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Structure and biosynthesis of free lipid A molecules that replace lipopolysaccharide in *Francisella novicida*

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Abstract

Francisella novicida U112 phospholipids, extracted without hydrolysis, consist mainly of phosphatidylethanolamine, phosphatidylglycerol, phosphatidylcholine, and two lipid A species, designated A1 and A2. These lipid A species, present in a ratio of 7:1, comprise 15 % of the total phospholipids, as judged by ³²P_i labeling. Although lipopolysaccharide is detectable in *F. novicida* U112, less than 5 % of the total lipid A is covalently linked to it. A1 and A2 were analyzed by electrospray ionization and matrix-assisted laser desorption ionization mass spectrometry, gas chromatography/mass spectrometry and NMR spectroscopy. Both compounds are disaccharides of glucosamine, acylated with primary 3-hydroxystearoyl chains at positions 2, 3, and 2', and a secondary palmitoyl residue at position 2'. Minor isobaric species and some lipid A molecules containing a 3-hydroxypalmitoyl chain in place of 3-hydroxystearate are also present. The 4'- and 3'-positions of A1 and A2 are not derivatized, and Kdo is not detectable. The 1-phosphate groups of both A1 and A2 are modified with an α -linked galactosamine residue, as shown by NMR spectroscopy and gas chromatography/mass spectrometry. An α -linked glucose moiety is attached to the 6'-position of A2. The lipid A released by mild acid hydrolysis of *F. novicida* lipopolysaccharide consists solely of component A1. *F. novicida* mutants lacking the *arnT* gene do not contain a galactosamine residue on their lipid A. Formation of free lipid A in *F. novicida* might be initiated by an unusual Kdo hydrolase present in the membranes of this organism.

Lipopolysaccharide (LPS) makes up the outer leaflet of the outer membranes of most Gram-negative bacteria (1-3). It consists of a hydrophobic moiety known as lipid A, a non-repeating core oligosaccharide, and a distal repeating oligosaccharide, termed the O-antigen (1-3). Lipid A of wild-type *E. coli* is a hexa-acylated disaccharide of glucosamine that is phosphorylated at the 1- and 4'-positions (1-3) (Fig. 1A). It is recognized by the TLR4/MD2 receptor of the innate immune system (4-8), which triggers an inflammatory response and helps to clear localized infections. However, a more generalized response to lipid A in the context of a systemic infection, accompanied by massive over-production of cytokines, can lead to Gram-negative septic shock and death (9,10).

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SUPPORTING INFORMATION AVAILABLE

The NOESY analysis of HA2 and the HMBC spectra of HA1 and HA2 are available free of charge via the Internet at <http://pubs.acs.org>.

Wild-type *E. coli* cells contain ~ 0.15 lipid A residues per glycerophospholipid molecule (11). All of the lipid A residues are covalently attached to LPS, whereas free lipid A and lipid A precursors are not usually detectable by $^{32}\text{P}_i$ labeling (12) or mass spectrometry (MS) (13). Heptose-deficient mutants of *E. coli* are viable but synthesize a minimal LPS substructure consisting of Kdo₂-lipid A, which is fully active as a TLR4/MD2 agonist (14). However, free lipid A and lipid A precursors do not accumulate in these mutants (15).

Francisella tularensis, a highly infectious Category A human pathogen, causes tularemia and is of concern because of its potential for bio-terrorism (16). The LPS of *F. tularensis* has been isolated (17), and the structures of its O-antigen, core and lipid A domains have been partially characterized (18-22). Some differences and ambiguities in the structures reported for *F. tularensis* lipid A are noteworthy (Fig. 1). For instance, Vinogradov *et al.* claimed that lipid A from the live vaccine strain (LVS) of *F. tularensis* is not phosphorylated at either the 1 or the 4' positions (Fig. 1B) (18). In contrast, Phillips *et al.* reported the presence of a phosphate group at the 1 position of *F. tularensis* LVS lipid A (Fig. 1C) and, in addition, described an unprecedented galactosamine phosphate substituent at the 1 position of lipid A isolated from *F. tularensis* 1547-57 (Fig. 1D) (22). Because NMR spectroscopy was not utilized by Phillips *et al.* (22), the anomeric configurations of the galactosamine and the proximal glucosamine residues could not be determined (Figs. 1C and 1D).

Strains of *Francisella* appear to make less LPS than *E. coli*, as judged by phenol-water extraction (18,23), and the bioactivity of *Francisella* LPS is very low (24,25). We now report that *F. tularensis* subsp. *novicida* U112, an attenuated mouse pathogen, does in fact synthesize lipid A in amounts comparable to *E. coli* (11) but that most of this material is not glycosylated with core or O-antigen sugars. Consequently, it is not recovered by the traditional phenol-water method for purifying LPS (18). Instead, the *F. novicida* glycerophospholipid fraction, extracted with chloroform/methanol according to the method of Bligh and Dyer (13,26), contains two prominent lipid A species, designated A1 and A2. These substances were purified and analyzed by MS and NMR spectroscopy. The ratio of A1 to A2 is about 7:1. Both A1 and A2 are disaccharides of glucosamine, acylated predominantly with 3-hydroxystearoyl chains at the 2, 3, and 2'-positions, and a secondary palmitoyl chain at the 2'-position (Fig. 1). Both A1 and A2 contain a phosphate moiety at the 1-position, which is further derivatized with a galactosamine residue, as confirmed by gas chromatography/mass spectrometry (GC/MS). NMR studies demonstrate unequivocally that the anomeric carbons of the galactosamine and proximal glucosamine units of A1 and A2 have the alpha configurations. We further show that the incorporation of the galactosamine residue into *F. novicida* lipid A is dependent upon its *arnT* gene orthologue (27,28). Neither A1 nor A2 contains a phosphate group at the 4'-position, presumably because of the periplasmic 4'-phosphatase encoded by *lpxF* in *Francisella* (29). The difference between A1 and A2 is the presence of an unusual α -linked glucose moiety at the 6'-position in A2. While compound A1 is probably identical to the lipid A species released by mild acid hydrolysis from *F. tularensis* LPS (Fig. 1D), a glucose-modified derivative of *F. tularensis* lipid A has not been described.

The discovery of large quantities of free lipid A associated with low levels of LPS in wild-type *F. novicida* is without precedent in LPS biochemistry and might play a role in the pathogenesis of *Francisella* infections. We show that lipid A from the LVS strain of *F. tularensis* also is present mostly in the free form. We propose that free lipid A may arise from nascent LPS by a novel system of extra-cellular remodeling enzymes, which include a Kdo hydrolase (15), the 4'-phosphatase *LpxF* (29), a galactosamine transferase encoded by *F. novicida* *arnT*, and a 3'-O-deacylase (Scheme 1).

MATERIALS AND METHODS

Materials and reagents

Glass backed 0.25 mm Silica Gel 60 thin layer chromatography plates were from E. Merck, Darmstadt, Germany. Chloroform, ammonium acetate, and sodium acetate were obtained from EMD Science, Gibbstown, NJ, while pyridine, methanol, and formic acid were from Mallinckrodt, Phillipsburg, NJ. Trypticase soy broth, yeast extract and tryptone were purchased from Difco, Detroit, MI. DEAE cellulose was purchased from Whatman, Florham Park, NJ. $^{32}\text{P}_i$ was purchased from NEN Life Science Products. Tri-Sil Reagent was from Pierce, Rockford, IL. The methanol-HCl kit was from Supelco, St. Louis, MO. D_2O , CD_3OD and CDCl_3 were from Sigma-Aldrich, St. Louis, MO. *F. novicida* U112 was obtained from Dr. F. Nano, University of Victoria, British Columbia, Canada. The *F. tularensis* LVS strain was purchased from the American Type Culture Collection, Rockville, MD.

Construction of a *F. novicida* mutant with a deletion in its *arnT* orthologue

The single orthologue of the *arnT* gene (27) present in the *F. novicida* genome (30) was amplified by PCR with 2 kb of flanking *F. novicida* genomic DNA. The forward primer (5'-CGTGCGGCCGCACTTATTACGACCTTCTTCGGTATTA-3') was designed with a clamp region, a NotI restriction site (*underlined*), and a match to the coding strand of chromosomal DNA about 2 kb upstream of *F. novicida* *arnT*. The reverse PCR primer (5'-ACGCTCGAGATGTTTGATATTGTTTTTATTAGCATATT-3') was designed with a clamp region, a XhoI restriction site (*underlined*), and a match to the anti-coding strand of chromosomal DNA about 2kb downstream of *F. novicida* *arnT*. The PCR was performed using *PfuTurbo* polymerase and *F. novicida* genomic DNA as the template. Amplification was carried out in a 100 μl reaction mixture containing 200 ng template, 200 ng primers and 2 units of polymerase. The reaction was started at 94 °C for 2 min, followed by 30 cycles of denaturation (30 seconds at 94 °C), annealing (30 seconds at 58 °C), and extension (5 min at 72 °C). After the 30th cycle, a 10 min extension time was used. The reaction product was analyzed on a 1% agarose gel. The desired band was excised and gel-purified. The PCR product was then digested using XhoI and NotI, and ligated into the expression vector pWSK29 (31), which had been similarly digested. The ligation mixture was electroporated into *E. coli* DY330 (32), and colonies were selected for those with the appropriate inserts on LB plates containing 100 $\mu\text{g/ml}$ ampicillin. Plasmid pWSK29-5000F was isolated in this manner, and the insert was confirmed by DNA sequencing.

Next, the *F. novicida* *arnT* orthologue (30) in the plasmid pWSK29-5000F was replaced with a kanamycin resistance gene. Tn903 from plasmid pWKS130 was amplified by PCR. It was flanked on its 5' and 3' ends by 50 bp of the chromosomal DNA that is immediately upstream or downstream of *F. novicida* *arnT*. The forward and reverse primers are 5'-TAATTC~~CCGCTAAATTTATTCCATACTTTTGCCAGATAAATACTACTATCATGAGCCATATTCAACGGGAAACG~~-3' and 5'-TAATCTAAAGATCATAAAAAAATAGTTAG-TTCTGTTTATTTTAAATCAGTTAGAAAAACT-CATCAAGCATCAAATG-3', respectively. The sequences belonging to the kanamycin resistance cassette are underlined. The PCR was performed using *Pfu* polymerase. Amplification was carried out in a 100 μl reaction mixture containing 100 ng of template, 250 ng primers and 2 units of *Pfu* polymerase. The reaction was started at 94 °C for 1 min, followed by 25 cycles of denaturation (45 seconds at 94 °C), annealing (45 seconds at 55 °C), and extension (60 seconds at 72 °C). After the 25th cycle, a 10 min extension time was used. The PCR product was resolved on a 1.0% agarose gel and purified. The purified PCR product was electroporated into *E. coli* DY330/pWSK29-5000F, which had been grown at 30 °C until A_{600} reached 0.5 and then induced at 42 °C for 20 min. After electroporation, recombination occurred between the homologous

sequences on the linear PCR product and the *arnT* gene on the plasmid, resulting in replacement of the *arnT* orthologue with the kanamycin resistance gene. The desired plasmid, designated pWSK29-5000K, was then isolated from kanamycin-resistant transformants, selected on LB agar containing 20 µg/ml kanamycin and 100 µg/ml ampicillin, and digested with *Apa*I and *Sac*II. About 100 ng of the linear insert DNA was gel-purified, and the replacement of *F. novicida arnT* with the kanamycin resistance gene was verified by DNA sequencing.

The final step was to replace the *arnT* orthologue present in the genome of *F. novicida* with the kanamycin resistance gene. *F. novicida* U112 cells were grown in 100 ml of Chamberlain's medium (33) to A₆₀₀ of 0.5. Cells were harvested by centrifugation and resuspended in 5 ml of transformation buffer (34). Next, 1 ml of the cell suspension was mixed with 100 ng of the linear insert DNA, prepared as described above, and shaken at 100 rpm at 37 °C for 30 min. Then 5 ml of Chamberlain's medium was added, and the mixture was shaken at 37 °C for 2 hrs at 250 rpm. The cells were harvested and resuspended in 1 ml of TSB-C medium (3% trypticase soy broth and 0.1% cysteine). Approximately 100 µl of the cell suspension was spread onto a TSB-C plate containing 10 µg/ml kanamycin. Genomic DNA was isolated from a kanamycin-resistant transformant, and the replacement of the *F. novicida arnT* gene with the kanamycin resistance gene was confirmed by DNA sequencing.

