

PREVALENCE OF MALARIA AND INTESTINAL PARASITE AMONG CHILDREN IN MUSLIM SCHOOLS, PORT HARCOURT, RIVERS STATE, NIGERIA

ABSTRACT

Introduction: malaria and intestinal helminthes infections major public health problem in low and middle-income countries affecting over 2 million people across the globe.

Objectives: This survey was done to determine the prevalence of malaria and intestinal parasites among Muslim pupils In Port Harcourt, Rivers State.

Methodolgy: Four Muslim schools: kab model school Rumuagolu, Manbaul hikma Eliozu, An-nur school Rumuodomaya and Al-ameen school Eneka. One hundred and fifty (150) blood and stool samples were collected randomly from the pupils (70 males and 80 female; age range 5-12years) and examined between the Months of May to July 2018.

Results: The overall prevalence of 50.7%, and 43.3% was recorded for malaria and intestinal helminthes respectively. The prevalence for malaria was 32.5%, 63.0%, 58.0% and 50.0%. Intestinal helminthes was 30%, 53.3%, 42.0% and 53.0% for kab, Manbaul hikma, An-nur and Al-ameen Schools Respectively. The frequency of parasites encountered was as follows, *Strongyloides stecoralis* 8.7%, *Ascaris lumbricoides* 14.0% ,*Enterobious vermicularis* 1.3%, *Necator americanus* 5.3%, *Ancylostoma duodenale* 4.7%, and *Trichuris trichuria* 9.3%. The prevalence for both malaria and intestinal helminthes infection was highest in females with 51.2% and 45.0% respectively. Pupils between ages 9-10 had the highest prevalence of 54.0% for malaria while Ages 7-8 had the highest prevalence of 47.7% for intestinal helminthes. Co-infection was 22.0%, 23.0%, 22.0% and 32.0% for kab, Manbaul hikma, An-nur and Al-ameen Schools Respectively.

Conclusions: The overall infection for malaria and intestinal helminthes was high. Therefore, regular de-worming of the pupils by parents, and health education are necessary to keep the prevalence in check. This study may also be used to predict risk for communities under consideration.

Key words: Malaria, helminthes, Muslim, Schools, Pupils

1. INTRODUCTION

Malaria is a life threatening disease caused by parasites that are transmitted to people through the bites of infected female anopheles mosquitoes. It is a vector borne infectious disease caused by an eukaryotic protista of the genus *Plasmodium*. Currently

there are five known species of *plasmodium* that are capable of infecting humans which are *Plasmodium falciparum*, *Plasmodium ovale*, *Plasmodium vivax*, *Plasmodium malariae*, and *Plasmodium knowlesi*. The latter causes malaria in humans but is also capable of infecting macaque monkeys. Of these species *P. falciparum* and *P. vivax* are the most virulent and were responsible for an estimated 660,000 deaths in 2016. *P. ovale* and *P. malariae* are generally less virulent than *P. vivax* and *P.falciparum*. Combined, these parasites had contributed to over 200 million reported cases of malaria infection in 2011 [1]. This is the most important of life threatening protozoan disease which is responsible for at least 750,000 deaths per year, mostly in young children in Africa [2, 3]. In 2013, there were 198 million cases and 584000 deaths as a result of malaria worldwide [4]. Over half of the world's population is at risk of malaria infection and as it stands, it's currently endemic in 109 countries in four continents, and of 500 million cases of malaria estimated to occur annually, approximately one million results in death. Global prevalence is concentrated around the equator in tropical and subtropical regions because of ample rainfall, warm temperatures and stagnant waters that provide ideal habitats for anopheles mosquito larva. Most of the fatalities are in children under the age of five years old and pregnant women [5]. Malaria accounts for at least \$12 billion in economic losses each year in Africa and a reduction in annual economic growth estimated at 1.3 percent [6]. In 2016, the region was home to 90% of malaria cases and 91% of malaria deaths. Total funding for malaria control and elimination reached an estimated US\$ 2.7 billion in 2016. Contributions from governments of endemic countries amounted to US\$ 800 million, representing 31% of funding. Malaria is the leading cause of child death in the country and around 250,000

Nigerian children die every year from the disease. While children under the age of five and pregnant women are particularly vulnerable, almost the entire population of Nigeria is at risk of contracting malaria [7].

