Derailing the aspartate pathway of *Mycobacterium tuberculosis* to eradicate persistent infection

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A major constraint for developing new anti-tuberculosis drugs is the limited number of validated targets that allow eradication of persistent infections. Here, we uncover a vulnerable component of *Mycobacterium tuberculosis* (*Mtb*) persistence metabolism, the aspartate pathway. Rapid death of threonine and homoserine auxotrophs points to a distinct susceptibility of *Mtb* to inhibition of this pathway. Combinatorial metabolomic and transcriptomic analysis reveals that inability to produce threonine leads to deregulation of aspartate kinase, causing flux imbalance and lysine and DAP accumulation. *Mtb*’s adaptive response to this metabolic stress involves a relief valve-like mechanism combining lysine export and catabolism via aminoadipate. We present evidence that inhibition of the aspartate pathway at different branch-point enzymes leads to clearance of chronic infections. Together these findings demonstrate that the aspartate pathway in *Mtb* relies on a combination of metabolic control mechanisms, is required for persistence, and represents a target space for anti-tuberculosis drug development.
**Mycobacterium tuberculosis** (*Mtb*), the causative agent of tuberculosis (TB), is the deadliest pathogen worldwide, with 1.8 million deaths in 2016, and is the source of severe societal and economic burdens. *Mycobacterium tuberculosis* is able to establish a devastating chronic infection in humans, where it can withstand host adaptive immunity and survive in granulomas, a phase commonly referred to as persistence. Many drugs are ineffective during persistence, most likely due to reduced penetration into the devascularized granuloma environment and enrichment of phenotypically drug-tolerant cells. As a result, treatment regimens require a lengthy 6-months of combination therapy to clear the infection, which greatly increases the risk of antibiotic resistance acquisition.

Recent efforts to target *Mtb* during persistence have focused on inhibiting previously ignored metabolic pathways. Several studies have demonstrated that central carbon metabolism (CCM) is required during persistence, and several targets have been identified as essential in both acute and chronic infections in mice. Importantly, biosynthetic pathways that utilize TCA cycle intermediates, such as amino acid biosynthesis, are also required for *Mtb* survival. However, none of the amino acid biosynthetic pathways of *Mtb* have yet been shown to be essential during chronic infection and it is largely unknown if such building blocks can be scavenged from the host during persistence.

We recently reported that methionine biosynthesis in *Mtb* is required to establish infections in immunocompetent and immunocompromised mice. Methionine, along with isoleucine, threonine, and lysine make up the family of essential amino acids synthesized from aspartate via the aspartate pathway. This pathway is also involved in the biosynthesis of the peptidoglycan building block, dianimonopimelate (DAP), and the cofactor, S-adenosyl methionine (SAM); both of which are required for survival in mycobacteria. Because of its branched nature and pivotal role in essential cellular processes (cell wall biosynthesis, translation, and one-carbon metabolism), metabolic regulation of the aspartate pathway is likely important to meet precursor requirements during growth and persistence in the host.

The apparent essentiality of the numerous metabolic products and absence of the aspartate pathway in humans and animals led us to investigate the effects of inhibition of the aspartate pathway in *Mtb* and its requirement during persistence in the host. We hypothesized that disruption of the aspartate pathway leads to a metabolic imbalance and eventual collapse of this essential biosynthetic network. To investigate this, we employed a combinatorial transcriptomic and metabolomics approach. mRNA transcript levels of the *ΔthrA* and *ΔthrB* mutant strains were measured by RNA-seq and microarray analysis, respectively, in time-course experiments in growth media lacking the amino acids required for survival.

**Threonine auxotrophy leads to accumulation of lysine and catabolites.** We next sought to elucidate the cellular consequences to inhibition at the threonine- homoserine-methionine branch-point. We employed a combinatorial transcriptomic and metabolomics approach. mRNA transcript levels of the *ΔthrA* and *ΔthrB* mutant strains were measured by RNA-seq and microarray analysis, respectively, in time-course experiments in growth media lacking the amino acids required for survival. The *ΔthrA* and *ΔthrB* strains have an expression profile similar to that previously observed in methionine starvation (Supplementary Data 2) and general nutrient starvation (Supplementary Data 2), as evidenced by the downregulation of lipid biosynthesis and respiration-related genes, and upregulation of pyruvate and succinate metabolism (Fig. 2). However, there were also significant differences in gene expression that indicated killing by threonine and homoserine auxotrophy occurs via a mechanism distinct from that mediated by methionine or general starvation. Threonine and homoserine auxotrophy each led to a particularly strong and early response in the absence of threonine supplementation (Fig. 1c). Death in this mutant could not be rescued by addition of isoleucine. Deletion of *thrA* was bactericidal and could be rescued by either the addition of homoserine or by a combination of methionine and threonine (Fig. 1d). Interestingly, the addition of either methionine or threonine alone in *ΔthrA* led to a statistically significant slowdown of cell killing (Fig. 1e).

