Non-bilayer Lipids Stimulate the Activity of the Reconstituted Bacterial Protein Translocase*

(Received for publication, September 29, 1999, and in revised form, November 2, 1999)

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*This work was supported by a PIONIER grant of the Netherlands Organization for Scientific Research (N.W.O.) and by EEC Biotech Grants BIO2 CT 930254 and BIO4 CT 960097. The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked "advertisement" in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.
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Complementary genetic and biochemical approaches have shown that the translocation of proteins across the inner membrane of Escherichia coli is mediated by the translocase (for reviews see Refs. 1–3). The essential subunits of the translocase are the dissociable peripheral ATPase termed SecA and integral membrane proteins SecY and SecE (4, 5). The SecYE complex forms a preprotein-conducting channel that may associate with SecG or the heterotrimeric SecDFYajC complex (6). SecA binds with high affinity to the SecYEG complex and functions as a molecular motor that utilizes the binding and hydrolysis of ATP to drive the stepwise translocation of a preprotein across the membrane (7). In addition, the proton motive force accelerates the translocation reaction (8).

Phospholipids have been shown to play an important role in protein translocation. The E. coli inner membrane mostly consists of the zwitterionic phospholipid phosphatidylethanolamine (PE, 70–75%) and two anionic phospholipids, phosphatidyglycerol (PG, 20–25%) and cardiolipin (CL, 5–10%). The membrane lipid composition is normally tightly regulated, but by the use of strains engineered in the expression of critical phospholipid biosynthetic enzymes, the in vitro manipulation of the bulk phospholipid composition has been achieved (9). The major anionic phospholipids PG and CL can be depleted in a strain in which the expression of the pgsA gene encoding the phosphatidylglycerophosphate synthase is controlled. At low PG and CL content (2–3%), protein translocation is severely compromised, whereas lack of only CL did not affect translocation (10). Only the negative charge of the polar lipid head group is important as many other anionic phospholipids are able to restore the protein translocation activity of PG and CL-depleted inner membrane vesicles (IMVs) or in reconstituted proteoliposomes (4, 11, 12). The efficiency of protein translocation is directly proportional to the amount of anionic phospholipids (13). Anionic phospholipids influence various steps in the preprotein translocation cascade as follows. (i) They promote the interaction of SecA with the membrane surface (14–16) and SecYEG (11) and are needed for the SecA translocation ATPase activity, i.e. the preprotein-stimulated ATPase activity of the SecYEG-bound SecA (12, 14). At low levels of anionic phospholipids, excess SecA can compensate for the reduced translocation activity (17), suggesting a role of these lipids in the targeting of preproteins and SecA to the membrane. In addition, the endogenous SecA ATPase activity at low Mg$^{2+}$ concentration is stimulated by the presence of anionic phospholipids, an activity termed SecA lipid ATPase (14). (ii) Membrane insertion of the positively charged signal sequence of a preprotein is dependent on anionic phospholipids (18–22). (iii) Anionic phospholipids stabilize the SecYEG complex during octyl glucoside solubilization (12, 23) and (iv) influence the acquisition of the correct topology of membrane proteins (24).

The cold-sensitive growth defect of a secG null strain (25–28) can be suppressed by various gene products involved in phospholipid biosynthesis along with the pgsA gene. Likewise, the cold-sensitive growth defect of the secAceR11 mutant strain is suppressed by overproduction of the PgsA protein (28). These effects have been attributed to an increase in the anionic phospholipid content that restores the secretion defect of these strains by facilitating the SecA catalytic cycle at low temperature (28). The exact mechanism of this activation, however, remains obscure as studies with photoreactive phospholipid analogues suggest that the SecYEG-bound, membrane-inserted form of SecA is not in contact with phospholipids (29, 30).

