Brucella abortus depends on pyruvate phosphate dikinase and malic enzyme but not on Fbp and GlpX fructose-1,6-bisphosphatases for full virulence in laboratory models
Zúñiga-Ripa, Amaia; Barbier, Thibault; Conde-Álvarez, Raquel; Martínez-Gómez, Estrella; Palacios-Chaves, Leyre; Gil-Ramírez, Yolanda; Grilló, María Jesús; Letesson, Jean-Jacques; Iriarte, Maite; Moriyón, Ignacio

Published in:
Journal of Bacteriology

DOI:
10.1128/JB.01663-14

Publication date:
2014

Document Version
Early version, also known as pre-print

Link to publication
Citation for published version (HARVARD):

General rights
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain.
- You may freely distribute the URL identifying the publication in the public portal.

Take down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Download date: 27. Dec. 2020
Brucella abortus depends on pyruvate phosphate dikinase and malic enzyme but not on Fbp and GlpX fructose-1,6-bisphosphatases for full virulence in laboratory models

Amaia Zúñiga-Ripa,1 Thibault Barbier,2 Raquel Conde-Álvarez,1 Estrella Martínez-Gómez,1 Leyre Palacios-Chaves,1,9 Yolanda Gil-Ramírez,1,∗ María Jesús Grilló,3 Jean-Jacques Letesson,2 Maite Iriarte,1,‡ and Ignacio Moriyón1,‡, *

1 Departamento de Microbiología e Instituto de Salud Tropical, Universidad de Navarra, 31008 Pamplona, Spain.
2 Research Unit in Biology of Microorganisms – URBM, NARILIS, UNAmur, B-5000 Namur, Belgium.
3 Grupo de Sanidad Animal, Instituto de Agrobiotecnología (CSIC-Universidad Pública de Navarra-Gobierno de Navarra). Campus de Arrosadía, 31006, Pamplona, Spain.

†Present address: Center for Infection and Immunity, Queen’s University Belfast Health Sciences Building, 97 Lisburn Rd. Belfast, UK BT9 7AE, UK
¥ Present address: Lev2050, CEIN, 31110, Noain, Spain.
‡These authors contributed equally to this work

*Corresponding author: Ignacio Moriyón. Mailing address, Departamento de Microbiología y Parasitología, Universidad de Navarra, C/ Irunlarrea 1, 31008 Pamplona, Spain, Phone: +34 948425600 (ext. 806356); Fax: +34 948 425649
Email address: imoriyon@unav.es

Key words: Brucella, metabolism, virulence, intracellular, pyruvate phosphate dikinase, malic enzyme, glyoxylate, gluconeogenesis, phosphoenolpyruvate carboxykinase, fructose bisphosphatase.

Running title: Brucella metabolism in virulence models.
The brucellae are the etiological agents of brucellosis, a worldwide-distributed zoonosis. These bacteria are facultative intracellular parasites, and thus are able to adjust their metabolism to the extra and intracellular environments encountered during an infectious cycle. However, this aspect of Brucella biology is imperfectly known and the nutrients available in the intracellular niche are unknown. Here, we investigated the central pathways of C metabolism used by Brucella abortus by deleting the putative fructose-1,6-bisphosphatases (fbp and glpX), phosphoenolpyruvate carboxykinase (pckA), pyruvate phosphate dikinase (ppdK) and malic enzyme (mae) genes. In gluconeogenic but not in rich media, growth of mutants ΔppdK and Δmae was severely impaired and growth of the double Δfbp-ΔglpX mutant was reduced. In macrophages, only ΔppdK and Δmae showed reduced multiplication, and studies with ΔppdK confirmed that it reached the replicative niche. Similarly, only ΔppdK and Δmae were attenuated in mice, the former being cleared by week 10 and the latter persisting longer than 12 weeks. We also investigated the glyoxylate cycle. Although aceA (isocitrate lyase) promoter activity was enhanced in rich medium, aceA disruption had no effect in vitro or on multiplication in macrophages or mouse spleens. The results suggest that B. abortus grows intracellularly using a limited supply of 6 C (and 5 C) sugars that is compensated by glutamate and possibly other amino acids entering the Krebs cycle without a critical role of the glyoxylate shunt.
INTRODUCTION

Brucella is a genus of gram-negative bacteria that groups the causative agents of brucellosis, a worldwide-extended zoonosis severely affecting animal production and human welfare. Three species, B. abortus, B. melitensis and B. suis, are the most common causes of brucellosis in domestic livestock and humans. These brucellae can grow both in vitro and within host cells, and their pathogenicity results largely from their capacity to escape a prompt detection by innate immunity and the use of a type IV secretion system to reach the replicative niche, an endoplasmic reticulum derived vacuole (1-6). In this compartment, these bacteria multiply extensively, which shows their ability to use efficiently substrates provided by the host. However, there is only sparse information on either the nature of these substrates or the metabolic pathways used in the replicative niche (7).

In vitro, most strains of B. abortus, B. melitensis and B. suis grow in several simple chemically defined media (8). Among these, Gerhardt’s medium contains glycerol, lactate, glutamate and mineral salts plus nicotinic acid, thiamine, pantothenic acid and biotin as growth factors. This medium supports growth better than other simple defined media, including those that provide glucose as the C source (8), and this has been attributed to the ability of these bacteria to use glutamate very efficiently through the tricarboxylic acid cycle (TCA) (9). Indeed, growth in this medium is clear proof of the ability of these bacteria to carry out gluconeogenesis in vitro. Concerning the catabolism of glucose, it has been accepted that it proceeds through the pentose phosphate pathway (in conjunction with TCA), rather than through the glycolysis or Entner–Doudoroff pathways (Fig. 1). This view, although supported by early radiorespirometric and enzymatic studies (10,11) performed with the attenuated B. abortus S19 vaccine, is not consistent with the presence of the genes putatively encoding all the enzymes of the Entner-Doudoroff pathway (12). It is possible that the multiple defects in S19 (13) and/or the
experimental conditions in those early experiments precluded detection of key enzymes of the Entner-Doudoroff pathway (12).

Concerning intracellular metabolism, some data come from randomly obtained mutants that show attenuation in human or mouse macrophages or in HeLa cells. Genes (putative functions) identified in this manner include, in *B. suis* 1330, a gguA homologue (*gluP*, presumably involved in sugar uptake), some erythritol catabolism genes, *gnd* (6-phosphogluconate dehydrogenase, required for using glucose through the pentose phosphate pathway), *rbsk* (ribose kinase), *pyc* (anaplerotic pyruvate carboxylase), *pgi* (phosphoglucose isomerase) and genes related to the biosynthesis of amino acids (14-16); in *B. melitensis* 16M, *dbsA* and *ugpA* (ribose and glycerol-3-phosphate transporters, respectively), *glpD* (glycerol-3-phosphate dehydrogenase) and some erythritol catabolism genes (17,18); and in *B. abortus* 2308, *gluP* (previously shown to encode an active glucose and galactose transporter in this species (19)), *gnd* (6-phosphogluconate dehydrogenase), *gltD* (glutamate synthase) and *gcvB* (glycine dehydrogenase) (20). Also in *B. abortus* 2308, *dxs* (an isoprenoid biosynthesis transketolase) and *mocC* (rhizopine or inositol catabolism) were identified as expressed in macrophages using a fluorescent reporter (21).

