Modeling synergistic drug inhibition of *Mycobacterium tuberculosis* growth in murine macrophages†

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We developed a metabolism-based systems biology framework to model drug-induced growth inhibition of *Mycobacterium tuberculosis* in murine macrophage cells. We used it to simulate *ex vivo* bacterial growth inhibition due to 3-nitropropionate (3-NP) and calculated the corresponding time- and drug concentration-dependent dose-response curves. 3-NP targets the isocitrate lyase 1 (ICL1) and ICL2 enzymes in the glyoxylate shunt, an essential component in carbon metabolism of many important prokaryotic organisms. We used the framework to *in silico* mimic drugging additional enzymes in combination with 3-NP to understand how synergy can arise among metabolic enzyme targets. In particular, we focused on exploring additional targets among the central carbon metabolism pathways and ascertaining the impact of jointly inhibiting these targets and the ICL1/ICL2 enzymes. Thus, additionally inhibiting the malate synthase (MS) enzyme in the glyoxylate shunt did not produce synergistic effects, whereas additional inhibition of the glycerol-3-phosphate dehydrogenase (G3PD) enzyme showed a reduction in bacterial growth beyond what each single inhibition could achieve. Whereas the ICL1/ICL2-MS pair essentially works on the same branch of the metabolic pathway processing lipids as carbon sources (the glyoxylate shunt), the ICL1/ICL2-G3PD pair inhibition targets different branches among the lipid utilization pathways. This allowed the ICL1/ICL2-G3PD drug combination to synergistically inhibit carbon processing and ultimately affect cellular growth. Our previously developed model for *in vitro* conditions failed to capture these effects, highlighting the importance of constructing accurate representations of the experimental *ex vivo* macrophage system.

Introduction

Tuberculosis (TB) remains a potential health threat to the general United States population, but the main burden of the disease is felt worldwide, with 9.4 million new cases and 1.8 million deaths in 2008.¹,² The causative agent of the disease, *Mycobacterium tuberculosis*, latently infects one-third of the world's human population.³ Current efforts to treat and eliminate TB are hindered by the complexity of drug regimens, the appearance of drug-resistant strains of *M. tuberculosis*, and the emergence of a patient population with compromised immune systems.³,⁴ Although current therapies include a combination of drugs that inhibits both metabolic and non-metabolic targets, these therapies will inevitably become less effective. The continued reliance on drugs to combat this disease necessitates a continuous search for new druggable targets and combination therapies.⁵

The search for novel bacterial drug targets and drug strategies is aided by recent genome-scale metabolic network reconstructions of several pathogenic organisms,⁶–¹⁰ including *M. tuberculosis*.¹¹,¹² These reconstructions can be used to understand species-specific differences among bacteria, e.g., mycolic acid utilization for Mycobacteria, the determination of nutrient requirements and metabolite processing steps, and the ability to probe metabolic enzymes/pathways for possible drug targets. Analyzing and probing these systems require systems biology tools to account for hundreds to thousands of metabolites and enzymes. Specifically, metabolic network reconstructions are primarily geared for studying metabolism processing and cell growth phenotypes. For example, given the availability of a specific set of nutrients, flux balance analysis (FBA) of metabolic networks can predict microbial growth rates.¹¹–¹⁵ Metabolic network analysis can also help identify the essential genes of an organism, *i.e.*, the genes required for the growth of the organism.⁸,¹¹–¹⁴,¹⁶ Essential genes constitute potential drug targets, especially if they do not have any homologous counterparts in the human genome.¹⁶,¹⁷
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One important application of metabolic network modeling is the ability to quantitatively model metabolic enzyme inhibition and predict bacterial growth inhibition within *in vitro* media.\[18\] By integrating enzyme inhibition kinetics, metabolic network analysis, and cellular growth dynamics, we previously developed a systems biology framework that allows us to quantitatively reproduce the dose response of 3-nitropropionate (3-NP) on the growth of *M. tuberculosis* in a medium with the fatty acid propionate used as its carbon source. Similarly, we modeled the growth inhibition of *M. tuberculosis* by 5′-O-(N-salicylsulfamoyl)adenosine in a medium with a low iron concentration. Both studies used *in vitro* media modified to capture some characteristic feature of the *in vivo* environment, i.e., restricted carbon sources and low ambient iron concentrations. One such specific *in vivo* environment is the macrophage.

In the early stages of infection, *M. tuberculosis* bacteria are typically localized in macrophage cells\[19\] where they survive host defense responses by multiple mechanisms, including blocking phagosomal maturation and countering host-induced antimicrobial peptides.\[20,22\] Part of the survival strategy involves metabolism, in particular, adaptation to the restricted availability of nutrients in the intracellular macrophage environment.\[19,22\]

When administering antibiotic treatments, it is essential to achieve the therapeutic antibiotic concentration at the site of action.\[23,24\] For TB treatments targeting intracellular *M. tuberculosis*, the drugs need to reach the immature phagosome in the macrophages, where the pathogen can persist for long times.\[19,23-26\] However, there are no studies that directly address the relationship between applied drug concentrations and the drug concentration in the cell compartment occupied by the pathogen. To understand the relationship between the applied drug dose and the actual intracellular concentration of the drug, researchers are exploring mechanisms that enable drug transport across cell boundaries\[27,29,30\] to ascertain the effective drug concentrations in macrophages.\[31,32\] If therapeutic concentrations at the site of action cannot be achieved at clinically relevant concentrations, the drug will not be efficacious and might instead speed up the appearance of drug-resistant pathogens.

Drug combinations aim to increase therapeutic efficacies and reduce pharmacological liabilities over single drug treatments.\[33,34\] For example, the addition of the proposed TB drug SQ109 to isoniazid (commercially available as Laniazid or Nydrazid) improves the growth inhibition of *M. tuberculosis* beyond what either drug can achieve singly.\[35\] Modeling studies of multiple inhibitions of metabolic reactions can potentially provide a rapid means to identify suitable targets that can be proposed for developing drug combination therapies.\[36\] In this spirit, Lehar and co-workers used *in silico* analyses to examine the effect of inhibiting multiple reactions in a small hypothetical network that included parallel and serial reactions as well as reaction feedback loops.\[37\] This group also examined shifts in synergy among drug combinations targeting metabolic pathways in *Escherichia coli* under fermentation and aerobic conditions.\[38\] However, to fully realize the potential of using systems biology tools to investigate drug combinations for intracellular pathogens, we need to develop quantitative models that can account for multiple metabolic inhibitors at different doses and to account for the host environment where the pathogen resides.

In this study, we have begun addressing these issues by creating a systems biology framework capable of simulating the inhibitory effects of 3-NP on the intracellular growth of *M. tuberculosis*. We developed a metabolic network that could mimic the growth of *M. tuberculosis* in murine macrophages, derived the effective relationship between the intracellular and extracellular 3-NP concentrations surrounding the pathogen, and quantitatively modeled 3-NP inhibition under different biological conditions. We used the developed framework to study the properties of potential drug combinations targeting metabolic enzymes/pathways and found that the different carbon-utilization strategies of the pathogen in the macrophage creates distinct patterns of synergistic drug targets. Optimal growth inhibition in this system could be achieved by jointly targeting isocitrate lyases in the glyoxylate shunt and the glycerol utilization pathway, an effect that could not be captured in an *in vitro* model. To our knowledge, this is the first attempt to use metabolic networks in a comprehensive analysis of the dose-dependent effects of drug combinations on a pathogen’s growth in host cells.

