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Stochastic Ensembles, Conformationally Adaptive Teamwork and Enzymatic Detoxification

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Abstract

It has been appreciated for a long time that enzymes exist as conformational ensembles throughout multiple stages of the reactions they catalyze, but there is renewed interest in the functional implications. The energy landscape that results from conformationlly diverse poteins is a complex surface with an energetic topography in multiple dimensions, even at the transition state(s) leading to product formation, and this represents a new paradigm. Nearly simultaneous with the renewed interest in conformational ensembles, a new paradigm concerning enzyme function has grown, wherein catalytic promiscuity has clear biological advantages in some cases. 'Useful', or biologically functional, promiscuity or the related behavior of 'multifunctionality', can be found in the immune system, enzymatic detoxification, signal transduction, and in the evolution of new function from an existing pool of folded protein scaffolds. Experimental evidence supports the widely held assumption that conformational heterogeneity promotes functional promiscuity. The common link between these co-evolving paradigms is the inherent structural plasticity and conformational dynamics of proteins that, on one hand, leads to complex but evolutionarily selected energy landscapes, and on the other hand promotes functional promiscuity. Here we consider a logical extension of the overlap between these two nascent paradigms: functionally promiscuous and multifunctional enzymes such as detoxification enzymes are expected to have an ensemble landscape with a greater number of states accessible on multiple time scales than substrate specific enzymes. Two attributes of detoxification enzymes become important in the context of conformational ensembles: these enzymes metabolize multiple substrates, often in substrate mixtures, and they can form multiple products from a single substrate. These properties, combined with complex conformational landscapes, lead to the possibility of interesting time-dependent, or emergent, properties. Here we demonstrate these properties with kinetic simulations of non-equilibrium steady state (NESS) behavior resulting from energy landscapes expected for detoxification enzymes. Analogous scenarios with other promiscuous enzymes may be worthy of consideration.

Keywords

Conformational ensemble; hysteresis; transition state; detoxification

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Background

Conformational Dynamics and the New Enzymology

Proteins are inherently flexible, and they exhibit time-dependent fluctuation on a wide range of time scales. Slow times scales of fluctuation naturally imply greater energy barriers between states and more 'rugged' energy landscapes. Rapidly interconverting conformations of ligand-free enzymes are clearly described by many experimental techniques (1 - 3), and there is little doubt that ligand binding alters the energy landscape, and therefore the relative population of states and their rates of interconversion. These atomic level motions are consistent with two limiting cases for ligand-dependent conformational change: a) the classic concept of induced fit and b) the thermodynamically equivalent but kinetically different 'conformational selection' model for molecular recognition (4 - 9). Recently, however, it has become increasingly appreciated that such conformational changes may be sufficiently slow to occur on time scales similar to ligand binding and dissociation, and on time scales relevant to catalytic steps.

Whereas heterogeneity of ground state conformations of free enzymes and ligand-bound enzymes is a 'certainty,' with mature experimental support, the analogous heterogeneity at the level of the catalytically relevant enzymatic transition states has only recently been appreciated. Many recent studies based on computation, NMR, and single molecule methods suggest enzyme hetrerogeneity 'during' transition states associated with a single elementary reaction (9 - 15). These refinements in theory, together with the more detailed experimental measurements, result in an energy landscape for enzyme catalysis as schematized in Figure 1, as already suggested in (13, 14). According to the strict definition from the theory of chemical kinetics, "the" transition state is located at the global energy with the lowest barrier height among all possible paths connecting the reactant to the product. The transition state ensemble perspective, however, articulates the existence of multiple saddle points with nearly equal barrier heights. For the promiscuous enzymes that we highlight here, we refer to transition state ensembles that result from distinct transition states for different substrates, or different products from a single susbtrate, that are well separated by significant energy barriers. For these cases, the heterogeneity of transition states refers to distinct elementary reactions, in the classic sense and we do not imply that there is conformational flux between transition states (no arrows between TS's in Figure 1A, even though Figure 1B suggests that possibility).

The specific case depicted in Figure 1 includes multiple rate limiting catalytic transition states, but conformational equilibria within the ensemble of the unbound enzyme may also include significant barriers. In short there are many populated 'states' for the ligand free enzyme (E1, E2, … En), various substrate complexes (E1S, E2S .. EnS), the transition state ensemble (TS1, TS2 .. TSn) and product complexes (E1P, E2P .. EnP). The extent to which these ground states equilibrate along the conformational axis on the time scale of flux through each transition state (the reaction coordinate) will have profound effects on the time-dependent behavior of the system. In general, if conformational sampling occurs on a time scale comparable with catalytic processes, then each successive round of catalysis for a single enzyme molecule can include participation from different states under certain conditions. The result could be a system that is highly heterogeneous and driven by probability and independent of its history at low turnover (stochastic), with increasingly deterministic behavior as it completes more catalytic cycles. Furthermore such a system can 'relax' back to its more stochastic state when not undergoing catalysis (15 - 16).

Transition State Ensembles and MultiFunctional Proteins

A related concept concerning enzyme function has recently developed, which acknowledges the utility of 'promiscuity' in biological function (18). Promiscuity is a functional property with distinct roles in: 1) protein interaction networks where 'hub' proteins regulate the function of multiple protein partners (19); 2) enzymatic detoxification by enzymes that metabolize an extraordinary range of structurally unrelated substrates (20); 3) the immune response, where promiscuous germline antibodies provide an efficient primary immune response (21); and 4) in maintaining a pool of structural scaffolds from which to evolve new enzymes (18, 22, 23). Tawfik and co-workers have attempted to articulate many aspects of 'promiscuity' and suggest the importance of distinguishing 'promiscuous' enzymes from 'multifunctional' enzymes, such as detoxification enzymes that have a clear function related to the metabolism of multiple substrates (18). Here, for convenience, we still use the terms interchangeably; the labels 'promiscuity' or 'multifunctionality' depend upon our knowledge of an enzyme's function. A presumed critical physical trait of promiscuous enzymes, or multifunctional ones, is flexibility, or the ability to sample a wide range of conformational space, and this may facilitate evolution of new function (18, 22, 23). In effect, functional promiscuity is likely to be correlated with structural plasticity (18, 22 - 24).

