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Behavioural divergence of sympatric *Anopheles funestus* populations in Burkina Faso

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Abstract

Background: In Burkina Faso, two chromosomal forms of the malaria vector *Anopheles funestus*, Folonzo and Kiribina, are distinguished by contrasting frequencies of shared polymorphic chromosomal inversions. Sympatric and synchronous populations of Folonzo and Kiribina mate assortatively, as indicated by a significant deficit of heterokaryotypes, and genetic associations among inversions on independently segregating chromosome arms. The present study aimed to assess, by intensive longitudinal sampling, whether sympatric Folonzo and Kiribina populations are characterized by behavioural differences in key malaria vectorial parameters.

Methods: The study was conducted in two adjacent villages near Ouagadougou, in the dry savanna of central Burkina Faso. Mosquito adult resting behaviour of both forms was compared based on parallel indoor/outdoor collections across six breeding seasons; 8,235 fully karyotyped samples of half-gravid females were analysed in total. Additionally, indoor/outdoor human biting behaviour, host selection, and *Plasmodium falciparum* sporozoite rate was assessed and compared between chromosomal forms.

Results: The Kiribina form was numerically predominant in the area. However, the Folonzo form was significantly over-represented in indoor resting collections and showed stronger post-prandial endophily, while Kiribina predominated outdoors. Neither form was statistically distinguishable in human biting behaviour, and both were more likely to seek human blood meals indoors than outside. The human blood index and sporozoite rate were comparably high in both chromosomal forms in indoor collections (>89% and >8%, respectively).

Conclusions: Both Kiribina and Folonzo chromosomal forms are formidable malaria vectors in Burkina Faso. However, the significantly greater tendency for the Kiribina form to rest outdoors despite its pronounced anthropophily suggests that uniform exposure of the overall *An. funestus* population to indoor-based vector control tools cannot be expected; Kiribina is more likely to evade indoor interventions and escape unharmed outdoors, reducing the efficacy of malaria control. Accordingly, more efficient methods to detect Kiribina and Folonzo, and a more complete understanding of their distribution and behaviour in Africa are advocated.

Keywords: *Anopheles funestus*, Anthropophily, Behavioural divergence, Chromosomal forms, Exophily, Folonzo, Kiribina, Malaria vector, West Africa

Background

The malaria vectorial system in tropical Africa is dominated by four species of major importance, *Anopheles gambiae*, *Anopheles coluzzii*, *Anopheles arabiensis* and *Anopheles funestus*, which are broadly codistributed across much of tropical Africa in close association with humans

[1,2]. The first three species belong to the same cryptic species complex (the *An. gambiae* complex) whose members cannot be distinguished morphologically at any developmental stage, although they differ in aquatic larval ecology and adult behaviours relevant to malaria transmission and control (e.g., degree of anthropophily and tendency to blood-feed or rest indoors) [3,4]. *Anopheles funestus* and its presently recognized closest relatives are classified into a group and subgroup [5,6] rather than a species complex, owing to slight morphological distinctions mainly at immature stages. However, further cryptic

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taxonomic complexities within the group have recently come to light and more can be anticipated as *An. funestus* research emerges from a period of neglect [7-11]. Malaria transmission by the Funestus Subgroup is overwhelmingly attributed to *An. funestus sensu stricto*, owing to its strong preference for human blood meals (see reviews by [7,12]).

Anopheles funestus s.s. is characterized by abundant genetic polymorphism, exemplified by at least 17 chromosomal re-arrangements segregating within and among populations across Africa [13,14]. Although this species is generally considered to be uniformly anthropophilic and endophilic throughout its range, complex and incompletely understood patterns of population structure based on cytogenetic and DNA markers have been detected [15-20]. In particular, two chromosomal forms designated "Folonzo" and "Kiribina" have been described in West Africa [16]. First discovered in Burkina Faso and most intensively characterized in that country, Folonzo and Kiribina populations carry markedly different frequencies of shared polymorphic chromosomal inversions, mainly involving arm 3R [16,17]. In localities where the chromosomal forms are synchronous and stably sympatric across successive breeding seasons and years, there are highly significant departures from Hardy Weinberg equilibrium and significant genetic associations among physically unlinked inversion systems; alternative homokaryotypes are more frequent than expected under random mating, and there are significant deficits of heterozygotes in virtually all population samples, consistent with assortative mating by form [16,17].

