FasR Regulates Fatty Acid Biosynthesis and Is Essential for Virulence of Mycobacterium tuberculosis

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Mycobacterium tuberculosis, the etiologic agent of human tuberculosis, is the world’s leading cause of death from an infectious disease. One of the main features of this pathogen is the complex and dynamic lipid composition of the cell envelope, which adapts to the variable host environment and defines the fate of infection by actively interacting with and modulating immune responses. However, while much has been learned about the enzymes of the numerous lipid pathways, little knowledge is available regarding the proteins and metabolic signals regulating lipid metabolism during M. tuberculosis infection.

In this work, we constructed and characterized a FasR-deficient mutant in M. tuberculosis and demonstrated that FasR positively regulates fas and acpS expression. Lipidomic analysis of the wild type and mutant strains revealed complete rearrangement of most lipid components of the cell envelope, with phospholipids, mycolic acids, sulfolipids, and phthiocerol dimycocerosates relative abundance severely altered. As a consequence, replication of the mutant strain was impaired in macrophages leading to reduced virulence in a mouse model of infection. Moreover, we show that the fasR mutant resides in acidified cellular compartments, suggesting that the lipid perturbation caused by the mutation prevented M. tuberculosis inhibition of phagolysosome maturation. This study identified FasR as a novel factor involved in regulation of mycobacterial virulence and provides evidence for the essential role that modulation of lipid homeostasis plays in the outcome of M. tuberculosis infection.

Keywords: tuberculosis, fatty acid biosynthesis, lipid homeostasis, mycobacterial virulence, mycobacterial cell envelope
INTRODUCTION

Tuberculosis (TB) is one of the top 10 causes of death and the leading cause from a single infectious agent, exceeding both malaria and HIV. In 2018, there were an estimated 10 million new cases of active TB, and 1.3 million patients died from active infection (WHO, 2019). The emergence of multidrug-resistant TB (MDR-TB) presents an increasingly difficult therapeutic challenge; consequently, MDR-TB is now the main cause of death due to antimicrobial resistance (WHO, 2019) thus transforming this long neglected disease into a global health priority.

The composition and complexity of the cell envelope is the most distinctive feature of Mycobacterium tuberculosis, the etiologic agent of human TB. Biochemical and microscopic data revealed that the mycobacterial envelope consists of three discrete entities: a plasma membrane surrounded by a complex cell wall and an outermost layer called the capsule in pathogenic species (Chiaradia et al., 2017; Daffe and Marrakchi, 2019). The M. tuberculosis cell wall consists of peptidoglycan covalently linked to arabinogalactan esterified by mycolic acids (MA), which form the inner leaflet of the outer membrane bilayer (mycomembrane). The outer leaflet of this membrane is composed of free, non-covalently bound lipids and glycolipids like trehalose monomycolates (TMM), trehalose dimycolates (TDM), glycerol monomycolates, glucose monomycolates, phthiocerol dimycocerosates (PDIM), poly acylated trehaloses (PAT), sulfolipids (SL), phosphatidylinositol mannosides (PIM), phenolic glycolipids (PGL), and mannose-capped lipoarabinomannans (Man-Lam; Brennan and Nikaido, 1995; Daffe and Reyrat, 2008).

During the course of infection, intracellular pathogenic bacteria use different strategies to survive, multiply, and persist inside the host (Russell et al., 2010). M. tuberculosis encounters heterogeneous host environments, including high oxygen tension in the lung alveolus, reactive oxygen and nitrogen species in activated macrophages, as well as nutrient depletion and hypoxia in tuberculous granulomas. How M. tuberculosis survives these diverse environments is not fully understood. M. tuberculosis has evolved a wide array of specific lipids and related metabolic pathways that actively interact with the immune system, and multiple reports from human and animal studies suggest an intimate link between M. tuberculosis lipid metabolism and outcome of infection (Neyrolles and Guilhot, 2011; Passemard et al., 2014; Vromman and Subtil, 2014). Of particular interest is the challenge that M. tuberculosis must face in order to adjust its own lipid metabolism while the macrophage is changing its lipid content (Teng et al., 2017). Little is known about the mechanism underlying the reorganization of M. tuberculosis envelope during infection (Dulberger et al., 2020). Understanding how this pathogen selects and organizes the envelope components to resist killing, and how these components make the envelope essential to M. tuberculosis survival, virulence, and pathogenesis, is still a major challenge. Therefore, characterization of the regulatory elements that control these processes should help improve understanding of the mechanisms involved in mycobacterial pathogenicity and could provide new potential targets for anti-TB chemotherapy.

Previous studies on the regulation of lipid biosynthesis in mycobacteria allowed us to identify FasR as the transcriptional activator of fatty acid biosynthesis in M. smegmatis (Mondino et al., 2013). Deletion of fasR from M. smegmatis was possible only in the presence of an extra copy of the gene, suggesting that fasR is essential for viability. Notably, we found that the affinity of FasR for its target promoter is modulated by long chain acyl-CoAs (≥C₁₆), the end products of the FAS I pathway. FasR, together with MabR, the transcriptional regulator of MA biosynthesis, is the only transcription factor involved in fatty acid biosynthesis regulation known to be essential for M. smegmatis survival, reflecting the relevance of lipid homeostasis for mycobacterial viability (Salzman et al., 2010; Mondino et al., 2013).

