

# Malaria Transmission and Asymptomatic Malaria during Rainy Season among Nomads, North-eastern Nigeria

## ABSTRACT

**Aim:** To determine the point prevalence of malaria infection and asymptomatic malaria during rainy season among some nomads of North Eastern Nigeria.

**Study Design:** A cross sectional observational study.

**Place and Duration of Study:** The study was conducted across 11 randomly selected nomads' camps around the Rivers Gongola and Benue basins spread over 3 Local Government Areas of Southern Adamawa State of North Eastern Nigeria. Data was collected during rainy season between July and September, 2016.

**Methodology:** Fifty-five randomly selected households (5 from each camp) were covered in the survey. One hundred and ninety two (192) consenting participants aged between 1 and 79 years (inclusive) were involved in the survey. Structured questionnaires were administered (care givers consented and responded on behalf of children) and blood samples collected. Blood samples were examined for malaria parasite using a microscopes and results of both survey and microscopy analysed.

**Results:** Overall malaria parasite prevalence was 87.5% and mean parasite density was 36,168 parasites per  $\mu$ l of blood. Thirty five (18.2%) of participant were of low parasite density, 35.5% were of moderate parasite density while 32.8% were of high parasite density. More than half (53%) of the malaria positive participants did not experience febrile symptoms within one month prior to the survey and were therefore asymptomatic. Tendency of manifestation of symptoms significantly increased with parasite density and decreased with age. The use of preventive measures against mosquito bite was 7.7% and only 16.1% of participants used antimalarial medicines or sought medical attention during their most recent fever episode.

**Conclusion:** The high prevalence of asymptomatic carriers with high parasite densities and abysmally low usage of preventive and curative measures among the study population represents an ideal condition for effective malaria transmission which is unlikely to abate unless control measures are intensified.

**Keywords:** [Malaria parasite transmission, asymptomatic malaria, parasite density, Nomadic Fulani]

## 1. INTRODUCTION

Malaria remains one of the most dreaded public health problems worldwide causing a death toll of 627000 in 2020 up from 558,000 in 2019 (1) and a loss of enormous economic resources which has been estimated at 12 billion USD annually in terms of direct cost and multiples of it in terms of loss in economic growth (2). About 95% of malaria burden is borne by Sub-Saharan Africa (3, 4). With renewed control efforts, malaria incidence, prevalence and mortality is declining globally prior to the COVID-19 pandemic (1). However, there is considerable disparity in the decline among and within countries (5, 6). While the estimated prevalence of malaria in Nigeria is below 31.4% (1), many localized studies have reported prevalence of between 50 - 80 percent in some foci (7-10). In highly

25 endemic areas, malaria prevalence could exceed 90% during peak transmission periods (11). During  
26 high transmission, individuals in malaria endemic regions often harbor high number of parasites -  
27 exceeding 100,000 parasites/ul of blood(12). Although parasitaemia has been found to positively  
28 correlate with severity of illness, repeated exposure to infection appear to gradually enhance  
29 mechanisms to limit the inflammatory response associated with febrile illness and hence large  
30 proportion of infected individuals remain asymptomatic carriers (13, 14).

31 There is an estimated 35 million Fulani nomads spread across West and Central African countries  
32 (15). Nigeria hosts a considerable proportion of the nomadic Fulani population in West Africa. It is  
33 estimated that there are about 15.3 million Fulani pastoralist in Nigeria and a considerable number of  
34 them are nomads (16, 17). Nomad groups usually have defined and fairly consistent pattern of  
35 migration following an annual cycle. Although most of their camps during their migration cycle is  
36 located in close vicinity to local sedentary communities, their culture and life style significantly differ  
37 from these neighboring sedentary communities (18).

38 Fulani nomads are by their life style more exposed to infectious diseases including malaria than  
39 sedentary populations. Although appearing generally healthier their sedentary rural neighbours, they  
40 seldom benefit from interventional programmes of the conventional health system (19).

