Centers for Disease Control Light Traps for Monitoring *Anopheles arabiensis* Human Biting Rates in an Area with Low Vector Density and High Insecticide-Treated Bed Net Use

Christen M. Fornadel,* Laura C. Norris, and Douglas E. Norris
The W. Harry Feinstone Department of Molecular Microbiology and Immunology, The Johns Hopkins Malaria Research Institute, Johns Hopkins Bloomberg School of Public Health, Baltimore, Maryland

Abstract. Human landing catches (HLCs) are currently the preferred method to determine vector human biting rates (HBRs), which are key determinants of entomologic inoculation rates and important measures for assessing the impact of vector control efforts. Although HLCs are the most direct means of establishing HBRs, they are labor-intensive, and their use is facing increasing ethical concerns. The relationship between Centers for Disease Control (CDC) light traps and HLC collections was evaluated in Macha, Zambia during the 2007–2008 and 2008–2009 rainy seasons. A CDC light trap captured on average 1.91 (95% confidence interval = 1.16–2.28) times as many *An. arabiensis* per night as an indoor HLC. Additionally, nets treated with deltamethrin did not affect the numbers of *An. arabiensis* collected. Our results suggest that in regions where use of vector control interventions is high and vector densities are low, CDC light traps can be used to monitor *An. arabiensis* HBRs.

INTRODUCTION

In the fight against malaria and the push toward eradication, interventions must be effectively used and accurately evaluated. Current strategies to reduce malaria transmission rely heavily on vector control, specifically the use of insecticide-treated bed nets (ITNs), indoor residual spraying, and source reduction. The most direct method for assessing these vector management measures is the entomologic inoculation rate (EIR) because it quantifies the tendency of a mosquito population to transmit infectious sporozoites to humans. The EIR, defined as the number of infectious bites received by an individual per unit time, is calculated by multiplying the proportion of mosquitoes in a vector population harboring sporozoite-stage parasites in their salivary glands by the nightly biting pressure of the vector on a human population (the human biting rate [HBR]). Human biting rates for particular vectors can be highly variable, even at a fine geographic scale.

The gold standard method for determining the HBR is the human landing catch (HLC) because mosquitoes are captured by aspiration as they land and attempt to feed on collectors. However, in many regions where vector control efforts are underway, extensive use of HLCs may not be practical. In addition to being non-standardized because of variability in the attractiveness and skill of collectors, HLCs are extremely labor-intensive, which limits the number of data points that may be simultaneously collected. Additionally, they require vigilance throughout the night by collectors and intense supervision to ensure that the information gathered is reliable. Furthermore, as has been noted by others, there are increasing ethical and worker safety concerns that this collection method increases the risk of exposure to infectious mosquitoes. The ethical dilemma is compounded in areas of drug-resistant malaria and when collectors would otherwise have the opportunity to be protected from infectious bites by sleeping under an ITN. Institutional Review Boards have begun to voice concerns about use of HLCs, going so far as to deem this method an occupational hazard.

Consequently, work has been conducted to evaluate alternative methods of determining HBRs that would be as sensitive as the HLC, and be cost-effective, exposure-free, and widely deployable. Indoor resting collections are largely unsuitable for this purpose because insecticides on walls or bed nets inherently reduce estimates of indoor biting by pyrethrum spray catches. Likewise, the recently developed Mbita traps have proven not to be sensitive enough for collections in areas of low mosquito densities. One promising alternative to the HLC is the Ifakara tent trap being developed in Tanzania. This collection method has been shown to correlate well with HLC collections independent of vector density, but it is still being modified and is not yet available as a commercial product.

Consequently, Centers for Disease Control (CDC) light traps hung beside occupied beds protected by bed nets remain a preferred alternative to an indoor HLC for collecting host-seeking vectors over a wide range of mosquito densities. The CDC light traps are affordable, easy to use, and have relatively high sampling efficiency, although the relationship between CDC trap and indoor HLC collections needs to be verified locally for each study area (Table 1) because vector species composition and intraspecific variation in feeding and resting behavior can have a significant impact on the quantitative association between the two methods.

