Abstract: Lipoproteins are important components of the mycobacterial cell envelope due to their function in cell wall homeostasis and bacterial virulence. They are post-translationally modified with lipid- and glycosyl-residues in various species and interference with acylation or glycosylation leads to reduced growth and attenuated virulence in Mycobacterium tuberculosis. Lipoproteins are also expressed in the emerging and highly drug resistant pathogen Mycobacterium abscessus which frequently affects the lungs of patients with chronic pulmonary disease or cystic fibrosis. We investigated post-translational modification, acylation and glycosylation, of heterologously expressed (M. tuberculosis LppX and Mpt83) and endogenous (SodC) lipoproteins at the molecular level in M. abscessus and identified MAB_{122} as protein O-mannosyltransferase (Pmt). Both, heterologous and endogenous lipoproteins carried characteristic lipid anchors with palmitic acid (C16), palmitoleic acid (C16 : 1), oleic acid (C18), or tuberculostearic acid (C19) modifications. Multiple hexose moieties were detected in the N-terminal region of the model lipoproteins expressed in M. abscessus. Conservation of lipoproteinglycosylation motifs for other regulatory mechanisms regarding this post-translational modification. Deletion of MAB_{122} translational modification of lipoproteins and the permeability of the mycobacterial cell envelope which stresses the importance.

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Lipoprotein Glycosylation by Protein-O-Mannosyltransferase (MAB_1122c) Contributes to Low Cell Envelope Permeability and Antibiotic Resistance of Mycobacterium abscessus

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Lipoproteins are important components of the mycobacterial cell envelope due to their function in cell wall homeostasis and bacterial virulence. They are post-translationally modified with lipid- and glycosyl-residues in various species and interference with acylation or glycosylation leads to reduced growth and attenuated virulence in Mycobacterium tuberculosis. Lipoproteins are also expressed in the emerging and highly drug resistant pathogen Mycobacterium abscessus which frequently affects the lungs of patients with chronic pulmonary disease or cystic fibrosis. We investigated post-translational modification, acylation and glycosylation, of heterologously expressed (M. tuberculosis LppX and Mpt83) and endogenous (SodC) lipoproteins at the molecular level in M. abscessus and identified MAB_1122c as protein O-mannosyltransferase (Pmt). Both, heterologous and endogenous lipoproteins carried a characteristic lipid anchor with palmitic acid (C16), palmitoleic acid (C16:1), oleic acid (C18), or tuberculostearic acid (C19) modifications. Multiple hexose-moieties were detected in the N-terminal region of the model lipoproteins expressed in M. abscessus.

Conservation of lipoprotein glycosylation in M. tuberculosis and M. abscessus was revealed and points toward the existence of an O-glycosylation motif or other regulatory mechanisms regarding this post-translational modification. Deletion of MAB_1122c prevented glycosylation and affected susceptibility to specific antibiotics which are large or target peptidoglycan synthesis and to lysozyme. Cell envelope permeability of M. abscessus Δpmt was increased and mutant bacteria showed reduced survival inside macrophages. The results provide a link between post-translational modification of lipoproteins and the permeability of the mycobacterial cell envelope which stresses the importance of lipoproteins as components of this complex structure.

Keywords: lipoprotein, glycosylation, Mycobacterium abscessus, post-translational modification, virulence, drug susceptibility
INTRODUCTION

The cell envelope of mycobacteria is comprised of a plasma membrane, the periplasm and a lipid-rich mycobacterial outer membrane which contains arabinogalactan and mycolic acids. This thick and low-permeable structure contributes to persistence and high intrinsic drug resistance of mycobacteria (Brennan and Nikaido, 1995). Lipoproteins are important components of the mycobacterial cell envelope and in mycobacterial physiology since they fulfill various functions in membrane homeostasis, transmembrane signaling and adhesion to host cells (Sutcliffe and Harrington, 2004; Becker and Sander, 2016). In *Mycobacterium tuberculosis*, approximately 100 putative lipoproteins are expressed. They have been identified as immunogenic antigens (for review see Becker and Sander, 2016) and virulence factors since interruption of the lipoprotein synthesis pathway led to increased killing of bacilli inside macrophages and improved survival of infected mice (Sander et al., 2004; Rampini et al., 2008). LpqW for instance plays a role in synthesis of crucial lipoglycans, the lipoarabinomannans (Rainczuk et al., 2012) and RptB (resuscitation promoting factor B) is involved in cleavage of peptidoglycan during resuscitation (Russell-Goldman et al., 2008). LppX mediates the transport of complex lipids, the phthiocerol-dimycocerosates, to the outer membrane where these serve as virulence factors (Suñenbacher et al., 2006). Mpt83 is described as an adhesion factor (Carr et al., 2006). Lipoproteins are synthesized from pre-pro-lipoproteins throughout the consecutive action of diacylglyceroltransferase (Lgt), pro-lipoprotein signal peptidase (LspA) and N-acyltransferase (Lnt) (Tschumi et al., 2009, 2012; Brulle et al., 2013). These membrane-bound enzymes assemble the N-terminal lipid anchor, comprised of a diacylglycerol- and an N-acyl moiety, onto the conserved cysteine in the characteristic lipobox (LVI/ASTVI/GAS/C) (Kovacs-Simon et al., 2011).

