Laboratory and field evaluation of MAÏA®, an ointment containing \(N,N\)-diethyl-3-methylbenzamide (DEET) against mosquitoes in Burkina Faso

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Abstract

Background: Malaria vector control relies upon the use of insecticide-treated nets and indoor residual spraying. However, as the emergency of insecticide resistance in malaria vectors grows, the effectiveness of these measures could be limited. Alternative tools are needed. In this context, repellents can play an important role against exophilic mosquitoes. This study evaluated the efficacy of MAÏA®, a novel repellent ointment, in laboratory and field conditions in Burkina Faso.

Methods: For laboratory and field assessment, 20 volunteers were enrolled and trained for nocturnal collection of mosquitoes using human landing catches (HLC). In the laboratory tests, 2 mg/sq cm of treatment (either MAÂ® or 20 % DEET) were used to assess median complete protection time (CPT) against two species: Anopheles gambiae and Aedes aegypti, following WHO guidelines. For both species, two strains consisting of susceptible and local strains were used. The susceptible strains were Kisumu and Bora Bora for An. gambiae and Ae. aegypti, respectively. For the field test, the median CPT of MAÏA® was compared to that of a negative (70 % ethanol) and positive (20 % DEET) after carrying out HLCs in rural Burkina Faso in both indoor and outdoor settings.

Results: Laboratory tests showed median Kaplan-Meier CPT of 6 h 30 min for An. gambiae (Kisumu), 5 h 30 min for An. gambiae (Goden, local strain), and 4 h for Ae. aegypti for both the local and sensitive strain. These laboratory results suggest that MAÏA® is a good repellent against the three mosquito species. During these field tests, a total of 3979 mosquitoes were caught. In this population, anophelines represented 98.5 %, with culicines (Aedes) making up the remaining 1.5 %. Among anopheline mosquitoes, 95 % belonged to the An. gambiae complex, followed by Anopheles funestus and Anopheles pharoensis. The median CPT of 20 % DEET and MAÏA® were similar (8 h) and much longer than that of the negative control (2 h).

Conclusions: Results from the present studies showed that MAÏA® offers high protection against anophelines biting indoors and outdoors and could play an important role in malaria prevention in Africa.

Keywords: Malaria, Mosquito, Anopheles gambiae, Aedes aegypti, Repellent, MAÏA®, Burkina Faso

Background

Malaria is one of the deadliest diseases in many low- and middle-income countries, affecting mainly children and pregnant women in sub-Saharan Africa [1]. Long-lasting
insecticide-treated nets (LLINs) have been regarded as the most effective method for controlling mosquitoes transmitting malaria parasites. Since 2000, about one billion nets have been distributed in Africa, resulting in a significant decline in malaria-related deaths on the continent between 2000 and 2015 [2–4].

However, the massive use of insecticides in public health in addition to that in agriculture causes concern regarding insecticide resistance [5–7] and changing behaviour [8, 9] of the malaria vectors. For example, a study conducted in Papua New Guinea showed a shift in mosquito biting from night to earlier hours in the evening after a nationwide distribution of LLINs [10]. Similar changes in the behaviour of Anopheles funestus have been observed in Benin and Senegal after LLIN distribution achieved a high level of coverage [9, 10]. Furthermore, studies suggest that the scaling up of LLIN distribution and indoor residual spraying (IRS) have led to more outdoor biting by Anopheles gambiae sensu lato (s.l.), commonly considered endophagic mosquitoes [11–13]. A recent study in the Cascades region of Burkina Faso showed a high level of insecticide resistance [14] where more than 50% of the major vector, An. gambiae s.l., were collected biting outdoors [15]. These altered patterns of outdoors, early evening and morning biting, by anophelines, combined with resistance to insecticides appear to be caused by the mass distribution of LLINs and imply the inexorable loss of efficacy of these interventions [16, 17]. A recent study highlighted that an increase in early evening biting could increase transmission not only because people are unprotected by nets, but also because there is a higher chance of malaria vectors becoming infectious [18]. The development of new vector control tools, in addition to LLINs, is therefore necessary to protect people, when they are not under a bed net.