In this study, a M. tuberculosis knockout strain in fasR was constructed and the mutant was used to analyze the implications of lipid homeostasis in M. tuberculosis virulence. Inactivation of fasR in M. tuberculosis was not lethal for in vitro growth but led to complete rearrangement of lipid composition of the envelope. Consequently, the mutant strain was less virulent in a mouse model and showed impaired replication in macrophages, thus implying that lipid homeostasis plays an important role during M. tuberculosis infection.

MATERIALS AND METHODS

Ethics Statement

All animal experiments were performed in compliance with the regulations of the Institutional Animal Care and Use Committee (CICUAE) of INTA with the authorization Number 20/2016.

Bacterial Strains, Culture, and Transformation Conditions

The E. coli strain DH5α (Hanahan, 1983) was used for routine subcloning and was transformed according to Sambrook et al. (1989). M. tuberculosis H37Rv (Cole et al., 1998), MtbΔfasR, and MtbΔfasR-cfasR were grown at 37°C in Middlebrook 7H9 broth (DIFCO) supplemented with 0.2% glycerol, 0.03% Tyloxapol and 10% OADC or Middlebrook 7H10 agar (DIFCO) supplemented with 0.2% glycerol and 10% OADC. Hygromycin (Hyg, 50 μg/ml), kanamycin (Kan, 15 μg/ml), and streptomycin (Str, 12 μg/ml) were added when needed.

Recombinant plasmids and strain genotypes are listed in electronic supplementary material (Supplementary Tables S1 and S2).

DNA Manipulation, Plasmid Construction and Mutant Generation

Isolation of plasmid DNA, restriction enzyme digestion, and agarose gel electrophoresis were carried out by conventional methods (Sambrook et al., 1989). Genomic DNA of M. tuberculosis was obtained as described previously (Connell, 1994). For plasmid generation purposes, all PCR products were ligated into the pCR®-Blunt II-TOPO® Vector (Invitrogen) according to the manufacturer’s recommendation.
For the construction of the fasR mutant allele, a fragment of 946 bp containing 58 bp of the fasR 5′ end plus 882 bp of the upstream region was PCR-amplified from genomic DNA using oligonucleotides 3208_1up and 3208_2down (Supplementary Table S3). The PCR product was cloned in the pCR BluntII TOPO vector (Invitrogen). The resulting plasmid was digested first with EcoRV and after with XhoI/NdeI. The 940 bp fragment obtained was cloned in the suicide vector pJG1100 (Gomez and Bishai, 2000; Muñoz-Elias and McKinney, 2006) digested with the same enzymes, generating plasmid pFR34. Additionally, a fragment of 883 bp containing 87 bp of the fasR 3′ end plus 784 bp of the downstream region was PCR-amplified from genomic DNA using oligonucleotides 3208_3up and 3208_4down (Supplementary Table S3). The PCR product was cloned in the pCR BluntII TOPO vector (Invitrogen). The resulting plasmid was digested first with EcoRV and after with NdeI/SpeI, and the fragment (877 bp) was cloned in a pFR34 vector digested with the same enzymes, generating plasmid pFR35.

For the construction of a fasR merodiploid strain, the fragment resulting from the digestion of the plasmid pFR3 (Mondino et al., 2013) with XbaI and EcoRI was cloned between the AvrII and MfeI sites in plasmid pGA44 (Kolly et al., 2014). The resulting plasmid, pFR33, is an integrative plasmid that harbors fasR expressed under the control of a Ptr promoter and the TetR/Pip OFF system necessary for its regulation. The deletion of fasR was accomplished by homologous recombination. After the transformation of M. tuberculosis with plasmid pFR35, the first recombination event was selected on 7H10 medium containing Kan and Hyg, and resistant colonies were screened by colony PCR. The merodiploid strain was generated by the integration of pFR33 at the L5 attB site, and transformants were plated on 7H10 with Hyg, Kan, and Str. Deletion of the wild-type gene by allelic replacement was accomplished by plating bacteria on 7H10 agar plates supplemented with Str and 2.5% sucrose. The HygR, KanR, StrR, and SucroseR colonies were tested by PCR, and the fasR conditional knockdown strain was named MtbΔfasR-cfasR. The generation of a fasR knockout strain (MtbdΔfasR) was obtained by swapping the pFR33 plasmid at the L5 attB site with the pND255 plasmid. Transformants were selected on 7H10 plates containing Hyg. The deletion of fasR was confirmed by Southern blot and colony PCR.

**Southern Blot**

Ten micrograms of genomic M. tuberculosis DNA was digested overnight with an excess of XhoI, and the fragments were separated by electrophoresis in a 0.7% agarose gel. Southern blotting was carried out using the Hybond-N+ membrane (Amersham) and DIG High Prime DNA Labeling and Detection Starter kit II (Roche), following the indications of the manufacturer. The probe consisted of a 594-bp fragment amplified with oligonucleotides Southern_1fasRRTB and Southern_2fasRRTB (Supplementary Table S3), corresponding to the upstream genomic region of fasR.

