Protonation and sugar binding to LacY

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The effect of bulk-phase pH on the apparent affinity (K_{app}^{pp}) of purified wild-type lactose permease (LacY) for sugars was studied. K_{app}^{pp} values were determined by ligand-induced changes in the fluorescence of either of two covalently bound fluorescent reporters positioned away from the sugar-binding site. K_{app}^{pp} for three different galactopyranosides was determined over a pH range from 5.5 to 11. A remarkably high pK_d of ~10.5 was obtained for all sugars. Kinetic data for thiodigalactoside binding measured from pH 6 to 10 show that decreased affinity for sugar at alkaline pH is due specifically to increased reverse rate. A similar effect was also observed with nitrophenylgalactoside by using a direct binding assay. Because affinity for sugar remains constant from pH 5.5 to pH 9.0, it follows that LacY is fully protonated with respect to sugar binding under physiological conditions of pH. The results are consistent with the conclusion that LacY is protonated before sugar binding during lactose/H^+ symport in either direction across the membrane.

lactose permease | membrane transporters | pH titrations | proton translocation | substrate affinity

The lactose permease of Escherichia coli (LacY), a paradigm for the major facilitator superfamily (MFS) of membrane transport proteins (1, 2), transports a galactosidic sugar and an H^+ across the cytoplasmic membrane (3, 4). This coupled translocation mechanism enables LacY to use free energy stored in an electrochemical H^+ gradient (ΔpH) to drive sugar accumulation against a concentration gradient (5–7).

X-ray crystal structures of LacY (8–10), all of which are in an inward-facing conformation, and a wealth of biochemical/biophysical data (7, 11–16) provide evidence for an alternating-access mechanism of action. By this means, upon sugar binding or imposition of a ΔpH (interior negative and/or alkaline), the inward-facing cavity closes with opening of an outward-facing cavity, thereby allowing alternative accessibility of the sugar-binding site, which is at the apex of the cavity in the middle of the molecule, to either side of the membrane. A similar model has been proposed for the glycerol phosphate/phosphate antiporter GlpT, a related MFS protein (17) and the ABC transporter Sav 1866 (18). The alternating-access model involves a global conformational change, which is consistent with the highly dynamic nature of LacY (7, 13, 14, 16, 19–23).

A simple kinetic model for lactose/H^+ symport (Fig. 1) has been proposed based on extensive studies of partial reactions (efflux, equilibrium exchange, and entrance counterflow) catalyzed by LacY and site-directed mutants defective in the symport mechanism (see ref. 5). Wild-type LacY is tightly coupled with respect to sugar and H^+ translocation and does not translocate H^+ without substrate or vice versa. The effect of ambient pH on ΔpH-driven symport is bell-shaped with an optimum at approximately pH 7.5 (24, 25), whereas the partial reactions exhibit different dependences on pH. For example, rates of lactose/H^+ symport down a concentration gradient (efflux) are sigmoidal with respect to pH. The rates are very slow at pH 5.5–7.5 and increase sharply between pH 8.0 and 9.5. Moreover, between pH 5.5 and 7.5, efflux is inhibited 3- to 4-fold by D_2O, whereas equilibrium exchange and ΔpH-driven symport are unaffected, indicating that the deprotonation is probably rate-limiting for downhill lactose/H^+ efflux (26). The rate of equilibrium exchange, a reaction that does not involve H^+ translocation, is independent of pH from pH 5.5 to 10 (26). Furthermore, sugar affinity on either side of the membrane is independent of ΔpH (11). Nevertheless, the observation that affinity for β-D-galactopyranosyl-1-thio-β-D-galactopyranoside (TDG) decreases at the alkaline pH (27), as well as numerous other observations (for review, see refs. 5 and 7), argues that protonation may precede sugar binding.