Proteomic analyses have also provided clues on the metabolism of brucellae in the host. Forty-eight h after infection of mouse macrophages with *B. suis* 1330, Al Dahouk et al. (22) found an important reduction of proteins putatively involved in energy, protein and nucleic acid metabolism. Some exceptions were ribitol kinase, glyceraldehyde-3-P-dehydrogenase and the isocitrate lyase (*AceA*) of the glyoxylate cycle. However, other studies in *B. suis* 1330 do not support the use of the glyoxylate cycle within host cells (15). Lamontagne et al. (23) analyzed *B. abortus* 2308 protein expression 3, 20 and 44 h after infection of RAW 264.7 macrophages. They found that multiple proteins associated with sugar uptake, TCA, the pentose phosphate shunt and the subsequent generation of pyruvate were down-regulated 3 h after infection. At 24 h, several proteins involved in sugar metabolism and transport were also reduced. Enzymes
associated with protein and amino acid catabolism were mainly increased early (3 h) but also 24 h after infection, when bacteria were already in vacuoles derived from the endoplasmic reticulum. This was also the case of enzymes involved in glutamate synthesis, suggesting conversion of amino acids into glutamate and \( \alpha \)-ketoglutarate. Accordingly, amino acid-based alternatives may be the preferred solution for *B. abortus* to derive precursors for the TCA cycle and ancillary routes during the midpoint time course of infection. At later times, the same authors observed an increase in proteins involved in transport, suggesting that the endoplasmic reticulum is able to supply at least some of the substrates required for bacterial growth. Likewise, the pentose phosphate shunt seemed to partially resume its functions.

Although the information given by these studies is valuable, the central metabolic pathways used by *Brucella* during infection remain unclear. The results are contradictory in some cases, as for the glyoxylate cycle or the metabolic activity in cells. In addition, some studies suggest the availability of sugars in the replicative niche whereas others indicate that amino acids could be the preferred C source \textit{in vivo}, which may require a gluconeogenic metabolism. Indeed, apparently conflicting data may result from the use of different host cell lines, different times of analysis, polarity of mutations and other experimental conditions. Moreover, there might be some variation among *B. suis* 1330, *B. melitensis* 16M and *B. abortus* 2308, as suggested by the known differences in oxidative rates of sugars and amino acids (24). In this work, we attempt to answer some aspects of the central metabolic pathways used by *B. abortus* in the host. For this purpose, we focused our research on genes putatively involved in classical gluconeogenesis, the anabolic pathways bridging TCA and the triose-phosphate pathway, and the glyoxylate cycle. We constructed in-frame mutants in genes coding for key enzymes and tested them in complex and chemically defined media and for multiplication within cultured cells and for persistence in the mouse model (25). Together with some of the previous analyses, our observations suggest a model of *B. abortus* metabolism in which, although TCA supplies...
molecules necessary for biosynthesis and subsequent growth, classical fructose-1,6-
bisphosphatases Fbp and GlpX are not necessary and 6 and/or 5 C molecules for polymer
biosynthesis are obtained mostly from the intracellular milieu.

MATERIAL AND METHODS

Bacterial strains and growth conditions. The bacterial strains and plasmids used in this study
are listed in Table 1S (supplemental material), and their origin and characteristics are described
in previous works (26-28). The strains resulting from the genetic manipulations described below
were characterized according to standard Brucella typing procedures: i.e. colonial morphology
after 3 days of incubation at 37°C, crystal violet-oxalate exclusion, urease, acriflavine
agglutination, sensitivity to Tb, Wb, Iz and R/C phages, agglutination with anti-A and anti-M
monospecific sera, CO₂ and serum dependence, and susceptibility to thionin blue, fuchsins and
safranin (24). Bacteria were routinely grown in standard Peptone-Yeast Extract-Glucose broth
(Biomerieux) or in this media supplemented with agar (TSA). The following antibiotics were used
at the indicated concentrations: kanamycin (Km; 50 μg/mL), nalidixic acid (Nal; 25 μg/mL),
chloramphenicol (Cm; 20 μg/mL) and/or gentamicin (Genta; 100 μg/mL or 25 μg/mL) (all from
Sigma). When needed, media was supplemented with 5% sucrose (Sigma). All strains were
stored at –80°C in skim milk (Scharlau). To study the phenotype of the metabolic mutants,
Peptone-Yeast Extract-Glucose or the medium of Gerhardt (henceforth Glutamate-Lactate-
Glycerol) were used (29). The components (for 1 L) of the latter were: glycerol (30 g), lactic acid
(5 g), glutamic acid (1.5 g), thiamine (0.2 mg), nicotinic acid (0.2 mg), pantothenic acid (0.04
mg), biotin (0.0001 mg), K₂HPO₄ (10 g), Na₂S₂O₅·5H₂O (0.1 g), MgSO₄ (10 mg), MnSO₄ (0.1
mg), FeSO₄ (0.1 mg), NaCl (7.5 g). The pH was adjusted to 6.8-7.

Growth measurements. Inocula pre-conditioned to the conditions in the test medium (Peptone-
Yeast Extract-Glucose, Glutamate-Lactate-Glycerol, Glycerol-Glutamate, Glycerol-Lactate or
Glutamate-Lactate) were prepared as follows. First, the strains to be tested were inoculated into

10 mL of Peptone-Yeast Extract-Glucose in a 50 mL flask and incubated at 37ºC with orbital shaking for 18 h. These exponentially growing bacteria were harvested by centrifugation, resuspended in 5 mL of the test medium at an optical density (O.D. 600nm) of 0.1 and incubated at 37ºC with orbital shaking for 18 h. Then, these pre-conditioned bacteria were harvested by centrifugation, resuspended at an O.D. 600nm of 0.1 (0.05 starting in the Bioscreen apparatus) in the same test medium in Bioscreen multi-well plates (200 µL/well) and cultivated in a Bioscreen C (Lab Systems) apparatus with continuous shaking at 37ºC. Absorbance values at 420-580 nm were automatically recorded at 0.5 h intervals over a 120 to 300 h period. All experiments were performed in triplicate. Controls with culture medium and no bacteria were included in all experiments.

**DNA manipulations.** Genomic sequences were obtained from the Kyoto Encyclopedia of Genes and Genomes (KEGG) database (http://www.genome.jp/kegg/). Searches for DNA and protein homologies were carried out using the National Center for Biotechnology Information (NCBI; http://www.ncbi.nlm.nih.gov/) and the European Molecular Biology Laboratory (EMBL) - European Bioinformatics Institute server (http://www.ebi.ac.uk/). Primers were synthesized by Sigma-Genosys (Haverhill, UK). 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).

In-frame deletion mutants in \( fbp \) and \( glpX \) were constructed by polymerase-chain reaction (PCR) overlap using genomic DNA of \( B. abortus \) 2308 as the DNA template. Primers were designed using the \( B. abortus \) 2308 sequences available in KEGG (http://www.genome.jp/kegg/). For the construction of the \( fbp \) mutant, two PCR fragments were
generated: oligonucleotides $\text{fbp-F1} (5'$-GTAGCCAAAAAGCCAGGT-3')$ and $\text{fbp-R2} (5'$-GCAACCAGAACCAGAGGA-3')$ were used to amplify a 203 bp fragment including codons 1–14 of the $\text{fbp}$ ORF as well as a 161 bp fragment upstream of the $\text{fbp}$ start codon, and oligonucleotides $\text{fbp-F3} (5'$-TCCTCTGTTCTGCTGGCGTGCCGAGGATGGATA-3')$ and $\text{fbp-R4} (5'$-CATTTGCCGCTTCCATGA-3')$ were used to amplify a 193 bp fragment including codons 327–341 of the $\text{fbp}$ ORF and a 148 bp fragment downstream of the $\text{fbp}$ stop codon. Both fragments were ligated by PCR using oligonucleotides $\text{fbp-F1}$ and $\text{fbp-R4}$ for amplification, and the complementary regions between $\text{fbp-R2}$ and $\text{fbp-F3}$ for overlapping. The resulting fragment, containing the $\text{fbp}$ deletion allele, was cloned into pCR2.1 (Invitrogen) to generate plasmid pAZI-1, sequenced to ensure that the reading frame was maintained, and subcloned into the BamHI and the XbaI sites of the suicide plasmid pJQKm (30). The resulting mutator plasmid (pAZI-2) was introduced into $\text{B. abortus}$ 2308 by conjugation (26). Integration of the suicide vector was selected by Nal and Km resistance, and the excisions (generating both the $\text{fbp}$ mutant $\text{BAB}\overline{\text{fbp}}$ and a sibling revertant strain carrying an intact gene $\text{BAB}\text{fbp}$-sibling revertant) were then selected by Nal and sucrose resistance and Km sensitivity. The resulting colonies were screened by PCR with primers $\text{fbp-F1}$ and $\text{fbp-R4}$, which amplified a fragment of 396 bp in the mutant and a fragment of 1332 bp in the sibling revertant strain. The mutation resulted in the loss of about 98% of the $\text{fbp}$ ORF and the mutant strain was called $\text{BAB}\overline{\text{fbp}}$.