The other main phospholipid of the E. coli inner membrane is

[phospho-l-serine]; DOPG, 1,2-dioleoyl-sn-glycero-3-phosphoglycerol; DOG, diolipoglycerol; OG, n-octyl β-glucopyranoside; DDM, dodecyl maltoside; IMVs, inner membrane vesicles; PAGE, polyacrylamide gel electrophoresis.
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The type II lipid PE. PE adopts a non-bilayer hexagonal II phase conformation (31). Deletion of the pssA gene (32), which encodes the phosphatidylserine (PS) synthase, renders cells devoid of the amino-based phospholipids PS and PE. This results in severe pleiotropic effects on membrane protein function and the inactivation of protein translocation (33–35). For growth, this strain is dependent on the presence of a high concentration of divalent cations (Ca$^{2+}$, Mg$^{2+}$, or Sr$^{2+}$) (32, 35). These cations stimulate the bilayer to non-bilayer transition of CL, which comprises 44% of the total phospholipid in this strain (36). It therefore appears that the polymorphic behavior of these lipids is essential for growth, a role normally fulfilled by PE. The requirement for non-bilayer lipids for protein translocation appears less strict than for anionic lipids (35), but the mechanism by which these lipids act on protein translocation is unknown. Non-bilayer lipids only marginally affect the SecA lipid ATPase activity (37) and appear not essential for the functional reconstitution of protein translocation using octyl glucoside (OG)-purified SecYEG complex (11).

The reconstitution of prepout protein translocation with only a limited set of purified Sec proteins provides a unique opportunity to assess systematically the lipid requirement for prepout translocation. We now report on a method that allows purification of the SecYEG complex in a delipidated state. The activity of the delipidated SecYEG can be restored after reconstitution into liposomes with a defined phospholipid composition. The data with the purified E. coli SecYEG complex not only confirm the hypothesis that anionic lipids are essential for activity but also show that non-bilayer lipids stimulate the activity of the reconstituted translocon. Strikingly, with the Bacillus subtilis SecYEG both lipid classes are essential for activity. Overall, optimal activity is observed when anionic and non-bilayer lipids are present at a concentration that matches that of the natural membrane.

EXPERIMENTAL PROCEDURES

Plasmids—pET610 (38) was partially digested with NotI/BamHI, and the large SecE fragment was cloned into pET302 (39) to yield pET720 that allows the overproduction of the E. coli SecYE. Materials—E. coli SecA (40), B. subtilis SecA (41), SecB (42), pre-OmpA (43), and pre-PhoB (44) were purified as described. ProOmpA and pre-PhoB were labeled by iodination with K$^{125}$I (44) and stored frozen in 6 M urea, 50 mM Tris-Cl, pH 7.5. After 30 min, 1,2-dioleoyl-sn-glycero-3-phosphocholine (DOPC), N-methyl-1,2-dioleoyl-sn-glycero-3-phosphoethanolamine (methyl-DOPE), N,N-dimethyl-1,2-dioleoyl-sn-glycero-3-phosphoethanolamine (dimethyl-DOPE), 1,2-dioleoyl-sn-glycerol-3-phosphoethanolamine (DOPE), 1,2-dimyristoyl-sn-glycero-3-phosphoethanolamine (DMPE), 1,2-dioleoyl-sn-glycerol-3-phosphoethanolamine (dimethyl-DOPE), 1,2-dioleoyl-sn-glycerol-3-phosphoethanolamine (DOPS), 1,2-dioleoyl-sn-glycerol-3-phosphoethanolamine (DOPG), and dioleoylglycerol (DOG) were from Avanti Polar lipids, Inc., Birmingham, AL. Polyclonal antibodies raised against purified His-tagged SecE and SecG against a synthetic peptide corresponding to a SecE domain were obtained as described previously (39).

