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**DYNAMIC STABILITY AND THERMODYNAMIC CHARACTERIZATION
IN AN ENZYMATIC REACTION AT THE SINGLE MOLECULE LEVEL**

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Abstract

In this work we study the thermodynamic and dynamic characteristics of an enzymatic reaction at the single molecular level. We investigate how the stability of the enzyme-state stationary probability distribution, the reaction velocity, and its efficiency of energy conversion depend on the system parameters. We employ in this study a recently introduced formalism for performing a multiscale thermodynamic analysis in continuous-time discrete-state stochastic systems.

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I. INTRODUCTION

The vast majority of processes taking pace inside a cell consist of, or at least involve to some extent, chemical reactions. Moreover, most, if not all of the chemical reactions taking place inside a cell are catalyzed by enzymes. Therefore, a profound understanding of enzymes' performance is necessary to better comprehend the processes of life.

Homeostasis, understood as the coordinated physiological processes which maintain most of the steady states in an organism, is regarded as a landmark concept in biology [1]. It can found, to some extent, in all living beings, and allows them to perform in optimal conditions despite ever-changing surroundings and inputs. From a dynamical standpoint, homeostasis implies the existence of a stable steady state [2–4]. Thus, the quality of homeostasis can be measured by the volume of the steady-state basin of attraction in phase space and/or the relaxation time with which the system returns to the steady state after a perturbation. Having a large basin of attraction is important because it allows the system to come back to the steady state even in the face of large deviations. On the other hand, a rapid relaxation time permits the system to quickly recover an optimal state after it is perturbed.

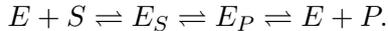
Recent studies on finite-time thermodynamic engines and heat pumps have shown that their stability and their thermodynamic performance are often governed by the same parameters. It has been observed that system stability usually weakens as its thermodynamic properties improve [5–17]. Consequently, these parameters need to be tuned to achieve an optimal trade-off between favorable thermodynamic and dynamic properties. Similar studies on the stretch-reflex regulatory pathway and on a simple Brownian motor have confirmed these findings [18, 19], in agreement with the notion that good design principles are usually shared by both artificial and biological systems [2]. If these results are of general applicability to a wide range of intracellular energy-converting processes, it would mean that the maintenance of the cell homeostatic state entails an expenditure of energy, which has to be taken into consideration to understand how organisms adapt to a constantly changing environment.

In the present work we study the thermodynamic and dynamic characteristics of an enzymatic reaction at the single molecular level. In particular, we investigate how the stability of the enzyme-state stationary probability distribution, the reaction velocity, and its efficiency of energy conversion depend on the system parameters. We employ in this study a recently introduced formalism for performing a multiscale thermodynamic analysis in continuous-time discrete-state stochastic systems.

II. MODELING ENZYMATIC REACTIONS

A simple but comprehensive enough model for an enzymatic reaction consists picturing the enzyme as undergoing the following series of chemical reactions [20]: first, a free enzyme E binds

the substrate S ; then, the bound substrate is converted into the product P to form the enzyme-product complex E_P ; and finally, the product is released leaving the enzyme free to catalyze another reaction. This process can be summarized as follows using the conventional notation for chemical reactions:



Of all these reactions, the conversion of the substrate S into the product P ($E_S \rightleftharpoons E_P$) is typically the rate limiting process. Taking this into account and assuming that the substrate ($[S]$) and product ($[P]$) concentrations remain constant along the catalytic reaction, we can visualize a single enzyme as going through a series of transitions that change the enzyme state cyclically during the catalytic process, as depicted in Figure 1. There, the enzyme state is represented as (i, j) , where index i denotes the i th cycle—that in which the i th product molecule is synthesized—while $j = 1, 2, 3$ respectively correspond to states E_P , $E + P$, and E_S . The assumption that $[S]$ and $[P]$ remain constant allows us to regard k_j^+ and k_j^- ($j = 1, 2, 3$) as pseudo first order reaction rates. Finally, the assumption that $E_S \rightleftharpoons E_P$ is the rate limiting process implies that vertical transitions in the scheme of Figure 1 are much faster processes than those involving changes in index i .