As part of the National Malaria Strategic Plan of Zambia, the Macha area of Southern Province received 4,800 ITNs in 2007 (Thuma PE, unpublished data), which largely eliminated use of insecticide spray catches to monitor *Anopheles arabiensis*, the principal vector of *Plasmodium falciparum* in the region. With both practical and ethical restrictions on the widespread use of HLCs to assess differences in EIRs across the Macha region, studies to evaluate the relationship between CDC light trap and indoor HLC collections were performed in Macha, an area with overall low vector densities, during the 2007–2008 and 2008–2009 rainy seasons. Investigations of the impact of insecticide treatment of bed nets on the numbers of *An. arabiensis* collected by CDC light trap were conducted in conjunction with these studies because of the high rate of ITN use in the research area.

*Address correspondence to Christen M. Fornadel, The W. Harry Feinstone Department of Molecular Microbiology and Immunology, Johns Hopkins Bloomberg School of Public Health, 615 North Wolfe Street, Baltimore, MD 21205. E-mail: cfornade@jhsph.edu*
MATERIALS AND METHODS

Mosquito collecting and handling. The Johns Hopkins Malaria Research Institute field station is located in Macha, Zambia at 16.39292S, 26.79061E. Mosquitoes were collected for this study in two village areas, Chidakwa and Namwalinda, both located <10 km from the field station. Collections were performed during the peak of the rainy season, January through April in 2008 and 2009.

For the comparison between indoor HLC and CDC light trap collections, three houses were chosen in Namwalinda and three houses were chosen in Chidakwa. The HLC and CDC light trap collections were performed in the same houses on alternating nights, with CDC light traps hung next to occupants sleeping under untreated bed nets. If the trapping room had more than one bed, the other occupants were instructed to use their existing bed nets, or if they had none, additional nets were provided. Six teams of two HLC collectors were randomly rotated among the houses on successive collection nights so that each HLC collection team collected at each house before the rotation began again. The HLC and CDC trap collections were performed from 7:00 pm to 7:00 am up to 3 times per week for a given house for a total of 176 trap nights for each collection method. Because HLC collections were only performed for the first half hour of every hour, the CDC light trap collections were compared with the total nights for each collection method. Whether the differences observed between the nightly means for this study yielded no mosquitoes, a negative binomial regression analysis was performed using STATA version 10 (StataCorp., College Station, TX). The negative binomial distribution has been used previously to model overdispersed mosquito count collections.

The difference in the sampling efficiency of CDC light traps relative to the HLC reference method was evaluated after controlling for month and year of collection and clustering on household. Using the negative binomial model, we assumed that total mosquito counts followed a Poisson distribution in which there is an overdispersion parameter to account for variance that is greater than that expected under a true Poisson distribution. The log of the expected counts is modeled as the function \( \log(E(Y_{ijk})) = \alpha + \beta X + \epsilon_{ijk} \), where \( Y_{ijk} \) is the monthly mosquito count for household \( i \) in month \( j \) and year \( k \) and \( X \) are the covariates of sampling method, collection year, and collection month. This method enabled standard errors to be adjusted for correlated observations within collection households. Density dependence was evaluated by generating a new variable to present the tertiles of the total combined numbers of An. arabiensis collected by CDC trap and HLC. Each tertile represented one of three parts of the ordered dataset distribution, and each contained one-third of the population. The above analysis was then repeated with the inclusion of interaction terms for sampling method and tertile density.

