Wolbachia in the Culex pipiens Group Mosquitoes: Introgression and Superinfection

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Abstract

Wolbachia bacteria in mosquitoes induce cytoplasmic incompatibility (CI), where sperm from Wolbachia-infected males can produce inviable progeny. The wPip strain in the Culex pipiens group of mosquitoes produces a complexity of CI crossing types. Several factors are thought to be capable of influencing the expression of CI including Wolbachia strain type and host genotype. In this study, the unidirectional CI that occurs between 2 C. pipiens complex laboratory strains, Col and Mol, was further investigated by nuclear genotype introgression. The unidirectional CI between Col and Mol was not found to be influenced by host genetic background, in contrast to a previous introgression study carried out using bidirectionally incompatible C. pipiens group strains. A line containing both wPip strain variants superinfection was also generated by embryonic cytoplasmic transfer. The same crossing type as the parental Col strain was observed in the superinfected line. Quantitative polymerase chain reaction demonstrated a low density of the injected wPipMol variant in the superinfected line after 18 generations, which was considered likely to be responsible for the crossing patterns observed. The Wolbachia density was also shown to be lower in the parental Mol strain males compared with Col strain males, and no inverse relationship between WO phage and Wolbachia density could be detected.

Key words: cytoplasmic incompatibility, Wolbachia
(Laven 1967), although there are questions concerning the sustainability of this control approach. Genetic replacement strategies targeting vector competence could also depend on CI and achieving a better understanding of how complex patterns of incompatibility between infected populations are generated would be important in this respect.

The mechanisms of Wolbachia-induced CI are not well characterized. CI is now generally accepted to involve sperm modification that interferes with the process of karyogamy and a rescue component provided by the *Wolbachia* present in the egg, which restores normal male and female pronuclear karyogamy in compatible crosses (Werren 1997). There are several factors that are thought to influence the expression of CI in insects including *Wolbachia* density (Bordenstein et al. 2006), *Wolbachia* strain type (Sasaki and Ishikawa 2000; Sakamoto et al. 2005), and host genotype (McGraw et al. 2001; Sasaki et al. 2005). The bidirectional CI between the Pel and Bei strains of *C. quinquefasciatus* was previously shown to be influenced by host modifying effects by introgressing the Pel nuclear genome into a Bei cytoplasmic background (Sinkins et al. 2005). In this comparative study, the unidirectional CI that occurs between 2 *wPip* infected *C. pipiens* complex laboratory strains, Mol (*C. molestus* from China) and Col (*C. quinquefasciatus* from Colombia), was investigated using a similar process of nuclear genotype introgression. In addition, a *wPip* strain variant superinfection was generated for the first time using these 2 mosquito strains as donor and recipient, to investigate the effects on CI/crossing type.

**Materials and Methods**

**Mosquito Colonies and Crossing Experiments**

*Culex pipiens* complex laboratory strains Mol (*C. molestus*, China) and Col (*C. quinquefasciatus*, Colombia) were reared using standard mosquito rearing procedures at low larval densities in insectary conditions (26 °C, 70% relative humidity) with a 12:12 h light:dark circadian cycle. Mass crossing experiments were carried out using 50 virgin individuals of each sex. Virgin male and female mosquitoes were obtained through isolation and sexing of pupae. The F$_1$ generation progeny from crosses were analyzed by calculating the percentage of hatched embryos from a minimum of 8 egg rafts, each containing between 50 and 110 eggs per raft, as a measure of the CI phenotype. Female spermathecae were examined for the presence of sperm if the hatch rate was zero to confirm insemination. Unidirectional incompatibility between Mol and Col strains allowed nuclear replacement by crossing Col males to females containing the Mol cytoplasmic background. In the backcrossing experiment, F$_1$ females from the cross between Mol females and Col males were backcrossed with Col males, and offspring females backcrossed with Col males for a further 4 generations. From the third backcross generation, males were crossed with Col females to examine whether crossing type was maintained.

**Discrimination of wPip Strain Variants**

The Mol and Col strains are infected with the *wPip* strain of *Wolbachia*. No sequence polymorphism has been found for *ftsZ* (Guillemaud et al. 1997) or the highly variable *Wolbachia* surface protein (*wp*) genes for the *Wolbachia* present in the Mol and Col strains (Sinkins et al. 2005). Further sequence analysis of *wPip* ankyrin repeat domain (ANK) genes revealed variation in both nucleotide sequence and predicted amino acid sequence for only 2 prophage-associated ANK genes. One of these genes, *pk1*, shows nucleotide sequence variability between the Col and Mol colonies, and discrimination of the *wPip*Mol variant in the Mol colony from the *wPip*Col variant in the Col colony was carried out using *pk1* primers previously described (Sinkins et al. 2005).

