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Adherence to artemether/lumefantrine treatment in children under real-life situations in rural Tanzania

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\textbf{A B S T R A C T}

A follow-up study was conducted to determine the magnitude of and factors related to adherence to artemether/lumefantrine (ALu) treatment in rural settings in Tanzania. Children in five villages of Kilosa District treated at health facilities were followed-up at their homes on Day 7 after the first dose of ALu. For those found to be positive using a rapid diagnostic test for malaria and treated with ALu, their caretakers were interviewed on drug administration habits. In addition, capillary blood samples were collected on Day 7 to determine lumefantrine concentrations. The majority of children (392/444; 88.3\%) were reported to have received all doses, in time. Non-adherence was due to untimeliness rather than missing doses and was highest for the last two doses. No significant difference was found between blood lumefantrine concentrations among adherent (median 286 nmol/l) and non-adherent [median 261 nmol/l; range 25 nmol/l (limit of quantification) to 9318 nmol/l]. Children from less poor households were more likely to adhere to therapy than the poor [odds ratio (OR) = 2.45, 95\% CI 1.35–4.45; adjusted OR = 2.23, 95\% CI 1.20–4.13]. The high reported rate of adherence to ALu in rural areas is encouraging and needs to be preserved to reduce the risk of emergence of resistant strains. The age-based dosage schedule and lack of adherence to ALu treatment guidelines by health facility staff may explain both the huge variability in observed lumefantrine concentrations and the lack of difference in concentrations between the two groups.

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1. Introduction

In 2006, Tanzania switched from sulfadoxine/pyrimethamine (SP) to artemether/lumefantrine (ALu) as the first-line treatment recommended for uncomplicated falciparum malaria. The limited availability of ALu, combined with its rapid onset of action and rapid clearance of severe clinical symptoms, might tempt caretakers to discontinue treatment prematurely\textsuperscript{1} and to keep the remaining medicine for an ‘anticipated’ next attack.\textsuperscript{2} This may accelerate the pace of development of drug resistance,\textsuperscript{3} thus making it important to maintain high adherence rates to preserve the efficacy of
Adherence levels as high as 90% have been reported in controlled clinical trials.\textsuperscript{5,6} These figures are likely to be an overestimation because in real-life situations caretakers are neither given detailed instruction on how to administer the treatment nor do they receive follow-up on the use of the drugs by the child.

Two community-based Tanzanian studies conducted in Kilombero/Ulanga Districts and Zanzibar Islands did report high levels of adherence to ACT treatment regimens (77\% and 99\%, respectively).\textsuperscript{7,8} However, more information on adherence to ALu is required in order to inform the implementation of drug policy change, given the low access to ALu in rural remote areas.\textsuperscript{9} Community-based data collected under real-life settings are also required in order to estimate the level of community effectiveness,\textsuperscript{10,11} which reflects the success of implementation of a drug policy change.

Several factors have been reported to influence adherence to treatment, including: education of caretakers and the use of pre-packaged co-blister packets;\textsuperscript{12,13} drug efficacy and rapid improvement of symptoms;\textsuperscript{1,3,14} local perceptions about the disease;\textsuperscript{15} and caretakers perceiving a child playing after a single dose as a sign of cure.\textsuperscript{16} Most of these findings, however, are based on studies performed in Asia or during the time when chlorquine (CQ) and SP were the first-line recommended drugs.\textsuperscript{7,8} This community-based study was therefore conducted to determine the level of and factors influencing adherence to ALu among under-fives in a real-life situation in rural Tanzania. Information from this study will be important in designing appropriate implementation programmes for ACT in Tanzania and elsewhere in Africa.\textsuperscript{5}

2. Materials and methods

2.1. Study site

A 12-month follow-up study was conducted in June 2008 in Kilosa District, Tanzania. A detailed description of the district is reported elsewhere.\textsuperscript{9} Malaria accounts for more than one-half of clinical presentations by patients attending the outpatient clinics and 60\% of deaths among under-five hospital admissions (district annual report, 2008). The health facilities are organised at three levels, with 61 dispensaries that are at the lowest contact level, seven health centres at the middle, and two hospitals as well as a district hospital at the highest level. Only hospitals and health centres have a functioning microscope for malaria diagnosis. Dispensaries have a catchment population of approximately 10 000 people, manned by a clinician and a public health nurse, and serve the majority of the children suspected to have uncomplicated malaria.

