

Demonstration of resistance to satyrization behavior in *Aedes aegypti* from La Réunion island

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Abstract – *Aedes aegypti* and *Aedes albopictus* are competent vectors of arboviruses such as dengue and chikungunya viruses which co-exist in some areas, including La Réunion island, Indian Ocean. A type of reproductive interference called satyrization has been described in sympatric species where dominant species mating fails to produce hybrids and thus reduces the fitness and tends to control the spread of the other species. Here, we investigated satyrization in laboratory experiments to provide insights on the potential impact on *Ae. aegypti* of a control campaign including a sterile insect technique component against *Ae. albopictus*. Different mating crosses were used to test sympatric, conspecific-interspecific and allopatric effects of irradiated and non-irradiated male *Ae. albopictus* on female *Ae. aegypti*, including in a situation of skewed male ratio. Our results suggest that there was only a low level of satyrization between sympatric populations of *Ae. aegypti* and *Ae. albopictus* colonized from La Réunion island. A male *Ae. albopictus* to female *Ae. aegypti* ratio of 3:1 did not increase the level of satyrization. Female *Ae. aegypti* previously mated to male *Ae. albopictus* were not prevented from being inseminated by conspecific males. A satyrization effect was not seen between allopatric *Ae. albopictus* and *Ae. aegypti* strains from La Réunion Island either. The tested *Ae. aegypti* strain from La Réunion island has therefore developed full resistance to satyrization and so releasing sterile male *Ae. albopictus* may not suppress *Ae. aegypti* populations if an overflowing of irradiated male *Ae. albopictus* leads to similar results. The management strategy of two competent species in a sympatric area is discussed.

Key words: *Aedes albopictus*, Satyrization, Sterile insect technique, Mating.

Résumé – Démonstration de la résistance au comportement de satyrisation chez *Aedes aegypti* de l'île de La Réunion. *Aedes aegypti* et *Aedes albopictus* sont des vecteurs compétents d'arbovirus, tels que les virus de la dengue et du chikungunya, qui coexistent dans certaines régions, y compris à La Réunion, dans l'Océan Indien. Une interférence sur la reproduction appelée satyrisation a été décrite chez les espèces sympatriques où l'accouplement d'une espèce dominante ne produit pas d'hybrides et donc réduit le succès reproducteur et tend à contrôler la propagation des autres espèces. Ici, nous avons étudié la satyrisation dans des expériences de laboratoire pour fournir des informations sur l'impact potentiel sur *Ae. aegypti* d'une campagne de lutte contre *Ae. albopictus* basée sur la technique de l'insecte stérile. Différents croisements d'accouplements ont été utilisés pour tester les effets sympatriques, conspécifiques-interspécifiques et allopatriques de mâles *Ae. albopictus* irradiés et non irradiés sur les femelles *Ae. aegypti*, y compris dans une situation de ratio masculin asymétrique. Nos résultats suggèrent qu'il n'y avait qu'un faible niveau de satyrisation entre les populations sympatriques d'*Ae. aegypti* et *Ae. albopictus* colonisées à La Réunion. Un rapport mâles *Ae. albopictus* sur femelles *Ae. aegypti* de 3 contre 1 n'a pas augmenté le niveau de satyrisation. Les femelles *Ae. aegypti* déjà accouplées à un mâle *Ae. albopictus* n'ont pas été empêchées d'être inséminées par des mâles conspécifiques. Aucun effet de satyrisation n'a été observé entre les souches allopatriques d'*Ae. albopictus* et d'*Ae. aegypti* de La Réunion. La souche testée d'*Ae. aegypti* de La Réunion a donc développé une résistance totale à la satyrisation et en conséquence la libération de mâles stériles d'*Ae. albopictus* peut ne pas supprimer les populations d'*Ae. aegypti* si des lâchers massifs de mâles irradiés d'*Ae. albopictus* conduisent à des résultats similaires. La stratégie de gestion de deux espèces compétentes dans une zone sympatrique est discutée.

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Introduction

Aedes albopictus (Skuse), the Asian tiger mosquito, has been invasive in many parts of the world since the 1980s [10]. *Aedes aegypti* (Linnaeus), also called the yellow fever mosquito and originating from Africa, is a highly invasive, medically important mosquito species that has received a considerable increase in attention after being linked to the Zika outbreak in Brazil in 2015 [19]. Both species transmit several arboviral diseases including dengue, chikungunya, yellow fever, and Zika [37]. Dengue alone is estimated to infect 390 million people per year, causing 96 million cases with clinical manifestations [11]. *Aedes albopictus* (Skuse) and *Ae. aegypti* are the most important vectors of the epidemic forms of dengue and chikungunya viruses to humans [36, 44]. *Aedes albopictus* is also responsible for the major chikungunya outbreak in the islands of the South-West Indian Ocean, including La Réunion island (an overseas department of France) between 2005 and 2007 [20] and for the more recent dengue outbreaks according to the local health authority (Agence Régionale de Santé Océan Indien) and the regional office of Santé publique France on La Réunion island [50].

The heavy reliance on insecticides to control adult *Aedes* mosquitoes, especially during disease outbreaks, has led to the emergence of widespread resistance to these chemicals, making traditional control strategies insufficient to achieve adequate reduction of vector populations. The use of insecticides is also inefficient against these container-breeding mosquito species with disseminated larval habitats. Therefore, complementary vector control methods are needed to enhance existing efforts [49]. Amongst those being advocated is the sterile insect technique (SIT) [12], a species-specific and environmentally-friendly pest population control method that relies on maintaining a continuous production and repeated release of overflooding numbers of sterile males [35] that can outcompete their wild counterparts within the target area [21], and induce sterility in wild females. A feasibility assessment of an area-wide integrated pest management (AW-IPM) program with an SIT component is ongoing on La Réunion island, where *Ae. albopictus* co-occurs with *Ae. aegypti*. The first releases of sterile male *Ae. albopictus* mosquitoes to study their behavior in urban areas were recently authorized by a prefectural order to the Institute for Research for Development (IRD) [2]. To successfully and cost-effectively apply the SIT in an area, it is recommended to target one species at a time [1, 21]. For example, where *Ae. aegypti* and *Ae. albopictus* are sympatric and are both competent vectors of human viruses, one would consider the best approach to guarantee successful suppression or elimination of both species. The best scenario would be the release of one species which was able to also readily mate with and induce sterility in both species [27]. This phenomenon is known as reproductive interference or satyrization, in which males of one species mate with and sterilize females of another species, and in this way contribute to its displacement from the shared area [45].