Neither inversions nor inversion combinations are diagnostic taxonomic characters. However, the Kiribina form is predominantly homokaryotypic for the standard chromosomal arrangements, while Folonzo, the more chromosomally polymorphic taxon, carries high frequencies of inversions 3Ra and 3Rb, and presumably corresponds to *An. funestus* from East Africa, where Kiribina has not been recorded [16]. Strongly reminiscent of the chromosomal forms of *An. gambiae* [21], these alternative karyotypes show cyclical patterns of seasonal variation in relative abundance linked to temperature and rainfall, likely reflecting differences in geographic distribution modulated by larval habitat utilization [22]. Although direct evidence is lacking, the Folonzo form is associated with natural larval habitats such as marshes, while Kiribina is associated with larval habitats created by the practice of agriculture, notably rice fields. Molecular genetic studies using mtDNA and microsatellite markers revealed very slight but significant divergence between sympatric samples of Folonzo and Kiribina across Burkina Faso, although nuclear divergence was not genome-wide and could be explained by loci on chromosome 3R inside and outside inversions [23,24]. These data are suggestive of an incipient process of ecological divergence and lineage splitting, similar to, but less advanced than, that responsible for the

divergence of *An. coluzzii* and *An. gambiae* (formerly recognized as chromosomal or molecular forms [25,26]).

Previous studies in Burkina Faso and Senegal have reported similarly high rates of anthropophily and comparable *Plasmodium falciparum* infection rates in sympatric Folonzo and Kiribina populations [23,24,27]. However, there were indications of differences in indoor resting behaviour, leading to the suggestion that the Kiribina form may be more easily diverted to outdoor resting and biting, particularly in localities where alternative hosts such as cattle outnumber the local human population [16]. If the ecological and genetic heterogeneities between Folonzo and Kiribina indeed extend to behavioural differences of importance to malaria epidemiology and control, these vectorial differences must be understood more deeply. Toward that end, resting and biting behaviour were assessed separately for sympatric and synchronous Folonzo and Kiribina populations in the rural villages of Kuiti and Koubri near Ouagadougou, Burkina Faso. Observations spanned six breeding seasons and 8,235 fully karyotyped Folonzo and Kiribina adult half-gravid females.

Methods

Study area

The study was carried out in the arid Sudan savanna vegetation belt of West Africa, in adjacent rural villages located 35 km south of Ouagadougou, the capital of Burkina Faso. Koubri (12°11'54 N; 1°23'43 W) and Kuiti (12°11'36 N; 1°23'11 W) lie about 1 km apart on opposite margins of an artificial lake bordered by permanent swamps. A detailed map and additional information about the study area can be found elsewhere [17,24]. In this region, the *An. funestus* breeding season commences at the end of the rainy season (September), extends throughout the cool dry season (October-February), and ends in April, mid-way through the hot dry season (March-May). Folonzo peaks in relative abundance following the rains, in October-December [22].

