# Short Report

## The bionomics of Anopheles funestus and its role in malaria transmission in a forested area of southern Cameroon

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The earliest studies on the bionomics of Anopheles in Cameroon were carried out some 40 years ago (ADAM, 1956; MOUCHET & GARIOU, 1961; MOUCHET, 1962) These studies showed that A. nili and A. moucheti were the principal malaria vectors in the rural forested areas of the country. Although A. gambiae s.l. was present, it had a minor role in malaria transmission in those areas. Very low densities of A. funestus were reported (LI-VADAS et al., 1958), indicating the limited epidemiological importance of this vector; hence, its bionomics in the forested zone were not well known. Recent studies on malaria vectors and transmission dynamics in the rural areas of southern Cameroon have confirmed these observations (CARNEVALE et al., 1992; NJAN NLOGA et al., 1993; MANGA et al., 1995). The discovery of high densities of A. funestus in some villages in the forested zone made it possible for the bionomics of this species to be clarified, and its involvement in malaria transmission in those villages to be studied.

Etoa, Simbock and Nsimalen are villages near Yaoundé, the capital city of Cameroon. They are irrigated by 2 rivers, Biyeme and Mefou, which flow through a network of swamps. The swampy areas of the villages are separated by about 100 m. The vegetation is semideciduous equatorial forest, and the climate is guinean with 2 rainy seasons and 2 dry seasons per year; the annual rainfall is about 1700 mm.

Mosquito larvae were searched for along the Biyeme and Mefou rivers banks, in the swamps, and in pools. Anopheles larvae collected were reared separately to the adult stage and then identified. Indoor, all-night catches on human bait were made at Etoa twice a month for 4 months, from February to May 1996 (from the end of the long dry season to the end of the short rainy season). The Anopheles mosquitoes collected were identified and dissected to determine their physiological age and to search for Plasmodium sporozoites. The gonotrophic cycle was studied in June 1996, during the short rainy season, and its duration was determined using the following mark-recapture technique. For 3 consecutive nights, blood-fed mosquitoes were caught on human bait, marked with fluorescent powder (a different colour was used each night), and then released inside the dwellings. The duration of stage I (the period between egg laying and taking the next blood meal) was evaluat-ed by examining follicle relics. The length of stage II (from blood meal to maturation of the oocytes) was determined by performing a series of delayed dissections; the oocyte stages were classified according to CHRISTO-PHERS (1911)

At Etoa, A. funestus larvae were collected from the small grassy bays in the Biyeme and Mefou rivers. Of 78 adults that were identified, 38 (49%) were A. funestus; the other species present were A. nili and A. moucheti. At Simbock, A. funestus were collected in smaller numbers: 4 specimens (6.5%) were collected from the Mefou river, again with A. moucheti and A. pili. A. funestus was



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also collected from the swamps together with A. obscurus and A. namibiensis. This is the first record of A. namibiensis from Cameroon. At Nsimalen, A. funestus was found in the Mefou river, once again together with A. moucheti and A. nili, mainly (in large numbers) in a weedy pool with A. namibiensis. A. funestus bred in a va-riety of sites: rivers (like A. nili and A. mouchen), swamps and pools. The presence of vegetation (not always erect) seemed to be the common factor in all these breeding sites

A total of 913 mosquitoes was caught indoors at night at Etoa. Anopheles spp. formed 97.3% of this sample, and A. funestus constituted 71.2% of the Anopheles col-lected. The other Anopheles species were A. nili, A. moucheti and A. gambiae s.l. The average biting rate was 31.7 bites per person per night for all species of Anopheles, with an average of  $22 \cdot 6$  for A. funestus.

A. funestus had a parity rate 83.1% (among 243 dissections) and its sporozoite index was 3.27% (336 dissections). The average entomological inoculation rate was 1.3 infective bites per person per night, representing 41 infective bites per person per month. A. funestus was responsible for 55% of these bites (0.7/night), while the 3 other species shared the remaining 45%. These findings led to the conclusion that A. funestus could be considered as a major local vector of malaria in forested areas, together with A. nili and A. moucheti.

A total of 158 A. funestus was marked and released, and 2 were recaptured the second night after release (a recapture rate of 1.3%). Thus, the duration of the gonotrophic cycle could be about 48 h. Dissection of 36 unfed female mosquitoes showed that 48% of them still had open follicle sacs, 36% had partially contracted follicle sacs, and only 17% had their follicle sacs completely contracted.

These findings indicated that most females took their next blood meal a few hours after egg laying. Twelve hours after the blood meal, 7 of 12 A. funestus had oocytes in stage III; at 24 h, 6 of 25 mosquitoes had oocytes at the beginning of stage IV; at 36 h, 60% (of a total of 33) had oocytes at stage V; and at 40 h, the maturation of the ovaries was complete: all 17 mosquitoes examined were at stage V. The duration of the gonotrophic cycle of A. funestus was therefore similar in the forested zone and in the savannah (BRENGUES & COZ, 1973).

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