STUDIES ON THE CHEMOTHERAPY OF THE HUMAN MALARIAS.
X. THE SUPPRESSIVE ANTIMALARIAL EFFECT
OF PALUDRINE^{1, 2}

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Paludrine, N₁-p-chlorophenyl-N₃-isopropylbiguanide acetate, was found by British workers to be the most active antimalarial of a series of synthetic guanidines and biguanidines (1, 2). It possesses high activity against the erythrocytic asexual parasites of both vivax and falciparum malaria and is practically non-toxic at therapeutic doses (3, 4). Previous study of the compound has included extensive prophylactic, curative and suppressive trials by the Australian Malarial Unit at Cairns in experimental sporozoite and trophozoite induced vivax and falciparum malaria (5) and curative trials in naturally acquired relapsing Southwest Pacific vivax malaria (6). The drug has a prophylactic action in falciparum malaria which is complete in single doses as low as 25 mgm. The prophylactic action in vivax malaria is less definite, although the incubation periods of patients in the Cairns studies were systematically prolonged, and subinoculations from infected patients showed that this was due to an effect upon the initial tissue phase (*cf.* 7) of the disease. It is also effective in the treatment of clinical attacks of malaria, being curative in falciparum but not in vivax malaria. The response of fever and symptoms in each infection is somewhat less prompt

than that obtained with high dosage of quinacrine or chloroquine (8).

The present paper is a report on the assay of the suppressive antimalarial activity of paludrine in standardized blood-induced malarial infections. The studies were designed to yield comparative data on the suppressive antimalarial activity of paludrine and other antimalarials now coming into general use.

MATERIALS AND METHODS

The therapeutic trials were performed in patients⁴ with blood-induced malaria in accordance with standard procedures previously outlined (7, 9). The infections utilized were due to the McCoy and Chesson strains of *P. vivax* and a relatively quinine-resistant strain of *P. falciparum* (Costa). The therapeutic results due to the administration of paludrine are classified in three groups: Class I, no certain effect; Class II, temporary suppression of parasitemia and/or fever; and Class III, a "permanent" effect, *i.e.*, absence of parasitemia for 14 days in McCoy vivax, or 21 days in Chesson vivax and Costa falciparum, followed by a positive reinoculation to demonstrate continuing host-susceptibility to the infection.

Paludrine⁵ was administered by mouth in each instance, and all doses are reported in terms of the free base. Plasma paludrine levels⁶ were estimated by a method that involves coupling of the compound with methyl orange to yield an organic-soluble colored complex (10). The method permits the valid estimation of

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⁵ Paludrine was generously furnished by The Imperial Chemical Industries, Ltd.

⁶ In many instances, especially, at the lower doses, when only one to four individual doses were used, a stable plasma drug level could not be maintained for the usual four or six days.

plasma paludrine levels as low as 10 $\mu\text{g.}$ per liter. However, considerably lower plasma paludrine concentrations resulted from some of the dosage regimens examined; these were not estimated. A more sensitive chemical method is now available (11).

RESULTS

Paludrine in varying dosage was administered to 28 white subjects with blood-induced McCoy strain vivax malaria (Table I). It was soon apparent that the drug is extremely effective against the erythrocytic phase of this strain of malaria. A permanent eradication of parasites, or Class III effect, was noted in all patients with a four-day mean plasma paludrine concentration of 10 or more $\mu\text{g.}$ per liter. Total doses of paludrine as low as 50 mgm. were sufficient to produce Class III effects with regularity. The lowest total dose investigated, 12.5 mgm., was given to ten patients. Even this minute amount of drug had a dramatic effect on the course of the disease, Class III effects being noted in three instances, while complete

TABLE I

The relationship between dosage and plasma concentration of paludrine and therapeutic effect in four-day tests against blood-induced McCoy vivax malaria

Patient	Total dose	Mean plasma paludrine concentration	Class of therapeutic effect	
			II	III
	mgm.	$\mu\text{g./L}$		
Bro	225	56		x
Con	225	51		x
Wit	238	48		x
Pow	150	38		x
Tay	150	30		x
Che	150	29		x
Bur	225	23		x
Jor	150	22		x
Sou	150	22		x
Fla	100	21		x
Ham	50	20		x
O'Br	50	17		x
Qui	100	17		x
Mor	50	13		x
Mar	100	10		x
Kli	25	—	x	
Cha	25	—	x	
Ott	25	—	x	
Tha	12.5	—		x
Fit	12.5	—	x	
McL	12.5	—	x	
Dic	12.5	—		x
Pur	12.5	—	x	
Lev	12.5	—	x	
Spe	12.5	—	x	
McI	12.5	—		x
Bre	12.5	—	x	
Moo	12.5	—	x	

TABLE II

The relationship between dosage and plasma concentration of paludrine and therapeutic effect in four-day tests against blood-induced Chesson vivax malaria

Patient	Total dose	Mean plasma paludrine concentration	Class of therapeutic effect	
			II	III
	mgm.	$\mu\text{g./L}$		
Mer	4000	688		x
Mur	4000	464		x
O'Be	500	70		x
Mon	500	59		x
Lyo	500	49		x
Mur	225	40		x
Pet	150	34		x
Pad	150	39		x
Gol	150	28		x
You	225	17		x
Lem	100	—		x
Sca	100	—	x	
Lev	25	—	x	
Sny	25	—		x
Sig	25	—	x	
Smi	25	—		x

temporary, or Class II, effects occurred in the other seven patients.