Merging Promiscuity with Conformational Ensembles; the Example of Detoxification Enzymes

Based on these nascent ideas about promiscuity, enzymatic conformational ensembles, and their common element of conformational dynamics, we have considered the possibility of time-dependent, emergent, properties of detoxification enzymes and transporters. Enzymes such as cytochrome P450s (CYPs), uridine diphosphate glucuronosyl transferases (UGTs), and transporters such as P-glycoprotein (P-gp) are known to be extraordinarily substrate promiscuous or 'multifunctional' and, as detoxification enzymes, they have evolved to decompose or transport a wide range of chemicals to which we are exposed. In the clinical realm, these enzymes dominate drug metabolism and thus are a critically important component of the therapeutic success or failure of drugs in development. We suggest that the energy landscapes of such enzymes lie near the limit of conformational breadth for transition state ensembles sampled by any enzymes.

The multi-conformational substrate nature of detoxification enzymes, coupled with the possibility of multiple "elementary reactions" connecting mixtures of reactants and products, has fascinating consequences when considered within the context of their functional niche, which includes interactions with multiple drugs or toxins in a mixture. In this paper, our aim is to reconsider possible sources of complex kinetic behavior frequently observed with detoxification enzymes and to suggest that researchers who study them consider these scenarios when interpreting their data. An important task for us is to demonstrate the thermodynamic feasibility of time-dependent behavior, as it shares some of the essence of "conformational drift theory" which has been rigorously disputed on thermodynamic grounds (25, 26). Our analysis is similar to the kinetic proofreading mechanism (27) which involves free energy expenditure, where catalysis itself provides the chemical free energy required for such "drift". Non-Michaelis-Menten (MM) kinetics are widely observed in detoxification enzyme reactions. One possible mechanism for the non-MM behavior comes from multiple substrates binding with allosterism, and we acknowledge that multiple binding is an established source of allosterism with these enzymes (28 - 32). In addition protein interactions between detoxification enzymes may contribute to complex kinetics (33). Another recently proposed possible mechanism of allosteric kinetics, with other enzymes, is dynamic cooperativity which is akin to the kinetic proofreading (27). The ideas

summarized here, however, provide the essential reminder that such kinetics are possible even in the absence of multple ligand binding or protein-protein oligomerization.

Examples of Complex Kinetics

The steady state and pre-steady state effects of conformational fluctuation on kinetically relevant time scales have been discussed in theoretical examples (15, 16), and these effects have been demonstrated experimentally with several enzymes, including at least two broad specificity enzymes (34 - 37). Specifically, butylcholinesterase is thought to be a detoxification enzyme and an epoxide hydrolase (SteEH1) metabolizes a wide range of plant epoxides. Both are well described by kinetic models that include conformational sampling coupled with hysteresis (lag times) in transient kinetics or apparent cooperativity in steady state turnover. In addition, for the case of the epoxide hydrolase, with (1R, 2R) *trans*-2-methyl-styrene oxide as substrate, two diastereomeric hydrolysis products are formed. The generation of two products from a single substrate (product promiscuity), by definition, requires an energy landscape with more than one transition state. Furthermore, product promiscuity is a hallmark of detoxification enzymes, in addition to their well-appreciated substrate promiscuity. We acknowledge that, for these cases, there may be no actual flux along the transition state barrier, i.e. the saddle points corresponding to the two transition states may be separated by large energy barriers. The essential point is that multiple saddle points are accessible to the ground state substrate ensemble {[E1•S] … [En•S]} and to the ground state product ensemble {[E1•P] … [En•P]}. Depending on the landscape between states within the substrate and product ensembles, fascinating things can happen during catalytic turnoer. We provide some simple models to demonstrate the relevant parameters of interest.

Results

Models of time-dependent catalytic behavior

To demonstrate the kinetic principles, one can simply consider two equilibrating conformational substates of an enzyme E in the ligand free form, E_1 and E_2, where the catalytic efficiency of E_2 is significantly higher. There are many variations on this type of enzyme kinetic model based on the work of C. Frieden and others in the 1960s and 1970s (38), including recent studies by Min et al. (39). Here, we propose a variation on the theme: we consider the possibility that a catalytic cycle can push the enzyme from the E_1 conformation to the E_2 conformation, as shown in Fig. 2A. In Figure 2 k_{cat}/K_m for E_1 is $k_1 k_2/(k_{-1} + k_2) = 10$ and k_{cat}/K_m for $E_2 = k_3 k_4/(k_{-3} + k_4) = 1000$. That is E_2 is a much better catalyst than E_1. In Fig. 2A, we have deliberately neglected the direct fluctuations between E_1S and E_2S: This is to accentuate the idea of multiple pathways with multiple transition states. This emphasizes the lack of flux between structurally distinct complexes, which is possible for the reactions of detoxification enzymes, that yield different products or that start with very different substrates. The similar case of rapid equilibrium between E_1S and E_2S has been studied in (14).

