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# A genome-scale metabolic model of a globally disseminated hyperinvasive M1 strain of *Streptococcus pyogenes*

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**ABSTRACT** *Streptococcus pyogenes* is responsible for a range of diseases in humans contributing significantly to morbidity and mortality. Among more than 200 serotypes of *S. pyogenes*, serotype M1 strains hold the greatest clinical relevance due to their high prevalence in severe human infections. To enhance our understanding of pathogenesis and discovery of potential therapeutic approaches, we have developed the first genome-scale metabolic model (GEM) for a serotype M1 *S. pyogenes* strain, which we name iYH543. The curation of iYH543 involved cross-referencing a draft GEM of *S. pyogenes* serotype M1 from the AGORA2 database with gene essentiality and autotrophy data obtained from transposon mutagenesis-based and growth screens. We achieved a 92.6% (503/543 genes) accuracy in predicting gene essentiality and a 95% (19/20 amino acids) accuracy in predicting amino acid auxotrophy. Additionally, Biolog Phenotype microarrays were employed to examine the growth phenotypes of *S. pyogenes*, which further contributed to the refinement of iYH543. Notably, iYH543 demonstrated 88% accuracy (168/190 carbon sources) in predicting growth on various sole carbon sources. Discrepancies observed between iYH543 and the actual behavior of living *S. pyogenes* highlighted areas of uncertainty in the current understanding of *S. pyogenes* metabolism. iYH543 offers novel insights and hypotheses that can guide future research efforts and ultimately inform novel therapeutic strategies.

**IMPORTANCE** Genome-scale models (GEMs) play a crucial role in investigating bacterial metabolism, predicting the effects of inhibiting specific metabolic genes and pathways, and aiding in the identification of potential drug targets. Here, we have developed the first GEM for the *S. pyogenes* highly virulent serotype, M1, which we name iYH543. The iYH543 achieved high accuracy in predicting gene essentiality. We also show that the knowledge obtained by substituting actual measurement values for iYH543 helps us gain insights that connect metabolism and virulence. iYH543 will serve as a useful tool for rational drug design targeting *S. pyogenes* metabolism and computational screening to investigate the interplay between inhibiting virulence factor synthesis and growth.

**KEYWORDS** *Streptococcus pyogenes*, metabolic modeling, genome-scale model, auxotrophy, carbon sources, essential gene

*Streptococcus pyogenes* is responsible for a wide range of diseases in humans, causing over 700 million infections and at least 517,000 deaths worldwide annually (1). Among the 200+ serotypes of *S. pyogenes*, serotype M1 is the most frequently isolated from streptococcal pharyngitis (2) and invasive diseases worldwide (3), and it is extensively studied to assess virulence mechanisms during infection (4–7). On the other hand, serotype M49 is commonly used by researchers due to the ease of genetic manipulation, despite generally possessing weaker disease phenotypes in infection models (8). Studies on particularly pathogenic *S. pyogenes* strains, including those of

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serotype M1, have revealed the significance of metabolic and transporter genes as virulence factors in necrotizing myositis (4) and pharyngitis (5). These genes, thus, hold potential as novel therapeutic targets. Genome-scale models (GEMs) play a crucial role in investigating bacterial metabolism, predicting the effects of inhibiting specific metabolic genes and pathways, and aiding in the identification of potential drug targets (9, 10). Although a GEM for *S. pyogenes* serotype M49 (strain 591) was previously developed by Levering et al. (11), there is currently no GEM available for predicting metabolism and identifying drug targets specifically for strains of the more virulent and clinically relevant M1 serotype.

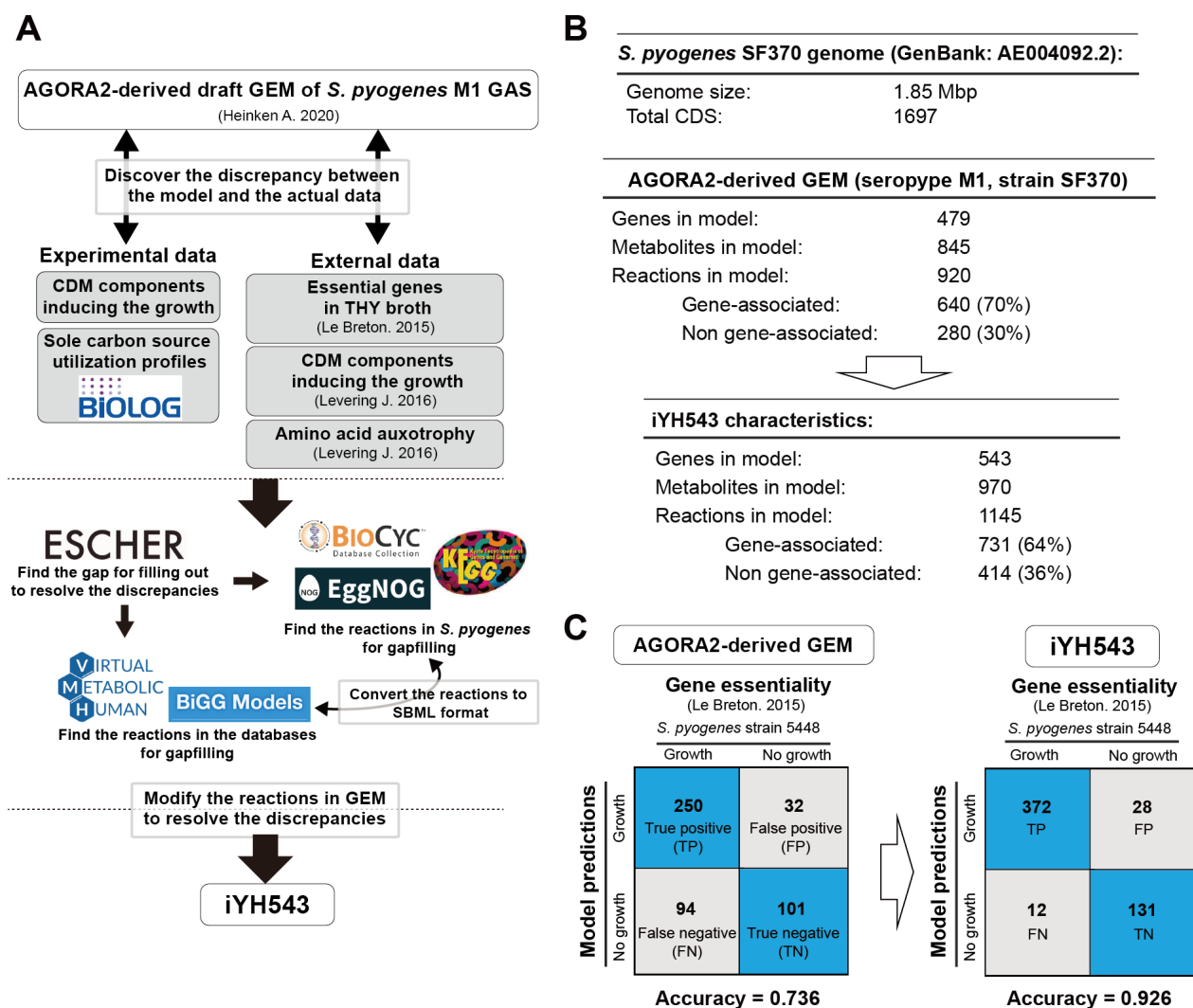
In this study, we developed and curated a highly accurate GEM for *S. pyogenes* serotype M1. To construct this model, we leveraged a draft GEM of *S. pyogenes* serotype M1 (strain SF370) from the AGORA2 computational resource (12), which we manually curated using information from various databases including BiGG (13), VMH (14), BioCyc (15), KEGG (16), and EC number information from EggNOG (17). We carefully examined newly generated experimental data (Biolog growth data) and external experimental data, including conditionally defined media (CDM) growth and transposon mutagenesis-based gene essentiality of *S. pyogenes* M1 (strain 5448) (18), to resolve discrepancies within the initial draft GEM derived from AGORA2. As a result, we significantly improved the accuracy of gene essentiality predictions from 73.6% (351/477 genes) in the draft GEM to 92.6% (514/543 genes) in the final curated GEM, iYH543. This accuracy surpasses the previously reported *S. pyogenes* GEM, which achieved 76.6% accuracy. Furthermore, we demonstrate that iYH543 can be effectively used to investigate the metabolic behavior of *S. pyogenes* on the genome scale using flux balance analysis (FBA) simulation (19). This provides a highly accurate platform to explore the metabolic behavior of *S. pyogenes* under diverse growth conditions, which may aid in the identification of novel therapeutic leads.

## RESULTS

### Overview of the AGORA2-derived draft GEM of *S. pyogenes* serotype M1 and iYH543

In this study, we present iYH543, a curated GEM of *S. pyogenes* serotype M1. The strategy used to produce iYH543 is summarized in Fig. 1. We started with a draft GEM of *S. pyogenes* serotype M1 strain SF370 derived from AGORA2, which contained 479 genes, 845 metabolites, and 920 reactions. To incorporate known gene essentiality (18) and auxotrophy (4, 11) information, we modified the draft GEM by adding 239 reactions, modifying 112 gene–protein–reaction (GPR) rules, deleting three reactions, and adjusting the biomass reaction (bio1; the objective of our flux balance analysis). Detailed modifications are provided in Table S1. To validate and further refine the draft GEM, we assessed sole carbon source utilization profiles and growth in various components of a CDM. Discrepancies between the experimental data and draft GEM were visualized using Escher (20) and addressed by referencing information from BiGG (13), VMH (14), BioCyc (15), KEGG (16), and EggNOG (17) (Fig. 1A). The manual curation processes for iYH543 are explained in subsequent sections. The final curated GEM, iYH543, consisted of 543 genes, 970 metabolites, and 1,145 reactions (Fig. 1B). The details of all reactions, metabolites, and GPRs in iYH543 can be found in Table S2 or DataS1\_iYH543.json at [http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirose_et_al.zip). Computing the essential genes in each model and comparing results with a published genome-scale mutant screen (18), the draft GEM achieved 73.6% accuracy, while iYH543 had a significantly improved accuracy of 92.6% (Fig. 1C).

