Comparative genome analysis of *Spiroplasma melliferum* IPMB4A, a honeybee-associated bacterium

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**Abstract**

**Background:** The genus *Spiroplasma* contains a group of helical, motile, and wall-less bacteria in the class Mollicutes. Similar to other members of this class, such as the animal-pathogenic *Mycoplasma* and the plant-pathogenic *Candidatus Phytoplasma*, all characterized *Spiroplasma* species were found to be associated with eukaryotic hosts. While most of the *Spiroplasma* species appeared to be harmless commensals of insects, a small number of species have evolved pathogenicity toward various arthropods and plants. In this study, we isolated a novel strain of honeybee-associated *S. melliferum* and investigated its genetic composition and evolutionary history by whole-genome shotgun sequencing and comparative analysis with other Mollicutes genomes.

**Results:** The whole-genome shotgun sequencing of *S. melliferum* IPMB4A produced a draft assembly that was ~1.1 Mb in size and covered ~80% of the chromosome. Similar to other *Spiroplasma* genomes that have been studied to date, we found that this genome contains abundant repetitive sequences that originated from plectrovirus insertions. These phage fragments represented a major obstacle in obtaining a complete genome sequence of *Spiroplasma* with the current sequencing technology. Comparative analysis of *S. melliferum* IPMB4A with other *Spiroplasma* genomes revealed that these phages may have facilitated extensive genome rearrangements in these bacteria and contributed to horizontal gene transfers that led to species-specific adaptation to different eukaryotic hosts. In addition, comparison of gene content with other Mollicutes suggested that the common ancestor of the SEM (*Spiroplasma*, *Entomoplasma*, and *Mycoplasma*) clade may have had a relatively large genome and flexible metabolic capacity; the extremely reduced genomes of present day *Mycoplasma* and *Candidatus Phytoplasma* species are likely to be the result of independent gene losses in these lineages.

**Conclusions:** The findings in this study highlighted the significance of phage insertions and horizontal gene transfer in the evolution of bacterial genomes and acquisition of pathogenicity. Furthermore, the inclusion of *Spiroplasma* in comparative analysis has improved our understanding of genome evolution in Mollicutes. Future improvements in the taxon sampling of available genome sequences in this group are required to provide further insights into the evolution of these important pathogens of humans, animals, and plants.

**Background**

The bacterial genus *Spiroplasma* contains a group of arthropod- and plant-associated endosymbionts [1-5]. Phylogenetically, the genus *Spiroplasma* belongs to the class Mollicutes, which are close relatives of Bacilli and other free-living Firmicutes [2,6,7]. Within Mollicutes, *Spiroplasma* are more closely related to the animal-pathogenic genus *Mycoplasma* than to the plant-pathogenic *Candidatus Phytoplasma*. However, *Spiroplasma* are more similar to *Candidatus Phytoplasma* in terms of the ecological niches occupied, because both groups have complex life cycles that involve insect and plant hosts [4]. For this reason, comparative analysis of gene content among these three groups of Mollicutes can provide insights into host adaptation in these parasites [8]. While most of the *Spiroplasma* species that have been characterized to date appeared to be harmless commensals,
some species have evolved pathogenicity toward their plant or arthropod hosts. Most of the pathogenic Spiroplasma species belong to the Citri-Chrysopicola-Mirum clade [3], notable examples include *S. citri* that causes the citrus stubborn disease [9], *S. kunkelii* that causes the corn stunt disease [10], *S. phoeniceum* that infects periwinkle [11], *S. penaei* that infects Pacific white shrimp [12,13], *S. eriocheiris* that infects Chinese mitten crab [14], and *S. melliferum* that infects honeybee [15].

To improve our understanding of these pathogenic Spiroplasma species, whole genome shotgun sequencing has been utilized as a powerful tool to investigate their metabolic capacity and possible interactions with hosts. The initial surveys of *S. kunkelii* genome revealed that this species has more genes involved in purine and amino acid biosynthesis, transcriptional regulation, cell envelope, and DNA transport than lineages from Mycoplasmataceae [16] and also a unique arrangement of genes in its ribosomal protein operon [17]. Furthermore, a comparative study within Mollicutes has identified several proteins that are shared between the phytopathogenic *S. kunkelii* and phytoplasma but absent in the animal-pathogenic *Mycoplasma* and *Ureaplasma* spp [8]. More recently, draft genome sequences have been published for *S. citri* GI13-3X [18] and *S. melliferum* KC3 [19]. These more comprehensive investigations of Spiroplasma genomes revealed that their chromosomes harbor a large amount of viral sequences and also allowed for detailed characterization of their metabolic capacities. Importantly, these draft genome sequences provided a valuable resource for identification of putative virulence factors and elucidation of possible pathogenicity mechanisms [18,19].

