Sustained Wolbachia-mediated blocking of dengue virus isolates following serial passage in Aedes aegypti cell culture

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Abstract

Wolbachia is an intracellular endosymbiont of insects that inhibits the replication of a range of pathogens in its arthropod hosts. The release of Wolbachia into wild populations of mosquitoes is an innovative biocontrol effort to suppress the transmission of arthropod-borne viruses (arboviruses) to humans, most notably dengue virus. The success of the Wolbachia-based approach hinges upon the stable persistence of the 'pathogen blocking' effect, whose mechanistic basis is poorly understood. Evidence suggests that Wolbachia may affect viral replication via a combination of competition for host resources and activation of host immunity. The evolution of resistance against Wolbachia and pathogen blocking in the mosquito or the virus could reduce the public health impact of the symbiont releases. Here, we investigate if dengue 3 virus (DENV-3) is capable of accumulating adaptive mutations that improve its replicative capacity during serial passage in Wolbachia wMel-infected cells. During the passaging regime, viral isolates in Wolbachia-infected cells exhibited greater variation in viral loads compared to controls. The viral loads of these isolates declined rapidly during passaging due to the blocking effects of Wolbachia carriage, with several being lost all together and the remainder recovering to low but stable levels. We attempted to sequence the genomes of the surviving passaged isolates but, given their low abundance, were unable to obtain sufficient depth of coverage for evolutionary analysis. In contrast, viral loads in Wolbachia-free control cells were consistently high during passaging. The surviving isolates passaged in the presence of Wolbachia exhibited a reduced ability to
replicate even in Wolbachia-free cells. These experiments demonstrate the challenge for dengue in evolving resistance to Wolbachia-mediated blocking.

Key words: dengue virus; Wolbachia; Aedes aegypti; evolution.

1. Introduction

The endosymbiont Wolbachia is being released into wild populations of the mosquito vector Aedes aegypti throughout the tropics as a strategy to reduce the transmission of viruses to humans (Hoffmann et al. 2011; Walker et al. 2011). In Ae. aegypti infected with Wolbachia, the replication of DENV—as well as other medically important positive single-stranded RNA viruses such as chikungunya, Zika, and yellow fever—is reduced, leading to the potential interruption in pathogen transmission (Walker et al. 2011; Van den Hurk et al. 2012; Dutra et al. 2016). A successful deployment in Townsville, Australia, has so far resulted in persistently high infection frequencies across the local mosquito population and elimination of domestic dengue transmission (O'Neill et al. 2018). Other studies of efficacy are underway in Southeast Asia and Latin America. The long-term stability of Wolbachia-mediated pathogen blocking in the field will be dependent on the complex selection pressures acting via the genomes of the three relevant organisms: Wolbachia, DENV, and the mosquito. Evolution of resistance either against Wolbachia or its effects in the viral or mosquito genomes could lead to reduced efficacy.

The mechanistic nature of Wolbachia-mediated blocking is currently poorly defined. The originally proposed putative mechanism was that of innate immune priming, whereby the presence of Wolbachia heightens the basal immune response of the insect, improving the insect’s ability to control any secondary pathogen it encounters. This theory originated from transcriptional profiling data in the mosquito demonstrating that Wolbachia induces transcription of the canonical innate immune pathways (Kambris et al. 2009, 2010; Moreira et al. 2009). Later work, including some comparative studies in the naturally infected Drosophila melanogaster and in Ae. aegypti (Rances et al. 2012; Terradas, Joubert, and McGraw 2017), demonstrated that the priming of these core pathways could not explain the entirety of the effect. Other studies have found evidence of competition between the symbiont and viruses for intracellular space (Moreira et al. 2009; Rainey et al. 2016) and cholesterol (Caragata et al. 2013), and still others cite the involvement of reaction oxygen species (Pan et al. 2012; Wong, Brownlie, and Johnson 2015) or Wolbachia-directed control of host gene expression by small RNA production (Zhang et al. 2013; Mayoral et al. 2014; Asad et al. 2018). Commonly, the strength of blocking is positively correlated with Wolbachia density (Osborne et al. 2009, 2012; Frentiu et al. 2010; Walker et al. 2011; Lu et al. 2012; Chrostek et al. 2013; Amuzu and McGraw 2016).

In wild mosquito populations where Wolbachia have been released for biocontrol and subsequently spread to high frequencies, a virus will be under strong pressure to increase its replicative fitness in the presence of the symbiont (Bull and Turelli 2013). Replication of the small, positive, single-stranded RNA genome of DENV is highly error prone, as it depends upon an RNA-dependent RNA polymerase that lacks proof-reading ability. As a consequence, the population of DENVs within a single host (human or insect) is highly genetically diverse and likely phenotypically heterogeneous (Wang et al. 2002; Parameswaran et al. 2017). Regardless, Bull and Turelli (2013) predicted that adaptive mutations allowing the virus to completely overcome Wolbachia-induced blocking would be difficult to achieve given the likely multifaceted nature of blocking mechanisms (Johnson 2015), coupled with the constraints of strong purifying selection necessarily resulting from replication in both vertebrate and invertebrate cells (Vasilakis et al. 2011).