Le Breton et al. conducted a study on the essentiality of 227 genes for *S. pyogenes* strain 5448, which belongs to the M1 serotype and is closely related to strain SF370. The study was performed under standard laboratory conditions (37°C, 5% CO<sub>2</sub>) using Todd–Hewitt yeast (THY) medium (18). Among the 227 essential genes identified in strain 5448, strain SF370 possesses 224 corresponding orthologs, as determined through



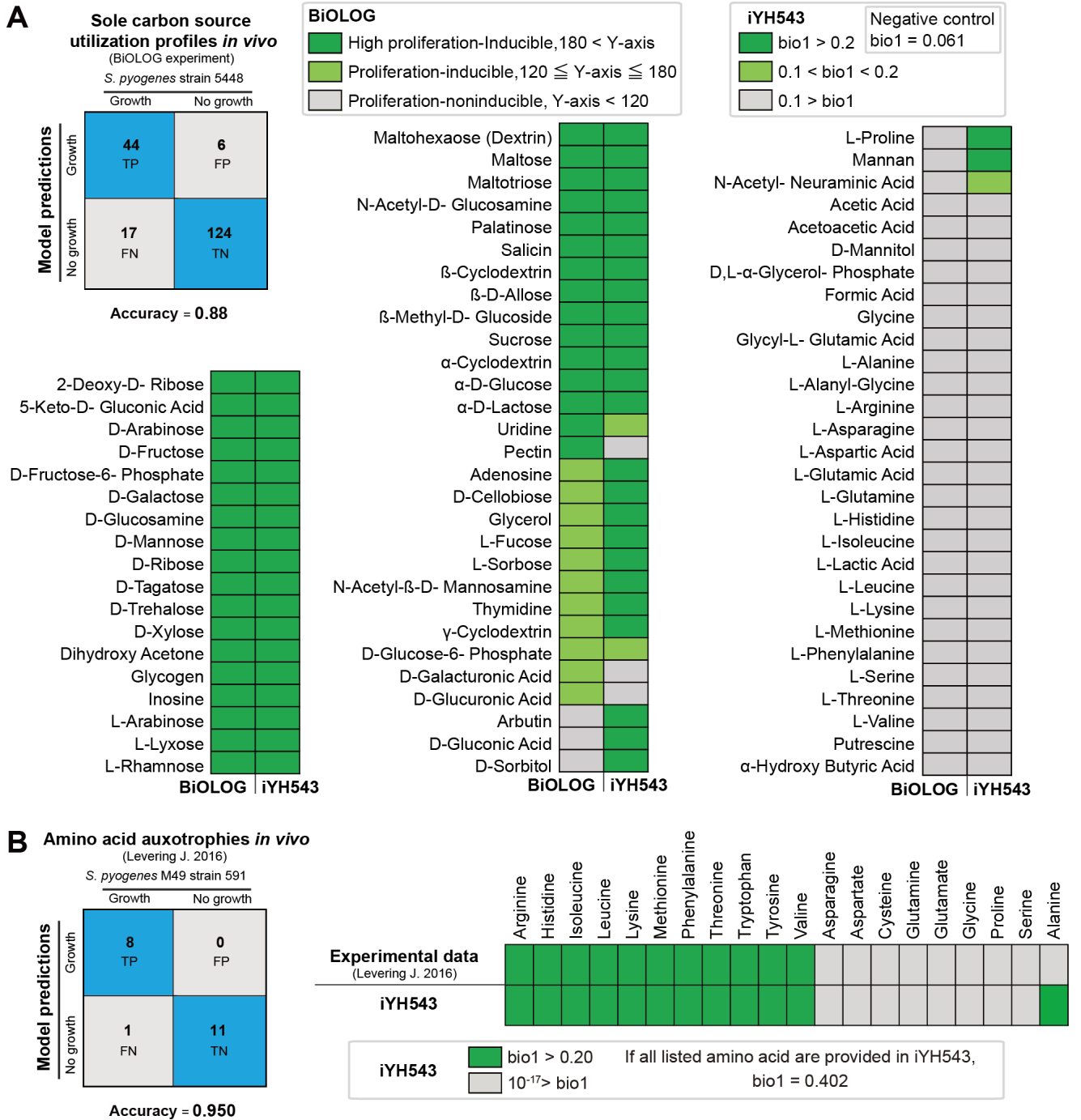
**FIG 1** Curation strategy of the AGORA2-derived genome-scale metabolic model (draft GEM) of *S. pyogenes* M1 GAS (strain SF370) to produce iYH543. (A) Schematic of the network reconstruction strategy for iYH543. The experimental data from this study and previously published data from other studies were compared with the results of the simulation in the AGORA2-derived draft GEM. Based on the identified discrepancies, the model was curated and reconstructed to produce iYH543. (B) Attributes of the AGORA2-derived draft GEM and iYH543. Details regarding the genes, metabolites, and reactions included in iYH543 are listed in Table S2. (C) Accuracy of essentiality predictions in the AGORA2-derived draft GEM and iYH543. Transposon mutagenesis-based gene essentiality in *S. pyogenes* strain 5448 (serotype M1) was used for validation. AGORA2 (12). Transposon mutagenesis-based gene essentiality (18). CDM components and amino acid auxotrophy (11). SBML, Systems Biology Markup Language. Accuracy: (TP + TN)/all genes.

bidirectional BLAST hits. Notably, the three unique proteins in strain 5448 are not metabolic enzymes or transporters. Of the 224 orthologous genes, 65 are not included in iYH543 (Fig. 2; Table S3). These genes mainly fall into clusters of orthologous groups (COGs) of protein categories S (function unknown), M (cell wall/membrane/envelope biogenesis), D (cell cycle control, cell division, and chromosome partitioning), U (intracellular trafficking, secretion, and vesicular transport), and O (posttranslational modification, protein turnover, and chaperones). Based on these findings, it can be inferred that iYH543 encompasses the majority of essential metabolic enzymes and transporters for *S. pyogenes* serotype M1.

### Sole carbon source utilization profiles in *S. pyogenes* serotype M1

We utilized the Biolog microarray system (21) to explore the growth capabilities of *S. pyogenes* strain 5448 in chemically defined medium 1 (CDM1) media (4) with 190





**FIG 3** Sole carbon source utilization profiles and amino acid auxotrophies in iYH543 versus experimental data. (A) Biolog validation. We tested 190 carbon sources to investigate the sole carbon source utilization profiles of *S. pyogenes* M1 serotype (strain 5448). The results of the Biolog experiment are visualized in Fig. S1. The details of the medium for the simulation in iYH543 are shown in Table S6 (CDM1). The results of the simulation in iYH543 are in Table S4 (Carbon\_sources). TP contains the carbon sources that allow *S. pyogenes* growth in both Biolog plates and DataS2\_iYH543\_CDM\_1.json. TN contains (i) the carbon sources that do not allow the growth of *S. pyogenes* in both Biolog plates and DataS2\_iYH543\_CDM\_1.json and (ii) the carbon sources that do not allow the growth of *S. pyogenes* in Biolog plates and are not exchanged in iYH543. FP contains the carbon sources that allow *S. pyogenes* growth in DataS2\_iYH543\_CDM\_1.json, but they do not allow *S. pyogenes* growth in Biolog plates. FN contains the carbon sources that allow *S. pyogenes* growth in Biolog plates but do not allow growth in DataS2\_iYH543\_CDM\_1.json due to the absence of the exchange reactions in BiGG or VMH. (B) Amino acid auxotrophies in living bacterial cells. Levering et al. reported the amino acid auxotrophies of *S. pyogenes* serotype M49 (strain 591) in living bacterial cells. We compared amino acid auxotrophies in iYH543 versus experimental data using DataS3\_iYH543\_CDM\_2.json. The details of the medium for these simulations are shown in Table (Continued on next page)

**FIG 3** (Continued)

S6 (CDM2). The json formats (iYH543\_CDM\_1.json and DataS3\_iYH543\_CDM\_2.json) can be downloaded at our lab website ([http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/04/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/04/Code_Model_Files_Hirose_et_al.zip)). The results of the simulations are in Table S4 (Amino\_acid\_Auxotrophy). CDM2 components and amino acid auxotrophy (11). Accuracy: (TP + TN)/all genes.

true negative: 124). Notably, the false positive group included the amino acids L-proline and L-serine. By calculating shadow prices in FBA with CDM1 medium, we determined that iYH543 utilized these amino acids for the production of cell wall-associated biomass components (reactants in biomass reaction listed in bio1 formula). Consequently, under CDM1 conditions, living bacterial cells may be unable to shift L-proline or L-serine from sources for cell wall synthesis to alternative carbon sources.

**Validation of amino acid auxotrophies in iYH543**

To assess the amino acid auxotrophies in iYH543, we used published experimental data from serotype M49 *S. pyogenes* strain 591 (11) (medium specified in Table S6, or see DataS3\_iYH543\_CDM\_2.json at [http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirose_et_al.zip)). We compared these data with simulations of iYH543 growth in CDM2, where each of the 20 amino acids was omitted individually (Fig. 3B; Table S4). Only alanine exhibited a discrepancy (false negative) between the experimental data and iYH543. In iYH543, alanine serves as a biomass component and is required for the production of peptidoglycan polymer (PGP\_c). In a nutrient-rich medium, iYH543 can obtain L-alanine from peptides or via exchange reactions for L-alanine (EX\_ala\_\_L\_e). However, it seems that in CDM2 (peptide-free CDM) without L-alanine, *S. pyogenes* serotype M1 is unable to synthesize alanine from other metabolites.

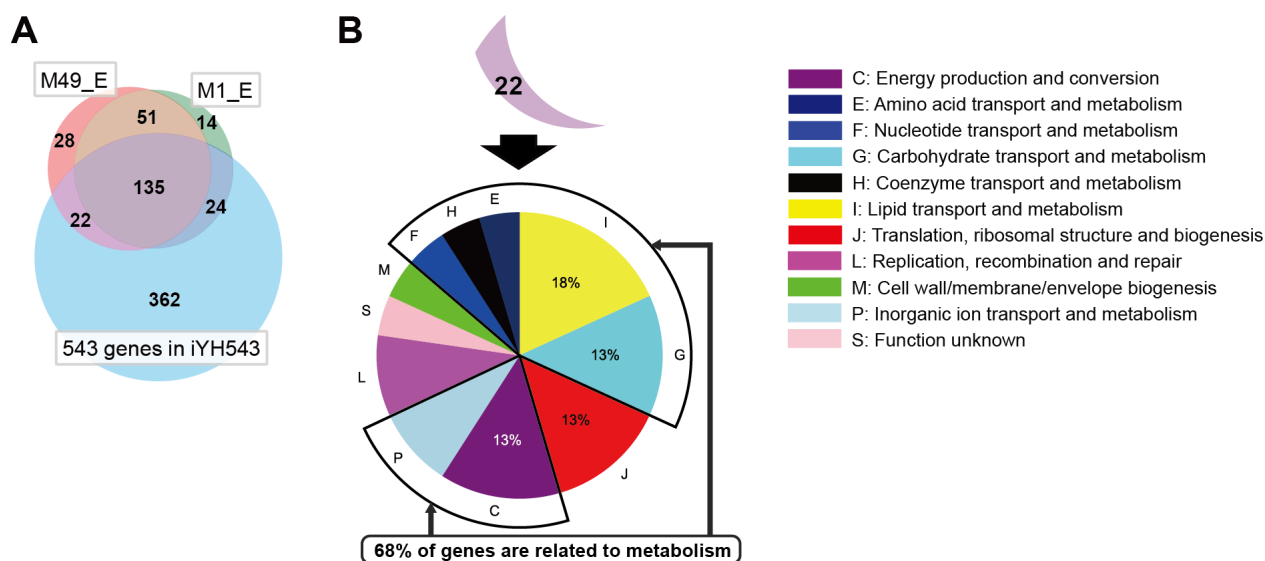
*S. pyogenes* serotype M1 and M49 share high sequence similarity, with 99.1% nucleotide identity covering 94% of the genome (serotype M49 NZ131 vs serotype M1 SF370, nBlast), which is the key reason that we used amino acid auxotrophies of serotype M49 for the curation of the serotype M1 GEM. On the other hand, Le Breton et al. reported differences in essential genes between serotype M1 and M49 when cultured in the same conditions (18). Comparing serotypes M1 and M49 among 543 genes in iYH543, 22 essential genes of serotype M49 are not essential for serotype M1. Of these 22 genes, 68% are related to metabolism (Fig. 4; Table S5). Thus, when performing curation based on the gene essentiality for GEM, it is necessary to use a serotype-specific draft. Therefore, in iYH543, we reflect amino acid auxotrophies of serotype M49 and the gene essentiality of serotype M1.