Purification of Delipidated SecYEG and SecYE Complexes—IMVs overexpressing His-tagged SecYEG (38) or pET720 (E. coli SecYE) (this study) as described (39). IMVs (60 mg of protein) were solubilized on ice for 30 min at 1 mg/ml in 10 mM Tris-Cl, pH 8.0, and 50 mM KCl in 0.03% (w/v) DDM. Non-solubilized proteins and aggregates were removed by centrifugation (30 min at 40,000 × g at 4 °C), and the supernatant was loaded onto a DEAE column (volume 60 ml) (Whatman, DE52) equilibrated with buffer A supplemented with 0.03% (w/v) DDM. The column was washed with 2 volumes of equilibration buffer, and proteins were eluted with a linear gradient of 0–300 mM KCl in 3 volumes of the same buffer, except the last 300 mM KCl and the last 50 ml of buffer A were incubated for 15 min at room temperature, and the activity of the reconstituted translocase was as active as complex purified from OG-solubilized membranes in the presence of added phospholipids (Table 1). The final amount of endogenous phospholipid present in the DDM-solubilized SecYEG complex was below the detection limit, which corresponds to less than 3 mol of phospholipid per mol of SecYEG complex, assuming an equimolar stoichiometry of phospholipids mixed in desired ratios in chloroform solution, dried under vacuum, washed with ethanol, and dried again. Next, the lipid film was slowly (1–2 h) hydrated at a final concentration of 10 mg/ml by incubation at room temperature under a nitrogen atmosphere in a buffer containing 10 mM Tris-Cl, pH 8.0, and 1 mM dithiothreitol. The buffer was then flushed with nitrogen, and the sample was allowed to equilibrate for 5 min on ice, and diluted rapidly with 8 ml of 50 mM Tris-Cl, pH 8.0, and 50 mM KCl. After 5 min, proteoliposomes were collected by centrifugation (30 min, Ti-70, 200,000 × g at 4 °C) and resuspended in 100 μl of 50 mM Tris-Cl, pH 8.0, and 50 mM KCl. Proteoliposomes were analyzed for the amount of incorporation of the DC Protein assay (Bio-Rad), by 15% SDS-PAGE stained with Coomassie Brilliant Blue, and by immunoblotting on polyvinylidene difluoride membranes using antibodies directed against SecY, SecE, and SecG. Reconstituted SecYEG(PE) proteoliposomes were frozen and stored in liquid nitrogen. Before use, samples were thawed at 37 °C and sonicated 3 times for 10 s in a bath sonicator. Control experiments showed that the SecA translocation ATPase activity after rapid dilution was linear with the amount of SecYEG added at lipid-to-protein ratios (w/w) above 10.

To reconstitute the B. subtilis SecYEG into liposomes of different lipid compositions, IMVs were isolated from E. coli. SF100 cells transformed with pET822 that directs the functional overexpression of the E. subtilis SecYEG complex (45). IMVs (100 μl; 0.1 mg of protein) were solubilized in 1.25% (w/v) OG in buffer A as described (39). After 30 min incubation on ice, the non-solubilized material was removed by centrifugation (TLA100.4, 30 min, 180,000 × g at 4 °C). The supernatant fraction (100 μl) was subsequently mixed with 20 μl of lipids (10 mg/ml), rapidly diluted, and isolated as described above. In control experiments, IMVs were used derived from E. coli SF100 cells transformed with pET810 that overproduce E. coli SecYEG (38).

Fusion of SecYEG Proteoliposomes—SecYEG (proteoliposomes (100 μl; containing 0, 10, or 2 μg of SecYEG protein and 0.2 mg of phospholipids) were mixed with an equal volume and amount of (proteoliposomes of different lipid composition. Samples were quickly frozen in liquid nitrogen and slowly thawed on ice. This procedure was repeated three times. Before use in activity assays, samples were sonicated 3 times for 10 s in a bath sonicator.

Translocation Assays—Translocation assays were performed in a final volume of 50 μl consisting of 50 mM HEPES-KOH, pH 7.6, 30 mM KCl, 5 mM Mg(Ac)$_2$, 2 mM ATP, 10 mM creatine phosphate, 0.5 μg of creatine kinase, 25 μg of bovine serum albumin, 1.6 μg of SecB, 1 μg of purified E. coli or B. subtilis SecA, and proteoliposomes containing 6.5 μg of SecYEG or SecYE complex. Samples were preincubated for 10 min at 37 °C, and the translocation reaction was started by the addition of 1 μl of $^{32}$P-labeled pro-OmpA or pre-PhoB (0.2 mg/ml in 6x urea, 50 mM Tris-Cl, pH 7.5). After 10 min, reactions were terminated by chilling on ice. Samples were treated with proteinase K (0.1 mg/ml) for 15 min, precipitated with ice-cold 10% (w/v) trichloroacetic acid, acetone-washed, and analyzed by SDS-PAGE on 10% (pre-PhoB) and 12% (pro-OmpA) polyacrylamide gels.