In this study, we isolated a novel strain of *S. melliferum* from a honeybee collected in Taiwan and investigated its genetic composition by whole-genome shotgun sequencing. Through comparative analyses with other available Mollicutes genome sequences, our aims were to characterize the genetic differences among the various lineages and to understand the evolutionary processes involved in shaping the genomes of these bacteria.

Results and discussion

Species identification and phylogenetic inference

Based on the molecular phylogeny inferred from the 16S ribosomal RNA gene, the two strains of *S. melliferum* (IPMB4A and KC3) form a monophyletic group and *S. citri* is the most closely related species (Figure 1).

![Figure 1 Phylogenetic placement of Spiroplasma melliferum IPMB4A](image)

The maximum likelihood phylogenetic tree was inferred using the 16S ribosomal RNA gene, GenBank accession numbers were listed in square brackets following the species names. The numbers on the internal branches indicated the percentage of bootstrap support based on 1,000 re-samplings.
Serological tests were confirmatory for the molecular phylogeny and supported the close association of *S. melliferum* IPMB4A with other *S. melliferum* strains. The deformation test showed a strong cross-reactivity between *S. melliferum* IPMB4A and the type strain BC-3$^T$ [15]. Positive deformation was observed up to 5,120-fold dilution. Against its own antisera, *S. melliferum* BC-3$^T$ showed positive deformation up to 10,240-fold dilution, but only up to 320-fold dilution with the closely related *S. citri* R8A2$^T$ [9]. This indicates that *S. melliferum* IPMB4A is a strong candidate for inclusion into *Spiroplasma* group I-2 with the other strains of *S. melliferum*.

**Genome assembly and annotation**

The whole genome shotgun sequencing of *S. melliferum* IPMB4A produced an assembly that contains 24 chromosomal contigs (Table 1). Consistent with the previous studies of *Spiroplasma* genomes [18,19], we found a large number of phage sequences in the chromosome of *S. melliferum* IPMB4A. Most of these phage sequences showed high similarity to plectrovirus SpV1-C74, SpV1-R8A2B, and SVTS2. The presence of these long repetitive sequences was the major obstacle in obtaining a complete genome assembly. Moreover, the divergence among different copies of these repetitive plectrovirus fragments generated many single-nucleotide polymorphisms (SNPs).

### Table 1 Genome assembly statistics.

|                  | *S. melliferum* IPMB4A | *S. melliferum* KC3 | *S. citri* GII3-3X |
|------------------|------------------------|--------------------|-------------------|
| Number of chromosomal contigs | 24                     | 4                  | 39                |
| Combined size of chromosomal contigs (bp) | 1,098,846              | 1,260,174          | 1,525,756         |
| Estimated chromosomal size (bp)       | 1,380,000              | 1,430,000          | 1,820,000         |
| Estimated coverage (%)                | 79.6                   | 88.1               | 83.8              |
| G+C content (%)                      | 27.5                   | 27.0               | 25.9              |
| Coding density (%)                    | 85.1                   | 83.0               | 80.2              |
| Protein-coding genes                  | 932                    | 1,222              | 1,905             |
| Length distribution (Q1/Q2/Q3) (a.a) | 176/280/440            | 119/233/376        | 83/149/286        |
| Plectrovirus proteins                 | 11                     | 132                | 375               |
| Hypothetical proteins                 | 337                    | 485                | 519               |
| Annotated pseudogenes$^1$             | 12                     | 12                 | 401               |
| rRNA operon                           | 1                      | 1                  | 1                 |
| tRNA                                  | 32                     | 31                 | 32                |
| Number of plasmids                    | 0                      | 4                  | 7                 |

For *S. melliferum* IPMB4A, putative pseudogenes were annotated with the 'pseudo' tag in gene feature as suggested by the NCBI GenBank guidelines and were not counted in the total number of protein-coding genes. For *S. melliferum* KC3 and *S. citri* GII3-3X, putative pseudogenes were annotated by adding the term ‘truncated’ in the CDS product description field and were included in the total number of protein-coding genes.

Most of these SNPs cannot be resolved confidently with the short 101-bp Illumina sequencing reads used in this study. To be conservative, ~135 kb of the plectrovirus-related regions that contained these SNPs were excluded from our final assembly. If these phage fragments were included, the assembly coverage would be similar to *S. melliferum* KC3 and higher than *S. citri* GII3-3X (Table 1). This result highlights the limitation of relying on short-read sequencing technologies for de novo assembly of highly repetitive genomes. However, the traditional approach of using clone libraries for Sanger shotgun sequencing has a much higher cost and may not be a good choice if the main focus is on the unique regions of the genome.

