



Contents lists available at ScienceDirect

Asian Pacific Journal of Tropical Biomedicine

journal homepage: www.elsevier.com/locate/apjtb



Document heading doi:10.1016/S2221-1691(14)60238-0 © 2014 by the Asian Pacific Journal of Tropical Biomedicine. All rights reserved.

Evidence of increasing L1014F *kdr* mutation frequency in *Anopheles gambiae* s.l. pyrethroid resistant following a nationwide distribution of LLINs by the Beninese National Malaria Control Programme

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PEER REVIEW

Peer reviewer

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Comments

The authors have clearly shown a widespread *kdr* frequency in the transect south–north Benin which have increased even after the distribution of LLINs. The effect of selection pressure after a large scale distribution of LLINs has confirmed previous reports by Czeher *et al.* (2008) on an increase Leu–Phe *kdr* mutation in Niger following a countrywide LLINs implementation. The study is important in its field. Details on Page 242

ABSTRACT

Objective: To determine the susceptibility status to pyrethroid in *Anopheles gambiae* s.l. (*An. gambiae*), the distribution of *kdr* “Leu–Phe” mutation in malaria vectors in Benin and to compare the current frequency of *kdr* “Leu–Phe” mutation to the previous frequency after long–lasting insecticide treated nets implementation.

Methods: Larvae and pupae of *An. gambiae* s.l. mosquitoes were collected from the breeding sites in Littoral, Zou, Borgou and Alibori provinces. CDC susceptibility tests were conducted on unfed females mosquitoes aged 2–5 d old. *An. gambiae* mosquitoes were identified to species using PCR techniques. Molecular assays were also carried out to identify *kdr* mutations in individual mosquitoes.

Results: The results showed that *An. gambiae* Malanville and Suru–lere populations were resistant to deltamethrin. Regarding *An. gambiae* Parakou and Bohicon populations, they were resistant to permethrin. PCR revealed 100% of mosquitoes tested were *An. gambiae* s.s. The L1014F *kdr* mutation was found in *An. gambiae* s.s. Malanville and Parakou at various allelic frequencies. The increase of *kdr* allelic frequency was positively correlated with CDC bioassays data.

Conclusions: Pyrethroid resistance is widespread in malaria vector in Benin and *kdr* mutation is the main resistance mechanism involved. More attention may be paid for the future success of malaria control programmes based on LLINs with pyrethroids in the country.

KEY WORDS

Resistance, Insecticide, Vectors, CDC bioassay, Leu–Phe *kdr* mutation

1. Introduction

There were an estimated 216 million episodes of malaria in 2010, with 149 million to 274 million cases. Approximately 81%, or 174 million cases, were in the African Region, with the Southeast Asian Region accounting for another 13%[1]. There were also an estimated 655 000 malaria deaths in 2010, of which 91% were in the African Region. Approximately 86% of malaria deaths globally were of children under 5 years of

age[1]. So, malaria remains one of the most critical public health challenges for Africa despite intense national and international efforts[2].

The intensive use of insecticide in the malaria control activities means that widespread mosquito insecticide resistance could have a devastating effect on the planned upscaling of the vector control activities[3].

In many African countries, malaria mosquitoes were already resistant to pyrethroids. For instance, pyrethroid

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Article history:

Received 12 Dec 2013

Received in revised form 20 Dec, 2nd revised form 25 Dec, 3rd revised form 30 Dec 2013

Accepted 20 Jan 2014

Available online 28 Mar 2014

resistance in *Anopheles gambiae* s.s. (*An. gambiae*) has already been described in Benin[4–6].

Malaria vector resistance to insecticides in Benin is conferred by two main mechanisms: (1) alterations at site of action in the sodium channel, *viz.* the *kdr* mutations and (2) an increase of detoxification and/or metabolism through high levels of multi-function oxidases, non-specific esterases and glutathione S-transferases[4–7].