The $\text{glpX}$ mutant was constructed in a similar way. Primers $\text{glpX-F1} (5'$-ACGGTGATTCTGGTGACACA-3')$ and $\text{glpX-R2} (5'$-CGAGCTCAGGTGAGGATG-3')$ were used to amplify a 576 bp fragment including 61 bp of the $\text{glpX}$ ORF as well as 515 bp upstream of the $\text{glpX}$ start codon, and primers $\text{glpX-F3} (5'$-CATTCTCACACTGGAGCTCGATACGACAGATCCGGACGAG-3')$ and $\text{glpX-R4} (5'$-CATCATACAGTTGCCCAGATGCGAG-3')$ were used to amplify a 574 bp fragment including 371 bp of the $\text{glpX}$ ORF and 203 bp downstream of the $\text{glpX}$ stop codon. Both fragments were ligated by...
overlapping PCR using primers glpX-F1 and glpX-R4, and the fragment containing the deletion allele was cloned into pCR2.1 to generate plasmid pAZI-3, sequenced to confirm that the glpX ORF had been maintained, and subcloned in pJQKm to produce the mutator plasmid pAZI-4. This plasmid was then introduced into B. abortus 2308 and the deletion mutant generated by allelic exchange was selected by Nal and sucrose resistance and Km sensitivity and by PCR using oligonucleotides glpX-F1 and glpX-R4, which amplified a fragment of 1150 bp in the deletion strain and a fragment of 1705 bp in the BABglpX-sibling revertant strain. The mutation resulted in the loss of approximately 56% of the glpX ORF, and the mutant was called BABΔglpX.

To construct the BABΔfbpΔglpX double mutant, the mutator plasmid pAZI-4 was introduced into strain BABΔfbp. After allelic exchange, the double mutant was selected as described above using primers glpX-F1 and glpX-R4. BABΔaceA was constructed using the same strategy. Oligonucleotides aceA-F1 (5'-TGACAAGATATCGCCAAAACAC-3') and aceA-R2 (5'-CGAAGGGATGAGGCTGTAAA-3') amplified a 238 bp fragment, including codons 1-10 of the aceA ORF and 208 bp upstream of the aceA start codon. Oligonucleotides aceA-F3 (5'-TTTACAGCCTCATCCCTTCGGAAACCGCACAGTTCAAGC-3') and aceA-R4 (5'-GGATCAAGAGATTCCCCAGT-3') amplified a 278 bp fragment including codons 420-430 of the ORF aceA and 245 bp downstream of the aceA stop codon. Both fragments were ligated by overlapping PCR using oligonucleotides aceA-F1 and aceA-R4. The PCR product was cloned into pCR2.1 to generate pAZI-7, sequenced and subcloned into pJQKm to produce the suicide plasmid pAZI-8. B. abortus 2308 mutants were selected by PCR using oligonucleotides aceA-F1 and aceA-R4. PCR products were 1743 bp in BABaceA-sibling revertant strain and 738 bp in BABΔaceA. This mutation eliminated 78% of the aceA ORF.
For the construction of the \textit{pckA} mutant, oligonucleotides \textit{pckA}-F1 (5’-TGTTTGCAGTTTTCCACACC-3’), \textit{pckA}-R2 (5’-AATCGAAGCGGCCTTATTGT-3’) and \textit{pckA}-R4 (5’-TCTTGCGAATAACAGCCAAAA-3’) were used. Primes \textit{pckA}-F1 and \textit{pckA}-R2 amplified a 219 bp fragment, which included codons 1-13 of the \textit{pckA} ORF and 180 bp upstream of the \textit{pckA} start codon.

Primers \textit{pckA}-F3 and \textit{pckA}-R4 amplified a 319 bp fragment including the last 37 codons of the \textit{pckA} ORF and 208 bp downstream of the \textit{pckA} stop codon. Both PCR products were ligated by overlapping PCR using \textit{pckA}-F1 and \textit{pckA}-R4, cloned into pCR2.1 to generate plasmid pAZI-5 and subsequently subcloned into the BamHI and the XbaI sites of the suicide plasmid pJQKm. The resulting mutator plasmid pAZI-6 was introduced into \textit{B. abortus} 2308, where it was integrated in the chromosome. A second recombination generated the excision of the plasmid.

The resulting colonies were screened by PCR (with \textit{pckA}-F1 and \textit{pckA}-R4) amplifying a fragment of 538 bp in the mutant and a fragment of 1864 bp in the sibling revertant strain. The mutant strain was called \textit{BAB\textbackslash\text{\textregistered}pckA} and lacked the 71.14\% of the \textit{pckA} ORF.

\textit{BAB\textbackslash\textregisteredppdK} was constructed using primers \textit{ppdK}-F1 (5’-CTCCCGATTCATTTTTCACG-3’) and \textit{ppdK}-R2 (5’-TGCTCATTTCCAGGCAGGT-3’) to amplify a 288-bp fragment including the first 103 bp of the \textit{ppdK} ORF, as well as 185 bp upstream of the \textit{ppdK} start codon and primers \textit{ppdK}-F3 (5’-AACTGGCTGAAATGAGCGGTCTGACTATGTGCC-3’) and \textit{ppdK}-R4 (5’-TCAACGCATCAAGCAGAG-3’) to amplify a 220 bp including the last 86 bp of the \textit{ppdK} ORF and 134 bp downstream of the \textit{ppdK} stop codon. Both fragments were ligated by overlapping PCR using primers \textit{ppdK}-F1 and \textit{ppdK}-R4 and the fragment obtained, containing the deletion allele, was cloned into pCR2.1 to generate pMZI-1, sequenced to confirm that the reading frame had been maintained, and subcloned into pJQKm to produce the mutator plasmid pMZI-2. This plasmid was introduced into \textit{B. abortus} 2308 and both the deletion mutant and the sibling revertant strain generated by allelic exchange was selected by Nal and sucrose resistance and
Km sensitivity, and by PCR using ppdK-F1 and ppdK-R4 which amplified a fragment of 508 bp in BABΔppdK and a fragment of 2983 bp in BABppdK-sibling revertant strain. The mutation generated resulted in the loss of the 93% of ppdK.

Primers ppdKII-F1 (5'-CTCCGATTCATTTTTCACG-3'), ppdKII-R2 (5'-CTGCTCATTTCAGCCAGGTT-3'), ppdKII-F3 (5'-AACCTGGCTGAAATGAGCAGCGGGTCTCGACTATGTGTCC-3') and ppdKII-R4 (5'-TCAACGCATCAAAGCAGAAG-3') were used to obtain the mutator plasmid pAZI-10. This plasmid was introduced into B. abortus 2308 to obtain a ppdK mutant that maintained only the 34 first amino acids of PpdK (BABΔppdK-II). This mutant had the same phenotype as the one previously described. Thus, the mutator plasmid pAZI-10 was also introduced into strain BABΔpckA carrying the pckA mutation to obtain the double mutant BABΔpckAΔppdK.

BABΔmae was constructed using primers mae-F1 (5'-TATGACGCGCAGTCTGCTAG-3') and mae-R2 (5'-TCGGATAGCGATGGAAGAAC-3') to amplify a 341 bp fragment including 76 bp of the mae ORF, as well as 265 bp upstream of the mae start codon, and primers mae-F3 (5'-GTTCTTCCATCGCTATCCGAGCGAAGCCAATCTTCTGGTA-3') and mae-R4 (5'-CGCCATAAAACGAACCTCAA-3') to amplify a 376 bp including 227 bp of the mae ORF and 149 bp downstream of the mae stop codon. Both fragments were ligated by overlapping PCR using primers mae-F1 and mae-R4 and the fragment obtained, containing the deletion allele, was cloned into pCR2.1 to generate pMZI-3, sequenced to confirm that the mae ORF had been maintained, and subloned into pJQKm to produce the mutator plasmid pMZI-4. This plasmid was then introduced into B. abortus 2308 and the mutant and sibling revertant strains generated by allelic exchange were selected by Nal and sucrose resistance and Km sensitivity and by PCR using oligonucleotides mae-F1 and mae-R4, which amplified a fragment of 717 bp in the deleted strain and a fragment of 2739 bp in the sibling revertant strain. The mutation generated resulted in the loss of the 87% of the mae ORF.
To check the different mutations, we used internal primers (gene-R5) hybridizing in the deleted regions.

