

Rain Season Malaria Parasite Transmission and Asymptomatic Malaria among Northeastern Nigerian Nomads

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 μl of blood. Thirty five (18.2%) of participant were of low parasite density, 35.5% were of moderate parasite density while 32.8% were in the category 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 1 million a year (1, 2) 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 (3). About 90% of malaria burden is borne by Sub-Saharan Africa (4, 5). With renewed control efforts, malaria incidence, prevalence and mortality is declining globally(5). However, there is considerable disparity in the decline among and within countries(6, 7). While the estimated prevalence of malaria in Nigeria is below 50%(6), many localized studies have reported prevalence of between 60 - 80 percent in some foci(8-10). In highly endemic areas, malaria prevalence could exceed 90% during peak

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

30 There is an estimated 35 million Fulani nomads spread across West and Central African
31 countries(15). Nigeria hosts a considerable proportion of the nomadic Fulani population in West
32 Africa. It is estimated that about 9% of its 160 million people are Fulani and a considerable number of
33 which are nomads (15, 16). Nomad groups usually have defined and fairly consistent pattern of
34 migration following an annual cycle. Although most of their camps during their migration cycle is
35 located in close vicinity to local sedentary communities, their culture and life style significantly differ
36 from these neighboring sedentary communities (17).

37 Fulani nomads are by their life style more exposed to infectious diseases including malaria than
38 sedentary populations. Although appearing generally healthier their sedentary rural neighbours, they
39 seldom benefit from interventional programmes of the conventional health system(18).

40 In Nigeria, nomads inhabiting the Gongola Benue basins in Adamawa State live in highly malaria
41 endemic region and prevalence of about 37% has been reported even during the dry season(17, 19).
42 Nomadic Fulani have for long held the belief that malaria which is locally identified as pabboje is an
43 inherent illness of the Fulani and does not need to be treated since it “visits for a while and goes
44 away”. It is believed that modern antimalarial treatment may aggravate subsequent malaria episodes
45 (17). The complacency which nomadic Fulani adopts in coping with malaria and anecdotal data
46 suggest that nomads could harbor substantial malaria parasite burden while remaining asymptomatic.
47 That, coupled with their reduced access to intervention services, positions the Nigerian nomadic
48 Fulani as reservoirs of malaria infection with the potential of upsetting current control and elimination
49 efforts. Determining the magnitude of this concealed prevalence among the nomads is important in
50 highlighting the need for targeted interventions. We report here the point prevalence of malaria
51 infection and asymptomatic malaria during rainy season (between July and September, 2006) among
52 the nomads of north western Adamawa State of North eastern Nigeria.

53 **2. MATERIAL AND METHODS / EXPERIMENTAL DETAILS / METHODOLOGY (ARIAL, 54 BOLD, 11 FONT, LEFT ALIGNED, CAPS)**

55 **2.1 Study Area**

56
57 The study was conducted in 11 nomadic Fulani camps spread across three Local Government areas
58 in the Southwestern part of Adamawa State, namely; Demsa, Numan and Mayo Belwa Local
59 government areas. These local areas are located between latitudes 8.400N - 9.500N and longitudes
60 11.50S - 12.35S. Along the border of the Local government areas (Demsa and Numan) is a
61 confluence of two important rivers - Gongola and Benue. The basins of these two rivers attract
62 substantial economic activities including fishing, cropping and animal grazing. Nomadic Fulani
63 pastoralists traditionally congregate along the river basin as soon as crops are harvested around
64 December. Many of the nomads stay along the river basins with their cattle devouring pasture – often
65 changing locations – until the rainy season sets in. In Mayo Belwa LGA, some nomads of the Goriji
66 clan spend their dry season close to two smaller rivers Mayo Inne and Mayo Sakanare mostly grazing
67 on harvested from lands and regularly visiting the rivers for their water need. In all the LGAs, there are
68 usually a smaller proportion of nomad pastoralists who spend the rainy season with a few cattle on
69 some uncultivated parcels of the land.
70

71 **2.2 Study Design**

72
73 A cross sectional survey design was conducted during the months of July and August targeting
74 nomads who spend the rainy season in the study area.

