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Continuing Intense Malaria Transmission in Northern Uganda

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Abstract. Recent reports of reductions in malaria transmission in several African countries have resulted in optimism that malaria can be eliminated in parts of Africa where it is currently endemic. It is not known whether these trends are global or whether they are also present in areas where political instability has hindered effective malaria control. We determined malaria parasite carriage and age-dependent antibody responses to Plasmodium falciparum antigens in cross-sectional surveys in Apac, northern Uganda that was affected by political unrest. Under-five parasite prevalence was 55.8% (115/206) by microscopy and 71.9% (41/57) by polymerase chain reaction. Plasmodium ovale alone, or as a co-infection, was detected in 8.6% (12/139) and Plasmodium malariae in 4.3% (6/139) of the infections. Age seroprevalence curves gave no indication of recent changes in malaria transmission intensity. Malaria control remains a tremendous challenge in areas that have not benefited from large-scale interventions, illustrated here by the district of Apac.

INTRODUCTION

Despite reported reductions in malaria transmission intensity in several African regions, malaria remains one of the most important public health problems in sub-Saharan Africa with an estimated 863,000 deaths annually. Widespread use of insecticide-treated nets (ITNs), effective vector control, increased urbanization, and treatment of uncomplicated falciparum malaria with artesinin-based combination therapies (ACT) have all been assumed to contribute to the reported reductions in malaria incidence, although in some areas these reductions were observed before control measures were scaled up. This widespread decline in the burden of malaria has resulted in optimism that malaria can be eliminated in parts of Africa where malaria is currently endemic. However, reports of reductions in malaria transmission mostly originate from areas that have been involved in intensive and effective malaria control programs and sustained implementation of health care, with some valuable findings from less well-controlled settings. It is uncertain if trends of declining transmission intensity are evident across Africa or whether they are apparent in areas where political instability and economic arrest have hindered effective control and surveillance of infectious diseases. Conflict and human insecurity pose considerable challenges by causing a breakdown in health delivery systems and a loss of human and financial resources for health programs. In those areas, malaria control is likely to have been less efficiently implemented and maintained and, as a result, malaria transmission intensity may have remained unaltered, or malaria may have re-emerged in areas where it was previously under control.

In this study, we determined the current level of malaria transmission intensity in the Apac district in northern Uganda, a remote region that was previously described as holoendemic for malaria. Northern Uganda has been involved in conflict since the early 1980s with the Lord’s Resistance Army as the main rebel group that continues to be a threat to the region up to the present day. This conflict hindered economic development in northern Uganda and resulted in a lower access to healthcare compared with other regions in Uganda. The area of Apac was affected by political unrest in the early 1990s. Although health facilities remained functioning throughout the conflict, serious supply shortages affected the quality of care.

The aim of this study was to determine the current prevalence of Plasmodium falciparum parasite carriage by microscopy and polymerase chain reaction (PCR) and to use age-dependent antibody responses to P. falciparum circumsporozoite (CSP) antigen and blood-stage antigens apical membrane antigen-1 (AMA-1) and merozoite surface protein-1 (MSP-1) to look for evidence of recent changes in transmission intensity.

METHODS

The study was conducted in Apac Sub-County, a rural district in Northern Uganda located between Kwania Lake and the Victoria Nile (latitude 1.985; longitude 32.535). Apac District covers an area of 6,684 square kilometers and ranges in altitude between 1,350 and 1,500 meters above sea level. The rainfall pattern is bimodal with a dry season from November to February and two short rainy seasons from April to May and from September to October. According to surveys conducted in 2001–2002, this area experiences perennial holoendemic malaria with parasite prevalence rates of 70–90% in children < 10 years of age.

The entomological inoculation rate was estimated at > 1,500 infective bites per person per year and the major vector responsible for transmission is Anopheles funestus. Plasmodium falciparum is the dominant parasite species, Plasmodium malariae being responsible for ~3% of the infections and Plasmodium ovale was previously not observed.

Ethical approval was obtained from the ethical review committee of the London School of Hygiene and Tropical Medicine (no. 5539), the ethical committee of the Medical Biotech Laboratory, and the national ethical committee of Uganda.

