Biogenesis of MalF and the MalFGK₂ Maltose Transport Complex in *Escherichia coli* Requires YidC*

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The polytopic inner membrane protein MalF is a constituent of the MalFGK₂ maltose transport complex in *Escherichia coli*. We have studied the biogenesis of MalF using a combination of *in vivo* and *in vitro* approaches. MalF is targeted via the SRP pathway to the Sec/YidC insertion site. Despite close proximity of nascent MalF to YidC during insertion, YidC is not required for the insertion of MalF into the membrane. However, YidC is required for the stability of MalF and the formation of the MalFGK₂ maltose transport complex. Our data indicate that YidC supports the folding of MalF into a stable conformation before it is incorporated into the maltose transport complex.

In the Gram-negative bacterium *Escherichia coli*, different inner membrane protein targeting and insertion pathways are operational (1). Although the biogenesis requirements of only a few membrane proteins have been studied thoroughly, most of them are targeted via the signal recognition particle (SRP) pathway to the inner membrane where they are delivered at the Sec translocon (2, 3). The core of the Sec translocon consists of the integral membrane proteins SecY and SecE and the peripheral ATPase SecA (4–6). SecY and -E form a protein-conducting channel that can mediate both translocation and membrane insertion (4, 7). It has been proposed that the translocon channel can open laterally toward the lipid bilayer thereby allowing the release of transmembrane segments (TMs) into the membrane (6, 8). The ATPase SecA is required for the translocation of large (≥60 amino acids) periplasmic domains of inner membrane proteins (1, 9).

The inner membrane protein YidC, which is essential for viability, has been identified as a factor that assists in the integration, folding, and assembly of inner membrane proteins both in association with the Sec translocon and separately (10, 11). Thus far, only a handful of inner membrane proteins that insert via the YidC-only pathway have been identified. All these proteins are small and do not contain a sizable periplasmic domain and more than two TMs. It is not clear how YidC assists the biogenesis of these proteins. During the biogenesis of SRP/Sec translocon-dependent inner membrane proteins, YidC specifically interacts with the TMs of these proteins (e.g. Refs. 12–15). It has been suggested that YidC mediates the transfer of TMs from the Sec translocon into the lipid bilayer (1). Indeed, YidC could be co-purified with the Sec translocon, suggesting a physical connection (14). A recent *in vitro* study using lactose permease (LacY) reveals a novel function for YidC in the co-translational folding of this inner membrane protein rather than its insertion into the membrane (16). The observation that YidC depletion leads to the induction of the Cpx and σE envelope stress responses, which both sense protein misfolding in the cell envelope, also points to a role of YidC in the folding of inner membrane proteins (17, 18).

Here, we have used the inner membrane protein MalF as a model protein to study the role of YidC in the biogenesis of polytopic inner membrane proteins, which are part of a heterooligomeric complex. MalF functions in maltose transport as a 1:1:2 complex with the integral inner membrane protein MalG and the peripheral inner membrane protein MalK (19, 20). The complex belongs to the ATP-binding cassette (ABC) transporter superfamily. MalF is a 514-amino acid-long inner membrane protein containing 8 TMs, with its N and C termini facing the cytosol (21) (see Fig. 1). A large periplasmic domain (~180 amino acids) is present between the third and the fourth transmembrane segments. This domain folds into a trypsin-resistant conformation when MalF is incorporated into the MalFGK₂ maltose transport complex (22).

The biogenesis of MalF was studied using a combination of *in vivo* and *in vitro* approaches. MalF was found to be targeted via the SRP pathway to the Sec/YidC insertion site. Despite close proximity of nascent MalF to YidC during insertion, YidC appeared dispensable for the insertion of MalF into the membrane. However, YidC is required for the stability of MalF and the formation of the MalFGK₂ maltose transport complex suggesting an important role of YidC in the assembly of oligomeric complexes in the inner membrane.
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**EXPERIMENTAL PROCEDURES**

**Materials**—Restriction endonucleases and the Expand Long Template PCR system were from Roche Molecular Biochemicals. The Megashort Script T7 transcription kit was from Ambion Inc. \[^{[35}S\]\text{Methionine and protein A-Sepharose were both from GE Healthcare. T4 ligase and T4 DNA polymerase were both from Epicenter Technologies.} \]p-\text{Dodecyl-}\beta\text{-}o-\text{maltoxyranoside was from Anactrace.} \text{Serva Blue G} was obtained from Serva. All other chemicals were obtained from Sigma. A polyclonal antibody against the c-Myc epitope tag was obtained from Abcam. Antisera against SecY, SecE, YidC, and Ffh were from our own collection. Antisera against MalF, -G, and -K, trigger factor (TF), and L23 were kind gifts from A. Davidson, W. Wickner, and R. Brimacombe, respectively.