Isolation of Phospholipids and Lipid A from ³²P Labeled Cells

F. novicida was grown in TSB-C medium, and *E. coli* was grown in LB broth (1% tryptone, 0.5% yeast extract and 1% NaCl). Typically, 20 ml cells, inoculated from an overnight culture to A₆₀₀ = 0.02, were grown in the presence of 5 µCi/ml ³²P_i to A₆₀₀ of 1.2. The cells were collected by centrifugation and washed with phosphate-buffered saline (35). The cell pellets were resuspended in 3 ml of a single-phase Bligh-Dyer mixture (26) consisting of chloroform, methanol and water (1:2:0.8, v/v/v), incubated at room temperature for 60 min, and centrifuged to remove insoluble debris. The supernatant (containing mostly glycerophospholipids and the free lipid A) was removed and converted to a two-phase Bligh-Dyer system (26) by adding chloroform and water to generate a mixture consisting of chloroform, methanol and water (2:2:1.8, v/v/v). The insoluble debris pellets from the initial extractions (containing intact LPS, proteins and nucleic acids) were resuspended in 3 ml of 12.5 mM sodium acetate, pH 4.5, with 1% SDS, heated at 100 °C for 30 min (36), cooled to room temperature and then converted to a two-phase Bligh-Dyer system by the addition of chloroform and methanol.

The solvent phases from both sets of extractions were individually mixed and separated by low-speed centrifugation. The upper phases were washed once with a fresh pre-equilibrated Bligh-Dyer lower phases. The appropriate lower phases were pooled and dried under a stream of nitrogen. The ³²P-labeled lipids were re-dissolved in chloroform and methanol (4:1, v/v) and spotted onto a Silica Gel 60 TLC plate (10,000 cpm per lane), which was developed with chloroform, pyridine, 88% formic acid and water (50:50:16:5, v/v/v/v). After drying, the plates were exposed to a PhosphorImager Screen overnight, and the ³²P-labeled lipids were detected with a Molecular Dynamics Storm PhosphorImager.

Large Scale Purification of Free Lipid A from *F. novicida* U112

Two liters of *F. novicida* U112 cells were grown at 37 °C in TSB-C, harvested by centrifugation, and washed with phosphate-buffered saline. About 10 g wet cells were obtained. The cells were extracted for 1 hr at room temperature with 1.8 l of a single-phase Bligh-Dyer mixture and centrifuged to remove insoluble debris. The initial supernatant (containing most of the glycerophospholipids and free lipid A) was converted to a two-phase Bligh-Dyer system. The debris pellet (containing LPS) was washed once with a single-phase Bligh-Dyer mixture and then resuspended in 360 ml of 12.5 mM sodium acetate, pH 4.5, containing 1% SDS (36). The suspension was heated in a boiling water bath for 30 min to release covalently linked

lipid A from intact LPS (36). The cooled suspension was converted to a two-phase Bligh-Dyer mixture. The lower phases from the initial supernatant and the hydrolyzed LPS were dried separately by rotary evaporation. About 260 mg lipids were obtained from the initial supernatant, and 4 mg of lipids were obtained from the hydrolyzed debris pellets.

The dried lipids from the initial supernatant and the debris pellets were dissolved in chloroform, methanol and water (2:3:1, v/v/v). Each sample was applied to a 4 ml DEAE-cellulose column in the acetate form equilibrated with the same solvent (36,37). The run-through was saved. Each column was washed with 10 column volumes of chloroform, methanol and water (2:3:1, v/v/v). The various lipid components were then eluted stepwise with 5 column volumes each of chloroform, methanol and ammonium acetate (2:3:1, v/v/v) with ammonium acetate concentrations of 60 mM, 120 mM, 240 mM, 360 mM and 480 mM (36). Fractions equal to one column volume were collected, and 20 μ l of each fraction was spotted onto a TLC plate to monitor the lipid A and phospholipid elution profile. The plates were developed in the solvent of chloroform, methanol, acetic acid and water (25:15:4:4 v/v/v/v), and the lipids were visualized by spraying the plates with 10% sulfuric acid in ethanol, followed by charring. All *F. novicida* lipid A species were found to emerge in the DEAE cellulose run-through together with phosphatidylethanolamine and phosphatidylcholine, as judged by TLC and ESI/MS. The run-through fractions were converted to two-phase Bligh-Dyer mixtures. The lower phases were recovered and dried by rotary evaporation. At this stage, 100 mg lipids were recovered from the initial supernatant, and 2 mg were obtained from the hydrolyzed debris pellets.

To purify the lipid A species further, preparative thin layer chromatography was employed. Lipids obtained from the run-through fractions of the DEAE column were dissolved in chloroform, methanol (4:1, v/v) and applied to a TLC plate, which was developed with chloroform, methanol, pyridine, acetic acid, water (25:10:5:4:3, v/v/v/v/v). While the plates were drying, the lipid A bands could be seen transiently as white zones. These bands were marked with a pencil and scraped off after the plates were dry. The silica chips were extracted with a single-phase Bligh-Dyer mixture for 1 hour at room temperature. The suspension was centrifuged. The supernatant was passed through a small column fitted with a small glass wool plug and converted into a two-phase Bligh-Dyer system. The two phases were separated by centrifugation, and the lower phase was dried. About 9.8 mg of lipid A component A1 and 1.4 mg of component A2 were purified from the initial supernatant, but only ~ 0.1 mg of A1 was recovered from the hydrolyzed debris pellets.

Lipid A species from the *arnT* knockout mutant were isolated by the same procedure described above, except that the cells were grown at 30 °C rather than 37 °C, and the lipid A was retained by the DEAE cellulose column, given the loss of the galactosamine substituent in the mutant.

To remove the ester-linked 3-hydroxyacyl chains of A1 and A2, the purified compounds were treated with 10% aqueous triethylamine for 4 hours at room temperature. The partially deacylated lipids, designated HA1 and HA2, were extracted from the reaction mixture by the Bligh-Dyer method and purified by preparative thin layer chromatography.

MALDI/TOF MS

Matrix-assisted laser desorption ionization time-of-flight (MALDI/TOF) mass spectra were acquired using an AXIMA-CFR from Kratos Analytical (Manchester, UK), equipped with a nitrogen laser (337 nm), 20 kV extraction voltage, and time delayed extraction. The samples were prepared for MALDI/TOF MS analysis by depositing 0.3 μ l of the lipid sample dissolved in chloroform and methanol (4:1, v/v), followed by adding 0.3 μ l of a saturated solution of 6-aza-2-thiothymine in 50% acetonitrile and 10% tribasic ammonium citrate (9:1, v/v) as the matrix. The samples were left to dry at room temperature before the mass spectra were acquired in linear mode. Each spectrum was the average of 100 laser shots.

Electrospray Ionization (ESI) MS and MS/MS Analysis

All MS/MS spectra were acquired on a QSTAR XL quadrupole time-of-flight tandem mass spectrometer (ABI/MDS-Sciex, Toronto, Canada) equipped with an ESI source. Lipid A samples were dissolved in chloroform and methanol (2:1, v/v) at about 25 $\mu\text{g}/\text{ml}$ and subjected to ESI/MS in the negative ion mode (14). Nitrogen was used as collision gas for MS/MS experiments (14). Data acquisition and analysis were performed using the instrument's Analyst QS software.

GC/MS Analysis

The purified lipid A species of *F. novicida* were hydrolyzed in acidic methanol, *N*-acetylated, and then converted to trimethylsilyl ethers. Mannose, galactose, glucosamine, galactosamine and glucose standards were processed in parallel with the lipid A samples. Typically, 0.5-1.0 mg sample was thoroughly dried in a Reacti-vial equipped with a Teflon-lined screw cap. Samples were hydrolyzed by adding 300 μl of 1 M HCl in methanol and heated at 80 $^{\circ}\text{C}$ for 15 hours. The reaction mixtures were then cooled, and the solvents were removed under a stream of nitrogen. Next, 200 μl of anhydrous methanol, 40 μl of pyridine, and 40 μl of acetic anhydride were added to the vial. The reaction mixtures were mixed and incubated overnight at room temperature. The solvents were evaporated under a stream of nitrogen. Finally, silylation of free OH groups was achieved by adding 200 μl of Tri-Sil reagent to the dried samples, mixing and incubating at room temperature for an hour. The samples were dried under a gentle stream of nitrogen, and re-dissolved in 100 μl of hexane and transferred to new vials for GC/MS analysis.

GC/MS was performed using a Finnigan Trace MS coupled with a Trace GC 2000 gas chromatography. The column is a 30 m RTX-5MS (0.25 mm internal diameter and 0.25 mm phase thickness) from Restek (Bellefonte, PA). The column oven temperature was held at 100 $^{\circ}\text{C}$ for 3 min, increased to 325 $^{\circ}\text{C}$ at a rate of 20 $^{\circ}\text{C}/\text{min}$, and then held at 325 $^{\circ}\text{C}$ for 3 min. The injector was operated in the split mode (1:20 split), and the temperature of the injection port was kept at 200 $^{\circ}\text{C}$. Helium was the carrier gas with a constant flow rate of 1 ml/min. The instrument was operated in the electron ionization mode with the electron energy set at 70 eV.

NMR spectroscopy

Typically, 1-2 mg samples of *F. novicida* lipid A were dissolved in 0.3 ml of CDCl_3 , CD_3OD and D_2O (2:3:1, v/v/v) in a 3 mm NMR tube. Proton and carbon chemical shifts are reported relative to internal tetramethylsilane (TMS) at 0.00 ppm. The ^2H signal of CD_3OD was used as a field frequency lock with the residual signal of CD_3OD serving as the secondary reference at 49.5 ppm for carbon spectra. ^1H NMR spectra were recorded at the Duke NMR Center on Varian Inova 800 or 600 NMR spectrometers, equipped with Varian cryogenic probes. ^1H NMR spectra at 800 MHz were obtained with a 7.2 kHz spectral window, a 67° pulse field angle (4.5 μsec), a 4.5 sec acquisition time, and a 1 sec relaxation delay. The spectra were digitized using 64k points to obtain a digital resolution of 0.225 Hz/pt. Two-dimensional NMR experiments (COSY, NOESY, TOCSY, HMQC and HMBC) were performed at 800 MHz as previously described (38-40). Zero-quantum correlation (ZQCOSY) 2D NMR was coded into Varian NMR software based on Mueller's sequence (41). ^1H -decoupled ^{31}P NMR spectra were recorded at 202.3 MHz on a Varian Inova 500 spectrometer with a spectral window of 12143.3 Hz digitized into 25,280 data points (digital resolution of 1Hz/point or ~ 0.005 ppm/point), a 60° pulse flip angle (8 μs), and a 1.6-s repeat time. ^{31}P chemical shifts were referenced to 85% H_3PO_4 at 0.00 ppm. Inverse decoupled difference spectra were recorded as ^1H -detected ^{31}P -decoupled heteronuclear NMR experiments, as previously described (38,40).

RESULTS

Free Lipid A Among the Phospholipids of *F. novicida*

F. novicida U112 and *E. coli* W3110 were labeled uniformly with $^{32}\text{P}_i$ until late log phase. Lipids were extracted directly from intact cells by the method of Bligh and Dyer, or after pH 4.5 hydrolysis at 100 °C of the intact LPS present in the Bligh-Dyer insoluble debris pellet, as previously described (36). Radioactive lipids were quantified by TLC and PhosphorImager analysis. *E. coli* phospholipids were recovered by the direct extraction (Fig. 2, lane 1), whereas *E. coli* lipid A was released from the insoluble debris pellet only after hydrolysis at pH 4.5 (36) (data not shown). Hydrolyzing the *F. novicida* debris pellet at pH 4.5 yielded very little lipid A (about 100 times less than from *E. coli*). However, several unknown radioactive substances were recovered together with the glycerophospholipids by direct Bligh-Dyer extraction of *F. novicida* cells (Fig. 2, lane 2). The rapidly-migrating lipids (Fig. 2, lane 2) are phosphatidylethanolamine, phosphatidylglycerol, cardiolipin and phosphatidylcholine, as judged by ESI/MS/MS (not shown). The slowly-migrating unknowns, designated compounds A1 and A2 (Fig. 2, lane 2), were purified on a larger scale and confirmed as lipid A derivatives by MS and NMR spectroscopy (see below). A1 and A2 make up 15 - 20% of the total ^{32}P -labeled lipids recovered by direct extraction of intact *F. novicida* cells, as judged by PhosphorImager analysis. This is about the same ratio of lipid A to glycerophospholipids as is found in *E. coli*, except that *E. coli* lipid A must first be released from LPS by mild acid hydrolysis (11,36). The fact that over 95 % of A1 and A2 is extracted directly from *F. novicida* with a single-phase Bligh-Dyer mixture demonstrates that *F. novicida* lipid A is not covalently linked to LPS. The amount of intact LPS present in *F. novicida* is low compared to *E. coli*, where it represents about 3.4 % of the dry weight (23). As in *F. novicida*, most of the lipid A in the *F. tularensis* LVS strain is in the free form (data not shown).

To purify larger amounts of A1 and A2, 2 l of *F. novicida* U112 were grown to late log phase. Lipids were extracted from intact cells with a single-phase Bligh-Dyer mixture or after pH 4.5 hydrolysis of the extracted, insoluble debris pellet. Portions of the crude lipids were separated on a TLC plate, which was sprayed with 10% sulfuric acid and charred on a hot plate. Consistent with the ^{32}P -labeling, almost all the lipids were recovered by direct extraction of the intact cells, and only a small amount was recovered after hydrolysis of the debris pellet. The pattern of lipids on the charred plate (not shown) was virtually the same as that determined by ^{32}P -labeling (Fig. 2, lane 2), showing that *F. novicida* does not synthesize lipid A species lacking phosphate groups, as is the case in *Rhizobium etli* and *Rhizobium leguminosarum* (39, 42, 43). Only compound A1 was observed amongst the lipids released by pH 4.5 hydrolysis of the debris pellet, indicating A1 as the sole hydrophobic anchor for the intact LPS in *F. novicida* cells.

