
Identification of \textit{lptA}, \textit{lpxE}, and \textit{lpxO}, Three Genes Involved in the Remodeling of \textit{Brucella} Cell Envelope

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The brucellae are facultative intracellular bacteria that cause a worldwide extended zoonosis. One of the pathogenicity mechanisms of these bacteria is their ability to avoid rapid recognition by innate immunity because of a reduction of the pathogen-associated molecular pattern (PAMP) of the lipopolysaccharide (LPS), free-lipids, and other envelope molecules. We investigated the \textit{Brucella} homologs of \textit{lptA}, \textit{lpxE}, and \textit{lpxO}, three genes that in some pathogens encode enzymes that mask the LPS PAMP by upsetting the core-lipid A charge/hydrophobic balance. \textit{Brucella} \textit{lptA}, which encodes a putative ethanolamine transferase, carries a frame-shift in \textit{B. abortus} but not in other \textit{Brucella} spp. and phylogenetic neighbors like the opportunistic pathogen \textit{Ochrobactrum anthropi}. Consistent with the genomic evidence, a \textit{B. melitensis} \textit{lptA} mutant lacked lipid A-linked ethanolamine and displayed increased sensitivity to polymyxin B (a surrogate of innate immunity bactericidal peptides), while \textit{B. abortus} carrying \textit{B. melitensis} \textit{lptA} displayed increased resistance. \textit{Brucella} \textit{lpxE} encodes a putative phosphatase acting on lipid A or on a free-lipid that is highly conserved in all brucellae and \textit{O. anthropi}. Although we found no evidence of lipid A dephosphorylation, a \textit{B. abortus} \textit{lpxE} mutant showed increased polymyxin B sensitivity, suggesting the existence of a hitherto unidentified free-lipid involved in bactericidal peptide resistance. Gene \textit{lpxO} putatively encoding an acyl hydroxylase carries a frame-shift in all brucellae except \textit{B. microti} and is intact in \textit{O. anthropi}. Free-lipid analysis revealed that \textit{lpxO} corresponded to \textit{olsC}, the gene coding for the ornithine lipid (OL) acyl hydroxylase active in \textit{O. anthropi} and \textit{B. microti}, while \textit{B. abortus} carrying the \textit{olsC} of \textit{O. anthropi} and \textit{B. microti} synthesized hydroxylated OLs. Interestingly, mutants in \textit{lptA}, \textit{lpxE}, or \textit{olsC}...
INTRODUCTION

Brucellosis is the collective name of a group of zoonotic diseases afflicting a wide range of domestic and wild mammals (Whatmore, 2009; Zheludkov and Tsirelson, 2010). In domestic livestock brucellosis is manifested mostly as abortions and infertility, and contact with infected animals and consumption of unpasteurized dairy products are the sources of human brucellosis, an incapacitating condition that requires prolonged antibiotic treatment (Zinsstag et al., 2011). Eradicated in a handful of countries, brucellosis is endemic or even increasing in many areas of the world (Jones et al., 2013; Ducrot et al., 2017; Lai et al., 2017).

This disease is caused by facultative intracellular parasites of the genus Brucella. Taxonomically placed in the α-2 Proteobacteria (Moreno et al., 1990), the brucellae are close to plant pathogens and endosymbionts such as Agrobacterium, Sinorhizobium, and Rhizobium and to soil bacteria such as Oscibacterium, the latter including some opportunistic pathogens, and comparative analyses suggest that soil bacteria of this group are endowed with properties that represent a first scaffold on which an intracellular life style develops (Velasco et al., 2000; Moreno and Moriyón, 2007; Barquero-Calvo et al., 2009). The brucellae owe their pathogenicity mainly to their ability to multiply within dendritic cells, macrophages, and a variety of other cells. Due to their ability to control intracellular trafficking and be barely detected by innate immunity, these bacteria are able to reach a safe intracellular niche before an effective immune response is mounted, and to multiply extensively (Gorvel and Moreno, 2002; Barquero-Calvo et al., 2007). A mechanism used by Brucella to escape from the host immune response is the interference with the toll-like receptor (TLR) signaling pathway by the injection of active effectors such as BtpA and BtpB through the Type IV secretion system T4SS. Both effector proteins contain a TIR domain that interferes with TLR signaling by directly interacting with MyD88 (Giré et al., 2008; Salcedo et al., 2008, 2013; Chaudhry et al., 2012) and contribute to the control of dendritic cell (DC) activation during infection. Moreover, Brucella has modified outer membrane (OM) components in order to reduce the pathogen-associated molecular patterns (PAMP) of the cell envelope. In Gram-negative bacteria, these PAMP are created by the conserved composition of the OM lipopolysaccharide (LPS) and the free lipids on which the topology of the OM also depends. However, in addition to free-lipid species present in most Gram-negative bacteria (i.e., cardiolipin, phosphatidyglycerol, and phosphatidylethanolamine), Brucella also possesses phosphatidylcholine and amino lipids. Phosphatidylcholine is a eukaryotic-type phospholipid required for Brucella full virulence (Comerci et al., 2006; Conde-Álvarez et al., 2006). Among the amino lipids, only the ornithine lipids (OL) have been investigated which unlike their counterparts in Bordetella, do not trigger the release of IL-6 or TNF-α by macrophages, possibly on account of their longer acyl chains that reduce the OL PAMP (Palacios-Chaves et al., 2011). Concerning the LPS, most bacteria carry C1 and C4' glucosamine disaccharides with C12 and C14 acyl and acyl-oxyacyl chains. This highly amphipathic structure, named lipid A, is adjacent to additional negatively charged groups of the core oligosaccharide, namely the heptose phosphates and 2-keto-3-deoxyoctulosonate carboxyl groups (Kastowsky et al., 1992; Moriyón, 2003). This lipid A-core PAMP is so efficiently detected by the innate immunity system that some pathogens partially conceal it by removing phosphate groups or substituting them with arabinosamine and/or ethanolamine, or by hydroxylating the acyl chains (Takahashi et al., 2008; Lewis et al., 2013; Moreira et al., 2013; Needham and Trent, 2013; Llobet et al., 2015; Trombley et al., 2015). In contrast, Brucella lipid A is a diaminoglucone disaccharide amide-linked to long (C16, C18) and very long (C28–C30) acyl chains (Velasco et al., 2000; Iriarte et al., 2004; Fontana et al., 2016). Furthermore, negative charges in lipid A phosphates and 2-keto-3-deoxyoctulosonate are counterbalanced by four glucosamine units present in the core (Kubler-Kielb and Vinogradov, 2013; Fontana et al., 2016). As illustrated by the unusually reduced endotoxicity of the Brucella LPS this structure is defectively detected by the innate immune response (Lapaque et al., 2005; Martirosyan et al., 2011; Conde-Álvarez et al., 2012). It remains unknown, however, whether Brucella LPS undergoes post-synthetic modifications that have been described for other bacteria that could alter its PAMP potential and contribution to virulence. In this work, we investigated in Brucella the role of gene homologs to phosphatases, phospho-ethanolamine (pEtN) transferases, and acyl hydroxylases (Figure 1) that have been shown in other Gram-negative pathogens to act on LPS and to contribute to overcoming innate immunity defenses.