**In Vitro Cross-linking**—E. coli strain MRE600 was used to prepare a lysate for translation of *in vitro* synthesized mRNA and suppression of UAG stop codons in the presence of (Tmd)-Phe-tRNA\textsuperscript{sup}. Strain MC4100 was used to isolate inverted membrane vesicles (IMVs). Strain JM110 was used to isolate pC4Meth-derived plasmids for *in vitro* transcription.

pC4Meth-derived plasmids for *in vitro* expression of truncated forms of MalF were constructed by PCR using pTAZFQ as template (23). These plasmids encode truncated MalF fused to a 4× methionine tag to improve labeling efficiency of nascent chains with \[^{[35}S\]\text{methionine.} \text{Immunoprecipitation of nascent chains was enabled by constructing a complementary set of constructs with a C-terminal c-Myc epitope tag (EQKLI-SEEDL) fused to the truncated MalF sequence. Furthermore, amber mutations (TAG) were introduced at the indicated positions to enable sup-tRNA photo-cross-linking, as described previously (14). The nucleotide sequence of all constructs was confirmed by DNA sequencing.

Truncated mRNAs were prepared from HindIII- or Clal-linearized pC4Meth-derived plasmids. *In vitro* translation, targeting to IMVs, photo-cross-linking, carbonate extraction (to separate membrane-bound material from non-membrane-bound material), and sample processing were performed as described previously (14).

**Strains, Plasmids, Growth Conditions, and Assay Used in MalF Membrane Targeting/Insertion Experiments in Vivo**—The 4.5 S RNA conditional strain FF283 was cultured in M9 minimal medium (for composition, see Ref. 24) supplemented with 1 mM IPTG as described previously (24). To deplete cells of 4.5 S RNA, cells were grown to mid-log phase in the absence of IPTG. The temperature-sensitive amber suppressor SecA deletion strain BA13 and the control strain DO251 were cultured in M9 minimal medium at 30 °C, as described previously (24). To deplete cells of SecA, they were grown to mid-log phase at 41 °C. The SecE depletion strain CM124 was cultured in M9 minimal medium supplemented with 0.2% glucose and 0.2% l-arabinose as described previously (24). To deplete cells of SecE, cells were grown to mid-log phase in the absence of l-arabinose. Depletion of SecA and SecE was checked by monitoring the accumulation of pro-OmpA during a short pulse-labeling step with \[^{[35}S\]\text{methionine.} \text{The SecG deletion strain KN370 was cultured in M9 minimal medium to mid-log phase, and plasmid pH\textsuperscript{+} was used to complement with SecG as described previously (24).} \text{The YidC depletion strain FTL10 was cultured in M9 minimal medium supplemented with 0.2% glucose and 0.2% l-arabinose, as described previously (25). To deplete cells of YidC, cells were grown to mid-log phase in the absence of l-arabinose.} 

MalF was expressed by arabinose induction from the pBAD24 vector (26) in strains FF283, BA13, DO251, and KN370 (\(z\text{pH}^+\)), by IPTG induction from the pEH1 vector (27) in strain CM124 and by IPTG induction from the pEH3 vector (27) in strain FTL10. Where appropriate, ampicillin (final concentration, 100 \(\mu\text{g/ml}\)), chloramphenicol (final concentration, 30 \(\mu\text{g/ml}\)), kanamycin (final concentration, 50 \(\mu\text{g/ml}\)), streptomycin (final concentration, 25 \(\mu\text{g/ml}\)), and tetracycline (final concentration, 12.5 \(\mu\text{g/ml}\)) were added to the medium.

For all experiments cells were grown to mid-log phase. Expression of the constructs was induced for 3 min with either IPTG (final concentration, 1 mM) or l-arabinose (final concentration, 0.2%). Cells were labeled with \[^{[35}S\]\text{methionine (100 Ci/ml, 1 Ci = 25 GBq) for 30 s.} \text{After labeling, cells were converted to spheroplasts. For spheroplasting, cells were collected at 14,000 rpm for 30 s in a microfuge, resuspended in ice-cold buffer (40% w/v sucrose, 33 mM Tris-HCl, pH 8.0), and incubated with lysozyme (final concentration, 5 \(\mu\text{g/ml}\) and 1 mM EDTA for 15 min on ice. Aliquots of the spheroplast suspension were incubated on ice for 1 h either in the presence or absence of proteinase K (final concentration, 0.3 mg/ml). Subsequently, phenylmethylsulfonyl fluoride was added to the spheroplast suspension (final concentration, 0.33 mg/ml) to inhibit the protease. After addition of phenylmethylsulfonyl fluoride, samples were precipitated with trichloroacetic acid (final concentration, 10%), washed with acetone, resuspended in 10 mM Tris-HCl, pH 8.0/2% SDS, immunoprecipitated with antisera to MalF, OmpA (a periplasmic control), and AraB/bandX (a cytoplasmic control), washed, and analyzed by standard SDS-PAGE (24). Gels were scanned in a Fuji FLA-3000 phosphorimaging device and quantitated using Image Gauge (version 3.4). All pulse experiments were repeated at least three times.

