Degenerative Minimalism in the Genome of a Psyllid Endosymbiont

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Psyllids, like aphids, feed on plant phloem sap and are obligately associated with prokaryotic endosymbionts acquired through vertical transmission from an ancestral infection. We have sequenced 37 kb of DNA of the genome of *Carsonella ruddii*, the endosymbiont of psyllids, and found that it has a number of unusual properties revealing a more extreme case of degeneration than was previously reported from studies of eubacterial genomes, including that of the aphid endosymbiont *Buchnera aphidicola*. Among the unusual properties are an exceptionally low guanine-plus-cytosine content (19.9%), almost complete absence of intergenic spaces, operon fusion, and lack of the usual promoter sequences upstream of 16S rDNA. These features suggest the synthesis of long mRNAs and translational coupling. The most extreme instances of base compositional bias occur in the genes encoding proteins that have less highly conserved amino acid sequences; the guanine-plus-cytosine content of some protein-coding sequences is as low as 10%. The shift in base composition has a large effect on proteins: in polypeptides of *C. ruddii*, half of the residues consist of five amino acids with codons low in guanine plus cytosine. Furthermore, the proteins of *C. ruddii* are reduced in size, with an average of about 9% fewer amino acids than in homologous proteins of related bacteria. These observations suggest that the *C. ruddii* genome is not subject to constraints that limit the evolution of other known eubacteria.

The insect suborder Sternorrhyncha (Hemiptera), which includes psyllids, also contains aphids, mealybugs, and whiteflies, all of which feed primarily or exclusively on plant phloem sap (5). All of these insects contain primary endosymbionts corresponding to different bacterial clades (11, 12, 18; P. Baumann, N. A. Moran, and L. Baumann, http://link.springer.de/link/service/books/10125/). The most extensively studied endosymbionts are those from aphids (3, 10; Baumann et al., website). The clade constituting these endosymbionts has been designated *Buchnera aphidicola*, and phylogenies based on several molecules indicate that the closest free-living relatives are members of the *Enterobacteriaceae*. Previously we have sequenced about 130 kb of DNA from *B. aphidicola* of the aphid *Schizaphis graminum* (Baumann et al., website), and recently the complete sequence of the 640-kb genome of the endosymbiont of the aphid *Acrithosiphon pisum* has been determined (24). These genetic studies indicate that in *B. aphidicola* there has been a major reduction in the gene content with a retention of the genes necessary for a variety of housekeeping functions as well as the synthesis of essential amino acids and riboflavin (3, 24; Baumann et al., website). The biosynthetic functions of *B. aphidicola* are also supported by numerous nutritional studies (10; Baumann et al., website).

Psyllids, or “jumping plant lice” (Hemiptera: Psyllidae), differ from aphids in many aspects of life history and biogeography but are similar in that they feed on phloem sap (5, 14) and possess maternally transmitted endosymbionts (6, 11, 26; Baumann et al., website). Phloem sap of most plants is deficient in essential amino acids (23), suggesting that *B. aphidicola* and the endosymbionts of psyllids have the same functions related to host nutrition. The primary endosymbionts of psyllids, designated *Carsonella ruddii*, constitute a unique lineage within the γ3 subdivision of the *Proteobacteria* (11, 26, 28). They are located within host cells called bacteriocytes, where they are enclosed by host-derived membrane vesicles; the multicellular structure containing the bacteriocytes is called a bacteriome (6, 7, 11, 31). In a recent study of 32 psyllid species, the phylogenetic tree derived from the 16S-23S rDNA of *C. ruddii* agreed with the tree derived from a host gene, a result consistent with a single infection of a psyllid ancestor and subsequent vertical transmission (cospeciation) of endosymbionts and hosts (28). Some psyllid species also contain morphologically diverse secondary (S) endosymbionts (6, 11, 26); molecular phylogenetic analyses indicate that S-endosymbionts result from multiple infections of hosts and possible horizontal transmission among them (29). Since *C. ruddii* appears to be present in all psyllids and the S-endosymbionts are absent in about one-third of the species (28), *C. ruddii* alone must be able to fulfill all of the necessary functions of the endosymbiotic association.

