**Bacillus wiedmannii biovar thuringiensis**: A Specialized Mosquitocidal Pathogen with Plasmids from Diverse Origins

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**Abstract**

*Bacillus cereus sensu lato* also known as *B. cereus* group is composed of an ecologically diverse bacterial group with an increasing number of related species, some of which are medically or agriculturally important. Numerous efforts have been undertaken to allow presumptive differentiation of *B. cereus* group species from one another. FCC41 is a *Bacillus* sp. strain toxic against mosquito species like *Aedes aegypti*, *Aedes (Ochlerotatus) albifasciatus*, *Culex pipiens*, *Culex quinquefasciatus*, and *Culex apicinus*, some of them responsible for the transmission of vector-borne diseases. Here, we report the complete genome sequence of FCC41 strain, which consists of one circular chromosome and eight circular plasmids ranging in size from 8 to 490 kb. This strain harbors six crystal protein genes, including *cry24Ca*, two *cry4*-like and two *cry52*-like, a *cry41*-like parasporin gene and multiple virulence factors. The phylogenetic analysis of the whole-genome sequence of this strain with molecular approaches places this strain into the *Bacillus wiedmannii* cluster. However, according with phenotypical characteristics such as the mosquitocidal activity due to the presence of Cry proteins found in the parasporal body and *cry* genes encoded in plasmids of different sizes, indicate that this strain could be renamed as *B. wiedmannii* biovar *thuringiensis* strain FCC41.

**Key words**: *Bacillus cereus* group, whole-genome, *cry* genes, mosquitocidal strain.

**Introduction**

The current description of the genus *Bacillus* integrates a great number of species genetically extremely heterogeneous. In the last century, the species in the genus were redefined many times resulting in the reclassification of its constituent species (Bhandari et al. 2013). In particular, *Bacillus cereus sensu lato*, also known as the *B. cereus* group, was initially formed by few species and in the last years several new species had been included in this classification (Bazinet 2017). Among them, *Bacillus anthracis* is the etiological agent of anthrax and its major virulence factors are the anthrax toxins and an antiphagocytic polyglutamic capsule, both encoded by two large plasmids pXO1 and pXO2 (Moayeri et al. 2015). *Bacillus cereus sensu stricto*, or *B. cereus* as it is usually called, is ubiquitous in nature and an emerging human pathogen that causes gastroenteritis, considered the third most important cause of food intoxication incidents in Europe and an opportunistic human pathogen associated with other local and systemic infections. *Bacillus cereus* strains encode a range of toxins and other extracellular virulence factors on the chromosome, and it has been shown that the same genes are present and actively expressed in *Bacillus thuringiensis* strains (Maughan and Van der Auwera 2011).

*Bacillus thuringiensis* has been studied for its properties as an insect pathogen useful as microbial agent for insect pests in agriculture and insect vectors control. This bacterium produces a parasporal inclusion body composed primarily by the insecticidal proteins Cry and Cyt that are mainly encoded in megaplasmids. These kinds of inclusions or crystals must be
ingested by a susceptible insect and after some biochemical processes, disrupt the cell osmotic balance, then the midgut integrity, resulting in gut paralysis and the insects’ death (Adang et al. 2014).

One of the most recently identified species of this bacterial group, *Bacillus wiedmannii*, produces hemolysin BL toxins and nonhemolytic enterotoxins and has been described as hydrolyze arginine negative being unable to produce acid from fermentation of sucrose and can be differentiated from the other *B. cereus* species by sequence comparison analysis (Miller et al. 2016).

In the past, species of *B. cereus* group were classified based on their habitats, morphological or physiological characteristics, 16S rRNA gene sequences and characteristics such as the presence or absence of virulence plasmids that confer pathogenicity against mammals or insects. Their classification according with comparative genomics is controversial and shows incongruent phylogenies (Rasko et al. 2005; Liu et al. 2015). In order to demonstrate diverse phylogenetic hypothesis of *B. cereus* group, different molecular approaches have been applied (Guinebretière et al. 2008; Kim et al. 2014; Miller et al. 2016; Bazinet 2017). The use of multiple molecular methods has shown that the species boundaries between members of this bacterial group are difficult to define, forcing to rethink the current descriptions of some *Bacillus* species (Maughan and Van der Auwera 2011).

