Comment on “Regularizing capacity of metabolic networks”

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In a recent paper, Marr, Müller-Linow and Hütter [Phys. Rev. E 75, 041917 (2007)] investigate an artificial dynamic system on metabolic networks. They find a less complex time evolution of this dynamic system in real networks, compared to networks of null models. The authors argue that this suggests that metabolic network structure is a major factor behind the stability of biochemical steady states. We reanalyse the same kind of data using a dynamic system modeling actual reaction kinetics. The conclusions about stability, from our analysis, are inconsistent with those of Marr et al. We argue that this issue calls for a more detailed type of modeling.

I. INTRODUCTION

Within living organisms, matter is constantly converted between different molecular species. It is often assumed that concentrations of metabolites tend to settle into steady states (rather than showing periodic or chaotic behavior) (1). Furthermore, experimental studies of metabolic pathways are typically performed under steady-state conditions (2). Although the steady state is, strictly speaking, a mathematical abstraction, it is nevertheless a useful reference state (3). The steady-state assumption is also fundamental to traditional metabolic control analysis (2). In a recent article (4) Marr, Müller-Linow and Hütter hypothesize that the network structure of metabolism is an important factor for promoting a steady state, rather than a complex, dynamics. The authors run an artificial dynamic system on the networks. In short, a vertex i can have two states, 0 or 1. If the sum of the state variables in the neighborhood of i exceeds a fixed threshold, then i changes to the other state. This scheme is very different from real biochemical dynamics and traditional models of reaction kinetics. First, the state variables in metabolic models are usually continuous (concentrations (2)) or sometimes discrete variables (molecule counts (5)), but never binary. Second, the sum of these variables is conserved (if in- and outflow is neglected); whereas, in the dynamics of Marr et al., this is not the case. This dynamics does in fact not reach a steady state; instead, the authors analyze the complexity of the output time series with entropy-like measures. They conclude that real metabolic networks give a less complex output than different ensembles of model networks, and argue that this implies that the structure of real metabolic networks may promote steady state dynamics. However, no theory for how the values of the “entropies” of the binary dynamics relate to the stability of metabolic flux is given. Many authors have used binary dynamics to model other processes of cellular biology, such as signal transduction (e.g. Ref. (6)) or genetic regulation (e.g. Ref. (7)). However, these models explicitly try to describe systems in which the components, at least under some circumstances, show switch-like, binary behavior (6). Moreover, the lack of detailed understanding of e.g. gene expression makes it more natural to use simplified dynamics to uncover underlying principles. Metabolic reactions follow known principles of chemistry and are described by well-established differential equation models. Therefore, in contrast to the mentioned information processes, there is no need for, or natural interpretation of, a binary description of the metabolism. Therefore the conclusions of Marr et al. can be validated by comparing their results to results obtained using such standard differential equation modeling. We follow this approach to find no significant difference between real and null-model network. Our simulations in this Comment are rather limited, but sufficient to make the point that Marr et al.’s study cannot rule out the possibility that stable steady-states can be explained more simply, as direct consequences of the reaction kinetics, rather than as a result from the interaction between the network topology and the dynamic system.

II. CONSTRUCTING A REACTION SYSTEM

Marr et al.’s examples of real metabolic networks are taken from a work of Ma and Zeng (MZ) (8). They are substrate graphs (9) where a substrate is linked to the products of a reaction (Fig 1(a)). MZ also preprocessed the data by omitting ubiquitous “currency metabolites” (10). We also use the MZ networks as the starting point for our simulations. In constructing the networks, information is lost; so if one wants to simulate the reaction system with a substrate graph as starting point, one needs (explicitly or implicitly) to recreate the reactions via a model. From two assumptions about reaction systems, we propose a simple scheme to create a plausible set of reactions that can be reduced to a given substrate graph. We first assume that all reactions are of a 2–2-form: A + B ↔ C + D, or 2–1-form: A + B ↔ C. These are the most com-

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**FIG. 1** (Color online.) Illustration of the construction of substrate graphs from chemical reactions (a), and our method for recreating a reaction system from a substrate graph (b).
mon forms of biochemical reactions. We do not include more complex reactions, mostly because it would significantly increase the computational complexity. Our second assumption is that the number of reactions creating an edge in the substrate graph is rather small (we confirm a posteriori that the average number of reactions per vertex is similar to that of the real substrate graph). The following algorithm is a simple way of fulfilling these assumptions:

1. Start with all edges unmarked.
2. Pick one unmarked edge (A, C).
3. Find the maximal, full bipartite subgraph \( K_{(A,C)} \) (a subgraph consisting of two vertex sets, and edges between every pair of vertices in the different sets, but no edge between vertices in the same set) that contains (A, C). (See Fig. 1(b).)
4. Pick one four-cycle of \( K_{(A,C)} \) including (A, C) (say (A, C, B, D, A)). (See Fig. 1(b).)
5. Add A + B \( \rightarrow \) C + D to the set of reactions and mark the edges (A, C), (C, B), (B, D), (D, A).
6. If there are unmarked edges go to step 2.

