presents to each.

ABCB10 lacking its 105-aa mTP efficiently localizes to the ER, raising the possibility that an isoform of ABCB10 is present in the ER under physiological conditions or that the start codon is mutated. A number of proteins have been described that employ alternate splicing or post-translational modification to direct ER or mitochondria-specific targeting as required. In such cases, the C-terminal end of ABCB10 is potential instability of EGFP-tagged proteins and the necessity to provide targeting information in addition to the mTP. This has been observed in the inner membrane protein cytochrome c1 but is a relatively rare feature of nuclear encoded mitochondrial proteins (34). Although the hydrophobically mutated mTP and the internal domain independently are incapable of targeting EGFP to the mitochondria, their union creates an efficiently targeted protein (without the introduction of a necessary targeting element at their junction). A similar phenomenon was demonstrated by Galanis et al. (35), who by doubling a mitochondrial presequence increased its targeting potency. Considering the challenge to mitochondrial import that ABCB10 presents, the protein is an ideal candidate for possessing redundant import features.

To determine whether aa 1–35 improves mitochondrial targeting by providing additive information to the targeting properties of aa 36–70, aa 1–70-EGFP was constructed; however, the resulting protein was unstable. This demonstrates the potential instability of EGFP-tagged proteins and the necessity to verify that unanticipated cleavage does not occur. Instead, we assessed the effect that the absence of aa 1–35 had on targeting efficiency by constructing \(\Delta35\)-ABCB10-EGFP/V5. Fluorescence microscopy found that the absence of aa 1–35 does not significantly reduce mitochondrial targeting of ABCB10-EGFP. However, protease K digestion of extramitochondrial proteins found that translocation of \(\Delta35\)-ABCB10-V5 across the mitochondrial membrane is reduced, indicating that the processes of mitochondrial targeting and import can be separated. aa 1–35, although necessary for proper mitochondrial import of ABCB10-V5, is not required for that of EGFP, a purely hydrophilic protein. Thus, the distinction between mitochondrial trafficking and import may be more easily observed in proteins that challenge the trafficking/import machinery. The mechanism by which aa 1–35 directs the mitochondrial import of ABCB10 is potentially related to a number of characteristics of ABCB10 and the import machinery. Since mitochondrial proteins are predicted to enter the mitochondria either through a linear or loop conformation, depending on the import pathway, aa 1–35 may affect the folding of ABCB10 and thereby influence the pathway selected (13, 36). Since each mitochondrial membrane possesses its unique import machinery, aa 1–35 may be specifically required for import across the inner membrane but not the outer. Mislocalization of imported proteins among the mitochondrial compartments has been demonstrated in cytochrome c oxidase subunit Va (37). Alternatively, aa 1–35 might simply add import elements needed to overcome a threshold, similar to the additive targeting properties observed in ABCB10. Further investigation will help to tease apart mitochondrial trafficking and import mechanisms and explain the obstacles that ABCB10 presents to each.

ABCB10 lacking its 105-aa mTP efficiently localizes to the ER, raising the possibility that an isoform of ABCB10 is present in the ER under physiological conditions or that the start codon is mutated. A number of proteins have been described that employ alternate splicing or post-translational modification to direct ER or mitochondria-specific targeting as required. In such cases, the C-terminal end of ABCB10 is
Mitochondrial Targeting, Import, and Dimerization of ABCB10

42963

artificial is unknown, for it was recently reported that oxidation during sample preparation may cause disulfide bond formation between dimeric units (42). Since no additional proteins intimately co-assemble with ABCB10, homo-oligomers of ABCB10 may themselves achieve functionality. Interestingly, homo-oligomerization of Δ105-ABCB10 appears to occur in the ER equally as efficiently as wild-type, mitochondria-localized ABCB10, suggesting that the process does not require the mTP of ABCB10 or any mitochondria-specific chaperones.

Acknowledgments—We thank Jim Pearson and Eric Cecil for important contributions; Stuart Orkin for valuable advice; Louis Kerr for technical support with confocal microscopy; Roland Lill for helpful discussions; and Dani Dagan, Eva Czerwiec, Mark Messeri, Jakob Wikstrom, Thorsten Schlaeger, and Shana Katzman for critical review of the manuscript.

REFERENCES


Downloaded from www.jbc.org at Marine Biological Laboratory on April 21, 2009