**Curation using discrepancies with experimental data**

We improved the AGORA2-derived draft GEM by incorporating reactions and modifying biomass components based on *S. pyogenes* growth in CDM1 (strain 5448) (4) and CDM2 (strain 591) (11). Table S1 provides full details of this curation process. We added reactions sourced from the BiGG or VMH databases and applied curation based on available physiological data, even in cases where the corresponding GPR was unknown. GPR rules were determined based on the *S. pyogenes* pathways suggested by BioCyc or EC number information obtained from EggNOG.

For instance, CDM2 exclusively contains Fe<sup>2+</sup> as an iron source, while the AGORA2-derived GEM lacks an enzyme to convert Fe<sup>2+</sup> (fe2 in GEM) to Fe<sup>3+</sup> (fe3 in GEM). However, both fe2 and fe3 are listed as biomass components in the AGORA2-derived draft GEM. To address this discrepancy, we introduced the Fe<sup>2+</sup> NAD oxidoreductase (FE2DH) reaction (Fig. 5A).

In *S. pyogenes* cultured in CDM2, L-ascorbate (ascb\_\_L in GEM) is consumed (11). However, the AGORA2-derived draft GEM did not include transport and metabolic reactions for L-ascorbate. Consequently, we added the necessary reactions to enable iYH543 to utilize L-ascorbate (Fig. 5B).



**FIG 4** The similarity and difference of essential genes between serotype M49 NZ131 and serotype M1 SF370. (A) Overlapping genes among M49 essential genes, M1 essential genes, and 543 genes in iYH543. M49\_E, 236 essential genes of serotype M49 NZ131. M1\_E, 224 essential genes of serotype M1 SF370. Transposon mutagenesis-based gene essentiality is referenced (18). (B) Details of 22 essential genes of M49 NZ131 in iYH543, which are not essential in M1 SF370. COGs are colored for each function. The details of genes are listed in Table S5.

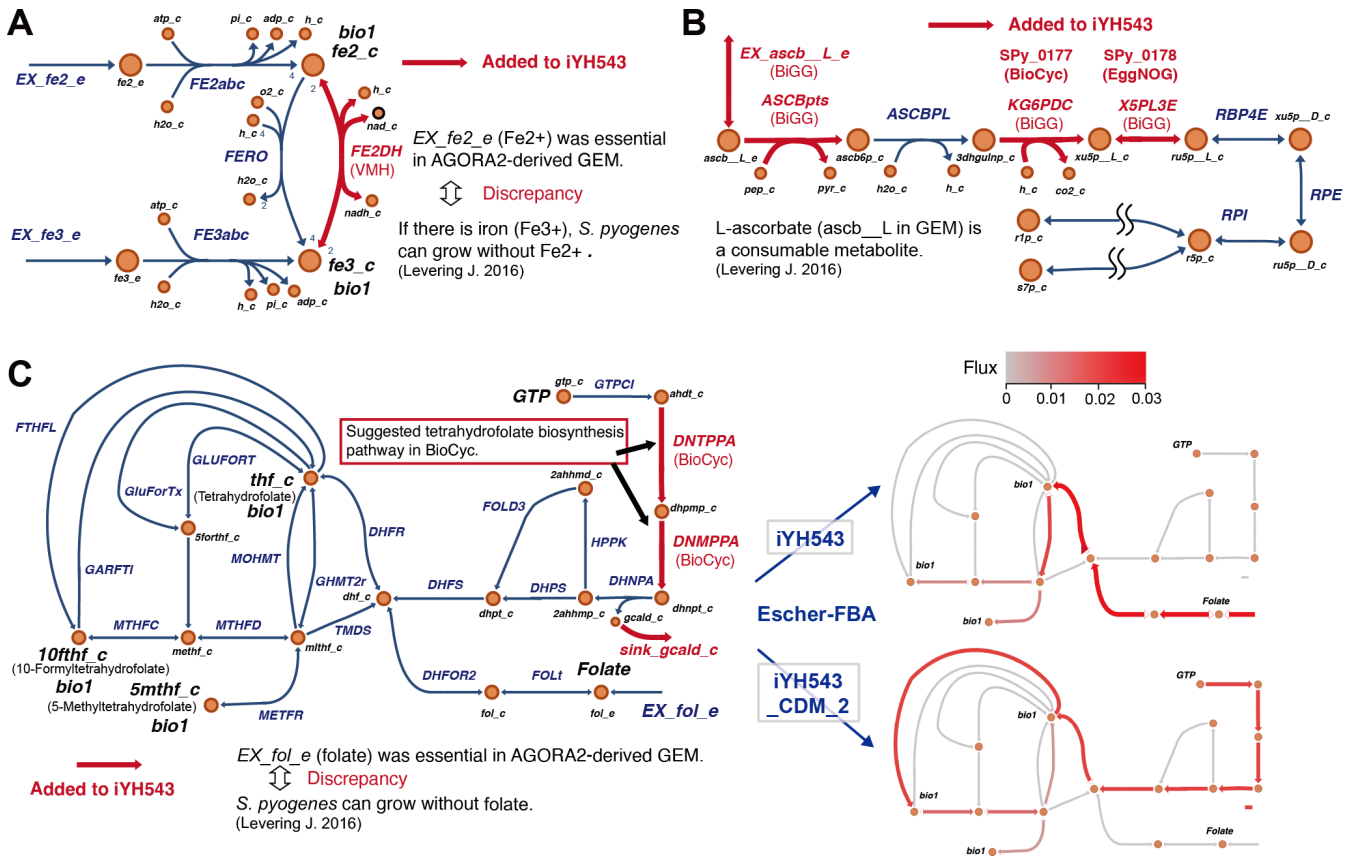
The AGORA2-derived draft GEM incorrectly predicted the requirement of folate (fol in GEM) for growth in CDM2. To rectify this, we added the DNTPPA, DNMPA, and the sink reaction for glycolaldehyde (sink\_gcald\_c) reactions to iYH543 (Fig. 5C). These reactions were identified by the BioCyc database as a part of the tetrahydrofolate biosynthesis pathway in *S. pyogenes*. Escher-FBA (22) was employed to confirm whether iYH543 and iYH543\_CDM\_2 (see json formats at [http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirose_et_al.zip)) utilized folate to generate biomass components (thf, 5,6,7,8-tetrahydrofolate; 10fthf, 10-formyltetrahydrofolate; 5mthf, 5-methyltetrahydrofolate). The calculated flux indicated that iYH543 utilized folate for biomass component generation, while iYH543\_CDM\_2 relied on the curated pathways due to the absence of folate in the growth medium.

In the AGORA2-derived draft GEM, cysteinylglycine (cgly in GEM) was used to generate glutathione (gthrd in GEM) as a biomass component. However, neither CDM1 nor CDM2 required cysteinylglycine (cgly in GEM) for growth, suggesting the presence of an alternative pathway for glutathione synthesis in *S. pyogenes*. In *Streptococcus agalactiae*, glutathione synthesis occurs through a protein homologous to *Escherichia coli* d-Ala, d-Ala ligase (23). We found structural similarity between *S. pyogenes* d-Ala, d-Ala ligase (encoded by SPy\_1421) and *E. coli* d-Ala, d-Ala ligase (DELTA-BLAST: query cover 99%, E-value 1.00E-126, Per. Ident 29.3%; Fig. S2). Consequently, we introduced the associated glutathione synthetase (GTHS) reaction to the model. Additionally, in *Saccharomyces cerevisiae*, form  $\gamma$ -glutamylcysteine ( $\gamma$ -L-glutamyl-L-cysteine, glucys in GEM) is formed by the condensation of  $\gamma$ -glutamyl phosphate (L-glutamate 5-phosphate, glu5p in GEM) and cysteine without the involvement of enzymes (24). Although the yeast *S. cerevisiae* is distant in evolutionary relationship to *S. pyogenes*,  $\gamma$ -glutamylcysteine synthesis from  $\gamma$ -glutamyl phosphate and cysteine also occurs in the closely related bacterium *S. agalactiae* (23). Therefore, we added the  $\gamma$ -glutamylcysteine synthesis (GLUCYSS) reaction to iYH543.

### Curation based on the gene essentiality

The draft GEM underwent cross-referencing with a transposon mutagenesis-based gene essentiality screen conducted in *S. pyogenes* M1 (strain 5448) for growth in a “rich”