Topical repellents could play an important role in addressing this problem if they are effective and accepted by the population. A systematic review of repellent interventions and mathematical modeling has shown that ‘user compliance’ is indeed one of the most decisive factors for the success of this intervention [19]. In sub-Saharan Africa, ointments are used primarily by mothers and children to moisturize their skins. In Burkina Faso, ointments are applied to 80% of children every evening, when mosquitoes start biting (Kadidia Ouedraogo et al., in prep). Maïa Africa, a company based in Burkina Faso, has developed a mosquito repellent ointment, MAÏA®, uniquely designed with local mothers, to be used daily within their families. The underlying idea is to leverage the existing habits of mothers to protect their families from infectious bites whenever they are not under a net. Affordability is a key criterion for the product’s adoption and use. MAÏA® has been industrially produced since June 2020 in Côte d’Ivoire and integrates a large share of ingredients sourced in West Africa. The ointment was officially launched in August 2020 in Burkina Faso and over 50,000 units were sold in the first four months. In March 2021, over 500 points of sales (mainly general stores, pharmacies and kiosks) distributed the product in the country. If MAÏA® proves that it is both effective and accepted by the population, it could play a key role in reducing the probability of children experiencing infectious bites during the evening and be positioned as a complementary intervention to LLINs.

The aim of this study is to evaluate the effectiveness of MAÏA® in both laboratory and field conditions, especially the median complete protection time (CPT) offered by the product. Results from these evaluations are important for validating how effective this new repellent is; behavioral responses to repellent differ between wild mosquito and laboratory-reared mosquito populations [20].

Methods

Study area

Laboratory tests were conducted in May 2019 in the insectary of Centre National de Recherche et de Formation sur le Paludisme (CNRFP) in Ouagadougou, Burkina Faso. Field tests were carried out at Goden (12°25'N, 1°20'W), a site located at 15 km northeast of Ouagadougou, the capital city of Burkina Faso (Fig. 1). Goden is a rural village with a Sudanian savanna climate and rainfall under 900 mm annually. The ~800 inhabitants mainly belong to the Mossi ethnic group, and are mostly devoted to agriculture and raising pigs, dogs, goats, and chickens within their compounds. LLINs were distributed in 2016 to ~90% of the population. Goden is known for its high density of malaria vectors due to its proximity to the Massili River. The field study was carried out during the rainy season (August to November 2019) corresponding to high vector density and high malaria transmission. A preliminary assessment of the mosquito density on the collection site was carried out using human landing catches (HLCs) before the tests started.

Human volunteer preparation

Healthy adult male volunteers aged between 18 and 40 years were enrolled in this study. The volunteers were instructed not to use fragranced soaps, perfume, tobacco, or alcohol 12 h before the start and throughout their participation. To establish the amount of repellent required for application, the surface area of the arm (for laboratory tests) or the leg (for field tests) of volunteers was determined using the following formula: Area = ½ (C_w + C_e) D_we.
Here $C_w$ is the circumference of the wrist or ankle in cm, $C_e$ is the elbow cubital fossa or the knee circumference in cm, and $D_{we}$ is the distance in cm between $C_e$ and $C_w$ [21]. The amount of ointment needed for each volunteer was determined depending on the area of their fore-arm or leg length. The quantity of product left in bottles was weighed using a precision weighing balance (KERN & SOHN GmbH, Balingen, Germany) to determine the amount applied by each volunteer.

**Repellents**

MAIA®, a shea butter-based ointment containing 15% DEET ($N, N$-diethyl-3-methylbenzamide) was received from Maia Africa SAS. It was tested against an ethanolic solution of 20% DEET as positive control and a negative control of 70% ethanol. DEET is known as the standard repellent reference.

**Strains of mosquitoes**

Four strains of mosquitoes were used in the laboratory tests, including Kisumu F57 and Bora bora F58 susceptible strains of respectively *An. gambiae* and *Aedes aegypti*. In addition, local strains laboratory-colonized from rural areas of Goden, in Burkina Faso, were used: hereafter named An-Goden (*An. gambiae* local strain, F418) and Loc-Aedes (local *Ae. aegypti*, F318). These species were maintained under a 12:12 h (light: dark) photoperiod. During rearing, larvae were fed on fish food while glucose was used for adults. The temperature and relative humidity in the rearing room were 25–28 °C and 60–80%, respectively. Individual mosquitoes used in these experiments were 5 to 10 days old nulliparous females starved from sugar solution for 12 h before the experiment.

**Evaluation in the laboratory**

The laboratory experiments were conducted following WHO guidelines for the arm-in-cage test [21]. Cages were $45 \times 45 \times 45$ cm screen enclosures. Two test cages were used, one for the repellent candidate and the other for the positive control. The test cages contained 200 females aged 5 to 10 days of one of the four mosquito strains.
strains: Kisumu, An-Goden, Bora bora and loc-Aedes. The experiment in the laboratory was carried out at temperatures ranging between 25 and 28 °C, with relative humidity between 60 and 80%.

