Comparative Susceptibility of *Aedes albopictus* and *Aedes aegypti* to Dengue Virus Infection After Feeding on Blood of Viremic Humans: Implications for Public Health

James Whitehorn,1,3 Duong Thi Hue Kien,3 Nguyet Minh Nguyen,3 Hoa L. Nguyen,3 Peter P. Kyrylos,5 Lauren B. Carrington,3,5 Chau Nguyen Bich Tran,3 Nguyen Thanh Ha Quyen,3 Long Vo Thi,3 Dui Le Thi,3 Nguyen Thanh Truong,4 Tai Thi Hue Luong,4 Chau Van Vinh Nguyen,4 Bridget Wills,2,3 Marcel Wolbers,2,3 and Cameron P. Simmons2,3,5

1London School of Hygiene and Tropical Medicine, and 2Oxford University, United Kingdom; 3Oxford University Clinical Research Unit, and 4Hospital for Tropical Diseases, Ho Chi Minh City, Vietnam; and 5University of Melbourne, Australia

(See the editorial commentary by Christofferson on pages 1177–9.)

*Aedes albopictus* is secondary to *Aedes aegypti* as a vector of dengue viruses (DENVs) in settings of endemicity, but it plays an important role in areas of dengue emergence. This study compared the susceptibility of these 2 species to DENV infection by performing 232 direct blood-feeding experiments on 118 viremic patients with dengue in Vietnam. Field-derived *A. albopictus* acquired DENV infections as readily as *A. aegypti* after blood feeding. Once infected, *A. albopictus* permitted higher concentrations of DENV RNA to accumulate in abdominal tissues, compared with *A. aegypti*. However, the odds of *A. albopictus* having infectious saliva were lower than the odds observed for *A. aegypti* (odds ratio, 0.70; 95% confidence interval, .52–.93). These results quantify the susceptibility of *A. albopictus* to DENV infection and will assist parameterization of models for predicting disease risk in settings where *A. albopictus* is present.

**Keywords.** Dengue; *Aedes aegypti*; *Aedes albopictus*; susceptibility; transmission.
Lambrechts et al reviewed vector competence literature and reported that, although *A. albopictus* were more susceptible to DENV midgut infection, rates of virus dissemination to other tissues were significantly lower in *A. albopictus* than in *A. aegypti* [18]. Additionally, laboratory-reared *A. albopictus* became increasingly susceptibility to DENV, which may have been a confounding variable in the literature. Furthermore, the comparative vector competence literature has been derived entirely from laboratory experiments using artificial blood meals and laboratory-passaged viruses. How well these laboratory conditions replicate the pathogenesis of DENV transmission from viremic humans to mosquitoes in the field is uncertain.

The benefits of understanding why *A. albopictus* is largely a secondary vector of DENV in settings of endemicity are numerous. First, existing risk-analysis models of the likelihood of dengue outbreaks in southern Europe and the United States could be improved with quantitative estimates of *A. albopictus* vector competence [19, 20]. Second, probability maps of dengue occurrence could be refined with better estimates of the relative vector competence of *A. albopictus* versus *A. aegypti*. Third, if the historical literature is correct and *A. albopictus* is more resistant than *A. aegypti* to disseminated DENV infection, then this provides an opportunity to identify species-specific antiviral defense mechanisms. Last, novel dengue control efforts using Wolbachia-infected *A. aegypti* will be better informed with an understanding of the vector competence of *A. albopictus* in candidate intervention settings [21]. To these ends, the current study compared the susceptibility of *A. aegypti* and *A. albopictus* to initial and disseminated DENV infection after direct blood-feeding episodes on viremic patients with dengue.

**METHODS**

**Ethics Statement**
All participants provided written informed consent to participate in the study. The study protocols were reviewed and approved by the Scientific and Ethical Committee of the Hospital for Tropical Diseases (CS/NĐ/12/15) and the Oxford Tropical Research Ethical Committee (OxTREC 29–12). All investigations were conducted in accordance with the principles expressed in the Declaration of Helsinki.

**Patient Cohorts**
The study was performed at the Hospital for Tropical Diseases in Ho Chi Minh City, Vietnam, between September 2012 and November 2013. The inclusion criteria were age of ≥15 years, fever duration of ≤96 hours, and clinical suspicion of dengue; positive result of an NS1 rapid test; and provision of written informed consent. The exclusion criteria were current pregnancy, determined by clinical examination or a urine dipstick test for β-human chorionic gonadotropin; current intensive care unit stay; intellectual disability; a history of severe reactions to mosquito bites; and severe dermatological conditions. Demographic and clinical information were recorded prospectively in a standard case report form.

**Mosquitoes for Blood-Feeding Experiments**
All of the mosquitoes (*A. aegypti* and *A. albopictus*) that fed on patients with dengue were F3 generation and derived from 2 independent pooled larval collections, each sampled from 3 locations within 5 km of each other in Ho Chi Minh City. *A. aegypti* used were distinct from those in our previous study [22]. Briefly, field-caught larvae were pooled and fed commercial dry fish and dog food. Mosquitoes were housed as described previously [22]. Briefly, adults (F1 generation) were kept in cages containing males and females in an environmental chamber with 12-hour cycles of light and dark at 27°C and 70% relative humidity. F1 females were provided blood meals by direct feeding on afebrile healthy human volunteers for multiple gonotrophic cycles over 45 days, with 15% sucrose provided freely in addition. Eggs from F1 females were hatched and reared and the subsequent F2 females provided with human blood meals as described above. When female F2 mosquitoes were 12 days old, they were killed and pooled into groups of 10 mosquitoes. Each pool was homogenized and tested, along with appropriate controls, by reverse transcription–polymerase chain reaction (RT-PCR) to confirm the absence of DENV, Japanese encephalitis virus, and chikungunya virus. Eggs collected and stored from F2 females were the source of F3 females that were used for direct feeding on patients with dengue.

