

RESEARCH

Open Access



# Use of novel lab assays to examine the effect of pyrethroid-treated bed nets on blood-feeding success and longevity of highly insecticide-resistant *Anopheles gambiae* s.l. mosquitoes

Priscille Barreaux<sup>1,2,3\*</sup> , Jacob C. Koella<sup>3</sup>, Raphael N'Guessan<sup>4,5</sup> and Matthew B. Thomas<sup>2,6</sup>

## Abstract

**Background:** There is a pressing need to improve understanding of how insecticide resistance affects the functional performance of insecticide-treated nets (ITNs). Standard WHO insecticide resistance monitoring assays are designed for resistance surveillance and do not necessarily provide insight into how different frequencies, mechanisms or intensities of resistance affect the ability of ITNs to reduce malaria transmission.

**Methods:** The current study presents some novel laboratory-based assays that attempt to better simulate realistic exposure of mosquitoes to ITNs and to quantify impact of exposure not only on instantaneous mortality, but also on blood-feeding and longevity, two traits that are central to transmission. The assays evaluated the performance of a standard ITN (Permanet<sup>®</sup> 2.0; Vestergaard Frandsen), a 'next generation' combination ITN with a resistance-breaking synergist (Permanet<sup>®</sup> 3.0) and an untreated net (UTN), against field-derived *Anopheles gambiae* sensu lato mosquitoes from Côte d'Ivoire exhibiting a 1500-fold increase in pyrethroid resistance relative to a standard susceptible strain.

**Results:** The study revealed that the standard ITN induced negligible instantaneous mortality against the resistant mosquitoes, whereas the resistance-breaking net caused high mortality and a reduction in blood-feeding. However, both ITNs still impacted long-term survival relative to the UTN. The impact on longevity depended on feeding status, with blood-fed mosquitoes living longer than unfed mosquitoes following ITN exposure. Exposure to both ITNs also reduced the blood-feeding success, the time spent on the net and blood-feeding duration, relative to the untreated net.

**Conclusion:** Although a standard ITN did not have as substantial instantaneous impact as the resistance-breaking net, it still had significant impacts on traits important for transmission. These results highlight the benefit of improved bioefficacy assays that allow for realistic exposure and consider sub- or pre-lethal effects to help assess the functional significance of insecticide resistance.

**Keywords:** *Anopheles gambiae*, Pyrethroid resistance, Sublethal effect, Vector control, Blood feeding inhibition, Delayed mortality

\*Correspondence: priscille.barreaux@lstm.ac.uk

<sup>1</sup> Liverpool School of Tropical Medicine, Liverpool, UK

Full list of author information is available at the end of the article



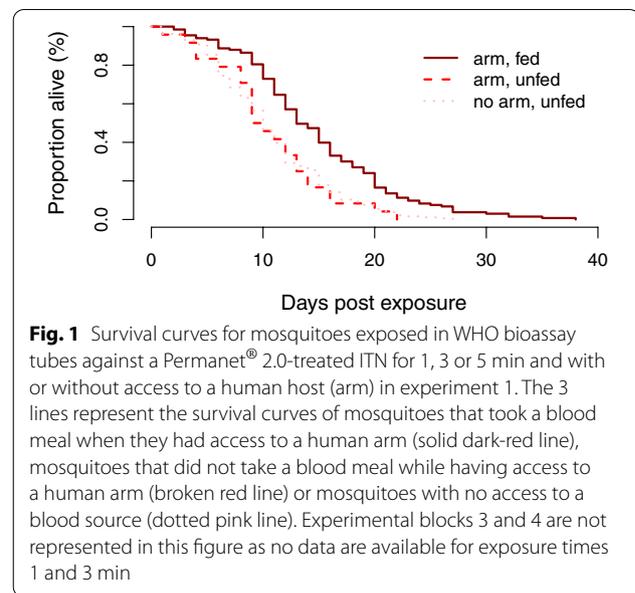
© The Author(s) 2022. **Open Access** This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>. The Creative Commons Public Domain Dedication waiver (<http://creativecommons.org/publicdomain/zero/1.0/>) applies to the data made available in this article, unless otherwise stated in a credit line to the data.

## Background

In recent decades, large-scale implementation of pyrethroid-based control tools that target the adult mosquito vectors have helped reduce the burden of malaria [1]. Insecticide-treated nets (ITNs) are the most widely distributed and perhaps most important tool to date [2]. However, their extensive use has led to the evolution of insecticide resistance in many mosquito populations [3–7], and there are now mounting concerns that resistance will render ITNs ineffective and lead to a resurgence of malaria [8–10]. As yet, however, the link between emergence of different resistance mechanisms and intensities of insecticide resistance and control failure remains unclear [11–15], which challenges the development of appropriate resistance mitigation strategies [16].

Mosquito populations are classified as resistant using standardized WHO testing procedures that measure the level of mortality within 24 h of exposure to a diagnostic dose of insecticide [17]. However, this focus on instantaneous mortality ignores possible pre- or sub-lethal effects of ITNs on longevity and blood-feeding success, two important parameters influencing malaria transmission potential [18]. Evidence suggests that ITNs can potentially reduce the transmission of malaria even in the absence of rapid knockdown and death, providing they reduce mosquito longevity and limit the number of mosquitoes that live long enough to enable the malaria parasite to complete its extrinsic incubation period [18–20]. This “sub-lethal” effect of insecticide might be minimal in some conditions [21] or may be enhanced by repeated exposures [18, 22]. In addition, previous studies with susceptible mosquito strains suggest that certain pyrethroid insecticides act on mosquito host-seeking and blood-feeding behavior, by irritating them upon contact or repelling them prior to net contact [23, 24]. While, over short distances, the repellent effect of pyrethroid seems to impact resistant mosquito strains [25–28], it is still unclear whether resistance alters the impact of ITNs on blood-feeding inhibition. In fact, some recent laboratory studies suggest that pyrethroids might even enhance host searching in resistant mosquitoes [29, 30].

In the study reported here we used two novel assay methods to explore the effects of ITN exposure on initial mortality, blood-feeding inhibition and longevity against field-derived populations of *Anopheles gambiae sensu lato* (s.l.) from central Cote d’Ivoire that are known to exhibit intense resistance to pyrethroids [4]. The primary aim was to examine the sub-/pre-lethal effects of ITNs considering different patterns of exposure. A secondary aim was to explore some novel assays that could possibly be used to supplement the standard WHO assays used to determine insecticide resistance and characterize the bioefficacy of ITNs, ultimately to provide a better



**Fig. 1** Survival curves for mosquitoes exposed in WHO bioassay tubes against a Permanet® 2.0-treated ITN for 1, 3 or 5 min and with or without access to a human host (arm) in experiment 1. The 3 lines represent the survival curves of mosquitoes that took a blood meal when they had access to a human arm (solid dark-red line), mosquitoes that did not take a blood meal while having access to a human arm (broken red line) or mosquitoes with no access to a blood source (dotted pink line). Experimental blocks 3 and 4 are not represented in this figure as no data are available for exposure times 1 and 3 min

assessment of the functional significance of insecticide resistance (Fig. 1).