Because there is a continuous S to P catalytic cycle, the substates of the enzyme are not in equilibrium, but rather they are in a non-equilibrium steady state (NESS). The steady state ratio of $[E_2]/[E_1]$ is

$$\frac{[E_2]^{ness}}{[E_1]^{ness}} = \frac{\widehat{a}(k_{-1}+k_2)+k_1 k_2[S]}{\overset{\prime}{a}(k_2+k_{-1})+k_{-1}k_{-2}[P]} = \frac{10^{-7}+10^3[S]}{10^{-2}+10^{-8}[P]}.$$

(1)

We see this ratio is very different from the ratio of $[E_2]/[E_1]$ in the absence of S or P, which is

$$\frac{[E_2]^{ness}}{[E_1]^{ness}} = \frac{\widehat{a}}{\acute{a}} = \frac{10^{-7}}{10^{-2}}.$$

(2)

Thus the ratio of E_2/E_1 is much greater in the presence of substrate than in its absence. The concentrations of free E_1 and E_2 differ from their equilibrium values when the enzyme is undergoing catalysis. Interestingly, however, for a reversible reaction the ratio of $[E_2]/[E_1]$ will be the same in the presence and absence of S, when the ratio of $[S]/[P] = (k_1 k_2 \alpha)/(k_1 k_2 \beta)$ is equal to their chemical equilibrium ratio. In this case, the ratio $[E_2]/[E_1]$ in equilibrium follows Boltzmann's law. It is determined by the internal energies of the E_1 and E_2 dictated by their molecular structures.

As long as $[S]/[P] \gg k_{-1}k_{-2}\beta/(k_1 k_2 \alpha) = 10^{-15}$, then the ratio in Eq. (1) $\gg \beta/\alpha$. So, the turnover of the enzyme with S keeps the ratio of $[E_2]/[E_1]$ high and pushes the enzyme population toward the more active state. The interesting result of this situation is observed upon calculating the turnover flux of S to P (J^{ness}), which is

$$J^{ness} = \frac{(1 + 10^7[S])\,(10^3[S] - 10^{-12}[P])}{10^2 + 10^7[S] + 10^6[S]^2 + 10^{-3}[S][P] + 10^{-14}[P]}$$

(3)

Using this equation, the substrate concentration dependence of the flux (Figure 2 C) and the time-dependence of product formation (Figure 2B) with a given initial concentration of S can be simulated. As can be seen there is an apparent cooperativity at low [S] (Inset of Figure 2C), or a sigmoidal response in the concentration [S] vs. flux curves and the progress curve includes a marked lag in the formation of P, which reflects the changing concentration of $([E_2] + [E_2 S])$ with increased turnover. These well-appreciated properties of slowly exchanging enzyme conformations are not typically observed with classic 'Michaelis-Menten enzymes.' Interestingly, however sigmoidal plots of velocity vs. [S] are commonly observed with detoxification enzymes (28 - 32). There is ample evidence for many detoxification enzymes that the non-Michaelis-Menten kinetics include contributions from multiple substrate binding (40, 41), but the behavior described above may also contribute. In fact, this type of model describes the behavior of StEH and butylcholinesterase, as already noted (34 - 37).

The important reminder that comes from this model, as already appreciated (15 - 17), is that ground state heterogeneity of enzymes, with corresponding heterogeneity at the level of the transition state in the form or distinct elementary reactions, can yield emergent properties wherein the apparent kinetic properties of a *population* of enzymes change with time. Simultaneous heterogeneity in both the ground state and the transition state, which implies parallel paths for enzyme turnover, provides energy driven conformational drift of enzyme states.

It is interesting to consider enzymes that might exhibit these properties. For the reasons described above, wherein catalytic promiscuity or multifunctionality is likely to be due to conformational flexibility, detoxification enzymes are among those likely to exhibit such hysteresis. The possibility for such behavior with detoxification enzymes becomes

significantly more intriguing when situations that are characteristic of detoxification catalysis are considered.

Case 1. Simultaneous Exposure to Multiple Substrates

An interesting implication of these ideas becomes obvious in the context of the functional promiscuity of detoxification enzymes. It is completely clear that a single isoform of, for example, a CYP can metabolize structurally unrelated drugs, including many that are simultaneously used in a single patient. This is a major source of drug-drug interactions. Extreme cases occur when one drug significantly inhibits the metabolism of another, but in many cases both drugs are metabolized efficiently. At the molecular level, however, simultaneous exposure to multiple drugs has fascinating consequences if the energy landscape has the properties we have considered above. We demonstrate this with the scheme in Figure 3A, which includes two substrates S_1 and S_2 that are converted to corresponding products P_1 and P_2, with differential kinetic properties for enzyme conformations E_1 and E_2.

In this case, the ratio of the rates of metabolism of S_1 and S_2 is a simple function of the dynamic ratio $[S_1](t)/[S_2](t)$:

$$\frac{J_{S_1 \to P_1}}{J_{S_1 \to P_1} + J_{S_2 \to P_2}} = \frac{\theta[S_1](t)}{\theta[S_1](t) + [S_2](t)},$$

4

where $\theta = k_{cat,1} K_{m2}/(k_{cat,2} K_{m1})$ is the ratio of the catalytic powers of the two states of the enzyme, for these respective substrates, and the J terms are the flux of S_1 to P_1 or S_2 to P_2 as denoted in the subscript. The Eq. (4) shows that if the ratio $[S_2]/[S_1]$ increases, then the reaction flux that metabolizes S_2 will increases with respect to that of S_1. Therefore, there is an "adaptation" in the teamwork of the enzyme ensemble. At high concentrations of both S_1 and S_2, the enzyme population adjusts to convert both substrates simultaneously; after several catalytic cycles no individual enzyme molecule alternates between S_1 and S_2. Rather, the population has divided itself into two teams which each specialize in the metabolism of S_1 or S_2. Importantly, upon partial depletion of the substrate that is more quickly eliminated, the enzyme population readjusts to metabolize the other substrate more efficiently (Figure 3B). The enzyme molecules previously dedicated to clearance of S_2 adapt to aid in the clearance of S_1, and vice versa. Such nongenetic adaptation has not been documented experimentally, yet it represents a fascinating possibility predicted by the emerging view of enzyme ensembles and the increasing evidence that detoxification enzymes are conformationally heterogeneous.

