Low bacterial community diversity in two introduced aphid pests revealed with 16S rRNA amplicon sequencing

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ABSTRACT

Bacterial endosymbionts that produce important phenotypic effects on their hosts are common among plant sap-sucking insects. Aphids have become a model system of insect-symbiont interactions. However, endosymbiont research has focused on a few aphid species, making it necessary to make greater efforts to other aphid species through different regions, in order to have a better understanding of the role of endosymbionts in aphids as a group. Aphid endosymbionts have frequently been studied by PCR-based techniques, using species-specific primers, nevertheless this approach may omit other non-target bacteria cohabiting a particular host species. Advances in high-throughput sequencing technologies are complementing our knowledge of microbial communities by allowing us the study of whole microbiome of different organisms. We used a 16S rRNA amplicon sequencing approach to study the microbiome of aphids in order to describe the bacterial community diversity in introduced populations of the cereal aphids, Sitobion avenae and Rhopalosiphum padi in Chile (South America). An absence of secondary endosymbionts and two common secondary endosymbionts of aphids were found in the aphids R. padi and S. avenae, respectively. Of those endosymbionts, Regiella insecticola was the dominant secondary endosymbiont among the aphid samples. In addition, the presence of a previously unidentified bacterial species closely related to a phytopathogenic Pseudomonad species was detected. We discuss these results in relation to the bacterial endosymbiont diversity found in other regions of the native and introduced range of S. avenae and R. padi. A similar endosymbiont diversity has been reported for both aphid species in their native range. However, variation in the secondary endosymbiont infection could be observed among the introduced and native populations of the aphid S. avenae, indicating that aphid-endosymbiont associations can vary across the geographic range of an aphid species. In addition, we discuss the potential role of aphids as vectors and/or alternative hosts of phytopathogenic bacteria.

Subjects Ecology, Entomology, Genomics

Keywords Sitobion avenae, Hamiltonella defensa, Pseudomonas, Regiella insecticola, Aphids, Bacterial community, Endosymbionts, Phytopathogens, Rhopalosiphum padi, 16S rRNA amplicon sequencing

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INTRODUCTION

Associations between bacterial endosymbionts and insects are widespread in nature (Gibson & Hunter, 2010). The microbial community inhabiting insects can be as diverse as the symbiotic associations that they maintain with their host insects. Mutualistic, pathogenic, and commensal relationships can take place concurrently and can significantly influence the insect host ecology (Toft & Andersson, 2010). For instance, ancient mutualistic relationships with primary or obligate bacterial endosymbionts that provide missing essential amino acids to phloem-based diets are common among plant sap-sucking insects (e.g., psyllids, whiteflies, mealybugs and aphids) (Baumann, 2005). Primary endosymbionts are usually found among the Betaproteobacteria and Gammaproteobacteria subgroups (Toft & Andersson, 2010). Contrary to primary endosymbionts, secondary or facultative endosymbiotic bacteria are not essential for host survival and reproduction and they are mainly found among the Alphaproteobacteria, Gammaproteobacteria (especially Enterobacteriaceae) and Bacteroidetes (Baumann, 2005; Moran, McCutcheon & Nakabachi, 2008). However, secondary endosymbionts may produce ecologically important phenotypic effects on their insect hosts. Specifically, they can establish facultative mutualistic associations with insects thus conferring beneficial traits such as protection against natural enemies (review by Oliver et al., 2010; Jaenike et al., 2010; Jiggins & Hurst, 2011), or they can establish parasitic associations that have deleterious effects on host fitness (Werren, Baldo & Clark, 2008).