Satyrization was proposed as a possible mechanism for the displacement of *Ae. aegypti* by *Ae. albopictus* in Florida in the late 1980s [43, 48]. Satyrization and other factors such as larval competition, climate, and socioeconomic factors have

influenced the distribution dynamics of the two species worldwide. A rapid decline in *Ae. aegypti* in the south eastern USA and Bermuda, for example, was associated with the invasions of *Ae. albopictus* [33, 38, 43]. The same mechanism was suggested to explain the spread of invasive *Ae. albopictus* in Athens, Greece and the reduction in distribution of the native *Ae. cretinus* [25]. Displacement of *Ae. albopictus* by *Ae. aegypti* has also occurred in certain tropical cities and regions in Asia including Bangkok, Kuala Lumpur, Manila, and southern Taiwan, and more recently in the Colombian port of Leticia (reviewed in [39]). However, it is suggested that resistance to satyrization may evolve more rapidly in female *Ae. aegypti* populations sympatric to *Ae. albopictus* than in females from allopatric populations [6, 40]. Bagny et al. [3] reported a progressive decrease in *Ae. aegypti* distribution on La Réunion island since the 1900s where it was mainly found in rock holes with *Ae. albopictus* in ravines located on the driest west coast of the island and was absent from artificial containers. The same study suggested that the dwindling *Ae. aegypti* densities observed during the 1950s was due to ecological factors including a competitive interaction between *Ae. aegypti* and *Ae. albopictus* combined with vector control campaigns during these years. However, to the best of our knowledge, the competitive interaction between La Réunion island strains has not yet been investigated.

With both *Ae. aegypti* and *Ae. albopictus* being competent vectors of dengue and several other arboviruses including chikungunya and Zika viruses [16, 38, 44], it is important to investigate whether or not the release of irradiated *Ae. albopictus* males would affect *Ae. aegypti* populations [17], in the framework of the SIT project on La Réunion island. Our satyrization experiments provide insights on the potential impact on *Ae. aegypti* of a control campaign against *Ae. albopictus* on La Réunion island. Different mating crosses were used to test sympatric, conspecific-interspecific, and allopatric effects of male *Ae. albopictus* on female *Ae. aegypti* including in a situation of skewed male ratio. A preprint of this work was published on February 12, 2020 [41].

Materials and methods

Mosquito strains, rearing, and irradiation conditions

The *Ae. aegypti* strain used in this study originated from field collections on La Réunion island. The strain was colonized in the laboratory by the Institute of Research for Development (IRD) for five generations (F5) before being transferred to the Insect Pest Control Laboratory (IPCL) of the joint FAO/IAEA division of Nuclear Sciences and Applications in 2016. The *Ae. albopictus* La Réunion island strain was maintained at the IPCL from 2009 for several generations before these experiments were performed in 2016. In order to perform different mating crosses, *Ae. albopictus* strains from China (Guangzhou wild type strain, provided by Wolbaki since 2015) and Italy (Rimini strain, provided by Centro Agricoltura Ambiente (CAA) since 2012) were maintained in parallel.

All the strains were reared in 30 × 40 × 10 cm trays at a density of 1 first instar larvae (L1) per mL under controlled

temperature, humidity, and lighting conditions ($T = 26 \pm 2$ °C, 70 ± 10 RH%, 12:12 h light: dark, including 1 h dawn and 1 h dusk). Larvae were fed with IAEA larval diet following the protocol described in the Guidelines for routine rearing [22]. Pupae were collected and separated using a glass plate sorter [23].

Male pupae of all strains were irradiated between 36 h and 44 h of age with 40 Gy using a Gamma Cell 220 (Nordion Ltd., Kanata, ON, Canada) emitting a dose rate of 90 Gy/min.

Experiment 1. Sympatric cross-mating between *Ae. aegypti* and *Ae. albopictus* strains from La Réunion island

Male *Ae. aegypti* and *Ae. albopictus* were irradiated and crossed with female *Ae. albopictus* and *Ae. aegypti*, respectively. Non-irradiated males of each species were also crossed with female mosquitoes of the other species. Non-irradiated male *Ae. aegypti* and *Ae. albopictus* were also crossed with virgin female *Ae. aegypti* and *Ae. albopictus*, respectively, and used as controls.

Three replicates were performed for each cross with 50 males and 50 females transferred to $15 \times 15 \times 15$ cm cages (MegaView Science Co. Ltd., Taiwan) when they were three days old for a period of seven days to ensure enough time was allowed for mating. Females were offered a defibrinated porcine bloodmeal using sausage casings (Grade Specification: 3 26 NC, EDICAS Co., Ltd.) for 1 h (2×30 min with 10 min reheating of the blood sausage in between feedings) on two consecutive days when they were 5–6 days old. Each of the females was transferred to an individual drosophila tube containing a cone of seed germination paper (Grade 6, Size: 580×580 mm, weight: 145 g/m^2 , Sartorius Stedim Biotech) and 10 mL of water. Females were given three days to lay eggs and then dissected to determine their insemination status under a stereomicroscope ($40\times$ magnification). Before dissection, females were kept in labelled 50 mL Falcon tubes (VWR International GmbH, Darmstadt, Germany) in a refrigerator at 4 °C, and samples held in a cold box containing ice to avoid desiccation while other samples were being dissected.

Virgin *Ae. aegypti* and *Ae. albopictus* females of the same batch were offered blood meals and were also placed in individual egg tubes to assess their egg capacity.

All eggs were collected, dried for seven days in the laboratory and allowed to hatch for 20 h with a hatching solution made of 0.25 g of CM 0001 Nutrient Broth (Oxoid, Hampshire, UK) and 0.05 g of yeast diluted in 0.7 L of deionized water [51].

The number of female mosquitoes still alive after seven days together with males was recorded for each replicate, and the survival rate was compared with survival in the conspecific *Ae. aegypti* mating control.

Experiment 2. Effect of male *Ae. albopictus* density (ratio) on female *Ae. aegypti* mating success

To assess whether an increase in male to female ratio would favour satyriation, female *Ae. aegypti* mosquitoes from La Réunion island were allowed to mate with male

Ae. albopictus in a male: female ratio of 3:1 corresponding to 75 male *Ae. albopictus* and 25 female *Ae. aegypti*. Three days-old males and females were held in $15 \times 15 \times 15$ cm BugDorm cages (MegaView Science Co. Ltd., Taiwan) for seven days. The crosses were performed using irradiated males (four replicates) and non-irradiated males (seven replicates). Females were then dissected to check their insemination status, as described above. This experiment was carried out in parallel with Experiment 1 and results of mating crosses could be compared to the insemination rates obtained with a 1:1 male: female ratio (50 males \times 50 females).

Experiment 3. Pre-exposure effect on mating success (interspecific-conspecific treatment)

To assess whether female *Ae. aegypti* that were pre-exposed to male *Ae. albopictus* could re-mate with their conspecific male *Ae. aegypti*, 4–5 days-old female *Ae. aegypti* (La Réunion island strain) were pre-exposed to irradiated and non-irradiated male *Ae. albopictus* (La Réunion island strain). Females were removed after seven days and were offered to irradiated and non-irradiated male *Ae. aegypti* (La Réunion island strain) for another seven days. Females were then blood fed for two consecutive days and eggs collected before females were dissected under a microscope to determine insemination status. We hypothesized that if female *Ae. aegypti* were inseminated by non-irradiated *Ae. albopictus* males, they would not be able to lay fertile eggs even if they had a blood meal.

Fifty adults were included in each cross in a 1:1 male: female ratio in Bugdorm cages ($15 \times 15 \times 15$ cm).

An experimental design was used which ensured that female *Ae. aegypti* that were pre-exposed to male *Ae. albopictus* had not been inseminated (Table 1), and female insemination and egg hatch rates were observed.

