Artemisinin Pharmacokinetics and Efficacy in Uncomplicated-Malaria Patients Treated with Two Different Dosage Regimens

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The immediate efficacies of two oral dosage regimens of artemisinin were investigated in 77 male and female adult Vietnamese falciparum malaria patients randomly assigned to treatment with either 500 mg of artemisinin daily for 5 days (group A; \( n = 40 \)) or artemisinin at a dose of 100 mg per day for 2 days, with the dose increased to 250 mg per day for 2 consecutive days and with a final dose of 500 mg on the fifth day (group B; \( n = 37 \)). Parasitemia was monitored every 4 h. The average parasite clearance time was longer in group B than in group A (means ± standard deviations, 50 ± 23 and 34 ± 14 h, respectively; \( P < 0.01 \)). Artemisinin concentrations in saliva samples obtained on days 1 and 5 were quantified by high-performance liquid chromatography. The average oral clearance, based on saliva drug concentrations in group B patients, was twofold higher than that in group A patients on day 1 (\( P < 0.01 \)), with no differences in drug half-lives (\( P = 0.40 \)), indicating a saturable first-pass metabolism. Female patients had higher oral clearance values on day 1. Artemisinin’s pharmacokinetic parameters were similar on day 5 in both groups, although a significant increase in oral clearance from day 1 to day 5 was evident. Thus, artemisinin exhibited both dose- and time-dependent pharmacokinetics. The escalating dose studied did not result in higher artemisinin concentrations toward the end of the treatment period.

Artemisinin and its semisynthetic family of derivatives are the most potent antimalarials available for treatment of falciparum malaria infections (15, 25, 26). Artemisinin has a short elimination half-life, a rapid onset of action, no major side effects, and markedly induces its own metabolic elimination, resulting in five- to sevenfold decreases in its concentrations over 5 to 7 days of administration in both malaria patients and healthy subjects (1, 3–5). A major drawback with artemisinin and its derivatives is the high recrudescence rates within 2 to 3 weeks after monotherapy (14, 17). It has proved difficult in field studies to determine whether the relapsed malaria episodes are due to new infections or the persistence of the previous one. However, true recrudescence has been described in patients kept in vector-free environments for longer periods of time (12, 13, 16). There is also evidence of true recrudescence, as determined by a PCR technique with patients treated with a combination of artesunate and benflumetol (10) as well as artesunate alone (18). It has been suggested that the occurrence of recrudescence may partly be due to the decreasing artemisinin concentrations toward the end of treatment (4). Thus, the time-dependent kinetics of the drug can be of importance in dosing suggestions.

Reports concerning dose-response studies with the artemisinin endoperoxides are lacking. Rectal administration of artemisinin was shown to have efficacy comparable to that of oral treatment in terms of the parasite clearance time (PCT) and the fever subsidence time (FST) in patients with uncomplicated falciparum malaria (4). This was despite the observation that artemisinin areas under the concentration-time curves (AUCs) after rectal administration were in general only 30% of those after oral intake. This raises doubts about how justified the standard dosage regimens have been and whether a 500-mg daily dose may be unnecessarily high to achieve the immediate effects of the drug.

Saliva artemisinin concentrations are highly correlated with the venous and capillary plasma artemisinin concentrations, making saliva a body fluid feasible for use in pharmacokinetic (PK) investigations of the compound (7, 21).

The aim of the present study was to investigate whether a dosing regimen consisting of low initial doses and then escalating doses of artemisinin was as efficacious as the standard high-dose regimen with respect to the immediate clinical end points. It was also of interest to evaluate if lower initial doses of artemisinin lead to a less pronounced induction of its elimination in patients with uncomplicated malaria. If this is the case, this would result in higher drug concentrations toward the end of the treatment period, which in turn could hypothetically reduce recrudescence rates.

MATERIALS AND METHODS

Study design. Seventy-seven male and female adults with uncomplicated falciparum malaria were included in this double-blind, comparative, parallel-group study. Block randomization was applied for treatment arm allocation. The study took place at the health care facilities of the Phú Rieng Rubber Plantation in Bình Phước Province, Vietnam, which is 130 km north of Hồ Chí Minh City, Vietnam. Written, informed consent was obtained from each patient prior to inclusion in the study. Permission for the study was obtained from the Ministry of Health, Hanoi, Vietnam; the Ethics Committee of the Medical Faculty of Uppsala University, Uppsala, Sweden; and the Medical Products Agency, Uppsala, Sweden.

