Glycopeptidolipid of *Mycobacterium smegmatis* J15cs Affects Morphology and Survival in Host Cells

Nagatoshi Fujiwara1,2,*, Naoya Ohara3☯, Midori Ogawa4☯, Shinji Maeda5, Takashi Naka2,6, Hatsumi Taniguchi4, Saburo Yamamoto7, Minoru Ayata8

1 Department of Food and Nutrition, Faculty of Contemporary Human Life Science, Tezukayama University, Nara City, Nara, Japan, 2 Department of Bacteriology, Osaka City University Graduate School of Medicine, Osaka City, Osaka, Japan, 3 Department of Oral Microbiology, Okayama University Graduate School of Medicine, Dentistry and Pharmaceutical Sciences, Okayama City, Okayama, Japan, 4 Department of Microbiology, School of Medicine, University of Occupational and Environmental Health, Kitakyushu City, Fukuoka, Japan, 5 Molecular Epidemiology Division, Mycobacterium Reference Center, The Research Institute of Tuberculosis, Japan Anti-Tuberculosis Association, Kiyose City, Tokyo, Japan, 6 MBR Co. Ltd., Toyonaka City, Osaka, Japan, 7 Japan BCG Laboratory, Kiyose City, Tokyo, Japan, 8 Department of Virology, Osaka City University Graduate School of Medicine, Osaka City, Osaka, Japan

* These authors contributed equally to this work.

**fujiwara@tezukayama-u.ac.jp**

Abstract

*Mycobacterium smegmatis* has been widely used as a mycobacterial infection model. Unlike the *M. smegmatis* mc2155 strain, *M. smegmatis* J15cs strain has the advantage of surviving for one week in murine macrophages. In our previous report, we clarified that the J15cs strain has deleted apolar glycopeptidolipids (GPLs) in the cell wall, which may affect its morphology and survival in host cells. In this study, the gene causing the GPL deletion in the J15cs strain was identified. The *mps1-2* gene (MSMEG_0400-0402) correlated with GPL biosynthesis. The J15cs strain had 18 bps deleted in the *mps1* gene compared to that of the mc2155 strain. The *mps1*-complemented J15cs mutant restored the expression of GPLs. Although the J15cs strain produces a rough and dry colony, the colony morphology of this *mps1*-complemented was smooth like the mc2155 strain. The length in the *mps1*-complemented J15cs mutant was shortened by the expression of GPLs. In addition, the GPL-restored J15cs mutant did not survive as long as the parent J15cs strain in the murine macrophage cell line J774.1 cells. The results are direct evidence that the deletion of GPLs in the J15cs strain affects bacterial size, morphology, and survival in host cells.

Introduction

Tuberculosis is an infectious disease worldwide, and the causative agent, *Mycobacterium tuberculosis*, is a slow-growing intracellular pathogen. To control mycobacterial infection, it is necessary to clarify the mechanisms of host–pathogen interactions and how the pathogen survives in host cells.
Mycobacterium smegmatis is a rapidly growing non-pathogenic mycobacterium. Its features make it useful as a tool for genetic analysis in the research on mycobacterial infection. In particular, the *M. smegmatis* mc²155 strain is widely used because of its high rate of transformation. We screened another *M. smegmatis* strain, J15cs, and constructed a convenient and safe host-vector system of a mycobacterial model. We previously reported that the J15cs strain replicated our original vector, pYT923, and survived in host cells longer than the mc²155 strain [1]. We clarified a further difference between J15cs and mc²155 strains. The J15cs strain lacks glycopeptidolipids (GPLs) on the cell surface, and this may affect its morphology and survival in host cells [2].

GPL, also called C-mycoside, is produced in the outer layer of some non-tuberculous mycobacteria, including *Mycobacterium avium*, *M. intracellulare*, *M. abscessus*, *M. chelonae*, and *M. smegmatis*. Structurally, GPLs comprise a lipotetrapeptide core, fatty acid, and oligosaccharide. Three amino acids (D-phenylalanine, D-allo-threonine and D-alanine) link a 3-hydroxy or 3-methoxy long-chain fatty acid (C26-34) to the N-terminal and an amino alcohol, alaninol, to the C-terminal. The 6-deoxy-talose is glycosidically linked to the D-allo-threonine and a methylated L-rhamnose to the alaninol to generate the non-specific core GPLs found in all members of GPL-producing mycobacteria. In *M. avium* and *M. intracellulare*, an oligosaccharide unit is elongated from 6-deoxy-talose, leading to the serotype-specific, so-called polar GPLs [3]. It is considered that GPLs are related to bacterial aggregation, sliding motility, biofilm formation, and cell wall integrity [4,5].