The exclusion of phage fragments reduced the number of annotated plectrovirus proteins in the *S. melliferum* IPMB4A assembly compared to the closely related *S. melliferum* KC3 (Table 1). With the elimination of these short plectrovirus proteins and other potential false-positive gene predictions (see Methods) from the annotation, we observed a shift in the length distribution of the annotated protein-coding genes compared to the other two *Spiroplasma* genomes (Table 1). Nonetheless, our conservative approach in assembly and annotation mainly removed viral proteins and short hypothetical proteins, thus was not expected to introduce biases in the downstream metabolic analysis. Intriguingly, in contrast to the abundance of plasmids in *S. melliferum* KC3 [19] and *S. citri* GII3-3X [18,20], we did not recover any plasmid-like contig in our draft assembly of *S. melliferum* IPMB4A genome (Table 1). This lack of plasmid-like contig is consistent with the negative result from our plasmid extraction tests. This finding may reflect the true absence of plasmids in this particular strain. Alternatively, it is possible that the plasmid(s) had been lost during the process of initial isolation or subsequent culturing in the laboratory.

Among the 932 annotated protein-coding genes in the *S. melliferum* IPMB4A genome, 392 have specific functional assignments according to the COG categories (Figure 2). Genes involved in translation and ribosomal structure (COG category J) represented the most abundant functional category, mostly due to the large number of ribosomal proteins and tRNA synthetases found in the genome. The second most abundant category contained genes involved in replication, recombination, and repair (COG category L). In addition to the DNA replication machinery such as DNA polymerases (e.g., dnaE, dnaN, dnaX, hollA, hollB, and polC) and other related proteins (e.g., dnaA, dnaB, dnaG, ligA, polA, ssbA, etc.), we also found genes involved in nucleotide excision repair (e.g., uvrA, uvrB, and uvrC) and several DNA topoisomerases (e.g., gyrA, gyrB, parC, and parE). However, the machineries for mismatch repair (e.g., mutS, mutL,
mutH, exol, exoX, recJ, etc.) and homologous recombination (e.g., recA, recB, recC, etc.) appeared to be missing.

**Comparative analysis with S. melliferum KC3 and S. citri**

The alignment among the three draft Spiroplasma genomes revealed extensive rearrangements between the closely related S. melliferum and S. citri (Figure 3A). This low level of conservation in synteny was unexpected for several reasons. First, the average nucleotide sequence identity between these two species was ~99%, suggesting that the divergence time was quite short [21]. For comparison, two other groups of related bacteria with similar (i.e., between-strain comparison in Mycoplasma hyopneumoniae; see Figure 3B) or even higher (i.e., between-species comparison in Bacillus; see Figure 3C) levels of divergence both had a higher level of conservation in their chromosomal organization. Second, in a previous study that compared the physical maps of S. melliferum and S. citri, the authors found that the overall organization was similar between these two species [22]. However, this discrepancy may be due to the higher resolution in detecting genome rearrangements provided by genome sequencing compared to pulsed-field gel electrophoresis (PFGE). Finally, these two species lack several genes that were involved in homologous recombination. For example, the recombinase A (encoded by recA) was reduced to 231 a.a. in S. melliferum by a premature stop codon and 130 a.a. in S. citri by one or more truncations of its coding region [18,23]; for comparison, the full length RecA protein in Mycoplasma mycoides contains 345 a.a. (GenBank accession number NP_975407). Although these Spiroplasma genomes contain a large number of direct and inverted repeats originated from the invasion of plecroviruses, the pseudogenization of recA was expected to promote genome stability [24,25]. One possible explanation of this paradox is that these extensive genome rearrangements were facilitated by the large number of repetitive sequences [26] and had occurred soon after the species divergence, while the independent losses of homologous recombination in these two species were relatively recent events. More extensive taxon sampling of Spiroplasma genomes, including additional strains from these two species and appropriate outgroup species, are required to investigate this hypothesis.

