

## Malaria among the Geriatric Population in Parts of South-Eastern Nigeria: Prevalence, Complications and Co-morbidity with Non-communicable Diseases

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### Abstract

This study was done to ascertain malaria prevalence, its complications and co-morbidity with non communicable diseases among the geriatric population in parts of South-Eastern, Nigeria. Ninety two (92) consenting subjects between the ages of 50 and 80 years were recruited in Ihiagwa, South-Eastern Nigeria for the study. Blood samples collected from them were Giemsa stained and examined microscopically for malaria parasites. Clinical examination in addition to health facility records analysis were done to determine malaria signs and symptoms, complications and co-morbidity with non- communicable diseases.

There was high malaria prevalence among study subjects. Respiratory involvement (28.2%) was the major complication associated with it while Diabetes Mellitus (44.5%) was found to be the most co-morbid non communicable disease. The older adults should be adequately included in malaria control programs. Further investigations into the pattern, dynamics and factors relating to malaria co morbidity with non communicable diseases among this age group should be given urgent attention.

**Keywords:** Malaria; Geriatric population; Prevalence; Complications; Co-morbidity

### Introduction

Malaria occurs almost exclusively in the tropics and subtropics and approximately 40% of the world's population, mostly those living in the world's poorest countries, are at risk of malaria. Every year, more than 500 million people become severely ill with malaria – most cases and deaths occur within sub-Saharan Africa [1].

The world health organization and the World bank ranked malaria as the largest single component of the disease burden in Africa, causing an annual loss of 35 million future life years from disability and premature mortality [1,2]. In Africa, malaria is responsible for about 20-30% of hospital admissions and about 30-50% of outpatient consultations [3].

Malaria is a major cause of morbidity and mortality in Nigeria. It is endemic throughout the country with seasonal variations in different zones of the country. At least 100% of the population suffers from one episode of malaria each year; it is also the commonest cause of outpatient hospital attendance [4].

Most efforts towards malaria management and control are aimed at under five children and pregnant women. Those in the older age categories are not adequately considered in most of these programs, amounting to some kind of neglect for those in the older age groups.

Although evidence suggests that clinical immunity to malaria develops after exposure to parasites as one progresses in age, this immunity is only partial and may be influenced by some variations such as variations in endemicity [5,6]. Secondly, there is increased advocacy for malaria prevention strategies such as the use of

Insecticide Treated Bed Nets and Indoor Residual Spraying etc. However, where these strategies are not well managed, factors that result in development of partial immunity may not be adequate enough to guarantee such immunity.

Furthermore, the recent upsurge in the prevalence and burden of non-communicable diseases in infectious disease endemic areas may be a threat to the integrity and sustenance of such partial immunity especially in the older age categories [7,8].

Evidence suggests also that some of the non-communicable diseases may exert reasonable impact on a person's immunity [9]. If so, this phenomenon may lead to compromise of partial immunity such as that acquired due to frequent exposure to an infectious disease agent like the malaria. These are therefore likely to put older people at risk of unexpected clinical burden of the illness with attendant morbidity and complications.

The public health burden and clinical implications of co morbidity of infectious and non-communicable diseases in infectious disease endemic areas should be a major cause for concern. Since Malaria is the most prevalent infectious disease in endemic areas, it is necessary that its dynamics among older adults especially with regards to the factors stated be reassessed. This will aid in ascertaining the true situation and ensure adequate as well as timely accommodation of people of this age group in malaria management and control programs.

This is very important so as not to shift the burden of the disease from paediatrics and pregnant women to the geriatric population. Again, the economic implications of this illness on the older adults would generally reflect on the economy of families, communities and malaria endemic Nations and should therefore be considered.

This study investigated the malaria prevalence, complications and co morbidity with non communicable diseases among the geriatric population in Ihiagwa, South-Eastern Nigeria.

## Materials and Methods

### Study setting and population

This study was carried out in Ihiagwa community health center, Imo State, South-Eastern Nigeria. The Area lies on Latitude 5.24N and Longitude 7.00E, has a total population of about 101,754 people as of the 2006 census and covers an area of 297 square kilometers.

It is located in the tropical rain forest with climatic and environmental conditions that support malaria endemicity. The target study populations were those above 50 years of age that presented at the health center. The subjects were recruited by simple balloting each day until the target population of 100 subjects was reached.

However in the course of the study 5 subjects died and 3 opted out leaving us with 92 subjects for the study. The course of death of the subjects were associated with complications arising from the ailments the presented with in the health facility.