In this study, we examined the GPL biosynthesis pathway and clarified the causative gene cluster that deletes the GPLs in the J15cs strain. We targeted the *mps* gene that involves the tetrapeptide synthesis in the GPLs [6]. Interestingly, recovery of the GPLs affected colony morphology and survival in host cells.

### Materials and Methods

#### Bacterial strains

The J15cs strain is diverged from the *M. avium* Jucho strain and maintained in our laboratory [1,2]. The mc²155 strain (ATCC 607) was purchased from the American Type Culture Collection (Manassas, VA). Both strains were grown on Middlebrook 7H11 agar (Difco Laboratories, Detroit, MI) with 0.5% glycerol and 10% Middlebrook OADC enrichment (Difco Laboratories) or nutrient agar at 37°C for 2–7 days. *Escherichia coli* was cultured in LB broth. Kanamycin (20 μg/ml) was added to the selective medium for pNN2-containing plasmids.

**Amplification and sequencing of DNA fragments in *M. smegmatis* J15cs genome**

DNA fragments containing the region covering the *mps1* and *mps2* genes were amplified by polymerase chain reaction (PCR) with specific primer pairs. The primers were prepared in reference to the reported sequence of the mc²155 strain (Table 1). KOD FX Neo DNA polymerase (Toyobo Co., Ltd, Osaka, Japan) was used for PCR amplification and the condition for PCR was 95°C-2 min, 5 cycles of 98°C-10 sec, 72°C-2.5 min, 5 cycles of 98°C-10 sec, 70°C-2.5 min, 30 cycles of 98°C-10 sec, and 68°C-2.5 min. The expected sizes of PCR products were 4,240 bps (primer pair: MSMEG_0400-F1 and MSMEG_0400-R2), 3,758 bps (primer pair: MSMEG_0400-F2 and MSMEG_0400-R1), 2,987 bps (primer pair: MSMEG_0401-F1 and MSMEG_0401-R1), 4,204 bps (primer pair: MSMEG_0402-F1 and MSMEG_0402-R2), and 3,776 bps (primer pair: MSMEG_0402-F2 and MSMEG_0402-R1). The DNA fragments were isolated and purified with a gel extraction kit (Qiagen, Germantown, MD) and were directly sequenced in both directions.
with a series of synthetic primers (not shown), a sequencing kit (BigDye terminator v3.1 cycle sequencing kit, Life Technologies, Carlsbad, CA), and automated sequencers (Prism 310 or 3130xl genetic analyzer, Life Technologies).

**Construction of plasmids expressing mps1 and transduction to M. smegmatis J15cs strain**

DNA fragments containing the whole mps1 gene of the mc²155 strain were amplified by PCR with specific primer pairs (Primer Nde-mps1 and Pac-mps1) in Table 1. LA Taq DNA polymerase (Takara Bio Inc., Shiga, Japan) was used for PCR amplification. The PCR conditions were 94°C for 1 min, followed by 98°C for 10 sec, 68°C for 15 min, which lasted for 30 cycles, and a final extension at 72°C for 10 min. The DNA fragments were purified and cloned into a pGEM-T Easy vector (Promega Corp., Fitchburg WI). A plasmid clone containing the authentic sequence was obtained (a single mutation was allowed because it is silent). The 10.3 kb of amplified DNA were double digested with NdeI-PacI and ligated into the same sites of pNPP, resulting in pNPP-MPS1. The pNPP is a pUC19 derivative containing the promoter region of aphII and the terminator of the antigen gene from *Mycobacterium kansasii* [7]. Then, pNPP-MPS1 was digested with XbaI and the inserted fragment was ligated into the same site of an *E. coli*-mycobacteria shuttle plasmid pNN2 to yield pNN-MPS1 [8]. The constructed plasmid, pNN-MPS1, was selected and its inserts were confirmed.

