Mycobacterium tuberculosis universal stress protein Rv2623 interacts with the putative ATP binding cassette (ABC) transporter Rv1747 to regulate mycobacterial growth

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Abstract

We have previously shown that the Mycobacterium tuberculosis universal stress protein Rv2623 regulates mycobacterial growth and may be required for the establishment of tuberculous persistence. Here, yeast two-hybrid and affinity chromatography experiments have demonstrated that Rv2623 interacts with one of the two forkhead-associated domains (FHA I) of Rv1747, a putative ATP-binding cassette transporter annotated to export lipooligosaccharides. FHA I domains are signaling protein modules that mediate protein-protein interactions to modulate a wide variety of biological processes via binding to conserved phosphorylated threonine (pT)-containing oligopeptides of the interactors. Biochemical, immunochemical and mass spectrometric studies have shown that Rv2623 harbors pT and specifically identified threonine 237 as a phosphorylated residue. Relative to wild-type Rv2623 (Rv2623WT), a mutant protein in which T237 has been replaced with a non-phosphorylatable alanine (Rv2623T237A) exhibits decreased interaction with the Rv1747 FHA I domain and diminished growth-regulatory capacity. Interestingly, compared to WT bacilli, an M. tuberculosis Rv2623 null mutant (ΔRv2623) displays enhanced expression of phosphatidyl-myo-inositol mannosides (PIMs), while the
ΔRv1747 mutant expresses decreased levels of PIMs. Animal studies have previously shown that ΔRv2623 is hypervirulent, while ΔRv1747 is growth-attenuated. Collectively, these data have provided evidence that Rv2623 interacts with Rv1747 to regulate mycobacterial growth; and this interaction is mediated via the recognition of the conserved Rv2623 pT237-containing FHA-binding motif by the Rv1747 FHA I domain. The divergent aberrant PIM profiles and the opposing in vivo growth phenotypes of ΔRv2623 and ΔRv1747, together with the annotated lipooligosaccharide exporter function of Rv1747, suggest that Rv2623 interacts with Rv1747 to modulate mycobacterial growth by negatively regulating the activity of Rv1747; and that Rv1747 might function as a transporter of PIMs. Because these glycolipids are major mycobacterial cell envelope components that can impact on the immune response, our findings raise the possibility that Rv2623 may regulate bacterial growth, virulence, and entry into persistence, at least in part, by modulating the levels of bacillary PIM expression, perhaps through negatively regulating the Rv1747-dependent export of the immunomodulatory PIMs to alter host-pathogen interaction, thereby influencing the fate of M. tuberculosis in vivo.

Introduction

Mycobacterium tuberculosis, the causative agent of tuberculosis (TB), remains a global public health problem, causing, in 2015 alone, over 10.4 million new cases and 1.8 million deaths worldwide [1]. M. tuberculosis is able to establish an asymptomatic latent infection that can later reactivate to cause active diseases [2–5]. In the latently infected immunocompetent host, the lifetime risk for reactivation is 10%. In those immunocompromised, the risk for recurrence of latent infection is 10% per year [2–5]. It has been estimated that one-third of the
world’s population is infected with *M. tuberculosis*, and it is generally believed that the majority of these individuals harbor latent bacilli [2–5]. The latently-infected thus constitute a significant reservoir for disease reactivation and transmission. Therefore, latent TB is a major hindrance to the control and eradication of *M. tuberculosis*. Understanding the mechanisms that regulate tuberculous latency and reactivation may lead to the design of strategies for better TB control.

We have previously demonstrated that *M. tuberculosis* Rv2623, a universal stress protein (USP) homolog, has the ability to regulate mycobacterial growth [6]. Results of transcriptome analysis have revealed that Rv2623 is among the most highly induced genes when *M. tuberculosis* is exposed to an environment of hypoxia or nitrosative stress [7–9], conditions that tubercle bacilli are likely to encounter in an infected host [6]. Rv2623 is also induced when *M. tuberculosis* is internalized by macrophages [10], in standing cultures [11], and in the lungs of infected mice during the chronic phase of tuberculous infection [7,12]. An *M. tuberculosis* Rv2623 deletion mutant (ΔRv2623) is unable to establish a chronic infection in mice and Guinea pigs, exhibiting a hypervirulence phenotype [6]. But the growth regulatory property of Rv2623 is not restricted to these *in vivo* systems. Thus, constitutive overexpression of Rv2623 in both *M. smegmatis* and *M. tuberculosis* attenuates bacillary growth *in vitro* [6]. Biochemical analysis of Rv2623 showed that it has the ability to bind ATP. Mutagenesis studies based on a 2.9 Å crystal structure yielded mutants defective in ATP binding. Analysis of these mutants revealed that the growth regulatory property of *M. tuberculosis* Rv2623 correlates with its ATP-binding capacity [6]. These observations prompted us to propose the possibility that *M. tuberculosis* Rv2623 may function as a signaling intermediate in a pathway that promotes latency [6].

Investigating the mechanisms by which this USP regulates bacillary growth, the present study provides evidence that (i) Rv2623 interacts with Rv1747, a putative ABC transporter annotated to export lipooligosaccharides [13], to negatively regulate *M. tuberculosis* growth; (ii) this interaction is mediated via the recognition of a conserved phosphothreonine-containing oligopeptide motif of the USP by the N-terminal FHA domain of Rv1747 (FHA I); and (iii) ΔRv2623, compared to WT bacilli, exhibits a morphological phenotype that is associated with increased levels of phosphatidylinositol mannosides (PIMs), immunologically active molecules that can modulate the host immune response [14,15]. Conversely, an Rv1747-deficient mutant produces lower levels of PIMs than WT *M. tuberculosis*. Further, while ΔRv2623 is hypervirulent in an infected host [6], ΔRv1747 is attenuated for growth *in vivo* [16,17]. Collectively, the interaction of Rv2623 with the FHA I of Rv1747 to negatively regulate *M. tuberculosis* growth and the divergent *in vivo* virulence and PIM phenotypes of ΔRv2623 and ΔRv1747 suggest that this USP may influence mycobacterial growth by modulating Rv1747’s functional activity, raising the possibility that Rv1747, which has been annotated as a lipooligosaccharide exporter [13], is implicated in the export of PIMs.

**Results**

**Interaction of Rv2623 with the Rv1747 FHA I domain: The yeast two-hybrid study**

Signaling pathways often mediate their biological functions through interactions of elements involved in the cascade [18–20]. Given the possibility that Rv2623 may be an intermediate in a signaling pathway that promotes persistence [6], we sought to identify its interacting partners. Indeed, evidence exists that various bacterial USPs interacts with specific interactors to regulate physiologically relevant processes [21,22]. Using the yeast two-hybrid screen with Rv2623 as bait, we have identified a set of *M. tuberculosis* proteins that interact with this USP, among which is the N-terminal FHA I of Rv1747, the focus of the present study (Fig 1A and 1B). The
rationale for focusing on the Rv1747 FHA I derives from the roles of forkhead-associated domains in mediating protein-protein interaction to facilitate a broad range of vital biological processes including signal transduction, transcription, cell cycle regulation, and protein transport [23–27]. Rv1747 is a putative ATP-binding cassette (ABC) transporter that harbors two FHA domains [13] (Fig 1A). The screen has identified a fragment spanning amino acids (a.a.) 46 to 142 of Rv1747, which harbors the majority of N-terminal FHA I (a.a. 29 to 92), to be an interactor of Rv2623 (Fig 1A and 1B). Yeast two-hybrid experiments using a re-cloned full-length Rv1747 FHA I domain validated its specific interaction with Rv2623 (Fig 1B).