Other Analytical Techniques—Translocation ATPase activity of urea-treated IMVs or SecYEG proteoliposomes was measured with pro-OmpA as substrate as described (46). Protein concentration was determined with the DC Protein assay (Bio-Rad). Phospholipid phosphorus was assayed after heat destruction of chloroform/methanol-extracted phospholipids using the method of Rouser et al. (47).

RESULTS

Purification and Delipidation of SecYEG Complex—For the purification of the OG-solubilized SecYEG complex, the presence of at least 0.2 mg/ml E. coli phospholipids is essential to retain activity (5, 12, 29). To assess the phospholipid dependence of the translocase in a systematic manner, it is necessary to deplete the purified SecYEG complex of endogenous lipids. When OG was replaced with DDM, the addition of phospholipids was no longer required to purify the SecYEG complex in a functional state. DDM-solubilized SecYEG complex was as active as complex purified from OG-solubilized membranes in the presence of added phospholipids (Table 1). The final amount of endogenous phospholipid present in the DDM-solubilized SecYEG complex was below the detection limit, which corresponds to less than 3 mol of phospholipid per mol of SecYEG complex, assuming an equimolar stoichiometry of phospholipids.
of the subunits. In contrast, the OG-purified enzyme retained about 200 mol of phospholipid per mol of SecYEG. These data demonstrate that the SecYEG complex can be purified in a delipidated and functional state with the detergent DDM.

Reconstitution of SecYEG and Lipid Dependence—Delipidated SecYEG complex was used to examine the phospholipid requirement of protein translocation. The phospholipid composition of the E. coli inner membrane corresponds to about 70 mol % of the non-bilayer lipid PE and 30 mol % of the acidic PG and CL. To mimic the native polar head group composition, the SecYEG complex was reconstituted into liposomes composed of 70 mol % of DOPE and 30 mol % of DOPG. The amount of DOPC was gradually replaced by DOPG to analyze the requirement for anionic phospholipids. Total amounts of SecY, SecE, and SecG reconstituted in the liposomes were determined by Western blotting and were equal for each of the proteoliposomes (data not shown). Proteoliposomes were supplemented with SecA and analyzed for the SecA ATPase activity in the absence and presence of the precursor pro-OmpA (Fig. 1B) and for the translocation of chemical amounts of 125I-labeled pro-OmpA (Fig. 1A). The activity was determined after 10 min of incubation, and within this time interval, the amount of translocated pro-OmpA increased linearly in time (data not shown). SecYEG was completely inactive when reconstituted into DOPC:DOPE (30:70, molar ratio) (Fig. 1, A and B) but became activated when the DOPC was replaced for DOPG or DOPS (Fig. 1A). A similar observation was made for the endogenous and pro-OmpA-stimulated SecA ATPase activity (Fig. 1B). The requirement for the other main lipid constituent of the E. coli inner membrane, PE, was determined by reconstitution of the SecYEG complex into liposomes composed of DOPG:DOPC (30:70, molar ratio) whereby the DOPC was gradually replaced for DOPE. Unlike DOPG, DOPE appears not to be essential for protein translocation activity. A low but significant translocation (Fig. 1C) and translocation ATPase (Fig. 1D) activity was observed with the SecYEG complex reconstituted in the absence of DOPE. However, DOPE markedly stimulated the activity of SecYEG to 3-fold. These results show that anionic phos-