41 In Nigeria, nomads inhabiting the Gongola Benue basins in Adamawa State live in highly malaria  
42 endemic region and prevalence of about 37% has been reported even during the dry season (18, 20).  
43 Nomadic Fulani have for long held the belief that malaria which is locally identified as pabboje is an  
44 inherent illness of the Fulani and does not need to be treated since it “visits for a while and goes  
45 away”. It is believed that modern antimalarial treatment may aggravate subsequent malaria episodes  
46 (18). The complacency which nomadic Fulani adopts in coping with malaria and anecdotal data  
47 suggest that nomads could harbor substantial malaria parasite burden while remaining asymptomatic.  
48 That, coupled with their reduced access to intervention services, positions the Nigerian nomadic  
49 Fulani as reservoirs of malaria infection with the potential of upsetting current control and elimination  
50 efforts. Determining the magnitude of this concealed prevalence among the nomads is important in  
51 highlighting the need for targeted interventions. We report here the point prevalence of malaria  
52 infection and asymptomatic malaria during rainy season (between July and September, 2016) among  
53 the nomads of north western Adamawa State of North eastern Nigeria.

## 54 55 **2. MATERIAL AND METHODS**

### 56 57 **2.1 Study Area**

58 The study was conducted in 11 nomadic Fulani camps spread across three Local Government areas  
59 in the Southwestern part of Adamawa State, namely; Demsa, Numan and Mayo Belwa Local  
60 government areas. These local areas are located between latitudes 8.400N - 9.500N and longitudes  
61 11.50S - 12.35S. Along the border of the Local government areas (Demsa and Numan) is a  
62 confluence of two important rivers - Gongola and Benue. The basins of these two rivers attract  
63 substantial economic activities including fishing, cropping and animal grazing. Nomadic Fulani  
64 pastoralists traditionally congregate along the river basin as soon as crops are harvested around  
65 December. A proportion of the nomads occupy parcels of uncultivated the hilly parts of the basin  
66 during the rainy season. Although the 2013 malaria epidemiological guide reported Adamawa State  
67 malaria risk model as mesoendemic (21), the Upper Benue Basin (also within Adamawa) is  
68 hyperendemic and a 2020 under-five malaria risk modeling report has designated it among the high  
69 malaria “hotspots” (22). Moreover, nomadic Fulani pastoralist in the Benue Basin have been known  
70 to be highly exposed and harbor high malaria infection rates (23)

### 71 72 **2.2 Study Design**

73 A cross sectional survey design was conducted during the months of July and September targeting  
74 nomads who spend the rainy season in the study area.

### 75 76 **2.3 Sampling**

77 Eleven camps were randomly selected from a list forty camps earlier identified for an interventional  
78 study by means of folded papers. Six (Anini, Chore, Dudel, Dwam, Marawo and Kadel) camps from  
the cluster of camps in Demsa and Numan LGAs and 5 (Korawa-Maccido, Korawa-Umaru, Korawa-

79 Burti, Korawa-Ahmadu and Liringo) camps from the Mayo Belwa LGA cluster. From each camp, 5  
80 households were randomly selected and all consenting members of the household who were 1 year  
81 or older were included in the study. In addition to children less than 1 year old, severely ill household  
82 members were also excluded from the study.

## 83 2.4 Data Collection

84 Structured questionnaires designed for oral interviews were administered for each participant either  
85 directly or through a child minder (in the case of children). The questionnaire was designed to collect  
86 demographic data, clinical manifestation of malaria, medicine usage (anti malaria medicine use was  
87 defined by either taking a prescription from a health personnel or use of any of the approved  
88 antimalarial medicines since they are easily recognizable) and use of barriers against mosquito bite  
89 (use of protective barrier was defined by use of insecticide, insect repellent or mosquito net). A  
90 section on blood collection and examination has been included in each questionnaire. The section  
91 was used to document results of microscopy for malaria parasite.

92 Blood sample collection was done concurrently with the interview. For each participant, a sterile lancet  
93 was used for blood collection. The lancet was opened by the collector (a research team member) and  
94 witnessed by the participant or community members (for children). The lancet was used to prick the  
95 ball of the finger after gently massaging and disinfecting the area with ethanol-soaked cotton wool. By  
96 squeezing the finger for free flow of blood, a drop of blood is collected on clean grease-free slide.  
97 The edge of another slide was used to make a thick smear and allowed to dry for laboratory staining  
98 and microscopy. Finally, the pricked area was cleaned with dry cotton wool.