For the MalF biotinylation-based membrane insertion assay, the genes encoding the MalF derivatives MalF\textsubscript{1} (MalF with a PBST biotinylation domain engineered in the periplasmic loop between TM3 and -4, Fig. 1) and MalF\textsubscript{L} (MalF with a PBST biotinylation domain engineered in the cytoplasmic loop between TM4 and -5, Fig. 1) were cloned into the expression vector pASK-IBA3 (IBA GmbH, Germany) (28). The expression vectors containing the two derivatives of MalF were transformed into the conditional YidC strain FTL10. Cells were grown overnight in LB medium containing arabinose. Cultures were then washed and transferred into fresh LB medium containing arabinose or glucose to deplete cells for YidC. Expression of MalF\textsubscript{1} and -L was induced by addition of 0.2 \(\mu\text{g/ml}\) anhydrotriacycline. After 4 h samples were taken and processed for SDS-PAGE and immunoblotting. Biotinylated proteins were detected using streptavidin-horseradish peroxidase (GE Healthcare).

**Analysis of the Accumulation Levels of the MalFGK\textsubscript{2} Maltose Transport Complex by Blue Native PAGE**—Overnight cultures of MC4100\textsuperscript{ara} \textsuperscript{7} and FTL10 cells were grown in LB medium supplemented with 0.2% glycerol. To induce the expression of.
The accumulation levels of MalF, MalG, and MalK in inner membranes were monitored by Western blot analysis. Sample preparation and running conditions were essentially as described previously (29). For blotting of blue native-PAGE gels, the cathode buffer containing 0.02% Serva Blue G was exchanged after one-third of the run to 0.02% Serva Blue G was exchanged after one-third of the run to prevent excessive binding of the dye to the polyvinylidene difluoride membrane. Blotting and decoration of blots with antisera to MalF, -G, -K, YidC, and SecY were performed as described previously (29).

Western Blotting Analysis—The accumulation levels of MalF, MalK, MalG, and YidC in inner membranes were monitored by immunoblot analysis. IMVs (3–5 million) were processed as described above under “Strains, Plasmids, Construction of malG and malK Deletion Strains and FTL10 Getting/Insertion Experiments” (29). After 3 h of growth, cells were collected by centrifugation and fractionated, and IMVs were isolated as described before (29). The accumulation levels of the MalFGK2 maltose transport complex were analyzed by blue native-PAGE (30) followed by Western blot analysis. Sample preparation and running conditions were essentially as described previously (29). For blotting of blue native-PAGE gels, the cathode buffer containing 0.02% Serva Blue G was exchanged after one-third of the run to cathode buffer containing 0.002% Serva Blue G. This was done to prevent excessive binding of the dye to the polyvinylidene difluoride membrane. Blotting and decoration of blots with antisera to MalF, -G, -K, YidC, and SecY were performed as described previously (29).

RESULTS

Nascent MalF Interacts with TF, Ffh, and the Ribosomal Component L23 and Inserts into the Inner Membrane in the Vicinity of YidC and the Sec Translocon—We initially analyzed the pathway of targeting and membrane insertion of MalF using an in vitro translation/targeting/photo-cross-linking approach. Radiolabeled nascent MalF of 88 amino acids was synthesized in a wild-type cell-free E. coli extract in the presence of inverted inner membrane vesicles to allow membrane targeting and insertion on the ribosome nascent chain complex (14). At this length, the ribosome nascent chain complex is expected to fully expose the first and second TMs of MalF, assuming that the ribosome covers ~30 amino acids (33). To enable the incorporation of a photoreactive probe in the middle of the first TM of MalF, a single TAG amber codon was introduced at position 26. Following translation in the presence of (Tmd)Phe-tRNAsup, nascent MalF interacts with TF, Ffh, and the ribosomal component L23 and inserts into the inner membrane vesicles to allow membrane targeting and insertion on the ribosome nascent chain complex (14). At this length, the ribosome nascent chain complex is expected to fully expose the first and second TMs of MalF, assuming that the ribosome covers ~30 amino acids (33). To enable the incorporation of a photoreactive probe in the middle of the first TM of MalF, a single TAG amber codon was introduced at position 26.
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from untargeted material. Cross-linked partners were identified by immunoprecipitation with specific antibodies.

