# ∂ OPEN ACCESS Scholars International Journal of Obstetrics and Gynecology

Abbreviated Key Title: Sch Int J Obstet Gynec ISSN 2616-8235 (Print) |ISSN 2617-3492 (Online) Scholars Middle East Publishers, Dubai, United Arab Emirates Journal homepage: https://saudijournals.com/journal/sijog/home

**Review Article** 

# Prevalence of Malaria Parasiteamia among Antenatal Pregnant Women Attending Selected Clinics in Hospitals within Abakiliki

Chukwurah Ejike Felix<sup>1,</sup> Obeagu Emmanuel Ifeanyi<sup>2\*,</sup> Okonkwo Chinaodili Edith<sup>3</sup> <sup>1</sup>Department of Haematology and Immunology, Faculty of Clinical Medicine, Ebonyi State University, Abakaliki, Nigeria <sup>2</sup>Department of Medical Laboratory Science, Faculty of Health Sciences, Imo State University, Owerri, Nigeria <sup>3</sup>Department of Medical Laboratory Science, Faculty of Health Sciences, Ebonyi State University, Abakaliki, Nigeria

\*Corresponding author: Obeagu Emmanuel Ifeanyi | Received: 18.03.2019 | Accepted: 26.03.2019 | Published: 31.03.2019 DOI: 10.36348/sijog.2019.v02i03.009

#### Abstract

Prevalence of malaria parasiteamia among pregnant women (age range 20-50 years) that attended mile four maternity hospital and Federal Teaching Hospital Abakiliki was analysed using standard laboratory procedure. The subjects were grouped based on age bracket, place of residence (rural, urban, and semi), occupation and prophylactic antimalaria drugs used. Thick film and giemsa staning was used for the malaria parasite identification and malaria parasite density calculation. Out of the 98 antenatal pregnant women blood sample analysed, 22(22%) of the blood film was positive for malaria parasite. Highest prevalence was seen in age group20-25 years of age range 5(5.1%) followed by 26-30 years of age range 9(9.2%) and 31-40 years of age range 5(5.1%). with the least prevalence in 41-50 years of age range 3(3.1%). At 95% confidence interval. X: cal 1.48 < X2tab7.815. P= 0.6869. By conventional criteria, the difference is considered not to be statistically significant. Parasite density count was done in all positive samples, which is compared and categorized according to settlement ranged from 01 to 400 parasite/ul. The highest parasite density was recorded among rural dwellers 3(01-100) parasite/ul. 2(101-200) parasite/ul 3(201-300) parasite/ul, 2(301-400) parasite/ul. this is followed by urban dwellers 1(01-100) parasite/ul. 1(101-200) parasite/ul 4(201-300) parasite/ul. 1(301-400) parasite/ul, with least occurrence among semi urban dwellers 1(01-100) parasite/ul, 2(101-200) parasite/ul and 3(301-400) parasite/ul respectively. Prevalence of malaria parasite based on socio demographic profiles, analysis of occupational based prevalence showed highest prevalence among farmers 7(30.4%). followed by unskilled labourer 8(25.8%). then civil servants 4(23.5%) with least occurrence among skilled labourers. Malaria parasite prevalence based on settlement indicated highest occurrence among rural dwellers 11(27.5%), followed by urban dwellers 5(20.8%) with least occurrence among semi urban dwellers 6(17.6%). Analysis of prophylaxis based parameter indicated a prevalence of 14(25.0%) (Those who are not on prophylaxis) compared to a prevalence of 6(19.0%) among those not on prophylaxis. Keywords: Prevalence, Malaria Parasiteamia, Antenatal Pregnant Women.

Copyright © 2019: This is an open-access article distributed under the terms of the Creative Commons Attribution license which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use (Non-Commercial, or CC-BY-NC) provided the original author and source are credited.

#### **INTRODUCTION**

Malaria is a major threat to public health and socioeconomic development in sub-Saharan Africa. Pregnant women and children are most affected, resulting in significant maternal, perinatal and fetal morbidity and mortality. About one to three million children die each year, while about 24 million pregnancies are threatened in Africa as a result of malaria [1]. The impact of malaria in pregnancy is most dramatic in populations, where pregnant women have little or no pre-existing immunity. However, in areas of high and stable malaria transmission, which predominate in sub-Saharan Africa, the effects of ill health are particularly apparent in first and second malaria-exposed pregnancies. In multigravidae, maternal morbidity due solely to malaria is uncommon,

and the major detrimental effects of the infection are low birth weight and maternal anaemia [2].