Data collection. Subjects were recruited in October 2009 in four parishes. Sampling was done in Apac District Hospital, two health facilities in the parishes of Abedi and Akere, and a primary school in the parish of Atopi. Before the sampling days, community meetings were organized to explain the purpose of the study and to invite people to attend sampling points.
At the health facilities and the hospital, all individuals attending the facilities for clinical care, antenatal visits, or who came specifically to benefit from the screening offered by this study were selected for enrollment together with accompanying family members or guardians. This approach was previously shown to provide an estimate of malaria-specific antibody prevalence that is comparable to that obtained through community surveys. Before sampling at the school in Atopi, a community meeting was organized and parents were informed of the survey through the school’s pupils. All inhabitants of Atopi who attended the sampling point, including pupils and their parents or guardians, were eligible for enrollment and were sequentially enrolled until the sample size was reached. We aimed to recruit 200–300 individuals per parish of whom half were <15 years of age. This sample size was based on a previous study where this number of participants was found to be sufficient for a reliable determination of transmission intensity by serological markers of malaria exposure. To ensure a balanced representation of all age groups, essential for determining the age-dependent seroconversion rates (SCRs), seven age categories were defined per parish (1–2 years, N = 45; 3–5 years, N = 45; 6–10 years, N = 40; 11–15 years, N = 40; 16–25 years, N = 40; 26–55 years, N = 40; and >30 years, N = 50) and questionnaires were printed in pre-defined quantities in different colors for each age group. Questionnaires contained clinical information, demographic data, information on the use of antimalarial drugs, and protective antimosquito measures. As soon as the sample size for an age category was reached, no further individuals were enrolled for this category but enrollment continued for other age categories.

Written informed consent or, in case of illiteracy, consent by thumb print, was obtained from each participant ≥15 years of age and from parents or guardians of younger individuals. Each individual enrolled in the study underwent a clinical examination, during which axillary temperature was measured twice using a digital thermometer and the higher of the two values was recorded. A single blood sample was obtained by finger prick (~0.3 mL) for thick and thin blood films, for filter paper blood collection (Whatman 3 mm, Maidstone, UK) and for Rapid Diagnostic Tests (RDT; Paracheck Orchid Biomedical Systems, Goai, India) for malaria. This RDT has an estimated detection rate of 97.5% for parasite densities >2,000 parasites/μL and 54.4% for parasite densities of 200/μL. Filter papers were air-dried and stored in plastic bags with silica desiccant gel type III (Sigma, Dorset, UK), stored at −20°C in the laboratory until further processing. Thick blood films were Giemsa-stained in the field and read after completion of the study. Clinical diagnosis was based on the result of the RDT; RDT-positive individuals with (reported) fever were treated with artesether-lumefantrine (Lonart; Bliss Gvs Pharma Ltd., India) according to national guidelines. The first dose was given under supervision; the remaining five doses were given to the participant/guardian for treatment at home.

**Parasite detection by microscopy and PCR.** Microscopic slides were examined for the presence of parasites in 100 high-power fields by two experienced microscopists; the average parasite density of the two readings was recorded and a third microscopist consulted in case of disagreement. Asexual parasites were counted against 200 white blood cells and converted to parasites/μL by assuming a density of 8,000 white blood cells/μL blood. We explored the value of PCR for parasite detection in a single parish, Abedi. DNA was extracted from all filter paper blood spots from Abedi, using the chelex method and tested for the presence of P. falciparum, P. vivax, P. malariae, and P. ovale in the nested PCR approach originally described by Snounou and others and Padley and others. Samples that were negative by PCR were rescreened by PCR using as a template DNA extracted using QIAamp DNA Mini Kit (QIAGEN, Hilden, Germany).