Chromosomal form identification

Adult Funestus Group females were sorted morphologically [28] in the field under a dissecting microscope. Ovaries from females at the appropriate gonotrophic stage were immediately dissected and preserved by individual mosquito in 1.5 ml microcentrifuge tubes using Carnoy's fixative (ethanol:glacial acetic acid, 3:1), while the associated carcass was placed in a correspondingly labelled microcentrifuge tube with desiccant. Molecular taxonomic identification of *An. funestus* based on DNA extracted from the carcass was performed with a modified rDNA-based PCR assay [24]. Ovaries in Carnoy's were held on ice until they could be stored at -20°C for later polytene chromosome analysis. Polytene chromosomes of *An. funestus* were spread [29] and examined

under a phase-contrast microscope. Karyotypes were assigned using the cytogenetic map of Sharakhov *et al.* [13]. Of all karyotyped samples, 92% were successfully scored for all inversions. Chromosomal form assignment followed the deterministic algorithm of Guelbeogo *et al.* [17]. Using a probabilistic assignment test as an alternative method of classification of karyotypes sampled from the same localities as the present study, these authors estimated the rate of mis-classification to be very low, about 0.7%.

Resting behaviour

An estimate of the odds of adult females of the Folonzo or Kiribina forms resting indoors was calculated by comparing the relative abundance of each form in resting collections that were conducted indoors and outdoors in parallel. Indoor resting mosquitoes were sampled in the afternoon inside multiple huts and compounds in both villages, by insecticide spray-sheet catches (ISC) three times per week; mosquitoes resting outdoors in the villages were sampled at least twice weekly from four Muirhead-Thompson style pit-shelters with manual aspirators [30]. In addition, an estimate of the odds of post-prandial indoor-resting by outdoor-biting Folonzo or Kiribina was calculated based on blood meal identifications performed on indoor/outdoor-resting collections made between 2005–2007 (described below). As cattle do not share the domestic environment with humans in the study area, mosquitoes with exclusively bovine blood meals must have fed outdoors on cattle. Accordingly, the numbers of each form that fed solely on cattle were compared between indoor-resting (ISC) and outdoor-resting (PIT) collections from the same time period.

Human biting behaviour

Human biting behaviour of Folonzo and Kiribina was assessed by human landing catches (HLC). Two teams of trained collectors worked in two different compounds in eight-hour shifts (21:00–05:00), twice per week from October 2002–January 2003. Each team consisted of a pair of collectors, one of whom performed an indoor landing catch while the other did the same outdoors, reversing positions on a subsequent night to control for collector-specific effects. To identify the chromosomal form of host-seeking *An. funestus* captured by HLC, mosquitoes were blood-fed on rabbits the same night of capture, and held in the insectary until they reached the stage of ovarian development appropriate for polytene chromosome analysis.

Blood meal identification and *Plasmodium falciparum* detection

Samples of blood-fed mosquitoes collected in the 2005–2006 and 2006–2007 breeding seasons were cut transversely between the thorax and the abdomen. The origin

of the blood meal (human, bovine, mixed) in the posterior portion was identified by an enzyme-linked immunosorbent assay (ELISA) using specific monoclonal antibodies. The presence of *P. falciparum* circumsporozoite protein (CSP) in the anterior portion (head + thorax) also was detected by ELISA in these samples.

Data analysis

The human blood index (HBI) of each chromosomal form was calculated as the proportion of human and mixed blood meals identified relative to all blood meals identified by ELISA in samples of that form. The sporozoite rate of each form was calculated as the proportion of mosquitoes in a sample that were positive for *P. falciparum* CSP by ELISA. The odds ratio (OR; [31]), the ratio of the odds of an event occurring in one group to the odds of it occurring in another, was used to compare vectorial parameters between the chromosomal forms. The precision of the OR was estimated using the 95% confidence interval (CI). *P*-values are reported based on the Pearson Chi-square test of association for 2x2 contingency tables, with *P* < 0.05 considered as significant.

Ethical approval

The study protocols were reviewed and approved by the institutional health ethical review board of Burkina Faso. Informed consent was obtained from participants.