**Microinjection of *C. pipiens* Embryos**

Microinjection of *Culex* embryos was carried out using a method developed for injection of *Aedes* embryos (Bossin and Benedict 2005). Donor Mol and recipient Col preblastoderm embryos were aligned against a thin hydrophilic blotting membrane in contact with moist filter paper. Microinjection was carried out under ×100 magnification using a FemtoJet microinjector system (eppendorf) with type II femtotip microinjection needles (eppendorf). A Narishige micromanipulator attached to a Nikon compound microscope was used to manipulate the microinjection needles. After breakage of the needle tip against the membrane, cytoplasm was withdrawn from the donor *wPip*Mol-infected embryos and subsequently injected into the posterior poles of the recipient *wPip*Col-infected embryos. G$_0$ females were mated to colony Col males and blood fed to establish isofemale lines.

**Quantitative Polymerase Chain Reaction**

Quantitative polymerase chain reaction (qPCR) analysis was carried out on 5 individual DNA extracts of *C. pipiens* colony adult mosquitoes. DNA was extracted using a modified version of the Livak buffer method with ethanol precipitation (Collins et al. 1987). Estimation of *Wolbachia* density was undertaken by comparing in vivo gene copy numbers: *ftsZ* gene copy counts were used to estimate the total *Wolbachia* abundance in the DNA extracts. The single copy *C. pipiens* gene was used to normalize the data, controlling for variation in the amount of DNA extracted or mosquito size. In order to estimate the relative density of the *wPip*Mol variant in the superinfected line compared with the parental Mol strain, relative copy numbers of the *wPip*Mol *pk1* gene variant were measured. The *pk1* gene is present in 3 of the integrated prophage copies in the *wPip* (Pel strain) genome sequence (Walker et al. 2007), and copy number was corrected for in the analysis. Bacteriophage WO density was estimated by comparing in vivo copy numbers of the phage capsid *orf7* gene (Masui et al. 2000) to *ftsZ* copy numbers.

Long oligonucleotide standards were designed along with corresponding primer sets, and standards were diluted
from $10^6$ copies to $10^4$ copies for each gene for generation of standard curves. Primer sequences were as follows: qS7F CGGACGCTGATCATCC, qS7R ATCGCTCCGAAAAGGAGATGCTA (standard = 120 bp); qftsZF TGG TTGGAAGGGTGACAG, qftsZR CCAGTACCCACCAACCATACC (ftsZ standard = 102 bp); qpk1MolF ATTT- TGCGACTGTTAATTGGA, qpk1MolR CATCGGTGTCCGATTTCGTG (standard = 101bp); qorf7F AAGTACCACTGATAAGATTATTT, qorf7R GCCAAAATATAGACCTGTT (standard = 98 bp).

The relative quantity of template DNA was estimated using an Opticon 2 Continuous Fluorescence Detection System (GRI) together with QuantiTect SYBR Green PCR Kit (Qiagen, Valencia, CA). Template DNA was PCR amplified using primers at 10 μM, and 1.0 μl of DNA was added in a total volume of 20 μl per reaction. qPCR cycling conditions consisted of an initial denaturation step of 95 °C for 15 min followed by 40 cycles of denaturing at 95 °C (15 s) and primer annealing at 50 °C (30 s). A melting curve was analyzed to check for any nonspecific amplification or primer dimers. Included in the assays were DNA extracts from the Wolbachia-uninfected Pel U colony and sterilized water controls to confirm the absence of contaminating DNA.

Results and Discussion

Crossing and Introgression

Crossing experiments between a PipMol-infected C. p;piens complex laboratory strains Mol and Col previously revealed unidirectional incompatibility (Walker et al. 2007). Col females mated to Mol males resulted in 0.44 ± 0.1% embryo hatch in contrast to 93.49 ± 1.17% hatch when Mol females mated with Col males. To investigate whether host modifying effects could influence this crossing pattern, as was previously observed in a cross between 2 bidirectionally incompatible Culex strains (Sinkins et al. 2005), 5 generations of backcrossing, introgressing the Col nuclear genotype into Mol (Sinkins et al. 2009:100(2)), was undertaken as shown in Table 1.