For complementation, a plasmid carrying ppdK was constructed using the Gateway cloning Technology (Invitrogen). Since the sequence of ppdK from B. abortus and B. melitensis is 99% identical, the clone carrying ppdK was extracted from the B. melitensis ORFEOMA and the ORF subcloned into plasmid pRH001 (31) to produce plasmid pAZI-19. This plasmid was introduced into BABΔppdK by mating with E. coli S17 λpir and the conjugants harboring this plasmid (designated as BABΔppdK pAZI-19) were selected by plating the mating mixture onto tryptic soy agar (TSA)-Nal-Km plates. For the construction of BABΔfbpΔglpX pAZI-21, gene fbp was amplified from BAB-parental using primers fbp-Fp (5'-GGGATCCATGCTTCTGAAAGGGTGGTACCG-3') and fbp-R4 (5'-CATTTGCCGCTTCCATGA-3') and cloned into pCR2.1. The resulting plasmid was sequenced, and the fbp gene was subcloned into the BamHI and XhoI sites of the vector pBBR1MCS1. The resulting plasmid (pAZI-21) was introduced into BABΔfbpΔglpX by conjugation (see above). BABΔfbpΔglpX pAZI-23 was constructed following the same strategy using primers glpX-F1 (5'-ACGGTGATTCTGGTGACACA-3') and glpX-R4 (5'-CATCATACAGTTGCCGATGG-3') to amplify glpX.

Gene expression studies. To determine whether aceA was expressed in vitro, its promoter was fused with the luciferase reporter gene. To this end, aceAp-F (5'-GGGATCCTAGTTGCGCTCGATCAGATT-3') and aceAp-R (5'-TTCTAGACATTTCGGTGTCCTCCTCGT-3') (respectively containing BamHI and XbaI sites; underlined) were used to amplify a 382bp region containing the ATG and the aceA promoter from B. abortus 2308 genomic DNA. This PCR product was verified by electrophoresis and ligated into the vector pGEM-T Easy (Promega), thereby originating plasmid pAZI-17. Then, the insert was digested with BamHI and XbaI and ligated to the pSKOriTKmluxAB to generate...
plasmid pAZI-18. This plasmid was introduced into E. coli S17λ pir Nal^R and then transferred by conjugation to B. abortus 2308 Nal^R Km^S. Cells were plated on TSA NalKm. Positive clones gave a 510 bp band when verified by PCR using aceA-F and luxAB-R. The resulting strain was called B. abortus pBABaceA-luxAB. To measure luciferase activity, fresh B. abortus pBABaceA-luxAB were adjusted (OD_{600nm}) to 0.4 in saline and finally, 50 or 200 μL were added to flasks with 10 mL of Peptone-Yeast Extract-Glucose or Glutamate-Lactate-Glycerol, respectively. Growth was followed by measuring absorbance at OD_{600nm} and 1 mL aliquots were taken at selected intervals to measure the luminescence in Relative Luminescence Units (RLU) after addition of 100 μL ethanol:decanal (1:1).

Cell infections and intracellular trafficking. In vitro infection assays were performed in RAW 264.7 macrophages (ATCC TIB-71) and HeLa cells (ATCC CCL-2) cultured in Dulbecco's Modified Eagle Medium (DMEM; Gibco) with 10% (v/v) heat-inactivated fetal bovine serum (Gibco), 1% (v/v) L-glutamine 200 nM (Sigma Aldrich) and 1% (v/v) non-essential amino acids (Gibco). Then, 24-well plates were seeded with 1 × 10^5 cells/well, and macrophages and HeLa cells were infected at a multiplicity of infection of 50:1 and 200:1 (bacteria:cell), respectively, by centrifuging the plates at 400 × g for 10 min at 4°C. After incubation for 15 min at 37°C under a 5% CO₂ atmosphere, extracellular bacteria were removed with four DMEM washes followed by Genta treatment (100 μg/mL) for 90 min. Then, fresh medium supplemented with 25 μg/mL of Genta was added and incubation carried on. Two, 24 and 48 h later, cells were washed three times with 100 mM phosphate buffered saline (pH 7.2), lysed with 0.1% (v/v) Triton X-100 in phosphate buffered saline, and plated on TSA to determine the number of intracellular bacteria. All experiments were performed in triplicate (32). Results were expressed as mean and standard error (n=3) of individual log_{10} CFU/well. Statistical comparison of means was performed by a one-way ANOVA followed by the Fisher’s Protected Least Significant Differences (PLSD) tests (33).
For immunofluorescence microscopy, RAW 264.7 macrophages and HeLa cells were grown on coverslips and inoculated with bacteria as described above. Cells were fixed in 3% paraformaldehyde in 100 mM phosphate buffered saline (pH 7.2) at 37°C for 10 min. Cells were washed twice with phosphate buffered saline and permeabilized with 0.1% (v/v) Triton X-100 and 3% bovine serum albumin (Sigma), for 30 min. Coverslips were incubated with primary antibodies for 45 min at room temperature, washed three times in the same phosphate buffered saline supplemented with 3% bovine serum albumin, and then incubated with the appropriate secondary antibodies. Coverslips were washed three times with phosphate buffered saline and once with H₂O and mounted onto glass slides using Mowiol 4–88. Samples were examined and images acquired using a Leica TCS SP5 laser scanning confocal microscope at the “UNAmur” (Namur, Belgium). The primary antibodies used for immunofluorescence microscopy were rabbit anti-calnexin (SPA-860, Stressgen) and a mouse anti-S-LPS monoclonal antibody (A76/12G12/F12). The secondary antibodies were donkey anti-rabbit IgG conjugated to Alexa Fluor 488, (Invitrogen), and goat anti-mouse IgG conjugated to Alexa Fluor 546 (Invitrogen). For lysosomal labeling, the primary antibody used was rat anti-mouse (DBHS) and the secondary antibody was goat anti-rat anti IgG conjugated to Alexa Fluor 633 (Invitrogen).

Assays in mice. Female BALB/c mice (Charles River, France) were kept in cages with water and food ad libitum, and accommodated under P3 biosafety containment conditions 2 weeks before and during the experiments, in the facilities of the "Instituto de Agrobiotecnología" (registration code ES/31-2016-000002-CR-SU-US). The animal handling and other 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 "Universidad Pública de Navarra", and authorized by the competent authority of "Gobierno de Navarra". To prepare inocula, TSA grown bacteria were harvested, adjusted spectrophotometrically (O.D.₆₀₀nm =0.170) in 10 mM phosphate buffered saline (pH 6.85) and diluted in the same diluent up to
approximately 5 × 10^5 colony forming units (CFU)/mL (exact doses were assessed retrospectively). For each bacterial strain, five mice were intraperitoneally inoculated with 0.1 mL/mouse and the CFU number in spleen was determined at different weeks post-inoculation as described previously (33). The identity of the spleen isolates was confirmed by PCR. The individual number of CFU/spleen was normalized by logarithmic transformation, and the mean log CFU/spleen values and the standard deviation were calculated for each group of mice (n=5). Statistical comparisons were performed by a one-way ANOVA followed by the Fisher’s Protected Least Significant Differences (PLSD) tests (33).

RESULTS

Dysfunction of *B. abortus* fbp, glpX, ppdK and mae but not of pckA or aceA homologues affects growth on gluconeogenic substrates *in vitro*. The conversion of fructose-1,6-bisP into fructose-6-P mediated by the cognate bisphosphatase(s) (FBPases) is the only irreversible step among those taking part in gluconeogenesis in *Brucella* (Fig. 1). Therefore, FBPase activity is strictly necessary to grow under gluconeogenic conditions.