**Analysis of GPL compositions**

The GPLs were detected by thin-layer chromatography (TLC) as described previously [2]. Briefly, the heat-killed bacteria were sonicated, and crude lipids were extracted with chloroform-methanol (2:1, vol/vol). The crude lipids were hydrolyzed with 0.2 N sodium hydroxide in methanol at 37°C for 2 h, and neutralized with 6 N hydrochloric acid. Alkaline-stable lipids extracted in the organic phase of a two-layer partition with chloroform-methanol (2:1, vol/vol) and water, were evaporated and precipitated with acetone to remove any acetone-insoluble components. The TLC on Silicagel G (Uniplate; 20 by 20 cm, 250 μm; Analtech, Inc., Newark, DE) was developed with a solvent system of chloroform-methanol (90:10, vol/vol). The TLC plate was sprayed with 20% sulfuric acid in ethanol and charred at 180°C for 3 min. The GPLs were detected as a brownish-yellow spot. The GPLs were purified by exposing the TLC plate to iodine vapor, and the GPL spot was marked, scraped off, and eluted with chloroform-methanol.
To assign the structures of the GPLs, the molecular weight of each GPL was measured by matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF MS) using an Ultraflex II (Bruker Daltonics, Billerica, MA) with 10 mg/ml 2,5-dihydroxybenzoic acid (DHB) in chloroform-methanol (1:1, vol/vol) as a matrix, and analyzed in Reflectron mode with an accelerating voltage operating in positive mode at 20 kV.

**Colony morphology and scanning electron microscopy**

Colony morphology was observed using an optical microscope after incubation for 10 days on nutrient agar plates. To prepare the bacteria for scanning electron microscopy (SEM), they were suspended with 0.1 M phosphate buffer (pH 7.2), passed through a 5 μm membrane filter (Millipore, Billerica, MA, USA) to remove aggregates, and then put on coverslips (Thermanox Plastic, 13 mm diameter; Thermo Fisher Scientific, Rochester, NY) coated with 0.1% (w/v) poly-L-lysine solution. The bacteria were fixed with 2.5% glutaraldehyde (TAAB Laboratories Equipment Ltd., West Berkshire, UK) in 0.1 M phosphate buffer for 2 h at room temperature, washed twice with 0.1 M phosphate buffer, and post-fixed with 1% osmium tetroxide (TAAB Laboratories Equipment) in 0.1 M phosphate buffer for 1 h at room temperature. The bacteria were then dehydrated in a graded ethanol series. After permutation with 3-methyl butyl acetate, the critical point drying was determined using a Hitachi HCP-2 Critical Point Dryer (Hitachi High-Technologies Corp., Tokyo, Japan). SEM was performed at ×5,000 on a Hitachi FE-SEM S-4700 instrument (Hitachi High-Technologies Corp.) using an accelerating voltage of 10 kV [9].

**Intracellular growth in J774.1 cells**

A murine macrophage, J774.1 cell line (Cell Resource Center for Biomedical Research Institute of Development, Aging and Cancer Tohoku University, Sendai, Miyagi, Japan), was cultured in RPMI 1640 medium (Nissui Pharmaceutical, Tokyo, Japan) supplemented with 10% inactivated fetal calf serum (FCS, GIBCO, Grand Island, NY) at 37°C in 5% CO₂. The J774.1 cells were grown to semi-confluence for 48 h on 24-well flat-bottom tissue culture plates (Becton Dickinson Labware, Oxnard, CA). *M. smegmatis* strains were pre-cultured on nutrient agar plates for 7 days. PBS added on agar plate. Bacterial cells were suspended, and were mixed slowly. After collecting and pumping by 23G syringe, the bacterial suspension was passed through a 5 μm membrane filter to remove aggregates. A single-cell suspension was checked by acid-fast staining, and was prepared at a concentration of approximately 1 × 10⁸ CFU/ml. A bacterial single-cell suspension of 1×10⁷ CFU/well was inoculated into the J774.1 cells at an approximate multiplicity of infection 10. After cocultivation for 3 h, the infected J774.1 cells were washed twice with PBS in order to remove non-infecting bacteria. To kill any remaining extracellular bacteria, the infected J774.1 cells were further incubated for 2 h with fresh culture medium supplemented with 5% inactivated FCS and 200 μg/ml of gentamicin (Sigma Chemicals, St. Louis, MO). The medium was replaced with fresh medium containing 2 μg/ml of gentamicin (this time point was day 0), and the infected cells were cultured for 8 days. The survival of mycobacteria was evaluated by counting colony-forming units (CFU). The adherent cells were treated with 1% Triton-X100/PBS, and were sonicated. The lysates plated on nutrient agar. The experiments were performed in triplicate, and statistical analysis was carried out using Dunnett’s test [1].
![Image of DNA sequence](image1.png)

