

Global characterization of in vivo enzyme catalytic rates and their correspondence to in vitro k_{cat} measurements

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Turnover numbers, also known as k_{cat} values, are fundamental properties of enzymes. However, k_{cat} data are scarce and measured in vitro, thus may not faithfully represent the in vivo situation. A basic question that awaits elucidation is: how representative are k_{cat} values for the maximal catalytic rates of enzymes in vivo? Here, we harness omics data to calculate $k_{\text{cat}}^{\text{in vivo}}$, the observed maximal catalytic rate of an enzyme inside cells. Comparison with k_{cat} values from *Escherichia coli*, yields a correlation of $r^2 = 0.62$ in log scale ($p < 10^{-10}$), with a root mean square difference of 0.54 (3.5-fold in linear scale), indicating that in vivo and in vitro maximal rates generally concur. By accounting for the degree of saturation of enzymes and the backward flux dictated by thermodynamics, we further refine the correspondence between $k_{\text{cat}}^{\text{in vivo}}$ and k_{cat} values. The approach we present here characterizes the quantitative relationship between enzymatic catalysis in vitro and in vivo and offers a high-throughput method for extracting enzyme kinetic constants from omics data.

kinetic constants | proteomics | turnover number | k_{cat} | flux balance analysis

The rates of enzyme catalysis are fundamental to the understanding of central cellular phenomena, such as metabolite concentrations and fluxes, cell responses to perturbations, and resource investment in enzymes (1–6). Many models of cellular metabolism include k_{cat} values, the maximal turnover rates of enzymes, as key inputs to predict the behavior of metabolic pathways and networks (7–9). However, most values have never been measured experimentally. *Escherichia coli* is the most intensely biochemically characterized organism, but k_{cat} values are available for only about 10% of its $\approx 2,000$ enzyme-reaction pairs (Dataset S1). Indeed, k_{cat} values are missing for several central metabolic enzymes. The scarcity of kinetic data limits the scope of models and necessitates generic parameter assignments that significantly reduce the predictive power of cellular models.

Even if a larger collection of k_{cat} values was made available, their current use poses a major difficulty: k_{cat} values are measured through in vitro enzyme assays, representing the initial rate of the reaction, i.e., full substrates saturation and negligible levels of products. Such assays may underrepresent factors like cellular metabolite concentrations, thermodynamic constraints, posttranslational modifications, chaperones, cellular crowding, and activating and inhibiting molecules, which can substantially alter enzyme kinetics in vivo. These omissions call into question the relevance of k_{cat} measurements in vivo (10–12). Furthermore, an effort to measure a large number of k_{cat} values under in vivo-like conditions presents a daunting challenge, given how many unknown biochemical factors might be involved.

Several studies grapple with missing k_{cat} values by sampling from the distribution of k_{cat} values measured in vitro or by using measurements of the same enzyme from related species (13–16). These approximations systematically ignore any errors resulting from the differences between in vitro and in vivo environments. Approximations of this sort may also introduce significant errors, as k_{cat} values can deviate by orders of magnitude between isoforms in the same organism as well as across organisms (17–19).

Here, we describe an alternative approach that addresses the above challenges by leveraging recent progress in omics studies. The in vivo catalytic rate of an enzyme can be inferred from the flux carried by the enzyme and the enzyme copy number. Proteomics methods now offer quantitative measurements of enzyme levels in several organisms and under a wide range of growth conditions. By dividing flux from computational flux predictions by enzyme abundances from proteomics, we calculate the rate of enzymes in vivo, denoted as k_{app} (20). We perform this analysis over a large set of growth conditions ($n = 31$). By taking the maximum value of k_{app} across many conditions, we obtain an estimate of the maximal turnover rate of an enzyme in vivo, which we define as $k_{\text{cat}}^{\text{in vivo}}$. What is the relationship between k_{cat} and $k_{\text{cat}}^{\text{in vivo}}$? We compare these values and quantitatively analyze the observed differences.

Significance

The k_{cat} values of enzymes are important for the study of metabolic systems. However, the current use of k_{cat} presents major difficulties, as values for most enzymes have not been experimentally measured, and experimentally available values are often measured under nonphysiological conditions, thereby casting doubt on the relevance of k_{cat} under in vivo conditions. We present an approach that utilizes omics data to quantitatively analyze the relationship between in vitro k_{cat} values and the maximal catalytic rate of enzymes in vivo. Our approach offers a high-throughput method to obtain enzyme kinetic constants, which reflect in vivo conditions, and are useful for more accurate and complete cellular metabolic models.