The TAG26 mutation in 88MalFTAG26 was efficiently suppressed by (Tmd)Phe-trNA\sup (data not shown), resulting in nascent MalF of the expected molecular weight. 88MalFTAG26 was efficiently targeted to IMVs, as judged by the relative amount of carbonate-resistant material (Fig. 2, compare lanes 1 and 2). Upon UV irradiation, several specific cross-linking products could be observed in the carbonate supernatant and pellet fractions (Fig. 2, lanes 3 and 7). An adduct of ~55 kDa could be immunoprecipitated with antisera to Ffh, the protein component of E. coli SRP (Fig. 2, lane 5). Substantial amounts of smaller adducts were also precipitated with antisera to Ffh, which most probably represent cross-linked degradation products of Ffh (34). A cross-linking product of ~65 kDa could be immunoprecipitated with antisera to trigger factor (TF), which is a cytosolic chaperone with general affinity for nascent polypeptides that resides on the ribosome close to the nascent chain exit site (35, 36) (Fig. 2, lane 4). A prominent ~20-kDa cross-link product was precipitated with antibodies directed toward the ribosomal protein L23 (Fig. 2, lane 6), which is located near the nascent chain exit site (37). Ffh and TF have both been shown to associate with L23 (34, 36, 38–40). In the carbonate pellet fraction a very strong cross-linking product of ~60 kDa becomes apparent after UV irradiation (Fig. 2, lane 7). We could immunoprecipitate this adduct with antisera to YidC (Fig. 2, lane 8) (35). A less prominent band of ~40 kDa could be immunoprecipitated with antisera to SecY (Fig. 2, lane 9) (35).

In summary, using this unbiased approach it was shown that, at a nascent chain length of 88 amino acids, TM1 of MalF interacts with TF, SRP, and L23 and inserts into the inner membrane in a carbonate-resistant state close to YidC and SecY. This strongly suggests that MalF is targeted via the SRP pathway to the SecY/YidC insertion site in the inner membrane.

TM1 of MalF Remains Close to YidC and the Sec Translocon during Insertion of TM2 and TM3 of MalF into the Membrane, and TM2 and TM3 Insert into the Membrane in a Molecular Environment Similar to That of TM1—To study the sequence of events taking place during the biogenesis of the three N-terminal TMs of MalF, nascent chains of increasing lengths were synthesized in vitro. The shortest nascent chain, 68MalF, is expected to fully expose the first TM segment from the ribosome nascent chain complex (see Fig. 3A). At a length of 100 amino acids, 100MalF exposes TM1 and TM2. 131MalF, the longest nascent chain, exposes TM1, TM2, and TM3. By placing a TAG codon at position 26 and by exploiting the photocross-linking procedure described above, the molecular environment of TM1 in each of these translation intermediates was studied. C-terminal c-Myc epitope tags were introduced to enable purification of nascent chains by immunoprecipitation. Previous work has shown that this approach can prevent misinterpretations of cross-linking experiments due to premature termination of translation (41).

For each of the constructs, the TAG codon was suppressed, and nascent MalF of the expected apparent molecular weight was produced (data not shown). As shown in Fig. 3B, all nascent chains could be immunoprecipitated using the anti-c-Myc antibodies. Upon UV irradiation, several adducts were resolved and co-immunoprecipitated with the nascent chains of desired length. 68MalFTAG26-Myc cross-linked YidC and SecY (Fig. 3B, lane 1). Also, an adduct of ~25 kDa was present that could be precipitated with anti-SecE (Fig. 3C, lane 3). As expected, the 100MalFTAG26-Myc nascent chain was cross-linked to YidC and SecY, similar to 88MalFTAG26 (compare Fig. 2, lane 7 to Fig. 3B, lane 2), because both constructs expose TM1 and TM2 and share a cross-link site at an identical position. Notably, the SecE cross-link was lost at this length. These cross-links represent the molecular environment of TM1 after both TM1 and TM2 are completely exposed outside the ribosome (Fig. 3A). Upon further extension of the MalF nascent chain to a length of 131 amino acids, the photoprobe in TM1 continued to cross-link YidC and SecY, whereas the SecE adduct reappeared (Fig. 3B, lane 4). These results show that TM1 of MalF initially

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FIGURE 2. Interactions of nascent 88MalFTAG26. 88MalFTAG26 was synthesized in vitro in the presence of (Tmd)Phe-trNA\sup and E. coli wild-type IMVs. After translation, samples were either kept in the dark (−UV) or, alternatively, cross-linking was induced by UV irradiation (+UV). A carbonate extraction was performed to separate membrane-integrated (p) from soluble and peripheral membrane proteins (s). Immunoprecipitations were performed using the indicated antibodies. Immunoprecipitated samples and carbonate pellets and supernatants were subjected to SDS-PAGE and phosphorimagintg. The position of the 88MalFTAG26 nascent chain in the gel is indicated. Peptidyl-tRNA band is marked with an asterisk.
The interactions of TM2 were mapped by establishing the cross-links of 100MalFTAG47-Myc and 131MalFTAG47-Myc nascent chains. With both constructs, cross-links to YidC, SecY, and SecE were co-immunoprecipitated with the nascent chains by anti-C-Myc antibody (Fig. 3B, lanes 3 and 5). 131MalFTAG85-Myc was also cross-linked to YidC, SecY, and SecE, indicating that the molecular environment of TM3 at this stage of biogenesis was comparable to that of TM1 and TM2 (Fig. 3B, lane 6). These data indicate that TM2 and TM3 initially insert into the membrane close to YidC, SecY, and SecE and therefore behave like TM1. TM2 also remains close to these proteins when TM3 has inserted into the membrane. This indicates that at least the first three TMs of MalF assemble simultaneously at the Sec/YidC insertion site in the inner membrane and do not partition completely into the lipids at this stage in biogenesis.