**Experimental Exposure of Patients to Mosquitoes**
Each patient was assigned a schedule of 2 exposures to mosquitoes on 2 different study days during the first 4 days after enrollment. The patient’s forearm was exposed to 30–40 *A. aegypti* and *A. albopictus* aged 3–7 days (using a 2:3 ratio of *A. aegypti* to *A. albopictus* because preliminary experiments indicated that a higher fraction of *A. aegypti* took a blood meal) contained in a mesh-covered plastic cup that was held against the patient’s forearm for 5 minutes. After 5 minutes, mosquitoes were returned to the insectary and subjected to cold anesthesia at 4°C for 45 seconds. Engorged mosquitoes were transferred to 500-mL plastic cups, separated by species, and maintained in an environmental chamber with 12-hour cycles of light and dark at 27°C and 70% relative humidity for 14 days. The number of dead mosquitoes was recorded daily.

**Clinical Adverse Events**
Postexposure severe adverse events were defined as any event that was clinically significant (ie, requiring clinical intervention, prolonged hospitalization, or admission to an intensive care unit) and possibly, probably, or definitely related to experimental exposure to mosquitoes.
Detection of DENV in Mosquito Tissues and Saliva

Technicians blinded to clinical and virological details of the participants performed laboratory assays of the mosquitoes. Mosquitoes were killed by cold exposure. The abdomen was dissected from the rest of the mosquito body and suspended in 0.5 mL of mosquito diluent (2% [v/v] heat-inactivated fetal calf serum [FCS] in Roswell Park Memorial Institute 1640 medium, antibiotics, and antifungotics). Individual mosquito abdomens were homogenized with 1-mm zirconia/silica beads for 15 minutes at 30 Hz by using a TissueLyser II system (Qiagen), as described previously [22]. Mosquitoes were scored as being infected with DENV, using a previously described quantitative RT-PCR analysis of homogenized tissue, with the results expressed as copies per abdomen [23]. Infectious virus in the saliva of individual mosquitoes was detected as described previously [22]. Briefly, the proboscis of dewinged and deglazed mosquitoes was inserted into the end of a micropipette tip containing 6 µL of filtered saliva medium (a 1:1 solution of 15% [v/v] sucrose in normal saline and inactivated FCS). After 30 minutes, the 6 µL of saliva medium was ejected and then drawn into a pointed glass capillary tube (tip diameter, <0.3 µm), after which the volume of saliva medium derived from 1 mosquito was injected into the thorax of 4–6 A. aegypti (age, 4–7 days; volume injected, approximately 1 µL per mosquito). These injected mosquitoes were maintained for 7 days in an environmental chamber as described above. After 7 days, the mosquitoes were killed and the bodies pooled, homogenized, and tested by quantitative RT-PCR for DENV infection, with saliva samples scored as positive or negative depending on this result.

Sequence Amplification and Sequencing of the Gene Encoding DENV Envelope Protein

Viral RNA was extracted as previously described from plasma and mosquito abdomen tissues [22]. Complementary DNA (cDNA) was synthesized from 8 µL of viral RNA by using the Superscript III First-Strand Synthesis System for RT-PCR according to manufacturers’ instructions. The genomic region spanning the genes encoding premembrane and envelope proteins was amplified from the cDNA in 16 amplifiers. Subsequent steps were performed according to the manufacturers instructions for 454 GS-Junior Next Generation Sequencing method (Roche). Briefly, the primers in this first PCR round contained universal tails at the 5′ end to allow the addition of 454 sequencing-specific nucleotides and isolate-specific multiplex identifiers (also known as “barcodes”) in a second PCR round. The first-round and second-round PCR analyses used FastStart High Fidelity polymerase (Roche). The long-read sequencing performance of the 454 GS-Junior (between 400 and 500 bases), in combination with a sample pooling strategy that uses barcoded amplicons, was used for parallel analysis of pooled samples. GS Mapping software was used for primer trimming and alignment of reads against a reference sequence DENV-1/VN/BID-V2732/2007 (GenBank accession number GQ199773.1). Sequence quality was measured using Phred (Q) scores with a minimum acceptable threshold of 95% of sequencing reads having Q scores of >20 (1/100 errors per base).

Dengue Diagnostic Tests

Serological responses were detected using immunoglobulin M (IgM) and immunoglobulin G (IgG) antibody-capture enzyme-linked immunosorbent assays in accordance with the manufacturer’s instructions (Panbio, Australia). In accordance with the manufacturer’s instructions, we classified serological profiles as “probable secondary” when >22 U of IgG were detected in either acute or early convalescent samples. If acute or early convalescent samples were IgM positive but IgG negative, serological profiles were classified as “probable primary.” When IgM or IgG tests result were equivocal, we classified the serological profile as “indeterminate.”

DENV plasma viremia levels were measured by a validated quantitative RT-PCR assay that has been described previously [23].

