Horizontal Gene Acquisition of *Liberibacter* Plant Pathogens from a Bacteriome-Confined Endosymbiont of Their Psyllid Vector

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Introduction

The Asian citrus psyllid *Diaphorina citri* (Hemiptera: Psyllidae) is an important agricultural pest that transmits the phloem-inhabiting alphaproteobacterial *Candidatus Liberibacter asiaticus* and allied plant pathogens, which cause the devastating citrus disease called Huanglongbing or greening disease. *D. citri* harbors two distinct bacterial mutualists in the symbiotic organ called bacteriome: the betaproteobacterium *Candidatus Profftella armatura* in the syncytial cytoplasm at the center of the bacteriome, and the gammaproteobacterium *Candidatus Carsonella ruddii* in uninucleate bacteriocytes. Here we report that a putative amino acid transporter LysE of *Profftella* forms a highly supported clade with proteins of *L. asiaticus*, *L. americanus*, and *L. solanacearum*. *L. crescens*, the most basal *Liberibacter* lineage currently known, lacked the corresponding gene. The *Profftella-Liberibacter* subclade of LysE formed a clade with proteins from betaproteobacteria of the order Burkholderiales, to which *Profftella* belongs. This phylogenetic pattern favors the hypothesis that the *Liberibacter* lineage acquired the gene from the *Profftella* lineage via horizontal gene transfer (HGT) after *L. crescens* diverged from other *Liberibacter* lineages. *Kₜ/Kₛ* analyses further supported the hypothesis that the genes encoded in the *Liberibacter* genomes are functional. These findings highlight the possible evolutionary importance of HGT between plant pathogens and their insect vector’s symbionts that are confined in the symbiotic organ and seemingly sequestered from external microbial populations.

Abstract

The Asian citrus psyllid *Diaphorina citri* is a notorius agricultural pest that transmits the phloem-inhabiting alphaproteobacterial *Candidatus Liberibacter asiaticus* and allied plant pathogens, which cause the devastating citrus disease called Huanglongbing or greening disease. *D. citri* harbors two distinct bacterial mutualists in the symbiotic organ called bacteriome: the betaproteobacterium *Candidatus Profftella armatura* in the syncytial cytoplasm at the center of the bacteriome, and the gammaproteobacterium *Candidatus Carsonella ruddii* in uninucleate bacteriocytes. Here we report that a putative amino acid transporter LysE of *Profftella* forms a highly supported clade with proteins of *L. asiaticus*, *L. americanus*, and *L. solanacearum*. *L. crescens*, the most basal *Liberibacter* lineage currently known, lacked the corresponding gene. The *Profftella-Liberibacter* subclade of LysE formed a clade with proteins from betaproteobacteria of the order Burkholderiales, to which *Profftella* belongs. This phylogenetic pattern favors the hypothesis that the *Liberibacter* lineage acquired the gene from the *Profftella* lineage via horizontal gene transfer (HGT) after *L. crescens* diverged from other *Liberibacter* lineages. *Kₜ/Kₛ* analyses further supported the hypothesis that the genes encoded in the *Liberibacter* genomes are functional. These findings highlight the possible evolutionary importance of HGT between plant pathogens and their insect vector’s symbionts that are confined in the symbiotic organ and seemingly sequestered from external microbial populations.

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Draft genome sequences have been determined for *L. asiaticus* [7], *L. solanacearum* [5], and *L. crescens* [6], whereas draft genome sequence is available for *L. americanus* [8].

In its abdomen, *D. citri* possesses a large yellow symbiotic organ called the bacteriome, where two distinct symbionts are harbored [9]. The betaproteobacterium *Candidatus Profftella armatura* is located in the syncytial cytoplasm at the center of the bacteriome, whilst the gammaproteobacterium *Candidatus Carsonella ruddii* is found in uninucleate bacteriocytes on the surface of the bacteriome. Our previous study revealed that *Profftella* is a toxin-producing defensive symbiont that potentially protects *D. citri* from natural enemies, whilst *Carsonella DC* is a nutritional symbiont that provides the host with essential amino acids, which are scarce in the psyllid’s diet of phloem sap [10].

Here we report that the *Liberibacter* lineage horizontally acquired a putative transporter gene from a bacterium closely related to the extant *Profftella*. 
Materials and Methods