Interaction of Rv2623 with the Rv1747 FHA I domain: The affinity chromatography study

To further validate the interaction between Rv2623 and the Rv1747 FHA I domain, affinity chromatography experiments were conducted (Fig 1C). For these studies, purified recombinant histidine (His$_6$)-tagged Rv2623 (Rv2623) and FLAG-tagged FHA I (FHA I: first 100 amino acids of Rv1747) were expressed in M. smegmatis mc$^{2}$155. Purified FLAG-tagged FHA I was passed over columns with or without Rv2623 immobilized onto the Nickel (Ni)-NTA resin. Western analyses of the appropriate elution fractions using anti-Rv2623 and anti-FLAG antibodies revealed that Rv2623 and Rv1747 FHA I co-eluted—upper and lower panels of lane 3 represent the results of probing eluents from column containing both (Ni)-NTA resin-immobilized (His$_6$)-tagged Rv2623 and FLAG-tagged Rv1747 FHA I with anti-Rv2623 and anti-FLAG antibody, respectively—thus demonstrating interaction of these two mycobacterial components. Lane 1: upper panel and lower panel represent results of reacting eluents from column with only Rv2623 with the appropriate antibody. The upper and lower panels of Lane 2 depict reactivity of eluents from column harboring only FHA I with the appropriate antibody. Lane 4 of upper and lower panels represent recombinant Rv2623 (+ve Rv) and FLAG-FHA I (+FHA); respectively, loaded as positive controls. α-Rv2623 and α-FLAG: anti-Rv2623 and anti-FLAG antibodies; respectively. Arrows indicated the molecular weight of His-tagged Rv2623 (~32.31 kDa) and FLAG-tagged Rv1747 FHA I (~12 kDa; expressed as the first 100 amino acids of Rv1747).
expressed in *M. smegmatis* mc²155 [28] via a pSD series acetamide-inducible expression system were used [29,30]. FLAG-FHA I was expressed as a tagged peptide spanning the first 100 a.a. of Rv1747 (Fig 1A). Western blot analyses of the various flow-through, wash, and elution fractions collected in the affinity chromatography studies using anti-FLAG and anti-Rv2623 antibodies revealed that Rv2623 and Rv1747 FHA I co-eluted, demonstrating the interaction of these two mycobacterial proteins (Fig 1C). This interaction was similarly and consistently observed when the affinity chromatography experiments employed recombinant Rv2623 derived from the previously described *E. coli*-based pQE80L-Rv2623 expression construct [6], and cMyc-FHA I harbored in the first 120 a.a. of Rv1747 and expressed in the LIC (ligation independent cloning) vector pMCSG7, as well as anti-His antibodies for the detection of Rv2623 (S1 Fig). Of note, results obtained from affinity chromatography experiments showed that Rv2623 does not interact with a pSD vector-expressed recombinant C-terminal Rv1747 FHA II domain (S2 Fig). Together, these results, in conjunction with that of the yeast two-hybrid experiments, strongly suggest that Rv2623 interacts specifically with the Rv1747 FHA I in the *in vitro* systems studied.

Bioinformatic analyses of the feasibility of *M. tuberculosis* Rv1747 FHA I to interact with conserved phosphothreonine-binding motifs of its interactors

The FHA domain, originally discovered in forkhead-type transcription factors [31], exists in a wide variety of proteins to mediate diverse functions via protein-protein interactions [23–27]. A characteristic property of FHA domains is that these interactions are based on recognition of a phosphorylated threonine (pT) residue of the binding partner [24,32,33]. One prototypic FHA domain that has been examined extensively is that of the *Saccharomyces cerevisiae* Rad53 protein kinase (Fig 2A and 2B). The Rad53 FHA1 mediates its functions in regulating cell cycle and DNA repair by binding to a pT residue that is embedded within a conserved oligopeptide motif [24,32,33]. Structural analysis of the Rad53 FHA1 has revealed that specific amino acids critical to the interaction with the pT motif are all located in the conserved region of this modular phosphopeptide recognition domain: Gly69, Arg70, Ser85, His88, and Asn107 (Fig 2A, 2B and 2D), with Asn107 and Arg70 implicated to play a role in interacting with the conserved phosphopeptide backbone. There is also evidence that Arg70, together with Ser85, play a role in hydrogen-bonding with the phosphorylated threonine of the conserved phosphothreonine-binding motif (Fig 2A and 2B) [24,32]. Parenthetically, the three amino acids critical in mediating binding with the conserved phosphopeptide motif (Asn 107, Arg70, and Ser85) are located on the surface of Rad53 FHA1 [24,32,33] (Fig 2A and 2B). The other two highly conserved non-surface FHA1 residues, Gly69 and His88, are thought to stabilize the structure of the binding site [24,32,33] (Fig 2A and 2B). The homology model of the FHA I domain of Rv1747 was generated utilizing the M4T server ver 3.0 [34] based on comparative modeling using a combination of 2 templates (PDB codes 2LC1 and 1UHT). The homology model of the Rv1747 FHA domain was then superimposed onto the Rad53 FHA domain [24,32,33] using Pymol (www.pymol.org) and displayed as shown in Fig 2C. This modeling studies of Rv1747 FHA I domain revealed a tertiary structure with 11-stranded β sandwich similar to that of the *S. cerevisiae* Rad53 protein kinase [24,32,33]. The conserved Gly, Arg, Ser, His, and Asn that play an important role in mediating the interaction of Rad53 with the pT motif are all present in Rv1747 FHA I (Gly32, Arg33, Ser47, His50, Asn69) with almost precisely matched spacing except for a one-amino acid difference between Rad53 FHA1 Arg70/Ser85 and Rv1747 FHA I Arg33/Ser47 (Fig 2C and 2D). In addition, tertiary structure modeling has revealed complete superposition of all the conserved a.a. between the *S. cerevisiae* Rad53 FHA1
and Rv1747 FHA I (Fig 2C and 2D). These results strongly support the feasibility of this M. tuberculosis forkhead-associated domain to interact with phosphorylated threonine-containing peptides. This observation prompted the initiation of study to determine whether putative phosphorylatable threonines are present in Rv2623.

Rv2623 harbors threonine residues that can potentially mediate interaction with FHA domains

Solvent accessibility analyses based on the crystal structure of Rv2623 revealed that the hydroxyl groups of five of the nine threonine residues of the USP studied are solvent-accessible and therefore potentially phosphorylatable [6,35,36] (Fig 3A and S3 Fig). The surface locale of these five threonine residues is further confirmed by Pymol display of Rv2623 (www.pymol.org; Fig 3B). Therefore, we examined the structure for additional elements described as part of the conserved pT peptide motif signature contributing to interactions with FHA domains. We
focused on the amino acid situated three residues from the corresponding phosphorylatable threonine toward the C-terminus (pT+3), because it has been observed, based on studies derived from peptide library screening experiments involving a subset of known FHA domains, that specific pT+3 residues contribute to mediating the interaction of this phosphoprotein-recognition module with phosphorylated threonine-containing oligopeptides by modulating binding and binding selectivity [24,32,33]. Sequence analysis revealed that four of the five Rv2623 threonines whose hydroxyl group is solvent-accessible (T90, T103, T212, T237) have residues in the pT+3 position that match those found in pT-containing peptides with known FHA domain-binding specificity (Fig S3A–S3C) [32].