Statistical Analysis

The probability of successful human-to-mosquito transmission was compared between A. aegypti and A. albopictus, using marginal logistic regression models. Models for assessment of abdomen samples were adjusted for the patient’s log10-transformed viremia level, and models for assessment of saliva samples were further adjusted for the abdominal tissue viremia level. Analyses were performed for all patients and stratified by serotype. To account for potential within-patient correlation, model-robust sandwich standard error estimates were used throughout to construct confidence intervals (CIs) and P values. We derived 50% mosquito infectious doses (MID50 values; defined as plasma viremia levels corresponding to a 50% probability of infection) for abdomen infection on the basis of marginal logistic regression coefficients for each mosquito type, and corresponding 95% CIs were calculated using the delta rule. Since the proportion of mosquitoes with infectious saliva was <50%, MID50 values were not estimated. In the initial analysis of the transmission of DENV to saliva, mosquitoes with uninfected abdomens were assigned a log value of 0. This analysis was also run with exclusion of mosquitoes with uninfected abdomen’s. The relationship between various covariates and the probability of successful human-to-mosquito transmission was assessed for each mosquito type, using a similar multivariable marginal regression model. The covariates assessed were day of illness, plasma viremia level, serotype, serological result, and abdominal viral burden. The number of mutations in the DENV-1 consensus sequence was compared between mosquito types, using a stratified version of the Wilcoxon test with stratification by patient [24]. All the analyses were performed using the R statistical software package, version 2.13.2 (R Foundation for Statistical Computing; Vienna, Austria).
RESULTS

Study Population Characteristics
Between September 2012 and November 2013, 120 patients with dengue were enrolled and experimentally exposed to field-derived *A. aegypti* and *A. albopictus* on 2 randomly allocated days within the first 4 study days. The patient enrollment flowchart is shown in Supplementary Figure 1. The final cohort for analysis comprised 118 DENV viremic patients with 232 independent mosquito exposure events. The baseline characteristics of the patients enrolled are shown in Table 1. Experimental exposure to mosquitoes was well tolerated, and no patient experienced a severe adverse event or required withdrawal from the study. DENV serotype 1 (DENV-1) and DENV-4 were responsible for 26% and 44% of cases, respectively, with DENV-2 (13%) and DENV-3 (16%) also represented.

Susceptibility of *A. aegypti* and *A. albopictus* to Acquisition of DENV Infection After Direct Feeding
There was a dose-response relationship between plasma viremia levels at the time of mosquito feeding and the proportion of mosquitoes of either species with DENV-infected abdomens 14 days later (Figure 1). The overall DENV MID$_{50}$ for *A. aegypti* was lower than that observed for *A. albopictus*, but this difference was not statistically significant (7.0 log$_{10}$ copies/mL [95% CI, 6.77–7.23] vs 7.1 log$_{10}$ copies/mL [95% CI, 6.9–7.3]; Table 2). Viral RNA concentrations of all 4 DENV serotypes were significantly higher in abdomen tissues from infected *A. albopictus* than in those from *A. aegypti* (Supplementary Table 1). Given that *A. albopictus* accommodated higher DENV RNA concentrations once infected, we examined whether this could result in greater virus sequence diversity. As a test case, the consensus nucleotide sequence of the gene encoding DENV-1 envelope protein was determined directly by analysis of plasma specimens from 10 patients and abdominal tissues from 20 *A. aegypti* and 20 *A. albopictus* that had taken blood meals from these cases (ie, 2 *A. aegypti* and 2 *A. albopictus* per patient). The median number of nucleotide differences between the consensus sequences of the gene recovered from plasma specimens and the gene recovered from abdominal tissue was 9 (interquartile range [IQR], 4–15) for *A. aegypti* and 17 (IQR, 11–18) for *A. albopictus* ($P = .02$, by the Wilcoxon test [stratified by patient]; Supplementary Figure 2).

Table 1. Characteristics of 118 Patients With Dengue Virus (DENV) Infection, by DENV Serotype

| Characteristic                      | DENV-1 (n = 31) | DENV-2 (n = 15) | DENV-3 (n = 19) | DENV-4 (n = 52) | Overall (n = 118)$^a$ |
|-------------------------------------|----------------|----------------|----------------|----------------|---------------------|
| Age, y                              | 23 (19–31)     | 23 (21–32)     | 25 (20–27)     | 27.5 (22–34)   | 26 (20–32)          |
| Sex                                 |                |                |                |                |                     |
| Male                                | 12 (38.7)      | 7 (46.7)       | 5 (26.3)       | 17 (32.7)      | 41 (34.8)           |
| Female                              | 19 (61.3)      | 8 (53.3)       | 14 (73.7)      | 35 (67.3)      | 77 (65.3)           |
| Illness duration at enrollment, d$^b$| 7 (22.3)       | 5 (33.3)       | 0 (0)          | 7 (13.4)       | 19 (16.2)           |
| Viremia level, log$_{10}$ copies/mL | 8.1 (6.9–8.7)  | 7.6 (7.1–8.9)  | 6.6 (5.5–7.1)  | 7.2 (6.5–7.8)  | 7.2 (6.6–8.1)       |
| Serological profile$^c$              |                |                |                |                |                     |
| Primary                             | 10 (32.3)      | 0 (0)          | 1 (5.3)        | 1 (2.0)        | 12 (10.3)           |
| Secondary                           | 17 (54.8)      | 12 (80.0)      | 16 (84.2)      | 48 (94.1)      | 94 (80.3)           |
| Indeterminate                       | 4 (12.9)       | 3 (20.0)       | 2 (10.5)       | 2 (3.9)        | 11 (9.4)            |
| Clinical classification             |                |                |                |                |                     |
| Dengue                              | 26 (83.9)      | 14 (93.3)      | 16 (84.2)      | 47 (90.4)      | 104 (88.1)          |
| Dengue with warning signs           | 5 (16.1)       | 1 (6.7)        | 2 (10.5)       | 4 (7.7)        | 12 (10.2)           |
| Severe dengue                       | 0 (0)          | 0 (0)          | 0 (0)          | 0 (0)          | 0 (0)               |
| Others                              | 0 (0)          | 0 (0)          | 1 (5.3)$^d$    | 1 (1.9)$^e$    | 2 (1.7)             |
| Transferred to ICU                  | 1 (3.2)        | 0 (0)          | 0 (0)          | 0 (0)          | 1 (0.9)             |