As shown in Table 1, close to complete incompatibility was observed when males from the introgressed line were crossed with Col females, showing no evidence for a host modifying effect in this particular example of CI. Host modifying effects on CI are thus not universal in this complex but dependent on the particular host strain variant combination being introgressed, and this pattern may contribute to the complexity of crossing type variation in the complex.

Introgression and replacement of the Col nuclear genotype into a PipMol-infected cytoplasm also confirm that the interstrain incompatibilities between Mol and Col are not controlled by host nuclear divergence but by Wolbachia. This introgression experiment has in effect created a novel crossing type in C. quinquefasciatus because the C. molestus nuclear genome has been largely replaced. The a PipMol variant would be expected to be able to spread through populations of C. quinquefasciatus of the Col crossing type

<table>
<thead>
<tr>
<th>Cross/backcross, female × male</th>
<th>Progeny name</th>
<th>Test cross, female × male</th>
<th>% hatch (no. rafts/embryos)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mol × Col</td>
<td>F1</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>F1 × Col</td>
<td>BCI</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>BCI × Col</td>
<td>BCI</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>BCI × Col × BCI</td>
<td>BCI</td>
<td>Col × BCI</td>
<td>0 (48/6427)</td>
</tr>
<tr>
<td>CIV × Col</td>
<td>BCI</td>
<td>Col × BCIIV</td>
<td>0 (42/5743)</td>
</tr>
<tr>
<td>BCI × Col</td>
<td>BCI</td>
<td>Col × BCI</td>
<td>0.65 (22/2780)</td>
</tr>
</tbody>
</table>

F1 females from the cross between Mol females and Col males were backcrossed with Col males, and offspring females backcrossed with Col males for a further 4 generation. From the third backcross generation (BCIII), males were crossed with Col females and embryo hatch rates counted. The resulting line was fully anautogenous (requiring a blood meal for egg laying, unlike the autogenous C. molestus).

wPip Strain Variant Superinfection

In order to create a line containing both the a PipMol and a PipCol variants, Col embryos were injected with a PipMol-infected preblastoderm embryo cytoplasm; of 350 injected embryos, there was a 15.1% hatch rate and 8.9% survival of injected embryos to eclosion (31/350). To detect the presence of the a PipMol variant in the Col G0 females and resulting isofemale lines, PCR analysis was carried out using primers that were designed to discriminate between variants of the WO prophage-associated pk1 gene present in the 2 a Pip strains (Sinkins et al. 2005). An isofemale line, termed a PipCol(a PipMol), was successfully established and maintained.

Crossing experiments were undertaken with generation 11 (G11) of the superinfected a PipCol(a PipMol) line to analyze the effect on CI crossing type. PCR analysis for the pk1Mol variant confirmed the presence of the a PipMol variant in all 5 G11 males assayed, suggesting no heterogeneity in the line. As shown in Figure 1, the very low embryo hatch resulting from crossing a PipCol(a PipMol) females to a PipMol males was similar to the low hatch when a PipCol females were crossed to a PipMol males. The presence of the a PipMol variant in the a PipCol(a PipMol) females would be expected to result in compatibility with Mol males, based on the model that the females contain the correct “rescue” factor for both singly and superinfected males (Sinkins et al. 1995). However, crossing analysis revealed that the a PipMol variant in females of the superinfected line was unable to rescue the modification factor in sperm produced by a PipMol males. Likewise, crossing the a PipCol(a PipMol)-infected G11 males with a PipCol-infected females produced a high hatch rate, very similar to the hatch seen when a PipMol females were crossed to a PipCol males. Thus, the presence of the a PipMol variant did not produce CI in this cross as had been expected.
and 50 virgin females (F), and the hatch rate of the F1 progeny was calculated from a minimum of 10 egg rafts, each containing 50–120 eggs per raft, as a measure of CI.

Crossing experiments were performed with 50 virgin males (M) and 50–120 eggs per raft, as a measure of CI.

Density of wPip Strain Variants

A possible explanation for the crossing data described is that the wPipMol variant did not reach a sufficiently high density in the superinfected line to be able to induce CI when superinfected males were crossed to Col females or to rescue CI when superinfected females were crossed to Mol males. qPCR density assays were carried out on the G18 generation to allow a stable equilibrium density to be reached. Although all individual G18 males assayed were positive for the wPipMol based on qPCR followed by gel electrophoresis, a very low comparative density of the wPipMol pk1 variant was detected by qPCR as shown by the pk1Mol:ftsZ gene ratios, equivalent to 1:250 (Table 2). Although naturally occurring Wolbachia strain superinfections such as that seen in Aedes aegypti can be stable despite significant density differences between strains (Dutton and Sinkins 2004), the density differences between them were lower than those seen here.