As in *E. coli* and *Salmonella*, the *B. abortus* 2308 genome presents two ORFs of putative FBPases: BAB2_0364 and BAB1_1292. The former is predicted to encode a protein of 340 amino acids that belongs to the class I FBPases (Fbp), and the latter a 328 amino acid protein ortholog to *E. coli* GlpX, a FBPase of class II (Fig. 1). In *E. coli*, Fbp is connected to the production of fructose-6-phosphate for nucleotide, polysaccharide and aromatic amino acid biosynthesis. GlpX, on the other hand, belongs to an operon (glpFKX) related to phospholipid biosynthesis (34). The putative *B. abortus glpX*, however, seems isolated and not part of any obvious operon.

We constructed mutants carrying in frame deletions in the putative *fbp* (*BABΔfbp* mutant) and *glpX* (*BABΔglpX*) as well as in both genes (*BABΔfbpΔglpX*), and tested their growth in a complex (Peptone-Yeast Extract-Glucose) medium and in the chemically defined medium of
Gerhardt (Glutamate-Lactate-Glycerol). In the complex medium, the three mutants grew with generation times and final yields similar to those of \textit{B. abortus} 2308 (BAB-parental) (Fig. 2A.1).

\textit{BAB}-parental showed reduced growth rates and final yields in Glutamate-Lactate-Glycerol, as expected, and we obtained identical results with either \textit{BAB}\textsubscript{Δ}\textit{fbp} or \textit{BAB}\textsubscript{Δ}\textit{glpX} (Fig. 2A.2). On the other hand, \textit{BAB}\textsubscript{Δ}\textit{fbp}\textsubscript{Δ}\textit{glpX} produced a markedly lower increase in turbidity in the minimal medium (Fig. 2A.2). Complementation with plasmid pAZI-21 carrying \textit{fbp} or with pAZI-23 carrying \textit{glpX} restored the ability to grow in Glutamate-Lactate-Glycerol to levels close to that of the parental strain (Supplemental Material; Fig. S3). These experiments strongly suggest that \textit{B. abortus} Fbp and GlpX are functional, an interpretation reinforced by their reciprocal complementation (i.e., Fbp complemented \textit{BAB}\textsubscript{Δ}\textit{glpX} and, conversely, GlpX complemented \textit{BAB}\textsubscript{Δ}\textit{fbp}). We then tested combinations of two C sources (Glycerol-Glutamate, Glycerol-Lactate or Glutamate-Lactate). Whereas \textit{BAB}\textsubscript{Δ}\textit{glpX} was not affected, \textit{BAB}\textsubscript{Δ}\textit{fbp} showed retarded growth only in the absence of glycerol (i.e. in Glutamate-Lactate; Fig. 3, upper panels). Since GlpX is the FBPase remaining in \textit{BAB}\textsubscript{Δ}\textit{fbp} and this mutant grew normally when glycerol was present, this result suggests that, as in \textit{E. coli} (34), GlpX is related to glycerol metabolism. Finally, although retarded and diminished, the double \textit{BAB}\textsubscript{Δ}\textit{fbp}\textsubscript{Δ}\textit{glpX} mutant still showed significant growth (Fig. 3, upper panels).

During these experiments we noticed that \textit{BAB}\textsubscript{Δ}\textit{fbp}\textsubscript{Δ}\textit{glpX} inoculated broths did not produce a homogeneous growth. This was clearly observed when the double mutant was grown in Glutamate-Lactate-Glycerol in side-arm flasks instead of the automated Bioscreen system. Under these conditions, the bacteria formed macroscopic aggregates (Fig. 4) settling on the bottom of the flasks that indicated a profound surface modification consistent with an altered biosynthesis of envelope molecules. In summary, although Fbp and GlpX deficiency did not abrogate bacterial multiplication, they were required not only for full growth but also for production of normal cells.
Next, we investigated the pathways that supply pyruvate or PEP for gluconeogenesis. For
this purpose, we carried out a genomic search for homologues of the genes encoding the
enzymes connecting the TCA cycle and the triose-phosphate pathway in bacteria (Fig. 1). We
identified homologues of *pdh* (pyruvate dehydrogenase), *pyk* (pyruvate kinase), *pckA*
(phosphoenolpyruvate carboxykinase), *ppdK* (pyruvate-phosphate dikinase), *pyc* (pyruvate
carboxylase), and *mae* (malic enzyme) but not of *pyrcK* (pyruvate carboxykinase), *pps*
(phosphoenolpyruvate synthase) or *ppc* (phosphoenolpyruvate carboxylase) (Fig. 1). Since Pyc,
Pdh and Pyk catalyze irreversible catabolic steps, we studied the *pckA*, *ppdK* and *mae*
homologues.

*B. abortus* 2308 BAB1_2091 is annotated as a pseudogene in some data bases
(http://www.genome.jp/kegg/) but not in others (http://biocyc.org/). This ORF encodes a protein
of 491 amino acids that bears a 77% similarity with the PckA of the phylogenetically related
*Agrobacterium tumefaciens*, a protein of 536 amino acids known to be functional (35). However,
*B. abortus* BAB1_2091 is separated by a stop codon from an intergenic region that together with
BAB1_2090 encodes the last 45 amino acids present in *A. tumefaciens* PckA. The frame-shift
that generates this stop codon is present also in all *B. abortus* strains sequenced so far
(http://www.genome.jp/kegg/). Nevertheless, *B. abortus* PckA conserves the IGGSYAGE-KKS
domain (amino acids 190 to 202) specifically required for the carboxykinase activity (36) as well
as the phosphate binding site (G--G-GKT; amino acids 236 to 243) and ATP and metal binding
sites, so that its functionality is not obviously compromised. In contrast, the putative *B. abortus*
2308 PpdK (encoded by BAB1_0525) represents a complete protein of 887 amino acids with the
PEP-binding (amino acids 19 to 376) and the TIM-barrel (amino acids 530 to 883) domains
characteristic of PEP-utilizing enzymes.

To test whether these ORFs encoded enzymes are required for growth under gluconeogenic
conditions, we constructed the non-polar mutants *BABΔpckA* and *BABΔppdK*. Moreover, since
both PckA and Ppdk catalyze reactions eventually leading to PEP (Fig. 1), we excluded their reciprocal complementation by constructing the double mutant BABΔpckAΔppdK. We then compared the growth of BAB-parental and the mutants in Peptone-Yeast Extract-Glucose and in Glutamate-Lactate-Glycerol. Whereas we did not observe differences in the growth of BABΔpckA and BAB-parental in these two media, both BABΔppdK and BABΔpckAΔppdK had a markedly reduced growth in Glutamate-Lactate-Glycerol (Fig. 2B). Complementation with plasmid pAZI-19 carrying ppdK restored the phenotype (Supplemental Material; Fig. S1), and the sibling revertant control (see Material and Methods) conserved the wild-type phenotype (not shown). These results strongly suggest that PpdK is functional and that, consistent with the frame shift in pckA, PckA does not synthesize PEP from oxaloacetate in B. abortus. The possibility that the role of B. abortus PckA is not detectable under these in vitro conditions seems less likely.

We followed a similar strategy to study the putative mae. BAB1_1036 encodes a protein of 774 amino acids annotated as a NADP (or NAD)-dependent enzyme involved in malate metabolism (http://www.genome.jp/kegg/; http://biocyc.org/). The N-terminal domain (amino acids 28-160) and the NADP (or NAD)-binding domain (amino acids 172-409) characteristic of the malic enzyme are conserved in the BAB1_1036 predicted protein. Mutant BABΔmae (deleted in the region encoding amino acids 26 to 673) displayed a small reduction in growth in Peptone-Yeast Extract-Glucose and a more marked one in Glutamate-Lactate-Glycerol (Fig. 2C). The impairment was not as accentuated as that of BABΔppdK (compare Fig. 2 panels B and C). In these experiments, mae mutants with the above-described phenotype were consistently obtained and the control sibling revertants of wild-type phenotype recovered after the last recombination event (see Material and Methods and Supplemental Material; Fig. S2).