## Methods

### Mosquito populations

The study used *An. gambiae* s.l. mosquitoes collected in natural breeding habitats around the villages of Yao Koffikro and M’be in central Côte d’Ivoire [31]. In this region, the resistance of mosquito populations to deltamethrin is > 1500-fold higher than that of a standard susceptible strain (Kisumu) [4, 32]. Among other resistance mechanisms, they carry the L1014F knockdown resistance (kdr) ( $\geq 90\%$  fixed) and N1575Y mutations and upregulate the *CYP6M2*, *CYP6P3* and *CYP9K1* genes [4]. The field-collected larvae were reared at  $27 \pm 2$  °C,  $60 \pm 20\%$  relative humidity and ambient light, in plastic boxes of 300 larvae with 1 l of deionized water and fed daily with fish food (TetraMin™ Baby; Tetra Werke, Melle, Germany) following a standardized “high food” regime described in [33]. Adult mosquitoes from both villages were combined and kept in  $32.5 \times 32.5 \times 32.5$ -cm mosquito cages and maintained on a 10% sugar solution.

### Human host preparation

The experimenter (PB) avoided tobacco, alcohol and the use of scented products for 12 h before and during testing. Her arms were washed with unscented soap and rinsed with water the morning before a test. PB’s temperature was monitored to reduce risk of exposing mosquitoes to any active pathogen infection (at no point during the study was PB infected with malaria).

### Bed nets

The experimental treatments consisted of three types of polyester bed nets: (i) unwashed PermaNet® 2.0-treated net (ITN; Vestergaard Frandsen, Lausanne, Switzerland); (ii) the roof of PermaNet® 3.0-treated net (ITN + PBO); (iii) and an untreated net (UTN) (Coghlan’s, Winnipeg, MB, Canada). The ITN is coated with deltamethrin at a target dose of 55 mg/m<sup>2</sup> (±25%). The ITN + PBO is coated with deltamethrin and piperonyl butoxide (PBO) at a target dose of 120 mg/m<sup>2</sup> (±25%) and 750 mg/m<sup>2</sup> (±25%), respectively [34]. It is worth noting that the difference between the standard ITN and the ITN + PBO product is not just the addition of PBO but also an increased concentration of the active ingredient, deltamethrin for the combination-treated net. Prior to testing, fully susceptible mosquitoes (Kisumu strain) were exposed to samples of netting in WHO tubes (see "General methods" section); all were killed within 24 h when exposed to the treated netting, while none of these exposed to the UTN were killed.

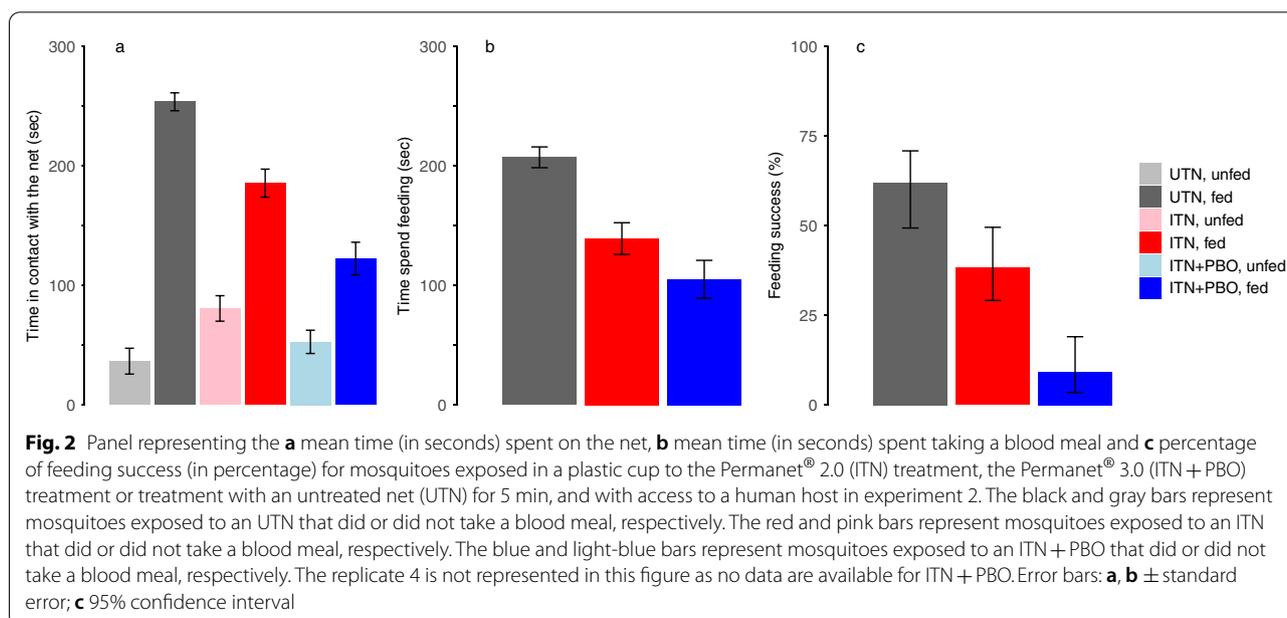
### General methods

The assays used 4- to 5-day-old adult female mosquitoes selected at random from the stock cages. Mosquitoes were assigned haphazardly to a net treatment to provide balanced sample sizes (Additional file 1: Table A). They were starved 4 h prior to testing, with assays conducted in the afternoon during daylight. Following exposure, feeding status was recorded; mosquitoes with a visible amount of bright red blood in their abdomen were considered as "fed". Following exposure, mosquitoes were

kept individually in transparent plastic cups covered with untreated netting, and mortality was recorded daily until all mosquitoes had died. Females had continuous access to a 10% sugar solution and to an egg-laying substrate, although egg numbers were not recorded in order to minimize daily mosquito handling.