The 16S and 23S rDNAs of *C. ruddii* have G+C contents lower than those of any other known bacteria (11, 26, 28). In addition, the 3' end of *C. ruddii* 16S rDNA lacks a sequence complementary to the ribosomal binding site (RBS) of mRNA (Shine-Dalgarno sequence) (28). These observations suggested that the genome of *C. ruddii* may also have unusual properties and led to the present study, in which we describe the results of a sequence analysis of 37 kb of *C. ruddii* DNA.

MATERIALS AND METHODS

General methods. Standard molecular biology methods were used in this study (2, 22). Additional methods have been described in our past publications and
include the isolation of total psyllid DNA, restriction enzyme and Southern blot analyses, and cloning into ZAP (Stratagene, La Jolla, Calif.) (28; Baumann et al., website). The nucleotide sequence of C. ruddii DNA was determined at the University of Arizona (Tucson) LMSE sequencing facility. Besides T3 and T7 primers, custom-made oligonucleotide primers were designed for sequencing. For sequence determination of some DNA fragments, a double-stranded DNA nested deletion kit (Pharmacia, Picataway, N.J.) was used.

Most of the sequence data involved the psyllids Pachypsylla venusta and P. celtidis; additional psyllids used were Acizzia uncatoides, Bactericera cockerelli, Ctenarytaina eucalypti, C. longicauda, C. spatulata, Calophya schini, Heteropsylla cubana, and Trioza eugenia. P. venusta and P. celtidis contained only C. ruddii and lacked the S-endosymbiont (28).

**General approach.** In our previous studies on Buchnera aphidicola (endosymbiont of aphids), we developed oligonucleotide primers based on conserved protein sequences, amplified the DNAs by PCR, cloned the fragments into pBluescript (Stratagene), and subsequently determined their nucleotide sequence. These DNA fragments were used as probes for restriction enzyme and Southern blot analyses of endosymbiont DNA. Based on these results, appropriate restriction enzyme-digested DNA fragments were cloned into ZAP (Stratagene) and sequenced (3). Initially this approach was suitable for the cloning of a 4.9-kb DNA fragment from C. ruddii-P. venusta which contained atp genes (Fig. 1B). Subsequent attempts to clone a portion of the 16S rDNA and the region upstream, as well as a portion of the 23S rDNA and the region downstream (Fig. 1C), using these methods failed due to instability of the recombinants or insertion of Escherichia coli sequences. Consequently, we used two approaches to generate sets of overlapping 1.5- to 3.8-kb fragments. In the first approach, we selected proteins that are highly conserved and tend to be clustered in single regulatory units (Fig. 1) (21). Synthetic oligonucleotide primers were designed based on conserved amino acid sequences of homologous proteins. These primers also contained, at the 5' ends, sequences for restriction enzymes. Following amplification by PCR, the DNA was digested with the appropriate restriction enzymes and cloned into pBluescript (Stratagene) or pWSK130 (a low-copy vector) (33), and the nucleotide sequence was determined. The sequences of the oligonucleotide primers and their positions on the sequenced DNA fragments will be made available on request. In some cases we were not able to extend the sequence using this approach and recourse was made to unidirectional PCR.

**Unidirectional PCR.** We used the Vectorette II system (Sigma-Genosys, St. Louis, Mo.) and the instructions provided by the manufacturer. From the nucleotide sequence of endosymbiont DNA fragments, we obtained a usable restriction enzyme site (close to the end which was to be extended) and the sequence for two oligonucleotide primers from which a probe was amplified by PCR. Restriction enzyme and Southern blot analyses were performed using this probe, the restriction enzyme that digests the site in the known sequence, and
RESULTS

General properties of the C. ruddii DNA. The three DNA fragments of C. ruddii DNA were 19,209, 10,049, and 7,806 bp in length (total, 37,057 bp) (Fig. 1A, B, and C, respectively). Searches in databases (September 2000) identified 38 ORFs as corresponding to known genes; 37 were represented in the GenBank accession numbers AF000398, AJ235269, and U00096, respectively). In comparative studies, sequences of B. aphidicola, R. prowazekii, and E. coli were also included (accession numbers AF211138, A232569, and U00096, respectively).