Nigeria is a malaria holoendemic country, and pregnant women are especially vulnerable to malaria, which can adversely affect the outcome of pregnancy. Malaria endemicity and parity are two of the factors that can influence the effects of malaria on a mother and her baby. In malaria-endemic regions, even the babies of asymptomatic pregnant women may show intrauterine fetal growth retardation and low birth weight [3].

Malaria can be diagnosed parasitologically, using microscopy or rapid diagnostic tests (RDTs) to rule out other causes of fever-like symptoms, particularly in areas of high prevalence of HIV where HIV-infected patients have a high incidence of febrile illness. More sophisticated tests like PCR and enzyme linked immuno-sorbent assays (ELISAs) are generally restricted to research studies or high income countries. One of the challenges in using peripheral blood tests such as RDTs or microscopy is that they will not detect sequestered parasites in the placenta. The range of antimalarial drugs available for treating malaria in pregnancy is more restricted than for non-pregnant adults as these drugs must first be proven to be safe and efficacious for both the mother and fetus, and yet pregnant women are systematically excluded from clinical trials. This is primarily clue to the risks, costs and complexities of undertaking clinical trials in pregnant women. The gold standard for the clinical evaluation of the efficacy of an antimalarial drug in pregnancy is the WHO 42- to 63-day follow-up protocol, whereby women are followed throughout pregnancy and at delivery, and. ideally, the infant is followed throughout the first year of life, to fully evaluate safety and clinical outcomes in both mother and infant. Nigeria has the highest malaria burden in the world, accounting for about one quarter of deaths due to the disease in Africa. Almost everyone is at risk of malaria but pregnant women and children under five years of age are particularly vulnerable to progression to severe disease [4].

#### Aim

This study therefore, was designed to evaluate the prevalence of malaria parasites and parasite density in antenatal pregnant women among hospitals in Abakaliki.

Ν

Where:

N= sample size  $X^2$  = chi square for the specified confidence interval level at 1 degree of freedom N = population size ME= desired margin of error

# **Ethical Approval**

Before the commencement of this study, ethical approval was obtained from the ethical committee of the various hospitals and also, informed consent from the subjects was appropriately sought for.

#### **Principle and procedure Principle**

A drop of blood is placed on the centre of a slide and rotated anticlockwise and clockwise to form a ring, the blood is not fixed and the diluted Giemsa stain used to stain the film contain water which causes rupture/lysis of red blood cells, allowing the parasites to be exposed and white blood cells to be seen [5].

#### Objectives

The specific objectives of this Research include:

- To evaluate the prevalence of malaria parasite among pregnant women.
- To determine the parasite density of the positive case and categorized based on urbanization.

# **MATERIALS AM) METHOD**

#### Study area

This cross-sectional study was carried out among two selected government hospitals in Abakaliki, Ebonyi State capital.

### METHOD

#### Subjects Used

The subjects chosen were antenatal pregnant women attending Mile 4 and Federal teaching hospital (1) in Abakaliki.

# Sampling/Sample Size

5mls of blood samples was obtained by venepuncture using sterile needle and syringe. It was placed into ethylene diamine tetra acetic acid (EDTA) container, thoroughly mixed by gentle inversion and taken to the laboratory for examination. A total of 98 blood samples from antenatal pregnant women among hospitals in Abakaliki were used for this study. Sample size of 98 was constructed using 95% confidence interval with a margin of almost 13%. The formular used for this calculation was

# Procedure

 $\frac{X^2 \times N \times (1 - P)}{(ME^2 \times (N - 1) + (X^2 \times P) (1 - P)}$ 

#### Thick Film Examination

The method for thick blood films for detection of malaria parasite as described by Ochie and Kolhatkar [6] and Cheesbrough [7] was employed. Briefly, the procedure is as outlined.

- The EDTA anticoagulated blood was mixed adequately
- A drop of blood was placed on a clean grease free slide using a Pasteur pipette
- With the aid of a test tube bulb, the drop of blood was mixed round into a circle to avoid the red cells from forming marked Rouleaux which can cause the blood to be easily washed from the slide during staining.
- The blood was allowed to air-dry for 2hours, with the slide placed in a horizontal position.
- The stock Giemsa to be used was diluted in 1:10 in 7.2 pH buffered water.
- The slide was covered with the diluted Giemsa stain, with the slide placed on the staining rack.
- It was allowed to stay for 30 minutes.

- The stain was then washed from the slide with clean water.
- The back of the slide was wiped clean and allowed to air-dry.
- When the thick film was completely dried, a drop of immersion oil was placed on the film.
- The slide was mounted on the microscope stage and focused with xIO objectives.