MATERIALS AND METHODS

Bacterial Strains and Growth Conditions

The bacterial strains and plasmids used in this study are listed in Supplementary Table S1. Bacteria were routinely grown in standard tryptic soy broth or agar either plain or supplemented with kanamycin at 50 μg/ml, or/and nalidixic acid at 5 or 25 μg/ml or/and 5% sucrose. All strains were stored in skim milk at −80°C.
DNA Manipulations

Genomic sequences were obtained from the Kyoto Encyclopedia of Genes and Genomes (KEGG) database\(^1\). Searches for DNA and protein homologies were carried out using the National Center for Biotechnology Information (NCBI\(^2\)) and the European Molecular Biology Laboratory (EMBL) - European Bioinformatics Institute server\(^3\). Primers were synthesized by Sigma-Genosys (Haverhill, United Kingdom). DNA sequencing was performed by the “Servicio de Secuenciación del Centro de Investigación Médica Aplicada” (Pamplona, Spain). Restriction–modification enzymes were used under the conditions recommended by the manufacturer. Plasmid and chromosomal DNA were extracted with Qiaprep Spin Miniprep (Qiagen) and Ultraclean Microbial DNA Isolation Kits (Mo Bio Laboratories), respectively. When needed, DNA was purified from agarose gels using the Qiack Gel Extraction Kit (Qiagen).

Mutagenesis

To obtain \(\text{Bme}\Delta\text{lptA}\), \(\text{Ba}\Delta\text{lpxE}\), and \(\text{Bmi}\Delta\text{olsC}\) in-frame deletion mutants, directed mutagenesis by overlapping PCR were performed using genomic DNA as template and pJQK (Scupham and Triplett, 1997) as the suicide vector. The corresponding gene was deleted using allelic exchange by double recombination as previously described (Conde-Alvarez et al., 2006).

For the construction of the \(\text{Bme}\Delta\text{lptA}\) mutant, we first generated two PCR fragments: oligonucleotides \(\text{lptA-F1 (5'}'-\text{GAA CGCGAGACTATGGAAAC-3}'}\) and \(\text{lptA-R2 (5'}'-\text{TGGT GAACGCCAGAAGATAGA-3}'}\) were used to amplify a 400-bp fragment including codons 1–26 of \(\text{BmelptA}\) ORF, as well as 324 bp upstream of the \(\text{BmelptA}\) start codon, and oligonucleotides \(\text{lptA-F3 (5'}'-\text{TCTATCTTTGGCGGTTCCACC GCACGCAATCTCTTGTC-3}'}\) and \(\text{lptA-R4 (5'}'-\text{AATATTTCCAT GGGCATTTC-3}'}\) were used to amplify a 472-bp fragment including codons 506–544 of the \(\text{lptA}\) ORF and 353-bp downstream of the \(\text{lptA}\) stop codon. Both fragments were ligated by overlapping PCR using oligonucleotides \(\text{lptA-F1 and lptA-R4}\) for amplification, and the complementary regions between

\(^{1}\)http://www.genome.jp/kegg/
\(^{3}\)http://www.ebi.ac.uk/
For pBME<sub>lpxE</sub> and pBME<sub>lptA</sub> construction we took advantage of the replicative plasmid pBBR1 MCS (Kovach et al., 1994). The resulting mutator plasmid (pRCI-32) was introduced in B. abortus S.17 strain (Simon et al., 1983).