### Forced exposure in modified WHO tubes

In this assay we examined the effect of a forced exposure to an ITN (PermaNet® 2.0) on mosquito mortality and capacity to blood-feed (Additional file 2: Dataset S1). The aim was partly to determine whether the highly resistant wildtype mosquitoes suffered obvious direct effects from forced contact with the ITN, but also to serve as something of a range finder to determine the effects of duration exposure and the relative influence of blood-feeding on survival following exposure, in order to inform the design of the following experiment. In two experimental blocks ( $n_1=219$  and  $n_2=122$ ), mosquitoes were individually exposed to the PermaNet® 2.0 for 1, 3 or 5 min using WHO tubes lined (inner wall and ends) with netting to force the mosquitoes to come into contact with the treated surface. During the exposure, half of the mosquitoes in each block ( $n_1=96$  and  $n_2=61$ ) had the opportunity to take a blood meal by feeding on PB’s arm through the netting; the other half were not allowed to feed, but PB held her arm 1 cm away from the tube to provide equivalent host cues. In two additional experimental blocks ( $n_3=84$  and  $n_4=56$ ), mosquitoes were exposed for 5 min only (Fig. 2).



### Variable exposure via individual feeding choice

This assay relaxed the forced contact experienced in the WHO tubes by placing mosquitoes individually in transparent 180-ml plastic cups with the top covered with netting, either ITN, ITN+PBO (roof of the Permanet<sup>®</sup>-3.0 treated netting) or UTN. PB's arm was placed onto the net at the top of the cup to attract mosquitoes and enable blood-feeding (Additional file 3: Dataset S2). The aim was to simulate more natural patterns and durations of contact during a 5-min exposure period. In principle, if mosquitoes were repelled by the netting or suffered from irritancy following initial contact, they might have only minimal contact. On the other hand, if they were motivated to feed and were unaffected by the presence of the netting, contact could last for up to 5 min. The time mosquitoes were in contact with the net and the duration of their blood meal were recorded, together with subsequent longevity. The ITN, ITN+PBO and a UTN were compared in three experimental blocks ( $n_1=61$ ,  $n_2=58$  and  $n_3=127$ ) while in a fourth experimental block ( $n_4=123$ ), females were not tested against the ITN+PBO because of limited mosquito numbers from the larval collection and the desire to increase the sample sizes (Additional file 4: Table C).

### Statistical analysis

All analyses and graphs were done in R version 3.6.1 (available in opensource: <https://www.R-project.org/>). Contrasts among treatments were assessed with the multcomp package version 1.4–10 and the function 'glht' with Tukey's honestly significant difference test. All complete statistical analyses can be found in Additional file 5: Table B.

### Forced exposure in modified WHO tube assays

Using a generalized linear model (GLM) with a binomial distribution, the blood-feeding success of mosquitoes given access to a blood source was analyzed to investigate whether the proportion that fed depended on the duration of insecticide exposure and the experimental block.

Survival post exposure was analyzed with a weighted Cox regression (using the R package coxphw due to the violation of the proportional hazards assumption in a Cox regression model) according to the blood-feeding categories (no access to a blood source; access to a blood source but unfed; access to a blood source and fed), the duration of insecticide exposure, the experimental blocks and their interaction. Given the complex interaction found in this analysis and the 5-min time needed for mosquitoes to engorge blood to repletion [35], we analyzed the survival post exposure for blood-fed and unfed mosquitoes separately with a weighted Cox regression including two exposure time categories (1- and 3-min exposure times

compared to 5-min exposure time), the experimental blocks and their interaction. In a preliminary model for unfed mosquitoes, whether mosquitoes had access to an arm or not during exposure did not influence longevity; thus, this factor it was not included in the final analysis.

In addition, mosquito survival post exposure in two additional replicates was analyzed together with the other experimental blocks considering mosquitoes exposed in WHO tubes for 5 min only. A weighted Cox model was used to investigate the effect of the blood-feeding categories, the experimental blocks and their interaction.

### Variable exposure via individual feeding choice

We analyzed the time spent on the net with a Gaussian GLM and an identity link function including the type of bed net (UTN, ITN, ITN+PBO) and the experimental blocks as nominal factors.

We analyzed the time spent feeding with a Gaussian GLM (for fed mosquitoes only) and the proportion of mosquitoes that fed with a binomial GLM, with both analyses including the type of bed nets, the time spent on the net, their interaction and the experimental blocks as nominal factors (the latter in interaction with the other parameters for the binomial GLM).

We analyzed the survival post exposure of the mosquitoes with a weighted Cox proportional hazards model, with the type of bed net, the feeding status, their interactions and the experimental blocks as factors. Considering fed mosquitoes alone, the same analysis was done without the feeding status and adding the time spent on the net and the time spent feeding. We then repeated that analysis for unfed mosquitoes alone with the time spent on the net (summary in Additional file 2: Table B.a.3).

The analysis for the time spent on the net, feeding success and survival post exposure were repeated with an additional experimental block for UTN and ITN treatments only.

## Results

### Forced exposure in modified WHO tube assays

Forced exposure to the ITN resulted in negligible mortality of highly resistant mosquitoes within 24 h regardless of exposure period (note this contrasts to 100% mortality of the susceptible Kisumu strain in pilot studies referred to in "Methods" section) (Fig. 1). In addition, 84.7% (95% confidence interval [CI]: 78.1–89.9%) of mosquitoes that were provided access to a blood meal were able to feed, irrespective of the duration of the exposure ( $\chi^2=1.32$ ,  $df=1$ ,  $P=0.25$ ). Blood-feeding increased mosquito longevity by approximately 4 days, with blood-fed mosquitoes having a mean ( $\pm$  standard error [SE]) survival time (post exposure at 4 days old) of  $14.7 \pm 0.59$  days post exposure, compared with  $10.5 \pm 1.04$  days post

exposure for unfed mosquitoes having access to the arm and  $10.7 \pm 0.39$  days post exposure for unfed mosquitoes having no access to the arm ( $\chi^2 = 29.80$ ,  $df = 1$ ,  $P < 0.001$ ) (Fig. 1). At exposure times of 1 and 3 min, fed mosquitoes lived an average of 2.8 more days post exposure than unfed mosquitoes (unfed:  $11.1 \pm 0.47$  days; fed:  $13.9 \pm 0.59$  days), and when exposed for 5 min, fed mosquitoes lived an average of 6.5 more days post exposure (unfed:  $9.8 \pm 0.58$  days; fed:  $16.3 \pm 1.36$  days) ( $\chi^2 = 14.64$ ,  $df = 1$ ,  $P < 0.001$ ). While the interaction between exposure time and blood-feeding was significant on longevity, there was no significant effect of exposure duration itself ( $\chi^2 = 0.56$ ,  $df = 1$ ,  $P = 0.45$ ). Subgroup analysis showed no influence of exposure duration on mean longevity for unfed mosquitoes ( $\chi^2 = 3.20$ ,  $df = 1$ ,  $P = 0.07$ ). However, longer exposure to insecticide led to a longer life post exposure for blood-fed mosquitoes ( $13.9 \pm 0.59$  days for 1- and 3-min exposure and  $16.3 \pm 1.36$  days for 5-min exposure) ( $\chi^2 = 5.78$ ,  $df = 1$ ,  $P = 0.02$ ).

