A Method to Study Complex Enzyme Kinetics Involving Numerical Analysis of Enzymatic Schemes

THE MANNITOL PERMEASE OF ESCHERICHIA COLI AS AN EXAMPLE*

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An analysis of complex kinetic mechanisms is proposed that consists of two steps, (i) building of a kinetic scheme from experimental data other than steady-state kinetics and (ii) numerical simulation and analysis of the kinetics of the proposed scheme in relation to the experimental kinetics. Procedures are introduced to deal with large numbers of enzymatic states and rate constants, and numerical tools are defined to support the analysis of the scheme.

The approach is explored by taking the mannitol permease of Escherichia coli as an example. This enzyme catalyzes both the transport of mannitol across the cytoplasmic membrane and the phosphorylation of mannitol. The challenge is to deduce the transport properties of this dimeric enzyme from the phosphorylation kinetics. It is concluded that (i) the steady-state kinetic behavior is largely consistent with the proposed catalytic cycle of the monomeric subunit, (ii) the kinetics provide no direct support but also do not disprove a coupled translocation of the binding sites on the two monomeric subunits. The approach reveals the need for further experimentation where the implementation of experimental results in the scheme conflict with the experimental kinetics and where specific experimental characteristics do not show up in the simulations of the proposed kinetic scheme.

The steady-state kinetic performance of an enzyme is a reflection of the mechanism by which the enzyme catalyzes a reaction. The kinetic behavior is a global property of a particular mechanism, and, therefore, this mechanism cannot be directly deduced from the kinetics. To learn about the mechanism from the experimental kinetics, the kinetic performance of hypothetical mechanisms are analyzed theoretically and the results are compared with the experimental behavior. Then, the mechanism for which the theory predicts similar behavior as observed experimentally is assigned to the enzyme under study. For relatively simple kinetic mechanisms there will be a unique relation between kinetic scheme and kinetic performance (e.g. "ping-pong," ternary complex, etc).

As kinetic schemes and performances become more complicated two problems arise in the approach outlined above, (i) the analysis of the schemes and (ii) degeneracy of the solution, i.e. more than one kinetic scheme may account for the same kinetic behavior. The theoretical analysis of a

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Scheme I shows schematically the method that consists mainly of two steps, (i) building of a hypothetical kinetic scheme from data other than steady-state kinetics (left) and (ii) numerical analysis of the kinetic performance of this scheme and, at the same time, comparing the results with the experimental steady-state kinetics (right). The method aims at building a kinetic scheme from many scattered pieces of evidence that is consistent with the steady-state kinetics.

Building of the Scheme—The steady-state kinetic performance of an enzyme is not enough to unravel a complex kinetic mechanism. Although the kinetic characteristics are a direct scheme results in a rate equation pertinent to a particular mechanism which analytically describes the turnover rate of the enzyme as a function of the concentrations of the ligands that interact with the enzyme, e.g. substrates, inhibitors, etc. The analysis involves the solution of sets of linear equations, the number of which equals the number of enzymatic states in the kinetic scheme. This becomes very tedious as the number of equations increases and although the King and Altman method (1) provides significant improvement of such analyses, complex mechanisms involving many states of the enzyme remain difficult to treat analytically. This limits the approach to relatively simple enzyme mechanisms and excludes mechanisms that, for instance, involve cooperativity, multiple pathways, or subunit interactions, that lead to rapidly increasing numbers of states of the enzyme.

The complexity of a kinetic mechanism is immediately evident from the experimental kinetic behavior of an enzyme when the relation between the turnover rate and the ligand concentrations cannot be described by single saturation curves. Especially in those cases, an understanding of the kinetics in terms of the underlying kinetic scheme would contribute significantly to the understanding of the functioning of the enzyme. In this paper, a method is described for the study of complex kinetic mechanisms. The degeneracy problem is dealt with by incorporating results from many different types of experiments other than steady-state kinetics into a hypothetical scheme. The kinetics of the hypothetical scheme are simulated and analyzed numerically. In this way the kinetics of enzyme mechanisms involving almost unlimited numbers of states can be analyzed.

The method is demonstrated by taking the mannitol permease of Escherichia coli as an example. This enzyme catalyzes both the transport of mannitol across the cytoplasmic membrane and the phosphorylation of mannitol. The mannitol phosphorylation kinetics of this transport protein can be measured much more accurately than the transport kinetics (2). The challenge will be to deduce the transport properties of this enzyme from the phosphorylation kinetics.