**Results**

**Mutation of mps1-2 genes in *M. smegmatis* J15cs strain**

The DNA fragments of MSMEG_0400–0402 were amplified by PCR and the sequence was determined. The nucleotide sequence MSMEG_0400–0402 of the J15cs strain was deposited in the NCBI GenBank database under accession number AB924538. The results showed a segment of DNA with a total length of 18,238 bps. The sequence was identical to that reported for the mc²155 strain (GenBank accession no. AY439015) except for a gap of 18 consecutive bps nucleotides (Fig 1). Two open reading frames, *mps1* and *mps2*, were encoded in this segment. The Mps1 of the J15cs strain was 6 consecutive amino acids shorter compared to that of the mc²155 strain, whereas the Mps2 of the J15cs strain was identical to that of the mc²155 strain.

**Phenotype of *M. smegmatis* J15cs mutant complemented with mc²155-mps1**

Next, the *mps1* of the mc²155 strain (mc²155-mps1) was inserted into a pNN2 vector (kanamycin-resistant, Fig 2), and the J15cs strain was transformed using this plasmid by

![Image of plasmid construction](image2.png)

**Fig 1. The mps1 gene of *M. smegmatis* J15cs strain was 18 bps shorter than that of *M. smegmatis* mc²155 strain.** The schematic gene map of the GPL biosynthetic cluster in *M. smegmatis* [6,13]. The sequence of MSMEG_0400–0402 was determined (GenBank accession no. AB924538). Two open reading frames, *mps1* and *mps2*, were encoded in this segment. The Mps1 of the J15cs strain was 6 consecutive amino acids shorter compared to that of the mc²155 strain, whereas the Mps2 of the J15cs strain was identical to that of the mc²155 strain.

**Fig 2. Construction of the mc²155-mps1 plasmid.** The *mps1* of the mc²155 strain was inserted into a pNN2 vector (kanamycin resistance). PaphII, the promoter of aphII; mc²155-mps1, the open reading frame of *mps1* from *M. smegmatis* mc²155; Ta1pha, the terminator region of the alpha antigen gene from *Mycobacterium kansasii*.
The promoter of *aphII* was employed to express *mps1*. Two mutants (3–5, 5–4) of the J15cs strain, which were complemented with the mc2155-*mps1*, were obtained. The phenotypes of GPLs were examined. The alkaline-stable lipids were analyzed by TLC and are shown in Fig 3. Although the parent strain and the pNN2-vector-inserted J15cs mutant lacked GPL spots, the five major GPL spots of the *mps1*-complemented J15cs mutants were restored with the same Rf (retention factor) values on TLC as those of the mc2155 strain. In addition to the comparative Rf value, the molecular weight of each spot was measured by MALDI-TOF/MS (Fig 4). The mass spectra of the *mps1*-complemented J15cs mutants showed the same masses as the GPL1-5 derived from the mc2155 strain previously reported by Miyamoto et al. [11]. As a result, it was clarified that mutation of the *mps1* gene directly caused the deletion of GPLs in the J15cs strain.

