

# Theoretical investigation of a genetic switch for metabolic adaptation

Kathrin S. ~~Laxhauber~~Laxhuber<sup>1</sup>, Muir J. Morrison<sup>2</sup>, Rob Phillips<sup>2,3\*</sup>

<sup>1</sup> ~~Dept???, ETHZ, Zurich, State???, Department of Chemistry and Applied Biosciences, ETH Zurich, 8093 Zurich, Switzerland~~

<sup>2</sup> Department of Physics, California Institute of Technology, Pasadena, CA, USA

<sup>3</sup> Department of Biology and Biological Engineering, California Institute of Technology, Pasadena, CA, USA

## Abstract

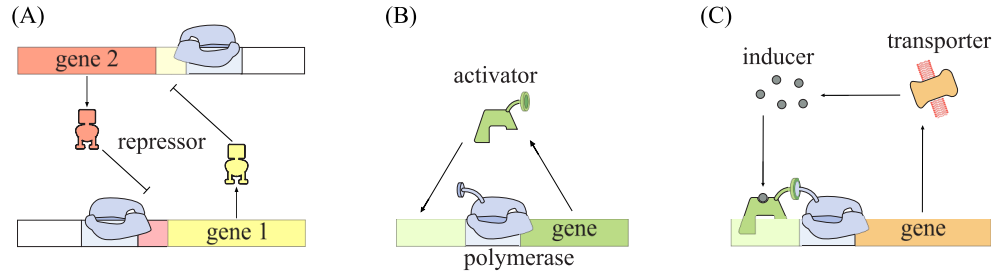
Membrane transporters carry important molecules through the cell membrane and, from a resource standpoint, should only be produced when necessary. The expression of membrane transporters in metabolic pathways is often upregulated by the transporter substrate. In *E. coli*, such systems include for example the *lacY*, *araFGH*, and *xylFGH* genes, which encode for lactose, arabinose, and xylose transporters, respectively. As a case study of a minimal system, we build a generizable physical model of the *xapABR* genetic circuit, which features a regulatory feedback loop through membrane transport (positive feedback) and enzymatic degradation (negative feedback) of an inducer. Dynamical systems analysis and stochastic simulations show that the membrane transport makes the model system bistable in certain parameter regimes. Thus, it serves as a genetic “on-off” switch, enabling the system to only produce a set of metabolic enzymes when the corresponding metabolite is present in large amounts. We find that the negative feedback from the degradation enzyme does not significantly disturb the positive feedback from the membrane transporter. We investigate hysteresis in the switching and discuss the role of cooperativity and multiple binding sites in the model circuit. AltogetherFundamentally, this work explores how a stable genetic switch for a set of enzymes is obtained from transcriptional auto-activation of a membrane transporter through its substrate.

## Introduction

remove  
images &  
todonotes  
everywhere  
for submission

Genetic regulatory circuits are fundamental building blocks of functioning cells and organisms. One abundant class of these circuits are genetic switches. Although their construction and function may differ, their common feature is bistability: their output gene expression will flow to and remain at one of two steady-state levels. The distribution of gene expression in a cell culture can then be bimodal. This is not to be confused with mere stochastic bimodality, where the system is not stable, and the gene expression in each cell can fluctuate between the two levels.

One classic example of a genetic switch is a system where two repressor proteins each regulate the transcription of the other [1, 2] (illustrated schematically in Fig 1). Here, one stable state is high expression of the first protein and low expression of the second, and the second stable state is the opposite. This switch enables the system to have a memory: if something induces expression of either one of the proteins, the



**Fig 1. A schematic of different genetic switches.** (A) and (B) show the two most well-known genetic switches: (A) two mutual repressors and (B) a self-activating gene. In (C), a very much simplified version of the circuit that we investigate in this paper can be seen, where the similarity to the switch in (B) ~~becomes obvious~~ is clear. A complete version of the model circuit can be found in Fig 2.

system will remain in this state until a significant perturbation occurs. Another well known and even simpler example is an auto-activating circuit in which a protein activates its own transcription [3]. This gives the system an “on-off” switch.

Through physical and mathematical modeling, we investigate a more complex ~~but common~~ switch system where the bistability is due, as we will show, to a membrane transport protein. ~~Its key feature~~ Such a switch is common for metabolic processes in biology, for reasons discussed below. So far, models we have found are either very extensive descriptions of specific, complicated systems, or they are very basic. To enable an intuitive and fundamental understanding of the behavior, we develop a simple model of a minimal system that only has the essential components and interactions for the questions we pose, while still modeling every component explicitly and discussing the necessary model complexity for a physically correct model.

The key feature of the type of system we investigate is the indirect activation of the transporter gene by the transporter substrate, leading to positive feedback similar to the aforementioned “on-off” switch. An example for such an architecture is the *lac* operon, where lactose indirectly activates the expression of lactose permease. Other examples in *E. coli* include the *araFGH* and *xylFGH* operons, which contain genes for arabinose and xylose transporters, respectively. For *lac* and *araFGH*, bistability has indeed been observed and attributed to such a positive feedback loop ~~[4, 6–11]~~ [4–11]. A eukaryotic example is the glucose transporter GLUT-2 in liver and  $\beta$ -cells [12, 13], though this system is much more complex than the following analysis.

It is quite conceivable that this auto-activation process is common to many substances that a cell would want to accumulate. Such a switch enables the cell to sense and respond to its environment: if the substrate enters the cell, it activates the production of membrane transporters. The cell then starts accumulating the substrate, thereby “testing” the substrate’s presence in the extracellular environment. If there is enough, the expression stabilizes at an “on” point and the cell has, in a short-term sense, adapted. When there is not enough substrate ~~anymore~~, the operon, which often encodes for a whole set of enzymes for this one metabolite, switches “off” again. Such a mechanism could be involved in various cases of short-term adaptation.

A key element of this mechanism is the presence of a transcription factor which is always expressed at a low level (often at copy numbers of order  $\sim 10$ ) and which binds to the transporter substrate. This is resource efficient for the cell, as this low copy number transcription factor acts as an “always on” sensor to detect the substrate, allowing high copy numbers of the membrane transporter and its attendant operon to be expressed only when their substrate is actually present. The transcription factors LacI, AraC, and XylR all appear to fill this role ~~[4, 6–11]~~ [4–11, 14, 15].

For our modeling, we focus on the *xapABR* genetic circuit from *E. coli* as a case study. It is similar to *lac*, but less complex. Instead of lactose, its purpose is to make use of the nucleoside xanthosine as an energy source [16, 17]. The circuit is made up of two operons: one that encodes for XapR and another that encodes for XapA and XapB. XapR appears to be a transcription factor that is induced by xanthosine and activates the *xapAB* promoter, in close analogy to ~~LacI, AraC, and XylR~~[AraC, XylR, and also LacI, although the latter acts via repression](#). The *xapAB* promoter has been suggested to have two binding sites for XapR [18], but the promoter architecture and function is not yet fully understood. The transcription of *xapR* seems to be constitutive and not auto-regulated [18]. Structural homology to other transcription factors suggests that XapR appears in dimers where one dimer can bind two xanthosine molecules [19]. The protein XapA is a purine nucleoside phosphorylase that degrades xanthosine into components (ribose and xanthine) that can be fed into metabolic pathways [16, 17]. XapB on the other hand is a membrane transporter of xanthosine [18, 20].

Experimentally it was found that the expression level of *xapAB* among cells follows a bimodal distribution and that the system seems to be bistable [21]. We aim to understand which of the circuit's features are necessary for bistability and investigate its behavior in different parameter regimes. In the following section, we will discuss the details of our model. After estimating the free parameters we then present the observations we made through phase diagrams, followed by the results from stochastic simulations.

## Model

### Step by step modeling of the system

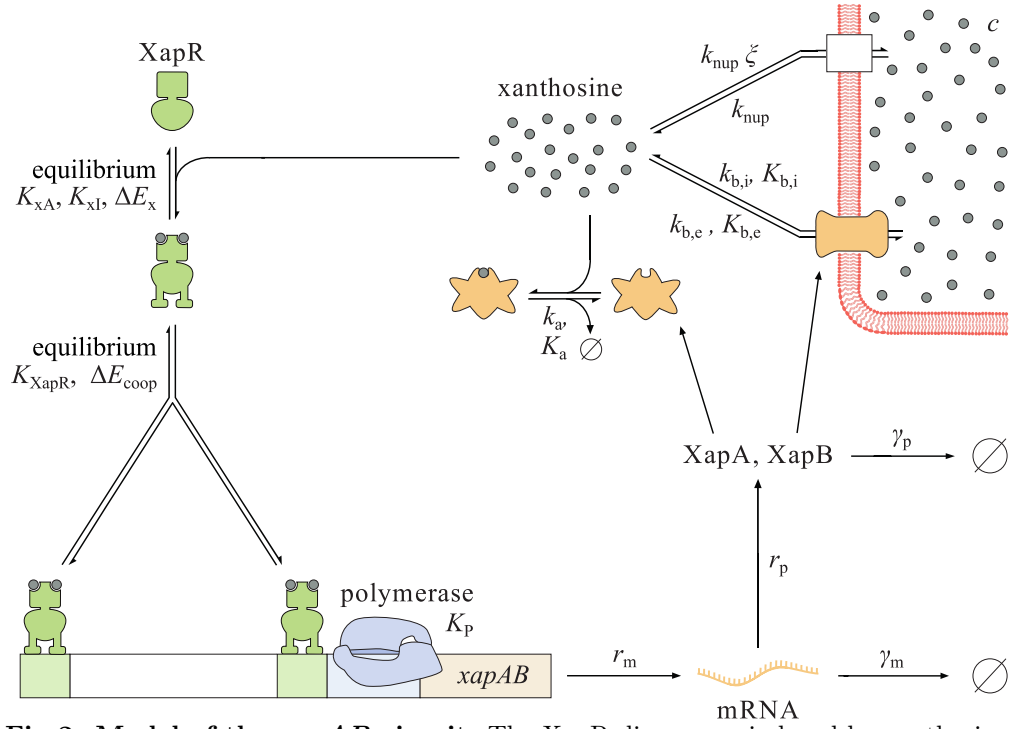
In this section, we present our model of the *xapAB* genetic circuit. Fig 2 shows an overview of this model. The qualitative picture of the circuit switching its state is as follows:

- In the initial absense of XapB, small amounts of xanthosine are nonspecifically transported into and out of the cell (discussed in more detail below).
- XapR is induced by xanthosine.
- The induced XapR binds to the *xapAB* promoter, leading to transcription.
- From this mRNA transcript, translation produces the two proteins XapA and XapB.
- XapB actively transports much larger amounts of xanthosine into the cell and XapA degrades xanthosine.
- Production of more XapA & XapB is balanced by decay or dilution through cell divisions.