TABLE I

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| Detergent        | Lipid added | Protein content | Phospholipid content | Specific SecA translocation ATPase activitya |
|------------------|-------------|-----------------|----------------------|-------------------------------------------|
|                  | mg/ml       | µmol phosphate/ml| nmol/mg of protein·min |
| Octyl glucoside  | 0.2         | 0.2             | 0.3                  | 625 ± 50 ND                               |
| Dodecyl maltoside| 0           | 0.2             | ND                   | ND                                        |

a SecA translocation ATPase was measured in the presence of pro-OmpA using SecYEG proteoliposomes reconstituted into E. coli phospholipids.

b ND, not detectable.
Phospholipids are essential for the reconstitution of a functional translocase using purified and delipidated SecYEG, whereas PE is stimulatory.

**Phospholipids Reversibly Influence Translocase Activity**—To exclude the possibility that the observed phospholipid requirement is due to a difference in reconstitution efficiency, the SecYEG complex was first reconstituted into proteoliposomes composed of a lipid mixture that does not support activity. Subsequently, other lipids were introduced into these inactive proteoliposomes by freeze-thawed induced fusion with liposomes of different phospholipid composition. SecYEG proteoliposomes composed of DOPC were essentially inactive (2% activity) for the pro-OmpA-stimulated SecA ATPase activity (Fig. 2). Activity could, however, be restored by fusion of these proteoliposomes with DOPG:DOPC:DOPE (30:20:50, molar ratio) liposomes yielding a final lipid composition of DOPG:DOPC:DOPE (15:60:25, molar ratio). The activity of these fused proteoliposomes was about 65% of a control in which the SecYEG complex was reconstituted directly into this lipid mixture (Fig. 2). In another experiment, SecYEG was reconstituted with DOPG:DOPC (30:70, molar ratio) yielding proteoliposomes that support a low SecA translocation ATPase activity. Re-introduction of DOPE into these proteoliposomes by fusion with DOPC:DOPE (50:50, molar ratio) liposomes, again yielding a final lipid composition of DOPG:DOPC:DOPE (15:60:25, molar ratio), resulted in an increase of the activity to about 60% of the control. These results demonstrate that the activity of the reconstituted SecYEG complex can be reversibly modulated by the bulk phospholipid composition.

**Non-bilayer Lipids Stimulate Protein Translocation**—To examine further the effect of the non-bilayer lipid PE on protein translocation, the head group and acyl chain properties were varied. In a lipid mixture of DOPC:DOPG (70:30, molar ratio), the DOPC was gradually replaced for either DOPE, methyl-DOPE, or dimethyl-DOPE. Although introduction of DOPE significantly stimulated the pro-OmpA translocation activity (Fig. 3A) and SecA translocation ATPase (Fig. 3B) of SecYEG proteoliposomes, this effect was largely abolished when the DOPE was mono- or dimethylated. When the DOPE was exchanged for dimyristoyl-PE (DMPE), a PE derivative with a shorter hydrophobic acyl chain, hardly any stimulation of the activity was observed (Fig. 3). Since the methylated forms of PE and DMPE are all bilayer lipids (31, 48), it appears that the stimulatory effect of DOPE is indeed due to its ability to form non-bilayer structures. To confirm this hypothesis further, the effect of dioleylglycerol (DOG), a lipid with strong non-bilayer forming properties, was examined. DOG markedly stimulated the activity of the SecYEG complex to the same extent as DOPE (Fig. 3). Taken together these data demonstrate that PE affects protein translocation by its ability to adopt a non-bilayer conformation.

**The Lipid Requirement of the SecYE Complex Is Not Affected by SecG**—To determine if the presence of SecG influences the lipid requirement of translocation, the SecYE complex was purified from an overexpressing strain and reconstituted into various lipid mixtures. Although some endogenous SecG co-purified with SecYE, Western blotting demonstrated that the amount of SecG in the SecYE proteoliposomes was at least 25-fold lower than in the SecYEG proteoliposomes (data not shown). The absence of SecG resulted in a dramatic reduction of the activity of the SecYE complex. However, the requirement for anionic phospholipids (DOPG) and non-bilayer lipids (DOPE) (Fig. 4) as observed with the SecYE complex was indistinguishable from that found for SecYEG. This indicates that there is no direct mechanistic relation between the SecG function and the activating effect of lipids.

