Distribution of ace-1R and resistance to carbamates and organophosphates in Anopheles gambiae s.s. populations from Côte d'Ivoire

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Abstract

Background: The spread of pyrethroid resistance in Anopheles gambiae s.s. is a critical issue for malaria vector control based on the use of insecticide-treated nets. Carbamates and organophosphates insecticides are regarded as alternatives or supplements to pyrethroids used in nets treatment. It is, therefore, essential to investigate on the susceptibility of pyrethroid resistant populations of An. gambiae s.s. to these alternative products.

Methods: In September 2004, a cross sectional survey was conducted in six localities in Côte d'Ivoire: Toumbokro, Yamoussoukro, Toumodi in the Southern Guinea savannah, Tiassalé in semi-deciduous forest, then Nieky and Abidjan in evergreen forest area. An. gambiae populations from these localities were previously reported to be highly resistant to pyrethroids insecticides. Anopheline larvae were collected from the field and reared to adults. Resistance/susceptibility to carbamates (0.4% carbosulfan, 0.1% propoxur) and organophosphates (0.4% chlorpyrifos-methyl, 1% fenitrothion) was assessed using WHO bioassay test kits for adult mosquitoes. Then, PCR assays were run to determine the molecular forms (M) and (S), as well as phenotypes for insensitive acetylcholinesterase (AChE1) due to G119S mutation.

Results: Bioassays showed carbamates (carbosulfan and propoxur) resistance in all tested populations of An. gambiae s.s. In addition, two out of the six tested populations (Toumodi and Tiassalé) were also resistant to organophosphates (mortality rates ranged from 29.5% to 93.3%). The M-form was predominant in tested samples (91.8%). M and S molecular forms were sympatric at two localities but no M/S hybrids were detected. The highest proportion of S-form (7.9% of An. gambiae identified) was in sample from Toumbokro, in the southern Guinea savannah. The G119S mutation was found in both M and S molecular forms with frequency from 30.9 to 35.2%.

Conclusion: This study revealed a wide distribution of insensitive acetylcholinesterase due to the G119S mutation in both M and S molecular forms of the populations of An. gambiae s.s. tested. The low cross-resistance between carbamates and organophosphates highly suggests involvement of other resistance mechanisms such as metabolic detoxification or F290V mutation.
the exploration of new tools or combination of existing ones.

One of these strategies, used in agriculture as well as in public health, consists to associate in the same treatment, several molecules having different modes of action. Although developed initially for agricultural use and for indoor residual spraying, carbamates and organophosphates constitute a new prospect to circumvent pyrethroid resistance in *An. gambiae* s.s.

In area of high prevalence of *kdr* in Côte d’Ivoire, experimental hut trials of carbamates or organophosphates alone and in combination with pyrethroids on mosquito nets showed very promising results [8,10-13]. However, little is known about the susceptibility status of pyrethroid resistance populations of *An. gambiae* to organophosphates and carbamates in Côte d’Ivoire, as well as potential resistance mechanisms.

Acetylcholinesterase (AChE) is a common target for carbamates and organophosphates. These insecticides blocks transmission of nerve impulses by irreversible inhibition of AChE at cholinergic synapses, causing insect death. Cross-resistance to carbamates and organophosphates can arise by insensitive AChE mechanism due to the glycine to serine substitution (G119S mutation) resulting from a single point mutation in the ace-1 gene [14]. The G119S mutation was selected independently in several mosquitoes species including *An. gambiae* s.s., the major malaria vector in Africa [11,14-17]. This mutation was found in both M and S molecular forms of *An. gambiae* from Côte d’Ivoire [16,17].

In the current study, the geographic extent of insensitive AChE mechanism in *An. gambiae* s.s. populations from Côte d’Ivoire according to molecular forms, as well as their susceptibility status to carbamates and organophosphates were investigated.

**Methods**

**Mosquito populations and sampling sites**

The study sites form a north-to-south transect across the Southern Guinea savannah, the semi-deciduous forest and the evergreen forest areas in Côte d’Ivoire. The last two zones are characterized by intensive human activities and agricultural land-degraded forest mosaic. Mosquitoes were collected during the rainy season from six localities: Toubokro (7°N; 5°35’ W), Yamoussoukro (6°82’ N; 5°28’ W) and Toumodi (6°55’ N; 5°03’ W) located in the Southern Guinea savannah, Tiassalé (5°88’ N; 4°38’ W) in a semi-deciduous forest area, then Nieky (5°20’N; 4°10’W) and Abidjan (5°33’N; 4°03’ W) in a evergreen forest area (Figure 1). Samples were collected from coffee and cocoa industrial plantations in Toubokro, banana cultivation fields in Nieky and in urban areas in Yamoussoukro, Toumodi, Tiassalé and Abidjan. Mosquitoes were collected at larval stage, brought to the laboratory and reared until for emergence of adults. A reference laboratory strain of *An. gambiae* s.s. named “Kisumu”, native from Kenya and susceptible to all insecticides was used as control.