Intestinal parasitic infections have been associated with problem such as vitamin deficiencies, inducing intestinal bleeding, protein energy malnutrition which in turn affects cognitive ability and intellectual development. Soil transmitted helminthes is a major public health problem in low and middle-income countries affecting about 2 million people across the globe [8]. Schistosomiasis along with other soil transmitted helminthes comprises over 40% of the illness caused by all tropical diseases apart from malaria [8]. Intestinal parasitic infections are particularly rampant in areas of the world where climate and poor sanitary conditions promote their survival, reproduction and transmission [9]. Intestinal parasitic infection may have serious consequences on human health, such as hepatomegaly, esophageal varices and bleeding [8]. Besides helminthes infected individuals could be susceptible to other infections such as malaria and HIV [10].

Intestinal parasites and malaria are diseases associated with poverty due to poor sanitary and environmental conditions, low level of education or illiteracy and low socio-economic class. They are among the common infection throughout the developing countries and have been associated with numerous deaths. The majority of deaths due to malaria is reported in Sub-Saharan Africa and occurs mostly in children below 15years [11].

This study was carried out to determine the prevalence of malaria and intestinal parasites among pupils of Muslim Schools in Port Harcourt Rivers State and to correlate the findings to child's hygiene status, habit and environmental sanitation.

2. METHODOLOGY

2.1 Study area

The study was conducted in Port Harcourt City; the capital of Rivers State, Nigeria. Obio/Akpor and Port Harcourt city Local governments are commonly referred to as Port Harcourt city Akpogomeh, [12]. It is situated on the bonny river and is located in the Niger delta. The study area lies within the latitude $4^{\circ}49'27\text{N}$ and longitude $7^{\circ} 2^{\circ}1\text{E}$ of the Greenwich meridian. It is situated some 60km from the open sea, which is immediately where the coastal marshes give way to the land of the interior [13]. Four (4) primary schools were randomly selected for this study; Al-ameen, kab model, An-nur and Man'baul hikma from Eneka, Rumuagholu, Rumuodomoya and Eliozu Communities.

2.2 Collection of stool sample

Children were given sterile vials containing 10% formalin for stool collection and adequately educated on how to get a little portion of their stool into clean and dried plastic container. They were advised that the stool specimen should be fresh, and not mix with urine. The collected samples were picked up between 8:00 to 10:00 in the morning. A total of 150 samples were collected from pupils through the assistance of the class teachers and was taken to the laboratory for examination.

2.3.1 Blood sample collection

The left thumb finger of each pupil was cleaned using a swab moistened with 70% alcohol and allowed to dry. Thereafter, it was pricked using a sterile lancet and

squeezed to obtain a small drop of blood which was smeared onto a slide to make thin blood smear.

2.4 Examination of Samples

2.4.1 Macroscopic examination of stool

This examination involves direct physical observation of the stool, the consistency of the stool and its general form such as hard stool (either formed or semi-formed stool), watery stool (unformed stool), bloody stool, diarrheic or stool containing mucus etc were observed.

2.4.2 Direct smear examination of stool

About 2grams of faeces was placed on a clean glass slide using an applicator stick. To it was added a saline solution to emulsify the specimen, so as to enhance the clarity of the cysts which may be observed. A cover slip was then placed on the preparation, avoiding the introduction of air bubbles and viewed with the light microscope using a x10 and x40 objectives respectively for egg/ova, of the parasites.