Overall, 2 mg of ointment were applied per sq cm on the left forearm of each volunteer, a concentration estimated from the average of 5 volunteers from the laboratory asked to apply ointment on their left forearm as they would normally do in real life. In all subsequent repellent trials, volunteers were then supplied with a total volume that would achieve this concentration over the surface area of their forearms and/or legs (determined as described above). A steel spatula was used to apply the ointment on the forearm of each volunteer prior to each experiment to avoid absorbing ointment on non-targeted areas of skin. Positive control consisted of 1 ml of 20% DEET solution applied to the right forearms of volunteers.

Negative controls and MAIA™ test arms were prepared by first washing left forearms with odorless soap, drying and rinsing with 70% ethanol solution and then drying again. All volunteers wore latex gloves to protect their hands from mosquitoes. To assess the readiness of the mosquitoes to land, both left and right cleaned forearms of volunteers were exposed in the experimental cages for 30 s (or until 10 landings of mosquito were counted). Then, for each volunteer, the right forearm was treated from wrist to elbow using 1 ml of the 20% DEET solution whilst the left forearm was treated from the wrist to elbow with MAIA™ ointment. Thirty mins after application of the repellents, the volunteer exposed their treated forearm in the test cage for 3 min. The procedure was repeated every 30 min until the first bite occurred and the elapsed time to the first bite was recorded. The test was performed three times for each volunteer per mosquito species. Considering the difference in the relative periods of biting activity of each mosquito species, the tests using *Ae. aegypti* strains were carried out between 09:00 and 18:00, whereas those for *An. gambiae* were conducted between 17:00 and 05:00 [21].

**Field evaluation**

The lower legs of volunteers were washed with neutral soap, rinsed with 70% of ethanol solution and naturally dried. Once their legs were treated volunteers were asked to avoid rubbing, touching or wetting the repellent-treated area. Two mg of MAIA™ per sq cm (2.4 ± 0.2 g per 1189 ± 79.2 sq cm) and 2 ml per sq cm of 20% DEET (2 ml ± 0.1 ml per 1189 ± 79.2 sq cm, as a positive control) were applied to volunteers’ lower legs, from knee to ankle. A total of 20 volunteers were recruited from Goden village and trained for nocturnal mosquito collection using HLC. Each volunteer was later randomly allocated to one of the five groups (2 for MAIA™, 2 for positive control, 1 for negative control) of four volunteers according to the repellent received. Each night of collection, the experiment took place at five different households, located at least 20 m apart as per WHO guidelines [21], in order to avoid biases in attractiveness to the mosquitoes.

Mosquito collection started 30 min following treatments. Volunteers acting as bait, sat on a chair in pairs (one indoor and one outdoor) and actively collected mosquitoes that landed on their treated lower leg using mouth aspirator and flash torch [22] for 45 min, followed by a 15-mins break. Volunteers wore long-sleeved shirts, buttoned at the wrist, long trousers, closed shoes and latex gloves with a hat on their head, but with the treated lower leg to be exposed to mosquitoes by rolling up trousers to the knee. During these experiments, mosquitoes were collected simultaneously indoors and outdoors between 19:00 and 06:00. To avoid biases introduced by individual attractiveness and skills [23, 24] volunteers at the same household rotated between indoors and outdoors hourly. In each household two groups of two people rotated collecting from 18:00 to 24:00 and from 00:00 to 06:00, following the Williams balanced Latin Square design.

Collected mosquitoes were transferred into plastic cups, covered with a piece of untreated net, with a small hole at the bottom to allow mosquitoes to be easily aspirated into them. After collection mosquitoes were brought to the entomological laboratory of CNRFP and morphologically identified using a stereo microscope and identification keys [25].

**Side effects**

No side effects were observed or reported by any of the volunteers throughout the period of tests both in the laboratory and in the field.

**Ethical clearance**

Written informed consent was obtained from all volunteers and household owners recruited in this study. The study was approved by the institutional ethic committee of CNRFP under 2019/000008/MS/SG/CNRFP/CIB.