75 **2.3 Sampling**

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

79 and Liringo) camps from the Mayo Belwa LGA cluster. From each camp, 5 households were randomly
80 selected and all consenting members of the household who were 1 year or older were included in the
81 study. In addition to children less than 1 year old, severely ill household members were also excluded
82 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 and use of barriers against
87 mosquito bite. A section on blood collection and examination has been included in each
88 questionnaire. The section was used to document results of microscopy for malaria parasite.

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

96 **2.5 Microscopy for Malaria Parasite**

97 Microscopy and parasite estimation were done as described by WHO and Adu-Gyasi and colleagues
98 (20, 21). The slides containing the thick smears were brought to the laboratory and stained using 10%
99 Giemsa stain. The slides were allowed to dry and observed under X100, oil immersion objective of a
100 light microscope. The number of asexual stage of parasites seen per oil immersion field and the
101 number of white blood cells (WBCs) were also counted for the same field. Counting continued until
102 100 parasites or 200 WBCs was counted and when less than 10 parasites were found after counting,
103 counting continued until 500 WBCs were counted. As many as 100 oil immersion fields were viewed
104 without identifying a malaria parasite before a slide is regarded negative. Microscopy of slides was
105 validated by a reference microscopist in Yola Specialist Hospital.

106 **2.6 Parasite Density Estimation**

107 Numbers of parasite and WBCs seen were recorded and entered into EpiInfo 6 along with the rest of
108 the data. Parasite densities per micro liter (μl) were subsequently computed in SPSS 16 as follows:

109 Parasite density per μl = (Number of parasite x 8000)/(Number of WBC)

110 Where 8000 is the assumed number of WBC per μl of blood.

111 Parasite densities across all the malaria positive participants were classified into three levels fairly
112 corresponding to classification by Kotepui and colleagues. (22). Participants whose slides had 1 –
113 3999 parasites per μl were classified as low parasite density, 4000 – 24999 parasites per μl classified
114 as moderate parasite density while 25000 or more parasites per μl were classified as high parasite
115 density. Those with 0 parasites per 100 oil immersion field were classified as not infected.

116 **2.7 Data Entry and Analysis**

117 All data were entered into EpiInfo 6 and transferred into SPSS 16 for analysis. Frequencies and
118 percentages were used to compute prevalence and proportions. Association between febrile
119 symptoms and parasite densities was explored and significance tested by Chi Square at 95%
120 confidence level. Similarly, Associations between asymptomatic infection and other demographic
121 variables were explored and significance tested at 95% confidence level using chi square.

122 **2.8 Ethical Consideration**

123 Ethical clearance was obtained from Adamawa State Ministry of Health and nomad community
124 leaders at various levels were approached and their consent to work among their community
125 members obtained. Informed consent was obtained from each adult participant and from child
126 minders (in the case of children). Standard aseptic techniques were strictly adhered to during sample
127 collection and data obtained were handled confidentially.

128

129 3. RESULTS AND DISCUSSION

130

131 3.1 Demographic characteristics and malaria parasite infection

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

143

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

Demographic characteristics and infection status of participants	Number in sample (N=192)	Percentage
Sex		
Males	131	68.2
Females	61	31.8
Clan		
Kiri	113	58.9
Goriji	79	41.1
Age group		
<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
Malaria infection status		
Not infected	24	12.5
Low parasite density	35	18.2
Moderate parasite density	70	36.4
High Parasite density	63	32.8

145

146 3.2 Parasite density and manifestation of symptoms

147 Table 2 shows the distribution of infection and parasite densities among participants with recent
148 (within one month) febrile symptoms and those without. Most (56.8%) of the participants (infected and
149 non-infected) did not experience febrile symptoms within one month prior to the date of collection of
150 blood sample. Proportions of participants with febrile symptoms were similar among non-infected
151 (16.7%) and those infected low parasite density (17.1%). Febrile symptoms experience shows
152 significant association with parasite density at $p=0.001$. More than half of the participants with
153 moderate parasite density (51.4%) and high parasite density (58.7%) had recent febrile symptoms.