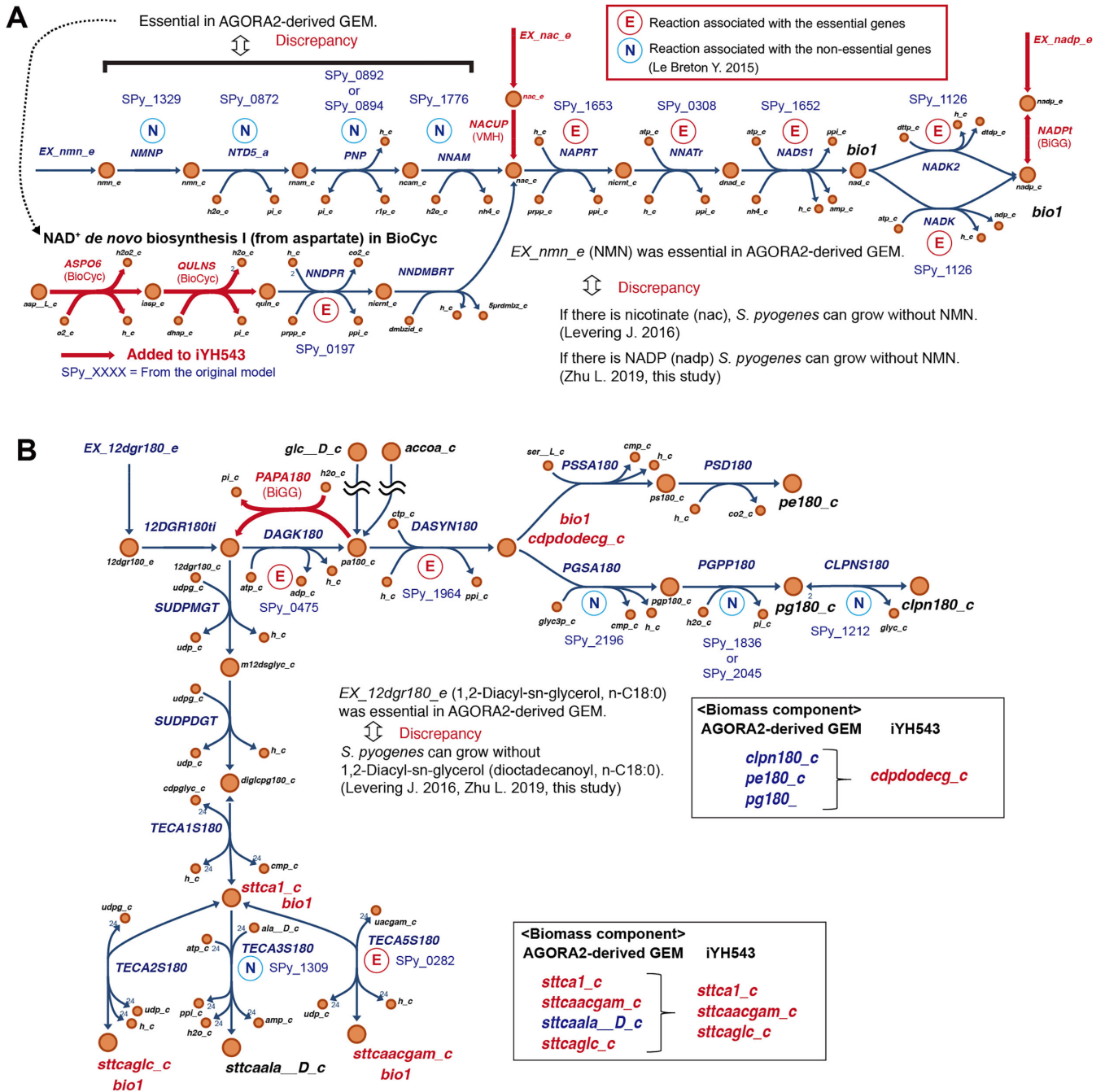




**FIG 5** Curation of iYH543 using discrepancies between experimental data and the AGORA2-derived draft GEM. (A) Rationale for adding the FE2DH reaction. Fe<sup>2+</sup> is a component of biomass in AGORA2-derived draft GEM; however, *S. pyogenes* can grow without Fe<sup>2+</sup> in CDM2 (11). FE2DH reaction was referred from the VMH database (14). (B) Curated pathways for L-ascorbate degradation. Since L-ascorbate (*ascb\_L* in GEM) is a consumable metabolite in CDM2 (11), L-ascorbate utilization reactions were added to the GEM referred from the BioCyc database (15). (C) Curated tetrahydrofolate biosynthesis pathway and its flux in iYH543 with a nutrient-rich media or CDM2. AGORA2-derived draft GEM requires folate (*fol* in GEM) in the medium, while folate is not contained in CDM2. Therefore, dihydroneopterin triphosphate pyrophosphatase (DNTPPA), dihydroneopterin monophosphate dephosphorylase (DNMPPA), and *sink\_gcaid\_c* reactions were added to the GEM referred from the BioCyc database. Escher-FBA (22) was used to confirm whether the curated pathways actually worked in iYH543. Added reactions were referred from databases, including VMH, BiGG, EggNOG, and BioCyc. CDM2 components that make *S. pyogenes* growth (11). *bio1*, a component of biomass (a reactant in the formula for biomass, *bio1* reaction); flux, estimated flux in each model.

THY medium (18). Discrepancies between the experimental data and reactions in the GEM were addressed through modifications. For example, the transport and metabolism pathway of nicotinate (*nac* in GEM) and nicotinamide adenine dinucleotide phosphate (*nadp* in GEM) were added to account for discrepancies with published experimental data (4, 11). Upon examining the gene essentiality (18) in the pathway depicted in Fig. 6A, a key point was identified between the nicotinamidase (NNAM) and nicotinate phosphoribosyltransferase (NAPRT) reactions, where the gene essentiality shifted from non-essential to essential. This observation suggests the existence of an alternative nicotinate synthesis pathway in *S. pyogenes*. To address this, we incorporated the L-aspartate oxidase (ASPO6) and quinolinate synthase (QULNS) reactions into the model by referencing NAD<sup>+</sup> *de novo* biosynthesis I (from aspartate) in *S. pyogenes* strain SF370 from BioCyc.

The mapping of essential genes and non-essential genes allowed us to make adjustments to the biomass composition. In the AGORA2-derived draft GEM, certain genes involved in the production of cardiolipin (n-C18:0) (*clpn180* in GEM) and phosphatidylglycerol (n-C18:0) (*pg180* in GEM), namely, phosphatidylglycerol phosphate phosphatase (n-C18:0) (PGPP180) and stearyl-cardiolipin synthase (CLPNS180), were



**FIG 6** Curation of iYH543 based on transposon mutagenesis-based gene essentiality. (A) The curation of NAD<sup>+</sup> de novo biosynthesis pathway. Visualization of transposon mutagenesis-based gene essentiality (18) allowed us to identify that there is no NAD<sup>+</sup> de novo biosynthesis pathway present in the AGORA2-derived draft GEM. Therefore, by tracing NAD<sup>+</sup> de novo biosynthesis I (from aspartate) of *S. pyogenes* in BioCyc (15), ASPO6 and QULNS reactions were added to the model. (B) Example for the modification of biomass components based on the transposon mutagenesis-based gene essentiality. Visualization of transposon mutagenesis-based gene essentiality (18) allowed us to distinguish between essential and non-essential biomass components of living bacteria. Biomass components associated with non-essential genes (*clpn180\_c*, *pe180\_c*, *pg180\_c*, and *sttcaala\_D\_c*) are deleted from the biomass reaction (*bio1* formula), and the coefficients are assigned to the precursor (*cdpdodecg\_c*, *sttca1\_c*) in the biomass reaction. Added reactions were referred from databases, including VMH, BiGG, and BioCyc. All reactions, metabolites, and GPR rules in iYH543 are detailed in Table S2 and can be found in the json format at our lab website ([http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/04/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/04/Code_Model_Files_Hirose_et_al.zip)). *bio1*, a component of biomass (a reactant in formula for biomass, *bio1* reaction). CDM2, components that make *S. pyogenes* growth (11); CDM1, components that make *S. pyogenes* growth (4); NMN, nicotinamide mononucleotide; NADP, nicotinamide adenine dinucleotide phosphate.

marked as essential. However, in the genome-scale mutant screen, they were found to be non-essential (18). Conversely, the genes responsible for diacylglycerol kinase (n-C18:0) (DAGK180) and CDP-diacylglycerol synthetase (n-C18:0) (DASYN180), which contribute to the production of CDP-1,2-dioctadecanoylglycerol (cdpdodecg in GEM), were essential in the mutant screen (Fig. 6B).

These findings indicate that *S. pyogenes* can survive even if one of the biomass components related to CDP-1,2-dioctadecanoylglycerol (pe180, pg180, or clpn180) is absent. However, the absence of CDP-1,2-dioctadecanoylglycerol (cdpdodecg in GEM) appears to induce cell death or cell cycle arrest according to the gene essentiality data obtained from the mutant screen (18). To maintain the stoichiometric balance in the biomass reaction (bio1) for our flux balance analysis, we consolidated clpn180, pe180, and pg180 as cdpdodecg, tripling their coefficients in the biomass reaction of iYH543 (Table S1). Similarly, the stearylphosphatidylcholine acid (linked, D-alanine substituted; sttcaala\_D in GEM) was removed; its coefficient was equally evenly distributed among the other three stearylphosphatidylcholine acids (sttca1: linked, unsubstituted; sttcaacgam: linked, N-acetyl-D-glucosamine; and sttcaagl: linked, glucose substituted) (Fig. 6B). The biomass related to isoheptadecanoylphosphatidylcholine acid (i17tca1 in GEM) and CDP-1,2-diisoheptadecanoylglycerol (cdpdihpdecg in GEM), as well as anteisoheptadecanoylphosphatidylcholine acid (ai17tca1 in GEM) and CDP-1,2-dianteisoheptadecanoylglycerol (cdpdaihpddecg in GEM), was also re-assigned by using the same approach (Fig. S3 and S4). The json formats for pathway maps (DataS4\_i17tca1\_cdpdihpddecg.json and DataS5\_ai17tca1\_cdpdaihpddecg.json) can be downloaded at our lab website ([http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirose_et_al.zip)).

In the draft GEM, the 1,2-diacyl-sn-glycerol (dioctadecanoyl, n-C18:0) exchange (EX\_12dgr180\_e) reaction was considered essential. However, live *S. pyogenes* can grow without 1,2-diacyl-sn-glycerol (dioctadecanoyl, n-C18:0) (12dgr180 in GEM) in both CDM1 (4) with glucose and CDM2 (11). In the AGORA2-derived draft GEM, the presence of 12dgr180\_e was required to produce stearylphosphatidylcholine acids (linked, unsubstituted; sttca1 in GEM)-related biomass components. To address this, we added the phosphatidate phosphatase (n-C18:0, PAPA180) reaction to the model without a GPR association. This modification allowed iYH543 to generate sttca1-related biomass components from D-glucose (glc\_D in GEM) or acetyl-CoA (accoa in GEM) (Fig. 6B).

## Drastically modified metabolic pathways from the AGORA2-derived draft GEM

Although the meso-2,6-diaminoheptanedioate exchange (Ex\_26dap\_M\_e) reaction is considered essential in the AGORA2-derived draft GEM, *S. pyogenes* can actually grow in CDM2 without meso-2,6-diaminoheptanedioate (26dap\_M in GEM) CDM2 (11). However, the genes associated with the peptidoglycan synthesis pathway from 26dap\_M\_c to PGP\_c (peptidoglycan polymer) are listed as essential genes (18), and there are no alternative pathways in iYH543 to produce PCP\_c. To address this, we added all the reactions of the meso-2,6-diaminoheptanedioate (26dap\_M in GEM) synthesis pathway from L-aspartate (asp\_L in GEM), based on the pathway found in the iYS854 model, a *Staphylococcus aureus* GEM (25) (Fig. S5A). It is worth noting that *S. aureus* and *S. pyogenes* are both Firmicutes and cause similar soft tissue infections (26, 27).

The lipid synthesis pathway underwent significant modifications (Fig. S5B). In *S. pyogenes*, exogenous biotin is necessary for growth (28). However, the AGORA2-derived draft GEM erroneously predicted that exogenous biotin is not required. Moreover, the pathway for lipid synthesis from biotin includes several essential genes in *S. pyogenes* serotype M1 (18). To rectify this inconsistency, we removed the acetyl coenzyme A carboxylase (ACCOAC) reaction and introduced the biotin--[biotin carboxyl-carrier protein] ligase (BBCCPLi) reaction, as well as the demand reactions for Apoc-Lysine (DM\_apoC\_Lys\_c) and apoprotein [acyl carrier protein] (DM\_apoACP\_c). Additionally, we adjusted the GPR for the biotin transport via ABC system (BTNabc) reaction, as studies

have shown that biotin transport in *S. pyogenes* is facilitated by *bioY* (SPy\_0207) or cobalt transporter homologs encoded by *cbiOQ* (SPy\_1787, SPy\_1788) (29, 30).