**Enzyme-linked immunosorbant assay (ELISA).** Antibodies were elicited from filter paper blood spots and assayed by ELISA, as described in the online protocol by Corran and others. Briefly, a 3.5-mm circle was cut from the spot and placed into 300 μL of phosphate buffered saline with 0.5% Tween 20 (PBS-T) and 0.05% sodium azide, approximately equivalent to a 1/200 serum dilution. Immunoglobulin G (IgG) antibodies against circumsporozoite protein (CSP), apical membrane antigen (AMA-1), and merozoite surface protein 1 (MSP-1) were detected by ELISA using standard methodology. Recombinant MSP-1, a synthetic peptide CSP (NANP4) were coated onto ELISA plates overnight at 4°C at a concentration of 0.5 mg/mL, respectively. Plates were washed using PBS plus 0.05% Tween 20 (PBS/T) and blocked with 1% (w/v) skimmed milk powder (Marvel, UK) in PBS/T. Samples were added in duplicate to each plate at a serum dilution of 1:200 for CSP, 1:2000 for AMA-1, and 1:750 for MSP-1 in 1% bovine serum albumin (BSA) in PBS/T. A positive control of pooled hyperimmune serum collected from adults resident in a malaria-endemic area was included in duplicates on each plate to allow standardization of day-to-day and plate-to-plate variation; serum from malaria-naive Europeans was included in each assay as negative controls. After overnight incubation at 4°C the plates were washed and horseradish peroxidase-conjugated rabbit anti-human IgG (Dako Ltd., High Wycombe, United Kingdom) (1/5,000 in PBS/T) was added to all wells. All plates were developed using OPD/H₂O₂ substrate solution and reactions were stopped with 2 M H₂SO₄. Plates were read immediately at 492 nm and optical density (OD) values recorded.

**Data analysis.** All data were double entered and validated in Microsoft Access (Redmond, WA); inconsistencies were verified against the original questionnaire. Data were imported into Stata 11.0 (Statata Statistical Software, StataCorp, College Station, TX) for statistical analysis. Fever was defined as a temperature ≥37.5°C, submicroscopic parasitemia was defined as parasitemia by PCR in the absence of microscopically confirmed parasite carriage. Parasite density was presented as geometric mean in microscopically positive parasite carriers only with the 25th and 75th percentile (interquartile range, IQR). Duplicate OD results in ELISA assays were averaged and normalized against the positive control sample on each plate. A cut-off above which samples were deemed antibody positive was defined using a mixture model as previously described. Briefly, the distribution of normalized OD values was fitted as the sum of two Gaussian distributions (a narrow distribution of seronegatives and a broader distribution of seropositives) using maximum likelihood methods. The mean OD of the Gaussian corresponding to the seronegative population plus three standard deviations was used as the cut-off for seropositivity. The seroconversion rate (SCR or λ) was estimated by fitting a simple reversible catalytic model to the measured seroprevalence, stratified into yearly age groups, using maximum likelihood methods. For
these models only individuals' ≥ 1 year of age were included to avoid the effect of maternally derived antibodies in infants. Evidence for temporal changes in SCR was explored by fitting models in which the SCR is allowed to change at a single time point. The significance of the change was identified using likelihood ratio tests against models with no change, and profile likelihoods were plotted to determine confidence intervals (CIs) for the estimated time of the change. The titer of the antibody response was estimated by using the formula dilution/(maximum OD/[OD test serum-minimum OD] − 1), where the maximum OD was the maximum value of the standard curve and the minimum OD the lowest value of the negative control. The titer was used as an indicator of antibody density in the analyses. Categorical variables were analyzed using the χ² test or χ² test for trend. Student’s t test, analysis of variance, or non-parametric equivalents were used when comparing continuous variables. Logistic and linear regression models were used to adjust binary and continuous variables for potential confounding. Titer was log₁₀ transformed for analysis and the exponentiated regression coefficients with 95% CI were presented.

RESULTS

Parasite carriage by microscopy and PCR. A total of 883 individuals were enrolled from four parishes (202–251 individuals per parish) with overall parasite prevalence by microscopy of 37.5% (219/851). Only *P. falciparum* was detected by microscopy. Fever was significantly more common among participants recruited at health facilities or the hospital compared with those recruited at school (odds ratio [OR] = 1.54; 95% CI = 1.03–2.31, *P* = 0.04); parasite prevalence by microscopy was not significantly different (OR = 0.77; 95% CI = 0.55–1.06, *P* = 0.11). Parasite prevalence by microscopy (*P* < 0.001) and parasite density (*P* < 0.001) decreased with age (Table 1). There was no statistically significant difference in the prevalence or density of parasites between parishes after adjusting for age (*P* = 0.15). There was a strong positive association between parasite prevalence by RDT and microscopy (*P* < 0.0001). Nevertheless, 19.6% (65/331) of individuals that were parasite positive by RDT were negative by microscopy; 52.3% (34/65) of these reported antimalarial drug use in the previous 2 weeks, possibly indicating HRP-2 antigen persistence after a successfully treated infection. Samples that were positive by microscopy but negative by RDT (*N* = 50) were characterized by a low parasite density, geometric mean density 238 parasites/μL (IQR = 80–600), significantly lower than that in RDT-positive parasite carriers (*P* < 0.001).