Recent work has suggested that Wolbachia interference occurs at the level of intracellular replication and assembly (Welsch et al. 2009; Zhang et al. 2013; Rainey et al. 2016; Bhattacharya, Newton, and Hardy 2017; Asad, Parry, and Asgari 2018; Asad et al. 2018). Hence, any adaptive mutations in viral proteins that improve efficiency at these stages may represent an evolutionary solution to resistance against Wolbachia. Improved functions of certain viral proteins could also relax any competition between Wolbachia and DENV for host resources, such as lipids and intracellular space, reducing the impact of the blocking phenotype. For example, the structural capsid protein primarily packages the viral genome and acts as a chaperone to promote correct folding, but it has also been reported to bind to lipid droplets (Samsa et al. 2009; Byk and Gamarnik 2016). Lipid droplets are intracellular vesicles containing triacylglycerides and cholesterol (Ducharme and Bickel 2008). The presence of DENV capsid proteins increases the abundance of lipid droplets in infected cells (Samsa et al. 2009). Although the underlying mechanism is unknown, increased lipid droplet enrichment by the capsid protein could lead to a higher availability of lipid sources of energy for viral replication. Furthermore, DENV nonstructural proteins are involved in assembling membrane-bound viral replication complexes, derived from the endoplasmic reticulum (ER) and enriched in cholesterol (Muller and Young 2013; Xie et al. 2017). As Wolbachia and DENV are both intimately associated with and compete for ER membrane structures (Welsch et al. 2009; White et al. 2017), changes in viral nonstructural proteins that improve the efficiency of replication complex formation may help to overcome this competition. Given the complex nature of blocking mechanisms, adaptation may only be possible through multiple beneficial mutations in viral protein encoding genes.

A variety of studies have attempted to model the coevolution of this mosquito-symbiont-pathogen system (Lively et al. 2005; Vavra and Charlat 2012; Bull and Turelli 2013), but none have empirically examined the adaptive potential of DENV in the presence of Wolbachia. In this work, we sought to test if adaptive evolution in DENV can be experimentally induced in vitro in response to the presence of Wolbachia. To achieve this, DENV-3 viral isolates were serially passaged in three distinct Ae. aegypti cell lines, namely (1) a naïve Wolbachia-free line, (2) a tetracycline-cured Wolbachia-free line, and (3) a Wolbachia-infected line. During the regime, we transferred only viruses that were able to successfully infect the cell types, replicate, and escape into the surrounding medium. We expected to produce virus isolates that, after accumulating adaptive mutations, exhibited increased replicative fitness relative to both the ancestral virus and those isolates passaged in Wolbachia-free cells. We then compared the replicative fitness of the ancestral and passaged virus isolates in parallel in all three cell types and attempted to characterize the sequence diversity generated in
the DENV isolates during the selection regime. Our results speak to the potential for DENV to adapt in response to Wolbachia in the short-term.

2. Materials and methods

2.1 Aag-2 cell lines

Three lines of Aag-2 cells were used in the virus evolution experiment: Aag-2, Aag-2Tet, and Aag-2uMel. The Aag-2 cell lineage was derived from Ae. aegypti embryos and is susceptible to arthropod-borne virus (arbovirus) infections (Peleg 1968). Aag-2 cells naturally do not carry any Wolbachia infection. Aag-2uMel cells were generated through transfection of the uMel Wolbachia strain isolated from its natural host, D. melanogaster (Terradas, Joubert, and McGraw 2017). To create the Aag-2Tet cell line, Aag-2uMel cells were cured of Wolbachia infection through treatment with the antibiotic tetracycline, as described in Terradas, Joubert, and McGraw (2017). All three cell lines were routinely maintained at 25 °C on complete medium consisting of equal parts Schneider’s medium and Mitsubishi and Maramorosch medium and supplemented with 10 per cent (v/v) heat-inactivated fetal bovine serum (FBS) (Life Technologies), 1× GlutaMAX (Gibco), and 20 mM HEPES (Sigma-Aldrich, St Louis, MO, USA). To propagate sufficient virus for this experiment, monolayers of C6/36 cells in two T175 flasks at 80 per cent confluency were inoculated with DENV-3. Inoculated cells were then incubated in medium supplemented with 2 per cent (v/v) FBS (Life Technologies). At seven days post-inoculation, clarified cell culture medium from the two flasks were collected and pooled. The pooled virus concentration was 10⁶ genome copies per ml, as quantified via methods described below. This ancestral virus (passage 0, P0) was then stored at −80 °C in single-use aliquots for use in initial inoculation of the sequential passaging regime, repeated blocking tests, and replicative fitness assays.

2.2 Virus

For this study, we used a DENV-3 isolated from Cairns (Queensland, Australia), during an outbreak in 2008/2009 (Ritchie et al. 2013). The strain had been passaged ten times post isolation in C6/36 Aedes albopictus cells maintained at 25 °C on RPMI 1640 medium supplemented with 10 per cent (v/v) FBS (Life Technologies), 1× GlutaMAX (Gibco), and 20 mM HEPES (Sigma-Aldrich, St Louis, MO, USA). To propagate sufficient virus for this experiment, monolayers of C6/36 cells in two T175 flasks at 80 per cent confluency were inoculated with DENV-3. Inoculated cells were then incubated in medium supplemented with 2 per cent (v/v) FBS (Life Technologies). At seven days post-inoculation, clarified cell culture medium from the two flasks were collected and pooled. The pooled virus concentration was 10⁶ genome copies per ml, as quantified via methods described below. This ancestral virus (passage 0, P0) was then stored at −80 °C in single-use aliquots for use in initial inoculation of the sequential passaging regime, repeated blocking tests, and replicative fitness assays.