Next to their three acyl moieties, some mycobacterial lipoproteins also carry glycosyl-residues. Protein glycosylation in *M. tuberculosis* is mediated by two enzymes, polyrenol-monophosphomannose-synthase (Ppm1) and protein-O-mannosyltransferase (Pmt). A mannose residue is transferred from the donor GDP-mannose to a polyrenol phosphate carrier in the plasma membrane by Ppm1. The mannosylated lipid carrier flips across the membrane so that the hexose is exposed at the periplasmic site. There, Pmt catalyzes the transfer of the mannose residue to a serine or threonine in the polypeptide chain (VanderVen et al., 2005). In mycobacteria to date only O-linked mannosylation has been described and the modification has been shown to be Sec-export dependent (Michell et al., 2003; VanderVen et al., 2005). Pmt might also be responsible for glycosylation of non-lipoproteins but it is clearly dispensable for synthesis of other crucial cell wall components like lipomannans or lipoarabinomannans (Horn et al., 1999; Liu et al., 2013). Lipoproteins are considered particularly important substrates of Pmt with regard to cell envelope homeostasis. In this study we investigate the effects of protein glycosylation deficiency on the mycobacterial cell envelope and therefore we focus on the lipoprotein glycosylating function of Pmt. In *M. tuberculosis*, *Mycobacterium smegmatis* or the closely related *Corynebacterium glutamicum* the lipoproteins LppX, Mpt83, LprI, LpqH, LprG, and SodC have been found with glycosyl-residues (Michell et al., 2003; Sartain and Belisle, 2009; Mohiman et al., 2012; Sethi et al., 2016; Alonso et al., 2017; Parra et al., 2017). The emerging pathogen *Mycobacterium abscessus*, a highly drug resistant fast-growing mycobacterium which infects the lungs of patients with chronic pulmonary disease or cystic fibrosis, also localizes lipoproteins in its cell envelope (Ripoll et al., 2009; Nessar et al., 2012). The bacteria’s capability to post-translationally modify these proteins and the effect of lipoprotein modification on *M. abscessus* homeostasis and virulence have not been investigated yet.

Here, we elucidated lipoprotein modification in *M. abscessus* at the molecular level. We characterized lipidation and glycosylation of heterologously expressed *M. tuberculosis* and endogenous *M. abscessus* lipoproteins by mass-spectrometry and revealed conservation of O-glycosylation throughout fast- and slow-growing mycobacterial species. The observation points to a universal but to date unknown motif for this post-translational modification. We identified MAB_1122c as Pmt, the lipoprotein glycosylating enzyme in *M. abscessus*. The role of lipoprotein glycosylation in *M. abscessus* cell envelope homeostasis, antibiotic resistance, and virulence in an in vitro model was assessed using a MAB_1122c deletion mutant. We show that lipoprotein glycosylation deficiency increases permeability of the *M. abscessus* cell envelope and alters susceptibility to large and bulky compounds and to antibiotics acting on the peptidoglycan layer. Furthermore, intracellular survival of glycosylation deficient *M. abscessus* is compromised.

MATERIALS AND METHODS

**Growth of Bacterial Strains**

*Mycobacterium abscessus* was grown at 37°C with daily manual shaking in LB or Middlebrook 7H9-OADC-Tween supplemented with apramycin (apra, 50 µg/ml), kanamycin (kan, 50 µg/ml), or isoniazid (32 µg/ml) if required. Growth was monitored by measuring OD 600 and bacteria were transformed by electroporation at 2.5 kV, 25 µF, 1000 Ω as previously described (Rominski et al., 2017a). Genomic DNA was isolated as described by Sander et al. (1995). For drug susceptibility assays cation-adjusted Müller-Hinton (CAMH) medium was used.

**Generation of M. abscessus ΔMAB_1122c**

For deletion of the putative protein-O-mannosyltransferase MAB_1122c, 1.5 kb upstream and downstream fragments of MAB_1122c were amplified from *M. abscessus* genomic DNA using primers 1122c_up_F/1122c_up_R and 1122c_down_F/1122c_down_R (see Supplementary Table S1 for list of primers). Amplified fragments were cloned stepwise into the pSE-apra-KatG vector (Rominski et al., 2017c) using PscI/KpnI and KpnI/XbaI linkers for the up and down fragment, respectively. After transformation of the pSE-MAB1122c-KO vector into *M. abscessus*, selection with apramycin (positive) and isoniazid (negative) led to the unmarked deletion of...
MAB_1122c. Southern Blot analyses with probes I and II that were amplified from M. abscessus genomic DNA using primers PI_1122c_F/PI_1122c_R and PII_1122c_F/PII_1122c_R, respectively, verified recombination. For complementation, the MAB_1122c gene including its native promoter region was amplified from M. abscessus genomic DNA with primers C_1122c_F and C_1122c_R and inserted into the pMV361-kan vector via KpnI thereby generating pMV361-MAB1122c. Integration of the complementation vector in the M. abscessus Δpmt genome was confirmed by Southern Blot.

Lipoprotein Expression

Genes encoding LppX (Rv2945c; 702 bp) and Mpt83 (Rv2873c; 663 bp) were amplified from M. tuberculosis genomic DNA with primers which introduce C-terminal, thrombin cleavable His- and HA- tags and allow fusion to selected promoters (LppX_F/LppX_R; Mpt83_F/Mpt83_R) (Supplementary Table S1). The 19 kDa and the pfbpB promoter were amplified from M. tuberculosis genomic DNA with primers 19 kDa_F/19 kDa_R and pfbpB_F/pfbpB_R. A fusion PCR with primers 19 kDa_F/LppX_R as well as pfbpB_F/Mpt83_R was performed to link the LppX and Mpt83 genes to the 19 kDa and pfbpB promoter, respectively. The fused pfbpB-Mpt83 sequence was cloned into the polyG-apra vector via HindIII and PstI. The fused 19 kDa-LppX sequence was first integrated into the pMV261-gentamicin vector then excised into the polyG-apra vector with HindIII and PstI. Gibson cloning (Bordat et al., 2015) was applied to insert endogenous M. abscessus SodC (MAB_4184c) into the polyG-apra vector. The gene (716 bp) was amplified from M. abscessus genomic DNA with primers SodC_F and SodC_R, introducing a Thrombin-cleavable His- and HA-Tag as well. The polyG-apra vector including the pfbpB promoter was amplified in two separate fragments from polyG-pfbpB-Mpt83 with primers polyG_a_F/polyG_a_R and polyG_b_F/polyG_b_R. Assembly was performed as described by Bordat et al. (2015).