The fatty acid biosynthesis pathway was enhanced by incorporating the SPy\_1746 (*fabZ*), SPy\_1748 (*fabF*), SPy\_1749 (*fabG*), SPy\_1751 (*fabK*), and SPy\_1758 (*fabM* or *phaB*) reactions (Fig. S6) based on pathways from the BioCyc database (15) and findings in *Streptococcus pneumoniae* (31). The AGORA2-derived draft GEM lacked unsaturated fatty acids in its biomass components, despite C16:1 and C18:1 monounsaturated fatty acids being constituents of the *S. pyogenes* cell membrane (32, 33). Although the draft GEM did not contain synthesis pathways for palmitate (C16:0) or unsaturated fatty acids, it did contain glucosyl diacylglycerol (Glc-DAG) synthesis pathways for CDP-1,2-dihexadecanoylglycerol (16:0/16:0) (*cdpdodec11eg\_c*) and CDP-1,2-dioctadec-11-enoylglycerol (18:1/18:1) (*cdpdhdecg\_c*). Hence, *cdpdodec11eg\_c* and *cdpdhdecg\_c* were added to the biomass reaction (bio1 formula). Notably, Glc-DAG, which can include C16:0 and C18:1 and is the predominant glycolipid among *S. pneumoniae*, *S. pyogenes*, and *S. agalactiae* (33, 34), exhibits specific recognition by invariant T cell receptors of natural killer T cells (34).

### Hypotheses for explaining the false positives and false negatives in iYH543

The curation of iYH543 incorporated available physiological data, even in cases with unknown GPRs. However, there were remaining genes in iYH543 that showed discrepancies in predicting gene essentiality, classified as FP or FN. These discrepancies led us to formulate several hypotheses (detailed in Table S3).

For example, SPy\_1746 (*fabZ*) and SPy\_1751 (*fabK*), involved in the fatty acid biosynthesis pathway (Fig. S6, or see [DataS6\\_fatty\\_acid\\_biosynthesis.json](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirase_et_al.zip) at [http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirase\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirase_et_al.zip)), were classified as FN, meaning they were deemed essential in iYH543 but non-essential in live bacterial cells cultured in the THY medium (18). This is likely because *S. pyogenes* is able to utilize exogenous fatty acids from rich media or the host via the fatty acid kinase system, in lieu of the *de novo* production of fatty acids via *FabZ* and *FabK* (35).

FPs refer to genes reported as non-essential by iYH543 but found to be essential in living bacterial cells cultured in the THY medium. One such FP is SPy\_1315 (ABC transporter permease subunit, *glnP*), which serves as a GPR for L-glutamine transport via the ABC system (GLNabc) reaction in iYH543. Since L-glutamine can be produced from other components in iYH543 (Fig. S7, or see [DataS7\\_core\\_metabolism.json](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirase_et_al.zip) at [http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirase\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirase_et_al.zip)), the GLNabc reaction (with *glnP* in its GPR) is considered non-essential in iYH543. However, a DELTA-BLAST search reveals that *S. pyogenes* *GlnP* (encoded by SPy\_1315) shares high similarity with a part of an ABC transporter involved in lysine, arginine, and ornithine transport in *Streptococcus infantarius* subsp. (accession: [EDT46715.1](https://pubmed.ncbi.nlm.nih.gov/15844441/)) (query cover 100%, *E*-value  $1.00E-126$ , Per. ident 72.7%). The L-lysine transporter (LYSt2r) reaction is essential and lacks a GPR in iYH543, suggesting that SPy\_1315 may be responsible for L-lysine transport. In summary, iYH543 and the provided json maps in this study serve as valuable resources for investigating the unknown metabolic mechanisms of *S. pyogenes*.

### Hypothesis and knowledge obtained by substituting actual measurement values for iYH543

We previously reported that the hemolytic activities of *S. pyogenes* M1 serotype are dramatically altered by different carbon sources in CDM1 (36). *S. pyogenes* has two hemolysins, streptolysin O (SLO) and streptolysin S (SLS), which contribute to the formation of necrotizing skin lesions (37, 38). In our previous study, glucose, maltose, and dextrin caused hemolysis in SLO-dependent, SLS-dependent, and both SLO- and SLS-dependent manners, respectively (36). However, the mechanism behind this

phenomenon remained unclear. Especially, it was difficult to explain why dextrin could upregulate both hemolysins.

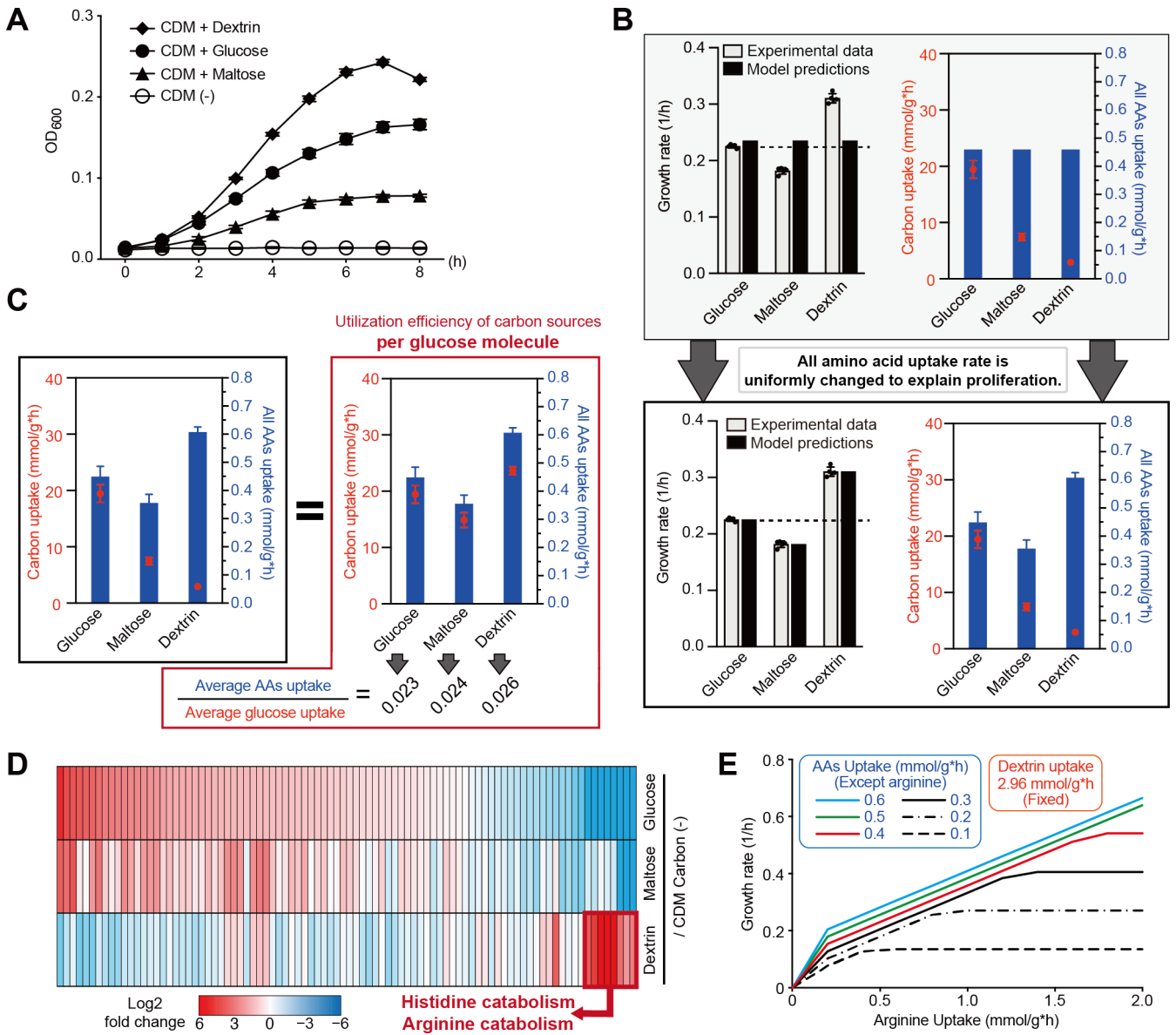
To investigate the metabolism involved in bacterial hemolytic activities, we used iYH543. First, we measured the growth parameters in CDM1 glucose (+), maltose (+), or dextrin (+) conditions (Fig. 7A). The specific growth rates on glucose, maltose, and dextrin during the exponential phase were 0.229, 0.182, and 0.311 h<sup>-1</sup>, respectively. The specific consumption rates of glucose, maltose, and dextrin were 19.40, 7.44, and 2.96 mmol g<sup>-1</sup> h<sup>-1</sup>, respectively. However, when substituting the measured specific consumption rate of glucose, maltose, and dextrin for iYH543, the predicted growth rates by FBA did not match the actual growth rate (Fig. 7B). When simulating the change of uptake efficiency (lower bounds) of medium components in iYH543, we found that the uptake efficiency of amino acids is directly related to the growth rate in FBA. In iYH543, the lower bounds for all amino acids (AAs) in the medium are set to one-tenth that of carbon sources (Table S6). Therefore, we uniformly modified the lower bounds for the uptake reaction for all AAs in iYH543 to match the actual growth and consumption rates (Fig. 7B). Maltose is a disaccharide of D-glucose. Dextrin is represented in iYH543 by eight linked polymers of D-glucose, because dextrin (starch) is consumed as  $\gamma$ -cyclodextrin (the eight glucose subunits are linked end-to-end via  $\alpha$ -1,4 linkages) in the model. When describing the utilization efficiency of all AAs and carbon sources per glucose molecule in each condition (Fig. 7C), iYH543 predicts that not only the difference in the uptake rate of carbon sources but also that of amino acids affects the growth rates.

To compare the gene expression associated with amino acid utilization in each condition, transcriptome data were extracted from our previous data set (36) (Fig. 7D; Table S7). Bacterial RNA samples from the CDM1 glucose (+), maltose (+), or dextrin (+) conditions were compared to those from the CDM1 carbon (-) condition. In the CDM1 maltose (+) condition, the expression levels of AA metabolic genes seemed to decrease overall compared to the CDM1 glucose (+) condition. In comparisons of the CDM1 dextrin (+) and glucose (+) conditions, gene expression profiles are significantly altered, with histidine and arginine utilization pathways being dramatically upregulated in the CDM1 dextrin (+) condition. As a result of FBA by using iYH543 in the CDM1 dextrin (+) condition, the growth rate is increased in proportion to the arginine uptake (flux of the exchange reaction of arginine) despite the low uptake (lower bounds of exchange reactions) of all other AAs (Fig. 7E). On the other hand, increasing histidine uptake did not affect the growth rate in FBA by using iYH543 in the CDM1 dextrin (+) condition. This suggests that enhanced arginine catabolism is associated with the high proliferation ability induced by dextrin. Additionally, we previously indicated that the upregulation of arginine catabolism is linked to the increased expression of both SLS and SLO-encoding genes (6). Together, the knowledge obtained by substituting actual measurement values for iYH543 helps us to gain unknown knowledge that connects metabolism and virulence.

## DISCUSSION

The fully curated GEM, iYH543, is the first GEM for *S. pyogenes* serotype M1 and achieved 92.6% accuracy in predicting gene essentiality. This accuracy is a significant advance upon the previously reported *S. pyogenes* serotype M49 GEM (76.6%), making iYH543 especially valuable for the study of serotype M1 strains, which are typically more pathogenic and clinically relevant. The GEM construction strategy employed here also serves as a template for curating highly accurate and consistent GEMs. The curation process provided unique insights that would have been difficult to obtain through any other means, establishing connections between previous knowledge and new discoveries in *S. pyogenes* metabolism.