The quantitative PCR assay is expected to amplify both integrated WO prophage copies of the pk1 genes in the Wolbachia genome and any lytic phage particles containing phage DNA that may be produced from those copies containing pk1. Thus, an alternative explanation is that the wPipMol variant may have infected a small subset of the wPipCol variant Wolbachia, rather than the transfer and transmission of whole wPipMol bacteria. There are no other markers outside of the phage regions available that can discriminate between these strains, if indeed any such markers exist. However, we believe this alternative hypothesis to be unlikely, based on a set of inter-Culex microinjection experiments using extracts filtered to exclude whole Wolbachia but not any phage particles, which did not result in any detectable conversions of pk1 type (data not shown). In any case, the same hypothesis would apply whether whole bacteria or just phage particles have in fact been transferred—that the titre reached was probably not high enough for crossing type conversion to occur—and either way the resulting line can be regarded as superinfected.

Table 2. qPCR analysis to estimate the wPipMol density in the superinfected wPipCol(wPipMol) line

<table>
<thead>
<tr>
<th>Strain/line</th>
<th>Overall wPip density (ftsZ: S7)</th>
<th>wPipMol relative density (pk1Mol:ftsZ)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mol</td>
<td>0.12 ± 0.04</td>
<td>1.278 ± 0.189</td>
</tr>
<tr>
<td>Col</td>
<td>0.39 ± 0.11</td>
<td>0 ± 0</td>
</tr>
<tr>
<td>wPipCol(wPipMol) G18</td>
<td>0.42 ± 0.12</td>
<td>0.004 ± 0.001</td>
</tr>
</tbody>
</table>

Wolbachia and Bacteriophage WO Density in Parental Col and Mol Strains

As shown in Table 2, the Wolbachia density in Mol males was significantly lower than for Col males (P = 0.03, Student’s t-test). The wPipMol variant may have a slower growth rate compared with wPipCol, and this could be responsible for it being unable to establish itself to a sufficiently high density when indirect intrahost competition with wPipCol. Mol males carrying the lower density wPipMol variant are incompatible with females carrying wPipCol (Walker et al. 2007). This pattern is the opposite of what would be expected if density differences were causally responsible for the different patterns of CI observed between these strains. The density of Wolbachia in C. pipiens testes was also found to be strain dependent and did not appear to influence CI in other C. pipiens strains (Duron et al. 2007).

A threshold level of Wolbachia density (and/or even distribution within the testes) may be required for induction of complete CI. However, if Wolbachia density differences influence the penetrance of CI only when the density falls below a threshold level, the varying wPip densities between the Mol and Col strains may all be above this threshold and therefore irrelevant with respect to CI induction/rescue. However, when the Wolbachia variants are superinfecting the same host, then density of the wPipMol variant may have fallen below this necessary threshold due to effects such as direct competition between strains.

The overall density of bacteriophage WO was also assessed in Col and Mol, measured as a ratio of the phage capsid orf7 gene copy number to Wolbachia ftsZ copy number (because not all prophage copies contain pk1), and was not significantly different between Mol males (8.04 ± 2.42) and Col males (7.07 ± 2.18) (P = 0.43, Student’s t-test). The densities of bacteriophage WO in males were not inversely correlated with Wolbachia density (Pearson r = −0.35) with no evidence for increased bacteriophage WO lytic activity in Mol males. These data suggest that the inverse relationship between phage density and Wolbachia density, and tripartite association as an explanation for unidirectional CI presented.
for *Nasonia* by Bordenstein et al. (2006), do not apply in this case. The observed phage copy numbers per *Wolbachia* are broadly consistent with the 5 prophage regions containing the *wPip* (Pel strain) genome, implying that bacteriophage *WO* is primarily temperate rather than lytic in *wPip*.

The development of intraspecific transfer of *wPip* strain variants by microinjection, coupled with the development of specific PCR assays to separate these variants and estimate their density (Sinkins et al. 2005; Walker et al. 2007) provide useful tools to investigate the control of CI in the *C. pipiens* complex. Future experiments can now be contemplated within the *Culex* genus involving other *wPip* strains with more similar growth rates/relative densities; the effects of introducing novel *Wolbachia* strains from other species can also be investigated. Meanwhile, the replacement of host genotype by introgression has provided a novel crossing type in *C. quinquefasciatus*, furthering understanding of CI in the complex and potentially providing a platform for CI-based mosquito control strategies.

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**References**


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