Using x100 objectives, malaria parasites and pigments were examined. Malaria diagnosis was based

> Parasite/ $\mu$ l of blood (parasite density) = N x total WBC counts/ $\mu$ l (8000) leucocyte count (200)

Where N = the number of asexual parasitic forms present in as many microscopic fields as possible to count 200 leucocytes.

#### **Data Analysis**

The data generated from this study was analyzed using Chi- square with the aid of a statistical package known as Statistical Package for Social Sciences (SPSS). Results were presented in Tables and a confidence interval of P<0.05 was considered significant [8].

# **RESULT**

Of the 98 blood samples that was assayed for the presence of malaria parasites in antenatal pregnant women among various hospitals in abakaliki within the age range 20 -50 years, 22(22.%) of the blood film were positive for malaria parasite. Analysis of age related prevalence showed highest parasitaemia in the 26 - 30 years of age range 9(9.2%). followed by the 20 -25 years of age range 5(5.1%) and the 31 - 40 years of age range 5(5.1 %), with least prevalence in the 41 -50 vears of age range 3(3.1%). At 95% confidence interval. X'cal  $1.48 < X^{2}$ tab7.815. P= 0.6869. By conventional criteria, the difference is considered not to be statistically significant (Table-1).

on identification of asexual stages of Plasmodium on the thick blood smears. Film was reported as "malaria parasite not seen" (i.e., negative) after examining about 100 fields. Parasite density was measured by the number of parasites per microlitre of blood (thick film). The number of asexual parasitic forms (trophozoites and schizoms) present in as many microscopic fields as possible necessary to count 200 leucocytes was recorded. The standard value of 8.000 WBCµl was assumed as a multiplier in the parasitaemia expression below:

Parasite density categorized according to settlement ranged from 01 to 400 parasite/ul. The highest parasite density was recorded among rural dwellers 3(01 - 100) parasite/ul. 2(101-200) parasite/ul 3(201-300) parasite/ul. 2(301-400) parasite/ul. This is followed by urban dwellers 1(01-100) parasite/ul. 1(101-200) parasite/ul 4(201-300) parasite/ul. 1(301-400) parasite/ul. With least occurrence among semi urban dwellers 1 (01-100) parasite/ul, 2(101-200) parasite/ul and 3(301-400) parasite/ul respectively (Table-2).

Prevalence of malaria parasite based on socio demographic profile is presented in Table-3. Result obtained based on occupation among the antenatal pregnant women showed highest prevalence among farmers 7(30.4%), followed by unskilled labourers 8(25.8%). then civil servants 4(23.5%) with least occurrence among skilled labourers. Malaria parasite prevalence based on settlement indicated highest occurrence among rural dwellers 11(27.5%). followed by urban dwellers 5(20.8%) with least occurrence among semi urban dwellers 6(17.6%). Analysis of prophylaxis based parameter indicated a prevalence of 14(25.0%) (those who are not on prophylaxis) compared to a prevalence of 6(19.0%) among those not on prophylaxis.

Age (years)	Malaria parasite not seen (%)	Parasitaemia (%)	Total (%)	$\mathbf{X}^2$	P- value
20-25	17(17.3)	5(5.1)	22(22) A		
26-30	23(23.5)	9(9.2)	32(32.7)		
31-40	27(27.6)	5(5.1)	32(32.7)		
41 -50	9(9.2)	3(3.1) '	12(12.2)		
Total	76(77.6)	22(22.4)	98(100)	1.48	0.6869

Table-1: Prevalence of malaria	parasites ar	nong antenatal	pregnant	t women	based o	on age

X2cal = 1.48,  $X^{2}tab = 7.815$ , degree of freedom (d.f) = 3, P - value = 0.689

Table-1 shows analysis of age related prevalence showed highest parasitaemia in the 26 -30 years of age range 9(9.2%), followed by the 20 -25 years of age range 5(5.1%) and the 31- 40 years of age range 5(5.1%), with least prevalence in the 41 - 50 years of age range 3(3.1%). At 95% confidence interval. X<sup>2</sup>cal  $1.48 < X^{2}$ tab7.815. P= 0.6869. By conventional criteria, the difference is considered not to be statistically significant.