Two additional experimental blocks, in which the exposure time of mosquitoes was 5 min only, corroborated these results. Combining all experimental blocks for the 5-min exposure time showed that a blood meal extended the lifespan of mosquitoes by around 7 days post exposure ( $9.0 \pm 0.73$  days for females with no access to blood source;  $9.7 \pm 0.43$  days for those with access to the blood source but unfed; and  $16.6 \pm 0.87$  days for those with access to the blood source and fed) ( $\chi^2 = 51.03$ ,  $df = 1$ ,  $P < 0.001$ ).

### Variable exposure via individual feeding choice

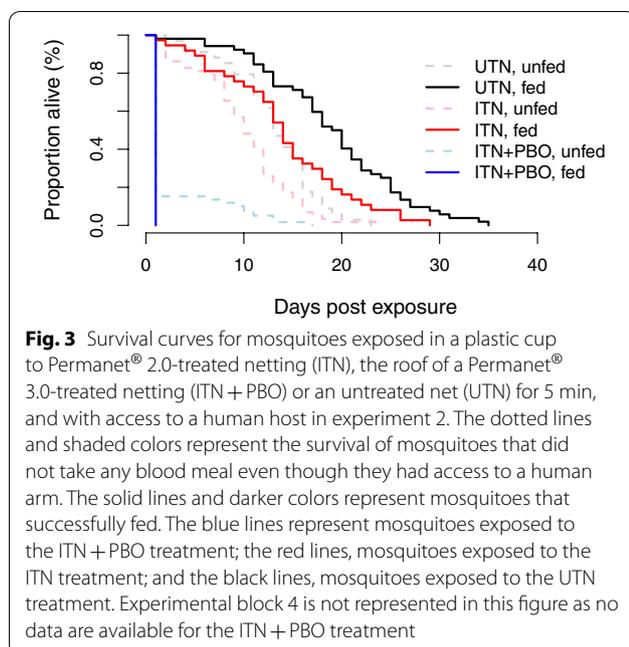
The presence of insecticide reduced the average contact time with the netting ( $F = 21.97$ ,  $df = 2$ ,  $P < 0.001$ ; Fig. 2a). Mosquitoes exposed to the UTN had an average contact time of  $167.7 \pm 13.06$  s, while those exposed to the ITN or ITN + PBO had average contact times of  $121.4 \pm 9.5$  and  $59.1 \pm 9.25$  s, respectively (Tukey pairwise comparisons:  $P_{ITN-UTN} = 0.005$ ,  $P_{ITN+PBO-UTN}$ ). The average contact time for the synergist-treated net (ITN + PBO) was lower than that for the ITN without PBO ( $P_{ITN+PBO-ITN} < 0.001$ ).

The feeding duration showed a similar pattern between net types ( $F = 45.30$ ,  $df = 2$ ,  $P < 0.001$ ; Fig. 2b), with average times of  $219.3 \pm 8.82$ ,  $135.4 \pm 9.80$  and  $105.0 \pm 15.85$  s for UTN, ITN, ITN + PBO, respectively (Tukey pairwise comparisons:  $P_{ITN-UTN}$  and  $P_{ITN+PBO-UTN} < 0.001$ ,  $P_{ITN+PBO-ITN} = 0.48$ ). Blood-feeding duration was longer with longer net contact time ( $F = 72.21$ ,  $df = 1$ ,  $P < 0.001$ ), independently of the net treatment ( $F = 2.36$ ,  $df = 2$ ,  $P = 0.10$ ).

The presence of insecticide reduced the percentage of mosquitoes that fed successfully. With the UTN, 60.5% (95% CI: 49.3–70.8%) of mosquitoes took a blood meal, while blood-feeding rates were only 38.9%

(95% CI: 29.1–49.5%) and 9.2% (95% CI: 3.46–19.0%) with the ITN and ITN + PBO treatments, respectively ( $\chi^2 = 45.70$ ,  $df = 2$ ,  $P < 0.001$ ; Fig. 2c). Blood-fed females spent 3.6-fold more time in contact with nets than unfed ones (unfed:  $59.8 \pm 6.22$  s; fed:  $218.7 \pm 7.46$  s) ( $\chi^2 = 110.22$ ,  $df = 1$ ,  $P < 0.001$ ). Contact times of the subset of mosquitoes that successfully took a blood meal showed a similar pattern between net types, with average contact times of  $253.4 \pm 7.49$ ,  $185.4 \pm 11.73$  and  $122.3 \pm 13.68$  s for the UTN, ITN, ITN + PBO treatments, respectively (Tukey pairwise comparisons:  $P_{ITN-UTN}$  and  $P_{ITN+PBO-UTN} < 0.001$ , and  $P_{ITN+PBO-ITN} = 0.29$ ). However, unfed mosquitoes exposed to an ITN spent more time in contact with the net compared to those exposed to an UTN, with an average contact time of  $36.5 \pm 10.82$  s for the UTN versus  $80.6 \pm 10.66$  s for the ITN and  $52.7 \pm 9.73$  s for the ITN + PBO ( $F = 11.97$ ,  $df = 2$ ,  $P = 0.002$ ; Tukey pairwise comparisons:  $P_{ITN-UTN} = 0.04$  and  $P_{ITN+PBO-UTN} = 0.88$ , and  $P_{ITN+PBO-ITN} = 0.24$ ). In one experimental block, the feeding success was slightly lower compared to the other blocks ( $F = 5.63$ ,  $df = 1$ ,  $P = 0.02$ ). Thus, while there was no difference in the time spent on the net for blood-fed mosquitoes, there was some variability between two experimental blocks for the contact time of unfed mosquitoes ( $F = 7.92$ ,  $df = 1$ ,  $P = 0.005$ ).