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To investigate if these substantial transcriptional changes also translate into metabolic changes, we characterized the intracellular and extracellular metabolites by ultra-performance liquid chromatography-mass spectrometry (UPLC-MS). In accordance with the upregulation of genes for lysine disposal, the *ΔthrA* mutant in media lacking homoserine displayed a significant increase (up to 100-fold) of all identifiable metabolites in the lysine biosynthesis pathway (Fig. 3a). Diaminopimelate, lysine, α-aminoadipate, and N-acetylysine were all found in at least 10-fold
Fig. 1 Inactivation of homoserine dehydrogenase and homoserine kinase is bactericidal. a The aspartate family amino acid biosynthesis pathway (aspartate pathway) in M. tuberculosis including the lysine degradation (aminoadipate) pathway. Ask: aspartate kinase, Asd: aspartate semialdehyde dehydrogenase, ThrA: threonine dehydrogenase, MetX: homoserine transacetylase, ThrB: homoserine kinase, ThrC: threonine synthase, MetK: S-adenosyl methionine synthase, DapE: Succinyl-diaminopimelate desuccinylase, LysE: lysine exporter, Pcd: aminoadipate semialdehyde dehydrogenase, Lat: lysine transacetylase. Two mutants were constructed, one threonine auxotroph \textit{Mtb} \textDelta \text{thrB} and \textit{Mtb} \textDelta \text{thrA}, a strain auxotrophic for both, threonine and methionine. b Deletion of \textit{thrB} can be chemically complemented by threonine (Thr) (blue inverted triangles) but not isoleucine (Ile). c Deletion of \textit{thrB} is bactericidal and mirrors the kill curve of \textit{Mtb} \textDelta \text{thrA} supplemented with methionine (Met) (open rhombus). d Deletion of \textit{thrA} can be chemically complemented by homoserine (Hse) (black circles) or a combination of threonine and methionine (blue triangles), but not by threonine, isoleucine (orange diamonds) and methionine alone (purple rhombus). e Deletion of \textit{thrA} is bactericidal. Cell death of \textit{Mtb} \textDelta \text{thrA} can be slowed by addition of threonine or methionine but not isoleucine, indicating that inhibition of each individual branch (threonine and methionine) contributes to killing. All values are the average of three biological replicates ($n=3$) ± s.d. (error bars show standard deviation) and are representative of a minimum of two independent experiments. *$p$-value $< 0.01$ in Student’s $t$-test. Source data are provided as a Source Data file.
greater abundance after 24 h of starvation of the ΔthrA mutant (Fig. 3a). Furthermore, lysine, but not other pathway intermediates, was found to accumulate extracellularly (Fig. 3b), suggesting increased activity of a lysine exporter. Over a period of 4 days the extracellular concentration of lysine increased to 160 µM after which it stayed constant (Supplementary Fig. 2). Analysis of the metabolomic response of threonine auxotrophy alone (Mtb ΔthrB) showed a similar pattern of intracellular metabolite abundances (Fig. 4), but in addition, Mtb ΔthrB also accumulated homoserine (Supplementary Fig. 3), the adduct of the thrB encoded threonine kinase (Fig. 1a), which nicely confirms the metabolic block at ThrB. In line with this homoserine accumulation, we observed increased expression of metX (Fig. 2b). Lysine export was also observed in the ΔthrB strain, albeit at a reduced rate compared to ΔthrA (Supplementary Fig. 4), which fits with the fact that carbon rerouting in ΔthrB occurs not only toward lysine but also homoserine. Homoserine was proposed to be toxic to M. tuberculosis20, and we have confirmed its toxicity for the MTB complex strain M. bovis BCG (Supplementary Fig. 5). We observed an adverse role of homoserine but not threonine on growth of M. bovis BCG at concentrations above 50 µg/ml (Supplementary Fig. 5), which is in excellent agreement with the work published earlier21,22.

Threonine is the main feedback regulator of Mtb’s aspartate pathway. Next, we wanted to understand at a regulatory level, how inhibition of ThrA or ThrB could lead to increased lysine production. Aspartate kinase (AK) encoded by ask (Rv3709c) catalyzes the first committed step in the aspartate pathway, converting aspartate to aspartyl phosphate (Fig. 1). In Escherichia coli, there are three AK isoforms that are separately sensitive to feedback regulation by threonine, lysine/leucine, or methionine, respectively23,24. The actinomycete Corynebacterium glutamicum regulates the aspartate pathway via a single AK that is feedback inhibited by lysine and threonine, and transcriptional repression of homoserine dehydrogenase (thrA) by methionine25,26. In M. tuberculosis, purified AK has been shown
to be sensitive to threonine but not lysine, yet no other amino acids (e.g., methionine or homoserine) were tested nor was it investigated if these results translated to whole-cell physiology. Methionine and SAM biosynthesis in Mtb are likely regulated at the transcriptional level. Consequently, addition of threonine should reverse lysine accumulation in Mtb ΔthrA. Indeed, threonine was capable of decreasing intracellular lysine accumulation, causing a concomitant accumulation of aspartate (Fig. 3a). Furthermore, addition of methionine as a control caused no change in lysine pathway metabolites (Fig. 4), indicating that the loss of threonine is the major driver for over-production of lysine. This argues that L-threonine inhibition of AK is likely a primary metabolic control mechanism of the aspartate pathway.