Upon transformation of polyG-apra-LppX, polyG-apra-Mpt83, and polyG-apra-SodC into M. abscessus strains (see list of all strains used in this Supplementary Table S2), clones were selected on LB plates supplemented with apramycin and tested for protein expression by Western Blot. In brief, cells were harvested from liquid culture and disrupted with acid washed glass beads (≤106 µm, Sigma-Aldrich, Switzerland) in the FastPep system (4x 1 min at 6 m/s; MP Bio, United States). The lysate was loaded on a 12.5% Tris-HCl SDS Page gel (Criterion, BioRad, Switzerland) and proteins were detected with primary and secondary goat anti-mouse-HRP (1:20,000, LabForce, Switzerland) antibodies.

Protein Purification

Mycobacterium abscessus strains constitutively expressing LppX, Mpt83, or SodC were harvested and disrupted by FastPrep. Lysates were incubated with 2% N-Lauroylsarcosine sodium salt (SLS; Sigma-Aldrich, Switzerland) for 1 h at room temperature and kept at 4°C overnight. Ultracentrifugation for 1 h at 100,000 × g was applied to separate membrane fragments and unbroken cells from the solubilized lipoproteins. The lipoprotein containing supernatant after ultracentrifugation was diluted with dilution buffer (20 mM NaH₂PO₄, 0.5 M NaCl, 0.2% SLS, pH = 7.4) in a 1:1 ratio and loaded onto a Ni²⁺-column (His-Trap, GE Healthcare; Germany). Proteins were gradually eluted with elution buffer (20 mM NaH₂PO₄, 0.5 M NaCl, 0.2% SLS, 0.5 M imidazole, pH = 7.4) by fast-protein-liquid-chromatography (FPLC) using the ÄKTA Prime System (Amersham Bioscience, United Kingdom).

Mass-Spectrometry

MALDI-TOF

Lipoprotein containing fractions after FPLC were precipitated with trichloroacetic acid (TCA) and separated on a 12.5% Tris-HCl SDS Page gel (Criterion, BioRad, Switzerland). Lipoprotein bands visualized with 0.3 M copper-chloride staining were excised and in-gel protein digestion with AspN was performed. Peptides were treated according to the Ujihara method as previously described (Ujihara et al., 2008) and samples were analyzed on an Ultraflex II MALDI-TOF/TOF instrument (Bruker, Germany) as noted by Tschumi et al. (2009). A mass tolerance of m/z±1 was applied for the selection of modified peptide and fragment ions.

LC-ESI-MS

Pure fractions of lipoproteins were precipitated with TCA and resuspended in NH₄HCO₃ buffer (pH = 8.3) supplemented with 15% methanol and 1% dithiothreitol (DTT). Disulfide bonds were reduced with DTT at 50°C for 30 min and cysteines were alkylated by addition of 20 mM iodoacetamide at room temperature in the dark. Lipoproteins were digested with trypsin or AspN overnight at 37°C. Sample clean-up was performed with C₁₈ ZipTips (Merck, Germany). Glycosylated peptide ions were identified by LC-ESI-MS/MS analysis with an Orbitrap Fusion mass-spectrometer (Thermo Fisher, Germany). Global settings included the method duration of 110 min and a spray voltage of 2000–2300 V for the positive ion and 600 V for the negative ion. MS-data were detected in the Orbitrap at a resolution of 120 k. The scan range was set to m/z = 400–2000 and the maximum injection time was 50 ms. The AGC Target value was 400,000. Peptide ions with a charge state of 2–7 were included and dynamic exclusion for 5 s after the first measurement was applied. A mass tolerance of 10 ppm was allowed. The minimum and maximum intensity thresholds were set to 50,000 and 1 × 10²⁰, respectively. MS/MS-data were obtained upon higher-energy collision dissociation (HCD) fragmentation. Collision Energy was set to 30% and fragment ions were detected in the Orbitrap at a resolution of 120 k as well. The maximum injection time for MS/MS was 250 ms. The AGC Target value was set to 200,000 and one microscan was performed. Results were interpreted with Mascot using a database of mycobacterial protein sequences and allowing two missed cleavages and a mass tolerance of 0.04 Da for MS/MS-data. Carbamidomethyl (C) as a fixed modification and Gln→pyro-Glu (N-term Q), Oxidation (M), Hex-CID/HCD (S), and Hex-CID/HCD (T) as variable modifications were selected. Identified glycopeptides were confirmed by manual analyses of
Minimal Inhibitory Concentration (MIC) Assay

Minimal inhibitory concentration (MIC) determination was performed according to the guidelines of the Clinical and Laboratory Standards Institute with the exception that incubation was at 37°C rather than at 30°C and as previously described (Rominski et al., 2017a). A fourfold difference in MIC values was considered significant. Ethambutol, isoniazid, bacitracin, vancomycin, rifampicin, and amikacin were dissolved in water and stored at −20°C until usage. Cefoxitin, ceftriaxone, penicillin G, and ampicillin were dissolved in water, aliquoted and stored as described by Rominski et al. (2017b). For the checkerboard MIC assay, twofold microdilutions of lysozyme (0.016–2 mg/ml) combined with bacitracin (0.5–64 mg/l) or vancomycin (0.008–1 mg/l) were produced in a 96-well plate format, basic fluorescence (emission: 530 nm/extinction: 590 nm) was measured at t−0 min, 20 µM ethidium bromide was added to each sample subsequently. 0.1 mM Carbonyl-Cyanide 3-Chlorophenylhydrazone (CCCP) was used to control the role of efflux pumps in ethidium bromide mediated fluorescence. Measurements were performed in triplicates and the median of the measurements is depicted.

Permeability Assay

*Mycobacterium abscessus* WT, Δpmt and Δpmt::pmt cultures in exponential growth phase were harvested, diluted to OD600 nm = 0.5 in uptake buffer (50 mM KH₂PO₄, 5 mM MgSO₄, pH = 7) and re-energized with 25 mM glucose. In a 96-well plate format, basic fluorescence (emission: 530 nm/extinction: 590 nm) was measured at t−2 min. 20 µM ethidium bromide was added to each sample subsequently. 0.1 mM Carboxyl-Cyanide 3-Chlorophenylhydrazone (CCCP) was used to control for the role of efflux pumps in ethidium bromide mediated fluorescence. Measurements were performed in triplicates and kinetics of ethidium bromide uptake were monitored from t−5 min on for 40 min at room temperature in a microplate reader (Bio Tek Synergy HT, Switzerland). Results were normalized against auto fluorescence of ethidium bromide and are depicted relative to the highest fluorescence measured. Statistical analysis of significance was performed by one-way ANOVA and Tukey’s Multiple Comparison Test (P < 0.05).