## 99 2.5 Microscopy for Malaria Parasite

100 Microscopy and parasite estimation were done as described by WHO and Adu-Gyasi and colleagues  
101 (24, 25). The slides containing the thick smears were brought to the laboratory and stained using 10%  
102 Giemsa stain. The slides were allowed to dry and observed under X100, oil immersion objective of a  
103 light microscope. The number of asexual stage of parasites seen per oil immersion field and the  
104 number of white blood cells (WBCs) were also counted for the same field. Counting continued until  
105 100 parasites or 200 WBCs was counted and when less than 10 parasites were found after counting,  
106 counting continued until 500 WBCs were counted. As many as 100 oil immersion fields were viewed  
107 without identifying a malaria parasite before a slide is regarded negative. Microscopy of slides was  
108 validated by a reference microscopist in Yola Specialist Hospital.

## 109 2.6 Parasite Density Estimation

110 Numbers of parasite and WBCs seen were recorded and entered into EpiInfo along with the rest of  
111 the data. Parasite densities per micro liter ( $\mu\text{l}$ ) were subsequently computed in SPSS 20 as follows:

112 Parasite density per  $\mu\text{l}$  = (Number of parasite x 8000)/(Number of WBC)

113 Where 8000 is the assumed number of WBC per  $\mu\text{l}$  of blood.

114 Parasite densities across all the malaria positive participants were classified into three levels fairly  
115 corresponding to classification by Kotepui and colleagues (26). Participants whose slides had 1 –  
116 3999 parasites per  $\mu\text{l}$  were classified as low parasite density, 4000 – 24999 parasites per  $\mu\text{l}$  classified  
117 as moderate parasite density while 25000 or more parasites per  $\mu\text{l}$  were classified as high parasite  
118 density. Those with 0 parasites per 100 oil immersion field were classified as not infected.

## 119 2.7 Data Entry and Analysis

120 All data were entered into EpiInfo 6 and transferred into SPSS 16 for analysis. Frequencies and  
121 percentages were used to compute prevalence and proportions. Association between febrile  
122 symptoms and parasite densities was explored and significance tested by Chi Square at 95%  
123 confidence level. Similarly, Associations between asymptomatic infection (those who did not report  
124 having fever in the last one month) and other demographic variables were explored and significance  
125 tested at 95% confidence level using chi square.

126

## 127 3. RESULTS AND DISCUSSION

128

### 129 3.1 Demographic characteristics and malaria parasite infection

130 One hundred and ninety two participants comprising of 131 males and 61 females took part in the  
131 study. Table 1 show that demographic characteristics of the participant and their malaria infection  
132 status. The participants belong to two major clans of nomadic Fulani frequently camping around the  
133 study area – the Kiri clan made up 58.9% of the participant while the remaining 41.1% were made up  
134 of Fulani from the Goriji clan. Malaria parasite count on thick blood smears from the participants  
135 showed that 24 (12.5%) of the participants were not infected while the remaining 168 (87.5%) were  
136 infected with varying densities of malaria parasite. Estimation of levels of parasitaemia showed that  
137 18.2% of participant were of low parasite density (1-3999 parasites per  $\mu$ l) category, 35.5% were of  
138 moderate parasite density (4000 – 24999 parasites per  $\mu$ l) while 32.8% were in the high parasite  
139 density (>24999 parasites per  $\mu$ l) category. The mean parasite density among the malaria positive  
140 participants was 36168 parasites per  $\mu$ l of blood.  
141

142 **Table 1: Demographic characteristics and malaria infection status of participants**

<b>Variables</b>	<b>Number</b>	<b>in Percentage</b>
<b>Sex</b>		
Males	131	68.2
Females	61	31.8
<b>Clan</b>		
Kiri	113	58.9
Goriji	79	41.1
<b>Age group</b>		
<5 years	25	13.0
5 – 14 years	93	48.4
15 – 29 years	26	13.5
30 – 44 years	27	14.1
45 – 59 years	16	8.3
60 years and above	5	2.6
<b>Malaria infection status</b>		
Not infected	24	12.5
Low parasite density	35	18.2
Moderate parasite density	70	36.5
High Parasite density	63	32.8