A lysine exporter responds to disruptions in lysine homeostasis. We next investigated the unexpected export of lysine as an alternative mechanism to cope with lysine overload in the cell. Our transcriptomic analysis of the ΔthrA and ΔthrB mutants showed a strong upregulation of a putative lysine exporter Rv1986 and arginine biosynthesis genes (Fig. 2a, b). Moreover, metabolomic data demonstrates that intracellular accumulation of Lys leads to export of both lysine and arginine (Arg) (Fig. 5a) but not other amino acids or metabolites (Supplementary Fig. 6). It has been shown in C. glutamicum that lysine accumulation is toxic, and that export of both Lys and Arg is carried out by the exporter LysE. A bioinformatic analysis showed that Mtb encodes several proteins with homology to LysE of C. glutamicum, yet only Rv1986 is located adjacent to its canonical transcriptional regulator LysG (Rv1985c) (Fig. 5b)9. Hence, we hypothesized that Rv1986 serves the purpose of an overflow mechanism to prevent intracellular lysine overload. To test the role of LysE in amino acid export of Mtb, we measured the ability of Mtb to export toxic amino acid analogs in the presence and absence of the LysE exporter. This is a commonly used strategy to characterize amino acid exporters in whole cells. We deleted the genes lysE (Rv1986) and lysG (Rv1985c) in Mtb H37Rv, resulting...
in mutants ΔlysE and ΔlysG. Using these strains, we determined the minimum inhibitory concentration (MIC) of toxic amino acid analogs of lysine (2,6-diaminohex-4-ynoic acid), arginine (canavanine) and histidine (β-2-Thiazolyl-DL-alanine) (Fig. 5c, d, e). These compounds (Supplementary Fig. 1) have previously been shown to be toxic analogs in other bacteria and act as substrates for their cognate amino acid exporters32–34. Toxicity arises from their aberrant incorporation into proteins during translation, and deletion of their analogs’ exporters should increase susceptibility (Fig. 5b). Indeed, we found that deletion of both lysE and lysG resulted in a greater than 10-fold reduction in the MIC of canavanine and 2,6-diaminohex-4-ynoic acid (Fig. 5c, d), but there was no change in the MIC of the histidine analog control (Fig. 5e). The lysE phenotype was rescued when genetically complemented (Fig. 5c, d) or when the cognate amino acid (e.g., arginine) was added as a competitive substrate (Supplementary Fig. 7). The results demonstrate that LysE exports lysine and arginine as predicted by its homology and similarity (32% identity determined by Basic Local Alignment Search Tool (BLAST)) to the C. glutamicum LysE permease and that LysG is the activator of LysE under the conditions tested (Fig. 5). We also found evidence that lysine disposal is relevant in Mtb during growth in standard growth medium under non-auxotrophic conditions. During the transition from exponential to stationary phase, lysine and aminoacidate are elevated in wild-type Mtb H37Rv (Supplementary Fig. 8) and the lysE deletion mutant shows a significant growth retardation under these conditions (Supplementary Fig. 9).

Taken together, these data indicate that Mtb utilizes lysine export and degradation pathways as a metabolic relief valve to prevent disruption of aspartate pathway homeostasis and maintain essential precursor production. Such a metabolic mechanism to maintain precursor availability for growth and survival is unprecedented in M. tuberculosis. It is tempting to speculate that prevention of lysine export, could synergize with inhibition of threonine production, and could be exploited to enhance bactericidal activity. To test this notion, we attempted to construct a double deletion strain ΔthrAΔlysE. However, multiple attempts were unsuccessful pointing to a synthetic lethal interaction between these two pathways.