Intracellular Survival Assay

Intracellular survival of *M. abscessus* strains in THP-1 macrophages was investigated as previously described (Roux et al., 2016). Macrophages were infected with bacteria at an MOI in the range of 0.2–2 for 2 h at 37°C. Extracellular bacteria were removed by extensive washes with PBS followed by incubation of cells with RPMI medium containing 250 µg/ml amikacin. After 1 h, amikacin concentration in the medium was reduced to 50 µg/ml for the remaining experiment. Statistical analysis of the results was performed by one-way ANOVA and Tukey’s Multiple Comparison Test (P < 0.001).

RESULTS

Lipoprotein Acylation in *M. abscessus*

*Mycobacterium tuberculosis* LppX (Rv2945c) and Mpt83 (Rv2873) were chosen as model lipoproteins for the analysis of acylation and glycosylation in *M. abscessus*. These proteins are stable, not toxic, and known to carry both modifications in other mycobacteria (Michell et al., 2003; Tschumi et al., 2009, 2012; Mohiman et al., 2012; Brulle et al., 2013). We also investigated SodC (MAB_4184c) as an endogenous *M. abscessus* protein which shows 61% homology to the lipidated and glycosylated *M. tuberculosis* protein (Rv0432) (Sartain and Belisle, 2009). Lipoprotein expression vectors polyG-apra-LppX, polyG-apra-Mpt83 and polyG-apra-SodC were transformed into *M. abscessus* for the expression of full-length His- and HA-tagged proteins. These were isolated by FPLC and analyzed by mass-spectrometry. The N-terminal peptide of heterologously expressed Mpt83 (AspN-digestion fragment: CSSTKPVSDQ) was modified with palmitic acid/oleic acid (C16/C18) diacylglycerol as indicated by the detected peptide ion at m/z = 1513.899 (Figure 1A). This peptide ion was also identified with one additional hexose at m/z = 1674.183 (Figure 1B). A palmitic acid (C16) N-acyl modification in addition to the diacylglycerol and the hexose was found on the peptide ion at m/z = 1909.970 (Figure 1C). Fragmentation of these peptide ions in the MALDI-TOF/TOF analysis revealed the elimination of various fragments which confirmed the modifications (Figure 1D). Also LppX and SodC were shown to carry the characteristic lipid anchor of lipoproteins. Heterologously expressed LppX was modified with palmitic acid/oleic acid (C16/C18) diacylglycerol and a palmitic acid (C16) N-acyl residue in *M. abscessus*. Indication for a tuberculostearic acid/tuberculostearic acid (C19/C19) diacylglycerol modification at the N-terminal peptide of LppX (AspN-digestion fragment: CSSPKP) was obtained as well (Table 1). The N-terminal peptide of SodC (AspN-digestion fragment: CSP) was detected with a palmitoleic acid/oleic acid (C16:1/C18) diacylglycerol modification and one additional hexose (Table 1). The enzymatic machinery to synthesize tri-acylated lipoproteins at the conserved cysteine in their lipobox is thus present in *M. abscessus*. In silico homology searches with *M. tuberculosis* Lgt, LspA, and Lnt protein sequences identified MAB_2642c, MAB_2700c, and MAB_2206 as *M. abscessus* homologs.

MAB_1122c Is a Putative Protein-O-Mannosyltransferase in *M. abscessus*

In order to identify the gene responsible for the second post-translational modification on lipoproteins, we searched for homologs of mycobacterial lipoprotein glycosylating enzymes in *M. abscessus*. We performed an in silico search with *M. tuberculosis* (Rv1002) and *M. smegmatis* (Msmeg_5447) Pmt in *M. abscessus* ATCC 19977 and identified MAB_1122c as putative Pmt with 68% homology to the *M. tuberculosis* and the *M. smegmatis* protein, respectively. In order to confirm gene annotation and to assess the role of lipoprotein

spectra. Detailed Mascot results of the analyses can be found in Supplementary Table S3.
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**FIGURE 1** | MALDI-TOF/TOF analysis of Mpt83 (Rv2873) acylation in Mycobacterium abscessus WT. (A) MS/MS spectrum of the N-terminal peptide ion of Mpt83 (CSSTKPVSQ) with palmitic acid/oleic acid (C16/C18) diacylglycerol modification (calculated m/z = 1513.356). Elimination of a palmitic acid-thiol-glyceryl (calculated m/z = 327.56) was observed. (B) MS/MS spectrum of the N-terminal peptide ion of Mpt83 with palmitic acid/oleic acid (C16/C18) diacylglycerol + one hexose modification (calculated m/z = 1674.6). Elimination of a palmitic acid-thiol-glyceryl (calculated m/z = 327.56), the hexose (calculated m/z = 162.05) and oleic acid-thiol-glyceryl (calculated m/z = 353.66) was observed. (C) MS/MS spectrum of the N-terminal peptide ion of Mpt83 with palmitic acid/oleic acid (C16/C18) diacylglycerol + palmitic acid (C16) N-acyl + one hexose modification (calculated m/z = 1910.0). Elimination of palmitic acid-thiol-glyceryl (calculated m/z = 327.56), the hexose (calculated m/z = 162.05), oleic acid-thiol-glyceryl (calculated m/z = 353.66), oleic acid (calculated m/z = 281.5) and the palmitic acid/oleic acid diacylglycerol (calculated m/z = 578.0) was observed. (D) Chemical structures of the eliminated fragments are depicted: (1) palmitic acid-thiol-glyceryl, (2) hexose, (3) oleic acid-thiol-glyceryl, (4) oleic acid, and (5) palmitic acid/oleic acid diacylglycerol.