**Western Blot**

Equal amounts of protein extracts were loaded on 4–12% NuPAGE gels (Invitrogen) and transferred onto nitrocellulose membranes using a dry electrophoresis transfer apparatus (Invitrogen). Membranes were incubated in TBS-Tween blocking buffer (20 mM Tris pH 7.5, 150 mM NaCl, 0.1% Tween 20) with 5% skimmed milk powder for 2 h at 4°C prior to overnight incubation with 1:1000 dilution of monoclonal anti-FasR primary antibody (Alere™). Membranes were washed in TBS-Tween three times at room temperature and then incubated with 1:100000 secondary antibody Anti-Mouse Kappa conjugated to Horseradish Peroxidase (Southern Biotech) for 2 h before washing again. Signals were detected using Chemiluminescent Peroxidase Substrate (Sigma-Aldrich). Monoclonal mouse anti-RpoB antibodies were purchased from NeoClone and used 1:20000.

**RNA Techniques**

RNA was extracted from M. tuberculosis strains grown for 8 days in complete 7H9 media using Trizol phenol based methods (Kolly et al., 2014) and treated, when needed, with RQ1 RNase-Free DNase (Promega). qRT-PCR was performed using second strand cDNA as the template, generated with SuperScript III Reverse Transcriptase (Invitrogen), random primers, and a green fluorochrome as the indicator dye (qPCR master mix, Biodynamics). The expression of fas, fasR, and acpS was quantitated after normalization of RNA levels to the expression of the sigA gene as previously described (Cabruja et al., 2017). qPCR data are presented as relative expression values normalized to sigA and to the wild-type strain H37Rv. qPCR cycling conditions were as follows: 95°C for 2 min followed by 40 cycles of 95°C for 15 s, 58°C for 15 s, and 68°C for 20 s. The sequences of all the primers used are listed in electronic supplementary material (Supplementary Table S3).

**Lipid Analysis**

Cell-associated lipids for lipidomic analyses were prepared as previously described (Cabruja et al., 2017). Briefly, cell pellets of cultures grown at 37°C in Middlebrook 7H9 broth (DIFCO) supplemented with 0.2% glycerol, 0.03% Tyloxapol, and 10% OADC were washed with 50 mM ammonium acetate, pH 7.8, and transferred to glass tubes containing 5 ml CHCl3:CH3OH (2:1, v/v). Samples were incubated overnight at 4°C with gentle agitation. After centrifugation, bacterial pellets were subjected to an additional extraction using CHCl3:CH3OH (1:2, v/v) for 2 h. Organic extracts were pooled and dried under nitrogen at 4°C. Lipids were resuspended in 3 ml of CHCl3, washed with 3 ml of H2O, and the organic phase was transferred to pre-weighted glass tubes, dried under nitrogen at 4°C, and re-weighted on a microbalance. Extracts were dissolved in CHCl3:CH3OH (1:1, v/v) at 1 mg ml−1 and centrifuged at 3000 x g for 5 min. Lipids were analyzed in an Agilent 1200 series HPLC system with a Reprospher (Dr. Maish) 100 C8 column (1.8 µm, 50 mm length, and 2 mm ID). The flow rate was 0.3 ml min−1 in binary gradient mode with the following elution program: the column was equilibrated with
100% mobile phase A [CH₃OH:H₂O (99:1, v/v), containing 0.05 mM AcNH₄], 2 μl of each sample was injected, and the same elution conditions continued for 1 min, followed by a 12-min gradient to 100% mobile phase B [isopropanol:hexane:H₂O (79:20:1, v/v), containing 0.05 mM AcNH₄], and finally holding this condition for 1 min. The column was equilibrated for 2 min with 100% mobile phase A before the injection of the next sample. An Agilent 6500 series Q-TOF instrument with a Dual AJS ESI was used for mass analysis. Ionization gas temperature was maintained at 200°C with a 14 l min⁻¹ drying gas flow, a 35 psig nebulizer pressure, and 3,500 volts. Spectra were collected in positive and negative mode from m/z 115 to 3,000 at 4 spectra s⁻¹. Automatic tandem-MS (MS-MS) was obtained in data-dependent acquisition mode at 15 MS-MS per cycle. Continuous infusion calibrants included m/z 121.051 and 922.010 in positive-ion mode and m/z 119.035 and 955.972 in negative-ion mode. CID-MS was carried out with energy of 50 volts. For the comparative analysis, the column is conditioned by four successive mock injections before randomized QCs and mycobacterial samples are analyzed. Lipid features were annotated based on the m/z database and fragmentation pattern published by Layre et al. (2011). The m/z values were assigned using MycoMass which is a mycobacteria-specific database. The fragmentation pattern of each lipid class was compared to the Supplementary Figure S4 of the mentioned work. All reported compounds were assigned to a specific lipid class based on exact m/z values and retention times. Retention times for all lipid classes were determined by analyzing the MS-MS information of several species within each lipid class.