In addition to Eq. (5), we also have

$$\frac{[E_1]+[E_1 S_1]}{[E_2]+[E_2 S_2]} = \left(\frac{\beta}{\alpha}\right) \frac{1+[S_1](t)/K_{m1}}{1+[S_2](t)/K_{m2}}.$$

(5)

We see that if both substrates are below their respective enzyme conformation K_m's, then the ratio of the two forms of the enzyme remains constant. However, if the concentration of one of the two substrates is greater than its corresponding K_m, then the enzyme population will shift toward the conformation needed to metabolize the remaining substrate. It is this dynamic adaption that "allocates" the enzyme to process the toxin at higher concentration relative to its K_M.

Dynamically, we also note that the S_2 really serves as a non-competitive inhibitor for the enzyme E_1, and S_1 is a noncompetitive inhibitor of E_2. That is, the two substrates will *not yield competitive inhibition* kinetics. If the rates of conversion of S_1 are monitored as a function of the concentration of S_2, then S_2 will not appear as a competitive inhibitor, but rather as a noncompetitive inhibitor of S_1. This is an important result in as much as many detoxification enzyme exposed simultaneously to multiple substrates *in vitro*, yield noncompetitive or mixed inhibition when analyzed by standard methods (42 - 45).

The striking result of the situation we consider here is that a population of equilibrating conformers, after exposure to multiple substrates, can distribute itself into 'teams' wherein one team specializes in the clearance of one substrate within the mixture, and other conformational team metabolizes other members of the substrate mixture. Such 'conformational teamwork', if it occurs, would represent a fascinating adaptability of the enzyme, ideally suited to perform its function of detoxification when faced with a mixture of toxins. In effect the ability of a single detoxification enzyme, as a single polypetide sequence, to metabolize a wide range of substrates simultaneously, without competition between substrates toward individual enzyme molecules, is enhanced by the teamwork achieved by discreet conformations.

Case 2: Formation of Multiple Products

In addition to their ability to metabolize multiple substrates, detoxification enzymes such as CYPs also frequently display product promiscuity, as noted above for StEH. That is, they can metabolize a single substrate to multiple products. With CYPs, the products formed from a single substrate often require dramatically different substrate orientations within the active site. Substantial rearrangement of the [E•S] complex would be required to convert the complex yielding one metabolite to the complex yielding the other metabolite.

This behavior also has interesting potential implications when the energy landscapes of the type we describe are operative. Specifically, for a single substrate being metabolized to multiple products via different conformations, after an initial pre-steady state period, at the level of the enzyme population, two 'teams' will emerge. For short time increments, or at saturating [S], one conformational team will generate exclusively one product and the other team will generate the second product, as long as each product leaves behind a conformation that is selective for the parent substrate orientation that yields that product, and as long as the conformational exchange among free enzyme states is sufficiently slow. However, the teams will redistribute in response to [S]. Thus, when [S] is high, a constant ratio of P2/P1 is formed, possibly to optimize clearance of total [S]. However, as [S] decreases the population shifts in favor of the conformation with higher affinity for S, with greater relative formation of the corresponding product (P_1).

A simple kinetic scheme to represent this scenario is shown in Fig. 4A in which the vertical transitions are sufficiently slow; thus each catalytic "pathway" essentially proceeds as an independent enzyme catalyzed reaction that produces different products P_1, P_2, .., Pn, with a slow conversion between the different forms of the enzyme.

Are there multiple interconverting ligand-dependent ground states and multiple transition states with detoxification enzymes?

Based on these speculative possibilities, which are supported by first principle simulations, it is important to seek or document experimental support for conformational exchange on relevant time scales for other detoxification enzymes. In addition to the examples of epoxide hydrolaase and butylcholinesterase mentioned above, other detoxification enzymes should be considered. Obviously, single molecule methods will be required to directly observe the

behavior we postulate here. To date there are few published single molecule experiments with substrates binding to CYPs (46) or any other detoxification enzyme, and none that include catalytic turnover. However, we propose these methods could be an essential component of future work required to fully understand these enzymes. In the absence of direct experimental support from single molecule studies, it is useful to consider other recent work that may suggest these behaviors. Indeed, several spectroscopic and structural studies suggest the presence of conformational heterogeneity of CYPs (47 - 55), possibly with slow exchange between states. Other spectroscopic methods that suggest ground state conformational heterogeneity include CO flash photolysis studies that reveal multiple kinetic components (56), and steady state spectral experiments that indicate the simultaneous presence of different [E • S] complexes from a single substrate (57). Together, the results demonstrate that the CYP fold is capable of functionally relevant, in some cases large-scale, conformational changes among ground state complexes that may exchange on relevant time scales. In short, the roles of conformational change in CYP function have been appreciated and well documented, including the possible role of distinct conformations in complex CYP kinetics (58), but they have not been treated as a source of time-dependent adaptation.

The concept we expose here is generally applicable to any protein that interacts with multiple partners or substrates via different conformations, and is not limited to detoxification enzymes. A population of promiscuous enzymes or proteins may work in temporally distinct conformational teams if simultaneously exposed to multiple substrates, and the teams may adapt to changing substrate concentrations. That is, such enzymes may exhibit stochastic behavior upon initial exposure to a mixture of substrates or to a substrate from which multiple products are generated, but after some catalytic turnover each enzyme molecule may exhibit temporary deterministic behavior, until the substrate environment changes and the population of molecules 'relaxes' to the stochastic situation again. Similarly, for regulatory proteins that interact with multiple partners, the conformational distribution could change in response to altered expression of some binding partners, and there could be interesting and important time-dependence to these processes. The possibility of such fascinating behavior, exemplified by the detoxification enzymes considered here, warrants consideration.