**Experimental**

We have previously developed the systems biology framework required to simulate *in vitro* drug-induced growth inhibition of *M. tuberculosis*. We applied this technology to model the growth inhibition due to 3-NP in medium containing propionate, an odd-chained fatty acid, as the primary carbon source.\[18\] Here, we describe the extensions of this framework to model drug-induced growth inhibition of *M. tuberculosis* in macrophages.\[39\]

**Mathematical framework**

Fig. 1 shows the general framework and relationship between our three model components, namely, the inhibition model, the metabolic network, and the population growth model. Each component deals with one specific aspect of modeling drug inhibition. The inhibition model (Inhibition Model) defines how a particular inhibitor affects the flux(es) of one or more metabolic reactions by creating inhibitor concentration-dependent constraints on each reactions, referred to as a “target reaction.” The metabolic network (Metabolic Network) component accounts for how the changes in the metabolite fluxes of the target reactions decrease the biomass production rate of the organism. The population growth model (Population Growth Model) uses the reduced biomass production rate to estimate the bacterial cell concentration under these conditions. With these model components in place, we can map a specific inhibitor concentration inside macrophages [I]<sub>i</sub> (the subscript “i” denotes intracellular concentration) to a cell concentration [X] as a function of time. This allows us to create dose-response curves and estimate minimum inhibitory concentrations in different media.

Although each of these components includes major differences compared to our previous *in vitro* model,\[18\] the main difference in the current framework formulation is that we did not explicitly account for the nutrient depletion in the macrophage. Instead, we assumed that the availability of
nutrients/substrates taken up from the host cell did not change
within the time frame of our study. In effect, this amounts to
assuming that nutrient depletion was not the major cause for
growth inhibition of the pathogen inside the macrophage.
Given the relatively short time frame modeled in this study
(less than 7 days) and that in the presence of an inhibitor
bacterial growth slows down, effectively reducing the
consumption of nutrients, this assumption should be reasonably
valid within this study. Thus, we did not dynamically change
the constraints on the bacterial uptake rates as a function
of time.

In order to capture the essential metabolic components of
*M. tuberculosis*-infected mouse macrophage cells growing in a
drug-infused medium, we assumed that the pathogen resides
in the phagosome compartment and uses lipids, composed of
fatty acids and glycerol, as carbon sources. The 3-NP drug
molecules in the medium are taken up by the macrophage,
enter the phagosome, and retard *M. tuberculosis* growth by
inhibiting two essential metabolic reactions required to use
fatty acids, isocitrate lyase (ICL) and methylisocitrate lyase
(MCL). The inhibition model describes the inhibitor
concentration-dependent constraints on the metabolic reaction fluxes in the metabolic network simulations. Since the enzymatic inhibition mechanism remains the same under *in vitro* and *ex vivo* conditions, we used the previous inhibition model except that the inhibitor concentration now refers to the intracellular phagosome 3-NP concentration. Thus, the ratio of inhibitor-present to inhibitor-free fluxes for the ICL reaction is as follows:

$$f_{ICL}(3-NP) = \frac{\nu_{ICL}}{\nu_{ICL,WT}}$$

$$= \frac{w_{ICL}}{1 + \frac{[SUC]}{K_{ICL1}} + \frac{[3-NP]}{K_{ICL1}}}$$

$$+ \frac{w_{ICL2}}{1 + \frac{[SUC]}{K_{ICL2}} + \frac{[3-NP]}{K_{ICL2}}}$$

where $\nu_{ICL}$ and $\nu_{ICL,WT}$ denote the inhibitor-present and
inhibitor-free reaction fluxes, respectively, the “$i$” in [3-NP] indicates intracellular concentrations, $w_{ICL}$ and $w_{ICL2}$ denote the fractions of the overall inhibitor-free ICL reaction flux for the reaction parts catalyzed by enzymes ICL1 and ICL2, respectively, SUC denotes the succinate substrate; $[SUC]$ indicates its concentration, and $K_{SUC,ICL1}$, $K_{SUC,ICL2}$, $K_{ICL1}$, and $K_{ICL2}$ denote Michaelis constants. Similarly, the ratio for the MCL reaction is as follows:

$$f_{MCL}(3-NP) = \frac{\nu_{MCL}}{\nu_{MCL,WT}}$$

$$= \frac{w_{MCL}}{1 + \frac{[SUC]}{K_{MCL1}} + \frac{[3-NP]}{K_{MCL1}}}$$

$$+ \frac{w_{MCL2}}{1 + \frac{[SUC]}{K_{MCL2}} + \frac{[3-NP]}{K_{MCL2}}}$$

where the variables and parameters have similar definitions to those in eqn (3). The parameter values in eqn (3) and (4) are the same as those used in the *in vitro* inhibition model.

### Metabolic network

The metabolic network model was used to calculate the biomass production rate $\mu$ (in h$^{-1}$), which was subject to the constraint in the flux of each target reaction modeled in eqn (3) and (4). We modified the metabolic network to account for the unique metabolic conditions that the pathogen encounters in the macrophage cells. As a starting point, we used the previously developed *inJ661v* model to represent the metabolic
Table 1 Modifications of the iNJ661v metabolic network to obtain the iNJ661i network. To develop a network compatible with the metabolism of *Mycobacterium tuberculosis* in cultivated mouse macrophage cells, we modified an existing *in vivo* network, iNJ661v,\(^4\) by reducing the inconsistencies between the predicted gene essentiality and the essentiality experimentally measured in mouse macrophage cells.\(^4\) We first corrected six inconsistencies between iNJ661v-predicted and experimental gene essentiality (for the *Rv0098, fadD21, plsC, Rv3588c, desA3*, and *kefB* genes) by restoring several enzyme functions and reactions disabled in iNJ661v and then, through a set of systematic optimization procedures,\(^4\) further corrected 13 inconsistencies.