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158 **Table 2: Distribution of malaria parasite densities among symptomatic and asymptomatic**
 159 **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 (51.3)	34 (48.6)
High Parasite density (63)	37 (58.7)	26 (42.3)
Total	83 (43.2)	109 (56.8)

160 $X^2=24.2691, df=3, p<0.001$

161

162 **3.3 Manifestation of symptoms febrile symptoms, demographic characteristics and**
 163 **use of control measures**

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

174

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

Demographic characteristics, antimalarial use and use of barriers against mosquito bite (N)	Recent (within one month) febrile symptoms		
	Symptomatic (%)	Asymptomatic (%)	p-value
Sex			0.086
Male (112)	48 (42.9)	64 (57.1)	
Female (56)	31 (55.4)	25 (44.6)	
Clan			0.084
Kiri (109)	56 (51.4)	53 (48.6)	
Goriji (59)	23 (39.0)	36 (61.0)	
Age group			>0.001**
<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)	
Antimalarial medicine use			0.224
Yes (27)	15 (55.6)	12 (44.4)	

No (141)	64 (45.4)	77 (54.6)
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Use of barriers against mosquito

bite		0.590
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Yes (13)	6 (46.2)	7 (53.8)
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No (155)	73 (47.1)	82 (52.9)
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178 **Significant at P=0.01

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180 **3.4 Discussion**

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

185 The results showed high prevalence (87%) of malaria infection during the rainy season which
 186 corresponds to season of high transmission similar to what has been reported in other studies in
 187 some parts of Nigeria (8, 23). This high prevalence of infection was expectedly, complemented by high
 188 parasite densities. Average parasite density among the malaria positive participants 36361/ul.

189 Considering those who were malaria parasite positive, there were more asymptomatic participants
 190 (53%) than the symptomatic ones (47%). This confirms high prevalence of asymptomatic malaria
 191 infection among the study population similar to what is being reported in many malaria endemic
 192 settings (24-27).

193 Although there appears to be no clear-cut threshold of parasite density that corresponds to febrile
 194 attacks, the risk of experiencing fever symptoms increased significantly with increasing parasite
 195 density. This finding is consistent with that of a cohort study in a Madagascan community with low
 196 malaria transmission where significant relationship between fever-risk and parasite density, among
 197 other variables, was reported but a threshold values of parasite density that corresponds to onset of
 198 fever could not be established (28).

199 The proportions of participants who reported recent febrile symptoms were similar among non-
 200 infected (16.7%) and those infected with low parasite density (17.1%) suggesting that, at low
 201 densities, parasitaemia might not have played a role in eliciting fever symptoms in the study area.
 202 Similar finding was reported in a study in West Bengal where none of the malaria positive participants
 203 with parasite densities up to 12,800 parasites per µl exhibited symptoms of malaria (24). Expectedly,
 204 significantly more of the participants with higher parasite densities (>3999) experienced recent febrile
 205 illness but even in those categories, a considerably large proportion (45%) of them was also
 206 asymptomatic. This is indicative of the existence of effective transmission in the study population even
 207 among apparently healthy individuals.

208 Similar to earlier reported findings from the same study area(17), the use preventive barriers against
 209 mosquito bites and the use of antimalarial medicines were as low as 7.7% and 16.1% respectively.
 210 Lower usage of preventive measure ensures exposure to malaria infection and hence transmission.
 211 Furthermore, the abysmally low antimalarial usage among both symptomatic and asymptomatic
 212 malaria positive participants during their most recent fever episode suggests that transmission could
 213 perpetuate uninterrupted.

214 When five socio- demographic attributes including sex, clan, age, use of antimalarial medicine and
 215 use of protective barriers against mosquito bite were explored for association with manifestation of
 216 symptoms among the malaria positive participants; it was found that only age showed significant
 217 association with manifestation of symptoms. More than 85% of malaria positive children younger than
 218 five years have experienced recent febrile symptoms while 53% of malaria positive participants within
 219 the age bracket of 5-14 years (inclusive) had recent symptoms and as few as 20% of malaria positive
 220 participants who were 45 years or older experienced recent febrile symptoms. This finding

221 corroborates the much reported age (hence exposure)-dependent acquired immunity among malaria
222 endemic populations (14, 29, 30).