**Data analysis**

All data were collected on standard forms and entered twice in a database by different people. Databases have been compared using Epi Info™ 3.5.3, and inconsistencies were verified using printed and corrected forms. The performance of the repellent was measured by calculating the repulsive efficiency and the median full protection time. A generalized linear mixed model (GLMM) was used to further analyse the effect of the location (indoors
vs. outdoors) on the performance of the treatments. Variation in the average number of bites received between treatments was also assessed.

The median CPT is defined as the interval of time between the beginning of collection/test and the first mosquito landing. To estimate the median CPT of each treatment, a Kaplan-Meier survival analysis was performed for each vector species and strain used in the laboratory experiments and on field data through ‘survival function’ from R software-version 3.5.0 (2018-04-23). However, for the field test, the analysis was performed on only An. gambiae s.l. as it was the most abundant species collected (~96% of the total collection). The analysis consisted of assessing the median CPT and the repulsive efficacy. The repulsive efficacy was calculated as a percentage of repulsion (% R) according to the formula 

\[ \% R = \left( \frac{C - T}{C} \right) \times 100 \]

where C is the number of mosquitoes collected on the treated legs of the two control treatments separately, and T is the total number of mosquito bite attempts on the volunteers’ legs treated with the test product [21].

**Results**

**Laboratory tests**

Overall, under laboratory conditions the relative repellency (median CPT) was higher for both MAÏA® and 20 % DEET against An. gambiae compared to Ae. aegypti (Table 1). MAÏA® performed well in repelling the four mosquito species used in this study. The median CPTs were, respectively, 6.5 h for An. gambiae (Kisumu), 5.5 h for An. gambiae (Goden, local strain) and 4 h for Ae. aegypti for both the local and sensitive strain. There was no significant difference between the two treatments for each of the experiment (Kisumu: \( \chi^2 = 2.1, p = 0.14 \); Goden: \( \chi^2 = 0.8, p = 0.36 \); Bora bora: \( \chi^2 = 1.7, p = 0.19 \); Ae. aegypti (local strain): \( \chi^2 = 0.9, p = 0.35 \) ) indicating that both MAÏA® and 20 % DEET have equal repellency for these strains. The Kaplan-Meier curves for MAÏA® and 20 % DEET, respectively, for Kisumu, An-Goden, Bora bora and Loc-Aedes are shown in Fig. 2.

**Repellency against mosquitoes**

Repellency against An. gambiae s.l. was stratified by time of collection. From 18:00 to 24:00 (6 h after application), the percentage of repellency varied from 100 to 90% for MAÏA® and DEET. Between 00:00 to 03:00 (9 h after application), the percentage was between 90 and 80% (Fig. 3). After 03:00 (10 h after application), this percentage was under 80% for 20 % DEET, but MAÏA® gave a high percentage of repellency throughout, however during the first 9 h after applications no difference in the repellency was observed between MAÏA® and the positive control.

When data were stratified by location of mosquito biting, the trend was the same for indoors and outdoors. No difference was observed during the first 9 h between MAÏA® and 20 % DEET. These results show that MAÏA® can protect both indoors and outdoors.

**Table 1**

| Mosquito Species | Median CPT (min) | Lower CI | Upper CI |
|------------------|-----------------|----------|----------|
| **An. gambiae**  | **Kisumu**      | **Goden** | **Bora bora** | **Local** |
| DEET             | 390             | 300      | 270      | 288      | 240     | 268     |
| MAÏA®            | 390             | 330      | 216      | 244      | 240     | 255     |

A total of 3,979 mosquitoes, stratified by treatment and species (Table 2), were caught using HLC. Anopheleslines represented 98.5 % of the total catch, with culicines (Aedes) making up the remaining 1.5 %. Among anopheline species, 99.6 % belonged to An. gambiae complex, followed by An. funestus (0.1 %), and Anopheles phar-oesis (0.3 %). The frequency of mosquitoes landing on treated collectors, compared with control subjects, varied according to the repellent used (Table 2). The hourly mosquito biting rate varied significantly between treatments (df = 2, \( \chi^2 = 426.22, p < 0.0001 \)). An average of 0.68 (95 % CI: 0.51–0.91) mosquito bites were received per person per hour for MAÏA® compared to 1.01 (95 % CI: 0.76–1.33) for 20 % DEET, and 8.98 (95 % CI: 6.56–12.29) for the 70 % ethanol. In addition, there was no variation between treatments according to location (indoors and outdoors, df = 2, \( \chi^2 = 1.703, p = 0.42 \)). Overall, the ratio outdoors:indoors biting was 1.26 (95 % CI: 1.25–1.27) showing that more bites were taking place outdoors compared to indoors (df = 1, \( \chi^2 = 5.79, p = 0.016 \)).