Aspartate pathway enzymes are required during chronic infections. Given the potent bactericidal phenotypes observed in several auxotrophs in vitro, we next evaluated the need for aspartate family amino acid biosynthesis during host infection. We first tested the ability of the threonine/methionine auxotroph Mtb ΔthrA to establish an infection in immunocompetent (C57BL/6) and immunocompromised (SCID) mice via aerosol infection. Similar to ΔmetX13, we found that ΔthrA mutant was incapable of establishing an infection in either C57BL/6 (Supplementary Fig. 10a, b) or SCID mice (Supplementary Fig. 10c, d). To test whether threonine auxotrophy alone prevents the establishment of infection, we also tested Mtb ΔthrB in C57BL/6 and found this strain to be avirulent too (Supplementary Fig. 11). However, as Mtb can maintain a persistent infection it is also necessary to test target vulnerability during an established, chronic infection35. As specific target inhibitors are not yet available, a genetic approach was employed. We constructed conditional knockdowns (cKD) of ThrA (ΔthrA-DUC) and MetX (ΔmetX-DUC), using the tetracycline-induced Dual-Control (DUC) system35. Both constructs are transcriptionally regulated by an anhydrotetracycline (ata) -inducible transcriptional repressor (T38), and are simultaneously repressed by atc-induced proteolytic degradation (DUC)36,37. The knockdown efficiency was validated in vitro by measuring the effect of increasing
concentrations of atc on growth of the recombinant strains in culture broth media (Fig. 6a, e). Growth of the DUC strains in medium without atc was indistinguishable from the parental strain, demonstrating genetic complementation to WT levels (Supplementary Fig. 12a). While the parental H37Rv strain showed no sensitivity to atc (Supplementary Fig. 12b), growth was inhibited in both knockdown strains (Fig. 6a, b). Next we tested the vulnerability of Mtb to inhibition of these enzymes during different stages of infection in C57Bl/6 mice. Cohorts of each infection group were started on doxycycline at 0, 1, 3, or 6 weeks after infection, which allowed us to study the in vivo essentiality of these enzymes during both the acute and chronic phases of infection. The growth of the knockdown strains was monitored over the course of the experiment (Fig. 6c–h). Gross lung pathology and Hematoxylin and Eosin (H&E) staining of lungs at day 189 are shown in (c, g). X-axis crosses Y-axis at limit of detection. Error bars show standard deviations from n = 4 mice. **p-value < 0.01, *p-value < 0.05 in Student’s t-test. Source data are provided as a Source Data file.

Fig. 6 Aspartate pathway enzymes are essential during acute and chronic mouse infections. Conditional knockdowns were constructed of two enzymes of the aspartate pathway: homoserine dehydrogenase (ΔthrA-DUC) (a–d) and homoserine transacetylase (ΔmetX-DUC) (e–f). a, e Effect of gene knockdown/protein degradation on growth was tested in vitro in unsupplemented 7H9 OADC with different concentrations of anhydrotetracycline (atc). b, f Mice were aerosol infected with either ΔthrA-DUC3 or Δmet-DUC5 to deliver ~100 CFU/lungs. Groups of mice were switched to doxycycline containing chow (2000 ppm) at day 0 (red square), day 7 (green diamond), day 21 (purple triangle) and day 42 (blue inverted triangle) to induce knockdown. One group (black) was left untreated throughout the whole experiment. At given time points, 4 mice per group were sacrificed and bacterial burden of lungs (b, f) and spleens (d, h) determined by CFU counts of homogenized, serially diluted tissue. Gross lung pathology and Hematoxylin and Eosin (H&E) staining of lungs at day 189 are shown under (c, g).

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stages of infection (Fig. 6b, f). For both cKD strains, administration of doxycycline immediately after infection resulted in a total bacterial clearance (below detection limit) within one week after treatment initiation. Initiation of doxycycline treatment at the later time points resulted in clearance (below detection limit) of the infection from both lungs and spleens. By the end of the experiments, bacilli could no longer be cultured in the ΔmetX-DUC (Fig. 6f) infected mice, and only ΔthrA-DUC infected mice that began doxycycline treatment at week 6 still showed detectable levels of bacteria (Fig. 6b). The total bacterial burden had been reduced by 4-logs in the lungs and >1-log in the spleens (Fig. 6d, h, respectively). Inspection of gross pathology and H&E staining of lungs harvested 189 days after infection (indicated by arrow P and dotted line in Fig. 6b, f) reveal typical tuberculous lesions in untreated mice with high bacterial burden and absence of lesions and inflammatory foci in mice that were switched to doxy chow (Fig. 6c, g). These experiments demonstrate the essentiality of the aspartate pathway in both acute and chronic Mtb infections in animals and show that Mtb is unable to scavenge nutrients from the host to complement these auxotrophies in vivo.

Discussion
Significant efforts to identify new drug targets in Mtb have focused on the metabolism of essential nutrients. Mtb is an amino acid prototroph that has developed nominal machinery to obtain nutrients from the host, unlike many other intracellular parasites. Consequently, Mtb appears to be particularly susceptible to inhibition of amino acid biosynthesis. Moreover, the biosynthetic enzymes for essential amino acids are largely absent in humans and mammals, making amino acid biosynthetic pathways in Mtb attractive drug targets.

To date, in vitro transposon sequencing (Tn-seq) screens have been used to predict gene essentiality in Mtb and have guided target selection in drug discovery. The usefulness of Tn-seq screens for the TB research field is undisputed, yet their limitations for drug discovery need consideration. A recent study highlighted the impact of growth medium on inferred gene essentiality and illustrated how in vitro Tn-seq screens can be limited predictors of in vivo essentiality and cannot differentiate between lethality, bacteriostasis, slow growth phenotypes or conditional essentiality, all information that is crucial for assessing druggability. Since M. tuberculosis is an intracellular pathogen, essentiality and druggability of targets have to be tested during chronic infection in order to prevent false targets being taken forward for drug development.