Nucleotide sequence accession numbers. The following are the GenBank accession numbers (in parentheses) for the sequences of the fragments obtained in this study: C. ruddii-P. venusta 19.2 kb (Fig. 1A), AF274444; C. ruddii-C. eucalypti 3.8 kb (Fig. 1A), AF250389; C. ruddii-P. celtidis 3.8 kb (Fig. 1A), AF250390; C. ruddii-P. venusta 10.0 kb (Fig. 1B), AF291051; and C. ruddii-P. celtidis 7.8 kb (Fig. 1C), AF211141. In addition, sequences of rRNA operons (complete or partial) for the following endosymbionts were deposited: C. ruddii-P. venusta, AF211143; C. ruddii-C. eucalypti, AF211133; C. ruddii-A. tenuoides, AF211124; C. ruddii-longicauda, AF211134; C. ruddii-C. spatulata, AF211135; C. ruddii-B. cockerelli, AF211126; C. ruddii-C. schini, AF211132; C. ruddii-H. cubana, AF211138; and C. ruddii-T. genigma, AF211151.

Amino acid composition and codon usage. In C. ruddii, the reduction in G+C content has had a major effect on polypeptide sequences, causing an increased frequency of amino acids that utilize high-A+T-containing codons. Although such a shift occurs in the endosymbiont B. aphidicola and the intracellular pathogen R. prowazekii, it is most drastic in C. ruddii, in which biased base composition shows a more extreme effect on polypeptide sequences than in any previously studied bacterium (Fig. 3). Within the 32 polypeptide sequences inferred for C. ruddii that have homologs in R. aphidicola, R. prowazekii, and E. coli, 50% of the residues consist of five amino acids for which the corresponding codons have maximum A and T content (phenylalanine, lysine, isoleucine, asparagine, and tyrosine). In comparison, these amino acids comprise only 29.3% of the amino acids of B. aphidicola, 29.4% of the amino acids of R. prowazekii, and 22.3% of the amino acids of E. coli, for the same genes. This shift also affects the charge of the proteins. The calculated isoelectric points of the combined proteins of C. ruddii, B. aphidicola, R. prowazekii, and E. coli were 10.1, 9.9, 9.6, and 9.2, respectively.

Intergenic spaces. An inspection of the putative coding regions of C. ruddii DNA indicates a major reduction or elimination of intergenic spaces. A summary of the nucleotide sequences found between 34 adjacent genes, which are transcribed in the same direction, is presented in Table 2. In 85% of the adjacent genes there are overlapping regions between the stop codons for the upstream genes and the initiating codons for the downstream genes. The most frequent overlap arrangement (44%) is ATGA, in which the last 3 nucleotides (nt) are used as a stop codon for the upstream protein and the first 3 nt are used as the initiating methionine for the downstream protein. In 35% of the cases, the overlapping regions are longer. Even in the 15% of the adjacent gene pairs that contain an intergenic space, it is short (7 nt or less). The elimination of intergenic spaces also results in operon fusion. In E. coli and many other organisms, the genes in Fig. 1A are part of five different transcription units (L11 operon, β operon, Str operon, S10 operon, and spe operon) (13, 16). In C. ruddii, these operons appear to be part of a single transcription unit. The 3.8-kb tuf-rpS3 DNA fragment designated by dashed double-headed arrows in Fig. 1A was also sequenced from C. ruddii of two additional psyllid species. Comparisons indicate that there is both conservation and variation in some of the protein initiation and termination arrangements. In C. ruddii-P. venusta, C. ruddii-P. celldis, and C. ruddii-C. eucalypti, the junctions between rpS10-rpL3, rpL3-rpL4, and rpL4-rpL2 were the same (ATGA: the initiating codon is overlined, and the stop codon is underlined). There were differences in the remaining junctions. Between tuf-rpS10 of C. ruddii-P. venusta and C. ruddii-P. celldis there was ATGTATAGA, while in C. ruddii-C. eucalypti the sequence was ATGA. Between tuf-rpS19 of C. ruddii-P. venusta and C. ruddii-P. celldis the sequence was ATGA, while in C. ruddii-C. eucalypti the sequence was ATGTATAGA. Between tuf-rpS19 of C. ruddii-P. venusta and C. ruddii-P. celldis the sequence was ATGA, while in C. ruddii-C. eucalypti the sequence was ATGTATAGA. Between rpL22-rpS3 of C. ruddii-P. venusta and C. ruddii-C. eucalypti the sequence was ATGGGAATTA, while in C. ruddii-P. celldis the sequence was ATGGGTAA. Similarly, the sequence of a 2.8-kb 23S-vaI3 C. ruddii DNA fragment (Fig. 2) indicates a correlation between the conservation of the amino acid sequence of a protein and the G+C content of the C. ruddii gene encoding the protein. Less highly conserved protein sequences are encoded by genes with lower G+C contents.
1C) was determined for C. ruddii of three additional psyllid species. In C. ruddii-P. celtidis, C. ruddii-C. eucalypti, C. ruddii-A. uncatoides, and C. ruddii-C. longicauda there was an overlapping region between the initiating ATG of aphC and the stop codon of valS, consisting of 23, 14, 8, and 14 nt, respectively. Previously it was found that the rpoB-rpoC junction (Fig. 1A, 2.7-kb fragment) of 13 strains of C. ruddii involved the sequence **ATGA** while in the related species