In a previous work, we characterized an Argentinian strain named as *B. thuringiensis* FCC41 that exhibited higher insecticidal activity against some mosquito species than *B. thuringiensis* subsp. *israelensis*, the reference strain used worldwide for vector control. This isolate produces a rounded crystal harboring two major proteins of ~70–80 kDa and until recently, we could only sequence the complete encoding gene of two of the crystal proteins (Cry), Cry24Ca (Berón and Salerno 2007) and one Cry52-like protein, even though we suspected the presence of more Cry proteins. To determine other toxins and virulence factors that may play a role in the activity of this strain and to understand fully this highly important toxic activity against mosquitos, we have therefore determined its complete genome sequence obtained by next-generation sequencing technologies as presented here. Additionally, we discuss the reclassification of FCC41 strain according with phenotypical and genotypical characteristics as well as the phylogenetic analysis with molecular approaches.

**Materials and Methods**

**Strain and Culture Condition**

The strain FCC41 was originally isolated from soil samples (Berón and Salerno 2006) and cultured in nutrient broth medium (Difco) at 28°C for 16 h under constant shaking at 180 rpm. In order to evaluate the psychrophilic nature of this strain, 100 µl of an overnight growth pre-inoculum were plated on nutrient broth agar plates and incubated at 5, 7, 10, and 12°C during 72 h.

**Genome Sequencing and Assembly**

Bacterial DNA was isolated using the GeneJET Genomic DNA Purification Kit (Thermo Fisher Scientific Inc.). Whole-genome sequencing was performed using Illumina 1500 HiSeq (INDEAR, Rosario, Argentina) and PacBio RSII (Macrogen Inc., Korea) platforms. The Illumina sequencing was conducted with 100 bp paired-end reads from a Nextera XT library (broad size between 150 and 1,500 bp), yielding 2.2 G of sequenced bases. The PacBio sequencing was run in one single-molecule real-time cell (SMRT) with a 20 kb library, which generated 80,337 reads (average read length 10,771 bp) and 865,518,541 read bases. The complete genome was obtained by assembling short and long reads through a hybrid assembly approach using Unicycler software (Wick et al. 2017). Sequencing coverage was calculated with SAMtools (Li et al. 2009) by aligning all reads to the final assembly using Bowtie 2 (Langmead and Salzberg 2012) and minimap (Li 2016). In order to manually verify the final assembly these alignments were subsequently visualized with Integrative Genomics Viewer (Thorvaldsdóttir et al. 2013).

**Genome Annotation**

Genome annotation was performed with the NCBI Prokaryotic Genome Annotation Pipeline (http://www.ncbi.nlm.nih.gov/genome/annotation_prok/) and Prokka (Seemann 2014). Clusters of Orthologous Groups (COG) annotations were computed using WebMGA server (Wu et al. 2011).

**Phylogenetic Analysis**

The sequences of *panC*, *rpoB* and the seven housekeeping genes of the MLST scheme proposed by Priest et al. (2004; http://www.pubmlst.org/bcereus/; Last accessed April 2018), were extracted of the whole-genome sequence of FCC41 strain and analyzed with BTyper software, which compares the sequences of the genes extracted to alleles present in specific online databases (Carroll et al. 2017). Single genes *panC* and *rpoB* sequence comparisons were performed using BLAST with NCBI nonredundant database and the Food Microbe Tracker database (http://www.foodmicrobetracker.com; Last accessed April 2018). For the MLST based phylogeny, as well as for the core genome and Average Nucleotide Identity (ANI) based phylogeny, we selected 35 *B. cereus sensu lato* genomes according to their distribution and representation along the phylogenetic tree previously reported by Bazinet (2017; see supplementary table S1, Supplementary Material online). These genomes were downloaded from NCBI RefSeq, and the sequence of the seven housekeeping genes of all genomes, including FCC41 strain, were extracted and concatenated, and the resulting sequences were aligned.
using ClustalO (Sievers and Higgins 2014). With this alignment we calculated the phylogenetic tree with RAxML performing 500 bootstraps (Stamatakis 2014).