Efficient Membrane Targeting and Insertion of MalF Requires SRP and the Sec Translocon—Next, targeting and insertion of full-length MalF expressed from a plasmid were monitored in vivo in SRP and Sec translocon mutant strains by analyzing the proteinase K accessibility of MalF in spheroplasts. OmpA, outer membrane protein A, served as a positive control for spheroplast formation by monitoring the proteinase K sensitivity of its periplasmic domain (42). Band X, a cytoplasmic protein, is a negative control. It is used to confirm the spheroplasts are intact (42).

Depletion of the essential SRP component 4.5 S RNA compromises the SRP-targeting pathway (43). Depletion of 4.5 S RNA strongly inhibited proteolysis of MalF in spheroplasts, as compared with spheroplasts made of non-depleted (control) cells (Fig. 4A). Notably, processing of the Sec-dependent OmpA protein was not affected upon 4.5 S RNA depletion, which shows that the Sec translocon was functional under these conditions. Assembly of MalF was further studied in a SecE depletion strain. Upon depletion of SecE, SecY is rapidly degraded by the FtsH protease resulting in the loss of the

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inserts into the membrane in a carbonate resistant form close to YidC, SecY, and SecE. At all nascent chain lengths studied here, the contacts of TM1 with YidC and SecY remain. The SecE contact is temporarily lost upon extension of the nascent chain, but reappears at a later stage, when TM1, TM2, and TM3 have all left the confinement of the ribosome.
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A. Proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in FF283 (4.5 S RNA depletion strain) spheroplasts, made from cells cultured in the presence/absence of IPTG (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SRP-depletion.

B. Proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in CM124 (SecE depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecE-depletion.

C. Proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in CM124 (SecE depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecE-depletion.

D. Proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in CM124 (SecE depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecE-depletion.

E. Proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in CM124 (SecE depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecE-depletion.

FIGURE 4. Membrane insertion of MalF is SRP/SecAYE-dependent and YidC-independent. A, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in FF283 (4.5 S RNA depletion strain) spheroplasts, made from cells cultured in the presence/absence of IPTG (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SRP-depletion. B, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in CM124 (SecE depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecE-depletion. C, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in BA13 and DO251 (SecA depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecA-depletion. D, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in KN370/KN370 harboring pH₇ (SecG deletion strain, pH₇ plasmid expressing SecG) spheroplasts. Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecG-depletion, D, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in FtL10 (YidC depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon YidC-depletion.

Taken together, these data indicate that MalF is targeted to the inner membrane via the SRP pathway and that insertion of MalF into the inner membrane is SecAYE-dependent. This is in keeping with the in vitro cross-linking data described above and previous in vivo studies (e.g. Refs. 45, 47, 48).

Insertion of MalF into the Inner Membrane Does Not Depend on YidC—The close proximity of nascent MalF to YidC during insertion in the membrane as shown by cross-linking (see Figs. 2 and 3) suggests that YidC plays an important role in the biogenesis of MalF. Therefore, insertion of plasmid-expressed MalF into the inner membrane was monitored in the YidC depletion strain FtL10 by analyzing the proteinase K accessibility of MalF in spheroplasts (Fig. 4E). Insertion of MalF into the inner membrane was not affected upon the depletion of YidC. Cells were efficiently depleted of YidC as determined by immunoblotting (data not shown), and the translocation of OmpA was, as expected, not affected upon YidC depletion.

To study the role of YidC in the insertion of MalF into the inner membrane in more detail, we used a very sensitive biotinylation assay (28). Two MalF derivatives were engineered, one containing the Propionibacterium shermanii transcarboxylase (PSBT) biotin-accepting domain from the 1.3 S subunit of PSBT in the large periplasmic loop between TM3 and TM4 (MalF-PSBT I), and the other containing the PBST domain in the small periplasmic loop between TM3 and -4 (MalF-PSBT L) (Fig. 1). The PSBT domain is biotinylated by the cytoplasmic enzyme biotin ligase. Hence, to be biotinylated, the PBST domain must at least temporarily be exposed to the cytoplasm (28). As expected, MalF-PSBT L was efficiently biotinylated irrespective of the presence/depletion of YidC (Fig. 5). MalF-PSBT L was neither in the presence nor upon the deletion of YidC biotinylation assay and control spheroplasts, made from cells cultured at 41 °C (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing is affected upon SecA-depletion. D, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in KN370/KN370 harboring pH₇ (SecG deletion strain, pH₇ plasmid expressing SecG) spheroplasts. Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon SecG-depletion. E, proteinase K accessibility of MalF (top) and OmpA/band X (bottom) in FtL10 (YidC depletion strain) spheroplasts, made from cells cultured in the presence/absence of arabinose (+, non-depleted cells; −, depleted cells). Cells were pulse-labeled and processed as described under "Experimental Procedures." OmpA secretion/processing was not affected upon YidC-depletion. MalF bands were quantified, and the band volumes of spheroplasts made of control cells not treated with proteinase K were set to 100%.

