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Aedes (Stegomyia) albopictus (Skuse): A Potential Vector of Zika Virus in Singapore

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Abstract

Background: Zika virus (ZIKV) is a little known arbovirus until it caused a major outbreak in the Pacific Island of Yap in 2007. Although the virus has a wide geographic distribution, most of the known vectors are sylvatic *Aedes* mosquitoes from Africa where the virus was first isolated. Presently, *Ae. aegypti* is the only known vector to transmit the virus outside the African continent, though *Ae. albopictus* has long been a suspected vector. Currently, *Ae. albopictus* has been shown capable of transmitting more than 20 arboviruses and its notoriety as an important vector came to light during the recent chikungunya pandemic. The vulnerability of Singapore to emerging infectious arboviruses has stimulated our interest to determine the competence of local *Ae. albopictus* to transmit ZIKV.

Methodology/Principal Findings: To determine the competence of *Ae. albopictus* to ZIKV, we orally infected local mosquito strains to a Ugandan strain virus. Fully engorged mosquitoes were maintained in an environmental chamber set at 29°C and 80–85%RH. Twelve mosquitoes were then sampled daily from day one to seven and on day 10 and 14 post infection (pi). Zika virus titre in the midgut and salivary glands of each mosquito were determined using tissue culture infectious dose₅₀ assay, while transmissibility of the virus was determined by detecting viral antigen in the mosquito saliva by qRT-PCR. High dissemination and transmission rate of ZIKV were observed. By day 7-pi, all mosquitoes have disseminated infection and 73% of these mosquitoes have ZIKV in their saliva. By day 10-pi, all mosquitoes were potentially infectious.

Conclusions/Significance: The study highlighted the potential of *Ae. albopictus* to transmit ZIKV and the possibility that the virus could be established locally. Nonetheless, the threat of ZIKV can be mitigated by existing dengue and chikungunya control program being implemented in Singapore.

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Introduction

The Asian tiger mosquito, *Aedes (Stegomyia) albopictus* (Skuse), is considered a vector or potential vector of several pathogens of human and veterinary importance. Viral isolation and vector competence studies have shown that this mosquito is an efficient vector of more than 20 arboviruses [1]. Due to its biological and ecological plasticity, this notoriously invasive species has a wide geographic distribution. At present, aside from its tropical Asian home, *Ae. albopictus* can be found in temperate Asian countries, in tropical and temperate Americas, Europe, Middle East, the Pacific islands, Australia and Africa [2,3,4,5]. The global spread of *Ae. albopictus* is mainly caused by human activities, such as increase in intercontinental trade, especially in the last three decades [1].

In places where *Ae. aegypti* and *Ae. albopictus* co-exist, *Ae. albopictus* was considered second only to *Ae. aegypti* in terms of importance as vector of dengue and chikungunya [6,7]. However, its notoriety as an important vector came to light during the recent unprecedented global outbreak of chikungunya. According to Tssetsarkin et al. [8], a mutation in the envelope gene of chikungunya virus (CHIKV) enhances the virus infectivity and transmissibility in *Ae.*

albopictus. The continual global expansion of *Ae. albopictus* is a growing concern as this mosquito may alter the transmission dynamics of arboviral diseases and increase the risks of humans to mosquito-borne viral infections [3,9]. This has stimulated increased interest to determine the extent of pathogens this mosquito can transmit.

Zika virus (ZIKV), a little known arbovirus, has gained prominence when it caused a large scale epidemic in the Pacific Island in 2007 [10,11]. It is a member of the genus *Flavivirus* of the Family Flaviviridae [12]. The virus is a positive single stranded RNA virus with a 10,794 nucleotide genome that is closely related to the Spondweni virus [13,14,15]. It was first isolated from a febrile sentinel monkey in Uganda in 1947 [15], but human ZIKV infection was first reported in 1964 [16]. The virus causes dengue-like syndromes such as rash, fever, arthralgia, headache and peri-orbital pain [11,16].