The Beninese National Malaria Control Programme has been implemented. Large-scale and free long-lasting insecticidal nets (LLINs) (OlysetNet) were distributed since July 2011 throughout the country to increase coverage of LLINs. Information on susceptibility to pyrethroid insecticides used in public health in Benin and the underlying mechanisms being investigated are crucial. This will properly inform control programs of the most suitable insecticides to use and facilitate the design of appropriate resistance management strategies.

The aim of this study is to determine the susceptibility status, pyrethroid resistance levels in *An. gambiae* Malanville, Parakou, Bohicon and Suru-lere populations to permethrin and deltamethrin using CDC (Center for Disease Control and Prevention) bottle bioassays, to evaluate the presence and extent of the distribution of the *kdr* mutation within and among these *An. gambiae* s.l. populations and to compare the current frequency of *kdr* “Leu-phe” mutation to the previous frequency in malaria vectors in the south-north transect Benin after LLINs implementation.

2. Materials and methods

2.1. Study area

The study was carried out in some localities, following a south–north transect. Four contrasting localities of Benin were selected for mosquito collection on the basis of variation in agricultural production, use of insecticides and/or ecological settings (Figure 1). The localities are as follows. Bohicon is located in the middle part of the country where the farmers used significant amounts of pyrethroids and organophosphates for cotton protection or to control agricultural pests. Parakou, an urban vegetable growing area located in the north of Benin. Malanville is a rice growing area located near the Niger River. Suru-lere is an urban locality in Cotonou district located in southern Benin.

The choice of the study sites took into account the economic activities of populations, their usual protection practices against mosquito bites, and peasant practices to control farming pests. The southern zone (Cotonou) is characterized by a tropical coastal Guinean climate with two rainy seasons (April–July and September–November). The mean annual rainfall is over 1 500 mm. The middle part of the country (Bohicon) is characterized by a Sudano–Guinean

climate with an average rainfall of 1 000 mm per year. The northern zone (Parakou and Malanville) is characterized by a Sudanian climate with only one rainy season per year (May to October) and one dry season (November–April). The temperature ranges from 22 to 33 °C with the mean annual rainfall of 1 300 mm.

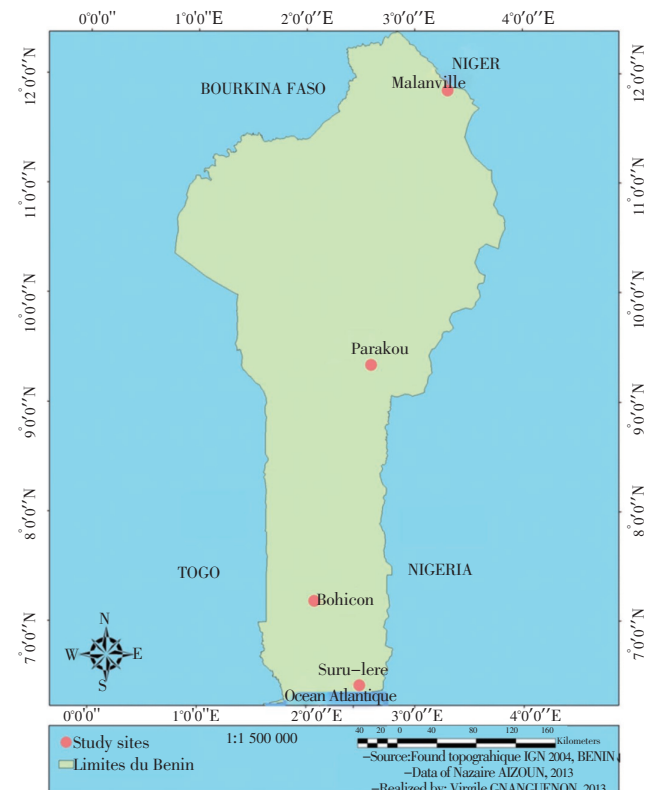


Figure 1. Map of the study area.

