Highly evolvable malaria vectors: The genomes of 16 Anopheles mosquitoes

Daniel E. Neafsey,*† Robert M. Waterhouse,* et al.

INTRODUCTION: Control of mosquito vectors has historically proven to be an effective means of eliminating malaria. Human malaria is transmitted only by mosquitoes in the genus Anopheles, but not all species within the genus, or even all members of each vector species, are efficient malaria vectors. Variation in vectorial capacity for human malaria among Anopheles mosquito species is determined by many factors, including behavior, immunity, and life history.

RATIONALE: This variation in vectorial capacity suggests an underlying genetic/genomic plasticity that results in variation of key traits determining vectorial capacity within the genus. Sequencing the genome of Anopheles gambiae, the most important malaria vector in sub-Saharan Africa, has offered numerous insights into how that species became highly specialized to live among and feed upon humans and how susceptibility to mosquito control strategies is determined. Until very recently, similar genomic resources have not existed for other anophelines, limiting comparisons to individual genes or sets of genomic markers with no genome-wide data to investigate attributes associated with vectorial capacity across the genus.

RESULTS: We sequenced and assembled the genomes and transcriptomes of 16 anophelines from Africa, Asia, Europe, and Latin America, spanning ~100 million years of evolution and chosen to represent a range of evolutionary distances from An. gambiae, a variety of geographic locations and ecological conditions, and varying degrees of vectorial capacity. Genome assembly quality reflected DNA template quality and homozygosity. Despite variation in contiguity, the assemblies were remarkably complete and searches for arthropod-wide single-copy orthologs generally revealed few missing genes. Genome annotation supported with RNA sequencing transcriptomes yielded between 10,738 and 16,149 protein-coding genes for each species. Relative to Drosophila, the closest dipteran genus for which equivalent genomic resources exist, Anopheles exhibits a dynamic genomic evolutionary profile. Comparative analyses show a fivefold faster rate of gene gain and loss, elevated gene shuffling on the X chromosome, and more intron losses in Anopheles. Some determinants of vectorial capacity, such as chemosensory genes, do not show elevated turnover but instead diversify through protein-sequence changes. We also document evidence of variation in important reproductive phenotypes, genes controlling immunity to Plasmodium malaria parasites and other microbes, genes encoding cuticular and salivary proteins, and genes conferring metabolic insecticide resistance. This dynamism of anopheline genes and genomes may contribute to their flexible capacity to take advantage of new ecological niches, including adapting to humans as primary hosts.

CONCLUSIONS: Anopheles mosquitoes exhibit a molecular evolutionary profile very distinct from Drosophila, and their genomes harbor strong evidence of functional variation in traits that determine vectorial capacity. These 16 new reference genome assemblies provide a foundation for hypothesis generation and testing to further our understanding of the diverse biological traits that determine vectorial capacity.

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Anopheles gambiae complex

An. gambiae
An. arabiensis
An. quadriannulatus
An. menes
An. mela
An. christyi
An. epirocticus
An. stephensi
An. maculatus
An. culicifacies
An. minimus
An. funestus
An. dirus
An. farauti
An. alatorpus
An. sinensis
An. albimanus
An. darlingi

Culiseta
Nyssorthiscus
Culex quinquefasciatus
Aedes aegypti

Geography, vector status, and molecular phylogeny of the 16 newly sequenced anopheline mosquitoes and selected other diptera.

The maximum likelihood molecular phylogeny of all sequenced anophelines and two mosquito outgroups was constructed from the aligned protein sequences of 1085 single-copy orthologs. Shapes between branch termini and species names indicate vector status and are colored according to geographic ranges depicted on the map. Ma, million years ago.
Highly evolvable malaria vectors: The genomes of 16 Anopheles mosquitoes

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Variation in vectorial capacity for human malaria among Anopheles mosquito species is determined by many factors, including behavior, immunity, and life history. To investigate the genomic basis of vectorial capacity and explore new avenues for vector control, we sequenced the genomes of 16 anopheline mosquito species from diverse locations spanning ~100 million years of evolution. Comparative analyses show faster rates of gene gain and loss, elevated gene shuffling on the X chromosome, and more intron losses, relative to Drosophila. Some determinants of vectorial capacity, such as chemosensory genes, do not show elevated turnover but instead diversify through protein-sequence changes. This dynamism of anopheline genes and genomes may contribute to their flexible capacity to take advantage of new ecological niches, including adapting to humans as primary hosts.