Aphids (Hemiptera: Aphididae) are phloem-feeding insects that reproduce by cyclical parthenogenesis (clonal) (Simon, Rispe & Sunnucks, 2002). They represent serious pests by reducing crop yields and quality, and can act as vectors of phytopathogenic viruses and bacteria (Dedryver, Le Ralec & Fabre, 2010; Ng & Perry, 2004; Nadarasa & Stavrinides, 2011). At least 15 aphid species are considered global crop pests of major agricultural importance (including the grain aphid Sitobion avenae, bird cherry-oat aphid Rhopalosipum padi and pea aphid Acrhythosiphon pisum), of which the majority are of Palaeartic origin (Eurasia) (Van Emden & Harrington, 2017). Symbiotic bacteria have been well studied in this insect group, becoming a model system of the insect-symbiont interactions (Oliver, Smith & Russell, 2014). Aphids have a well-known obligate nutritional relationship with the primary endosymbiont Buchnera aphidicola, which confers essential nutrients to the aphid host (Douglas, 1998). At least nine common secondary endosymbionts have been reported among aphid species, including six Gammaproteobacteria; Hamiltonella defensa, Serratia symbiotica, Regiella insecticola, PAXS (Pea aphid X-type symbiont), Rickettsiella viridis and Arsenophonus sp., and two Alphaproteobacteria of the genera Wolbachia and Rickettsia, as well Spiroplasma from Mollicutes (reviewed in Zytynska & Weisser, 2016). These secondary endosymbionts have diverse effects on the aphid phenotype, such as conferring protection against natural enemies (parasitoids and fungal pathogens) (Oliver et al., 2003; Oliver, Moran & Hunter, 2005; Vorburger, Gehrer & Rodriguez, 2009; Scarborough, Ferrari & Godfray, 2005; Parker et al., 2013), providing resistance to heat stress (Montllor, Maxmen & Purcell, 2002), influencing insect-plant interactions (Tsuchida, Koga & Fukatsu, 2004; Tsuchida et al., 2011;
Ferrari, Scarborough & Godfray, 2007), as well as manipulating aphid reproduction (Simon et al., 2011). These heritable bacterial endosymbionts are mainly maintained in aphid populations through vertical transmission (i.e., maternal) and to a lesser extent by horizontal transmission (e.g., sexual) (Vorburger, 2014; Peccoud et al., 2014). Although, the aphid–endosymbiont interactions have received considerable attention, much of this research has been focused in the model pea aphid, A. pism. Accordingly, there is a lack of data for some aphid species across different regions particularly at the continental scale (e.g., South America) (Zytynska & Weisser, 2016). Therefore, it is necessary to make greater efforts to other aphid species in order to have a better understanding of the role of endosymbionts in aphids as a group. In addition, aphid endosymbionts have frequently been studied by PCR-based approaches, using species-specific primers. In spite of increasing the ease of testing for specific symbionts, and being useful for detecting target endosymbiont groups, this approach may omit other non-target bacteria cohabiting a particular host species. Regarding this, advances in high-throughput sequencing technologies are now complementing our previous knowledge of microbial endosymbiont communities (Riesenfeld, Schloss & Handelsman, 2004). A greater understanding of the microbiome of aphid species through next-generation sequencing could allow the identification of novel bacterial associations and their potential effects on the ecology and phenotype of aphid species. Such knowledge could be instrumental for understanding the role of the bacterial interactions on the invasive potential of economically important aphid species.

We used a 16S rRNA amplicon sequencing approach to study the microbiome of aphids, in order to describe the bacterial community diversity in introduced populations of the cereal aphids, Sitobion avenae and Rhopalosiphum padi in Chile (South America). Then we discuss whether the bacterial community diversity found in these introduced populations of cereal aphids is similar to the previously estimated in native populations of these aphid species (Europe).

**MATERIALS AND METHODS**

**Sample collection and DNA extraction**

A total of 80 individuals of the aphid S. avenae and 52 individuals of the aphid R. padi were collected from oat (Avena sativa) and wheat (Triticum aestivum) crops in two different agroclimatic regions (Maule and Los Ríos regions) in Chile (Table 1). In addition, the field experiments performed in this study were approved for Ethical scientific committee of the Universidad de Talca in Chile (FONDECYT project 3140299). DNA extraction was individually performed for each aphid specimen using the “Salting out” method described by Sunnucks & Hales (1996). The quantification and quality of the extracted DNA was examined by absorbance using Infinite 200 PRO NanoQuant (TECAN) and by electrophoresis in 0.8% agarose gels. Each individual DNA extraction was normalized to a concentration of 5 ng/ul and kept at −20 °C until later 16 S library preparation.