Data are no. (%) of patients or median value (interquartile range).

Abbreviation: ICU, intensive care unit.

$^a$ The viremia level for 1 patient was below the limit of detection.

$^b$ Data for 1 patient were missing.

$^c$ Data for 1 patient with DENV-4 infection were missing.

$^d$ One patient was categorized as having a “viral infection” but was actually infected with DENV-3.

$^e$ One patient was categorized as having a “viral infection” but was actually infected with DENV-4.
Acquisition of Infectiousness Among *A. aegypti* and *A. albopictus*

As was observed in abdomen tissues, there was a dose-response relationship between the plasma viremia level and the proportion of mosquitoes with infectious saliva 14 days after blood feeding (Figure 2). However, the likelihood of detecting infectious saliva differed by mosquito species and DENV serotype. Table 3 summarizes the odds of abdomen and saliva infection and demonstrates that the detection of infectious saliva was less likely in blood-fed *A. albopictus*, compared with blood-fed *A. aegypti*, in both unadjusted analysis and analysis that adjusted for plasma viremia level (adjusted odds ratio [OR], 0.70; 95% CI, .52–.93). By serotype, the odds of *A. albopictus* having infectious saliva were significantly lower for blood meals involving uptake of DENV-2 and DENV-4, compared with those involving uptake of DENV-1 or DENV-3 (Table 3). These data identified the odds of *A. albopictus* becoming infectious as lower than the odds of *A. aegypti* becoming infectious after feeding on the blood of viremic patients. This analysis used all engorged mosquitoes as the denominator. Supplementary Table 2 shows the odds of having infectious saliva, using mosquitoes with DENV-positive abdomens as the denominator. This analysis confirms that the odds of *A. albopictus* becoming infectious were lower than those for *A. aegypti* after viremic blood feeding (OR, 0.69; 95% CI, .49–.96).

**Covariates and Their Association With Successful Human-to-Mosquito Transmission**

In multivariable regression analysis, a higher plasma viremia level at the time of exposure was independently associated with a greater likelihood of DENV transmission to abdominal tissue for both mosquito types. Each 1-log increase in the

Table 2. 50% Mosquito Infectious Doses (MID₅₀ Values) for *Aedes albopictus* and *Aedes aegypti* Abdomen Infection, by Dengue Virus (DENV) Serotype

| Serotype | *A. aegypti* | *A. albopictus* | Absolute Difference | P Value |
|----------|--------------|----------------|--------------------|---------|
| DENV-1   | 6.62 (6.10–7.13) | 6.74 (6.27–7.21) | 0.12 (−.11 to .35) | .29     |
| DENV-2   | 6.96 (6.54–7.38) | 7.03 (6.62–7.43) | 0.07 (−.17 to .30) | .59     |
| DENV-3   | 6.49 (5.69–7.30) | 6.70 (5.86–7.56) | 0.21 (−.39 to .82) | .49     |
| DENV-4   | 7.37 (7.04–7.70) | 7.38 (7.07–7.69) | 0.01 (−.21 to .24) | .89     |
| Overall  | 7.00 (6.77–7.23) | 7.10 (6.90–7.30) | 0.10 (−.04 to .24) | .15     |

MID₅₀ values (defined as plasma viremia levels corresponding to a 50% probability of infection) were derived on the basis of marginal logistic regression coefficients for each mosquito type, and corresponding 95% confidence intervals were calculated using the delta rule.
The log10 plasma viremia level was associated with a 4.65-fold increase in the odds of abdominal tissue infection in \textit{A. aegypti} (95% CI, 3.34–6.48) and a 5.3-fold increase in the odds of abdominal tissue infection in \textit{A. albopictus} (95% CI, 3.80–7.43; Table 4). Each 1-log increase in the log10 plasma viremia level was associated with 2.16-fold increase in the odds of saliva infection in \textit{A. aegypti} (95% CI, 1.68–2.78) and a 2.79-fold increase in the odds of saliva infection in \textit{A. albopictus} (95% CI, 2.17–3.60; Table 4). These data highlight plasma viremia level as a risk factor for infectiousness among \textit{A. aegypti} and \textit{A. albopictus}. Supplementary Table 3 shows the association between these covariates and successful transmission, with mosquitoes with DENV-negative abdomens excluded from the model.

