Characterization of Purified, Reconstituted Site-directed Cysteine Mutants of the Lactose Permease of Escherichia coli*

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lac permease mutated at each of the 8 cysteinyl residues in the molecule was solubilized from the membrane, purified, and reconstituted into proteoliposomes. The transport activity of proteoliposomes reconstituted with each mutant permease relative to the wild-type is virtually identical with that reported for intact cells and/or right-side-out membrane vesicles. Moreover, a double mutant containing Ser in place of both Cys14 and Cys154 exhibits significant ability to catalyze active lactose transport. The results provide strong confirmation for the contention that cysteinyl residues in lac permease do not play an important role in the transport mechanism. The effect of sulfhydryl oxidant 5-hydroxy-2-methyl-1,4-naphthoquinone on lactose transport in proteoliposomes reconstituted with wild-type or mutant permeases was also investigated, and the results indicate that inactivation is probably due to formation of a covalent adduct with Cys14 and/or Cys154 rather than disulfide formation. Thus, it seems unlikely that sulfhydryl-disulfide interconversion functions to regulate permease activity.

The lactose (lac) permease of Escherichia coli is a hydrophobic polytopic cytoplasmic membrane protein that catalyzes concomitant translocation of β-galactosides and H+ with a stoichiometry of 1:1 (i.e. lactose/H+ cotransport or symport) (cf. Refs. 1–4 for reviews). Encoded by the lacY gene that has been cloned and sequenced, this prototype membrane transport protein has been solubilized from the membrane, purified to homogeneity, reconstituted into proteoliposomes, and demonstrated to be solely responsible for β-galactoside transport. Based on circular dichroic measurements with purified permease and hydropathy analysis of the primary amino acid sequence (5), a secondary structure has been proposed in which the protein consists of 12 hydrophobic domains in α-helical conformation that traverse the membrane in zig-zag fashion, connected by hydrophilic segments with the N and C termini on the cytoplasmic surface (cf. Fig. 1). The general features of the model are consistent with other spectroscopic measurements (6), chemical modification (7), limited proteolysis (8, 9), and immunological studies (10–16). Furthermore, recent studies (17) on an extensive series of lac permease-alkaline phosphatase (lac-Y-phaA) fusion proteins provide strong and exclusive support for the topological predictions of the 12-helix model.

As demonstrated initially by Fox et al. (18) and Kennedy et al. (19), the lac permease is irreversibly inactivated by N-ethylmaleimide, and protection is afforded by substrates such as β-D-galactopyranosyl-β-D-thiogalactopyranoside. On the basis of these findings, it was postulated (18) that a cysteinyl residue is at or near the substrate-binding site of lac permease. Beyreuther et al. (20) later showed that the substrate-protectable residue is Cys154. In addition, the permease is reversibly inactivated by other sulfhydryl reagents such as p-chloromercuribenzenesulfonate or by sulfhydryl oxidants such as diamide (21) or plumbagin (22), and β-D-thiogalactopyranosyl-β-D-thiogalactopyranoside prevents inactivation by these reagents. Because of the latter observations, it was suggested (22, 23) that permease activity may be regulated by a redox process involving sulfhydryl-disulfide interconversion.

Given the importance attributed to sulfhydryl groups in lac permease (cf. Ref. 24 in addition to arguments presented above), particularly Cys154, site-directed mutagenesis was used to replace Cys154 with Gly (25, 26) or Ser (27, 28). Surprisingly, although Cys154 is required for substrate protection against alkylation by N-ethylmaleimide, it is not important for lactose/H+ symport. Subsequently, it was shown (2, 29) that replacement of Cys154 with Gly leads to complete loss of transport activity, although the permease retains the ability to bind the high affinity ligand p-nitrophenyl-β-D-galactopyranoside. Moreover, replacement of Cys154 with Ser or Val yields permease with 10% or 30%, respectively, of wild-type activity, indicating that, although Cys154 is important for full activity, it is not mandatory. Finally, Brooker and Wilson (30) replaced Cys340 with Cys340 with Ser, and Menick et al. (31) replaced Cys340, Cys340 or Cys340 and Cys340 with Ser and observed little or no effect on permease activity. Therefore, out of a total of 8 cysteinyl residues in the permease, only Cys154 appears to be important for transport, and even this residue is not absolutely essential. Although the results from site-directed mutagenesis provide strong evidence that cysteinyl residues in the permease do not play a direct role in the mechanism, alkylation of Cys154

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1 In addition to circular dichroism and laser Raman spectroscopy, Fourier transform infrared studies also show that purified lac permease is largely helical (P. D. Roepe, H. R. Kaback, and K. J. Rothschild, unpublished work).