<table>
<thead>
<tr>
<th>Index</th>
<th>Gene locus</th>
<th>Gene name</th>
<th>Essentiality measured in experiment</th>
<th>Essentiality predicted from iNJ661v</th>
<th>Essentiality predicted from iNJ661i</th>
<th>Modification to correct the inconsistency between the experimental and predicted gene essentiality</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Rv0098</td>
<td>Rv0098</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Restored the disabled ability of the product of the <em>fabG1</em> gene to catalyze mycolic acid synthesis</td>
</tr>
<tr>
<td>2</td>
<td>Rv1185c</td>
<td>fadD21</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Restored the disabled ability of the products of the <em>fadD9, fadD24</em>, and <em>fadD23</em> genes to function as fatty acid-CoA ligase</td>
</tr>
<tr>
<td>3</td>
<td>Rv2483c</td>
<td>plsC</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Restored the disabled ability of the product of the <em>Rv2182c</em> gene to function as 1-hexadecanoyl-sn-glycerol 3-phosphate O-acyltransferase</td>
</tr>
<tr>
<td>4</td>
<td>Rv3588c</td>
<td>Rv3588c</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Restored the disabled ability of the product of the <em>Rv3273c</em> gene to catalyze carboxylic acid dissociation and association</td>
</tr>
<tr>
<td>5</td>
<td>Rv3229c</td>
<td>desA3</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Restored the disabled synthesis of hexadecenoate</td>
</tr>
<tr>
<td>6</td>
<td>Rv3236c</td>
<td>kefB</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Restored the disabled ion transports by the potassium ABC transporter and sodium proton antipporter</td>
</tr>
<tr>
<td>7</td>
<td>Rv0992c</td>
<td>coaA</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Deleted pantetheine 4'-phosphate from the biomass objective function</td>
</tr>
<tr>
<td>8</td>
<td>Rv1653</td>
<td>argJ</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Allowed the uptakes of ornithine and arginine</td>
</tr>
<tr>
<td>9</td>
<td>Rv2210c</td>
<td>ilvE</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Allowed the uptake of leucine</td>
</tr>
<tr>
<td>10</td>
<td>Rv2288c</td>
<td>hemN</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Deleted protoheme from the biomass objective function</td>
</tr>
<tr>
<td>11</td>
<td>Rv2573</td>
<td>Rv2573</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Deleted pantetheine 4'-phosphate from the biomass objective function</td>
</tr>
<tr>
<td>12</td>
<td>Rv2702</td>
<td>ppgK</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Allowed fluxes through the reactions catalyzed by maltose trehalose isomerase and α-glucosidase</td>
</tr>
<tr>
<td>13</td>
<td>Rv2945c</td>
<td>lppX</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Deleted extracellular phthiocerol dimycocerosate A and phenol phthiocerol dimycocerosate from the biomass objective function</td>
</tr>
<tr>
<td>14</td>
<td>Rv2949c</td>
<td>Rv2949c</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Allowed the uptake of phenol palmitic acid</td>
</tr>
<tr>
<td>15</td>
<td>Rv2977c</td>
<td>thiL</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Deleted thiamin from the biomass objective function</td>
</tr>
<tr>
<td>16</td>
<td>Rv2995c</td>
<td>leuB</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Added the uptakes of leucine</td>
</tr>
<tr>
<td>17</td>
<td>Rv3340</td>
<td>mcmC</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Changed the reaction catalyzed by the cystathionine β-synthase enzyme into reversible</td>
</tr>
<tr>
<td>18</td>
<td>Rv0503c</td>
<td>cmaA2</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Blocked the ability of the product of the <em>mmaA2</em> gene to catalyze mycolic acid cyclopeptanolation</td>
</tr>
<tr>
<td>19</td>
<td>Rv0820</td>
<td>phoT</td>
<td>Essential</td>
<td>Non-essential</td>
<td>Essential</td>
<td>Blocked the phosphate uptake through diffusion and blocked the ability of the product of the <em>pstA2</em> gene to catalyze the phosphate uptake via the ABC system</td>
</tr>
</tbody>
</table>

state of *M. tuberculosis* in whole animals\(^4\) (see Section S1 in the ESI‡ for details on the network model selection). Next, we used gene essentiality data to modify\(^4\) this network by reducing the inconsistencies between the network-predicted gene essentiality and those experimentally validated for *M. tuberculosis* cells growing in murine macrophages.\(^4\) We first corrected six discrepancies between the iNJ661v-predicted and experimental essentiality by restoring several enzyme functions and reactions disabled in iNJ661v and then, through a set of systematic optimization procedures,\(^4\) further corrected 13 wrong predictions (see Table 1). The resultant network, iNJ661i, did not change the role of fatty acids and glycerol as major carbon sources, consistent with the experimental observation that these molecules are the main carbon sources for *M. tuberculosis* in both whole animals and cultivated macrophages.\(^39\)
We then refined the constraints on the substrate uptakes of iNJ661i, focusing on the role of carbon metabolism. Based on the experimental observation that the limitations associated with carbon metabolism determine the growth of *M. tuberculosis*, we left all non-carbon-containing metabolite uptakes unconstrained. As fatty acids and glycerol are the major carbon sources, we allowed all uptakes of these metabolites. The upper limits of the fatty acid uptakes (\(U_{fad}\)) and the glycerol uptake (\(U_{Glyc}\)) were determined by matching our model calculation with experimentally determined *M. tuberculosis* cell concentrations. As for other carbon-containing substrates (arginine, cytidine, isoleucine, leucine, ornithine, phenol palmic acid, tyrosine, valine, and xylose), we decreased the upper limits of their uptakes (\(U_C\)) to a small value [0.005 mmol h\(^{-1}\) g dry wt\(^{-1}\) (i.e., 0.005 millimoles per hour per gram dry weight of *M. tuberculosis*)] below which they generated new false predictions of gene essentiality. We provide the developed iNJ661i network in Systems Biology Markup Language format in the ESI. Compared with the network used in our previous *in vitro* study, the iNJ661i network included the uptakes of more substrates, including glycerol, fatty acids other than propionate, nitrate, and the above listed other carbon-containing substrates.

Finally, we performed FBA of the iNJ661i metabolic network using the COBRA Toolbox to calculate *M. tuberculosis* biomass production rates (\(\mu\)) with different sets of constraints on the target reaction fluxes. We calculated the biomass production rate of wild-type *M. tuberculosis* in the absence of the 3-NP inhibitor by leaving the ICL and MCL reaction fluxes unconstrained. In this case, after FBA, we also used the COBRA Toolbox to minimize the sum of reaction fluxes while keeping the calculated maximal biomass growth rate. This procedure allowed us to obtain a unique set of minimum inhibitor-free fluxes for all reactions, including \(v_{ICL,WT}\) and \(v_{MCL,WT}\) for the ICL and MCL reactions, respectively, corresponding to the most parsimonious flow of metabolites through the network.

We calculated the biomass production rate in the presence of 3-NP by constraining the flux of the ICL and MCL reactions, respectively, to be no more than the product of the flux ratios \(f_{ICL}\) and \(f_{MCL}\), calculated from eqn (3) and (4), and the inhibitor-free fluxes \(v_{ICL,WT}\) and \(v_{MCL,WT}\). That is, the upper limit of the flux through the ICL (or MCL) reaction was \(f_{ICL}v_{ICL,WT}\) (or \(f_{MCL}v_{MCL,WT}\)). We calculated the biomass production rate of the deletion mutant \(\Delta icl1\Delta icl2\) of *M. tuberculosis* by setting the fluxes of the target reactions ICL and MCL to zero. In cases where we explored potential drug combinations with 3-NP, we not only limited the ICL and MCL reaction fluxes but also constrained the reaction flux associated with the other putative target enzyme.

Under each particular set of conditions and constraints, we calculated \(\mu\) and used the population growth model to estimate the time-dependent cell concentrations of *M. tuberculosis*.

**Population growth model**

We used a population growth model to calculate the *M. tuberculosis* concentration \([X]\) as a function of time \(t\), using the biomass production rate \(\mu\) determined from FBA of the metabolic network. This model was an extension of the earlier *in vitro* population model in that it accounted for the elimination of pathogen cells by the macrophage. It also accounted for a delay in the onset of bacterial growth by including an initial lag phase in which bacterial cells are still adapting to the environment and not multiplying. These features were captured by the population growth model by including the macrophage bacterial lysis rate \((k_d)\) and two different equations for the bacterial growth within and after the lag stage, respectively.