Experiment 4. Effect of geographic origin on mating success (allopatric crosses)

Since we observed resistance to satyriation against *Ae. albopictus* in the sympatric cross experiment, we explored the allopatric effect to better understand the mating behavior of the La Réunion island strain of *Ae. aegypti*. Several crosses were performed using irradiated and non-irradiated males of the *Ae. albopictus* strains from La Réunion island, China (Guangzhou wild type strain) and Italy (Rimini strain).

Female *Ae. aegypti* mosquitoes from La Réunion island were crossed for seven days with non-irradiated or irradiated male *Ae. albopictus* mosquitoes from either La Réunion island, China or Italy, in 14 combinations. In each interspecific treatment (*Ae. aegypti* female \times *Ae. albopictus* male), 30 males and 30 females, three-days old, from each strain were housed in $15 \times 15 \times 15$ cm BugDorm cages. In addition, conspecific control crosses (*Ae. albopictus* female \times *Ae. albopictus* male, and *Ae. aegypti* female \times *Ae. aegypti* male, La Réunion island strain) were carried out using the same number of males and females for each strain. The number of females that were successfully inseminated was recorded (as described for Experiment 1) and results were compared to the interspecific

Table 1. Experimental design of the interspecific-conspecific treatments.

Cross	Status	Number of replicats	Male	Female	Irradiated males	Fecundity and fertility checks	Dissection for insemination status check
C1	Control	5	AEG	AEG	Yes	No	Yes
C2	Control	4	AEG	AEG	No	No	Yes
C3	Control	4	ALBO	ALBO	Yes	No	Yes
C4	Control	4	ALBO	ALBO	No	No	Yes
C5	Pre-exposed	4	ALBO	AEG	Yes	No	Yes
C6	Pre-exposed	4	ALBO	AEG	No	No	Yes
C7	Control	5	AEG	AEG	No	Yes	Yes
C8	Pre-exposed	5	ALBO	AEG	No	Yes	Yes

AEG and ALBO stand for *Ae. aegypti* and *Ae. albopictus*, respectively; C for cross, and numbers 1–8 for cross identity.

crosses and to the conspecific control crosses. Egg hatch was performed as described for Experiment 1 for batches that were collected from females mated with non-irradiated males.

Statistical analysis

Statistical analyses were performed and graphs drawn using RStudio Team [46]. The proportion of inseminated females was calculated as the number with at least one spermathecae filled with sperm divided by the number of dissected females. The proportion of egg hatch was calculated from the number of hatched eggs divided by the total number of eggs laid per individual female (Experiment 1) or per group of females in a cage (Experiments 3 and 4). The proportions were transformed following the Freeman–Tukey method (arcsine transformed data). An analysis of variance (ANOVA) was then performed followed by a Tukey multiple comparison to compare means of each crossed pair. A paired *t*-test was performed to compare egg hatch between pre-exposed and non-exposed females (pre-exposure effect on mating success experiment).

The survival rate (Experiment 1) was analyzed using a generalized binomial linear mixed-effects model fit by maximum likelihood (Laplace approximation) with logit link with the survival rate, defined as dependent variable, and type of mating pairs (cross) as fixed effects, and replicates as a random effect. The best model was selected based on the lowest corrected Akaike information criterion (AICc), and the significance of fixed effects was tested using the likelihood ratio test [14, 29].

The statistical significance for all experiments was determined at $\alpha = 0.05$ level.

Results

Experiment 1. Sympatric cross-mating between *Ae. aegypti* and *Ae. albopictus* strains from La Réunion island

There was no insemination between sympatric species from La Réunion island strains either when non-irradiated or irradiated male *Ae. aegypti* or *Ae. albopictus* were caged with female *Ae. albopictus* ($n = 82$, mean (\pm SE) = $0 \pm 0\%$ and $n = 72$, mean (\pm SE) = $0 \pm 0\%$) or *Ae. aegypti* ($n = 28$, mean (\pm SE) = $0 \pm 0\%$ and $n = 41$, mean (\pm SE) = $0 \pm 0\%$), respectively, compared to the conspecific mating controls (*Ae. albopictus*: $n = 58$, mean

(\pm SE) = $99.2 \pm 0.01\%$; *Ae. aegypti*: $n = 71$, mean (\pm SE) = $100 \pm 0\%$) (Fig. 1, $F_{(5,12)} = 735.6$, $p < 0.0001$). A Tukey multiple comparison of means did not show any difference in insemination rate regardless of the cross type between irradiated or non-irradiated males of *Ae. albopictus* or *Ae. aegypti* and non-irradiated females of *Ae. aegypti* and *Ae. albopictus*, respectively ($p > 0.05$).

Regardless of the male status (irradiated or non-irradiated) the females exposed to interspecific mating, and virgin females, laid a lower number of eggs per female than the females of the conspecific mating control crosses (Interspecific: (C3-6): 146 eggs ($n = 223$); virgin females (Faeg-Falbo): 92 eggs ($n = 77$), controls (C1-2): 2047 eggs ($n = 129$)). None of the eggs from interspecific mating crosses and virgin females hatched compared to the controls (Fig. 2, $F_{(7, 46)} = 84.73$, $p < 0.0001$).

Interspecific mating was more detrimental to female *Ae. aegypti* survival than to *Ae. albopictus* (Table 2, $p < 0.0001$).

Experiment 2. Effect of male *Ae. albopictus* density (ratio) on female *Ae. aegypti* mating success

Increasing the ratio of male: female to 3:1 in favor of male *Ae. albopictus* did not significantly increase the satyriation effect on female *Ae. aegypti* (La Réunion island) judged by insemination rates for both non-irradiated (*t*-test, ratio 1:1: mean (\pm SE) = $0 \pm 0\%$, $n = 28$, ratio 3:1: mean (\pm SE) = $2 \pm 0.1\%$, $n = 169$, $p = 0.26$) and irradiated (*t*-test, ratio 1:1: mean (\pm SE) = $0 \pm 0\%$, $n = 41$, ratio 3:1: mean (\pm SE) = $0 \pm 0\%$, $n = 101$, $p = 1$) male *Ae. albopictus*. All these crosses had negligible insemination rates compared to conspecific mating (controls) ($F_{(7, 21)} = 272.8$, $p < 0.0001$).

Experiment 3. Pre-exposure effect on mating success

When female *Ae. aegypti* were pre-exposed to non-irradiated or irradiated male *Ae. albopictus*, there was no insemination (Fig. 3, C5: mean (\pm SE) = $0 \pm 0\%$, $n = 90$, and C6: mean (\pm SE) = $0 \pm 0\%$, $n = 121$), whereas groups of females that were pre-exposed to male *Ae. albopictus* were inseminated by their male *Ae. aegypti* conspecifics (Fig. 3, C7: mean (\pm SE) = $100 \pm 0\%$, $n = 106$, and C8: mean (\pm SE) = $95.99 \pm 0\%$,