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Results

Integrating Enzyme Abundances and Fluxes to Calculate the Catalytic Rate of Enzymes in Vivo. Kinetic models of metabolic pathways and networks often describe the maximal rate of an enzyme-catalyzed reaction with the V_{\max} parameter:

$$v = \overbrace{E \cdot k_{\text{cat}}}^{V_{\max}} \cdot \eta, \quad [1]$$

where v is the flux through the reaction in the considered system, E is the overall number of enzyme active sites in the system, and η is a condition-dependent function, ranging between 0 and 1, which describes the decrease in the catalytic rate (relative to the maximum – k_{cat}), due to, for example, incomplete saturation with substrate and backward flux from thermodynamic effects (21).

Here we examine *E. coli* proteomic data measured via mass spectrometry to disentangle k_{cat} from V_{\max} (Eq. 1). We begin by calculating k_{app} , a previously introduced measure of the in vivo catalytic rate of enzymes (20) (Eq. 2; Fig. 1A). For a metabolic reaction catalyzed by a unique enzyme, there is almost always proportionality between v and E , given condition C . As such, we define:

$$k_{\text{app}}(C) \equiv \frac{v(C)}{E(C)}. \quad [2]$$

We can thus explicitly see the relationship between V_{\max} and k_{app} :

$$v(C) = \underbrace{E(C) \cdot k_{\text{cat}} \cdot \eta(C)}_{k_{\text{app}}} \quad [3]$$

Note, that because k_{cat} values are measured in vitro, such projection requires the consideration of effects caused by the shift between in vitro and in vivo, as discussed below.

Proteomic measurements provide abundances of individual polypeptides in cells, but enzymes are often composed of multiple subunits. Therefore, to infer the catalytic rate of enzymes per active site (as traditionally defined), we collected data on subunit and active site stoichiometry for *E. coli* enzymes (22) (Dataset S1). When the enzyme contains a single active site per

subunit, E equals the measured abundance of the polypeptide. To calculate k_{app} for multimeric enzymes, we divide the copy number of the polypeptide by the number of chains required to make an active site.

As discussed above, we focus our analysis on reactions catalyzed by unique homomeric enzymes, which constitute about 60% of the enzymatic reactions in iJO1366, the recent genome-scale reconstruction of *E. coli* metabolism (23). Reactions that are catalyzed by multiple distinct enzymes are difficult to analyze in this framework because we do not know how flux is partitioned across isozymes. Similarly, heteromeric enzymes composed of multiple distinct polypeptides complicate the analysis because it is often unclear which subunits contain active sites. To circumvent the challenge posed by heteromeric enzymes, one may consider k_{app} to be the catalytic rate per milligram of enzyme complex rather than the rate per active site. This definition corresponds to the notion of “specific activity” (as opposed to turnover number, k_{cat}). All analyses below give similar results when the maximal specific activity is used instead of k_{cat} (Dataset S1 contains the values of all such cases). Generalization of k_{app} for reactions catalyzed by isozymes is addressed in the [Supporting Information](#).

We use polypeptide abundances from recent proteomic studies of *E. coli* in 31 conditions (20, 24, 25), comprising various carbon sources, stress conditions and glucose-limited chemostats (Dataset S1). Given polypeptide abundances, flux measurements or computational flux predictions can be used to calculate k_{app} . To get comprehensive coverage of the metabolic network across conditions, here we use flux balance analysis (FBA) to calculate intracellular fluxes (26, 27). FBA is a mathematical approach for analyzing the rates and stoichiometry of transformations of metabolites through a metabolic network. The reactions are integrated into a stoichiometric matrix to be evaluated through linear programming (28). We chose to use a variant of FBA that minimizes the total flux through the network, and constrains the model by the growth rate and media composition, according to the measurements reported for each condition in the proteomics experiments (see [Materials and Methods](#) and [Supporting Information](#) for further details). A parallel analysis using experimental flux measurements (29) is shown in the [Supporting Information](#), and is in good agreement with FBA derived fluxes (log correlation of $r^2 = 0.85, p < 10^{-5}$), however, with much smaller coverage than FBA (only 13 enzyme-reaction pairs in four growth conditions). Experimental flux measurements are