Questionnaires were used to collect sociodemographic data from the consenting study subjects

### Sample collection and parasitologic examination

Blood samples were collected from finger prick of each consenting subject. Thick and thin smears were immediately made on clean grease free slide and stained with 5 % Giemsa. The stained slides were taken to the Federal Medical Centre (FMC), Owerri, South Easten Nigeria where Parasitological examinations were made independently by two microscopists, with discrepancies resolved by a third reader.

A slide was declared negative if parasites were absent after examining 200 high power fields. Parasite density was quantified

against 200 leukocytes on an assumed leukocyte count of 8000 per  $\mu$ L of blood.

## Clinical Examination and Analysis of Health Facility Records

Clinical examination by an experienced clinician and health records analysis were done to ascertain signs, symptoms and complications associated with malaria as well as other reported non communicable diseases.

### Data Analysis

Data was analysed with descriptive statistics in the Statistical Package for Social Sciences (SPSS).

### Ethical Considerations

The study protocol and instruments were approved by the ethical committee of the Department of Public Health Technology, Federal University of Technology, Owerri, Nigeria.

Informed oral consent was sought and obtained from study participants after the objectives and purpose of the study were clearly explained to them.

The limitation of this study was in recruiting more patients as the health facility was a small one that just served the community.

Furthermore it was a bit difficult to convince the patients to enroll in the study as some of them were skeptical.

## Results

Table 1 represents the socio-demographic characteristics of respondents.

Variables	Frequency	Percentage
Gender of the respondents		
Females	34	36.9
Males	58	63
Total	92	100
Age of the respondents		
51-60	11	11.9
61-70	49	53.2
71-80	27	29.3
81 above	5	5.4
Total	92	100
Educational status of the respondents		
Primary	31	33.6
Secondary	36	39.1

Tertiary	25	27.1
Total	92	100
<b>Occupational Status</b>		
Farmer	25	27.1
Driver	30	32.6
Civil servant	17	18.4
Unemployed	20	21.7
Total	92	100

**Table 1:** Socio-demographic characteristics of the respondents.

The study encountered more males 58(63%) than females 34(36.9%). Most were 61-70 years of age (53.2%).

Age and gender distribution of malaria infections is depicted in (Table 2).

Variables	Positive to malaria parasite	Negative to malaria parasite	Total
<b>51-60 yrs</b>			
Male	2 (2.1%)	2 (2.1%)	4 (4.3%)
Female	4 (4.3%)	3 (3.2%)	7 (7.6%)
<b>61-70 yrs</b>			
Male	13 (14.1%)	6 (6.5%)	19 (20.6%)
Female	23 (25.0%)	7 (7.6%)	30 (32.6%)
<b>71-80 yrs</b>			
Male	6 (6.5%)	3 (3.2%)	9 (9.7%)
Female	16 (7.3%)	2 (2.1%)	18 (19.5%)
<b>81 above</b>			
Male	1 (1.0%)	1 (1.0%)	2 (2.1%)
Female	2 (2.1%)	1 (1.0%)	3 (3.2%)
Total	67 (72.8%)	25 (27.1%)	92 (100%)

**Table 2:** Distribution of malaria parasitemia.

Variables	Intensity			Total
	(+++)	(++)	(+)	
<b>51-60 yrs</b>				
Male	2 (2.1%)	3 (3.2%)	2 (2.1%)	7 (7.6%)
Female	1 (1.0%)	3 (3.2%)	1 (1.0%)	5 (5.4%)
<b>61-70 yrs</b>				
Male	4 (14.3%)	6 (6.5%)	8 (8.6%)	18 (19.6%)
Female	7 (7.6%)	19 (20.6%)	6 (6.5%)	32 (34.8%)
<b>71-80 yrs</b>				

Male	3 (3.2%)	7 (7.6%)	4 (4.3%)	14 (15.2%)
Female	2 (2.1%)	8 (8.6%)	3 (3.2%)	13 (14.1%)
<b>81 above</b>				
Male	-	1 (1.0%)	1 (1.0%)	2 (2.2%)
Female	-	2 (2.1%)	1 (1.0%)	3 (3.3%)
Total	19 (20.6%)	49 (53.2%)	24 (26%)	92 (100%)

**Table 3:** Distribution of malaria parasite intensity.

More females (7.6%; 32.6%; 19.5%) were infected than males (7.6%; 20.6%; 9.7%) with respect to the different age categories of 51-60 years; 61-70 years and 71-80 years.