2.2. Sample collection

From March 2012 to November 2012, larvae and pupae of *An. gambiae* mosquito were collected several times per year, *i.e.* at the beginning and the end of rainy season from a wide range of breeding sites (puddles, shallow wells, gutters and rice fields) in Malanville, Parakou, Bohicon districts and in Suru-lere locality of Cotonou district. All larvae were brought back to laboratory of Centre de Recherche Entomologique de Cotonou for rearing. Emerging adult female mosquitoes were used for insecticide susceptibility tests. A susceptible strain of *An. gambiae* Kisumu was used as reference strain for bioassays.

2.3. Insecticide susceptibility tests

Females *An. gambiae* aged 2 to 5 d old were exposed to CDC diagnostic dosage of various insecticides according to the CDC protocol[8]. The following insecticides were tested: permethrin (21.5 µg per bottle) and deltamethrin (12.5 µg per bottle). Mosquitoes were exposed for 2 h to insecticide-treated bottles and monitored at different time intervals (15, 30, 35, 40, 45, 60, 75, 90, 105 and 120 min). This allowed us to

determine the total percent mortality (Y axis) against time (X axis) for all replicates using a linear scale. All susceptibility tests were conducted in the Centre de Recherche Entomologique de Cotonou laboratory at (25±2) °C and 70 to 80% relative humidity.

Dead and surviving mosquitoes were separately stored in individual tubes with silicagel and preserved at –20 °C in the laboratory for further molecular characterization. The choice of permethrin was justified by the insecticide used on OlysetNets that were distributed free by the national malaria control program in July 2011 across all the country whereas we used deltamethrin, an insecticide of same class as permethrin to assess its cross–resistance with this product.

2.4. PCR detection of species and the *kdr* mutation

At the end of CDC bioassays, PCR tests for species identification was performed to identify the members of *An. gambiae* complex collected from each site[9]. PCR for the detection of the *kdr* “Leu–phe” mutation was carried out on dead and alive *An. gambiae* mosquitoes as described by Martinez–Torres *et al*[10].

2.5. Statistical analysis

The resistance status of mosquito samples was determined according to the CDC criteria[8,11]. The susceptibility thresholds at the diagnostic time of 30 min for pyrethroids are shown below:

Mortality rate=100%: the population is fully susceptible.

Mortality rate<100%: the population is considered resistant to the tested insecticides.

Molecular results (*kdr* frequencies) were correlated with the results of insecticide susceptibility tests performed with CDC method from each of the districts surveyed.

ANOVA test was performed with mortality rate as the dependent variable and the localities as a covariate. ANOVA test was also performed with *kdr* frequency as the dependent variable and the localities as a covariate.

3. Results

3.1. Susceptibility of *An. gambiae* s.l. populations to permethrin and deltamethrin

Table 1 shows that Kisumu strain (control) confirmed its susceptibility status as a reference strain. All females mosquitoes of *An. gambiae* Kisumu which were exposed to CDC bottles treated with permethrin 21.5 µg/bottle and deltamethrin 12.5 µg/bottle were dead and none of them can fly after 30 min, which represented susceptibility threshold time or diagnostic time clearly defined by CDC

protocol. That confirmed this strain was fully susceptible to these products. A non neglected proportion of *An. gambiae* Malanville and Suru–lere populations, 100% and 13.93% respectively, after 30 min exposure to CDC bottles treated with deltamethrin, continued to fly in these bottles. That showed these populations were resistant to this product. *An. gambiae* Parakou and Bohicon populations after 30 min exposure to CDC bottles treated with permethrin 7.5% and 12.5% respectively, continued again to fly in these bottles. That also showed these populations were resistant to this product (Table 1).

Table 1

Mortality of *An. gambiae* from the districts of Malanville, Parakou, Bohicon and Suru–lere locality of Cotonou district after 2 h exposure to permethrin (21.5 µg/bottle) and deltamethrin (12.5 µg/bottle).