**Susceptibility test**

Bioassays were carried out using WHO test kits for adults mosquitoes [18] with four insecticides of technical grade quality: two carbamates (0.4% carbosulfan, 0.1% propoxur) and two organophosphates (0.4% chlorpyrifos-methyl and 1% fenitrothion). Filter papers were impregnated according to WHO specifications by the Institut Pierre Richet de Bouaké. Papers were stored at 4°C and were not used more than three times.

Tests were performed with batches of 25 unfed females of *An. gambiae* s.s., 2-5 days old, four replicates per insecticide. Mosquitoes were exposed to the insecticide treated papers for 60 min at 27 ± 1°C and 80% relative humidity. After exposure, mosquitoes were kept in observation tubes, supplied with 10% honey solution and held for 24 h before scoring mortality. Batches exposed to untreated papers were used as control.

**M/S taxon determination**

According to previous studies, *An. gambiae* complex in Côte d’Ivoire was only represented by *An. melas* on the Atlantic littoral area and *An. gambiae* s.s., the most widespread all over the country [19-21]. So the PCR analysis in this study was carried directly on the molecular forms of *An. gambiae* s.s. Genomic DNA was extracted from individual mosquitoes according to Collins et al [22] and used for PCR analysis to determine M/S taxon according to Favia et al [23]. The PCR conditions were 10 min at 94°C as initial step, followed by 29 cycles (94°C for 30 seconds, 63°C for 30 seconds and 72°C for 30 seconds). After the last cycle the products were finally extended for 7 min at 72°C. Primers used in the PCR were: R5 5’GCCAGTCTGATAGCCG3’, R3 5’CAGATCTAGGAGCTCCAG3’, Mopint 5’GCCCTTCTCGATGGCAT3’, B/S 5’ACACAGATGTCTGTGGC3’. Amplified fragments were analysed on a 1.5% agarose gel.

**DNA diagnostic test for insensitive acetylcholinesterase G119S mutation**

Genomic DNA extracted from the field samples and used for PCR was also used to determine the phenotypes for insensitive AChE G119S mutation according to Weill et al [16]. The DNA was PCR amplified with the primers Ex3Agdir 5’AGGATGGCCCGCTGGAACAG3’ and Ex3Agrev 5’GATCGTGGACACCGTGTTCG3’. Amplified fragments were analysed on a 1.5% agarose gel.
Figure 1 Map of Côte d’Ivoire showing the localities in the different ecological zones where anopheline mosquitoes were collected
and fractionated on a 2% agarose gel. The two primers produced a 403 bp fragment, which is undigested by AluI for susceptible homozygous mosquitoes (SS), and cut into two fragments (253 bp and 150 bp) for homozygous resistant (RR). Heterozygous individuals (RS) display a combined pattern.

Data analysis
Mortality data were analysed according to WHO [18]. To compare the status of insecticide resistance, Fisher’s exact test was performed to determine if there was any significant difference between mortality rates of two given populations of An. gambiae s.s. using Statistica 6.0. Allelic frequencies of G119S mutation were analysed using the version 3.2a of Genepop [24]. To assess if the mutation frequencies was identical across populations, the test of genotypic differentiation was performed [25].

Results
Susceptibility to carbamates and organophosphates
Mortality rates of the Kisumu reference strain to all insecticides was 100% (Table 1). Conversely, all the field samples were resistant to carbamates, with mortality rates less than 83%. Susceptibility to chlorpyrifos-methyl was assessed on five populations except on the Yamoussoukro population. Chlorpyrifos-methyl resistance was detected in Toumodi and Tiassalé, with 82-94% mortality rates, while it was suspected in Toumbokro with 97% mortality rate. The two other populations were fully susceptible to this organophosphate. Fenitrothion resistance was observed in five out of the six populations tested (Toumbokro, Toumodi, Tiassalé, Nieky, Abidjan). Only the Yamoussoukro population was fully susceptible to this insecticide. Overall the two populations from Toumodi and Tiassalé were resistant to all insecticides used. Tiassalé sample was the most resistant to carbamates and organophosphates with mortality rates of 3% and 12% for carbosulfan and propoxur and 83% and 30% for chlorpyrifos-methyl and fenitrothion, respectively.