2 The abbreviations used are: plumbagin, 5-hydroxy-2-methyl-1,4-naphthoquinone; DTT, dithiothreitol.
or replacement of Cys\textsuperscript{14} with Gly leads to inactivation. Furthermore, Cys\textsuperscript{14} and Cys\textsuperscript{15} are both located in putative transmembrane helix V (Fig. 1) and are predicted to be on the same face. Therefore, an attractive explanation for inactivation by sulfhydryl oxidants is catalysis of disulfide formation between these 2 residues. In order to test this possibility, we have solubilized and purified each site-directed Cys mutant in the permease, as well as a double mutant containing Ser in place of both Cys\textsuperscript{14} and Cys\textsuperscript{15}, and studied transport activity and inactivation by plumbagin in reconstituted proteoliposomes. The results indicate that permease inactivation by the sulfhydryl oxidant results from formation of a covalent adduct with either Cys\textsuperscript{14} or Cys\textsuperscript{15} and not from disulfide formation.

EXPERIMENTAL PROCEDURES

Materials

[\textsuperscript{14}C\textsubscript{1}]-Lactose (57 mCi/mmole) was obtained from Amersham (Buckinghamshire, U. K.). All other reagents were obtained as described (32, 33) and were reagent grade.

Methods

Bacterial Strains and Plasmids—\textit{E. coli} T206, \textit{E. coli} T184, and plasmid pGM21 encoding wild-type \textit{lac} Y (34) or \textit{lac} Y with site-directed mutants in single Cys codons (25–31) have been described.

Construction of C148S/C154S Double Mutant—Starting with the M13 mp18 (\textit{lac} Y) DNA containing a Ser codon at position 148 (25), the double mutant containing C148S/C154S was constructed as described by Menick et al. (29). Subsequently, mutations were confirmed by sequencing the entire lac-Y gene in single-stranded M13 mp18 DNA and the region containing the mutations in plasmid pGM21. In both cases, with the exception of the mutations described, the remainder of the lac-Y sequence was identical with that reported by Biechei et al. (35).

Cell Growth—\textit{E. coli} T184 transformed with a given plasmid was grown and induced with isopropyl-1-thio-\textgreek{p}-D-galactopyranoside as described by Teather et al. (34), with the exception that 0.2–0.5% casamino acid was used in place of methionine and threonine in the growth medium (36).

Purification and Reconstitution of lac Permease—Wild-type and mutant lac permeases were solubilized, purified, and reconstituted into proteoliposomes as described (32), except that the C148S and C148G permeases were reconstituted into proteoliposomes by overnight dialysis instead of detergent dilution. In addition, after freeze-thawing, the proteoliposomes were not sonified, but extruded through polycarbonate filters of 100-nm pore size, using an extrusion device (Lipex Biomembranes, Vancouver). This procedure yields proteoliposomes of a relatively uniform size distribution (37). The final preparations contained proteoliposomes at 37.5 mg of \textit{E. coli} phospholipids/ml and 40–90 mg/ml of protein, as indicated, in 50 mM KP, pH 7.5, 1.0 mM DTT.

Transport Assays—Membrane potential (\textgreek{A}\textgamma)-driven lactose transport in proteoliposomes reconstituted with purified permeases was assayed in the presence of a potassium diffusion potential (interior negative) as described (32).

Protein Determinations—Protein was determined by the method of Schaffer and Weissmann (38).