To date, only *Aedes* mosquitoes have been known to transmit ZIKV. In Africa, the virus was isolated from both sylvatic and peri-domestic mosquitoes: *Ae. africanus*, *Ae. apicocoxargenteus*, *Ae. luteocephalus*, *Ae. fuscifer*, *Ae. vitattus* and *Ae. aegypti* [17,18,19,20,21,22]. In Asia, ZIKV was only isolated from a pool

Author Summary

The prominence of *Ae. albopictus* as a major vector came to light during the global pandemic of chikungunya. The continual global expansion of *Ae. albopictus* coupled with data from competence studies that show its ability to transmit more than 20 arboviruses has fuelled growing concerns that this mosquito may alter the epidemiology of arbovirus worldwide and increase the risks of humans to mosquito-borne diseases. This has stimulated our interest to determine the extent of pathogens *Ae. albopictus* can transmit, especially arboviruses with high epidemic potential. The current study showed that local *Ae. albopictus* are susceptible to ZIKV. By day 7 pi, all mosquitoes that were orally infected were found to have disseminated infections and 73% have ZIKV in their saliva. This study highlighted the potential of *Ae. albopictus* to transmit ZIKV. The local transmission of ZIKV in Singapore is plausible due to the presence of susceptible vectors, an immunologically naive and vulnerable human population, and the travel hub status of Singapore. Nonetheless, the threat of ZIKV can be mitigated by existing dengue and chikungunya control program being implemented in Singapore.

of *Ae. aegypti* caught from shop houses in the State of Pahang in Peninsular Malaysia [22]. During the ZIKV outbreak in Yap Island in 2007, no virus was isolated from any of the mosquitoes caught. However, based on epidemiological evidences, *Ae. hensilli* was suspected to be the vector responsible for the outbreak [11]. In 2010, a case of ZIKV involving a three year old child was reported in Kampong Speu Province in Cambodia, however, the vector responsible was not known [23].

Identification of vectors and potential vectors of ZIKV or any other mosquito-borne diseases in a given geographical area has important implication when it comes to disease outbreak control. It is imperative that vectors are identified, so that a holistic and sound vector control program can be formulated. To date, *Ae. aegypti* is the only vector of ZIKV identified in Southeast Asia [22] and data on mosquito-ZIKV interactions have also been confined to this mosquito [24,25]. Our recent study has also showed that local *Ae. aegypti* strains are highly susceptible to ZIKV and viral dissemination rates reflect that observed for a local, highly epidemic DENV-2 strain [26].

In the light of continuous global niche expansion of *Ae. albopictus*, coupled with its catholic feeding behaviour, ecological adaptability and propensity to support a wide range of arboviruses, it is important to determine its competence to transmit pathogens with high epidemic potential such as ZIKV. The current study describes ZIKV infection in our local *Ae. albopictus*.

Materials and Methods

Mosquitoes

Aedes albopictus, used for the experimental infection, was derived from eggs collected during weekly ovitrap surveillance study as previously described [26]. Mosquitoes were colonized under standard insectary conditions as described by Li and colleagues [26]. In order to obtain enough number of mosquitoes for the study, F₃ generations were used.

Virus

Ugandan MR766 ZIKV strain obtained from the American Type Culture Collection (Manassas, VA, USA) was used to infect

the mosquitoes. The stock virus used in the current study was passaged thrice in Vero cell line prior to the infectious feed [26].

Oral infection of mosquitoes

Five to seven-day old female mosquitoes ($n=120$) were transferred to 0.5 L containers and starved for 24 hours prior to the infectious blood meal. The blood meal consisted of 1:1 100% swine-packed RBC (Innovative Research, USA) and a fresh ZIKV suspension, at a final concentration of 7.5 Log₁₀ tissue culture infectious dose₅₀/mL ((Log₁₀TCID₅₀/ml). Adenosine Triphosphate (Fermentas, USA), at a final concentration of 3 mM, was added to the blood meal as a phagostimulant. Mosquitoes were fed with an infectious blood meal that was warmed to 37°C using a Hemotek membrane feeding system (Discovery Workshops, Lancashire, United Kingdom) housed. After thirty minutes, mosquitoes were cold anesthetized and fully engorged females were transferred to 300 ml ca. paper cups and were maintained in an environmental chamber (Sanyo, Japan) at 29°C and 70–75% RH with a 12 h/12 h L:D cycle and provided with 10% sugar/vitamin B complex *ad libitum*. All experiments were carried out in an arthropod containment level 2 (ACL-2) facility.