<table>
<thead>
<tr>
<th>Gene</th>
<th>Protein</th>
<th>G+C content (mol%)</th>
<th>% Amino acid identity <em>b</em></th>
</tr>
</thead>
<tbody>
<tr>
<td><em>tal</em></td>
<td>Transaldolase</td>
<td>16.4</td>
<td>36.1</td>
</tr>
<tr>
<td><em>tkt</em></td>
<td>Transketolase</td>
<td>17.6</td>
<td>33.2</td>
</tr>
<tr>
<td><em>atpA</em></td>
<td>ATP synthase, α subunit</td>
<td>24.3</td>
<td>51.6</td>
</tr>
<tr>
<td><em>atpB</em></td>
<td>ATP synthase, subunit a</td>
<td>12.8</td>
<td>31.4</td>
</tr>
<tr>
<td><em>atpD</em></td>
<td>ATP synthase, β subunit</td>
<td>25.8</td>
<td>68.7</td>
</tr>
<tr>
<td><em>atpE</em></td>
<td>ATP synthase, subunit c</td>
<td>23.9</td>
<td>52.0</td>
</tr>
<tr>
<td><em>atpF</em></td>
<td>ATP synthase, subunit b</td>
<td>9.9</td>
<td>22.1</td>
</tr>
<tr>
<td><em>atpG</em></td>
<td>ATP synthase, γ subunit</td>
<td>13.2</td>
<td>33.3</td>
</tr>
</tbody>
</table>