For the construction of the <i>Bmi</i>Δ<em>olsC</em> mutant, we first generated two PCR fragments: oligonucleotides <em>lpxE</em>-<em>F1</em> (5′-CCGCTGGATCCTAGGTATATT-3′) and <em>lpxE</em>-<em>R2</em> (5′-TATAGG CAGGGCCGAGA-3′) were used to amplify a 334-bp fragment including codons 1–15 of <em>lpxE</em> ORF, as well as 289 bp upstream of the <em>lpxE</em>-1 start codon, and oligonucleotides <em>lpxE</em>-<em>F3</em> (5′-TTCTG CGCCCTGCGCTATAGTTGTTICCGCATGTT-3′) and <em>lpxE</em>-<em>R4</em> (5′-CCAAATACCCGTCATGAGA-3′) were used to amplify a 577-bp fragment including codons 226–255 of the <em>lpxE</em> ORF and 488-bp downstream of the <em>lpxE</em> stop codon. Both fragments were ligated by overlapping PCR using oligonucleotides <em>lpxE</em>-<em>F1</em> and <em>lpxE</em>-<em>R4</em> for amplification, and the complementary regions between <em>lpxE</em>-<em>R2</em> and <em>lpxE</em>-<em>F3</em> (5′-TTCTG CGCCCTGCGCTATAGTTGTTICCGCATGTT-3′) and <em>lpxE</em>-<em>R4</em> (5′-CCAAATACCCGTCATGAGA-3′) were used to amplify a 320-bp fragment including codons 286–313 of the <em>lpxE</em> ORF and 289 bp downstream of the <em>lpxE</em> stop codon. Both fragments were ligated by overlapping PCR using oligonucleotides <em>olsC</em>-<em>F1</em> and <em>olsC</em>-<em>R4</em> for amplification, and the complementary regions between <em>olsC</em>-<em>R2</em> and <em>olsC</em>-<em>F3</em> (5′-TGCTGGATCCTAGGTATATT-3′) and <em>olsC</em>-<em>F3</em> (5′-CCGCTGGATCCTAGGTATATT-3′) were used to amplify a 334-bp fragment including codons 1–15 of <em>olsC</em> ORF, as well as 289 bp upstream of the <em>olsC</em> start codon, and oligonucleotides <em>olsC</em>-<em>F3</em> (5′-TA GTTCCCATCGCTATAGTTGTTICCGCATGTT-3′) and <em>olsC</em>-<em>R4</em> (5′-AACCGCCGAGGCTACCCACAC-3′) were used to amplify a 320-bp fragment including codons 286–313 of the <em>olsC</em> ORF and 289 bp downstream of the <em>olsC</em> stop codon. Both fragments were ligated by overlapping PCR using oligonucleotides <em>olsC</em>-<em>F1</em> and <em>olsC</em>-<em>R4</em> for amplification, and the complementary regions between <em>olsC</em>-<em>R2</em> and <em>olsC</em>-<em>F3</em> for overlapping. The resulting fragment, containing the <em>lptA</em> deleted allele, was cloned into pCR2.1 (Invitrogen, Barcelona, Spain), sequenced to ensure maintenance of the reading frame, and subcloned into the <em>BamHI</em> and the <em>XbaI</em> sites of the suicide plasmid pJQK (Scupham and Tripplett, 1997). The resulting mutator plasmid (pRCI-65) was introduced in <i>B. microti</i> CM445 by conjugation using the <i>E. coli</i> S.17 strain (Simon et al., 1983).

Deletion of each gene was checked with oligonucleotides gene-F1 and gene-R4 and internal primers hybridizing in the non-deleted regions.

**Complementation of Deleted Genes**

For pBME<em>lpxE</em> and pBME<em>lptA</em> construction we took advantage of the <i>Brucella</i> ORFeome constructed with the Gateway cloning Technology (Invitrogen) (Dricot et al., 2004). The clones carrying Bm<em>lpxE</em> or Bm<em>lptA</em> were extracted and the DNA containing the corresponding ORF was subcloned in plasmid pRH001 (Hallez et al., 2007) to produce pBME<em>lpxE</em> and pBME<em>lptA</em>. For pBMlolsC, <em>olsC</em> was amplified using genomic DNA of Bmi-parental as DNA template. The primers used were <em>olsC</em>-<em>F6</em> (5′-GCTTTCCGAACAAGCAGACTGA-3′) and <em>olsC</em>-<em>R7</em> (5′-GCCCTCCCTTCACCGGTATT-3′). The resulting PCR product, containing the ORF from 342 bp upstream to 84 bp downstream, was then cloned into pCR2.1 TOPO (Invitrogen) plasmid by “TA cloning” (Life Technologies). The resulting plasmid was sequenced to ensure that the gene was correctly cloned. Then, the gene was subcloned into the <em>BamHI</em> and the <em>XbaI</em> sites of the replicative plasmid pBBR1 MCS (Kovach et al., 1994) pBME<em>lpxE</em>, pBME<em>lptA</em>, and pBMlolsC were introduced into <i>Brucella</i> by conjugation using <i>E. coli</i> S.17-1 strain and the conjugants harboring corresponding plasmid were selected by plating onto TSA-Nal-Cm plates.

**Sensitivity to Cationic Peptides**

Exponentially growing bacteria were adjusted to an optical density equivalent to one of the McFarland scale and the minimal inhibitory concentrations (MICs) of polymyxin B were determined by the <i>e</i>-test method on Müller–Hinton agar (Izasa) or by the serial dilution method in a similar broth.