To compare the gene content between S. melliferum and S. citri, we identified the homologous genes among the three available genomes from these two species and classified these genes based on their patterns of presence and absence. The functional categorization of each protein-coding gene was classified according to the COG assignments, genes that did not have any inferred COG annotation were assigned to a custom category X. The number of protein-coding genes in each set was labeled in the center of each pie chart. (A) All 932 annotated protein-coding genes in the S. melliferum IPMB4A genome. (B) The 392 protein-coding genes that have specific functional category assignments.
Figure 3. Extensive level of genome rearrangement between closely related Spiroplasma spp. The color blocks represent regions of homologous backbone sequences without rearrangement among the genomes compared. The vertical red bars indicate the boundaries of individual contigs. The average nucleotide sequence identities were calculated based on single-copy genes that were conserved among the three genomes compared in each group. (A) Comparison among Spiroplasma melliferum IPMB4A, S. melliferum KC3, and S. citri GI3-3X. (B) Comparison among Mycoplasma hyopneumoniae strains 7448, J, and 232. (C) Comparison among Bacillus anthracis str. Ames, B. cereus ATCC 10987, and B. thuringiensis serovar konkukian str. 97-27.
on the patterns of presence and absence. When the ongoing genome sequencing of *S. kunkelii* [8,16,17] reaches completion in the future, combined analysis using a phylogenetic framework will provide more insights into the gene content evolution of these pathogenic *Spiroplasma* species.

Despite these potential shortcomings, inspection of the gene lists suggested that this comparison of gene content provided results that were consistent with our expectations. We found that both *S. melliferum* KC3 and *S. citri* GII3-3X contained large numbers of genome-specific genes and also shared 85 homologous gene clusters that were absent in *S. melliferum* IPMB4A. The majority of these genes encoded hypothetical proteins and proteins of viral origin. These observations were expected due to our exclusion of most plectrovirus fragments from the assembly of *S. melliferum* IPMB4A genome (see above). Furthermore, examination of these gene lists provided insights into the biology of these bacteria. Both species have phosphotransferase system (PTS) transporters and enzymes for the uptake and utilization of trehalose, glucose, and fructose (i.e., *treB*, *ptsG*, *fruA*, and *crr*; see Figure 5A and Additional file 1). These findings were consistent with our current knowledge of spiroplasma biology. For example, trehalose exists in higher concentration than glucose or fructose in insect haemolymph [27] and may be an important carbon source for spiroplasmas during their life stage in insect vectors. In addition, both species were known to utilize glucose and fructose [9,15] and the preferential uptake of fructose by *S. citri* is linked to its phytopathogenicity [28,29].

Intriguingly, compared to the plant-pathogenic *S. citri*, the insect-pathogenic *S. melliferum* has additional genes involved in the utilization of cellulose, chitin, and *N*-acetylmuramic acid (MurNAc) (i.e., *celA*, *celB*, *chia*, *nagE*, and *murP*; see Figure 5B and Additional file 1). The presence of these genes may be linked to its adaptation to the honeybee host. For example, the ability to degrade chitin can facilitate its invasion of the host tissues (e.g., through the chitinous peritrophic matrix, gut epithelium, or exoskeleton, etc.) and the ability to utilize cellulose may enhance its fitness by allowing access to the partially hydrolyzed pollen cell walls in honeybee diet for additional carbon sources. More interestingly, several of these *S. melliferum*-specific genes (e.g., *chia*, *celA*, and *celB*) were adjacent to plectrovirus fragments on the chromosome, suggesting that these genes may have been acquired through phage-mediated horizontal gene transfers. The sequence similarity search against the GenBank database revealed that the chitinase in *S. melliferum* has no putative homolog within the class Mollicutes and the most similar genes were found in the order Lactobacillales, including the genera *Lactococcus* and *Enterococcus*.

**Comparative analysis with mycoplasma and phytoplasma**

For comparative genomics analysis at a deeper divergence level (Figure 1), we compared the gene content of
S. melliferum IPMB4A with two other representative Mollicutes species: Mycoplasma genitalium G37 (GenBank accession number NC_000908) and ‘Candidatus Phytoplasma asteris’ OY-M (NC_005303). The inclusion of the M. genitalium, which has the smallest genome among sequenced Mollicutes, is useful for defining the most conserved core gene set in these bacteria. Among these three genomes, S. melliferum IPMB4A is the largest one and contains the highest number of genome-specific genes (Figure 4B and Additional file 2). Examination of these genes revealed that Spiroplasma possesses more flexible metabolisms and consequently a lower degree of host dependence. In the initial isolation and characterization, S. melliferum was reported to ferment glucose and to hydrolyze arginine [15]. Our pathway analysis revealed that these two features may be important in its physiology. In addition to the PTS transporter for glucose uptake, S. melliferum possesses all genes required for glycolysis to convert glucose-6-phosphate into pyruvate (Figure 5 and Figure 6). Pyruvate may be used for the production of cysteine by an aminotransferase (patB) and the biosynthesis of Fe-S clusters by other downstream genes.
(i.e., sufS, sufU, sufB, sufC, and sufD; see Figure 6). Alternatively, pyruvate may also be converted into acetyl-CoA by the pyruvate dehydrogenase complex (i.e., pdhA, pdhB, and pdhC). Other than glycolysis, S. melliferum also has the genes in the pentose phosphate pathway (e.g., pgk, tkt, rpe, rpiB, and prs) for producing 5-phosphoribosyl diphosphate (PRPP), which is a precursor for pyrimidine metabolism (Figure 6).