“permeate”  
wouldn’t  
include ac-  
tive trans-  
port (NupC  
& NupG),  
would it?

Because xanthosine induces the transcription factor XapR, we have positive and negative feedback loops due to XapB and XapA, respectively. The remainder of this subsection discusses each of the above steps in detail, leading us to a set of two coupled ODEs. More in-depth explanations can be found in S1 Text.

**Induction of XapR.** We treat dimers as the only form of XapR that appears in the cell. Each dimer can bind two xanthosine molecules. The Monod-Wyman-Changeux



**Fig 2. Model of the *xapAB* circuit.** The XapR dimers are induced by xanthosine and the induced XapR binds cooperatively as an activator to the *xapAB* promoter. For these two steps, quasi-equilibrium is assumed. If both XapR binding sites are occupied and the polymerase is bound, the gene is transcribed at rate  $r_m$ . The mRNA decays at rate  $\gamma_m$ , and both proteins are translated at rate  $r_p$  and decay at rate  $\gamma_p$ . XapA degrades xanthosine with Michaelis-Menten parameters  $k_a$ ,  $k_a$  and  $K_a$ ,  $K_a$ . Similarly, XapB imports and exports xanthosine with parameters  $(k_{b,i}, K_{b,i})$  and  $(k_{b,e}, K_{b,e})$ , respectively. Furthermore, xanthosine enters and leaves the cell through non-specific transport, proportional to rates  $k_{nup}$  and  $\xi k_{nup}$ , respectively.

(MWC) model is used to describe the fraction of XapR dimers in the active state, which has the form

$$[\text{XapR}]_A = [\text{XapR}]_{\text{tot}} \frac{\left(1 + \frac{[x]}{K_{xA}}\right)^2}{\left(1 + \frac{[x]}{K_{xA}}\right)^2 + e^{\beta \Delta E_x} \left(1 + \frac{[x]}{K_{xA}} \frac{K_{xA}}{K_{xI}}\right)^2}. \quad (1)$$

Here,  $[x]$  is the xanthosine concentration, and  $[\text{XapR}]_A$  and  $[\text{XapR}]_{\text{tot}}$  denote the concentration of active and total XapR dimers, respectively. Furthermore,  $K_{xI}$  and  $K_{xA}$  are the dissociation constants of xanthosine to the inactive and the active XapR dimer, respectively, and  $\Delta E_x$  stands for the energy difference between the inactive and the active state. We expect  $\Delta E_x > 0$  and  $K_{xA} < K_{xI}$  for inducible activation. This corresponds to XapR being mainly inactive in the absence of xanthosine and becoming mostly active at high concentrations of xanthosine. A detailed discussion of the MWC model can be found in [22].

**Transcription.** Transcription and translation of the *xapAB* gene, regulated by the induced XapR, produce the two proteins XapA and XapB. We start with transcription and assume that the binding of XapR and polymerase to the promoter is at quasi-equilibrium. The polymerase binding is modeled as independent of that of XapR,

State	Weight
	1
	$\frac{[\text{XapR}]_A}{K_{\text{XapR}}}$
	$\frac{[\text{XapR}]_A}{K_{\text{XapR}}}$
	$\left(\frac{[\text{XapR}]_A}{K_{\text{XapR}}}\right)^2 e^{-\beta \Delta E_{\text{coop}}}$
	$\frac{[\text{P}]}{K_{\text{P}}}$
	$\frac{[\text{P}]}{K_{\text{P}}} \frac{[\text{XapR}]_A}{K_{\text{XapR}}}$
	$\frac{[\text{P}]}{K_{\text{P}}} \frac{[\text{XapR}]_A}{K_{\text{XapR}}}$
	$\frac{[\text{P}]}{K_{\text{P}}} \left(\frac{[\text{XapR}]_A}{K_{\text{XapR}}}\right)^2 e^{-\beta \Delta E_{\text{coop}}}$
XapR Polymerase	

**Fig 3. The promoter states.** We consider only the completely occupied state as active and all other states (faded out in the figure) as completely inactive. The parameters are the interaction energy of the two XapR dimers  $\Delta E_{\text{coop}}$  and the dissociation constants  $K_{\text{XapR}}$  and  $K_{\text{P}}$  of XapR and polymerase to the promoter, respectively. The concentrations of polymerase and active XapR are denoted by  $[\text{P}]$  and  $[\text{XapR}]$ .

and all influence of the activator is pushed into the transcription rate. Furthermore, the binding energy of XapR to each of its two sites is assumed to be the same. A discussion of these simplifications can be found in S1 Text.

In Fig 3, all possible states of the promoter in our model and their corresponding thermodynamic weights are shown.  $[\text{P}]$  denotes the polymerase concentration, and  $\Delta E_{\text{coop}}$  stands for the interaction energy of the two XapR dimers. If cooperativity in transcription factor binding is neglected, this is set to zero. Furthermore,  $K_{\text{XapR}}$  and  $K_{\text{P}}$  denote the dissociation constant of XapR and polymerase to the promoter, respectively. In statistical mechanics language these dissociation constants are equivalent to  $\frac{N_{\text{NS}}}{V} e^{\beta \Delta E_{\text{XapR}}}$  and  $\frac{N_{\text{NS}}}{V} e^{\beta \Delta E_{\text{P}}}$ , respectively, with  $N_{\text{NS}}$  being the number of non-specific binding sites on the DNA,  $V$  the volume of the cell, and  $\Delta E_{\text{XapR}}$  and  $\Delta E_{\text{P}}$ , respectively, the interaction energies of XapR or polymerase with the promoter.

enumerate  
states in  
figure

We consider only the state where both XapR binding sites are occupied as active and all other states as inactive, meaning they have transcription rate equal zero. Experiments show that the expression becomes very weak when one of the XapR binding sites is removed from the promoter, suggesting that this simplification is reasonable [21]. Furthermore, we find that in the bistable parameter range, considering the single occupancy states as active instead has almost no influence on the results (see also S1 Text).

With  $[\text{m}]$  being the mRNA concentration,  $r_{\text{m}}$  the transcription rate,  $\gamma_{\text{m}}$  the mRNA decay rate, and  $p_{\text{active}}$  the probability of the promoter being in the active state, we

obtain

$$\frac{d[m]}{dt} = r_m p_{\text{active}} - \gamma_m [m] \quad (2)$$

$$p_{\text{active}} = \frac{w_8}{\sum_{i=1}^8 w_i} = \frac{[P]}{K_P + [P]} \frac{\left(\frac{[XapR]_A}{K_{XapR}}\right)^2 e^{-\beta \Delta E_{\text{coop}}}}{1 + 2 \frac{[XapR]_A}{K_{XapR}} + \left(\frac{[XapR]_A}{K_{XapR}}\right)^2 e^{-\beta \Delta E_{\text{coop}}}} \quad (3)$$

Here,  $w_i$  stands for the thermodynamic weight of the  $i$ th state in the order in which they are listed in Fig 3. As written above, the partition function factorizes into a polymerase and a XapR term because of our assumption of independent binding, which is further discussed in S1 Text. Note that because  $r_m$  implicitly contains the gene copy number per cell, it has units of  $M^{-1} s^{-1}$  and not just  $s^{-1}$ . This rate equation gives the mean mRNA concentration  $\langle [mRNA] \rangle = \frac{r_m}{\gamma_m} p_{\text{active}}$ , which we will need in the next paragraph. ~~In, The mean can also be found from the full chemical master equation is shown, which leads to a Poisson distribution of the mRNA concentration with the same mean, which is shown in S1 Text.~~

**Translation.** The next step in our modeling progression is translation. As a simplification, we write  $[p] = [XapA] = [XapB]$  for the general protein concentration. This assumes that the rates of transcription, mRNA decay, translation, and protein decay are the same for both proteins, which, as discussed in S1 Text, does not have a significant influence on the results. We write the following rate equation for the protein concentration, where  $r_p$  denotes the translation rate,  $\gamma_p$  the protein decay rate, and  $\langle [mRNA] \rangle$  the mean mRNA concentration:

$$\frac{d[p]}{dt} = r_p \langle [mRNA] \rangle - \gamma_p [p]. \quad (4)$$

**Xanthosine dynamics.** Having described how xanthosine activates the synthesis of XapA and XapB through XapR, we now close the feedback loop by setting up a xanthosine rate equation.