For the core genome-based phylogeny, the bcgTree pipeline automatically extracts 107 essential housekeeping genes found in most bacteria using hidden Markov models, and then performs a partitioned maximum-likelihood analysis using the multiple alignment of the concatenated sequences (Ankenbrand and Keller 2016). The phylogenetic tree was constructed with RAxML (Stamatakis 2014) under a maximum likelihood model. To evaluate node support 100 bootstrap replicates were computed and the consensus tree visualized by FigTree 1.4.2 (http://tree.bio.ed.ac.uk/software/figtree/).

The selected genomes were analyzed by ANI using PyANI (https://github.com/widdowquinn/pyani) in ANIm mode, which uses MUMmer aligner (Kurtz et al. 2004).

**Sequences of Interest Overview**

Virulence factors and antibiotic resistance genes were predicted using BLASTP against the data set of Virulence Factor Database (VFDB; http://www.mgc.ac.cn/VFs/; Last accessed April 2018) and the Comprehensive Antibiotic Resistance Database (CARD; https://card.mcmaster.ca/; Last accessed April 2018), respectively.

Cry protein coding DNA sequences (CDS) were detected using Hidden Markov Modeler (HMMER, Eddy 1998). For that purpose, we constructed a HMMER profile with an alignment of all known deduced amino acid sequences of the Cry haplotype proteins (https://www.btnomeclature.info; Last accessed April 2018) and used it to search through the translated CDS obtained in the automated annotation step. To confirm the results, the positive matches were compared with known Cry proteins with a BLASTP search (Altschul et al. 1990).

**Plasmid Analysis**

Nucleotide sequences of plasmids of FCC41 strain were used to search for similar and complete plasmids DNA sequences in NCBI nonredundant nucleotide database, these plasmids were downloaded and used to analyze regions of shared homology between FCC41 strain plasmids and then were compared with BLAST Ring Image Generator (BRIG) and EasyFig software (Sullivan et al. 2011). On the other hand, in order to identify plasmid minireplicons from FCC41 strain BLASTP analysis was performed against known replication proteins as previously described (Zheng et al. 2013).

**Results and Discussion**

**Whole-Genome Sequencing and Assembly**

To characterize the mosquitocidal FCC41 strain and to find all putative Cry protein coding genes, its whole-genome was sequenced and analyzed. The complete genome was obtained by two different next-generation sequencing methodologies: Illumina HiSeq providing high fidelity, accurate short reads and higher coverage, resulting in an assembly with multiple gaps; and, PacBio SMRT providing longer but less precise reads. By combing both reads with the Unicycler assembly pipeline (Wick et al. 2017) the complete ungapped genome was obtained. The final assembly is comprised of one circular chromosome and eight circular plasmids, with a total genome size of 6,873,481 bp as described in table 1 and visualized in figure 1. This complete genome has been deposited in INSDC (International Nucleotide Sequence Database Collaboration: www.insdc.org) under the GenBank/EMBL/DDBJ accession numbers CP024684 to CP024692.

**Phylogenetic Analysis**

It has been discussed that the B. cereus group should be considered to form a single species with different ecotypes and pathotypes and according with that, the main phenotypical properties originally used to distinguish B. cereus sensu stricto, B. thuringiensis and B. anthracis were related to the presence or absence of large plasmids, which carry the genetic