In this study, we described the pathway map for cell wall synthesis and nucleotide metabolism (Fig. S8 and S9, respectively) (see DataS8\_cell\_wall.json and DataS9\_nucleotide.json at [http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirose_et_al.zip)). The predicted pathways are based on actual



**FIG 7** Utilization efficiency of amino acid is changed by the difference of supplemented carbon sources in CDM1. (A) Growth curves in CDM1 supplemented with indicated carbon sources. (B) Calculated differences in the efficiency of AA uptake (mmol/g\*h) in CDM1 glucose (+), maltose (+), or dextrin (+) conditions. Assuming that the efficiency of AA uptake is constant regardless of carbon sources, iYH543 predicts the same growth rates in all CDM1 glucose (+), maltose (+), or dextrin (+) conditions (above, gray). The efficiency of AA uptake is adjusted according to the actual values of growth rate (1/h) and carbon uptake (mmol/g\*h) (bottom). (C) Calculated differences in the efficiency of AA uptake per glucose molecule uptake in CDM1 glucose (+), maltose (+), or dextrin (+) conditions. The simulation suggested that maltose utilization tends to show inefficient amino acid uptake as compared to glucose utilization. On the other hand, the simulation suggested that glucose utilization tends to show efficient amino acid uptake as compared to glucose utilization. (D) The expression of selected genes in iYH543, associated with amino acid transport and metabolism. Bacterial RNA samples from CDM1 glucose (+), maltose (+), or dextrin (+) conditions are compared to those from CDM1 carbon (-) conditions. These data are acquired from a previous study (36). The color scale indicates enrichment (red) and depletion (blue). Genes are arranged in descending order of log2 fold change values in CDM1 glucose (+) condition. The detail of the expression levels is shown in Table S7. (E) The growth rate in proportion to the arginine uptake in iYH543 with CDM dextrin (+) condition. The actual consumption rates of dextrin 2.96 mmol g<sup>-1</sup> h<sup>-1</sup> are used as the fixed value. AA uptake, lower bounds of exchange reactions of other all AAs; arginine uptake, flux of the exchange reaction of arginine.

experimental data such as the essential gene information or the nutrient uptake information, making iYH543 more accurate than pathway maps based on genome information alone (39).

Interestingly, among the eight genes present in iYH543, but not M49 strains, is an operon for maltose or dextrin utilization (*malCDX* or *amyAB*) (Fig. S10B). Our previous research showed an *S. pyogenes* M1 strain upregulated both *nga-ifs-slo* operons and an operon for maltose or dextrin utilization at the same time. These co-upregulated gene sets (called as MalR2 iModulon) contributed to the upregulation of *nga-ifs-slo* operons in the CDM1 maltose (+) or dextrin (+) conditions (Fig. S10C) (36). The *nga-ifs-slo* operon encodes an NAD glycohydrolase (NADase), immunity factor, and pore-forming cytolytic toxin SLO that contribute to enhanced virulence of *S. pyogenes* (40, 41).

To utilize dextrin, *S. pyogenes* must break it down into glucose molecules. Even though dextrin utilization requires more enzyme effort than glucose utilization, our results showed that the actual growth efficiency was higher in the CDM1 dextrin (+) condition as compared to the CDM1 glucose (+) condition. The simulations using iYH543 suggested that differences in the utilization efficiency of amino acids may be a contributing factor. Specifically, we suggest that arginine catabolism is associated with the high proliferation ability induced by dextrin. This implies that utilizing simulations with iYH543 to interpret omics data is more effective than exploring it without guidance. Our previous study suggested that *S. pyogenes* increases the utilization of various polysaccharides, peptides, and amino acids when sensing glucose depletion in the host organism (7). This increased utilization of polysaccharides and amino acids may contribute to the expression of hemolytic toxins and the proliferation of bacteria when *S. pyogenes* is exposed to the host's nutritional immunity *in vivo*. However, the limitation of this study is that actual amino acid consumption and enzyme costs were not reflected in the simulation. Therefore, further investigation is needed to demonstrate the additional benefits of iYH543.

Mismatches between essential genes predicted by the GEM and those observed in living bacterial cells highlighted areas of uncertainty in our current understanding of *S. pyogenes* metabolism. Further research is required to address these discrepancies. Since gene functions in both iYH543 and living *S. pyogenes* cells are inferred from database annotations, it remains unclear if these predicted functions align with actual enzymatic activities. Therefore, as in any genome-scale annotation effort, continuous efforts are required for the network reconstruction process of *S. pyogenes*. Despite these limitations, iYH543 represents a useful tool for rational drug design targeting *S. pyogenes* metabolism (10) and computational screening to investigate the interplay between inhibiting virulence factor synthesis and growth (42).

## MATERIALS AND METHODS

### Strains

The draft GEM for *S. pyogenes* serotype M1 (strain SF370) (accession no. [AE004092](#)) was obtained from AGORA2 (termed AGORA2-derived GEM) created by Heinken et al. (12). In laboratory experiments, *S. pyogenes* M1 strain 5448 (accession no. [CP008776](#)), isolated from a patient with toxic shock syndrome and necrotizing fasciitis (43), was used, because a transposon mutagenesis-based gene essentiality screen was performed using this strain (18). Orthologs between 5448 and SF370 strains were determined using bidirectional BLAST hits.

*S. pyogenes* was grown at 37°C in a screw-cap centrifuge tube (BD Biosciences, San Jose, CA, USA) filled with 10 mL Todd–Hewitt broth supplemented with 0.2% yeast extract (THY) (Hardy Diagnostics, Santa Maria, CA, USA) or CDM1 (4) in an ambient atmosphere and standing cultures. To obtain cultures for experiments, overnight cultures of *S. pyogenes* were back diluted 1:50 into fresh THY broth and grown at 37°C, with growth monitored by measuring optical density at 600 nm (OD<sub>600</sub>). For the experiments using CDM1, overnight culture of *S. pyogenes* in THY broth was centrifuged, resuspended to fresh CDM, and diluted 1:20 into fresh CDM1 supplemented with indicated carbon sources (4.5 g/L). Growth kinetic experiments were performed in a Tecan Infinite Pro 200

plate reader (Tecan, Mannedorf, Switzerland). Cell concentration was measured as OD<sub>600</sub> every 30 min. CFUs were determined by plating diluted samples on THY agar.

## Biolog phenotyping experiments

Biolog's Phenotype MicroArray technology (21) was used to investigate the sole carbon source utilization profiles of *S. pyogenes* strain 5448 during growth in 190 carbon sources (PM1- and PM2A-Microplates, Biolog, Hayward, CA, USA). *S. pyogenes* was grown to the mid-log phase in THY (OD<sub>600</sub> = 0.3 ~ 0.5), then was centrifuged and resuspended in CDM1 (Table S6) (4). The suspension was adjusted to OD<sub>600</sub> = 0.09 and mixed with Redox Dye Mix H (Biolog). One hundred microliters of suspension was added into each well of the microplate and incubated at 37°C for 48 hours. The colorimetric assay was considered as positive when the average of the y-axis was 1.2 times higher than that of the negative control.

## Construction of iYH543

We constructed the iYH543 model based on the AGORA2-derived draft GEM of *S. pyogenes* M1 (strain SF370). The AGORA2-derived draft GEM of *S. pyogenes* M1 serotype was generated in the AGORA2 pipeline (12, 44). Briefly, we downloaded draft genome-scale metabolic reconstructions using KBase (45). In the platform, the genome sequence of an organism is automatically annotated, and a metabolic reconstruction is assembled based on the identified metabolic functions. The imported assemblies were annotated using RAST subsystems (46). All reactions and metabolites of these draft metabolic reconstructions were translated into the VMH (14). Gaps in the draft reconstruction are automatically filled, building a metabolic reconstruction whose condition-specific models can carry flux through a defined biomass objective function. We refined the draft reconstructions using rBioNet (47) and performed quality control and quality assurance tests.

The additional curation in this study was conducted by using the protocol made by Orth et al. (19, 48). In short, the AGORA2-derived draft GEM was intensively manually curated using the information from BiGG (13), VMH (14), BioCyc (15), KEGG (16), and EggNOG-generated EC number information (17). To extract metabolite and reaction information from BiGG, all VMH database formats of the AGORA2-derived draft GEM were converted to the BiGG database format. Where possible, reactions were verified with experimental data or external data. For gap filling, pathways were visualized using Escher (20), and the gaps were filled with reactions shown in BioCyc or other BiGG models containing iB21\_1397 (49), iJO1366 (50), iMM904 (51), iYO844 (52), iYS854 (25), and Recon3D (53). Escher-FBA was used to confirm the fluxes in GEMs (22). To identify homologous genes, BLASTp homology searches were done using the NCBI webserver with default settings or using DELTA-BLAST (54). Demand reactions are added to allow the accumulation of compounds in iYH543, and sink reactions are added to provide the network with metabolites. Some reactions were added based on literature information. A fully detailed rationale for each curation is listed in Table S1.

The proton/element balance in the reactions and the molecular weight of the biomass (which ideally should be 1 g/mmol) were checked by BiomassMW (55).

## Simulations using COBRApy

The reconstruction was converted to a mathematical model, using Constraint-Based Metabolic Modeling in Python (COBRApy) version 0.22.1 (56). COBRApy (solver: glpk) was also used for simulating with FBA and simulating the knockout of single genes and reactions.

All simulations on gene essentiality, amino acid auxotrophy, and carbon sources were done using FBA while optimizing for the biomass reaction. For all metabolites present in the media, the upper and lower bounds were set to -1,000 and 1,000 mmol/g/h,



respectively. In FBA simulations in nutrient-rich media, we used a value of  $10^{-8}$  as a growth/no growth cutoff.

In cases/conditions where the FBA model predicted a no-growth phenotype, while the experimental or external data suggested growth, we calculated the shadow price (57), allowing the prediction of specific metabolite accumulation or depletion preventing the growth of the FBA model (58).

### Measurement of a conversion coefficient between OD<sub>600</sub> and dry cell weight

Overnight culture of *S. pyogenes* in THY broth was centrifuged, resuspended to fresh CDM1, and diluted 1:20 into 1 L of fresh CDM1 supplemented with dextrin (4.5 g/L). A screw-cap bottle was filled with the mixture and incubated in an ambient atmosphere and standing cultures. During the exponential phase, 1,000 mL of bacterial culture was filtered through a membrane filter (Cat. H050A090C, pore size: 0.5  $\mu\text{m}$ ) (Advantec, Tokyo, Japan) with measurement of its OD<sub>600</sub>. The filter with cells was dried at 60°C for 16 hours in an oven and was weighted. The conversion coefficient between OD<sub>600</sub> and dry cell weight was calculated as  $0.713 \text{ g L}^{-1} \text{ OD}_{600}^{-1}$ .

### Measurement of extracellular metabolites and specific rates

To evaluate the time-course of extracellular substrate concentration, the culture supernatants were collected every hour for 8 hours by centrifugation at 12,000 rpm for 5 min. Enzymatic assay kits were used for the measurement of D-glucose (Cat. 716251), maltose (Cat. E1268), and dextrin (Cat. 1113950) (Boehringer Mannheim, Mannheim, Germany). Specific rates for growth and substrate consumptions were determined by linear regression using the culture profile during the exponential growth phase. The calculation procedure was described elsewhere (59).