In the present work, we investigate the effect of pH on substrate binding to LacY. Galactoside binding is readily detected and quantified by using V331C LacY labeled with 2-(7-dimethylamino-4-methylcoumarin-3-yl)maleimide (DACM), which places the reporter away from the sugar-binding site (Fig. 2A), and in this position, the fluorophore is sensitive to long-range conformational changes. This approach has been used to measure affinity for TDG (28), for detection of sugar binding to LacY mutants (29, 30), and for stopped-flow kinetic studies (31). It is noteworthy that binding is measured indirectly, but the fluorescence changes observed are the immediate result of sugar binding. Therefore, reference is made here to apparent affinity (K_{app}^{pp}) only. We used purified V331C LacY labeled with either MIANS or N-(7-dimethylamino-4-methylcoumarin-3-yl)maleimide (DACM) to study the effect of pH on K_{app}^{pp}. The findings demonstrate that sugar binding exhibits an unusually high-affinity on either side of the membrane is independent of ΔpH (11). Nevertheless, the observation that affinity for β-D-galactopyranosyl-1-thio-β-D-galactopyranoside (TDG) decreases at the alkaline pH (27), as well as numerous other observations (for review, see refs. 5 and 7), argues that protonation may precede sugar binding.

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parent pK_a of ≈10.5, thereby indicating that protonation is essential for sugar binding under physiological conditions.

Results

**TDG Binding.** In the presence of galactopyranosides specifically, MIANS-labeled V331C LacY displays a decrease in fluorescence intensity (28, 30–32). Several other fluorophores were tested, and DACM-labeled V331C LacY was also found to be sensitive to sugar binding. Saturating concentrations of TDG alter the fluorescence of DACM- or MIANS-labeled LacY in inverse fashion (Fig. 2). Thus, TDG binding to DACM-labeled V331C LacY causes an increase in DACM fluorescence with a clear blue shift in the spectrum (Fig. 2B), whereas MIANS fluorescence decreases under the same conditions with no shift (Fig. 2C). The inverse response of the fluorophores is likely caused by different interactions with the local environment induced by sugar binding. Although both fluorophores report a sugar-induced conformational change (31), the reporters may sample different local environments because DACM is more hydrophobic relative to negatively charged MIANS. In contrast, large maleimide-based fluorophores such as different Alexa dyes, rhodamine, and Oregon Green, as well as smaller fluorophores like acrylodan and pyrene derivatives or surprisingly another coumarin derivative [7-diethylamino-3-(4'-maleimidylphenyl)-4-methylcoumarin] do not act as reporters for sugar binding with V331C LacY.

TDG affinity measured with DACM- or MIANS-labeled LacY (Fig. 3A: 21.4 and 1.2 μM, respectively) is significantly higher than the affinity of unlabeled LacY, which is characterized by a K_d^{pp} in the low mM range (ref. 32 and supporting information (SI) Fig. S1A and B). Replacement of Val-331 with Cys itself does not affect ΔμHL-driven lactose/H^+ symport (33) or alter affinity for sugar (see Fig. S1C). Thus, the covalently bound fluorophore is responsible for the change in affinity. Analysis of kinetic parameters for TDG binding to DACM- or MIANS-labeled LacY (Fig. 3B and C) demonstrates that k_{reverse} (k_r) for MIANS-labeled LacY is ≈100 times lower than that for DACM-labeled LacY, whereas k_{forward} (k_f) is ≈3 times lower, resulting in a much lower K_d^{pp} for MIANS-labeled LacY relative to DACM-labeled LacY.

**pH Dependence of TDG Binding.** The emission spectra of V331C LacY labeled with either DACM or MIANS do not change significantly from pH 5.5 to 11.0 (see Fig. S2). However, specific sugar binding markedly affects fluorescence and allows titration measurements over a wide pH range (see Figs. S2 and S3). Values for K_d^{pp} estimated from TDG concentration dependence at each pH (see Fig. S4) are plotted as a function of pH in Fig. 4. Remarkably, no significant change in K_d^{pp} is observed for LacY labeled with either fluorophore in the pH range 5.5–9.0, whereas a sharp increase in K_d^{pp} (i.e., decreased affinity) is observed between pH 9.5 and 11.0. The estimated pK_a value is at least 10.5 for protein labeled with either DACM or MIANS (for details, see Fig. S5). Extremely high pH results in a decrease in fluorescence with time. An experiment testing protein stability at alkaline pH shows that MIANS-labeled LacY after preincubation at pH 11 for 12 min has the same protein stability at alkaline pH shows that MIANS-labeled LacY after preincubation at pH 11 for 12 min has the same protein stability at alkaline pH shows that MIANS-labeled LacY after preincubation at pH 11 for 12 min has the same