Acknowledgments

References

1. Zwanzig R. Diffusion in a rough potential. Proc Natl Acad Sci. 1988; 85:2029–2030. [PubMed: 3353365]

2. Frauenfelder H, Sligar SG, Wolynes PG. The energy landscapes and motions of proteins. Science. 1991; 254:1598–1603. [PubMed: 1749933]

3. Frauenfelder H, Parak F, Young RD. Conformational substates in proteins. Annu Rev Biophys Biophys Chem. 1988; 17:451–479. [PubMed: 3293595]

4. Boehr DD, Nussinov R, Wright PE. The role of dynamic conformational ensembles in biomolecular recognition. Nat Chem Biol. 2009; 5:789–796. [PubMed: 19841628]

5. Hammes GG, Chang YC, Oas TG. Conformational selection or induced fit: a flux description of reaction mechanism. Proc Natl Acad Sci U S A. 2009; 106:13737–13741. [PubMed: 19666553]

6. Weikl TR, von Deuster C. Selected-fit versus induced-fit protein binding: kinetic differences and mutational analysis. Proteins. 2009; 75:104–110. [PubMed: 18798570]

7. Ma B, Nussinov R. Enzyme dynamics point to stepwise conformational selection in catalysis. Curr Opin Chem Biol. 2010; 14:652–659. [PubMed: 20822947]

8. Cui Q, Karplus M. Allostery and Cooperativity Revisited. Protein Sci. 2008; 17:1295–13076. [PubMed: 18560010]

9. Roca M, Messer B, Hilvert D, Warshel A. On the relationship between folding and chemical landscapes in enzyme catalysis. Proc Natl Acad Sci U S A. 2008; 105:14237–14238. [PubMed: 18799750]

10. Vamvaca K, Vögeli B, Kast P, Pervushin K, Hilvert D. An enzymatic molten globule: efficient coupling of folding and catalysis. Proc Natl Acad Sci U S A. 2004; 101:12860–12864. [PubMed: 15322276]

11. Rissin DM, Gorris HH, Walt DR. Distinct and long-lived activity states of single enzyme molecules. J Am Chem Soc. 2008; 130:5349–5353. [PubMed: 18318491]

12. Eisenmesser EZ, Millet O, Labeikovsky W, Korzhnev DM, Wolf-Watz M, Bosco DA, Skalicky JJ, Kay LE, Kern D. Intrinsic dynamics of an enzyme underlies catalysis. Nature. 2005; 438:117–121. [PubMed: 16267559]

13. Benkovic SJ, Hammes GG, Hammes-Schiffer S. Free-energy landscape of enzyme catalysis. Biochemistry. 2008; 47:3317–3321. [PubMed: 18298083]

14. Swint-Kruse L, Fisher HF. Enzymatic reaction sequences as coupled multiple traces on a multidimensional landscape. Trends Biochem Sci. 2008; 33:104–112. [PubMed: 18261912]

15. Qian H. Cooperativity and specificity in enzyme kinetics: a single-molecule time-based perspective. Biophys J. 2008; 95:10–17. [PubMed: 18441030]

16. Zhou HX. From induced fit to conformational selection: a continuum of binding mechanism controlled by the timescale of conformational transitions. Biophys J. 2010; 98:L15–L17. [PubMed: 20303846]

17. Min W, Xie S, Bagchi B. Role of conformational dynamics in kinetics of an enzymatic cycle in a nonequilibrium steady state. J Chem Phys. 2009; 131:065104. [PubMed: 19691414]

18. Khersonsky O, Tawfik DS. Enzyme Promiscuity: A Mechanistic and Evolutionary Perspective. Annu Rev Biochem. 2010; 79:471–505. [PubMed: 20235827]

19. Patil A, Kinoshita K, Nakamura H. Hub promiscuity in protein-protein interaction networks. Int J Mol Sci. 2010; 11:1930–1943. [PubMed: 20480050]

20. Ekins S. Predicting undesirable drug interactions with promiscuous proteins in silico. Drug Discov Today. 2004; 9:276–285. [PubMed: 15003246]

21. Sethi DK, Agarwal A, Manivel V, Rao KV, Salunke DM. Differential epitope positioning within the germline antibody paratope enhances promiscuity in the primary immune response. Immunity. 2006; 24:429–438. [PubMed: 16618601]

22. Meier S, Ozbek S. A biological cosmos of parallel universes: does protein structural plasticity facilitate evolution? Bioessays. 2007; 29:1095–1104. [PubMed: 17935152]

23. Tokuriki N, Tawfik DS. Protein dynamism and evolvability. Science. 2009; 324:203–207. [PubMed: 19359577]

24. Hou L, Honaker MT, Shireman LM, Balogh LM, Roberts AG, Ng KC, Nath A, Atkins WM. Functional promiscuity correlates with conformational heterogeneity in A-class glutathione S-transferases. J Biol Chem. 2007; 282:23264–23274. [PubMed: 17561509]

25. Weber G. Stability of oligomeric proteins and its bearing on their association equilibria (a reply). Proc Natl Acad Sci USA. 1983; 80:5303–5304. [PubMed: 6577427]

26. Berg OG. Time-averaged chemical potential of proteins and the detailed-balance principle (an alternative viewpoint). Proc Natl Acad Sci USA. 1983; 80:5302–5303. [PubMed: 6577426]

27. Goldstein B, Coombs D, Faeder JR, Hlavacek WS. Kinetic proofreading model. Adv Exp Med Biol. 2008; 640:82–94. [PubMed: 19065786]

28. Atkins WM. Non-Michaelis-Menten kinetics in cytochrome P450-catalyzed reactions. Annu Rev Pharmacol Toxicol. 2005; 45:291–310. [PubMed: 15832445]

29. Tracy TS. Atypical cytochrome p450 kinetics: implications for drug discovery. Drugs R D. 2006; 7:349–363. [PubMed: 17073518]

30. Ueng YF, Kuwabara T, Chun YJ, Guengerich FP. Cooperativity in oxidations catalyzed by cytochrome P450 3A4. Biochemistry. 1997; 36:370–381. [PubMed: 9003190]

31. Korzekwa KR, Krishnamachary N, Shou M, Ogai A, Parise RA, Rettie AE, Gonzalez FJ, Tracy TS. Evaluation of atypical cytochrome P450 kinetics with two-substrate models: evidence that multiple substrates can simultaneously bind to cytochrome P450 active sites. Biochemistry. 1998; 37:4137–4147. [PubMed: 9521735]

32. Atkins WM. Current views on the fundamental mechanisms of cytochrome P450 allosterism. Expert Opin Drug Metab Toxicol. 2006; 2:573–579. [PubMed: 16859405]

33. Davydov DR. Microsomal monooxygenase as a multienzyme system: the role of P450-P450 interactions. Expert Opin Drug Metab Toxicol. 2011 Epub.