Age and gender distribution of malaria parasite intensity is shown in Table 3.

Males between the age group of 51-60 years had 7.6% parasite intensity while their female counterparts had 5.4%. Among those aged 61-70 years, males had 19.6% parasite intensity while the females had 34.8%.

Age/parasite intensity	Cerebral (CI) involvement	Respiratory involvement (RSI)	Renal involvement (RI)	Anaemia	Coma
51-60 yrs	1 (1.0%)	4 (4.3%)	2 (2.1%)	3 (3.2%)	1 (1.0%)
+++	0	2 (2.1%)	0	0	0
++	0	1 (1.0%)	1 (1.0%)	1 (1.0%)	0
+	1 (1.0%)	1 (1.0%)	1 (1.0%)	2 (2.1%)	1 (1.0%)
61-70 yrs	12 (13.0%)	11 (11.9%)	5 (5.4%)	13 (14.1%)	8 (8.6%)
+++	4 (4.3%)	3 (3.2%)	2 (2.1%)	4 (4.3%)	2 (2.1%)
++	6 (6.5%)	4 (4.3%)	2 (2.1%)	4 (4.3%)	2 (2.1%)
+	2 (2.1%)	4 (4.3%)	1 (1.0%)	5 (5.4%)	4 (4.3%)
71-80 yrs	3 (3.2%)	10 (10.8%)	7 (7.6%)	4 (4.3%)	3 (3.2%)
+++	0	2 (2.1%)	2 (2.1%)	0	1 (1.0%)
++	2 (2.1%)	3 (3.2%)	3 (3.2%)	2 (2.1%)	1 (1.0%)
+	1 (1.0%)	5 (5.4%)	2 (2.1%)	2 (2.1%)	1 (1.0%)
81 yrs and above	1 (1.0%)	1 (1.0%)	1 (1.0%)	0 (0%)	2 (2.1%)
+++	0	0	0	0	0
++	0	0	1 (1.0%)	0	1 (1.0%)
+	1 (1.0%)	1 (1.0%)	0	0	1 (1.0%)
Total	16 (16.3%)	26 (28.2%)	15 (16.3%)	20 (21.7%)	14 (15.2%)
Grand total = 92(100%), +++ = 1-10 parasites in every high power field; ++ = 11-100 parasites per 100 high power fields; + = 1-10 parasites per 100 high power fields					

**Table 4:** Malaria related complications.

In Table 4 indicates that the most common complication associated with malaria was respiratory involvement (RSI) (28.2%) among others. In the Table 5 it is shown that the most common co-morbid disease

recorded was diabetes mellitus (25%) which was found more among those aged 61-70 years.

Age/Malaria density	Diabetes mellitus	Hypertension	Nephropathy	Other Non-communicable diseases
51-60 yrs	5 (5.4%)	3 (3.2%)	2 (2.1%)	1 (1.0%)
+++	3 (3.2%)	1 (1.0%)	0	0
++	1 (1.0%)	1 (1.0%)	1 (1.0%)	0
+	1 (1.0%)	1 (1.0%)	1 (1.0%)	1 (1.0%)
61-70 yrs	23 (25.0%)	16 (17.3%)	6 (6.5%)	4 (4.3%)
+++	12 (13.0%)	8 (8.6%)	2 (2.1%)	2 (2.1%)
++	7 (7.6%)	7 (7.6%)	1 (1.0%)	1 (1.0%)
+	4 (4.3%)	1 (1.0%)	1 (1.0%)	1 (1.0%)
71-80 yrs	11 (11.9%)	9 (9.7%)	1 (1.0%)	6 (6.5%)
+++	5 (5.4%)	5 (5.4%)	1 (1.0%)	2 (2.1%)
++	5 (5.4%)	0	2 (2.1%)	2 (2.1%)
+	1 (1.0%)	1 (1.0%)	0	2 (2.1%)
81 yrs and above	2 (2.1%)	2 (2.1%)	0 (0%)	1 (1.0%)
+++	1 (1.0%)	0	0	0
++	1 (1.0%)	1 (1.0%)	0	1 (1.0%)
+	0	1 (1.0%)	0	0
Total	44(44.5%)	30(32.6%)	9 (9.7%)	12 (13.0%)
Grand total = 92(100%), +++ = 1-10 parasites in every high power field; ++ = 11-100 parasites per 100 high power fields; + = 1-10 parasites per 100 high power fields				