143

### 144 3.2 Parasite density and manifestation of symptoms

145 Table 2 shows the distribution of infection and parasite densities among participants with recent  
146 (within one month) febrile symptoms and those without. Most (56.8%) of the participants (infected and  
147 non-infected) did not experience febrile symptoms within one month prior to the date of collection of  
148 blood sample. Proportions of participants with febrile symptoms were similar among non-infected  
149 (16.7%) and those infected low parasite density (17.1%). Febrile symptoms experience shows  
150 significant association with parasite density at  $p=0.001$ . More than half of the participants with  
151 moderate parasite density (51.4%) and high parasite density (58.7%) had recent febrile symptoms.  
152 Symptoms most frequently reported includes hot-body (71 cases) followed by headache (66 case).  
153 Least frequently reported symptoms were diarrhea (4 cases), cough (3 cases) and chest pain (2  
154 cases) which were all categorized as “other” symptoms. Majority (109) of participants reported  
155 experiencing no symptoms in the last one month prior to data collection (Figure 1).

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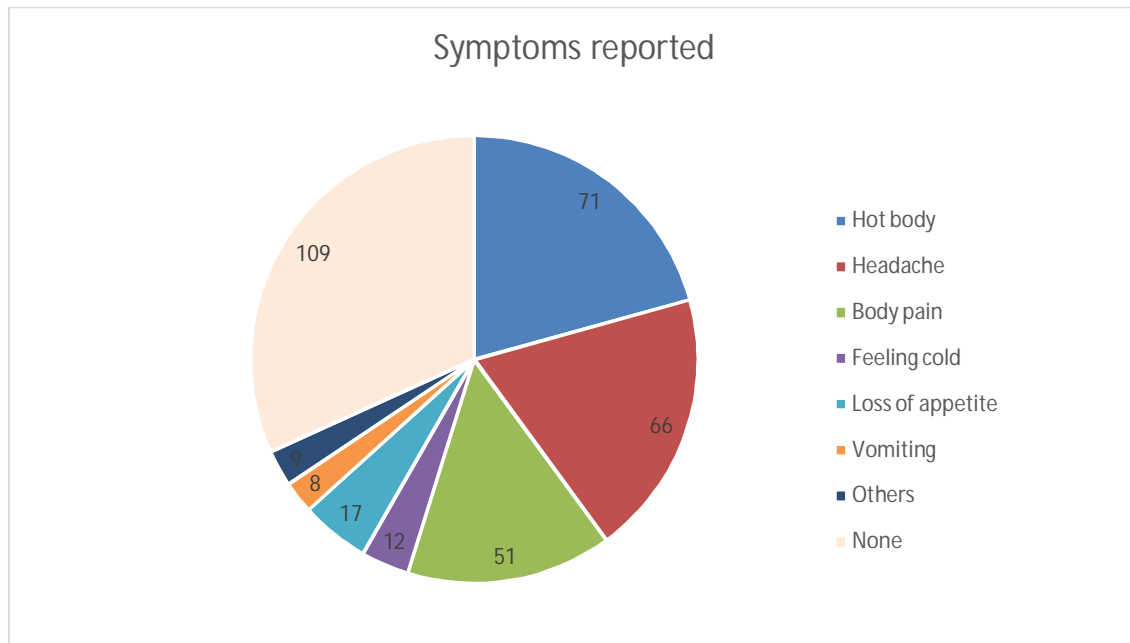
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160 **Table 2: Distribution of malaria parasite densities among symptomatic and asymptomatic**  
 161 **participants**

Malaria infection status (N)	Recent (within one month) febrile symptom	
	Symptomatic (%)	Asymptomatic (%)
Not infected (24)	4 (16.7)	20 (83.3)
Low parasite density (35)	6 (17.1)	29 (82.9)
Moderate parasite density (70)	36 (51.4)	34 (48.6)
High Parasite density (63)	37 (58.7)	26 (42.3)
Total	83 (43.2)	109 (56.8)

162  $\chi^2=24.2691, df=3, p<0.001$   
 163



164

165 **Figure 1: Symptoms (in including “no symptoms”) as reported by participants**

166 **3.3 Manifestation of symptoms febrile symptoms, demographic characteristics and**  
 167 **use of control measures**