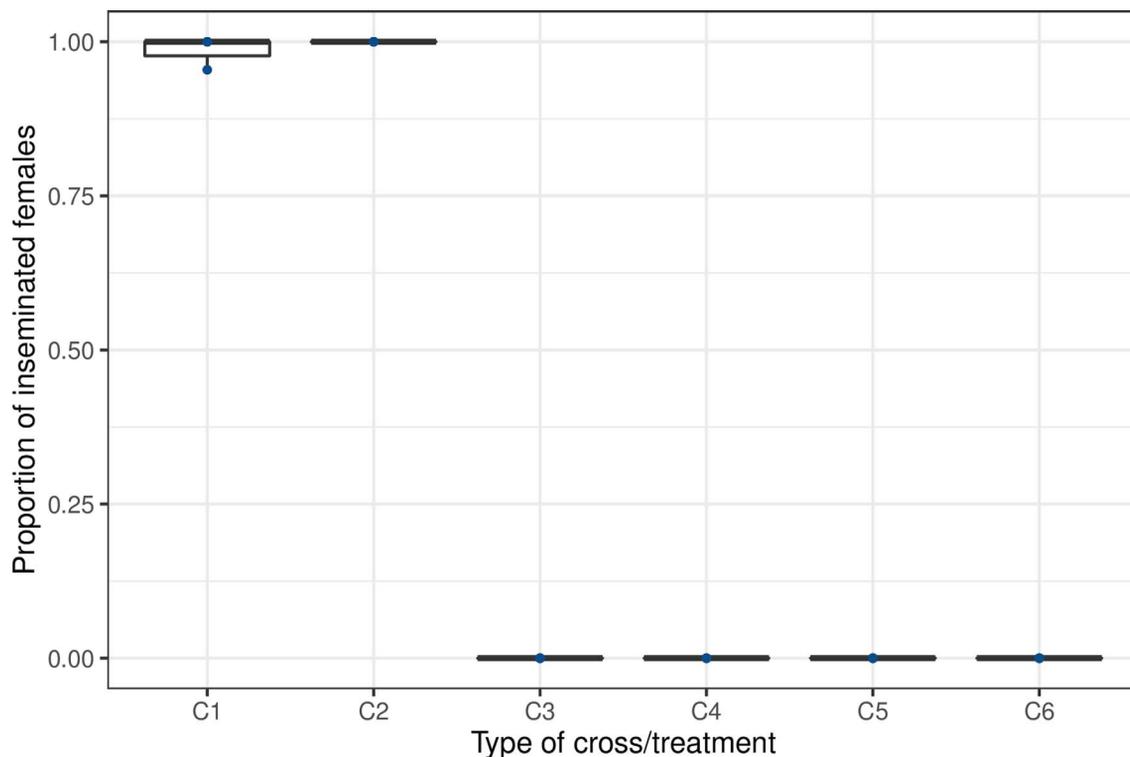


Figure 1. Sympatric cross-mating between *Ae. aegypti* and *Ae. albopictus* strains from La Réunion island. C denotes the cross, and numbers (1–6) are related to the cross identity. The number of dissected females (n) per cross were C1 = control *Ae. albopictus* (non-irradiated), $n = 58$; C2 = control *Ae. aegypti* (non-irradiated), $n = 71$; C3 = male *Ae. albopictus* (non-irradiated) \times female *Ae. aegypti*, $n = 28$; C4 = male *Ae. albopictus* (irradiated) \times female *Ae. aegypti*, $n = 41$; C5 = male *Ae. aegypti* (non-irradiated) \times female *Ae. albopictus*, $n = 82$; C6 = male *Ae. aegypti* (irradiated) \times female *Ae. albopictus*, $n = 72$.

$n = 146$). No difference in female insemination rates was observed between pre-exposed and non-exposed females ($p = 0.27$) or between pre-exposed females and controls (Fig. 3, C1-3 vs. C7-8) ($p > 0.05$).

Female *Ae. aegypti* that were pre-exposed to male *Ae. albopictus* successfully laid fertile eggs when they then mated with their male *Ae. aegypti* conspecifics, and mean egg hatch was lower but not significantly different between the non-exposed (mean (\pm SE) = $95.99 \pm 0.008\%$) and pre-exposed females (mean (\pm SE) = $91.61 \pm 0.02\%$) (t -test: $t = 2.0576$, $df = 4$, $p = 0.1$).

Experiment 4. Effect of geographic origin on mating success (allopatric crosses)

A greater female insemination rate was observed when conspecific mating was compared to the interspecific mating groups, regardless of the male origin or irradiation status (Table 3, $F_{(13, 23)} = 229.1$, $p < 0.0001$). Neither geographic origin nor irradiation status of male *Ae. albopictus* had an impact on female *Ae. aegypti* insemination rates, confirming the conclusion that allopatric and sympatric effects were similar when the La Réunion island strain of *Ae. aegypti* was used in these experiments (Table 3, $p > 0.05$).

We observed a significantly lower egg hatch in interspecific than conspecific crosses, irrespective of the male origin (Table 3, $F_{(13, 23)} = 182.9$, $p < 0.0001$).

Discussion

The objective of this study was to assess the level of reproductive interference between male *Ae. albopictus* and female *Ae. aegypti*, and to discuss the management strategy for the two vector species in a sympatric area. In laboratory conditions, almost no satyrization effect was observed between sympatric populations of *Ae. aegypti* and *Ae. albopictus* colonized from La Réunion island, even when the male *Ae. albopictus* to female *Ae. aegypti* ratio was increased to 3:1. Female *Ae. aegypti* previously exposed to male *Ae. albopictus* were not prevented from being inseminated by conspecific male *Ae. aegypti*. Satyrization did not occur between allopatric *Ae. albopictus* and *Ae. aegypti* strains from La Réunion island either. An *Ae. aegypti* strain from La Réunion island has therefore developed full resistance to satyrization (anti-satyrization behavior).

Many reasons could explain the observed low level of satyrization. Bargielowski et al. [6, 7] have previously described sympatric field populations that have co-existed long enough to evolve resistance to cross-mating and showed that *Ae. aegypti* females allopatric to *Ae. albopictus* are more susceptible to interspecific insemination by *Ae. albopictus* [40]. Cage experiments and field observations indicate that *Ae. albopictus* males are capable of satyrizing females of other species of the *Stegomyia* subgenus, potentially leading to competitive displacement, and even extinction, especially of endemic species