We here focused on the aspartate pathway, because it produces multiple essential building blocks for vital cellular functions in Mtb, such as cell wall biosynthesis, translation, and one-carbon metabolism. The rapid cell death phenotypes observed for auxotrophs of threonine (this study), homoserine (this study) and methionine, points to a distinct susceptibility of Mtb in inhibition of this pathway. The mechanism of action for the lethality of threonine and homoserine auxotrophy appear to be multilayered. Cell death kinetics due to the loss of proteinogenic amino acids is likely accelerated by the accumulation of intermediates of lysine biosynthesis and homoserine (only in thrB mutant).

By studying the metabolic response to aspartate pathway inhibition, we discovered that Mtb employs a combination of feedback control, overflow metabolism and catabolic action to prevent flux imbalances and ensure balanced precursor production. We show here corroborating evidence that the committed step of the aspartate pathway in Mtb, aspartate kinase, is controlled by allosteric feedback of threonine, but not by lysine or methionine. As Mtb lacks the ubiquitously used lysine-AK feedback loop, it must employ other regulatory mechanism to control production of proteinogenic lysine and the cell-wall building block diaminopimelate. By studying the adaptive metabolism of homoserine and threonine auxotrophic strains, we discovered two compensatory mechanisms that help Mtb to control the flux through this pathway branch: lysine degradation and export (Fig. 7b).

The lysine catabolic pathway catalyzed by Lat and Pcd is a unique feature of mycobacteria (Fig. 7). In a survey of 1362 bacteria, the aminoadipate branch was found to exist in only 36 microorganisms, predominately species of mycobacteria. This catabolic pathway seems to be a remnant of the cephapycin C biosynthetic pathway found in Streptomyces, yet mycobacteria have only retained the first two reactions (mycobacteria do not produce cephapycin C). Another unique feature of this pathway in mycobacteria is the localization of lat in an operon with the alternative sigma factor sigF (Rv3286c) and lrp (Rv3291c), the putative transcriptional regulator of lat. SigF is important under various stress conditions, including osmotic stress, pH, heat-shock, oxidative stress, and during infection. Expression of lrp and lat genes is enriched in a persistence model, as part of the stringent response, and whenever Mtb transitions from replicating to non-replicating conditions.

Fig. 7 Schematic of aspartate pathway regulation. a Wildtype levels of threonine lead to proper feedback regulation of Ask, and subsequent wildtype levels of lysine and its degradation (Lat/Pcd) and export (LysE) pathway enzymes and metabolites. b Disruption of threonine biosynthesis causes derepression of Ask leading to an accumulation of Lysine, which in turn activates the lysine degradation and export pathways. Nomenclature: Metabolites in rounded boxes: Asp: aspartate, Asp-P: aspartate-phosphate, ASA: aspartate-semialdehyde, Hse: Homoserine, Hse-P: homoserine-phosphate, Thr: Threonine, HSe-O:Ac: O-acetyl-homoserine, Met: methionine, Lys: lysine, AAA: aminoadipate, Enzymes: Ask: aspartate kinase, Asd: aspartate semialdehyde dehydrogenase, ThrA: homoserine dehydrogenase, ThrB: threonine kinase, ThrC: threonine synthase, MetX: homoserine O-acetyl-transferase, LysE: lysine exporter, Lat: lysine aminotransferase, Pcd: aminoadipate-semialdehyde dehydrogenase. Genes in square boxes. Color coding: green: wildtype levels, blue: reduces levels, red: increased levels.
inhibition). These observations fit well with the increased abundance of lysine and amino acid observed in stationary phase cells (Supplementary Fig. 8). It is tempting to speculate that Mtb has lost lysine feedback control at the AK level in order to allow increased lysine flux to act as a signal for cellular stress. In this model, elevated intracellular lysine levels would cause upregulation of the Lnt degradation pathway, which generates aminoacidopete and leads to the activation of the stringent response and persistence. Alternatively, lysine catabolism and its potential toxic intermediate(s) may serve as a means to control growth under conditions where replication could be lethal, similar to the role of Toxin-Antitoxin modules or methylglyoxal metabolism in persist cell formation. Potential candidates for toxic intermediates are aminoacidemia and piperidine-6-carboxylate (Fig. 1a), as both have been shown to be cytotoxic.

The second mechanism that mycobacteria employ to compensate for a missing lysine-AK feedback loop is catalyzed by a dedicated lysine/arginine permease LysE, encoded by Rv186. We show that LysE in Mtb is used as a metabolic relief valve to clear excess cytoplasmic lysine, as a result of the stringent response. This exporter also is a role under non-auxotrophic conditions as evidenced by the retarded exit from lag phase and slow growth phenotype of the Mtb ΔlysE mutant. Current literature shows that Rv186 is induced during infection of resting and activated macrophages, as well as in a lysosomal exposure model and it plays a role during latent TB infection in humans. The exporter is a dominant target of IL-2 secreting memory T cells and is proposed to contribute to protective immunity in humans. In fact, amino acids, specifically arginine, were shown to modulate human immunometabolism and T-cell responses. Together, these data suggest that LysE-mediated export of arginine, and possibly lysine, might play a role in shaping Mtb’s niche during infection.