As observed in the first experiment, there was negligible mortality within 24 h in the UTN and ITN treatments. The ITN + PBO treatment, however, caused a substantial mortality within 24 h of exposure of 86.1% (95% CI: 75.3–93.5%). Beyond the instantaneous effects, insecticide exposure led to a reduction in long-term survival ( $\chi^2 = 146.87$ ,  $df = 2$ ,  $P < 0.001$ ) (Fig. 3). Mosquitoes exposed to an UTN had an average survival time (post exposure at 4–5 days old) of  $16.7 \pm 0.74$  days; those exposed to the ITN,  $11.5 \pm 0.62$  days; and those exposed to the ITN + PBO, just  $2.3 \pm 0.44$  days. Blood-feeding increased overall longevity ( $\chi^2 = 24.53$ ,  $df = 1$ ,  $P < 0.001$ ) by approximately 6 days for the UTN treatment and 4 days for the ITN treatment. Nonetheless, blood-fed females died more quickly after an exposure to insecticide than those exposed to a UTN ( $14.0 \pm 1.10$  days for the ITN vs  $19.0 \pm 1.00$  days for the UTN). For the ITN + PBO treatment, blood-fed mosquitoes had a marginally shorter lifespan than their nonblood-fed counterparts (average survival time post exposure of  $1.0 \pm 0$  days and  $2.5 \pm 0.49$  days, respectively;  $\chi^2 = 10.21$ ,  $df = 2$ ,  $P = 0.006$ ). The time that blood-fed mosquitoes spent on the net and feeding duration did not influence longevity. However, fed mosquitoes exposed to an UTN had a longer lifespan when they spend more time blood-feeding, which was not the case for fed mosquitoes exposed to insecticide ( $\chi^2 = 6.95$ ,  $df = 2$ ,  $P = 0.03$ ).



One additional experimental block was added to an analysis comparing the effects of the ITN against the UTN only (providing 4 blocks in total for this comparison (Additional file 1: Table A)). The presence of insecticide reduced the mean time spent on the net ( $119.6 \pm 7.73$  s for the ITN vs  $173.3 \pm 10.11$  s for the UTN) ( $F=18.02$ ,  $df=1$ ,  $P<0.001$ ). Exposure to the ITN led to a significant reduction in blood-feeding ( $\chi^2=17.20$ ,  $df=1$ ,  $P<0.001$ ), with 38.2% (95% CI 30.6–46.3%) blood-fed mosquitoes in the ITN treatment compared with 61.9% (95% CI: 53.6 to 69.8) in the UTN treatment. Mosquitoes that spent a longer period on the net were proportionally more successful in taking a blood meal ( $\chi^2=206.56$ ,  $df=1$ ,  $P<0.001$ ). When mosquitoes spent  $< 1$  min on the net, the feeding rate did not differ between the ITN and UTN treatments. However, once contact time exceeded 1 min, blood-feeding increased with contact time for the UTN treatment but did not for the ITN treatment ( $\chi^2=11.31$ ,  $df=1$ ,  $P<0.001$ ). There was an overall effect of insecticide exposure on longevity ( $\chi^2=29.34$ ,  $df=1$ ,  $P<0.001$ ). The mean survival time of unfed mosquitoes was  $14.0 \pm 0.83$  days post exposure for the UTN treatment and  $10.0 \pm 0.50$  days post exposure for the ITN treatment, while for fed mosquitoes it was  $18.2 \pm 0.78$  days post exposure and  $15.0 \pm 0.88$  days post exposure, respectively ( $\chi^2=42.09$ ,  $df=1$ ,  $P<0.001$ ) (Fig. 3).

## Discussion

In the forced exposure assays, a high percentage of mosquitoes were able to blood-feed, and increased contact time with the ITN for up to 5 min was not obviously a

factor that would result in death of the mosquitoes compared to a contact time of 1 min. On the contrary, those mosquitoes that did blood-feed lived longer, and a longer contact time with the ITN increased their survival post-exposure. We expected that a longer exposure to insecticide would decrease longevity, but instead we found a clear benefit of having more time to complete a blood meal for survival post-exposure. Other studies have also shown that ITNs fail to cause instantaneous mortality [3, 5] or fully prevent blood-feeding [3, 24, 32] against highly resistant mosquitoes, suggesting a loss of personal protection due to resistance. However, the feeding choice exposure assays provide a slightly more nuanced picture and showed that the ITN did reduce the proportion of mosquitoes that fed successfully. Unfed mosquitoes were found to spend more time in contact with the ITN than their unfed counterparts exposed to the UTN, yet fewer mosquitoes exposed to the ITN ultimately fed. This result suggests that reduced feeding was not due to the mosquitoes avoiding the net or being repelled by it, but more likely because contact with the insecticide reduced feeding capacity. In turn, insecticide exposure reduced the time spent on the net feeding. Spending less time on a net during blood-feeding does not necessarily mean that the size of the blood meal is smaller and/or insufficient for malaria transmission [36]. The presence of insecticide might reduce the capacity to engorge blood [24], or it could be hypothesized that mosquitoes take the same amount of blood in a shorter period in order to minimize their contact time with the treated net. A shorter time in contact with the treated net could lower the insecticide dosage received by blood-feeders, and this could be a behavioral adaptation of mosquitoes living in areas with a high use of ITNs. Whether similar results are observable in mosquitoes infected with malaria parasites is unclear. It is known that malaria infection alters feeding rates and blood-seeking behaviors [37, 38], but more research is needed to understand whether insecticide exposure impacts vector competence [39–42] and/or whether the presence of malaria parasites affects the expression of insecticide resistance [29, 43].

Regardless of the exposure pattern/duration there was negligible mosquito mortality within 24 h of contact with an ITN. However, this standard 24-h assessment [17] misses potential long-term effects of exposure. Data from both assays show reduced long-term survival following exposure to a standard ITN, a result consistent with the delayed mortality for highly insecticide-resistant mosquitoes reported elsewhere [18]. The experiments also highlight that the standard WHO test procedures for evaluating resistance and measuring the bio-efficacy of ITNs [17, 44] are weak indicators of how ITNs ultimately determine malaria transmission risk

and, hence, of understanding the functional significance of insecticide resistance. The WHO resistance assay uses tubes (as used in the initial assays here) to force mosquitoes into contact with filter paper treated with diagnostic doses of insecticide for 1 h [45–47], and the WHO ITN bioefficacy assay uses cones to force mosquitoes into contact with ITNs for 3 min [48–50]. Neither assay simulates how mosquitoes contact ITNs during host-searching and blood-feeding in nature [27]. According to data acquired by Diop et al. [51], mosquitoes exposed to insecticide tend to bounce on the ITN until they decide to probe and take a blood meal. How long they choose to stay on the net depends on the level of toxicity of the net as well as on the presence of a host [52]. The cup assay used in the present study provides a potential method to allow for more realistic patterns of contact with an ITN. The data reveal that the repellent and irritant effects of deltamethrin [53, 54], especially in combination with the synergist PBO [55], reduce the time spent on the treated net but do not completely prevent mosquitoes from biting through it. Interestingly, limited repellency may help maximize the sublethal toxic effects of insecticide against them, which would increase ITN efficacy [18, 20].