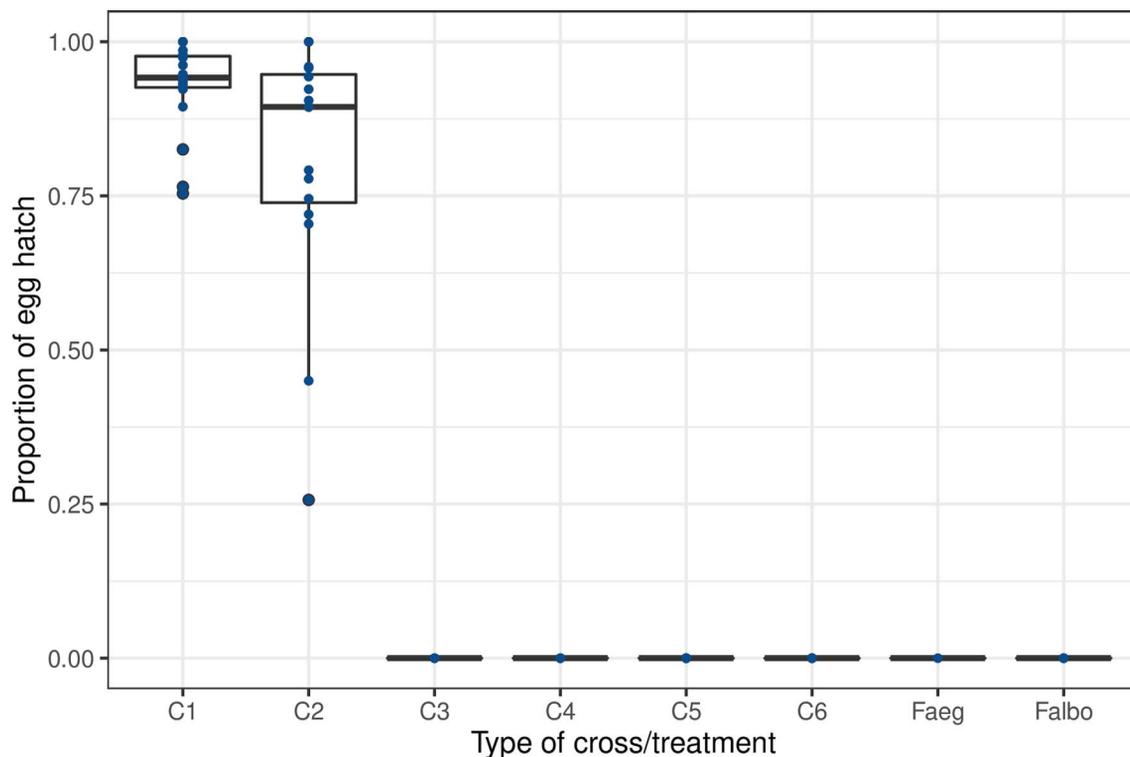


Figure 2. Egg hatch in sympatric crosses between *Ae. aegypti* and *Ae. albopictus* strains from La Réunion island. C denotes the cross and numbers (1–6) are related to the cross identity. C1 = control *Ae. albopictus* (non-irradiated), C2 = control *Ae. aegypti* (non-irradiated), C3 = male *Ae. albopictus* (non-irradiated) × female *Ae. aegypti*, C4 = male *Ae. albopictus* (irradiated) × female *Ae. aegypti*, C5 = male *Ae. aegypti* (non-irradiated) × female *Ae. albopictus*, C4 = male *Ae. aegypti* (irradiated) × female *Ae. albopictus*.

Table 2. Comparison of survival rate between female *Ae. aegypti* in the cross-mating between sympatric *Ae. aegypti* and *Ae. albopictus* strains from La Réunion island.

	Estimate	Std. error	z value	$p(> z)$
(Intercept)	5.93E−16	1.63E−01	0	1
crossctrlalbo	−2.96E−01	2.32E−01	−1.272	0.2032
crossMaegFalbo	5.04E−01	2.02E−01	2.492	0.0127*
crossMalboFaeg	−1.05E+00	2.10E−01	−4.987	6.13E−07***

Std.: standard. Ctrlalbo stands for control *Ae. albopictus* cross; MaegFalbo = male *Ae. aegypti* × female *Ae. albopictus*; MalboFaeg = male *Ae. albopictus* × female *Ae. aegypti*. * $p < 0.05$ and *** $p < 0.001$.

on islands [7]. In contrast, in our study even an increase in ratios in favor of male *Ae. albopictus* did not significantly increase insemination of female *Ae. aegypti*. One study has pointed to the implication of population density on mating behavior [42], and found that male density significantly increased swarming behavior, number of mating pairs, and egg production of hetero-specific females, but not female insemination. They also found that *Ae. aegypti* males mate more readily with hetero-specific females than do *Ae. albopictus* males, and so if *Ae. aegypti* were released into the field they may mate with both *Ae. aegypti* and *Ae. albopictus* females, and reduce populations of both species by preventing offspring. There is no evidence that this would be the case on La Réunion island since we

observed low reproductive success when crosses were performed in both directions. In addition, we observed that interspecific mating was detrimental to female *Ae. aegypti* survival. It has been previously documented that interspecific copulation and hybridization can reduce female reproductive success, but interspecific sexual harassment, which does not always result in interspecific copulation, can also adversely affect individual reproductive success and longevity by suppressing both sugar and blood feeding [47]. In white butterflies (*Leptidea* spp.), for example, the prolonged mating ritual of hetero-specific males decreases the number of opportunities that females have to mate with conspecific males even when interspecific copulation does not take place [24]. Similarly, in bean weevils (*Callosobruchus* spp.), males persistently chase hetero-specific females, causing reductions in the oviposition rate and shortened longevity of these females [34].

Ae. aegypti females pre-exposed to male *Ae. albopictus* were able to be inseminated by their conspecific male *Ae. aegypti* demonstrating that the La Réunion island *Ae. aegypti* strain has developed a resistance to satyriation. Carrasquilla and Lounibos [15] have shown that *Ae. aegypti* females, previously exposed to *Ae. albopictus* males, were rendered refractory to subsequent conspecific mating even when their spermathecae contained no hetero-specific sperm. Additional experiments demonstrated transfer of labelled semen from *Ae. albopictus* males to *Ae. aegypti* females and low production of viable eggs of females housed with conspecific males, following exposure to *Ae. albopictus* males, and