223

224

225 **4. CONCLUSION**

226

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

236

237

238 **CONSENT (WHERE EVER APPLICABLE)**

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240 All participants were duly informed about the study and they signed informed consent form before
241 participation. Adult care givers consented on behalf of minors.

242

243

244 **ETHICAL APPROVAL (WHERE EVER APPLICABLE)**

245

246 Ethical clearance was obtained from Adamawa State Ministry of Health.

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249 **REFERENCES**

250

251 1. Onkoba NW, Chimbari MJ, Mukaratirwa S. Malaria endemicity and co-infection with tissue-
252 dwelling parasites in Sub-Saharan Africa: a review. *Infect Dis Poverty*. 2015;4:35.

253 2. Casares S, Richie TL. Immune evasion by malaria parasites: a challenge for vaccine
254 development. *Curr Opin Immunol*. 2009;21(3):321-30.

255 3. Centers for Disease Control and Prevention. Impact of Malaria 2016 17 September, 2016.
256 Available from: https://www.cdc.gov/malaria/malaria_worldwide/impact.html.

257 4. World Health Organization. Malaria Fact sheet 2016 25/05/2016. Available from:
258 <http://www.who.int/mediacentre/factsheets/fs094/en/>.

259 5. World Health Organization. World malaria report 2015. Geneva: World Health Organization; 2015.
260 242 p.

261 6. World Health Organization. World malaria report 2013. Geneva: World Health Organization; 2013.
262 xxviii, 253 p. p.

263 7. World Health Organization. World malaria report 2012. Geneva: World Health Organization; 2012.
264 xxxiv, 249 p. p.

265 8. Olasehinde GI, Ajayi AA, Taiwo SO, Adekeye BT, Adeyeba OA. Prevalence and management of
266 falciparum malaria among infants and children in Ota, Ogun State, Southwestern Nigeria. *Afr J Clin
267 Exper Microbiol*. 2010;11(3):159-63

268 9. Olasehinde GI, Ojorongbe DO, Akinjogunla OJ, Egwari LO, Adeyeba AO. Prevalence of Malaria
269 and Predisposing Factors to Antimalarial Drug Resistance in Southwestern Nigeria. *Research Journal
270 of Parasitology*. 2015;10: 92-101.