The crude lipids extracted directly from the intact cells were subjected to anion exchange chromatography on a DEAE-cellulose column (36). A1 and A2 eluted together with phosphatidylethanolamine, lyso-phosphatidylethanolamine and phosphatidylcholine in the run-through, indicating that they possess no net negative charge. Compounds A1 and A2 were then further purified by preparative TLC (Fig. 3A). From 2 l of cells, about 9.8 mg of A1 and 1.4 mg of A2 were recovered by direct extraction, but only ~ 0.1 mg of A1 was released from the Bligh-Dyer insoluble debris pellet by hydrolysis at pH 4.5.

Negative Ion MALDI/TOF MS of Compounds A1 and A2

A1 of *F. novicida* was analyzed by MALDI/TOF MS in the negative ion mode (Fig. 3B). The peaks at m/z 1665.4 and 1637.4 are interpreted as $[\text{M-H}]^-$ molecular ions, which differ by 28 amu, i.e. two methylene units. The ion at m/z 1504.4 may be derived from the base peak ion

at m/z 1665.4 by loss of a 161 amu moiety, possibly a hexosamine unit (22). The m/z 1383.3 ion is derived from the m/z 1665.4 ion by loss of a 3-hydroxystearoyl group.

The negative ion MALDI/TOF mass spectrum of A2 (Fig. 3C) shows a similar pattern as A1. The peaks at m/z 1827.5 and 1799.5 could be interpreted as molecular ions $[M-H]^-$, differing by 28 mass units. The peaks at m/z 1666.4 and 1545.7 might be derived from these parent ions by loss of a hexosamine or a 3-hydroxystearoyl moiety, respectively. All the peaks in the spectrum of A2 are 162 amu larger than the corresponding peaks of A1, suggesting that A2 contains an additional 162 amu substituent, possibly a hexose residue.

GC/MS Analysis of Compounds A1 and A2

The labile hexosamine unit, observed in the MALDI/TOF mass spectra of A1 and A2, could be attached to the 1-phosphate group, as proposed for lipid A of *F. tularensis* 1547-57 (Fig. 1D). To test this idea, A1 was hydrolyzed in 0.1 M HCl at 100 °C for 30 min. A hexosamine moiety attached to the 1-phosphate group of a lipid A molecule would be cleaved under these conditions. The hydrolysis mixture was then converted into a two-phase Bligh/Dyer system by the addition of chloroform and methanol. The phases were separated and dried. The materials in the dried upper and lower phases were hydrolyzed in acidic methanol, *N*-acetylated, converted to trimethylsilyl ethers and analyzed by GC/MS. Glucose, mannose, galactose, glucosamine and galactosamine standards were processed in parallel. The products recovered from the upper phase of the A1 hydrolysis mixture yielded peaks at 11.46 and 11.52 min, as did a galactosamine standard, confirming that the labile hexosamine unit of A1 is indeed galactosamine (data not shown). The products recovered from the lower phase yielded peaks at 10.59, 11.52 and 12.40 min (data not shown); their electron impact mass spectra were searched against the National Institute of Standards and Technology (NIST) library of mass spectra and identified as derivatives of palmitate, 3-hydroxypalmitate and 3-hydroxystearate, respectively. The presence of both 3-hydroxypalmitate and 3-hydroxystearate is consistent with the fatty acid chain length heterogeneity seen in the negative ion MALDI/TOF mass spectrum of A1 (Fig. 3B).

To characterize the additional hexose moiety present in A2, intact A1 and A2 were hydrolyzed in acidic methanol, *N*-acetylated, and converted to trimethylsilyl ethers, followed by GC/MS analysis (Fig. 4). Standards of glucose, mannose, galactose, glucosamine and galactosamine were processed in parallel. Peaks at 11.46 and 11.52 min were observed in the profiles of both A1 and A2, confirming that they both contain galactosamine. Peaks at 10.59, 11.52 and 12.40 min were observed with both A1 and A2 (Fig. 4A and B), the electron impact mass spectra of which matched with derivatives of palmitate, 3-hydroxypalmitate and 3-hydroxystearate, respectively. However, the peaks seen at 10.46 and 10.53 min in A2 (Fig. 4B) were not present in A1 (Fig. 4A), suggesting that they were derived from the additional hexose unit present in A2. Identical peaks at 10.46 and 10.53 min were seen with the glucose standard (Fig. 4C), but not with the other standards, showing that the additional hexose unit in A2 must be glucose.

ESI/MS/MS of Compounds A1 and A2

The molecular ions $[M-H]^-$ of A1 and A2 (at m/z 1665.18 and 1827.23, respectively) were further analyzed by high resolution ESI/MS/MS (Fig. 5). The peaks at m/z 78.96 $[PO_3]^-$ and 96.97 $[H_2PO_4]^-$ confirm the presence of a phosphate group. The prominent peak at m/z 258.04, observed in both spectra (Fig. 5), has the ion mass expected for a phosphorylated hexosamine, suggesting the galactosamine moiety is connected to the phosphate group (Fig. 1D). Fragment ions derived from the parent ions by neutral loss of palmitic acid were observed at m/z 1408.96 for A1 and at m/z 1570.98 for A2 (Fig. 5), respectively. Fragment ions derived by neutral loss of both hydroxystearic acid and the hexosamine unit were observed at m/z 1203.83 for A1 and m/z 1365.89 for A2, and fragment ions arising from neutral loss of palmitic acid, hydroxystearic

acid, and the hexosamine unit were observed at m/z 947.60 for A1 and m/z 1109.65 for A2. The further cleavage of an *N*-linked hydroxyacyl chain could generate the fragment ions at m/z 665.34 for A1 and m/z 827.40 for A2. Taken together with the GC/MS data (Fig. 4), the ESI/MS/MS spectra show that A1 and A2 are very similar in structure, both containing phosphate, galactosamine, palmitate and hydroxystearate residues. The phosphate and galactosamine units are connected to each other, as demonstrated by the peak at m/z 258.34.

Mild Alkaline Hydrolysis of A1 and A2

The ester-linked 3-hydroxyacyl chains of lipid A are very susceptible to mild alkaline hydrolysis (44). To determine the locations of the ester-linked fatty acids, A1 and A2 were treated with 10% aqueous triethylamine for 4 h at 25 °C. The products, HA1 and HA2, were purified and analyzed by negative ion MALDI/TOF MS. The molecular ions of HA1 and HA2 are each 282 amu smaller than those of A1 and A2, respectively (not shown), demonstrating that only one base-labile 3-hydroxystearoyl moiety is present in each compound. When analyzed by TLC in chloroform, pyridine, 88% formic acid and water (50:50:16:5, v/v/v/v), HA1 and HA2 migrate with R_f s that are similar to those of A1 and A2, respectively. These findings suggest that the base-labile 3-hydroxystearoyl moiety is attached to proximal glucosamine 3-position in both A1 and A2 (Fig. 1), as deacylation of the 3'-position usually causes a larger shift in the R_f than deacylation of the 3-position (44).

Positive Ion MALDI/TOF MS

Compounds A1, HA1, A2 and HA2 were analyzed by positive ion MALDI/TOF MS (Fig. 6). Parent ions, interpreted as $[M+H]^+$, were observed at m/z 1667.0 for A1, m/z 1384.6 for HA1, m/z 1829.0 for A2 and m/z 1547.7 for HA2 (Fig. 6). Variable amounts of sodium adducts $[M+Na]^+$ were also detected (not labeled in Fig. 6). A prominent B_1^+ ion peak (45) near m/z 683, derived from the distal glucosamine unit, was present in the spectra of both A1 and HA1 (Fig. 6A and 6B), confirming that the ester-linked 3-hydroxystearoyl moiety is attached to the proximal glucosamine unit at position 3 in A1. Prominent B_1^+ ions near m/z 845.0 were observed for both A2 and HA2 (Fig 6C and 6D), demonstrating that the base-labile 3-hydroxystearoyl acid moiety is connected to the 3-position in A2 as in A1. However, the difference in the masses of the B_1^+ ions of A1 versus A2 or HA1 versus HA2 is 162 amu, showing that the additional glucose residue present in A2 is linked to the distal glucosamine unit.

Neutral loss of the galactosamine moiety from the 1-phosphate group of each of the four compounds gave rise to product ions mainly as sodium adducts at m/z 1527.8 for A1, m/z 1245.5 for HA1, m/z 1689.9 for A2, m/z 1408.7 for HA2 (Fig. 6). The B_2^+ ions (45), which are derived from the parent ions by loss of the galactosamine 1-phosphate group, were observed at m/z 1407.8 for A1, m/z 1125.5 for HA1, m/z 1569.9 for A2, and m/z 1287.7 for HA2. The peaks at m/z 1125.4 in the A1 spectrum and at m/z 1287.5 in the A2 spectrum could arise by loss of the labile, ester-linked hydroxystearoyl moiety from the B_2^+ ion of each compound. As in the negative ion spectra (Fig. 4), some fatty acid chain length heterogeneity was observed in the positive ion spectra, though less on the distal unit, as judged by the masses of the B_1^+ ions (Fig. 6). The proposed structures of A1, HA1, A2 and HA2, which are based on the MS analysis together with the NMR data discussed below, are shown in Fig. 7.

1H and ^{31}P NMR Analysis of Compounds A1, A2, HA1 and HA2

The 800 MHz 1H NMR spectra of compounds HA1, HA2, A1 and A2 dissolved in $CDCl_3$, CD_3OD and D_2O (2:3:1, v/v/v) reveal relatively well-resolved resonances in the sugar (3.2-5.8 ppm) (Fig. 8) and acyl chain regions (0.8-2.7 ppm) (shown for HA1 in Fig. 9), similar to previous NMR spectra for *E. coli* and *R. leguminosarum* lipid A in the same solvent (38-40). The numbering scheme for the relevant protons is shown in Fig. 7D. The full assignments of

the NMR chemical shifts and coupling constants for HA1 and HA2 (Table 1) were made from 2D COSY, ZQCOSY, TOCSY, HMQC and HMBC spectra.

^{31}P NMR spectroscopy at 202 MHz of HA1, HA2, A1 and A2 revealed a single ^{31}P resonance around -2 ppm (not shown), indicating that each contains a single phosphorus atom. ^1H -observed, ^{31}P -decoupled experiments (38,40) were implemented to identify the linkage site of the phosphate group on the glucosamine disaccharide backbone. The difference ^1H NMR spectrum derived from selective on- and off-resonance decoupling of the -2.18 ppm ^{31}P signal of HA1 revealed the simultaneous simplification of the H-1 and H-1⁰ signals (at 5.46 and 5.72 ppm respectively) into two doublets, and the H-2 and H-2⁰ resonances (at 3.90 and 3.49 ppm respectively) into two double-doublets (Fig. 8A). This indicates that the single phosphate group, present in HA1, bridges the proximal glucosamine and the capping galactosamine residue, consistent with the fragments ions seen by MS. Difference ^1H NMR spectra confirmed that the single phosphate moiety present in HA2, A1 and A2 likewise connected the proximal glucosamine and capping galactosamine residue (not shown).

The 1D ^1H NMR spectra of HA1 (Fig. 8B) and A1 (Fig. 8C) reveal three resolved anomeric resonances, consistent with the presence of three sugars. However, the NMR spectra of HA2 and A2 (Fig. 8D and E, respectively) show four resolved anomeric signals, consistent with the presence of the additional glucose moiety in A2 and HA2. The α CH₂ signals of HA1 (Fig. 9) and HA2 (not shown) integrate to six protons, while the α CH₂ resonances of A1 and A2 integrate to eight protons (not shown). The integration is consistent with the presence of three acyl chains in HA1 and HA2, and four acyl chains in A1 and A2, respectively (Fig. 7). All four lipids show one esterified β -hydroxymethine proton near 5.2 ppm (Fig. 8B, 8C, 8D and 8E), consistent with the presence of a single acyloxyacyl group (Fig. 7) (38,39). The downfield shift of the H-3 signal to near 5.16 ppm in A1 and A2 (Fig. 8C and 8E) confirms the presence of an acyl chain at position 3 in these compounds (Fig. 7).