Malaria is a complex disease, mediated by obligate eukaryotic parasites with a life cycle requiring adaption to both vertebrate hosts and mosquito vectors. These relationships create a rich coevolutionary triangle. Just as Plasmodium parasites have adapted to their diverse hosts and vectors, infection by Plasmodium parasites has reciprocally induced adaptive evolutionary responses in humans and other vertebrates (I) and has also influenced mosquito evolution (2). Human malaria is transmitted only by mosquitoes in the genus Anopheles, but not all species within the genus, or even all members of each vector species, are efficient malaria vectors. This suggests an underlying genetic/genomic plasticity that results in variation of key traits determining vectorial capacity within the genus.

In all, five species of Plasmodium have adapted to infect humans and are transmitted by ~60 of the 450 known species of anopheline mosquitoes (3). Sequencing the genome of Anopheles gambiae, the most important malaria vector in sub-Saharan Africa, has offered numerous insights into how that species became highly specialized to live among and feed upon humans and how susceptibility to mosquito control strategies is determined (4). Until very recently (5-7), similar genomic resources have not existed for other anophelines, limiting comparisons to individual genes or sets of genomic markers with no genome-wide data to investigate attributes associated with vectorial capacity across the genus.

Thus, we sequenced and assembled the genomes and transcriptomes of 16 anophelines from Africa, Asia, Europe, and Latin America. We chose these 16 species to represent a range of evolutionary distances from An. gambiae, a variety of geographic locations and ecological conditions, and varying degrees of vectorial capacity (8) (Fig. 1, A and B). For example, An. quadrimaculatus, although extremely closely related to An. gambiae, feeds preferentially on bovines rather than humans, limiting its potential to transmit human malaria. An. merus, An. melas, An. farauti, and An. albimanus females can lay eggs in salty or brackish water, instead of the freshwater sites required by other species. With a focus on species most closely related to An. gambiae (9), the sampled anophelines span the three main subgenera that shared a common ancestor ~100 million years ago (Ma) (10).

Materials and methods summary

Genomic DNA and whole-body RNA were obtained from laboratory colonies and wild-caught specimens (tables S1 and S2), with samples for nine species procured from newly established isofemale colonies to reduce heterozygosity. Illumina sequencing libraries spanning a range of insert sizes were constructed, with ~100-fold paired-end 101-base pair (bp) coverage generated for small (180 bp) and medium (1.5 kb) insert libraries and lower coverage for large (38 kb) insert libraries (table S3). DNA template for the small and medium input libraries was sourced from single female mosquitoes from each species to further reduce heterozygosity. High-molecular-weight DNA template for each large insert library was derived from pooled DNA obtained from several hundred mosquitoes. ALLPATHS-LG (11) genome assemblies were produced using the “haplodbility” option to reduce haplotype assemblies caused by high heterozygosity. Assembly quality reflected DNA template quality and homogygosity, with a mean scaffold N50 of 3.6 Mb, ranging to 18.1 Mb for An. albimanus (table S4). Despite variation in contiguity, the assemblies were remarkably complete and searches for arthropod-wide single-copy orthologs generally revealed few missing genes (fig. S1) (12).

Genome annotation with MAKER (13) supported with RNA sequencing (RNAseq) of transcriptomes (produced from pooled male and female larvae, pupae, and adults) (table S5) and comprehensive noncoding RNA gene prediction (fig. S2) yielded relatively complete gene sets.
Rapidly evolving genes and genomes
Orthology delineation identified lineage-restricted and species-specific genes, as well as ancient genes found across insect taxa, of which universal single-copy orthologs were employed to estimate the molecular species phylogeny (Fig. 1, B and C, and fig. S4). Analysis of codon frequencies in these orthologs revealed that anophelines, unlike drosohils, exhibit relatively uniform codon usage preferences (fig. S5).