**Colony-Forming Units (CFU) Analysis**

*M. tuberculosis* H37Rv, *MtbΔfasR*, and *MtbΔfasR-cfasR* strains were used to infect THP-1 cells (MOI = 5). The day before infection, 4 × 10⁶ THP-1 cells were plated in a 24-well plate in RPMI Medium with 4.5 g/L glucose, 10% heat-inactivated fetal calf serum (Internegocios), and 2 mM L-glutamine (full medium) plus 50 nM phorbol 12-myristate 13-acetate (PMA) to allow differentiation of the cells to macrophages. On the infection day, cells were washed and a suspension of the different strains of *M. tuberculosis* in RPMI medium was added. After 2 h of uptake, cells were washed three times with PBS, and fresh full RPMI medium plus gentamicin 10 μg ml⁻¹ was added. Cells were incubated at 37°C in 5% CO₂ atmosphere. Subsequently, at the indicated time points, cells were lysed in sterile water and serially diluted in PBS-Tween 80 0.05%. Dilutions were plated on Middlebrook 7H10 complete agar medium (Difco Laboratories). Colonies were counted after 3–4 weeks to determine CFU.

**Mouse Infections**

Bacillary suspensions of the different strains were adjusted to 100 viable cells in 100 ml phosphate-buffered saline (PBS). Each animal was anesthetized and intratracheally inoculated with *M. tuberculosis* strains (*MtbH37Rv, MtbΔfasR,* and *MtbΔfasR-cfasR*). Infected mice were kept in cages fitted with microisolators connected to negative pressure. Groups of 10 mice were infected with the different *M. tuberculosis* strains. The inoculum doses were determined by enumerating CFU recovered from the lungs of three mice per strain, 24 h post-infection. Seven mice per group were killed at 60 days after infection, and lungs were removed and homogenized. Four dilutions of each homogenate were spread onto triplicate plates of Middlebrook 7H10 complete agar medium. The number of colonies was counted after 3–4 weeks.

**Infection of THP-1 Cells and Indirect Immunofluorescence**

*M. tuberculosis* strains were covalently stained with FITC (Sigma-Aldrich). Briefly, 1 × 10⁶ bacteria were washed twice with 0.1 M PBS and suspended in 1 ml of the same solution. FITC was added to a final concentration of 5 μg/ml and incubated in the dark for 1 h at 37°C. Bacteria were washed gently with PBS until unincorporated dye was eliminated, and were then used to infect THP-1 cells. THP-1 cells were maintained in complete full RPMI medium. The day before infection, 2 × 10⁵ THP-1 cells per well were plated in a 24-well plate with coverslips in full RPMI medium plus 50 nM PMA to allow differentiation of the cells to macrophages. On the day of infection, cells were washed and infected with the different stained *M. tuberculosis* strains (*MtbH37Rv, MtbΔfasR,* and *MtbΔfasR-cfasR*) prepared in RPMI medium, at a MOI of five. After 2 h of uptake, cells were washed and incubated with 10 μM Lysotracker Red DND-99 for 1 h to allow the staining of acidic compartments, fixed with 4% paraformaldehyde solution in PBS for 30 min and subjected to immunofluorescence analysis.

Cells were quenched by incubation with 50 mM NH₄Cl solution in PBS for 30 min. and permeabilized with 0.05% saponin/1% BSA in PBS for 15 min, followed by overnight incubation with goat anti-LAMP-3 (Santa Cruz Biotechnology, Santa Cruz, CA) primary antibody diluted 1:50 in PBS. Secondary anti-goat antibody conjugated to Cy3 (Jackson Immuno Research Labs, Inc.) was diluted 1:600 in PBS. Cells were washed and incubated with 10 μM Lysotracker Red DND-99 or LAMP-3 association of the cells to macrophages. The day before infection, cells were washed and infected with the different stained *M. tuberculosis* strains (*MtbH37Rv, MtbΔfasR,* and *MtbΔfasR-cfasR*) prepared in RPMI medium, at a MOI of five. After 2 h of uptake, cells were washed and incubated with 10 μM Lysotracker Red DND-99 for 1 h to allow the staining of acidic compartments, fixed with 4% paraformaldehyde solution in PBS for 30 min and subjected to immunofluorescence analysis. Cells were incubated with 50 mM NH₄Cl solution in PBS for 30 min. and permeabilized with 0.05% saponin/1% BSA in PBS for 15 min, followed by overnight incubation with goat anti-LAMP-3 (Santa Cruz Biotechnology, Santa Cruz, CA) primary antibody diluted 1:50 in PBS. Secondary anti-goat antibody conjugated to Cy3 (Jackson Immuno Research Labs, Inc.) was diluted 1:600 in PBS. Cells were mounted with mounting medium (Dako, Denmark) and analyzed by confocal microscopy using an SP5 AOBs confocal microscope (Leica Microsystems, Germany). Mycobacterial internalization was monitored by following FITC fluorescence (green). Lysotracker Red DND-99 or LAMP-3 association of the cells to macrophages. The experiments were performed in duplicates in three independent experiments. Data were analyzed using the Brown-Forsythe and Welch ANOVA tests with Games-Howell's multiple comparisons test.