**16S rRNA amplicon sequencing library preparation**

In order to produce DNA pools that represent the genetic diversity of aphids from different species, locations and host-plants; four DNA pools of 20 S. avenae aphids and
Table 1  Summary of collection details and 16S rRNA gene sequencing results for aphid samples. Host plant, locality, date, total number of reads, and Shannon diversity index for each sample of S. avenæ (SA-1, SA-2, SA-3 and SA-4) and R. padi (RP-1, RP-2, RP-3, RP-4, RP-5 and RP-6).

| Sample ID | Host plant | Locality | Date      | Numbers of reads after filtering | Shannon index |
|-----------|------------|----------|-----------|---------------------------------|---------------|
| SA-1      | Oat        | Los Ríos | 10/2014   | 165,703                         | 2.39          |
| SA-2      | Wheat      | Los Ríos | 11/2014   | 317,806                         | 1.13          |
| SA-3      | Wheat      | Maule    | 10–12/2013| 500,906                         | 1.2           |
| SA-4      | Wheat      | Maule    | 10–12/2013| 343,371                         | 1.96          |
| Total     |            |          |           | 1,327,786                       | 1.51 (0.61)   |
| RP-1      | Wheat      | Maule    | 10/2013   | 210,118                         | 0.16          |
| RP-2      | Wheat      | Maule    | 10/2013   | 372,013                         | 0.06          |
| RP-3      | Wheat      | Maule    | 10/2013   | 402,514                         | 0.06          |
| RP-4      | Wheat      | Maule    | 10/2013   | 351,336                         | 0.06          |
| RP-5      | Wheat      | Maule    | 10/2013   | 556,558                         | 0.07          |
| RP-6      | Wheat      | Maule    | 10/2013   | 203,033                         | 0.06          |
| Total     |            |          |           | 2,095,602                       | 0.07 (0.04)   |

six DNA pools of 9 R. padi aphids were used (Table 1). Pools of the genomic DNA were generated in two steps using the Illumina MiSeq protocol for 16S amplicon sequencing (https://support.illumina.com/downloads/). The V3 and V4 variable regions of the 16S ribosomal RNA gene were amplified using the primer pair S-D-Bact-0341-b-S-17 and S-D-Bact-0785-a-A-21 (Klindworth et al., 2012); these regions have a total length of approximately 460 bp. Specific Illumina Adapter sequences were added to the 5' region of the forward (5'-TCGTCGGCAGCGTCAGATGTGTATAAGAGACAG) and reverse primers (5'-GTCTCGTGGGCTCGGAGATGTGTATAAGAGACAG). PCR reactions were performed in a volume of 15 µl; 1.5 µl of normalized DNA (5 ng/µl), each primer at 0.2 µM plus 7.5 µl of 2x KAPA Hifi HotStart Ready Mix (KAPA Biosystems, Wilmington, MA, USA). The PCR program consisted of initial denaturation at 95 °C for 3 min, followed by 25 cycles of: 95 °C for 30 s, 55 °C for 30 s and 72 °C for 30 s, and then a final extension of 72 °C for 5 min. The amplicons were purified using a PCR clean-up protocol including 10 µl of PCR product, 8 µl of AMPureXP (Beckman Coulter, Brea, CA, USA), and 200 µl of 80% ethanol. The mixture was incubated on a magnetic stand and diluted in 22.5 µl of Tris (10 mM), pH 8.5. The expected size of PCR amplicons was verified using the DNF-900 Kit for Fragment Bioanalyzer (Advanced Analytical Technologies, Inc., Ankeny, IA, USA); the quantity of amplicons was estimated by fluorescence using the Quant-iT PicoGreen dsDNA Assay kit (Invitrogen, Carlsbad, CA, USA) and a HOEFER DQ300 fluorometer (Hoefer Inc., Holliston, MA, USA). Each DNA pool was constructed by mixing 5 µl of each of the amplicons (Table 1). Then DNA pools were subjected to a second PCR where dual indices and Illumina sequencing adapters were attached using a NEXTERA XT index Kit (Illumina, San Diego, CA, USA). This second PCR was conducted in a total volume of 50 µl which contained 5 µl of each pooled DNA, 5 µl of each Nexetera XT index primer, 25 µl of 2x KAPA Hifi HotStart Ready Mix, and 10 µl of PCR grade water. PCR program consisted of initial denaturation at 95 °C for 3 min, followed by eight cycles of: 95 °C for 30 s, 55 °C
for 30 s and 72 °C for 30 s, and 72 °C for 5 min. The PCR product was corroborated using a Fragment Analyzer and the DNF 479 kit. Finally, each DNA pool was normalized to a concentration of 4 nM and then pooled. The mix DNA pool was prepared for sequencing using the Denature and Dilute Libraries Guide. Paired-end sequencing was performed using the Miseq Reagent Kit v3 (2 × 300 cycles) on the MiSeq Illumina sequencing platform in the AUSTRAL-omics Core-Facility (Facultad de Ciencias, Universidad Austral de Chile).