RESULTS

\textgreek{A}\textgamma-Driven Lactose Accumulation—As demonstrated previously (39–42), proteoliposomes reconstituted with purified \textit{lac} permease rapidly accumulate lactose against a concentration gradient in the presence of the potassium diffusion gradient (\textgreek{K}_\text{in} \rightarrow \textgreek{K}_\text{out}) and the ionophore valinomycin (Fig. 2). Under identical conditions, proteoliposomes reconstituted with purified C117S, C148S, C176S, C234S, C333S, or C353S/C355S transport lactose at initial rates to steady-state levels of accumulation that are comparable with proteoliposomes reconstituted with wild-type permease (cf. Table 1 for quantitation of initial rates relative to wild-type). In contrast, proteoliposomes reconstituted with C148G, C154S, or C154V transport lactose at initial rates that are about 15, 10, and 35% respectively, of wild-type proteoliposomes (Fig. 2 and Table I). In each instance, the level of accumulation continues to increase as the time course of the experiments is extended (Fig. 2; data not shown). Finally, proteoliposomes reconstituted with C148S/C154S permease transport at about 5% the rate of wild-type but continue to accumulate the disaccharide with time, whereas proteoliposomes reconstituted with C154G
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Figure 2. Membrane potential (ΔΨ)-driven lactose transport in proteoliposomes reconstituted with purified wild-type (●), C148G (○), C154G (▲), C154S (■), C154V (□) or C148S/C154S (△) lac permease. lac permease was purified from membranes of E. coli T206 or T184 transformed with the appropriate plasmid and reconstituted into proteoliposomes at a protein concentration of 85 µg/ml (wild-type), 75 µg/ml (C148G), 45 µg/ml (C154G), 70 µg/ml (C154S), 65 µg/ml (C154V), or 40 µg/ml (C148S/C154S). Lactose transport was measured as described (32) by diluting 1 µl of proteoliposomes containing 20 µg valinomycin into 200 µl of 50 mM sodium phosphate (pH 7.5) containing 35 µM [1-14C]lactose (57 mCi/mmol). Control experiments (▲) were performed by diluting 1 µl of proteoliposomes into 50 mM potassium phosphate (pH 7.5).

Table I

<table>
<thead>
<tr>
<th>Permease</th>
<th>Initial rate of transport</th>
<th>% of wild-type activity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild-type □</td>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>C117S</td>
<td>75</td>
<td>□</td>
</tr>
<tr>
<td>C148S</td>
<td>85</td>
<td>□</td>
</tr>
<tr>
<td>C148G</td>
<td>15</td>
<td>□</td>
</tr>
<tr>
<td>C154S</td>
<td>10</td>
<td>□</td>
</tr>
<tr>
<td>C154G</td>
<td>&lt;1</td>
<td>□</td>
</tr>
<tr>
<td>C154V</td>
<td>35</td>
<td>□</td>
</tr>
<tr>
<td>C148S/C154S</td>
<td>5</td>
<td>□</td>
</tr>
<tr>
<td>C176S</td>
<td>100</td>
<td>□</td>
</tr>
<tr>
<td>C234S</td>
<td>90</td>
<td>□</td>
</tr>
<tr>
<td>C333S</td>
<td>95</td>
<td>□</td>
</tr>
<tr>
<td>C353S/C355S</td>
<td>61</td>
<td>□</td>
</tr>
</tbody>
</table>

*a Initial rates of transport were calculated from the initial linear portion of transport curves (cf. Fig. 2).

Effect of Plumbagin—Treatment of right-side-out membrane vesicles with the sulfhydryl oxidant plumbagin at a concentration of 0.5 mM leads to complete inactivation of lactose transport in a manner that can be restored by 10 mM DTT (22). Similar phenomena are observed in proteoliposomes reconstituted with purified lac permease. Thus, ΔΨ-driven lactose transport is progressively inactivated by relatively low concentrations of plumbagin, and maximal inactivation is observed at about 0.5 mM (Figs. 3 and 4A). Moreover, when 10 mM DTT is added after treatment of the proteoliposomes with plumbagin, transport is restored to a great extent (Fig. 4A; Table II). Parenthetically, when the proteoliposomes are washed and resuspended in buffer without addition of DTT, lac permease is apparently slightly inactivated by air oxidation, as evidenced by the observation that addition of DTT by itself causes a small stimulation of transport activity. Results of similar experiments carried out on proteoliposomes reconstituted with each mutant permease are summarized in Table II. The initial rate of ΔΨ-driven lactose transport in proteoliposomes reconstituted with C117S, C148S, C154V, C176S, C234S, C333S, or C353S/C355S is essentially completely inactivated by treatment with 0.5 mM plumbagin, and activity is restored by addition of 10 mM DTT in each instance. In contrast, C148G or C154S permease is inactivated by only about 50%, and strikingly, C148S/C154S permease...
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Inhibition of the initial rate of $\Delta\Psi$-driven lactose transport with 0.5 mM plumbagin in proteoliposomes reconstituted with mutant permeases and the restoration of activity by DTT

Initial rates of lactose transport were measured as in Table I. Proteoliposomes were treated with plumbagin and DTT as described in the legend to Fig. 4.