HGT candidates in the Profftella genome were extracted by BLASTP searches [11] against NCBI nr database, using deduced amino acid sequences of all protein coding genes on the Profftella genome as queries. Amino acid sequences were aligned using MAFFT 6.847 [12], followed by manual refinement. Amino acid sites corresponding to alignment gap(s) were omitted from the data set. The best fitting amino acid substitution model for the alignment was estimated using ProtTest3 [13]. For the present analysis, ProtTest selected LG with a gamma distribution (+G), a proportion of invariable sites (+I) and empirical base frequencies (+F) as the best fitting substitution model, followed by WAG with the options +I +G +F. Phylogenetic trees were inferred by the Maximum Likelihood (ML) [14] and the Bayesian Inference (BI) [15] methods. ML trees were constructed using RAxML7.2.1 [16] with LG + G + I + F model. The support values for the internal nodes were inferred by 1,000 bootstrap replicates. In the BI, we used the program MrBayes 3.1.2 [15]. Since the LG model is not implemented in MrBayes, WAG as the next best available model was used with the options +I +G +F. In total, 18,000 trees were obtained (Nruns = 2, Ngen = 900000, Samplefreq = 100), and the first 2,000 of each run were considered as the “burn in” and discarded. The posterior probability of each node was used as the support value of the node. We checked that the potential scale reduction factor was approximately 1.00 for all parameters and that the average standard deviation of split frequencies converged towards zero.

\[ K_S \] and [\( K_A \)] values were calculated as described previously [17].

Statistical significance of the obtained [\( K_S/K_A \)] values was tested against a bootstrap distribution of [\( K_S/K_A \)] values, which was generated by 10,000 bootstrap resamplings of codons from the original alignment. When [\( K_S \)] values calculated from resampled alignments were close to saturation values (larger than 2.0 per site), the [\( K_S \)] values was set as 2.0 for the estimation of [\( K_S/K_A \)].

To analyze the structural organization, the genomic sequences of [\( L. asiaticus \)] strain psy62 [accession no. NC_012985], [\( L. asiaticus \)] strain gxpsy [NC_020549], [\( L. solanacearum \)] CLso-ZC1 [NC_014774], [\( L. americus \)] PW_SP [AOFG01000001-22], and [\( L. crescens \)] BT-1 [NC_019907] were obtained from GenBank.

Results

BLASTP searches against the NCBI nr database demonstrated that the putative LysE protein (accession no: YP_003065395) of [\( P. syringae \)] and [\( L. solanacearum \)] were saturated (\( > 3.00 \)), the [\( K_A \)] values were still as low as 0.42 and 0.39, respectively. These results support the hypothesis that the [\( lye \)] genes of [\( L. solanacearum \)] spp. are under purifying selection and thus are functional.

[Figure 1. Alignment of amino acid sequences of LysEs. Residues conserved in all lineages, three lineages, and two lineages are shaded black, dark gray, and light gray, respectively. doi:10.1371/journal.pone.0082612.g001]
Figure 2. Phylogenetic position of Profftella LysE in related transporter proteins. A total of 185 unambiguously aligned amino acid sites were subjected to the analysis. A maximum likelihood phylogeny is shown, whereas a Bayesian analysis inferred essentially the same result. On each node, support values of maximum-likelihood analysis/Bayesian posterior probabilities are shown. Scale bar indicates substitutions per site. Source organisms are shown with higher bacterial taxa in brackets. α, β, and γ indicate classes of the Proteobacteria, respectively. Accession numbers of proteins are shown in parentheses. The Profftella-Liberibacter cluster is highlighted in red.

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Figure 3. Gene order of lysE flanking regions in the Liberibacter genomes. Pentagons indicate genes with coding directions. Colored pentagons are conserved among the three Liberibacter genomes. Genes with the same color indicate gene clusters. lysE-type protein genes are indicated by arrowheads. Gray pentagons are phage related genes. For L. asiaticus, structural organization of the str. psy 62 genome is shown, whereas that of str. gypsy was essentially the same.

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Discussion

The present study demonstrated that the Liberibacter lineage horizontally acquired a lysE-type transporter gene from the Profftella lineage, an endosymbiont of their vector insect. Ks/Ka analyses further supported the hypothesis that the genes encoded in the Liberibacter genomes are functional. Although their true functions are yet to be identified, LysE superfamily proteins of various bacteria are generally involved in exporting substrates, playing important roles in resistance to toxic substances, in maintenance of optimum intracellular concentration of metabolites, and in excretion of regulatory molecules [20,21]. Thus, it is probable that Liberibacter have acquired novel functions through this HGT. Whereas HGTs are rampant among bacteria [22,23], such transfers of genes are rare in intracellular bacteria that are harbored in insects’ symbiotic organ and are seemingly sequestered from external microbial populations [24,25,26]. Apparently, Profftella, the putative donor lineage of the lysE gene, is this type of endosymbiont. In this context, infection style of Liberibacter, the putative accepter of the gene, would be noticeable. As Liberibacter spp. are transmitted by psyllids in a persistent manner, exhibiting near systemic infection of various organs and tissues [27], they may also intrude into the bacteriome of the vector psyllids, having opportunity of HGT with endosymbionts therein. The present findings highlight the previously unrecognized possible evolutionary importance of HGT between plant pathogens and their vector’s mutualists that are confined in symbiotic organs.

Author Contributions

Conceived and designed the experiments: AN NN. Performed the experiments: AN NN. Analyzed the data: AN NN. Contributed reagents/materials/analysis tools: KO HI MO YH SM MH. Wrote the paper: AN NN TF.

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