2.3 Virus passaging

Six-well cell culture plates were seeded with Aag-2, Aag-2Tet, and Aag-2uMel cells at a density of 1 × 10⁶ cells per well. Each well was inoculated with 2 ml of P0 and passed five days later. In total, twelve virus lines were generated and transferred for nine sequential passages with five days of growth in between: three passaged in Aag-2 cells, three in Aag-2Tet cells, and six in Aag-2uMel cells. Twice as many Aag-2uMel lines were passaged to account for potential loss of virus due to complete blocking by Wolbachia. At each sequential passaging, cell supernatants were collected and clarified via centrifugation at 200 × g for 10 min. A total of 1 ml of this viremic supernatant was transferred to a fresh set of uninfected cells, followed by an addition of 800 μl of complete medium containing 2 per cent (v/v) FBS per well. The remaining supernatants were stored at −80 °C for later RNA extraction and sequencing. At every passage, Wolbachia was quantified and the uninfected status of Aag-2 and Aag2Tet cells was verified using the methods below. The DENV-blocking capability of the uMel cells relative to Aag-2 and Aag2Tet cells was also confirmed at each passage in a parallel assay by inoculating all three cell lines with P0. At five days post-inoculation, supernatants from this blocking test and from passaged isolates were sampled, and DENV load was quantified using methods described below.

2.4 Wolbachia density quantification

A multiplex quantitative real-time polymerase chain reaction (qRT-PCR) was used to detect the presence of Wolbachia and quantify its density in our cell lines. This method has been previously described in Ferguson et al. (2015). In our study, all qRT-PCR assays were performed with a LightCycler 480 instrument II (Roche, Basel, Switzerland). Sample DNA was prepared by resuspending cell pellets in 50 μl of extraction buffer (10 mM Tris pH 8.2, 1 mM EDTA, 50 mM NaCl, and 1.25% (v/v) proteinase K [Bioline, Memphis, TN]) and incubating this suspension in a thermal cycler for 5 min at 56 °C, 5 min at 95 °C, and held at 4 °C (Yeap et al. 2014). A total of 1 μl of DNA sample was used in a single qRT-PCR reaction combined with 5 μl of 2× LightCycler 480 probes master mix (Roche), 0.25 μM of rps17 forward and reverse primers, 0.1 μM of rps17 probe, 0.3 μM of W0513 forward and reverse primers and probes, and RNase-free water to make up a total volume of 10 μl. Primer and probe sequences are shown in Supplementary Table S1. Cycling conditions were as follows: initial denaturation at 95 °C for 5 min; 45 cycles of amplification consisting of 95 °C for 10 s, 60 °C for 15 s, and 72 °C for 1 s; and a cooling step at 40 °C for 10 s. Each sample was run in triplicate. Relative Wolbachia density was calculated as the ratio of the Wolbachia gene W0513 to the housekeeping Ae. aegypti ribosomal protein gene rps17 by using the Q-gene method (Simon 2003).

2.5 Fluorescent in situ hybridization for Wolbachia

The presence of Wolbachia in cells was also validated through fluorescent in situ hybridization (FISH) at every passage of the virus evolution experiment. Aag-2, Aag-2Tet, and Aag-2uMel cells were seeded onto Nunc Lab-Tek 8-well chamber slides (Thermo Fisher Scientific, Waltham, MA, USA). When the cells were confluent, the cell culture medium was removed and cells were fixed, hybridized with rhodamine-labeled probes specific to Wolbachia 16S rRNA and DAPI, then mounted as per previously published methods (Moreira et al. 2009). Images were captured using a Leica DM compound fluorescent microscope with a Leica DC300 camera through the appropriate channels, keeping exposure times uniform across samples.

2.6 Viral quantification

The viral genome copy number in cell medium supernatant was quantified using TaqMan fast virus 1-step master mix (Applied Biosystems, Foster City, CA, USA). For every supernatant sample to be quantified, 10 μl of supernatant was mixed with extraction buffer in a 1:1 ratio and incubated as per above. Each qRT-PCR reaction consisted of 2.5 μl of 4× master mix, 0.5 μM of DENV forward and reverse primers, 0.25 μM of probe (Warrilow et al. 2002), 2 μl of viral RNA sample in extraction buffer, and RNase-free water to make up a total volume of 10 μl. Primer and probe sequences are shown in Supplementary Table S1. Cycling conditions were as follows: reverse transcription at 50 °C for 10 min; denaturation at 95 °C for 20 s; 50 cycles of amplification consisting of 95 °C for 3 s, 60 °C for 60 s, and 72 °C for 1 s; and a cooling
at 40 °C for 10 s. Absolute DENV-3 copy numbers were obtained by extrapolating to a standard curve of DENV genome copies ranging from 10^8 to 10^10 in a tenfold serial dilution, created as described in Ye et al. (2014). Each sample was run in triplicate. Copy number was expressed as genome copies per μl of supernatant. Loads were compared using analysis of variance (ANOVA) and post hoc tests performed on log-transformed values with GraphPad Prism 7 software (San Diego, CA, USA).

2.7 Replicative fitness assay
After experimental evolution through sequential passaging, each of the twelve virus lines was assayed for their replicative fitness in Aag-2, Aag-2Tet, and Aag-2uMel cells (n = 5 per isolate per cell line). Ninety-six-well plates were seeded with 5 × 10^4 cells per well with 200 μl of complete medium with 10 per cent (v/v) FBS (Life Technologies) and incubated for two days prior to inoculation. Cells were washed once with 200 μl of sterile 1× PBS before being inoculated with 1.5 × 10^4 genome copies per well, followed by incubation for 2 h at 25 °C with rocking at every 15 min. Viral inoculants were then removed and 200 μl of complete medium with 2 per cent (v/v) FBS (Life Technologies) was added to each well. Post-inoculation, 10 μl of cell medium was sampled every day for seven days. Virus accumulation curves were obtained by sampling 10 μl of cell medium every day for seven days and quantifying number of virus genome copies via qRT-PCR as per methods described above.