THE METHOD

Scheme I shows schematically the method that consists mainly of two steps, (i) building of a hypothetical kinetic scheme from data other than steady-state kinetics (left) and (ii) numerical analysis of the kinetic performance of this scheme and, at the same time, comparing the results with the experimental steady-state kinetics (right). The method aims at building a kinetic scheme from many scattered pieces of evidence that is consistent with the steady-state kinetics.
fraction \( \phi_i \) may be defined for every transition within the scheme
\[
\phi_i = \frac{\nu_i}{\nu} = \frac{k_j E_j - k_i E_i}{\nu}
\]  
(Eq. 1)

Rate constants \( k_i \) and \( k_j \) are either first order or pseudo-first order. The flux fraction is the tool to determine the major pathway through the scheme by connecting subsequent transitions with high flux fractions. A related parameter is the flux fraction of a subclass of transitions (see below), \( \phi_F \), which sums the flux fractions of the transitions in the subclass.

Balance \( B_{ij} \) is defined as the ratio of the distribution over states \( E_i \) and \( E_j \) in the kinetic steady-state and at thermodynamic equilibrium:
\[
B_{ij} = \frac{E_j/E_i}{k_i/k_j}
\]  
(Eq. 2)

Balance indicates how far a transition is out of thermodynamic equilibrium. When \( B_{ij} = 1 \) the transition is at thermodynamic equilibrium.

The friction coefficient \( f_{ij} \) of the transition between states \( E_i \) and \( E_j \) measures the sensitivity of the turnover rate to a change in the free energy of the transition state between \( E_i \) and \( E_j \) without affecting the free energies of the two states themselves. Friction \( F_{ij} \) in the transition between \( E_i \) and \( E_j \) may be defined as:
\[
F_{ij} = \frac{\nu_i}{\nu_j} = \frac{1}{k_i} \sqrt{K_{ij}}
\]  
(Eq. 3)
in which \( K_{ij} \) is the ratio of \( k_i \) and \( k_j \), i.e. the equilibrium constant. It then follows for the friction coefficient.
\[
f_{ij} = -\frac{d\nu_i}{dF_{ij}/F_{ij}}
\]  
(Eq. 4)

The friction coefficient indicates to which extent the transition between states \( E_i \) and \( E_j \) determines the flux through the enzyme. In the numerical treatment the friction coefficient is determined by calculating the relative increase in the turnover rate of the enzyme when both \( k_i \) and \( k_j \) are increased by 1%. If the increase in the turnover rate would be 1%, the transition between states \( E_i \) and \( E_j \) is completely rate-determining. The friction coefficient is similar to the sensitivity function defined by Ray (3). A more detailed treatment of the friction and the friction coefficient as defined in Equations 3 and 4, respectively, will be published elsewhere.

The software—CACES is a menu-driven, interactive computer program that both simulates and analyzes steady-state kinetics of a predefined enzymatic scheme. The sets of linear equations are solved by Gaussian elimination or matrix inversion which are standard computational techniques (4). The enzyme is defined by the number of states and numerical values for the first and second order rate constants. An experimental condition is defined by numerical values for the ligand concentrations. The rate of change of the state is calculated by a combination of concentration changes from the ligand while keeping other constants and outputting the data in Lineweaver-Burk plots. The flux fraction, balance, and friction coefficient (see above) have been implemented into the software as tools for the analysis of the major pathways, the equilibrium positions, and rate-determining steps in the enzymatic scheme, respectively. Calculated turnover rates may be fitted to known rate equations by a nonlinear fitting procedure using the simplex algorithm (6) followed by analysis of the residuals (7, 8). CACES was written and compiled using the Borland Pascal 7.0 IDE (Borland Inc.) and runs on any AT-type personal computer. The CACES program is available on request.