The prevalence of infections was also determined by PCR in the parish of Abedi, where 241 DNA samples were analyzed for the presence of different malaria species. Parasites were detected by PCR in 107 samples after Chelex extraction; the QIAGEN DNA Mini Kit (QIAGEN) was used to extract DNA from 131 samples that were negative after Chelex extraction and yielded another 32 PCR-positive samples, giving parasite prevalence by PCR of 57.7% (139/241). The vast majority of infections detected by PCR were *P. falciparum* mono-infection (87.8%, 122/139), whereas 8.6% (12/139) of the infections were with *P. ovale*, either as mono-infection (*N* = 6) or as co-infection with *P. falciparum* (*N* = 5) or as co-infection with *P. falciparum* and *P. malariae* (*N* = 1). Five mixed infections with *P. falciparum* and *P. malariae* were detected (3.6% of all infections, 5/139). *Plasmodium vivax* was not detected in any of the 241 samples analyzed by PCR. Individuals with non-falciparum malaria species detected by PCR, either as mono- or mixed-infection with *P. falciparum*, were on average younger than those with *P. falciparum* mono-infection by PCR (*P* = 0.035). Overall, 35.1% (52/148) of the samples negative by microscopy were positive by PCR; four samples were positive by microscopy but negative by PCR. *Plasmodium falciparum* parasite prevalence by PCR was highest in children < 15 years of age and showed an overall negative association with age (Figure 1, *P* < 0.001). The proportion of infections that was below the microscopic threshold for detection increased from 16.4% to 19.3% in children < 15 years of age to 48.7–53.1% in older age groups (Figure 1, *P* < 0.001).

<table>
<thead>
<tr>
<th>Characteristics of enrolled individuals*</th>
<th>N</th>
<th>Abedi</th>
<th>Akere</th>
<th>Apac town</th>
<th>Atopi</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, median (IQR)</td>
<td></td>
<td>15 (5.3–25)</td>
<td>13 (4.3–29)</td>
<td>19 (5.5–32)</td>
<td>13 (5.6–30)</td>
</tr>
<tr>
<td>Fever, % (n/N)</td>
<td></td>
<td>&lt; 5 years: 30.0 (18/60)</td>
<td>27.1 (16/59)</td>
<td>46.9 (23/49)</td>
<td>14.3 (6/42)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>5–14 years: 12.9 (8/62)</td>
<td>17.7 (9/51)</td>
<td>36.6 (15/41)</td>
<td>9.1 (6/66)</td>
</tr>
<tr>
<td>Parasite prevalence by RDT, % (n/N)</td>
<td></td>
<td>&lt; 5 years: 56.7 (34/60)</td>
<td>73.3 (44/60)</td>
<td>71.4 (35/49)</td>
<td>81.0 (34/42)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>5–14 years: 59.4 (38/64)</td>
<td>44.2 (23/52)</td>
<td>51.2 (21/41)</td>
<td>64.2 (43/67)</td>
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<tr>
<td></td>
<td></td>
<td>≥ 15 years: 16.8 (21/125)</td>
<td>17.3 (18/104)</td>
<td>14.1 (17/121)</td>
<td>15.4 (14/91)</td>
</tr>
<tr>
<td><em>P. falciparum</em> parasite prevalence by microscopy, % (n/N)</td>
<td></td>
<td>&lt; 5 years: 54.4 (31/57)</td>
<td>50.9 (29/57)</td>
<td>59.6 (28/47)</td>
<td>63.4 (26/41)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>5–14 years: 59.7 (37/62)</td>
<td>42.0 (21/50)</td>
<td>51.3 (20/39)</td>
<td>65.7 (44/67)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>≥ 15 years: 20.2 (25/124)</td>
<td>22.0 (22/100)</td>
<td>11.3 (12/106)</td>
<td>23.3 (21/90)</td>
</tr>
<tr>
<td><em>P. falciparum</em> parasite density, GM (IQR)</td>
<td></td>
<td>&lt; 5 years: 6,063 (480–77,880)</td>
<td>2,844 (400–32,640)</td>
<td>8,102 (1,380–43,720)</td>
<td>2,874 (520–11,120)</td>
</tr>
<tr>
<td></td>
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<td>5–15 years: 753 (120–3,320)</td>
<td>1,235 (440–2,120)</td>
<td>1,404 (460–3,740)</td>
<td>918 (310–2,740)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>≥ 15 years: 215 (80–520)</td>
<td>453 (80–1,320)</td>
<td>232 (60–540)</td>
<td>456 (160–800)</td>
</tr>
<tr>
<td><em>P. falciparum</em> parasite prevalence by PCR, % (n/N)</td>
<td></td>
<td>&lt; 5 years: 69.4 (43/62)</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td></td>
<td></td>
<td>5–15 years: 42.6 (52/122)</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
</tbody>
</table>