^1H - ^1H COSY and NOESY Analysis

The anomeric 5.46 ppm double-doublet (H-1) and the 4.47 ppm doublet (H-1') of compound HA1 are at similar shift positions to those of *E. coli* lipid A (38,40). The low field position and double-doublet character of H-1 are typical of an α anomeric proton in a pyranose 1-phosphate (38-40). The anomeric double-doublet at 5.72 ppm (designated H-1⁰) (Figs. 8 and 9) is not seen in *E. coli* lipid A (38). H-1⁰ connects to six other sugar resonances (Table 1 and Fig. 9), consistent with the presence of a galactosamine residue. The low field position and double-doublet character of H-1⁰ indicates that the 1⁰-position is phosphorylated (Figs. 7-9). H-4' of compound HA1 (3.33 ppm) (Table 1) resonates 0.8 ppm upfield of H-4' of *E. coli* lipid A (4.17 ppm) (38), consistent with the proposal that the 4'-position of compound HA1 is not phosphorylated (Fig. 7). The presence of *O*-acyl chains of positions 3 and 3' in *E. coli* lipid A (Fig. 1A) shifts the corresponding H-3 and H-3' signals downfield to ~5.2 ppm (38,40). In contrast, the H-3 and H-3' signals of compound HA1 appear in the sugar region at 3.72 and 3.46 ppm (Table 1), respectively, indicating that there is no *O*-acyl substituent at positions 3 and 3' of HA1. This pattern is very similar in HA2 (Table 2).

The small $J_{1,2}$, $J_{1^0,2^0}$ and $J_{1'',2''}$ couplings (3.2 - 3.7 Hz) (Table 1) provide further evidence that the proximal glucosamine unit, the galactosamine residue and the glucose residue of HA2 (Fig. 7) are in the α -anomeric configuration. Likewise, the proximal glucosamine and galactosamine units of HA1 are α -anomers. The large $J_{1',2'}$ coupling (8.6 Hz) (Table 1) shows that the distal glucosamine unit of both HA1 and HA2 is in the β -configuration (Fig. 7). The NOE enhancements from H-1' to H-6a and H-6b (and from H-1'' to H6a' and H6b') in the NOESY spectrum of HA2 (Supplementary Fig. 1) provide independent evidence for the proposed 1'-6 and 1''-6' glycosidic linkages in HA2 (Fig. 7D). The NOE enhancement from H-1' to H6a and H6b is also seen with HA1 (data not shown). The NOE enhancement from

H-1 to H-1⁰ in both HA2 (Supplementary Fig. 1) and HA1 (not shown) confirms the proposed monophosphodiester linkage between the proximal glucosamine and the capping galactosamine residues (Fig. 7C and D).

Two sets of α/β cross-peak pairs are seen in the COSY (Fig. 9) and TOCSY (not shown) spectra of *F. novicida* compound HA1, indicating the presence of only two β -hydroxyacyl chains (Fig. 7C). The cross-peak pairs near 2.3 ppm and 4.0 ppm (designated $\alpha 2/\beta 2$) arise from the α -protons adjacent to the β -oxymethine of the unsubstituted β -hydroxyacyl chain. The other α/β cross-peak pairs near 2.5 ppm and 5.2 ppm (designated $\alpha 2'/\beta 2'$) are considerably more downfield, consistent with a 2' acyloxyacyl moiety in HA1 (Fig. 7C). Our results are consistent with the acylation patterns deduced from previous NMR studies of the lipid A obtained from LPS of the LVS strain of *F. tularensis* (18).

¹³C-NMR Analysis of HA1 and HA2

The HMQC spectra of HA1 and HA2 (Fig. 10) reveal the following key features: 1) The HMQC analysis confirms the presence of three anomeric protons in HA1 and four anomeric protons in HA2 (Fig. 10A and 10B). The C-1⁰, C-1 and C-1' chemical shift positions support the proposed α -, α - and β -configurations respectively for the capping galactosamine, proximal glucosamine and distal glucosamine units in both HA1 and HA2 (46). 2) Nitrogen-substituted carbons of amino sugars usually resonate around 52 to 54 ppm (46). The HMQC spectra of compounds HA1 and HA2 (Fig. 10C and 10D) reveal three methine resonances at 54.5 ppm (C-2), 56.0 ppm (C-2') and 52.4 ppm (C-2⁰), indicating that HA1 and HA2 contain three amino sugars (Fig. 7C and D). 3) The HMQC of HA1 (Fig. 10C) confirms the presence of a glycosylated C-6 (at 72 ppm) and a nonglycosylated C-6' and C-6⁰ (signals at 61.3 and 62 ppm, respectively) (46-48). However, in the HMQC of HA2 (Fig. 10D), C6' shifts from a nonglycosylated position to a glycosylated position at 66.7 ppm, in agreement with the NOESY data (Supplementary Fig. 1), which likewise indicates that the glucose moiety present in HA2 is linked to the 6'-position.

HA1 and HA2 gave excellent HMBC responses in both the sugar and acyl chain regions (Supplementary Fig. 2). Strong cross peaks from H-1'' to C-6' and from H-6a' to C-1'' were observed in the HMBC of compound HA2 (Supplementary Fig. 2B), thus confirming the attachment of the glucose unit to the 6' position of HA2 (Fig. 7D). Linkage of a glucose residue to position 6' is without precedent in lipid A biochemistry, as Kdo is normally attached at this important site.

Origin of the Galactosamine Group in Francisella Lipid A

D-galactosamine is similar in structure and charge to 4-amino-4-deoxy-L-arabinose (Fig. 1D). The latter is synthesized and attached to lipid A in polymyxin-resistant mutants of *E. coli* and *Salmonella* (49,50). The inner membrane enzyme ArnT transfers the 4-amino-4-deoxy-L-arabinose residue from the donor substrate undecaprenyl phosphate-4-amino-4-deoxy-L-arabinose to the 1 and/or 4' phosphate groups of lipid A (27,28). We reasoned that a membrane protein with sequence similarity to ArnT might be responsible for the attachment of the galactosamine residue to lipid A in *F. novicida*. A deletion mutant of the single *arnT* orthologue present in *F. novicida* (30) synthesizes lipid A molecules lacking the galactosamine moiety found in the wild-type, as judged by the loss of the 161 amu substituent (Fig. 11A versus 11B). Loss of the galactosamine modification causes the cells to grow more slowly above 30 °C and is associated with a shift to shorter fatty acyl chains on lipid A (more of the molecular species at m/z 1476.014 versus m/z 1504.045) (Fig. 11B). Interestingly, the structure of lipid A present in this *arnT* deletion mutant of *F. novicida* is nearly identical to that reported by Phillips *et al.* for the LVS strain of *F. tularensis* (Fig. 1C) (22).

DISCUSSION

Virtually all Gram-negative bacteria, including sequenced strains of *Francisella*, encode orthologues of the enzymes that assemble Kdo₂-lipid A (3,30), the outer membrane lipid to which the other core and O-antigen sugars of LPS are attached. It is therefore remarkable that over 95 % of the lipid A present in *F. novicida* is recovered in the “free” form, *i.e.* lacking the usual Kdo, core and O-antigen sugars. Because it is not extensively glycosylated, *F. novicida* lipid A is extracted together with the glycerophospholipids by the method of Bligh and Dyer. Small amounts of LPS are indeed detectable in *F. novicida*, as well as other strains of *Francisella* (18,22), and their lipid A has been partially characterized by other investigators (Fig. 1). However, the possibility that *Francisella* lipid A was mostly present in the “free” form was not considered (17,18,22), and the glycerophospholipid fraction was not examined.¹ As shown in Fig. 2, compounds A1 and A2 make up about 15 % of the total phospholipid, the same ratio of lipid A to glycerophospholipids found in *E. coli* (11). The key distinction is that all the lipid A of *E. coli* is covalently linked to LPS and can only be released by mild acid hydrolysis at 100 °C (11,36). In wild-type *E. coli*, none of the lipid A is present in the “free” form (36), as is also the case in all other, well-characterized Gram-negative bacteria.

Component A1 (Fig. 7) of the free lipid A in *F. novicida* has the same structure as proposed by Phillips *et al.* for the lipid A moiety covalently linked to LPS in *F. tularensis* 1547-57 (Fig. 1D) (22). Compound A2 (Fig. 7) was not reported by Phillips *et al.*, consistent with our observation that it is not present in the lipid A released from *F. novicida* LPS by mild acid hydrolysis (data not shown). Given the unprecedented glucose substituent linked to the 6' position of A2, attachment of the Kdo and core sugars would be precluded (3). The GC/MS analysis (Fig. 6) validates the presence of galactosamine in both A1 and A2, and an additional glucose substituent in the latter. Our NMR data (Figs. 8-10 and Supplementary Figs.) unequivocally demonstrate the alpha anomeric configurations for the galactosamine, the glucose and the proximal glucosamine units (Fig. 7). The axial disposition of the phosphate group attached to the galactosamine moiety of A1 and A2 is the same as that seen with the 4-amino-4-deoxy-L-arabinose substituent (Fig. 1D, inset), which is attached to lipid A in polymyxin-resistant mutants of *E. coli* and *Salmonella* (40,50). Both galactosamine and 4-amino-4-deoxy-L-arabinose may neutralize the negative charge of the proximal phosphate group, thereby conferring resistance to cationic anti-microbial peptides.

The presence of all the genes encoding the enzymes that synthesize Kdo₂-lipid A (30), in conjunction with the proposed structures of A1 and A2, raises the possibility that *Francisella* cells first assemble conventional core-lipid A molecules on the inner surface of their inner membrane (Scheme 1), and then dismantle and modify them, perhaps after they are flipped to the periplasmic surface of the inner membrane by the ABC transporter MsbA (51, 52). A Kdo hydrolase (Scheme 1) that might initiate this process is present in *F. novicida* (15). The active site of the 4'-phosphatase (Scheme 1), encoded by *lpxF*, is located on the outer surface of the inner membrane (29). Mutants of *F. novicida* lacking the 4'-phosphatase² synthesize free lipid A molecules containing the 4'-phosphate residue, demonstrating that LpxF alone accounts for its removal in living cells. How the 3' acyl chain is removed (Scheme 1) to generate A1 and A2 remains unclear. In *Salmonella*, the 3' acyl chain can be cleaved by the outer membrane enzyme LpxR (53), but an LpxR orthologue sequence is not present in the *Francisella* genome. The absence of the 4'-phosphate and 3'-hydroxyacyl group in *Francisella* lipid A (Figs. 1 and 7), in conjunction with the relatively long acyl chains, may account for the low bioactivity of *Francisella* LPS (24,25).

¹Vinogradov *et al.* (18) noted the presence of a “minor amount”... of “free lipid A” in their smooth LPS preparation but did not examine the phospholipid fraction.

²X. Wang, A. A. Ribeiro, Z. Guan, S. N. Abraham and C. R. H. Raetz, in preparation.

The *Francisella* genome (30) encodes one orthologue of ArnT, the enzyme that uses an undecaprenyl phosphate linked donor substrate to transfer 4-amino-4-deoxy-L-arabinose to lipid A on the outer surface of the inner membrane in *E. coli* and *Salmonella* (27,28). Given the structural similarity of galactosamine and 4-amino-4-deoxy-L-arabinose (Fig. 1D), we inactivated the *F. novicida* *arnT* gene and examined the structure of its lipid A. As shown in Fig. 11, the galactosamine residue is absent in the lipid A of the *F. novicida* *arnT* mutant. In fact, this construct synthesizes free lipid A molecules with the structure proposed for LVS lipid A by Phillips *et al.* (Fig. 1C) (22), suggesting that the LVS strain may also harbor a mutation in its *arnT* gene. As in *F. novicida*, most of the lipid A in the LVS strain is recovered in the free form (data not shown). The absence of the galactosamine moiety might sensitize the LVS strain to some cationic anti-microbial peptides. The identification of the endogenous galactosamine donor, possibly a polyisoprene phosphate derivative by analogy to the 4-amino-4-deoxy-L-arabinose pathway (28), and the development of an *in vitro* assay for *F. novicida* ArnT (Scheme 1) should shed light on the mechanism of galactosamine transfer to lipid A. An *in vitro* assay for glucose addition to the distal residue to make A2 (Fig. 7) also remains to be developed.

Given that *F. novicida* contains a distinct lipid A 1-phosphatase, encoded by LpxE (15), it is surprising that all the lipid A molecules present in our *arnT* deletion mutant retained their 1-phosphate group (Fig. 11). Heterologous expression of *lpxE* in *E. coli* removes over 90 % of the lipid A 1-phosphate moiety (15). However, LpxE, unlike LpxF, requires the presence of the Kdo disaccharide for catalytic activity (15,29). If the rate of Kdo removal by the *Francisella* Kdo hydrolase (15) is much faster than the rate of cleavage of the 1-phosphate group by LpxE, one might not expect to see much lipid A 1-dephosphorylation. Although easily assayed *in vitro* (15), the gene encoding the *F. novicida* Kdo hydrolase and its sub-cellular localization are unknown. A Kdo hydrolase has also recently been reported to be present in strains of *Helicobacter*, but the gene encoding this enzyme is unknown (54).

An alternative explanation for the failure of LpxE to cleave the 1-phosphate group is the possibility that MsbA exports nascent lipid A more rapidly than it can be glycosylated by KdtA (3) in *F. novicida*. Accordingly, the active site of LpxE, which faces the outer surface of the inner membrane (15), would not be presented with its requisite substrate. If this scenario is correct, it should be possible to delete *kdtA* in *F. novicida* (55). This idea is consistent with the recent discovery that over-expression of MsbA in *E. coli* bypasses the normal requirement for Kdo biosynthesis (56). In such constructs, only the tetra-acylated lipid A precursor, lipid IV_A, is synthesized (56) and then flipped to the outer surface of the inner membrane by the excess MsbA (56). Consequently, no core domain is incorporated, as is the case with compounds A1 and A2 (Fig. 7) of *F. novicida*. Whatever the explanation, some kind of lipid A molecule is nevertheless required for the growth of *F. novicida*, because it is sensitive to the LpxC inhibitor CHIR-090, which blocks the second step of lipid A biosynthesis (57).