2.2. Study design and sample size

This was a follow-up observational study. Stratified cluster sampling was carried out. All 30 rural villages listed as rural in the National Census Report, 2003, located in the wet-lowland areas, were grouped into those with and without a health facility. One village among 12 villages with a health facility and two villages among 18 villages without a health facility were randomly selected using EPI Info v.6 (CDC, Atlanta, GA, USA). The rationale of 1:2 was based on the proportion of villages with health facilities in the district. During the course of data collection, it was necessary to compare between rural and semi-urban villages. Thus, the ward headquarters of the three rural villages located in semi-urban areas were included in the study. Two of the initial villages shared the same ward; therefore, two semi-urban villages were added making the total number of study villages equal to five (Figure 1).

The unadjusted sample size was calculated as 280 based on the assumption that 24\% of caretakers would adhere to treatment, as reported in a study in Uganda.\textsuperscript{11} Although this study was based on CQ treatment, this was the most recent adherence study in a setting similar to Tanzania. A maximum 95\% CI of ±5\% was required, and adding 10\% for possible loss to follow-up and another 27\% as a correction factor for clustering effect (intracluster correlation coefficient = 0.01), the sample size was calculated as 391, approximated to 400. Based on Tanzanian studies, it was assumed that approximately 80–90\%\textsuperscript{17,18} of children with fever would be treated at health centres or dispensaries. Data collection was planned to be completed in a 4-month period. However, during the study it was found that most caretakers residing in villages without a health facility sought care from drug vendors and ordinary shops instead.\textsuperscript{9} Consequently, it took 12 months to recruit the required sample size.

2.3. Data collection

Three research assistants (RA) were purposely recruited both to administer the questionnaire and to take blood samples. They were taught for 6 days on how to administer the questionnaire and take blood samples.

At the time of the study, there were only three health facilities in the study area that provided ALu, all of which were public facilities. Therefore, these health facilities acted as enrolment centres where RAs identified the names of children who met the enrolment criteria from the laboratory and outpatient registers.

The enrolment criteria of children for the study were:

- child aged 3–59 months;
- child diagnosed at the health facility as positive for malaria using a rapid diagnostic test (RDT);
- child prescribed with the correct dose of ALu according to national treatment guideline, which is based on weight and age, twice daily for 3 days\textsuperscript{19}; and
- child resided in one of the five selected villages.

RAs traced the children home on Day 7 after the first dose of ALu. Recruitment of children into the study was performed after a caretaker had provided written informed consent. Caretakers were not given prior information about the follow-up. Caretakers were interviewed regarding the instructions received at the facility, the day and time they administered the drugs, whether food was taken with the drugs, whether food was cooked with oil, and whether
the child vomited, and if so, how soon after the drug intake. (The questionnaire can be obtained from the corresponding author upon request.) Two caretakers could not remember the events and were excluded from the analysis. In addition, baseline data on demographic and socioeconomic status (SES) and knowledge about malaria treatment were obtained. The questionnaire was based on a literature review of previous adherence studies.6,11 A pilot study was carried out for 1 week in two villages to validate the questionnaire and to gain field experiences in blood sampling techniques.

2.3.1. Definition of adherence to artemether/lumefantrine treatment schedule

Adherence to treatment with ALu was defined as a caretaker reported to have administered two doses daily for 3 days; if the first dose was started in the evening, then one dose on the first day, two doses for 2 days and the last dose on the fourth day, in the morning. The dose was one tablet of ALu (20 mg artemether and 120 mg lumefantrine) for a child aged <3 years and two tablets (artemether 40 mg and lumefantrine 240 mg) for a child aged ≥3 years.