SecY/E core of the translocon (44). Depletion of SecE strongly inhibited proteolysis of MalF in spheroplasts, as compared with non-depleted (control) cells (Fig. 4B and Ref. 45). As expected, processing and translocation of the Sec-dependent OmpA was strongly affected upon SecE depletion. Upon depletion of SecA, which does not affect the rest of the Sec translocon (46), proteolysis of MalF in spheroplasts was strongly reduced compared with the control sample (Fig. 4C). This indicates that SecA is required for the biogenesis, presumably by energizing the translocation of the periplasmic loop between TM3 and TM4. Processing and translocation of OmpA were, as expected, also affected upon SecA depletion. Finally, the insertion of MalF was not affected in the absence of the non-essential Sec translocon component SecG (Fig. 4D).
YidC is Required for the Stability of MalF—Subsequently, the steady-state level of endogenously expressed MalF in the inner membrane of cells depleted of YidC and control cells was monitored by immunoblotting. Upon YidC depletion, the accumulation level of full-length MalF was strongly reduced, and two putative MalF degradation products appeared (Fig. 7A). YidC depletion did not affect the level of MalG, whereas the level of MalK attached to the membrane and in whole cells was affected (~70% down) upon YidC depletion (Fig. 7A).

The immunoblotting experiments could mean that YidC is required for the stability of MalF. To monitor the stability of MalF upon YidC depletion in more detail, a pulse-chase approach was used. Cells depleted of YidC and control cells were cultured and labeled with [35S]methionine as described under “Experimental Procedures.” After chases of 3, 30, and 60 min, MalF was isolated by means of immunoprecipitation, and precipitated material was subsequently analyzed by SDS-PAGE and phosphorimaging (Fig. 7B). After a chase of 3 min, the amount of MalF was similar in cells expressing and depleted of YidC. However, in cells depleted of YidC the MalF signal strongly decreased in time, whereas it remained stable in the control cells. Because the level of MalK in cells depleted of YidC was lower than in wild-type cells, MalF stability was also monitored in cells not expressing its complex partners MalK and MalG (Fig. 7B). In both malK and malG deletion strains MalF stability was not affected, which is in keeping with observations made by Traxler and Beckwith (22). This indicates that YidC is important for MalF stability rather than its complex partners MalF and MalG.

DISCUSSION

So far, the role of YidC in the folding of complex membrane proteins has only been studied in detail for the monomeric membrane protein lactose permease (LacY). It was shown,
Biogenesis of MalF and MalFGK2 in E. coli Requires YidC

![Image]

**FIGURE 7. YidC is required for the stability of MalF.** A, MC4100ΔmalG and FTL10 cells were cultured in LB medium in the presence of maltose. Inner membranes were isolated and analyzed by SDS-PAGE and immunoblotting with antibodies directed against YidC, MalK, MalF, and -G. The band in the αMalG blot that is marked with an asterisk is a background band, which indicates that equal amounts of membranes were loaded. B, MC4100ΔmalG, FTL10, SW1242(ΔmalG), and SW1282(ΔmalK) cells were cultured in LB medium in the presence of maltose and labeled with [35S]methionine as described under “Experimental Procedures.” 3, 30, and 60 min after an excess of cold methionine was added to the cultures, cells were precipitated with trichloroacetic acid, and MalF was subsequently isolated by immunoprecipitation. Immunoprecipitations were analyzed by SDS-PAGE and phosphorimaging (upper panel). Relative band volumes are shown in the graph in the lower panel.

using *in vitro* reconstitution approaches and a folding assay based on monoclonal antibodies against conformational epitopes of LacY, that YidC is required for the co-translational folding of LacY (16). As a consequence, the stability of LacY was affected upon YidC depletion (16). Here, we have studied the role of YidC in the folding of a polytopic inner membrane protein that is part of a hetero-oligomeric complex. Using a combination of an *in vitro* cross-linking approach, *in vivo* biogenesis assays, and blue native-PAGE we have monitored the role of YidC at different stages in the biogenesis of MalF, which is a constituent of the MalFGK2 maltose transport complex. First, however, we re-evaluated the SRP/Sec dependence of MalF, because not all studies on its SRP/Sec dependence were conclusive.