The ability to uptake and metabolize arginine provides S. melliferum with a source of nitrogen and also precursors for de novo nucleotide biosynthesis. The possession of the entire gene set (i.e., arcA, arcB, arcC, pyrC, pyrD, pyrE, and pyrF; see Figure 6) for converting arginine to uridine monophosphate (UMP) is rare among Mollicutes. Other than S. melliferum, we only found that Mycoplasma penetrans [30] contains all these genes as well. However, result from database searches demonstrated sporadic distribution patterns of individual genes among different Mycoplasma species (e.g., some of these genes were found in M. capricolum, M. hominis, M. putrefaciens, etc.). In addition to the de novo biosynthesis of UMP, S. melliferum may acquire xanthine and uracil by permeases (e.g., pbuG and pyrP). These nucleobase transporters were also rare among Mollicutes. Other than S. melliferum, these permeases were only found in S. citri and Mesoplasma florum. Although these findings may indicate independent gene acquisitions by horizontal gene transfers, a more plausible explanation is that the common ancestor of the SEM (Spiroplasma, Entomoplasma, and Mycoplasma) clade may have maintained these genes for nucleotide transport and metabolism following its divergence from the AAA (Asteroeplasma, Anaeroplasma, and Acholeplasma; including phytoplasmas) clade. Subsequently, the nutrient-rich environments provided by the eukaryotic hosts reduced the selective pressure for maintaining these genes in the genome and allowed parallel gene losses in most SEM lineages. In addition to the genes discussed above, we also found several Spiroplasma-specific genes that are associated with its unique biology among Mollicutes. For example, S. melliferum IPMB4A has five copies of mreB (encoding for the cell shape determining protein MreB) and two copies of spiralin (which are likely to be involved in its interactions with the insect host [31]). Furthermore, S. melliferum IPMB4A has several genes involved in lipid metabolism, such as the non-mevalonate pathway for terpenoid backbone biosynthesis (i.e., dxs, dcr, ispd, ispf, ispg, and ispH; see Additional file 2).

Among the three genomes included in this comparative analysis, M. genitalium contained the fewest number of protein-coding genes (i.e., 475 compared to 749 in ‘Candidatus Phytoplasma asteris’ and 932 in S. melliferum). Given these numbers, one might expect that S. melliferum would shared more genes with ‘Candidatus Phytoplasma asteris’ than with M. genitalium. However, the homologous gene cluster identification showed that S. melliferum and M. genitalium actually shared more genes to the exclusion of ‘Candidatus Phytoplasma asteris’ (Figure 4B and Additional file 2). This finding is consistent with the close evolutionary relationship between Spiroplasma and Mycoplasma. Many of the genes in this list represented gene losses in the phytoplasma lineage. For example, the absence of F0F1-type ATP synthase (e.g., atpA, atpB, atpD, atpF, atpE, atpG, and atpH) and pentose phosphate pathway (e.g., tkt, rpiB, prs, deoC) were noted in the initial genome sequencing of phytoplasmas [32,33]. Furthermore, the phytoplasma-specific losses of purine salvage pathway (apt and hprT), pyrimidine metabolism (trxB), formylation of methionyl-tRNA (mnt and folD), and potassium ion uptake (ktrA and ktrB) were noted in our previous large-scale comparative analysis between mycoplasmas and phytoplasmas [34].

According to the phylogenetic relationship among these three lineages, the genes that are shared between S. melliferum and ‘Candidatus Phytoplasma asteris’ but absent in the M. genitalium genome would be of particular interest (Additional file 2). Although some of these genes may reflect accelerated sequence divergence or gene losses in M. genitalium, others may provide insights into the adaptation of spiroplasmas and phytoplasmas to their similar ecological niches. Examination of the 30 homologous gene clusters identified in this study mostly confirmed the results of a previous study that investigated this topic [8] and also provide additional candidate genes for future characterization.