Rv2623 is post-translationally modified and harbors phosphorylated threonines

The above-described experiments provide strong evidence that *M. tuberculosis* Rv1747 FHA I can interact with Rv2623 through recognition of and binding with potentially phosphorylatable threonine residues in the context of a conserved phosphopeptide motif including the pT+3 residues (Figs 1–3). To further evaluate the interaction of these two mycobacterial components, we investigated whether Rv2623 is phosphorylated. Lysates of BCG which encodes an Rv2623 orthologue that is 100% identical to the *M. tuberculosis* counterpart [tuberculist.epfl.ch].
ch], were resolved by two-dimensional (2-D) gel electrophoresis and probed using a monoclonal antibody specific for Rv2623. This revealed the USP to be detected as three isoforms (Fig 4A), which have comparable approximate molecular mass, but differ based on their isoelectric points, with a shift toward the acidic direction, suggesting that Rv2623 can be post-translationally modified via phosphorylation. Further, Rv2623 affinity-purified from \( M. \) \( \text{tuberculosis} \) and BCG lysates is reactive with antibodies relatively specific for phosphorylated threonines using both a mouse monoclonal antibody and a rabbit polyclonal antibody (Fig 4B). Similarly, affinity purified \( M. \) \( \text{smegmatis} \)-expressed recombinant Rv2623 is also reactive with the anti-phosphothreonine monoclonal antibody (Fig 4B). Together, these results suggest that Rv2623

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**Fig 4.** Rv2623 is post-translationally modified and phosphorylated at T237. (A) Immunoblot of 2D-gel electrophoretically-resolved BCG lysates with anti-Rv2623 monoclonal antibody revealed three isoforms with differing isoelectric pH values, thus providing evidence for post-translational modification of the USP. (B) Dot-blot analysis of affinity-purified Rv2623 from \( M. \) \( \text{smegmatis} \) mc \(^{25}315 \) (Top panel); \( M. \) \( \text{tuberculosis} \) Erdman (Mid panel), and \( M. \) \( \text{bovis} \) BCG (Bottom panel) demonstrating immunoreactivity with an anti-pT antibody. "++" indicate positive pT control; soy bean trypsin inhibitor serves as negative controls (S9 Fig). (C) Mass spectrometry-based phosphomapping of Rv2623. The sequence coverage of the protein was ~90%. Manual examination of the appropriate MS/MS spectra was conducted to verify the phosphopeptides identified via software programs. The graph shown represents MS/MS spectra of m/z 507.24(2+) corresponding to peptide spanning amino acid residues 231–238 of Rv2623. Labelled are b- and y- fragment ions produced from Collision Induced Dissociation (CID) in an LTQ-Orbitrap Elite LC MS/MS instrument. Phosphorylation on T237 of Rv2623 was determined based on the MS/MS fragmentation patterns; in particular are the observed loss of phosphoric acid (indicated as "y(n)–98Da") for fragment ions y(3), y(4), y(5), y(6), y(7), and b(7) under CID conditions [76]. The "***" sign indicates that the assigned fragment ion is doubly charged. The graph shown represents results derived from analysis of \( \textit{in vitro} \) phosphorylated \( M. \) \( \text{smegmatis} \)-expressed recombinant Rv2623. Analysis of \( \textit{in vitro} \) phosphorylated \( E. \) \( \text{coli} \)-expressed recombinant Rv2623 yielded the same results. (D) Molecular docking was employed to depict the interaction of the pTRVV-containing FHA domain-binding motif of Rv2623 with Rv1747 FHA I [38] (https://life.bsc.es/servlet/pydock/home). This was performed with the [amino acids 152 to 294] domain of Rv2623 (with T237 phosphorylated) on the [amino acids 1 to 90] domain of Rv1747 FHA I, using PyDockWeb, with the d(N64-V240) constrained. The phosphate was built into T237 by the PyTMs plugin in Pymol prior to the docking. The PDB code of Rv2623 is 3cis. The structure of the FHA domain of Rv1747 is taken from Fig 2 (B and C). The Threonine residue and the pT+3 residues in the binding motifs of Rv2623 are labeled in yellow and cyan, respectively.

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potentially can be phosphorylated at threonine residues, thus further supporting the possibility that a specific pT-containing peptide of this USP interacts with the FHA I domain of Rv1747.

Identification of potentially phosphorylated threonine residues of Rv2623

Results of 2-D gel electrophoretic analyses of Rv2623 and its reactivity with anti-phosphothreonine antibodies (Fig 4B) led us to initiate studies to directly examine if any of the threonine residues of Rv2623 can be phosphorylated. For these studies, recombinant Rv2623 purified from M. smegmatis or E. coli that had undergone an in vitro phosphorylation reaction with recombinant M. tuberculosis protein kinase G (PknG) was subjected to mass spectroscopic analysis. In pilot experiments, PknG was found to be among a set of serine/threonine protein kinases (STPKs) capable of phosphorylating Rv2623 to varying degrees. This observation is in agreement with a previous report noting that M. tuberculosis STPKs are relatively promiscuous in terms of substrate use [37]. PknG was subsequently used for our studies because of its availability (gift of Dr. J. Blanchard, Albert Einstein College of Medicine). The kinased samples were resolved by sodium dodecyl sulfate-polyacrylamide (SDS-PAGE) gel electrophoresis, transferred onto nitrocellulose membrane and probed with anti-phosphothreonine and anti-Rv2623 monoclonal antibodies to evaluate for the presence of pT residues in the USP. The results revealed that Rv2623’s reactivity to anti-phosphothreonine monoclonal antibody increases with time as the kinase reaction progresses (S4 Fig). For phosphomapping, the band corresponding to phosphorylated Rv2623 was excised from a Coomassie-stained gel containing electrophoretically-resolved kinasing reaction, washed, destained and then subjected to in-gel trypsin digestion. Phosphorylated species in the digest were then TiO$_2$-enriched and analyzed by Liquid Chromatography-Tandem mass spectrometry (LC-MS/MS) to identify specific phosphorylated residues. This phosphomapping study identified the Rv2623 threonine at position 237, one of the solvent accessible residues that also has a conserved pT+3 amino acid known to facilitate binding of FHA domains, as a phosphorylatable residue (Fig 4C). Having demonstrated that the M. tuberculosis Rv1747 FHA I possesses structural attributes and conserved determinants that mediate interactions with interactor proteins, and that the signature forkhead-associated domain-interacting phosphothreonine oligopeptide (together with the appropriate pT+3 residue) exists in Rv2623, in silico docking analysis was conducted [38]. Results of the analysis further support the interaction of these two mycobacterial components (Fig 4D).