The hypothesis that CDC light traps collect the same number of mosquitoes whether suspended next to people sleeping under deltamethrin-treated or untreated bed nets was examined by using paired t-tests. For the 2009 data, the significance

Table 1

<table>
<thead>
<tr>
<th>Study</th>
<th>Relative sensitivity of CDC trap versus that of HLC (95% CI)</th>
<th>Density dependent</th>
<th>Species composition</th>
<th>Location</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.33 (0.24–0.46)</td>
<td>Yes</td>
<td>Not reported</td>
<td>Ulanga District, Tanzania</td>
<td>Okumu and others, 2008(^2)</td>
</tr>
<tr>
<td>2</td>
<td>0.56 (0.49–0.66)</td>
<td>No</td>
<td>An. arabiensis</td>
<td>Ahero, Kenya</td>
<td>Mathenge and others, 2005(^3)</td>
</tr>
<tr>
<td>3</td>
<td>1.06 (0.88–1.26)</td>
<td>No</td>
<td>An. gambiae s.s.</td>
<td>Bo District, Sierra Leone</td>
<td>Mbagi and others, 2002(^4)</td>
</tr>
<tr>
<td>4</td>
<td>1.09 (0.98–1.29)</td>
<td>No</td>
<td>&gt;90% An. gambiae s.s.</td>
<td>Muheza District, Tanzania</td>
<td>Lines and others, 1991(^5)</td>
</tr>
<tr>
<td>5</td>
<td>1.08</td>
<td>No</td>
<td>~70% An. gambiae s.s.</td>
<td>Nounjou, Burkino Faso</td>
<td>Costantini and others, 1998(^6)</td>
</tr>
<tr>
<td>6</td>
<td>1.10</td>
<td>No</td>
<td>&lt;40–100% An. gambiae s.s.</td>
<td>Bagamoyo District, Tanzania</td>
<td>Davis and others, 1995(^7)</td>
</tr>
<tr>
<td>7</td>
<td>1.3</td>
<td>Yes</td>
<td>12% An. gambiae s.s.</td>
<td>Ulanga District, Tanzania</td>
<td>Govella and others, 2009(^8)</td>
</tr>
<tr>
<td>8</td>
<td>1.86 (1.73–2.00)</td>
<td>No</td>
<td>55% An. arabiensis</td>
<td>Suba District, Kenya</td>
<td>Mathenge and others, 2004(^9)</td>
</tr>
</tbody>
</table>

\(^{*}\) CDC = Centers for Disease Control; HLC = human landing catch; CI = confidence interval.
\(^{†}\) Comparison between 1 CDC trap collection and 2 HLC collectors.
\(^{‡}\) Comparison between 1 CDC trap collection and 2 HLC collectors.

DNA isolation and polymerase chain reaction. Collected mosquitoes were identified morphologically\(^13\) at the field station and individually placed in tubes containing silica gel desiccant (Fisher Scientific, Fair Lawn, NJ) and cotton for stable storage until they were processed for molecular analysis. Heads/thoraces were separated from abdomens before homogenization and rehydrated at room temperature in 20 \( \mu L \) of double-distilled water for 10 minutes. The DNA was extracted from mosquito heads/thoraces and abdomens by a modified salt procedure as previously described.\(^16\) DNA pellets were resuspended in 50 \( \mu L \) of double-distilled water. Host source of blood fed mosquitoes was determined by polymerase chain reaction (PCR) on abdominal DNA.\(^14,16\) The DNA from head/thorax extractions was used to confirm species by the PCR assay of Scott and others.\(^17\) Additionally, head/thorax DNA was screened for \( P. falciparum \) by nested PCR.\(^18\)

Data analysis. As an exploratory step, the An. arabiensis totals for all trap nights for both the CDC light trap and HLC methods were graphed and Pearson’s correlation coefficient for linear association was calculated. Initially, we used the Bland-Altman analysis\(^19\) used in previous studies\(^5,8,20,21\) to assess whether the differences observed between the nightly means calculated for each collection method were density dependent and to estimate the bias associated with the collection methods. However, handling non-normally distributed data by using a logarithmic transformation with the addition of one to count totals of zero may bias the results.\(^22\) Therefore, to avoid artificially converting zeros to non-zero values when many sampling occasions in this study yielded no mosquitoes, a negative binomial regression analysis was performed using STATA version 10 (StataCorp., College Station, TX). The negative binomial distribution has been used previously to model overdispersed mosquito count collections.