2.4. Blood sampling, handling and drug sample analysis

RAs collected capillary blood samples from the children on Day 7 through a finger-prick using the method previously described by Ntale et al.20 After drying the filter paper, the specimen was stored in a plastic envelope to avoid contamination and was collected by the supervisors weekly. Filter paper samples were stored at room temperature for 3–15 months at the Pharmacology Laboratory of Muhimbili University (Dar es Salaam, Tanzania) before transfer to the Clinical Pharmacology Laboratory at Karolinska Institutet (Stockholm, Sweden) where blood lumefantrine concentrations were determined by HPLC as described by Ntale et al.20 Samples were assayed with an intra-assay and inter-assay coefficient of variation of <9.2% and with a limit of quantification (LoQ) of 25 nmol/l.

At the time of the study, several pilot studies were ongoing in the country aimed at rolling out RDTs for malaria. We therefore introduced RDTs for malaria diagnosis in the three facilities, since Kilosa was not one of the pilot districts. Health workers in each facility and village health workers in two dispensaries that were understaffed were trained in performing the RDT for malaria. A 1-day training course was held followed by 2 days of work under supervision. The tests were performed routinely for all children under five with a suspected malaria diagnosis irrespective of whether they resided in the five study villages or not. The RDT called Paracheck (Orchid Biomedical Systems, Goa, India) was obtained through donation from the National Malaria Control Programme.

2.5. Data analysis

Data were double-entered into EpiData 3.0 (EpiData Association, Odense, Denmark) before being transferred to Stata v.10.1 (StataCorp., College station, TX, USA). Bivariate analysis was performed on variables thought to influence adherence to treatment with ALu. Variables found to be associated were introduced in a stepwise multiple regression model using a cut-off point of 0.10. The variables were: knowledge of treatment of severe malaria; SES; and intake of food with oil. Marital status and age of the child were also included. The variables SES and intake of food with oil remained in the final regression model; these were tested for interaction and no association was found. SES was calculated using principal component analysis in which

Figure 1. Overview of the process to recruit 5 out of 161 villages and study subjects (n = 446) in Kilosa District, Tanzania.
Table 1
Background demographic characteristics of children under five enrolled in the study and their caretakers by reported adherence status (n = 444)

<table>
<thead>
<tr>
<th>Characteristic of parent or child</th>
<th>Reported non-adherence</th>
<th>Reported adherence</th>
<th>Total</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n (%)</td>
<td>n (%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Caretaker’s age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;30</td>
<td>32 (12.3)</td>
<td>228 (87.7)</td>
<td>260</td>
<td>0.642</td>
</tr>
<tr>
<td>≥30</td>
<td>20 (10.9)</td>
<td>164 (89.1)</td>
<td>184</td>
<td></td>
</tr>
<tr>
<td>Child’s age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;2</td>
<td>26 (14.1)</td>
<td>159 (85.9)</td>
<td>185</td>
<td>0.195</td>
</tr>
<tr>
<td>≥2</td>
<td>26 (10.0)</td>
<td>233 (90.0)</td>
<td>259</td>
<td></td>
</tr>
<tr>
<td>Child’s sex</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>30 (13.7)</td>
<td>189 (86.3)</td>
<td>219</td>
<td>0.199</td>
</tr>
<tr>
<td>Female</td>
<td>22 (9.8)</td>
<td>203 (90.2)</td>
<td>225</td>
<td></td>
</tr>
<tr>
<td>Caretaker’s marital status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Not married</td>
<td>34 (14.3)</td>
<td>204 (85.7)</td>
<td>238</td>
<td>0.070</td>
</tr>
<tr>
<td>Married</td>
<td>18 (8.7)</td>
<td>188 (91.3)</td>
<td>206</td>
<td></td>
</tr>
<tr>
<td>Caretaker’s education (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;7</td>
<td>25 (13.4)</td>
<td>162 (86.6)</td>
<td>187</td>
<td>0.354</td>
</tr>
<tr>
<td>≥7</td>
<td>27 (10.5)</td>
<td>230 (89.5)</td>
<td>257</td>
<td></td>
</tr>
<tr>
<td>Residence</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rural</td>
<td>40 (12.5)</td>
<td>281 (87.5)</td>
<td>321</td>
<td>0.428</td>
</tr>
<tr>
<td>Semi-urban</td>
<td>12 (9.8)</td>
<td>111 (90.2)</td>
<td>123</td>
<td></td>
</tr>
<tr>
<td>Village status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Without a health facility</td>
<td>22 (13.0)</td>
<td>147 (87.0)</td>
<td>169</td>
<td>0.502</td>
</tr>
<tr>
<td>With a health facility</td>
<td>30 (10.9)</td>
<td>245 (89.1)</td>
<td>275</td>
<td></td>
</tr>
<tr>
<td>Overall</td>
<td>52 (11.7)</td>
<td>392 (88.3)</td>
<td>444</td>
<td></td>
</tr>
</tbody>
</table>