Results

Resting behaviour by sympatric populations of the two chromosomal forms of *An. funestus* was assessed by parallel indoor/outdoor collections across six breeding seasons in two adjacent rural villages located in the dry savanna of central Burkina Faso. The results from the 8,235 fully karyotyped samples are presented in Table 1. Samples of the Kiribina form were generally larger than corresponding samples of the Folonzo form both indoors and out, particularly for the outdoor collections. However, during most breeding seasons and for the pooled samples, the Folonzo form was more likely than Kiribina to rest inside human dwellings rather than outside in pit-shelters. For 50% of the seasons, particularly when the numbers of outdoor-resting mosquitoes were sufficiently large, a Chi-square test of association for a 2x2 contingency table indicated that the stronger indoor resting tendency of Folonzo relative to Kiribina was statistically significant (Table 1).

A measure of post-prandial resting behaviour by outdoor-feeding *An. funestus* was estimated by focusing on those mosquitoes with exclusively bovine-derived blood meals and comparing their numbers between indoor-resting and outdoor-resting samples of each chromosomal form. Based on this measure, the Folonzo form was significantly more

Table 1 Resting behaviour of *Anopheles funestus* chromosomal forms in Burkina Faso

Season	Sample	Total	Folonzo	Kiribina	OR	0.95 CI	P
1999-2000	ISC/indoor	1154	377	777	1.94	1.29-2.93	0.001
	PIT/outdoor	155	31	124			
2000-2001	ISC/indoor	1164	208	956	1.58	0.55-4.54	NS
	PIT/outdoor	33	4	29			
2001-2002	ISC/indoor	2733	659	2074	0.95	0.49-1.84	NS
	PIT/outdoor	48	12	36			
2002-2003	ISC/indoor	485	301	184	3.03	1.70-5.37	<0.0001
	PIT/outdoor	57	20	37			
2005-2006	ISC/indoor	99	63	36	2.33	0.49-11.02	NS
	PIT/outdoor	7	3	4			
2006-2007	ISC/indoor	932	343	589	11.87	8.93-15.76	<0.0001
	PIT/outdoor	1368	64	1304			
Pooled	ISC/indoor	6567	1951	4616	4.84	4.02-5.82	<0.0001
	PIT/outdoor	1668	134	1534			

OR, odds ratio of indoor/outdoor resting by Folonzo *versus* Kiribina; CI, confidence interval; ISC, insecticide spray-sheet catch; PIT, Muirhead-Thomson pit-shelter [30].

likely than Kiribina to rest indoors following a bovine blood meal taken outdoors (Table 2).

HLC, conducted in parallel indoors and outdoors during the 2002–2003 breeding season, were used to compare human-biting behaviour between the chromosomal forms. Based on the >1,000 female mosquitoes captured, karyotyped and assigned to chromosomal form in 2002–3, human biting behaviour indoors *versus* outdoors was indistinguishable between chromosomal forms (Table 3). For both, the proportion of mosquitoes seeking human blood meals indoors *versus* outdoors was higher and of a similar magnitude. Importantly, the absolute numbers of the Folonzo form captured by HLC, both indoors and out, were larger than those from the corresponding Kiribina samples. These observations suggest that the Kiribina form may be more opportunistic, and the Folonzo form more anthropophilic, in host-seeking behaviour.

Host selection was assessed by blood meal identification during the 2005–2006 and 2006–2007 breeding seasons. The indoor resting samples of Folonzo and Kiribina both had a relatively high human blood index, 95.9 and 89.3%, respectively (Table 4). Folonzo was the form more likely

Table 2 Post-prandial resting behaviour of outdoor-feeding *Anopheles funestus* chromosomal forms in Burkina Faso (2005–2007)

Sample*	Total	Folonzo	Kiribina	OR	0.95 CI	P
ISC/indoor	38	9	29	5.64	2.43-13.10	0.0003
PIT/outdoor	518	27	491			

OR, odds ratio of post-prandial indoor resting by Folonzo *versus* Kiribina for outdoor-feeding *An. funestus* (2005–2007 season); CI, confidence interval; PIT, Muirhead-Thomson pit-shelter [30]; ISC, insecticide spray-sheet catch.