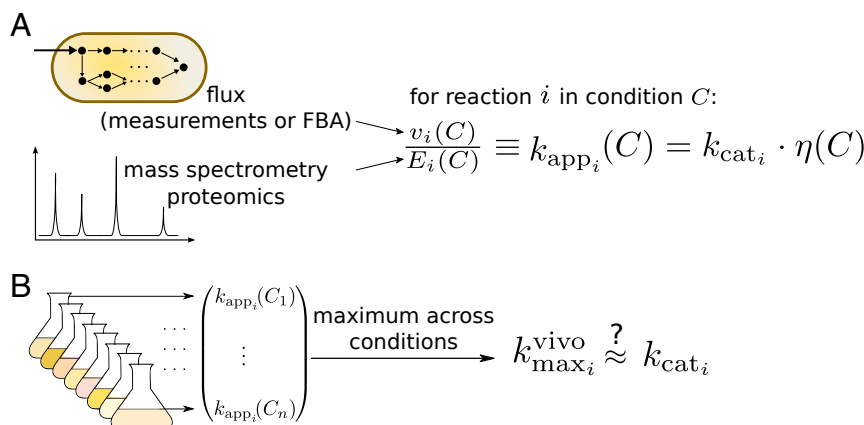


Fig. 1. Using omics data to estimate the catalytic rate of enzymes in vivo. (A) Flux (v) and enzymatic active site abundance (E) are integrated to calculate the rate of a single enzyme. Because v and E will change between conditions, the catalytic rate, k_{app} , is defined per condition (C). From Eq. 1, k_{app} equals the maximal rate (k_{cat}) times a condition-dependent function η . (B) For each metabolic reaction in the cell, there exists a condition in which the catalytic rate of enzyme i is maximal. $k_{\text{max}_i}^{\text{vivo}}$ is the maximal value of k_{app} , over many growth conditions ($N=31$ in this study), and represents a lower bound estimate of the maximal catalytic capacity of enzyme i in vivo. The relation between in vivo $k_{\text{max}_i}^{\text{vivo}}$ and in vitro k_{cat_i} , i.e., the maximal value of η , is investigated in this study.

currently limited in their applicability, and could be used in the future as depicted here, when data becomes available.

We calculate k_{app} values for 436 enzymes. To minimize stochastic effects we analyzed only enzymes with at least 10 copies of the polypeptide chain per femto liter (fL) of cytoplasm [roughly the volume of *E. coli* (30); *Materials and Methods*]. Notably, the turnover rate of an enzyme can vary between the forward and backward direction of the reaction and so k_{app} is defined for a specific enzyme-reaction-direction combination.

In Vivo and in Vitro Maximal Rates Show High Similarity. The catalytic capacity of *E. coli* cells varies with their physiology, i.e., with the growth rate and nutrient conditions (20, 31, 32). Given such variation across conditions, how can we infer the maximal catalytic rates of enzymes in vivo? We obtain a proxy for the maximal catalytic rate of enzymes in vivo by calculating k_{app} values in many conditions and taking the maximum value (Fig. 1B). Effectively, we look for the condition in which η is maximal (Eq. 3), and denote the value of k_{app} in this condition as k_{max}^{vivo} .

In vitro k_{cat} measurements are often used to describe the maximal catalytic rate of enzymes in metabolic models. How close are k_{max}^{vivo} values to in vitro k_{cat} measurements? To perform a global analysis, we mined the literature for k_{cat} values of *E. coli* enzymes. We collected a set of 146 reported k_{cat} values from the BRENDA database (33) and other publications, each pertaining to a specific enzyme-reaction pair. Manual curation of the data was required to ensure values were reported per active site and to verify that the literature value was correctly entered into the database (with errors mostly resulting from incorrect unit conversion; list of all curated k_{cat} values with references is provided in Dataset S1).

We calculated k_{max}^{vivo} predictions for 132 of the 146 enzyme-reaction pairs, for which k_{cat} values were available (the other 14 were expressed in too low copy numbers or supported no flux). These 132 enzymatic reactions span a wide range of metabolic pathways, including glycolysis, pentose-phosphate pathway, amino acid metabolism, nucleotide biosynthesis, biotin biosynthesis, and heme biosynthesis. As shown in Fig. 2, we find a correlation of $r^2 = 0.62$ in log scale ($p < 10^{-10}$) between in vivo and in vitro values. Both k_{max}^{vivo} and k_{cat} values span a similar range of roughly six orders of magnitude, with a root mean square difference (RMSD) of 3.5-fold in linear scale. Analysis using the second largest value of k_{app} instead of k_{max}^{vivo} shows similar correlation to k_{cat} ($r^2 = 0.59$, $p = 10^{-10}$), indicating that our k_{max}^{vivo} approximation is quite robust to the specific conditions measured.