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Yujiro Hirose, Conceptualization, Data curation, Formal analysis, Funding acquisition, Investigation, Methodology, Validation, Visualization, Writing – original draft, Writing – review and editing | Daniel C. Zielinski, Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Software, Supervision, Validation, Visualization | Saugat Poudel, Conceptualization, Data curation, Formal analysis, Validation, Visualization, Writing – original draft, Writing – review and editing | Kevin Rychel, Conceptualization, Data curation, Formal analysis, Validation, Visualization, Writing – review and editing | Jonathon L. Baker, Data curation, Formal analysis, Validation, Visualization, Writing – original draft, Writing – review and editing | Yoshihiro Toya, Data curation, Validation, Visualization, Writing – original draft, Writing – review and editing | Masaya Yamaguchi, Software, Writing – review and editing | Almut Heinken, Data curation, Formal analysis, Visualization, Writing – review and editing | Ines Thiele, Data curation, Formal analysis, Visualization, Writing – review and editing | Shigetada Kawabata, Supervision, Writing – original draft, Writing – review and editing.

## DATA AVAILABILITY

The new metabolic network reconstructions for *S. pyogenes* serotype M1, iYH543, are provided in a spreadsheet format in Table S2. The json formats of iYH543, iYH543\_CDM\_1, iYH543\_CDM\_2, and AGORA2\_derived\_draft\_GEM can be downloaded at our lab website ([http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code\\_Model\\_Files\\_Hirose\\_et\\_al.zip](http://www.dent.osaka-u.ac.jp/wp-content/uploads/2024/03/Code_Model_Files_Hirose_et_al.zip)). Curation notes are detailed in Table S1. The lab website also includes files detailing the development of models, simulation codes for

FBA, and calculation codes for shadow price, biomass molecular weight, and growth rate in proportion to the arginine uptake.

## ADDITIONAL FILES

The following material is available [online](#).

### Supplemental Material

**Supplemental information (mSystems00736-24-s0001.docx).** Figures S1 to S10.

**Table S1 (mSystems00736-24-s0002.xlsx).** All information of the curation.

**Table S2 (mSystems00736-24-s0003.xlsx).** All reactions, metabolites, and GPR rules in iYH543.

**Table S3 (mSystems00736-24-s0004.xlsx).** Details of essential genes in both iYH543 and a published genome-scale mutant screen.

**Table S4 (Supplementary\_Table\_4.xlsx).** Sole carbon source utilization profiles and amino acid auxotrophies in both the experimental data and iYH543.

**Table S5 (mSystems00736-24-s0006.xlsx).** Details of genes that M49 NZ131 or M1 5448 shows essential among iYH543 genes.

**Table S6 (mSystems00736-24-s0007.xlsx).** Details of nutrient-rich medium, CDM1, and CDM2 in iYH543 and actual recipe for CDM1.

**Table S7 (mSystems00736-24-s0008.xlsx).** Expression of selected genes in iYH543, associated with the amino acid transport and metabolism.

## REFERENCES

- Carapetis JR, Steer AC, Mulholland EK, Weber M. 2005. The global burden of group A streptococcal diseases. *Lancet Infect Dis* 5:685–694. [https://doi.org/10.1016/S1473-3099\(05\)70267-X](https://doi.org/10.1016/S1473-3099(05)70267-X)
- Shulman ST, Tanz RR, Kabat W, Kabat K, Cederlund E, Patel D, Li Z, Sakota V, Dale JB, Beall B, US Streptococcal Pharyngitis Surveillance Group. 2004. Group A streptococcal pharyngitis serotype surveillance in North America, 2000–2002. *Clin Infect Dis* 39:325–332. <https://doi.org/10.1086/421949>
- Steer AC, Law I, Matatolu L, Beall BW, Carapetis JR. 2009. Global *emm* type distribution of group A streptococci: systematic review and implications for vaccine development. *Lancet Infect Dis* 9:611–616. [https://doi.org/10.1016/S1473-3099\(09\)70178-1](https://doi.org/10.1016/S1473-3099(09)70178-1)
- Zhu L, Olsen RJ, Beres SB, Eraso JM, Saavedra MO, Kubiak SL, Cantu CC, Jenkins L, Charbonneau ARL, Waller AS, Musser JM. 2019. Gene fitness landscape of group A streptococcus during necrotizing myositis. *J Clin Invest* 129:887–901. <https://doi.org/10.1172/JCI124994>
- Zhu L, Olsen RJ, Beres SB, Saavedra MO, Kubiak SL, Cantu CC, Jenkins L, Waller AS, Sun Z, Palzkill T, Porter AR, DeLeo FR, Musser JM. 2020. *Streptococcus pyogenes* genes that promote pharyngitis in primates. *JCI Insight* 5:e137686. <https://doi.org/10.1172/jci.insight.137686>
- Hirose Y, Yamaguchi M, Sumitomo T, Nakata M, Hanada T, Okuzaki D, Motooka D, Mori Y, Kawasaki H, Coady A, Uchiyama S, Hiraoka M, Zurich RH, Amagai M, Nizet V, Kawabata S. 2021. *Streptococcus pyogenes* upregulates arginine catabolism to exert its pathogenesis on the skin surface. *Cell Rep* 34:108924. <https://doi.org/10.1016/j.celrep.2021.108924>
- Hirose Y, Yamaguchi M, Okuzaki D, Motooka D, Hamamoto H, Hanada T, Sumitomo T, Nakata M, Kawabata S. 2019. *Streptococcus pyogenes* transcriptome changes in the inflammatory environment of necrotizing fasciitis. *Appl Environ Microbiol* 85:e01428–19. <https://doi.org/10.1128/AEM.01428-19>
- Le Breton Y, Mclver KS. 2013. Genetic manipulation of *Streptococcus pyogenes* (the group A *Streptococcus*, GAS). *Curr Protoc Microbiol* 30:9D. <https://doi.org/10.1002/9780471729259.mc09d03s30>
- McCloskey D, Palsson BØ, Feist AM. 2013. Basic and applied uses of genome-scale metabolic network reconstructions of *Escherichia coli*. *Mol Syst Biol* 9:661. <https://doi.org/10.1038/msb.2013.18>
- Chavali AK, D'Auria KM, Hewlett EL, Pearson RD, Papin JA. 2012. A metabolic network approach for the identification and prioritization of antimicrobial drug targets. *Trends Microbiol* 20:113–123. <https://doi.org/10.1016/j.tim.2011.12.004>
- Levering J, Fiedler T, Sieg A, van Grinsven KWA, Hering S, Veith N, Olivier BG, Klett L, Hugenholtz J, Teusink B, Kreikemeyer B, Kummer U. 2016. Genome-scale reconstruction of the *Streptococcus pyogenes* M49 metabolic network reveals growth requirements and indicates potential drug targets. *J Biotechnol* 232:25–37. <https://doi.org/10.1016/j.jbiotec.2016.01.035>
- Heinken A, Hertel J, Acharya G, Ravcheev DA, Nyga M, Okpala OE, Hogan M, Magnúsdóttir S, Martinelli F, Nap B, Preciat G, Edirisinghe JN, Henry CS, Fleming RMT, Thiele I. 2023. Genome-scale metabolic reconstruction of 7,302 human microorganisms for personalized medicine. *Nat Biotechnol* 41:1320–1331. <https://doi.org/10.1038/s41587-022-01628-0>
- King ZA, Lu J, Dräger A, Miller P, Federowicz S, Lerman JA, Ebrahim A, Palsson BO, Lewis NE. 2016. BiGG models: a platform for integrating, standardizing and sharing genome-scale models. *Nucleic Acids Res* 44:D515–D522. <https://doi.org/10.1093/nar/gkv1049>
- Noronha A, Modamio J, Jarosz Y, Guerard E, Sompairac N, Preciat G, Daniëlsdóttir AD, Krecke M, Merten D, Haraldsdóttir HS, et al. 2019. The virtual metabolic human database: integrating human and gut microbiome metabolism with nutrition and disease. *Nucleic Acids Res* 47:D614–D624. <https://doi.org/10.1093/nar/gky992>
- Karp PD, Billington R, Caspi R, Fulcher CA, Latendresse M, Kothari A, Keseler IM, Krummenacker M, Midford PE, Ong Q, Ong WK, Paley SM, Subhraveti P. 2019. The BioCyc collection of microbial genomes and metabolic pathways. *Brief Bioinform* 20:1085–1093. <https://doi.org/10.1093/bib/bbx085>
- Kanehisa M, Sato Y, Kawashima M, Furumichi M, Tanabe M. 2016. KEGG as a reference resource for gene and protein annotation. *Nucleic Acids Res* 44:D457–D462. <https://doi.org/10.1093/nar/gkv1070>
- Huerta-Cepas J, Szklarczyk D, Heller D, Hernández-Plaza A, Forslund SK, Cook H, Mende DR, Letunic I, Rattei T, Jensen LJ, vonMering C, Bork P. 2019. eggNOG 5.0: a hierarchical, functionally and phylogenetically annotated orthology resource based on 5090 organisms and 2502 viruses. *Nucleic Acids Res* 47:D309–D314. <https://doi.org/10.1093/nar/gky1085>
- Le Breton Y, Belew AT, Valdes KM, Islam E, Curry P, Tettelin H, Shirtliff ME, El-Sayed NM, Mclver KS. 2015. Essential genes in the core genome of the human pathogen *Streptococcus pyogenes*. *Sci Rep* 5:9838. <https://doi.org/10.1038/srep09838>