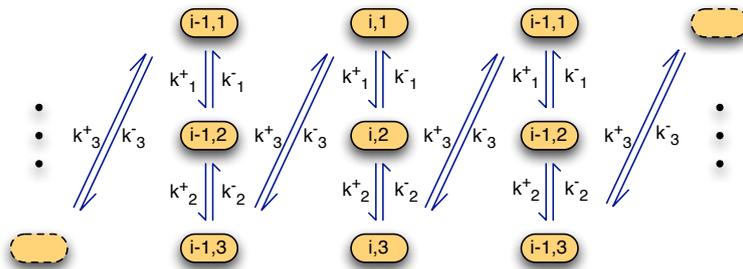


FIG. 1: Schematic representation of the various states available for an enzyme molecule and the transitions rates among them, while the enzyme is catalyzing the synthesis of molecules P . The states are denoted as (i, j) , where $j = 1, 2, 3$ respectively correspond to states E_P , $E + P$, and E_S , while the index i denotes the i th enzyme cycle: that in which the i th product molecule is synthesized and released.

III. PROBABILISTIC DESCRIPTION AND TIME-SCALE SEPARATION

Let us introduce a probabilistic description for an enzyme reaction at the single molecule level and follow the approach in [21, 22] to simplify the model taking advantage of the assume separation of time scales. Let $P(i, j; t)$ denote the probability that the enzyme is in state (i, j) at time t . From the scheme in Figure 1, the chemical master equation for $P(i, j; t)$ consists of the

following set of coupled differential equations:

$$\frac{dP(i, 1; t)}{dt} = k_3^+ P(i-1, 3; t) + k_1^- P(i, 2; t) - (k_3^- + k_1^+) P(i, 1; t), \quad (1)$$

$$\frac{dP(i, 2; t)}{dt} = k_1^+ P(i, 1; t) + k_2^- P(i, 3; t) - (k_1^- + k_2^+) P(i, 2; t), \quad (2)$$

$$\frac{dP(i, 3; t)}{dt} = k_2^+ P(i, 2; t) + k_3^- P(i+1, 1; t) - (k_2^- + k_3^+) P(i, 3; t). \quad (3)$$

The probability that the enzyme is in state (i, \cdot) at time t is given by

$$P(i; t) = \sum_{j=1}^3 P(i, j; t), \quad (4)$$

while it follows from the definition of conditional probability that

$$P(i, j; t) = P(j|i; t)P(i; t). \quad (5)$$

Add Equations (1)-(3) and use (4) and (5) to obtain

$$\begin{aligned} \frac{dP(x; t)}{dt} &= (k_3^+ P(3|i-1; t)) P(i-1; t) + (k_3^- P(1|i+1; t)) P(i+1; t) \\ &\quad - (k_3^+ P(3|i; t) + k_3^- P(1|i; t)) P(i; t). \end{aligned} \quad (6)$$

Differentiate (5) and assume a time-scale separation so that the transitions between states $(i-1, 1)$ and $(i, 3)$ are much slower than all the other (that is, they are the rate limiting steps along the reaction chain). Then,

$$\frac{dP(1|i; t)}{dt} = k_1^- P(2|i; t) + k_1^+ P(1|i; t), \quad (7)$$

$$\frac{dP(2|i; t)}{dt} = k_1^+ P(1|i; t) - k_1^- P(2|i; t) + k_2^- P(3|i; t) - k_2^+ P(2|i; t), \quad (8)$$

$$\frac{dP(3|i; t)}{dt} = k_2^+ P(2|i; t) - k_2^- P(3|i; t). \quad (9)$$

If we invoke once more the separation of time scales to assume that the fast dynamics rapidly reach an equilibrium distribution while slow dynamics have not changed noticeably: $dP(j|i; t)/dt \approx 0$, $j = 1, 2, 3$, we have that

$$P(1|i; t) \simeq P^*(1|i),$$

$$P(2|i; t) \simeq P^*(2|i) = K_1 P^*(1|i),$$

$$P(3|i; t) \simeq P^*(3|i) = K_2 P^*(2|i) = K_1 K_2 P^*(1|i),$$

where $K_j = k_j^+ / k_j^-$, $j = 1, 2, 3$. Finally, the normalization condition ($\sum_{j=1}^3 P^*(j|i) = 1$) implies that

$$P^*(1|i) = \frac{1}{1 + K_1 + K_1 K_2}, \quad (10)$$

$$P^*(2|i) = \frac{K_1}{1 + K_1 + K_1 K_2}, \quad (11)$$

$$P^*(3|i) = \frac{K_1 K_2}{1 + K_1 + K_1 K_2}. \quad (12)$$

Notice that $P^*(j|i)$ ($j = 1, 2, 3$) are all independent of i . Furthermore, it is straightforward to prove that this stationary distribution satisfies the following relations:

$$k_1^+ P^*(1|i) = k_1^- P^*(2|i), \quad \text{and} \quad k_2^+ P^*(2|i) = k_2^- P^*(3|i).$$