~~Transport of~~ There are two significant ways of transport of xanthosine across the cell membrane happens in two steps which we will discuss now. All statements in this paragraph are based on data from [20]. In a first transport step, xanthosine passes the outer membrane through porins like Tsx, OmpF and OmpC. This does not seem to be a rate-limiting step. Then, XapB actively transports it across the inner membrane, powered by the proton motive force. There are two other active nucleoside transporters that seem to be transporting xanthosine with. In the induced system, the main transporter is XapB, whereas in the uninduced system, there is almost no XapB. Instead, xanthosine can enter the cell through the two nucleoside transporters NupC and NupG, which have a very low affinity for NupC and NupG. It was found that  $\Delta nupC \Delta nupG$  strains cannot grow on xanthosine. Hence, these seem to be necessary to “start” the system by importing small amounts of xanthosine that then activate *xapB*, and there appears to be no other significant way in which xanthosine can enter the cell.

~~We assume the kinetic scheme of all three transporters for xanthosine [20]. All these transporters, XapB, NupC, and NupG, are powered by the proton gradient across the membrane [20], which is why we assume their kinetic scheme to be similar to that of the *lac* permease (as it is described in [23]). One transporter can then do two things to change the intracellular xanthosine concentration  $[x]$ : it can actively transport one xanthosine into the cell, powered by the proton gradient, or it can transport one out of the cell, against the proton gradient. For net transport There can be import and~~

export of xanthosine, and which one dominates depends on the proton and xanthosine concentrations on the two sides of the membrane. In both cases, a proton and a substrate need to bind to the transporter on one side of the membrane and detach from the transporter it on the other side of the membrane before the transporter goes before the empty transporter moves back to the first side again (see also other side again. We refer the reader to S1 Text ). Because of the proton gradient, influx is overall thermodynamically more likely than export at low intracellular xanthosine concentrations, which leads to a net influx. For much higher intra- than extracellular xanthosine concentrations, the difference in the chemical potential of xanthosine across the membrane can dominate that of the protons and there is net efflux.

for a detailed description of the transport and its modeling, and just state the result here. We model influx and efflux separately and. For XapB, we use Michaelis-Menten kinetics with parameters  $k_{b,i}$ ,  $K_{b,i}$  for influx and  $k_{b,e}$ ,  $K_{b,e}$  for XapB. For the turnover rate and Michaelis constant for influx we write  $k_{b,i}$  and  $K_{b,i}$ , respectively, and  $k_{b,e}$ ,  $K_{b,e}$  for efflux. These parameters implicitly include the proton gradient. For NupC and NupG, the Michaelis constant is very large, due to their low xanthosine affinity. Thus, the general  $k_{cat}[E]_0 \frac{[S]}{K_M + [S]}$  can be approximated as  $\frac{k_{cat}}{K_M} [E]_0 [S]$ . In the following,  $\frac{k_{cat}}{K_M} [E]_0$  is denoted by  $\xi$  for the Nup transporters, we can linearize because of their low affinity for xanthosine and are left with the rate parameters  $k_{nup}$  for influx, and the efflux rate is written as  $\xi k_{nup}$ . For the XapB kinetics, no approximations can be made, because the xanthosine concentration in the dynamic system can range from far below the respective  $K_M$  value to far above. The final terms for Nup and XapB are shown below for efflux.

After transport into the cell, XapA degrades xanthosine. We model this using standard Michaelis-Menten kinetics, with parameters  $k_a$ ,  $K_a$  (corresponding to turnover rate and Michaelis constant, respectively). All the above Transport and degradation then leads to the xanthosine rate equation

$$\frac{d[x]}{dt} = \underbrace{\left( k_{b,i} \frac{c}{K_{b,i} + c} - k_{b,e} \frac{[x]}{K_{b,e} + [x]} \right)}_{\text{XapB}} - \underbrace{k_a \frac{[x]}{K_a + [x]}}_{\text{XapA}} [p] + \underbrace{k_{nup} (c - \xi [x])}_{\text{NupC \& NupG}}. \quad (5)$$

We recall Recall that  $[x]$  is the intracellular xanthosine concentration, while  $c$  denotes the extracellular. Because  $k_{b,i} > k_{b,e}$  and  $K_{b,i} < K_{b,e}$ , influx dominates at low intracellular xanthosine concentrations. At much higher intra- than extracellular xanthosine concentrations, the efflux term takes over. More details on the aforementioned steps and a discussion of passive diffusion can be found in S1 Text.

## Nondimensionalization

We have now formulated the behavior of the system in terms of the rate equations for mRNA, protein, and xanthosine. These equations can be nondimensionalized, which reduces the dimension of parameter space. We measure time in units of  $\gamma_p^{-1}$  and concentrations in units of  $K_a$  (except XapR, where the equations make it more natural to use  $K_{XapR}$ ). In Table 1, all the nondimensional parameters and their definitions are listed. Furthermore, we define  $[m]_a := \frac{[m]}{K_a}$ ,  $[p]_a := \frac{[p]}{K_a}$ ,  $[x]_a := \frac{[x]}{K_a}$ , and  $\tau := \gamma_p t$ . Using



these definitions, the following equations are obtained

$$\frac{d[m]_a}{d\tau} = \rho_m \frac{[XapR]_{R,A}^2 e^{-\Delta\epsilon_{coop}}}{1 + 2[XapR]_{R,A} + [XapR]_{R,A}^2 e^{-\Delta\epsilon_{coop}}} - \gamma_{mp}[m]_a \quad (6)$$

$$\frac{d[p]_a}{d\tau} = \rho_p[m]_a - [p]_a \quad (7)$$

$$\frac{d[x]_a}{d\tau} = \left( k_{\beta,i} \frac{[c]_a}{K_{\beta,i} + [c]_a} - k_{\beta,e} \frac{[x]_a}{K_{\beta,e} + [x]_a} - k_{\alpha} \frac{[x]_a}{1 + [x]_a} \right) [p]_a + k_{\eta} ([c]_a - \xi[x]_a) \quad (8)$$

$$\text{with } [XapR]_{R,A} = [XapR]_R \frac{\left(1 + \frac{[x]_a}{K_{\chi A}}\right)^2}{\left(1 + \frac{[x]_a}{K_{\chi A}}\right)^2 + e^{\Delta\epsilon_x} \left(1 + \frac{[x]_a}{K_{\chi A}} \frac{1}{K_{IA}}\right)^2}$$

Very little is known about the *xap* system, and thus, there are almost no measured values for the free parameters. Nevertheless, we were able to estimate a reasonable range by using values from similar, well studied systems and by finding physical constraints or relations between parameters. The results of these estimates are shown in Table 1. They are based on a choice of  $\gamma_p = 5 \cdot 10^{-4} \text{ s}^{-1}$  and  $K_a = 5 \cdot 10^{-5} \text{ M}$ . A detailed derivation can be found in S1 Text.

## Results and discussion

In the modeling process in the previous section, we have obtained three coupled differential equations. In this section, we will analyze these equations with deterministic methods and stochastic simulations. Analytical closed-form solutions could not be obtained and would, if they existed, probably not be helpful due to their large complexity. Finding such solutions requires solving a fifth order algebraic equation.

### Deterministic phase portraits

A standard way to analyze dynamical systems deterministically is to plot phase portraits. In the following, we present such plots where the state variables are the mRNA, the protein, and the xanthosine concentration.

**From a 3D to a 2D system.** Fig 4A shows the 3D phase portrait for a representative set of parameters (shown in Table 1), whose choice is explained below. The plot looks rather complicated at first but can be understood intuitively. The three ~~planes are the nullclines~~ surfaces are the nullcline surfaces and the gray lines point in the direction in which the dynamical system moves at each point. The ~~planes~~ surfaces intersect in three points, which are the steady-state solutions of the dynamical system. For this choice of parameters, the system first flows towards the mRNA nullcline, then it moves along that ~~plane~~ surface to the intersection with the protein nullcline, and lastly, it moves along that intersection line to one of the three intersection points of all three ~~planes~~ surfaces.

It is important to point out that, for a different set of parameters, the dynamics can be quite different. There are, for example, scenarios where the xanthosine kinetics are roughly as fast as the mRNA kinetics and the dynamics unfolds in two steps: first to the intersection of the mRNA and the xanthosine nullcline, then along that line to the protein nullcline and thereby to a fixed point.