Mosquito processing

To determine the ZIKV infection and dissemination rates in *Ae. albopictus*, 12 mosquitoes were sampled daily from day one to seven, and subsequently on day 10 and 14 post infection (pi). Saliva was collected using the forced salivation technique as previously described [27] with modification. The proboscis of each mosquito, with its legs and wings, was inserted into a micropipette tip containing 10 µl of M199 and allowed to salivate. After 45 minutes, the Medium 199 containing the saliva from each mosquito was transferred into microcentrifuge tubes containing 100 µl of M199. The midgut and salivary glands of each mosquito were processed as described by Li and colleagues [26]. Briefly, the midgut and the salivary glands were homogenized using stainless steel grinding balls (Retsch, Germany) in a MM301 mixer mill (Retsch, Germany) set at frequency of 12/sec for 1 min. The supernatant of the homogenate was applied in the viral titre assay. All dissecting needles were dipped in 80% ethanol and cleaned before being re-used. All experiments were conducted inside an ACL-2 facility.

Quantitative reverse transcriptase-polymerase chain reaction (qRT-PCR) assay

Total RNA was isolated from saliva using the QIAamp Viral Mini Kit (Qiagen, Germany) following manufacturer's recommendations. ZIKV in saliva was detected using a one-step real-time reverse transcriptase-polymerase chain reaction (qRT-PCR) as previously described [10].

Tissue Culture Infectious Dose₅₀ (TCID₅₀) assay

Viral titres from the midgut and salivary glands were determined with tissue culture infectious dose₅₀ assay, an endpoint dilution technique, using Vero cells as described by Higgs et al. [28].

Data analysis

The infection rate at each sampling day was determined by the percentage of infected midguts, while dissemination rate was calculated by dividing the number of infected salivary glands by the total number of mosquitoes with infected midguts. On the other hand, transmission rate was calculated by dividing the number of positive saliva by the number of infected salivary

glands. Kolmogorov-Smirnov tests indicated that the data did not conform to conditions of normality, hence non-parametric analyses were performed. Kruskal-Wallis tests were used to determine differences in viral titres in midguts and salivary glands between days post-infection. When a significant difference was detected, Mann-Whitney *U*-tests were performed to determine which day differed. All analyses were performed in Minitab.

Results

Oral susceptibility of *Ae. albopictus* to ZIKV

From day 3-pi onwards, when blood had been completely digested, all midguts were positive for ZIKV, except for day 7- and 10-pi (Table 1). The presence of viable ZIKV in the salivary glands was first observed on day 3-pi in three mosquitoes. By day 5-pi, half of the mosquitoes sampled showed disseminated infection. From day 7-pi onwards, all mosquitoes were found to have ZIKV in their salivary glands (Table 1).

Transmission was first observed on day 4 after the infectious blood meal and transmission rates were observed to increase at each sampling days. By day 10-pi onwards, ZIKV RNA was found in all saliva tested (Table 1).

ZIKV midgut and salivary gland titres

Figure 1 presents ZIKV midgut and salivary gland titres at different days pi. A significant difference in midgut ZIKV titres was observed between different days pi (Kruskal-Wallis test, $P=0.017$). Midgut viral titres at days 3-, 4-, and 5-pi ($>5.15 \text{ Log}_{10}\text{TCID}_{50}/\text{ml}$) were found to be significantly higher when compared to viral titre at day 14-pi (Mann-Whitney test, $P<0.05$) (Figure 1a). From days 6-pi onwards, viral titres fluctuated between $4.94 \text{ Log}_{10}\text{TCID}_{50}/\text{ml}$ to $4.21 \text{ Log}_{10}\text{TCID}_{50}/\text{ml}$ with no significant difference observed (Kruskal-Wallis test: $P=0.81$).