**Diversity of morphology by GPLs restoration**

After being cultured on nutrient agar for 10 days at 37°C, the J15cs strain formed rough, dry, and irregular colonies. In contrast, the colonies of the mc2155 strain were smoothly curved and wet. Thus, the morphology of the colonies was strain-specific. The J15cs strain carrying the pNN2 vector showed the same morphology as the parent J15cs strain, and the *mps1*-complemented J15cs mutant formed smoothly curved and wet colonies like the mc2155 strain (Fig 5A). The hydrophobicity of the *mps1*-complemented J15cs mutant was reduced, compared to the parent J15cs strain. The SEM images showed that the bacterial length of the J15cs strain was longer than that of the mc2155 strain, and the *mps1*-complemented J15cs mutant was significantly shorter than the parent J15cs strain. There was no significant difference in their widths (Fig 5B and 5C). The restoration of GPLs in the *mps1*-complemented J15cs mutant may modify the cell wall construction. As a result, the colony morphology and bacterial size were changed to those of the mc2155 strain. It was concluded that the existence of GPLs affected bacterial size and morphology.
Effect of GPLs on intracellular survival of host cells

To determine the effect of GPLs on bacterial survival in host cells, J774.1 cells were infected with the mps1-complemented J15cs mutant. Surprisingly, this GPL-restored J15cs mutant did not survive as long as the parent J15cs strain. On day 2 after infection, the survival rate of the GPL-restored J15cs mutant was significantly lower than that of the parent strain, and the survival curve was almost the same as that of the mc2155 strain that expressed GPLs (Fig 6).

Discussion

*M. smegmatis* is a widely analyzed mycobacterial infection model because of its nonpathogenicity and rapid growth. The J15cs strain survives in host cells longer than the mc2155 strain. In our previous reports, it was shown that the difference between these two strains was the presence or absence of an apolar GPL. GPLs are synthesized by some nontuberculous mycobacteria. Because of their location on the cell surface, their presence is considered to correlate with the pathogenicity of the bacteria by interfering with the host immune responses. We predicted that the GPLs play crucial roles in the intracellular survival of the J15cs strain. The GPL was composed of a tetrapeptide-linked acyl chain and a sugar. Recently, the biosynthesis pathway of GPL is well understood, and more than fifteen genes are involved [6,12]. In this study, we...
Fig 5. Change of morphology due to expression of GPLs. The colony morphology was observed by an optical microscope after culturing. The J15cs strain formed rough, dry, and irregular colonies. The mc²155 strain and the mps I-complemented J15cs mutant colonies were smoothly curved and wet (a). Scanning electron microscopy (b) and a comparison of bacterial cell length (c) are shown. The bacteria were grown on nutrient agar plates for 10 days at 37°C, collected, and filtered through a 5 μm membrane to remove aggregates. The average bacterial cell length was 2.11±0.46 μm, the mc²155 strain; 5.70±1.65, the J15cs strain; 2.18±0.62 μm, the mps I-complemented J15cs mutant. The data are means±standard deviations (SD) for 20 bacteria. The separate experiments were done in duplicate.

Fig 6. Survival of mps I-complemented J15cs mutant in J774.1 cells. The J774.1 cells were grown on 24-well flat-bottom tissue culture plates. A bacterial single-cell suspension of approximately 1×10⁷ CFU/well was inoculated into the J774.1 cells at an approximate multiplicity of infection 10. After cocultivation for 3 h, the infected J774.1 cells were washed, and the infected J774.1 cells were further incubated for 2 h with the medium plus 200 μg/ml gentamicin. The medium was replaced with fresh medium containing 2 μg/ml of gentamicin (this time point was day 0), and the infected cells cultured for 8 days. At various time intervals after inoculation, the adherent cells were treated with 1% Triton-X100/PBS, and were sonicated. The bacterial burden was evaluated by counting CFU. The separate experiments were done in triplicate, and statistical analysis was performed using Dunnett’s test. * and ** indicate a p value of <0.05, compared to the J15cs strain inserted with a pNN2 vector only respectively.
focused on the mps1 and mps2 genes, which belong to the non-ribosomal protein synthesis family (nrp), and are responsible for synthesis of the peptidyl core of the GPLs [13,14]. Comparison of the sequences shows that 18 bps encoding 6 amino acids were deleted in the mps1 in the J15cs strain, and insertion of mc2155-mps1 into the J15cs mutant restored expression of the GPLs. The MALDI-TOF/MS analysis showed that the detailed structure, modifying groups of acyl chains, the sugar of GPLs, and the GPLs of the mps1-complemented J15cs mutant were completely identical to those of the mc2155 strain. These results implied that the mps1 mutation directly caused the deletion of GPLs in the J15cs strain. Because the deletion of 18 bps did not cause a frame-shift mutation, these deleted 6 amino acids in the Mps1 protein are considered to be important for the secondary, tertiary protein structures and active domain. Based on the phenotype of the mps1-complemented J15cs mutant, the deletion of GPLs affects two prominent features in the J15cs strain. One is bacterial size. The J15cs strain is longer than mc2155, and that of the mps1-complemented mutant was almost the same length as mc2155 rather than the parent J15cs strain. The other is the colony morphology. The J15cs strain was the rough type, and the complemented mutant shifted to the smooth type. These phenomena were due to the presence of GPLs.