\( K_{(A,C)} \) gives all reactions of the 2–2-form that induce the edge (A, C) in a substrate graph. If \( K_{(A,C)} \) is empty at step 3 then (A, C) must have been generated by a reaction of a 2–1-form. Therefore, instead of \( K_{(A,C)} \), consider the set \( L_{(A,C)} \) of (A, C) and its adjacent edges, and deduce reactions of the 2–1-form at step 5.

III. REACTION KINETICS

Once we have a set of reactions, derived from a substrate graph, we simulate the biochemical dynamics by standard Michaelis–Menten (MM) kinetics (11) supplemented by noise in the enzyme and substrate concentrations. MM kinetics are used to model enzyme-catalyzed reactions, which are common in metabolism. The MM description builds on the principle of mass-action (11)—the forward rate of a reaction is proportional to the product of the concentrations of the substrates—and uses additional assumptions to simplify the resulting rate laws. For the reaction \( i, A + B \rightarrow C + D \) (which is assumed to be enzyme-catalyzed), we use a version of the two-substrate MM rate law to calculate the flux

\[
r_i = \frac{E_i + \eta_E}{k_0^{-1} + 1/k_1[A] + 1/k_2[B] + 1/k_3[A][B]}.
\]  

where \( \eta_E \) is a random variable modeling fluctuations in the enzyme concentration. The time evolution of the concentration of a substance A is then determined by

\[
\frac{d[A]}{dt} = \eta_{conc.} + \sum_i \pm r_i,
\]

where the sum is over all A’s reactions, the sign in front of \( r_i \) is positive (negative) if A is a product (substrate) of i, and

\[
\eta_{conc.}
\]

is a random noise term modeling fluctuations due to inflow and outflow of A. We seek to use the same information as in Ref. (4), therefore we do not use empirical parameter values (which are, anyway, hard to obtain). Instead, we assign parameters in arbitrary units, but we choose them to be reasonable relative to one another. \( \eta_E \) and \( \eta_{conc.} \) are normally distributed \( N(0,0.002) \) (the first argument is the mean; the second is the standard deviation). New values for \( \eta_E \) and \( \eta_{conc.} \) are drawn every time step of the integration. The initial values of substance concentrations, enzyme concentrations and reaction coefficients (\( k_0, k_1, k_2 \) and \( k_{12} \)) are drawn from \( N(1,1), N(0.1,0.1) \) respectively. We choose \( \mu_k = 1 \times 10^{-3}, \sigma_k = 5 \times 10^{-4}, \mu = 1 \) and \( \sigma = 0.5 \). The equations are integrated with a second order Runge–Kutta–Helfand–Grenside scheme with time step 0.1, total running time 2500 time units, and 20 averages over different sets of initial configurations. These runs are, for each network, averaged over 20 realizations of the reaction-system construction. For comparison, we also run the dynamics on 100 samples of one Marr et al.’s null-model networks—random graphs with the same degree sequence as the original network. These are obtained by random rewiring—we go through all edges and for each edge (i, j) randomly pick an edge \( (i’, j’) \) and replace these edges with \( (i’, j) \) (unless this would introduce a multiple edge or a self-edge, in which case a new random edge \( (i’, j’) \) is selected).