The Mannitol Permease of E. coli

Mannitol transport in \( E. coli \) is catalyzed by a phosphoenolpyruvate-dependent phosphotransferase system (for reviews, see Refs. 9 and 10). The physiological relevant reaction catalyzed by enzyme II'\textsuperscript{M} the mannitol-specific transport protein of this system, is
\[
\text{Mannitol}_{\text{out}} + \text{P-HP} \xrightarrow{\Pi^{\text{M}}} \text{mannitol-P}_{\text{n}} + \text{HP} 
\]  
(Eq. 5)
In addition to transport, the permease catalyzes phosphorylation of mannitol; mannitol appears as mannitol-P inside the cell. The phosphoryl group donating substrate is a small cytoplasmic protein, termed P-HPr. The enzyme II^\text{mt} molecule consists of three well defined domains (11–14). Domains IIA and IIB protrude into the cytoplasm while domain IIC is situated in the membrane (15). The phosphoryl group of substrate P-HPr is transferred to mannitol by subsequent phosphorylation of phosphoryl group binding sites on domains IIA and IIB (16). Transmembrane domain IIC contains the mannitol binding site (11, 17).

Experimental Data Used to Build the Scheme

1) Both physical and kinetic measurements indicate that enzyme II^\text{mt} is a homodimer (18–26). 2) Dissociated monomeric enzyme II^\text{mt} catalyzes P-HPr-dependent phosphorylation of mannitol (Equation 5) with a different specific activity (28). 3) Phosphorylation of domain IIA does not effect the mannitol binding properties of domain IIC (27). 4) NMR studies indicate that P-HPr does not bind to phosphorylated domain IIA (28). 5) Translocator domain IIC is activated by phosphorylation of enzyme II^\text{mt} (29). 6) Mannitol can bind to unphosphorylated enzyme II^\text{mt} in the absence of P-HPr, and P-HPr can phosphorylate enzyme II^\text{mt} in the absence of mannitol (17, 19, 24). 7) Membrane bound domain IIC constitutes a mannitol translocator that exposes a single binding site alternately to either side of the membrane (17, 30, 31). 8) Phosphorylation and transport of mannitol are, mechanistically, separate steps (29, 30). 9) Phosphorylated enzyme II^\text{mt} catalyzes efflux of mannitol from cells (32, 10). At 4 °C, mannitol binds to membrane vesicles with an inside-out orientation with an affinity constant of 35 nM. Binding to vesicles with a right-side-out orientation indicated a affinity of the periplasmic binding site that was about a factor of 10 lower (30). Binding to purified enzyme II^\text{mt} solubilized in detergent revealed two different binding sites per dimer (24). 11) Enzyme II^\text{mt} catalyzes phosphorylation of cytoplasmic mannitol (31). 12) The fraction of mannitol bound to the cytoplasmic binding site that is phosphorylated in a single turnover upon the addition of P-HPr is 30–40% ("the mannitol-P burst"). This condition will be referred to as the mannitol-P burst efficiency (29, 13). In the unphosphorylated state, the association/dissociation of mannitol at the cytoplasmic side of the membrane is in the order of minutes, whereas this is considerably faster at the periplasmic side (17, 30). 14) The affinity of enzyme II^\text{mt} for the substrate analogue peresitol that cannot be phosphorylated does not change dramatically upon phosphorylation of the enzyme (33). 15) Size exclusion chromatography of the separate domain IIC, and domains IIA and IIB together, demonstrate that the sites of interaction that keep the enzyme II^\text{mt} dimer together are exclusively located in membrane bound domain IIC (34).

Properties of the Kinetic Scheme

A Cooperative Dimer—Enzyme II^\text{mt} is a dimer (see above number 1). Dissociated monomeric enzyme II^\text{mt} possesses all the machinery necessary to catalyze P-HPr-dependent mannitol phosphorylation. Therefore, the turnover of each monomeric subunit within the dimer constitutes a complete catalytic cycle (number 2). In the associated state the two subunits interact; the two cycles are coupled at one or more steps (number 2).

The Monomeric Cycle—Scheme II shows the catalytic cycle of the monomeric subunit. The scheme has been set up to simulate initial rate measurements. The concentration of the two products, mannitol-P and HPr, is zero. Consequently, the enzyme-product complexes and the product association steps may be omitted from the scheme. The internal phosphoryl group transfer between the cytoplasmic domains IIA and IIB is ignored. This implies that the different states of phosphorylation of the monomeric subunit do not effect the kinetic behavior of the enzyme (number 3). P-HPr does not bind to phosphorylated enzyme II^\text{mt} (number 4).