Hence, the fast dynamics are in equilibrium only if each one of the underlying chemical reactions is in equilibrium itself. Finally, substitution of (10)-(12) into (6) allows us to conclude that, when time-scale separation is possible and the enzyme states are grouped as sketched in Figure 1, the system dynamics is that of a biased one-dimensional random walk:

$$\frac{dP(x;t)}{dt} = k^+ P(i-1;t) + k^- P(i+1;t) - (k^+ + k^-) P(i;t), \quad (13)$$

where

$$k^+ = \gamma \frac{K_1 K_2 K_3}{1 + K_1 + K_1 K_2}, \quad \text{and} \quad k^- = \gamma \frac{1}{1 + K_1 + K_1 K_2}, \quad (14)$$

while $\gamma = k_3^-$ and $K_3 = k^+ 3/k_3^-$.

From (13), the slow-dynamics stationary distribution obeys

$$P^*(i) = P^* \equiv \text{constant}.$$

Observe that this distribution does not fulfill detailed balance since $k^+ P^*(i-1) - k^- P^*(i) \neq 0$, unless $k^+ = k^-$.

IV. THERMODYNAMIC STATE VARIABLES AND RELAXATION TO THE STATIONARY STATE

Following [22, 23], the enzyme internal energy, entropy, and Helmholtz free energy can be respectively defined as follows:

$$\begin{aligned} U &= -k_B T \sum_{i,j} P(i,j;t) \log P^*(i,j), \\ S &= -k_B \sum_{i,j} P(i,j;t) \log P(i,j;t), \\ F &= U - TS = k_B T \sum_{i,j} P(i,j;t) \log \frac{P(i,j;t)}{P^*(i,j)}. \end{aligned}$$

By substituting (5) into the above equations they can be rewritten as

$$U = -k_B T \sum_i P(i;t) \log P^*(i) - k_B T \sum_i P(i;t) \sum_j P(j|i;t) \log P^*(j|i), \quad (15)$$

$$S = -k_B \sum_i P(i;t) \log P(i;t) - k_B \sum_i P(i;t) \sum_j P(j|i;t) \log P(j|i;t), \quad (16)$$

$$F = k_B T \sum_i P(i;t) \log \frac{P(i;t)}{P^*(i)} + k_B T \sum_i P(i;t) \sum_j P(j|i;t) \log \frac{P(j|i;t)}{P^*(j|i)}. \quad (17)$$

Observe that this way of writing the thermodynamic state variables renders a natural separation of contributions from the fast and slow dynamics.

The first term in the right hand side of Equation (17) is nothing else but the Kullback-Leibler divergence between distributions $P(i; t)$ and $P^*(i)$ and so it is positive defined and only equals zero when the two distributions are identical. Similarly, the sum over j in the second term is the Kullback-Leibler divergence between $P(j|i; t)$ and $P^*(j|i)$, it is positive defined, and only equals zero when $P(j|i; t) = P^*(j|i)$. From these considerations, the value of F can be used as an indicator of how far the system is from the stationary distribution, and $dF/dT \leq 0$ can be understood as the rate of relaxation to the stationary distribution.

After differentiating Equation (17) and imposing the separation of time scales we obtain

$$\frac{dF}{dt} = Q_{hk} - T\sigma, \quad (18)$$

where Q_{hk} is known as the housekeeping heat and is given by

$$\begin{aligned} Q_{hk} = & k_B T \sum_i (P(i; t)k^+ - P(i+1; t)k^-) \log \frac{P^*(i)k^+}{P^*(i+1)k^-} \\ & + k_B T \sum_i P(i; t) \sum_{j=1}^2 \left(P(j|i; t)k_j^+ - P(j+1|i; t)k_j^- \right) \log \frac{P^*(j|i)k_j^+}{P^*(j+1|i)k_j^-}, \end{aligned} \quad (19)$$

while σ is the entropy production rate:

$$\begin{aligned} \sigma = & k_B \sum_i (P(i; t)k^+ - P(i+1; t)k^-) \log \frac{P(i; t)k^+}{P(i+1; t)k^-} \\ & + k_B \sum_i P(i; t) \sum_{j=1}^2 \left(P(j|i; t)k_j^+ - P(j+1|i; t)k_j^- \right) \log \frac{P(j|i; t)k_j^+}{P(j+1|i; t)k_j^-}. \end{aligned} \quad (20)$$