2.8 RNA isolation and sequencing
Material was collected for RNA sequencing at generations 0, 5, and 9 for the uMel-infected and tetracycline-cured control cells. Viral RNA in the cell medium supernatant was extracted using the TRIzol LS reagent (Invitrogen, Carlsbad, CA, USA) according to manufacturer’s protocol. Each sample of viral RNA was solubilized in 44 μl of RNase-free water. To each sample, 1 μl of DNase I recombinant enzyme (Roche) and 5 μl of buffer was added and incubated at 37 °C for 50 min to ensure no contamination of genomic DNA. To clean up the isolated RNA for sequencing use, we used an RNase kit (Qiagen, Hilden, Germany) according to the manufacturer’s protocol and eluted purified RNA into 30 μl of RNase-free water. Samples were prepared with the Illumina whole-transcriptome library prep kit with Ribo-Zero Gold rRNA depletion, and sequencing was performed on the Illumina NextSeq 500 platform at the Australian Genome Research Facility (AGRF).

2.9 Sequence data analysis
Illumina paired-reads were assembled in Geneious 11.1.3 (https://www.geneious.com, last accessed 12 January 2019) to the DENV-3 reference genome (GenBank accession number JN406515). Poor quality reads were removed. The resulting assemblies were manually inspected for genome coverage and potential genetic variations within the virus population. Notably, 54,972,167 reads mapped to the reference DENV-3 genome for generation 0, with an average coverage of 375,511 reads per nucleotide position. No genetic difference was observed between the sequenced sample and the reference sequence. Thus, the DENV-3 sequence was suitable to be used as reference for generations 5 and 9. However, read count and, thus, coverage, for these two generations was very low: less than 300 reads assembled in total for either generation. This reduced the median coverage to two reads per nucleotide position across the genome, although parts of the genome were not covered with any reads. Such limited coverage prevented analysis of virus genetic diversity or evolution in the genomes of the Aag-2uMel-passaged isolates, and thus isolates from Wolbachia-free control lines were not sequenced. PCR amplification of the genome to increase material for sequencing was not considered as the very low viral load would make amplification of full genomes challenging. Additionally, we mapped reads to the cell fusing agent virus (CFAV) reference genome to check for its presence as this insect-specific flavivirus is known to persistently infect Aag2 cells and enhance DENV infection (Stollar and Thomas 1975; Zhang et al. 2017). Assembly was conducted as described for DENV-3 and also using Bowtie 2 v2.3.4.3 (Langmead et al. 2018).

3. Results
3.1 Viral loads throughout serial passaging
During serial passaging, Aag-2- and Aag-2Tet-passaged isolates showed little variation in viral loads, whereas variation among the viral loads of Aag-2uMel-passaged isolates increased dramatically with higher passage numbers (Fig. 1A). Viral copy number of isolates passaged in Aag-2uMel cells showed a decreasing trend from P1 but began to stabilize from P4 onward. At P10, three of these isolates were lost, whereas three others (isolates 7, 11, and 12) remained at 10^10 genome copies per μl. On average, the loads of Aag-2uMel-passaged isolates at the end of serial passaging were 3–4 log_{10} -folds lower than the levels of isolates passaged in Wolbachia-free cells.

To verify that the Wolbachia-induced blocking phenotype was present throughout the experiment, cells used at each passage were challenged in parallel with an aliquot of P0 at multiplicity of infection (MOI) of 0.1, and the viral copy number present in the supernatant was quantified at five days post-inoculation (Fig. 1B). Throughout the passaging regime, DENV-3 genome copies in Aag-2uMel cells were always lower than those in Aag-2 or Aag-2Tet cells by at least one log_{10} -fold, confirming the effect of the pathogen blocking phenotype (P < 0.0001, Tukey’s multiple comparison test). The reduced loads in Aag-2Tet cells relative to Aag-2 cells can be explained by adverse side effects from tetracycline treatment on mitochondrial function, potentially dampening viral replication efficiency in these cells (Moullan et al. 2015). Wolbachia densities were quantified via qRT-PCR at every passage (Fig. 1C). While they vary substantially, high infection frequencies throughout were validated through FISH (Supplementary Fig. S1).

3.2 Replicative fitness following selection
To detect fitness gains as a result of serial passaging, we assessed the replicative fitness of the nine virus isolates that remained at P10 in Aag-2, Aag-2Tet, and Aag-2uMel cells. As the viral copy number of Aag-2uMel-passaged isolates (isolates 7, 11, and 12) was much lower than that of Aag-2- or Aag-2Tet-passaged isolates (isolates 1–6), all inocula were standardized to 1.5 × 10^4 virus copies per 5 × 10^4 cells for all isolates in this assay—the highest virus copy-to-cell ratio possible while maintaining consistency across all isolates. We compared the replication kinetics of passaged isolates to the ancestral virus in all three cell lines used in this study to detect any fitness gains obtained from serial passaging. Supernatants were collected at two-day intervals during the fitness assay, but due to the lack of virus replication in the Aag-2uMel cells, only samples collected at six days post-infection were quantified via qRT-PCR.
We did not find evidence of an increase in the replicative fitness relative to P₀ in any of our passaged isolates, even when reinoculated onto the cell line in which they had been serially passaged. In Aag-2 and Aag-2Tet cells, the replication of Aag-2-passaged isolates was comparable to the P₀ (Fig. 2A and B). Interestingly, Aag-2Tet-passaged isolates showed decreased replicative ability compared to P₀ (\( P < 0.001 \), Dunnett’s multiple comparison test) (Table 1), with the exception of isolate 6 in Aag-2Tet cells. In Aag-2uMel cells, none of the passaged isolates produced detectable viral copies (Fig. 2C).