Next, we were interested in whether these phenotypes correlated with bacterial survival in host cells or not. Surprisingly, the mps1-complemented J15cs mutant did not survive as long as the parent J15cs strain in host cells. This result showed that the survival of the J15cs strain is correlated with the lack of GPLs. Toll-like receptor (TLR) 2 recognizes GPLs [2,15,16,17]. The GPL-recognition via TLR2 does not work when the J15cs strain infects to host cell. As a result, the J15cs strain evades one of the major host-pathogen recognition systems and may survive longer than the other strain. In addition, the colony morphology and bacterial size may affect host-bacteria interaction. Mycobacterium abscessus is a rapidly growing mycobacteria closely related to M. smegmatis. This species is recognized as a pulmonary pathogen in humans. Recently, Bernut et al. reported that the rough variant of M. abscessus, which is devoid of GPLs, causes more severe clinical disease than the smooth variant. The mechanism of the increased virulence of the M. abscessus rough variant was explained to be the massive production of serpentine cords, which are absent in smooth variant infection. In apoptotic macrophages, the rough variant of M. abscessus initiated cord formation and grew so large that the macrophages and neutrophils could not phagocytize them [18,19]. In addition, Davidson et al. reported that GPLs mask underlying bioactive cell wall lipids, and the GPL-lacking M. abscessus rough variant is immunostimulatory and invasive in macrophage infection [15]. Roux et al. reported that the M. abscessus rough variant is associated with synthesis/exposure of the lipoprotein at the cell surface [20]. Etienne et al. reported that the GPL-deficient mc2155 mutant strain exhibited rough colony morphology, and indicated that GPLs are involved in the cell wall permeability barrier of M. smegmatis [21,22]. These previous reports show the possibility of the GPL-host relationship, and we summarize the reasons for the survival of the J15cs strain. By the deletion of GPLs, the J15cs strain lost prominent features that are correlated with host-bacterial interaction and host responses, in addition to the direct host immune response to GPLs via TLR2.

As for the GPLs, the major nontuberculous pathogens M. avium and M. intracellulare produce a serotype-specific polar GPL that plays a crucial role in the host immune responses. According to our previous report, the acetyl or methyl modifications of oligosaccharides are necessary for host cells to recognize polar GPLs via TLR2. The activities of the polar GPLs are greatly decreased by the deletion of acetyl groups from oligosaccharides [16]. The major epitopes for the host immune system may be oligosaccharides in the polar GPLs. The production of an anti-GPL IgA antibody is increased in patients and useful in the serodiagnosis of nontuberculosis infection using the GPL core antigen [23,24]. In the nontuberculous mycobacteria infection, the GPLs were one of the key components on the cell wall related to their virulence.
and pathogenicity. We should pay attention to the functions of two types of GPLs, the apolar types in *M. abscessus* and the polar types in the *M. avium-intracellulare* complex. On the other hand, GPLs are proposed to confer the capacity for sliding motility, biofilm formation, and dormancy (to be published elsewhere) [25,26].

In this study, direct evidence was shown that the *mps1* mutation causes the deletion of GPLs in the J15cs strain, and affects bacterial size, morphology, and survival in host cells.

**Acknowledgments**

We are grateful to Hideki Nakagawa, Kazuki Matsushita and Takashi Wada (Osaka City University Graduate School of Medicine, Osaka), and Kazumasa Fukuda (School of Medicine, University of Occupational and Environmental Health, Kitakyushu) for providing technical assistance.