*Includes only *An. funestus* having blood fed exclusively on cattle, based on blood meal identification.

to have fed on humans in whole or in part, rather than solely on cattle ($P < 0.006$), but this trend may reflect differences between forms in post-prandial resting behaviour rather than differences in the intrinsic preference for human hosts (i.e., host choice). While the size of indoor samples was balanced between Folonzo and Kiribina, there was a large skew in outdoor resting sample size between the forms, only 30 for Folonzo compared to 529 for Kiribina, reflecting the greater tendency for the latter form to rest outdoors. Both outdoor resting samples had drastically lower human blood indices, 10 and 7% for Folonzo and Kiribina, respectively. Folonzo remained the form more likely to have fed on humans than cattle (OR, 1.44), although this trend was not statistically significant.

During the same 2005–2007 seasons that host selection was evaluated, samples were analysed for *P. falciparum* infection by testing for the presence of the CSP in indoor and outdoor resting samples of the two chromosomal forms. Infection rates in the indoor resting samples did not differ significantly between forms, being similarly high in both (8.5–8.8%; Table 5). Among the outdoor resting mosquitoes, the small Folonzo sample contained no sporozoite-positives, while the much larger Kiribina sample contained 32 (3%) sporozoite positives.

Discussion

Intensive longitudinal sampling of *An. funestus* from adjacent villages in the Sudan savanna of Burkina Faso, West Africa, affirms and extends the previous findings by Costantini *et al.* [16] of behavioural divergence between sympatric and synchronous chromosomal forms known as Folonzo and Kiribina. The high rate of anthropophagy by both forms (>89% of indoor samples),

Table 3 Human biting behaviour of *Anopheles funestus* chromosomal forms in Burkina Faso (2002–2003)

Month	Sample	Total	Folongo	Kiribina	OR	0.95 CI	P
October	HLC/indoor	363	263	100	0.87	0.54-1.40	NS
	HLC/outdoor	117	88	29			
November	HLC/indoor	218	151	67	1.18	0.74-1.87	NS
	HLC/outdoor	131	86	45			
December	HLC/indoor	70	42	28	0.62	0.30-1.30	NS
	HLC/outdoor	58	41	17			
January	HLC/indoor	80	45	35	1.63	0.73-3.65	NS
	HLC/outdoor	34	15	19			
Pooled	HLC/indoor	731	501	230	1.04	0.79-1.37	NS
	HLC/outdoor	340	230	110			

OR, odds ratio of indoor/outdoor human biting by Folongo versus Kiribina; CI, confidence interval; HLC, human landing catch.

coupled with comparably high rates of *P. falciparum* infection (>8% of indoor samples) emphasize the fact that Folongo and Kiribina both are formidable malaria vectors in this part of Africa. The Kiribina form often outnumbered Folongo. Yet, Folongo was disproportionately represented in indoor versus outdoor resting samples and was more inclined to post-prandial endophily, while Kiribina was over-represented outdoors in pit shelters. This suggests that the overall *An. funestus* population is not uniformly exposed to indoor-based malaria interventions such as insecticide-treated nets and house spraying by residual insecticides, and that those indoor interventions are less effective against the Kiribina form.

There is precedence for chromosomal inversion-associated heterogeneity in mosquito resting behaviour in the West African savanna, uncovered by Coluzzi and colleagues through polytene chromosome analysis of *An. gambiae* and *An. arabiensis* populations during the Garki Project in Nigeria [3,32]. Such behavioural heterogeneity was responsible for the failure to interrupt malaria transmission during the course of the Project, despite rigorous insecticide applications and simultaneous administration of anti-malarial drugs to the human population [33]. Indeed,