**RESULTS**

**Construction and Characterization of *M. tuberculosis fasR* Mutant Strain**

To investigate the function of FasR in *M. tuberculosis*, and considering that *fasR* is essential for *M. smegmatis* viability,
we first constructed an *M. tuberculosis* conditional knockdown strain in fasR (Figure 1A). Plasmids pFR33, providing a second copy of fasR under the control of the TET-PIP OFF system at the chromosomal L5 phage attachment site attB (Boldrin et al., 2010) and pGA80, were simultaneously transformed in *M. tuberculosis* (Supplementary Table S1). Disruption of fasR was then obtained by two-step allelic exchange using pFR35, which deleted the gene at the native locus. Homologous recombination was confirmed by colony PCR and Southern blot analysis, and the resulting conditional knockdown strain was named MtbΔfasR-cfasR (Figure 1B). In this strain, expression of fasR was down-regulated by addition of anhydrotetracycline (ATc) to the culture medium, thus leading to protein depletion. However, we were not able to detect any growth defect after addition of ATc, suggesting that fasR was not essential for *M. tuberculosis* survival. This result prompted us to try to obtain a fasR knockout (KO) mutant strain by replacing the fasR complementing plasmid with the empty vector pND255. Consistent with the ability of the conditional mutant to grow in the non-permissive condition, plenty of colonies were obtained in the presence of Hyg, and the replacement was successfully confirmed by colony PCR (Supplementary Figure S1A). We selected two Hyg-resistant clones and confirmed the depletion of FasR by Western blot using total protein extracts (Supplementary Figure S1B). The genotype of this strain was further analyzed by Southern Blot (Figure 1B), thus validating the genomic deletion of fasR and the generation of the fasR KO mutant strain, named MtbΔfasR.

We have previously demonstrated that FasR is a transcriptional activator of fas in *M. smegmatis* (Mondino et al., 2013). Thus, to validate the role of FasR in fatty acid biosynthesis in *M. tuberculosis*, we analyzed expression of the fas-acpS operon genes in MtbΔfasR and in the complemented strain MtbΔfasR-cfasR. The relative amounts of fas and acpS mRNAs were measured by quantitative RT-PCR after 8 days of growth in complete 7H9 medium and compared to those of MtbH37Rv wild-type strain. As shown in Figure 1C, transcription of fas and acpS was reduced by 70 and 40%, respectively, when cells were deprived of FasR, confirming that FasR is a transcriptional activator of the fas-acpS operon in *M. tuberculosis*.

Once the non-essentiality of fasR for *M. tuberculosis* survival was confirmed, we then hypothesized that deletion of this transcriptional regulator could be deleterious for bacterial growth under specific conditions. In particular, we analyzed the growth of the MtbΔfasR mutant using different culture media. As shown in Figure 2A, growth of MtbΔfasR in complete 7H9 medium (7H9 + OADC) was similar to that of the wild type and the complemented strain (MtbΔfasR-cfasR grown without ATc), although the KO mutant reached stationary phase earlier and at lower OD₆₀₀. Interestingly, if the medium was not supplemented with oleate and BSA (i.e., 7H9 without OADC), growth defects appeared to be more pronounced, suggesting that the presence of lipids in complete 7H9 medium could partially rescue the absence of FasR (Figure 2B). Moreover, the MtbΔfasR mutant showed severe growth defects in Sauton minimal medium, reaching a maximal OD₆₀₀ of 0.3 (Figure 2C). Altogether, these experiments showed that FasR is not essential for *M. tuberculosis* survival, although it is required for optimal bacterial growth, especially under lipid-limited conditions.

**FIGURE 1** | Construction and characterization of *M. tuberculosis* fasR mutant. (A) Schematic representation of the genetic organization of the chromosomal region of fasR in the wild-type H37Rv strain (MtbH37Rv), fasR conditional knockdown strain (MtbΔfasR-cfasR), and fasR KO strain (MtbΔfasR). (B) Southern blot analysis of the different strains: chromosomal DNA was digested with XhoI and probed for hybridization with the probe indicated with a black bar, corresponding to the 5' end of the Rv3207 gene. (C) Relative expression levels of fasR, fas, and acpS mRNAs measured by quantitative RT-PCR. Values were normalized to sigA and to the wild-type strain (MtbH37Rv). Samples for RNA extraction were collected at OD₆₀₀≈0.3. qPCR reactions were performed in technical duplicates, and three independent biological replicates were used for quantitative analysis.
fasR Knockout Strongly Impacts Composition of M. tuberculosis Cell Envelope

Previous characterization of M. smegmatis fasR (Mondino et al., 2013) and fas (Cabruja et al., 2017) conditional mutant strains revealed that reduced FAS I levels had a strong impact on fatty acid and phospholipid biosynthesis. Interestingly, MA remained being synthesized in the FAS I mutant for several hours after addition of ATc (non-permissive conditions), although with different abundance of the various species (Cabruja et al., 2017). Considering the relevance that lipids have in M. tuberculosis survival and virulence, the impact of the absence of FasR on the global lipid profile of M. tuberculosis was further studied. We performed comparative lipidomic analyses between Mtb H37Rv, Mtb ΔfasR, and the complemented strain (Supplementary Table S4). Briefly, we extracted cell-associated lipids as previously described (Cabruja et al., 2017) and analyzed the lipid content by high-performance liquid chromatography-mass spectrometry with data-dependent tandem MS (HPLC-MS/MS).