The difference in the sampling efficiency of CDC light traps relative to the HLC reference method was evaluated after controlling for month and year of collection and clustering on household. Using the negative binomial model, we assumed that total mosquito counts followed a Poisson distribution in which there is an overdispersion parameter to account for variance that is greater than that expected under a true Poisson distribution. The log of the expected counts is modeled as the function \( \log(E(Y_{ijk})) = \alpha + \beta X + \epsilon_{ijk} \), where \( Y_{ijk} \) is the monthly mosquito count for household \( i \) in month \( j \) and year \( k \) and \( X \) are the covariates of sampling method, collection year, and collection month. This method enabled standard errors to be adjusted for correlated observations within collection households. Density dependence was evaluated by generating a new variable to present the tertiles of the total combined numbers of An. arabiensis collected by CDC trap and HLC. Each tertile represented one of three parts of the ordered dataset distribution, and each contained one-third of the population. The above analysis was then repeated with the inclusion of interaction terms for sampling method and tertile density.

The hypothesis that CDC light traps collect the same number of mosquitoes whether suspended next to people sleeping under deltamethrin-treated or untreated bed nets was examined by using paired t-tests. For the 2009 data, the significance
of the difference in the proportion of unfed mosquitoes collected in light traps hung next to treated or untreated nets was tested by using a two-proportion z-test.

RESULTS

For the CDC light trap/HLC comparison collections, 497 An. gambiae complex mosquitoes were captured: 19 An. quadrimaniatus and 478 An. arabiensis. However, 113 of the An. arabiensis were caught during a single trap night (108 from the CDC trap and 5 from the corresponding HLC). This trap night was an extreme outlier because the next largest number of An. arabiensis caught in a trap night was 47. Because inclusion of this data point significantly skewed the results obtained it was dropped from the analysis, leaving a total of 365 An. arabiensis (225 from CDC traps and 140 from HLCs) collected during 175 trap nights. All mosquitoes caught by light trap or landing catch were negative for P. falciparum sporozoites.

Preliminary analysis showed that CDC light traps tended to catch more mosquitoes overall (Figure 1). As expected, there was a statistically significant correlation between the numbers of An. arabiensis caught by CDC trap and indoor HLC collection ($r = 0.51, P < 0.001$). The negative binomial regression analysis performed on the nightly collection data revealed that a CDC light trap captured on average 1.91 (95% confidence interval = 1.61–2.28, $P < 0.001$) times as many An. arabiensis per night as an indoor HLC pair, after controlling for month and year. All coefficients for month and year of collection were negative for $P. falciparum$ infection interval $= 1.61–2.28$, and 78.8% (Table 2) of An. arabiensis captured by light trap were unfed when analyzed in 2008 ($P = 0.36$), and 96 pairs were examined in 2009 ($P = 0.26$). No statistically significant difference was observed in the proportion of unfed mosquitoes caught by traps suspended next to treated or untreated nets during the 2009 season ($P = 0.355$) (Table 2). Of those mosquitoes that were blood fed, 98% of blood meals were taken from human hosts.