The current environment of the nutritional composition of the Mtb microenvironment in the host is still incomplete. How that environment evolves over the course of an infection is even less understood. Auxotrophic bacterial strains are essential tools for identifying metabolic drug targets required during infection and probing the host environment for specific metabolites. However, because most Mtb auxotrophs fail to establish infection, their requirement for late-stage persistence cannot be investigated. To date, no data is available on the vulnerability of amino acid biosynthetic pathways during Mtb persistence in vivo. By utilizing conditional knockdowns of two branch-point enzymes (metX and thrA), we uncovered the aspartate pathway as being broadly required for persistent infection in mice. These data also clearly demonstrate the inability of Mtb to scavenge sufficient methionine, homoserine or threonine from the host throughout the course of infection. The observations presented here lay an important foundation for future drug development. We propose that the numerous essential enzymes identified in this pathway and their absence in eukaryotic cells make it particularly amenable to drug discovery efforts.

**Methods**

**Mycobacterial strain and growth conditions.** All strains and plasmids used in this study are listed in Supplementary Tables 1. Mycobacterial strains were grown in Middlebrook 7H9 medium (Difco) supplemented with 10% (vol/vol) OADC enrichment (0.5% oleic acid, 50 g albumin, 20 g dextrose, 0.04 g catalase, 8.5 g sodium chloride in 1 L water), 0.2% (vol/vol) glycerol, and 0.05% (vol/vol) tryptophan (Sigma). Antibiotic selection media contained 75 µg/mL hygromycin B, 20 µg/mL kanamycin, and/or 25 µg/mL streptomycin, as required. Supplemented media for culturing auxotrophic mutants contained 50 µg/mL methionine, 500 µg/mL threonine, and/or 50 µg/mL homoserine. For time-course starvation and supplement growth curve experiments, strains were grown in the presence of the required supplements to an OD₆₀₀ of 0.5, washed three times with PBS, and diluted into supplement free medium to an OD₆₀₀ of 0.01 (growth/kill curves) or 0.2 (metabolomics and transcriptomics). Samples from three independent biological replicates were harvested on days 0, 2, 4, and 7 (ΔΔHRA) or days 0, 1, 2 (ΔΔHRB) for RNA extractions, and days 0, 1, and 2 for metabolite extractions (see below). Pathologically we studied the infection in a C3H/Fla background model in which the main transcriptomic and metabolic changes are happening in the first 2 days. Hence, we focused on this timeframe for the rest of the study. Anhydrotetracycline (Acros) was dissolved in DMSO and used at the specified concentrations.

**Gene Knockout and complementation.** The genes thrA (Rv1294), thrB (Rv1296), lysE (Rv186), and lysG (Rv1865c) were deleted in Mtb H37Rv by specialized transduction. The Mtb H37Rv ΔmetX strain (previously known as ΔmetA) was constructed previously. Transductants were recovered on selective medium containing hygromycin (75 µg/mL) and appropriate amounts of supplements when needed: homoserine (50 µg/mL) for Mtb ΔΔHRA and threonine (500 µg/mL) for Mtb ΔΔHRB. Mutations were confirmed by three-primer PCR using respective L, R, and Universal_upg primer pairs, listed in Supplementary Table 2. The H37Rv ΔΔHRA and ΔΔHRB strains were cultured using the tetracycline-inducible dual control (DUC) knockdown plasmid. Briefly, each strain was complemented with pGMCJK-q19-T358S38-P750-6yg-DAS+4 (pGMCJK-6yg). The gene sequences were PCR-amplified using the respective P1 and P2 primers (Supplementary Table 2) and recombined into pD012A using BP cloning (Invitrogen), to generate the entry vectors pEN12A-ygf-DAS+. The final construct was generated by 3′-way gateway cloning with pEN23A-ygf-DAS+4, pEN41A-T358S38, pEN12A-P750, and pDE43-MCK using LR clonase II (Invitrogen), to generate the final construct. This was transformed into the respective mutant strain to generate DAS complements whose expression of the complemented genes was under a TetOFF promoter. The absence of either thermosensitive (ΔmetX-DAS and thrA-DAS) or the inducible (ΔΔHRA, ΔΔHRB) Mtb strains was confirmed with a copy of Rv186 (Supplementary Table 2). This DNA fragment was PCR amplified using primers Rv186_fl_EcoRI and Rv186_fl_HindIII (Supplementary Table 2) and was cloned into pMV361 using EcoRI and HindIII restriction sites, resulting in plasmid pMV361-LysE.