**LPS Preparation**

Lipopolysaccharide was obtained by methanol precipitation of the phenol phase of a phenol–water extract (Leong et al., 1970). This fraction [10 mg/ml in 175 mM NaCl, 0.05% NaN<sub>3</sub>, 0.1 M Tris–HCl (pH 7.0)] was then purified by digestion with nucleases [50 µg/ml each of DNase-II type V and RNase-A (Sigma, St. Louis, MO, United States), 30 min at 37°C and three times with proteinase K (50 µg/ml, 3 h at 55°C), and ultracentrifuged (6 h, 100,000 × g) (Aragón et al., 1996). Free lipids (OLs and phospholipids) were then removed by a fourfold extraction with chloroform–methanol [2:1 (vol/vol)] (Velasco et al., 2000).

**Infections in Mice**

Seven-week-old female BALB/c mice (Charles River, Elbeuf, France) were kept in cages with water and food ad libitum and accommodated under biosafety containment conditions 2 weeks before the start of the experiments. To prepare inocula, tryptic soy agar (TSA) grown bacteria were harvested and suspended in 10 mM phosphate buffered saline (pH 6.85), and 0.1 ml/mouse containing approximately 5 × 10<sup>8</sup> colony forming units (CFU) for <i>B. melitensis</i> or <i>B. abortus</i> and 1 × 10<sup>4</sup> CFU for <i>B. microti</i> was administered intraperitoneally. The exact doses assessed retrospectively by plating dilutions of the inocula. Number of CFU in spleens was determined at different time after inoculation. For this, the spleens were aseptically removed and individually weighed and homogenized in 9 volumes of PBS. Serial 10-fold dilutions of each homogenate were performed and each dilution was plated in triplicate. Plates were incubated at 37°C for 5 days. At several points during the infection process, the identity of the spleen isolates was confirmed by PCR. The individual data were normalized by logarithmic transformation, and the mean log CFU/spleen values and the standard deviations (n = 5) were calculated.
Intracellular Multiplication Assays
Bone marrow cells were isolated from femurs of 7–8-week-old C57Bl/6 female and differentiated into dendritic cells [bone-marrow derived dendritic cells (BMDCs)] as described by Inaba et al. (1992). Infections were performed by centrifuging the bacteria onto the differentiated cells (400 x g for 10 min at 4°C; bacteria:cells ratio of 30:1 followed by incubation at 37°C for 30 min under a 5% CO₂ atmosphere). BMDCs were gently washed with medium to remove extracellular bacteria before incubating in medium supplemented with 50 µg/ml gentamicin for 1 h to kill extracellular bacteria. Thereafter, the antibiotic concentration was decreased to 10 µg/ml. To monitor Brucella intracellular survival at different time-points post-infection, BMDC were lysed with 0.1% (vol/vol) Triton X-100 in H₂O and serial dilutions of lysates were plated onto TSA plates to enumerate the CFU.

Flow Cytometry
To assess activation and maturation, BMDC were analyzed for surface expression of classical maturation markers at 24 h post-treatment with the different Brucella strains and derived mutants. Cells were labeled with fluorochrome-conjugated antibodies specific for mouse CD11c:APC-Cy7 (clone N418), IA-IE:PE (MHC class II clone M5/114.15.2) (PE), CD86:FITC (Clone GL-1), CD40:APC (clone 3/23), and CD80:PE-Cy5 (clone 16-10A1), all from BioLegend. Labeled cells were then subjected to multi-color cytometry using a LSR II UV (Becton Dickinson) and the data analyzed using FlowJo Software by first gating on the CD11c+ population (100,000 events) prior to quantifying expression of receptors. Cells were stimulated with E. coli LPS (055:B5) as a positive control.

Lipid A Extraction
Five milligrams of LPS was hydrolyzed in 5 ml 1% acetic acid by sonication, heating to 100°C for 30 min, and cooling to room temperature. Concentrated HCl was added to the mixture until the pH was 1–2. The solution was converted to a two-phase acidic Bligh–Dyer mixture by adding 5.6 ml of chloroform and 5.6 ml of methanol. Phases were mixed by inverting the tubes and separated by centrifugation at 4000 g for 20 min. The lower phases containing lipid A were collected, washed two times with water, and dried under a stream of nitrogen. Extraction was repeated, and the lower phases (11.2 ml) were combined and neutralized with a drop of pyridine. Samples were evaporated to dryness under a stream of nitrogen.

Mass Spectrometry
Mass spectrometa were acquired on a Bruker Autoflex®Speed TOF/TOF Mass Spectrometer (Bruker Daltonics Inc.) in negative reflective mode with delayed extraction. The ion-accelerating voltage was set at 20 kV. Each spectrum was an average of 300 shots. A peptide calibration standard (Bruker Daltonics Inc.) was used to calibrate the Matrix Assisted Laser Desorption/Ionization Time-of-Flight (MALDI-TOF), and lipid A extracted from E. coli strain MG1655 grown in LB medium at 37°C.

Extraction and Analysis of Envelope Lipids
The free-lipid fraction was extracted as described by Bligh and Dyer (1959), and analyzed on a silica gel 60 high-performance thin layer chromatography (HPLC) plates (Merck, Darmstadt, Germany). Chromatography was performed either monodimensionally with chloroform–methanol–water [14:6:1 (volume)] or bidimensionally with chloroform–methanol–water [14:6:1 (volume)] first and chloroform–methanol–acetic acid [13:5:2 (volume)] in the second dimension (Weissenmayer et al., 2002). Plates were developed with 0.2% ninhydrin in acetone at 180°C or 15% sulfuric acid in ethanol at 180°C.