168 Among the 168 malaria positive participants, proportion of those who were asymptomatic vary  
 169 significantly with age group but insignificantly with sex, clan, antimalarial medicine use and use of  
 170 protective barriers against mosquito bite at  $p=.05$  (Table 3). Sixty four (57.1%) of malaria positive  
 171 males were asymptomatic while 25 (44.6%) of females were asymptomatic. However, Chi-square test  
 172 showed that the difference in the proportion of asymptomatic participants between to two genders  
 173 was not statistically significant at 95% confidence level. Use of preventive measures against mosquito  
 174 bite and use of antimalaria medicine during the most recent fever episode were 7.7% and 16.1%  
 175 respectively among the participants. Manifestation of febrile symptoms did not significantly differ  
 176 between those who use protective barriers against mosquito bite and those who do not neither did it  
 177 differ between those who between took antimalarial medicines and those who did not.

178

179 **Table 3: Comparison of symptomatic (N=79) and asymptomatic (N=89) malaria positive**  
 180 **participants by demographic characteristics, antimalarial use and use of barriers against**  
 181 **mosquito bite**

Demographic characteristics, antimalarial use and use of barriers against mosquito bite (N)	Recent (within one month) febrile symptoms		
	Symptomatic (%)	Asymptomatic (%)	p-value

<b>Sex</b>			0.086
Male (112)	48 (42.9)	64 (57.1)	
Female (56)	31 (55.4)	25 (44.6)	
<b>Clan</b>			0.084
Kiri (109)	56 (51.4)	53 (48.6)	
Goriji (59)	23 (39.0)	36 (61.0)	
<b>Age group</b>			<b>&gt;0.001**</b>
<5years (21)	18 (85.7)	3 (14.3)	
5-14 years (84)	45 (53.6)	39 (46.4)	
15 – 29 years (24)	7 (29.2)	17 (70.8)	
30 – 44 years (24)	6 (25.0%)	18 (75.0%)	
45 years and above (15)	3 (20.0)	12 (80.0)	
<b>Antimalarial medicine use</b>			0.224
Yes (27)	15 (55.6)	12 (44.4)	
No (141)	64 (45.4)	77 (54.6)	
<b>Use of barriers against mosquito bite</b>			0.590
Yes (13)	6 (46.2)	7 (53.8)	
No (155)	73 (47.1)	82 (52.9)	

182 \*\*Significant at P=0.01

183

### 184 3.4 Discussion

185 We conducted a cross sectional survey to determine the point prevalence of asymptomatic malaria  
 186 and its relationship with other relatable factors among the Fulani nomads who inhabit some of the less  
 187 swampy uncultivated parcels of land around the basins of Rivers Benue and Gongola in South  
 188 Western part of Adamawa State during the rainy season.

189 The results showed high prevalence (87%) of malaria infection during the rainy season which  
 190 corresponds to season of high transmission similar to what has been reported in other studies in  
 191 some parts of Nigeria (10, 11). This high prevalence of infection was expectedly, complemented by  
 192 high parasite densities, agreeing with findings from other similar studies (27,28). Average parasite  
 193 density among the malaria positive participants 36361/ul.

194 Considering those who were malaria parasite positive, there were more asymptomatic participants  
 195 (53%) than the symptomatic ones (47%). This confirms high prevalence of asymptomatic malaria  
 196 infection among the study population similar to what is being reported in many malaria hyper-endemic  
 197 settings (29-31).

198 Although there appears to be no clear-cut threshold of parasite density that corresponds to febrile  
 199 attacks, the likelihood of experiencing fever symptoms increased significantly with increasing parasite  
 200 density. This finding is consistent with a number of studies reporting significant relationship between  
 201 fever-risk and parasite density, among other variables (32, 33).

202 The proportions of participants who reported recent febrile symptoms were similar among non-  
203 infected (16.7%) and those infected with low parasite density (17.1%) suggesting that, at low  
204 densities, parasitaemia might not have played a role in eliciting fever symptoms in the study area.  
205 Similar finding was reported in a study in West Bengal where none of the malaria positive participants  
206 with parasite densities up to 12,800 parasites per µl exhibited symptoms of malaria (34). Expectedly,  
207 significantly more of the participants with higher parasite densities (>3999) experienced recent febrile  
208 illness but even in those categories, a considerably large proportion (45%) of them was also  
209 asymptomatic. This is indicative of the existence of effective transmission in the study population even  
210 among apparently healthy individuals.