Let \(t\) represent time and \(\tau\) the time point at which the lag stage ends. Thus, \(t \leq \tau\) indicates time within the lag stage, and \(t > \tau\) indicates time after the lag stage. Therefore, the growth population model is as follows:

\[
\frac{d[X]}{dt} = -24k_d[X] \text{ at } t \leq \tau
\]

\[
\frac{d[X]}{dt} = 24(\mu - k_d)[X] \text{ at } t > \tau
\]

where \(t\) and \(\tau\) are measured in units of days, \([X]\) represents the cell concentration [in colony-forming units (CFUs)], the factor of 24 converts the units of time from hours to days, and \(k_d\) represents the bacterial lysis rate (in h\(^{-1}\)). We set the value of \(k_d\) to 0.015 h\(^{-1}\), which is compatible with the largest experimental decline of cell concentrations of *M. tuberculosis* in the mouse macrophage. Eqn (5) indicates that cell concentrations could decrease as a function of time during the lag stage. This allowed us to model a possible “pause” in bacterial growth brought on by the change in environment, i.e., bacterial entry into a macrophage, yet still permitted macrophage-induced cell killing. Such initial decrease in cell concentrations of intracellular bacteria during the lag stage has also been experimentally observed in other intracellular pathogens, e.g., for *Legionella pneumophila* taken up by *Acanthamoeba castellanii* cells. To make the calculation results comparable with the experimental data, we integrated eqn (5) and (6) and converted the resultant natural logarithms into common logarithms as follows:

\[
\log_{10}[X](t) = \log_{10}[X](t_0) - 24k_d(t - t_0)/2.303 \text{ at } t \leq \tau
\]

\[
\log_{10}[X](t) = \log_{10}[X](\tau) + 24(\mu - k_d)(t - \tau)/2.303 \text{ at } t > \tau
\]

where \(t_0\) represents the initial time, which based on the experimental data was set to 1 day. The calculated cell concentrations \([X]\) in eqn (7) and (8) were then directly compared with the experimental data.

**Sensitivity analysis**

The presence of a number of parameters in our mathematical framework warranted a sensitivity analysis to ascertain how the assigned parameter values affected the final computational results. Table 2 shows a summary of all parameters used to construct the computational framework. To address sensitivity issues for different types of parameters, we classified the parameters into three groups: (1) those obtained from the literature (group I), (2) those determined by matching experimental data (fitted parameters) (group II), and (3) those that, by
definition, were derived from other parameters, i.e., dependent parameters (group III).

We estimated the relationship between the computational result and the corresponding parameter by calculating the sensitivity coefficient for the parameter. For example, if log$_{10}$[X] represents the common logarithm of cell concentration (in CFUs) and $p$ represents the parameter being analyzed, the sensitivity coefficient ($S_p$) is defined as follows:

$$S_p = \frac{\partial(\log_{10}[X])/\log_{10}[X]}{\partial p/p} \quad (9)$$

We numerically estimated the sensitivity coefficient $S_p$ for parameter $p$ by starting from $\partial p = +0.5p$ and repeating the process of reducing $\partial p$ and calculating $S_p$ until the value of $S_p$ converged. We then repeated the process starting from $\partial p = -0.5p$ until convergence. In the calculations performed here, both processes converged to the same numerical value.

We only performed sensitivity analysis for the parameters in groups I and II, as the variations in the dependent parameters in group III were, by definition, implicitly considered in these analyses.

### Results and discussion

#### Simulation of growth inhibition of M. tuberculosis in macrophage cells

Munoz-Elias and McKinney used $M. tuberculosis$-infected mouse macrophage cells to quantify intracellular pathogen growth during a six-day period in defined media with and without 3-NP. The Munoz-Elias and McKinney study consists of three independent experiments that compare: (1) growth of the Δicl1Δicl2 deletion mutant strain with wild-type $M. tuberculosis$; (2) growth of $M. tuberculosis$ with 10.0 mM 3-NP present in the medium compared with the inhibitor-free condition; and (3) growth of $M. tuberculosis$ with 0.2, 1.0, and 5.0 mM 3-NP in the medium compared with the inhibitor-free condition. Note that the cited 3-NP concentrations in the medium correspond to extracellular concentrations rather than the intracellular concentrations present in the phagosome compartment.

We used the mathematical framework shown in Fig. 1 to attempt to reproduce the results from the above experiments. We performed eight distinct simulations to mimic the growth data for eight different conditions in the above experimental study (see Table 3). For each condition, we simulated the growth of a particular strain of $M. tuberculosis$ (wild-type or the Δicl1Δicl2 deletion mutant) under a distinct extracellular 3-NP concentration by setting the initial cell concentration [X]$_{t_0}$ to the measured value at day 1 and fitting growth data to the experimental cell concentrations at days 4 and 7 to estimate one or two other parameter values. These parameters included the time $\tau$ at which the lag stage ends, the upper limit of the fatty acid uptake ($U_{\text{glyc}}$), the upper limit of fatty acid uptake ($U_{\text{fat}}$), and the intracellular 3-NP concentration ([3-NP]). We determined the value for $\tau$ for each condition, set the uptake limits $U_{\text{glyc}}$ and $U_{\text{fat}}$ in the simulations for Conditions 1 and 2, respectively, and obtained the intracellular 3-NP concentrations [3-NP] in the simulation for Conditions 4–7 (see Section S2 in the ESI† for details).

Fig. 2A–C, shows the simulated cell concentrations of (1) wild-type $M. tuberculosis$ and the Δicl1Δicl2 deletion mutant; (2) $M. tuberculosis$ with and without 10.0 mM 3-NP in the medium; and (3) $M. tuberculosis$ with and without 0.2, 1.0, and 5.0 mM 3-NP in the medium. Table 3 shows the root mean squared error (RMSE) between the simulated and experimental cell concentrations in log$_{10}$ units. In general, the simulated cell concentrations were in close agreement with the experimental data, with a RMSE of less than 0.07. The exception is shown in Fig. 2C, where, in the absence of 3-NP, the simulated cell concentration at day 7 was higher than the corresponding experimental data, with a RMSE of 0.20. The predicted higher cell concentrations stemmed from our assumption that the substrate uptake rates of $M. tuberculosis$ inside the phagosome do not change with time. For drug-induced growth-retarded bacteria, this is an adequate assumption for the time period modeled, but for unrestricted growth, this assumption can introduce errors at days 5 to 7.
Our approach used a static description of nutrient uptakes from the host cell, i.e., the potential dynamic responses of the host were not included. Eventually, we would like to include the time-dependent responses of host cells that influence the metabolism and growth of intracellular pathogens. The recent development of an integrated *M. tuberculosis*-macrophage metabolic model ([A/B]-IAMO-1410-M1-661)\textsuperscript{56} provides the foundation for this effort, although much additional development will be needed to account for all relevant interactions between the host and the pathogen. Further extensions of the present work to incorporate these models and other biological network descriptions of signaling and regulatory networks will allow us to construct a more comprehensive description of drug-induced growth inhibition of intracellular pathogens.