The Aag-2uMel-passaged isolates performed poorly in all tested cell lines. Very few biological replicates produced detectable viral loads, all of which were significantly lower than those of P₀ (\( P < 0.0001 \), Dunnett’s multiple comparison test) (Table 1). Data points below the qRT-PCR detection limit were excluded from statistical analysis. The poor ability of these isolates to
replicate in either Wolbachia-free cell line was surprising, as Aag-2- and Aag-2Tet-passaged isolates were able to produce replicated virions from similar inoculum concentrations.

However, we believe the few Aag-2wMel-passaged replicates with detectable viral loads (Fig. 2A and B) consisted of true products of replication during the assay and not remnants of the inoculum, as the supernatants of all replicates at one-day post-inoculation did not contain detectable numbers of genome copies (data not shown).

3.3 DENV-3 sequence diversity generated during the selection regime

Post-selection we prepared the following set of samples for sequencing; the ancestral P₀ isolate and isolates 7, 11, and 12 (from Aag-2wMel cell lines) at both P₅ and P₀. The resulting depth of sequencing coverage was very low in the passaged isolates: the total number of reads mapped to the DENV-3 genome averaged ~290 across the passaged lines, compared to the ancestral isolate with ~55,000. Approximately 80 per cent of the genome was only represented by 1× read coverage in the passaged isolates. Hence, these data reveal the rarity of virus in the culture and necessarily prevented statistical analysis of shifts in allele frequencies relative to the ancestor required to demonstrate evolutionary change. Thus, it is inconclusive whether evolution occurred in Aag-2wMel-passaged isolates. To ensure that CFAV, which persistently infects Aag2 (Stollar and Thomas 1975), is not confounding DENV infection, we also assembled reads against CFAV reference genome. No contigs covering the CAFV genome were found, indicating that CFAV likely absent from these isolates.

Table 1. Summary of one-way ANOVAs and multiple comparisons tests comparing the replicative fitness of each passaged isolate to that of P₀.

| Test cell line | ANOVA P-value | Multiple comparisons test | P-value |
|----------------|---------------|---------------------------|---------|
| Aag-2          | <0.0001       | P₀ vs. isolate 1, n.s.    |         |
|                |               | P₀ vs. isolate 2, n.s.    |         |
|                |               | P₀ vs. isolate 3, n.s.    |         |
|                |               | P₀ vs. isolate 4, 0.0001  |         |
|                |               | P₀ vs. isolate 5, 0.0001  |         |
|                |               | P₀ vs. isolate 6, 0.0003  |         |
|                |               | P₀ vs. isolate 7, 0.0001  |         |
|                |               | P₀ vs. isolate 12, 0.0001 |         |
| Aag-2Tet       | <0.0001       | P₀ vs. isolate 1, n.s.    |         |
|                |               | P₀ vs. isolate 2, n.s.    |         |
|                |               | P₀ vs. isolate 3, n.s.    |         |
|                |               | P₀ vs. isolate 4, 0.0004  |         |
|                |               | P₀ vs. isolate 5, 0.0001  |         |
|                |               | P₀ vs. isolate 6, n.s.    |         |
|                |               | P₀ vs. isolate 7, 0.0001  |         |

n.s., not significant.

Figure 2. Replicative fitness assay of the nine remaining isolates following sequential passaging. Log₁₀-transformed viral genome copy numbers in the supernatants of (A) Aag-2, (B) Aag-2Tet, and (C) Aag-2wMel cells were quantified via qRT-PCR at six days post-inoculation by each isolate. Each data point represents one biological replicate. Data points at 0 log₁₀ copies/μl indicate that viral copies were below detection limits at ten genome copies/μl (dotted line) and were excluded from data analysis. Mean and SEM are shown in graphs (n = 5 per isolate).
4. Discussion

The higher variation in viral load in Aag-2uMel-passaged isolates relative to those passaged in Wolbachia-free lines may reflect an underlying variation in fitness or random noise, which may in turn be linked to the substantial population bottlenecks during passage. A similar phenomenon was observed in foot-and-mouth-disease virus in which sequential population bottlenecks were experimentally induced during plaque-to-plaque transfers (Lazaro et al. 2003). This notwithstanding, the failure of viral copy numbers to revive substantially after their initial decline when passaged in Aag-2uMel cells, the eventual loss of three of these isolates at P9, and the very low sequence coverage in the passaged isolates at P9 suggests that adaptive mutations against Wolbachia-induced blocking may be hard to achieve and maintain in viral populations. Indeed, the viral loads of Aag-2uMel-passaged isolates by the end of serial passaging were 3–4 log10-folds lower than isolates passaged in Wolbachia-free cells. Assuming that the viral RNA polymerase error rate is constant regardless of cell line, the total opportunities for mutational events in these populations were therefore much reduced compared to Wolbachia-free controls, lessening the chance for an evolutionary solution to blocking. The lack of increased replicative fitness in the passaged isolates relative to P0, even when reinoculated into the original cell line in which they had been passaged, was unexpected given that recent studies have demonstrated that serial passaging of RNA viruses in a single host leads to rapid (<10 passages) fitness gains in said host (Coffey et al. 2008; Vasilakis et al. 2009; Grubaugh et al. 2015). Most of the Aag-2Tet-passaged isolates showed decreased replicative ability compared to P0. It is possible that the suboptimal conditions for viral replication as a side effect of tetracycline treatment (Moullan et al. 2015) induced some unexpected selective pressures, producing genotypes that are generally less fit. The Aag-2uMel-passaged isolates performed poorly in all tested cell lines. In the Wolbachia-infected cell lines the inoculum may have been too low to recover in the presence of the symbiont. However, the inability of these isolates to replicate in Wolbachia-free cell lines was surprising and unlike that of the Aag-2- and Aag-2Tet-passaged isolates. This difference may suggest that while a subset of viruses was able to escape the Wolbachia effect, they may be less fit in a wildtype context.