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure2.png}
\caption{Dose response scatterplot and curve (derived from logistic regression models) of plasma viremia versus the proportion of mosquitoes with infectious saliva after feeding on 118 dengue virus (DENV)–infected patients, showing curves for \textit{Aedes aegypti} and \textit{Aedes albopictus}. Each dot point represents the proportion of mosquitoes that took a blood meal during an exposure event and had saliva containing infectious DENV 14 days later and the corresponding plasma viremia level in the patient at the time of mosquito exposure, stratified by DENV serotype.}
\end{figure}

\begin{table}[h]
\centering
\begin{tabular}{|l|l|l|l|l|l|l|}
\hline
Specimen, Serotype & A. albopictus & A. aegypti & Unadjusted & Adjusted & & \\
& Infected/Tested, No. (%) & Infected/Tested, No. (%) & OR (95% CI) & OR (95% CI) & P Value & P Value \\
\hline
\hline
Abdomen & & & & & & \\
DENV-1 & 27 & 291/411 (70.8) & 278/379 (73.4) & 0.88 (.62–1.25) & .48 & 0.79 (.52–1.10) & .28 \\
DENV-2 & 13 & 108/185 (58.4) & 143/220 (65.0) & 0.76 (.52–1.09) & .13 & 0.97 (.52–1.82) & .93 \\
DENV-3 & 16 & 67/173 (38.7) & 83/208 (39.8) & 0.95 (.55–1.66) & .86 & 0.76 (.39–1.48) & .42 \\
DENV-4 & 49 & 229/777 (29.5) & 232/751 (30.9) & 0.93 (.70–1.24) & .64 & 0.95 (.69–1.31) & .77 \\
Overall & 105 & 695/1546 (45.0) & 736/1558 (47.2) & 0.91 (.78–1.07) & .27 & 0.85 (.69–1.06) & .16 \\
\hline
Saliva & & & & & & \\
DENV-1 & 27 & 177/411 (43.1) & 142/379 (37.5) & 1.26 (.89–1.80) & .20 & 1.31 (.90–1.93) & .16 \\
DENV-2 & 13 & 22/185 (11.9) & 91/220 (41.4) & 0.19 (.11–.33) & <.001 & 0.17 (.09–.31) & <.001 \\
DENV-3 & 16 & 17/173 (9.8) & 35/208 (16.8) & 0.54 (.21–1.37) & .19 & 0.44 (.19–1.00) & .051 \\
DENV-4 & 49 & 60/777 (7.7) & 92/751 (12.3) & 0.60 (.36–.99) & .044 & 0.60 (.37–.98) & .041 \\
Overall & 105 & 276/1546 (17.9) & 360/1558 (23.1) & 0.72 (.55–.95) & .022 & 0.70 (.52–.93) & .014 \\
\hline
\end{tabular}
\caption{Comparison of the Odds of Abdomen and Saliva Infection Among \textit{Aedes albopictus} Versus the Odds Among \textit{Aedes aegypti}, by Dengue Virus (DENV) Serotype}
\end{table}

*Abbreviations: CI, confidence interval; OR, odds ratio.

* Marginal logistic regression models adjusted for plasma viremia.
Ilness duration at enrollment
MID50 values for systemic infection [33]. Our R80 proteins may act as midgut receptors leading to subsequent pathways. There are few insights into the nature of these recep-
ters species difference of epidemiological importance.

However, blood-fed A. albopictus was significantly less likely to become infectious with DENV-2 and DENV-4, suggesting an interspecies difference of epidemiological importance.

A first step in the establishment of DENV infection in mosquitoes is attachment of virus particles to luminal receptors on the midgut epithelium, followed by virus entry via endocytic pathways. There are few insights into the nature of these receptors for DENV, although it has been proposed that the R67 and R80 proteins may act as midgut receptors leading to subsequent systemic infection [33]. Our finding that the plasma viremia MID50 values for A. aegypti and A. albopictus were broadly similar suggests that DENV uses common mechanisms to attach and initially infect the midgut epithelium in these specimens. The MID50 values for DENV-1 and DENV-4 measured in this study were concordant with those observed previously, whereas the MID50 values differed for DENV-2 and DENV-3 [22]. We speculate that this reflects the smaller numbers of patients with DENV-2 and DENV-3 in this current study and hence decreased accuracy.

Our results identified that A. albopictus and A. aegypti were equally likely to develop infectious saliva containing DENV-1 and DENV-3. However, DENV-2 and DENV-4 were detected in saliva from A. aegypti less frequently than in saliva from A. aegypti. This might simply reflect intrinsic differences between the virus types in their virulence for A. albopictus, with DENV-1 and DENV-3 being fitter in this instance. However, it is interesting to note that, while DENV-1 was the dominant serotype in the United States, where A. albopictus is prevalent [34].

We explored the relationship between various covariates and the likelihood of successful human-to-mosquito transmission. We confirmed the importance of the plasma viremia level in success-
ful human-mosquito transmission: each 1-log increase in plasma viremia level was associated with an approximately 5-fold increase in the odds of successful human-to-mosquito transmission for both A. aegypti and A. albopictus [22]. This finding again suggests that interventions that reduce DENV viremia during natural infection could have both an individual benefit and a public health benefit through reducing human infectiousness and thus reducing the risk of further transmission [19].