RESULTS

The Brucella lptA Orthologs Encode a Lipid A Phosphate-Ethanolamine Transferase
A genomic search in the KEGG database revealed that all Brucella spp. carry an ORF (BMEI0118 in B. melitensis) homologous to Neisseria meningitidis lptA, a pEtN transferase that modifies lipid A (Cox et al., 2003). Strikingly, in B. abortus but not in other Brucella spp., all genomic sequences available at KEGG show a deletion of a thymine in position 774 that should result in a truncated protein lacking the amino acids related to the enzymatic activity (Naessan et al., 2008; Figure 1 and Supplementary Figure S1 and Supplementary Table S2). In addition to LptA, two other pEtN transferases have been identified in N. meningitidis: Lpt-3 and Lpt-6, which, respectively, modify the LPS core at the third and sixth position of heptose II (Mackinnon et al., 2002; Wright et al., 2004). By multiple sequence alignment, the B. melitensis putative pEtN transferase showed highest homology with Neisseria LptA and also displayed the LptA membrane-associated domains not present in Lpt-3 and Lpt-6 (ORFs NMB1638, NMB2010, and NMA0408, respectively). Accordingly, it can be predicted that ORF BMEI0118 (henceforth BMe/lptA) encodes a pEtN transferase that acts on lipid A, a hypothesis fully consistent with the absence of heptose in the Brucella LPS core (Iriarte et al., 2004; Fontana et al., 2016).

To test this hypothesis, we constructed a B. melitensis non-polar mutant (BmeΔlptA) lacking the LptA enzymatic domain (amino acids 26–506), which as expected maintained a smooth (S) phenotype (negative crystal violet test and positive coagglutination with anti-S-LPS antibodies). As a consequence of the increased positive charge of the amino group, pEtN has been shown to decrease binding of the polycationic lipopeptide polymyxin B to LPS, and to increase resistance to this antibiotic in a variety of bacteria (Needham and Trent, 2013; Trombley et al., 2015; Herrera et al., 2017). In keeping with this possibility, the BmeΔlptA mutant was more sensitive to polymyxin B than the parental strain B. melitensis 16M (Bme-parental) (Figure 2A). In contrast, and consistent with the frame-shift in its lptA homolog, B. abortus 2308 (Ba-parental) displayed polymyxin B sensitivity similar to that of BmeΔlptA. Moreover, complementation of BmeΔlptA with the multi-copy plasmid
The Brucella lptA orthologs are involved in polymyxin B resistance and code for a phosphate-ethanolamine transferase acting on lipid A. (A) Polymyxin B sensitivity of B. melitensis wild-type (Bme-parental), B. melitensis non-polar lptA mutant (BmeΔlptA), the cognate complemented mutant (BmeΔlptA pBMElptA), B. abortus wild-type (Ba-parental), and B. abortus wild-type carrying a plasmid with the B. melitensis lptA gene (Ba-parental pBMElptA) (the results are representative of three independent experiments). (B) MALDI-TOF analysis of the lipid A of Bme-parental, BmeΔlptA, and Ba-parental.

The Brucella lpxE Orthologs Encode a Phosphatase Involved in the Remodeling of the OM

As described above, MALDI-TOF analyses showed the presence of molecular species with a mass compatible with monophosphorylated lipid A. Since lipid A synthesis produces C1 and C4′ bisphosphorylated disaccharide backbones (Qureshi et al., 1994), a possible explanation could be its dephosphorylation by a phosphatase such as LpxE, an inner membrane enzyme that in the phylogenetic neighbor Rhizobium leguminosarum removes the lipid A phosphate at C1 (Raetz et al., 2009). A search in KEGG showed that all Brucella spp. carry an ORF homologous to R. leguminosarum lpxE (Supplementary Table S2). However, the start codon in the B. melitensis 16M homolog (BMEI1212) is annotated to a position different from that determined for other brucellae (Supplementary Table S2), including other B. melitensis strains. Thus, whereas the B. abortus homolog (BAB1_0671) is predicted to encode a protein of 255 amino acids, the B. melitensis
one could encode a protein of either 235 or 255 amino acids (Figure 1). Both proteins conserve the consensus sequence of the lipid phosphate superfamily [KX6RP-(X12–54)-PSGH-(X31–54)-SRX5HX3D] (Stukey and Carman, 2008) which is also present in LpxE from R. leguminosarum, Sinorhizobium melloti, and Agrobacterium tumefaciens (Karbarz et al., 2003). Although BAB1_0671 and BMEI1212 code for proteins that contain the three motifs conserved in the LpxF phosphatase from Francisella, they lack two amino acids of the central motif, NCSFX2G, which seems LpxF specific (Wang et al., 2006, 2007). Thus, the Brucella proteins were named BaLpxE and BMElpxE.

To study whether BALpxE actually acts as a lipid A phosphatase, we constructed a non-polar mutant (BaΔlpxE) and tested it against polymyxin B, since the permanence of a phosphate group in an OM group should increase sensitivity to this antibiotic. Mutant BaΔlpxE was eight times more sensitive than the parental strain (MIC 0.2 and 1.6 µg/mL, respectively). Moreover, when we introduced a plasmid containing the BMElpxE ortholog into BaΔlpxE, the resistance to polymyxin B was restored (MIC 1.6 µg/mL). Although final confirmation of this interpretation would require to assay the enzymatic activity of the protein, these results are consistent with the predicted role of lpxE as a phosphatase and its functionality in both B. abortus and B. melitensis 16M, a strain where the annotation of the start codon was a source of ambiguity.