* IQR = interquartile range; RDT = rapid diagnostic test; GM = geometric mean; ND = not done; PCR = polymerase chain reaction.
The occurrence of fever for different parasite densities in children < 10 years of age. The prevalence of fever (temperature ≥ 37.5°C) is given for children < 10 years of age with no parasites (N = 172), with < 400 parasites/μL (N = 50), 400–1,500 parasites/μL (N = 46), 1,500–5,000 parasites/μL, 5,000–15,000 parasites/μL (N = 39), 15,000–80,000 parasites/μL (N = 35), and ≥ 80,000 parasites/μL (N = 35). Error bars indicate the upper and lower limits of the 95% confidence interval around the proportion.

DISCUSSION

In this work, we describe the current malaria situation in Apac, northern Uganda. *Plasmodium falciparum* parasite prevalence was high, ≥ 50% by microscopy in children. ...
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8 years of age, and infections with *P. malariae* and especially *P. ovale* were common. Parasite carriage often occurred asymptptomatically, at densities ranging from submicroscopic concentrations to densities > 5,000 parasites/μL. These parameters are consistent with continuing holoendemic malaria transmission and, unsurprisingly, the age-dependent prevalence of malaria-specific antibodies did not indicate recent changes in transmission intensity.

In our surveys, *P. falciparum* parasite prevalence in children 2–9 years of age was 58%. This was slightly lower than the previously reported parasite prevalence of 70–90% in the same age group living in neighboring parishes in Apac District in 1995–1999, but within the 50–100% range reported in world malaria maps. When PCR was used for parasite detection, overall *P. falciparum* parasite prevalence was 55.2% in all age groups (37.5% by microscopy) and 69.3% in children 2–9 years of age (57.3% by microscopy). The prevalence of submicroscopic parasite carriage is in perfect agreement with the recent meta-analysis by Okell and colleagues, who reported a median PCR parasite prevalence of 58.6 (IQR = 51.4–74.0%) for areas where the microscopical parasite prevalence is 25–50%. The relative proportion of parasite carriers that harbored parasites at submicroscopic densities increased with age; this is likely to be a reflection of acquired immunity that allows adults to control infections more effectively. However, even in the youngest age group a substantial proportion of infections were not detected by microscopy, as was previously shown in areas of intense and low endemicity. Microscopy will be sufficiently sensitive to detect clinically relevant parasite densities, although our data clearly indicate that not every episode of parasitemia with fever equals malaria. One-quarter of children < 10 years of age presented with fever in the absence of parasites and the proportion of febrile children did not change considerably until malaria parasite densities exceeded 10,000 parasites/μL. Our data were insufficient to define a pyrogenic threshold parasite density or malaria-attributable fraction of fever episodes; only 77 children had a parasite density ≥ 10,000 parasites/μL. One could argue that a single blood film per individual is insufficient for calculating the pyrogenic threshold density or

![Figure 3](image-url)

**Figure 3.** Age-seroprevalence plots for circumsporozoite protein (CSP), merozoite surface protein-1 (MSP-1), and apical membrane antigen-1. Dots indicate the observed antibody prevalence for different age groups, the solid line the best fit based on age as a continuous variable, the dotted line the upper and lower limit of the 95% confidence interval (CI). The seroconversion rate λ was estimated at 0.025 (95% CI = 0.019–0.033) for CSP, 0.056 (95% CI = 0.044–0.072) for MSP-1, and 0.260 (95% CI = 0.208–0.326) for AMA-1.