glycosylation in the bacteria’s physiology we generated an unmarked MAB_1122c deletion mutant by transformation of *M. abscessus* with the allelic replacement vector pSE-MAB1122c-KO (**Figure 2A**). Single crossover transformants were selected on agar plates containing apramycin and identified by Southern Blot analysis (**Figures 2A,B**). Transformants were subsequently subjected to counter selection in the presence of isoniazid and also analyzed by Southern Blot (**Figures 2A,B**). The *M. abscessus Δpmt* mutant was transformed with the complementation vector pMV361-MAB1122c. Genomic DNA was used to confirm integration of the vector by Southern Blot (**Figures 2A,B**).
TABLE 1 | Lipidation of heterologous and native lipoproteins in *M. abscessus*.

| Protein | Peptide modification | MS | MS/MS |
|---------|----------------------|----|-------|
|         |                      |    |       |
| LppX (Rv2945c) | CSSPKP + Diacylglycerol (C16/C18) | 1194.812 | 1195.202 |
|         |                      | C16/C18 diacylglycerol/A crude diacylglycerol | 616.82 | 616.287 |
|         |                      | C16/C18-thio-glycerol | 585.72 | 586.134 |
|         |                      | 1 Hex | 1433.691 | 1434.299 |
|         |                      | 1 Hex | 1080.031 | 1080.294 |
|         |                      | 616.82 | 616.287 |
| SodC (MAB_4184c) | CSP + Diacylglycerol (C16:1/C18) | 881.274 | 881.022 |
|         |                      | C16:1/C18 diacylglycerol/ C18-thio-glycerol ester | 305.274 | 305.100 |
|         |                      | 1 Hex | 305.274 | 305.100 |
|         |                      | 718.051 | 717.495 |
|         |                      | 305.274 | 305.100 |

Protein-O-Mannosyltransferase (MAB_1122c) Glycosylates Lipoproteins in *M. abscessus*

Heterologous expression of LppX (Rv2945c) and Mpt83 (Rv2873) in *M. abscessus* WT, Δpmt and Δpmt::pmt and subsequent LC-ESI-MS analysis revealed that in *M. abscessus* WT the lipoprotein LppX is glycosylated with one to two hexoses (Figure 3A). Mpt83 was more extensively glycosylated with up to six hexoses (Table 2). Glycosylated peptide ions eluted earlier than non-glycosylated peptide ions from the liquid-chromatography column due to increased hydrophilicity [see retention time (RT); Table 2]. All hexoses, presumably mannoses as described for other mycobacteria (Dobos et al., 1996; Michell et al., 2003), were identified in the tryptic N-terminal peptides (LppX: SSPKPDAAEQGVPVSPTASDPLAELIR/ Mpt83: PAT SPAAPVTTAAMADPAADLIGR). Analysis of lipoprotein glycosylation in *M. abscessus* WT compared to *M. abscessus* Δpmt and *M. abscessus* Δpmt::pmt revealed that MAB_1122c encodes for Pmt and is essential for lipoprotein glycosylation in *M. abscessus*. LppX purified from *M. abscessus* Δpmt was not glycosylated (Figure 3B) but glycosylation was restored in the complemented strain (Figure 3C). Mpt83 glycosylation was also abolished in *M. abscessus* Δpmt but was fully restored in the complemented *M. abscessus* Δpmt::pmt strain (Table 2). The identity of modified peptide ions was confirmed by manual analysis of corresponding MS/MS-spectra (Figure 3D). Due to the loss of glycosylation during HCD-fragmentation, the MS/MS-spectrum depicted in Figure 3D is representative both for the glycosylated and non-glycosylated N-terminal peptide ion of LppX (see Supplementary Table S4 for predicted fragment ions). In order to confirm that also endogenous *M. abscessus* proteins are glycosylated, we recombinantly expressed and subsequently analyzed *M. abscessus* SodC (MAB_4184c). LC-ESI-MS
FIGURE 3 | Glycosylation of LppX (Rv2945c) in M. abscessus. (A–C) Mass-spectra representative for LC-ESI-MS analyses of LppX purified from M. abscessus WT, Δpmt and Δpmt::pmt are shown. The N-terminal, tryptic peptide ion (SSPKPDAEQGVPVSPTASDPALLAEIR) of LppX was detected unmodified (calculated m/z = 954.489, 3+), with one hexose (calculated m/z = 1008.505, 3+) and with two hexoses (calculated m/z = 1062.522, 3+) in M. abscessus WT (A) and M. abscessus Δpmt::pmt (C). Differences of m/z = 162.05/3 = 54 between the peptide ion peaks reflect hexose modification (green dot). In M. abscessus Δpmt only the unmodified peptide ion (calculated m/z = 954.489, 3+) was detected (B). (D) The MS/MS spectrum of the unmodified peptide ion found in M. abscessus WT (depicted) is also representative for the N-terminal peptide ions of LppX from M. abscessus Δpmt and Δpmt::pmt since glycosylation is eliminated during higher-energy collision dissociation (HCD) fragmentation. Detection of predicted fragment ions (y- and b-ions; Supplementary Table S3) confirms the identity of modified and unmodified peptide ions.

analysis indicated that this protein is glycosylated with up to nine hexoses in M. abscessus WT. Glycosylation sites were located in the N-terminal peptide (AspN-digestion fragment: DQASTTPTGTPPYWTGSTAPATAEHGGHGGQPVPA GEKLNATLKL; Table 3). Identification of glycosylation of M. abscessus SodC indicated that also endogenous lipoproteins are glycosylated by Pmt (MAB_1122c) in M. abscessus.