**Data analysis**

Removal of adapters and quality filtering of the data were conducted using the Trimmomatic and PRINSEQ software (Bolger, Lohse & Usadel, 2014; Schmieder & Edwards, 2011). To assemble the overlapping Illumina Paired-end reads PANDAseq was used (Masella et al., 2012). In order to determine operational taxonomic units (OTUs), sequences sharing 97% identity were assembled as suggested by Kunin et al. (2010); this was done using the software QIIME (Caporaso et al., 2010). The OTUs were aligned using the GreenGenes database (http://greengenes.lbl.gov). Bacterial diversity was studied using the Shannon diversity index calculated for each DNA pool. The relative abundance of each OTU was estimated by examining the number of reads for each sequence and each sample as recommended by Jousselin et al. (2016). Taking into account that bacterial DNA contaminants can be commonly found in DNA extraction kits and other laboratory reagents or could enter samples during analysis (Salter et al., 2014), reads from taxa accounting for <1% of all the reads of a given sample were excluded from the data analysis (“unrepresented reads”). Regarding this, Jousselin et al. (2016) found that the removal of low frequency sequences (<1%) excluded the most DNA contaminants allowing for increased repeatability and reliability of results. They showed that by using this method, DNA contaminants have little impact on the analysis of aphid endosymbionts when using 16S rRNA Illumina sequencing. While, reads for which significant BLAST hits with known taxon could not be found are detailed as “unassigned reads”.

**Identifying *Pseudomonas* species by 16S Sanger sequencing**

From the 16S rRNA amplicon sequencing, a species of *Pseudomonas* was encountered (see ‘Results’). In order to characterize the *Pseudomonas* species from the 16S rRNA sequences identified, a portion of the 16S and 23S ribosomal genes (~1,500 bp) was amplified and sequenced in 20 aphids collect from field and used to prepare sample SA-1 (Table 1); this was done using the universal bacterial primers 10F and 35R (Sandström et al., 2001; Russell & Moran, 2005). These primers were selected because they target the intergenic spacer between the 16S and 23S genes, which can be used to avoid amplifying the aphid primary endosymbiont, *B. aphidicola* as both genes are not contiguous in this endosymbiont (Russell & Moran, 2005).

The PCR reactions were performed in a total volume of 25 µl including: 2.5 µl of 10× buffer, 0.2 mM dNTP’s, 2 mM MgCl₂, 0.3 µl of Taq (5 U/µl), each primer at 0.5 uM, and 3 µl of DNA (10 ng/µl). The PCR conditions consisted of initial denaturation at 94 °C for 5 min, followed by 35 cycles of 94 °C for 40 s, 57 °C for 40 s and 72 °C for 45 min and a final extension at 72 °C for 7 min. The resulting amplicons were sequenced in an
ABI PRISM® 310 Genetic Analyzer (Applied Biosystems, Foster City, CA, USA). The alignment of the sequences with known 16S rRNA sequences from species of the genus *Pseudomonas* was conducted in Geneious v.8.1 (Drummond et al., 2011). All sequences of the genus *Pseudomonas* were obtained from GenBank, including sequences from the seven *Pseudomonas* clusters reported by Anzai et al. (2000): “*Pseudomonas syringae* group”, “*Pseudomonas chlororaphis* group”, “*Pseudomonas fluorescens* group”, “*Pseudomonas putida* group”, “*Pseudomonas stutzeri* group”, “*Pseudomonas aeruginosa* group”, and “*Pseudomonas pertucinogenae* group” (Data S1). A phylogenetic tree of the *Pseudomonas* sequences was constructed using the HKY genetic distance model and the neighbor-joining method implemented in Geneious v.8.1 (Drummond et al., 2011). Branch significance was calculated using bootstrap values of 1,000 replications.