To highlight the difference between our previously in vitro-derived framework\textsuperscript{18} and the deployed macrophage-based framework derived here, we compared the abilities of the two models to reproduce the ex vivo wild-type and mutant growth data. Fig. 3 shows that, although the in vitro simulation results for wild-type *M. tuberculosis* could roughly match the experimental data, the simulated cell concentrations of the Δicl1Δicl2 deletion mutant were inconsistent with the experimental data. To account for the macrophage action in the in vitro framework, we added the lysis rate (*k_d*) to the population growth model and re-simulated the growth of the Δicl1Δicl2 deletion mutant. Fig. 3 shows that the simulation results from this modification improved the correspondence with the experimental data, but the results neither fully captured the experimental data nor the results derived from the intracellular macrophage model simulations, because the in vitro framework failed to predict a non-zero growth rate for the Δicl1Δicl2 deletion mutant.

### Sensitivity analysis of the model parameters

To quantitatively measure how the parameter values affected the simulation results, we calculated the sensitivity coefficients $S_p$ for each of the parameters in *groups I* and *II* (Table 2) at different 3-NP concentrations in the medium. For this analysis, we used the simulation results shown in Fig. 2C at day 4 as references, and, accordingly, $\log_{10}[X]$ in eqn (9) refers to the common logarithms of the calculated cell concentrations at that day. Table 4 shows the calculated sensitivity coefficient for each parameter at different extracellular 3-NP concentrations (0.0, 0.2, 1.0, and 5.0 mM). By construction, the parameters associated with the inhibition model had zero sensitivity in the absence of an inhibitor. Among the parameters in the metabolic network, the upper limit of fatty acids uptake ($U_{fat}$) had a significantly higher sensitivity coefficient than that of glycerol uptake ($U_{glyc}$), suggesting that, among major carbon sources, fatty acids were more important than glycerol for the growth of *M. tuberculosis*. The relatively low sensitivity associated with glycerol uptake was consistent with the experimental observation that in the presence of other carbon sources *M. tuberculosis* glycerol metabolism is not altered.\textsuperscript{57} Thus, the observed fixed metabolic fate of glycerol indicated that

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<table>
<thead>
<tr>
<th>Condition Index</th>
<th>Strain</th>
<th>[3-NP] (mM)</th>
<th>Parameters from the literature</th>
<th>Parameters determined in this condition</th>
<th>Parameters from previous conditions</th>
<th>Figure</th>
<th>RMSE</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Δicl1Δicl2</td>
<td>0.0</td>
<td>$<a href="t_0">X</a> = 10^{3.95}$ CFU, $k_d$ τ = 4.0 day, $U_{glyc} = 0.041$ mmol h$^{-1}$ g dry wt$^{-1}$</td>
<td>—</td>
<td>$U_{glyc}$ from Condition 1</td>
<td>2A</td>
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<tr>
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<td>$<a href="t_0">X</a> = 10^{3.95}$ CFU, $k_d$ τ = 2.2 day, $U_{fat} = 0.014$ mmol h$^{-1}$ g dry wt$^{-1}$</td>
<td>$U_{glyc}$ from Condition 1</td>
<td>$U_{fat}$ from Condition 1</td>
<td>2A</td>
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</tr>
<tr>
<td>3</td>
<td>Wild-type</td>
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<td>$<a href="t_0">X</a> = 10^{4.10}$ CFU, $k_d$ τ = 1.5 day</td>
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<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
<td>2B</td>
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<tr>
<td>4</td>
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<td>$<a href="t_0">X</a> = 10^{4.07}$ CFU, $k_d$ and 11 parameters of inhibition model τ = 2.4 day, $[3-NP]_i = 0.118$ mM</td>
<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
<td>2C</td>
<td>0.001</td>
</tr>
<tr>
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<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
<td>2C</td>
<td>0.032</td>
</tr>
<tr>
<td>6</td>
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<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
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<td>0.015</td>
</tr>
<tr>
<td>7</td>
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<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
<td>2C</td>
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</tr>
<tr>
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<td>$U_{glyc}$ from Condition 1, $U_{fat}$ from Condition 2</td>
<td>2C</td>
<td>0.204</td>
</tr>
</tbody>
</table>
**M. tuberculosis** might be unable to adjust its glycerol metabolism to optimize growth. Instead, the organism might attempt to optimize its growth by adjusting its fatty acid metabolism, reinforcing the notion that, compared with glycerol, fatty acids might play a more important role in the growth of **M. tuberculosis** under limiting nutrient conditions.

The relatively small sensitivity coefficient of the upper limit of the uptakes of other carbon-containing substrates ($U_C$) also indicated that fatty acids were more important than these substrates. For the parameters in the population growth model, the initial cell concentration, $\log_{10}X(t_0)$, had a large sensitivity coefficient, as the initial cell concentration directly affects the concentrations at later time points. The lysis rate ($k_d$) had relatively small coefficients, suggesting that the lysis of the **M. tuberculosis** cells had a small influence on cell concentration. This lack of influence also propagated to small sensitivity coefficients for the lag time ($\tau$), as before this time, only lysis affected the concentration of **M. tuberculosis**.

**Analysis of the relation between the intracellular and extracellular inhibitor concentrations**

In the experiments, the extracellular 3-NP concentrations were set to 0.2, 1.0, 5.0, and 10.0 mM, while, correspondingly, we

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**Fig. 2** Simulation results and experimental data for the growth of *Mycobacterium tuberculosis* in the absence and presence of 3-nitropropionate (3-NP) inhibitor. Growth curves were compared between (A) wild-type *M. tuberculosis* cells and the Δicl1Δicl2 deletion mutant, (B) wild-type cells in medium with and without 10 mM 3-nitropropionate (3-NP), and (C) wild-type cells in medium with 3-NP concentrations of 0.2, 1.0, and 5.0 mM and without 3-NP. The experimental data were taken from the literature. [X] represents the cell concentration of *M. tuberculosis*; $t$ represents time. CFU, colony-forming units.

**Fig. 3** Simulation results of the growth of wild-type *Mycobacterium tuberculosis* cells and the Δicl1Δicl2 deletion mutant using the previous *in vitro* framework. The solid line shows cell concentrations of wild-type cells calculated using the *in vitro* framework, the dotted-dashed line shows cell concentrations of the Δicl1Δicl2 deletion mutant calculated using the *in vitro* framework, and the dashed line shows cell concentrations of the Δicl1Δicl2 deletion mutant calculated using the *in vitro* framework with the lysis rate ($k_d$). The experimental data were taken from the literature.
used our model framework to determine the effective intracellular 3-NP concentrations to be 0.012, 0.023, 0.063, and 0.118 mM, respectively. This was done by adjusting the intracellular 3-NP concentrations to values that reproduced the experimentally measured cell concentrations at different extracellular 3-NP concentrations (see Table 3) under fixed extracellular 3-NP concentrations ([3-NP]_e). The solid line shows the linear regression for the four data points. The coefficient of determination (R^2) indicates how well a linear relation fits the four data points. A F-test was used to estimate the statistical significance in the form of a p-value of the linear relation.

performed a more robust analysis of the linearity by increasing and decreasing each parameter in group I (Table 2) by 50%, except for those parameters whose values cannot exceed one, and recalculating the intracellular 3-NP concentrations. Table 5 shows the calculated intracellular 3-NP concentrations, coefficients of determination (R^2), and the p-values from the F-test associated with changing the parameter values. With increased or decreased parameter values, the obtained intracellular 3-NP concentrations could be very different from the original ones, e.g., the determined intracellular inhibitor concentrations were quite sensitive to changes in \( w_{\text{C1L1}} \). However, the R^2 and p-values were always above 0.990 and below 0.050, respectively, indicating that the linear relation between the intracellular and extracellular 3-NP concentrations was robust to the changes in parameter values.