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**Table 1**

| Features | Size (bp) | G + C Content (%) | No. of Genes | No. of tRNAs | No. of rRNAs | No. of cry-like Protein Genes | Accession Number | Illumina Sequencing Coverage | PacBio Sequencing Coverage |
|----------|-----------|-------------------|--------------|--------------|-------------|-----------------------------|------------------|-----------------------------|---------------------------|
| Chromosome | 55,13,449 | 36.09             | 5,893        | 108          | 39          | –                           | CP024684         | 209.8                       | 76                         |
| pFCC41-1-490k | 4,90,693 | 32.65             | 467          | 1            | –           | –                           | CP024685         | 306.9                       | 100.5                      |
| pFCC41-2-313k | 3,13,432 | 32.46             | 287          | –            | –           | 1                           | CP024686         | 307.4                       | 100.8                      |
| pFCC41-3-257k | 2,57,021 | 34.49             | 259          | –            | –           | 4                           | CP024687         | 607.3                       | 199.1                      |
| pFCC41-4-144k | 1,44,434 | 33.82             | 143          | –            | –           | –                           | CP024688         | 539.3                       | 168.4                      |
| pFCC41-5-125k | 1,24,955 | 34.10             | 134          | –            | –           | 1                           | CP024689         | 727.3                       | 188.4                      |
| pFCC41-6-12k  | 11,912   | 31.51             | 18           | –            | –           | –                           | CP024690         | 309.5                       | 5.6                       |
| pFCC41-7-9k   | 9,217    | 35.20             | 13           | –            | –           | –                           | CP024691         | 947.6                       | 46.8                       |
| pFCC41-8-8k   | 8,368    | 29.36             | 8            | –            | –           | –                           | CP024692         | 732.1                       | 19.2                       |
| Total       | 68,73,481| 35.05             | 7,222        | 109          | 39          | 6                           |                  |                             |                            |
determinants that are responsible for those phenotypes. Despite these differences, the 16S rRNA sequences of most members of B. cereus group have a homology >98%, providing insufficient resolution at the species level (Cohan 2002; Liu et al. 2015; Wang and Ash 2015). In addition to ribosomal genes, panC and rpoB genes are currently being used to differentiate among the most important Bacillus spp. as a more accurate system for phylogenetic analysis (Caamaño-Antelo et al. 2015; Kovac et al. 2016). In particular, the panC sequence comparison analysis performed with BTyper reveals that the FCC41 strain belongs to Group II of the phylogeny proposed by Guinebretière et al. (2008) since it has 99% similarity to several B. wiedmannii strains, including strain FSL W8-0169. This group is comprised by B. wiedmannii,

**FIG. 1.**—Genome map of FCC41 strain chromosome and plasmids. Rings from the outermost to the center: 1) scale marks, 2) protein-coding genes on the forward strand, 3) protein-coding genes on the reverse strand, 4) tRNA (black) and rRNA (red), 5) Illumina sequencing coverage, 6) PacBio sequencing coverage, 7) GC skew, 8) GC content. Protein-coding genes are color coded according to their COG categories. cry-like coding genes are indicated by red lines. The genome maps were drawn using the software Circos v0.69 (Krzywinski et al. 2009).
and some strains of B. cereus and B. thuringiensis; and most of its members are psychrotolerant and able to grow at 7 °C, since they carry one or more copies of the cold shock protein A gene (cspA). FCC41 strain has seven copies of cspA and the culture’s growth was detected at temperatures of 10°C, or higher. On the other hand, BTyper tool showed that rpoB allelic type (AT) of this strain has 100% identity to AT 410, which is associated to 12 B. cereus sensu lato strains deposited in the Food Microbe Tracker database (http://www.foodmicrobetracker.com; Last accessed April 2018.); additionally, there are 115 identical matches of the rpoB FCC41 sequence in NCBI nonredundant database, 94 of them corresponding to recently sequenced B. wiedmannii strains.

Moreover, different MLST schemes of housekeeping genes have been used in several works in order to make a greater effort in the clarification of B. cereus group phylogeny (Ko et al. 2004; Sorokin et al. 2006; Tourasse and Kolstø 2008). To compare FCC41 strain using a MLST approach, seven housekeeping genes from the B. cereus PubMLST typing scheme were used. Although the AT of the seven housekeeping genes already existed in the database, the combination of this seven AT represents a novel sequence type (ST). The resulting phylogenetic tree indicated that strain FCC41 is placed in the same group as other B. wiedmannii strains (see supplementary fig. S1, Supplementary Material online).

Additionally, the core genome phylogenetic analysis was performed with bcgTree which shows that strain FCC41 and the others B. wiedmannii strains are grouped together and close to Clade I in a similar way as described in Bazinet (2017; fig. 2).

ANI has been used to compare prokaryotic genome sequences and is recognized as one of the most robust measurements of relatedness between strains. An ANI value of 95% is accepted as the cutoff value for the delimitation of bacterial species. The ANI analysis value of 96.4% between FCC41 and B. wiedmannii FSL W8-0169 indicates that this strain belongs to the recently described species (fig. 3; see supplementary table S2, Supplementary Material online), and the results were highly related to the grouping obtained with bcgTree and previously reported phylogenetic trees (Bazinet 2017).