Taken together, these results support the hypothesis that B. abortus Mae supplies pyruvate for PpdK to produce PEP for gluconeogenesis, lactate being a complementary source of pyruvate in
Glutamate-Lactate-Glycerol (Fig. 1). We tested this hypothesis further by using media containing only two of the three C substrates of this chemically defined medium. BABΔmae did not grow in Glycerol-Glutamate, showed optimal growth in Glycerol-Lactate, and reduced growth in Glutamate-Lactate (Fig. 3, lower panels). These results are fully consistent with the predicted role of Mae (Fig. 1) and suggest that, although B. abortus 2308 can use glycerol, glutamate and lactate, provision of the latter cannot completely replace the Mae pathway. This interpretation is also supported by the fact that whereas PpdK dysfunction in BABΔppdK severely impaired growth in Glycerol-Lactate and Glutamate-Lactate, this mutant grew in Glycerol-Glutamate (Fig. 3, lower panels), two substrates able to act as sources of PEP and pyruvate through the triose-P and Mae pathways, respectively (Fig. 1). Indeed, all these results are in agreement with the early studies that led to the formulation of the simple medium of Gerhardt as well as with the nutritional studies that showed the preferential use of glutamate and the complementary role of glycerol and lactate in B. abortus (9). Also, the demonstration by Marr et al. (37) of the ability of B. abortus to generate pyruvate (and alanine) from glutamate supports an activity of Mae in this bacterium (Fig. 1).

Since the putative Mae was active in B. abortus 2308, we investigated whether malate replenishment could occur through the classical glyoxylate pathway (Fig. 1) or the PEP-glyoxylate cycle. The latter cycle combines the operation of PckA with the glyoxylate cycle enzymes and operates in E. coli under conditions of glucose limitation (38). In these pathways, isocitrate lyase (AceA) cleaves isocitrate to yield glyoxylate and succinate, and a malate synthase (AceB) condenses glyoxylate and acetyl-CoA to produce malate (39-41). The genome of B. abortus 2308 carries only one putative aceA (ORF BAB1_1631) and one putative aceB (ORF BAB1_1663). The predicted AceA is a protein of 429 amino acids with 61% identity and 76% similarity to E. coli AceA, and it conserves the amino acids required for the enzymatic activity and the assembly of the tetrameric enzyme (42-47). The predicted AceB has 728 amino
acids with 59% identity and 74% similarity to *E. coli* malate synthase G, and conserves the catalytic site and the amino acids interacting with acetyl-CoA (48,49). Accordingly, we constructed a non-polar BABΔaceA mutant truncated in the 409 central amino acids. This mutant did not show growth differences with BAB-parental in Peptone-Yeast Extract-Glucose and Glutamate-Lactate-Glycerol (data not shown), even though these media contain acetogenic substrates (glucose, glycerol, lactate and serine, threonine and alanine) (50). Since the genomic analysis strongly suggests the presence of the glyoxylate cycle, we examined this point further by constructing a luciferase reporter under the control of the AceA promoter. Although growth curves were similar, luciferase activity was considerably higher in Peptone-Yeast Extract-Glucose than in Glutamate-Lactate-Glycerol (Fig. 5), as expected if the glyoxylate cycle becomes active on dependence of the abundance of acetogenic substrates. In such a case, the lack of phenotype in complex media could be explained if the glyoxylate cycle plays only a subsidiary role in this rich medium, and the experiments do not rule out that the possibility that it becomes important under other nutritional conditions.

*B. abortus* mutants in *ppdK* and *mae* but not in *fbp, glpX, pckA* or *aceA* show lower multiplication rates in macrophages. *B. abortus* is characteristically able to multiply intracellularly in professional phagocytes (51). We thus investigated the ability of the above-described mutants to multiply in macrophages using the parental strain and the attenuated virB mutant (unable to reach the replicating vacuole) as controls. Figure 6A shows that BABΔfbpΔglpX replicated like BAB-parental, even though these bacteria differed in growth under gluconeogenic conditions *in vitro* (see above). BABΔppdK (Fig. 6B) and BABΔmae (Fig. 6C) multiplied in macrophages, although at lower overall rates than BAB-parental both 24 (p<0.0001) and 48h (p<0.0001) after infection. On the other hand, mutation of *pckA* had no effect either by itself or combined with the *ppdK* deletion (Fig. 6B).
These results suggest that, albeit impaired in growth, BABΔppdK and BABΔmae are still able to reach the replicative intracellular niche. Since Mae and PpdK belong to the same pathway, we selected BABΔppdK (the mutant blocked in the upper step of the pathway; Fig. 1) to confirm that the metabolic dysfunction did not prevent these bacteria from reaching the endoplasmic reticulum-derived replicating niche. Figure 7A shows that, in contrast to the virB mutant, BABΔppdK and the parental bacteria were similar in intracellular distribution.

*B. abortus* can also penetrate and multiply in epithelial cells (51). We found that the behavior of BABΔppdK in macrophages was reproduced in HeLa cells (Fig 7A and B). In addition, we found that the CFU of the mutant and BAB-parental in HeLa cells did not differ 2 h after infection (not shown), indicating that they were similar with regards to penetration.

*B. abortus* mutants in *ppdK* and *mae* but not in *pckA*, *fbp*, *glpX* or *aceA* are attenuated in mice. Virulent *B. abortus* is able to establish spleen infections in mice that characteristically develop in four phases: (i), onset phase (spleen colonization; first 48 h); (ii), acute phase (from the third day to weeks 2-4 when bacteria reach maximal numbers); (iii), chronic steady phase (weeks 2-4 to 12), where the bacterial numbers plateau; and (iv), chronic declining phase, during which brucellae are eliminated. The ability to induce a marked splenomegaly is also a characteristic of virulent brucellae. (25). Using this model, we first studied the BABΔfbpΔglpX mutant and found that it did not differ from BAB-parental in either the CFU/spleen profile or the splenomegaly induced (Fig. 8A). Similarly, we did not observe attenuation for BABΔpckA (Fig. 8B), which is remarkable because pckA expression increases in *B. abortus* mutants in BvrR/BvrS, a master regulator of *B. abortus* virulence (52). Nevertheless, this result is consistent with the genomic features of pckA and with the above-described experiments in vitro and in macrophages.

BABΔppdK failed to reach the chronic steady phase typical of virulent brucellae yielding significantly lower CFU counts after week 2 (Fig. 8B). Clearly indicative of the strong attenuation
of BABΔppdK, we did not recover any bacteria from the spleens of 3 out of the 5 mice examined at post-infection week 12, and this mutant induced less splenomegaly than the virulent bacteria (Fig. 8B). Consistent with the observations that showed no additive effect of the mutations of ppdK and pckA in macrophages or in vitro, the results of BABΔpckAΔppdK in mice paralleled those of BABΔppdK (Fig. 8B). BABΔlmae produced a CFU/spleen profile that differed from that of either BAB-parental or BABΔppdK. Although not affected in the first 48 h (onset phase; not shown) this mutant showed a lower multiplication rate during the acute phase (Fig. 8C) that was reminiscent of the lower multiplication rates observed in macrophages. Strikingly, even though CFU/spleen numbers were lower than those of the wild type strains, BABΔlmae produced a chronic steady phase with reduced splenomegaly.

Finally, we tested BABΔaceA in mice. In a first experiment, the mutant did not show attenuation at weeks 2, 8 and 12 (not shown). It has been reported that isocitrate lyase is essential for Salmonella persistence in mice during chronic infection but dispensable for acute lethal infection (53). Similarly, isocitrate lyase is dispensable in the acute phase of Mycobacterium tuberculosis infection in lung macrophages of mice but facilitates persistence during the chronic phase (54). Accordingly, we repeated the mouse infections and determined the number of BABΔaceA CFU in spleens 16 and 24 weeks later. However, we did not find any difference between this mutant and the parental strain (not shown). Therefore, even though we did not rule out the possibility that the glyoxylate cycle plays a role under conditions different from those tested in vitro, we concluded that this shunt is not essential for B. abortus multiplication and persistence in the laboratory models used. Since the PEP-glyoxylate pathway relies on both AceA and PckA, this conclusion can be extended to this cycle.

