

Literature Cited

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COMPARATIVE INFECTIVITY OF *PLASMODIUM FALCIPARUM* (COLOMBIA STRAIN) TO *ANOPHELES QUADRIMACULATUS* SAY AND *ANOPHELES ALBIMANUS* (WIED.).

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INTRODUCTION. Studies on the comparative susceptibility of *Anopheles quadrimaculatus* and *A. albimanus* to *Plasmodium falciparum* have been made previously by several workers. Boyd *et al.* (1938) and Reid and Jobbins (1940) found that *A. albimanus* from Cuba and Panama were highly susceptible to *P. falciparum* from their own region, but distinctly refractory to strains of the same parasite species in the Nearctic region. *A. quadrimaculatus* mosquitoes from Florida exhibited a high degree of susceptibility to infection by strains of *P. falciparum* from both the tropical and the Nearctic regions. Jones and Young (1950), in a similar study using a South Carolina strain of *P. falciparum*, a Panama strain of *A. albimanus*, and the Q-1 strain of *A. quadrimaculatus*, found that the latter species is more susceptible to infection. Jeffery *et al.* (1950) showed that *A. albimanus* from Panama (A-2 strain) was markedly superior to *A. albimanus* from the Florida strains (A-3 strain) and the Q-1 strain in susceptibility to infection with a Panama strain of *P. falciparum*. Because of

these previous reported differences, studies were made on the comparative infectivity of a Colombia, South America, strain of *P. falciparum* to the Q-1 strain of *A. quadrimaculatus* and a Central American strain of *A. albimanus*.

METHODS AND PROCEDURES. The Colombia strain of *Plasmodium falciparum* is a chloroquine-resistant strain described by Young and Moore (1961) and was originally from the Magdalena Valley of Colombia, South America.

The *A. quadrimaculatus* (Q-1 strain) mosquitoes were originally from the southeastern United States and have been maintained in the laboratory since 1941. The *A. albimanus* (A-9 strain) mosquitoes were originally from El Salvador and were obtained through the courtesy of Dr. H. G. Simkover, Shell Development Company, Modesto, California. The colony has been maintained since 1960.

The patients were adult Negro males being treated for neurosyphilis. Patients A and B were infected by intravenous inoculation of parasitized blood which had been stored frozen in a dry ice chest

(-78° C.) for 346 and 384 days respectively. In patient A, parasites were first demonstrated 6 days post inoculation and gametocytes 12 days later. In patient B, parasites appeared on day 13 and gametocytes 16 days later. Patient A received no antimalarial drugs prior to or during the feeding period whereas patient B received 10 grains of quinine 12, 10, 9, and 8 days prior to the appearance of gametocytes. No other antimalarials were given during the study period.

Three- to five-day old adult female mosquitoes were caged in lots of 100 in pint ice cream carton cages and allowed to feed, through the screened top, on the patient's leg. Engorged mosquitoes were incubated in these cages at 78° to 80° F. and fed 5 percent honey water daily in a cellulose sponge. Eight to 10 days after feeding, the mosquitoes were dissected and the guts microscopically examined for the presence of oocysts.

The feedings were designed to study

(1) the initial period when the *A. quadrimaculatus* mosquitoes could be infected and (2) the comparative susceptibility of the *A. quadrimaculatus* (Q-1) and *albimanus* (A-9) mosquitoes to infection by the Colombia strain of *P. falciparum*.

RESULTS. The results of initial infectivity feedings using *A. quadrimaculatus* are shown in Table 1. Patient A demonstrated a rapid increase in gametocyte density reaching a peak of 2790 per cmm on day 5 after which the number slowly declined. Patient B demonstrated a slower rate of increase reaching a peak of 1110 per cmm on day 8 followed by a slow decrease in density. The mosquitoes were first infected on gametocyte day 1 with patient A (gametocyte count of 10 per cmm) and gametocyte day 5 with patient B (gametocyte count of 200 per cmm). Both patients were infectious 1 day after reaching a gametocyte count greater than 100 per cmm.