- 271 10. Ezugbo-Nwobi IK, Obiukwu MO, Umeanato PU, Egbuche CM. Prevalence of Malaria Parasites
272 among Nnamdi Azikwe University Students and Anti-Malaria Drug Use. *African Research Review*.
273 2011;5(4):135-44.
- 274 11. Kalu Mong Kalu, Obasi NA, Nduka FO, Glory Otuchristian. A Comparative Study of the
275 Prevalence of Malaria in Aba and Umuahia Urban Areas of Abia State, Nigeria. *Research Journal of*
276 *Parasitology* 2012;7:17-24.
- 277 12. Odongo-Aginya E, Ssegwanyi G, Kategere P, Vuzi PC. Relationship between malaria infection
278 intensity and rainfall pattern in Entebbe peninsula, Uganda. *Afr Health Sci*. 2005;5(3):238-45.
- 279 13. Weiss GE, Traore B, Kayentao K, Ongoiba A, Doumbo S, Doumtable D, et al. The Plasmodium
280 falciparum-specific human memory B cell compartment expands gradually with repeated malaria
281 infections. *PLoS Pathog*. 2010;6(5):e1000912.
- 282 14. Boisier P, Jambou R, Raharimalala L, Roux J. Relationship between parasite density and fever
283 risk in a community exposed to a low level of malaria transmission in Madagascar highlands. *Am J*
284 *Trop Med Hyg*. 2002;67(2):137-40.
- 285 15. Mission Africa. Belfast: Mission Africa. 2016. [cited 2016]. Available from:
286 <http://www.missionafrica.org.uk/ministries/14/engaging-the-nomadic-fulani-in-nigeria>.
- 287 16. Anter T. Pointblank news. 2013. [cited 2016]. Available from:
288 <http://pointblanknews.com/pbn/exclusive/special-report-who-are-the-fulani-people-and-their-origins/>.
- 289 17. Akogun OB, Gundiri MA, Badaki JA, Njobdi SY, Adesina AO, Ogundahunsi OT. Febrile illness
290 experience among Nigerian nomads. *International journal for equity in health*. 2012;11:5.
- 291 18. Sheik-Mohamed A, Velema JP. Where health care has no access: the nomadic populations of
292 sub-Saharan Africa. *Tropical medicine & international health : TM & IH*. 1999;4(10):695-707.
- 293 19. Akogun OB, Adesina AO, Njobdi S, Ogundahunsi O. Nomadic Fulani communities manage
294 malaria on the move. *Int Health*. 2012;4(1):10-9.
- 295 20. World Health Organization. Basic laboratory methods in medical parasitology. Geneva: World
296 Health Organization; 1991. 114 p. p.
- 297 21. Adu-Gyasi D, Adams M, Amoako S, Mahama E, Nsoh M, Amenga-Etego S, et al. Estimating
298 malaria parasite density: assumed white blood cell count of 10,000/mul of blood is appropriate
299 measure in Central Ghana. *Malar J*. 2012;11:238.
- 300 22. Kotepui M, Piwkhram D, PhunPhuech B, Phiwklam N, Chupeerach C, Duangmano S. Effects of
301 malaria parasite density on blood cell parameters. *PLoS One*. 2015;10(3):e0121057.
- 302 23. Adefioye OA, Adeyeba OA, Hassan WO, Oyeniran OA. Prevalence of Malaria Parasite Infection
303 among Pregnant Women in Osogbo, Southwest, Nigeria
304 *American-Eurasian Journal of Scientific Research*. 2007;2(1):3.
- 305 24. Ganguly S, Saha P, Guha SK, Biswas A, Das S, Kundu PK, et al. High prevalence of
306 asymptomatic malaria in a tribal population in eastern India. *J Clin Microbiol*. 2013;51(5):1439-44.
- 307 25. Alves FP, Durlacher RR, Menezes MJ, Krieger H, Silva LH, Camargo EP. High prevalence of
308 asymptomatic Plasmodium vivax and Plasmodium falciparum infections in native Amazonian
309 populations. *Am J Trop Med Hyg*. 2002;66(6):641-8.
- 310 26. Dal-Bianco MP, Koster KB, Kombila UD, Kun JF, Grobusch MP, Ngoma GM, et al. High
311 prevalence of asymptomatic Plasmodium falciparum infection in Gabonese adults. *Am J Trop Med*
312 *Hyg*. 2007;77(5):939-42.

- 313 27. Singh R, Godson, II, Singh S, Singh RB, Isyaku NT, Ebere UV. High prevalence of asymptomatic
314 malaria in apparently healthy schoolchildren in Aliero, Kebbi state, Nigeria. *J Vector Borne Dis.*
315 2014;51(2):128-32.
- 316 28. Boisier P, Jambou R, Raharimalala L, Roux J. Relationship between parasite density and fever
317 risk in a community exposed to a low level of malaria transmission in Madagascar highlands. *Am J*
318 *Trop Med Hyg.* 2002;67(2):4.
- 319 29. Rogier C, Commenges D, Trape JF. Evidence for an age-dependent pyrogenic threshold of
320 *Plasmodium falciparum* parasitemia in highly endemic populations. *Am J Trop Med Hyg.*
321 1996;54(6):613-9.
- 322 30. Bodker R, Msangeni HA, Kisinza W, Lindsay SW. Relationship between the intensity of exposure
323 to malaria parasites and infection in the Usambara Mountains, Tanzania. *The American journal of*
324 *tropical medicine and hygiene.* 2006;74(5):716-23.