Virulence Factors and Antibiotic Resistance Genes
The virulence factors predicted by the BLASTP analysis of FCC41 strain reveals proteins that are widely distributed in B. cereus sensu lato such as phospholipases, proteases (inhA), enhancins, chitinases, among others, which have been suggested as virulence assistance factors, to aid B. thuringiensis in the invertebrate host and tissue invasion (Raymond et al. 2010; see supplementary table S3,
This strain carries coding genes of hemolysin BL, enterotoxin, cereolysin and other common chromosomal toxin genes mainly related with *B. cereus*. Additionally, FCC41 harbors two copies of the complete operon of the nonhemolytic enterotoxin, nheABC, one of them encoded in the chromosome and the other in plasmid pFCC41-5-125k, probably because of a horizontal gene transfer event. In particular, operon nheABC is abundant within *B. cereus* strains isolated from infections and from the environment, and it is usually found in the chromosome. It has been described that the duplication of nheABC is extremely infrequent and appears to be phylogenetically unstable (Bohm et al. 2015). The operon present in plasmid pFCC41-5-125k is highly divergent from the chromosome copy and does not have the PlcR recognition sequence, which is commonly found in nheABC operon and other virulence factors coding genes. Moreover, the enterotoxin C encoded by nhec has an additional C-terminal domain with homology to ricin domain.
Although this strain has pXO1-like and pXO2-like plasmids it does not carry any of the virulence factors associated with anthrax-like symptoms (protective antigen, and edema and lethal factors), nor capsule coding genes (capABCD), typical characteristics of B. anthracis.

Furthermore, we detected antibiotic resistance genes which may help this bacterium to survive and compete with other gut bacteria. Most of these virulence factors might contribute to insect pathogenicity (Raymond et al. 2010) of this strain and are shown in supplementary table S3, Supplementary Material online.

**Cry-like Protein Coding Genes**

We identified nine CDS with conserved Cry-like domains using hmmrsearch and further BLASTP analysis. The translated CDS were identified as two Cry4-like (both have 51% identity with Cry4Ca2, and 48% identity between them); two Cry52-like (82% identity with Cry52Aa1 and 63% identity with Cry52Ca1, respectively) with their corresponding downstream Orf2-like regions; the previously described Cry24Ca with its Orf2-like; and three contiguous pseudogenes within the same operon with Cry proteins domains homology. Cry4-like and Cry52-like proteins may contribute to the mosquitocidal activity since Cry4 and Cry52 proteins previously described have insecticidal activity against some species of *Aedes* sp., *Anopheles* sp., and *Culex* sp. (Ben-Dov 2014; Zheng et al. 2017). As well, a Cry41-like sequence with 50% identity to Cry41Ba2 and 42% identity to Cry41Aa1 was detected. In particular, this Cry41-like sequence contains a conserved ricin domain also detected in Cry41Aa. It has been previously reported that similar Cry proteins, called parasporins, cause cytotoxicity in vitro towards cancerous cells (Krishnan et al. 2017). All these sequences are encoded in three plasmids (fig. 1; table 2). In contrast with *B. thuringiensis* mosquitocidal strains like the subsp. *israelensis*, FCC41 does not have coding sequence genes for the previously described Cyt proteins (Soberón et al. 2013; Ben-Dov 2014).

**Plasmid Analysis**

Some of the most important *B. cereus* group phenotypic determinants are encoded in plasmids >100 kb. In most cases, these megaplasmids contain two or more different putative minireplicons, which are the minimum regions necessary to permit plasmid replication and comprise the origin of replication and the replication proteins coding genes (Zheng et al. 2013). In megaplasmids of the *B. cereus* group at least six types of minireplicons have been described, and the diversity and wide distribution are related with the evolution of those plasmids. FCC41 strain has eight plasmids, and five of them are megaplasmids (table 1) that harbor minireplicons previously found in *B. cereus* group (Zheng et al. 2013; Wang et al. 2018). In particular, plasmids pFCC41-1-490k and pFCC41-4-144k have a copy of a repX-like gene (see supplementary figs. S2 and S4, Supplementary Material online), defined as replication initiator proteins involved in the replication of megaplasmids. pFCC41-4-144k has additionally the pXO1-14/pXO-16-like proteins genes, initially described in *B. anthracis* pXO1 (Tinsley and Khan 2006), also present in other *B. cereus* group strains plasmids but not found in those with only one minireplicon (Zheng et al. 2013). It has been suggested that the presence of multiple minireplicons of different types could avoid the incompatibility between plasmids that possess the same minireplicon (Hüller et al. 2017). The lack of incompatibility between plasmids suggests that there are other replications proteins involved. Additionally, plasmid pFCC41-3-257k has a repA-like gene, and plasmids pFCC41-2-313k and pFCC41-5-125k have also different variants of the replication associated protein gene *orf44*-like (RepFR55), as shown in supplementary figures S3 and S5, Supplementary Material online.