RESULTS

16S rRNA amplicon sequencing

A total of 1,327,786 reads were obtained after filtering the four DNA pools of the aphid *S. avenae* (SA-1, SA-2, SA-3 and SA-4) (Table 1). The mean Shannon diversity index was 1.51 (SD = 0.61) and ranged from 1.13 to 2.39 for the aphid *S. avenae* (Table 1). Of the total reads for *S. avenae*, 98% were classified as Gammaproteobacteria and included mostly bacteria from the Enterobacteriaceae (94.7% of the total reads) (*Buchnera aphidicola*, *Regiella insecticola* and *Hamiltonella defensa*) and to a lesser extent from the Pseudomonadaceae (*Pseudomonas*) families (3.3% of the total reads). The Gammaproteobacteria, the aphid primary endosymbiont, *Buchnera aphidicola*, was the most common endosymbiont in the four samples (84.4% of the total reads). The second most common taxon was the aphid secondary endosymbiont *R. insecticola* (9.3% of the total reads), which was also found in all studied samples (Fig. 1). Whilst another well-known aphid endosymbiont, *H. defensa*, represented an average of 0.9% of the total reads in two of the four samples studied (SA-1 and SA-3) (Fig. 1 and Data S2). *Pseudomonas* sp. sequences were well represented in two aphid samples (SA-1 and SA-4), making up an average of 3.3% of the total reads (Fig. 1 and Data S2). Unrepresented reads (i.e., reads of taxa accounting for <1% of all the reads; see methods) were found in an average of 1.5% of the total reads. Also, the four DNA pools of *S. avenae* had a low proportion of unassigned reads (i.e., reads for which no significant BLAST hits with known taxon were found); unassigned reads ranged from 0.5% to 0.6% (average of 0.5% of the total of reads).

For the aphid *R. padi*, a total of 2,095,602 reads were obtained from the six DNA pools analyzed (RP-1, RP-2, RP-3, RP-4, RP-5 and RP-6) (Table 1). A lower bacterial diversity than *S. avenae* was observed with an estimated mean Shannon diversity index of 0.07 (SD = 0.04) (Table 1). *B. aphidicola* was found in a percentage >98.5% in all DNA pools, however no additional bacteria were found in *R. padi* (Fig. 1). Finally, a low proportion of unassigned reads was detected among the six DNA pools (<0.01%), as well the proportion of unrepresented reads was found in an average of 0.24% of the total reads, being the highest proportion of unrepresented reads detected in the DNA pool RP-1 (1.3% of the total of reads) (Fig. 1). Sequencing data generated on Illumina were submitted to GenBank
Proportion of taxa assignments for each DNA pool of *S. avenae* (SA-1, SA-2, SA-3 and SA-4) and *R. padi* (RP-1, RP-2, RP-3, RP-4, RP-5 and RP-6). The proportion of sequences assigned to <1% of the total reads per sample are identified as ‘unrepresented’, and sequences that did not cluster with any known sequences are identified as ‘unassigned’.

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(accession numbers: MG958610, MG958611, MG958612, MG958613 and MG958614) (Data S2).