there are hints that this same phenomenon has been witnessed previously with respect to *An. funestus* in the West African savanna, where Kiribina co-exists with Folongo. In the absence of Kiribina in eastern and southern Africa, historical house spraying campaigns not only locally eliminated *An. funestus*, but the effect was maintained for several years following the cessation of spraying, due to the apparent inability of *An. funestus* to recolonize some areas [34]. Likewise, *An. funestus* was eliminated from humid forest and degraded forest areas in West Africa where malaria is meso- or hypo-endemic [34], an environment where Folongo is predicted to dominate [16,35-37]. However, in the savannas of West Africa where malaria is holo- or hyperendemic, similar historical indoor spraying campaigns failed to eliminate the species [34]. Exophilic populations persisted which, despite marked anthropophily, continued to feed outdoors on cattle as well as humans, and also entered sprayed houses to bite humans, but escaped unharmed to rest outdoors. These exophilic populations likely represented what would now be recognized as the Kiribina form of *An. funestus*.

More recently, further epidemiologically significant behavioural heterogeneities in *An. funestus* from the same

Table 4 Host selection of *Anopheles funestus* chromosomal forms in Burkina Faso (2005–2007)

Sample	Blood meal	Folongo	Kiribina	OR	0.95 CI	P
ISC/indoors	Human + Mixed	191 + 21 (212)	232 + 11 (243)	2.81	1.30-6.07	0.006
	Bovine	9	29			
	Total	221	272			
	HBI	95.9%	89.3%			
PIT/outdoors	Human + Mixed	2 + 1 (3)	13 + 25 (38)	1.44	0.42-4.95	NS*
	Bovine	27	491			
	Total	30	529			
	HBI	10.0%	7.2%			

*NS by Fisher Exact test. Chi-square not calculated due to an expected cell frequency below 5.

OR, odds ratio of human/bovine blood meal by Folongo versus Kiribina; CI, confidence interval; ISC, insecticide spray-sheet catch; PIT, Muirhead-Thomson pit-shelter [30]; Mixed, blood meal of human and bovine origin; HBI, human blood index.

Table 5 *Plasmodium falciparum* sporozoite rate of *Anopheles funestus* chromosomal forms in Burkina Faso (2005–2007)

Sample	CSP	Folonzo	Kiribina	OR	0.95 CI	P
ISC/indoor	+	27	45	0.97	0.59-1.59	NS
	-	291	469			
	Total	318	514			
	%CSP+	8.5%	8.8%			
PIT/outdoor	+	0	32	0.00	0.00-2.14	NS*
	-	61	1035			
	Total	61	1067			
	%CSP+	0.0%	3.0%			

*NS by Fisher Exact test.

CSP, circumsporozoite protein; OR, odds ratio of sporozoite infection in Folonzo versus Kiribina; CI, confidence interval; ISC, insecticide spray-sheet catch; PIT, Muirhead-Thomson pit-shelter [30].

biogeographical area have been recognized following large-scale implementation of indoor-based vector control interventions. After mass deployment of insecticide-treated bed nets, the biting cycle of *An. funestus* shifted from its usual peak between 02:00 and 04:00 toward a later peak between dawn and early morning hours, when human hosts are less likely to be protected by nets [38]. Unfortunately, it is not known whether this behavioural shift was associated with a change in the chromosomal composition of the local *An. funestus* population.

The Folonzo and Kiribina chromosomal forms have been well characterized across several hundred kilometres and all ecozones of Burkina Faso [16,23]. However, their broader geographical distribution in Africa is poorly known. Certainly, they occur as far west as Senegal [15,27,39]. A recent study of sympatric populations of these forms, the first of its kind in Senegal, found stable co-existence of the forms across three successive breeding seasons and concluded, in accord with the present study, that Kiribina predominated, and rates of anthropophagy and sporozoite infection were comparable between forms, although both metrics were considerably lower in Senegal (~30 and ~3%, respectively) than they were in Burkina Faso [27]. Unfortunately, due to very low outdoor resting sample size (five total, of which only three could be identified chromosomally as Kiribina), indoor/outdoor resting behaviour was difficult to compare between forms, and thus, between studies. Cameroon is the most easterly country in which *An. funestus* chromosomal forms have been reported [36], but their vectorial heterogeneities (if any) are essentially uncharacterized. Available cytogenetic data suggest that these forms are largely allopatric in Cameroon, with Folonzo occurring in the mesic, forested south and Kiribina to the north in the dry savannas, except for a central contact zone at the forest-savanna transition, where stable sympatric co-existence of the two forms has not been clearly resolved [35-37]. In another