State 9 is the "productive" state. After dissociation of mannitol-P, the binding site on state 1 does not reorient because the translocator is not activated (number 5). The binding of P-HPr and mannitol is random (number 6). States 5–8 built a facilitated diffusion cycle for mannitol (numbers 7 and 8). It is only active in the phosphorylated state (numbers 5 and 9). Mannitol binds to the cycle at either side of the membrane (numbers 7 and 10). The two orientations of the binding site will be referred to as the periplasmic and cytoplasmic binding site.

The scheme provides two different kinetic pathways for mannitol phosphorylation in a noncompartamentalized system. Mannitol may bind to the periplasmic binding site (steps 6 to 7) followed by translocation to the inside (state 8) and phosphoryl group transfer to mannitol (state 9). Alternatively, the translocation step may be omitted by direct binding of mannitol to the cytoplasmic-facing binding sites (states 1, 4, and 5) followed by phosphorylation (number 11). State 5 is a switch between the two pathways that is controlled by the mannitol concentration. At low mannitol concentrations translocation to state 6 will be favored over binding of mannitol (state 8).

The rate constant of transition 8 to 5 is about twice as large as the combined rate constant for the combined transitions 8 to 9 to 1 (number 12). Since the latter has to be faster than the turnover number of the enzyme, in the scheme, dissociation of mannitol from phosphorylated enzyme II^\text{mt} (step 8 to 5) is much faster than dissociation from the unphosphorylated enzyme (step 1 to 2)(number 13). The affinity constants for mannitol of the periplasmic binding site on the phosphorylated and unphosphorylated enzyme are not very different (number 14).

Functional Coupling of the Translocator Domains—The two monomeric catalytic cycles are coupled at the level of the translocation cycles (states 5 through 8, Scheme II) (number
The membrane the binding site on the other subunit faces the
that when the binding site on one subunit faces one side of
the opposite direction. This property together with those assigned
to the catalytic cycle of the monomer leads to three states of
activation of the translocator. The translocator is inactive
when both subunits are not phosphorylated and fully active
when both are phosphorylated. In between, when only one of
the subunits is phosphorylated the translocation activity of
the dimer as a whole is diminished since the activated subunit
has to drag along the binding site on the nonactivated subunit.
The state of activation of the translocator under turnover
conditions will depend on the rates of phosphorylation and
dephosphorylation of the enzyme and, therefore, on the man-
nitol and P-HPr concentrations.

The Two Subunits Are Indistinguishable—The state in
which, for instance, subunit A is phosphorylated and subunit
B has bound mannitol is in the same pool of molecules as the state
in which subunit A has bound mannitol and subunit B is phosphorylated.

Defining the Scheme

The properties assigned to enzyme II* result in a total of
36 states (Fig. 1). There are 168 transitions between these
states with non-zero rate constants. The first step in control-
ing this large number of transitions is to classify them ac-
cording to the type of transition in which they are involved.
Two types of main classes are presented in Table I, those that
involve binding of a ligand (mannitol binding, P-HPr binding,
and mannitol-P binding) and those that do not (enzyme II*-
phosphorylation and translocation). The two rate constants
pertinent to each main class are denoted by two successive
high case characters (e.g. A and B, C and D, etc). Within each
of the main classes, transitions are grouped together with
identical rate constants. These groups make up the subclasses
within a main class. For instance, mannitol binding may be
subclassified into binding to the two orientations of the site
on the unphosphorylated enzyme (cys and per, Table I) and
on the phosphorylated enzyme (cytP and perP). This results
in subclasses A1 to A4 for mannitol dissociation and sub-
classes B1 to B4 for mannitol association. In case the binding
sites on the two subunits interact cooperatively, binding to
the enzyme with the other site occupied results in four addi-
tional subclasses (cytS, cytFS, perS, and perFS). Each step
in the scheme is characterized by a code consisting of a
character for the main class and an integer for the subclass
(e.g. A4). Each code represents a numerical value for the rate
constant(s) of the subclass. The main classes (A and B, C
and D, etc) are defined by the different states of the enzyme
and indicate all possible transitions. The subclassification is
a variable within a certain scheme; it assigns values to the rate
constants.

The complete kinetic scheme is defined by the transition
matrix in Fig. 2. Columns and rows represent the states of the
enzymes and the elements indicate the transitions by the
code of the subclass.