19 assets adopted from the Tanzania Demographic Health Survey were used. Variables that had >90% of the study children in one category were excluded. The final list included household construction material composing walls and the roof; the source of drinking water and type of light; and ownership of a watch or clock, radio, mattress, a bed-net and a bicycle. The first component explained 30% of the variability and was used to construct a SES index. The SES index was categorised in five quintiles, and households in the first and second quintiles were classified as poor and the others as less poor. Crude and adjusted odds ratios (OR) with 95% CIs were reported.

Blood lumefantrine concentrations were found to have a positive skewed distribution. Hence, log-transformed values of lumefantrine concentrations were used for statistical testing of difference in mean levels (t-test) between children who adhered to therapy and those who did not. The original concentration levels were also described using medians and box plot. Before and after comparison of blood lumefantrine concentrations could not be done because this was an observational study that aimed at determining adherence to ALu under real-life conditions. Blood sampling for determination of lumefantrine concentrations was not done routinely at the health facilities. Permission to conduct the study was obtained from the regional and district authorities. Written informed consent was obtained prior to interviews from each adult caretaker during baseline data collection. All caretakers approached provided informed consent for the study, except 18 who refused to have a blood sample taken from their children.

3. Results

A total of 544 names of eligible children were identified at the health facilities, 467 of whom were found on Day 7 and their caretakers interviewed; 77 (14.2%) of the children were not included because they were not found at home (70), they were >5 years of age (2), they did not receive ALu (4) or they had died (1). Blood samples for lumefantrine concentrations were collected on Day 7 from 446 children; blood samples could not be collected from 21 children because their caretakers refused (18), the RAs failed to draw a second sample after the first attempt failed (2) or the capillary tube broke (1). Two caretakers could not recall most of the information; therefore, analysis was carried out based on the 444 children who had complete data.

Table 1 demonstrates no difference in sociodemographic characteristics between children who adhered to ALu treatment and those who did not. More than one-half of the children (58.3%) were aged ≥2 years and over one-half of the caretakers (57.9%) had at least primary level education.

The level of reported adherence to ALu was high (392/444; 88.3%). Figure 2 shows that the number of
children who were reported to have taken all doses, in time, was lowest for the last two doses.

When controlling for other factors, children from less poor households were more likely to report adherence to Alu treatment compared with the poor (OR = 2.45, 95% CI 1.35–4.45; adjusted OR = 2.23, 95% CI 1.20–4.13) (Table 2). Children who took food with oil were also more likely to adhere to Alu treatment compared with those who did not (OR = 4.62, 95% CI 2.53–8.41; adjusted OR = 4.60, 95% CI 2.49–8.50). Caretakers from the third SES quintile were most likely to adhere to treatment compared with the first quintile, however there was no observed trend.

Although the majority of caretakers (409/444; 92.1%) reported being advised on the frequency and duration for Alu administration, the knowledge did not significantly influence the reported adherence. Basic education, offered for 7 years in the country, did not influence adherence to the treatment schedule.