34. Lindberg D, de la Fuente Revenga M, Widersten M. Temperature and pH dependence of enzyme-catalyzed hydrolysis of trans-methylstyrene oxide. A unifying kinetic model for observed hysteresis, cooperativity, and regioselectivity. Biochemistry. 2010; 49:2297–2304. [PubMed: 20146441]

35. Lindberg D, Ahmad S, Widersten M. Mutations in salt-bridging residues at the interface of the core and lid domains of epoxide hydrolase StEH1 affect regioselectivity, protein stability and hysteresis. Arch Biochem Biophys. 2010; 495:165–173. [PubMed: 20079707]

36. Lindberg D, Gogoll A, Widersten M. Substrate-dependent hysteretic behavior in StEH1-catalyzed hydrolysis of styrene oxide derivatives. FEBS J. 2008; 275:6309–6320. [PubMed: 19016837]

37. Masson P, Schopfer LM, Froment MT, Debouzy JC, Nachon F, Gillon E, Lockridge O, Hrabovska A, Goldstein BN. Hysteresis of butyrylcholinesterase in the approach to steady-state kinetics. Chem Biol Interact. 2005; 157-158:143–152. [PubMed: 16256969]

38. Frieden C. Slow Transitions and Hysteretic Behavior in Enzymes. Ann Rev Biochemistry. 1979; 48:471–489.

39. Min W, Gopich IV, English BP, Kou SC, Xie XS, Szabo A. When does the Michaelis-Menten equation hold for fluctuating enzymes? J Phys Chem B. 2006; 110:20093–20097. [PubMed: 17034179]

40. Ekroos M, Sjögren T. Structural basis for ligand promiscuity in cytochrome P450 3A4. Proc Natl Acad Sci U S A. 2006; 103:13682–13687. [PubMed: 16954191]

41. Davydov DR, Halpert JR. Allosteric P450 mechanisms: multiple binding sites, multiple conformers or both? Expert Opin Drug Metab Toxicol. 2008; 4:1523–1535. [PubMed: 19040328]

42. Wang LQ, Falany CN, James MO. Triclosan as a substrate and inhibitor of 3′-phosphoadenosine 5′-phosphosulfate-sulfotransferase and UDP-glucuronosyl transferase in human liver fractions. Drug Metab Dispos. 2004; 32:1162–1169. [PubMed: 15269185]

43. Zhang ZY, Wong YN. Enzyme kinetics for clinically relevant CYP inhibition. Curr Drug Metab. 2005; 6:241–257. [PubMed: 15975042]

44. Si D, Wang Y, Zhou YH, Guo Y, Wang J, Zhou H, Li ZS, Fawcett JP. Mechanism of CYP2C9 inhibition by flavones and flavonols. Drug Metab Dispos. 2009; 37:629–634. [PubMed: 19074529]

45. Volak LP, Ghirmai S, Cashman JR, Court MH. Curcuminoids inhibit multiple human cytochromes P450, UDP-glucuronosyltransferase, and sulfotransferase enzymes, whereas piperine is a relatively selective CYP3A4 inhibitor. Drug Metab Dispos. 2008; 36:1594–1605. [PubMed: 18480186]

46. Nath A, Koo PK, Rhoades E, Atkins WM. Allosteric effects on substrate dissociation from cytochrome P450 3A4 in nanodiscs observed by ensemble and single-molecule fluorescence spectroscopy. J Am Chem Soc. 2008; 130:15746–15747. [PubMed: 18980315]

47. Gay SC, Sun L, Maekawa K, Halpert JR, Stout CD. Crystal structures of cytochrome P450 2B4 in complex with the inhibitor 1-biphenyl-4-methyl-1H-imidazole: ligand-induced structural response through alpha-helical repositioning. Biochemistry. 2009; 48:4762–4771. [PubMed: 19397311]

48. Zhao Y, Sun L, Muralidhara BK, Kumar S, White MA, Stout CD, Halpert JR. Structural and thermodynamic consequences of 1-(4-chlorophenyl)imidazole binding to cytochrome P450 2B4. Biochemistry. 2007; 46:11559–11561. [PubMed: 17887776]

49. Zhang H, Kenaan C, Hamdane D, Hoa GH, Hollenberg PFJ. Effect of conformational dynamics on substrate recognition and specificity as probed by the introduction of a de novo disulfide bond into cytochrome P450 2B1. J Biol Chem. 2009; 284:25678–25686. [PubMed: 19605359]

50. Lepesheva GI, Seliskar M, Knutson CG, Stourman NV, Rozman D, Waterman MR. Conformational dynamics in the F/G segment of CYP51 from Mycobacterium tuberculosis monitored by FRET. Arch Biochem Biophys. 2007; 464:221–227. [PubMed: 17585868]

51. Muralidhara BK, Negi S, Chin CC, Braun W, Halpert JR. Conformational flexibility of mammalian cytochrome P450 2B4 in binding imidazole inhibitors with different ring chemistry and side chains. Solution thermodynamics and molecular modeling. J Biol Chem. 2006; 281:8051–8061. [PubMed: 16439365]