Analysis of the role of Rv2623 T237 in regulating mycobacterial growth

The identification of a phosphorylatable threonine in Rv2623 with a pT+3 residue that can potentially mediate an interaction with Rv1747 FHA I (Figs 3 and 4) prompted studies designed to determine the significance of T237 in the growth-regulatory function of Rv2623. Because the phosphomapping experiment revealing that T237 is phosphorylatable used in vitro phosphorylated Rv2623 protein (Fig 4), we investigated, in addition to T237, the potential contributions of the other four solvent-accessible threonines of Rv2623 (T90, T103, T212, and T280 (Fig 3A)) to the growth-regulatory function of the USP. All solvent-accessible threonines of Rv2623 were individually mutagenized to alanine to yield five T→A mutants. The ability of the five mutants to attenuate growth in M. smegmatis mc$^\text{2}155$ upon pMV261-based overexpression in recipient cells was evaluated using the in vitro BACTEC system as previously described [6]. The results of these studies revealed that, relative to the WT Rv2623, the ability of the Rv2623$^{T103A}$ and Rv2623$^{T237A}$ mutants to attenuate mycobacterial growth when overexpressed in M. smegmatis mc$^\text{2}155$ was significantly reversed (Fig 5A). By contrast, overexpression of the Rv2623$^{T90A}$, Rv2623$^{T212A}$, and Rv2623$^{T280A}$ mutants in M. smegmatis mc$^\text{2}155$
displayed growth attenuating capacity comparable to that of the $Rv2623_{\text{WT}}$. These results suggest that of the five solvent-accessible threonines, T103 and T237 are the two residues that...
most likely contribute significantly to the growth-regulatory property of Rv2623. Expression study, however, revealed that while the levels of overexpression of T237A mutant and WT Rv2623 proteins in M. smegmatis were comparable, Rv2623_T103A was poorly expressed for reasons that are currently unknown. This latter result has led us to focus solely on role of Rv2623 T237 in regulating mycobacterial growth (Fig 5A). The observation that the melting temperatures of Rv2623_T237A (42.96˚C) and Rv2623_T237A (41.63˚C) are comparable effectively excludes the possibility that the loss of growth-regulatory ability of these threonine mutants is due to protein misfolding or instability (Fig 5B).

The ability of Rv2623 to regulate mycobacterial growth correlates with its ability to interact with Rv1747 FHA I and requires Rv2623 T237

The fact that mutating the phosphorylatable T237 to a non-phosphorylatable alanine results in the loss of the ability of Rv2623 protein to attenuate growth upon overexpression in M. smegmatis strongly suggests that this threonine residue plays a significant role in regulating mycobacterial growth in a phosphorylation-dependent manner. This, together with the above-described evidence supporting the ability of Rv2623 to interact with the phosphopeptide-recognition FHA I of Rv1747, has prompted us to conduct affinity chromatography experiments to examine the capacity of Rv2623_T237A to interact with Rv1747 FHA I. The results of these studies using purified differentially-tagged recombinant Rv2623 WT, Rv2623 T237A, and Rv1747 FHA I revealed that, relative to the WT Rv2623, the capacity of the T237A mutant USP to bind the M. tuberculosis Rv1747 FHA I, is significantly reduced (Fig 5C). Of note, although results derived from the affinity chromatography experiments for the study of protein-protein interaction are not quantitative, the incomplete ablation of the Rv2623-Rv1747 FHAI interaction (Fig 5C) upon T→A mutation of Rv2623 raises the possibility that additional factors may, in concert with T237 phosphorylation, regulate the interaction between these two mycobacterial proteins. Collectively, these results strongly suggest that mycobacterial growth can be regulated through the interaction of the phosphorylated T237 residue of Rv2623 and the FHA I of Rv1747.

To examine whether the Rv2623-Rv1747 interaction plays a role in regulating growth in virulent M. tuberculosis as demonstrated in M. smegmatis, the ability of Rv2623 T237A mutant protein to attenuate growth in the Erdman strain upon overexpression was examined using the in vitro BACTEC system [6]. As in the M. smegmatis study, the T237A but not T90A mutation of Rv2623 significantly reversed the ability of WT USP to regulate M. tuberculosis growth (Fig 5D). M. tuberculosis overexpressing the Rv2623 T237A mutant protein exhibited growth kinetics comparable to that of the WT Erdman strains lacking pMV261 or harboring the vector control (pMV261 with no Rv2623 sequence). Rv2623 WT, Rv2623 T90A, and Rv2623 T237A proteins were overexpressed to comparable levels in M. tuberculosis Erdman via pMV261 (S5 Fig), thus excluding the possibility that the observed inability of the T237A mutant protein to retard growth of the tubercle bacillus is due to inadequate expression. The growth regulatory property of Rv2623 WT, Rv2623 T90A, and Rv2623 T237A was further examined in M. tuberculosis by growth curve analysis monitoring OD_{600nm} (Fig 5E). In agreement with the BACTEC study, growth curve analysis also indicated attenuated growth in strains expressing Rv2623 WT and Rv2623 T90A, while that expressing Rv2623 T237A mutant grew at a rate similar to Erdman and the Erdman+pMV261 control (Fig 5E). Taken together, these results strongly suggest that T237 of Rv2623 is critical for the growth-regulatory function of this USP through mediating, in its phosphorylated form, the interaction with Rv1747: Binding of pT237-containing Rv2623 to the FHA I domain of Rv1747 negatively regulates M. tuberculosis growth.
**M. tuberculosis** Rv2623- and Rv1747-deficient mutant exhibit opposing and aberrant cell wall expression of phosphatidyl-myo-inositol mannosides (PIMs)

During the course of manipulation of the *M. tuberculosis* Rv2623-deficient mutant, it was observed that this strain displays a sedimentation phenotype when grown in Middlebrook 7H9 (M7H9) medium with Tween 80 (Fig 6A). Under this culture condition, WT Erdman and ΔRv2623 grow with similar kinetics monitored over a 2 week-period [6]. Upon standing of suspension of individual 10-day-old bacterial cultures, the ΔRv2623 strain formed a loosely-packed, fluffy sediment compared to WT (Fig 6A). In addition, when plated onto M7H10 agar, these two strains display distinct colony morphotypes, with the Rv2623-deficient strain forming colonies with an apparently smoother, less ruffled appearance (Fig 6B and 6C), which is restored to that of WT with complementation, thus demonstrating Rv2623-specificity of this phenotype (Fig 6B–6D). This observation, together with the functional assignment of Rv1747 as a putative exporter of lipooligosaccharides, prompted us to begin characterizing cell envelope components of *M. tuberculosis* ΔRv2623. We initiated studies to evaluate the expression of PIMs, one of the most abundant and bioactive glycolipid families in the *M. tuberculosis* cell wall [14,15]. Two-dimensional thin-layer chromatography (TLC) analyses of total polar lipid extracts (normalized by total protein content) revealed that the ΔRv2623 mutant displays a PIM profile distinct from that observed in WT Erdman (Fig 6E and 6F). Specifically, ΔRv2623 expresses higher amounts of PIMs, particularly tri- and tetra-acylated PIM₂ and PIM₆, compared to WT bacilli (Fig 6E & 6F). In a separate and independent series of experiments, Rv2623-specificity for the hyper-producing phenotype for all the PIM types examined was demonstrated by complementation—using the integrating pMV306 construct that expresses

![Fig 6. PIM expression by M. tuberculosis ΔRv2623 and ΔRv1747.](https://doi.org/10.1371/journal.ppat.1006515.g006)
Rv2623 under the control of its native promoter [6]—except for AC1PIM1 (Fig 6G). The apparent unique attribute of this latter PIM species is unclear. Given the observation that Rv2623 interacts with Rv1747 to negatively regulate \textit{M. tuberculosis} growth, the enhanced PIM levels of \textDelta{}Rv2623 suggest that Rv2623 may affect \textit{in vivo} \textit{M. tuberculosis} growth by modulating the production of PIMs, raising the possibility that Rv1747 (which has been annotated to be an exporter of lipooligosaccharides [13]), could be involved in PIM transport. Consequently, we evaluated the amount of PIMs in an Rv1747-deficient mutant. The results have revealed that \textit{M. tuberculosis} \textDelta{}Rv1747 is a hypo-producer of PIMs relative to WT Erdman (Fig 6H and S6 Fig). This latter observation lends support to the notion that Rv1747 can indeed be involved in the transport of PIMs, thereby influencing their levels in the \textit{M. tuberculosis} cell wall.