RESULTS

Biphasic Kinetics—Enzyme II* solubilized in detergent
shows biphasic kinetics with respect to the mannitol concen-
tration. The biphasicity manifests itself most strongly at
saturating P-HPr concentrations (Fig. 1 in Ref. 2). A quali-
tative explanation for the biphasicity is provided by the
different possible pathways through the proposed scheme. At
low mannitol concentrations, the high affinity regime, man-
nitol would bind predominantly to the periplasmic binding
site followed by translocation and phosphorylation group transfer
to mannitol (route 1 → 3 → 2 → 5 at high P-HPr, Fig. 1). At
high mannitol concentrations, the low affinity regime, the
translocation step would be short-circuited by direct binding
of mannitol to the cytoplasmic binding site (route 3 → 4 →
7). In order for the latter to show up as a second kinetic phase
the translocation step should be rate-determining in the high
affinity regime.

The set of rate constants listed in Table I termed “free
access” contains all experimental data available on the rate
constants. Rate constants for which no data was available
were chosen such that they do not add new properties to the
scheme. Simulation of the mannitol-dependent kinetics with
this set of rate constants results in a single kinetic phase. The
free access set was arbitrarily constructed to match the high
affinity regime of the experimental mannitol-dependent ki-
netics. However, no set of rate constants that fulfilled all the
conditions set by the experiments could be constructed that
resulted in the biphasic kinetics observed in the experiments.
The failure to show biphasic kinetics is caused by the
condition set by the mannitol-P burst efficiency. The high
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Analysis of Complex Kinetic Schemes

FIG. 1. The 36 states of dimeric enzyme II\textsuperscript{mut}. The top and bottom half of each state represent the two subunits. For further explanation, see the legend of Scheme II.

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<thead>
<tr>
<th>Table I</th>
<th>Three sets of rate constants used in the simulations of the kinetic behavior of the proposed scheme</th>
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<td>Groups and codes are explained in the text. The subclassification of the main classes mannitol binding and translocation relate to each other via thermokinetic balancing. The transitions are defined in Fig. 2. ( k_{\text{off}} ) and ( k_{\text{on}} ) refer to the association and dissociation rate constants, respectively, and ( k_{\text{fw}} ) and ( k_{\text{bw}} ) to the forward and backward rate constants, respectively. ( k_{\text{on}} ) is in ( \mu \text{M}^{-1} \text{s}^{-1} ), the other rate constants are ( \text{s}^{-1} ).</td>
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cytoplasmic binding site is overcome by binding to the periplasmic binding site followed by translocation to form productive state 2 (Scheme IIIB). The high friction coefficient of the translocation step indicates that the kinetic regime is mainly determined by this step. In the low affinity regime, the high concentrations of mannitol compensate for the low accessibility of the cytoplasmic binding site. The flux is through the cytoplasmic binding sites, thereby, bypassing the friction in the translocation step (Fig. 3). In the cooperativity model, product formation follows both in the high and low affinity regime predominantly from binding to the cytoplasmic binding sites (Fig. 3). After binding of mannitol to the periplasmic site (Scheme IIIC), binding to state 3 to form productive state 4 is unfavorable because of the negative cooperativity between the two binding sites. Translocation to state 2 is unfavorable because the translocation equilibrium \( (3 \leftrightarrow 2) \) is far to state 3, a consequence of thermokinetic balancing (Table I). Instead, mannitol dissociates from the periplasmic binding site followed by binding to the cytoplasmic-oriented binding site which is no longer hindered by the negative cooperativity. High concentrations of mannitol compensate for the low affinity of the cytoplasmic site when the periplasmic site is occupied and the flux will be through the cytoplasmic binding sites as well (Fig. 3).