**Table 5:** Malaria co-morbidity with non-communicable diseases.

## Discussion

This study has shown a high prevalence of malaria among the geriatrics, in South- Eastern, Nigeria. The prevalence was high among both sexes and across all the age groups with slightly higher infections observed among women and those between 61-70 years. Malaria in the study area is hyper endemic, with all year round transmission [10]. Most malaria management and control programmes are concentrated on under five children and pregnant women [11,12]. This has also made the older populations to care less about the illness since it is not generally considered as serious. Albeit, everyone is equally exposed to transmission through the bite of the mosquito vector, but in situations where people in the older populations do not make efforts at prevention as is done in younger children, prevalence is expected to be high. In general therefore, gender infection of malaria also reflects access of mosquito to the people as well as the failure of vulnerable populations to use proven and effective interventions for prevention, control and treatment [13]. The mistake is that the geriatric population in malaria endemic areas are not seriously considered to be vulnerable to the disease.

As people enter old ages it seems their health and self- definitions of old age becomes decreasingly multifaceted and increasingly related to health status [14]. Therefore a lot of changes begin to set in that may make them vulnerable to complications or more severe outcomes of infectious diseases. It is evident in this study that malaria infections

were associated with complications such as cerebral involvement, renal involvement and Anaemia. Previous studies have also reported similar complications among people of older age groups [15,16]. Respiratory disorder has also been associated with malaria in older people [17]. In most malaria endemic areas, such complications are not regarded to be as a result of malaria infections; therefore their aetiologies are relatively unknown. But it is important to note that high parasite intensity could lead to degradation of the haemoglobin which that could lead to anaemia. Sequestration could also result in cerebral and respiratory damages. One of the reasons why older people are not particularly thought to be vulnerable to malaria complications is because it is believed that as one advances in age, partial immunity is developed against malaria. However, it is known that immune response to malaria comes along with products like cytokines involved in its pathology [18]. Furthermore the increased hormonal activities in these age groups may also be involved in severe malaria outcomes [19].

Recently, there is increased prevalence of geriatric non communicable diseases in tropical malaria endemic areas particularly diabetes, hypertension etc. These diseases are associated with possible immunosuppressive functions likely to affect the integrity and sustenance of partial immunity like that developed against malaria [20]. This study confirms the prevalence of co-morbid malaria and such non communicable diseases. Similar report has also been made elsewhere [21]. This situation is rather complex because it may go both ways. While it is possible that impaired immunity as a result of non-

communicable disease pathologies could increase susceptibility to severe infections including malaria, it is also possible that high malaria prevalence and intensity could make one susceptible to non-communicable diseases. The aetiology of these non-communicable diseases in tropical endemic areas is largely unknown [21]. However, it is known that malaria parasite activity is associated with oxidative stress in the blood and other associated tissues. Frequent exposure to the parasite without control and increased parasite intensity will only translate to oxidative stress. Increased oxidative stress on the other hand triggers inflammation, contributing significantly to the pathology of non-communicable diseases by inducing a lot of metabolic disturbances as a result of the activities of reactive oxygen species [22]. The incidence of diseases related to such metabolic disturbances for instance diabetes, renal diseases; cancers are rising and are mostly age dependent. In addition immune response to malaria is associated with the release of pro inflammatory cytokines that are associated with the compromise of normal body metabolism that could result in non-communicable disease conditions [23]. Therefore it is plausible to suppose that changes resulting from constant exposure to infectious diseases such as malaria without adequate management and control could therefore have pathophysiological impacts associated with non communicable disease conditions and outcomes.

## Conclusion

Malaria remains a major public health problem in endemic areas and exposure to it is not limited to under-fives and pregnant women alone. In fact due to neglect, the elderly ones are even more exposed now. The older adults undergo physiological changes that may make them very susceptible to malaria complications. Increased intensity of the disease may also make them susceptible to non-communicable diseases. People in these age groups play very important roles in the family and economy of developing countries of which many are malaria endemic. Frequent malaria illness among them will also hinder them from playing their roles in the family and economic development of their societies. It is therefore important that malaria control programmes should be all inclusive, embracing all age categories including the older adults. Serious attention should be paid on the aetiologies of non-communicable diseases in infectious disease endemic areas, in order to ascertain possible relationships between them.