DISCUSSION

In this study, CDC light traps in Macha, Zambia caught on average 1.91 times as many mosquitoes as a pair of indoor HLC collectors. Although this relative sampling efficiency is higher than those previously published for the An. gambiae complex (Table 1), the difference might be largely explained by the sibling species composition at each locality and the considerable variation in An. arabiensis foraging behavior throughout Africa. There has been only one other CDC/HLC comparison study conducted in an area where An. arabiensis was the sole sibling species present. In Ahero, Kenya, a CDC light trap captured approximately 60% of the number of An. arabiensis as an HLC.$^{11}$ However, unlike in Mcha where An. arabiensis is highly anthropophilic, the rice irrigation region of Kenya where the previous work was undertaken, An. arabiensis is known to be predominate zoophilic.$^{24}$ The CDC traps in this study might also have caught more mosquitoes because the population of An. arabiensis in Macha displays higher post-prandial endophily. The light traps may be attracting not only host-seeking An. arabiensis, but also some proportion of indoor resting mosquitoes.$^{20}$ In our comparison of CDC light trap efficiency by bed net type, on average only 78.8% (Table 2) of An. arabiensis captured by light trap were unfed. It has been shown that the added stimulus of light from incandescent bulbs increases the numbers of An. gambiae s.l. caught by CDC light trap by approximately 2.5 times.$^{5}$ Perhaps this added stimulus was enough to attract a portion of non-host-seeking An. arabiensis. Furthermore, the positioning of the CDC light trap in relation to the host acting as bait has been shown to have a significant impact on catch sizes.$^{35}$ It is therefore possible that some of the observed difference in the sampling efficiency of CDC traps between this study and others might be caused by trap placement.

Although it has been reported that the efficiency of CDC light traps as a substitute for HLCs may vary as a function of vector abundance,$^{7,20}$ most studies in Africa have shown the correlation between the two methods to be independent of density.$^{7,6,8,10,11,21}$ Importantly, the relative sampling efficiency of CDC light traps in this study was not density dependent. Therefore, even with the relatively low numbers of An. arabiensis

![Figure 1](image)

**Figure 1.** Nightly scatter plot of paired *Anopheles arabiensis* collections for 175 trap nights. The solid line indicates the regression line of the dataset and is shown in comparison with the dashed identity line. Each point represents the collections from one household per night.  

**Table 2**  

<table>
<thead>
<tr>
<th>Year (no. mosquitoes)</th>
<th>Trap or net</th>
<th>No. <em>An. quadrimaniatus</em></th>
<th>No. <em>An. arabiensis</em> (% unfed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2008 (120)</td>
<td>Bayer K-O Tab</td>
<td>11</td>
<td>60</td>
</tr>
<tr>
<td></td>
<td>(deltamethrin)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Untreated bed net</td>
<td>6</td>
<td>78</td>
</tr>
<tr>
<td>2009 (96)</td>
<td>Permanet</td>
<td>2</td>
<td>210</td>
</tr>
<tr>
<td></td>
<td>(deltamethrin)</td>
<td></td>
<td>80.8</td>
</tr>
<tr>
<td></td>
<td>Untreated bed net</td>
<td>0</td>
<td>171</td>
</tr>
</tbody>
</table>

* CDC = Centers for Disease Control; ITNs = insecticide-treated bed nets.
collected in Macha, we should be able to predict what the results of an HLC collection would have been from the catch total of a CDC light trap. Because neither trapping method yielded sporozoite-positive mosquitoes, we were unable to assess the influence of trapping method on sporozoite prevalence. Similar studies have found conflicting results regarding the influence of trapping method on infection rates. Some studies have shown higher sporozoite rates in *An. gambiae* s.l. collected by light trap than in those collected by HLCs, whereas other investigations have found the methods to yield similar rates. Further investigation will need to be undertaken in areas with higher *P. falciparum* transmission rates to determine if trapping method has an effect on sporozoite prevalence in southern Zambia.

It has been suggested that the exicto-repellent properties of ITNs can reduce the numbers of mosquitoes that enter sleeping huts or cause those that do to exit more quickly. However, in our study, the presence of ITNs did not affect the utility of CDC traps. This is a critical finding because surveys completed during the 2008 and 2009 rainy seasons in Macha found that 72–86% of the population, depending on village area, reported sleeping under a bed net treated with deltamethrin the previous night (unpublished data). If CDC light traps are used as a replacement for indoor HLCs, it would be impractical and unethical in an area with high ITN use to replace all existing nets with untreated bed nets for monitoring purposes. Previous studies have shown a slight decrease in or no effect on the sampling efficiency of CDC traps in the presence of ITNs. However, although surrounding occupied beds were covered with ITNs, the CDC traps in both of these investigations were hung next to untreated bed nets. Our data are the first to directly compare the catch totals of *An. arabiensis* by CDC trap hung beside each bed net type. These results will have to be confirmed elsewhere, but the finding is encouraging in that control programs throughout southern Africa might benefit from the ability to use existing ITNs in conjunction with CDC traps as part of their surveillance efforts.