By MALDI-TOF analysis, the Ba-parental lipid A spectrum showed three of the four predominant clusters of ions (A, B, and C) found in B. melitensis (Figure 2B and Supplementary Table S3). Cluster A (m/z 2173) was consistent with an hexaacylated bisphosphorylated dianaminogluucose disaccharide (C120H332N4O32P2) and the signal at 2093 m/z, which differed in the mass of one phosphate group (i.e., 80), was consistent with the cognate monophosphorylated lipid A (C120H332N4O35P) (A-Pi, Figure 2). Other signals differing in a mass of 14 or 28 units should result from the heterogeneity in acyl chain length that is typical of lipid A. The B and C clusters also contained signals differing in 80 mass units that could correspond to bis- and mono-phosphorylated species. The mass spectrum of BaΔlpxE lipid A (not shown) did not differ significantly from that of Ba-parental, and again showed acyl chain heterogeneity in the A, B, C clusters, as well as the ~80 m/z signals indicative of mono- and bisphosphorylated lipid A species. As mutation of LpxE is concomitant with an increase in polymyxin B sensitivity, it is tempting to speculate that LpxE directly or indirectly modulates Brucella cell envelope by removing an accessible phosphate group from a substrate different from lipid A. Further studies need to be performed to clarify the role of LpxE.

The Brucella lpxO Orthologs Encode an Acyl Hydroxylase Acting on Ornithine Lipids

The genomes of all Brucella species available at KEGG contain an ORF homologous to Salmonella lpxO (Gibbons et al., 2000), which encodes an enzyme hydroxylating the 3’-secondary acyl chain of lipid A. In all Brucella spp. except B. microti and B. vulpis this ORF presents a frame-shift leading to a truncated protein that lacks the consensus of the aspartyl/asparaginyl β-hydroxylases family to which LpxO belongs (Figure 1 and Supplementary Table S2). These characteristics are consistent with chemical studies that previously failed to observe S2 hydroxylated fatty acids in B. abortus lipid A (Velasco et al., 2000). Moreover, a lpxO homolog is present in Ochrobactrum anthropi where S2 hydroxylated fatty acids were also not observed in the lipid A (Velasco et al., 2000), indicating that a role similar to that of Salmonella LpxO is unlikely. Thus, the lpxO homologs present in these B. microti and O. anthropi could be acting on a free lipid and, in fact, it has been reported that the corresponding R. tropici homolog is a β-hydroxylase acting on OLs (Vences-Guzmán et al., 2011). If this were the case in O. anthropi and the brucellae, the end product [a hydroxylated OL (OH-OL) of the pathway described previously in members of the Rhizobiaceae (Figure 3A) should be observed in O. anthropi and B. microti (and B. vulpis) but not in other Brucella spp.

To investigate these hypotheses, we compared the free lipids of B. abortus, B. melitensis, B. suis, B. ovis, B. microti, and O. anthropi. As can be seen in Figure 3B, B. microti but not B. abortus produced an amino lipid with the migration pattern predicted for OH-OL (Vences-Guzmán et al., 2011), and results similar to those of B. microti were obtained for O. anthropi but not for the other Brucella spp. tested (not shown). These observations support the interpretation that O. anthropi and B. microti LpxO are OL hydroxylases and are fully consistent with the aforementioned genomic and chemical evidence. Accordingly, Brucella lpxO should be named olcC. To confirm this, we examined the amino lipids of a non-polar olcC mutant in B. microti (BmiΔolcC). As predicted, this mutant did not synthesize OH-OL and complementation with a plasmid containing B. microti olcC restored the wild-type phenotype (Figure 3C). Furthermore, introducing this plasmid or a plasmid carrying O. anthropi olcC into B. abortus resulted in the synthesis of OH-OL (Figure 3C and Supplementary Figure S2). No difference in polymyxin sensitivity was observed in these constructs or the mutant BmiΔolcC when compared to the corresponding parental strains.

LptA, LpxE, and OlsC Are Not Required for Brucella Virulence in Laboratory Models

Brucella abortus, B. melitensis, and B. suis have been shown to multiply in murine and human monocyte-derived dendritic cells while interfering with their activation and maturation and reducing both antigen presentation and an effective adaptive response (Billard et al., 2007; Martirosyan et al., 2011; Conde-Álvarez et al., 2012; Gorvel et al., 2014; Papadopoulos et al., 2016). To assess whether LptA, LpxE, and OL β-hydroxylase (OlsC) were involved, we compared parental and mutant strains of B. melitensis, B. abortus, and B. microti in mouse BMDCs. As shown in Figure 4, the kinetics of multiplication of the mutants and wild-type strains were similar. We also performed a phenotypic characterization of MHC II and co-stimulatory
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FIGURE 3 | The Brucella lpxO orthologs encode an acyl hydroxylase acting on ornithine lipids. (A) Pathway of synthesis of ornithine lipids in α-2 Proteobacteria (adapted from Geiger et al., 2010); the ORFs of B. abortus and B. microti are indicated, whereas B. microti, B. vulpis, and O. anthropi contain an intact olsC acyl hydroxylase gene, B. abortus and other Brucella spp. carry a frame-shift in the olsC homolog. (B) Lipid profile of B. abortus wild-type (Ba-parental) and B. microti wild-type (Bmi-parental) showing the absence or presence, respectively, of OH–OL. (C) Amino lipid profile of B. abortus wild-type (Ba-parental), B. microti wild-type (Bmi-parental), B. microti deleted in olsC (Bmi ΔolsC), the cognate reconstituted mutant (Bmi ΔolsCpOlsC), and B. abortus wild-type carrying a plasmid with the B. microti olsC gene (Ba-parentalpOlsC).

receptors CD86 and CD80 (Figure 5). In agreement with previous studies, these analyses showed that activation and maturation was only partially induced in BMDC infected with B. melitensis and B. abortus (Martirosyan et al., 2011). In addition, a similar partial-activation profile was evident both for B. microti, for which no previous studies exist in infected BMDC, and all of the tested mutants obtained for each of the three Brucella spp.