M. abscessus Δpmt Is Susceptible to Bulky Antibiotics and to Compounds Acting on Peptidoglycan

The effect of lipoprotein glycosylation deficiency on antibiotic susceptibility of M. abscessus was assessed in a MIC assay. M. abscessus Δpmt showed increased susceptibility to all tested β-lactams (cefoxitin, ceftriaxone, penicillin G, and ampicillin) as well as to the high molecular weight antibiotics (Supplementary Figure S1) bacitracin (MIC = 256 mg/l), vancomycin (MIC = 4 mg/l), and rifampicin (MIC = 64 mg/l). Also susceptibility to lysozyme (MIC = 0.5 mg/ml) was elevated (Table 4). In contrast, susceptibility to smaller antibiotics interfering with the synthesis of other cell wall components (ethambutol, isoniazid) or affecting translation (amikacin) was not altered. Tolerance for the β-lactams, bacitracin, vancomycin, rifampicin, and lysozyme comparable to that seen in WT bacteria was restored in M. abscessus Δpmt::pmt. In order to investigate potential synergistic effects between the bulky and peptidoglycan affecting antibiotics bacitracin/vancomycin and lysozyme we conducted a checkerboard titration assay. Specifically, we tested subinhibitory concentrations of these antibiotics in combination with the peptidoglycan cleaving enzyme. Highest antibiotic concentrations tested were fourfold below MIC values determined for M. abscessus Δpmt (Table 4). The maximum tolerated concentrations of antibiotic and lysozyme are depicted in Figure 4. Susceptibility of M. abscessus WT and M. abscessus Δpmt::pmt to the antibiotics and lysozyme was unaffected by the combination of these compounds (Figures 4A,B). However, M. abscessus Δpmt became more susceptible to lysozyme with increasing concentrations of bacitracin. At the highest bacitracin concentration, which is still fourfold below the MIC value for M. abscessus Δpmt, as little as 0.02 mg/ml lysozyme were tolerated (Figure 4A). This observation indicates synergistic effects of lysozyme and bacitracin against M. abscessus Δpmt. Tolerated concentrations of lysozyme remained at 0.25 mg/ml until vancomycin concentrations rose to 0.5 and 1 mg/l (Figure 4B). Since only higher concentrations of vancomycin affected susceptibility of M. abscessus Δpmt to lysozyme, vancomycin and lysozyme exhibit weaker synergistic effects.

Increased Cell Wall Permeability in M. abscessus Δpmt

Based on the susceptibility of M. abscessus Δpmt to large antibiotics which affect peptidoglycan and the observed
TABLE 2 | Glycosylation of Mpt83 (Rv2873) in M. abscessus WT, Δpmt and Δpmt::pmt.

| Glycosylation | Calculated m/z | Observed m/z | z | RT | Observed m/z | z | RT | Observed m/z | z | RT |
|---------------|----------------|--------------|---|----|--------------|---|----|--------------|---|----|
| No hexose     | 761.05         | 761.05       | 3+| 46.3| 761.05       | 3+| 46.25| 761.05       | 3+| 46.2|
| 1 Hexose      | 815.07         | 815.072      | 3+| 44.69| 815.07       | 3+| 44.56| 815.07       | 3+| 44.56|
| 2 Hexoses     | 869.09         | 869.09       | 3+| 42.18| 869.08       | 3+| 42.23| 869.08       | 3+| 42.23|
| 3 Hexoses     | 923.10         | 923.10       | 3+| 41.32| 923.1        | 3+| 41.28| 923.1        | 3+| 41.28|
| 4 Hexoses     | 977.12         | 977.12       | 3+| 40.9 | 977.12       | 3+| 40.08| 977.12       | 3+| 40.08|
| 5 Hexoses     | 1031.14        | 1031.14      | 3+| 39.53| 1031.14      | 3+| 39.47| 1031.14      | 3+| 39.47|
| 6 Hexoses     | 1085.15        | 1085.15      | 3+| 38.76| 1085.15      | 3+| 38.96| 1085.15      | 3+| 38.96|

SodC (MAB_4148c)

| Glycosylation | Calculated m/z | Observed m/z | z | RT |
|---------------|----------------|--------------|---|----|
| No hexose     | 970.67         | 970.485      | 5+| 65.97|
| 1 Hexose      | 1003.08        | 1002.895     | 5+| 64.33|
| 2 Hexoses     | 1035.49        | 1035.305     | 5+| 63.32|
| 3 Hexoses     | 10967.90       | 1067.715     | 5+| 62.77|
| 4 Hexoses     | 1100.32        | 1100.125     | 5+| 61.06|
| 5 Hexoses     | 1132.72        | 1132.535     | 5+| 59.69|
| 6 Hexoses     | 1165.13        | 1164.945     | 5+| 59.02|
| 7 Hexoses     | 1197.54        | 1197.357     | 5+| 58.87|
| 8 Hexoses     | 1229.95        | 1229.767     | 5+| 57.96|
| 9 Hexoses     | 1262.36        | 1262.175     | 5+| 57.63|

synergistic effects with lysozyme, we hypothesized that the permeability of the cell envelope of this strain might be increased due to structural effects of lipoprotein glycosylation deficiency. We therefore conducted an ethidium bromide uptake assay to test for altered permeability in the MAB_1122c deletion mutant. Relative fluorescent intensity measurements indicated that M. abscessus Δpmt takes up ethidium bromide more efficiently than M. abscessus WT and M. abscessus Δpmt::pmt (Figure 5A). This elevated level of ethidium bromide uptake in the mutant strain was also observed upon inhibition of efflux processes with 0.1 mM CCCP while overall fluorescent intensity in all strains was higher due to increased ethidium bromide accumulation. The total increase in fluorescent intensity after 40 min was significantly higher in M. abscessus Δpmt as compared to WT and the complemented strain both with and without CCCP treatment (Figure 5B). Combined, these results point toward an increased permeability of the M. abscessus Δpmt cell envelope.