211 Similar to earlier reported findings from the same study area (18), the use of preventive barriers  
212 against mosquito bites and the use of antimalarial medicines were as low as 7.7% and 16.1%  
213 respectively. Lower usage of preventive measure ensures exposure to malaria infection and hence  
214 transmission. Furthermore, the abysmally low antimalarial usage among both symptomatic and  
215 asymptomatic malaria positive participants during their last fever episode suggests that transmission  
216 could perpetuate uninterrupted.

217 When five socio- demographic attributes including sex, clan, age, use of antimalarial medicine and  
218 use of protective barriers against mosquito bite were explored for association with manifestation of  
219 symptoms among the malaria positive participants; it was found that only age showed significant  
220 association with manifestation of symptoms. More than 85% of malaria positive children younger than  
221 five years have experienced recent febrile symptoms while 53% of malaria positive participants within  
222 the age bracket of 5-14 years (inclusive) had recent symptoms and as few as 20% of malaria positive  
223 participants who were 45 years or older experienced recent febrile symptoms. This finding  
224 corroborates the much reported age (hence exposure)-dependent acquired immunity among malaria  
225 endemic populations (35-37).

226  
227

#### 228 4. CONCLUSION

229

230 The prevalence of malaria parasitaemia is high in the study area with most of the study participants  
231 hosting thousands of parasites per µl of blood. However, in most cases, clinical symptoms do not  
232 manifest. This veiled parasitaemia is more common among the older members of the population than  
233 in the younger ones apparently because of the relative immunity acquired by the older ones through  
234 repeated exposure. The use of preventive measures against mosquito bites and antimalarial  
235 medicines are abysmally low among the study population. The scenario represents an ideal condition  
236 for effective malaria transmission which is unlikely to abate unless control measures are intensified.  
237 We therefore recommend the application of tailor-made control strategies to this population with  
238 exceptional lifestyle.

239

#### 240 Ethical Approval and Consent

241

242 Ethical approval was obtained from Adamawa State Ministry of Health via a letter referenced  
243 S/MOH/1131/1/06. Nomad community leaders at various levels were approached and their consent to  
244 work among their community members obtained. Informed consent was obtained from each adult  
245 participant and from child minders (in the case of children). Standard aseptic techniques were strictly  
246 adhered to during sample collection and data obtained were handled confidentially. Field team  
247 included a health personnel who managed minor illnesses using essential medicines in the team's kit.  
248 Cases were referred to the health facility when necessary.

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#### 251 REFERENCES

252

253 World Health Organization. World Malaria Report; 2021. Geneva: World Health  
254 Organization; 2021.

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2. Centers for Disease Control and Prevention. Impact of Malaria Worldwide 2021  
(Accessed 1 September 2022). Available:  
[https://www.cdc.gov/malaria/malaria\\_worldwide/impact.html](https://www.cdc.gov/malaria/malaria_worldwide/impact.html).