In contrast, increasing ZIKV titre was observed in the salivary glands of *Ae. albopictus* (Figure 1b) (Kruskal-Wallis test, $P<0.0001$). Although the differences in ZIKV salivary gland titres from day 3-pi ($2.54 \text{ Log}_{10}\text{TCID}_{50}/\text{ml}$) to day 6-pi ($3.26 \text{ Log}_{10}\text{TCID}_{50}/\text{ml}$) were comparable (Mann-Whitney test, $P\geq 0.09$), the viral load increased significantly (Mann-Whitney test, $P<0.05$) by day 10-pi onwards ($>5.96 \text{ Log}_{10}\text{TCID}_{50}/\text{ml}$).

Discussion

Emerging and re-emerging mosquito-borne diseases are considered to be major threats to global health in both developing and

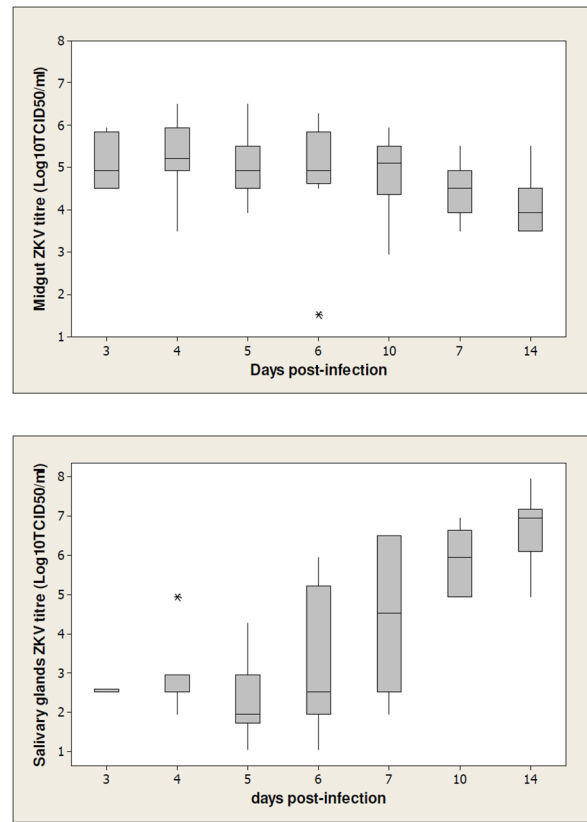


Figure 1. Box plots of ZIKV midgut and salivary glands titres at different days post-infection. Each box represents the median (horizontal bar), the interquartile range (box) and the full range (“whiskers”). Outliers are denoted by asterisk (*). doi:10.1371/journal.pntd.0002348.g001

developed countries. Their tendency of spreading outside their known geographic range and causing large scale epidemics has been clearly demonstrated during the recent global epidemic of CHIKV [29].

Zika is a neglected tropical disease, and like CHIKV, interest in ZIKV epidemiology was limited until recently, when its high epidemic potential was demonstrated during a large-scale outbreak in the Pacific Island of Yap in 2007 [10,11,13,30].

Table 1. Infection, dissemination and transmission rates for *Ae. albopictus* orally fed with ZIKV and held at 29°C at various days post-infection.