In the initial survey of an 85-kb segment of the S. kunkelii CR2-3X genome, it was reported to have a unique arrangement of ribosomal protein genes [17]. Compared to M. capricolum, Bacillus subtilis, and Escherichia coli, the intergenic regions are much shorter in S. kunkelii. When we compared this region across the available Mollicutes genomes, we found that the compactness of this operon is not necessarily a defining feature of Spiroplasma genomes. For example, the highly reduced M. genitalium genome has similar or even shorter intergenic regions between neighboring genes. However, we noted that the three Spiroplasma species examined (i.e., S. kunkelii, S. citri, and S. melliferum) all have the same set of three genes (i.e., secY, adk, and infA) located downstream of ribosomal protein L15 (rplO). In contrast, several phytoplasma and Mycoplasma species, including M. mycoides that is nested within the Spiroplasma clade (Figure 1), all have an additional methionine aminopeptidase (map) located between adk and infA. As the taxon sampling of available Spiroplasma genomes continues to improve, it will be interesting to investigate if this relocation of the map gene is specific to the Citri clade or shared by other Spiroplasma lineages.
Conclusions
The genome survey by shotgun sequencing revealed that several *Spiroplasma* species harbor abundant plectrovirus fragments in their genome. Because of the repetitive nature of these viral sequences and the limitations of current sequencing technologies, obtaining the complete genome sequence of these bacteria has remained extremely challenging. Nonetheless, our analysis of the partial genome sequences of *S. melliferum* and *S. citri* provided glimpses into the genome architecture and metabolic capacity of these bacteria. Furthermore, comparative analysis with other Mollicutes demonstrated that the availability of *Spiroplasma* genome sequences is of fundamental importance in improving our knowledge of the evolutionary history in this class. Through comparison with the animal-pathogenic *Mycoplasma* and the plant-pathogenic ‘*Candidatus Phytoplasma,*’ results from this study suggested that the extensive gene losses observed in these bacteria may have occurred independently among different lineages, rather than mainly in their common ancestor as previously thought [34]. For future study, additional taxon sampling of *Spiroplasma* genomes would further improve the reconstruction of ancestral states and the inference of evolutionary history in these important pathogenic bacteria.

Methods
Strain isolation and DNA preparation
The bacterial strain *S. melliferum* IPMB4A was isolated from a honeybee collected at the main campus of the National Taiwan University (Taipei, Taiwan) on April 13, 2011 following the protocol described in a previous study [35]. Briefly, each honeybee was homogenized in 2 ml of the R2 medium [36] using a ceramic mortar and pestle set. The homogenized samples were allowed to sit for 20 minutes before passing through a 0.45 um filter (Millipore, USA) by gravity without added pressure. The filtered medium was diluted 20X and incubated at 31°C without shaking. The culture tubes were checked daily for color change, samples that turned yellow were checked using dark field microscopy to confirm the presence of helical-shaped spiroplasma cells and triply cloned by colony isolation on 1.5% agar plates. The pathogenicity of the IPMB4A strain was unknown. For PCR and whole-genome shotgun Illumina sequencing (see below), we isolated the genomic DNA of *S. melliferum* IPMB4A using the Wizard Genomic DNA Purification Kit (Promega, USA) according to the manufacturer’s protocol.

Pulsed-field gel electrophoresis
To determine the genome size of *S. melliferum* IPMB4A, we performed pulsed-field gel electrophoresis (PFGE) based on established protocols [37,38]. Briefly, 40 ml of saturated cell culture was obtained by incubation at 30°C without shaking for 72 hours. Cells were collected by centrifugation at 12,000 xg for 35 minutes. The resulting pellet was gently re-suspended in 400 ul of Tris buffer (20 mM Tris/Cl, 1% sucrose, 0.05 M EDTA, pH 8.0) and warmed to 30°C before mixing with an equal volume of pre-melted 2% low melting temperature agarose (Lonza, Switzerland). The mixture was quickly transferred into a cold gel plug mold and allowed to solidify at 4°C for 30 minutes. The gel plugs were incubated in lysis buffer (0.5 M EDTA, 1% N-lauroylsarcosine, pH 8.0, 1 mg/ml proteinase K added freshly) at 55°C for three days with three buffer changes daily. The PFGE step was carried out using a CHEF Mapper XA System (Bio-Rad, USA) under the following conditions: 24-hour run, 120° included angle, 24.03–120 sec switch-time ramp, 6 V/cm, 0.5X TBE (0.045 M Tris/borate, 1 mM EDTA), 14°C, 1.0% pulsed-field certified agarose (Bio-Rad, USA). The *Saccharomyces cerevisiae* chromosomal DNA (Bio-Rad, USA) was used as the size standards to estimate the molecular weight of the *S. melliferum* IPMB4A chromosome.