**Kinetics of Sugar Binding.** Rates of sugar-induced changes in MIANS fluorescence were measured from pH 6 to 10, and the data exhibit a linear dependence of observed rate on TDG concentration from 0.5 to 15.0 μM at all pH values tested (Fig. S4). The slope, which represents k_f, does not depend on pH, but intercept with the y axis (k_i) is clearly dependent on pH. The estimated values of the kinetic parameters from Fig. S4 are plotted as a function of pH in Fig. 5B. Strikingly, k_i is constant as a function of pH, whereas k_f is constant from pH 6.0 to 7.5 and increases sharply above pH 9.0. Hence, the effect of pH on...
apparent affinity for sugar is mediated by an increase in \( k_\text{on} \) at high pH. Data from control experiments with protein preincubated at pH 11 and then shifted back to pH 6.1 are also presented (open symbols) and practically indistinguishable from the data for untreated LacY. Furthermore, \( K_d^{\text{app}} \) values determined from kinetic data (\( K_d^{\text{app}} = k_d/k_r \)) are practically identical to those obtained from the titration studies and can be superimposed when plotted as a function of pH (see Fig. S6).

A similar effect of pH on the kinetic parameters was observed with 4-nitrophenyl-\( \alpha \)-D-galactopyranoside (NPG) by using a direct binding method based on Trp→NPG FRET (31). Rates of binding to unlabeled LacY were measured by stopped-flow at pH 5.5, 7.5, and 9.5. Kinetic parameters estimated from the NPG concentration dependence of \( k_\text{obs} \) are presented in Table 1. The calculated \( k_\text{on} \) and \( k_\text{off} \) values are at least 3 orders of magnitude larger, but the effect of pH is very similar to that observed with TDG (Fig. 5). Alkaline pH decreases the affinity of LacY for NPG by increasing \( k_\text{off} \) with little effect on \( k_\text{on} \). Unfortunately, it is not possible to measure rapid kinetics of NPG binding at higher pH accurately because of increased rates and decreased amplitudes of the fluorescence changes.

Fig. 3. TDG binding to DACM-labeled or MIANS-labeled V331C LacY. (A) Dependence of fluorescence on TDG concentration as percentage of initial emission level before TDG addition for each fluorophore. Titration traces were recorded as a function of time at 0.4 \( \mu \)M protein with excitation and emission wavelengths of 397 and 440 nm for DACM-labeled or 330 and 415 nm for MIANS-labeled LacY, respectively. Solid lines are hyperbolic fits to the data with estimated \( K_d^{\text{app}} \) values of 21.4 ± 0.4 and 1.2 ± 0.1 \( \mu \)M for DACM- and MIANS-labeled LacY, respectively. ○, DACM; ▽, MIANS. (B) Kinetics of TDG binding to DACM-labeled LacY. Concentration dependence of the rate (\( k_\text{obs} \)) of the fluorescence changes was measured at 0.4 \( \mu \)M protein in 50 mM NaPi, (pH 7.5), 0.02% DDM. Data were collected after rapid mixing of equal volumes of TDG and labeled protein by stopped-flow. Final concentrations of TDG are 0.05 to 4 mM and from time traces at TDG concentrations from 0.45 to 15 \( \mu \)M. Estimated parameters are \( k_\text{on} = 0.011 \pm 0.001 \text{s}^{-1} \); \( k_\text{off} = 10.81 \pm 0.03 \text{mM}^{-1} \text{s}^{-1} \); and \( K_d^{\text{app}} = 1.0 \pm 0.1 \text{\mu M} \).

![Fig. 3](image)

Fig. 4. Effect of pH on \( K_d^{\text{app}} \) for TDG with DACM-labeled or MIANS-labeled V331C LacY. Time traces from TDG titration experiments were recorded at indicated ambient pH values with excitation and emission wavelengths at 397 and 440 nm for DACM- or 330 and 415 nm for MIANS-labeled LacY, respectively. Data collection and fitting details were as described in Fig. 3A (see also Figs. S3–S5). \( K_d^{\text{app}} \) values estimated from titrations at each pH were plotted versus \( pH \) concentration and fitted with a hyperbolic equation (see Fig. S5); they are presented as a function of pH. Estimated \( pK_a \) values are 10.9 and 10.7 for DACM-labeled (○) and MIANS-labeled (▽) V331C LacY.