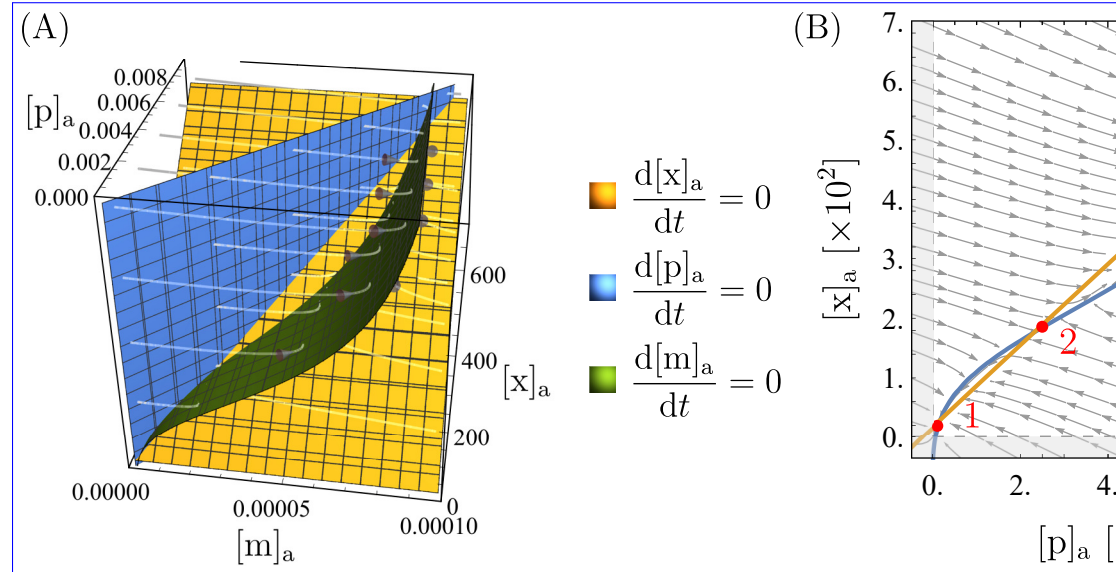
A usual simplification with genetic circuits like this is to assume the mRNA concentration to be at steady-state, i.e., to write  $\frac{d[m]_a}{d\tau} = 0$  and solve this for



**Table 1. Nondimensional parameters and their estimated values.**

Param.	Definition	Estimated range	Value used
$\rho_m$	$:= \frac{r_m}{\gamma_p K_a} \frac{[P]}{K_P + [P]}$	$\approx 10^{-3 \pm 2}$	$10^{-3}$
$\gamma_{mp}$	$:= \frac{\gamma_m}{\gamma_p}$	$\approx 10^{1 \pm 0.5}$	$10^1$
$\rho_p$	$:= \frac{r_p}{\gamma_p}$	$\approx 10^{2 \pm 0.5}$	$10^2$
$[XapR]_R$	$:= \frac{[XapR]_{tot}}{K_{XapR}}$	$\approx 10^{0 \pm 2}$	$10^0$
$[c]_a$	$:= \frac{c}{K_a}$	$(\in [0, 10^3])$	13
$k_{\beta,i}$	$:= \frac{k_{b,i}}{\gamma_p}$	$\approx 10^{4 \pm 1}$	$5 \cdot 10^4$
$k_{\beta,e}$	$:= \frac{k_{b,e}}{\gamma_p}$	$\approx 10^{3 \pm 2}$	$10^3$
$k_\alpha$	$:= \frac{k_a}{\gamma_p}$	$\approx 10^{2 \pm 0.8}$	$10^2$
$k_\eta$	$:= \frac{k_{nup}}{\gamma_p}$	$\approx 10^{0 \pm 3}$	$5 \cdot 10^{-1}$
$\xi$	$= \xi$	$\approx 0.8 \pm 0.1$	0.8
$K_{\beta,i}$	$:= \frac{K_{b,i}}{K_a}$	$\approx 10^{1 \pm 2}$	$10^1$
$K_{\beta,e}$	$:= \frac{K_{b,e}}{K_a}$	$\approx 10^{2 \pm 2}$	$10^2$
$K_{\chi A}$	$:= \frac{K_{\chi A}}{K_a}$	$\approx 10^{2 \pm 1} \cdot 10^{\Delta\epsilon_x - 5}$	$10^2$
$K_{IA}$	$:= \frac{K_{\chi I}}{K_{\chi A}}$	$\approx 10^{2 \pm 1}$	$10^2$
$\Delta\epsilon_x$	$:= \beta \Delta E_x$	$\approx 2 \text{ to } 2 (\ln(K_{IA}) - 1) < 12$	5
$\Delta\epsilon_{coop}$	$:= \beta \Delta E_{coop}$	$\approx 0 - 10$	5

The left column shows all nondimensional parameters that appear in the final equations. In the middle are their definition and estimated values. They are based on  $\gamma_p = 5 \cdot 10^{-4} \text{ s}^{-1}$  and  $K_a = 5 \cdot 10^{-5} \text{ M}$ . Note that the range of the three MWC parameters depends on each other, but they can still be chosen independently. The range given for  $[c]_a$  denotes the estimated “interesting” range in which switching happens, but  $[c]_a$  can of course exceed these values. Details on the parameters and their estimation can be found in S1 Text. Finally, the last column shows the value that we use for the rest of this paper, unless otherwise noted. An explanation of this choice will follow in the next section.



**Fig 4. Phase portraits showing bistability.** 3D and 2D phase portraits for one set of parameters that leads to bistability. The parameter values are listed in Table 1. Note that all the concentrations ( $[m]_a$ ,  $[p]_a$ ,  $[x]_a$ ) are measured in units of  $K_a = 5 \cdot 10^4$  nM. The ~~planes~~ surfaces in (A) and the lines in (B) are the nullclines of the state variables, and their intersection points, marked in red in (B), are the steady-state solutions of the system. The region shaded in gray in (B) leads to negative concentrations and is unphysical. A vector plot of (B) that also shows the magnitude of flow at each point can be found in S1 Text.

$[m]_a([p]_a, [x]_a)$  to simplify the 3D to a 2D system. This restricts the dynamics to the green ~~plane~~ surface in our plot, which is reasonable here because as explained above, the system first flows towards that ~~plane~~ surface before either the protein or the xanthosine concentration changes significantly. However, as already pointed out, this is different for other parameter values, and thus, this assumption does not hold in general. If the xanthosine dynamics are faster than the mRNA dynamics, the system first flows towards the xanthosine nullcline. In that case, forcing it onto the mRNA nullcline leads to significant changes in the dynamics.

Nevertheless, the steady-state solutions and the qualitative features that we address in this paper remain the same. Because the 3D plots are rather hard to read, we will, in the following, make the compromise to show a 2D version of the phase portraits but ensure that all of our statements also hold true in 3D space. As explained above, it makes the most sense here to do this by setting  $\frac{d[m]_a}{d\tau} = 0$ . The resulting equations can be found in S1 Text. In particular, we define  $\rho := \frac{\rho_m \rho_p}{\gamma_{mp}}$  for everything that follows.

**Bistability.** ~~From the~~ We map the mRNA nullcline surface (green in Fig 4A) onto a plane to show it as the 2D plot in Fig 4B. From this 2D plot, it can clearly be seen that for the chosen parameters, there are three steady-state solutions. Because the ~~2D plane that we are in now is~~ system is restricted to the mRNA nullcline, ~~these surface, these steady-state~~ these steady-state solutions are the same ~~fixed points as as those~~ in the 3D plot ( $\frac{d[m]_a}{d\tau} = 0$  on the nullcline and  $\frac{d[p]_a}{d\tau} = 0$ ,  $\frac{d[x]_a}{d\tau} = 0$  for the 2D fixed points). One can see from the vector field that the two outer fixed points (labeled 1 and 3) are stable and the middle one (labeled 2) is unstable and serves as a sort of “switch-point” between the other two. This means that there are two stable states the cell can be in, one at high

(point 3) and one at low (point 1) expression. As a result, there is bistability and the distribution of expression among cells can be bimodal, depending on initial conditions.

~~This~~ The bistability corresponds to the experimental observations [21], so the model passes this sanity check. Furthermore, the xanthosine and protein concentrations at the upper fixed point have the expected order of magnitude: the xanthosine concentration is roughly 10 – 100 mM, and there are roughly 500 proteins, which is just a bit lower than what was measured for the number of Nup transporters [24] which fulfill a similar purpose. We do not have well founded expectations for the other fixed points, so no comparison can be made here. Nevertheless, the orders of magnitude at the lower fixed point – roughly 1 – 10 nM of xanthosine and around 5 proteins – seem ~~perfectly~~ quite reasonable. Note that  $[x]_a \approx [c]_a$  at the lower fixed point because there is only weak accumulation due to Nup and a few XapB transporters.