**Field test**

**Mosquito species composition and biting behaviours**

The vector species used in this study were Anopheles gambiae and Ae. aegypti. The determination of the species was performed using the molecular dataset from the strain-specific primers. The analysis of the data was performed using the R software version 3.5.0 (2018-04-23) and the survival function from the survival package. The analysis was performed on only An. gambiae s.l. as it was the most abundant species collected (~96% of the total collection). The analysis consisted of assessing the median CPT and the repellence efficacy. The repellence efficacy was calculated as a percentage of repulsion (% R) according to the formula

\[ \% R = \left( \frac{C - T}{C} \right) \times 100 \]

where C is the number of mosquitoes collected on the treated legs of the two control treatments separately, and T is the total number of mosquito bite attempts on the volunteers' legs treated with the test product [21].

Overall, under laboratory conditions the relative repellency (median CPT) was higher for both MAÏA® and 20 % DEET against An. gambiae compared to Ae. aegypti (Table 1). MAÏA® performed well in repelling the four mosquito species used in this study. The median CPTs were, respectively, 6.5 h for An. gambiae (Kisumu), 5.5 h for An. gambiae (Goden, local strain) and 4 h for Ae. aegypti for both the local and sensitive strain. There was no significant difference between the two treatments for each of the experiment (Kisumu: \( \chi^2 = 2.1, p = 0.14 \); Goden: \( \chi^2 = 0.8, p = 0.36 \); Bora bora: \( \chi^2 = 1.7, p = 0.19 \); Ae. aegypti (local strain): \( \chi^2 = 0.9, p = 0.35 \) ) indicating that both MAÏA® and 20 % DEET have equal repellency for these strains. The Kaplan-Meier curves for MAÏA® and 20 % DEET, respectively, for Kisumu, An-Goden, Bora bora and Loc-Aedes are shown in Fig. 2.
Complete protection time

The overall median CPTs of 20% DEET and MAÏA® (Table 3) were estimated at 480 min (8 h) against 120 min (2 h) for the negative control. For outdoor collections, the median CPTs were 480, 450 and 120 min for 20% DEET, MAÏA® and negative control respectively (Table 4). For indoor collections, these estimates were 480, 480 and 60 min, respectively, for 20% DEET, MAÏA® and ethanol (Table 4). Statistical analyses showed that there was no difference in the median CPT between 20% DEET and MAÏA® (df = 1, χ² = 0.2, p = 0.7). However, there was a significant difference between median CPT as estimated for MAÏA® and the negative control (df = 2, χ² = 106, p < 0.0001, Fig. 4). Even when the collection was stratified by location this difference still occurred in both indoor (Fig. 5A; df = 2, χ² = 41.6, p < 0.0001) and outdoor collections; df = 2, χ² = 66.7, p < 0.0001 (Fig. 5B).

Discussion

The results of this study demonstrated that MAÏA®, a shea butter-based ointment with 15% DEET provides high protection against mosquitoes in Goden, a rural area of Burkina Faso. Both tests under field and laboratory conditions suggested that MAÏA® has equal repellency effect as 20% DEET ethanolic solution during the period of collection. Similar results were also found both indoors and outdoors. The percentage of repellency when using MAÏA® varied between 100 and 80% over the major malaria vector biting period, which occurs between 18:00 and 06:00. The median CPT were also similar and estimated around 480 min. Both MAÏA® and the 20% DEET ethanolic solution were found to provide up to 90% of repellency during the first 6 h after their applications. Furthermore, results suggested that using MAÏA®, the average hourly bites received was significantly lower
(less than 1 bite per hour) compared to that of both 20% DEET and negative control. However, one of the limitations is that biting activity on control arms was not checked at the end of laboratory experiments to validate consistency of mosquito avidity, but this did not affect the overall result. Overall, it can be argued that MAÏA® could provide protection to people before they go to bed.

Previous studies in the same locality comparing repellency between three different repellents found that DEET, IR3535 and KBR 3023 were effective against An. gambiae s.l. and other Afrotropical vector mosquitoes [26]. In this study authors showed that protection from KBR 3023, DEET and IR3535 were still high against anophelines for up to 10 h post-exposure. In contrast, results from the current study indicated that the relative repellency was 100% for ~8 h. Results were similar to that from a recent study in Ethiopia comparing DEET \((N, N\text{-diethyl-1,3-methylbenzamide})\) and MyggA \((p\text{-methane diol})\) and other laboratory products (20% neem oil and 20% chinaberry oil), where the mean CPT was 8 h for DEET whilst an estimated 6 h for MyggA [27]. Eight hours of repellency may suffice to protect against earlier vector biting both indoors and outdoors before residents take protection from insecticide-treated nets deployed indoors.