DISCUSSION

B. abortus lacks the genes necessary for the metabolism of glycogen or poly-beta-hydroxyalkanoates, the two C reserve materials used by prokaryotes (55). Thus, these bacteria
depend on nutrients provided by the host to multiply intracellularly and, accordingly, they need a supply of (at least) 6 C skeletons or to carry out gluconeogenesis. To investigate these possibilities, we deleted ORFs that could be encoding enzymes of critical steps of gluconeogenesis, or of steps providing the necessary precursors. The genomic characteristics of the ORFs analyzed and the phenotypes observed in vitro support the hypothesis that they encode FBPases, a pyruvate phosphate dikinase and a malic enzyme of B. abortus. Moreover, the analyses in cells show that the proteins coded for by ppdK and mae become necessary once the replicative niche is reached, as expected from metabolic mutants.

Since FBPases are essential for gluconeogenesis, the observation that growth in gluconeogenic media was not abolished in the Fbp-GlpX double mutant is intriguing and several hypotheses can be considered to explain these results. The existence of a third FBPase is the first and more obvious possibility. Up to now, five different types of FBPases (I to V) have been described in prokaryotes (56). Whereas FBPases of classes IV and V are restricted to Archaea and their close hyperthermophilic Aquifex bacterial group, many bacteria have dual combinations of class I (Fbp homologues), class II (GlpX homologues), and class III FBPases (34,56). However, a genomic search for Bacillus subtilis YydE homologues (the prototype of class III FBPases (57)) in the Brucellaceae only revealed an imperfect match (a hypothetical protein of 218 amino acids with a 32% identity with the 671 amino acids in YydE) in Ochrobactrum anthropi and none in Brucella. This is in agreement with the fact that no bacterial genome has been described to carry a combination of classes I and III (34). Also, it has been described that some E. coli carry two class II FBPases (GlpX and YggF (56)) but genomic analysis of all phosphatases in B. abortus fail to identify clear candidates for any phosphatase close to GlpX and Fbp (Supplemental Material: Fig. S4). This genomic evidence and the Fbp-GlpX reciprocal complementation (i.e. the fact that only the double mutant shows phenotype in vitro in the absence of glycerol) suggest that these are the main and possibly the only FBPases
in *B. abortus*. An alternative to the third FBPase hypothesis is the existence of an atypical

 gluconeogenesis less efficient than the classical one. Hypothetically, a fructose-6-P aldolase
could take part in gluconeogenesis. This has been described in *E. coli* K-12 where ORF b0825
encodes an enzyme that catalyzes the reversible conversion of fructose-6-P to
dihydroxyacetone and glyceraldehyde-3-P (58). This aldolase is different from that operating in
the pentose-phosphate cycle and its physiological role is uncertain. However, the only
homologous in *B. abortus* (ORF BAB1_1813) is annotated as transaldolase and, in addition to
the fact that it should represent the enzyme of the pentose-phosphate cycle, the identity (30%) is
below the threshold considered to be significant (59). Obviously, a rigorous analysis of these two
hypotheses requires enzymatic analysis of double deficient cells. Finally, the significance of the
mucoid aggregates produced by the double FBPase mutant cannot be disregarded since this
phenotype suggests that the growth observed does not correspond to a natural condition.

Bearing in mind that enzymatic analyses are necessary to reach definite conclusions, our
results and those of previous works with *B. abortus* 2308 offer insight into some global models of
the metabolism of these bacteria during intracellular life in the host.

A first model (gluconeogenic model) can be proposed on the ability of *B. abortus* to grow in
the defined medium of Gerhardt and on the proteomic studies in macrophages that suggest that
*B. abortus* shifts to an amino acid-based metabolism in which the glutamate pool is increased
(23). According to this model, molecules like glycerol, lactate or pyruvate and amino acids
channeled to oxaloacetate, keto-glutarate or pyruvate are the main substrates, and molecules of
6 and 5 C are derived from the latter. In this regard, it is remarkable that dysfunction of two
major FBPases did not bring about any perceptible attenuation either in cells or mice. Although
the reduced growth of the *BABΔfbpΔglpX* mutant *in vitro* precludes a clear-cut conclusion, the
contrasting *in vivo* and *in vitro* multiplication and the mucoid phenotype of the double mutant in
the minimal medium are more consistent with models alternative to the gluconeogenic one.
Moreover, two lines of evidence indicate that glucose (or closely related hexoses) is available in the host. First, a *B. abortus* 2308 (and *B. suis* 1330) GluP (glucose/galactose transporter) mutant has been identified as attenuated in signature tagged experiments in (15,20,20). Second, it has been reported recently that the multiplication of *B. abortus* in alternatively activated macrophages increases when the intracellular glucose levels are artificially increased (60). Based on these observations, an almost opposite model proposes a main role for 6 C sugars in the replicative niche (and 5 C sugars if we assume that the evidence obtained in *B. melitensis* also applies to *B. abortus*; see Introduction). These sugars would provide trioses-phosphate through the pentose phosphate cycle and serve also as precursors for biosynthesis of envelope polymers. This second model, however, does not account for the attenuation observed for the *ppdK* and *mae* mutants, which strongly suggests that molecules necessary for growth are derived from the TCA in vivo.

A third model proposes that there is a limited supply of 6 C (and 5 C) sugars that is compensated by glutamate, alanine and other amino acids. Those sugars would be used mostly or exclusively for biosynthesis of envelope polymers and for the pentose-phosphate cycle-dependent biosynthetic reactions. This model is consistent with the results of this and previous works. First, the different phenotype of the *B. abortus* BABΔfbpΔgplx double mutant in vivo and in vitro is more coherent with the hypothesis that classical gluconeogenesis is not extensively used in vivo. In addition, the *B. abortus* 2308 gluP mutant identified in signature-tagged mutagenesis studies is not clearly attenuated at two weeks post-infection and manifests its attenuation at times (8 weeks) that correspond to the chronic phase (20). This suggests that the bacteria do not depend totally on hexoses for intracellular biosynthetic processes, and that this dependence is manifested at late times perhaps as the result of changes in the replicative vacuole (see below). Indeed, the infection experiments performed in alternatively activated macrophages suggest that, although available, glucose is a limiting factor for *B. abortus* growth.
at least during the chronic phase (60). Proteomic studies carried out with *B. abortus* 2308 show that expression of two proteins of the dihydroxyacetone kinase complex (Dha) of the PEP-carbohydrate phosphotransferase system (PTS) is reduced throughout infection in macrophages (23). Although this has been interpreted to mean that reduced PTS expression may be the result of a short supply of sugars within the replicative niche, the *Brucella* PTS lacks the sugar permease unit and is likely to act as a regulatory system coordinating C and N metabolism (61). On the other hand, the signature-tagged mutagenesis and proteomic studies show attenuation of a *gltD* (putatively encoding the small subunit of glutamate synthase) mutant (20) and an increment of enzymes involved in increasing the pool of glutamate (23) that are consistent with the model. Finally, this third model accounts for the attenuation of the *ppdK* and *mae* mutants observed in mice (Fig. 8). Since according to the model sugars are used mostly to construct envelope polymers and for pentose-phosphate cycle derived precursors, additional molecules for biosynthesis must be derived from TCA. PpdK works to produce PEP, which is used to synthesize phenylalanine, tyrosine and tryptophan, glycerolipids and other PEP-derived molecules. Mae supplies pyruvate for PpdK in vivo but TCA would not be the only source of pyruvate. This is suggested by the fact that the *mae* mutant was both delayed in reaching the chronic phase of infection and in lower numbers in the spleen during this phase, which contrasts with the inability of the *ppdK* mutant to generate chronic infections. Interestingly, it has been shown recently that, when provided with multiple carbon sources, *Mycobacterium tuberculosis* differentially catabolizes each carbon source through the glycolytic, pentose phosphate, and/or TCA pathways to distinct metabolic fates, and it has been suggested that this ability reflects an adaptation to pathogenicity (62). Indeed, such ability could also be necessary for *B. abortus* to coordinately use the different substrates proposed for this model.
Although the last model fits the experimental evidence, it is obvious that it represents only a first approach to the situation in the natural hosts because we cannot assume that the niche is static during a chronic infection or uniform among different types of cells. In the above-mentioned signature-tagged mutagenesis studies, Hong et al. (20) have presented evidence for the hypothesis that different set of genes are required during the onset-acute phases and the chronic steady phase. This study identified three putative metabolic genes (\textit{gluP}, \textit{gID} [see above] and \textit{gcvP}) required during the chronic phase but not markedly during the acute phase of infection. Indeed, both the need of a functional \textit{ppdK} for the infection to progress during the acute phase and the different phenotype of \textit{mae} add further support to the hypothesis that different genes are required to a different extent during the course of infection. Moreover, at least macrophages and trophoblastic cells have been clearly associated with \textit{B. abortus} infection in cattle (63) and the physiological characteristics of these cells are different. Also, different spleen cells become colonized at different times after intraperitoneal inoculation of mice (64). Clearly, research is necessary to investigate these aspects of the relationship between metabolism and intracellular multiplication in \textit{B. abortus} and in other species of the genus.