2.4.3 Formaldehyde-ether concentration method

This method was adopted by [14]. About 1 gram of faeces was emulsified in about 4ml of 10% formol water contained in a test tube. The formol water was prepared by mixing 50ml of strong formaldehyde solution with 150ml of distilled water. About 4ml of the formol water was added to the solution and mixed properly by shaking. The mixture was filtered into a test tube using a cloth guage and about 3-4ml of diethyl ether was added and shaken vigorously and allowed to stand for 2minutes. The mixture was then centrifuged at 1000revolutions per minute (1000rpm) for 3 minutes. Using a glass rod, the faecal debris from the side of the tube was loosened and the supernatants poured

off by inverting the tube. The tube was returned to its original upright position and the fluid from the side of the tube allowed draining to the bottom. The deposit was mixed by tapping the tube with the finger and using a Pasteur pipette, a drop of the sediment was applied on a microscope slide, mixed with Lugol's iodine and covered with a cover slip and examined under the microscope using X10 and X40 objectives respectively. The Ova and vermiform of the helminthes were identified with reference to Atlas of Parasitology [14].

2.4.4 Examination of blood

The blood was allowed to air dry with the slide in a horizontal position on a staining rack and placed in a safe place to avoid contaminants or flies. A small drop of methanol is added to the thin film, the thin film was fixed with methyl alcohol for 3-5 minutes and allowed to dry, and then it was stained with 10% Giemsa stain and was then taken to the Department of Animal and Environmental Biology laboratory, University of Port Harcourt for examination, where it was examined under oil immersion lens for detection of malaria parasites.

2.5 Statistical Analysis:

The statistical analysis was done using chi square data analyzing software.

3. RESULTS

A total of 150 blood and stool samples were obtained from four (4) Muslim primary schools during the study as follows; 40, 30, 50, and 30 for Kab model, Al-ameen, An-nur, and Manbal hikma respectively. Seventy-six (50.7%) was positive for malaria parasites as follows; 13 (32.5%), 15 (50.0%), 29 (58.0%), and 19 (63.3) for Kab model, Al-ameen, An-nur, and Manbal hikma respectively, while 65 (43.3%) was

positive for gastrointestinal helminthes as follows; 12 (30.0%), 16 (53.3%), 21 (42.0%), and 16 (53.3%) for Kab model, Al-ameen, An-nur, and Manbal hikma respectively (Table 1).

Table 1. Prevalence of malaria and gastrointestinal helminthes in the study area.

P-value= .87

One hundred and fifty pupils were examined for malaria and gastrointestinal helminthes in the study area, 76(50.7%) was positive for malaria as follows; 35(50.0%) and

Schools	No.examined	(%) No. positive for Malaria parasites	(%) No. positive for Gastrointestinal helminthes
Kab model	40	13 (32.5)	12 (30.0)
Al-ameen	30	15 (50.0)	16 (53.3)
An-nur	50	29 (58.0)	21 (42.0)
Manbal hikma	30	19 (63.3)	16 (53.3)
Total	150	76 (50.7)	65 (43.3)

41(51.2%) for males and females. While

65(43.3) was positive for gastrointestinal helminthes as follows; 29(41.1%) and 36(45.0%) for males and females respectively (Table 2).

Table 2. Sex related prevalence of malaria and gastrointestinal helminthes in the study area.

P-value= .04

Sex	No. examined	(%) No. positive for Malaria parasites	(%) No. positive for Gastrointestinal helminthes
-----	--------------	---	--

UNDER PEER REVIEW

Male	70	35(50.0)	29(41.1)
Female	80	41(51.2)	36(45.0)
Total	150	76(50.7)	65(43.3)

Age related prevalence of malaria parasites were as follows; 27(50.9%), 22(50.0%),

Age	No. examined	(%) No. positive for Malaria parasites	(%) No. positive for Gastrointestinal helminthes
-----	--------------	--	--

13(54.2%) and 14(48.3%) for ages 5-6, 7-8, 9-10 and 11-12, while prevalence for gastrointestinal helminthes were as follows; 24(45.3%), 21(47.7%), 10(41.7%) and 10(34.5%) (Table 3).