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TABLE 1. Artemisinin doses in the two treatment arms

<table>
<thead>
<tr>
<th>Dosing regimen</th>
<th>Day 1</th>
<th>Day 2</th>
<th>Day 3</th>
<th>Day 4</th>
<th>Day 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>Standard</td>
<td>2 × 250</td>
<td>250 × 2</td>
<td>250 × 2</td>
<td>250 × 2</td>
<td>2 × 250</td>
</tr>
<tr>
<td>Escalating</td>
<td>2 × 50</td>
<td>50 × 2</td>
<td>125 × 2</td>
<td>125 × 2</td>
<td>2 × 250</td>
</tr>
</tbody>
</table>

*On days 1 and 5 the drug was given as a single dose (in milligrams) of two capsules. On the other days the dose (in milligrams) was divided and given on two occasions.*

Each patient had a peripheral axonal parasite density of more than 1,000 parasites/μl of blood and a sublingual body temperature above 37.5°C upon admission. Parasite counts were determined on Giemsa-stained thick blood films and were recorded as the number of parasites per 300 white blood cells. The following were exclusion criteria: current medication with other drugs; evidence of a cardiovascular, endocrine, gastrointestinal, hematological, hepatic, renal, or respiratory disorder; a hemoglobin level below 80 g/liter; symptoms of cerebral malaria; and antimalarial treatment 3 weeks prior to study inclusion, according to the patients when they were asked. Moreover, patients unable to communicate well with the health care workers were excluded from the study.

Artemisinin was purchased from the Institute of Materia Medica (Hanoi, Vietnam). Three different doses (50, 125, and 250 mg) were prepared in hard gelatin capsules, with cellulose used as bulk material and with all doses having identical appearances (The National Corporation of Swedish Pharmaceuticals [Apo- teek A/B], Stockholm, Sweden). All patients received the same total number of capsules on each occasion. All capsules were administered under the supervision of study team members, with ingestions followed by the intake of 100 ml of water and rinsing of the mouth with another 100 ml of water. Solid food was avoided 2 h before and 2 h after drug intake on the 2 sampling days.

Patients enrolled in group A (standard-dose group; n = 40) received the standard dose of artemisinin, while those enrolled in group B (escalating-dose group; n = 37) were treated according to an escalating-dose scheme (Table 1). On the first and the last two days all doses were given as single doses to both groups to ensure detectable saliva drug levels, while on the second, third, and fourth days the doses were administered in the morning and evening, as generally implemented in clinical settings at the time of the study. All patients were hospitalized for the duration of drug treatment. Patients were asked to return to the hospital 3 weeks after the start of treatment for a parasite test.

The target number of patients included in each group (n = 36) was calculated to enable the detection of an 8-h difference in the PCT between the two treatment groups at a significance level of 0.05 and with a power of 0.80. Each patient not completing the 5-day treatment course was replaced by two new patients, who were randomly allocated to each study arm in order to maintain the blindness of the study.

**Clinical assessments.** The parasite count in finger capillary blood was determined immediately before and every 4 h after the start of treatment until three consecutive smears were negative. The time until the first (of three) negative smear was defined as the PCT. Body temperature was simultaneously recorded sublingually until it was normal. PCT was defined as the time required for the largest number of parasites observed to decrease by 50%. Parasite mean residence time (PMT) was calculated by dividing the last predicted concentration by k. The AUC from time zero to infinity (AUC∞) was calculated as the sum of AUC0–t and AUCt–∞. The oral clearance based on the concentrations in saliva (CLs/F) for each subject was derived as the dose divided by AUC∞. The mean residence time (PMT) for each patient was calculated as the AUMC/AUC∞, where AUMC is the area under the first moment-versus-time curve and was calculated by the linear trapezoidal rule with the residual area from the last sampling point extrapolated by CLsal−k2, where CLsal is the calculated concentration.

**Results**

**Demographics.** No significant differences between the two treatment groups with respect to the initial number of parasites, body weight, age, and sex distribution were evident (Table 2). Artemisinin was well tolerated by all patients. Data for two patients in the escalating-dose group were excluded from the data analysis due to either incomplete treatment or missing data for a shorter interval than that permitted in the study protocol, respectively. These patients were replaced by four new patients, randomly allocated in pairs to either dosing regimen.

**Efficacy.** The patients in the standard-dose group had significantly shorter mean PCTs and PMRTs compared to those for the patients in the escalating-dose group ($P < 0.01$) (Fig. 1; Table 2). No statistically significant differences in FST, PCT50, or PAUC were found between the two groups. One patient in the escalating-dose group did not clear the parasites up to 104 h after intake of the first dose. In the statistical tests, the...
PCT for this patient was given an arbitrary value of 120 h. In a sensitivity analysis, the outcomes of the tests did not differ with or without the inclusion of the data for this patient.