### Amino acid biosynthesis

**Glutamate family**

- *argI* Ornithine transcarbamylase 13.1 28.7

**Aromatic amino acid family**

- *aroQ* 3-Dehydroquinate dehydratase 14.6 33.8

### rRNA

- *rrf* 5S rRNA 26.8 50.0
- *rrl* 23S rRNA 33.1 68.7
- *rrs* 16S rRNA 35.6 73.2

### Ribosomal proteins

- *rpS3* Ribosomal protein S3 (rpS3) 17.5 29.5
- *rpS7* Ribosomal protein S7 (rpS7) 18.5 30.8
- *rpS8* Ribosomal protein S8 (rpS8) 13.4 21.7
- *rpS10* Ribosomal protein S10 (rpS10) 12.8 27.6
- *rpS12* Ribosomal protein S12 (rpS12) 25.1 63.3
- *rpS14* Ribosomal protein S14 (rpS14) 14.2 31.6
- *rpS17* Ribosomal protein S17 (rpS17) 14.7 35.4
- *rpS19* Ribosomal protein S19 (rpS19) 20.6 45.5
- *rpL1* Ribosomal protein L1 (rpL1) 11.3 23.8
- *rpL2* Ribosomal protein L2 (rpL2) 25.4 44.6
- *rpL3* Ribosomal protein L3 (rpL3) 20.2 40.8
- *rpL4* Ribosomal protein L4 (rpL4) 14.8 29.0
- *rpL5* Ribosomal protein L5 (rpL5) 15.0 31.5
- *rpL6* Ribosomal protein L6 (rpL6) 14.6 19.8
- *rpL7/12* Ribosomal protein L7/12 (rpL7/12) 17.0 31.3
- *rpL10* Ribosomal protein L10 (rpL10) 11.1 13.3
- *rpL11* Ribosomal protein L11 (rpL11) 17.5 30.7
- *rpL14* Ribosomal protein L14 (rpL14) 21.1 54.9
- *rpL16* Ribosomal protein L16 (rpL16) 21.8 40.6
- *rpL22* Ribosomal protein L22 (rpL22) 11.4 25.0

### Amino acid tRNA synthetases

- *trpS* Tryptophanyl-tRNA synthetase 12.2 20.3
- *valS* Valyl-tRNA synthetase 10.6 24.8

### DNA replication

- *gidA* Cell division protein 17.8 38.3

### Protein translation

- *fus* Elongation factor G 22.2 46.9
- *tau* Elongation factor Tu 28.3 71.6

### RNA synthesis

- *rpoB* RNA polymerase, β subunit 17.1 34.3
- *rpoC* RNA polymerase, β′ subunit 18.9 36.8

### Detoxification

- *ahpC* Alkyl hydroperoxide reductase 17.2 32.8

### Unidentified

- ORF-A 8.0
- ORF-B 11.1
- ORF-C 9.7

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* Unless otherwise indicated, all genes are from *C. ruddii-P. venusta.*

* Unless otherwise indicated, all comparisons are to *E. coli* proteins.

* Genes from *C. ruddii-P. celtidis.*

* Comparison to *H. influenzae* protein.

* Partial sequence.
C. ruddii-H. cubana and C. ruddii-H. texana, the sequence in the junctions was TAAAAAATG (30). Similarly, in a previous study of the atpAGD region (Fig. 1B, 1.8-kb fragment) from 31 strains of C. ruddii, it was found that in all cases the junction region was ATGA (30).

For 22 pairs of genes transcribed in the same direction, the flanking genes are the same in C. ruddii as in B. aphidicola, R. prowazekii, and E. coli, allowing direct comparison of the homologous intergenic space. For every one of these 22 cases, the space between genes in C. ruddii is absent or smaller than in any of these other three organisms (Fig. 4). This indicates that C. ruddii differs from all of these organisms in the elimination or reduction of intergenic spaces throughout the genome.

The same tendency to minimize sequence length is evident in the two cases in which adjacent genes are transcribed in opposite directions, with end points flanking each other. In argI-ORFC (Fig. 1B) there is an overlap of 26 nt involving the sequence TCAN 18TAG, in which the last triplet is the stop codon for argI and the complement of the first triplet is the stop codon of the putative ORFC. In the ahpC-tal region (Fig. 1C) of C. ruddii-P. celtidis, C. ruddii-P. venusta, C. ruddii-C. eucahpti, C. ruddii-C. longicauda, and C. ruddii-C. spatulata, translation termination involves the sequence TTATAA, in which the last triplet is the stop codon for tal while the complement of the first triplet is the stop codon of ahpC. In C. ruddii-A. uncatoides, termination involves the shorter sequence TTAA, in which the last 3 nt are the stop codon for tal while the complement of the first 3 nt (two of which overlap with the stop codon for tal) are the stop codon for ahpC.