The lack of explicitly measured 3-NP concentrations in the phagosome compartments of the macrophage prevents a direct experimental verification of the linear relation proposed here. Instead, we used indirect approaches to investigate the possible mechanism that could underlie the linear relation between the intracellular and extracellular 3-NP concentrations. When we assumed that the inhibitor molecules entered the mouse macrophages through diffusion and that, at some intracellular concentration threshold, the macrophages initiated active efflux transport of the inhibitor, we found that the steady-state intracellular and extracellular 3-NP concentrations could indeed be expressed by a linear relation (see Section S3 in the ESI† for details):

\[
[3\text{-NP}]_i = \alpha[3\text{-NP}]_e + B \tag{10}
\]

where \([3\text{-NP}]_i\) denotes the extracellular 3-NP concentration, the subscript “e” indicates extracellular concentration, and \(\alpha\) and \(B\) are constants independent of the value of \([3\text{-NP}]_e\). The linear relation between the intracellular and extracellular 3-NP concentrations is thus plausible from a cellular pharmacodynamics and pharmacokinetics standpoint. Although direct evidence for 3-NP is as yet unavailable, experimentally...

![Fig. 4 Relation between intracellular 3-nitropropionate (3-NP) concentrations and extracellular 3-NP concentrations. Diamonds indicate data points of the modeled relationship between intracellular and extracellular 3-nitropropionate (3-NP) concentrations. Intracellular 3-NP concentrations ([3-NP]_i) were determined by adjusting their values in our model to obtain the experimentally measured cell concentrations (see Table 3) under fixed extracellular 3-NP concentrations ([3-NP]_e). The solid line shows the linear regression for the four data points. The coefficient of determination (R^2) indicates how well a linear relation fits the four data points. A F-test was used to estimate the statistical significance in the form of a p-value of the linear relation.](image-url)
measured average concentrations in macrophages exist for some inhibitors, such as β-lactams, and the linear relationship between drug concentrations inside and outside macrophages has been experimentally verified for a β-lactam, penicillin.

### Combinational effects of 3-NP and other enzyme inhibitions

The strategy of combining multiple drugs is promising from considerations of efficacy, safety, and potential reduction in emerging drug resistance. Although not explicitly treated here, the emergence of drug resistance through escape mutants could be reduced if different drugs affect different targets. Previous studies have shown that modeling the inhibition of metabolic pathways can help identify potential synergistic inhibitory effects of drug combinations and suggest novel drug combination therapies.

We used the mathematical framework to computationally study the growth inhibition of *M. tuberculosis* by different combinations of two sets of “drugs” separately targeting (1) the ICL1/ICL2 and malate synthase (MS) enzymes and (2) the ICL1/ICL2 and glycerol-3-phosphate dehydrogenase (G3PD) enzymes. MS was chosen as it is considered a possible drug target for *M. tuberculosis* and is part of the ICL1/ICL2 pathway. We selected G3PD because it is not part of the same pathway as ICL1/ICL2 yet is crucial for the utilization of glycerol, an intracellularly available carbon source other than fatty acids. We also examined whether the *in vitro* framework used to study *M. tuberculosis* growth on propionate medium was capable of exhibiting synergistic inhibition. Here, we used synergy in the sense that two or more drugs working together produce a result not obtainable by any of them independently. In the following sections, we considered the actual drug concentrations for 3-NP, but for the second inhibition we mimicked the drug action by directly reducing the flux of materials through the corresponding enzyme-catalyzed reaction.

### ICL1/ICL2-MS inhibition

The first drug combination study analyzed the effect of 3-NP and concomitant inhibition of the MS enzyme on the growth of *M. tuberculosis*. With the assigned parameter values from Table 3, we used the mathematical framework to calculate the cell concentrations of *M. tuberculosis* after six days of growth at different 3-NP concentrations and at different degrees of inhibition of the MS enzyme. Fig. 5A shows the calculated dose-response curves when we constrained the flux through the MS enzyme to 100%, 67%, 33%, and 0% of the wild-type flux value.

In general, these curves indicated that MS inhibition was able to reduce the dose requirement of 3-NP to achieve a particular inhibitory effect. We can quantify this effect by calculating the reduction in 3-NP concentration that can be achieved by the combination. For example, to achieve 80% of the greatest possible 3-NP inhibitory effect...
functions of 3-nitropropionate (3-NP) concentrations ([3-NP]) obtained from: (A) the intracellular framework and (B) the in vitro framework, where the flux through the malate synthase (MS) enzyme was constrained to 100%, 67%, 33%, and 0% of its wild-type flux. The “80% effect” indicates 80% of the greatest decrease in log_{10}[X] that could be achieved with 3-NP, where specifically the “greatest decrease” indicates the difference between log_{10}[X] without 3-NP and at the highest 3-NP concentration. [3-NP]_{WT} represents the 3-NP concentration required to achieve the “80% 3-NP effect” without constraint on the flux through the MS enzyme, and [3-NP]_{AMS} represents the required 3-NP concentration with zero flux through the MS enzyme. Δlog_{10}[3-NP] = log_{10}[3-NP]_{WT} − log_{10}[3-NP]_{AMS}. CFU, colony-forming unit.

Fig. 5 Effects of the combined inhibition of 3-nitropropionate (3-NP) and the malate synthase (MS) enzyme. Mycobacterium tuberculosis cell concentrations after six days of growth ([X]) as functions of 3-nitropropionate (3-NP) concentrations ([3-NP]) obtained from: (A) the intracellular framework and (B) the in vitro framework, where the flux through the malate synthase (MS) enzyme was constrained to 100%, 67%, 33%, and 0% of its wild-type flux. The “80% effect” indicates 80% of the greatest decrease in log_{10}[X] that could be achieved with 3-NP, where specifically the “greatest decrease” indicates the difference between log_{10}[X] without 3-NP and at the highest 3-NP concentration. [3-NP]_{WT} represents the 3-NP concentration required to achieve the “80% 3-NP effect” without constraint on the flux through the MS enzyme, and [3-NP]_{AMS} represents the required 3-NP concentration with zero flux through the MS enzyme. Δlog_{10}[3-NP] = log_{10}[3-NP]_{WT} − log_{10}[3-NP]_{AMS}. CFU, colony-forming unit.

Thus, with a complete inhibition of the MS enzyme, the 80% inhibition effect could be achieved by only using 13.4% (10^{Δlog_{10}[3-NP]} = 10^{−0.874} = 0.134) of the original concentration of 3-NP.

The effect of MS inhibition was not synergistic, because MS inhibition could not further reduce the pathogen cell concentration below the maximum inhibition achieved by 3-NP. This is because both the MS enzyme and 3-NP-targeted ICL1/ICL2 enzymes are all part of the glyoxylate pathway that processes isocitrate. ICL1/ICL2 converts isocitrate to glyoxylate, which, in turn, is converted by MS into S-malate. This pathway is required for M. tuberculosis to use fatty acids, the major carbon sources for the pathogen in macrophages. In essence, targeting both ICL1/ICL2 and MS enzymes inhibits the same flux through the glyoxylate pathway. Thus, one could either target ICL1/ICL2 or MS to inhibit this pathway, but no additional reduction in the growth of the pathogen would be possible using this drug combination strategy. However, it may be important to explore a different druggable target in the same pathway under certain conditions. For example, an existing drug for a particular target in the pathway might be associated with non-optimal pharmacological properties, making it desirable to either develop alternative drugs against that target or explore other druggable targets in the same pathway. Another condition could be that, if the known target enzyme were capable of mutating under drug pressure and potentially giving rise to a drug-resistant pathogen, it would be desirable to have two druggable targets in the same pathway as the likelihood of two enzyme mutations arising at the same time is considerably lower than that of any single one.