Molecular phylogenetic inference
To perform molecular phylogenetic inference, the partial 16S ribosomal RNA gene was amplified by using the primer pair 8F (5’-agagtttgatctcggctcag-3’) and 1492R (5’-ggtacccttgaactctag-3’). Each 50 ul reaction mix contained 1 ul of PfuUltra Fusion HotStart DNA polymerase (Agilent, USA), 5 ul of the supplied 10X buffer, 2 mM MgCl₂, 0.2 uM of each primer, 250 uM of each dNTP, and 100 ng of DNA template. The cycling conditions were: (1) an initial activation step at 95°C for 3 min; (2) 25 cycles of 95°C for 40 sec, 55°C for 40 sec, and 72°C for 40 sec; and (3) a final extension step for 5 min. The PCR product of expected size was isolated using agarose gel electrophoresis and purified using QIAquick Gel Extraction Kit (Qiagen, Germany) according to the manufacturer’s protocol. The purified PCR product was cloned using the Zero Blunt TOPO PCR Cloning Kit (Invitrogen, USA) and sequenced using the BigDye Terminator v3.1 Cycle Sequencing Kit on an ABI Prism 3700 Genetic Analyzer (Applied Biosystems, USA). The consensus sequence from 11 individual clones (1,442-bp, excluding the PCR primers) was used as the representative sequence and deposited in NCBI GenBank under the accession number JQ347516.

This partial 16S rRNA gene sequence was used as the query to run BLASTN [39,40] against the NCBI nt db [41] to identify homologous sequences from related species. These sequences (see Figure 1 for accession numbers) were aligned using MUSCLE [42] with the default settings. The resulting multiple sequence alignment was used to infer the species phylogeny using the maximum likelihood program PhyML [43]. The transition/
transversion ratio, proportion of invariable sites, and the gamma distribution parameter (with four categories of substitution rates) were estimated from the alignment in the maximum likelihood framework. To estimate the level of support for each internal branch, we generated 1,000 non-parametric bootstrap samples of the alignment by using the SEQBOOT program in the PHYLIP package [44] and repeated the phylogenetic inference as described above.

Serology test
The deformation test was performed using *S. melliferum* IPMB4A against the hyperimmune antisera of *S. melliferum* BC-3T (ATCC 33219) as described previously [45]. Positive reactions were indicated by the visual appearance of grape-like clustering and clumping of the spiroplasma cells, indicative of a reaction between the cell membrane and the antisera. The appearance of normal helical-shaped spiroplasmas indicated no reaction to the antisera. A dilution is designated as positive when over 50% of the spiroplasma cells present show a positive reaction. The test was performed three times.

Whole-genome shotgun sequencing
To obtain the genome sequence of *S. melliferum* IPMB4A, we used a commercial sequencing provider (Yourgene Bioscience, Taipei, Taiwan) for whole-genome shotgun sequencing based on the Illumina HiSeq 2000 platform (Illumina, USA). Two separate libraries were prepared and sequenced, including one paired-end library (insert size = 365 bp, 79,924,629 read-pairs, approximately 16 Gb of raw data) and one mate-pair library (insert size = ~4.5 kb, 20,463,824 read-pairs, approximately 4 Gb of raw data).

Genome assembly and annotation
The paired-end Illumina reads were used for *de novo* genome assembly by the program Velvet-SC [46]. The raw reads were trimmed from the 5'-end at the first position that has a quality score of lower than 20, all reads that were shorter than 70 bp after this quality trimming step were discarded. This resulted in 14,129,347 un-paired and 80,693,442 paired reads. The k-mer, expected coverage, and coverage cutoff were set to 79, 4,000, and 100, respectively.

The 25 contigs that were longer than 1 kb after this initial assembly step were used as the starting point for our iterative assembly correction process. For each iteration, we mapped all raw reads (including paired-end and mate-pair reads) to the assembled contigs using BWA [47] and visualized the results using IGV [48]. Neighboring contigs with mate-pair support for continuity were merged into scaffolds and polymorphic sites were examined using the MPILEUP program in the SAMTOOLS package [49]. For regions that appear to have been mis-assembled (often associated with aberrant coverage patterns), we extracted the raw reads that were mapped to the corresponding regions and re-assemble these regions by PHRAP [50]. These re-assembled regions were incorporated into the assembly and verified in the next iteration. We repeated this iterative process until no improvement could be made with the method described.