IV. SIMULATION RESULTS

In the simulations, a vast majority of the substance concentrations converge to steady states. A few subnetworks of oscillating or chaotic concentrations may exist (12), but in this Comment we focus on bulk properties. To study the approach to equilibrium, and fluctuations, we measure the average standard deviation of the flux through the substances,

\[
\Sigma = \sqrt{\frac{1}{n} \sum_i \Phi_i^2 - \left( \frac{1}{n} \sum_i \Phi_i \right)^2}, \quad \text{where} \quad (3a)
\]

\[
\Phi_i = \frac{1}{2} \sum_j |r_j|, \quad \text{and} \quad (3b)
\]

The sums in Eq. (3a) are over all substances; the sum in Eq. (3b) is over i’s reactions; the factor 1/2 comes from the double
count of mass-flow in and out of substance $i$, and $n$ is the number of substances. In a plot of $\Sigma(t)$, the stability of the steady state can be monitored by two quantities. First, when the system approaches an equilibrium, $\Sigma(t)$ will decrease—a faster decrease implies a more stable system. Second, a lower equilibrium level means that the system has less cyclic or chaotic components, and responds faster to perturbations from the noise, and therefore has more stable steady states. In Fig. 2 we plot $\Sigma(t)$ for MZ’s human metabolic network and the null-model networks. The two curves almost overlap. For different null-model realizations the curves may deviate slightly from those of the real network, but there the null model cannot be rejected with any high level of significance. The equilibrium level is of the order of the noise, which means that periodic and chaotic behavior is almost fully suppressed. To be more systematic, we note that $\Sigma(t)$ display two aspects of stability—how fast equilibrium is reached and the height of the equilibrium level. Since the curves do not fit any simple functional form we measure the half-time (the time to reach midway between $\Sigma(0)$ and $\Sigma(\infty)$) using spline interpolation. The p-value (fraction of null-model observations lower than the real value) of the half-time is 67%; the corresponding value of the equilibrium level is 65%. The effect of the difference in network structure between the real and null-model networks is thus, in this case, negligible. We test a few other sets of parameter values, organisms (E. coli and M. musculus), and a more straightforward mass action kinetics; and, in all cases, arrive at the above conclusion. A full scan of the parameter space would be interesting, however, in this Comment we just make the point that the conclusion of Marr et al. can be inconsistent with more realistic simulations, and leave the insignificance of network structure to the stability of metabolic steady states as a conjecture.

V. CONCLUSIONS

To conclude; simple, stylized dynamic systems—“dynamic probes”—are valuable tools for studying complex biological systems. We believe that these should be designed to model the real dynamics as closely as possible. Marr et al. [4] propose a dynamic probe to study metabolic steady states that violates many of the known features of reaction kinetics; it does not even reach a steady state—the objective of the study as given in their Abstract. By the collective effort of researchers, our understanding of the metabolism continuously advances. This does, however, not improve the approach of Marr et al. as their dynamics does not make use of biochemical information. Relative to biological information processes, metabolism is a rather simple system, and is believed to be well-described by simple differential equation models. For the simulations we carry out, the null-model networks are not less efficient than the real networks in suppressing complex dynamics. If this holds in general, then steady-state stability is a fundamental property of chemical reaction systems. The study of Marr et al. does not rule out such a simple explanation of steady-state behavior—an explanation that is a common opinion in biochemical literature (cf. Hofmeyr and Cornish-Bowden’s dictum “mass-action is the intrinsic driving force for self-organization of reaction networks” [3]). If the reaction kinetics is the sole cause of metabolic steady-states, the steady-state dynamics is a constraint to, rather than an outcome of, natural selection. This situation is reminiscent of the power-law degree distribution of metabolic networks. Such distributions can also be seen in astrochemical networks that are not subject to natural selection [13]. We believe the question of dynamic stability in metabolism should be studied at a more detailed level than networks, avoiding the reduction of database information to substrate graphs. This is not to say that graph theory is useless in the study of metabolism. On a large scale, metabolic networks are different from the null-model networks. This is a valid conclusion from Ref. [4]—the authors manage to separate the real metabolic networks from both the null-model network we use, and various other types of model networks derived from the original graph. We believe much information about the organization of metabolism lies in the answer to how this separation occurs. On smaller scales, network theory can be used to find e.g. functional modules [10] and functions of individual metabolites [14].

In a reply [15] to this comment the Marr et al. argue that (with reference to our reconstruction of the reaction system) the fact that reaction coefficients are larger in one direction effectively break the network into disconnected components with respect to the dynamics. This is, of course, true in real reaction systems too, and a possible stabilizing mechanism for metabolism. However, it is irrelevant to our point that Marr et al., in Ref. [4], by measuring an “entropy” of a time series from a dynamic system unrelated to reaction kinetics, cannot draw conclusions about the effect of metabolic network structure on biochemical dynamics. To make this point, we use the same starting point as Marr et al., it should be easy to check that our conclusions also hold for the MM dynamics on full reaction systems from metabolic databases.

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