**Mannitol Phosphorylation by Inside-out Membrane Vesicles**—In a compartmentalized system like an inside-out vesicle, mannitol inside the vesicles and mannitol outside the vesicles are different substrates. The steady-state concentration of internal mannitol will be reached very rapidly because of the very small internal volume of these vesicles. The internal concentration of mannitol is set by enzyme \( E^{in} \) at a value that makes the rates of association and dissociation to

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**FIG. 2. Definition of the kinetic scheme; the transition matrix.** The states in Fig. 1 are listed from left to right and from top to bottom. Each element represents a rate constant according to the code given in Table I. Groups A5 through A8 and B5 through B8 are identical to groups A1 through A4 and B1 through B4, respectively, when no cooperativity is assumed between the two mannitol binding sites.
Analysis of Complex Kinetic Schemes

FIG. 3. Flux fractions of the mannitol binding equilibria (A&B) and the high activity translocation (I&J high). In the case of the mannitol binding equilibria, the open left part of the bar and the solid right part of the bar indicate the flux fractions through the periplasmic and cytoplasmic binding steps, respectively. \( \Sigma \phi_{\text{per}^*} \) and \( \Sigma \phi_{\text{cyt}^*} \) indicate the flux fractions through the periplasmic and cytoplasmic binding steps, respectively. Note that \( \Sigma \phi_{\text{per}^*} + \Sigma \phi_{\text{cyt}^*} = 1 \). The P-HPr concentration was 24 \( \mu \text{M} \) and from the internal, periplasmic binding sites equal in the steady-state. Using this criteria, the value of the internal mannitol concentration in the steady-state may be found by an iterative procedure. At a fixed external concentration, the internal concentration is varied until the flux fractions of the periplasmic binding equilibria equal zero. The rate of mannitol phosphorylation is calculated from the distribution under this condition. This rather time consuming procedure can be circumvented by setting the periplasmic association/dissociation rate constants to zero.

It was argued in the accompanying paper (2) that the mannitol-dependent phosphorylation kinetics of enzyme II\textsuperscript{Im} embedded in the membrane of vesicles with an inside-out orientation is characterized by a single low affinity phase. The affinity for mannitol would be in the same order of magnitude as the affinity in the low affinity regime of the kinetics of the solubilized enzyme. The maximal rate would be identical to the maximal rate observed with the solubilized enzyme.

Simulation of the mannitol phosphorylation kinetics catalyzed by inside-out vesicles discriminates strongly between the restricted access model and the cooperativity model (Fig. 5). In the inside-out vesicles the flux is forced through the cytoplasmic binding sites. With the solubilized enzyme in the cooperativity model, the major pathway was through the cytoplasmic binding sites over the whole range of mannitol concentrations and, therefore, the kinetic behavior is roughly the same with the inside-out vesicles (Fig. 5B). In contrast, with the solubilized enzyme in the restricted access model...
**Fig. 4.** Simulations of the bi-phasic kinetics. The top inset shows the residual analysis after a nonlinear fit of the data to the sum of two saturation curves. The bottom inset shows the kinetic characteristics of the low affinity regime that was computed from the difference between the turnover rate and the extrapolated turnover rate from the high affinity regime. The P-HPr concentration was 24 μM.

The major pathway was via the periplasmic binding sites in the high affinity regime, resulting in a much lower rate at the lower mannitol concentrations with the inside-out vesicles. A single phase with a low affinity results (Fig. 5A, +). At the higher mannitol concentrations, the pathway is identical to that observed with the solubilized enzyme, resulting in the same maximal rate. In conclusion, the restricted access model gives a good fit to the experimental behavior of enzyme IImut embedded in the membrane of inside-out vesicles, whereas the cooperativity model fails to predict the experimental behavior.

**Conformational Coupling between the Binding Sites**—The combination of the activation of the translocator by phosphorylation of the enzyme and the coupled movement of the two binding sites resulted in the high and low activity translocation modes depending on the degree of phosphorylation of the enzyme. Qualitatively, it may be argued that this property may explain the drop in the phosphorylation rate in the high affinity regime upon lowering the P-HPr concentration (Figs. 2 and 4 in Ref. 2). In the high affinity regime of the restricted access model the rate is at high P-HPr concentrations determined by the translocation between states 2 and 3 that is of the high activity type. Decreasing the P-HPr concentration and, thereby, the steady-state degree of phosphorylation of the enzyme would shift the rate-determining step from the high to the low activity translocation modes.