**Author Contributions**

Conceived and designed the experiments: NF. Performed the experiments: NF NO MO SM TN HT SY MA. Analyzed the data: NF MA. Contributed reagents/materials/analysis tools: NF SM. Wrote the paper: NF MA.

**References**

1. Nomoto M, Ogawa M, Fukuda K, Miyamoto H, Taniguchi H (2009) A host-vector system for molecular study of the intracellular growth of *Mycobacterium tuberculosis* in phagocytic cells. Microbiol Immunol 53: 550–558. doi: 10.1111/j.1348-0421.2009.00158.x PMID: 19780968
2. Fujiwara N, Naka T, Ogawa M, Yamamoto R, Ogura H, Taniguchi H (2012) Characteristics of Mycobacterium smegmatis J15cs strain lipids. Tuberculosis (Edinb) 92: 187–192. doi: 10.1016/j.tube.2011.10.001 PMID: 22056691
3. Chatterjee D, Kho KH (2001) The surface glycopeptidolipids of mycobacteria: structures and biological properties. Cell Mol Life Sci 58: 2018–2042. PMID: 11814054
4. Howard ST, Rhoades E, Recht J, Pang X, Alsup A, Kolter R, et al. (2006) Spontaneous reversion of *Mycobacterium abscessus* from a smooth to a rough morphotype is associated with reduced expression of glycopeptidolipid and reacquisition of an invasive phenotype. Microbiology 152: 1581–1590. PMID: 16735722
5. Kocincova D, Singh AK, Beretti JL, Ren H, Euphrasie D, Liu J, et al. (2008) Spontaneous transposition of IS1096 or ISMsm3 leads to glycopeptidolipid overproduction and affects surface properties in *Mycobacterium smegmatis*. Tuberculosis (Edinb) 88: 390–398. doi: 10.1016/j.tube.2008.02.005 PMID: 18439873
6. Ripoll F, Deshayes C, Pasek S, Laval F, Beretti JL, Biet F, et al. (2007) Genomics of glycopeptidolipid biosynthesis in *Mycobacterium abscessus* and *M. chelonae*. BMC Genomics 8: 114. PMID: 17490474
7. Matsuo K, Yamaguchi R, Yamazaki A, Tasaka H, Terasaka K, Yamada T (1990) Cloning and expression of the gene for the cross-reactive alpha antigen of *Mycobacterium kansasii*. Infect Immun 58: 550–556. PMID: 2404875
8. Ohara N, Nishiyama T, Ohara-Wada N, Matsumoto S, Matsuo T, Yamada T (1997) Characterization of the transcriptional initiation regions of genes for the major secreted protein antigens 85C and MPB51 of *Mycobacterium bovis* BCG. Microb Pathog 23: 303–310. PMID: 9405208
9. Naka T, Maeda S, Niki M, Ohara N, Yamamoto S, Yano I, et al. (2011) Lipid phenotype of two distinct subpopulations of *Mycobacterium bovis* Bacillus Calmette-Guerin Tokyo 172 substrains. J Biol Chem 286: 44153–44161. doi: 10.1074/jbc.M111.310537 PMID: 22033395
10. Sonden B, Kocincova D, Deshayes C, Euphrasie D, Khayat L, Laval F, et al. (2005) Gap, a mycobacterial specific integral membrane protein, is required for glycolipid transport to the cell surface. Mol Microbiol 58: 426–440. PMID: 16194230
11. Miyamoto Y, Mukai T, Nakata N, Maeda Y, Kai M, Naka T, et al. (2006) Identification and characterization of the genes involved in glycosylation pathways of mycobacterial glycopeptidolipid biosynthesis. J Bacteriol 188: 86–95. PMID: 16352824
12. Gokhale RS, Saxena P, Chopra T, Mohanty D (2007) Versatile polyketide enzymatic machinery for the biosynthesis of complex mycobacterial lipids. Nat Prod Rep 24: 267–277. PMID: 17389997
13. Biet F, Bay S, Thibault VC, Euphrasie D, Grayon M, Ganneau C, et al. (2008) Lipopentapeptide induces a strong host humoral response and distinguishes Mycobacterium avium subsp. paratuberculosis from M. avium subsp. avium. Vaccine 26: 257–268. PMID: 18068277