The logarithm of the highest level of parasitemia had a significant effect on PCT and PAUC when it was used as covariate in the MANCOVA ($P < 0.01$). It also exhibited an influence on FST, with a $P$ value close to the level of significance ($P < 0.07$), whereas PCT50 and PMRT were unaffected ($P > 0.76$, and 0.74, respectively). No significant effect of age was found in any of the variables used in the PD analysis (PCT, PCT50, PMRT, PAUC, and FST).

A total of 31 patients returned for parasite control within 3 weeks after treatment termination. Among these 31 patients, *Plasmodium falciparum* parasites were detected in 6 of 17 patients in the standard-dose group and 7 of 14 patients in the escalating-dose group. No other plasmodium types were detected in any of the returning patients. All recurring infections were treated with mefloquine (Lariam; Roche, Basel, Switzerland).

**Pharmacokinetics.** Due to low saliva artemisinin concentrations, pharmacokinetic parameters were based on data for 43 male patients (24 and 19 patients in the standard- and escalating-dose groups, respectively) and 16 female patients (7 and 9 in the standard- and escalating-dose groups, respectively). Artemisinin concentrations in samples taken 30 min after drug intake were abruptly high in several patients. In the final analysis, all concentration data from this time point were considered unreliable and were not used in the data evaluation.

A MANCOVA with $CL_{\text{T}}/F, t_{1/2},$ dose-adjusted $C_{\text{max}}, T_{\text{max}},$ and MRT as the dependent variables and dosing group as the independent variable indicated significant differences between the two groups on the first treatment day ($P < 0.01$). This overall $P$ value was due to differences in $CL_{\text{T}}/F$ values between the two groups ($P < 0.01$), with higher values obtained for those in the escalating-dose group. A $P$ value close to the level of significance was obtained for MRT ($P = 0.05$), with larger values observed for patients in the standard-dose group (Table 3). No significant differences were found with respect to other

![Graph](http://aac.asm.org/)

**FIG. 1.** Number of asexual *P. falciparum* parasites per microliter of blood in adult Vietnamese patients treated with either a standard oral regimen of 500 mg of artemisinin for 5 days (group A) or an escalating-dosage oral regimen of 100 mg on days 1 and 2, 250 mg on days 3 and 4, and a final dose of 500 mg on day 5 (group B).
The extrapolated fractions of AUC_{0-\infty} were, on average, 10 and 15% for the standard- and escalating-dose groups, respectively, on day 1 and 12% for both groups on day 5.

Analysis by canonical correlation indicated no evidence for any correlation between the PD variables (PCT, PCT_{50}, PMRT, PAUC, and FST) and any of the PK variables (CL_{s/F}, C_{max}, day 1 \text{, MRT}_{day 1}) with a P value equal to 0.21. For the variable C_{max}, day 1, two outliers were removed from the data set because they were out of the range of the calibration curve. Additionally, there were three outliers in the data set, and these were removed due to the abnormal distribution that they produced. These consisted of one PMRT datum point, one PAUC datum point, and one MRT datum point. Inclusion or exclusion of these datum points did not alter the overall conclusions (P = 0.35 and 0.21, respectively). No trends for any correlation could be detected upon visual inspection of scattergrams of PD versus PK variables.

**DISCUSSION**

With improvements in analytical systems for the detection of artemisinin and its derivatives in recent years, increased numbers of reports on the PK properties of this family of antimalarials have been published (25). However, the available information has not been used in any dose-finding studies for any of these compounds.

In the present study, the PCT following the use of an escalating-dose regimen was compared with that following the use of standard artemisinin therapy. Moreover, the PKs of artemisinin were studied with saliva samples, with the hypothesis being that lower initial doses of artemisinin would result in less pronounced autoinduction of drug metabolism. The escalating-dose regimen was chosen to somewhat counter the unusual time-dependent PKs of artemisinin. The starting dose chosen for the escalating-dose regimen was based on earlier findings that suggested that lower artemisinin concentrations suffice for the clearance of the parasites (4).

PCT was prolonged in the escalating-dose group by a mean of 16 h compared to that in the group receiving the standard treatment. This is explained by the inadequate artemisinin concentrations in the former group, as manifested by 10-fold lower salivary AUCs and 7-fold lower salivary C_{max} values. In general, the prolongation of PCT is undesirable, resulting in an increased level of gametocyto genesis and an increased level of parasite transmission (6, 19).

The number of recurring infections appeared to be similar in the two treatment arms. However, this study was not dimensioned to detect any such difference. One male patient in the
escalating-dose group had not cleared the parasites until the last treatment day, although he was symptom-free. The final parasite count in his blood smears was done after his release from the study, and we were unable to locate him after we found parasites in his blood smears. This patient, however, returned to the hospital 10 days after the end of the treatment presenting with renewed symptoms. Falciparum malaria was detected in his blood, and he was successfully treated with melfloquine.