Bio-climatic areas	Localities	Insecticides	Number tested	% Mortality	Resistance status
	Kisumu	Permethrin	37	100.00	S
		Deltamethrin	25	100.00	S
Sudanian	Malanville	Deltamethrin	21	0.00	R
	Parakou	Permethrin	40	92.50	R
Sudano–guinean	Bohicon	Permethrin	64	87.50	R
Guinean	Suru–lere	Deltamethrin	79	86.07	R

S: Sensitive, R: Resistant.

Univariate logistic regression performed with mortality rate as the dependent variable and localities as a covariate with ANOVA test showed that the phenotypic resistance to permethrin and deltamethrin was associated with the localities (P<0.05) on the one hand. Univariate logistic regression performed with *kdr* frequency as the dependent variable and localities as a covariate with ANOVA test showed that high *kdr* frequency was associated with the localities (P<0.05) on the other hand.

3.2. Species of *An. gambiae*

Mosquitoes from CDC bioassay were analysed by PCR for identification of sibling species among *An. gambiae* s.l. complex. PCR revealed 100% of mosquitoes tested were *An. gambiae* s.s. (Table 2).

Table 2

Species identification and *kdr* frequency in *An. gambiae* s.l. from CDC bioassays.

Localities	Number tested	Species Ag	<i>Kdr</i> mutation			
			RR	RS	SS	F(<i>kdr</i>)
Malanville	48	48	38	10	0	0.90
Parakou	40	40	21	17	2	0.74

Ag: *An. gambiae* s.s.

3.3. Detection of resistance genes

The L1014F *kdr* mutation was found in *An. gambiae* s.s. Malanville and Parakou at various allelic frequencies (Table 2). The increase of *kdr* allelic frequency was positively correlated with CDC bioassays data.

4. Discussion

The resistance level to deltamethrin observed in *An. gambiae* Malanville was higher than the one observed with *An. gambiae* Suru-lere. It was also higher than the resistance level to permethrin observed in *An. gambiae* Parakou and Bohicon. Similar results were already reported by Aïzoun *et al.*[7]. In fact, Malanville is a rice growing area where no insecticidal products are generally used to control agricultural pests comparatively to the vegetable growing area of Parakou and cotton growing area of Bohicon where various insecticidal products are used for this purpose.

The alterations at site of action in the sodium channel, *viz.* the *kdr* mutations in *An. gambiae* s.l. mosquitoes from Malanville were not the same resistance mechanism involved in these mosquitoes populations. A previous study carried out by Djègbé *et al.* showed higher oxidase activity in *An. gambiae* s.l. Malanville populations compared to the Kisumu susceptible strain in 2009[5]. In addition, a recent study carried out by Aïzoun *et al.* on *An. gambiae* Kandi in Alibori province including Malanville district showed that the synergist assay with piperonyl butoxide, an inhibitor of cytochrome P450 monooxygenases, played a role in the deltamethrin resistance observed in Kandi[7].

In the current study, *kdr* frequencies recorded in *An. gambiae* Parakou and Malanville were 74% and 90% respectively. The *kdr* frequencies recorded in *An. gambiae* Parakou and Malanville in 2007 by Corbel *et al.* were 20% and 6% respectively[12]. These results showed that *kdr* frequencies in these *An. gambiae* populations had significantly increased after five years. This is consistent with previous observation reporting an increase of the *kdr* 1014F frequency in *An. gambiae* following a nationwide distribution of LLINs in Niger[13].

In the rice field area of Malanville, the increase in resistance to deltamethrin can be attributed to the augmentation of the 1014F *kdr* prevalence even if deltamethrin is a cyanopyrethroid and detoxifying enzymes such as cytochrome P450 monooxygenases. The sudden increase in *kdr* frequency and pyrethroid resistance is worrying considering the relatively low amount of insecticide use in Malanville. It is possible that *An. gambiae* populations carrying the *kdr* mutation might have migrated through active or/and passive ways from bordering countries (*e.g.* Niger, Nigeria) due to intense traffic and exchanges in this locality[5].