Further elucidation of the enzymology and genetics of lipid A modification in *Francisella* (Scheme 1) may prove useful for the development of effective vaccines, which are not currently available for this important pathogen (16). As noted above, *F. novicida* 4' phosphatase mutants synthesize free lipid A molecules containing the 4'-phosphate moiety.² Such mutants are hypersensitive to polymyxin, are greatly attenuated in virulence in a mouse infection model², and might provide protection against tularemia.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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ABBREVIATIONS

COSY, correlation spectroscopy
 ESI/MS, electrospray ionization/mass spectrometry
 GC/MS, gas chromatography/mass spectrometry
 HMQC, heteronuclear multiple quantum coherence spectroscopy
 HMBC, heteronuclear multiple bond coherence spectroscopy
 Kdo, 3-deoxy-D-manno-octulosonic acid
 LPS, lipopolysaccharide
 MALDI/TOF, matrix-assisted laser desorption ionization/time-of-flight
 MS, mass spectrometry
 NMR, nuclear magnetic resonance
 NOESY, nuclear Overhauser effect spectroscopy
 TLC, thin layer chromatography
 TOCSY, total correlation spectroscopy

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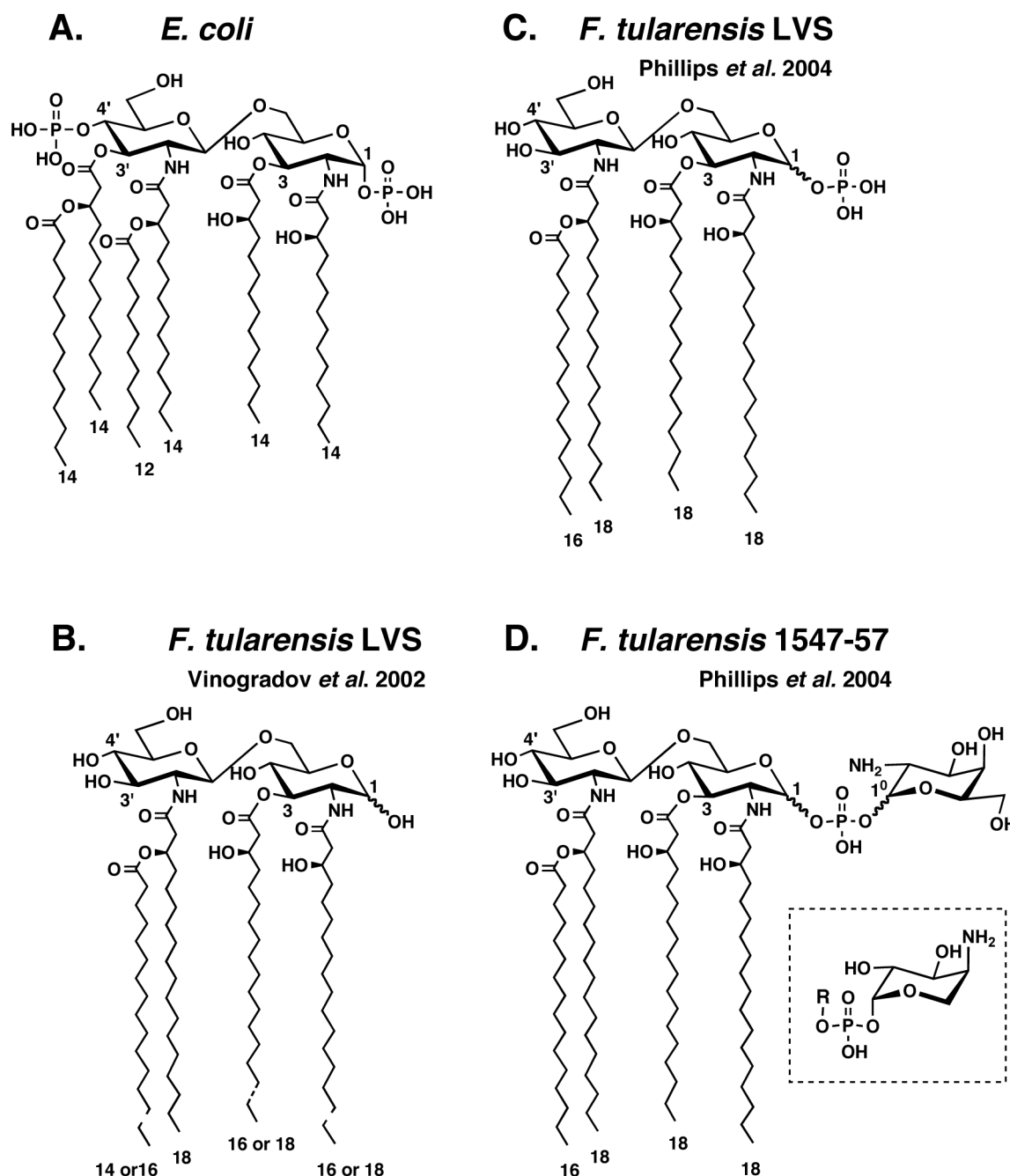


Figure 1. *E. coli* lipid A compared to structures proposed for *F. tularensis* lipid A

Panel A. The major lipid A species of wild-type *E. coli* K-12 is a hexa-acylated disaccharide of glucosamine, substituted with phosphate groups at the 1- and 4'-positions (3,38). Key positions are numbered. **Panel B.** *F. tularensis* LVS lipid A was first reported as a tetra-acylated disaccharide of glucosamine lacking both phosphate residues and the 3' acyl chain (18). Dotted lines indicate possible acyl chain length heterogeneity. **Panel C.** A more recent study of LVS lipid A demonstrated the presence of a phosphate group at the 1 position but not the anomeric configuration (22). **Panel D.** In the same study (22), the lipid A of *F. tularensis* 1547-57 was shown to be a tetra-acylated disaccharide of glucosamine substituted with a novel galactosamine-phosphate group at position 1. The anomeric configurations of the proximal

glucosamine and galactosamine-1-phosphate units were not determined (22). The inset shows the structure of the 4-amino-4-deoxy-L-arabinose-1-phosphate substituent found on lipid A in polymyxin-resistant mutants of *E. coli* and *Salmonella* (36,40,50).

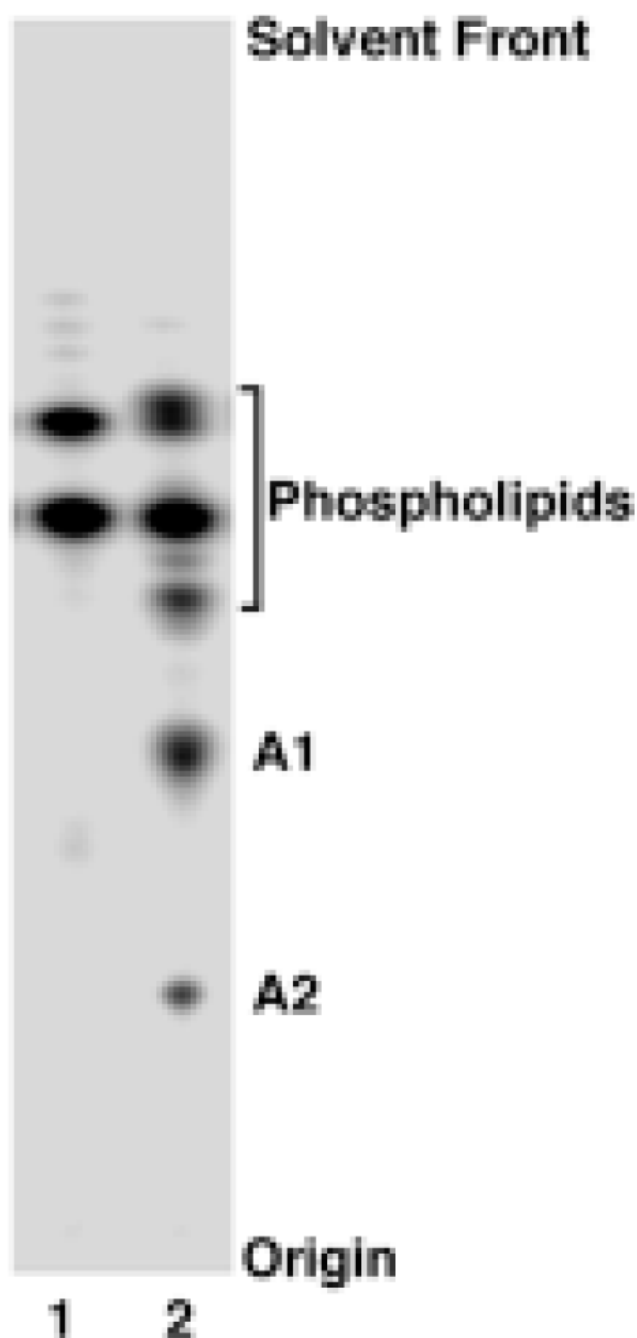


Figure 2. Phospholipids extracted from $^{32}\text{P}_i$ labeled *E. coli* versus *F. novicida* U112

Lipids were extracted from uniformly labeled *E. coli* (lane 1) or *F. novicida* (lane 2) by the method of Bligh and Dyer (26) with the solvent chloroform, methanol and water (1:2:0.8, v/v/v). Lipids were separated by TLC using chloroform, pyridine, 88% formic acid and water (50:50:16:5, v/v/v/v) and visualized with a PhosphorImager. About 10,000 cpm were loaded. Glycerophospholipids were identified by migration with standards and by ESI/MS/MS (not shown). A1 and A2, which did not match known standards, are novel free lipid A molecules, as discussed in the text.

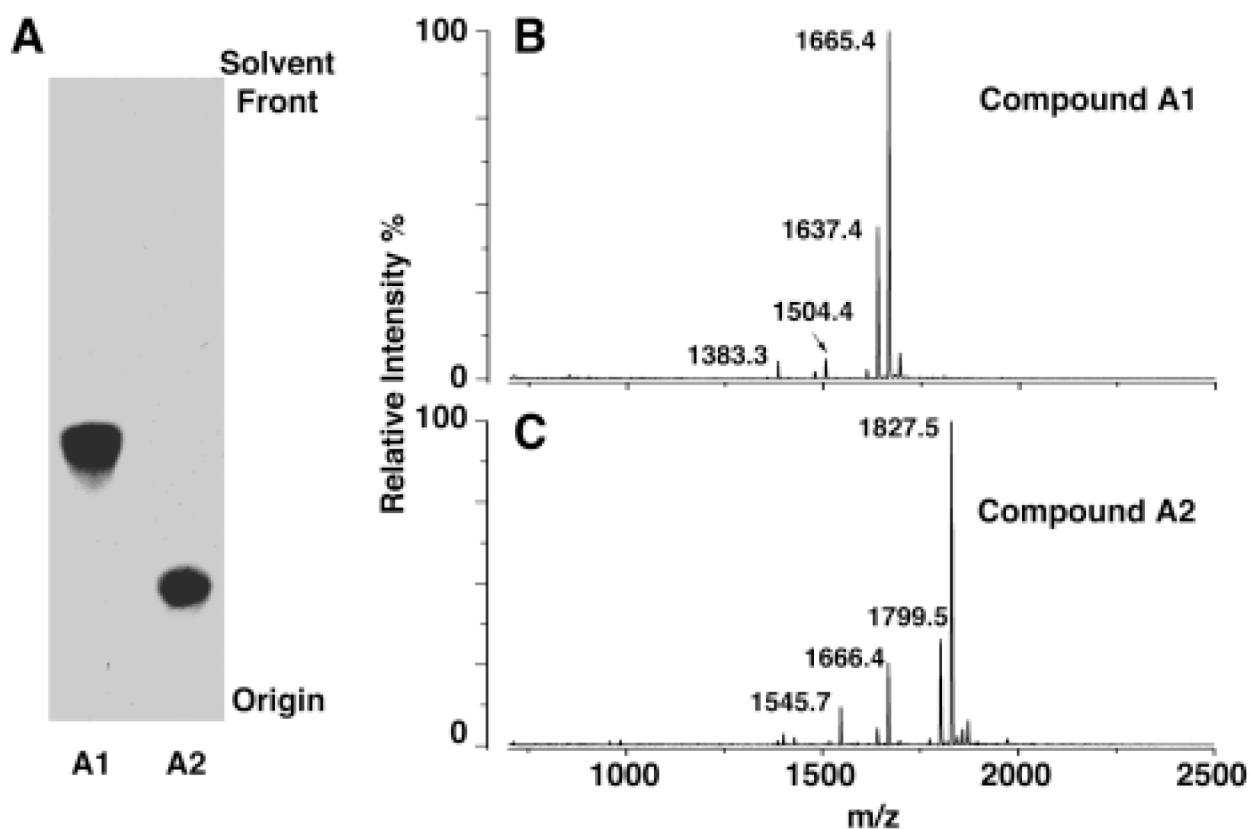


Figure 3. Purification and negative ion MALDI/TOF MS of A1 and A2

Panel A. Compounds A1 and A2 were purified from a 2 l culture of *F. novicida*, as described in the Experimental Procedures. About 10 μ g A1 and 5 μ g A2 were loaded onto a TLC plate and separated using the solvent chloroform, methanol, acetic acid and water (25:15:4:4, v/v/v/v). Lipids were detected by spraying the plate with 10 % sulfuric acids in ethanol, followed by charring. **Panel B.** Negative ion MALDI/TOF mass spectrum of compound A1. **Panel C.** Negative ion MALDI/TOF mass spectrum of A2. Some fatty acid chain length heterogeneity and partial fragmentation is evident in both spectra.