It is possible that the lack of replication by Aag-2uMel-passaged isolates in Wolbachia-free cells could be explained by a higher than expected proportion of defective virus genome copies being produced during the later stages of serial passaging, leading to an overestimation of inoculum size used in this fitness assay. To replicate, defective virus particles rely on the presence of coinfecting complete viral genome copies for functional complementation and hence may interfere with the replication of functional virus copies through competitive inhibition (Huang and Baltimore 1970; Stauffer Thompson, Rempala, and Yin 2009). They have been reported to exist in both in vitro and in vivo systems and are capable of being carried through many transmission cycles (Wang et al. 2002; Aaskov et al. 2006; Li et al. 2011; Juarez-Martinez et al. 2013). Our qRT-PCR method of quantifying viral load detects viral genome copies instead of infectious virions. Genome copies are often used as a proxy for infectious particles because the two are well correlated, but the presence of immature virus or defective genomes can inflate estimations of viral load by as much as 2–5 log10-folds (Choy et al. 2013). If Aag-2uMel-passaged isolates contain a higher proportion of defective genome copies, the number of functional virus particles may be too low to cause infection even in Wolbachia-free cells. One way to validate this hypothesis would be to conduct plaque assays on viral isolates and measure the concentration of infectious virions. Unfortunately, the DENV-3 isolate utilized does not plaque well. This isolate was selected after pilot experiments (also including DENV-2) showed that DENV-3 had the greater ability of replicating in the presence of Wolbachia.

Assuming our theory of defective genome copy enrichment in Aag-2uMel-passaged isolates is correct, there are two possible mechanisms behind the population crashes observed during the serial passaging regime. Wolbachia may be creating an intracellular environment that interferes with the accurate and efficient replication of the DENV genome, increasing the ratio of defective to non-defective genomes. Wolbachia infection in insect cells is known to increase ROS production (Pan et al. 2012; Wong, Brownlie, and Johnson 2015) that negatively affects RNA virus replication (Choi et al. 2004). Alternatively, Wolbachia-induced blocking may simply have caused viral population sizes to fall below a threshold necessary to sustain functional complementation, given an unaltered number of defective genome copies constantly present in cell culture (Juarez-Martinez et al. 2013).

Replicated virus in Aag-2uMel-passaged isolates could be the product of low levels of non-Wolbachia-infected cells in the cell lines rather than survivors of selection. While Wolbachia prevalence rates appeared high via FISH throughout passaging, we cannot rule out the presence of low numbers of uninfected or very slowly infected cells. Viral variants that are the products of these cells, therefore, would not have had the opportunity to face purifying selection imposed by the blocking phenotype. These variants may also outnumber true products of uMel-infected cells and may explain why the Aag-2uMel-passaged isolates did not exhibit adaptation. Care must, therefore, be taken when investigating variant sequences of these isolates, as the consensus sequence may not reflect the adapted variants. Combined with the inability to analyze sequences of Aag-2uMel-passaged isolates, it is difficult to confirm from our study whether evolution has occurred during the serial passaging regime.

Due to obligate replication in two disparate host systems, the adaptive evolution of arboviruses is thought to be slower relative to other RNA viruses transmitted through a single host species. In particular, beneficial mutations in one host may be antagonistic in the alternate host, incurring a fitness tradeoff (Holmes 2006), in turn increasing the strength of purifying selection. Purifying selection has also been found to act at the intracellular and extracellular levels (Stapledford et al. 2014; Grubaugh et al. 2016; Lequime et al. 2016). The presence of Wolbachia adds an additional layer of constraint on top of the strong purifying selective forces shaping DENV evolution, considerably impeding its evolutionary potential. Given that half of our Aag-2uMel-passaged isolates experienced population extinction and the surviving half failed to exhibit significant population expansion or fitness gains, our experiment has demonstrated that Wolbachia-induced blocking poses a great challenge for DENV-3 (strain 08/09) adaptation.

Regardless, the sheer scale of the release of Wolbachia into global populations of mosquitoes significantly amplifies the opportunity for evolutionary events to occur, over time, that could allow the emergence of viral escape variants. Additionally, viral replication in the human host or in a Wolbachia-free mosquito (in a mosquito population where Wolbachia infection is not
Data availability
All raw data are available via Figshare: http://doi.org/10.26180/5c2f68dd897005. Sequence data have not been made available given poor coverage of the genome.

Supplementary data
Supplementary data are available at Virus Evolution online.