Higher mortality rates were observed with permethrin compared to DDT in *An. gambiae* Parakou and Bohicon populations and may be explained by the presence of an additional resistance mechanism in Benin (*e.g.* “Leu-Ser” mutation) which might confer higher resistance to DDT than to permethrin[14].

Deltamethrin resistance in *An. gambiae* Suru-lere may

be explained by increased use of household insecticide and availability of xenobiotics for larval breeding sites in the urban. It was one of the possible factors selecting for pyrethroid resistance in *An. gambiae* in urban areas.

In conclusion, pyrethroid resistance is widespread in malaria vector in Benin and *kdr* mutation is the main resistance mechanism involved. The L1014F *kdr* mutation frequency in *An. gambiae* s.l. which is resistant to pyrethroid from the south–north transect Benin may increase after a nationwide LLINs implementation. More attention may be paid for the future success of malaria control programmes based on LLINs with pyrethroids in the country.

Conflict of interest statement

We declare that we have no conflict of interest.

Acknowledgements

The authors thank the Ministère de l’Enseignement Supérieur et de la Recherche Scientifique, Benin for supporting the doctoral training of Nazaire. We also thank Dr. William C. BROGDON from CDC Atlanta who supplied us the reagents used for CDC bioassays. The authors would also like to thank Frédéric OKE-AGBO for statistical analysis, Damien TODJINOU for providing technical assistance and Virgile GNANGUENON who contributed to the mapping. This research was funded by the President’s Malaria Initiative of the U.S. Government through USAID and the Ministère de l’Enseignement Supérieur et de la Recherche Scientifique, Benin.

Comments

Background

The spread of pyrethroid resistance in malaria vectors is increasingly becoming a cause of concern especially in endemic countries where insecticide treated nets were the basic tool for vector control. Several published reports on pyrethroid resistance in *Anopheles* mosquitoes in Benin have raised an alarm to the threat it causes to the future usefulness of insecticide treated nets for effective malaria control.

Research frontiers

The present study evaluated the status of pyrethroid resistance, the distribution of *kdr* mutation and compared the present *kdr* allelic frequencies to that of previous reports before the distribution of LLINs took place. Significant increase frequencies of *kdr* mutation have been evidenced in areas of Malanville and Parakou previously with remarkably

low frequencies.

Related reports

Pyrethroid resistance has been documented in *An. gambiae* s.l. in South Benin with Leu–Phe *kdr* mutation as the main resistance mechanism with 80% frequency (Corbel *et al.*, 2007). The experimental huts reducing efficacy of ITNs and IRS against pyrethroid resistant mosquitoes was reported (N'Guessan *et al.*, 2007). Another report in South Benin showed evidence of loss of household protection by insecticide treated nets in areas of pyrethroid resistance (Asidi *et al.*, 2012).

Innovations and breakthroughs

Pyrethroid resistance is a serious concern for *Anopheles* control in South Benin. In the present paper, the authors showed evidence that those populations of *An. gambiae* from Malanville (North Benin) had lost their susceptibility to deltamethrin. And the *kdr* mutation is rapidly spreading with increased frequency in the northern and central (Parakou) parts of Benin despite a scaling up of LLINs distributions nationwide.

Applications

Good knowledge on the status of insecticide resistance would help to prevent vector control failure and the right choice of insecticides. The findings from this study suggested that resistance monitoring would be best advised and helpful to the national malaria control program. In the meantime, investigations on alternatives to pyrethroids would be an indicative route to develop effective tools for insecticide resistance management strategies.

Peer review

The authors have clearly shown a widespread *kdr* frequency in the transect south–north Benin which have increased even after the distribution of LLINs. The effect of selection pressure after a large scale distribution of LLINs has confirmed previous reports by Czeher *et al.* (2008) on an increase Leu–Phe *kdr* mutation in Niger following a countrywide LLINs implementation. The study is important in its field.

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