**Lipid Requirement of the B. subtilis SecYEG Complex**—To determine if the lipid requirement found for the *E. coli* SecYEG complex extends to other bacterial species, the lipid specificity of the *B. subtilis* SecYEG complex was determined. Unlike *E. coli*, anionic phospholipids are the major constituents of the *B. subtilis* cytoplasmic membrane, i.e. 70% PG, 4% CL, and 12% PE (49). To obtain functional *B. subtilis* translocase, the SecY, SecE, and SecG proteins were overproduced in *E. coli* (45).
IMVs derived from cells overexpressing either the *B. subtilis* or *E. coli* SecYEG were solubilized in OG without the addition of exogenous lipids and directly reconstituted into a 20-fold excess of synthetic lipids by rapid dilution. A mixture of DOPG:DOPE (70:30, molar ratio) was used to mimic the lipid composition of *B. subtilis*. Either the DOPG or DOPE was replaced for DOPC to analyze the requirement for anionic and non-bilayer lipids, respectively. The translocation activity of the *B. subtilis* SecYEG complex was assayed in the presence of purified *B. subtilis* SecA and 125I-labeled pre-PhoB, a *Bacillus*-specific pre-cursor (44, 45). 125I-Pro-OmpA was used with the *E. coli* SecYEG proteoliposomes. Translocation of pro-OmpA by the *E. coli* SecYEG proteoliposomes again showed the strict requirement for PG and stimulation by PE (Fig. 5). The presence of 25 mol % of PG and 30 mol % of PE in the *E. coli* SecYEG proteoliposomes was already sufficient to saturate the activity (see also Fig. 1, C and D). Therefore, it seems that the *E. coli* SecYEG is most active in a synthetic lipid mixture that resembles the polar head group composition of the *E. coli* inner membrane. Remarkably, translocation of pre-PhoB by *B. subtilis* SecYEG proteoliposomes appeared much more critical with respect to the lipid composition. Maximal translocation activity required a very high DOPG concentration (Fig. 5) and was optimal in a lipid mixture corresponding closely to the polar head group composition of the *B. subtilis* membrane, i.e. DOPG:DOPE (70:30, molar ratio) (Fig. 5). The data suggest that the requirement for anionic and non-bilayer lipids for preprotein translocation is a general feature of bacterial translocase complexes and further show that in *B. subtilis* non-bilayer lipids are essential for activity.

**DISCUSSION**

The *E. coli* preprotein translocase depends for its activity on specific classes of phospholipids. Previous *in vivo* and *in vitro* studies have shown that anionic phospholipids are essential for activity, whereas non-bilayer lipids are stimulatory (for review see Ref. 51). A systematic study of the phospholipid requirement of the purified translocase has not been reported despite the fact that the lipid composition of the reconstituted proteoliposomes can be manipulated in a convenient and systematic manner. We now show that functional reconstitution of the purified, delipidated *E. coli* SecYEG requires anionic phospho-
lipids, whereas non-bilayer lipids stimulate translocation. These results confirm the studies performed in the crude membrane system but, in addition, extend these observations to the B. subtilis SecYEG. Remarkably, the activity of the B. subtilis SecYEG strictly depends on the presence of non-bilayer lipids. For optimal activity, anionic and non-bilayer lipids need to present at amounts corresponding to the phospholipid composition of the native inner membrane.