Observe that both Q_{hk} and σ have contributions from the slow (first term on the right hand side) and fast dynamics (second term). However, the fast dynamics contribution to Q_{hk} vanishes because $P^*(j|i)$ complies with detailed balance, and so

$$Q_{hk} = k_B T \sum_i (P(i; t)k^+ - P(i+1; t)k^-) \log \frac{P^*(i)k^+}{P^*(i+1)k^-}. \quad (21)$$

This result is in complete agreement with the interpretation of Q_{hk} as the energy that has to be pumped into the system to drive the stationary state out from equilibrium (detailed balance).

In the present formalism, the enzyme molecule is implicitly assumed to be in equilibrium with a thermal bath. Concomitantly, the proper thermodynamic description is that of the Helmholtz free energy. As seen in Equation (17), $F \geq 0$ and it only equals zero in the steady state. In other words, the value of F can be used as a measure of how distant the system state is from the stationary one, as we have previously asserted. Furthermore, it is not hard to prove from (18), (20), and (21) that $dF/dt \leq 0$, and that $dF/dt = 0$ only when $P(i; t) = P^*(i)$ and $P(j|i; t) = P^*(j|i)$. Therefore, dF/dt can be interpreted as the rate of relaxation to the stationary distribution.

We can see from (18), (20), and (21) that dF/dt can be decomposed into contributions from the slow and fast dynamics:

$$\frac{dF}{dt} = \dot{F}_{\text{slow}} + \dot{F}_{\text{fast}}, \quad (22)$$

where

$$\dot{F}_{\text{slow}} = k_B \sum_i (P(i;t)k^+ - P(i+1;t)k^-) \log \frac{P^*(i)P(j+1|i;t)}{P^*(i+1)P(j|i;t)}, \quad (23)$$

and

$$\dot{F}_{\text{fast}} = k_B \sum_i P(i;t) \sum_{j=1}^2 (P(j|i;t)k_j^+ - P(j+1|i;t)k_j^-) \log \frac{P(j|i;t)k_j^+}{P(j+1|i;t)k_j^-}. \quad (24)$$

The relaxation of the fast dynamics subspace to the corresponding quasi-stationary state, given by the slow-dynamics probability distribution, is determined by \dot{F}_{fast} . According to the assumed separation of time scales, fast dynamics relaxation takes place without any noticeable change in the slow dynamics probability distribution ($P(i;t)$) and, in the long run, it is the slow dynamics which governs the system relaxation to the steady state. That is the system relaxation rate is well approximated by \dot{F}_{slow} .

Under the assumption that the probability distribution $P(i;t)$ is slightly different from the stationary distribution P^* : $P(i;t) = P^* + \epsilon(i;t)$, with $\epsilon(i;t) \ll 1$. Then, it is straightforward to see from (23) that the system relaxation rate in the neighborhood of the stationary distribution (ξ) is proportional to—see equation (14):

$$\xi \propto k^+ - k^- = \gamma \frac{K_1 K_2 K_3 - 1}{1 + K_1 + K_1 K_2}. \quad (25)$$

To better understand the result in Equation (25) let us analyze the significance of parameters K_i ($i = 1, 2, 3$). From their definition, these parameters are nothing else but the association constants of the following reactions, respectively: $E_P \rightleftharpoons E + P$ (K_1), $E + S \rightleftharpoons E_S$ (K_2), and $E_S \rightleftharpoons E_P$ (K_3). On the other hand, a reaction's association rate (K_A) is related to its free energy change (ΔG) by $K_A = \exp(-\Delta G/RT)$, with R the ideal gas constant and T the absolute temperature. These considerations allow us to visualize an enzymatic reaction as a process occurring along an energy profile like the one pictured in Figure 2. In such scheme, the enzyme states correspond to the local minima of the energy profile, and the transition probabilities are determined by the height of the energy barriers.