In the face of these high global correlations, we also observe large deviations between k_{max}^{vivo} and k_{cat} for specific enzymes. A remarkable example is the enzyme malate dehydrogenase (*mdh*), whose k_{max}^{vivo} is $\approx 10 \text{ s}^{-1}$ compared with a k_{cat} of $\approx 1,000 \text{ s}^{-1}$. Indeed, it has been suggested that *mdh* evolved a high k_{cat} value to compensate for a thermodynamically unfavorable reaction in the direction of the tricarboxylic acid cycle, which causes much backward flux in vivo. This may result in reduction in the k_{max}^{vivo} , which is not accounted for under in vitro conditions (21). Other outliers serve as good candidates for in-depth biochemical analysis. In the Supporting Information we also discuss the relation between the number of allosteric regulators and the residual between k_{max}^{vivo} and k_{cat} .

Saturation and Thermodynamics Improve the Correspondence Between in Vitro and in Vivo Rates. To rationalize the difference between k_{max}^{vivo} and k_{cat} values, we consider η , the condition dependent term in the rate law (Eq. 1), as the product of (i) the degree of saturation of the enzyme, (ii) an effect of the thermodynamic driving force of the reaction, leading to backward flux, and (iii) vivo-vitro effects, which account for regulation and other ambient factors including pH, temperature, crowding agents, cofactors, chaperones, etc. (Fig. 3). Saturation and thermodynamic effects are functions of

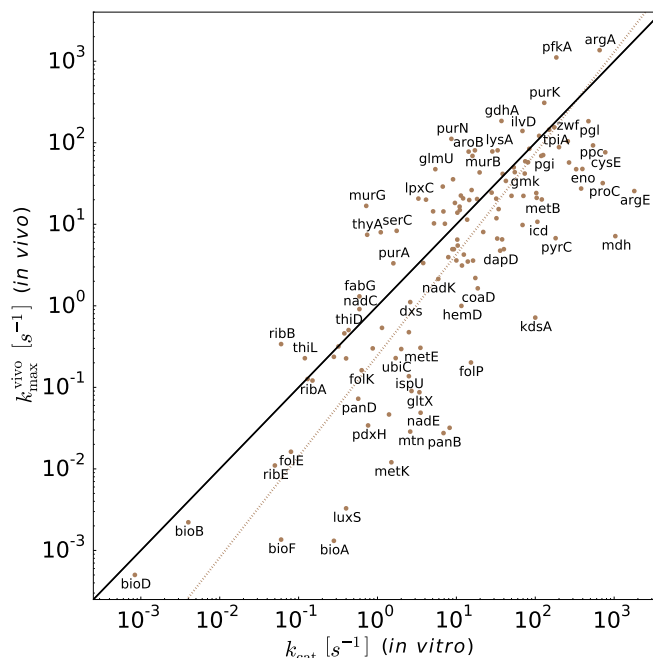


Fig. 2. In vivo and in vitro maximal catalytic rates. Log-log plot of k_{max}^{vivo} values versus in vitro k_{cat} values for all measured enzymes in *E. coli* ($N = 132$). Each data point represents an enzyme-reaction-direction combination and is labeled by the name of the enzyme (if not overlapping with other labels; data labels appear just above the data points for points that are above the line), and just below the data points for points that are below the line. The $y = x$ line is shown in black. Values show a correlation of $r^2 = 0.62$ ($p < 10^{-10}$) and span a similar range of 6 orders of magnitude. Dashed brown line represents the best fit by total least squares in log₁₀ scale with slope = 1.23 ± 0.07 , intercept = -0.6 ± 0.1 , and an RMSD of 0.54.

metabolite concentrations and will range between 0 and 1, representing how much of the in vitro capacity (i.e., k_{cat}) is used (34–36). Regulatory and environmental effects, however, may result in higher rates in vivo compared with in vitro (e.g., due to absence of chaperones in the test tube) or vice versa (e.g., suboptimal pH in the cytosol). Fig. 3 shows how one can apply this rationale to interpret the residual between k_{cat} and k_{max}^{vivo} .

Although the vivo-vitro effects may represent key characteristics of enzyme catalysis inside the cell, current knowledge is not enough to fully account for them. We can, however, calculate the decrease in rate caused by nonsaturating substrate levels and backward flux, thereby assessing these differences attributed to the cellular environment. For this we use metabolite concentrations measured in *E. coli* (37, 38) (Dataset S1). For each enzyme, we sought metabolite concentrations measured in the same conditions as the proteomics data ultimately used to calculate k_{max}^{vivo} .