Molecular forms and frequencies of the G119S mutation
All PCR analysis to determine M/S molecular forms realized in this study were positive, showing either the form M or the form S. So it was not necessary to make the PCR analysis for species identification [26].

Three hundred twenty-eight mosquitoes were identified to molecular forms and analyzed for the G119S mutation; results are shown in Table 2. The M and S molecular forms of An. gambiae s.s. occurred in sympatry in two of the six localities, namely Toumbokro and Toumodi in the savannah area. However, the M-form was predominant in the six areas, representing 91.8% of the whole sample (n = 328). In sympatric areas, the frequencies of the S-form were 41.9% (n = 62) and 1.3% (n = 76) respectively in Toumbokro and Toumodi. However, no M/S heterozygote was found.

The G119S mutation was detected in all the six populations tested, but only at heterozygote state, either in the M or in the S form. The highest mutation frequency was observed in the M form from the Tiassalé urban area located in semi-deciduous forest (50%) and the lowest in the M form from the Nieky banana cultivation fields in evergreen forest (12%). No significant difference was seen between G119S mutation frequencies in M and S forms from Toumbokro (p = 0.9153).

| Localities    | Carbosulfan (0.4%) | Propoxur (0.1%) | Chlorpyrifos-methyl (0.4%) | Fenitrothion (1%) |
|--------------|-------------------|----------------|-----------------------------|------------------|
| Kisumu       | 100 (101)         | S              | 100 (104)                   | S                |
| Toumbokro    | 21.4b (98)        | R              | 40.9b (88)                  | R                |
| Yamoussoukro | 42.3d (104)       | R              | 96.9 (97)                   | S                |
| Toumodi      | 28.4d (95)        | R              | 93.3d (89)                  | R                |
| Tiassalé     | 2.8a (217)        | R              | 82.8a (192)                 | R                |
| Nieky        | 28.2c (103)       | R              | 82.8c (99)                  | S                |
| Abidjan      | 39.0d (100)       | R              | 100 (102)                   | S                |

Number of tested mosquitoes in parentheses; NT: no tested; Mort: Mortality rate 24 h post exposure; S: indicates susceptibility; R: suggests resistance.

NB: Numbers in the same column with the same superscript do not differ significantly by Fisher’s exact test (p > 0.05)
Discussion

The distribution of M and S molecular forms of An. gambiae s.s. in the study agrees with previous findings that reported both M and S forms in Guinea savannah areas and only the M form in the forest areas [27-30]. This geographic distribution seems to follow more the global environment than the breeding sites nature. Both forms are involved in carbamate and organophosphate resistance, although at different level according to insecticides. Indeed, in this study, An. gambiae s.s. displayed large variations in resistance level to carbamates and organophosphates. Although the wild populations were all resistant to carbamates, resistance was less marked to propoxur than to carbosulfan at WHO diagnostic concentrations.

All these populations were as resistant to carbosulfan as the population of Yaokoffikro in surrounding area of Bouaké [11]. The resistance reported in Bouaké was attributed to agricultural or domestic hygiene or public health use of carbamates. In Burkina-Faso, Diabaté et al [31] attributed An. gambiae s.s. pyrethroid resistance in cotton field areas to their use in agriculture.

The observed cross-resistance to organophosphates and carbamates in Tiassalé and Toumodi highlights implication of their common target site: the AChE-1. Although the mutation ace-1 G119S provided cross-resistance to organophosphates and carbamates, the resistance level greatly varied between both insecticide families. This difference observed in resistance level could be the consequence of their difference observed in dominance level relied on insecticide specificity. According to Djogbénon et al [32], dominance status of ace-1 G119S varied between semirecessivity with fenitrothion and chlorpyrifos methyl to semidominance with propoxur and carbamates. The fact that low cross-resistance was observed in the other populations, suggests and confirms potential involvement of metabolic resistance mechanisms and/or alternative mutation associated to G119S. This may explain why mortality rates to organophosphates among samples from Nieky, Abidjan and Yamoussoukro were so strong despite confirmed resistance level to carbamates in bioassays.

Such result could also be explained by possible cross-resistance between organophosphates and pyrethroids based on an increased detoxification mechanism were as suggested for other anopheline species selected for pyrethroid resistance [33].