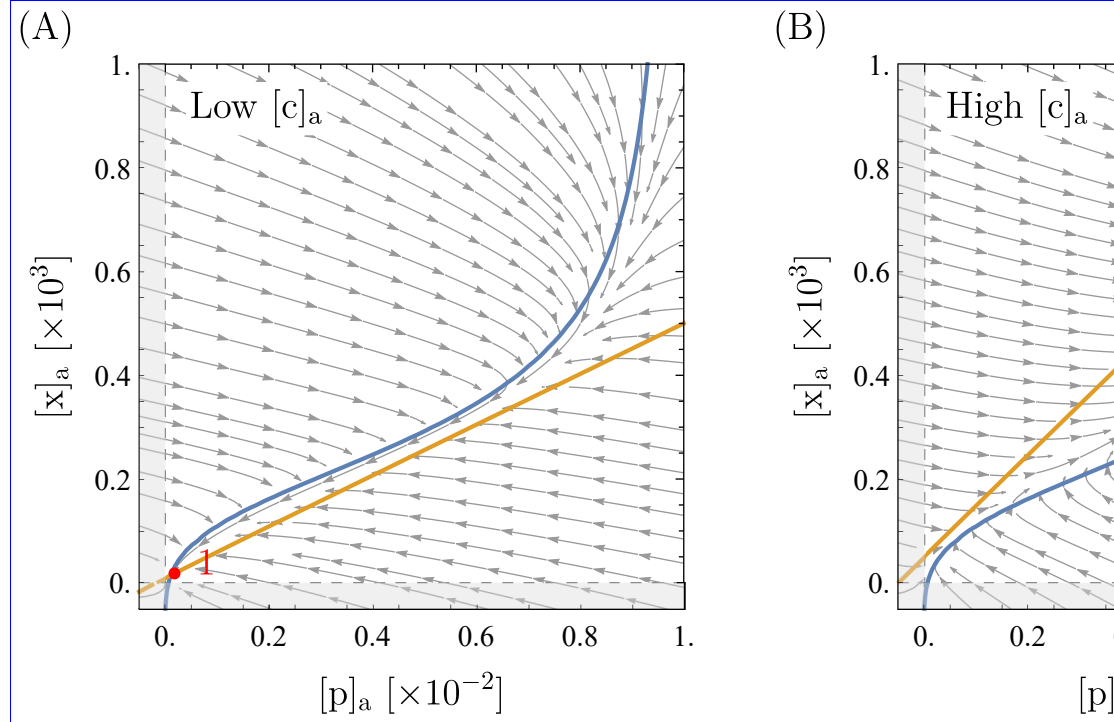
As already mentioned, we are working with one specific set of parameters here and we will now explain this choice of values. Firstly, they were picked such that they are well within ~~their estimated range~~ the parameter range that was estimated beforehand (see Table 1). Secondly, we ~~wanted clear bistability to occur~~ chose parameters that allow clear bistability in the phase portraits as well as in the stochastic simulations (see later), which, of course, is not the case for any possible choice of parameters. Thirdly, by the corresponding choice of parameters it was ensured that the mRNA number per cell at the “switch-point” is around 1: this is large enough to enable the system to clearly resolve the two stable fixed points (as we will see from the stochastic simulations later on), but is low enough to lead to mean mRNA numbers that are very reasonable (see [25] for the average mRNA numbers in cells, ~~which are rather low~~). The protein and xanthosine concentrations followed from this, but with some variation in the parameters they could still be tuned to a certain extent.

As a remark we point out that we have not observed any oscillations in the system. Intuitively, they might be expected when the XapA rate is significantly larger than the XapB rate, but it turns out that oscillations cannot be obtained. ~~This~~ Why they do not occur can be understood when looking at the regions that are bounded by all three nullclines: on these boundaries, the streamlines point into the bounded regions, so deterministically, they serve as trapping regions from which the system cannot escape. Once inside, the only possible trajectory is non-oscillatory flow towards the stable fixed point.

For a different set of parameters, the orders of magnitude in the plots and even the qualitative behavior can change. In the following, we will discuss some interesting features of the system that can be observed through the phase portraits.

**The extracellular xanthosine concentration.** The parameter that is the experimentally most easily tunable and biologically the most relevant is the extracellular xanthosine concentration. When it is increased in experiments, the cells go from (1.) all being in the low expression state to ~~being in either the low or the~~ (2.) the population being in a mixed state with some cells in a low expression state and others in a high expression state (all-or-none phenomenon) ~~to~~ and then to (3.) all being in the high expression state [21]. If our model is correct, it should exhibit the same qualitative behavior. Indeed we find exactly this: as can be seen in Fig 5, increasing  $[c]_a$  makes the high stable fixed point appear and then, for even higher  $[c]_a$ , the lower one ~~disappear~~ disappears. Thus, for low  $[c]_a$  the only stable point of the system is at low expression, and for high  $[c]_a$  there is only high expression. In between, there are two deterministically stable expression levels.

Furthermore, we found that in the absence of xanthosine, i.e., setting  $[c]_a = 0$  (not shown here), there are roughly 2-3 ~~proteins~~ copies of XapA and XapB, which agrees very well with measurements, where around 2 ~~proteins~~ copies per cell were found [24].



**Fig 5. Phase portraits for different extracellular xanthosine concentrations.** All parameters but  $[c]_a$  are as presented in Table 1. The extracellular xanthosine concentration in these plots is  $[c]_a = 10^1$  in (A) and  $[c]_a = 5 \cdot 10^2$  in (B) (recall that  $[c]_a := \frac{c}{K_a}$  ; with  $K_a = 5 \cdot 10^{-5} \text{ M}$  is dimensionless). Tuning  $[c]_a$  moves the orange line (xanthosine nullcline) up or down, the blue line-curve (mRNA nullcline) is unchanged (see also S1 Text). It can clearly be seen that in (A) there is only the lower fixed point (fixed point number 1), whereas in (B) there is only the upper one (fixed point number 3). In between lies the bistable case that was shown in Fig 4.

In addition, the parameter  $K_{\chi A}$  (dissociation constant of xanthosine from active XapR) can be tuned such that the extracellular xanthosine concentration  $[c]_a$  in the switching-regime is similar to that in the experiment. It was found that the cell only adapts at very high xanthosine concentrations of almost a millimolar [20] which is not completely unexpected when recalling that for *lac*, cells also limit themselves to glucose as long as possible. Interestingly, because there is no parameter other than  $K_{\chi A}$  that tunes the critical value of  $[c]_a$ , this tells us that  $K_{\chi A}$  is large as argued in the estimation of  $K_{\chi A}$  in S1 Text. Thus, we predict that the interaction between xanthosine and XapR should be weak.

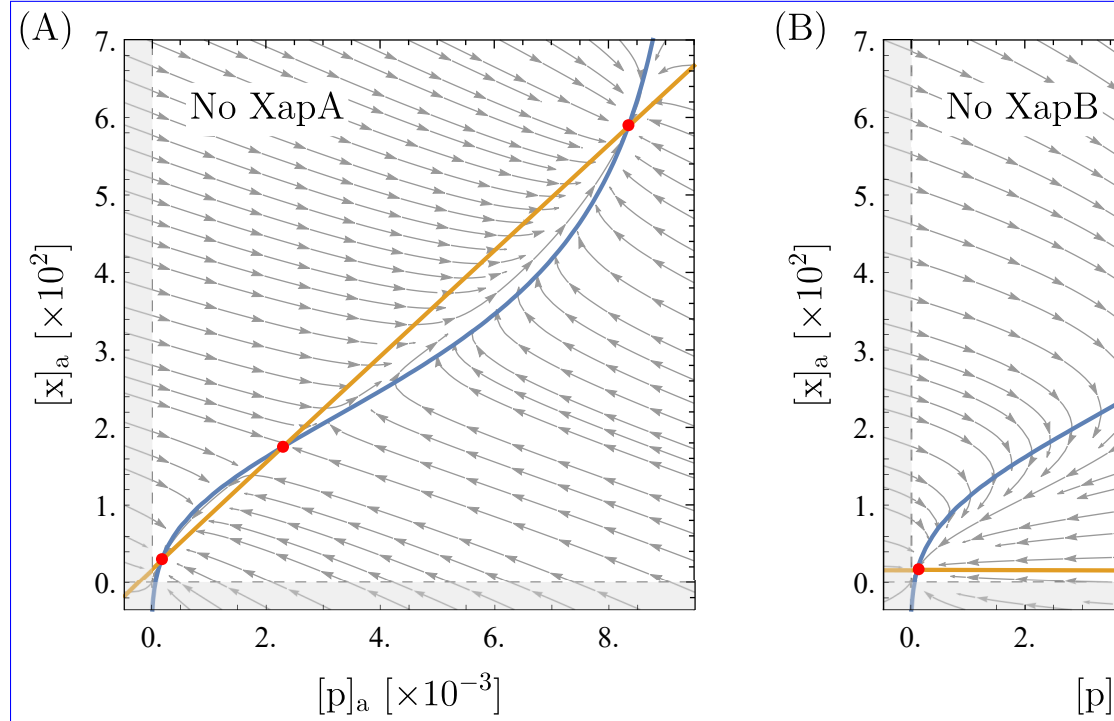
**The roles of XapA and XapB.** While it is clear that the bistability in the model system is due to the feedback loop from XapA and XapB, it is not obvious if both XapA and XapB are necessary. It turns out that the bistability is due to XapB only. XapA is neither sufficient nor necessary and, within the estimated parameter regime, does not even have a significant influence on the system. This can be seen from the plots in figure 6. The degradatory effect of XapA lowers the upper solution xanthosine and protein concentration at the upper fixed point by a small amount and could, in principle, thereby make the high-expression solution vanish. For our choice of all other parameters, this requires it only vanishes for  $k_\alpha > 10^4$  which is far from what has been measured. However, a higher effective rate could, in principle, be achieved by different translation rates of XapA and XapB (see simplifications of the model in S1 Text). Hence, we cannot exclude the possibility that XapA becomes so strong that it makes bistability impossible, but this is an extreme case. XapB, on the other hand, is essential; without it the system only has the one fixed point at low expression.<sup>1</sup>

For a cell, the above insignificance of the effect of XapA is a useful feature: by coupling XapA and XapB on an operon, XapA is switched on and off together with XapB but it does not significantly disturb this adaptation mechanism, while its kinetic parameters and expression levels can be chosen somewhat freely as necessary for metabolism. By having a membrane transporter gene on an operon whose expression is activated by the transporter substrate, the expression of a whole set of enzymes can be turned on and off depending on the presence of the substrate. It seems likely that this mechanism of short-term adaptation of a single cell to its environment may be used by cells for many metabolic processes.