Polypene chromosomes have provided a glimpse into anopheline chromosome evolution (14). Our genome-sequence–based view confirmed the cytological observations and offers many new insights. At the base-pair level, ~90% of the non-gapped and nonmasked An. gambiae genome (i.e., excluding transposable elements, as detailed in table S7) is alignable to the most closely related species, whereas only ~13% aligns to the most distant (Fig. 1D, fig. S6, and table S8), with reduced alignability in centromeres and on the X chromosome (Fig. 1D). At chromosomal levels, mapping data assigned 35 to 76% of the An. stephensi, An. funestus, An. atroparvus, and An. albimanus genome assemblies to chromosomal arms (tables S15 to S19). Detailed analysis of unchokep regions showed that symmetry at the whole-arm level is highly conserved, despite several whole-arm translocations (Fig. 2A and table S13). In contrast, small-scale rearrangements disrupt genome cleanliness within arms over time, leading to extensive shuffling of gene order over a time scale of 29 million years or more (10, 15) (Fig. 2B and fig. S7). As in Drosophila, rearrangement rates are higher on the X chromosome than on autosomes (Fig. 2C and tables S14 to S16). However, the difference is significantly more pronounced in Anophelipse, where X chromosome rearrangements are more frequent by a factor of 2.7 than autosomal rearrangements; in Drosophila, the corresponding ratio is only 1.2 (t test, t = 7.3; P < 1 × 10^{-12}) (fig. S8). The X chromosome is also notable for a significant degree of observed gene movement to other chromosomes relative to Drosophila (one-sample proportion test, P = 2.2 × 10^{-5}) (Fig. 2D and tables S17 and S18), as was previously noted for Anophelipsea relative to Aeles (16), further underscoring its distinctive evolutionary profile in Anophelipsea compared with other dipteran genera.

Such dynamic gene shuffling and movement may be facilitated by the multiple families of DNA transposons and long terminal repeat (LTR) and non-LTR retroelements found in all genomes (table S7), as well as a weaker dosage compensation phenotype in Anophelipsea compared with Drosophila. These transposing genes and flanking genomic loci of orthologs can be successfully employed to reconstruct ancestral chromosomal arrangements (fig. S9) and to confidently improve assembly contiguity (tables S19 to S21).

Copy-number variation in homologous gene families also reveals striking evolutionary dynamics. Analysis of 11,636 gene families with CAFE 3 (18) indicates a rate of gene gain/loss higher by a factor of at least 5 than that observed for 12 Drosophila genomes (19). Overall, these Anophelipsea genomes exhibit a rate of gain or loss per gene per million years of 3.12 × 10^{-3} compared with 5.90 × 10^{-4} for Drosophila, suggesting substantially higher gene turnover within anophelines relative to fruit flies. This fivefold greater gain/loss rate in anophelines holds true under models that account for uncertainty in
gene family sizes at the tips of the species tree due to annotation or assembly errors and is not sensitive to inclusion or exclusion of taxa affecting the root age of the tree nor to the exclusion of taxa with the poorest assemblies and gene sets (fig. S10 and tables S22 and S23). Examples include expansions of cuticular proteins in An. arabiensis and neurotransmitter-gated ion channels in An. albimanus (table S24).

The evolutionary dynamism of Anopheles genes extends to their architecture. Comparisons of single-copy orthologs at deeper phylogenetic depths showed losses of introns at the root of the true fly order Diptera and revealed continued losses as the group diversified into the lineages leading to fruit flies and mosquitoes. However, anopheline orthologs have sustained greater intron loss than drosophilids, leading to a relative paucity of introns in the genes of extant anophelines (fig. S11 and table S25). Comparative analysis also revealed that gene fusion and fission played a substantial role in the evolution of mosquito genes, with apparent re-arrangements affecting an average of 10.1% of all genes in the genomes of the 10 species with the most contiguous assemblies (fig. S12).

Because molecular evolution of protein-coding sequences is a well-known source of phenotypic change, we compared evolutionary rates among different functional categories of anopheline orthologs. We quantified evolutionary divergence in terms of protein sequence identity of aligned orthologs and the $d_{K}/d_{S}$ statistic (ratio of nonsynonymous to synonymous substitutions) computed using PAML (22, 20). Among curated sets of genes linked to vectorial capacity or species-specific traits against a background of functional categories defined by Gene Ontology or InterPro annotations, odorant and gustatory receptors show high evolutionary rates and male accessory gland proteins exhibit exceptionally high $d_{K}/d_{S}$ ratios (Fig. 3, figs. S14 and S15, and tables S27 to S29). Rapid divergence in functional categories related to malaria transmission and/or mosquito control strategies led us to examine the genomic basis of several facets of anopheline biology in closer detail.