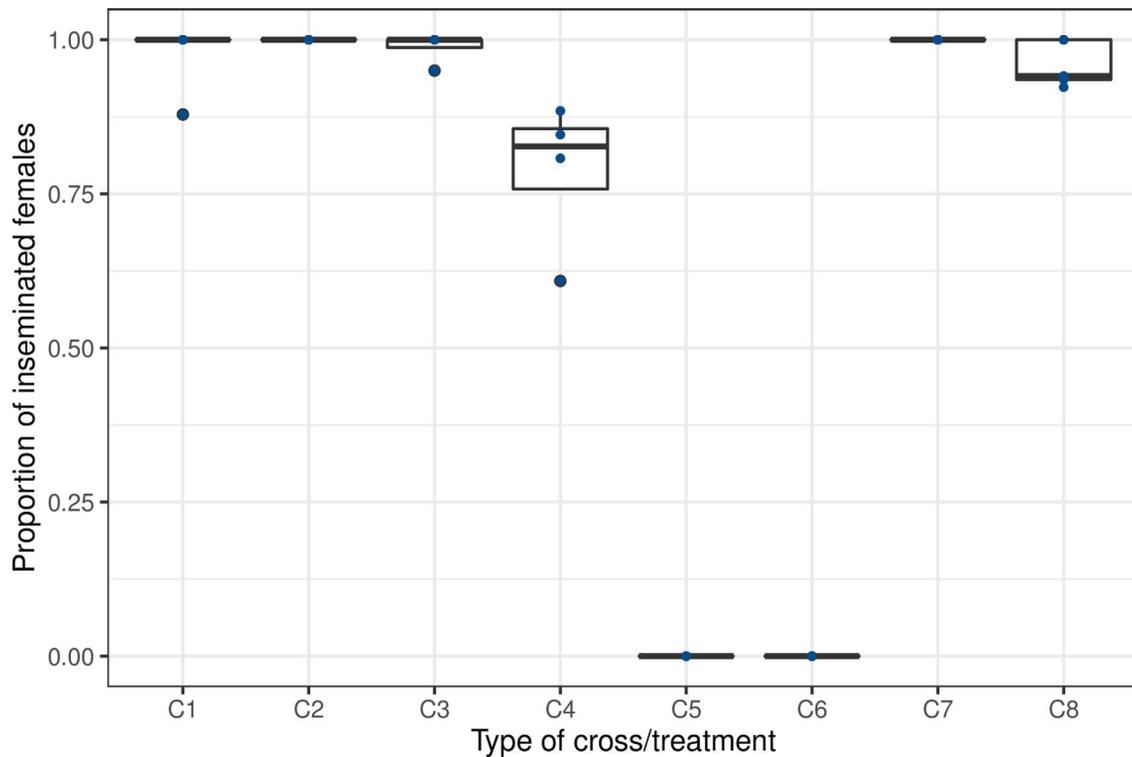


Figure 3. Pre-exposure effect on mating success. C denotes the cross, and numbers (1–8) are related to the cross identity. C1 = control *Ae. aegypti* (non-irradiated), C2 = control *Ae. aegypti* (irradiated), C3 = control *Ae. albopictus* (non-irradiated), C4 = control *Ae. albopictus* (irradiated), C5 = male *Ae. albopictus* (non-irradiated) × female *Ae. aegypti*, C6 = male *Ae. albopictus* (irradiated) × female *Ae. aegypti*, C7 = male *Ae. aegypti* (non-irradiated) × female *Ae. aegypti* (non-exposed to male *Ae. albopictus*), C8 = male *Ae. aegypti* (non-irradiated) × female *Ae. aegypti* (pre-exposed to male *Ae. albopictus*).

Table 3. Mean insemination and egg hatch rates (±SE) in crosses between female *Ae. aegypti* (La Réunion island strain) and *Ae. albopictus* males from China, Italy (allopatric), and La Réunion island (sympatric).

Mating type	Male		Female	Insemination rates (%)	Hatch rates (%)
	Irradiated	Non-irradiated			
Interspecific	China	China	Réunion	0.92 ± 0.92 ^a (95)	NA
		China	Réunion	0 ± 0 ^a (82)	0 ± 0 ^a
Conspecific	China	China	China	100 ± 0 ^b (24)	NA
		China	China	100 ± 0 ^b (48)	100 ± 0 ^b
Interspecific	Réunion	Réunion	Réunion	1.92 ± 1.11 ^a (94)	NA
		Réunion	Réunion	1.33 ± 1.33 ^a (74)	0 ± 0 ^a
Conspecific	Réunion	Réunion	Réunion	100 ± 0 ^b (57)	NA
		Réunion	Réunion	100 ± 0 ^b (54)	71.8 ± 2.7 ^c
Interspecific	Italy	Italy	Réunion	0 ± 0 ^a (99)	NA
		Italy	Réunion	5.5 ± 3.68 ^a (77)	0 ± 0 ^a
Conspecific	Italy	Italy	Italy	100 ± 0 ^b (50)	NA
		Italy	Italy	100 ± 0 ^b (57)	83.5 ± 1.55 ^c
Conspecific	Réunion*	Réunion*	Réunion	100 ± 0 ^b (72)	NA
		Réunion*	Réunion	100 ± 0 ^b (54)	89.8 ± 2.11 ^c

“Interspecific” stands for crosses between female *Ae. aegypti*, La Réunion island strain and *Ae. albopictus* (strains from China, Italy, La Réunion island) and “conspecific” for control mating between male and female of the same species. Réunion*= male *Ae. aegypti* from La Réunion island. Numbers in parentheses represent the number of dissected females for insemination assessment. Different letters show significant differences between groups.

confirmed higher incidence of satyriation than expected, based on hetero-specific insemination rates. We did not observe this result after pre-exposing *Ae. aegypti* females to *Ae. albopictus* for seven days before replacing male *Ae. albopictus* by male

Ae. aegypti mosquitoes. It has been shown that interspecific pairs needed more time together before mating occurred. Bargielowski et al. [7] found that when female *Ae. aegypti* were exposed for up to three weeks, interspecific insemination

increased significantly from 1% after one day, to 10% after one week, and to more than 50% after three weeks. However, assuming that most released sterile males will survive around one week in the field, it is unlikely that most released males would be able to find and mate with females after three weeks in the wild [9, 30]. These results indicate that in areas where *Ae. aegypti* and *Ae. albopictus* co-occur, releasing sterile male *Ae. albopictus* may not suppress *Ae. aegypti* populations. It would be more beneficial to suppress the species with the smallest population, before further planning to control the second species, assuming the epidemiological impact of each species was equal.

In our study, female *Ae. aegypti* pre-exposed to male *Ae. albopictus* produced eggs which had similar egg hatch when mated with their conspecifics, meaning that females had not been inseminated by the *Ae. albopictus* males. The females that later mated with their conspecifics and laid eggs were apparently fully fertilized by conspecific sperm. However, it has been shown that the satyrization effect could be underestimated when evaluation of mating status of females is based on whether the spermathecae were filled with sperm or not [15]. It therefore cannot be ruled out that some females might have been inseminated when pre-exposed to *Ae. albopictus* based on the variation observed in egg hatch. Bargielowski et al. [7] demonstrated that multiple inseminations can occur in older female *Ae. aegypti* when the effects of accessory gland proteins have worn off, and in females mated to sperm-depleted males. In any case, hetero-specific sperm is known to be stored in separate spermathecae [7] and so was presumably not significantly used for egg fertilization.

Allopatric *Ae. albopictus* males did not perform better than sympatric males and anti-satyrization effects seem to protect against allopatric populations. This shows that resistance to one strain confers resistance to others. Honório et al. [28] demonstrated that only some populations of *Ae. albopictus* are capable of satyrization. Female *Ae. aegypti* from populations allopatric to *Ae. albopictus* in the field were more susceptible to interspecific mating than females from sympatric populations, and selection experiments in cages confirmed the rapid development of resistance to satyrization in the laboratory, as well as changes in behavior toward conspecifics associated with increased satyrization resistance [5]. The fact that the *Ae. aegypti* populations persist in La Réunion island ravines as opposed to urban environments could be due to some genetic differentiation from domestic subspecies. Lounibos and Juliano [39] have recently pointed out that the feral subspecies *Ae. aegypti formosus* is expected to behave differently than the domestic subspecies but populations of this species from Madagascar, La Réunion island and Mayotte have not been tested yet for genetic distinctiveness from *Ae. aegypti (aegypti)* to the best of our knowledge. In any case, a signature of selection in the *Ae. aegypti* genome to a specific type of interspecific interaction (mating) was found by Burford Reiskind et al. [13] allowing the identification of its genetic basis.

When considering a regional approach for *Aedes* control using the SIT, compatibility of strains as well as species may be important as it would allow strains to be imported for release from nearby countries where they can be more easily reared and/or irradiated. Damiens et al. [18] demonstrated that male

Ae. albopictus from Mauritius and Seychelles islands, about 50-200 km away from La Réunion island, were compatible and could successfully inseminate female *Ae. albopictus* regardless of their origin. A regional SIT mass-rearing program could therefore be considered, with a good transportation method, but the release of sterile *Ae. albopictus* males may not have the added benefit of satyrizing the local *Ae. aegypti* population if an overflowing of irradiated male *Ae. albopictus* leads to similar results.