parallel with the *An. gambiae* chromosomal forms, there is no evidence for the co-occurrence of *An. funestus* chromosomal forms in East Africa [40]; existing populations of *An. funestus* in eastern Africa are hypothesized to be allied with the Folonzo form [16], although that proposal has yet to be tested genetically.

Ample indication now exists of the practical importance of population structure and behavioural heterogeneities hidden within *An. funestus*, for malaria epidemiology and control in West Africa, if not beyond. In this light, the dearth of information about the wider geographic distribution and associated bionomics and vectorial parameters of the Folonzo and Kiribina forms is a problem that must be remedied as a matter of priority. The polytene chromosomes of *An. funestus* are considerably more difficult to spread and analyse than those of *An. gambiae*, a factor that has impeded past research on *An. funestus*. The demanding and specialized task of polytene chromosome-based identification, the restrictive sex and life stage from which favourable chromosomes are obtained, and the absence of any known DNA-based diagnostics to distinguish the chromosomal forms, all but prohibit deeper field investigation of Folonzo and Kiribina, particularly studies of their larval biology which is presumed to be a driver of their ecological and behavioural divergence. Genome sequencing of *An. gambiae* in 2002 [41], and the discovery of molecular forms of *An. gambiae* detectable by a simple PCR assay [26], greatly transformed understanding of the complexities of *An. gambiae* population structure and its impacts on malaria transmission. Recent whole genome sequencing and a newly available reference assembly for *An. funestus* [42] offer a platform that will support a more detailed understanding of *An. funestus* population structure across Africa, as well as an efficient means to discover genomic sequences potentially useful for molecular taxonomy of Folonzo and Kiribina.

For decades, patterns of chromosomal inversion polymorphism have guided discovery of population structure and even species boundaries hidden inside otherwise morphologically indistinguishable groups of anopheline mosquitoes i.e., [16,43-46]. Such an association of inversions with population substructure could be an incidental consequence of genetic drift owing to reduced gene flow, or the result of demographic history, but the observation that polymorphic inversions are often clinally distributed with respect to environmental gradients and subject to repeating seasonal fluctuations in frequency suggests that they are subject to strong selective forces [47]. In anopheline mosquitoes, as in many animal and plant species, chromosomal inversions are implicated in local adaptation to environmental heterogeneities [35,48-51]. To the extent that speciation may occur as a by-product of adaptive divergence, chromosomal inversions may also be instrumental in lineage splitting, as proposed by Coluzzi for

anopheline mosquitoes [52]. That Kiribina and Folonzo forms are characterized by alternative arrangements of chromosomal inversions, and that these alternative arrangements shift in relative frequency according to geography, season, and larval habitat availability, suggests a direct role for chromosomal rearrangements in adaptation to heterogeneous and changing environments (see also [35,50]). Thus, beyond simply serving as markers for epidemiologically relevant population structure, alternative chromosomal arrangements somehow condition different physiological and behavioural responses to the environment. A mechanistic understanding of what the adaptations are and how they evolved could prove instrumental in predicting how *An. funestus* may be capable of responding to future environmental challenges, including anthropogenic changes to climate and landscape, and exposure to new means of vector control.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

CC, NFS and NB conceived the study. WG performed the field collections. NFS provided logistical support throughout the study. WG performed the karyotyping. WG, CC and FL analysed results. WG, CC and NB wrote the manuscript. All authors read and approved the final manuscript.

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