No significant difference in mean blood lumefantrine concentrations was found between children whose caretakers reported adherence and those who reported non-adherence. The corresponding median levels were 286 nmol/l and 261 nmol/l, respectively (Figure 3). However, there was a wide variation in blood lumefantrine concentrations, ranging from 25 nmol/l (LOQ) to 9318 nmol/l for the total group, and from 25 nmol/l to 1382 nmol/l for those with reported non-adherence and from 25 nmol/l to 9318 nmol/l for those with reported adherence. Extreme values (outliers) are not shown in Figure 3.

4. Discussion

This study has shown a high (88%) level of reported adherence to Alu treatment under real-life settings, contrary to fears that efficacy observed in controlled trials might not be reflected in real-life situations. Similar findings were reported in studies performed in Kilombero/Ulanga Districts and in Zanzibar Islands, Tanzania, despite some methodological differences. Whereas these studies included patients residing closer to health facilities (5–6 km from health facilities), mostly in urban and semi-urban areas and applied presumptive malaria diagnosis, the current included children from rural remote areas (up to 25 km away) where the burden of malaria is the highest. 

Therefore, the current study confirms a high adherence to Alu among under fives diagnosed to have malaria using a RDT even in most rural settings remote from health facilities.

No significant difference in blood lumefantrine concentrations was found between children reported to adhere to therapy and those who did not, in line with a study in Malawi but contrary to a study in Uganda. One of the reasons for the lack of any difference might be due to the lack of a clear cut-off point. Although a previous study in Thailand reported a Day 7 blood lumefantrine

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Table 2
Logistic regression analysis of factors influencing adherence to artemether/lumefantrine (Alu) among children under five in Kilosa District, Tanzania (n = 444)

<table>
<thead>
<tr>
<th>Factor</th>
<th>OR</th>
<th>95% CI</th>
<th>Adjusted OR</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Access to healthcare services</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Child’s age ≥ 2 years</td>
<td>1.47</td>
<td>0.82–2.61</td>
<td>1.80</td>
<td>0.90–3.58</td>
</tr>
<tr>
<td>Married</td>
<td>1.74</td>
<td>0.95–3.19</td>
<td>1.87</td>
<td>0.90–3.88</td>
</tr>
<tr>
<td>Caretaker’s basic education ≥ 7 years</td>
<td>1.32</td>
<td>0.74–2.35</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Residing in semi-urban area</td>
<td>1.32</td>
<td>0.67–2.60</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Using own or hired bicycle to go to health facility</td>
<td>1.16</td>
<td>0.58–2.29</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Distance &lt;5 km from a government facility</td>
<td>1.46</td>
<td>0.81–2.63</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Knowledge on malaria</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Knowledge that mosquitoes cause malaria</td>
<td>1.65</td>
<td>0.65–4.17</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Knowledge that high fever causes convulsion in children</td>
<td>1.41</td>
<td>0.76–2.60</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Knowledge of treatment of severe malaria</td>
<td>0.45</td>
<td>0.20–1.01</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Knowledge that dawa mseto (Alu) is the recommended drug for the treatment of malaria ya kavaida (uncomplicated malaria).</td>
<td>0.73</td>
<td>0.39–1.37</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Socioeconomic status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Less poor (quintiles 3–5)</td>
<td>2.45</td>
<td>1.35–4.45</td>
<td>2.23</td>
<td>1.20–4.13</td>
</tr>
<tr>
<td>Other factors</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>History of vomiting during malaria episode</td>
<td>0.68</td>
<td>0.22–2.06</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Taken food with oil</td>
<td>4.62</td>
<td>2.53–8.41</td>
<td>4.60</td>
<td>2.49–8.50</td>
</tr>
<tr>
<td>Taken adequate amount of food</td>
<td>1.08</td>
<td>0.59–1.98</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sought treatment same or next day</td>
<td>0.97</td>
<td>0.51–1.85</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

OR: odds ratio.
concentration cut-off point of 331 nmol/l (175 ng/ml). 25 Another study in Malawi reported patients were cured even below this level. 24 The difference might be explained by possible variations in food intake, 26 as the levels of lumefantrine blood concentrations may increase 16-fold in patients who took the drug with fatty food compared with without fat. 27, 28 Age-based dosage schedules of ALu might also explain the unpredictable blood concentrations of lumefantrine, since the dose per kg may vary widely between patients, as reported for the CQ/SP combination. 29 This underscores the need to monitor the rate of adherence to treatment guidelines by providers as well as treatment schedules among patients in rural areas when implementing ALu treatment on a wide scale in Africa.