As illustrated in Figure 3, lipid analysis in the negative mode indicated that the relative abundance (expressed as percentage of lipid MS signals) of phospholipids (PL) and MA in the wild-type and complemented strains was approximately 75 and 25%, respectively. These values changed to 50% for both PL and MA in the fasR mutant strain, reflecting the strong impact that lower levels of de novo fatty acid biosynthesis had in PL and MA relative abundance. The relative composition of PL also showed a biased effect towards phosphatidylinositol (PI) and cardiolipins (CL), which were significantly reduced in the mutant strain (Figure 3; Supplementary Figure S2). Furthermore, there was a strong reduction in the content of sulfolipids (Figure 3; Supplementary Figure S2), and the acyl chain substitutions of this lipid were consistently shorter in the mutant strain (Supplementary Figure S3). A relevant change observed in the qualitative composition of the PL was the significant increase in the double bonds of the acyl residues in the mutant strain, with those of the CL, phosphatidylglycerol (PG), and PI being the most significant (Supplementary Figure S4).

Additionally, results obtained in the positive mode analysis showed that the M. tuberculosis Mtb ΔfasR mutant strain displayed a significant reduction in the relative content of PDIM (Figure 4; Supplementary Figure S2), with acyl substitutions that are shorter in the mutant strain, compared with the wild-type or the complemented strain (Supplementary Figure S3). Most of the changes observed in the deletion mutant were reversed in the complemented strain Mtb ΔfasR-cfasR, indicating that fasR is needed to maintain the global lipid composition of M. tuberculosis cell envelope.

FasR Is Required for M. tuberculosis Replication in Macrophages

To investigate whether FasR is required for the intracellular survival of M. tuberculosis in macrophages, we infected THP-1-derived macrophages with exponentially growing cultures of Mtb H37Rv, Mtb ΔfasR, and the complemented strain Mtb ΔfasR-cfasR. To quantify bacterial uptake and intracellular replication, we determined the number of intracellular viable bacteria at different time points post-infection. As shown in Figure 5A, infection with a similar number of bacteria of each strain resulted in recovery of a similar number of CFUs for all of the strains at 24 and 48 h post-infection. However, 120 and 144 h after the infection, intracellular bacterial replication of Mtb ΔfasR strain was significantly reduced compared to that...
**FIGURE 3** | Lipid profile in negative ion mode. LC-MS analyses (negative mode) of the phospholipids and mycolic acids present in *M. tuberculosis* wild type strain (*Mtb* H37Rv), fasR complemented strain (*MtbΔfasR-cfasR*), and fasR KO strain (*MtbΔfasR*). The values indicated represent the relative abundance of the MS signals corresponding to the different lipid classes found in the samples. Results are the means of three independent experiments. CL, cardiolipin; PI, phosphatidylinositol; PIM, phosphatidylinositol mannosides; PG, phosphatidylglycerol; PE, phosphatidylethanolamine; SL, sulfolipids; MA, mycolic acids.

**FIGURE 4** | Lipid profile in positive ion mode. LC-MS analyses (positive mode) of lipids present in *M. tuberculosis* wild type strain (*Mtb* H37Rv), fasR complemented strain (*MtbΔfasR-cfasR*), and fasR KO strain (*MtbΔfasR*). The values indicated represent the relative abundance of the MS signals corresponding to the different lipid classes found in the samples. Results are the means of three independent experiments. TAG, triacylglycerol; DAG, diacylglycerol; MMDAG, monomycolyl DAG; PDIM, phthiocerol dimycocerosates; GroMM, glycerol monomycolate; TMM, trehalose monomycolate.
of the wild-type strain. The wild-type phenotype was partially restored in the MtbΔfasR-cfasR complemented strain at day 5 (120 h) and completely restored at day 6 (144 h), indicating that the intracellular growth defect was due to the deletion of fasR and that this regulator is necessary for optimal intracellular multiplication of M. tuberculosis in macrophages.

**FasR Regulates Phagosome Maturation in Macrophages**

We have shown that the absence of FasR and, consequently, reduced levels of fas-acps have a strong impact on the composition of the M. tuberculosis cell envelope, and also on bacterial multiplication within macrophages. We hypothesized that the altered bacterial cell envelope of the MtbΔfasR mutant may affect M. tuberculosis-driven phagosome maturation during infection. To investigate this, we analyzed the acidification of the M. tuberculosis MtbΔfasR phagosome during infection of human macrophages by staining with Lysotracker dye and measuring the phagosomal acquisition of LAMP-3 (late endosomal/lysosomal markers). Interestingly, while MtbH37Rv prevented phagosome acidification, the MtbΔfasR mutant strain resided in phagosomes that were strongly stained by Lysotracker dye and that acquired LAMP-3, suggesting that this strain cannot avoid phagosomal maturation. The observed mutant phenotype was rescued in the MtbΔfasR-cfasR complemented strain, indicating that the increased phagosome acidification was due to the absence of FasR (Figures 5B,C).