To estimate the degree of saturation, we collected K_M values of enzymes from the BRENDA (33) and EcoCyc (39) databases (Dataset S1), and used a relation derived from the irreversible Michaelis-Menten rate law (40, 41) for multiple substrates (42):

$$\text{saturation term} = \frac{\prod_j \left(\frac{s_j}{K_j} \right)^{m_j}}{1 + \prod_j \left(\frac{s_j}{K_j} \right)^{m_j}}, \quad [4]$$

where s_j is the concentration of substrate j , and K_j and m_j are its K_M and stoichiometric coefficient, respectively.

None of *E. coli* enzymes have fully measured (i) k_{cat} values, (ii) K_M values for both substrates and products, and (iii) substrate and product concentrations. Hence, we calculate the saturation

which may prove useful for better understanding the regulation of metabolic networks (54–56).

The $k_{\text{max}}^{\text{vivo}}$ value is obtained directly from flux prediction and enzyme abundance data and requires no parameter fitting. Large-scale absolute proteome quantification is advancing in great strides and becoming ubiquitous. It may soon be possible to consider hundreds of conditions in calculating the maximum k_{app} value. Genomewide estimation of maximal rates could prove invaluable for metabolic engineering applications, for example, to select the fastest enzyme from a collection of orthologs (57, 58), or compare between alternative synthetic pathways (59–61). In addition, tuning the expression levels of enzymes in a recombinantly expressed pathway will necessarily require information on their specific activities. As demonstrated here, the availability of absolute proteome quantification provides the missing information required to accelerate the collection of enzyme kinetic data.

Materials and Methods

Computational Flux Predictions and Proteomics Data as Inputs for k_{app} Estimates.

Flux estimations were performed using parsimonious FBA (pFBA) (62, 63) using the COBRApy package (64). The pFBA analysis minimizes the total flux through the network and enforces reaction directionality constraints that exist in the model, and was shown to be consistent with measured fluxes (63, 65). We constrained the model by the growth rate and media composition, as was measured in the proteomics experiment. For each reaction, we analyzed the variation in flux via Flux Variability Analysis [FVA (66)]. For 99% of reactions the flux variability was negligible, representing less than 1% of the net flux through the reaction (Supporting Information). The flux through the ATP maintenance reaction was also shown to have negligible effect on the results (Supporting Information). For integration of proteomics data we used gene annotations from the iJO1366 model to map polypeptide chains to metabolic reactions. In Valgepea et al. (20) and Peebo et al. (25) data were reported in copies per fL cytoplasm, given condition specific cell size measurements. In Schmidt et al.'s dataset (24), values were given in copies per cell, hence we converted to copies per fL using a characteristic value of $180 \text{ fg}/\mu\text{m}^3$ protein concentration in *E. coli* (67). Though this value may vary on the order of 10% across conditions, there is currently no comprehensive information on this variation across conditions, and in any case this will be a minor effect given the

variation in expression levels. Out of 436 homomeric enzymes expressed at a level of at least 10 polypeptide chains per fL, 257 support flux in at least one condition, resulting in $k_{\text{app}} > 0$. Notably, the 179 apparently idle enzymes might be an artifact of minimizing the total flux, and in reality may carry flux. By using a flux prediction method that suggests many zero values, we restrict ourselves to values that are relatively reliable. Enzymes that are expressed and do not support flux are not included in the analysis. All numerical data are available at Dataset S1. Codes are open-source and available at <https://github.com/milo-lab/in-vivo-catalytic-rates> (implemented in Python and tested mainly on Linux).

Data Collection and Unit Conversions for Comparison Between k_{cat} and $k_{\text{max}}^{\text{vivo}}$

Values. We collected a comprehensive set of k_{cat} values from BRENDA and primary literature. Where multiple values were reported for the same k_{cat} we took the maximal reported value. The units of k_{cat} are 1/s, hence flux values were converted from millimole per gram cell dry weight per hour (mmol/gCDW/h) to millimole per gram cell dry weight per second (mmol/gCDW/s). Abundance was converted from molecules/fL to mmol/gCDW, assuming dry weight fraction of 30% [BNID:109838 (68, 69), and a cell density of 1.1 g/mL (BNID:103875 (69, 70)]. Thus, dividing flux by abundance results in rate in units of 1/s. For specific activities, we used the molecular weight of enzymes to infer the rate in units of μmol product per mg enzyme per minute (see Supporting Information for further details).

Calculating the Backward Flux of Reactions. To calculate $\Delta_r G^\circ$ we used the component contribution method, which integrates measured thermodynamic data to calculate the formation energies of metabolites (43, 44). By including measured metabolite concentrations we approximate the in vivo Gibbs free energy of a reaction, which is a more accurate way to assess reversibility and the thermodynamic driving force than examining the standard Gibbs energy (43).

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