The high reported adherence found in this study can be explained by the availability of ALu in pre-packaged packets with pictorial instructions, which have been reported to improve caretakers’ understanding of the schedule administration of the drug 30 and hence they are more likely to follow the provider’s advice. 12, 22, 31 Studies conducted when CQ resistance was high reported low adherence levels, 10, 11 thus the highly efficacious ALu used in the present study might also explain the high reported level of adherence to treatment. 22

Non-adherence to ALu treatment found in this study was mostly attributed to untimely dosing rather than missing doses, contrary to concerns that caretakers might discontinue treatment and save the remaining drugs for later use. 1, 2 The high level of non-adherence in the last two doses has also been reported in previous studies in Tanzania and Malawi. 3, 24 This might be due to some caretakers perceiving that a child playing after an initial dose is cured. 16 Encouragingly, the low non-adherence level might have little impact on the treatment strategy; however, there is a potential risk of building up pressure for resistant strains over time. 32

Children from the less poor households were found to adhere to treatment better than poor patients. This can partly be explained by the higher level of education among caretakers in the less poor households and the tendency to own radios compared with the poor. 9 This might have exposed them to more information disseminated by the National Malaria Control Programme through radios and posters since the introduction of ALu.

4.1. Methodological issues

Caretakers’ reports are subjective and were likely to overestimate the level of reported adherence owing to possibilities of obtaining socially desirable responses. 24 This was minimised by explaining to the caretaker the objective of the study prior to the interviews. The fact that most children were no longer sick on Day 7 reduced the likelihood of receiving desirable responses. This is in contrast to reports of limited validity of caretakers’ reports when interviewed at the health facilities. 33 Pill count could not be performed on Day 7 since most of the caretakers had lost the packets. This limited the opportunity to triangulate study findings. However, the fact that the results conform to other community-based studies conducted in Tanzania where pill count was performed indicates that misreporting might be minimal.

Recalling detailed information about how the drug was administered after 7 days might result in data inaccuracy as some details might be forgotten. We tried to minimise this by following-up children weekly when the caretakers could still remember the details. In addition, a sick child tends to draw the attention of caretakers in a household, thus increasing the likelihood of remembering events in more details.

Blood lumefantrine concentrations would be useful when studying adherence. However, the lack of a clear cut-off point 24 and baseline levels limited a detailed analysis. There were also other challenges resulting from applying the method in a real-life situation. Several methodological issues were found in this study that could limit the usefulness of the capillary blood sampling method for lumefantrine in evaluating ALu effectiveness under real-life conditions. These included the lack of baseline lumefantrine concentration, inability to control or record food intake, and inconsistency in determining ALu dosage where some prescribers based the dosage on child weight while others used age. In addition, some prescribers used child weight measured on the same day while others used the record found in the growth monitoring card, which could be outdated by several weeks or months.

5. Conclusion

This study showed that the reported rate of adherence to ALu treatment was encouragingly high in rural villages with or without a health facility. The large variability in lumefantrine concentrations cast doubt on the rationale of using aged-based dosage schedules of ALu as presently used in some facilities in Tanzania. Non-adherence was mainly attributed to off-schedule dosing rather than taking a fewer number of doses and was more pronounced in the last two doses. Despite the high adherence, advocacy is still needed to improve caretakers’ understanding of the importance of adhering to the dose schedule in order to reduce the risk of building up resistant strains.

Authors’ contributions: DOS, DK, GT, ZP, MP, MM and LLG designed the study protocol; DOS and DK supervised the data collection; DOS, MP and LLG analysed the data; all authors participated in drafting and revising the manuscript critically for intellectual content. All authors read and approved the final manuscript. DOS is guarantor of the paper.

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