Days post- infection	Infection rate		Dissemination rate		Transmission rate	
	No. positive MG (number sampled)	Percent	No. positive SG (number sampled)	Percent	No. positive saliva (number sampled)	Percent
3	12 (12)	100	3 (12)	25	0 (3)	0
4	12 (12)	100	7 (12)	58.3	1 (7)	14.3
5	12 (12)	100	6 (12)	50	2 (6)	33.3
6	12 (12)	100	9 (12)	75	4(9)	44.4
7	11 (12)	91.7	11(11)	100	8 (11)	72.7
10	10 (12)	83.3	10(10)	100	10(10)	100
14	12 (12)	100	12(12)	100	12(12)	100

MG = midgut; SG = salivary gland.
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During the outbreak that lasted four months, more than 70% of the island's population was affected [11]. Like many vector-borne diseases, the absence of vaccines and specific treatment against ZIKV means prevention and control relies heavily on vector control. Therefore, key information such as the identity of the vector, its bionomics, distribution, and density are needed in order to develop and implement sound mosquito control program. To date, little is known about the vectors of ZIKV outside Africa, except for *Ae. aegypti*.

The overlapping geographic distribution of *Ae. albopictus* with that of ZIKV has stimulated our interest to determine the potential of this mosquito to transmit ZIKV. Furthermore, the widespread distribution of *Ae. albopictus* in Singapore and large-scale local outbreaks of chikungunya in 2008–09 attest to the country's vulnerability to emerging mosquito-borne diseases [31]. The potential of these diseases to be established locally is accentuated by the country's reputation as a popular commercial and tourist hub, high dependency on migrant workers, tropical climate, dense human population, and the presence of potential mosquito vectors. Recently, we have shown the potential for ZIKV outbreak in Singapore by *Ae. aegypti* [26]. The study showed local *Ae. aegypti* are highly susceptible to ZIKV, with a short extrinsic incubation period (EIP) of five days.

Our current study showed that Singapore's *Ae. albopictus* mosquitoes are susceptible to ZIKV, with high dissemination and transmission rates observed. By day 4-pi, 58% ($n = 7/12$) of the infected mosquitoes have disseminated infection and of these, three (43%) had ZIKV in their saliva. By day 7-pi, all infected mosquitoes are capable of transmitting the virus. A short EIP and high ZIKV salivary gland titres were also observed when we infect our local *Ae. aegypti* with ZIKV [26]. In that study, it took five and ten days post infectious blood meal to achieve a 62% and 100% dissemination rate, respectively. However, it does not mean that *Ae. albopictus* is more susceptible to ZIKV than *Ae. aegypti*, rather it could be due the amount of virus used to infect *Ae. aegypti* was lower (6.95 Log₁₀ TCID₅₀/mL) compared to the current study (7.52 Log₁₀ TCID₅₀/mL).

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Although *Ae. albopictus* has long been a suspected vector of ZIKV [32], to our knowledge, this is the first report on the potential of this mosquito species to transmit ZIKV. However, further studies (such as entomological surveillance in endemic areas) are needed to validate our findings to ascertain the vectorial status of *Ae. albopictus* for ZIKV in nature especially when both *Ae. Aegypti* and *Ae. albopictus* are geographically sympatric. Nonetheless, data from our previous [26] and current studies suggest that both *Ae. aegypti* and *Ae. albopictus* will have significant roles in the transmission of ZIKV, should the virus be introduced in Singapore or in places where these mosquitoes abound. Recent phylogenetic analysis has identified two major lineages of ZIKV, an African and Asian lineage [17]. The strain used in our current study is the prototype MR766 Ugandan strain belonging to the former lineage. It will also be very interesting to study the strains belonging to the Asian lineage; unfortunately, at the time of the study, we only had access to the prototype strain.

Given the vulnerability of Singapore to ZIKV and presence of the virus in neighbouring countries, the Environmental Health Institute screened febrile cases not attributable to DENV and CHIKV for ZIKV and other arboviruses. To date, none was found positive for arboviruses other than DENV and CHIKV. Based on the information gathered from this study, the threat of ZIKV can be addressed by existing dengue and chikungunya control program being implemented in Singapore.

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Author Contributions

Conceived and designed the experiments: CHT LCN. Performed the experiments: CHT PSJW MZIL CSC. Analyzed the data: CHT PSJW MZIL CSC. Contributed reagents/materials/analysis tools: CHT PSJW MZIL CSC. Wrote the paper: CHT PSJW.

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