**Discussion**

Given the remarkable number of people world-wide estimated to harbor dormant tubercle bacilli [2–5], which physiological state renders treatment challenging [39], tuberculous latency and reactivation poses a formidable problem for the effective control of \textit{M. tuberculosis}. The mechanisms by which tuberculosis latency and reactivation are regulated remain incompletely defined [2–5]. We have previously reported that Rv2623, which encodes a USP and is highly induced by hypoxia and nitrosative stress, can regulate bacillary growth \textit{in vitro} and \textit{in vivo} [6]. Importantly, virulent \textit{M. tuberculosis} deficient in Rv2623 fails to establish a chronic persistent infection and is hypervirulent in an infected host [6]. This latter observation suggests that Rv2623 plays a role in regulating \textit{in vivo} \textit{M. tuberculosis} growth, particularly in the context of tuberculous latency [6].

The present study seeks to understand the mechanisms by which Rv2623 regulates \textit{M. tuberculosis} growth. The results, derived from yeast two-hybrid screen, affinity chromatography studies, crystallographic, as well as bioinformatics and modeling analyses [6,24,32,33], have provided evidence that this USP interacts with Rv1747, a putative ABC lipooligosaccharide exporter [13], to negatively regulate \textit{M. tuberculosis} growth in concert. This interaction is mediated via the recognition by the Rv1747 FHA I domain, an Rv2623 conserved oligopeptide motif harboring a phosphorylated threonine at position 237, as assessed by mass spectrometry-based phospho-mapping study. Compared to the WT USP, the interaction of Rv1747 FHA I domain with a mutant Rv2623 protein, which T237 has been mutated to a non-phosphorylatable alanine (Rv2623\textsubscript{T237A}), is significantly compromised. Significantly, \textit{in vivo} analyses of specific Rv2623 mutants have shown that the interaction of the USP with Rv1747 has functional consequences: relative to Rv2623\textsubscript{WT}, the growth regulatory capacity of the mutant Rv2623\textsubscript{T237A} is markedly attenuated. Together, these results have provided strong evidence that specific elements of Rv2623 and the FHA I domain of Rv1747 mediate the interaction between these two mycobacterial components to negatively regulate \textit{M. tuberculosis} growth in a phosphorylation-dependent manner.

In support of the observation on Rv2623-Rv1747 interaction, evidence exist that bacterial USPs mediate biological functions in concert with interacting molecules. For example, \textit{Escherichia coli} UspC functions as a scaffolding protein of the KdpD/KdpE two-component salt sensing signaling pathway, and via phosphorylation-dependent mechanisms, regulates the expression of the high-affinity K\textsuperscript{+} transporter KdpFABC [21]. There is evidence that the \textit{Halomonas elongate} USP TeaD may regulate the activity of the ectoine transporter TeaABC to maintain osmotic equilibrium [22]. The KdpD/KdpE and TeaD studies thus provide evidence for the interaction between bacterial USP and transporters, supporting our observation that the \textit{M. tuberculosis} USP Rv2623 interacts with the putative ABC transporter Rv1747.
M. tuberculosis Rv1747, a 92-kDa integral membrane protein, has been annotated as an ABC transporter [13]. The Rv1747's nucleotide-binding domain (NBD) and its membrane spanning domain (MSD), two major components of a typical ABC transporter, are fused [13,40]. This structural organization suggests that the transporter operates as a homodimer [13]. Although its function remains to be formally characterized, Rv1747 harbors elements strongly suggesting that it is an ABC transporter [40]. In addition to the NBD and MSD domains, Rv1747 carries the Walker A and B motifs, which make up the ATP-binding pocket of ABC transporters [40]. Rv1747 also harbors the ABC transporter family signature sequence, a characteristic 12-residue segment located between the two Walker motifs, and the 6-amino acid Linton and Higgins motif downstream of Walker B [40]. In vitro studies have revealed that the Rv1747 NBD displays ATPase activity [41]. A unique feature of Rv1747 is the presence in the NBD of two FHA domains, phosphopeptide-recognizing signaling modules that mediate diverse biologically important processes via protein-protein interactions [23–27]. Of the various M. tuberculosis ABC transporters, Rv1747 is the sole FHA domain-containing member [13]. And Rv1747, one of the 6 FHA domain-containing M. tuberculosis proteins (S7 Fig), is the only one in this group to harbor two such domains [23,26].

The FHA domain, together with the bacterial counterparts of eukaryotic STPKs, is an important component of reversible phosphoregulation pathways that mediates a wide range of biological processes [23–27]. Characterization of FHA-containing M. tuberculosis proteins including EmbR (Rv1267c) [42–46], FhaB (Rv0019c) [47], and GarA (Rv1827, glycogen-accumulation regulator A) [43,48–50] has revealed that these molecules, together with other mycobacterial components and STPKs, play important roles in regulating biologically highly significant processes, including synthesis of cell wall components, cell division, drug resistance, virulence levels, and metabolism. The observation that Rv2623 interacts with Rv1747 to regulate bacterial growth adds to the gravity of M. tuberculosis FHA domain-containing components. Rv1747 is required for optimal survival of M. tuberculosis in macrophages, dendritic cells, and in mice [16,17], and this virulence attribute is dependent on its phosphoregulation by PknF [16,17]. Thus, a ΔRv1747 M. tuberculosis mutant, while exhibiting no in vitro growth phenotype, is hypovirulent in mice [16,17]. This latter hypovirulence phenotype is in stark contrast to the in vivo hypervirulence of ΔRv2623 [6]. Given the observation that Rv2623 and Rv1747 interact to negatively regulate M. tuberculosis growth, the opposing in vivo virulence phenotype of ΔRv2623 and ΔRv1747 suggests that the USP may modulate the growth of the tubercle bacillus in vivo by attenuating the function of the putative transporter. This scenario, together with the divergent PIM phenotype of the Rv2623- and Rv1747-deficient mutants (with ΔRv2623 and ΔRv1747 being a hyper- and hypo-producer of PIMs; respectively), raises the possibility that the putative ABC transporter Rv1747 serves to transport PIM(s), and that these glycolipids may, at least in part, impact mycobacterial growth in vivo through their immune-regulatory attributes [14,15].