Glycosylation Deficiency Reduces Intracellular Survival of M. abscessus Δpmt

In vitro growth of M. abscessus strains was assessed in Middlebrook 7H9-OADC-Tween medium at 37°C with regular shaking and no difference in growth rate was observed between M. abscessus WT, Δpmt and Δpmt::pmt (Figure 5C). In order to investigate potential effects of glycosylation deficiency on virulence of M. abscessus we conducted an intracellular survival assay with the human monocyte-macrophage THP-1 cell line. M. abscessus WT and M. abscessus Δpmt::pmt survived inside the macrophages during a 3 days course of infection (Figure 5D). CFU counts for M. abscessus Δpmt significantly decreased by 0.8 log within 3 days of infection as compared to day 0 (Figure 5D). This indicates killing of M. abscessus Δpmt inside infected macrophages and represents virulence attenuation of the MAB_1122c deletion mutant as compared to M. abscessus WT and M. abscessus Δpmt::pmt.
DISCUSSION

In this study we investigated post-translational modification of lipoproteins and the function of lipoprotein glycosylation in the emerging pathogen *M. abscessus*. We show that *M. abscessus* tri-acylates lipoproteins with fatty acids obtained from the phospholipids in its plasma membrane. The observed lipid modifications include palmitic acid (C16), oleic acid (C18), palmitoleic acid (C16:1), and tuberculostearic acid (C19). Consistently with observations in *M. smegmatis* and *M. bovis* BCG, the membrane anchor of lipoproteins in *M. abscessus* is comprised of a heterogeneous combination of fatty acids which can vary per protein-molecule (Brulle et al., 2013). We demonstrate that lipoproteins can be glycosylated with a variable number of hexoses in *M. abscessus*. This modification is mediated by MAB_1122c which encodes for Pmt and is essential for lipoprotein glycosylation in the emerging pathogen. Modification by Pmt marks the first step in lipoprotein glycosylation in which the mannose is attached to the exported protein (Liu et al., 2013). Consequently, lipoprotein glycosylation in *M. abscessus Δpmt* is abolished completely. All glycosyl-residues in *M. abscessus* were detected in the N-terminal region of the model lipoproteins as described for lipoproteins expressed in other mycobacteria as well (Herrmann et al., 1996; Michell et al., 2003; Sartain and Belisle, 2009; Mohiman et al., 2012; Sethi et al., 2016; Parra et al., 2017). This observation indicates the existence of a conserved motif which is recognized by Pmt from various mycobacterial species and directs site-specific glycosylation of lipoproteins in fast- and slow-growing mycobacteria (Sartain and Belisle, 2009; Smith et al., 2014). Our notion, that heterologously expressed lipoproteins like Mpt83 carry glycosyl-residues in the same N-terminal region and probably at identical residues as when expressed in their native environment supports this hypothesis (Michell et al., 2003). Despite intensive investigations by Smith et al. (2014) the potential consensus sequence for O-glycosylation sites has to be further defined and verified in mycobacteria. Another possible explanation for the exclusive and conserved localization of glycosyl-moieties could be that only serine or threonine residues in the N-terminal region of lipoproteins are accessible for Pmt during Sec-export. The SodC protein was detected with up to nine mannoses in *M. tuberculosis* (Sartain and Belisle, 2009) and carried the same number of hexoses in *M. abscessus*. However, a mechanism for glycosyl-chain length control in mycobacteria is unknown to date. It is also not clear yet whether detected mannoses are attached to one amino acid each or whether oligosaccharide chains are formed in *M. abscessus*. Extensive glycosylation as seen for Mpt83 and SodC, however, suggests that at least short mannose chains occur. Elongation of these would be performed not by Pmt but by PimE (Rv1159), an enzyme also involved in the synthesis of other membrane components such as LMs and LAMs where it mediates the addition of α-(1,2) linked mannose to PIM molecules (Liu et al., 2013). In *M. abscessus* MAB_1304 has 64% homology to *M. tuberculosis* PimE.

Next to the structural analysis of lipoprotein modification we show that glycosylation deficiency in *M. abscessus* alters bacterial susceptibility to large and bulky compounds and to drugs and enzymes which affect peptidoglycan. Due to their thick and almost impermeable outer membrane like structure and the expression of β-lactamases mycobacteria are usually not susceptible to antibiotics acting on the peptidoglycan layer (Brennan and Nikaido, 1995; Nessar et al., 2012). Nevertheless, we observed synergistic effects of both bacitracin and vancomycin with lysozyme in the Δpmt strain. We confirmed by *in silico* analysis that *M. abscessus* does not express a homolog of the lysozyme inhibitor LprI which has been described in *M. tuberculosis* recently and could affect lysozyme MIC-testing (Sethi et al., 2016). Particularly the antimicrobial activity of the bacitracin-lysozyme combination was strong. The observed effects on antibiotic susceptibility and intracellular survival might occur due to structural alterations in the cellular envelope of *M. abscessus Δpmt* upon lipoprotein glycosylation deficiency. This hypothesis is in line with the increased cell envelope permeability seen in the ethidium bromide uptake assay and could explain why particularly large antibiotics like bacitracin, vancomycin, and rifampicin (Supplementary Figure S1) can affect the mutant and why the β-lactams can better reach their target, the peptidoglycan layer. Also for the *M. tuberculosis* RpPnull mutant deficient for all five mycobacterial RPs it was described that the permeability of the outer membrane like structure is increased which led to a change in susceptibility to various β-lactams (Wivagg et al., 2014). This observation supports the link between an altered cell wall homeostasis and increased susceptibility to a broad range of large and bulky as well as to peptidoglycan affecting compounds which we describe in this study for the *M. abscessus Δpmt* mutant. The details of how lipoprotein glycosylation deficiency affects cell permeability and drug susceptibility of *M. abscessus*, however, remain elusive. One possibility is that specific lipoproteins lose function without their post-translational modification but to date there is no indication for a functional requirement of glycosylation on lipoproteins. For SodC it was shown that glycosylation deficiency affects localization of the lipoprotein but is dispensable for its function (Sartain and Belisle, 2009). Also for Lpr1 in *M. tuberculosis* it was reported that glycosylation is crucial for correct localization of the protein (Sethi et al., 2016). It is therefore likely that the

| **TABLE 4** | Antibiotic susceptibility of *M. abscessus* WT, Δpmt and Δpmt:pmt. |
| --- | --- | --- |
| **Compound (mg/l)** | **Strain** | **Strain** | **Strain** |
|  | WT | Δpmt | Δpmt:pmt |
| Ethambutol | 64 | 64 | 64 |
| Isoniazid | 512 | 1024 | 512 |
| Bacitracin | >2048 | 256 | >2048 |
| Vancomycin | 128 | 4 | 128 |
| Rifampicin | 256 | 64 | 256 |
| Amikacin | 2 | 2 | 2 |
| Cefoxitin | 64 | 16 | 32 |
| Ceftriaxone | 128 | 32 | 128 |
| Penicillin | >512 | 256 | >512 |
| Ampicillin | >512 | 128 | >512 |
| Lysozyme | >2000 | 500 | >2000 |
Becker et al. Lipoprotein Glycosylation in M. abscessus Resistance