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References
Aaskov, J. et al. (2006) ‘Long-Term Transmission of Defective RNA Viruses in Humans and Aedes Mosquitoes’, Science (New York, N.Y.), 311: 236–8.
Amuzu, H. E., and McGraw, E. A. (2016) ‘Wolbachia-Based Dengue Virus Inhibition Is Not Tissue-Specific in Aedes aegypti’, PLoS Neglected Tropical Diseases, 10: e0005145.
Asad, S., Parry, R., and Asgari, S. (2018) ‘Upregulation of Aedes aegypti Vago1 by Wolbachia and Its Effect on Dengue Virus Replication’, Insect Biochemistry and Molecular Biology, 92: 45–52.
Byk, L. A., and Gamarnik, A. V. (2016) ‘Properties and Functions of the Dengue Virus Capsid Protein’, Annual Review of Virology, 3: 263–81.
Caragata, E. P. et al. (2013) ‘Dietary Cholesterol Modulates Pathogen Blocking by Wolbachia’, PLoS Pathogens, 9: e1003459.
Choi, J. et al. (2004) ‘Reactive Oxygen Species Suppress Hepatitis C Virus RNA Replication in Human Hepatoma Cells’, Hepatology, 39: 81–9.
Choy, M. M. et al. (2013) ‘Comparison of the Mosquito Inoculation Technique and Quantitative Real Time Polymerase Chain Reaction to Measure Dengue Virus Concentration’, The American Journal of Tropical Medicine and Hygiene, 89: 1001–5.
Chrostek, E. et al. (2013) ‘Wolbachia Variants Induce Differential Protection to Viruses in Drosophila melanogaster: A Phenotypic and Phylogenomic Analysis’, PLoS Genetics, 9: e1003896.
Coffey, L. L. et al. (2008) ‘Arbovirus Evolution In Vivo Is Constrained by Host Alternation’, Proceedings of the National Academy of Sciences of the United States of America, 105: 6970–5.
Ducharme, N. A., and Bickel, P. E. (2008) ‘Lipid Droplets in Lipogenesis and Lipolysis’, Endocrinology, 149: 942–9.
Dutra, H. L. et al. (2016) ‘Wolbachia Blocks Currently Circulating Zika Virus Isolates in Brazilian Aedes aegypti Mosquitoes’, Cell Host & Microbe, 19: 771–4.
Ferguson, N. M. et al. (2015) ‘Modeling the Impact on Virus Transmission of Wolbachia-Mediated Blocking of Dengue Virus Infection of Aedes aegypti’, Science Translational Medicine, 7: 279ra37.
Frentiu, F. D. et al. (2010) ‘Wolbachia-Mediated Resistance to Dengue Virus Infection and Death at the Cellular Level’, PLoS One, 5: e13398.
Gubraith, N. D. et al. (2015) ‘Experimental Evolution of an RNA Virus in Wild Birds: Evidence for Host-Dependent Impacts on Population Structure and Competitive Fitness’, PLoS Pathogens, 11: e1004874.
Huang, A. S., and Baltimore, D. (1970) ‘Defective Viral Particles’, Science, 169: 1324–5.
Hygiene’, 326: 134–6.
Kambris, Z. et al. (2009) ‘Immune Activation by Life-Shortening Wolbachia and Reduced Filarial Competence in Mosquitoes’, Science, 326: 134–6.
Johnson, K. N. (2015) ‘The Impact of Wolbachia on Virus Infection in Mosquitoes’, Viruses, 7: 5705–17.
Juarez-Martinez, A. B. et al. (2013) ‘Detection and Sequencing of Defective Viral Genomes in C6/36 Cells Persistently Infected with Dengue Virus 2’, Archives of Virology, 158: 583–99.
Lazaro, E. et al. (2003) ‘Resistance of Virus to Extinction on Bottleneck Passages: Study of a Decaying and Fluctuating Pattern of Fitness Loss’, Proceedings of the National Academy of Sciences of the United States of America, 100: 10830–5.
Lequime, S. et al. (2016) ‘Genetic Drift, Purifying Selection and Vector Genotype Shape Dengue Virus Intra-Host Genetic Diversity in Mosquitoes’, PLoS Genetics, 12: e1006111.

Li, D. et al. (2011) ‘Defective Interfering Viral Particles in Acute Dengue Infections’, PLoS One, 6: e19447.

Lively, C. M. et al. (2005) ‘Competitive Co-Existence of Vertically and Horizontally Transmitted Parasites’, Evolutionary Ecology Research, 7: 1183–90.

Lu, P. et al. (2012) ‘Wolbachia Induces Density-Dependent Inhibition to Dengue Virus in Mosquito Cells’, PLoS Neglected Tropical Diseases, 6: e1754.

Mayoral, J. G. et al. (2014) ‘Wolbachia Infection Modifies the Profile, Shuttling and Structure of microRNAs in a Mosquito Cell Line’, PLoS One, 9: e96107.

Moreira, L. A. et al. (2009) ‘A Wolbachia Symbiont in Aedes aegypti Limits Infection with Dengue, Chikungunya, and Plasmodium’, Cell, 139: 1268–78.

Moullan, N. et al. (2015) ‘Tetracyclines Disturb Mitochondrial Function across Eukaryotic Models: A Call for Caution in Biomedical Research’, Cell Reports, 10: 1681–91.

Muller, D. A., and Young, P. R. (2013) ‘The Flavivirus NS1 Protein: Molecular and Structural Biology, Immunology, Role in Pathogenesis and Application as a Diagnostic Biomarker’, Antiviral Research, 98: 192–208.

O’Neill, S. et al. (2018) ‘Scaled Deployment of Wolbachia to Protect the Community from Dengue and Other Aedes Transmitted Arboviruses’, Gates Open Research, 2: 36.

Osborne, S. E. et al. (2009) ‘Variation in Antiviral Protection Mediated by Different Wolbachia Strains in Drosophila simulans’, PLoS Pathogens, 5: e1000656.

et al. (2012) ‘Antiviral Protection and the Importance of Wolbachia Density and Tissue Tropism in Drosophila simulans’, Applied and Environmental Microbiology, 78: 6922–9.

Pan, X. L. et al. (2012) ‘Wolbachia Induces Reactive Oxygen Species (ROS)-Dependent Activation of the Toll Pathway to Control Dengue Virus in the Mosquito Aedes aegypti’, Proceedings of the National Academy of Sciences of the United States of America, 109: E23–31.