The effect of paludrine against the Chesson strain of vivax malaria was examined in 16 patients (Table II). This strain had previously been demonstrated to be more resistant than the McCoy strain of vivax to the suppressive anti-malarial action of quinine, quinacrine, and chloroquine (7, 12, 13). Total doses of 150 mgm. or more of paludrine resulted in Class III effects. One of two patients given a total dose of 100 mgm. had a Class III effect, the other, a Class II effect. At the lowest total dose studied, 25 mgm., two patients had Class III effects, and two had Class II effects. It is likely that paludrine is only slightly less effective against the Chesson strain than against the McCoy strain.

In contrast, the quinine-resistant Costa strain of falciparum malaria was also relatively resistant to the suppressive action of paludrine (Table III). Total doses ranging from 338 to 750 mgm. were administered to nine patients. The six-day mean plasma paludrine levels varied from 55 to 106 $\mu\text{g.}$ per liter. Class III or permanent effects were noted in two patients with mean plasma drug levels of 78 and 95 $\mu\text{g.}$ per liter, the remainder of the patients having Class II or complete temporary effects, the parasite-free intervals ranging from two to 14 days.

TABLE III

The relationship between dosage and plasma concentration of paludrine and therapeutic effect in six-day tests against blood-induced Costa falciparum malaria

Patient	Total dose	Mean plasma paludrine concentration	Class of therapeutic effect	
			II	III
	<i>mgm.</i>	<i>μg./L</i>		
McC	500	106	x	
O'Ne	750	103	x	
Sca	750	102	x	
Dav	750	95		x
Gib	500	82	x	
Wil	550	78		x
Sha	500	74	x	
Vel	338	68	x	
Ros	500	55	x	

In no instance of treatment with paludrine was an adverse side reaction noted that could be ascribed to the drug.

DISCUSSION

The doses and plasma drug levels of paludrine, quinacrine (12) and chloroquine (13) required to achieve Class III effects in McCoy and Chesson strain vivax malaria and Costa strain falciparum are summarized in Table IV.

It is apparent that paludrine has extraordinary effectiveness against the erythrocytic phase of vivax malaria. The peripheral trophozoites of two strains of vivax can be permanently eradicated with oral doses much smaller than are necessary with other antimalarials such as quinacrine or chloroquine. Permanent eradication of the peripheral trophozoites of Costa falciparum was not regularly achieved at the dosage tested. It should be noted, however, that the dosage used was far

TABLE IV

Total doses and mean plasma drug levels of paludrine, quinacrine and chloroquine required to achieve Class III effects in two strains of vivax malaria and one strain of falciparum malaria

	Paludrine		Quinacrine		Chloroquine	
	Total dose	Plasma level	Total dose	Plasma level	Total dose	Plasma level
	<i>mgm.</i>	<i>μg./L</i>	<i>mgm.</i>	<i>μg./L</i>	<i>mgm.</i>	<i>μg./L</i>
Vivax, McCoy strain	50	10*	700	25*	300	10*
Vivax, Chesson strain	150	17*	1100	25†		
Falciparum, Costa strain	>750	>100†	>1100	65†	650	30†

* 4 day mean plasma drug level.

† 6 day mean plasma drug level.

below the toxic dose (5, 6) and the dose recommended for treatment of clinical attacks of malaria (6).

The study of paludrine was undertaken at too late a date in the malaria program to collect definitive data on the relation between oral dosage, plasma paludrine levels, and antimalarial effects in either vivax or falciparum malarias. As far as falciparum malaria was concerned, there was no apparent correlation between effect and drug levels in the range of 55 to 106 $\mu\text{g.}$ per liter.

The observations upon which these summarizing statements are based do not define the potentialities of paludrine in the routine suppression and treatment of malaria. It seems likely that routine suppression in falciparum malaria is accomplished by a combination of the very high order of the drug's prophylactic action and the moderate order of its suppressive action; whereas, in vivax malaria, it is accomplished by the high order of its suppressive action, perhaps aided by a minor degree of prophylactic action. Routine treatment of the clinical attack will usually involve the administration of high dosage as compared to the dosage utilized in these studies. The effectiveness of such high dosage is best illustrated by the experimental work of others (3, 4, 5, 6).

The high order of antimalarial activity shown by paludrine against more than a single phase of the malarias, *i.e.*, primary tissue phase of falciparum and erythrocytic phases of vivax and falciparum, places the drug in a unique position among the synthetic antimalarials developed in recent years.

SUMMARY AND CONCLUSIONS

1. Paludrine is the most active suppressive agent in vivax malaria yet described, exceeding quinacrine or chloroquine to a considerable extent in this respect.

2. It is less active as a suppressive in falciparum malaria, routine suppression at low dosage being due presumably to its high order of prophylactic action in this infection.

3. Paludrine is apparently non-toxic in therapeutic doses.

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