The global free energy change (ΔG_T) of an enzymatic reaction is given by $\Delta G_T = \Delta G_1 + \Delta G_2 + \Delta G_3 = -RT \log(K_1 K_2 K_3)$. Therefore, since the presence of an enzyme does not change ΔG_T :

$$K_1 K_2 K_3 = \exp\left(-\frac{\Delta G_T}{RT}\right) \equiv \text{constant}.$$

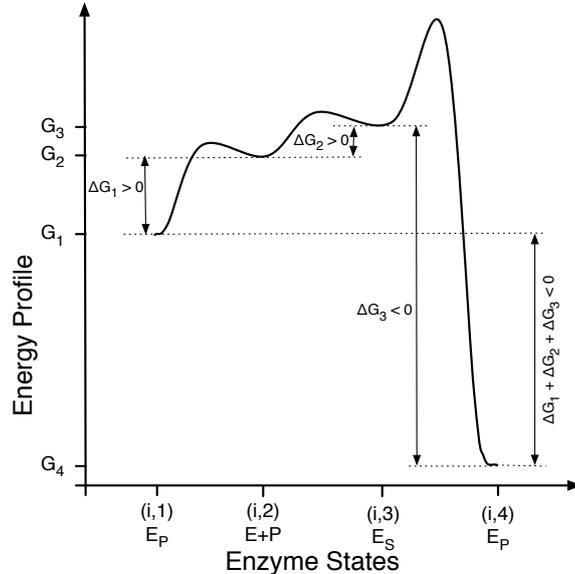


FIG. 2: A cartoon representation of the Gibbs free energy profile for an enzymatic reaction. The minima correspond to the states the enzyme goes through, while the transition probabilities are determined by the height of the energy barriers. The individual free energy changes between adjacent states can be either positive or negative, but the global free energy change has to be negative in order for the reaction to proceed forward.

This restriction further implies that only two of the three K_i constants are independent. Without loss of generality we shall consider that K_1 and K_2 are determined by the nature of the enzyme catalyzing the reaction, while $K_3 = \exp(-\Delta G/RT)/K_1K_2$. With this, Equation (25) can be rewritten as

$$\xi \propto \gamma \frac{\exp(-\Delta G_T/RT) - 1}{1 + K_1 + K_1K_2}. \quad (26)$$

Note that the relaxation rate is a monotonic decreasing function of both K_1 and K_2 . If we further take into consideration that $K_1, K_2 > 0$, it follows that the relaxation rate can be increased by making K_1 and K_2 as close to zero as possible, and therefore by making ΔG_1 and ΔG_2 positive and as large as possible. In particular, the maximum value of ξ is attained when $K_1 = 0$, regardless the value of K_2 . Thus, in order to increase the value of ξ it is more important to increase the value of ΔG_1 than that of ΔG_2 .

On the other hand, we can see from Equation (26) that the relaxation rate is a monotonic decreasing function of ΔG_T . That is, the more energetically favorable the global function is, the more stable the stationary distribution becomes. Similarly, ξ is proportional to $\gamma = -k_3$. Therefore, the system stationary distribution is more strongly stable when the substrate-to-product conversion process is more rapid.

V. REACTION VELOCITY AND EFFICIENCY OF ENERGY CONVERSION

As we have seen, by exploiting the time-scale separation to simplify the reaction scheme, an enzyme can be modeled as a biased one-dimensional random walk with forward and backward transition probabilities k^+ and k^- , respectively—see Equations (13) and (14). Therefore, by taking into consideration that the stationary probability distribution P^* is constant, the reaction velocity can be defined as the forward minus the backward fluxes:

$$\nu = P^*(k^+ - k^-) = \gamma P^* \frac{\exp(-\Delta G_T/RT) - 1}{1 + K_1 + K_1 K_2}. \quad (27)$$

On the other hand, if we consider that a certain amount of energy is consumed during each forward step, and that this energy is wasted during backward steps, the system efficiency in the stationary state can be define as

$$\eta = 1 - \frac{k^-}{k^+} = 1 - \frac{1}{\exp(-\Delta G_T/RT)}. \quad (28)$$

Observe that $\nu > 0$ is a decreasing function of ΔG_T , and that $\nu = 0$ when $\Delta G_T = 0$. In other words $\Delta G_T < 0$ in order to have a positive reaction velocity. The efficiency η is also a decreasing function of ΔG_T , and $\eta = 0$ when $\Delta G_T = 0$. That is, both the reaction velocity and the its efficiency can be increased by making ΔG_T more negative.