The data also highlight the importance of blood-feeding in the evaluation of insecticide resistance. In general, those mosquitoes that blood-fed survived insecticide exposure better than those that did not (the exception being in the ITN+PBO exposure). Whether this is because those mosquitoes able to feed during contact with an insecticide are the most resistant and robust individuals, or whether blood-feeding itself enhances expression of insecticide resistance is unclear. The observation that the longevity of mosquitoes having no access to a blood source was similar to that of mosquitoes failing to blood-feed despite the access to a blood source suggests the effect is more to do with blood-feeding than individual variation (i.e. it would be expected that the overall survival of mosquitoes with no access to blood should be greater as these mosquitoes represent a mixed population that includes the potentially more robust individuals). In addition, ingestion of a blood meal induces oxidative stress, leading to increased metabolic activity [56, 57], which could in turn result in the higher expression of detoxification enzymes [58] and perhaps also influence the toxic dose received by mosquitoes exposed to insecticide.

The PBO-treated net was found to have a clear advantage in comparison to the standard ITN, yielding reduced blood-feeding success and increased mortality rate, yet it is unclear whether these effects are due to differences in the concentrations of the active ingredient (the net with PBO has a higher deltamethrin concentration than

the standard ITN) or the action of the synergist (PBO) alone. Whatever the mechanism, the results provide an interesting perspective on the significance of resistance relative to different comparators; i.e. the impact depends on whether you consider resistance to be the difference between an ITN and an untreated net (which is like asking how bad an ITN has become), or between an ITN and a resistance-breaking net (which is like asking how much better could an ITN be).

We acknowledge that the present study used only one human host and that attraction and blood meal quality can vary between hosts. It would be interesting for future work to explore the extent to which different hosts lead to variation in patterns of feeding and subsequent lethal and sub-lethal effects of exposure (our expectation is that the qualitative differences between human hosts will likely yield less variation than the difference between the presence or absence of a host). It would also be interesting to further examine possible effects of *Plasmodium* infection on feeding behavior and exposure rates since it is the infected mosquitoes that we ultimately care about.

## Conclusion

Overall, the combination of delayed mortality and anti-feedant effects suggests that ITNs retain at least some functionality above and beyond a simple physical barrier, even against mosquitoes with 1500-fold higher resistance. These effects do not mean that ITNs are as effective against resistant mosquitoes as they are against susceptible ones. Moreover, the Permanet<sup>®</sup> 3.0 (ITN + PBO) treatment induced much greater mortality and feeding inhibition than did the standard ITN treatment, suggesting improved control potential of ‘resistance-breaking’ nets in areas of high insecticide resistance. Nonetheless, the results provide further evidence to support why ITNs might continue to contribute to reduced malaria transmission in the face of insecticide resistance. This is likely to be especially relevant in areas with high effective coverage (i.e. high ownership and use) of ITNs, as even small effect sizes at the individual level can lead to large overall effect sizes when multiplied up to community level [9, 32]. As such, the current study supports the need for further research to fully understand the epidemiological significance of resistance.

## Abbreviations

CI: Confidence Interval; GLM: Generalized linear model; ITN: Insecticide-treated net (Permanet<sup>®</sup> 2.0); ITN + PBO: Insecticide-treated net with PBO (Permanet<sup>®</sup> 3.0 roof section); PB: Priscille Barreaux; PBO: Piperonyl butoxide; UTN: Untreated net.

## Supplementary Information

The online version contains supplementary material available at <https://doi.org/10.1186/s13071-022-05220-y>.

**Additional file 1: Table A.** Sample size used for each experimental blocks. Experiment a: forced exposure in modified WHO tubes; Experiment b: variable exposure via individual feeding choice.

**Additional file 2: Dataset 1.** Forced exposure in modified WHO tube assay.

**Additional file 3: Dataset 2.** Variable exposure via individual feeding choice.

**Additional file 4: Table C.** Scripts of the statistical analysis in Rstudio.

**Additional file 5: Table B.** Statistical analysis summaries.

### Acknowledgements

The authors thanks N'Guessan Brou and the entomological team at the Institute Pierre Richet in Côte d'Ivoire for collecting the larvae in the field and helping with the rearing before experimentation.

### Authors' contributions

PB and MBT conceived the study. PB developed the methodology, performed all the behavioral assays, monitored the mosquitoes for longevity, performed the statistical analysis, prepared the figures and wrote the draft. RNG gave access to the laboratory space, and his team collected the larvae in the field. All authors reviewed and edited the draft. MBT provided financial support. All authors read and approved the final manuscript.

### Funding

This work was supported in part by the Bill & Melinda Gates Foundation, Grant No. OPP1131603.

### Availability of data and materials

All datasets generated, used and analyzed in this study are included in this published article.

### Declarations

#### Ethics approval and consent to participate

Not applicable.

#### Consent for publication

Not applicable.

#### Competing interests

The authors declare that they have no competing interests.

#### Author details

<sup>1</sup>Liverpool School of Tropical Medicine, Liverpool, UK. <sup>2</sup>Pennsylvania State University, State College, PA, USA. <sup>3</sup>University of Neuchâtel, Neuchâtel, Switzerland. <sup>4</sup>London School of Tropical Medicine, London, UK. <sup>5</sup>Vector Control Product Evaluation Centre, Institute Pierre Richet, Bouaké, Côte d'Ivoire. <sup>6</sup>University of York, York, UK.