The presence of two or more minireplicons in *B. cereus* group megaplasmids suggests that some of them may have resulted from the integration of two or more smaller plasmids (Zheng et al. 2013). pFCC41-3-257k shows a clear confirmation of such an integration event: About one fourth of the plasmid sequence has similarity with plasmids pMC183 from

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**Table 2**

Homologous Sequences to cry Genes Present in FCC41 Strain

| Gene ID | Location | From | To  | Length (bp) | Closest Cry Protein Homology | Identity (%) | Coverage (%) |
|---------|----------|------|-----|-------------|-----------------------------|--------------|--------------|
| cry41-like | Plasmid 2 | 75,380 | 77,956 | 2,577 | Cry41Ba2 (ZP_04099652) | 50 | 99 |
| cry4-like1 | Plasmid 3 | 72,700 | 76,215 | 3,516 | Cry4Ca2 (AJW76682) | 51 | 99 |
| cry24Ca | Plasmid 3 | 1,13,304 | 1,15,364 | 2076 | Cry24Ca1 (CAJ43600) | 99 | 95 |
| cry24Ca orf2 | Plasmid 3 | 1,11,767 | 1,13,236 | 1,470 | Cry39Aa1 orf2 (BAB782017) | 85 | 99 |
| cry52-like1 | Plasmid 3 | 1,13,768 | 1,18,751 | 2094 | Cry52Aa1 (ABU96490) | 82 | 100 |
| cry52-like1 orf2-like | Plasmid 3 | 1,76,911 | 1,78,590 | 1,680 | Cry39Aa1 orf2 (BAB782017) | 87 | 100 |
| cry52-like2 | Plasmid 3 | 1,43,701 | 1,45,767 | 2067 | Cry52Aa1 (AJW76683) | 63 | 100 |
| cry52-like2 orf2-like | Plasmid 3 | 1,45,835 | 1,47,508 | 1,674 | Cry39Aa1 orf2 (BAB782017) | 82 | 99 |
| cry4-like2 | Plasmid 5 | 77,853 | 81,524 | 3,572 | Cry4Ca2 (AJW76682) | 51 | 99 |
**B. thuringiensis** MC28 and pXO2 from *B. anthracis*, and the remainder of the plasmid shares regions of partial homology with plasmids pAM65-52-4-128k from *B. thuringiensis israelensis* AM65-52, pMC189 from *B. thuringiensis* MC28, and pH518-2 from *B. thuringiensis* HS18-1. However, we only detected one minireplicon in this plasmid. The regions of
homology shift between related plasmids correspond with regions of GC skew sign change, which is known to be associated to DNA replication origins or terminus (Tillier and Collins 2000).