Table 3. Age related prevalence of malaria and gastrointestinal helminthes in the study area.

P-value= .87

5-6	53	27(50.9)	24(45.3)
7-8	44	22(50.0)	21(47.7)
9-10	24	13(54.2)	10(41.7)
11-12	29	14(48.3)	10(34.5)
Total	150	76(50.7)	65(43.3)

Sixty-five (43.3%) of the pupils were positive for gastrointestinal helminthes as follows; 21 (14.0%), 13 (8.7%), 2 (1.3), 8 (5.3%), 7 (4.7%), and 14 (9.3%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Enterobius vermicularis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. Twelve (30.0%) pupils were positive in Kab school as follows; 3(7.5%), 3(7.5%), 1(2.5%), 3(7.5%) and 2(5.0%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. Sixteen (53.3%) pupils were positive in Al-ameen as follows; 6(20.0%), 3(10.0%), 2(6.7%), 5(16.7%) and 5(16.7%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus* and *Trichuris trichuria* respectively. Twenty-one (42.0%) pupils were positive in An-nur school as follows; 6(12.0%), 6(12.0%), 1(2.0%), 3(6.0%), 3(6.0%) and 2(4.0%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Enterobius vermicularis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively, while 16 (53.3%) pupils were positive in Manbaul hikma school as follows; 6(20.0%), 1(3.3%), 1(3.3%), 2(6.7%), 1(3.3%), and

5(16.7%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Enterobius vermicularis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively (Table 4).

Table 4. Distribution of gastrointestinal helminthes species in the study area.

	No. examined	no. infected (%)	A.l	S.s	E.v	N.a	A.d	T.t	p-value
Location									
Kab	40	12(30.0)	3(7.5)	3(7.5)	(0.0)	1(2.5)	3(7.5)	2(5.0)	
Al-ameen	30	16(53.3)	6(20.0)	3(10.0)	(0.0)	2(6.7)	(0.0)	5(16.7)	.14
An-nur	50	21(42.0)	6(12.0)	6(12.0)	1(2.0)	3(6.0)	3(6.0)	2(4.0)	
Manbaul hikma	30	16(53.3)	6(20.0)	1(3.3)	1(3.3)	2(6.7)	1(3.3)	5(16.7)	
Total	150	65(43.3)	21(14.0)	13(8.7)	2(1.3)	8(5.3)	7(4.7)	14(9.3)	

Key:

A.l *Ascaris lumbricoides*

N.a *Necator americanus*

A.d *Ancylostoma duodenale*

E.v *Enterobium vermicularis*

T.t *Trichuris trichuria*.

S.s *Strongyloides stecoralis*

Thirty-seven (24.7%) pupils had co-infection of malaria and gastrointestinal parasites in the study area as follows; 12(8.0%), 8(5.3%), 6(4.0%), 7(4.7%) and 4(2.7%) for malaria and *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. Kab model school had co-infection of malaria and gastrointestinal parasites of 9 (22.0%) as follows; 2(5.0%), 2(5.0%),

1(2.5%), 2(5.0%), and 2(5.0%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. Al-ameen school had co-infection of malaria and gastrointestinal parasites of 5 (16.7%) as follows; 2(6.7%), 2(6.7%), 1(3.3%) and 1(3.3%) for *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus* and *Ancylostoma duodenale* respectively.

An-nur school had co-infection of malaria and gastrointestinal parasites of 16 (32.0%), as follows; 3(6.0%), 4(8.0%), 3(6.0%), 4(8.0%) and 2(4.0%) for malaria and *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. Manbaul hikma school had co-infection of malaria and gastrointestinal parasites of 7 (23.3%), as follows; 5(16.7%),1(3.3%) and 1(3.3%) for malaria and *Ascaris lumbricoides*, *Necator americanus* and *Trichuris trichuria* respectively.

Table 5. Distribution of co-infection of malaria and intestinal parasites in the study area.