52. Gamage NU, Tsvetanov S, Duggleby RG, McManus ME, Martin JL. The structure of human SULT1A1 crystallized with estradiol. An insight into active site plasticity and substrate inhibition with multi-ring substrates. J Biol Chem. 2005; 280:41482–41486. [PubMed: 16221673]

53. Lampe JN, Brandman R, Sivaramakrishnan S, Ortiz de Montellano PR. 2D NMR and All-atom molecular dynamics of cytochrome P450 CYP119 reveal hidden conformational substates. J Biol Chem. 2010; 285:9594–9603. [PubMed: 20097757]

54. Lampe JN, Floor SN, Gross JD, Nishida CR, Jiang Y, Trnka MJ, Ortiz de Montellano PR. Ligand-induced conformational heterogeneity of cytochrome P450 CYP119 identified by 2D NMR spectroscopy with the unnatural amino acid (13)C-p-methoxyphenylalanine. J Am Chem Soc. 2008; 130:16168–16169. [PubMed: 18998650]

55. Roberts AG, Díaz MD, Lampe JN, Shireman LM, Grinstead JS, Dabrowski MJ, Pearson JT, Bowman MK, Atkins WM, Campbell AP. NMR studies of ligand binding to P450(eryF) provides insight into the mechanism of cooperativity. Biochemistry. 2006; 45:1673–1684. [PubMed: 16460014]

56. Koley AP, Buters JTM, Robison RC, Markowitz A, Friedman FK. CO binding kinetics of human Cytochrome P4503A4. J Biol Chem Soc. 1995; 270:5014–5018.

57. Hutzler JM, Melton RJ, Rumsey JM, Schnute ME, Locuson CW, Wienkers LC. Inhibition of cytochrome P450 3A4 by a pyrimidineimidazole: Evidence for complex heme interactions. Chem Res Toxicol. 2006; 19:1650–1659. [PubMed: 17173379]

58. Atkins WM, Wang RW, Lu AY. Allosteric behavior in cytochrome p450-dependent in vitro drug-drug interactions: a prospective based on conformational dynamics. Chem Res Toxicol. 2001; 14:338–347. [PubMed: 11304120]
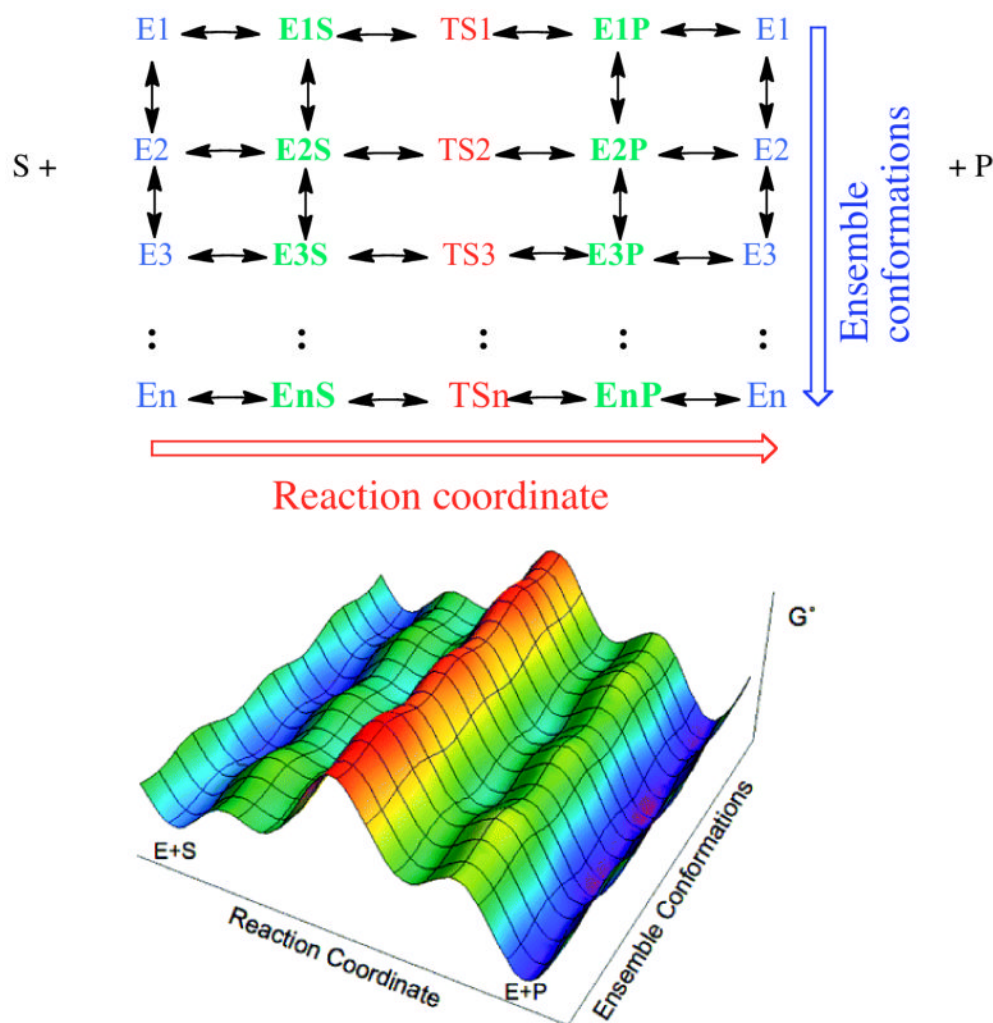
Figure 1.
Top: Two-dimensional matrix of enzyme states reflecting ligand free forms (blue) on the left (E1 through EN), substrate complexes (green) with the corresponding states (E1S through ENS), Transition state complexes (red, TS1 through TSN), and product complexes (blue, E1P through ENP) for each state, with return to the initial ensemble of ligand free states. Note, in our model the transition state are not connected via arrows – we do not imply conformation rearrangement at the transition state 'ridge' (red). The multiple conformations present along this ridge are separated by significant barriers. Bottom: A three dimensional free energy (G′) landscape representation of the system, when the catalytic transition states (TS) are rate limiting, color coded to as the matrix above. The lower free energy surface is adapted from reference 13.
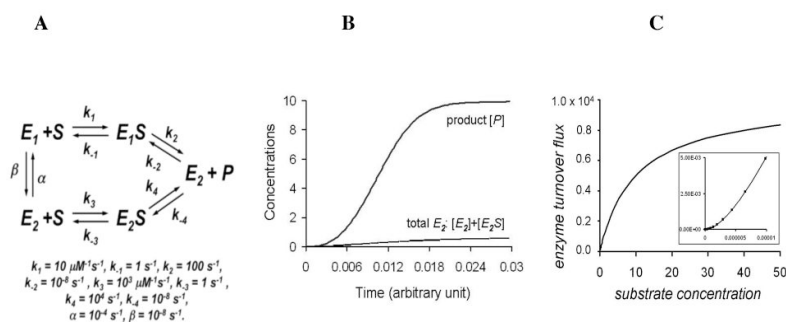
A   **B**   **C**