**Disruption of Lipid Homeostasis Attenuates Mycobacterial Virulence in Mice**

To characterize the role of FasR in an experimental model of infection, three groups of BALB/c mice (10 per group) were infected intratracheally with 10^2 CFU of MtbH37Rv, MtbΔfasR, and MtbΔfasR-cfasR strains. Mice were killed 60 days post-infection, and bacterial loads in lungs were determined. CFU counts were significantly reduced for the MtbΔfasR strain compared to MtbH37Rv, while the wild-type phenotype was restored in the MtbΔfasR-cfasR strain (Figure 6), indicating that FasR is essential for survival and multiplication of M. tuberculosis in mice.

**DISCUSSION**

In a previous investigation we identified FasR, a TetR-like transcriptional regulator that specifically binds to the fas promoter region to activate its expression, and found that
FasR was essential for in vitro growth of M. smegmatis (Mondino et al., 2013). However, its role in M. tuberculosis viability was controversial. Using Himar 1 transposon mutagenesis, it was initially proposed that FasR was essential for M. tuberculosis survival (Sassetti et al., 2003; Griffin et al., 2011). However, more recently, De Jesus et al. found that FasR was dispensable for growth, by assessing the composition of a highly saturated library of transposon mutants of M. tuberculosis by deep sequencing (DeJesus et al., 2017). Moreover, the relevance of FasR for mycobacterial survival in intracellular environments or animal models had not been investigated. In this work, by constructing a deletion for M. tuberculosis viability, although it is required for the optimal growth of M. tuberculosis in vitro and for virulence in macrophages and the mice model of infection.

We had shown that FasR binds to three inverted repeats present in the promoter region of the fas-acpS operon, thereby activating fatty acid biosynthesis in M. smegmatis (Mondino et al., 2013). Furthermore, we also demonstrated, in vitro and in vivo, that FasR binding to its target promoter is regulated by long-chain acyl-CoAs, the main products of FAS I (Mondino et al., 2013). The fas and acpS genes form a bicistronic operon, coding for the FAS I synthase and the 4-phosphopantetheinyl transferase, respectively, the latter being an essential enzyme to produce functional acyl carrier protein (holo ACP), central to all fatty-acid biosynthesis systems. Transcriptional studies carried out in this work showed that expression of the fas-acpS operon genes in M. tuberculosis ΔfasR was reduced by 70% and 40%, respectively (Figure 1C). This result confirmed that FasR is a transcriptional activator of the fas-acpS operon and is in agreement with the results observed in the M. smegmatis fasR knockdown strain, where 90% reduction of fasR mRNA levels led to 40% reduction in the expression of the fas-acpS operon (Mondino et al., 2013). Moreover, we have previously shown that in the M. smegmatis fasR mutant, a 90% reduction in fas mRNA levels led to reduced FAS I activity (Cabruja et al., 2017). Therefore, we can anticipate a similar reduction in FAS I activity in the M. tuberculosis fasR mutant, although further experiments are required to unequivocally demonstrate this. The activator nature of FasR highlights a remarkable difference with most transcriptional regulators of fatty acid biosynthesis, which are repressor proteins (Zhu et al., 2009). Moreover, FasR is a member of the TetR family of regulators, and the vast majority of these proteins are transcriptional repressors, with few acting as activators (Cuthbertson and Nodwell, 2013).

Since fatty acids are major constituents not only of phospholipids but also of several complex lipids, inactivation of FasR was expected to modify the synthesis of several components of the cell envelope essential for mycobacterial viability and/or virulence. Indeed, lipidomic analysis of MtbΔfasR and comparison with wild-type and complemented strains proved this hypothesis. As expected, reduced levels of fatty acid biosynthesis in MtbΔfasR had a strong impact on relative abundance of phospholipids and their relative and qualitative composition (Figure 3; Supplementary Figure S2). Phosphatidylinositol (PI) and cardiolipins (CL) were significantly reduced in the mutant strain, while the acyl chains present in these phospholipids exhibited a significant increase in their double bond content (Supplementary Figure S4). Mycobacterial phospholipids have been implicated in immunomodulation. For example, CL is processed into lysocardiolipin by the lysosomal phospholipase A2 during infection, probably generating active lipid derivatives that play important roles during infection or persistence within the host (Fischer et al., 2001). The MtbΔfasR mutant strain also displayed a strong reduction in the content of sulfolipids (Figure 4; Supplementary Figure S2) and PDIM (Figure 4; Supplementary Figure S2). Interestingly, the acyl chain substitutions of both lipids were consistently shorter compared to the wild type MtbH37Rv strain (Supplementary Figure S3). On this regard, a M. tuberculosis mutant strain deficient in sulfolipids
and PDIM has been shown to be attenuated in macrophages and in mice models of infection (Passemar et al., 2014). Moreover, during infection, genes involved in the biosynthesis of sulfolipids and PDIM are up-regulated (Graham and Clark-Curtiss, 1999; Rodriguez et al., 2013) and the mass of these lipids increases (Yang et al., 2009; Griffin et al., 2012). Altogether, our results showed that *M. tuberculosis fasR* mutant exhibits an extensive rearrangement of the lipid components of the cell envelope, which are reversed in the complemented strain *MtbΔfasR-cfasR*, confirming that FasR is needed to maintain the lipid homeostasis of the *M. tuberculosis* cell envelope. As the mycobacterial cell envelope is the interface with the host, FasR modulation of its structure and consequent responsiveness to immune factors might be important in determining the state of an infection.