**Metabolite extractions.** Samples of bacterial cultures in biological triplicates were harvested at given time points. For intracellular metabolites, an equivalent of 10 mL culture at an OD₆₀₀ of 0.2 was rapidly filtered on a 0.22 µm nitrocellulose filter (Millipore). The filter paper containing the collection pad immediately placed into 1.5 mL screw cap tubes of 1 mL extraction solvent containing 20%/40%/40% (vol/vol) water/acetone/methanol (Fisher) with approximately 500 µL of silica beads at −20 °C. The samples were homogenized twice in a Precellys Evolution (Bertin) at 4,500 rpm for 45 seconds, cooled to 5 °C using a Cryolys (Bertin), and with 5 min rest between cycles. Samples were centrifuged, and 750 µL of the extract was filtered through 0.22-µm uniformly concentrated filter (Corning, Life Technologies) at 10,621 g for 3 min, and stored at −80 °C. For extracellular metabolites, 1 mL of culture was collected and centrifuged, and 750 µL of supernatant was filtered twice through 0.22-µm Spin-X centrifuge filters (Corning, Life Technologies) at 10,621 g for 3 min, and stored at −80 °C. Metabolite extractions were performed using Pierce NAP-5 desalting columns (ThermoFisher), according to manufacturer instructions. Metabolites were extracted by adding 200 µL sample to 800 µL of 50%/50% (vol/vol) acetonitrile/methanol (Fisher), the samples were spun down, as before, and the supernatant and collected at −80 °C until analysis.

**Metabolomics.** Metabolomics analysis was performed using an Acquity UPLC system coupled with a Synapt G2 quadruple–time of flight hybrid mass spectrometer (Waters, Massachusetts, USA). Column eluents were delivered via electrospray ionization. UPLC was performed in a hydrophilic interaction liquid chromatography-mode (HILIC) gradient elution using an Acquity 1.7-µm amide column (2.1 × 100 mm). The flow rate is 0.5 mL/min with mobile phase A (100% acetonitrile) and mobile phase B (100% water), both containing 0.1% formic acid. In both positive and negative mode, the gradient began with 1% B 1 min until 1 min, ramped 0% B by 8 min at 17% B, held at 17% B for 6 min, then ramped to 1% B by 19 min and held at 1% B to the end of run at 20 min. The mass spectrometer was operated in V mode for high sensitivity using a capillary voltage of 2 kV and a cone voltage of 17 V. The desolvation gas flow rate was 500 L/h, and the source and desolvation gas temperature were 120 °C and 325 °C, respectively. MS spectra were acquired in centroid mode from m/z 50 to 1,000,000 with a scan width of 0.5 s. Leucine enkephalin (2 ng/µL) was used as lock mass (m/z 556.2615 and 554.2615 in positive and negative experiments, respectively). Analysis was performed using MarkerLynx (Waters) with extended statistic function to identify statistically significant differences of metabolite abundances. A Targetlynx (Waters) database was compiled by running standards of metabolites of interest to determine retention times and m/z ratios. Calibration curves of compound mixtures are run routinely to determine the linear range of metabolite detection. All samples were analyzed using Targetlynx with manual curation of peak areas where necessary. With abundances we refer to raw intensity values for the area under the curve for each metabolite peak. Peak areas were used to calculate fold changes of metabolites relative to day 0. Metabolite abundances, retention times and m/z values can be found in Supplementary Data 1. Threonine and homoserine have the...
same molecular mass. The retention time of homoserine is 0.1–0.2 min slower than threonine, however differentiation of the two peaks is difficult. Fragmentation pattern of the two compounds in positive mode is nearly identical while in negative mode there is one distinction (m/z 72.04). We have detected high amounts of m/z = 120.0655 in the ThrB strain but not in Rv or ThrA. Since ThrB is downstream of the Thra strain, as expected from a strain that cannot produce homoserine nor threonine, the m/z 120.0655 was undetectable.

RNA extraction. Samples of bacterial cultures were harvested in triplicate at appropriate times. 10 ml of culture was pelletted and the supernatant decanted. The cell pellet was resuspended in 1 ml of TRIzol (Invitrogen) and incubated overnight at 4 °C, before storage at −80 °C. The suspension was transferred to Fast-Prep Blue Cap tubes and processed twice for 45 s at speed 6 in a Fast-Prep apparatus (MP Biomedicals). After a brief incubation on ice, the debris was spun down, and the supernatant (750 µL) was processed for purification (described below).

Microarray (Mtbb Δthra). Total RNA was purified from the cell lysate using Direct-zol RNA MiniPrep (Zymo Research) with on-column DNAse I according to the manufacturer’s protocol. Total RNA (~5 µg) was used to create cDNA labeled with aminoallyl dUTP (Sigma). Fluorescent Cy3 and Cy5 dyes (GE Healthcare) were then covalently attached to the aminoallyl tags. Each pair of differentially labeled probes was resuspended in 60 µL of hybridization buffer (500 µL for-mamide, 250 µL 20 x SSC, 5 µL 10% (wt/vol) SDS, 245 µL ultrapure water) and hybridized to the microarray slide overnight in a 42 °C water bath. Slides then were washed for 3 minutes under stringent wash conditions (three times for 5 minutes each) washing in 1 x SSC 0.1% SDS; three times 5 min each washing in 0.1 x SSC 0.1% SDS; three times for 5 min each washing in 0.1 x SSC, and a final dip in 0.22-µm-filtered Milli-Q water (Millipore). Arrays were scanned in a Genepix 4000A scanner, and spots were quantified with TIGR SpotFinder. The data then were processed in TIGR MIDAS. Two-color tiff images of the microarrays were inspected visually for defects and obvious spatial biases. Before normalization, pin- tip (block) intensity box plots were used to detect slides exhibiting spatial and/or pin biases; after normalization, intensity box plots and MA plots were used to assess the effectiveness of normalization in correcting these biases. Radio-intensity plots were used to identify outlier slides and to assess the effectiveness of the normalization process, and Z-score histograms were used to look for slides with abnormal intensity distributions. Slides that did not pass the quality-control process were rejected, and repeat hybridizations were performed. After normalization (total intensity and LOWESS) the in-slide replicate spots were averaged before expression ratios were calculated. The results from four independent biological replicates including two dye swaps then were subjected to a t-test without false discovery correction in TIGR MeV software. The analysis was used as a ranking method. For a general overview, genes with expression ratios >2 and -0.5 and a P-value <0.05 were used for data interpretation.