![Fig. 4](image)

Table 1. Kinetic parameters for NPG binding to LacY measured by stopped-flow at different pH using direct sugar binding method based on Trp→NPG FRET (31)

<table>
<thead>
<tr>
<th>pH</th>
<th>( k_\text{off}, \text{s}^{-1} )</th>
<th>( k_\text{on}, \text{M}^{-1} \text{s}^{-1} )</th>
<th>( K_d^{\text{app}}, \text{\mu M} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>5.5</td>
<td>35</td>
<td>0.50 \times 10^6</td>
<td>70</td>
</tr>
<tr>
<td>7.5</td>
<td>18</td>
<td>0.43 \times 10^6</td>
<td>42</td>
</tr>
<tr>
<td>9.5</td>
<td>114</td>
<td>0.50 \times 10^6</td>
<td>228</td>
</tr>
</tbody>
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![Table 1](image)
pH Dependence of Lactose or Melibiose Binding. The increase in sugar affinity observed for MIANS-labeled V331C LacY allows measurement of the effect of pH on binding of natural substrates that bind with low affinity. For example, a $K_d^{app}$ of $\approx 10$ mM was estimated for lactose by protection of Cys-148 against alkylation at pH 7.4 (34). Changes in fluorescence of MIANS-labeled V331C LacY were measured with lactose or melibiose at concentrations ranging from 0.05 to 30 mM at different pH values (see Fig. S7), and $K_d^{app}$ values were plotted as a function of pH (Fig. 6). The data exhibit marked similarity to those obtained for TDG (Fig. 4) despite the difference in affinity for the two disaccharides ($K_d^{app}$ values for lactose and melibiose at neutral pH are $\approx 1.5$ mM and 0.14 mM, respectively). There is no significant change in $K_d^{app}$ for either lactose or melibiose from pH 5.5 to 9.0, and a sharp increase in $K_d^{app}$ (i.e., decreased affinity) is observed between pH 9.5 and 11; $pK_a$ values are estimated to be 10.1 and 10.5 for lactose and melibiose, respectively.

Discussion

In the kinetic model originally proposed for lactose/H$^+$ symport (35) and supported by many findings (for reviews, see refs. 5 and 7), LacY binds a H$^+$ before sugar on either side of the membrane to initiate translocation in either direction across the membrane (Fig. 1). Numerous studies demonstrate that sugar binding at neutral pH induces a conformational change with LacY in the membrane (12, 15) or with purified protein in detergent (13, 14, 28). Furthermore, rates of NPG binding measured by stopped-flow demonstrate that rapid sugar binding is followed by a slower conformational change detected with covalently bound MIANS as a reporter (31). This technique is used here to study the effect of pH on sugar binding to test the assertion that protonation is a prerequisite for sugar binding.

V331C LacY labeled with fluorescent reporter exhibits the same change in $K_d^{app}$ as a function of ambient pH with TDG, lactose, and melibiose (Figs. 4 and 6) as well as unlabeled LacY with NPG (Table 1). In each instance, $K_d^{app}$ is constant from pH 5.5 to 9.0 and increases dramatically from pH 9.5 to 11.0. Kinetic studies provide a clear explanation for the decrease in affinity at alkaline pH. Sugar binds with a forward rate that is independent of pH; however, the reverse rate increases dramatically at high pH (Fig. 5 and Table 1). Both rates are complex kinetic functions that include conformational changes. The $pK_a$ values approximate 10.5 as a lower limit because a complete sigmoidal dependence of $K_d^{app}$ on pH cannot be obtained because the protein denatures too rapidly above pH 11.0. Because the $K_d^{app}$ is constant over the physiological range of pH and increases only above pH 9.0, the data indicate that H$^+$ binding is required for sugar binding. It has also been shown (J. Vazquez-Ibar, S. Schuldiner, and H.R.K., unpublished data) that addition of TDG to a concentrated solution of purified, detergent-solubilized LacY induces no change in pH. In contrast, a positive control with the antiporter EmrE under identical conditions releases 1 H$^+$ per mol of EmrE upon addition of tetraphenylphosphonium (36). Because LacY exhibits no change in pH upon addition of TDG under conditions where EmrE releases an H$^+$ upon substrate binding, the findings are consistent with the argument that LacY is already protonated before sugar binding.