Received: 19 September 2021 Accepted: 24 February 2022

Published online: 28 March 2022

### References

- Bhatt S, Weiss DJ, Cameron E, Bisanzio D, Mappin B, Dalrymple U, et al. The effect of malaria control on *Plasmodium falciparum* in Africa between 2000 and 2015. *Nature*. 2015;526:207–11.
- Strode C, Donegan S, Garner P, Enayati AA, Hemingway J. The impact of pyrethroid resistance on the efficacy of insecticide-treated bednets against African Anopheline mosquitoes: systematic review and meta-analysis. *PLoS Med*. 2014;11:1.
- Glunt KD, Abílio AP, Bassat Q, Bulo H, Gilbert AE, Huijben S, et al. Long-lasting insecticidal nets no longer effectively kill the highly resistant *Anopheles funestus* of southern Mozambique. *Malar J*. 2015;14:298.
- Oumbouke WA, Pignatelli P, Barreaux AMG, Tia IZ, Koffi AA, Ahou Alou LP, et al. Fine scale spatial investigation of multiple insecticide resistance and underlying target-site and metabolic mechanisms in *Anopheles gambiae* in central Côte d'Ivoire. *Sci Rep*. 2020;10:15066.
- Ranson H, Lissenden N. Insecticide resistance in African *Anopheles* mosquitoes: a worsening situation that needs urgent action to maintain malaria control. *Trends Parasitol*. 2016;32:187–96.
- Hancock PA, Hendriks CJM, Tangena JA, Gibson H, Hemingway J, Coleman M, et al. Mapping trends in insecticide resistance phenotypes in African malaria vectors. *PLoS Biol*. 2020;18:1–23.
- Riveron JM, Huijben S, Tchappa G, Tchouakui M, Wondji MJ, Tchoupo M, et al. Escalation of pyrethroid resistance in the malaria vector *Anopheles funestus* induces a loss of efficacy of piperonyl butoxide-based insecticide-treated nets in Mozambique. *J Infect Dis*. 2019;220:467–75.
- Hemingway J, Ranson H, Magill A, Kolaczinski J, Fornadel C, Gimnig J, et al. Averting a malaria disaster: will insecticide resistance derail malaria control? *Lancet*. 2016;387:1785–8.
- Churcher TS, Lissenden N, Griffin JT, Worrall E, Ranson H. The impact of pyrethroid resistance on the efficacy and effectiveness of bednets for malaria control in Africa. *Elife*. 2016;5:1.
- Barbosa S, Kay K, Chitnis N, Hastings IM. Modelling the impact of insecticide-based control interventions on the evolution of insecticide resistance and disease transmission. *Parasit Vectors*. 2018;11:1–21.
- Lindblade KA, Mwandama D, Mzilahowa T, Steinhart L, Gimnig J, Shah M, et al. A cohort study of the effectiveness of insecticide-treated bed nets to prevent malaria in an area of moderate pyrethroid resistance Malawi. *Malar J*. 2015;14:31.
- Ochomo E, Chahilu M, Cook J, Kinyari T, Bayoh NM, West P, et al. Insecticide-treated nets and protection against insecticide-resistant malaria vectors in Western Kenya. *Emerg Infect Dis*. 2017;23:758–64.
- Alout H, Labbé P, Chandre F, Cohuet A. Malaria vector control still matters despite insecticide resistance. *Trends Parasitol*. 2017;33:610–8.
- Alout H, Dabiré RK, Djogbénou L, Abate L, Corbel V, Chandre F, et al. Interactive cost of *Plasmodium* infection and insecticide resistance in the malaria vector *Anopheles gambiae*. *Sci Rep*. 2016;1:1–11.
- Kleinschmidt I, Bradley J, Knox TB, Mnzava AP, Kafy HT, Mbogo C, et al. Implications of insecticide resistance for malaria vector control with long-lasting insecticidal nets: a WHO-coordinated, prospective, international, observational cohort study. *Lancet Infect Dis*. 2018;18:640–9.
- Lindsay SW, Thomas MB, Kleinschmidt I. Threats to the effectiveness of insecticide-treated bednets for malaria control: thinking beyond insecticide resistance. *Lancet Glob Heal*. 2021;1:1–7.
- World Health Organization. Test procedures for insecticide resistance monitoring in malaria vector mosquitoes, 2nd ed. 2016. <https://apps.who.int/iris/handle/10665/250677>. Accessed 12 Mar 2022.
- Viana M, Hughes A, Matthiopoulos J, Ranson H, Ferguson HM. Delayed mortality effects cut the malaria transmission potential of insecticide-resistant mosquitoes. *Proc Natl Acad Sci USA*. 2016;113:8975–80.
- Read AF, Lynch PA, Thomas MB. How to make evolution-proof insecticides for malaria control. *PLoS Biol*. 2009;7:0001–00010.
- Thomas MB, Read AF. The threat (or not) of insecticide resistance for malaria control. *Proc Natl Acad Sci USA*. 2016;113:8900–2.
- Hughes A, Lissenden N, Viana M, Toé KH, Ranson H. *Anopheles gambiae* populations from Burkina Faso show minimal delayed mortality after exposure to insecticide-treated nets. *Parasit Vectors*. 2020;13:17.
- Barreaux AMG, Oumbouke WA, Tia IZ, Brou N, Koffi AA, Nguessan R, et al. Semi-field evaluation of the cumulative effects of a "Lethal House Lure" on malaria mosquito mortality. *Malar J*. 2019;18:298.
- Siegert PY, Walker E, Miller JR. Differential behavioral responses of *Anopheles gambiae* (Diptera: Culicidae) modulate mortality caused by pyrethroid-treated bednets. *J Econ Entomol*. 2009;102:2061–71.
- Hauser G, Thiévent K, Koella JC. The ability of *Anopheles gambiae* mosquitoes to bite through a permethrin-treated net and the consequences for their fitness. *Sci Rep*. 2019;9:8141.
- Bowman NM, Akialis K, Cave G, Barrera R, Apperson CS, Meshnick SR. Pyrethroid insecticides maintain repellent effect on knock-down resistant populations of *Aedes aegypti* mosquitoes. *PLoS ONE*. 2018;13:1–14.