<table>
<thead>
<tr>
<th>Permease</th>
<th>Residual activity after plumbagin inhibition</th>
<th>Restoration of activity by DTT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild-type</td>
<td>%</td>
<td>%</td>
</tr>
<tr>
<td>C117S</td>
<td>0</td>
<td>90</td>
</tr>
<tr>
<td>C148S</td>
<td>8</td>
<td>85</td>
</tr>
<tr>
<td>C148G</td>
<td>0</td>
<td>95</td>
</tr>
<tr>
<td>C154S</td>
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<td>C154V</td>
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<td>80</td>
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<tr>
<td>C148S/C154S</td>
<td>7</td>
<td>100</td>
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<td>C176S</td>
<td>80</td>
<td>100</td>
</tr>
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<td>C234S</td>
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<td>90</td>
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<td>C333S</td>
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<tr>
<td>C353S/C355S</td>
<td>0</td>
<td>78</td>
</tr>
<tr>
<td>C148G/C154S</td>
<td>0</td>
<td>100</td>
</tr>
</tbody>
</table>

In contrast, proteoliposomes reconstituted with wild-type permease or any of the other mutants exhibit no accumulation of lactose after treatment with plumbagin (Fig. 4B).

**DISCUSSION**

The data presented in this paper provide strong confirmation of the conclusion that cysteiny1 residues in lac permease do not play a direct role in the mechanism of lactose/H$^+$ symport (2, 25–31). Proteoliposomes reconstituted with purified permease molecules containing mutations at each individual cysteinyl residue exhibit transport activities relative to wild-type proteoliposomes that are virtually identical with those reported for intact cells and/or right-side-out membrane vesicles. Specifically, purified permease with C117S, C148S, C176S, C234S, C333S, or C353S/C355S exhibits small if any defects in $\Delta\Psi$-driven lactose accumulation, permease with C148G, C154S, or C154V exhibits reduced rates of lactose uptake, and C154G permease is devoid of transport activity. Importantly, the double mutant C148S/C154S exhibits low but significant activity. Clearly, therefore, although Cys$^{148}$ is responsible for inactivation by plumbagin, thereby demonstrating directly that Cys$^{148}$ and/or Cys$^{154}$ are the residues involved.

The data presented in this paper provide strong confirmation of the conclusion that cysteiny1 residues in lac permease do not play a direct role in the mechanism of lactose/H$^+$ symport (2, 25–31). Since modification of either Cys$^{148}$ or Cys$^{154}$ alone can lead to inactivation, it is possible that disulfide formation between either of these residues and another cysteiny1 residue in the permease causes inactivation. Finally, sulfhydryl oxidants may undergo "half-reactions" with cysteinyl residues (21, 43, 44), thereby forming sulfhydryl adducts without catalyzing disulfide formation. Our results favor the last possibility for the following reasons. (i) Plumbagin treatment leads to a marked inactivation of C117S, C176S, C234S, C333S, or C353S/C355S permease, suggesting that none of these cysteiny1 residues is involved in the phenomenon. Consequently, it is unlikely that mixed disulfide formation between Cys$^{148}$ and Cys$^{154}$ and any of the other cysteiny1 residues in the permease is responsible for inactivation. (ii) The double mutant C148S/C154S, which has a low but significant activity, is resistant to inactivation by plumbagin, thereby demonstrating directly that Cys$^{148}$ and/or Cys$^{154}$ are the residues involved. (iii) The single mutants C148S and C154V are essentially completely inactivated by the sulfhydryl oxidant, whereas C148G or C154S permease are partially inactivated. Thus, a single cysteiny1 residue at either of these positions is sufficient for inactivation by plumbagin, a result not expected if disulfide formation between the residues is required for inactivation. In conclusion, since it is unlikely that permease inactivation by sulfhydryl oxidants involves the formation of disulfide bonds, sulfhydryl-disulfide interconversion probably does not have a regulatory function in the activity of this transport protein.

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