**16S rRNA sequencing and phylogenetic analysis of *Pseudomonas* species**

Of the sequences generated for the 20 aphid samples of *S. avenae*, only one DNA sample corresponded to a *Pseudomonas* species (GenBank accession number MF536106). Sequences of the other DNA samples were observed as belonging to some of the other aphid secondary endosymbionts (*R. insecticola* and *H. defensa*), as it was identified by the 16S rRNA sequencing. The phylogenetic tree constructed show the seven clusters previously described for the genus *Pseudomonas* (Fig. 2). The *Pseudomonas* sequence generated from aphid DNA was located into the “*P. fluorescens* group”, being closely related to *Pseudomonas palleroniana* with an identity percentage >95% (Fig. 2).

**DISCUSSION**

**Secondary endosymbionts in the introduced aphid populations**

A low bacterial diversity in the introduced populations of the cereal aphids *S. avenae* and *R. padi* was revealed by 16S rRNA amplicon sequencing in Chile. Gammaproteobacteria was the most common class identified and as expected the aphid primary endosymbiont, *B. aphidicola*, was the most common bacterial species detected in *S. avenae* and *R. padi*. In all DNA pools of both aphid species, *Buchnera* made up a large percentage of all of
the reads (ranged between 84.4% and 99% respectively). In contrast to our systems, a greater diversity of secondary endosymbionts can be found in other aphid species (Zytynska & Weisser, 2016). For instance, the well-studied pea aphid, A. pisum, hosts at least eight secondary endosymbionts (Serratia symbiotica, R. insecticola, H. defensa, Rickettsiella, PAXS, Spiroplasma, Rickettsia and Wolbachia) that are highly abundant according to two 16 rRNA amplicon sequencing studies (Russell et al., 2013; Gauthier et al., 2015). A similar study in several aphid species of the genus Cinara has detected the presence of four dominant secondary endosymbions (S. symbiotica, H. defensa, R. insecticola and Wolbachia) (Jousselin et al., 2016). Our results show an absence of secondary endosymbionts in the introduced populations of the aphid R. padi. Whilst the diversity of secondary endosymbionts observed in the aphid S. avenae was low; only two well-known aphid endosymbionts were encountered, R. insecticola and H. defensa. A similar bacterial diversity has been found in other geographic populations of these aphid species. For instance, aphid samples screened for a set of endosymbionts (H. defensa, R. insecticola, S. symbiotica, Rickettsia, Rickettsiella and Spiroplasma) have found only three secondary endosymbionts (H. defensa, R. insecticola and S. symbiotica) in the native range.
of *S. avenae* (U.K. and Germany) (*Łukasik et al.*, 2013; *Henry et al.*, 2015; *Alkhedir et al.*, 2015). In particular, a positive association between *H. defensa* and *S. avenae* was found, being this the most common endosymbiont followed by *R. insecticola*, whilst *S. symbiotica* was reported in a lower frequency (≤6%) in the aphid populations (*Łukasik et al.*, 2013; *Henry et al.*, 2015). Differently, higher infection rates of *R. insecticola* and *S. symbiotica* were found in Chinese populations of *S. avenae* (*Luo et al.*, 2016), as well a high prevalence of *R. insecticola* (75% of infected aphids) was found in introduced populations of *S. avenae* in Morocco (*Fakhour et al.*, 2018). In this study, we found that *R. insecticola* was the dominant secondary endosymbiont in *S. avenae*, while *H. defensa* was observed at lower prevalence among DNA samples studied. However, the read abundance should be interpreted carefully when it is used as an estimate of the infection frequency of endosymbions, because PCR amplification bias can be introduced by primer specificity (*Klindworth et al.*, 2012). Despite this, our results from the deep sequencing of 16S rRNA gene are consistent with previous PCR-based studies on Chilean populations of *S. avenae*, in which ∼50% of the aphids harbored *R. insecticola* and a lower proportion of aphids harbored *H. defensa* (between 4% and 15%) (*Sepúlveda et al.*, 2016; *Zepeda-Paulo, Villegas & Lavandero*, 2017), suggesting that the aphid-endosymbiont associations can vary across geographic range of aphid species.