Lending credence to the above notions, there exist precedents that ABC transporters function as exporters of phospho- and glycolipids [51–53] and that protein phosphorylation serves as a regulatory mechanism of this transport system [52,54–56]. As well, it has been reported recently that among the most differentially up-regulated genes in an M. tuberculosis ΔRv1747 strain, relative to WT bacilli, are iniA and iniB [57], members of the iniBAC operon which are induced by a variety of agents that inhibit mycobacterial cell wall synthesis [58,59]. Thus, the differentially enhanced expression of genes of this operon in ΔRv1747 could reflect the bacterial reaction to certain dysregulated cell wall biosynthesis pathways. This latter idea is consistent with the diminished level of expression in ΔRv1747 of PIMs, molecules that are important components of the M. tuberculosis cell envelope, which also serve as precursors for the synthesis of lipomannan (LM) and mannose-capped lipoarabinomannan (LAM), the latter a major
mycobacterial surface lipoglycan that is essential in the tubercle bacillus [14,15,60]. Collectively, the ini operon observations, together with annotation of Rv1747 as an ABC transporter that exports lipooligosaccharides, and the diametrically opposed growth and PIM phenotypes of ΔRv2623 and ΔRv1747, have provided evidence suggesting that Rv1747 may function as a PIM exporter, perhaps translocating certain member(s) of this family of glycolipids from the cytoplasmic side to the periplasmic side of the plasma membrane [14,60].

The enzymes for PIM biosynthesis and the subcellular locales in which PIMs are produced are incompletely defined [14,60]. Evidence exists, however, that synthesis of PIM is compartmentalized, with the lower (PIM$_{1/2}$) and higher (PIM$_{3/6}$) order PIMs produced in the cytosolic and periplasmic side of the inner membrane, respectively [14,60]. This paradigm suggests that translocation of PIMs (PIM$_2$, PIM$_3$, and/or PIM$_4$) across the inner membrane must occur [14]. Nonetheless, bona fide transporter(s) of PIMs, their substrate specificity, and how many there are, are unknown [14]. Results of the present study suggest that Rv1747 may function as a PIM transporter. Due to the essentiality of PIM$_1$, PIM$_2$, and PIM-derived LM/ManLAM for M. tuberculosis survival, the biosynthesis of PIMs (including their translocation across the inner membrane) and related lipoglycans is likely complex and may involve compensatory and/or redundant mechanisms [14]. Indeed, the results of a study on PIM$_4$ channeling protein LpqW suggest that compensatory mechanisms are in place to ensure the maintenance of adequate levels of the essential LAM at the expense of higher order PIMs [61,62]. It is possible that this apparent LAM-preserving attribute of M. tuberculosis could be operative in the PIM-hypo-producing ΔRv1747. This would explain the results of a report noting that a ΔRv1747 strain, despite being a hypo-producer of PIMs (the precursors for LAM biosynthesis), the mutant displays WT level of LAM/ManLAM, notwithstanding the use of a non-quantitative immuno-detection approach and the lack of evaluation of PIM expression by the deletion mutant [57].

The present study has provided evidence that M. tuberculosis Rv2623 negatively modulates the transport function of Rv1747 to regulate mycobacterial growth through phosphorylation-dependent mechanisms (Fig 7). Further, the opposing PIM expression phenotype of the ΔRv2623 and ΔRv1747 mutants suggests that Rv1747 may be a transporter for PIMs, immunologically active molecules that have been shown in vitro to impact M. tuberculosis-host interaction to influence the immune response [14,15]. The immune-regulatory properties thus could contribute, at least in part, to the ability of the Rv2623-Rv1747 interaction to regulate M. tuberculosis growth in the host by influencing the anti-tuberculous response [14,15]. It is also plausible that the altered levels of expression of these important cell enveloped glycolipids can have an intrinsic effect on mycobacterial growth. These two possibilities are not mutually exclusive. PIMs are downregulated in stationary phase [14], and it has recently been observed that the production of PIMs by M. tuberculosis is enhanced during infection in primary human macrophages (S8 Fig). This latter result reinforces the concept that the tubercle bacillus can adapt to environmental signals during infection by modulating the cell envelope, including lipid components [63–67]. Whether the level of PIMs in the M. tuberculosis cell envelope can influence in vivo mycobacterial growth and virulence is currently unclear; that it can is supported by our findings that the PIM-hyperproducing ΔRv2623 is hypervirulent in vivo and is unable to establish a chronic infection, while ΔRv1747, a hypoproducer of PIMs, is attenuated for growth in mice [16,17].

Much remains to be learned regarding the precise mechanisms by which Rv2623 modulates mycobacterial growth and by which this USP is regulated. The presence of three apparent Rv2623 isoforms suggests that the regulation of this protein is complex, and that phosphorylation of T237 might not be the only mode of post-translational modification involved. The phosphorylated Rv2623 that was subjected to mass spectrometric analysis was generated via an in vitro phosphorylation reaction mediated by recombinant PknG. Hence it is possible that in
vivo, other threonines, in addition to T237, could serve as additional targets of phosphorylation, thus accounting for the three Rv2623 isoforms observed. It is also possible that other mode of post-translational modification is involved. Equally unclear is the mechanism underlying the previously described dependency of the \textit{M. tuberculosis} growth-regulating attribute of Rv2623 on its ATP-binding capacity [6]. Is this dependency related to the Rv2623-Rv1747 interaction? Finally, the apparent function of Rv1747 as a PIM transporter and the mechanisms underlying its regulation remain to be characterized. This will likely not be a straightforward endeavor given the transmembrane nature of Rv1747. Nevertheless, the collective results of the present study have provided a framework for understanding the mechanisms by which Rv2623 interacts with the putative PIM transporter Rv1747 (Fig 7) to regulate \textit{M. tuberculosis} growth, particularly in the context of tuberculous persistence, and for potentially advancing knowledge of the biosynthetic pathways of mycobacterial glycolipids and lipoglycans.

Materials and methods

Bacterial strains, culture conditions, and molecular cloning

Primers, oligonucleotides (IDT DNA technologies) and vectors used for generating various constructs are listed in \textit{S1 Table}. \textit{M. tuberculosis} strain Erdman and H37Rv and \textit{M. smegmatis}...
The Mc2155 were cultured in supplemented Middlebrook 7H9 (M7H9) medium (Becton Dickinson, Sparks, MD) as previously described [6]. Mycobacterial strains overexpressing WT Rv2623 and various mutants of the USP were maintained in supplemented M7H9 broth containing 40 μg/ml of kanamycin. Mycobacterial growth was monitored by measuring absorbance at 600 nm or by the in vitro BACTEC 9000 system as previously described (Becton Dickinson, Sparks, MD) [6]. The latter approach involved inoculating stationary phase *M. tuberculosis* or *M. smegmatis* in triplicates into BD Myco/F lytic vials (final bacterial suspension of 10^4 colony forming units (CFU) per ml) (BD Bosciences, Sparks, MD), whose liquid medium is supplemented with a compound that fluoresces as a result of oxygen depletion due to bacterial growth. The time to detection of fluorescence signals thus reflects the rate of bacterial growth.

The pSD series vectors, which direct expression of proteins of interests in *M. smegmatis* under the control of the acetamide-inducible promoter, were used to generate N-terminal His_6-tagged and N-terminal FLAG-tagged recombinant Rv2623 (His-Rv2623 and FLAG-Rv2623) in mc2155 [29,30]. Fragments of Rv1747 representing the first 120 amino acids and harboring the FHA I domain (Fig 1A) were PCR-amplified to contain the cMyc or FLAG tag and cloned into the pSD vectors for the expression of an N-terminal cMyc-tagged FHA I (Myc-FHA I) and N-terminal FLAG-tagged FHA I (FLAG-FHA I) [29,30]. An N-terminal FLAG-FHA II (spanning amino acids 201–310) fusion (FLAG-FHA II) was similarly expressed in the pSD system [29,30].