DENV RNA concentrations were significantly higher in abdo-
minal tissues of A. albopictus than A. aegypti for all serotypes.

Table 4. Covariates and Their Association With Successful Dengue Virus (DENV) Transmission Among Aedes albopictus and Aedes aegypti

| Variable                      | A. aegypti | A. albopictus | A. aegypti | A. albopictus |
|-------------------------------|------------|---------------|------------|---------------|
|                                | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value |
| Illness duration at enrollment | 0.58 (.36–.93) | .025 | 0.60 (.39–.92) | .018 | 0.87 (.61–1.24) | .45 | 0.91 (.62–1.34) | .63 |
| Viremia level (per 1-log10 copies/mL increase) | 4.65 (3.34–6.48) | <.001 | 5.32 (3.80–7.43) | <.001 | 2.16 (1.68–2.78) | <.001 | 2.79 (2.17–3.60) | <.001 |

DENV serotype

| DENV | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value |
|------|----------------------|---------|----------------------|---------|----------------------|---------|----------------------|---------|
| DENV-1 | 1.00 (reference) | 1.00 (reference) | 1.00 (reference) | 1.00 (reference) |
| DENV-2 | 0.57 (.18–1.78) | .34 | 0.74 (.24–2.34) | .59 | 1.43 (.69–2.95) | .34 | 0.18 (.08–.39) | <.001 |
| DENV-3 | 1.29 (.30–5.49) | .73 | 1.31 (.31–5.47) | .71 | 1.08 (.41–2.86) | .88 | 0.40 (.12–1.38) | .15 |
| DENV-4 | 0.27 (.10–.71) | .008 | 0.36 (.15–.88) | .026 | 0.48 (.24–.99) | .046 | 0.23 (.12–.44) | <.001 |

Serological profile

| Type | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value | Adjusted OR (95% CI) | P Value |
|------|----------------------|---------|----------------------|---------|----------------------|---------|----------------------|---------|
| Primary | 1.00 (reference) | 1.00 (reference) | 1.00 (reference) | 1.00 (reference) |
| Secondary | 0.84 (.18–3.86) | .83 | 0.73 (.18–2.96) | .66 | 0.76 (.29–1.99) | .58 | 1.01 (.45–2.26) | .99 |
| Indeterminate | 0.27 (.06–1.16) | .08 | 0.51 (.11–2.43) | .40 | 0.37 (.13–1.09) | .07 | 0.31 (.09–1.02) | .054 |

DISCUSSION

There is speculation on the arboviral disease risks posed by the invasion of A. albopictus into southern Europe and the United States [25]. Central to calibrating these risk assessments is an understanding of A. albopictus vector competence. There is conflicting literature on whether A. albopictus is less susceptible than A. aegypti to disseminated DENV infection [26–32]. Furthermore, all of the previous work in this area has used laboratory methods that do not mimic natural infection. Here, in experiments involving direct feeding on viremic patients with dengue, we provide evidence that A. albopictus had a susceptibility to ac-
quiring DENV infection that was similar to that of A. aegypti. However, blood-fed A. albopictus was significantly less likely to become infectious with DENV-2 and DENV-4, suggesting an interspecies difference of epidemiological importance.
When examined for DENV-1, this was also associated with higher levels of sequence drift away from the consensus sequence of the gene encoding envelope protein in the patient’s plasma sample. Plausibly, by being more permissive to DENV replication in its abdominal tissues, *A. albopictus* might contribute to the genetic diversity that exists within DENV populations. Despite *A. albopictus* harboring higher viral burdens in abdominal tissues, significantly fewer *A. albopictus* had infectious saliva resulting from DENV-2 or DENV-4 infections. These data are broadly consistent with previous observations demonstrating that *A. albopictus* was more susceptible to fulminant DENV infection of the mosquito body but that viral dissemination to saliva was less than in *A. aegypti* [18]. That such nuanced interspecies differences can exist is perhaps not surprising, given the literature reporting variation in the susceptibility of *A. aegypti* populations to DENV infection, albeit under laboratory conditions [35–39]. Future studies to understand the basis for these phenotypes, particularly when using field-derived mosquitoes and direct feeding on human viremic hosts, will require large sample sizes to overcome the intrinsic variance in this system.

While our findings have identified interspecies differences in susceptibility to DENV infection, the absolute differences between the mosquito types are nonetheless small. This points to factors other than outright susceptibility to DENV infection as important reasons why *A. albopictus* is a marginal contributor to dengue transmission in urban and peri-urban settings in dengue-endemic countries. Behavioral attributes are also important for vector competence, and thus we speculate that the tendency of *A. aegypti* to live proximate to humans is the major reason why it is the primary DENV vector.

Our study has limitations [18]. First, for practical reasons we measured the phenotype of blood-fed, field-derived mosquitoes at 14 days after feeding; data acquisition at other time points might have led to additional insights. Second, the results were limited in scope to the virus serotypes and genotypes in circulation during the study period, and consequently we acquired sparse data on DENV-2 and DENV-3. Additionally, there may be specific interactions between DENV types and mosquito genotypes that influence infection outcome [40]. Studies in other settings should be encouraged to understand the generalizability of this work.