### Table 2

Antibody prevalence and density in relation to parasite carriage by microscopy

<table>
<thead>
<tr>
<th></th>
<th>CSP</th>
<th>AMA-1</th>
<th>MSP-1</th>
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<tbody>
<tr>
<td></td>
<td>Prevalence, % (n/N)</td>
<td>Titer, median (IQR)</td>
<td>Prevalence, % (n/N)</td>
</tr>
<tr>
<td>1–5 years</td>
<td>Parasite-negative</td>
<td>2.5 (2/79)</td>
<td>31.1 (13.2–52.4)†</td>
</tr>
<tr>
<td></td>
<td>Parasite carrier</td>
<td>7.8 (8/103)</td>
<td>37.9 (21.4–61.5)†</td>
</tr>
<tr>
<td>5–14 years</td>
<td>Parasite-negative</td>
<td>22.7 (20/88)</td>
<td>71.2 (45.4–111.8)</td>
</tr>
<tr>
<td></td>
<td>Parasite carrier</td>
<td>21.4 (25/117)</td>
<td>60.5 (28.9–97.4)</td>
</tr>
<tr>
<td>≥ 15 years</td>
<td>Parasite-negative</td>
<td>36.4 (112/308)</td>
<td>93.0 (51.3–147.2)</td>
</tr>
<tr>
<td></td>
<td>Parasite carrier</td>
<td>34.7 (25/72)</td>
<td>79.6 (49.3–152.9)</td>
</tr>
</tbody>
</table>

*Median titer includes all individuals, also seronegatives. The prevalence or (log-transformed) titer were compared between parasite-positive and parasite-negative individuals by microscopy; these analyses were done for different age categories and adjusting for age within that category.

† P = 0.05.
‡ P < 0.01.
malaria-attributable fraction, because the density of peripheral parasitemia within a single individual can fluctuate widely between times of the day and between days.

The prevalence and density of antibody responses was influenced by microscopically detectable parasite densities. In older age groups immune responses were not influenced by concurrent parasitemia. Contrary to previous studies, we did not find evidence for a boosting of immune responses by submicroscopic parasite carriage. We observed few submicroscopic infections in children 1–5 years of age \((N = 11)\), which will have affected our power to detect such an immune-boosting effect. These findings confirm previous indications that parasitization status can be an important consideration in longitudinal assessments of the protective role of immune responses.

Age-seroprevalence plots can reveal recent reductions in transmission intensity when the age-seroprevalence curve shows an improved fit to the data when more than one SCR rate is assumed for specific time periods. We did not find any evidence for more than one force of infection and therefore have no reason to conclude a reduction in transmission intensity since the last surveys in the area.

The used methods may not have picked up a steady, gradual decline in transmission intensity, but this was not suggested by our findings in relation to previous surveys. Our approach of convenient sampling at health facilities and a school where surrounding villagers were mobilized for screening has some drawbacks. Although it was previously shown that this approach is valid in obtaining an estimate of antimalarial antibody prevalence, it may have resulted in an overestimation of parasite carriage because of a selection of symptomatic or overly exposed individuals. Although this implies that some caution is required in extrapolating the results to the general population, we feel that the symptomatic nature of the vast majority of infections makes it likely that our estimates are informative for the general population. The observation of persisting intense transmission are in agreement with a recent review that concluded that reductions in transmission intensity are not evident in all African settings and that transmission intensity may have remained unchanged or even increased in northwestern Uganda and neighboring countries in East and Central Africa. The failure to reduce the burden of malaria could reflect sub-optimal implementation of malaria control measures. Even in areas of intense malaria transmission intensity, considerable gains can be achieved by vector control and effective antimalarial treatment, as was illustrated by successful malaria control on Bioko island.

Malaria control efforts in Apac were not reliably monitored in the last decade and affected by political unrest in preceding years. The ACTs were officially available in Apac from the year 2006, but especially the smaller health facilities were affected by supply shortages that affected their implementation; recently, malaria initiatives in Apac have been intensified. Indoor residual spraying (IRS) with DDT was banned in Apac in 2008 by a court injunction launched by organic farmers, but IRS with pyrethroids (Fendona, BASF, Midrand, South Africa) started again in May 2010 (i.e., after the current survey was completed). Community-wide distribution of long lasting ITNs took place in early 2009, reaching the majority of households. Our study illustrates that these control efforts are still profoundly needed in areas in Africa where transmission remains intense and malaria control continues to be a tremendous challenge.

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