**Insights into mosquito biology and vectorial capacity**

Mosquito reproductive biology evolves rapidly and presents a compelling target for vector control. This is exemplified by the An. gambiae male accessory gland protein (Acp) cluster on chromosomal 3R (21, 22), where conservation is mostly lost outside the An. gambiae species complex (fig. S16). In Drosophila, male-biased genes such as Acps tend to evolve faster than loci without male-biased expression (23–25). We looked for a similar pattern in anophelines after assessing each gene for sex-biased expression using microarray and RNAseq data sets for An. gambiae (27). In contrast to Drosophila, female-biased genes show dramatically faster rates of evolution across the genus than male-biased genes (Wilcoxon rank sum test, $P = 5 \times 10^{-4}$; fig. S17).

Differences in reproductive genes among anophelines may provide insight into the origin and function of sex-related traits. During
Fig. 2. Patterns of anopheline chromosomal evolution. (A) Anopheline genomes have conserved gene membership on chromosome arms (“elements”; colored and labeled 1 to 5). Unlike Drosophila, chromosome elements reshuffle between chromosomes via translocations as intact elements and do not show fissions or fusions. The tree depicts the supported molecular topology for the species studied. (B) Conserved synteny blocks decay rapidly within chromosomal arms as the phylogenetic distance increases between species. Moving left to right, the dot-plot panels show gene-level synteny between chromosome 2R of An. gambiae (x axis) and inferred ancestral sequences (y axes; inferred using PATHGROUPS) at increasing evolutionary time scales (million years ago) estimated by an ultrametric phylogeny. Gray horizontal lines represent scaffold breaks. Discontinuity of the red lines/dots indicates rearrangement. (C) Anopheline X chromosomes exhibit higher rates of rearrangement (P < 1 × 10^-5), measured as breaks per Mb per million years, compared with autosomes, despite a paucity of polymorphic inversions on the X. (D) The anopheline X chromosome also displays a higher rate of gene movement to other chromosomal arms than any of the autosomes. Chromosomal elements are labeled around the perimeter; internal bands are colored according to the chromosomal element source and match element colors in (A) and (C). Bands are sized to indicate the relative ratio of genes imported versus exported for each chromosomal element and the relative allocation of exported genes to other elements.

copulation, An. gambiae males transfer a gelatinous mating plug, a complex of seminal proteins, lipids, and hormones that are essential for successful sperm storage by females and for reproductive success (26–28). Coagulation of the plug is mediated by a seminal transglutaminase (TG3), which is found in anophelines but is absent in other mosquito genera that do not form a mating plug (26). We examined TG3 and its two paralogs (TG1 and TG2) in the sequenced anophelines and investigated the rate of evolution of each gene (Fig. 4A). Silent sites were saturated at the whole-genus level, making dS difficult to estimate reliably, but TG1 (the gene presumed to be ancestral owing to broadest taxonomic representation) exhibited the lowest rate of amino acid change (dS = 0.20), TG2 exhibited an intermediate rate (dS = 0.93), and the anopheline-specific TG3 has evolved even more rapidly (dS = 1.50), perhaps because of male/male or male/female evolutionary conflict. Note that plug formation appears to be a derived trait within anophelines, because it is not exhibited by An. albimanus and intermediate, poorly coagulated plugs were observed in taxa descending from early-branching lineages within the genus (table S30). Functional studies of mating plugs revealed identical spatial expression patterns for CLPCW and CPLCG group A gene pairs suggestive of coregulation (fig. S19). For these five gene clusters, complete grouping by organizational lineage was observed for most deep nodes as well as for many individual species outside the shallow An. gambiae species complex (Fig. 4B), consistent with a relatively rapid (less than 20 million years) homogenization of sequences via concerted evolution. The emerging pattern of anopheline CP evolution is thus one of relative stasis for a majority of single-copy orthologs, juxtaposed with consistent concerted evolution of a subset of genes.

Anophelines identify hosts, oviposition sites, and other environmental cues through specialized chemosensory membrane-bound receptors. We examined three of the major gene families that encode these molecules: the odorant receptors (ORs), gustatory receptors (GRs), and variant ionotropic glutamate receptors (iGRs). Given the rapid chemosensory gene turnover observed in many other insects, we explored whether varying host preferences of anopheline mosquitoes could be attributed to chemosensory gene gains and losses. Unexpectedly in light of the elevated genome-wide rate of gene turnover, we found that the overall size and content of the chemosensory
gene repertoire are relatively conserved across the genus. CAFE 3 (38) analyses estimated that the most recent common ancestor of the anophelines had approximately 60 genes in each of the OR and GR families, similar to most extant anophelines (Fig. 4C and fig. S20). Estimated gain/loss rates of OR and GR genes per million years (error-corrected $\lambda = 1.5 \times 10^{-8}$ for ORs and $2.0 \times 10^{-4}$ for GRs) were much lower than the overall level of anopheline gene families. Similarly, we found almost the same number of antennae-expressed IRs (~20) in all anopheline genomes. Despite overall conservation in chemosensory gene numbers, we observed several examples of gene gain and loss in specific lineages. Notably, there was a net gain of at least 12 ORs in the common ancestor of the *An. gambiae* complex (Fig. 4C).