On average, $\text{CL}/F$ was significantly higher in the patients in the escalating-dose group than in the standard-dose group on the first treatment day, with no evidence for any difference in $t_{1/2}$. The difference observed here might be an indication of a saturable first-pass metabolism, such that the higher dose would have a higher oral bioavailability. The presumably lower bioavailability on day 1 in patients receiving the 100-mg dose resulted in concentrations lower than expected, which probably contributed to the prolonged PCT in this group. With arteisinin exhibiting the characteristics of a highly extracted drug (5), there would be no saturation of the systemic clearance and therefore no differences in $t_{1/2}$. Indications of saturable arteisinin kinetics after oral administration have previously been found in healthy subjects (2).

Artemisinin $\text{CL}/F$ values on day 5 did not differ between the two groups. However, these values were significantly higher compared to those on day 1 in both groups (Table 3). The increase is in accordance with earlier findings based on concentrations in plasma in both malaria patients and healthy volunteers (1, 3–5). The extent of the increase in arteisinin $\text{CL}/F$ values has been reported to be six- to sevenfold after standard treatment (1, 4, 22). In our study, the average increase in $\text{CL}/F$ from day 1 to 5 in the patients in the standard-dose group was 5-fold, whereas the average increase in the escalating-dose group was 2.5-fold. However, both groups seem to have reached the maximum induction level, resulting in similar $\text{CL}/F$ values on day 5. MANCOVA detected a significant difference between the groups with respect to the increases in these values. However, the seemingly smaller increase in the patients allocated to the escalating-dose regimen is probably due to the higher $\text{CL}/F$ values on day 1 for this group.

In a population kinetic study, lower $\text{CL}/F$ values and volumes of distribution for arteisinin were found in female children (22). However, due to the small number of male and female children involved, sex was not concluded to influence the pediatric arteisinin kinetics. Our results suggest a significant effect of sex on the kinetics of the drug on the first treatment day, with higher $\text{CL}/F$ values for female patients.

Arteisinin metabolism in human liver microsomes is mediated primarily by CYP2B6, with a probable secondary contribution of CYP3A4 in individuals with low levels of CYP2B6 expression (23). Reports on differences in the activities of these enzymes by sex are scarce, although indications of higher levels of activity in female subjects have been reported for CYP2A6 (11, 20). Another explanation for the observed effect of sex could be the imbalance in the number of male and female patients. Although MANCOVA analysis is, in general, robust enough to handle such an inequality, a possible statistical misinterpretation could not be ruled out. No effect of sex on $\text{CL}/F$ was in evidence on the last treatment day.

An alternative means for the achievement of higher arteisinin concentrations at the end of a treatment period would be to start with the standard dose of 500 mg and then increase the dose toward the end of the treatment. However, counteracting of the time-dependent PKs of arteisinin would require doses as high as 2 to 3 g toward the end of the treatment period. No data on the abilities of patients to tolerate arteisinin or its metabolites at such high doses are available, however.

Combination therapies for the treatment of malaria with an arteisinin analogue and other antimalarials with longer $t_{1/2}$ values have been proposed (17, 26). Such combinations are believed to reduce the risk for the development of resistance. Our findings support previous suggestions on the use of combination therapy because of the complex PK of arteisinin, which are both dose and time dependent. However, arteisinin is probably the least suitable of its class for use in combination chemotherapy due to the potential for drug-drug interactions caused by the induction of drug-metabolizing enzymes.

This is the first reported study on the PKs of arteisinin based solely on saliva samples. The agreement of the results of the study with those presented in previous publications on the nonlinearity of the kinetics of the compound and its autoinduction capacity encourages the use of saliva as a feasible sampling matrix. The utility of this approach lies primarily with the easier sampling procedure compared to that for blood sampling and patients’ more ready acceptance of saliva sampling. However, care must be taken in rinsing the mouth properly in order to avoid contamination of the early samples with residual arteisinins, which was the reason that the samples obtained at 30 min were discarded from the PK analysis. Moreover, due to low concentrations in saliva, no arteisinin could be detected in samples from several patients. Analytical systems more sensitive than the present one are a necessity for further investigations based on saliva sampling.

In conclusion, the kinetics of arteisinin were found to exhibit both time and dose dependencies. The escalating-dose regimen with lower initial doses of arteisinin did not result in higher levels of exposure to the drug toward the end of the treatment. This dose regimen resulted in prolonged PCTs, and its clinical implementation is not recommended.

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REFERENCES