The results of eleven comparative feedings

TABLE 1.—Infectivity of *Plasmodium falciparum* (Colombia strain) to *Anopheles quadrimaculatus* (Q-1 strain) and *A. albimanus* (A-9 strain)

Patient	Day of gametocytemia	Gametocytes per cmm	No. mosq. dissected		Percent infection		Infection ratio	Gut infection index		Gut infectivity index
			Q-1	A-9	Q-1	A-9		Q-1/A-9	Q-1	
A	1	30	25	..	0	0
	2	110	25	..	0	0
	3	680	25	..	52	132
	4	1640	25	..	48	220
	5	2790	25	25	56	8	7.0	448	44	10.2
	6	2697	25	25	80	20	4.0	1984	116	17.1
	7	2697	25	25	72	36	2.0	792	688	1.1
	8	2536	25	25	88	36	2.4	1892	1244	1.5
	9	2247	25	25	44	12	3.7	636	160	4.0
B	1	10	25	..	0	0
	2	40	25	..	0	0
	3	30	25	..	0	0
	4	140	25	..	0	0
	5	200	25	..	16	44
	6	450	25	19	72	0	∞	388	0	∞
	7	810	25	..	64	433
	8	1110	25	15	56	0	∞	220	0	∞
	9	940	25	22	60	32	1.9	312	182	1.7
	10	1100	35	25	89	12	7.4	1174	56	21.0
	12	850	25	25	32	0	∞	164	0	∞
	13	900	25	25	56	8	7.0	132	16	8.2
	* Totals			285	256	65	16	4.2	755	243

* Totals are given for comparative feedings only.

ings are also shown in Table 1. The infection rates with the *A. quadrimaculatus* ranged from 44 to 88 percent with patient A and 32 to 89 percent with patient B. In contrast, the infection rates with *A. albimanus* ranged from 8 to 36 percent with patient A and 0 to 32 percent with patient B. In the total of 11 feedings, the *A. quadrimaculatus* had an infection rate of 4.2 times that of the *A. albimanus*.

The intensity of infection, as measured by the average number of oocysts per gut 100 (gut infection index) was in all instances higher for the *A. quadrimaculatus* than comparable lots of *A. albimanus*. The ratios showed a wide variation but the average gut infection index ratio was 1.

These results indicate that with the Colombia strain of *P. falciparum*, the Q-1 strain of *A. quadrimaculatus* is more susceptible to infection than is a strain of *A. albimanus* from El Salvador (A-9). In this case, neither of the mosquitoes was coindigenous with the strain of *P. falciparum*. This adds support to reports in the literature that *A. quadrimaculatus* is susceptible to infection by *P. falciparum* from widely separated geographical areas.

The *A. albimanus* (A-9 strain), although not refractory to the Colombia strain of *P. falciparum*, did not demonstrate the marked susceptibility that would be expected if the two were coindigenous and they are therefore considered as belonging to separate distributional populations.

The susceptibility relationship between *A. albimanus* and *P. falciparum* appears to be markedly affected by the geographical origin of the strains of parasite and vector. Thus, this vector does not seem ideally suited to infectivity studies on new strains of *P. falciparum*.

These results and previous studies strongly suggest the desirability of using established strains of mosquitoes in infectivity studies involving new strains of *P. falciparum*. The most desirable mosquito would be one which not only had been used in comparable studies, but which showed a susceptibility to infection by non-coindigenous strains of malaria,

thus making it useful for the study of malarias from widely separated geographical areas. It appears that the Q-1 strain of *A. quadrimaculatus*, which was used by Eyles and Young (1950), Jeffrey *et al.* (1950) and in the present study as well as having had wide use in other related studies, is one of the mosquito strains which could be considered as a standard for comparison.

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SUMMARY. The Q-1 strain of *Anopheles quadrimaculatus* proved to be more susceptible than an El Salvador strain of *A. albimanus*, to a Colombia strain of *Plasmodium falciparum*, when the two species of mosquito were fed simultaneously. The gut infection index in *A. quadrimaculatus* was 3.1 times that in *A. albimanus*. The average infection rate in *A. quadrimaculatus* was 4.2 times that in the *A. albimanus*.

Initial infection of *A. quadrimaculatus* occurred one day after the gametocyte count exceeded 100 per cmm in each of the two patients studied.

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