26. Agramonte NM, Bloomquist JR, Bernier UR. Pyrethroid resistance alters the blood-feeding behavior in Puerto Rican *Aedes aegypti* mosquitoes exposed to treated fabric. *PLoS Negl Trop Dis*. 2017;11:e0005954.
27. Parker JE, Angarita-Jaimes N, Abe M, Towers CE, Towers D, McCall PJ. Infrared video tracking of *Anopheles gambiae* at insecticide-treated bed nets reveals rapid decisive impact after brief localised net contact. *Sci Rep*. 2015;5:1.
28. Porciani A, Diop M, Moiroux N, Kadoke-Iambi T. A pyrethroid-treated bed net increases host attractiveness for *Anopheles gambiae* s.s. carrying the *kdr* allele in a dual-choice olfactometer. *PLoS ONE*. 2017;12(7):e0164518. <https://doi.org/10.1371/journal.pone.0164518>.
29. Mulatier M, Pennetier C, Porciani A, Chandre F, Dormont L, Cohuet A. Prior contact with permethrin decreases its irritancy at the following exposure among a pyrethroid-resistant malaria vector *Anopheles gambiae*. *Sci Rep*. 2019;9:8177.
30. Diop MM, Chandre F, Rössignol M, Porciani A, Chateau M, Moiroux N, et al. Sub-lethal insecticide exposure affects host biting efficiency of *Kdr*-resistant *Anopheles gambiae*. *BioRxiv*. 2019. <https://doi.org/10.24072/pjournal.15>.
31. N'Guessan R, Darriet F, Guillet P, Carnevale P, Traore-Lamizana M, Corbel V, et al. Resistance to carbosulfan in *Anopheles gambiae* from Ivory Coast, based on reduced sensitivity of acetylcholinesterase. *Med Vet Entomol*. 2003;17:19–25.
32. Glunt KD, Coetzee M, Huijben S, Koffi AA, Lynch PA, N'Guessan R, et al. Empirical and theoretical investigation into the potential impacts of insecticide resistance on the effectiveness of insecticide-treated bed nets. *Evol Appl*. 2017;11:431–41.
33. Kulma K, Saddler A, Koella JC. Effects of age and larval nutrition on phenotypic expression of insecticide-resistance in *Anopheles* mosquitoes. *PLoS ONE*. 2013;8:8–11.
34. Guessan RN, Asidi A, Boko P, Odjo A, Akogbeto M, Pigeon O, et al. An experimental hut evaluation of PermaNet<sup>®</sup>3.0, deltamethrin–piperonyl butoxide combination net, against pyrethroid-resistant *Anopheles gambiae* and *Culex quinquefasciatus* mosquitoes in southern Benin. *Trans R Soc Trop Med Hyg*. 2010;104:758–65.
35. MacDougall C. Effect of blood meal size on mosquito response to disturbance while blood feeding on a simulated host. Master thesis. Simon Fraser University; 2005.
36. O'Donnell AJ, Rund SSC, Reece SE. Time-of-day of blood-feeding: effects on mosquito life history and malaria transmission. *Parasit Vectors*. 2019;12:1–16.
37. Thiévent K, Zilio G, Hauser G, Koella JC. Malaria load affects the activity of mosquito salivary apyrase. *J Insect Physiol*. 2019;116:10–6.
38. Cator LJ, Pietri JE, Murdock CC, Ohm JR, Lewis EE, Read AF, et al. Immune response and insulin signalling alter mosquito feeding behaviour to enhance malaria transmission potential. *Sci Rep*. 2015;5:11947.
39. Alout H, Ndam NT, Sandeu MM, Djégbé I, Chandre F, Dabiré RK, et al. Insecticide resistance alleles affect vector competence of *Anopheles gambiae* s.s. for *Plasmodium falciparum* field isolates. *PLoS ONE*. 2013;8:1.
40. Alout H, Dabiré RK, Djogbénou LS, Abate L, Corbel V, Chandre F, et al. Interactive cost of *Plasmodium* infection and insecticide resistance in the malaria vector *Anopheles gambiae*. *Sci Rep*. 2016;6:1–11.
41. Alout H, Djégbé I, Chandre F, Djogbénou LS, Dabiré RK, Corbel V, Cohuet A. Insecticide exposure impacts vector–parasite interactions in insecticide-resistant malaria vectors. *Proc R Soc B*. 2014;281:20140389.
42. Alout H, Yameogo B, Djogbénou LS, Chandre F, Dabiré RK, Corbel V, et al. Interplay between *Plasmodium* infection and resistance to insecticides in vector mosquitoes. *J Infect Dis*. 2014;210:1464–70.
43. Thiévent K, Hauser G, Elaïan O, Koella JC. The interaction between permethrin exposure and malaria infection affects the host-seeking behaviour of mosquitoes. *Malar J*. 2019;18:79.
44. WHO. Guidelines for laboratory and field testing of long-lasting insecticidal mosquito nets. 2013. <https://apps.who.int/iris/handle/10665/80270>. Accessed 12 Mar 2022.
45. Tchouakui M, Mugenzi LMJ, Menze B, Khauxha JNT, Tchappa W, Tchoupo M, et al. Pyrethroid resistance aggravation in Ugandan malaria vectors is reducing bednet efficacy. *Pathogens*. 2021;10:1.
46. Muhammad A, Ibrahim SS, Mukhtar MM, Irving H, Abajue MC, Edith NMA, et al. High pyrethroid/DDT resistance in major malaria vector *Anopheles coluzzii* from Niger-Delta of Nigeria is probably driven by metabolic resistance mechanisms. *PLoS ONE*. 2021;16:1–16.
47. Foster GM, Coleman M, Thomsen E, Ranson H, Yangalbé-Kalnone E, Moundai T, et al. Spatial and temporal trends in insecticide resistance among malaria vectors in Chad highlight the importance of continual monitoring. *PLoS ONE*. 2016;11:1–15.
48. Musa JJ, Moore SJ, Moore J, Mbuba E, Mbeyela E, Kobe D, et al. Long-lasting insecticidal nets retain bio-efficacy after 5 years of storage: implications for malaria control programmes. *Malar J*. 2020;19:1–12.
49. Yewhalaw D, Asale A, Tushune K, Getachew Y, Duchateau L, Speybroeck N. Bio-efficacy of selected long-lasting insecticidal nets against pyrethroid resistant *Anopheles arabiensis* from South-Western Ethiopia. *Parasit Vectors*. 2012;5:1.
50. Allossogbe M, Gnanguenon V, Yovogan B, Akinro B, Anagonou R, Agossa F, et al. WHO cone bio-assays of classical and new-generation long-lasting insecticidal nets call for innovative insecticides targeting the knock-down resistance mechanism in Benin. *Malar J*. 2017;16:1–11.
51. Diop MM, Moiroux N, Chandre F, Martin-herrou H. Behavioral cost & overdominance in *Anopheles gambiae*. *Plos ONE*. 2015;1–12. <https://doi.org/10.1371/journal.pone.0121755>.
52. Takken V, Verhulst NO. Chemical signaling in mosquito-host interactions: the role of human skin microbiota. *Curr Opin Insect Sci*. 2017;20:68–74.
53. Dufour I, Achee NL, Roberts DR, Grieco JP. Contact irritancy and spatial repellency behaviors in *Anopheles albimanus* Wiedemann (Diptera: Culicidae) collected in Orange Walk, Belize, C.A. *J Vector Ecol*. 2009;34:232–7.
54. Achee NL, Sardelis MR, Dufour I, Chauhan KR, Grieco JP. Characterization of spatial repellent, contact irritant, and toxicant chemical actions of standard vector control compounds. *J Am Mosq Control Assoc*. 2009;25:156–67.
55. Martin JL, Moshia FW, Lukole E, Rowland M, Todd J, Charlwood JD, et al. Personal protection with PBO-pyrethroid synergist-treated nets after 2 years of household use against pyrethroid-resistant *Anopheles* in Tanzania. *Parasit Vectors*. 2021;14:1–8.
56. Graça-Souza AV, Maya-Monteiro C, Paiva-Silva GO, Braz GRC, Paes MC, Sorgine MHF, et al. Adaptations against heme toxicity in blood-feeding arthropods. *Insect Biochem Mol Biol*. 2006;36:322–35.
57. Oliver SV, Brooke BD, Sies H, Martindale J, Holbrook N, Ahmad S, et al. The role of oxidative stress in the longevity and insecticide resistance phenotype of the major malaria vectors *Anopheles arabiensis* and *Anopheles funestus*. *PLoS ONE*. 2016;11:e0151049.
58. Spillings BL, Coetzee M, Koekemoer LL, Brooke BD. The effect of a single blood meal on the phenotypic expression of insecticide resistance in the major malaria vector *Anopheles funestus*. *Malar J*. 2008;7:226.

## Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more [biomedcentral.com/submissions](https://biomedcentral.com/submissions)