Secondary endosymbionts make up an important component of the bacterial community of aphids and several studies have indicated that they have important effects on the host phenotype. Specifically, aphid secondary endosymbionts can protect the host from natural enemies, can provide tolerance to heat shock and can facilitate the colonization of new host plants (*Oliver et al.*, 2010). Although recent studies have not found evidence that the endosymbionts *R. insecticola* nor *H. defensa* can confer defense against parasitoid wasps in *S. avenae* (*Łukasik et al.*, 2013; *Zepeda-Paulo, Villegas & Lavandero*, 2017), at least one strain of *R. insecticola* has shown to provide protection to *S. avenae* against the pathogenic fungus *Pandora neoaphidis* (*Łukasik et al.*, 2015). This symbiont-mediated advantage could explain the higher prevalence of *R. insecticola* in the populations of *S. avenae* here studied; however, this is not consistent with the lower prevalence of this endosymbiont reported in native regions of *S. avenae*. An explanation for this observation could be the founder effect and drift experienced by aphid populations introduced in a new region (*Desneux et al.*, 2018). During the invasive process only a subset of symbiont-harboring aphid clones may have been introduced from the native regions, resulting in particular aphid—endosymbiont associations in the novel established populations. Indeed, variation in the associations between aphid clones and endosymbionts can be found in field populations, suggesting that they could be relevant for understanding of aphid—symbiont populations dynamics (*Zepeda-Paulo, Villegas & Lavandero*, 2017). In addition, we cannot rule out the effect of sampling method (e.g., number and distribution of sampling in a season) on the infection rates observed in aphid populations, since the frequency of endosymbionts can increase and/or fluctuate during the course of a season (*Henry et al.*, 2015). In this regard, our aphid sampling would be considered representative of the endosymbiont diversity, as it was performed during the period of highest abundance of aphids (*Raymond, Ortiz-Martínez & Lavandero*, 2015; *Ortiz-Martínez & Lavandero*, 2018) and endosymbionts of the populations in the field (F Zepeda-Paulo & B Lavandero, 2018, unpublished data).
Unlike *S. avenae*, there is little knowledge on the diversity of bacterial endosymbionts in the aphid *R. padi*. Despite this, the existing data are consistent with our results in showing an absence of secondary endosymbionts in aphid samples from the native range of *R. padi* (Europe) analyzed using species-specific primers developed for three aphid endosymbionts (*H. defensa, R. insecticola* and *S. symbiotica*) (Henry et al., 2015; Desneux et al., 2018) nor their introduced range (Morocco) using 16S rRNA gene sequencing (Fakhour et al., 2018). The bacterial diversity could be non-randomly distributed throughout host species. In this sense, it has been raised that the prevalence of secondary endosymbionts in a particular insect host may depend on the balance between the costs and benefits of harboring symbionts (Oliver, Smith & Russell, 2014). Indeed, the lack of an important protective phenotype providing direct benefits, fitness costs on symbiont-harboring host and the transmission rates of endosymbionts are some of the factors that could explain the low occurrence of endosymbionts in a particular host species (Oliver, Smith & Russell, 2014; Dykstra et al., 2014). Another factor that may influence bacterial diversity of aphids are the symbiont-symbiont interactions, such as competition between primary and secondary endosymbionts. Regarding this, several studies have shown that the density of the aphid primary endosymbiont, *B. aphidicola*, could be affected by the coexistence with secondary endosymbionts in the same host (Koga, Tsuchida & Fukatsu, 2003; Sakurai et al., 2005; Leclair et al., 2017). A negative effect on *Buchnera* abundance may be detrimental to the fitness of aphids and could significantly affect some aphid species. Aphids species can vary in their ability to increase the amino acid concentration in the phloem, in response to chlorotic damage induced by them (Sandström, Telang & Moran, 2000). This increase may reduce the nutritional dependence of aphids on *Buchnera* for the synthesis of essential amino acids, which could affect the aphid-symbiont associations. For instance, *R. padi* could show a high dependence on *Buchnera* for the synthesis of essential amino acids, since this does not affect the phloem composition of the host plant, compared to a higher amino acid concentration induced by other aphid species (Sandström & Moran, 1999; Sandström, Telang & Moran, 2000). A greater dependence in *Buchnera* could limit the infection of secondary endosymbionts, if they affect the abundance of the primary endosymbiont of hosts and thus explain the absence of secondary endosymbionts in some aphid species. However, the association between *Buchnera*-dependent aphids and the prevalence of secondary endosymbionts still have to be studied for a better understanding of the role of symbiont–symbiont interactions on the bacterial diversity of aphid species.