The development of resistance to satyrization in the *Ae. aegypti* strain shows that strong competition between the sympatric *Ae. aegypti* and *Ae. albopictus* probably occurs on La Réunion island. An SIT project against *Ae. albopictus* would not have an effect on *Ae. aegypti* populations, and other mechanisms such as larval competition probably explain the current geographical retraction of *Ae. aegypti* to the ravines. Bagny et al. [4] observed that this narrow distribution of *Ae. aegypti* was due to its poorer ability to cope with unfavorable temperatures and to its lower competition between larvae for resources compared to *Ae. albopictus* [31]. The two species may co-exist as long as the dominant *Ae. albopictus* is present and the resistance could be maintained by satyrization pressure [8]. Global climate change may favor an increase in the population size of *Ae. aegypti* [32], which is a greater vector of arboviral diseases including dengue, chikungunya, yellow fever, and Zika. Therefore, suppressing or eliminating *Ae. albopictus* will likely promote expansion of *Ae. aegypti* [26]. Whilst it may be important to target the most epidemiologically important vector first [1], considering its limited distribution, the eradication of the *Ae. aegypti* population may be seen as a first priority.

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Author contributions

HM carried out the lab experiments, data analysis, and contributed to the design of the study and drafting of the manuscript. JG, RL, and HY contributed to the design of the study and drafting of the manuscript. JB helped in data analysis and drafting of the manuscript. JG and JB supervised this work. HY, RL, JG, and JB critically revised the manuscript. All authors gave final approval for publication.

Competing interests

The authors declare that they have no competing interests.

Data availability

All data generated or analyzed during this study are included in this published article.

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References

- Alphey L, Benedict M, Bellini R, Clark GG, Dame DA, Service MW, Dobson SL. 2010. Sterile-insect methods for control of mosquito-borne diseases: an analysis. *Vector Borne and Zoonotic Diseases*, 10, 295–311.
- Arrêté No 2019-2213. Autorisant l'IRD à procéder à des lâchers de moustiques stériles en vue d'études entomologiques au sein du quartier Duparc à Sainte-Marie – 13/06/2019.
- Bagny L, Delatte H, Quilici S, Fontenille D. 2009. Progressive decrease in *Aedes aegypti* distribution in Reunion Island since the 1900s. *Journal of Medical Entomology*, 46, 1541–1545.
- Bagny Beilhe L, Delatte H, Juliano SA, Fontenille D, Quilici S. 2013. Ecological interactions in *Aedes* species on Reunion Island. *Medical and Veterinary Entomology*, 27, 387–397.
- Bargielowski I, Lounibos LP. 2014. Rapid evolution of reduced receptivity to interspecific mating in the dengue vector *Aedes aegypti* in response to satyriation by invasive *Aedes albopictus*. *Evolutionary Ecology*, 28, 193–203.
- Bargielowski IE, Lounibos LP, Carrasquilla MC. 2013. Evolution of resistance to satyriation through reproductive character displacement in populations of invasive dengue vectors. *Proceedings of the National Academy of Sciences of the United States of America*, 110, 2888–2892.
- Bargielowski IE, Blosser E, Lounibos LP. 2015. The effects of interspecific courtship on the mating success of *Aedes aegypti* and *Aedes albopictus* (Diptera: Culicidae) males. *Annals of the Entomological Society of America*, 108, 513–518.
- Bargielowski I, Honório NA, Blosser EM, Lounibos LP. 2019. Rapid loss of resistance to satyriation in invasive mosquitoes and the effects of age on interspecific mating frequency. *Journal of Medical Entomology*, 56, 329–333.
- Bellini R, Albieri A, Balestrino F, Carrieri M, Porretta D, Urbanelli S, Calvitti M, Riccardo MR, Maini S. 2010. Dispersal and survival of *Aedes albopictus* (Diptera: Culicidae) males in Italian urban areas and significance for sterile insect technique application. *Journal of Medical Entomology*, 7, 1082–1091.
- Benedict MQ, Levine RS, Hawley WA, Lounibos LP. 2007. Spread of the tiger: global risk of invasion by the mosquito *Aedes albopictus*. *Vector Borne and Zoonotic Diseases*, 7, 76–85.
- Bhatt S, Gething PW, Brady OJ, Messina JP, Farlow AW, Moyes CL, Drake JM, Brownstein JS, Hoen AG, Sankoh O, Myers MF, George DB, Jaenisch T, Wint GRW, Simmons CP, Scott TW, Farrar JJ, Hay SI. 2013. The global distribution and burden of dengue. *Nature*, 496, 504–507.
- Bouyer J, Yamada H, Pereira R, Bourtzis K, Vreysen MJB. 2020. Phased conditional approach for mosquito management using sterile insect technique. *Trends in Parasitology*, S1471–4922(20), 30014-3.
- Burford Reiskind MO, Labadie P, Bargielowski I, Lounibos LP, Reiskind MH. 2018. Rapid evolution and the genomic consequences of selection against interspecific mating. *Molecular Ecology*, 27, 3641–3654.
- Burnham KP, Anderson DR. 2003. Model selection and multimodel inference: a practical information-theoretic approach. Berlin: Springer Science & Business Media.
- Carrasquilla MC, Lounibos LP. 2015. Satyriation without evidence of successful insemination from interspecific mating between invasive mosquitoes. *Biology Letters*, 11, 20150527.
- Chouin-Carneiro T, Vega-Rua A, Vazeille M, Yebakima A, Girod R, Goindin D, Dupont-Rouzeyrol M, Lourenço-de-Oliveira R, Failloux AB. 2016. Differential Susceptibilities of *Aedes aegypti* and *Aedes albopictus* from the Americas to Zika Virus. *PLoS Neglected Tropical Diseases*, 10, e0004543.
- Conseil scientifique de l'Agence française pour la biodiversité – Séance des 26–27 avril 2018 – Délibération n° CS/2018-01.
- Damiens D, Lebon C, Wilkinson DA, Dijoux-Millet D, Le Goff G, Bheecarry A, Gouagna LC. 2016. Cross-mating compatibility and competitiveness among *Aedes albopictus* strains from distinct geographic origins – implications for future application of SIT Programs in the South West Indian Ocean Islands. *PLoS One*, 11, e0163788.
- de Araújo TVB, Rodrigues LC, de Alencar Ximenes RA, de Barros Miranda-Filho D, Montarroyos UR, de Melo APL, Valongueiro S, de Albuquerque MFPM, Souza WV, Braga C, Filho SPB, Cordeiro MT, Vazquez E, Di Cavalcanti Souza Cruz D, Henriques CMP, Bezerra LCA, da Silva Castanha PM, Dhalia R, Marques-Júnior ETA, Martelli CMT, Investigators from the Microcephaly Epidemic Research Group, Brazilian Ministry of Health, Pan American Health Organization, Instituto de Medicina Integral Professor Fernando Figueira, State Health Department of Pernambuco. 2016. Association between Zika virus infection and microcephaly in Brazil, January to May, 2016: preliminary report of a case-control study. *Lancet Infectious Diseases*, 16, 1356–1363.
- Delatte H, Dehecq JS, Thiria J, Domerg C, Paupy C, Fontenille D. 2008. Geographic distribution and developmental sites of *Aedes albopictus* (Diptera: Culicidae) during a Chikungunya epidemic event. *Vector Borne and Zoonotic Diseases*, 8, 25–34.
- Dyck VA, Hendrichs JP, Robinson AS. 2005. The sterile insect technique: principles and practice in area-wide integrated pest management. Dordrecht: Springer.
- FAO/IAEA. 2017. Guidelines for routine colony maintenance of *Aedes* mosquito species – version 1.0 [Internet]. p. 18. Available from: <https://www.iaea.org/resources/manual/guidelines-for-routine-colony-maintenance-of-aedes-mosquito-species-version-10>.
- Focks DA. 1980. An improved separator for the developmental stages, sexes, and species of mosquitoes (Diptera: Culicidae). *Journal of Medical Entomology*, 17, 567–568.
- Friberg M, Leimar O, Wiklund C. 2013. Heterospecific courtship, minority effects and niche separation between cryptic butterfly species. *Journal of Evolutionary Biology*, 26, 971–979.
- Giatropoulos A, Papachristos DP, Koliopoulos G, Michaelakis A, Emmanouel N. 2015. Asymmetric mating interference between two related mosquito species: *Aedes (Stegomyia) albopictus* and *Aedes (Stegomyia) cretinus*. *PLoS One*, 10, e0127762.
- HCB Scientific Committee. 2017. Scientific Opinion of the High Council for Biotechnology concerning use of genetically modified mosquitoes for vector control in response to the referral of 12 October 2015 (Ref. HCB-2017.06.07). Paris: HCB. p. 142. Available online: <http://www.hautconseildes-biotechnologies.fr>.
- Honma A, Kumano N, Noriyuki S. 2019. Killing two bugs with one stone: a perspective for targeting multiple pest species by incorporating reproductive interference into sterile insect technique. *Pest Management Science*, 75, 571–577.