Settlement	Malaria	Malaria Parasite density (parasite/ul)				
	parasite not seen	parasite seen	U1-10U	101-200	2U1-3UU	3U1-40U
Rural	29(29.6)	11(11.2)	3	3	3	2
Urban	19(19.4)	5(5.1)	114	1		
Semi urban	28(28.6)	6(6.1)	1	2	-	3
Total	76(77.6)	22(22.4)	557	6		

Table-2: Parasite density level by settlement among antenatal pregnant women

Table-2 showsParasite density categorizedaccording to settlement ranged from 01 to 400parasite/ul. The highest parasite density was recordedamong rural dwellers 3(01-100) parasite/ul. 2(101-200)parasite/ul 3(201-300) parasite/ul. 2(301-400)parasite/ul. this is followed by urban dwellers 1(01-100)parasite/ul. 1(101-200) parasite/ul 4(201-300)parasite/ul. 1(301 - 400) parasite/ul. With least

occurrence among semi urban dwellers 1(01-100) parasite/ul. 2(101-200) parasite/ul and 3(301-400) parasite/ul respectively. The settlementment categorization was based on the census bureau's entertainment for classifying areas (www.census/main/www/pdf.htm accessed 18th November 2015.

Age (years)	Number examined	Number infected	Prevalence	
20-25	22	5	22.7	
26-30	32	9	28.1	
31-40	32	5	15.6	
41 -50	12	3	25.0	
Occupation				
Farmers	23	7	30.4	
Civil servants	17	4	23.5	
Skilled labour	29	-t J	10.3	
Unskilled labour	31	8	25.8	
Settlement				
Rural	40	11	27.5	
Urban	24	5	20.8	
Semi urban	34	6	17.6	
Prophylaxis				
Yes	42	8	19.0	
No	56	14	25.0	

Prevalence of malaria parasite based on socio demographic profile is presented in Table-3. Result obtained based on occupation among the antenatal pregnant women showed highest prevalence among farmers 7(30.4%). followed by unskilled labourer 8(25.8%). then civil servants 4(23.5%) with least occurrence among skilled labourers. Malaria parasite prevalence based on settlement indicated highest occurrence among rural dwellers 11(27.5%), followed by urban dwellers 5(20.8%) with least occurrence among semi urban dwellers 6(17.6%). Analysis of prophylaxis based parameter indicated a prevalence of 14(25.0%) (Those who are not on prophylaxis) compared to a prevalence of 6(19.0%) among those not on prophylaxis.

#### DISCUSSION

This study examined the prevalence of malaria parasites in antenatal pregnant women among hospitals in abakaliki. Overall prevalence of 22.4% was recorded in this study. Of the 98 blood samples that was assayed for the presence of malaria parasites in antenatal pregnant women among various hospitals in Abakaliki within the age range 20 -50 years, 22(22.%) of the blood film were positive for malaria parasite. Analysis of age related prevalence showed highest parasitaemia

in the 26 - 30 years of age range 9(9.2%). followed by the 20-25 years of age range 5(5.1%) and the 31 -40 years of age range 5(5.1%). with least prevalence in the 41 - 50 years of age range 3(3.1%). At 95% confidence interval. X<sup>2</sup> cal 1.48 < X<sup>2</sup> tab7.815. P= 0.6869. By conventional criteria, the difference is considered not to be statistically significant.

This however disagree with the report of Singh *et al.*, [9] who reported prevalence of peripheral parasitaemia was low: 1.3% (35/2696) among women at ANCs and 1.9% at DUs (19/1025). The higher prevalence of malaria and its sequelae in pregnancy in this study therefore may be due to associated low health literacy, lack of access to vector control methods, or limited access to antimalarial drugs, or it may represent some local ecology unable to be identified in this study.

Parasite density categorized according to settlement ranged from 01 to 400 parasite/ul. The highest parasite density was recorded among rural dwellers 3(01-100) parasiie/ul, 2(101-200) parasite/ul 3(201-300) parasite/ul. 2(301-400) parasite/ul, this is followed by urban dwellers 1(01-100) parasite/ul. 1(101-200) parasite/ul 4(201-300) parasite/ul. 1(301-400) parasite/ul, with least occurrence among semi urban dwellers 1(01-100) parasite/ul, 2(101-200) parasite/ul and 3(301-400) parasite/ul respectively. Result obtained based on occupation among the antenatal pregnant women showed highest prevalence among farmers 7(30.4%). followed by unskilled labourer 8(25 8%). then civil servants 4(23.5%) with least occurrence among skilled labourers.

The age of the subjects in the study was between 20 - 50 years which is similar to earlier finding in Kano. Nigeria [10]. The prevalence of malaria parasitaemia is usually higher in the rural areas where mosquito breeding and transmission is intense. The fact that majority of the study population do not have tertian' education may have contributed to the high parasitaemia in this study. High standard of education usually affect health awareness and therefore has a positive impact on health [11]. This may also have contributed to the high level of parasitaemia obtained in this study since they were probably not better informed about vector control such as the use of insecticide treated nets [11].