RNA-seq (Mtbb Δthra). RNA from the clarified cell lysate samples were purified by Triozol-chloroform precipitation according to the manufacturer’s instructions (Invitrogen). After purification remaining DNA was removed from RNA samples using the DNA-Free Turbo DNase Kit (Ambion) and RNA integrity was checked on a Bioanalyzer RNA 6000 Pico chip (Agilent). The ribosomal RNA (rRNA) was removed using the Bacterial RiboZero kit (Illumina), and removal was confirmed by Bioanalyzer as above. RNA-seq library construction was conducted with the TrueSeq Stranded mRNA Library Prep (Illumina) according to manufacturer’s instructions and the library was sequenced on a NextSeq 500 (Illumina) using the NextSeq 500/550 HO V2 (75 cycles) (Illumina). Illumina reads were mapped to the M. tuberculosis reference genome from NCBI (NC_000962.3) in the form of a SAM file using Bowtie2 (http://bowtie-bio.sourceforge.net/index.shtml), the SAM files were converted to coordinate-sorted BAM files using samtools, and reads over genes were counted using featureCounts [64]. Differential expression analysis was then conducted using R and the DESeq2 package.

Mouse experiments. Mouse studies were performed in accordance with the National Institutes of Health guidelines following the recommendations in the Guide for the Care and Use of Laboratory Animals [65]. The protocols used in this study were approved by the Institutional Animal Care and Use Committee of Albert Einstein College of Medicine (Protocols #2012114 and #20180612). Female SCID mice and female C57BL/6 mice (Jackson Laboratories) were infected via the aerosol route using a 1 × 10^6 cfu/mL mycobacterial suspension in PBS containing 0.05% tryptosol and 0.004% antifoam. Infected yielded approximately 1000 bacilli per lung as determined by quantification of bacillary loads at 24 h post-infection (four mice per group). Subsequently, four mice from each group were killed at days 1, 7, 17, 21, 63, 84, 98 (C57BL/6) or 1, 14, 22, and 42 (SCID) to determine the bacterial burden in the lung and spleen. Six SCID mice per group were kept for survival experiments. For conditional knockdown experiments, groups of mice infected with each strain were started on 2000 ppm doxycycline-treated chow (TestDiet, Missouri, USA) at 0, 1, 3, and 6 weeks. Four mice from each group were sacrificed at given times, the lungs and spleens were harvested, homogenized and plated on 7H10 media containing the appropriate supplement to determine the bacterial burden. All mice infected with Mtbb were maintained under appropriate conditions in an animal biosafety level 3 laboratory.

Pathology. Lung samples were fixed in 10% neutral buffered formalin for 48–72 h and the subjected to paraffin embedment. Tissues were sectioned at 5 µm and stained with Hematoxylin and Eosin (H&E).

MIC determination of toxic metabolite analogs. Cultures of Mtbb H7Rv, ΔlysE, ΔlysGcomp, and ΔlysG were grown to log-phase in un-supplemented media. Each toxic analog (Supplementary Fig. 1) was two-fold serially diluted in a sterile 96 well plate. In triplicate, each strain was diluted to an OD_{590} of 0.001 in wells containing each of the drugs to a final volume of 0.2 ml. The plates were incubated at 37 °C for 7 days and the growth was measured by optical density in an Epoch plate reader (Biotek, Vermont, USA). Percent growth was calculated based on the density in the untreated wells. For arginine rescue experiments, arginine was added to each well at a concentration of 50 µg/ml.

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability
All microarray and RNAseq raw data were deposited in Gene Expression Omnibus (GEO) (GSE191105 (Δthra), GSE191106 (Δthrb), GSE191107 (SuperSeries)). Data underlying Figs. 1–6 are provided as Source Data files. All other data are available from the corresponding author upon reasonable requests.

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Author contributions

E.H., G.M.C., W.R.J. and M.B. designed the research. E.H., D.R.S., L.B.M., S.I., J.M.T., T.F. and M.B. performed research. E.H. and M.B. analyzed data. E.H., G.M.C. and M.B. wrote the paper.