OR and GR gene repertoire stability may derive from their roles in several critical behaviors. Host preference differences are likely to be governed by a combination of functional divergence and transcriptional modulation of orthologs. This model is supported by studies of antennal transcriptomes in the major malaria vector *An. gambiae* (35) and comparisons between this vector and its morphologically identical sibling *An. quadriannulatus* (36), a very closely related species that plays no role in malaria transmission (despite vectorial competence) because it does not specialize on human hosts. Furthermore, we found that many subfamilies of ORs and GRs showed evidence of positive selection (19 of 53 ORs; 17 of 59 GRs) across the genus, suggesting potential functional divergence.

Several blood-feeding–related behaviors in mosquitoes are also regulated by peptide hormones (37). These peptides are synthesized, processed, and released from nervous and endocrine systems and elicit their effects through binding appropriate receptors in target tissues (38). In total, 39 peptide hormones were identified from each of the sequenced anophelines (fig. S21). Notably, no ortholog of the well-characterized head peptide (HP) hormone of the culicine mosquito *Aedes aegypti* was identified in any of the assemblies. In *Ae. aegypti*, HP is responsible for inhibiting host-seeking behavior after a blood meal (39). Because anophelines broadly exhibit similar behavior (40), the absence of HP from the entire clade suggests they may have evolved a novel mechanism to inhibit excess blood feeding. Similarly, no ortholog of insulin growth factor 1 (IGF1) was identified in any anophelines even though IGF1 orthologs have been identified in other dipterans, including *D. melanogaster* (41) and *Ae. aegypti* (42). IGF1 is a key component of the insulin/insulin growth factor 1 signaling (IIS) cascade, which regulates processes including innate immunity, reproduction, metabolism, and life span (43). Nevertheless, other members of the IIS cascade are present, and four insulin-like peptides are found in a compact cluster with gene arrangements conserved across anophelines (fig. S22). This raises questions regarding the modification of IIS signaling in the absence of IGF1 and the functional importance of this conserved genomic arrangement.

Epigenetic mechanisms affect many biological processes by modulation of chromatin structure, telomere remodeling, and transcriptional control. Of the 215 epigenetic regulatory genes in *D. melanogaster* (44), we identified 169 putative *An. gambiae* orthologs (table S32), which suggested the presence of mechanisms of epigenetic control in *Anopheles* and *Drosophila*. We find, however, that retrotransposition may have contributed to the functional divergence of at least one gene associated with epigenetic regulation. The ubiquitin-conjugating enzyme E2D (orthologous to effete (45) in *D. melanogaster*) duplicated via retrotransposition in an early anopheline ancestor, and the retrotransposed copy is maintained in a subset of anophelines. Although the entire amino acid sequence of E2D is perfectly conserved between *An. gambiae* and *D. melanogaster*, the retrogenes are highly divergent (Fig. 5A) and may contribute to functional diversification within the genus.

Saliva is integral to blood feeding; it impairs host hemostasis and also affects inflammation and immunity. In *An. gambiae*, the salivary proteome is estimated to contain the products of at least 75 genes, most being expressed solely in the adult female salivary glands. Comparative analyses indicate that anophele salivary proteins are subject to strong evolutionary pressures, and these genes exhibit an accelerated pace of evolution, as well as a very high rate of gain/loss (Fig. 3 and fig. S23). Polymorphisms within *An. gambiae* populations from limited sets of salivary genes were previously found to carry signatures of positive selection (46). Sequence analysis across the anophelines shows that salivary genes have the highest incidence of positively selected codons among the seven gene classes (fig. S24), indicating that coevolution with vertebrate hosts is a powerful driver of natural selection in salivary proteomes. Moreover, salivary proteins also exhibit functional diversification through new gene creation. Sequence similarity, intron-exon boundaries, and secondary structure prediction point to the birth of the *SG7/SG7-2* inflammation-inhibiting (47) gene family from the genomic region encoding the C terminus of the 30-kD protein (Fig. 5B), a collagen-binding platelet inhibitor already present in the blood-feeding ancestor of mosquitoes and black flies (48). Based on phylogenetic representation, these events must have occurred before the radiation of anophelines but after separation from the culicines.