So what becomes protonated? Tyr-236 (helix VII), Glu-269 (helix VIII), Arg-302 (helix IX), His-322 (helix X), and Glu-325 (helix X) are the residues in LacY that play a central role in H$^+$ translocation (for review, see ref. 7). As shown by x-ray crystallography (8–10), these residues are in the C-terminal 6-helix bundle and positioned in the approximate middle of the molecule across from the sugar-binding site. The distances between the OH group of Tyr-236 and Arg-302, and His-322, or Glu-325 are estimated to be 3–4 Å. It is also noteworthy that remarkable conservation of these residues has been found in many other
members of the MFS (37, 38). Although the high pH could be caused by Arg-302 or Tyr-236, it is unlikely that any single residue is responsible for the unique pH dependence of sugar binding. Moreover, several single site-directed mutants in side chains involved in H+ translocation (for example, Y236A, H322A, and R302A) display a shift in pH of ~2 pH units in the acidic direction. In contrast, E325Q LacY maintains high affinity for sugar that remains constant from pH 5.5 to 11.0 (unpublished observations). Therefore, the observed affinity change at alkaline pH results specifically from H+ interaction with a site that directly alters affinity for sugar.

H+ binding and translocation across membranes are thought to involve movement of H+ through pathways that run perpendicular to the plane of the membrane and parallel to the transmembrane helices. However, this situation does not appear to involve movement of H+ through pathways that run perpendicular to the plane of the membrane and parallel to the transmembrane helices. Such a scenario would at least explain the long-standing problem of how sugar gradient-driven H+ translocation can occur in either direction across the membrane with the same residues.

**Experimental Procedures**

**Materials.** Oligonucleotides were synthesized by Integrated DNA Technologies. Restriction enzymes were purchased from New England Biolabs. The QuikChange II kit was from Stratagene. All sugars were purchased from Sigma. Fluorophores were obtained from Molecular Probes/Invitrogen. Talon superflow resin was purchased from BD Clontech. All other materials were of reagent grade obtained from commercial sources.

**Construction of Mutants and LacY Purification.** Construction of mutants and purification of LacY were carried out as described in ref. 14. Purified proteins (10–15 mg/ml) in 50 mM NaPi, 0.01% dodecyl-β-D-maltopyranoside (DDM) (pH 7.5) were frozen in liquid nitrogen and stored at −80 °C until use.

**Labeling with Fluorophores.** Purified V331C LacY (40–50 μM) was labeled with an equimolar concentration of fluorophore in 50 mM NaPi, pH 7.0, 0.02% DDM in the presence of 15 mM TDG to protect Cys-148 against alkylation (13, 14). Reactions were carried out for 15 min at room temperature in the dark. TDG and unreacted label were removed by buffer exchange on an Amicon Ultra-15 concentrator with a 30-kDa cutoff (Millipore). The extent of labeling estimated from absorption spectra was 0.7–0.8 mol/mol of protein. Control experiments with wild-type LacY exhibited essentially no labeling by MIANS under the same conditions.

**Fluorescence Measurements.** Fluorescence was measured at room temperature in steady-state or stopped-flow modes by using an SLM-Aminco 8100 spectrofluorometer modified by OLIS as described in ref. 31. With excitation and emission wavelengths as indicated. Titration traces were recorded for 10–20 min with sequential additions of 5–10 μl of concentrated sugars to 2 ml of protein solution (0.4 μM) in 0.02% DDM and 50 mM buffers with overlapping pH ranges. Buffers used were: citrate-Ph for pH 5.5–7.0; NaPi for pH 7.0–8.0; sodium borate for pH 8.0–9.5; sodium glycine for pH 9.0–10.5, and Na-CAPS for pH 9.5–11.0. Titration data were corrected for dilution and fluorescence drift at high pH. Data analysis was done by using SigmaPlot 10 (Systat Software). Stopped-flow time traces were fitted by using OLIS GlobalWorks software.

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