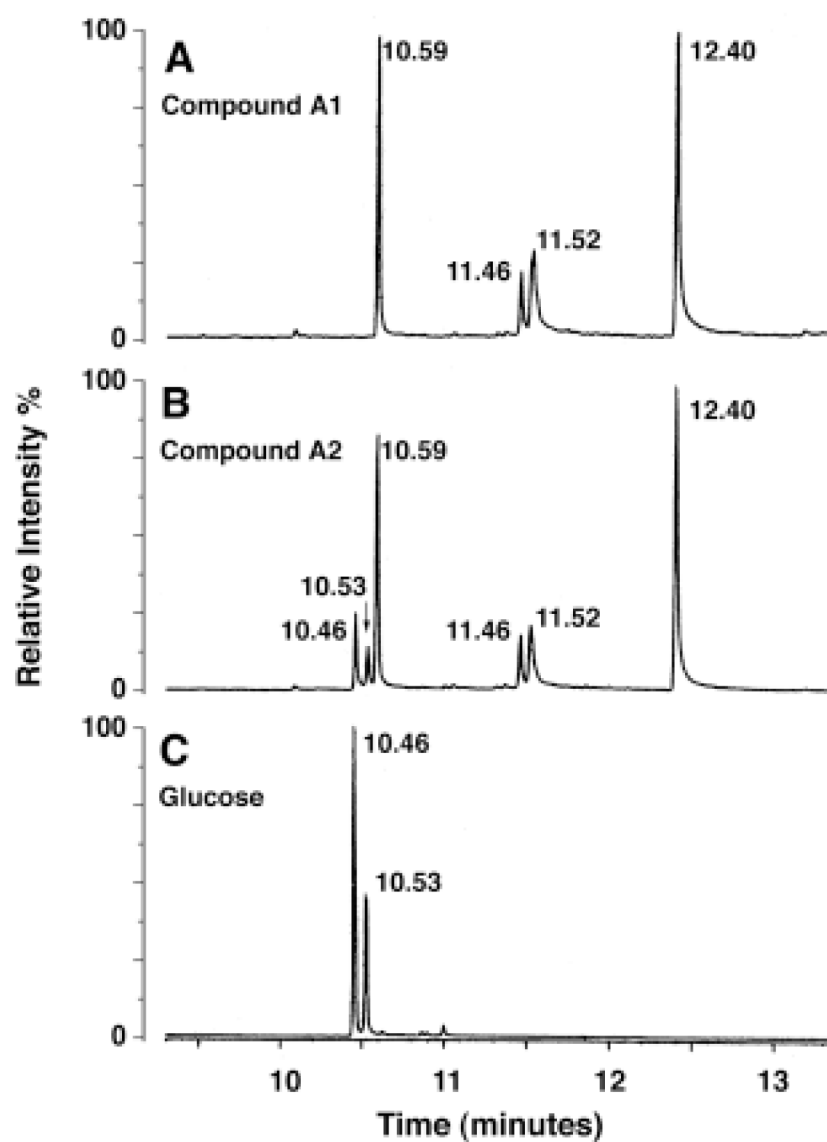


Figure 4. GC/MS analysis of compounds A1 and A2

The trimethylsilyl ether derivatives of the *N*-acetylated methyl glycosides and the fatty acid methyl esters, obtained by methanol-HCl hydrolysis of compounds A1 (**Panel A**), A2 (**Panel B**) and a glucose standard (**Panel C**), were separated and detected by electron impact ionization MS.

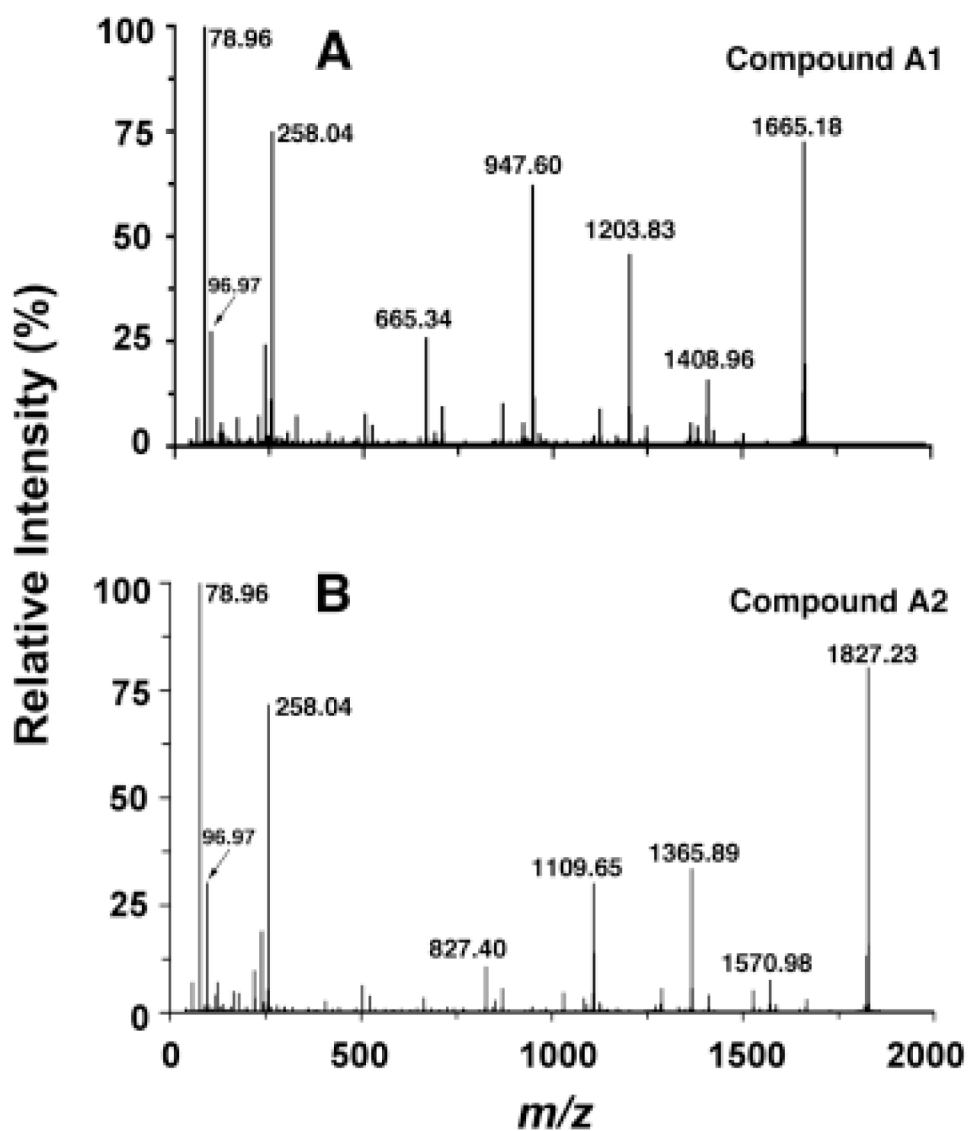


Figure 5. Negative ion ESI/MS/MS analysis of A1 and A2
Collision-induced dissociation mass spectra of $[M-H]^-$ at m/z 1665.18 (Panel A) and m/z 1827.33 (Panel B) were obtained using an ABI QSTAR-XL quadrupole time-of-flight tandem mass spectrometer (14).

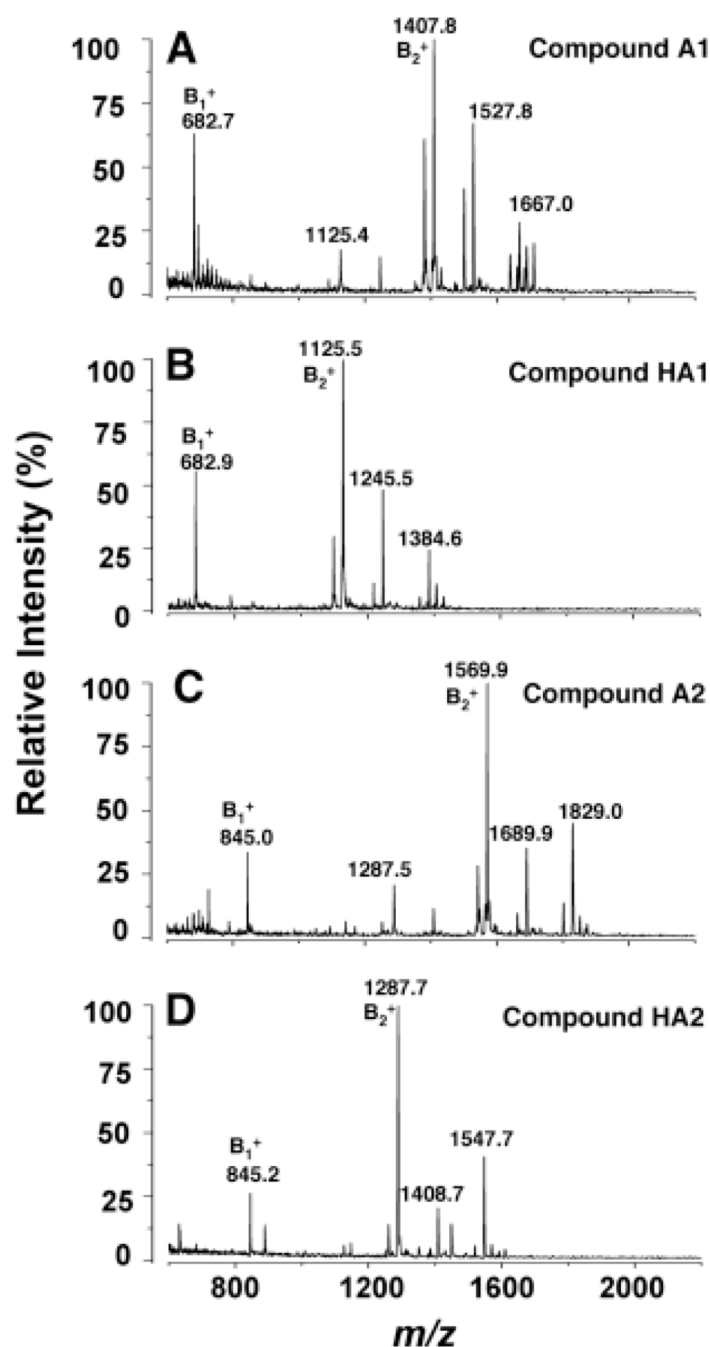


Figure 6. Positive ion MALDI/TOF MS of A1, HA1, A2 and HA2

Mild alkaline hydrolysis with triethylamine was used to generate HA1 and HA2 from A1 and A2, respectively (44). This procedure removes only the unsubstituted ester-linked 3-hydroacyl chains. The proposed structures of these compounds and the characteristic cleavages that give rise to the B_1^+ and B_2^+ ions are summarized in Fig. 7. Some fatty acid chain length heterogeneity was observed, as in the negative ion spectra (Fig. 3).