## References

1. Hay SI, Guerra CA, Tatem AJ (2004) The global distribution and population at risk of malaria: past, present and future. *Lancet Infectious Diseases* 4: 327-336.
2. Greenwood B (1999) Malaria mortality and morbidity in Africa. *Bulletin of the World Health Organization* 77: 617-618.
3. Fairhurst RM, Nayyar GM, Breman JG (2012) Artemisinin-resistant malaria: Research challenges, opportunities, and public health implications. *American Journal of Tropical Medicine and Hygiene* 87: 231-241.
4. Federal Ministry of Health (FMOH) (2012) National anti-malarial treatment policy and Guidelines. Federal Ministry of Health, National Malaria and Vector Control Division Abuja-Nigeria.
5. Drakeley CJ, Corran PH, Coleman PG (2005) Estimating medium and long-term trends in malaria transmission by using serological markers of malaria exposed. *Proc Natl Acad Sci U S A* 102: 5108-5113.
6. Adefioye OA, Adeyeba OA, Hassan WO, Oyeniran OA (2007) Prevalence of Malaria Parasite Infection among Pregnant Women in Osogbo, Southwest, Nigeria. *American-Eurasian Journal of Scientific Research* 2: 43-45.
7. Johnson RC, Scheni RF (2011) Early-life origins of adult disease: National longitudinal population, Based study of the United States. *American Journal of Public health* 101: 2317-2324.
8. Imam TS, Indabawa JI (2011) Prevalence of Anemia associated with malaria among patients in the extreme age groups attending three selected hospitals in Kano. *Biological and Environmental Science Journal* 4: 137-147.
9. Idowu OA, MafianaLaye O (2008) Perceptions and home management practices of malaria in some rural communities in Abeokuta, Nigeria. *Travel Medicine and Infectious Disease* 6 : 210-214.
10. Ogbusu FI, Nwoke BE, Njoku AJ, Anosike JC, Uwaezuoke JC (2004) Prevalence of malaria among pregnant women in Owerri municipality, Imo State, Nigeria. *African Journal of Applied Zoology and Environmental Biology* 6: 31-35.
11. Kay H (2009) *Aging by the Book: The Emergence of Midlife in Victorian Britain*. H-Albion.
12. Brubaka TH (2006) The stereotype of "old": A review of an alternative approach. *Journal of Comparative Family Studies* 31: 441-447.
13. Rogerson SJ, Chaluluka E, Kanjala S (2001) Intermittent sulphadoxine pyrimethamine in pregnancy: effectiveness against malaria morbidity in Blantyre, Malawi, in 1997-99. *Trans R Soc Trop Med Hyg* 94: 549-553.
14. Breman JG, Egan A, Keusch G (2001) The intolerable burden of malaria: a new look at the numbers. *American Journal of Tropical Medicine and Hygiene* 64: 4-7.
15. Higgias SJ, Kain KC, Liles WC (2011) Immunopathogenesis of falciparum malaria: Implications for adjunctive therapy in the management of severe and cerebral malaria. *Expert Review on Anti-infective Therapy* 9: 803-819.
16. Lampietti J (2012) Gender and preferences for malaria prevention in Tigray, Ethiopia. Policy and research report on gender and development working paper series.
17. Lena S (2011) The case for the use of oxidant antagonists as an adjunctive therapy for cerebral malaria. PPAR Research.
18. Muxel SM, Freitas-do-Rosario AP, Zago CA, Castillo-Mendez SI (2011) The spleen CD4 T cell response to blood stage Plasmodium chabaudi malaria develops in two phases characterized by different properties. *PLoS ONE*. 6: e22434.
19. Klein SL (2004) Hormonal and immunological mechanisms mediating sex differences in parasite infection. *Parasite Immunology* 26: 247-264.
20. Hunter DJ, Srinath R (2013) Noncommunicable Diseases. *New England Journal of Medicine* 369: 1336-1343.
21. Remais JV, Zeng G, Li G, Tian L, Engelgau MM (2013) Convergence of non-communicable and infectious diseases in low- and middle-income countries. *International Journal of Epidemiology* 42: 221-227.
22. Gottlieb MA, Morassutti AL, da Cruz IM (2011) Epidemiological transition, oxidative stress and chronic non-communicable diseases. *Scienza Medica* 21: 132-141.
23. Camps J, Garcia-Heredia A (2014) Introduction: oxidation and inflammation, a molecular link between non-communicable diseases. *Adv Exp Med Biol* 824: 1-4.