Targeting of MalF via the SRP pathway had been studied before using genetic approaches. In the Slo (synthetic lethality upon overexpression) screen, which was developed by Bernstein and co-workers (49), MalF was not identified as an SRP substrate. Furthermore, blocking of the SRP-targeting pathway by overexpression of a non-functional variant of the SRP-receptor FtsY only marginally affected the insertion of a MalF alkaline phosphatase fusion protein (49). However, in a genetic screen designed to isolate *E. coli* mutants affected in membrane protein assembly, a strain with a mutation in the *fss* gene, which encodes the SRP component 4.5 S RNA, was isolated that affected the targeting of a MalF LacZ fusion protein (47). In the present study we show that Ffh, a component of the SRP, can be readily cross-linked to TM1 of nascent MalF *in vitro* and that the *in vivo* targeting of full-length MalF is strongly hampered in cells depleted of the SRP component 4.5 S RNA. It should be mentioned that the observed cross-linking to Ffh is highly significant considering the extremely low abundance of Ffh in wild-type cells and hence in the translation lysate used in this assay (50). This strongly suggests that, at least a considerable fraction of, MalF is targeted to the inner membrane via the SRP pathway. In short MalF nascent chains, TM1 not only interacts with Ffh, but also with the ribosomal protein L23, which is located near the exit of the ribosomal tunnel, and the chaperone TF (37). SRP and the chaperone TF bind simultaneously to L23 (51, 52). TF generically binds nascent peptides exiting the ribosome (53). Neither TF depletion nor overproduction has any notable effect on MalF membrane targeting/insertion. It is in keeping with the notions that TF and SRP sample nascent chains on the ribosome in a nonexclusive fashion, and that binding of the SRP receptor, FtsY, to ribosome-bound SRP excludes TF from the ribosome allowing the docking of the ribosome to the Sec translocon (51).

Most SRP-dependent proteins studied thus far are targeted to the Sec translocon in the inner membrane (1, 3). Indeed, by means of *in vitro* cross-linking using MalF nascent chains of different length, we showed that the first three TMs of MalF insert and remain in the vicinity of the Sec translocon core components SecY and SecE during the biogenesis of MalF. Consistently, *in vivo* assembly of MalF is affected upon SecE depletion (our data and Ref. 45). This is also in keeping with the observation that the membrane insertion of a MalF alkaline phosphatase fusion is affected in a strain with a mutation in SecY that specifically affects the biogenesis of inner membrane proteins (45, 54). In a SecG null mutant background there was no effect on the insertion of MalF into the membrane.

Thus far, azide, an inhibitor of SecA, has been used to show that SecA is involved in the biogenesis of MalF (45). To examine the involvement of SecA in the biogenesis of MalF more directly, we monitored the assembly of MalF in a SecA depletion strain. Upon depletion of SecA, which does not affect the rest of the Sec translocon (46), insertion of MalF into the inner membrane was strongly hampered. Taken together, biogenesis of MalF appears to be SRP/SecAYE-dependent.

Using two different assays, it was shown here that *in vivo* insertion and topogenesis of MalF into the membrane were not affected upon YidC depletion. However, *in vitro* cross-linking experiments demonstrated that the first three TMs of MalF not only insert into the membrane close to SecY and SecE but also to YidC. Notably, they remain in this environment at least until the first three TMs of MalF have emerged from the ribosome in the co-translational insertion process. Correspondingly, it has been shown that the first TMs of the nascent polytopic inner membrane protein MtlA assemble simultaneously at the Sec/YidC insertion site (15). Recently, docking of the projection maps of YidC and the SecYEG translocon generated a potential contiguous pathway for translocating TMs (55). This pathway could provide an environment for the TMs where their folding and assembly can occur prior to their release into the lipid bilayer. The integration of aquaporin-4 into the endoplasmic

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reticulum membrane by the Sec61 translocon has been shown to involve sequential triage of TMs from their initial portal of entry into multiple secondary sites within the translocon (56). It was proposed that this mechanism provided a means to facilitate early folding events before release into the lipid bilayer. It has been speculated that the endoplasmic reticulum-localized TRAM protein has a similar function as YidC, and it has also been shown that there are specialized chaperones in the endoplasmic reticulum mediating the co-translational folding of specific membrane proteins (57–59).

In contrast to the TMs of MalF, MtlA, and aquaporin-4, the two TMs of the inner membrane protein leader peptidase (Lep) partition one by one into the lipid bilayer via the Sec/YidC interface according to the linear insertion model of TMs (41). A possible explanation for this discrepancy may relate to the role of the TMs in the respective proteins. The TMs of Lep merely act as a membrane anchor for the periplasmic P2 domain of Lep to which the catalytic activity of Lep is confined, whereas the TMs of MtlA, MalF, and aquaporin-4 play key roles in the functioning of these proteins. Proper folding and interaction of the TMs of MalF, MtlA, and aquaporin-4 is an absolute requirement for the functioning of these proteins and may also be a prerequisite for complex formation.