The mouse model has been widely used for testing Brucella virulence (Grilló et al., 2012). In this model, the LptA and LpxE mutants and the parental strains behaved identically (Figure 6 upper panels). Deletion of olsC in B. microti did not alter the CFU/spleen profile produced by this species which is characterized by a lower lethal dose in mice as well as a faster clearance from mouse spleens (Jiménez de Bagüés et al., 2010; Figure 6, lower left panel). Moreover, when we tested whether the expression of B. microti olsC in B. abortus could affect virulence, we found no differences between the B. microti olsC-carrying and the wild-type B. abortus strains (Figure 6, lower right panel).
FIGURE 5 | LptA, LpxE, and OlsC deletions do not significantly impact the intrinsic immunogenicity of Brucella. Each point represents the mean ± standard error of the median intensity of surface receptor expression in dendritic cells treated with Brucella strains or derived mutants. E. coli LPS was used as a positive control for dendritic cell activation.

DISCUSSION

In this work we investigated three Brucella ORFs that according to homologies with genes of known function in other pathogens could modify the lipid A and contribute to further altering the LPS PAMP of representative Brucella species. The results show that, whereas Brucella LptA modifies the lipid A, this is not the case for lpxE and lpxO (redesignated olsC), the former encoding a putative phosphatase acting on an unidentified OM molecule and the latter for an enzyme with OlsC activity.

Our data strongly suggest that B. melitensis LptA is involved in the addition of pEtN to lipid A, homologous proteins carrying out this function are not uncommon in Gram-negative pathogens and modulate the properties of lipid A. In Salmonella Typhimurium, Shigella flexneri, E. coli, Vibrio cholerae, Helicobacter pylori, Haemophilus ducreyi, N. gonorrhoeae, and N. meningitidis pEtN reduces the binding of cationic bactericidal peptides by balancing the negative charge of lipid A (Needham and Trent, 2013; Trombley et al., 2015). Conversely, pEtN promotes binding to N. gonorrhoeae lipid A of factors that downregulate the complement cascade and thwart building of the membrane-attack complex and opsonophagocytosis (Lewis et al., 2013). N. meningitidis pEtN also promotes adhesion of non-encapsulated bacteria to endothelial cells (Takahashi et al., 2008). Indeed, properties that parallel some of those observed for the above-listed pathogens can also be attributed to the pEtN transferase counterpart in Brucella. An intact lptA was related to polymyxin B resistance in B. melitensis and the introduction of B. melitensis lptA into B. abortus increases polymyxin B resistance to the level of B. melitensis, suggesting that LptA function is severely impaired in B. abortus. This is in agreement with the presence of a frame-shift in B. abortus lptA encompassing the consensus sequence, which makes likely that it codes for a protein with no or residual enzymatic activity. Previous analyses are contradictory with regard to the presence (Casabuono et al., 2017) or absence (Moreno et al., 1990) of ethanolamine in B. abortus lipid A but the materials analyzed differ in methods of extraction and presence of B. abortus lipid A markers, such as...
very long chain fatty acids (VCLFA). Although further chemical and enzymatic analyses are necessary for a definite conclusion, our results strongly suggest that, if present, pEtN is in much less amounts in *B. abortus* than in *B. melitensis* lipid A. It is also worth noting that such genetic and phenotypic differences in the lipid A of *B. abortus* and *B. melitensis* could relate to differences in biological properties. The LPS of *B. abortus* and *B. melitensis* is a poor activator of the complement cascade, and this property has been traced to the core and lipid A structure (Moreno et al., 1981; Conde-Álvarez et al., 2012; Fontana et al., 2016). Since *B. abortus* is less resistant than *B. melitensis* to normal serum (González et al., 2008), it is tempting to suggest that, like in *N. gonorrhoeae*, *B. melitensis* pEtN could sequester regulatory elements enhancing complement resistance in this species.

Concerning LpxE, phosphatases acting on lipid A have at least been shown in *Francisella tularensis*, *H. pylori*, *Porphyromonas gingivalis*, and *Capnocytophaga canimorsus*, bacteria where lipid A dephosphorylation is involved both in resistance to bactericidal peptides and the reduction of TLR-4-dependent recognition (Needham and Trent, 2013). Although these properties are displayed by the LPS of *B. abortus* and *B. melitensis* (Martínez de Tejada et al., 1995; Lapaque et al., 2006; Conde-Álvarez et al., 2012), our results do not support a role for BALpxE as a lipid A phosphatase. This is consistent with genomic analysis showing that, whereas in bacteria where LpxE acts on lipid A the gene is located together with *lptA* in an operon (Tran et al., 2006; Renzi et al., 2015), *Brucella* lpxE is instead located upstream of three sequences annotated as pseudogenes and downstream, but in the opposite direction, of a cystathionine beta-lyase. On the basis of the data shown here, the origin of monophosphoryl lipid A in *Brucella* remains to be explained. Further, we believe it unlikely to be an artifact resulting from the hydrolytic steps used to obtain lipid A and instead favor the hypothesis of the existence of an as yet unidentified lipid A phosphatase.