In summary, we are the first to demonstrate that *A. albopictus* are less likely than *A. aegypti* to develop an infectious phenotype 14 days after direct blood feeding on viremic patients with dengue due to DENV-2 or DENV-4 infection. These results will enable more-accurate parameterization of DENV transmission models in regions where dengue is endemic and those that are at risk for endemicity. In addition, we have confirmed the central importance of plasma viremia in determining the likelihood of mosquito tissue infection, suggesting that interventions that attenuate viremia will have both individual and community benefits.

---

**Supplementary Data**

Supplementary materials are available at *The Journal of Infectious Diseases* online (http://jid.oxfordjournals.org). Supplementary materials consist of data provided by the author that are published to benefit the reader. The posted materials are not copyrighted. The contents of all supplementary data are the sole responsibility of the authors. Questions or messages regarding errors should be addressed to the author.

**References**

1. Simmons CP, Farrar JJ, Nguyen vV, Wills B. Dengue. *N Engl J Med* 2012; 366:1423–32.
2. Bhatt S, Gething PW, Brady OJ, et al. The global distribution and burden of dengue. *Nature* 2013; 496:504–7.
3. Qiu FX, Gubler DJ, Liu JC, Chen QQ. Dengue in China: a clinical review. *Bull World Health Organ* 1993; 71:349–59.
4. Mitchell CJ. The role of Aedes albopictus as an arbovirus vector. *Parasitologia* 1995; 37:109–13.
5. Msetelaar D, Grainger CR, Oei KG, et al. An outbreak of type 2 dengue fever in the Seychelles, probably transmitted by Aedes albopictus (Skuse). *Bull World Health Organ* 1980; 58:937–43.
6. Peng HJ, Lai HB, Zhang QL, et al. A local outbreak of dengue caused by an imported case in Dongguan China. *BMC Public Health* 2012; 12:83.
7. Xu G, Dong H, Shi N, et al. An outbreak of dengue virus serotype 1 infection in Cixi, Ningbo, People’s Republic of China, 2004, associated with a traveler from Thailand and high density of Aedes albopictus. *Am J Trop Med Hyg* 2007; 76:1182–8.
8. Issack MI, Pursem VN, Barkham TM, Ng LC, Inoue M, Manraj SS. Re-emergence of dengue in Mauritius. *Emerg Infect Dis* 2010; 16:716–8.
9. Leroy EM, Nkoghe D, Ollomo B, et al. Concurrent chikungunya and dengue virus infections during simultaneous outbreaks, Gabon, 2007. *Emerg Infect Dis* 2009; 15:591–3.
10. ProMed-mail. Dengue-Japan. ProMed-mail, 2014.
11. Reitter P, Sprenger D. The used tire trade: a mechanism for the worldwide dispersal of container breeding mosquitoes. *J Am Mosq Control Assoc* 1987; 3:494–501.
12. Paupy C, Ollomo B, Kamgang B, et al. Comparative role of Aedes albopictus and Aedes aegypti in the emergence of Dengue and Chikungunya in central Africa. *Vector Borne Zoonotic Dis* 2010; 10:259–66.
13. Medlock JM, Hansford KM, Schaffner F, et al. A review of the invasive Aedes albopictus mosquito in Europe: ecology, public health risks, and control options. *Vector Borne Zoonotic Dis* 2012; 12:435–47.
14. Scolte E-J, Schaffner F. Waiting for the tiger: establishment and spread of the Aedes albopictus mosquito in Europe. *In: Takken W, Knols B, eds. Emerging pests and vector-borne diseases in Europe. Vol 1: Wage-ningen: Wageningen Academic Publishers, 2007;241–60.
15. Hawley WA, Pumpani CB, Brady RH, Craig GB Jr. Overwintering survival of Aedes albopictus (Diptera: Culicidae) eggs in Indiana. *J Med Entomol* 1989; 26:122–9.
16. Thomas SM, Obermayr U, Fischer D, Kreyling J, Beierkuhnlein C. Low-temperature threshold for egg survival of a post-diapause and non-diapause European aedine strain, *Aedes albopictus* (Diptera: Culicidae). *Parasit Vectors* 2012; 5:100.
17. Romi R, Severini F, Toma L. Cold acclimation and overwintering of female Aedes albopictus in Roma. J Am Mosq Control Assoc 2006; 22:149–51.
18. Lambrechts L, Scott TW, Gubler DJ. Consequences of the expanding global distribution of Aedes albopictus for dengue virus transmission. PLoS Negl Trop Dis 2010; 4:e4646.
19. Christofferson RC, Mores CN, Wearing HI. Characterizing the likelihood of dengue emergence and detection in naive populations. Parasit Vectors 2014; 7:282.
20. Ruiz-Moreno D, Vargas IS, Olson KE, Harrington LC. Modeling dynamic introduction of Chikungunya virus in the United States. PLoS Negl Trop Dis 2012; 6:e1937.
21. Wolbers M, Kleinschmidt I, Simmons CP, Donnelly CA. Considerations in the design of clinical trials to test novel entomological approaches to dengue control. PLoS Negl Trop Dis 2012; 6:e1937.
22. Nguyet MN, Duong TH, Trung VT, et al. Host and viral features of human dengue cases shape the population of infected and infectious Aedes aegypti mosquitoes. Proc Natl Acad Sci U S A 2013; 110:9072–7.