**Protein size.** In Fig. 5 the sizes of 32 C. ruddii proteins are compared with those of homologous proteins of B. aphidicola, R. prowazekii, and E. coli. In 29 of the 32 cases, the C. ruddii proteins were shorter than any of the homologs. The numbers of amino acids in proteins of B. aphidicola, R. prowazekii, and E. coli averaged 9.2, 9.1, and 9.5%, respectively, greater than those in the C. ruddii proteins. The major size difference appears in ribosomal protein L3, which has a substantial deletion

**TABLE 2. Nucleotide sequences between adjacent genes**

<table>
<thead>
<tr>
<th>Sequence</th>
<th>Nucleotide space</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATGNTAA</td>
<td>-8 to -32</td>
<td>35</td>
</tr>
<tr>
<td>ATGA</td>
<td>-2</td>
<td>44</td>
</tr>
<tr>
<td>TAATG</td>
<td>-1</td>
<td>6</td>
</tr>
<tr>
<td>TAANATG</td>
<td>+1, +2, +7</td>
<td>15</td>
</tr>
</tbody>
</table>

* Based on comparisons of 34 adjacent genes that are transcribed in the same direction.
* - , nucleotide overlap between start and stop codons; +, space between start and stop codons.

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at the N terminus. We have also cloned and sequenced the 3.8 kb tuf-rpS3 fragment from *C. ruddii-P. celtidis* and *C. ruddii-C. eucalypti* (Fig. 1A). A comparison of the lengths of the proteins in *C. ruddii* from these three psyllid species indicated that they are identical or nearly identical in size. The number of amino acids is listed in parentheses following the protein designation in the order *C. ruddii-P. venusta, C. ruddii-P. celtidis, and C. ruddii-C. eucalypti* (where the size is constant only one value is given): Tuf, partial 312, 312, 310; RpS10, 98; RpL3, 136; RpL4, 172; RpL2, 235, 235, 234; RpS19, 88; RpL22, 104, 101, 105; RpS3, partial 148, 148, 150.

**rRNA operon.** Sequence comparison of the DNA region upstream of rRNA operons of *B. aphidicola* indicated that a sequence resembling the −35, −10 promoter region is conserved and is followed by downstream conserved sequences resembling boxA and boxC (3; Baumann et al., website). In addition, inverted repeats resembling rho-independent terminators following the rRNA operons were found. We have used a similar comparative approach to look for sequences conserved upstream of 16S rDNA which might correspond to putative promoter regions. The results are presented in Fig. 6A and indicate conservation of a sequence resembling boxA; no
sequences resembling the ~35, ~10 promoter region were found upstream of the putative boxA, in the last 200 nt of trpS. Similarly, there is no sequence conservation following 5S rDNA (Fig. 6B) and no significant inverted repeats resembling rho-independent terminators (25). These results suggest that the 16S-23S-5S rRNA genes are part of a larger transcription unit that includes upstream and downstream genes (Fig. 1C).

**DISCUSSION**

**General properties.** *C. ruddii* has a unique combination of properties distinguishing it from other known bacteria. These properties consist of (i) extremely low G+C content (18% for the protein-coding regions); (ii) elimination or reduction of intergenic spaces, sometimes resulting in fusion of operons; (iii) absence of the complement of an RBS at the 3' end of 16S rDNA (28) and an RBS preceding any structural genes; (iv) a reduction in protein size; (v) absence of conserved sequences corresponding to the ~35, ~10 promoter region preceding 16S rDNA; and (vi) absence of inverted repeats characteristic of rho-independent terminators following 5S rDNA. The absence of intergenic spaces and the complement of the RBS at the 3' end of 16S rDNA suggest that long polycistronic mRNAs are made and translational coupling occurs during protein synthesis (25). The elimination or reduction of intergenic spaces and the reduction in the protein sizes indicate a history of mutational pressure and/or selection favoring reduced sequence length. One interpretation of the tendency toward elimination of intergenic spacers and reduction in gene length in *C. ruddii* is that mutations are biased in favor of deletions relative to insertions and that selection to oppose these deletions is often weak or ineffective (due to genetic drift). An analysis of mutational patterns in pseudogenes of *Rickettsia* species indicated a mutational bias favoring deletions (1). It is possible that *C. ruddii* experiences similar mutational bias but relaxed selection for conservation of function (or less efficient selection due to increased genetic drift). An analysis of mutational patterns in pseudogenes of *Rickettsia* species indicated a mutational bias favoring deletions (1). It is possible that *C. ruddii* experiences similar mutational bias but relaxed selection for conservation of function (or less efficient selection due to increased genetic drift) results in even greater shrinkage. Alternatively, selection might have favored the reduction of sequence length for some unknown reason. However, the length reduction occurs both in spacers (Fig. 4) and in protein-coding sequences (Fig. 5), which are generally subject to different kinds of selection. The bias toward A+T in DNA sequences is most readily explained as the result of mutational pressure. The pattern in Fig. 2 suggests that this mutational pressure is opposed by selection for conservation of function, with the result that genes encoding proteins which are poorly conserved have a higher A+T content. Conservative selection opposing the mutational bias appears to be weaker or less efficient than in other bacteria, with the result that *C. ruddii* has the most extreme bias toward A+T known for bacteria (15). A similarly low G+C content has been found in the genome of the eukaryote *Plasmodium* (20).