LpxE belongs to the type 2 family of phosphatases that can act on lipid A but also on phosphatidylglycerol phosphate, phosphatidic acid, sphingosine phosphate, and lysophosphatidic acid (Brindley and Waggoner, 1998; Sciarr and Morris, 2002). Significantly, LpxE from *Agrobacterium*, although predicted to be a lipid A phosphatase, dephosphorylates...
phosphatidyl glycerophosphate (Karbarz et al., 2009) to generate phosphatidylglycerol, a cell envelope phospholipid. Indeed, a hypothetical phosphatidyl glycerophosphate phosphatase activity of Brucella LpxE could account for both the polymyxin B sensitivity of the mutated bacteria and the unaltered mass spectra of the lipid A of the mutant. Such a modification of a phospholipid could be meaningful by itself on account of the LpxE-dependent bactericidal peptide resistance but there are other possibilities. In some bacteria (i.e., Rhizobium) phosphatidylglycerol is a precursor for the synthesis of amino lipids such as l-ysyl-phosphatidylglycerol. This synthesis is induced by acid pH and brings about resistance to daptomycin and polymyxin B (Sohlenkamp et al., 2007; Ernst and Peschel, 2011; Arendt et al., 2012). Interestingly, whereas the BaΔlpxE mutant is impaired for growth at pH 6, the parental B. abortus becomes more resistant to cationic peptides (L. Palacios-Chaves and R. Conde-Álvarez, Unpublished observations). These observations suggest the existence in Brucella of pH-dependent envelope modifications that require a functional LpxE. Research is in progress to elucidate the mechanisms behind the increased resistance at acid pH and the implication regarding a role for LpxE.

In S. Typhimurium, Pseudomonas aeruginosa, Bordetella bronchiseptica, Legionella pneumophila, and Klebsiella pneumoniae, LpxO is a Fe2+/α-ketoglutarate-dependent dioxygenase that catalyzes the hydroxylation of the 3′-secondary acyl chain of lipid A. LpxO has been implicated indirectly in stress responses at the envelope level (Needham and Trent, 2013) and, in K. pneumoniae, it has been shown to be relevant in vivo by increasing bactericidal peptide resistance and reducing the inflammatory responses (Llobet et al., 2015). However, as discussed above, previous chemical analysis (Velasco et al., 2000) of lipid A and the evidence presented here indicate that the Brucella lpxO homolog is not a lipid A hydroxylase but rather an OlsC whose mutation, in contrast with LpxO, does not result in increased sensitivity to polymyxin B. This absence of an effect on polycation resistance is in keeping with both the lack of activity on lipid A and the fact that OL do not play a major role in resistance to polycationic bactericidal peptides in B. abortus (Palacios-Chaves et al., 2011). At the same time, it would also appear to rule out, the involvement of this protein in the metabolism of succinate in B. microti as has been previously suggested (Audic et al., 2009).

Previous data showing lpta, lpxE, and lpxO to be involved in modulating the properties of the OM in a way that in some cases confers in vitro resistance to innate immunity bactericidal peptides, complement, and cytokine responses (Needham and Trent, 2013) have been drawn upon as evidence for a role in virulence. However, to the best of our knowledge, a role in vivo has thus far been shown only for lpxO from K. pneumoniae (Llobet et al., 2015). Moreover, contrasting results have been obtained with mutants both showing bactericidal peptide sensitivity in vitro and no phenotype in vivo have been reported for at least H. ducreyi (Trombley et al., 2015) and may reflect the complexities of the infection processes and/or the inadequacies of the currently available in vivo models. Despite their effect on the envelope, our results show that Brucella lpta, lpxE, or olsC do not play a role in the ability of Brucella to replicate in BMDC and do not modulate the activation and maturation profile in these cells. Similarly, the mouse model did not reveal any effect on its ability to colonize and multiply in the spleen. However, further experimental work in the natural hosts and alternative routes of infection might provide evidence on the role in virulence of these genes. The fact that lpta and olsC are not functional in all Brucella spp. must therefore be considered in the context of the models used. While the absence of a functional lpta in B. abortus suggests that the gene is not essential for the virulence of this species we cannot conclude it to be totally irrelevant. Differences between B. melitensis and B. abortus related to lpta could explain the higher invasiveness of the former species noted by early researchers in studies carried out in guinea pigs, animals that are highly susceptible to brucellosis (Braude, 1951). This possibility together with the presence of intact lpta and olsC in Ochrobactrum and B. microti is also compatible with the hypothesis that they represent ancestral characters that are liable to be lost in the absence of a selective pressure during the intracellular life cycle or, in the case of lpta, that is no longer present in the ruminant host species (i.e., cattle) to which B. abortus is characteristically associated.

ETHICS STATEMENT

Female BALB/c mice (Charles River, France) were kept in cages with water and food ad libitum under P3 biosafety conditions in the facilities of “Centro de Investigación Médica Aplicada” (registration code ES31 2010000132) 2 weeks before and during the experiments. The procedures were in accordance with the current European (directive 86/609/EEC) and Spanish (RD 53/2013) legislations, supervised by the Animal Welfare Committee of the University of Navarra, and authorized by the “Gobierno de Navarra” [CEEA045/12 and E36-14 (045-12E1)].

AUTHOR CONTRIBUTIONS


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