23. Hue KD, Tuan TV, Thi HT, et al. Validation of an internally controlled one-step real-time multiplex RT-PCR assay for the detection and quantitation of dengue virus RNA in plasma. J Virol Methods 2011; 177:168–73.
24. Hothorn T, Hornik K, van de Wiel MA, Zeileis A. A Lego system for conditional inference. Am Stat 2006; 60:257–63.
25. Morens DM, Fauci AS. Dengue and hemorrhagic fever: a potential threat to public health in the United States. JAMA 2008; 299:214–6.
26. Sunarto J, Gubler DJ, Nalim S, Eram S, Sulianti Saroso J. Epidemic dengue hemorrhagic fever in rural Indonesia. III. Entomological studies. Am J Trop Med Hyg 1979; 28:717–24.
27. Rosen L, Roseboom LE, Gubler DJ, Lien JC, Chaniotis BN. Comparative susceptibility of mosquito species and strains to oral and parenteral infection with dengue and Japanese encephalitis viruses. Am J Trop Med Hyg 1985; 34:603–15.
28. Alto BW, Reiskind MH, Lounibos LP. Size alters susceptibility of vectors to dengue virus infection and dissemination. Am J Trop Med Hyg 2008; 79:688–95.
29. Vazeille M, Rosen L, Mousson L, Failloux AB. Low oral receptivity for dengue type 2 viruses of Aedes albopictus from Southeast Asia compared with that of Aedes aegypti. Am J Trop Med Hyg 2003; 68:203–8.
30. Whitehead RH, Yull TM, Gould DJ, Simasathien P. Experimental infection of Aedes aegypti and Aedes albopictus with dengue viruses. Trans R Soc Trop Med Hyg 1971; 65:661–7.
31. Chen WJ, Wei HL, Hsu EL, Chen ER. Vector competence of Aedes albopictus and Ae. aegypti (Diptera: Culicidae) to dengue 1 virus on Taiwan: development of the virus in orally and parenterally infected mosquitoes. J Med Entomol 1993; 30:524–30.
32. Higgs S, Vanlandingham DL, Klingler KA, et al. Growth characteristics of ChimeriVax-Den vaccine viruses in Aedes aegypti and Aedes albopictus from Thailand. Am J Trop Med Hyg 2006; 75:986–93.
33. Mercado-Curiel RF, Esquinca-Aviles HA, Tovar R, Diaz-Badillo A, Camacho-Nuez M, Munoz Mde L. The four serotypes of dengue recognize the same putative receptors in Aedes aegypti midgut and Ae. albopictus cells. BMC Microbiol 2006; 6:85.
34. Richards SL, Anderson SL, Alto BW. Vector competence of Aedes aegypti and Aedes albopictus (Diptera: Culicidae) for dengue virus in the Florida Keys. J Med Entomol 2012; 49:942–6.
35. Bennett KE, Olson KE, Munoz Mde L, et al. Variation in vector competence for dengue 2 virus among 24 collections of Aedes aegypti from Mexico and the United States. Am J Trop Med Hyg 2002; 67:85–92.
36. Lozano-Fuentes S, Fernandez-Salas I, de Lourdes Munoz M, et al. The neovolcanic axis is a barrier to gene flow among Aedes aegypti populations in Mexico that differ in vector competence for Dengue 2 virus. PLoS Negl Trop Dis 2009; 3:e468.
37. Sylla M, Bosio C, Urdaneta-Marquez L, Ndiaye M, Black WC 4th. Gene flow, subspecies composition, and dengue virus-2 susceptibility among Aedes aegypti collections in Senegal. PLoS Negl Trop Dis 2009; 3:e408.
38. Gubler DJ, Nalim S, Tan R, Saipan H, Sulianti Saroso J. Variation in susceptibility to oral infection with dengue viruses among geographic strains of Aedes aegypti. Am J Trop Med Hyg 1979; 28:1045–52.
39. Sim S, Jupatanakul N, Ramirez JL, et al. Transcriptomic profiling of diverse Aedes aegypti strains reveals increased basal-level immune activation in dengue virus-refractory populations and identifies novel virus-vector molecular interactions. PLoS Negl Trop Dis 2013; 7:e2295.
40. Lambrechts L, Chevillon C, Albright RG, et al. Genetic specificity and potential for local adaptation between dengue viruses and mosquito vectors. BMC Evol Biol 2009; 9:160.
Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:
Whitehorn, J; Duong, THK; Nguyet, MN; Nguyen, HL; Kyrylos, PP; Carrington, LB; Chau, NBT; Nguyen, THQ; Long, VT; Dui, LT; Nguyen, TT; Tai, THL; Chau, VVN; Wills, B; Wolbers, M; Simmons, CP

Title:
Comparative Susceptibility of Aedes albopictus and Aedes aegypti to Dengue Virus Infection After Feeding on Blood of Viremic Humans: Implications for Public Health

Date:
2015-10-15

Citation:
Whitehorn, J; Duong, THK; Nguyet, MN; Nguyen, HL; Kyrylos, PP; Carrington, LB; Chau, NBT; Nguyen, THQ; Long, VT; Dui, LT; Nguyen, TT; Tai, THL; Chau, VVN; Wills, B; Wolbers, M; Simmons, CP, Comparative Susceptibility of Aedes albopictus and Aedes aegypti to Dengue Virus Infection After Feeding on Blood of Viremic Humans: Implications for Public Health, JOURNAL OF INFECTIOUS DISEASES, 2015, 212 (8), pp. 1182 - 1190

Persistent Link:
http://hdl.handle.net/11343/56666

File Description:
Published version