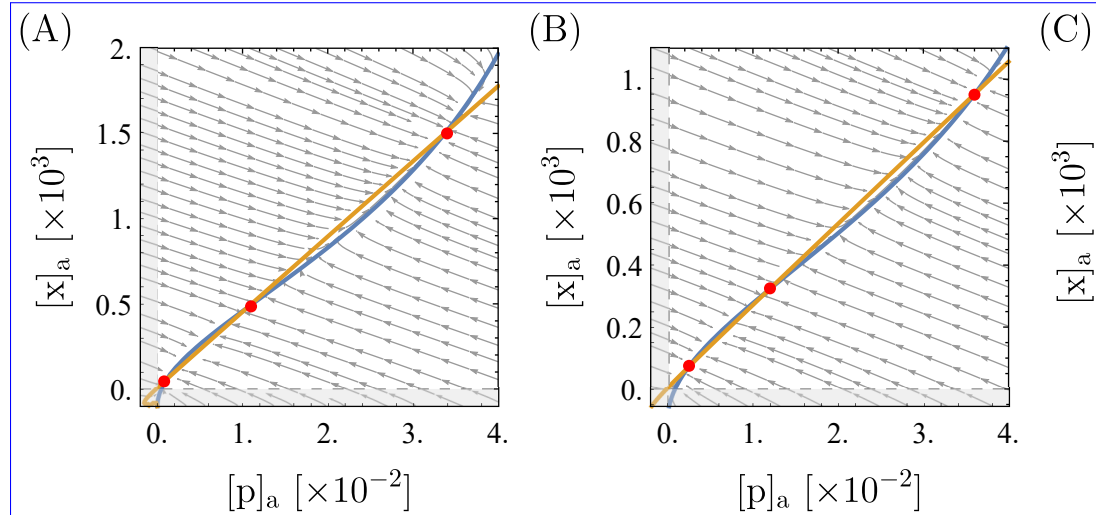
**The role of cooperativity.** The model has two cooperatively interacting binding sites for XapR on the *xapAB* promoter and two cooperative binding sites on XapR for xanthosine. An interesting question to pose is whether the cooperativity is a necessary feature for bistability. This question is motivated by the importance of cooperativity in “typical” genetic switches [2, 26].

If, as a purely theoretical consideration, we remove either the second xanthosine binding site on XapR or the second XapR binding site on the promoter, leaving cooperativity in one place of the system, we find that the system still has a bistable parameter regime. However, this is only a rather narrow parameter range bistable parameter range is only narrow, which makes the system less bistable. When less stable: small stochastic fluctuations in the parameter values can collapse the system to monostability, possibly leaving it in the wrong state and without its ability to adapt. But only when the second binding site is removed in both places, the leaving no cooperativity in the system, the equations are insufficiently non-linear to produce

<sup>1</sup> Seeger et. al. [18] observed that  $\Delta xapB$  mutants could survive, but grew extremely slowly, with xanthosine as the only carbon source, which makes sense in light of Fig 6. With *xapB* removed, the switch never activates and the cells are forced to survive with an extremely meager quantity of XapA to metabolize the abundant xanthosine.



**Fig 6. Phase portraits without XapA or XapB.** All parameters are as presented in Table 1. In (A), the XapA term was removed from the kinetic equations. In (B), the equations lack the two terms from XapB. These plots clearly show that XapA has almost no influence on the qualitative behavior of the system (i.e. bistability and the order of magnitudes), but XapB is the essential feature for bistability.



**Fig 7. Phase portraits for less or no cooperativity.** Most parameters are as presented in Table 1, changes are mentioned below. Fixed points are marked in red. In (A), there is only one xanthosine binding site on XapR and everything unchanged for the XapR-promoter binding. Two parameters are changed:  $\rho = 0.07$  and  $[c]_a = 6$ . This is necessary to compensate for the weaker induction such that the system is bistable. In (B), there is only one XapR binding site on the promoter and everything is unchanged for the xanthosine-XapR binding. Two parameters are changed:  $\rho = 0.13$  and  $[c]_a = 3$ . In (C), there is only one xanthosine binding site on XapR and also only one XapR binding site on the promoter. Two parameters are changed:  $\rho = 0.1$  and  $[XapR]_R = 5$ . Whereas bistability is retained in (A) and (B), it cannot be obtained anymore in (C).

bistability. An example of ~~these three scenarios~~ the three scenarios (only cooperative XapR, only cooperative promoter, no cooperativity) can be seen in Fig 7. It follows that there need to be either two xanthosine binding sites on XapR or two XapR binding sites on the promoter (or both) in order to obtain a switch-like behavior.

One can also ask how much cooperative interaction is needed between the two binding sites. For the promoter, ~~this~~ the amount of cooperativity is given by  $\Delta E_{coop}$  in our model, and we find that setting  $\Delta E_{coop} = 0$  has almost no influence on the phase diagrams. For XapR, we cannot test how much interaction is needed: the two binding sites interact indirectly, because the active state is much likelier if two xanthosine molecules are bound. ~~This cooperative interaction can, however, not be tuned like,~~ and thus there is no tuning parameter for the cooperative interaction like  $\Delta E_{coop}$  in the case of the promoter ~~with  $\Delta E_{coop}$ .~~

Note that we are not writing Hill equations and measuring cooperativity in terms of the Hill coefficient. If Hill equations were to be used for the modeling, the Hill coefficient could have values between 1 and 2, which would yield bistability for large enough values, but not for lower ones. This could be investigated more rigorously similar to the analysis of a simple genetic switch in [26]. However, we refrain from looking for a minimal Hill coefficient in our system, because we do not find this very insightful. Hill equations only describe ~~scenarios with multiple, non-interacting binding sites where any other state but the fully occupied and active or the fully empty and inactive state is not significantly populated.~~ In that sense, the Hill coefficient simply denotes the number of binding sites and does not reflect how cooperative these binding sites are some specific limit cases of cooperative systems, but for example do



not account for interaction energies and assume the partially bound states (e.g. only one XapR bound to the promoter) to never be populated. We suggest that cooperativity should be explored more in-depth and a more rigorous analysis of the role of cooperativity in simple genetic switches should be done before returning to more complex systems like this one.

## Stochastic simulations

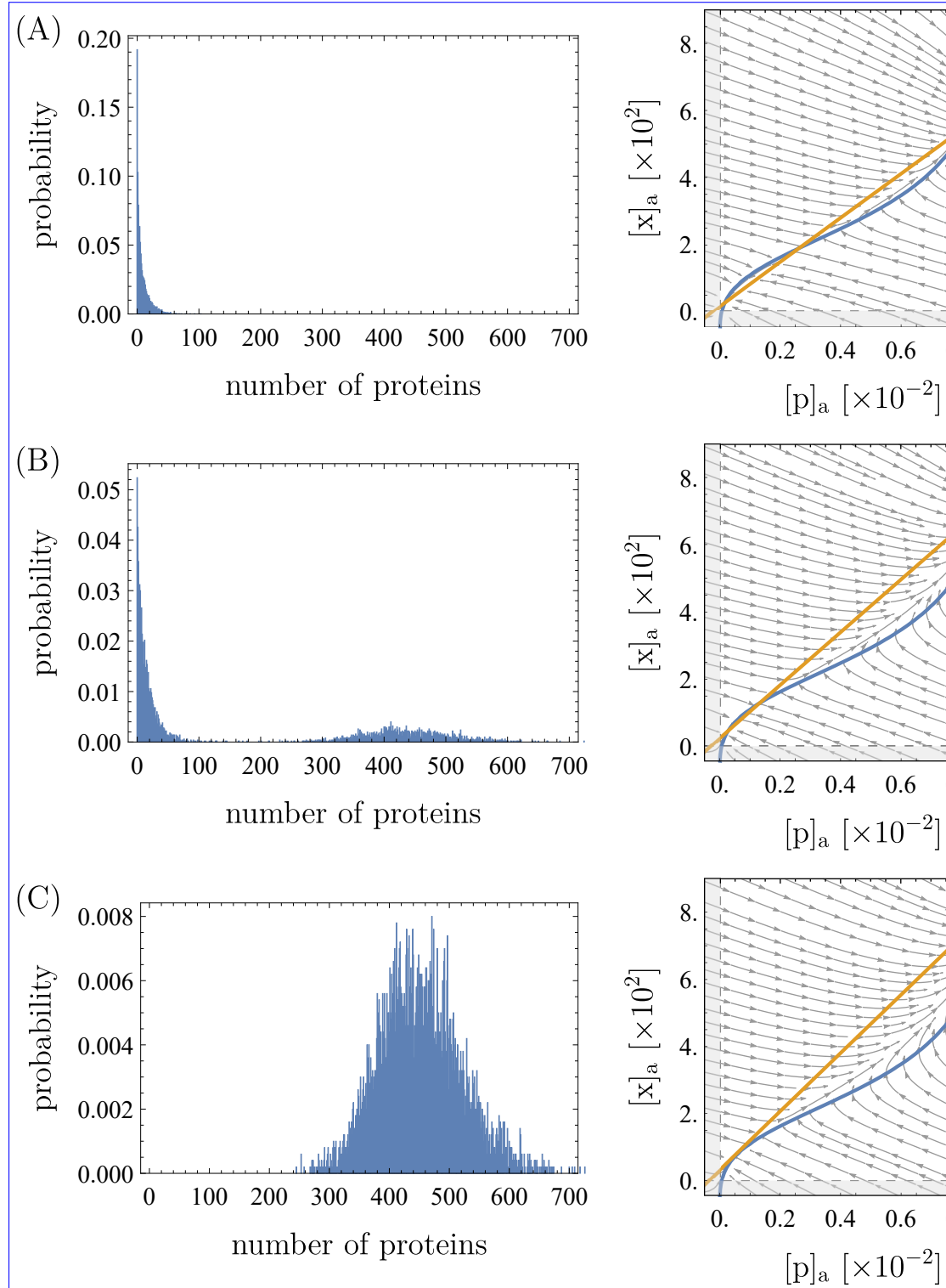
Stochastic simulations of the full 3-dimensional system of mRNA, protein, and xanthosine were run for comparison with the deterministic results. In S1 Text, we present the underlying chemical master equation of the system. Because of the two different fixed points at low and high expression, the copy numbers in the problem vary from very low to very high. For large amounts of mRNA, protein, and xanthosine, the number of reactions is large too, and thus, Gillespie’s classical algorithm has a large computational cost and his  $\tau$ -leap algorithm would be ideal. On the other hand,  $\tau$ -leaping cannot be used for the small copy numbers. For these reasons, we chose to work with the algorithm described in [27], a hybrid form between Gillespie’s classical and his  $\tau$ -leap algorithm. We gratefully worked with the Python implementation of this algorithm in *StochPy*, version 2.3 [28].