The pQE80L (Qiagen, Inc.)-based plasmid pQE-Rv2623 containing an N-terminal His_6-tagged Rv2623 fusion construct was also used for the expression of His-Rv2623 [6]. Various *M. tuberculosis* Rv2623 threonine residue mutants were derived from pQE-Rv2623 via site-directed mutagenesis and similarly expressed (see below). As well, Myc-FHA I and FLAG-FHA I were also expressed in *E. coli* via the LIC (Ligation Independent Cloning) vector pMCSG7 [68] (pMCSG7-His-TEV-cMyc-FHA I; pMCSG7-His-TEV-FLAG-FHA I) (TEV: Tobacco Etch Virus cleavage site).

**Expression and purification of recombinant *M. tuberculosis* proteins**

The plasmid pSD1-Rv2623 was electroporated into *M. smegmatis* mc2155 for the expression of His-Rv2623 in this acetamide-inducible system [29,30]. Log-phase cultures were induced with acetamide (final concentration: 0.2%) overnight at 37˚C with shaking. Expression of *M. tuberculosis* Rv2623 based on the pQE80L system was carried out as previously described, following isopropyl beta-D-thiogalactoside (IPTG; final concentration, 0.3mM) induction in transformed BL21 *E. coli* E. coli [6]. The threonine residue mutants of Rv2623 were similarly expressed. Protein purification was carried out as described with modification [6]. Autoinduction medium was used for the expression of recombinant proteins via the LIC vector pMCSG7 [68]. Cells were disrupted using hydraulic press or the Matrix B and Fast Prep apparatus (MP Biomedicals, CA) for Mycobacterium and sonication for *E. coli*. Clarified bacterial cell lysates were filter-sterilized and expressed proteins were purified by affinity chromatography using appropriate antibodies against specific tags or Rv2623 followed by gel-filtration chromatography as described previously [6] (S1 Text).

**Yeast two-hybrid identification of Rv2623 interacting proteins**

*M. tuberculosis* Erdman strain chromosomal DNA and pGADT7 AD (Clontech, Mountain View, CA) cloning vector DNA were used to generate the yeast library for the yeast two-hybrid screen. Construction of the library was carried out by DNA Technologies Inc. (Gaithersburg, MD). Two *E. coli* libraries (> 2 x 10^6 clones per library) were generated by cloning partially digested, size-fractionated chromosomal DNA consisting of fragments spanning 0.5 to 2.5 kb as well as those ranging from 2.5 to 5.0 kb into the BamHI site of the pGADT7AD vector.
These two DNA libraries, which were used to screen for Rv2623-interacting prey proteins were cloned in pGADT7AD harboring the GAL4 activation domain. pGADT7AD has the Leu2 nutritional marker for selection in yeast. The gene for the bait protein Rv2623 was cloned into the pGBK7 vector to generate an Rv2623-GAL4 DNA-binding domain translational fusion. pGBK7 has the TRP1 nutritional marker for selection in yeast. The screens were conducted using Matchmaker Gold Yeast Two-Hybrid System, according to manufacturer's instructions. To confirm the interaction between the prey hits with Rv2623, the putative interactor sequence harbored in pGADT7AD was rescued. pGADT7AD harboring DNA fragments expressing the putative interactor were then individually co-transformed with pGBK7 containing the full-length Rv2623 DNA into an auxotrophic reporter strain Y2H Gold (his, ade) for validation of the interaction. The positive control was Rv2623 interacting with itself, as it forms a dimer in solution [6].

Affinity chromatography and immunoblotting experiments

*E. coli* and/or *M. smegmatis*-expressed His-tagged full-length WT Rv2623 (Rv2623<sub>WT</sub>), the Rv2623<sub>T237A</sub> mutant, and cMyc-tagged FHA I, and FLAG-FHA II were purified by affinity chromatography using the appropriate antibody followed by gel filtration as described for use in the affinity chromatographic study [6]. Spin columns were packed with 50 μl of Ni-NTA agarose bead slurry (~25 μl resin; Qiagen) and equilibrated with Buffer A (S1 Text). Resins in the equilibrated columns were either left untreated or allowed to react with 150 μg each of either His-tagged Rv2623<sub>WT</sub> or Rv2623<sub>T237A</sub>, for 2 hours at 4˚C. The columns were then washed to rid of unbound proteins using Buffer B (S1 Text). Appropriate columns were allowed to react with 400 μg each of recombinant cMyc or FLAG-tagged FHA I for 2 hours at 4˚C and then thoroughly washed to remove unbound proteins using Buffer A and Buffer B sequentially. The columns were then eluted for bound proteins with 100 ml of Buffer C (S1 Text). The eluates were electrophoretically resolved and subjected to Western blot analysis using antibodies specific for the specific epitope tag or to Rv2623, followed by an appropriate Horseradish peroxidase-conjugated secondary antibody, and then subjected to detection by chemiluminescence (Amersham). Proteins were densitometrically quantified (S1 Text).

Two-dimensional gel electrophoretic and Western blot analysis of *M. tuberculosis* Rv2623

The BioRad system was used for the 2-D gel electrophoresis analysis. To prepare samples for isoelectric focusing, 150 μg total protein (1 μg for purified protein) was processed using a ReadyPrep 2-D Cleanup Kit (Bio-Rad, Hercules, CA) according to the manufacturer’s instructions. The prepared protein (final volume: 125 μl) was allowed to react with a ReadyStrip IPG, pH range 4–7 (Bio-Rad) in a rehydration tray for ~16 hours at 20˚C. The IPG strip thus prepared was subjected to isoelectric focusing for three hours (8kV for 10kvh (50 μA/strip)). The focused IPG strip was equilibrated and washed according to the manufacturer’s instructions, and loaded onto a 1.5 mm-thick 14% SDS polyacrylamide gel with a 4% stacking gel, along with a Precision Plus Protein Dual Color Standard (Bio-Rad, Hercules, CA). The gel was run at 155 V until the dye front migrated to the bottom of the gel (~60 min). In some cases, the gel was then stained with Sypro Ruby (Bio-Rad, Hercules, CA) according to the manufacturer’s instructions. In other cases, the SDS gel was transferred to a PVDF membrane and analyzed for protein separation and detection of Rv2623 by standard Western blotting techniques. The PVDF membrane was reacted with a primary monoclonal antibody against *M. tuberculosis* Rv2623 (Advanced Immunochemicals, Inc., Long Beach, CA; 5-Rv2623-A10) followed by a secondary horseradish peroxidase-conjugated anti-mouse IgG antibody. Protein was
visualized by chemiluminescence using a Super Signal West Pico Chemiluminescent Substrate (ThermoFisher Scientific, Waltham, MA) according to the manufacturer’s instructions and exposed to Kodak scientific imaging film (BioMax XAR, Eastman Kodak Company, Rochester, NY). Relative abundance of protein was quantified using ImageQuant densitometry software (Molecular Dynamics, Inc; Sunnyvale, CA). In some experiments, the PVDF membrane was also stained with Coomassie Brilliant Blue G-250 (Bio-Rad, Hercules, CA) according to the manufacturer’s instruction.