To date, 11 countries across the world are classified as having a high burden of malaria [2]. In these countries malaria vector control is still based on the use of insecticides, either in the form of indoor spraying or by promoting the large-scale distribution of LLINs. These strategies can effectively reduce the number of malaria cases [28], however the major challenge is the resistance of malaria vectors to different classes of insecticides and

### Table 3

| Anopheles gambiae s.l. | DEET | Ethanol | MAÏA® |
|------------------------|------|--------|-------|
| Median CPT             | 480  | 120    | 480   |
| Lower CI               | 454  | 91     | 448   |
| Upper CI               | 506  | 149    | 512   |

### Table 4

| Anopheles gambiae s.l. | Indoors | Outdoors |
|------------------------|---------|----------|
| 20% DEET               | Ethanol | MAÏA®    | 20% DEET | Ethanol | MAÏA®    |
| Median CPT             | 480     | 60       | 480     | 120     | 450      |
| Lower CI               | 440     | < 60     | 440     | 95      | 428      |
| Upper CI               | 521     | NA       | 521     | 145     | 472      |
the shifts in their feeding and resting behaviours, with the tendency of biting and resting outdoors. For example, a study in the Cascades region in Burkina Faso indicates that in addition to insecticide resistance, more than 50% of the malaria vector biting were taking place outdoors. Therefore, new and supplementary methods are urgently needed to complement these tools in the perspective of malaria elimination [29]. In accordance with the spirit of locally adapted-integrated vector and disease control [30], repellents can usefully complement existing control strategies and provide an additional tool in the management of insecticide resistance. In the context of widespread resistant vectors to insecticide and the tendency of mosquitoes to bite outside houses, there is a need to add MAÏA® ointment to vector control tools in sub-Saharan, malaria-burdened countries.

The originality of MAÏA® comes from its formulation based on local butter extensively used in rural areas of West African, and which contributes to women's economic income. Promotion of use of local endogenous strategies can sustain malaria control, and also improve the economic situation for African women. Additionally, it has been shown that shea butter is a source of anti-inflammatory and anti-tumour promoting compounds [31]. Another interesting compound in shea butter is cinnamic acid which is known for its antibacterial, antifungal and antiviral properties [32]. Shea butter both moisturizes and heals the skin. Clinical studies have shown it to be safe for skin [33]. MAÏA® ointment will not only protect from mosquito-borne diseases but also against other micro-organisms.

Besides the level of protection offered by repellents, daily compliance and appropriate use seem to be major obstacles to achieving the potential impact on malaria [34]. An efficacy study carried out in Tanzania has shown that volunteers preferred MAÏA® ointment to a more classical 20% DEET solution [35]. Sales by Maïa Africa SAS of 50,000 units in local stores in Burkina Faso between August and November 2021 further illustrate its desirability. However, more data are needed to understand who is likely to use the product and whether its usage is appropriate in terms of frequency and application in order to have an impact on mosquito-borne diseases, such as malaria.

**Conclusions**

MAÏA®, a novel ointment formulated with shea butter widely used in West Africa to moisturize the skin of children, has shown high repellency against laboratory-reared and wild malaria vectors. In the context of widespread vector resistance to insecticide and growing tendency of mosquitoes to bite outside houses, there is a
need to add MAID® ointment to the vector control tools used in sub-Saharan countries with high malaria burden.

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Authors’ contributions
GWM, NFS and AS designed the study, AT and GN conducted the laboratory and field data collection and drafted the manuscript. AS and GWM performed the data analysis, interpreted the results and participated in manuscript writing. NFS and AG contributed to the design of the study and provided comments upon the manuscript. FL provided comments upon the manuscript. FL and GN sought the funding to support all laboratory and field experiments. All authors read and approved the final manuscript.

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Availability of data and materials
Not applicable.

Declarations

Ethics approval and consent to participate
The study was approved by the institutional ethic committee of CNRP under 2019/000008/MS/SG/CNRP/CIB. All volunteers involved in this study were recruited upon a written informed consent form and they were informed that they are free to leave the study at any time without any repercussion.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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