A major drawback to using CDC light traps as a substitute for HLCs is that although HLCs can be performed inside and outside, CDC light traps are known to be ineffective for collecting *An. gambiae* complex mosquitoes when hung outdoors. Consequently, light traps can only be used to sample the indoor-biting fraction of the vector population. Although *An. arabiensis* in Macha has been shown to be predominately exophagic, CDC light traps were efficient at collecting the indoor host-seeking mosquitoes that are responsible for most bites after people have gone to bed. We acknowledge that HLCs remain the best sampling technique to glean information on the degree of exophagy in a population and vector biting times. However, when HLCs become prohibitive because of logistical issues and ethical concerns, the results presented here indicate that CDC light traps may be used as a stopgap measure to sample indoor, host-seeking *An. arabiensis* until better tools are developed, proven effective and cost-efficient, and become widely available.

This is the first study to report that the relationship between a CDC light trap and indoor HLC collection can be established for *An. arabiensis* in an area of low vector density where ongoing vector control interventions are being used. The situation in Macha is one that will be seen more and more as the malaria map begins to shrink. Although the Southern Province of Zambia has historically had hyperendemic transmission of *P. falciparum*, there has been a significant decrease in the number of malaria cases in the Macha region since 2003 when Zambia adopted artemisinin combination therapy as the national standard for the treatment of uncomplicated malaria. As malaria rates in Macha and similar sites throughout Africa are reduced further through vector control efforts, it will become even more important to monitor EIRs over a wider region to identify remaining foci of malaria transmission.

To calculate HBRs for this purpose, we show that as long as known caveats are kept in mind and the relationship between catch methods is established locally, CDC light traps can be used next to treated bed nets as a substitute for indoor HLCs.

Received February 9, 2010. Accepted for publication June 29, 2010.

Acknowledgments: We thank Dr. Marie Diener-West for her statistical insights; Shadrack Habbanti for his time and effort spent coordinating field team operations in Zambia; Musapa Mulenga for managing our collections in Macha; and our field mosquito collectors (Corrence Munsanje, Nathan Phiri, Clement Mwaanga, Fines Mwaanga, Twaambo Moono, Gift Shapamani, Miyanda Moono, Guide Hansumo, Mathias Muleka, Chaltone Munsanje, Pathias Chibambo, Malony Mulota, Haggard Musyutilia, Cliff Singanga, Paul Haakaloba, and Ojukwu Himunwe) for their assistance.

Financial support: This study was supported by the Johns Hopkins Malaria Research Institute (Douglas E. Norris), National Institutes of Health training grant T32AI007417 (Laura C. Norris), and a Simpson Student Award from the Tropical Medicine Dinner Club of Baltimore and a Johns Hopkins Bloomberg School of Public Health Summer Scholarship (Christen M. Fornadel).

Authors’ addresses: Christen M. Fornadel, Laura C. Norris, and Douglas E. Norris, The W. Harry Feinstone Department of Molecular Microbiology and Immunology, Johns Hopkins Bloomberg School of Public Health, 615 North Wolfe Street, Baltimore, MD 21205, E-mails: cfornade@jhsph.edu, lnorris@jhsph.edu, and dnorris@jhsph.edu.

REFERENCES

9. Mirabello L, Vineis JH, Yanoviak SP, Scarpassa VM, Povoa MM, Padilla N, Achec NL, Conn JE, 2008. Microsatellite data suggest...
significant population structure and differentiation within the malaria vector *Anopheles darlingi* in Central and South America. *BMC Ecol* 8:3.