Figure 2.
A. A variation of the general fluctuting enzyme kinetic model with two conformational substates E_1 and E_2, which yield the same product from a single substrate with different kinetic parameters. In this example, we specifically assume that E_2 is a better catalyst than E_1. The parameters used for the simulations are shown. In particular, we have assumed that the enzyme is inevitably in the state E_2 after accomplishing a catalysis step. **B**. The concentration of product (P) or all forms of E_2 ($[E_2] + [E_2S]$) as a function of time. There is marked hysteresis (lag) in the formation of product as the population of enzyme states shifts toward a greater fraction of E_2 with increasing turnover. **C**. The rate of product formation vs. [S]. The inset is a close up of the low [S] range, and demonstrates the sigmoidal dependence of rate on [S].
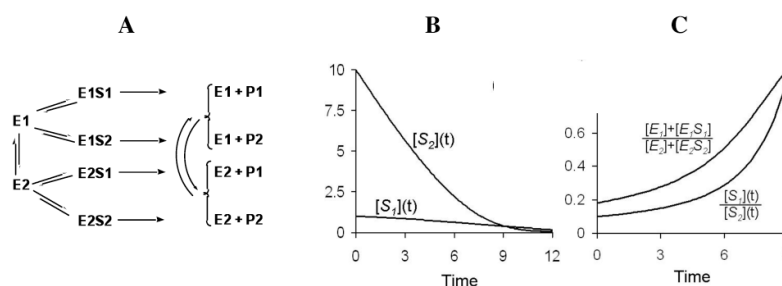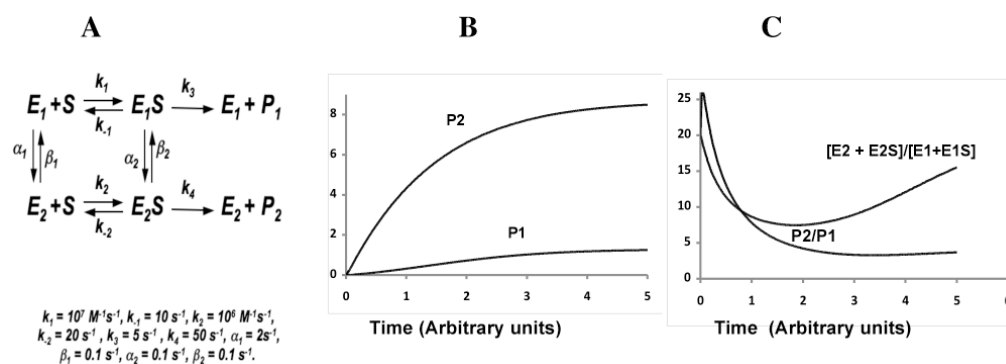
Figure 3.
Dynamic adaptation of a stochastic enzyme ensemble in processing two different substrates S_1 and S_2, using "teamwork". **A**. Minimal kinetic scheme with equilibrating E1 and E2, each contributing to clearance of both S_1 and S_2. However, as simulated, the two dynamically exchanging forms E_1 and E_2 have relative specificities for S_1 and S_2 respectively, and the reactions are assumed to be irreversible. The ratio of the two enzyme catalytic powers, $\theta = k_{cat,1} K_{m2}/(k_{cat,2} K_{m1}) = 0.3$ in the simulations shown. In equilibrium in the absent of substrates, the ratio of the two enzyme forms is $\alpha/\beta = 1$. The initial concentrations for $[S_1]/K_{m1} = 1$ and $[S_2]/K_{m2} = 10$. **B**. Concentrations of S_1 and S_2 vs. time. The enzyme ensemble immediately adjusts its distribution to 1:10 in order to process the dominant S_2. **C**. Ratios of E_1 and E_2 forms or S_1 and S_2 vs. time. With increasing time the ratio of $[S_1](t)$ to $[S_2](t)$ changes from 0.1 to 1, and the enzyme distribution 'adapts' accordingly.

Figure 4.
A. Minimal kinetic scheme for two enzyme conformations generating a multiple products from a single substrate. In the simulations E_2 is a more efficient catalyst than E_1. **B.** Rate of formation of P_1 and P_2. Note the hysteresis in the formation of P1. **C.** Ratios of all enzyme forms E_2 vs. E_1 ($[E_2+E_2S]/[E_1+E_1S]$) or the product ratio (P_2/P_1) vs. time. The enzyme forms 'adapt' from the initial preference for E_2 in the absence of substrate to a more 'balanced' ratio of all E_2 forms/all E_1 forms, and increases again as substrate is depleted.