\textbf{ACKNOWLEDGEMENTS}

We are grateful to Professor E. Van Schaftingen for helpful discussions and suggestions. This research was supported by grants from the "Ministerio de Economía y Competitividad" of Spain (AGL2011-30453-C04-00), "Fundación para la Investigación Médica Aplicada" (FIMA) and, in part, by a grant from the "Fond National de la Recherche Scientifique" (FNRS) (convention FRFC N° 2452110, Belgium) and by the Interuniversity Attraction Poles Programme initiated by the Belgian Science Policy Office. A.Z.-R., was supported by a post-doctoral grant from FIMA, T.B. has a PhD grant as "Aspirant du FNRS" and L.P.-C. and R.C-A. were funded in part by "Departamento de Educación del Gobierno de Navarra" ("Programa ANABASID").
REFERENCES


phospholipid, is necessary for full virulence of the intracellular bacterial parasite *Brucella abortus*. Cell Microbiol. 8:1322-1335.


**FIGURE LEGENDS**

**FIG. 1.** Conventional central metabolic pathways (glycolysis, gluconeogenesis, Entner-Doudoroff, pentose-phosphate, TCA and glyoxylate) of bacteria. Dashed arrows indicate steps for which no putative genes can be identified in *B. abortus*. Red arrows indicate the steps studied in this work. Abbreviations used in the figure are: AcCoA (acetyl-CoA), AceA (isocitrate lyase), AceB (malate synthase), Acn (aconitate hydratase), AKG (α-ketoglutarate), Akgdh (α-ketoglutarate dehydrogenase), Cgs (cyclic-glucan synthesis), CIT (citrate), DHAP (dihydroxyacetone phosphate), Eda (keto-deoxy-phosphogluconate aldolase), Edd (6-phospho-D-gluconate dehydratase), E4P (erythrose-4-phosphate), Fba (fructose bisphosphate aldolase), Fbp (fructose-1,6-bisphosphatase), Fum (fumarase), FUM (fumarate), F1,6dP (fructose-1,6-bisphosphate), F6P (fructose-6-phosphate), GaIE (UDP-glucose-4 epimerase), GAP (glyceraldehyde-3-phosphate), Gnd (6-phosphogluconate dehydrogenase), Glk (glucokinase), GlpX (fructose-1,6-bisphosphatase), GltA (citrate synthase), GLX (glyoxylate), G1P (glucose-1-phosphate), G6P (glucose-6-phosphate), ICIT (isocitrate), Idh (isocitrate dehydrogenase), KDPG (2-keto-3-deoxyphosphogluconate), LPS (lipopolysaccharide), Mae (malic enzyme), MAL (malate), Mdh (malate dehydrogenase), OAA (oxaloacetate), Pfk (phosphofructokinase), PckA (phosphoenol pyruvate carboxikinase), Pdh (pyruvate dehydrogenase), PEP (phosphoenol pyruvate), Pgi (phosphoglucone isomerase), Pgl (lactonase), Pgm (phosphoglucomutase), Ppc (phosphoenol pyruvate carboxylase), PpdK (pyruvate phosphate dikinase), Pps (phosphoenol pyruvate synthase), Pyc (pyruvate carboxylase), Pyk (pyruvate kinase), PYR (pyruvate), Pyrck (pyruvate carboxikinase), RIB5P (ribulose-5-phosphate), Sdh (sucinate dehydrogenase), Stk (succinyl-CoA synthethase), SUC (succinate), SucCoA (succinyl-CoA), Tkt (transketolase), TpiA (triose phosphate isomerase), X5P (xylulose-5-phosphate), Zwf (glucose-6-phosphate dehydrogenase), 6PG (6-phosphogluconate), 6PGL (6-phosphogluconolactone).
FIG. 2. Growth curves in Peptone-Yeast Extract-Glucose (A.1, B.1 and C.1) and Glutamate-Lactate-Glycerol (A.2, B.2 and C.2) of BAB-parental and mutants BABΔfbp, BABΔglpX, BABΔfbpΔglpX, BABΔpckA, BABΔppdK, BABΔpckAΔppdK and BABΔmae. Each point represents the mean ± standard error (error bars are within the size of the symbols) of optical density (O.D.) values of triplicate samples. The experiment was repeated three times with similar results.

FIG. 3. Growth curves in Glycerol-Glutamate, Glycerol-Lactate and Glutamate-Lactate of BAB-parental and mutants BABΔfbp, BABΔglpX, BABΔfbpΔglpX, BABΔppdK and BABΔmae. Each point represents the mean of triplicate samples (error bars are within the size of the symbols). The experiment was repeated three times with similar results.

FIG. 4. Immunofluorescence analysis of BAB-parental and BABΔfbpΔglpX grown in Glycerol-Lactate-Glutamate. Cells were labeled with anti-smooth-lipopolysaccharide mouse monoclonal antibody A76/12G12/F12 and Alexa Fluor 546 conjugated goat anti-mouse immunoglobulin.

FIG. 5. Luciferase expression under the control of B. abortus aceA promoter in Peptone-Yeast Extract-Glucose and Glutamate-Lactate-Glycerol. The results are representative of three experiments (RLU, relative luminescence units).

FIG. 6. Intracellular multiplication in RAW 264.7 macrophages of BAB-parental and mutants BABΔfbpΔglpX, BABΔpckA, BABΔppdK, BABΔpckAΔppdK and BABΔmae (virB is an attenuated mutant used as a control). Values are the mean ± standard error of triplicate infections and the results shown are representative of three independent experiments.
FIG. 7. Trafficking of BAB-parental, BAB∆ppdK and virB in macrophages and HeLa cells. (A), confocal images of infected RAW 264.7 macrophages and HeLa cells labeled with Moabs to either calnexin (in green) or LAMP-1 (in blue) 24 hours after infection (bacteria are immunostained in red). (B), Percentage of calnexin-positive vacuoles of HeLa cells that contain bacteria at 24 and 48 hours post-infection.

FIG. 8. Bacterial multiplication (upper panels) and spleen weights (lower panels) generated in BALB/c mice by BAB parental and mutants BAB∆fbp∆glpX (A); BAB∆pckA, BAB∆ppdK and BAB∆pckA∆ppdK (B); and BAB∆mae (C). Each point is the mean ± standard deviation (n=5) of the Log of CFU or grams per spleen. Statistical differences with BAB parental were significant (p < 0.001) from week 4 onwards for BAB∆ppdK and from week 2 onwards for BAB∆mae. BAB∆pckA∆ppdK was also attenuated at weeks 8 and 12 (p < 0.001) (weeks 4 and 6 were not tested).