Consistent with this hypothesis, we have shown that FasR is necessary for optimal intracellular multiplication of *M. tuberculosis* in macrophages, therefore confirming the role of cell wall components in the ability of the pathogen to survive and multiply intracellularly (García-Vilanova et al., 2019). In addition, the lack of FasR impaired the normal *in vitro* replication of *M. tuberculosis* in minimal medium devoid of lipids. This finding indicates that FasR is a crucial regulator of lipid metabolism in *M. tuberculosis* rather than a virulence factor. Persistence of the tubercle bacillus inside cells relies upon the inhibition of phagosome maturation and acidification (Rohde et al., 2007). In this context, several cell wall lipids have been shown to modulate host endocytic pathways. This is the case for Man-Lam, TDM, PIM, and PDIM, which, besides their role in phagocytosis, also mediate intracellular trafficking and vacuole maturation arrest induced by *M. tuberculosis* (Passemar et al., 2014). More recently, PDIM and SL were shown to be involved in controlling autophagy-related pathways in human macrophages (Bah et al., 2020). Considering the changes observed in the PDIM content of the *MtbΔfasR* mutant, we analyzed the acidification of phagosome during infection of human macrophages. Interestingly, while the wild-type *MtbH37Rv* and the complemented strains prevented phagosome acidification, the *MtbΔfasR* mutant was impaired in avoiding phagosomal maturation, clearly indicating that increased phagosomal acidification was due to absence of FasR (Figure 5). These results suggested that although FasR is not essential for *in vitro* growth, regulation of lipid biosynthesis, mediated by FasR, is critical for macrophage infection. Moreover, using a mouse model of infection we demonstrated the essentiality of FasR for *M. tuberculosis* virulence *in vivo* (Figure 6), most probably as a consequence of the loss of various mycobacterial lipids that contribute to pathogenicity.

Co-evolution with the human immune system prompted *M. tuberculosis* to generate exquisite mechanisms of immunomodulation and immune evasion, many of which depend on its cell envelope. In fact, a comparison of cell envelope components between *M. tuberculosis* and ancestral species revealed a restructuring of the mycomembrane glycolipid components, including mycolic acids and free lipids of the outer leaflet (Dulberger et al., 2020). The observed phenotype of the FasR mutant is probably a consequence of additive effects, as several components of the cell envelope were altered. However, we cannot rule out that other general metabolic changes non-related with lipid homeostasis may cooperate with the observed phenotype. *M. tuberculosis* interferes with innate immune signaling pathways, for example, by masking pathogen-associated molecular patterns with PDIM (Cambier et al., 2014) and inhibiting toll-like receptor 2 with sulfolipidolipids (Blanc et al., 2017). PDIM is also crucial for other steps of infection: it contributes to cell envelope properties, such as the permeability barrier (Camacho et al., 2001), improves the invasiveness of *M. tuberculosis* and its capacity to arrest phagosome maturation, possibly by inducing changes in the plasma membranes of human macrophages during infection (Astarie-Dequeker et al., 2009), and directly protects against reactive nitrogen species (Rousseau et al., 2004). Therefore, the reduced content of these molecules (and/or changes in their chemical composition) may contribute to sensitivity to the bactericidal activities of phagocytes.

Altogether, our data indicate an important role for FasR in *M. tuberculosis* pathogenesis and encourage further investigations into the contribution of lipid regulation to virulence. It is well known that *M. tuberculosis* is able to modulate the expression of its cell wall components in response to environmental changes; however, little is known about their reorganization during infection and, to a lesser extent, about the regulatory mechanisms involved in this process (Dulberger et al., 2020). The dynamism of the bacterial cell envelope should be a tightly regulated phenomenon, ensuring *M. tuberculosis* survival during infection. Thus, specific regulatory components involved in this process, as we have demonstrated for FasR, would constitute interesting targets for the development of new treatment strategies.

Along this line of thinking, we recently solved the crystal structures of *M. tuberculosis* FasR in complex with DNA and the acyl effector ligands, which provided insights into the molecular sensory and transmission mechanisms of this regulatory protein (Lara et al., 2020). This knowledge could lead now to perform structure-guided drug discovery strategies to identify antimycobacterial molecules with novel mechanisms of action.

**DATA AVAILABILITY STATEMENT**

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

**ETHICS STATEMENT**

All animal experiments were performed in compliance with the regulations of the Institutional Animal Care and Use Committee (CICUAE) of INTA with the authorization Number 20/2016.

**AUTHOR CONTRIBUTIONS**

GG and HG contributed to conception and design of the study. SM, CV, MC, GS, and FCB performed the experiments. SM, CV, and MC prepared the figures. SM and CS constructed and analyzed mutant growth. CV and FCB performed infection experiments.
FB supervised and coached infection experiments. MC performed and analyzed the HPLC-MS/MS measurements for lipid analysis and AC-G, MW, and GG supervised the resulting data. HG, FB, SC, CS, and GG supported the design of the study. All authors contributed to the interpretation of the data. GG designed the study, supervised the experimental work, and wrote the final version. All authors contributed to manuscript revision, read, and approved the submitted version.

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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2020.586285/full#supplementary-material

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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