Note from equations (26) and (28) that $\eta \propto \xi$. Therefore, the discussion regarding the dependence of ξ on K_1 and K_2 applies as well to η . In particular, we want to emphasize that the reaction velocity can be increased by making K_1 and K_2 as close to zero as possible (and thus by making ΔG_1 and ΔG_2 as large as possible). However, varying K_1 is more important since the maximum velocity can be achieved by setting $K_1 = 0$, regardless the value of K_2 .

Interestingly, the reaction efficiency is independent of K_1 and K_2 . Thus, given that an enzyme does not alter the global free energy change of the reaction it catalyzes, this result implies that a reaction efficiency is the same regardless whether it is catalyzed or not.

VI. CONCLUDING REMARKS

In this work we have investigated the dynamic stability, as well as velocity and efficiency, of an enzymatic reaction at the single molecule level. For this, we followed the ideas of Ge and Qian [23], and used a recently developed formalism for performing multiscale thermodynamic analysis on discrete-state, continuous-time, Maekivian stochastic processes [22]. Our results can be summarized as follows:

1. The dynamic and thermodynamic characteristics associated to the stationary probability distribution are completely determined by the the Gibbs free energy changes of the enzymatic reaction steps: ΔG_1 ($E_P \rightleftharpoons E + P$), ΔG_2 ($E + S \rightleftharpoons E_S$), and ΔG_3 ($E_S \rightleftharpoons E_P$).

2. The energies ΔG_i ($i = 1, 2, 3$) are not all independent because $\sum_{i=1}^3 \Delta G_i = \Delta G_T$, and ΔG_T (the global free energy change) is not modified by the enzyme.
3. The stationary probability distribution is stable and the corresponding relaxation rate (γ) is directly proportional to the global reaction velocity (ν).
4. Both γ and ν are decreasing functions of ΔG_T , and $\gamma, \nu = 0$ when $\Delta G_T = 0$. Thus, the global reaction accelerates and the stationary probability distribution turns more strongly stable as ΔG_T is more negative.
5. Both γ and ν are increasing functions of ΔG_1 and ΔG_2 . The relaxation rate and the reaction velocity achieve their maximum value in the limit $\Delta G_1 \rightarrow \infty$, regardless the value of ΔG_2 . Contrarily, γ and ν increase as ΔG_2 increases and converge to a value that depends on ΔG_1 as $\Delta G_2 \rightarrow \infty$.
6. The efficiency (η) is a function of ΔG_T but is independent of ΔG_1 and ΔG_2 . In particular, η is a decreasing function of ΔG_T , and $\eta = 0$ when $\Delta G_T = 0$. That is, the reaction efficiency increases as ΔG_T becomes more negative.

Although in an enzymatic reaction the global free energy change is predetermined by the substrate, the products, and the nature of their reaction, the actual shape of the energy profile along the reaction coordinate depends on the enzyme structure. Our results demonstrate that although the efficiency is not affected by the shape of the energy profile, the reaction velocity and the strength of the stationary distribution stability can be highly improved by properly shaping this profile. This behavior is contrary to that observed in other systems (like thermal engines) in which variation in some parameters makes the velocity and the efficiency change in opposite directions. In other words, in this case no trade-off between efficiency and reaction velocity (and stability strength) need to be looked for, regarding the energy profile. We believe this may be one of the reasons why it has been possible for evolution to drive the structure of enzymes so the corresponding reaction velocity is increased by several orders of magnitude.

Another feature worth noticing is the fact that the more energetically unfavorable reactions $E_P \rightleftharpoons E + P$ and $E + S \rightleftharpoons E_S$ are, the faster the global enzymatic reaction is. This behavior can be understood by looking at Figure (2). We can see there that large, positive ΔG_1 and ΔG_2 values imply a very negative ΔG_3 . This further means that the backward reaction $E_S \leftarrow E_P$ is much less probable than the forward reaction $E_S \rightarrow E_P$. That is, the strategy to accelerate the global enzymatic reaction seems to be making the rate limiting step almost unidirectional, even though this implies that, in the rapid processes, the backward reactions are more probable than the corresponding forward reactions.

Acknowledgments

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