**Bimodality and the extracellular xanthosine concentration.** This stochastic approach results in the same bimodal distributions that were already seen in the deterministic investigation and experimental studies. Fig 8 shows the distribution of protein expression found in the simulations for different values of the extracellular xanthosine concentration. The parameters that were used are the same as in the previous section (listed in Table 1). To obtain the distributions, we ran the simulation 5000 times for a simulated time of  $10^6$  s each and started at a mRNA, protein and intracellular xanthosine count of 0.

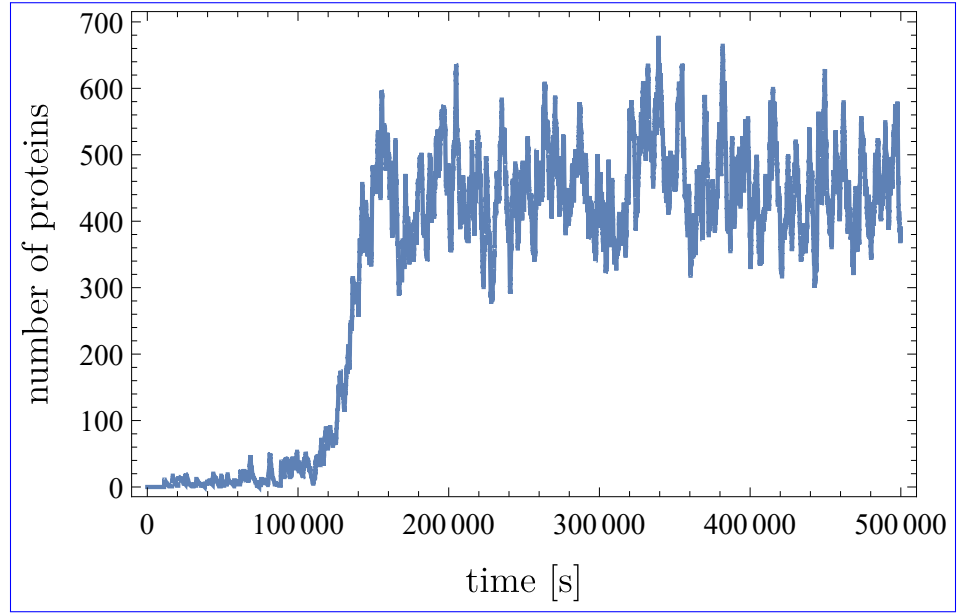
The results agree very well with the deterministic fixed points and experiments: the mean numbers of mRNA, protein, and xanthosine in the stochastic results are as predicted from the phase portraits. It does, however, become clear that the phase portraits do not tell whether the cells will actually populate both the high and the low expression state: ~~as can be seen in~~, because they do not show the barrier height between the two states. In Fig 8(A), a deterministically bistable scenario is shown where the cells never switched to the high expression state during the run time of our simulations.

We found that the two lower fixed points (marked as 1 and 2 in Fig 4) need to be very close like in Fig 8(B) to give bimodality. For lower  $[c]_a$ , meaning larger distance between the first and second fixed point, ~~no or~~ almost no switching was observed. Of course, ~~this switching~~ is also a matter of the waiting time and stochastic effects: if one waits for long enough, ~~switching it~~ should eventually occur. However, switching times of more than several hours are not at the center of this investigation and would mean that switching is extremely unlikely. There are two aspects that become relevant in this context that we neglect in our analysis but shortly mention here: transcription and translation bursts lead to higher stochasticity and cell division leads to some discontinuity in the process.

Note that while the deterministic analysis assumes the variables to be continuous, the simulations work with discrete numbers of mRNA, protein, and xanthosine. This per se is no problem, because the deterministic analysis describes the mean values and the simulation fluctuates around this mean. If, however, the mRNA number at the third (high) fixed point is too low, the system will not be able to resolve the two points anymore. ~~This becomes a problem~~ It is too low when the distance between the first and



**Fig 8. Distributions from stochastic simulations and the corresponding phase portraits.** Apart from  $[c]_a$ , the parameters are the same as in Table 1. For the distributions, the simulations were run 5000 times for  $10^6$  s each (simulated time) and started at a mRNA, protein and intracellular xanthosine count of 0. We show the two cases of unimodality (low expression in (A)) and high expression in (C)) as well as the case of bimodality in (B). The values of  $[c]_a$  are 12 in (A), 18.5 in (B), and 25 in (C) (recall  $[c]_a := \frac{c}{K_a}$ ,  $K_a = 5 \cdot 10^{-5}$  M). The output from the stochastic simulations is in good agreement with the concentrations at the fixed points in the deterministic phase portraits.



**Fig 9. Time evolution of protein (XapA/XapB) from one run of the simulation.** The simulation was run once for  $5 \cdot 10^5$  s and started at an mRNA, protein and intracellular xanthosine count of 0. In this case the system switched into the high-expression state after  $10^5$  s. The parameters that were used are the same as in Table 1, the only exception being the extracellular xanthosine concentration, which was chosen to be  $[c]_a = 25$  (recall  $[c]_a := \frac{c}{K_a}$ ,  $K_a = 5 \cdot 10^{-5}$  M) just as in Fig 8(B).

the third fixed point becomes as low as the stochastic fluctuations in the system, which is around 3 mRNA molecules. In Fig 8, the mRNA number at the low fixed point was fluctuating between 0 and 2 with a mean significantly smaller than 1, and at the high fixed point it was varying between 2 and 6.

**Time evolution and switching times.** Fig 9 shows the time evolution of ~~one specific example~~ the protein concentration for one exemplary run of the simulation. Again, the simulation was started with a mRNA, protein and intracellular xanthosine count of 0 and was run for a simulated time of  $5 \cdot 10^5$  s. In this specific example, switching occurred after  $10^5$  s.

Comparing this to the experimentally expected timescales [21] is difficult, because the switching time strongly depends on the extracellular xanthosine concentration. Experiments were always stopped after a few hours, and in this time, the cell population might not reach its steady-state expression distribution. Hence, the distribution could be bimodal when the experiment is stopped but become unimodal after further waiting. That way, extracellular xanthosine concentrations that are too high for deterministic bistability could lead to experimental bimodality if the experiment is stopped too early. In this case, the observed switching time would be shorter, which makes the comparison to our simulations even harder. Thus, we cannot say if it is problematic that the  $10^5$  s is larger than what was found in the experiment.

Nevertheless, we do warn the reader that the timescales in the simulations should be taken with reservations. Cell divisions are not considered here, and neither is the burstiness of transcription and translation. This means that stochasticity may be larger in the real system which should have an influence on the timescales and probably shorten the time until switching occurs.

**Hysteresis.** In Figs 8 and 9, the simulations were started at initial intracellular concentrations of 0 to investigate what happens if xanthosine is suddenly added to the cell's environment. We can now ask the opposite question: what happens when xanthosine is removed from the extracellular environment? To answer this, the simulation was started with initially fully induced cells, i.e. at the mRNA, protein and intracellular xanthosine counts of the high fixed point in the corresponding phase portrait.

The resulting distributions can be found in S1 Text. They show clear hysteresis effects: there exist extracellular xanthosine concentrations where initially uninduced cells remained uninduced and initially induced cells remain induced. Only when the second and the third fixed point are very close can initially induced cells switch to the uninduced state. Interestingly, this behavior is symmetric to the “switching on” in the previous paragraphs, where the lower two fixed points needed to be about as close as the upper two need to be now for “switching off”.

In other words, cells only change their metabolism to xanthosine if enough of the latter is around, but after they have switched, this metabolic state is stable even if the xanthosine concentration decreases to a certain extent. This [stability](#) explains what has been observed by Novick and Weiner [4] for the *lac* operon: when induced cells were transferred into lower concentrations of lactose, they remained induced, even though uninduced cells could not become induced at these concentrations.

## Conclusion

**A genetic switch for metabolic adaptation with remarkable features.** In this paper, we propose a [simple](#) model for genetic circuits containing a membrane transporter whose gene expression is, directly or indirectly, activated by its substrate. We have shown that such a system can be bistable and thus work as a genetic switch which reacts to the extracellular concentration of the relevant metabolite. This switch has very useful biological features. First, coupling of the transporter with, for example, an enzyme which metabolizes the transporter substrate creates a genetic switch that enables short-term adaptation of the cell's metabolism to its environment. Second, the switch is stabilized by hysteresis effects when the extracellular substrate concentration decreases, which explains previous experimental findings.