**Dot-blot analysis for Rv2623 phosphorylation**

Recombinant Rv2623 derived from *M. smegmatis*, as well as the native USP immunoaffinity-purified from *M. tuberculosis* and BCG cell lysates were analyzed for phosphorylation using a dot blot assay. Briefly, increasing concentrations of Rv2623 protein derived from the various sources were spotted onto nitrocellulose membrane. Phosphothreonine (Sigma-Aldrich; St. Louis, MO) served as positive control. The membrane was allowed to dry, followed by blocking with 5% BSA in Tris-HCl (20 mM Tris-HCl; pH 7.5) containing 150 mM NaCl, 0.05% Tween 20 (TBST). The membrane was probed with anti-phosphothreonine antibodies: clone #42H4 mouse monoclonal or rabbit polyclonal antibodies (Cell Signaling Technology; Danvers, MA), washed thrice with TBST and then reacted with appropriate horseradish peroxidase (HRP)-conjugated secondary antibody (Sigma-Aldrich; St. Louis, MO). The blot was developed using ECL reagent (Amersham; Piscataway, NJ).

**Mass spectrometry analysis of Rv2623 phosphorylation**

Purified recombinant Rv2623 (5 μg) [6] was subjected to *in vitro* phosphorylation by *M. tuberculosis* PknG as described in S1 Text. The reaction product, electrophoretically resolved in a 10% SDS PAGE, was subjected to Western blot analysis using anti-phosphothreonine antibody (Cell Signaling; Danvers, MA). In parallel, a separate gel of electrophoretically resolved reaction mixture was stained with Coomassie Blue to identify the Rv2623 band based on molecular mass. The Coomassie Blue-stained gel band harboring the phosphorylated Rv2623 protein was excised, and the sample sent in chloroform-treated microcentrifuge tubes (Eppendorf; Westbury, NY) to the Mass Spectrometry & Proteomics Resource of the W.M. Keck Foundation Biotechnology Resource Laboratory (Yale School of Medicine, New Haven, CT) for further processing and mass spectrometric analysis for phosphorylated threonine based on published protocol [69]. Briefly, the gel band was SpeedVac-dried, then solubilized in appropriate buffer. The protein sample was then subjected to dithiothreitol (DTT) reduction, IAN-mediated alkylation, and the urea content brought down to 2 M with water prior to trypsin digestion. The digested sample was acidified with 0.5% trifluoroacetic acid (TFA), 50% acetonitrile and then subjected to titanium dioxide enrichment using the Top Tips system (Glygen Corp; Columbia, MD). The resulting phosphopeptide-enriched sample, dissolved in 70% formic acid and diluted with 0.1% TFA, was then subjected to LC-MS/MS analysis using the LTQ Orbitrap Elite that is equipped with a Waters nanoACQUITY UPLC system, and which uses a Waters Symmetry C18 180 μm x 20 mm trap column and a 1.7 μm, 75 μm x 250 mm nanoACQUITY UPLC column for peptide separation. The acquired data was peak-picked and searched using the Mascot Distiller and the Mascot search algorithm, respectively; and quantitatively processed with Progenesis LCMS (Nonlinear Dynamics, LLC) as previously described [69] (S1 Text). Protein identification was achieved using Mascot search algorithm (Matrix Science; Boston, MA) as described [69] (S1 Text). Manual examination of the MS/MS spectra was conducted to verify the phosphopeptides identified via software programs.
Generation of strains of *M. smegmatis* mc²155 and *M. tuberculosis* Erdman overexpressing WT Rv2623 and various Rv2623 threonine mutants

Wild-type Rv2623, with and without a His₆ tag, was cloned into the pMV261 vector under the control of the hsp60 promoter using 5'-EcoRI and the 3'-HindIII sites [30]. This construct served as template for the generation of specific Rv2623 single amino acid substitution threonine mutants by mismatched PCR priming using primers designed to direct the incorporation of specific mutations into the Rv2623 coding region following Stratagene Quikchange (La Jolla, CA) protocol as described [6]. All constructs were confirmed by sequence analysis. Appropriate constructs were transformed into *M. smegmatis* mc²155 and *M. tuberculosis* Erdman to generate mycobacterial strains over-expressing WT Rv2623 and the desired threonine mutants. Transformants were screened by PCR targeting the kanamycin resistance marker (for *M. smegmatis* and *M. tuberculosis*) and/or Rv2623 (for *M. smegmatis*). PCR product from *M. smegmatis* was gel purified (Qiagen gel extraction kit; Valencia, CA) and the sequence of the WT and mutant Rv2623 verified. Following sequence verification, expression of Rv2623 and its mutants in *M. smegmatis* was probed by Western blot, using a mouse anti-Rv2623 monoclonal antibody (Advanced Immunochemical; Long Beach, CA). To distinguish between the native and pMV261-based expression of the USP and its mutants in *M. tuberculosis*, a mouse anti-His tag monoclonal antibody (Sigma-Aldrich; St. Louis, MO) was used. Finally, thermal denaturation curves were determined for purified WT and mutant Rv2623 using an IQ5 Real Time PCR Detection System (Bio-Rad; Hercules, CA) following incubation with SYPRO Orange protein gel stain (ThermoFisher Scientific, Waltham, MA) as previously described [6] (S1 Text).

Lipid analysis of *M. tuberculosis* strains

For analysis of PIMs located in the cell wall of *M. tuberculosis* strains, bacteria were harvested 12 days after inoculation on M7H11 agar plates and lysed [70]. Protein quantification was performed by the BCA method following the manufacturer’s instructions (Bio-Rad, Hercules, CA). Lysates totaling 10 mg of protein from each strain were delipidated at 37˚C for 12 h using CHCl₃:CH₃OH (2:1, v/v) followed by CHCl₃:CH₃OH:H₂O (10:10:3, v/v/v/v) for an additional 12 h. For 2D-TLC analyses, total crude lipids from each strain were loaded (100 μg) onto the origin of a 10 cm × 10 cm TLC silica gel 60 F₂₅₄ aluminum plate (EMD Millipore, Temecula, CA) based upon equal amounts of protein content from the bacterial lysates and run in the first dimension using CHCl₃:CH₃OH:H₂O (60:30:6, v/v/v) as a solvent system. The TLC plate was then rotated 90˚ to the left and run in the second dimension using CHCl₃:CH₃COOH:CH₃OH: H₂O (40:25:3:6, v/v/v/v) as a solvent system. Plates were dried and sprayed with 10% concentrated sulfuric acid in absolute ethanol and heated at 110˚C until lipid bands appeared [71]. The NIH ImageJ program (http://rsb.info.nih.gov/ij/) was used to perform PIM densitometry analysis by calculating mean spot intensities from three independent experiments.

Supporting information

S1 Text. Supplementary materials and methods
(DOCX)

S1 Fig.
(DOCX)
S2 Fig.  
(DOCX)

S3 Fig.  
(DOCX)

S4 Fig.  
(DOCX)

S5 Fig.  
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S6 Fig.  
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S7 Fig.  
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S8 Fig.  
(DOCX)

S9 Fig.  
(DOCX)

S1 Table.  
(XLSX)

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