Resistance to insecticides and other xenobiotics has arisen independently in many anopheline species, fostered directly and indirectly by anthropogenic environmental modification. Metabolic resistance to insecticides is mediated by...
Fig. 4. Phylogeny-based insights into anopheline biology. (A) Maximum-likelihood amino acid–based phylogenetic tree of three transglutaminase enzymes (TG1, green; TG2, yellow; and TG3, red) in 14 anopheline species with Culex quinquefasciatus (Cxqu), Aedes aegypti (Aeae), and D. melanogaster (Dmel) serving as outgroups. TG3 is the enzyme responsible for the formation of the male mating plug in An. gambiae, acting upon the substrate Plugin, the most abundant mating plug protein. Higher rates of evolution for plug-forming TG3 are supported by elevated levels of $d_n$. Mating plug phenotypes are noted where known within the TG3 clade. (B) Concerted evolution in CPFL cuticular proteins. Species symbols used are the same as in (A). In contrast to the TG1/TG2/TG3 phylogeny, CPFL paralogs cluster by subgeneric clades rather than individually recapitulating the species phylogeny. Gene family size variation among species may reflect both gene gain/loss and variation in gene set completeness. (C) OR observed gene counts and inferred ancestral gene counts on an ultrametric phylogeny. At least 10 OR genes were gained on the branch leading to the common ancestor of the An. gambiae species complex, although the overall number of OR genes does not vary dramatically across the genus.

Having multiple gene families, including cytochrome P450s and glutathione S-transferases (GSTs), which serve to generally protect against all environment stresses, both natural and anthropogenic. We manually characterized these gene families in seven anophelines spanning the genus. Despite their large size, gene numbers (87 to 104 PER genes, 27 to 30 GST genes) within both gene families are highly conserved across all species, although lineage-specific gene duplications and losses are often seen (tables S33 and S34). As with the OR and GR dafaction-related gene families, P450 and GST repertoires may be relatively constant due to the large number of roles they play in anopheline biology. Orthologs of genes associated with insecticide resistance either via up-regulation or coding variation (e.g., Cyp6m2, Cyp6o9 in An. funestus, Gste2, and Gste4) were found in all species, which suggested that virtually all anophelines likely have genes capable of conferring insecticide resistance through similar mechanisms. Unexpectedly, one member of the P450 family (Cyp18a1) with a conserved role in ecyclodysteroid catabolism [and consequently development and metamorphosis (49)] appears to have been lost from the ancestor of the An. gambiae species complex but is found in the genome and transcriptome assemblages of other species, which indicates that the An. gambiae complex may have recently evolved an alternate mechanism for catabolizing ecyclodysterone.

Susceptibility to malaria parasites is a key determinant of vectorial capacity. Dissecting the immune repertoire (50, 51) (table S35) into its constituent phases reveals that classical recognition genes and genes encoding effector enzymes exhibit relatively low levels of sequence divergence. Signal transducers are more divergent in sequence but are conserved in representation across species and rarely duplicated. Cascade modulators, although also divergent, are more lineage-specific and generally have more gene duplications (Fig. 3 and fig. S25). A rare duplication of an immune signal transduction gene occurred through the retrotransposition of the signal transducer and activator of transcription STAT2 to form the intronless STAT1 in the An. funestus lineage. In An. gambiae, STAT1 controls the expression of STAT2 and is activated in response to bacterial challenge (52, 53), and the STAT pathway has been demonstrated to mediate immunity to Plasmodium (53, 54), so the presence of these relatively new immune signal transducers may have allowed for rewiring of regulatory networks governing immune responses in this subset of anophelines.

Conclusion

Since the discovery over a century ago by Ronald Ross and Giovanni Battista Grassi that human malaria is transmitted by a narrow range of blood-feeding female mosquitoes, the biological basis of malarial vectorial capacity has been a matter of intense interest. Inasmuch as previous successes in the local elimination of malaria have always been accomplished wholly or in part through effective vector control, an increased understanding of vector biology is crucial for continued progress against malarial disease. These 16 new reference genome assemblies provide a...
foundation for additional hypothesis generation and testing to further our understanding of the diverse biological traits that determine vectorial capacity.

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SUPPLEMENTAL MATERIALS

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Materials and Methods

Supplementary Text Figs. S1 to S25

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