The unique combination of properties found in *C. ruddii* is also illustrated by a comparison of the G+C contents of homologous rRNA subunits of bacteria, plastids, and mitochondria with the G+C contents of their genomes (or representa-
In addition, we detected many of the components of the ATP synthase. In E. coli and many other organisms, the genes order of the ATP synthase is \textit{atpBEFHAGDC} (9). In \textit{C. ruddii}, part of this order is maintained (Fig. 1B). However, we have not been able to detect \textit{atpH} or \textit{atpC}. ORFA and ORFB are found in the locations where these genes are expected, but they show no significant similarity to \textit{atpH} or \textit{atpC}. The presence of genes encoding ATP synthase suggests that \textit{C. ruddii} may be able to generate ATP from the proton motive force. Genes for transaldolase (Fig. 1B) and transketolase (Fig. 1C) were detected, suggesting a functional nonoxidative pentose phosphate cycle. In addition, genes encoding proteins involved in the synthesis of amino acids of the glutamate and aspartate family were found. These results suggest that \textit{C. ruddii} may be able to synthesize essential amino acids for the plastid host as in the case of \textit{B. aphidicola}, the endosymbiont of aphids (10, 24; Baumann et al., website).

Comparison with \textit{B. aphidicola}. Both \textit{C. ruddii} and \textit{B. aphidicola} are associated with insect hosts that feed on plant phloem sap and thus have similar nutritional needs. These endosymbiotic associations result from infections by two different bacterial ancestors, both from the \(\gamma\) division of the proteobacteria (11, 18, 28; Baumann et al., website). It is likely that \textit{C. ruddii} retains genes for essential amino acid pathways, as in \textit{B. aphidicola}. Some of the same genomewide features appear in both species and can be interpreted as convergences; these include increased A+T content (18), accelerated sequence evolution (18, 26, 28), and shortened proteins (8). Although a slight reduction in protein length has been noted previously for \textit{B. aphidicola} relative to \textit{E. coli} (8), the reduction is much more severe in \textit{C. ruddii}, as is evident in Fig. 5. \textit{B. aphidicola} has intergenic spaces similar to those found in other bacteria (Fig. 4), and the 3'end of its 16S rDNA has the complement of the RBS (3, 24; Baumann et al., website). The G+C contents of both the \textit{B. aphidicola} genome and the \textit{B. aphidicola} rRNA are comparable to those of other bacteria at the lower end of the known scale (Fig. 7). In \textit{C. ruddii}, the modifications are considerably more drastic, resulting in an even lower G+C content of the DNA and almost complete elimination of intergenic spaces. \textit{B. aphidicola} and some other bacteria with low G+C contents have very small genomes, having lost most of the genes found in ancestors (1, 24); future studies determining the \textit{C. ruddii} genome size will be of interest.

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**References**


FIG. 7. Comparison of the G+ C content of 23S and 23S-like rDNA with the G+ C content of the genome. , bacteria; , mitochondria; +, plastids; arrow a, \textit{B. aphidicola}; arrow b, \textit{Reclinomonas americana}; arrow c, Cr-Pve; arrow d, \textit{Toxoplasma gondii}; arrow e, \textit{Plasmodium falciparum}. Data compiled in January 2000 from GenBank and data in reference 15.