Moreover an alternative mutation in ace-1 gene was described in the Culex pipiens strain originating from Cyprus. This mutation is F290V substitution and it confers cross-resistance to OP and carbamate insecticides [34]. Because C. pipiens and An. gambiae s.s. share G119S, it is possible that they share also this other mutation. Asidi et al [13] had noted that G119S mutation did not confer effective resistance to chlorpyrifos-methyl. Yet, the G119S mutation involved certainly a high resistance to carbamate but could enhance organophosphate hydrolysis. Similar mutations in a homologous position to G119S are known to alter substrate specificity in Drosophila melanogaster and enhanced hydrolysis of some organophosphates [35].

The presence of G119S mutation in both M and S forms of An. gambiae s.s. has already been reported by Weill et al [16] and Djogbénon et al [17] and was suggested to result from introgression between forms. The wide distribution of ace-1R reported here could result

Table 2: Acetylcholinesterase phenotypes and frequency of G119S mutation in the molecular M and S forms of Anopheles gambiae s.s.

| Locality   | M Form | S Form | F(G119S)% (%) | S | RS | R | F(G119S)% (%) | S | RS | R |
|------------|--------|--------|---------------|---|----|---|---------------|---|----|---|
| Toumbokro  | 12     | 24     | 0             | 33.3 b | 58.1 | 8 | 18 | 0 | 34.6 | 41.9 |
| Yamoussoukro | 7     | 24     | 0             | 38.7 b | 100 | 0 | 0 | 0 | _ | 0 |
| Toumodi    | 47     | 28     | 0             | 18.6 a | 98.7 | 0 | 1 | 0 | 50.0 | 1.3 |
| Tiassalé   | 0      | 82     | 0             | 50.0 c | 100 | 0 | 0 | 0 | _ | 0 |
| Nieky      | 23     | 7      | 0             | 11.7 a | 100 | 0 | 0 | 0 | _ | 0 |
| Abidjan    | 26     | 21     | 0             | 22.3 a | 100 | 0 | 0 | 0 | _ | 0 |
| Total      | 115    | 186    | 0             | 30.9 | 91.8 | 8 | 19 | 0 | 35.2 | 8.2 |

*: Percentage of each molecular form in the population tested

a,b Values sharing a superscript letter are not significantly different at the 5% level for G119S mutation distribution

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The presence of G119S mutation in both M and S forms of An. gambiae s.s. has already been reported by Weill et al [16] and Djogbénon et al [17] and was suggested to result from introgression between forms. The wide distribution of ace-1R reported here could result
from an unique event that then spread as reported in C. pipiens amplified esterase B2 genes through the world [36].

The absence of homozogous resistant individuals might be related to high fitness cost of the G119S mutation, resulting on death of the homozogous resistant [13,16,17]. Indeed, greater mortality of resistant individuals during pupation relative to their sensitive counterparts was reported. There was also evidence for costs to adult fitness as resistant individuals were smaller than sensitive adults [37]. Consequently, in area where the resistant allele ace-1R is present, resistant mosquitoes will mainly at heterozygote state (ace-1RS). Because of this fitness cost, at least one duplication combining resistant and susceptible alleles of the ace-1 locus has recently appeared, started to spread and replace ace-1R in treated areas [17,38-41]. Duplications lead to an excess of heterozygotes in natural populations because that heterozygotes involving either ace-1S or ace-1R alleles do not exhibit deleterious side effects. To date, no specific test is available for detecting specifically ace-1 duplications as mosquitoes carrying duplications appear as heterozygous for ace-1R mutation.

Further investigation is needed to tackle the origin of the difference of resistance between carbamates and organophosphates.

Conclusion
Data from this study complemented resistance to carbamates and organophosphates in An. gambiae s.s. populations from Côte d’Ivoire and the wide distribution of G119S mutation in both molecular forms. The low cross-resistance between carbamates and organophosphates through susceptibility tests in most of the populations suggests the involvement of other resistance mechanisms, probably a metabolic detoxification or an alternative mutation such as the F290V substitution. These results must be carefully considered while elaborating malaria control programs in Côte d’Ivoire.

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

Authors’ contributions
LPAA, AAK designed the study, conducted the field work, genotyping, summarized the data and drafted the manuscript. MAA, ET jointly carried out PCR assays, and interpreted the results. PWK and MK supervised LPAA and AAK and contributed to the manuscript. FC contributed to design of the study and manuscript drafting. All authors read and approved the final manuscript.

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