Synergistic effects of bacitracin and vancomycin with lysozyme. Antibiotic dilution series starting with concentrations fourfold below the MIC value for bacitracin (A) and vancomycin (B) determined for M. abscessus Δpmt were combined with lysozyme concentrations of 0.016–2 mg/ml. Maximum tolerated concentrations of both compounds are depicted for M. abscessus WT (*), Δpmt (•) and Δpmt::pmt (_). Results of three independent experiments are represented and the median of tolerated concentrations with the range of all measurements is shown.

Permeability and virulence of M. abscessus WT, Δpmt and Δpmt::pmt. (A) Fluorescent intensity of M. abscessus cultures at t = 2 min and after subsequent addition of ethidium bromide was measured to detect uptake and incorporation of the fluorescent molecule into DNA. Kinetics of ethidium bromide uptake from t = 0 min onward were measured for 40 min at room temperature. Results of three independent experiments are shown. Fluorescent intensities were normalized against autofluorescence of ethidium bromide in uptake buffer and are depicted relative to the maximal intensity measured. (B) The relative fluorescent intensity increase after 40 min was calculated for all strains with and without CCCP treatment and statistically analyzed by one-way ANOVA and Tukey’s Multiple Comparison Test (**P < 0.05; ***P < 0.001). Error bars reflect the standard deviation in three independent experiments. (C) Growth curve in Middlebrook 7H9-OADC-Tween at 37°C with regular shaking for M. abscessus WT, Δpmt and Δpmt::pmt. Results of three independent experiments with standard deviations are depicted. (D) Survival of M. abscessus WT, Δpmt and Δpmt::pmt in THP-1 macrophages 3 days after infection. Results of multiple independent experiments are shown (WT: n = 3; Δpmt: n = 3; Δpmt::pmt: n = 2). One-way ANOVA and Tukey’s Multiple Comparison Test statistical analysis were performed (**P < 0.01).

Localization of specific lipoproteins is altered upon glycosylation deficiency. This could affect the cell envelope structure directly or interfere with the homeostasis through indirect alterations in membrane composition. In M. tuberculosis, lipoproteins such as LpqW (conversion of phosphoinositol mannosides to lipomannans and lipoarabinomannans) or LppX (transport
of phthiocerol dimycocerosates) for instance are important players in the synthesis and transport of cell wall components, respectively (Sulzenbacher et al., 2006; Rainczuk et al., 2012). Homologs of LpqW (MAB_1315) and the LppX_LprAFG superfamily (MAB_2806) are encoded in M. abscessus. Incorrect localization of these or other lipoproteins in M. abscessus Δpmt could affect cell wall homeostasis because their substrates might not be synthesized or located correctly.

Based on the obtained results it cannot be excluded that Pmt possess an additional function independent of lipoprotein glycosylation that leads to the observed phenotypes. The enzyme might be involved in the glycosylation of other (non-lipo-) proteins but does not play a role in lipomannan or lipoarabinomannan synthesis (Liu et al., 2013). Due to the specific localization of lipoproteins in the cell envelope, however, it is plausible that the observed phenotypes are assigned to a deficiency in lipoprotein glycosylation. Pmt deletion mutants of M. smegmatis and M. tuberculosis have been investigated (Liu et al., 2013). The fast-growing, non-pathogenic M. smegmatis Δpmt strain showed WT-like growth and showed reduced virulence in immunocompromised mice (Liu et al., 2013). The observed phenotype was very recently attributed to increased lipoarabinomannan production in the Δpmt strain and secretion of this molecule partly bound to LprG. This led to a stronger inflammatory response via TLR-2 activation and secretion of this molecule partly bound to LprG. This led to a stronger inflammatory response via TLR-2 activation and could be a rationale for the attenuated virulence of M. tuberculosis Δpmt (Alonso et al., 2017) and thus potentially also of M. abscessus Δpmt. However, in the fast-growing pathogenic M. abscessus we see an intermediate phenotype: attenuated survival in macrophages but no in vitro growth defect. It is tempting to speculate that the growth rate of the mycobacterial species could affect the severity of the glycosylation deficient phenotype and that lipoprotein glycosylation is more crucial in pathogenic mycobacteria but this requires further investigation.

It will be of interest to explore the detailed downstream effects of lipoprotein glycosylation deficiency on the cell envelope permeability of M. abscessus in order to elucidate how the observed phenotypes arise on a molecular level. Furthermore, it remains an open question how glycosylation is exclusively directed to the N-terminal region of lipoproteins in all mycobacterial species investigated so far and whether mannose chain length is regulated and conserved. The results of this study contribute to a more detailed understanding of the fundamental biology and the drug resistance of the emerging pathogen M. abscessus. They can be used as a foundation for further exploration of lipoproteins in the context of the complex mycobacterial cell envelope in non-tuberculous but pathogenic mycobacteria.

**AUTHOR CONTRIBUTIONS**

KB and PSa conceived and designed the study. KB, KH, PS, and LR acquired and analyzed the data. KB drafted the manuscript and PSa critically revised it. All authors approved the final version of the manuscript.

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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.2017.02123/full#supplementary-material

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Conflict of Interest Statement: 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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