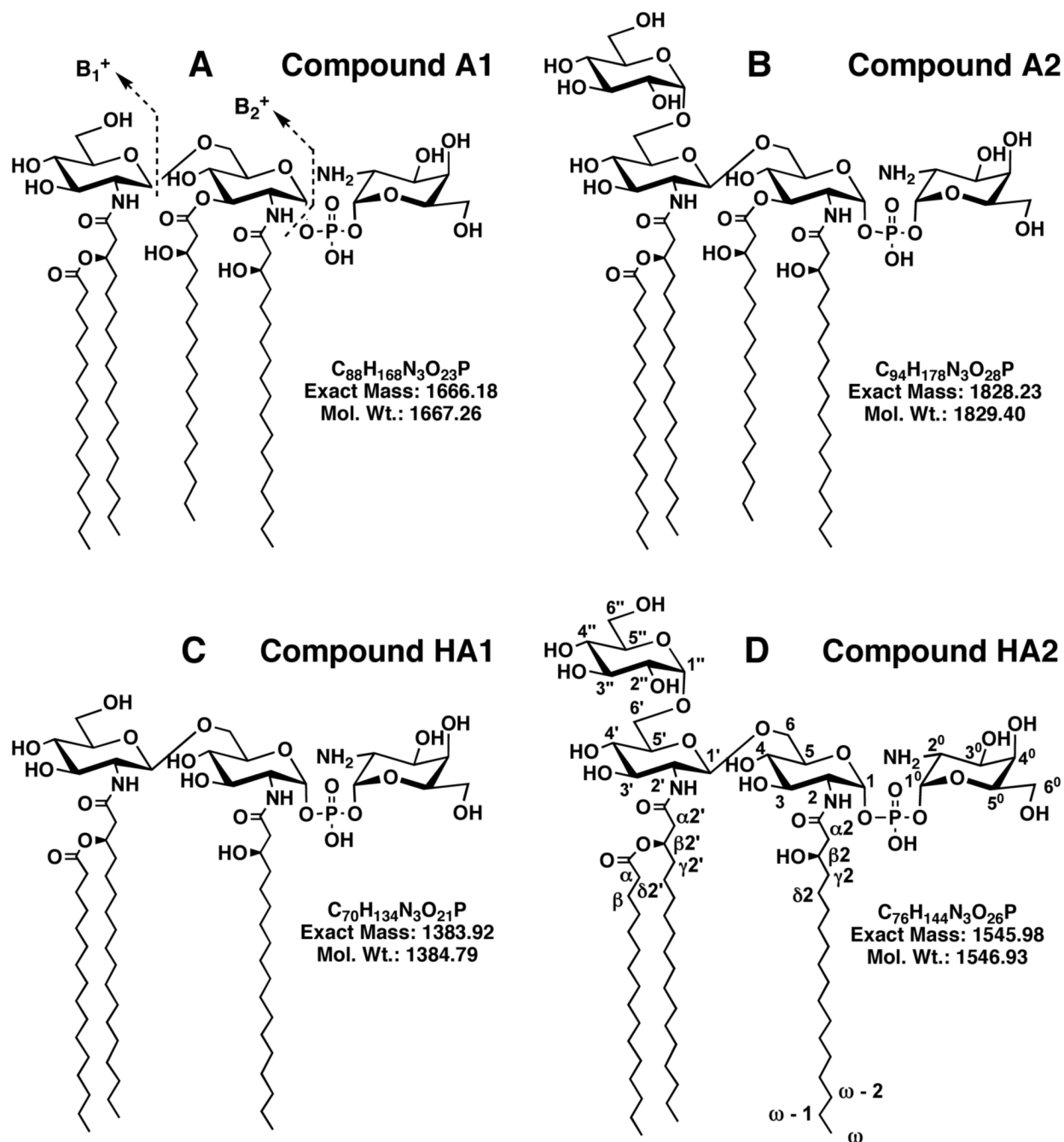


Figure 7. Proposed structures of free lipid A species purified from *F. novicida* U112
Compounds A1 (**Panel A**) and A2 (**Panel B**) are converted by mild alkaline hydrolysis with aqueous triethylamine to compounds HA1 (**Panel C**) and HA2 (**Panel D**) respectively. The anomeric configurations and the location of glucose residue at position 6' in A2 and HA2 were determined by NMR spectroscopy, as discussed in the text. Positions are labeled according to the scheme shown in **Panel D**. The characteristic cleavages that give rise to the B_1^+ and B_2^+ ions during MS (45) are shown in **Panel A**.

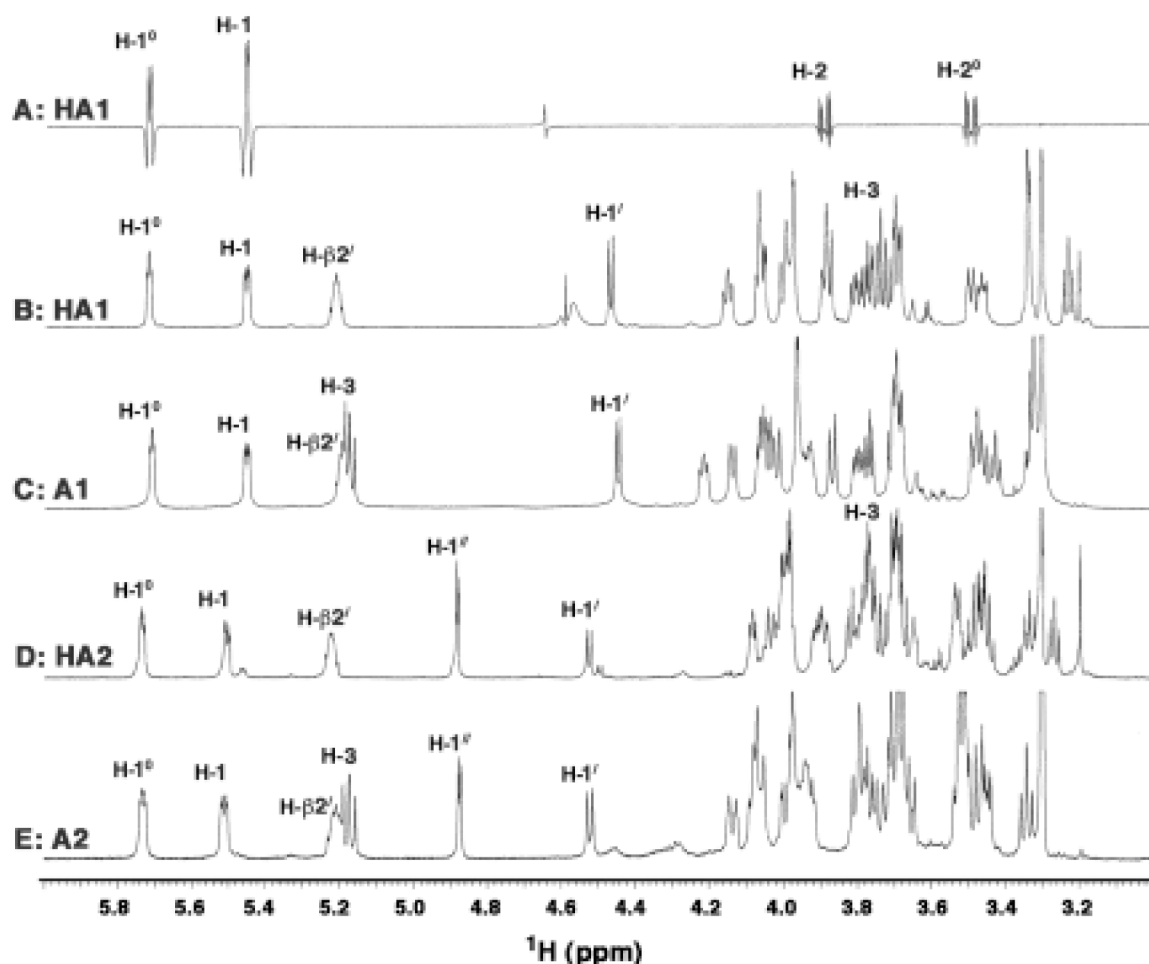


Fig. 8. ^1H NMR spectra of HA1, A1, HA2 and A2 and ^{31}P -decoupled ^1H NMR difference spectrum of HA1

Panel A. Difference 500 MHz ^1H NMR spectrum (38,40) of *F. novicida* HA1 obtained by subtraction of ^1H spectra obtained with selective on- and off-resonance decoupling of the ^{31}P NMR signal at -2.18 ppm. The difference spectrum reveals the simultaneous appearance of doublets at 5.45 ppm (H-1) and 5.72 ppm (H-1') and double-doublets at 3.90 ppm (H-2) and 3.49 ppm (H-2'). This pattern demonstrates that the single phosphorus group in compound HA1 is a monophosphodiester linkage that connects C-1 of the proximal glucosamine unit with C-1' of the capping galactosamine (Fig. 7). The observation of the H-2 and H-2' double-doublets in the difference spectrum allows estimates of the $J_{1,2}$ and $J_{2,3}$ coupling constants (Table 1). **Panels B to E.** Partial 1D 800 MHz ^1H NMR spectra of compounds HA1, A1, HA2 and A2 respectively in $\text{CDCl}_3:\text{CD}_3\text{OD}:\text{D}_2\text{O}$ (2:3:1, v/v/v), highlighting key sugar and β -hydroxymethine proton resonances. The numbering scheme for each position is defined in Fig. 7 and Table 1.

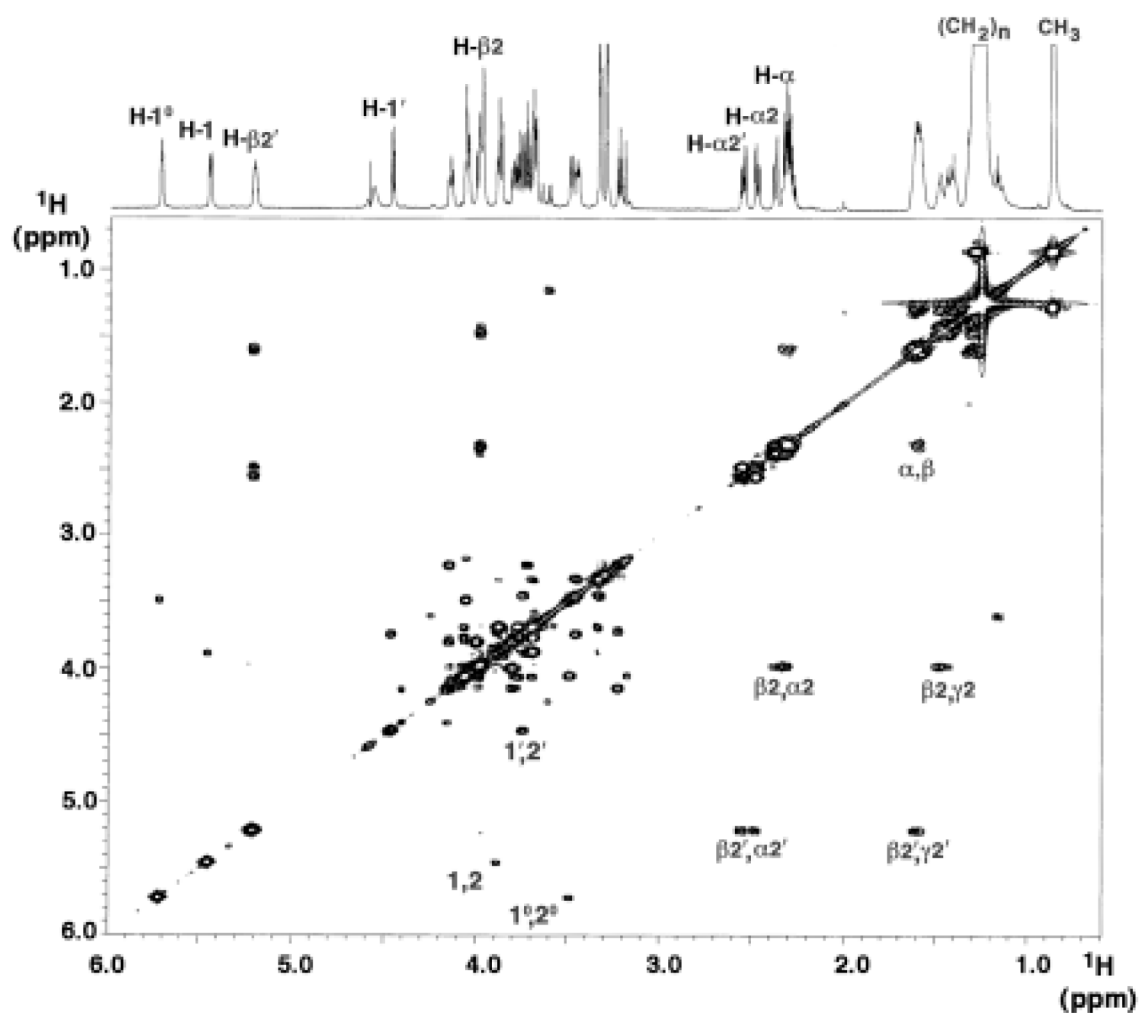


Figure 9. An 800 MHz ^1H - ^1H COSY analysis of compound HA1

Key correlations within the sugar rings and acyl chains are labeled. The acyl chain cross-peaks distinguish the unsubstituted β -hydroxyacyl chain from the esterified β -hydroxyacyl (acyloxyacyl) chain. The numbering scheme for each position is defined in Fig. 7 and Table 1.