- 257 3. World Health Organization. Malaria Fact sheet 2022 (Accessed 2 September 2022).  
258 Available: <https://www.who.int/news-room/fact-sheets/detail/malaria>.
- 259 4. World Health Organization. World malaria report 2020. Geneva: World Health  
260 Organization; 2020.
- 261 5. Severe Malaria Observatory. Malaria Facts: Nigeria. Knowledge sharing for Malaria  
262 [Internet]. 2020 (Accessed 1 September 2022). Available:  
263 <https://www.severemalaria.org/countries/nigeria>.
- 264 6. World Health Organization. World malaria report 2013. Geneva: World Health  
265 Organization; 2013.
- 266 7. Oladeinde B, Omoregie R, Olley M, Anunibe J, Onifade A, Oladeinde O. Malaria and  
267 Anemia among Children in a Low Resource Setting In Nigeria. *Iran J Parasitol.*  
268 2012;7(3):31-7.
- 269 8. Awosolu OB, Yahaya ZS, Farah Haziqah MT, Simon-Oke IA, Fakunle C. A cross-  
270 sectional study of the prevalence, density, and risk factors associated with malaria  
271 transmission in urban communities of Ibadan, Southwestern Nigeria. *Heliyon.* 2021;7(1):  
272 e05975.
- 273 9. Ajayi IO, Afonne C, Dada-Adegbola H, Falade CO. Prevalence of Asymptomatic  
274 Malaria and Intestinal Helminthiasis Co-infection among Children Living in Selected Rural  
275 Communities in Ibadan Nigeria. *Am J Epid Infect Dis.* 2015;3(1):15-20.
- 276 10. Eze CN, Amadi EC. Survey of haemoparasitic infections among Fulani pastoralists in  
277 Rivers State. *J. Appl SCI Environ Manag.* 2016;20(3):790-4.
- 278 11. Awosolu OB, Yahaya ZS, Farah Haziqah MT, Simon-Oke IA, Olanipekun IT, Oniya  
279 MO. Epidemiology of falciparum malaria among residents of some rural and periurban  
280 communities in Ekiti State, Southwestern Nigeria. *Trop Biomed.* 2021;38(1):14-21.
- 281 12. Samdi LM, Ajayi JA, Oguiche S, Ayanlade A. Seasonal variation of malaria parasite  
282 density in paediatric population of Northeastern Nigeria. *Glob J of Health Sci.* 2012;4(2):103-  
283 9.
- 284 13. Mangal P, Mittal S, Kachhawa K, Agrawal D, Rath B, Kumar S. Analysis of the  
285 Clinical Profile in Patients with Plasmodium falciparum Malaria and Its Association with  
286 Parasite Density. *J. Glob Infect. Dis.* 2017;9(2):60-5.
- 287 14. Gonzales SJ, Reyes RA, Braddom AE, Batugedara G, Bol S, Bunnik EM. Naturally  
288 Acquired Humoral Immunity Against Plasmodium falciparum Malaria. *Front. Immunol.*  
289 2020;11.
- 290 15. Mission Africa. Belfast: Mission Africa. 2016. [Accessed 3 December 2016].  
291 Available: [http://www.missionafrica.org.uk/ministries/14/engaging-the-nomadic-fulani-in-](http://www.missionafrica.org.uk/ministries/14/engaging-the-nomadic-fulani-in-nigeria)  
292 [nigeria](http://www.missionafrica.org.uk/ministries/14/engaging-the-nomadic-fulani-in-nigeria).
- 293 16. Ducrot MJ, Revie CW, Shaw AP, Musa UB, Bertu WJ, Gusi AM, et al. Wealth,  
294 household heterogeneity and livelihood diversification of Fulani pastoralists in the Kachia