We examined how robust these observations were with respect to parameter variation in the model framework. We increased and decreased the value of each model parameter by 50% (except for those parameters whose values cannot exceed one) and calculated the resultant values for Δlog_{10}[3-NP]. Table 6 shows that the values for Δlog_{10}[3-NP] ranged from 0.498–1.320, indicating that the required 3-NP concentration was 4.8–31.8% (=10^{−1.320−10^{−0.498}}) of [3-NP]_{WT}. This suggests that the observation was robust with respect to variations in the parameters of the model.

We also examined whether the previously developed in vitro framework was associated with quantitative or qualitative differences compared with our macrophage-based model. Fig. 5B shows the calculated dose-response curves and indicates that the inhibition of the MS enzyme did not influence the dose-response curves as much as in Fig. 5A. This decreased effect stems from the ICL reaction not playing a crucial role under the in vitro condition. In the simulated in vitro conditions, propionate, a fatty acid, is the main carbon source, and the utilization of propionate mainly depends on the MCL reaction and not on the ICL reaction.

ICL1/ICL2-G3PD inhibition. The second drug combination used 3-NP as the primary drug and modeled the inhibition of the G3PD enzyme as the second target. We used the same mathematical framework and parameters used in the ICL1/ICL2-MS study above to calculate the cell concentrations of M. tuberculosis after six days of growth at different 3-NP concentrations and at different degrees of inhibition of the G3PD enzyme. Fig. 6A shows the calculated dose-response curves when we constrained the flux through the G3PD enzyme to 100%, 67%, 33%, and 0% of the wild-type flux values.

One can note two key features from this curve: (1) fully inhibiting the G3PD enzyme by itself at insignificant 3-NP concentrations did not significantly reduce M. tuberculosis growth; and (2) at higher 3-NP concentrations, inhibition of G3PD further reduces growth beyond what was possible with
Table 6 Effects of the parameter values on the drug combinations. \( \Delta \log_{10}[3\text{-NP}] \) indicates the difference between \( \log_{10}[3\text{-NP}]_{\text{WT}} \) and \( \log_{10}[3\text{-NP}]_{\text{AMS}} \), where \( [3\text{-NP}]_{\text{WT}} \) represents the 3-nitropropionate (3-NP) concentration required to obtain an 80% inhibitory effect of 3-NP without the inhibition of the malate synthase (MS) enzyme and \( [3\text{-NP}]_{\text{AMS}} \) represents the required 3-NP concentration with full inhibition of the enzyme. \( \Delta \log_{10}[X] \) indicates the difference between \( \log_{10}[X]_{\text{WT}} \) and \( \log_{10}[X]_{\text{AMS}} \), where \( [X]_{\text{WT}} \) represents the cell concentrations of \( M.\) \text{tuberculosis} at a high 3-NP concentration (10^2 mM) in the absence of inhibition of the glycerol-3-phosphate dehydrogenase (G3PD) enzyme and \( [X]_{\text{AMS}} \) represents the cell concentration under full inhibition of the G3PD enzyme.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Change</th>
<th>Inhibition of MS</th>
<th>Inhibition of G3PD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Original values</td>
<td></td>
<td>( \Delta \log_{10}[3\text{-NP}] )</td>
<td>( \log_{10}[X]_{\text{WT}} )</td>
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<tr>
<td>([\text{SUC}])</td>
<td>+ 50%</td>
<td>0.874</td>
<td>13.4</td>
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<tr>
<td>(-50% )</td>
<td>0.844</td>
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<tr>
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<tr>
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<td>0.874</td>
<td>13.4</td>
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<tr>
<td>(-50% )</td>
<td>0.874</td>
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3-NP alone. Thus, we have the situation where inhibition of a non-essential enzyme potentiates the action of 3-NP in a synergistic manner.

We quantified this effect at a 3-NP concentration of 10^2 mM by calculating the additional reduction in cell concentration \( \Delta \log_{10}[X] \) when the G3PD flux was constrained to zero:

\[
\Delta \log_{10}[X] = \log_{10}[X]_{\text{WT}} - \log_{10}[X]_{\text{AMS}}
\]

where \( [X]_{\text{WT}} \) denotes the calculated cell concentrations in the presence of \( [3\text{-NP}] = 10^2 \) mM and \( [X]_{\text{AMS}} \) denotes cell concentrations in the presence of both “drugs.” The additional reduction in cell growth beyond what was achievable with 3-NP alone was 38.3% \( (10^{\log_{10}[X]} = 10^{-0.417} = 0.383) \). The synergistic effect stems from the utilization of both fatty acids (targeted by 3-NP) and glycerol (affected by G3PD inhibition) as carbon sources for \( M.\) \text{tuberculosis} in macrophages.\(^{11,19,30}\)

The pathways processing these carbon sources are not directly connected, and the fluxes through these enzyme reactions are much more independent than what was the case for ICL1/ICL2-MS enzymes in the glyoxylate shunt.

Similar to the ICL1/ICL2-MS inhibition, we examined whether the parameter values in the intracellular framework affected the observed synergistic effect in the ICL1/ICL2-G3PD inhibition combination. Table 6 shows that the changes of most parameters did not affect the values of \( \Delta \log_{10}[X] \). Only the variation in the upper limit of glycerol uptake \( (U_{\text{Glyc}}) \) significantly changed the value of \( \Delta \log_{10}[X] \), suggesting that the synergistic effect of 3-NP and G3PD inhibition mainly depended on the uptake of glycerol. Conversely, the observed effect on \( M.\) \text{tuberculosis} would only be present when glycerol is an important nutrient source in the macrophage.

We also compared the modeled effects with the previously developed \textit{in vitro} framework.\(^{18}\) Fig. 6B shows that the calculated dose-response curves did not show any synergistic effects in that the additional inhibition of the G3PD flux could not decrease the growth of \( M.\) \text{tuberculosis} beyond 3-NP inhibition. The reason for this behavior is that the \textit{in vitro} medium contains different nutrient sources. Thus, when constraining the G3PD flux to 33% of its wild-type value, the cell concentration \( [X] \) was independent of 3-NP concentrations.
lower than 0.024 mM, indicating that the primary inhibition mechanisms under these conditions were through the G3PD catalyzed reaction. At a 3-NP concentration higher than 0.024 mM, the dose-response curve indicates that the inhibition was independent of the G3PD reaction and solely driven by 3-NP inhibition. In the *in vitro* propionate medium, *M. tuberculosis* synthesizes glycerol-3-phosphate from propionate through a serial set of reactions, including the MCL reaction (inhibited by 3-NP) and the G3PD-catalyzed reaction. When we placed constraints on the fluxes of a set of serially connected reactions, only the strictest constraint affected the calculated cell concentrations of *M. tuberculosis*, and, hence, no synergistic effects were possible.