Malaria parasite prevalence based on settlement indicated highest occurrence among rural dwellers 11(27.5%). followed by urban dwellers 5(20.8%) with least occurrence among semi urban dwellers 6(17.6%). Analysis of prophylaxis based parameter indicated a prevalence of 14(25.0%) (those who are not on prophylaxis) compared to a prevalence of 6(19.0%) among those not on prophylaxis. In Enugu South eastern Nigeria reported positive influence among pregnant women attending antenatal care have been reported. However, it is contrary to finding by Arnolu et al., [5] who reported that previous use of antimalarial drug before booking had no significant effect on the prevalence of malaria parasitaemia.

In a study conducted in Lagos of formal education on the use of maternity services. Some of the women took antimalarial since they were learned and knew some of the antimalarial available over the counter in Nigeria. This may explain why this group of women had lower parasitaemia as noted in this'study. This is corroborated by the finding by Gajida *et al.*, [10] who reported that self-treatment of malaria led to a reduction in prevalence of parasitaemia.

# **CONCLUSION**

Given the overall high prevalence of malaria, a strategy of enhanced anti-vector measures coupled with intermittent screening and targeted treatment during pregnancy should be considered for preventing malariaassociated morbidity in central India.

#### REFERENCES

1. Collins, W. E. (2012). *Plasmodiinn knowlesi:* a malaria parasite of monkeys and humans. *Annals Review of Entomolog*, 57(115):107-121.

- Bouyou-Akotet, M. K., Ionete-Collard, D. E., Mabika-Manfoumbi, M., Kendjo, E., Matsiegui, P. B., Mavoungou, E., & Kombila, M. (2003). Prevalence of Plasmodium falciparum infection in pregnant women in Gabon. *Malaria journal*, 2(1), 1-18.
- 3. Enato, E. F., Okhamafe, A. O., Okpere, E. E., & Oseji, F. I. (2007). Prevalence of malaria during pregnancy and antimalarial intervention in an urban secondary health care facility in Southern Nigeria. *Medical Principles and Practice*, *16*(3), 240-243.
- World Health Organization. (2014). WHO Policy 4. Brief for the implementation of intermittent preventive treatment of malaria in regnancy using stilfadoxine-pyrimethamine (IPTp-SP). WHO Global Malaria Programme. WHO Department of Reproductive Health and Research, and WHO Department of Maternal. Newborn. Child and Adolescent Health. Available at: www.who.int/maIaria/publications/atoz/Policy bri ef\_IPTpP\_implementation\_l lapr il2013.pdf.pdf. Available at: Accessed 12 November 2015.
- Anorlu, R. I., Odum, C. U., & Essien, E. E. (2001). Asymptomatic malaria parasitaemia in pregnant women at booking in a primary health care facility in a periurban community in Lagos, Nigeria. *African journal of medicine and medical sciences*, 30, 39-41.
- 6. Ochei, J., & Kolhatkar, A. (2000). Medical Laboratory Science. Theory and Practices. Tata McGraw-Hill Publishers. New Delhi.
- 7. Cheesbrough, M. (2005). District Laboratory Practice in Tropical Countries. Part 1. Cambridge University Press. New York.
- Ekpuka, B. M., Okogun, G. R., Obodo, B. N., Itua, E. E., Olagboye, J. A., & Obhakhan, J. O. (2013). The Prevalence of Malaria Parasitic Infections in Cord Blood: Association with Some Socio Demographic Profile. *International Journal of Basic, Applied and Innovative Research*, 2(2), 40-45.
- Singh, N., Singh, M. P., Wylie, B. J., Hussain, M., Kojo, Y. A., Shekhar, C., ... & Hamer, D. H. (2012). Malaria prevalence among pregnant women in two districts with differing endemicity in Chhattisgarh, India. *Malaria journal*, 11(1), 274-277.
- 10. Gajida, A. U., Iliyasu, Z., & Zoakah, A. I. (2010). Malaria among antenatal clients attending primary health care facilities in Kano state, Nigeria. *Annals* of *African medicine*, 9(3):188-193.
- 11. Isah, A. Y., Amanabo, M. A., & Ekele, B. A. (2011). Prevalence of malaria parasitemia amongst asymptomatic pregnant women attending a Nigerian teaching hospital. *Annals of African medicine*, 10(2).