28. Honório NA, Carrasquilla MC, Bargielowski I, Nishimura N, Swan T, Lounibos LP. 2018. Male origin determines satyri- zation potential of *Aedes aegypti* by invasive *Aedes albopictus*. *Biological Invasions*, 20, 653–664.
29. Hurvich CM, Tsai CL. 1995. Model selection for extended quasi-likelihood models in small samples. *Biometrics*, 51, 1077–1084.
30. Iyaloo DP, Damiens D, Sunita F, Elahee KB, Bheecarry A. 2019. Dispersal and survival of radio-sterilised male *Aedes albopictus* Skuse (Diptera: Culicidae) and estimation of the wild populations in view of an sterile insect technique programme in Pointe des Lascars, Mauritius. *International Journal of Tropical Insect Science*, 39, 63.
31. Juliano SA. 2009. Species interactions among larval mosqui- toes: context dependence across habitat gradients. *Annual Review of Entomology*, 54, 37–56.
32. Juliano SA, Lounibos LP, O’Meara GF. 2004. A field test for competitive effects of *Aedes albopictus* on *Aedes aegypti* in South Florida: differences between sites of coexistence and exclusion? *Oecologia*, 139, 583–593.
33. Kaplan L, Kendell D, Robertson D, Livdahl T, Khatchikian C. 2010. *Aedes aegypti* and *Aedes albopictus* in Bermuda: extinction, invasion, invasion and extinction. *Biological Inva- sions*, 12, 3277–3288.
34. Kishi S, Nishida T, Tsubaki Y. 2009. Reproductive interference determines persistence and exclusion in species interactions. *Journal of Animal Ecology*, 78, 1043–1049.
35. Knippling EF. 1959. Sterile-male method of population control: successful with some insects, the method may also be effective when applied to other noxious animals. *Science*, 130, 902–904.
36. Kyle JL, Harris E. 2008. Global spread and persistence of dengue. *Annual Review of Microbiology*, 62, 71–92.
37. Levy-Blitchein S, Del Valle-Mendoza J. 2016. Zika virus is arriving at the American continent. *Asian Pacific Journal of Tropical Medicine*, 9, 1019–1021.
38. Lounibos LP. 2002. Invasions by insect vectors of human disease. *Annual Review of Entomology*, 47, 233–266.
39. Lounibos LP, Juliano SA. 2018. Where Vectors Collide: the importance of mechanisms shaping the realized niche for modeling ranges of invasive *Aedes* Mosquitoes. *Biological Invasions*, 20, 1913–1929.
40. Lounibos LP, Bargielowski I, Carrasquilla MC, Nishimura N. 2016. Coexistence of *Aedes aegypti* and *Aedes albopictus* (Diptera: Culicidae) in Peninsular Florida two decades after competitive displacements. *Journal of Medical Entomology*, 53, 1385–1390.
41. Maïga H, Gilles JRL, Lees RS, Yamada H, Bouyer J. 2020. Demonstration of resistance to satyri- zation behavior in *Aedes aegypti* (Linnaeus) from La Réunion island. *bioRxiv* 2020.02.10.942839. <https://doi.org/10.1101/2020.02.10.942839>.
42. Marcela P, Abu Hassan A, Hamdan A, Dieng H, Kumara TK. 2015. Interspecific cross-mating between *Aedes aegypti* and *Aedes albopictus* laboratory strains: implication of population density on mating behaviors. *Journal of the American Mosquito Control Association*, 31, 313–320.
43. Nasci RS, Hare SG, Willis FS. 1989. Interspecific mating between Louisiana strains of *Aedes albopictus* and *Aedes aegypti* in the field and laboratory. *Journal of the American Mosquito Control Association*, 5, 416–421.
44. Paupy C, Ollomo B, Kamgang B, Moutailler S, Rousset D, Demanou M, Hervé JP, Leroy E, Simard F. 2010. Comparative role of *Aedes albopictus* and *Aedes aegypti* in the emergence of dengue and Chikungunya in central Africa. *Vector Borne and Zoonotic Diseases*, 10, 259–266.
45. Ribeiro JMC. 1988. Can satyrs control pests and vectors? *Journal of Medical Entomology*, 25, 431–440.
46. RStudio Team. 2016. RStudio: Integrated Development for R. Boston, MA: RStudio Inc. URL: <http://www.rstudio.com/>.
47. Soghigian J, Gibbs K, Stanton A, Kaiser R, Livdahl T. 2014. Sexual harassment and feeding inhibition between two invasive dengue vectors. *Environmental Health Insights*, 8, 61–66.
48. Triplet F, Lounibos LP, Robbins D, Moran J, Nishimura N, Blosser EM. 2011. Competitive reduction by satyri- zation? Evidence for interspecific mating in nature and asymmetric reproductive competition between invasive mosquito vectors. *Journal of the American Mosquito Control Association*, 85, 265–270.
49. World Health Organization. 2017. Global vector control response 2017–2030.
50. World Health Organization. 2019. Dengue fever – Réunion, France. <https://www.who.int/csr/don/01-may-2018-dengue-reunion/en/> (Assessed on 11 Sep 2019).
51. Zheng ML, Zhang DJ, Damiens DD, Lees RS, Gilles JR. 2015. Standard operating procedures for standardized mass rearing of the dengue and chikungunya vectors *Aedes aegypti* and *Aedes albopictus* (Diptera: Culicidae). II. Egg storage and hatching. *Parasites & Vectors*, 8, 348.

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