**Overall assessment of ICL1/ICL2 inhibition partners**

To examine other possible non-intuitive synergistic inhibition partners with ICL1/ICL2 among the metabolic enzymes and pathways in *M. tuberculosis*, we calculated $\Delta \log_{10}[X]$ for the inhibition of each reaction in the *iNJ661* metabolic network (provided in the ESI† as Delta_log10X_all_011311.xls). The top seven reactions with the largest $\Delta \log_{10}[X]$ values were all associated with the central carbon metabolism pathways, highlighting the importance of these pathways in providing possible synergistic targets. Fig. 7 shows the predicted synergistic level of each reaction in combination with 3-NP in the central carbon metabolism of *M. tuberculosis*. The dotted arrows indicate reactions that are catalyzed by the isocitrate lyase 1 (ICL1) and isocitrate lyase 2 (ICL2) enzymes and thus targeted by 3-nitropropionate (3-NP). Solid arrows represent other reactions in the central carbon metabolism, and the color intensity indicates the synergy levels of the corresponding reactions with 3-NP. For each reaction RXN, the synergy level was computed using the value for $\Delta \log_{10}[X] = \min(\log_{10}[X]_{WT}, \log_{10}[X]_{0,\text{ARXN}}) - \log_{10}[X]_{\text{ARXN}}$, where $[X]_{WT}$ represents the cell concentration at $10^2$ mM 3-NP, $[X]_{0,\text{ARXN}}$ represents the cell concentration at the zero 3-NP concentration level with the flux through the reaction RXN constrained to zero, and $[X]_{\text{ARXN}}$ represents the cell concentration at $10^2$ mM 3-NP with the flux through the reaction RXN constrained to zero. FUM, fumarase; G3PD, glycerol-3-phosphate dehydrogenase; GAPD, glyceraldehyde-3-phosphate dehydrogenase; GK, glycerol kinase; ICDH, isocitrate dehydrogenase; MS, malate synthase; PGK, phosphoglycerate kinase; TPI, triose phosphate isomerase.

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Fig. 6 Effects of the combined inhibition of 3-nitropropionate (3-NP) and the glycerol-3-phosphate dehydrogenase (G3PD) enzyme. *Mycobacterium tuberculosis* cell concentrations after six days of growth ([X]) as functions of 3-NP concentrations ([3-NP]) obtained from: (A) the intracellular framework and (B) the *in vitro* framework, where the flux through the glycerol-3-phosphate dehydrogenase (G3PD) enzyme was constrained to 100%, 67%, 33%, and 0% of its wild-type flux. $[X]_{\text{WT}}$ represents the cell concentration at $10^2$ mM 3-NP, when there was no constraint on the flux through the G3PD enzyme, and $[X]_{\text{G3PD}}$ represents the cell concentration when the flux through the G3PD enzyme was constrained to zero. $\Delta \log_{10}[X] = \log_{10}[X]_{\text{WT}} - \log_{10}[X]_{\text{G3PD}}$. CFU, colony-forming unit.

Fig. 7 Calculated level of synergy between 3-NP and each reaction in the central carbon metabolism of *Mycobacterium tuberculosis*. The dotted arrows indicate reactions that are catalyzed by the isocitrate lyase 1 (ICL1) and isocitrate lyase 2 (ICL2) enzymes and thus targeted by 3-nitropropionate (3-NP). Solid arrows represent other reactions in the central carbon metabolism, and the color intensity indicates the synergy levels of the corresponding reactions with 3-NP. For each reaction RXN, the synergy level was computed using the value for $\Delta \log_{10}[X] = \min(\log_{10}[X]_{\text{WT}}, \log_{10}[X]_{0,\text{ARXN}}) - \log_{10}[X]_{\text{ARXN}}$, where $[X]_{\text{WT}}$ represents the cell concentration at $10^2$ mM 3-NP, $[X]_{0,\text{ARXN}}$ represents the cell concentration at the zero 3-NP concentration level with the flux through the reaction RXN constrained to zero, and $[X]_{\text{ARXN}}$ represents the cell concentration at $10^2$ mM 3-NP with the flux through the reaction RXN constrained to zero. FUM, fumarase; G3PD, glycerol-3-phosphate dehydrogenase; GAPD, glyceraldehyde-3-phosphate dehydrogenase; GK, glycerol kinase; ICDH, isocitrate dehydrogenase; MS, malate synthase; PGK, phosphoglycerate kinase; TPI, triose phosphate isomerase.
Curves for jointly inhibiting ICL1/ICL2-GK were similar to those in the ICL1/ICL2-G3PD study (results not shown), suggesting that the entire glycerol-processing pathway and possible regulatory mechanisms of this pathway could be considered synergistic drug-targets with 3-NP. The reactions, catalyzed by triose phosphate isomerase (TPI), glyceraldehyde 3-phosphate dehydrogenase (GAPD), and phosphoglycerate kinase (PGK), had moderately high synergistic levels. These reactions were necessary for the synthesis of serine precursors from glycerol when we blocked the synthesis routes from fatty acids by 3-NP inhibition. The last two reactions associated with synergistic inhibition in the central carbon metabolism were catalyzed by isocitrate dehydrogenase (ICDH) and fumarase (FUM) as part of the tricarboxylic acid (TCA) cycle. While inhibiting the MS enzyme in the glyoxylate shunt pathway did not provide synergistic benefits, inhibition of these additional targets blocked the TCA cycle through which the bacterium utilizes fatty acids as carbon sources. Thus, inhibition of the glyoxylate shunt in the macrophage environment highlighted the importance of M. tuberculosis pathways converting glycerol to other metabolites and selected parts of the TCA cycle for the growth and viability of the pathogen. It is necessary, however, to verify our predictions of synergy and validate their importance in different experimental settings before proceeding with a targeted drug-development effort against pairs or combinations of metabolic pathway targets.

Conclusion

The major promise of a systems biology approach lies in the ability to determine and describe emergent properties of a system from an underlying description of each of its components. Attractive as a concept and highly relevant to complex problems, such as cancer, traumatic brain injury, and infectious diseases, systems biology can provide the framework to understand the relationships among a multitude of cellular components. Translating these concepts into quantitative and qualitative computational models impacts both rational target selection and understanding systemic effects of drugs. Our work, focusing on the metabolic adjustments required of intracellular pathogens when colonizing within in vitro, ex vivo, and in vivo environments, presents a novel construct to realize both quantitative and qualitative aspects of systems biology modeling with the availability of experimental data and parameters. Here, we mitigated this problem by performing extensive parameter sensitivity analyses to verify that our results and insights were robust to changes in the parameter values. We feel that the promise of systems biology will be fulfilled by both being able to use complex models to create very specific quantitative predictions as well as in creating qualitative predictions that yield specific well-defined testable hypotheses. This is the balance we have tried to strike in the current work: specific quantitative model of the inhibitory effect of 3-NP on M. tuberculosis growth inside murine macrophages and a more qualitative discussion on possible synergistic targets based on metabolic considerations, e.g., the relative importance of the glycerol-processing pathway under limiting nutritional conditions and when targeting the glyoxylate shunt with drugs.

Ultimately, the ability to model the dynamics of host-pathogen interactions under in vivo conditions by incorporating more extensive metabolic, signaling, and transcriptional regulatory network models will provide a platform for both host and pathogen drug-target identification that optimally selects potent and safe drug combinations.

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References