Plasmid pFCC41-1-490k has regions of shared homology with plasmids pMC429 and pH518-1 from mosquitoicidal B. thuringiensis MC28 and HS18 strains, respectively, and pAR460 from plant pathogen control agent and growth promoter B. cereus AR156 strain; and has homology with contigs 14 and 17 from B. wiedmannii strain AFS023094 (see supplementary fig. S2, Supplementary Material online). Plasmid pFCC41-2-313k is very similar to the previously described B. thuringiensis subsp. israelensis strain AM65-52 and to contigs from strains ATCC35646, IBL4222 and IBL4200 of B. thuringiensis subsp. israelensis. This plasmid contains metabolite transporter genes, FtsK-like DNA transport protein, and type IV secretion system protein VirDIV. It has been previously mentioned that this type of plasmids appeared to be restricted to B. thuringiensis species (Bolotin et al. 2017). However, in recent times there has been an increasing number of published genomes of B. cereus and B. wiedmannii strains that have plasmids or contigs homologous to pFCC41-2-313k (see supplementary fig. S3, Supplementary Material online). Additionally, plasmid pFCC41-3-257k lacks the region of pXO2 that has the coding sequences for B. anthracis capsid proteins. This plasmid has the highest density of transposons and contains most of the Cry protein coding genes of this strain, which are all flanked by transposase coding genes (fig. 4). Nucleotide BLAST results indicate that plasmid pFCC41-4-144K shows high homology with pXO1 from B. anthracis; however, it lacks the coding genes of anthrax virulence factors (see supplementary fig. S4, Supplementary Material online). Plasmid pFCC41-5-125K has homology regions with plasmids from Bacillus mycoides and multiple strains of B. thuringiensis (see supplementary fig. S5, Supplementary Material online). Smaller plasmids, which range in size from 8 to 12 kb, have conserved regions present in plasmids from other strains of B. cereus group (see supplementary figs. S6–S8, Supplementary Material online).

Horizontal gene transfer is a driving force in bacterial evolution, in which genetic exchange occurs, mainly caused by transposons, phages or integrons, and can take place between highly divergent strains (Zheng et al. 2015). FCC41 chromosome has high similarity to B. wiedmannii strains sequences; however, its plasmids appear to be from different origins. The presence of cry genes, a high number of transposons particularly in highly divergent plasmids (fig. 4A) and large plasmid regions similar to plasmid sequences present in other Bacillus species suggest strong evidence of horizontal gene transfer events, considering that the involved species belong to different clades (Bazinet 2017). Recent studies in comparative genomics suggest that these lateral processes result in phylogenetic networks that explain more adequately the complexity of the evolutionary route (Zheng et al. 2015). Guinebretière et al. (2008) discussed that the global evolution of the B. cereus group seems to be strongly determined by ecological adaptations, being the temperature tolerance an important mechanism by which strains of the different groups adapted to novel environments and by which these groups have diverged. Most recently, Méric et al. (2018) reinforce this hypothesis proposing that the strains of different clades which share an ecological niche had increased opportunities for plasmid transfer between them. Most of the plasmids from strain FCC41 share homologous regions with plasmids of distant lineages from other clades, such as specialized insect pathogens from clade 2, which could indicate recent horizontal transfer. Strain FCC41 must have shared the ecological niche with other members of the B. cereus group. Our results support the hypothesis of cooperative infection, which predicts that conjugal plasmids carrying social genes such as Cry toxins could be widely distributed across clades (Méric et al. 2018).

The classification criteria for the B. cereus group division are widely discussed, the main problem has been to decide between evolutionary criteria or the use of phenotypical traits and traits of clinical and biotechnological importance, which are not clearly clustered in current phylogenetic trees (Raymond and Federici 2017; Zheng et al. 2017). The complete genome of FCC41 enriches the information about the genomic diversity and evolution among this group and proposes new questions about the position of B. thuringiensis strains that apparently belong to different lineages of evolution.

Conclusion

FCC41 was previously described as B. thuringiensis according to the phenotypic traits and the characterization of at least one Cry-toxic protein (Berón and Salerno 2007). This strain has mosquitoicidal activity against Ae. aegypti, Ae. (Ochlerotatus) albifasciatus, Culex pipiens, Culex quinquefasciatus, and Culex apicinus, showing higher activity against Ae. aegypti than the B. thuringiensis subsp. israelensis reference strain. In this work we present the complete genome of FCC41 strain which includes one circular chromosome and eight circular plasmids. The genome-wide screening of this strain reveals new putative cry genes, its regulatory regions and some virulence factors potentially useful for the development of biotechnological applications for the control of vector-borne diseases important in human health. According with the bioinformatics analysis this strain is placed into the B. wiedmannii cluster, but due to the phenotypical characteristics it could be renamed as B. wiedmannii Biovar thuringiensis strain FCC41, being the first reported B. wiedmannii specialized mosquitoicidal pathogen strain that has Cry
proteins with insecticidal activity, a phenotypic characteristic that is commonly associated with \textit{B. thuringiensis}.

**Supplementary Material**

Supplementary data are available at Genome Biology and Evolution online.

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