The cross-linking data are consistent with the notion that YidC functions in co-translational folding of complex inner membrane proteins. The importance of YidC in membrane protein folding and assembly is also suggested by the observation that YidC depletion leads to the induction of Cpx and σE envelope stress responses that sense protein abnormalities in the inner and outer membranes as well as in the periplasm (17, 18). Misfolding or misassembly of membrane proteins is often linked to decreased stability (60). Although MalF was rapidly degraded upon YidC depletion, the absence of the other components of the MalFGK₂ complex did not affect the stability of the protein as judged by pulse-chase analysis. However, trypsin accessibility experiments showed that the folding of the large periplasmic loop of MalF is not completed before it is incorporated into the MalFGK₂ complex (22). Possibly, YidC mainly chaperones the initial folding of MalF by acting on its integral membrane part, which is likely to be critical for MalF stability and hence for assembly of MalFGK₂ complexes.

In the mal operon, the gene encoding MalG is located downstream of the gene encoding MalF; i.e., MalG is synthesized after MalF and most likely will insert after MalF into the membrane. It is tempting to speculate that MalF is kept by YidC in a conformation that allows a smooth MalF-MalG complex formation. However, we did not manage to isolate Sec/YidC-MalF complexes (results not shown). In contrast, the YidC homolog in the thylakoidal membrane, Albino3, could be isolated in a complex within the membrane-inserted D1 protein, before it was incorporated into the photosystem II complex (61). The Sec and/or YidC-MalF complexes, if they indeed exist, may be too fragile or transient to purify.

Summarizing, we have shown that the polytopic integral membrane protein MalF, which is a constituent of the MalFGK₂ complex, is targeted to the inner membrane via the SRP pathway to the Sec/YidC insertion site. Despite close proximity of nascent MalF to YidC during insertion, YidC is not required for the insertion of MalF into the membrane. However, YidC is required for the stability of MalF and the formation of the MalFGK₂ maltose transport complex. Our data indicate that YidC supports the folding of MalF into a stable conformation before it is incorporated into the maltose transport complex.

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The compound concentrations (EC50) evoking half-maximal estrogenic activity in transcription activation, the dissociation inhibition constants (Kᵢ) calculated from the IC₅₀ values of the ES2 displacement curves (Fig. 1), and the corresponding Ki values calculated using the 6D-QSAR model are compared. For Polysantol and Javanol, only the Kᵢ values calculated using the 6D-QSAR model are reported. The IC₅₀ values for the remaining stereoisomers are as follows: 140 ± 173 µM Polysantol (S,S), 216 ± 122 µM Polysantol (S,R), 2.86 ± 4.45 mM Polysantol (R,R), 2.41 ± 2.64 mM Javanol (R,S), 176.7 ± 567 µM Javanol (S,S), and 40.73 ± 23.33 µM Javanol (S,R). The Kᵢ values for the remaining stereoisomeric forms are as follows: 299 µM Polysantol (S,S), 83 µM Polysantol (S,R), 1 mM Polysantol (R,R), 931 µM Javanol (R,S), 68 µM Javanol (S,S), and 15 µM Javanol (S,R).

<table>
<thead>
<tr>
<th>Compound</th>
<th>EC₅₀ of ER activation</th>
<th>Kᵢ of ER competitive binding</th>
<th>Kᵢ of ER predicted</th>
</tr>
</thead>
<tbody>
<tr>
<td>Estradiol</td>
<td>0.16 ± 0.01 nM</td>
<td>1.6 ± 1 nM</td>
<td>2 nM</td>
</tr>
<tr>
<td>MC</td>
<td>86 ± 8 µM</td>
<td>76 ± 6 µM</td>
<td>21 µM</td>
</tr>
<tr>
<td>Polysantol (R,S)</td>
<td>98 ± 23 µM</td>
<td>56 ± 5 µM</td>
<td>26 µM</td>
</tr>
<tr>
<td>Javanol (R,S)</td>
<td>50 ± 4 µM</td>
<td>ca. 50 µM</td>
<td>5 µM</td>
</tr>
<tr>
<td>Androstenol</td>
<td>52 ± 3 µM</td>
<td>ca. 25 µM</td>
<td>10 µM</td>
</tr>
</tbody>
</table>

* The indicated stereoisomers were used for predicting Kᵢ values, whereas experiments were performed using mixtures of stereoisomers.

On page 36212, the sequences of PCR primers of IFN-α are incorrect.

The correct sequences are as follows: IFN-α, 5’-ATGGCTAGGCTCTGT-GCTTTCT-3’ and 5’-AGGGCTTCCAGACTTCTGCTTG-3’. We suggest that subscribers photocopy these corrections and insert the photocopies in the original publication at the location of the original article. Authors are urged to introduce these corrections into any reprints they distribute. Secondary (abstract) services are urged to carry notice of these corrections as prominently as they carried the original abstracts.