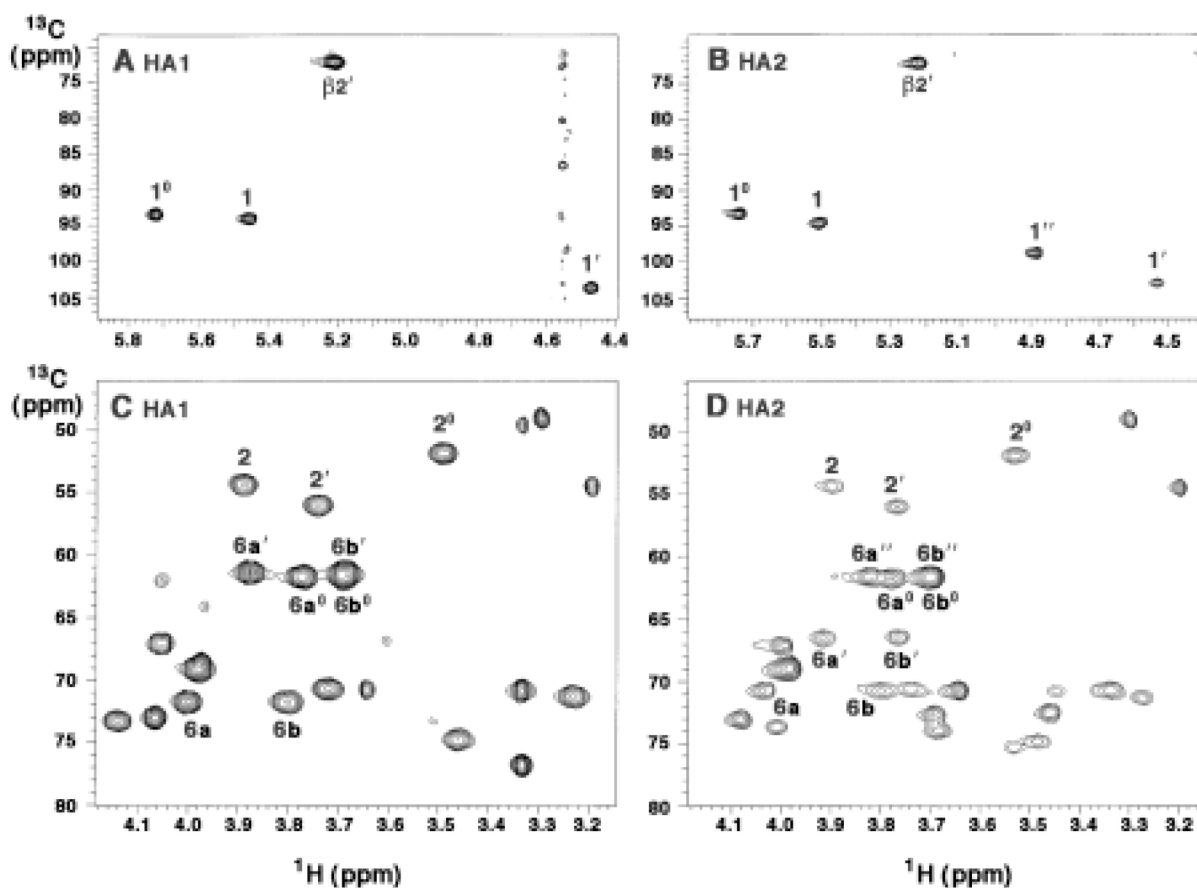


Figure 10. Partial 800 MHz HMQC spectra of HA1 and HA2

Panels A and B show the $^1\text{H}/^{13}\text{C}$ cross peaks in the anomeric region for HA1 and HA2 respectively. The numbering for each position is defined in Fig. 7 and Table 1. This analysis confirms the presence and anomeric linkage assignments of the three sugars in HA1 and four sugars in HA2. **Panels C and D** show the remaining sugar region of HA1 and HA2 respectively. The numbering scheme for the C-2⁰, C-2 and C-2' aminomethines (demonstrating the presence of three amino sugars in each compound) and for the C6 methylene cross peaks is shown in Fig. 7.

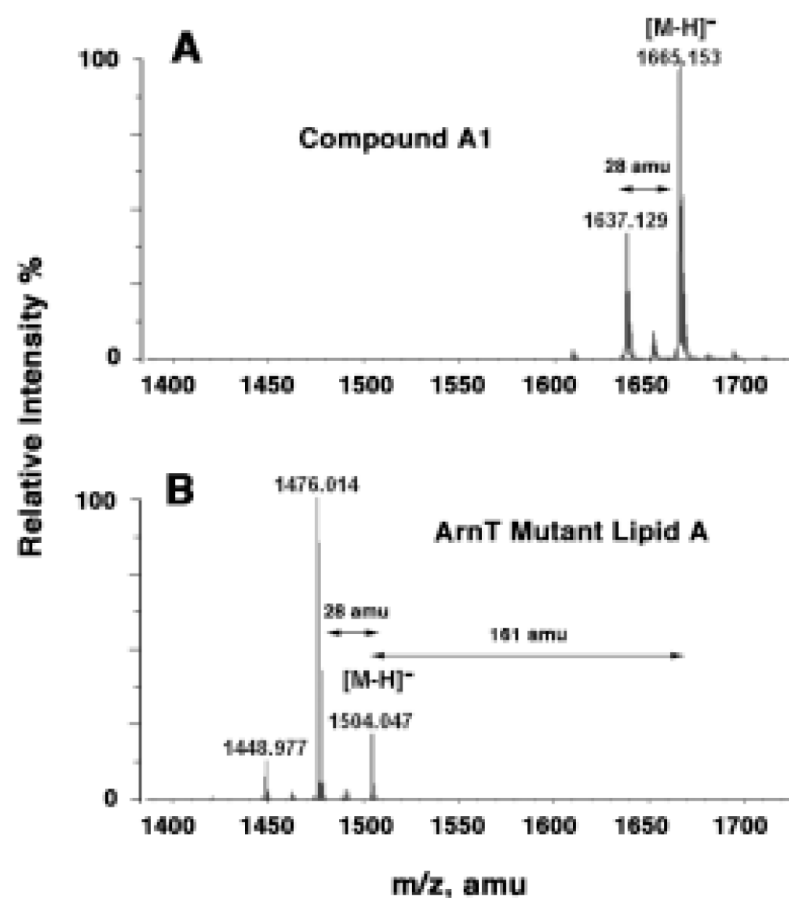
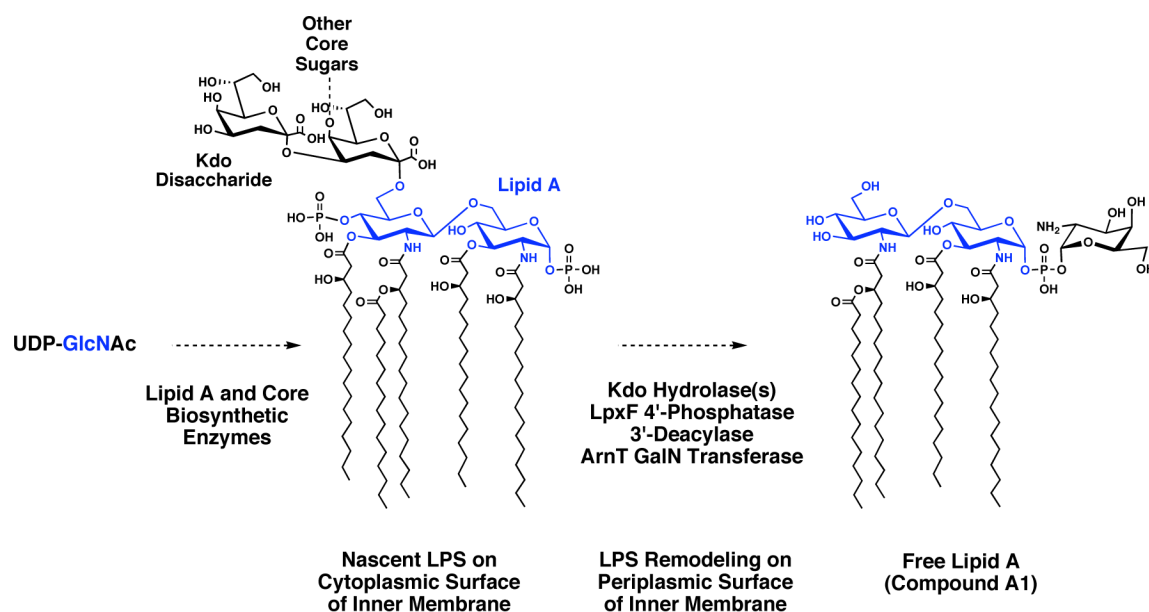


Figure 11. Absence of the galactosamine substituent on lipid A in the *arnT* mutant of *F. novicida*
Panel A. Negative ion ESI mass spectrum of compound A1 from wild-type *F. novicida* U112.
Panel B. Negative ion ESI mass spectrum of the major free lipid A species isolated from the *arnT* knockout of *F. novicida*. For ESI/MS, lipids were dissolved in chloroform and methanol (2:1, v/v) at about 25 $\mu\text{g/ml}$.



Scheme 1. A proposed enzymatic pathway for the processing of nascent LPS to generate free lipid A molecules in *Francisella novicida*

The glucosamine residues are shown in *blue*. The galactosamine unit incorporated by ArnT is shown in *black*. The genes encoding the Kdo hydrolase and 3'-deacylase, as well as their sub-cellular localization, are unknown.

Table 1¹H and ¹³C NMR Assignments of Compounds HA1 and HA2 from *F. novicida*¹H and ¹³C chemical shifts at 25 °C in CDCl₃, CD₃OD, D₂O (2:3:1, v/v/v) are relative to an internal TMS standard and are assigned based on 2D NMR experiments as discussed in the text. The numbering scheme is shown in Fig. 7

Position	HA1	HA2	HA1	HA2
	δ H [(mult), J(Hz)]	δ H [(mult), J(Hz)]	δ C [mult]	δ C [mult]
1 ⁰	5.72 [dd,3.5,5.4]	5.74 [dd,3.6,5.8]	93.5 [dd]	93.3 [dd]
2 ⁰	3.49 [dt,3.3,10.7]	3.53 [dt,3.3,10.7]	52.4 [dd]	52.3 [dd]
3 ⁰	4.06 [m]	4.00 [m]	67.0 [d]	67.5 [d]
4 ⁰	3.98 [m]	3.98 [m]	69.3 [d]	69.3 [d]
5 ⁰	4.07 [m]	4.01 [m]	73.0 [d]	74.0 [d]
6a ⁰	3.77 [m]	3.78 [m]	62.6 [t]	62.2 [t]
6b ⁰	3.69 [m]	3.70 [m]		
1	5.46 [dd,3.3,7.4]	5.51 [dd,3.1,7.9]	94.0 [dd]	94.8 [dd]
2	3.90 [m]	3.90 [m]	54.5 [dd]	54.7 [dd]
3	3.72 [m]	3.74 [m]	70.5 [d]	70.9 [d]
4	3.23 [dd]	3.2 [dd]	71.5 [d]	71.6 [d]
5	4.15 [m]	4.08 [m]	73.4 [d]	73.4 [d]
6a	4.00 [m]	4.04 [m]	72.0 [t]	71.1 [t]
6b	3.80 [dd,7.3,12.0]	3.80 [dd,7.3,12.0]		
1'	4.47 [d,8.6]	4.54 [d,8.6]	103.6 [d]	103.1 [d]
2'	3.75 [m]	3.77 [m]	56.0 [d]	56.1 [d]
3'	3.46 [m]	3.49 [m]	74.5 [d]	75.2 [d]
4'	3.33 [m]	3.45 [m]	70.7 [d]	71.0 [d]
5'	3.34 [m]	3.54 [m]	77.0 [d]	75.6 [d]
6'a	3.88 [m]	3.92 [m]	61.3 [t]	66.7 [t]
6'b	3.70 [m]	3.77 [m]		
1''		4.88 [d,3.7]		98.8 [d]
2''		3.47 [m]		72.9 [m]
3''		3.69 [m]		74.5 [m]
4''		3.34 [m]		71.4 [d]
5''		3.70 [m]		72.9 [d]
6''a		3.82 [m]		62.1 [t]
6''b		3.70 [m]		
α 2a	2.39 [dd,3.1,14.2]	2.39 [dd,3.1,14.2]	44.2 [t]	44.2 [t]
α 2b	2.35 [m]	2.35 [m]		
β 2	3.98 [m]	3.98 [m]	69.0 [d]	69.3 [d]
γ 2	1.45 [m]	1.45 [m]	37.4	
δ 2	1.3 [m]	1.3 [m]		
CO2			175.1	174.8
α 2'a	2.5 [dd,7.6,14.5]	2.56 [dd,7.6,14.5]	41.8 [t]	41.6 [t]
α 2'b	2.49 [dd,5.6,14.5]	2.49 [dd,5.6,14.5]		
β 2'	5.22 [m]	5.23 [m]	72.0 [d]	71.8 [d]
γ 2'	1.61 [m]	1.61 [m]	34.5	
δ 2'	1.3	1.3		
CO2'			173.4	173.2
α CH ₂	2.31 [m]	2.31 [m]	35.2 [t]	35.3 [t]
β CH ₂	1.61 [m]	1.61 [m]	25.5	
γ CH ₂	1.3	1.3		
CO			175.7	175.6
(CH ₂) _n	1.24 [m]	1.24 [m]	30.4 [m]	30.4 [m]
ω -2 CH ₂	1.26 [m]	1.26 [m]	32.7 [t]	32.7 [t]
ω -1 CH ₂	1.26 [m]	1.26 [m]	23.4 [t]	23.4 [t]
ω CH ₃	0.88 [t, ~7]	0.88 [t, ~7]	14.0 [q]	14.5 [q]