In order to analyze the phospholipid requirement of the purified translocase, it is desirable to first delipidate the enzyme and subsequently restore its activity by reconstitution in liposomes with a defined phospholipid composition. The SecYEG complex has been purified from IMVs after solubilization with OG. However, to obtain a functional SecYEG complex it is necessary to include phospholipids in the buffers used during the purification (Refs. 5, 12, and 23 and this study). Delipidation of OG-purified SecYEG leads to the irreversible inactivation of the enzyme (Table 1). We now show that SecYEG can be purified in a delipidated and functional form when DDM is used as a detergent instead of OG. OG is a detergent with a rather short acyl chain of only eight carbon moieties. The acyl chain of DDM is longer and thus may more closely resemble the interaction of the phospholipid acyl chain with the SecYEG complex present in detergent micelles. In this respect, it was previously noted that OG-solubilized SecYEG is thermolabile (39, 51, 52). DDM-purified SecYEG complex is less susceptible to such thermal inactivation. The DDM-purified SecYEG complex, like the OG-purified, lipid-supplemented SecYEG complex (39), supports the high affinity binding of pre-OmpA as indicated.

FIG. 5. Comparison of the lipid dependence of B. subtilis and E. coli SecYEG-mediated preprotein translocation. IMVs containing highly over-produced B. subtilis or E. coli SecYEG complex were solubilized in octyl glucoside and reconstituted in proteoliposomes with the indicated lipid compositions. Proteoliposomes were supplemented with purified B. subtilis or E. coli SecA protein, and analyzed for the ATP-dependent translocation of 125I-labeled pre-PhoB or pro-OmpA as indicated.

Although the purification method in the absence of phospholipids was developed to be able to examine the effects of small quantities of phospholipids on the activity of the translocase, the data show that high concentrations of anionic and non-bilayer lipids are needed to saturate the translocation activity. It thus appears that phospholipids act on protein translocation in a more global sense. The E. coli SecYEG complex is maximally active in a mixture of 30% DOPG and 70% DOPE. As far as the ratio between anionic and non-bilayer lipids concerns, this mixture “more or less” corresponds to the lipid composition of the E. coli inner membrane. The activity in the optimal synthetic mixture is about 75% of that found with natural E. coli phospholipids, showing that substantial activity can be recovered with the defined system. Reconstituted B. subtilis SecYEG complex is maximally active in 70% DOPG and 30% DOPE, i.e. at a ratio that closely matches that of anionic to non-bilayer lipids in the native B. subtilis inner membrane. It is remarkable that both systems differ in their quantitative lipid requirement and are most active at their physiological lipid conditions. In this respect, the lipid requirement of the E. coli SecYEG complex did not change significantly when pre-PhoB translocation was assayed instead of pro-OmpA.

Anionic phospholipids have been shown to be able to fulfill at least a dual role, i.e. they promote SecA membrane binding and insertion and stimulate the interaction of the signal sequence of preproteins and the membrane (50). These lipids may indirectly stimulate targeting of SecA to the SecYEG complex, for instance by promoting the low affinity membrane binding of SecA. This may result in a membrane-bound pool of SecA protein that could have a kinetic advantage relative to the cytosolic pool to associate with SecYEG complexes that have accomplished a translocation reaction. Experimental evidence for such mechanism is, however, difficult to obtain. Another possible role for anionic phospholipids may be found in the putative assembly of the SecYEG complex into larger functional oligomers. The collective cold-sensitive secretion defect of the secG null and secAcsR11 strain (28) and many other Sec mutants may be found in a compromised channel assembly activity.

The reconstitution studies with the purified SecYEG complex provide compelling evidence that non-bilayer lipids stimulate translocation. The activating effect of DOPE can be mimicked with dioleoylglycerol, a lipid that like DOPE adopts a non-bilayer conformation, whereas bilayer-forming PE derivatives fail to stimulate translocation. These data provide strong evidence that the lipid shape is the major factor for activation rather than the amino group of PE. In contrast to the E. coli SecYEG, the activity of the B. subtilis SecYEG strictly required the presence of non-bilayer lipids. In contrast, the amino group of PE is needed for the functional reconstitution of the lactose permease of E. coli (55, 56) and leucine permease of Lactococ-
lipid requirement of E. coli preprotein translocase.

Acknowledgments—We thank Jeanine de Keyzer, Erik Manting, Gert Moll, and Andreas Veenendaal for technical assistance and fruitful discussions.

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