- 295 Grazing Reserve, northern Nigeria, during a period of social transition. *PLoS One*.  
296 2017;12(3):e0172866.
- 297 17. The Editors of Encyclopaedia Britannica. Fulani. Encyclopaedia Britannica 2022.
- 298 18. Akogun OB, Gundiri MA, Badaki JA, Njobdi SY, Adesina AO, Ogundahunsi OT.  
299 Febrile illness experience among Nigerian nomads. *Int J Equity Health*. 2012;11:5.
- 300 19. Sheik-Mohamed A, Velema JP. Where health care has no access: the nomadic  
301 populations of sub-Saharan Africa. *Trop Med Int Health*. 1999;4(10):695-707.
- 302 20. Akogun OB, Adesina AO, Njobdi S, Ogundahunsi O. Nomadic Fulani communities  
303 manage malaria on the move. *Int Health*. 2012;4(1):10-9.
- 304 21. National Malaria Control Programme, suMAP, World Health Organization, The  
305 INFORM Project. A description of the epidemiology of malaria to guide the planning of  
306 control in Nigeria: A report prepared for the Federal Ministry of Health, Nigeria, the Roll  
307 Back Malaria Partnership and the Department for International Development, UK. Abuja,  
308 Nigeria: 2013.
- 309 22. Chigozie Louisa Jane U, Temesgen Z. Evaluating the Effects of Climate and  
310 Environmental Factors on Under-5 Children Malaria Spatial Distribution Using Generalized  
311 Additive Models (GAMs). *J Epidemiol Glob Health*. 2020;10(4):304-14.
- 312 23. Gundiri MA, Lombonyi CA, Akogun OB. Malaria in an obligate nomadic Fulani  
313 camps in Adamawa State, north-eastern Nigeria. *Niger J Parasitol*. 2007;28(2):87-9.
- 314 24. World Health Organization. Basic laboratory methods in medical parasitology.  
315 Geneva: World Health Organization; 1991. 114 p. p.
- 316 25. Adu-Gyasi D, Adams M, Amoako S, Mahama E, Nsoh M, Amenga-Etego S, et al.  
317 Estimating malaria parasite density: assumed white blood cell count of 10,000/mul of blood  
318 is appropriate measure in Central Ghana. *Malar J*. 2012;11:238.
- 319 26. Kotepui M, Piwkhram D, PhunPhuech B, Phiwklam N, Chupeerach C, Duangmano S.  
320 Effects of malaria parasite density on blood cell parameters. *PLoS One*.  
321 2015;10(3):e0121057.
- 322 27. Mayengue PI, Kouhounina Batsimba D, Niama RF, Ibara Ottia R, Malonga-Massanga  
323 A, Fila-Fila GPU, et al. Variation of prevalence of malaria, parasite density and the  
324 multiplicity of *Plasmodium falciparum* infection throughout the year at three different health  
325 centers in Brazzaville, Republic of Congo. *BMC Infect Dis*. 2020;20(1):190.
- 326 28. Awosolu OB, Yahaya ZS, Farah Haziqah MT. Prevalence, Parasite Density and  
327 Determinants of *Falciparum* Malaria Among Febrile Children in Some Peri-Urban  
328 Communities in Southwestern Nigeria: A Cross-Sectional Study. *Infect Drug Resist*.  
329 2021;14:3219-32.
- 330 29. Fogang B, Biabi MF, Megnekou R, Maloba FM, Essangui E, Donkeu C, et al. High  
331 Prevalence of Asymptomatic Malarial Anemia and Association with Early Conversion from  
332 Asymptomatic to Symptomatic Infection in a *Plasmodium falciparum* Hyperendemic Setting  
333 in Cameroon. *Am J Trop Med Hyg*. 2021;106(1):293-302.

- 334 30. Starzengruber P, Fuehrer H-P, Ley B, Thriemer K, Swoboda P, Habler VE, et al. High  
335 prevalence of asymptomatic malaria in south-eastern Bangladesh. *Malar J.* 2014;13(1):16.
- 336 31. Rose UN, Nwabueze US, Emmanuel OI, Kalu OA. Asymptomatic Plasmodium  
337 falciparum malaria infection in University undergraduate students. *Asian J Biomed Pharm.*  
338 2019;9(68).
- 339 32. Dollat M, Talla C, Sokhna C, Diene Sarr F, Trape JF, Richard V. Measuring malaria  
340 morbidity in an area of seasonal transmission: Pyrogenic parasitemia thresholds based on a  
341 20-year follow-up study. *PLoS One.* 2019;14(6):e0217903.
- 342 33. Plucinski MM, Rogier E, Dimbu PR, Fortes F, Halsey ES, Aidoo M, et al.  
343 Performance of Antigen Concentration Thresholds for Attributing Fever to Malaria among  
344 Outpatients in Angola. *J Clin Microbiol.* 2019;57(3).
- 345 34. Ganguly S, Saha P, Guha SK, Biswas A, Das S, Kundu PK, et al. High prevalence of  
346 asymptomatic malaria in a tribal population in eastern India. *J Clin Microbiol.*  
347 2013;51(5):1439-44.
- 348 35. Rogier C, Commenges D, Trape JF. Evidence for an age-dependent pyrogenic  
349 threshold of Plasmodium falciparum parasitemia in highly endemic populations. *Am J Trop*  
350 *Med Hyg.* 1996;54(6):613-9.
- 351 36. Bodker R, Msangeni HA, Kisinza W, Lindsay SW. Relationship between the intensity  
352 of exposure to malaria parasites and infection in the Usambara Mountains, Tanzania. *Am J*  
353 *Trop Med Hyg.* 2006;74(5):716-23.
- 354 37. Zhao Y, Zeng J, Zhao Y, Liu Q, He Y, Zhang J, et al. Risk factors for asymptomatic  
355 malaria infections from seasonal cross-sectional surveys along the China-Myanmar border  
356 *Malar J.* 2018;17(1):247.