19. Orth JD, Thiele I, Palsson BØ. 2010. What is flux balance analysis? *Nat Biotechnol* 28:245–248. <https://doi.org/10.1038/nbt.1614>
20. King ZA, Dräger A, Ebrahim A, Sonnenschein N, Lewis NE, Palsson BO. 2015. Escher: a web application for building, sharing, and embedding data-rich visualizations of biological pathways. *PLoS Comput Biol* 11:e1004321. <https://doi.org/10.1371/journal.pcbi.1004321>
21. Bochner BR, Gadzinski P, Panomitros E. 2001. Phenotype microarrays for high-throughput phenotypic testing and assay of gene function. *Genome Res* 11:1246–1255. <https://doi.org/10.1101/gr.186501>
22. Rowe E, Palsson BO, King ZA. 2018. Escher-FBA: a web application for interactive flux balance analysis. *BMC Syst Biol* 12:84. <https://doi.org/10.1186/s12918-018-0607-5>
23. Janowiak BE, Griffith OW. 2005. Glutathione synthesis in *Streptococcus agalactiae*. One protein accounts for gamma-glutamylcysteine synthetase and glutathione synthetase activities. *J Biol Chem* 280:11829–11839. <https://doi.org/10.1074/jbc.M414326200>
24. Tang L, Wang W, Zhou W, Cheng K, Yang Y, Liu M, Cheng K, Wang W. 2015. Three-pathway combination for glutathione biosynthesis in *Saccharomyces cerevisiae*. *Microb Cell Fact* 14:139. <https://doi.org/10.1186/s12934-015-0327-0>
25. Seif Y, Monk JM, Mih N, Tsunemoto H, Poudel S, Zuniga C, Broddrick J, Zengler K, Palsson BO. 2019. A computational knowledge-base elucidates the response of *Staphylococcus aureus* to different media types. *PLoS Comput Biol* 15:e1006644. <https://doi.org/10.1371/journal.pcbi.1006644>
26. Bowen AC, Tong SYC, Chatfield MD, Carapetis JR. 2014. The microbiology of impetigo in indigenous children: associations between *Streptococcus pyogenes*, *Staphylococcus aureus*, scabies, and nasal carriage. *BMC Infect Dis* 14:727. <https://doi.org/10.1186/s12879-014-0727-5>
27. Thapaliya D, O'Brien AM, Wardyn SE, Smith TC. 2015. Epidemiology of necrotizing infection caused by *Staphylococcus aureus* and *Streptococcus pyogenes* at an Iowa hospital. *J Infect Public Health* 8:634–641. <https://doi.org/10.1016/j.jiph.2015.06.003>
28. Pancholi V, Caparon M. 2016. *Streptococcus pyogenes* metabolism. In Ferretti JJ, Stevens DL, Fischetti VA (ed), *Streptococcus pyogenes: basic biology to clinical manifestations*. Oklahoma City (OK).
29. Hebbeln P, Eitinger T. 2004. Heterologous production and characterization of bacterial nickel/cobalt permeases. *FEMS Microbiol Lett* 230:129–135. [https://doi.org/10.1016/S0378-1097\(03\)00885-1](https://doi.org/10.1016/S0378-1097(03)00885-1)
30. Hebbeln P, Rodionov DA, Alfandega A, Eitinger T. 2007. Biotin uptake in prokaryotes by solute transporters with an optional ATP-binding cassette-containing module. *Proc Natl Acad Sci U S A* 104:2909–2914. <https://doi.org/10.1073/pnas.0609905104>
31. Marrakchi H, Choi KH, Rock CO. 2002. A new mechanism for anaerobic unsaturated fatty acid formation in *Streptococcus pneumoniae*. *J Biol Chem* 277:44809–44816. <https://doi.org/10.1074/jbc.M208920200>
32. Cohen M, Panos C. 1966. Membrane lipid composition of *Streptococcus pyogenes* and derived L form. *Biochemistry* 5:2385–2392. <https://doi.org/10.1021/bi00871a031>
33. Joyce LR, Guan Z, Palmer KL. 2021. *Streptococcus pneumoniae*, *S. pyogenes* and *S. agalactiae* membrane phospholipid remodeling in response to human serum. *Microbiology (Reading)* 167:001048. <https://doi.org/10.1099/mic.0.001048>
34. Kinjo Y, Illarionov P, Vela JL, Pei B, Girardi E, Li X, Li Y, Imamura M, Kaneko Y, Okawara A, et al. 2011. Invariant natural killer T cells recognize glycolipids from pathogenic Gram-positive bacteria. *Nat Immunol* 12:966–974. <https://doi.org/10.1038/ni.2096>
35. Parsons JB, Broussard TC, Bose JL, Rosch JW, Jackson P, Subramanian C, Rock CO. 2014. Identification of a two-component fatty acid kinase responsible for host fatty acid incorporation by *Staphylococcus aureus*. *Proc Natl Acad Sci U S A* 111:10532–10537. <https://doi.org/10.1073/pnas.1408797111>
36. Hirose Y, Poudel S, Sastry AV, Rychel K, Lamoureux CR, Szubin R, Zielinski DC, Lim HG, Menon ND, Bergsten H, Uchiyama S, Hanada T, Kawabata S, Palsson BO, Nizet V. 2023. Elucidation of independently modulated genes in *Streptococcus pyogenes* reveals carbon sources that control its expression of hemolytic toxins. *mSystems* 8:e0024723. <https://doi.org/10.1128/msystems.00247-23>
37. Fontaine MC, Lee JJ, Kehoe MA. 2003. Combined contributions of streptolysin O and streptolysin S to virulence of serotype M5 *Streptococcus pyogenes* strain Manfredo. *Infect Immun* 71:3857–3865. <https://doi.org/10.1128/IAI.71.7.3857-3865.2003>
38. Datta V, Myskowski SM, Kwinn LA, Chiem DN, Varki N, Kansal RG, Kotb M, Nizet V. 2005. Mutational analysis of the group A streptococcal operon encoding streptolysin S and its virulence role in invasive infection. *Mol Microbiol* 56:681–695. <https://doi.org/10.1111/j.1365-2958.2005.04583.x>
39. Ferretti JJ, Stevens DL, Fischetti VA, eds. *Streptococcus pyogenes: basic biology to clinical manifestations*. 2nd ed. Oklahoma City (OK).
40. Zhu L, Olsen RJ, Nasser W, Beres SB, Vuopio J, Kristinsson KG, Gottfredsson M, Porter AR, DeLeo FR, Musser JM. 2015. A molecular trigger for intercontinental epidemics of group A *Streptococcus*. *J Clin Invest* 125:3545–3559. <https://doi.org/10.1172/JCI82478>
41. Zhu L, Olsen RJ, Nasser W, de la Riva Morales I, Musser JM. 2015. Trading capsule for increased cytotoxin production: contribution to virulence of a newly emerged clade of emm89 *Streptococcus pyogenes*. *mBio* 6:e01378-15. <https://doi.org/10.1128/mBio.01378-15>
42. Bartell JA, Blazier AS, Yen P, Thøgersen JC, Jelsbak L, Goldberg JB, Papin JA. 2017. Reconstruction of the metabolic network of *Pseudomonas aeruginosa* to interrogate virulence factor synthesis. *Nat Commun* 8:14631. <https://doi.org/10.1038/ncomms14631>
43. Kansal RG, McGeer A, Low DE, Norrby-Teglund A, Kotb M. 2000. Inverse relation between disease severity and expression of the streptococcal cysteine protease, SpeB, among clonal M1T1 isolates recovered from invasive group A streptococcal infection cases. *Infect Immun* 68:6362–6369. <https://doi.org/10.1128/IAI.68.11.6362-6369.2000>
44. Magnúsdóttir S, Heinken A, Kutt L, Ravcheev DA, Bauer E, Noronha A, Greenhalgh K, Jäger C, Baginska J, Wilmes P, Fleming RMT, Thiele I. 2017. Generation of genome-scale metabolic reconstructions for 773 members of the human gut microbiota. *Nat Biotechnol* 35:81–89. <https://doi.org/10.1038/nbt.3703>
45. Arkin AP, Cottingham RW, Henry CS, Harris NL, Stevens RL, Maslov S, Dehal P, Ware D, Perez F, Canon S, et al. 2018. KBase: the United States department of energy systems biology knowledgebase. *Nat Biotechnol* 36:566–569. <https://doi.org/10.1038/nbt.4163>
46. Aziz RK, Bartels D, Best AA, DeJongh M, Disz T, Edwards RA, Formsma K, Gerdes S, Glass EM, Kubal M, et al. 2008. The RAST Server: rapid annotations using subsystems technology. *BMC Genomics* 9:75. <https://doi.org/10.1186/1471-2164-9-75>
47. Thorleifsson SG, Thiele I. 2011. rBioNet: a COBRA toolbox extension for reconstructing high-quality biochemical networks. *Bioinformatics* 27:2009–2010. <https://doi.org/10.1093/bioinformatics/btr308>
48. Thiele I, Palsson BØ. 2010. A protocol for generating a high-quality genome-scale metabolic reconstruction. *Nat Protoc* 5:93–121. <https://doi.org/10.1038/nprot.2009.203>
49. Monk JM, Charusanti P, Aziz RK, Lerman JA, Premyodhin N, Orth JD, Feist AM, Palsson BØ. 2013. Genome-scale metabolic reconstructions of multiple *Escherichia coli* strains highlight strain-specific adaptations to nutritional environments. *Proc Natl Acad Sci U S A* 110:20338–20343. <https://doi.org/10.1073/pnas.1307797110>
50. Orth JD, Conrad TM, Na J, Lerman JA, Nam H, Feist AM, Palsson BØ. 2011. A comprehensive genome-scale reconstruction of *Escherichia coli* metabolism—2011. *Mol Syst Biol* 7:535. <https://doi.org/10.1038/msb.2011.65>
51. Mo ML, Palsson BO, Herrgård MJ. 2009. Connecting extracellular metabolomic measurements to intracellular flux states in yeast. *BMC Syst Biol* 3:37. <https://doi.org/10.1186/1752-0509-3-37>
52. Oh YK, Palsson BO, Park SM, Schilling CH, Mahadevan R. 2007. Genome-scale reconstruction of metabolic network in *Bacillus subtilis* based on high-throughput phenotyping and gene essentiality data. *J Biol Chem* 282:28791–28799. <https://doi.org/10.1074/jbc.M703759200>
53. Brunk E, Sahoo S, Zielinski DC, Altunkaya A, Dräger A, Mih N, Gatto F, Nilsson A, Preciat Gonzalez GA, Aurich MK, Prlić A, Sastry A, Danielsdóttir AD, Heinken A, Noronha A, Rose PW, Burley SK, Fleming RMT, Nielsen J, Thiele I, Palsson BO. 2018. Recon3D enables a three-dimensional view of gene variation in human metabolism. *Nat Biotechnol* 36:272–281. <https://doi.org/10.1038/nbt.4072>
54. Boratyn GM, Camacho C, Cooper PS, Coulouris G, Fong A, Ma N, Madden TL, Matten WT, McGinnis SD, Merezchuk Y, Raytselis Y, Sayers EW, Tao T, Ye J, Zaretskaya I. 2013. BLAST: a more efficient report with usability

- improvements. *Nucleic Acids Res* 41:W29–W33. <https://doi.org/10.1093/nar/gkt282>
55. Chan SHJ, Cai J, Wang L, Simons-Senftle MN, Maranas CD. 2017. Standardizing biomass reactions and ensuring complete mass balance in genome-scale metabolic models. *Bioinformatics* 33:3603–3609. <https://doi.org/10.1093/bioinformatics/btx453>
56. Ebrahim A, Lerman JA, Palsson BO, Hyduke DR. 2013. COBRApy: COstraints-based reconstruction and analysis for Python. *BMC Syst Biol* 7:74. <https://doi.org/10.1186/1752-0509-7-74>
57. Reznik E, Mehta P, Segrè D. 2013. Flux imbalance analysis and the sensitivity of cellular growth to changes in metabolite pools. *PLoS Comput Biol* 9:e1003195. <https://doi.org/10.1371/journal.pcbi.1003195>
58. Töpfer N, Scossa F, Fernie A, Nikoloski Z. 2014. Variability of metabolite levels is linked to differential metabolic pathways in *Arabidopsis*'s responses to abiotic stresses. *PLoS Comput Biol* 10:e1003656. <https://doi.org/10.1371/journal.pcbi.1003656>
59. Matsuda F, Toya Y, Shimizu H. 2017. Learning from quantitative data to understand central carbon metabolism. *Biotechnol Adv* 35:971–980. <https://doi.org/10.1016/j.biotechadv.2017.09.006>