At the final step of our assembly process, we removed regions that contained unresolved polymorphisms (all corresponded to repetitive plectrovirus fragments) and un-filled gaps from the scaffolds. The resulting contigs were processed using RNAmer [51], rRNAscan-SE [52], and PRODIGAL [53] for gene prediction. The gene name and description for the protein-coding genes were assigned based on the orthologous genes identified by OrthoMCL [54] in other Mollicutes genomes and BLASTP [39,40] searches against NCBI nr db [41]. For functional categorization, all protein sequences were annotated by utilizing the KAAS tool [55] provided by the KEGG database [56,57] using the bi-directional best hit method. In addition to the default Prokaryotes representative set, we included selected Tenericutes and Firmicutes in the reference genomes. The complete list of abbreviated genome ids selected includes: *lisa*, *dme*, *ath*, *sce*, *pfa*, *eco*, *sty*, *hin*, *pae*, *nme*, *hpy*, *rpr*, *mlo*, *bsu*, *sau*, *lla*, *spn*, *cac*, *nme*, *mtu*, *ctr*, *bbu*, *syn*, *aae*, *mja*, *afu*, *aph*, *ape*, *mrt*, *mny*, *mmo*, *maa*, *uur*, *poy*, *acl*, *mfl*, *bsu*, *lca*, and *cho*. The KEGG orthology assignments were further mapped to the COG functional category assignment [58,59] to generate summary statistics (Figure 2). Genes that did not have any COG functional category assignment were assigned to a custom category (category X: no COG assignment). Finally, the annotation was manually curated to incorporate evidence from each of the approaches described above and to ensure the consistency of gene descriptions.

To correct for possible over-annotation, predicted protein-coding genes that were shorter than 100 amino acids, had low confidence scores as inferred by PRODIGAL [53], and lacked any functional predictions or putative homologous sequences were eliminated from the final annotation. To correct for possible under-annotation, genes that were conserved between the closely related *S. melliferum* KC3 and *S. citri* GII3-3X genomes but were absent in our initial gene prediction were used as queries to search against the contigs.

This Whole Genome Shotgun project has been deposited at DDBJ/EMBL/GenBank under the accession AMGI000000000 (BioProject PRJNA80357). The version described in this paper is the first version, AMGI01000000.

Comparative analysis with other genomes
To examine the extent of genome rearrangement in *Spiroplasma* and other related bacteria, we used
MAUVE [60] to compare S. melliferum IPMB4A and the following genomes (see Figure 3): S. melliferum KC3 (GenBank accession numbers AGBZ01000001-AGBZ01000004), S. citri GII3-3X (AM285301-AM285339), Mycoplasma hypopneumoniae 7448 (NC_007332), M. hypopneumoniae J (NC_007295), M. hypopneumoniae 232 (NC_006360), Bacillus anthracis str. Ames (NC_003997), B. cereus ATCC 10987 (NC_003909), and B. thuringiensis serovar konkukian str. 97-27 (NC_005957). The average sequence identities between genomes were calculated by the DNASIST program in the PHYLIP package [44] using conserved single-copy genes identified by OrthoMCL [54] within each group.

To identify lists of shared and genome-specific genes at different levels of evolutionary divergence, we compared the gene content of S. melliferum IPMB4A with other representative Mollicutes genomes (Figure 4). Because of the varying levels of sequence divergence, the sequence similarity search step in the OrthoMCL [54] analysis was conducted at the nucleotide level using BLASTN [39,40] for comparison with other representative Mollicutes genomes and at the protein level using BLASTP [39,40] for comparison with mycoplasma and phytoplasma genomes.

Additional files

Additional file 1: Table S1. Lists of shared and unique homologous gene clusters among S. melliferum IPMB4A (Sml), S. melliferum KC3 (SmK), and S. citri Sc). The genes in each worksheet were sorted by (1) COG categories, (2) Genome ID, (3) gene name, and (4) gene description.

Additional file 2: Table S2. Lists of shared and unique homologous gene clusters among S. melliferum IPMB4A (Sml), Mycoplasma genitalium G37 (Mg), and Candidatus Phytoplasma asteris OY-M (PaO). The genes in each worksheet were sorted by (1) COG categories, (2) Genome ID, (3) gene name, and (4) gene description.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

WSL carried out the genome assembly, annotation, and comparative analysis. LLC handled the biological materials and carried out DNA extraction, cloning, and sequencing. WCC carried out the pulsed field gel electrophoresis and participated in genome assembly and annotation. GEG carried out the serology test and participated in the writing of the manuscript. CHK conceived of the study, isolated the bacterial strain, carried out the serology test and participated in the writing of the manuscript. WCC carried out the pulsed field gel analysis. LLC handled the biological materials and carried out DNA extraction, cloning, and sequencing. WCC carried out the pulsed field gel electrophoresis. Funding for this work was provided by research grants from Academia Sinica and the National Science Council of Taiwan (NSC 101-2621-B-001-004-MY3) to CHK. The funders had no role in the design of the study, the collection and analysis of the data, the writing of the manuscript, or the decision to submit the manuscript for publication.

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