**Presence of Pseudomonas sp. in cereal aphids**

In addition to the most common aphid endosymbionts, the results from 16S rRNA sequencing showed the occurrence of *Pseudomonas* sp. in two DNA pools analyzed of the aphid *S. avenae*. However, of the sequences generated for 20 aphid samples of *S. avenae*, only one DNA sample corresponded to a *Pseudomonas* species. The phylogenetic analysis incorporating known *Pseudomonas* sequences showed clustering with the “*P. fluorescens* group”; the *Pseudomonas* sp. sequence generated here was closely related to the bacteria *P. palleroniana* and *P. tolassi*. These bacterial species are known phytopathogenic Pseudomonads, which have been found in rice (*Oryza sativa*) and garlic.
(Allium sativum), respectively (Gardan et al., 2002; Höfte & De Vos, 2007). Others studies based on 16 rRNA amplicon sequencing have identified phytopathogenic Pseudomonas sp. in the pea aphid (Pseudomonas syringae) and R. padi (P. viridiflava and P. veronii) (Gauthier et al., 2015). Moreover, the pea aphid has previously proven capable of acting as both a vector and a non-plant host for P. syringae (Stavrinides, McCloskey & Ochman, 2009). Some strains of P. syringae could be pathogenic to aphids causing death by bacterial sepsis (Stavrinides, McCloskey & Ochman, 2009; Hendry, Clark & Baltrus, 2016). The finding of Pseudomonas sp. in different aphid species suggests that these types of phytopathogen-vector associations may be more common than previously thought among aphid species. Secondary endosymbionts can also influence the interactions between phytopathogens and insects. Hendry, Hunter & Baltrus (2014) reported that secondary endosymbionts can influence interactions between whiteflies and the phytopathogen (P. syringae); whiteflies harboring Rickettsia have decreased their mortality from P. syringae (Hendry, Hunter & Baltrus, 2014). This latter finding might suggest that similar interactions among endosymbiotic and phytopathogenic bacteria may also occur in other host insects (Gonzalez et al., 2016). However, there are currently no studies on the extent of phytopathogen-vector/host associations or the effect of secondary endosymbionts on the interactions between aphids and phytopathogenic bacteria.

CONCLUSIONS

The study presented employing 16S rRNA gene sequencing indicates that the bacterial diversity of the introduced populations of the aphid pests, S. avenae and R. padi, is low. A similar endosymbiont diversity has been reported for both aphid species in their native range. However, variation in the secondary endosymbiont infection could be observed among the introduced and native populations of the aphid S. avenae, indicating that aphid-endosymbiont associations can vary across the geographic range of an aphid species. Our results showed that R. insecticola was the dominant secondary endosymbiont of the introduced populations, while this endosymbiont could be less important in the native range of S. avenae; where H. defensa is the most common endosymbiont reported. Interestingly, the presence of a Pseudomonas sp. closely related to phytopathogenic Pseudomonad species was detected in the aphid samples. As has been observed for other aphids, the detection of Pseudomonas sp. could suggest that aphids can act as a potential vector of phytopathogenic bacteria. However, further studies are necessary to determine the role of aphid species as vectors and/or alternative hosts of important phytopathogenic bacteria.

ADDITIONAL INFORMATION AND DECLARATIONS

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**Author Contributions**
- Francisca Zepeda-Paulo conceived and designed the experiments, performed the experiments, analyzed the data, contributed reagents/materials/analysis tools, prepared figures and/or tables, authored or reviewed drafts of the paper, approved the final draft.
- Sebastián Ortiz-Martínez contributed reagents/materials/analysis tools, authored or reviewed drafts of the paper.
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