14. Billman-Jacobe H, McConville MJ, Haites RE, Kovacevic S, Coppel RL (1999) Identification of a peptide synthetase involved in the biosynthesis of glycopeptidolipids of Mycobacterium smegmatis. Mol Microbiol 33: 1244–1253. PMID: 10510238

15. Davidson LB, Nessar R, Kemppainen P, Perkins DJ, Byrd TF (2011) Mycobacterium abscessus glycopeptidolipid prevents respiratory epithelial TLR2 signaling as measured by H\(\beta\)D2 gene expression and IL-8 release. PLoS One 6: e29148. doi: 10.1371/journal.pone.0029148 PMID: 21856857

16. Naka T, Nakata N, Maeda S, Yamamoto R, Doe M, Mizuno S, et al. (2011) Structure and host recognition of serotype 13 glycopeptidolipid from Mycobacterium intracellulare. J Bacteriol 193: 5766–5774. doi: 10.1128/JB.05412-11 PMID: 21856857

17. McCarthy TR, Torrelles JB, MacFarlane AS, Katabcik M, Kutzbach B, Desjardins LE, et al. (2011) Overexpression of Mycobacterium tuberculosis manB, a phosphomannomutase that increases phosphatidylinositol mannoside biosynthesis in Mycobacterium smegmatis and mycobacterial association with human macrophages. Mol Microbiol 58: 774–790. PMID: 16238626

18. Bernut A, Herrmann JL, Kiss K, Dubremetz JF, Gaillard JL, Lutfalla G, et al. (2014) Mycobacterium abscessus cording prevents phagocytosis and promotes abscess formation. Proc Natl Acad Sci USA 111: E943–952. doi: 10.1073/pnas.1321390111 PMID: 24567393

19. Catherinot E, Clarissou J, Etienne G, Ripoll F, Emile JF, Daffe M, et al. (2007) Hypervirulence of a rough variant of the Mycobacterium abscessus type strain. Infect Immun 75: 1055–1058. PMID: 17145991

20. Roux AL, Ray A, Pawlik A, Medjahed H, Etienne G, Rottman M, et al. (2011) Overexpression of proinflammatory TLR-2-signalling lipoproteins in hypervirulent mycobacterial variants. Cell Microbiol 13: 692–704. doi: 10.1111/j.1462-5822.2010.01565.x PMID: 21143571

21. Etienne G, Villeneuve C, Dinadayala P, Abouwarda A, Zerbib D, et al. (2005) The cell envelope structure and properties of Mycobacterium smegmatis mc\(^{+}\)55: is there a clue for the unique transformability of the strain? Microbiology 151: 2075–2086. PMID: 15942014

22. Etienne G, Villeneuve C, Billman-Jacobe H, Astarie-Dequeker C, Dupont MA, Daffe M (2002) The impact of the absence of glycopeptidolipids on the ultrastructure, cell surface and cell wall properties, and phagocytosis of Mycobacterium smegmatis. Microbiology 148: 3089–3100. PMID: 12368442

23. Jeong BH, Kim SY, Jeon K, Lee SY, Shin SJ, Koh WJ (2013) Serodiagnosis of Mycobacterium avium complex and Mycobacterium abscessus complex pulmonary disease by use of IgA antibodies to glycopeptidolipid core antigen. J Clin Microbiol 51: 2747–2749. doi: 10.1128/JCM.00702-13 PMID: 23740728

24. Kitada S, Levin A, Hiserote M, Harbeck RJ, Czaja CA, Huit G, et al. (2013) Serodiagnosis of Mycobacterium avium complex pulmonary disease in the USA. Eur Respir J 42: 454–460. doi: 10.1183/09031936.00098212 PMID: 23105006

25. Anuchin AM, Mulyukin AL, Suzina NE, Duda VI, El-Registan GI, Krapelyants AS (2009) Dormant forms of Mycobacterium smegmatis with distinct morphology. Microbiology 155: 1071–1079. doi: 10.1099/mic.0.029302-0 PMID: 19326099

26. Schorey JS, Sweet L (2008) The mycobacterial glycopeptidolipids: structure, function, and their role in pathogenesis. Glycobiology 18: 832–841. doi: 10.1093/glycob/cwn076 PMID: 18723691