**Required properties for this system.** We have found that no bistability can emerge from the genetic circuit unless at least one component has two binding sites for its activator. Additional binding sites or cooperativity seem to increase the stability of the switch. In addition, simply knowing the experimental switching concentration of xanthosine let us, for example, infer the approximate value of the dissociation constant between the transcription factor XapR and the inducer xanthosine. We found that their interaction needs to be rather weak [compared to usual interactions between an inducer and its target](#).

**Possibilities for further investigation.** Phase diagrams, showing for which parameters the system is bistable and for which there is only the lower or the upper stable fixed point, could be calculated from arguments made in [26]. However, the simulations have shown that deterministic bistability does not mean that bimodality occurs, which is why we have refrained from showing such diagrams. Furthermore, the timescales in the problem could be investigated more thoroughly, for example the dependence of the switching time on  $[c]_a$ , but such an analysis would probably need to account for the burstiness of transcription and translation as well as for cell divisions, which is anything but straightforward. ~~Lastly, some rigorous work could be done to~~

identify the most and least relevant parameters of the system to create a simpler, minimal model that captures the most relevant features.

**General lessons learned for the modeling of other systems**A successful generalizable model. We point out again that assuming the mRNA concentration to be at steady-state can have a significant impact on the apparent dynamics of a system, as we have seen in the 3-dimensional phase portraits. Moreover, we remind the reader that deterministic phase portraits do not show which states are actually populated, and thus hide essential effects like hysteresis. Lastly, we have exemplified issues with the common picture of measuring cooperativity in terms of the Hill coefficient: the Hill coefficient simply denotes the number of binding sites and does not reflect how cooperative these binding sites are.

**A successful model.** All model parameters could be reasonably estimated despite the paucity of experimental knowledge about the model system. ~~In the results, every variable and parameter has its expected order of magnitude~~The concentrations of mRNA, protein, and xanthosine at the fixed points as well as all qualitative features are as expected from similar systems and the few experiments on the *xap* circuit, which suggests that the model captures ~~all relevant features~~the relevant components of the system correctly and is able to describe ~~the dynamics of the *xap* system~~its dynamics. Furthermore, the modeling results let us, to some extent, understand why the biological system is constructed the way it is. By keeping the model as minimal as possible, but still modeling every part explicitly with an appropriate complexity, we can investigate the interesting features while still being able to understand the influence of all parameters and their interplay intuitively.

With the framework given in this text, it should be straightforward to model other promoters, regulatory pathways or enzymes and thereby adapt the model to other genes and metabolites. Examples include *lac*, *ara*, and *xyz*, but we suspect that many if not most metabolic processes involve the adaptation mechanism that we have investigated here, and that much can be understood about them through our model. This apparent success demonstrates once more that even for broadly unknown systems, rigorous physical modeling can potentially offer an efficient way to gain a very thorough understanding of the behavior of the system.

## Supporting information

**S1 Text. The aforementioned further information.** Discussion of simplifications in the model, parameter estimation, elaborations on the results, and the chemical master equation of this circuit.

## Acknowledgments

We thank Griffin Chure, Charlotte Strandkvist, and Kyle Naughton for providing their data on the *xap* system and pointing out the bistability. We also thank Griffin Chure for answering the many biological questions that arose, Nathan Belliveau for bringing similar systems to our attention, and Jane Kondev and Jin Wang for interesting discussions. For assistance with illustrations, we also thank Nigel Orme and Griffin Chure. K.S.L. acknowledges financial support from the Werner Siemens-Foundation through the Swiss Study Foundation. This material is based upon work supported by the National Science Foundation Graduate Research Fellowship under Grant No. DGE-114446.

MJM: this kinda reads like GC should be in the author list...

MJM: need to double-check my NSF #

remove fund-

## References

1. Jacob F, Monod J. Genetic regulatory mechanisms in the synthesis of proteins. *Journal of Molecular Biology*. 1961;3(3):318–356.
2. Gardner TS, Cantor CR, Collins JJ. Construction of a genetic toggle switch in *Escherichia coli*. *Nature*. 2000;403:339.
3. Wolf DM, Eeckman FH. On the Relationship between Genomic Regulatory Element Organization and Gene Regulatory Dynamics. *Journal of Theoretical Biology*. 1998;195(2):167 – 186. doi:<https://doi.org/10.1006/jtbi.1998.0790>.
4. Novick A, Weiner M. Enzyme induction as an all-or-none phenomenon. *Proceedings of the National Academy of Sciences of the United States of America*. 1957;43(16590055):553–566.
5. [Santillan M, Mackey MC. Quantitative approaches to the study of bistability in the lac operon of Escherichia coli. \*Journal of The Royal Society Interface\*. 2008;5\(suppl\\_1\):S29–S39. doi:10.1098/rsif.2008.0086.focus.](#)
6. Narang A, Pilyugin SS. Bistability of the lac Operon During Growth of *Escherichia coli* on Lactose and Lactose + Glucose. *Bulletin of Mathematical Biology*. 2008;70(4):1032–1064.
7. Choi PJ, Cai L, Frieda K, Xie XS. A Stochastic Single-Molecule Event Triggers Phenotype Switching of a Bacterial Cell. *Science*. 2008;322(5900):442–446. doi:10.1126/science.1161427.
8. Ozbudak EM, Thattai M, Lim HN, Shraiman BI, van Oudenaarden A. Multistability in the lactose utilization network of *Escherichia coli*. *Nature*. 2004;427:737.
9. Fritz G, Megerle JA, Westermayer SA, Brick D, Heermann R, Jung K, et al. Single Cell Kinetics of Phenotypic Switching in the Arabinose Utilization System of *E. coli*. *PLOS ONE*. 2014;9(2):e89532. doi:10.1371/journal.pone.0089532.
10. Jenkins A, Macauley M. Bistability and Asynchrony in a Boolean Model of the l-arabinose Operon in *Escherichia coli*. *Bulletin of Mathematical Biology*. 2017;79(8):1778–1795.
11. Siegele DA, Hu JC. Gene expression from plasmids containing the araBAD promoter at subsaturating inducer concentrations represents mixed populations. *Proceedings of the National Academy of Sciences of the United States of America*. 1997;94(9223333):8168–8172.
12. Bae JS, Kim TH, Kim MY, Park JM, Ahn YH. Transcriptional Regulation of Glucose Sensors in Pancreatic beta-Cells and Liver: An Update. *Sensors*. 2010;10(5):5031–5053. doi:10.3390/s100505031.
13. Tiedge M, Lenzen S. Regulation of glucokinase and GLUT-2 glucose-transporter gene expression in pancreatic beta-cells. *The Biochemical journal*. 1991;279 (Pt 3)(1953686):899–901.
14. [Song S, Park C. Organization and regulation of the D-xylose operons in Escherichia coli K-12: XylR acts as a transcriptional activator. \*Journal of Bacteriology\*. 1997;179\(22\):7025–7032. doi:10.1128/jb.179.22.7025-7032.1997.](#)

15. [Schmidt A, Kochanowski K, Vedelaar S, Ahrne E, Volkmer B, Callipo L, et al. The quantitative and condition-dependent \*Escherichia coli\* proteome. Nature Biotechnology. 2015;34:104.](#)
16. Buxton RS, Hammer-Jespersen K, Valentin-Hansen P. A second purine nucleoside phosphorylase in *Escherichia coli* K-12. Molecular and General Genetics MGG. 1980;179(2):331–340.
17. Hammer-Jespersen K, Buxton RS, Hansen TD. A second purine nucleoside phosphorylase in *Escherichia coli* K-12. Molecular and General Genetics MGG. 1980;179(2):341–348.
18. Seeger C, Poulsen C, Dandanell G. Identification and characterization of genes (xapA, xapB, and xapR) involved in xanthosine catabolism in *Escherichia coli*. Journal of bacteriology. 1995;177(7559336):5506–5516.
19. Jorgensen C, Dandanell G. Isolation and characterization of mutations in the *Escherichia coli* regulatory protein XapR. Journal of bacteriology. 1999;181(10400599):4397–4403.
20. Norholm MH, Dandanell G. Specificity and topology of the *Escherichia coli* xanthosine permease, a representative of the NHS subfamily of the major facilitator superfamily. Journal of bacteriology. 2001;183(11466294):4900–4904.
21. Chure G, et al. Measurements on the *xapABR* system in *E. coli*; Unpublished.
22. Marzen S, Garcia HG, Phillips R. Statistical Mechanics of Monod–Wyman–Changeux (MWC) Models. Journal of Molecular Biology. 2013;425(9):1433 – 1460. doi:<https://doi.org/10.1016/j.jmb.2013.03.013>.
23. Kaback HR. A chemiosmotic mechanism of symport. Proceedings of the National Academy of Sciences. 2015;112(5):1259–1264. doi:10.1073/pnas.1419325112.
24. Li GW, Burkhardt D, Gross C, Weissman J. Quantifying Absolute Protein Synthesis Rates Reveals Principles Underlying Allocation of Cellular Resources. Cell. 2014;157(3):624 – 635. doi:<https://doi.org/10.1016/j.cell.2014.02.033>.
25. Milo R, Phillips R. In: Cell biology by the numbers. Garland Science. 2016; 217.
26. Cherry JL, Adler FR. How to make a Biological Switch. Journal of Theoretical Biology. 2000;203(2):117–133.
27. Cao Y, Gillespie DT, Petzold LR. Efficient step size selection for the tau-leaping simulation method. J Chem Phys. 2006;124(4):044109. doi:10.1063/1.2159468.
28. Maarleveld TR, Olivier BG, Bruggeman FJ. StochPy, Stochastic Modeling in Python. 2015. Available from: <http://stochpy.sourceforge.net/>.