Simulating the effects of a decreasing P-HPr concentration within the restricted access model does not show the expected behavior. Lineweaver-Burk plots of the rate of mannitol phosphorylation as a function of the mannitol and P-HPr concentrations show sets of parallel lines in the high affinity regime, as is the case for a classical ping-pong mechanism. Analysis of the friction coefficients in the scheme shows that when the P-HPr concentration is decreased, the friction in the scheme shifts from the high activity translocation step to the steps that lead to the phosphorylation of the enzyme (Fig. 6). The degree of phosphorylation of the enzyme is only decreased when the enzyme phosphorylation steps become rate-determining. The expected increase of the flux fraction through the low activity translocation steps is only limited (2σlowSS
Fig. 6. The friction coefficient of the subclasses in the high affinity regime at 24 μM (A), 1 μM (B), and 0.1 μM P-HPr (C) in the restricted access model. The subclassifications of mannitol binding and translocation is according to Table I.

The key property of the scheme responsible for this behavior is the slow association step of mannitol to the cytoplasmic binding site which makes the friction coefficient in this step high over a wide range of mannitol concentrations. The plateau in each line indicates the region in which the steps leading to the phosphorylation of the enzyme become rate-determining. The experiments indicated a maximal rate of mannitol phosphorylation that was a factor of two higher for enzyme Ilm at below the cmc as compared to above the cmc. The simulations do not show this increase.

DISCUSSION

Sets of Rate Constants—Much of the experimental data that was used to build the hypothetical scheme for enzyme

1 The abbreviation used is: cmc, critical micellar concentration.

+ lowSP + lowS + low = 0.0017 and 0.22 at P-HPr = 24 and 0.1 μM, respectively) but, more importantly, the friction coefficients of these steps are small (Fig. 6). The P-HPr dependence of the phosphorylation rate in the high affinity regime provides no direct support for the property of the coupled translocation of the binding sites.
binding sites were not trapped as mannitol-P upon the addition of P-HPr but seemed to dissociate into the cytoplasm. It was concluded that phosphorylation of the enzyme enhanced the dissociation rate constant of bound mannitol to the same order of magnitude as the overall rate constant for the transfer of the phosphoryl group to mannitol (27). The simulations show that this conclusion is at variance with the biphasic kinetics exhibited by enzyme II\textsuperscript{mut}. It seems to be essential that the experimentally determined slow association/dissociation of mannitol at the cytoplasmic binding site in the unphosphorylated state (17, 30) is not dramatically affected by phosphorylation of the enzyme.

**Monomer or Dimer**—Up to now, no reference has been made to the discussion on the coupled movement of the binding sites or even to the dimeric structure. The restricted access model does not provide direct support for these properties. In fact, implementation of the essential features of the restricted access model listed above in the monomeric cycle described in Scheme II results in a kinetic scheme with the same kinetic behavior. A pronounced structural property like the conformational coupling between two subunits does not show up in the functioning of the enzyme at the kinetic level. It should be noted that the steady-state kinetics does not disprove the dimeric structure. A dimer with coupled binding sites is consistent with the kinetics to the extent described above. The dimer was introduced in the scheme based upon other types of experiments. The failure of the proposed scheme to explain the 2-fold increase in maximal rate after dilution of the solubilized enzyme under the cmc of the detergent might be a manifestation of the dimeric structure of the enzyme. Possibly, the dilution results in two identical subunits, both with the binding site fixed at the cytoplasmic side of the enzyme, resulting in a 2-fold increase of catalytic units.

The Approach—The approach described in this paper results in a catalytic cycle for the monomeric subunit of enzyme II\textsuperscript{mut} that is supported by a lot of experimental data. The coupled movement of the binding sites on the dimer are consistent with the kinetics, but the kinetics provides no proof for the correctness of this property. It may serve as a working hypothesis for future experimentation. The interpretation of the biphasic kinetics of P-HPr burst efficiency needs to be re-evaluated and requires further experimentation. The inability of the scheme to explain the P-HPr-dependent kinetics may reflect properties of the enzyme for which no experimental evidence is available to date. Possibly, the simplifying contraction of the two phosphorylation sites on one subunit may not be justified. Recently, domains IIB and IIC were subcloned from the gene coding for enzyme II\textsuperscript{mut} and expressed as one protein that appears to be fully active in mannitol phosphorylation when phosphorylated domain IIA is used as substrate.\textsuperscript{2} In this assay system the internal phosphoryl group transfer on enzyme II\textsuperscript{mut} is omitted and the kinetic characteristics may show whether the P-HPr dependence relates to different degrees of phosphorylation of the monomeric subunits of enzyme II\textsuperscript{mut}.

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