Parameswaran, P. et al. (2017) ‘Intra-host Selection Pressures Drive Rapid Dengue Virus Micro-evolution in Acute Human Infections’, Cell Host & Microbe, 22: 400–10.

Peleg, J. (1968) ‘Growth of Arboviruses in Primary Tissue Culture of Aedes aegypti Embryos’, The American Journal of Tropical Medicine and Hygiene, 17: 219–23.

Rainey, S. M. et al. (2016) ‘Wolbachia Blocks Viral Genome Replication Early in Infection without a Transcriptional Response by the Endosymbiont or Host Small RNA Pathways’, PLoS Pathogens, 12: e1005536.

Rances, E. et al. (2012) ‘The Relative Importance of Innate Immune Priming in Wolbachia-Mediated Dengue Interference’, PLoS Pathogens, 8: e1002548.

Ritchie, S. A. et al. (2013) ‘An Explosive Epidemic of DENV-3 in Cairns, Australia’, PLoS One, 8: e68137. ‘An Samsa, M. M. et al. (2009) ‘Dengue Virus Capsid Protein Usurps Lipid Droplets for Viral Particle Formation’, PLoS Pathogens, 5: e1000632.

Simon, P. (2003) ‘Q-Gene: Processing Quantitative Real-Time RT-PCR Data’, Bioinformatics, 19: 1439–40.

Stapleford, K. A. et al. (2014) ‘Emergence and Transmission of Arbovirus Evolutionary Intermediates with Epidemic Potential’, Cell Host & Microbe, 15: 706–16.

Stauffer Thompson, K. A., Rempala, G. A., and Yin, J. (2009) ‘Multiple-Hit Inhibition of Infection by Defective Interfering Particles’, Journal of General Virology, 90: 888–99.

Stollar, V., and Thomas, V. L. (1975) ‘An Agent in the Aedes aegypti Cell Line (Peleg) Which Causes Fusion of Aedes albopictus Cells’, Virology, 64: 367–77.

Terraradas, G., Joubert, D. A., and McGraw, E. A. (2017) ‘The RNAI Pathway Plays a Small Part in Wolbachia-Mediated Blocking of Dengue Virus in Mosquito Cells’, Scientific Reports, 7: 43847.

Van den Hurk, A. F. et al. (2012) ‘Impact of Wolbachia on Infection with Chikungunya and Yellow Fever Viruses in the Mosquito Vector Aedes aegypti’, PLoS Neglected Tropical Diseases, 6: e1892.

Vasilakis, N. et al. (2011) ‘Fever from the Forest: Prospects for the Continued Emergence of Sylvatic Dengue Virus and Its Impact on Public Health’, Nature Reviews Microbiology, 9: 532–41.

et al. (2009) ‘Mosquitoes Put the Brake on Arbovirus Evolution: Experimental Evolution Reveals Slower Mutation Accumulation in Mosquito than Vertebrate Cells’, PLoS Pathogens, 5: e1000467.

Vavre, F., and Charlat, S. (2012) ‘Making (Good) Use of Wolbachia: What the Models Say’, Current Opinion in Microbiology, 15: 263–8.

Walker, T. et al. (2011) ‘The uMel Wolbachia Strain Blocks Dengue and Invades Caged Aedes aegypti Populations’, Nature, 476: 450–3.

Wang, W. K. et al. (2002) ‘Dengue Type 3 Virus in Plasma Is a Potential’, Journal of Virology, 76: 4662–5.

Warrell, D. et al. (2002) ‘Single Rapid TaqMan Fluorogenic Probe Based PCR Assay That Detects All Four Dengue Serotypes’, Journal of Medical Virology, 66: 524–8.

Welsch, S. et al. (2009) ‘Composition and Three-Dimensional Architecture of the Dengue Virus Replication and Assembly Sites’, Cell Host & Microbe, 5: 365–75.

White, P. M. et al. (2017) ‘Reliance of Wolbachia on High Rates of Host Proteolysis Revealed by a Genome-Wide RNAi Screen of Drosophila Cells’, Genetics, 205: 1473–88.

Wong, Z. S., Brownlie, J. C., and Johnson, K. N. (2015) ‘Oxidative Stress Correlates with Wolbachia-Mediated Antiviral Protection in Wolbachia-Drosophila Associations’, Applied and Environmental Microbiology, 81: 3001–5.

Xie, Q. et al. (2017) ‘Structure and Function of the Non-Structural Protein of Dengue Virus and Its Applications in Antiviral Therapy’, Current Topics in Medicinal Chemistry, 17: 371–80.

Ye, Y. H. et al. (2014) ‘Comparative Susceptibility of Mosquito Populations in North Queensland, Australia to Oral Infection with Dengue Virus’, The American Journal of Tropical Medicine and Hygiene, 90: 422–30.

Yeap, H. L. et al. (2014) ‘Assessing Quality of Life-Shortening Wolbachia-Infected Aedes aegypti Mosquitoes in the Field Based on Capture Rates and Morphometric Assessments’, Parasites & Vectors, 7: 58.

Zhang, G. et al. (2013) ‘Wolbachia Uses a Host microRNA to Regulate Transcripts of a Methyltransferase, Contributing to Dengue Virus Inhibition in Aedes aegypti’, Proceedings of the National Academy of Sciences of the United States of America, 110: 10276–81.

Zhang, G. M. et al. (2017) ‘Cell Fusing Agent Virus and Dengue Virus Mutually Interact in Aedes aegypti Cell Lines’, Scientific Reports, 7: 6935.