Location	No.examined	No.infected (%)	MP+A.I	MP+S.s	MP+N.a	MP+A.d	MP+T.t	p-value
Kab model	40	9(22.5)	2(5.0)	2(5.0)	1(2.5)	2(5.0)	2(5.0)	
Al-ameen	30	5(16.7)	2(6.7)	2(6.7)	1(3.3)	1(3.3)	0(0)	.45
An-nur	50	16(32.0)	3(6.0)	4(8.0)	3(6.0)	4(8.0)	2(4.0)	
Manbaul hikma	30	7(23.3)	5(16.7)	0(0)	1(3.3)	0(0)	1(3.3)	
Total	150	37(24.7)	12(8.0)	8(5.3)	6(4.0)	7(4.7)	4(2.7)	

A.l *Ascaris lumbricoides*.

N.a *Necator americanus*

A.d *Ancylostoma duodenale*

E.v *Enterobium vermicularis*

T.t *Trichuris trichuria*

S.s *Strongyloides stecoralis*

Thirty-seven (24.7%) pupils had co-infection of malaria and gastrointestinal parasites in the study area as follows; 12(8.0%), 8(5.3%), 6(4.0%), 7(4.7%) and 4(2.7%) for malaria and *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. Twelve (17.1%) males had co-infection as follows; 4(5.7%), 2(2.9%), 2(2.9), 3(4.3), 1(1.4) for malaria and *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. While females had 25(31.3%) co-infection as follows; 8(10.0), 6(7.5), 4(5.0), 4(5.0) and 3(3.8) for malaria and *Ascaris lumbricoides*, *Strongyloides stecoralis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* respectively. (table 6).

Table 6. Sex related distribution of co-infection of malaria and intestinal parasites.

Sex	No.examined	No.infected (%)	MP+A.l	MP+S.s	MP+N.a	MP+A.d	MP+T.t	p-value
Male	70	12(17.1)	4(5.7)	2(2.9)	2(2.9)	3(4.3)	1(1.4)	0.04
Female	80	25(31.3)	8(10.0)	6(7.5)	4(5.0)	4(5.0)	3(3.8)	
Total	150	37(24.7)	12(8.0)	8(5.3)	6(4.0)	7(4.7)	4(2.7)	

A.l *Ascaris lumbricoides*.

N.a *Necator americanus*

A.d *Ancylostoma duodenale*

E.v *Enterobium vermicularis*

T.t *Trichuris trichuria*

S.s *Strongyloides stecoralis*

4. Discussion

The prevalence of malaria and gastrointestinal helminthes among pupils of four Muslim schools in Port Harcourt Rivers State was 50.7% and 43.3% respectively. This is finding on malaria is higher when compared to [15] who recorded a prevalence of (43.1%) malaria parasitemia in school children from two districts of Ghana but lower than the findings of [7] who recorded a prevalence of 63.3%. The 43.3% prevalence of gastrointestinal helminthes in line with [16,8 and 17]. But lower when compared to 72.2% finding by [18].

Manbaul hikma School had the highest prevalence of (63.0%) and (53.8%) for malaria and gastrointestinal helminthes respectively, this might be as a result of the surrounding bushes and water puddles which aid the proliferation of the vector agent mosquito. The high occurrence of intestinal helminthes infection in Manbaul hikma was not unusual because the area is rural. Parasitic diseases are known to be common in rural areas due to poverty, ignorance and low sanitary conditions [19, 20]. This is similar to the reports of [21] that carried out research to determine Risk factors and Socio-Demographic determinants of intestinal helminthes among children in school.

The least prevalence was recorded in Kab model school. The low prevalence may be attributed to high level of hygienic practices and regular deworming among the pupils which reduced transmission.

This is also in agreement with the works of [16, 15 and 17].

Males had higher prevalence of (33.3%) to females who had (27.8%) in gastrointestinal helminthes. This might be as a result of male children getting involved in activities of such as playing in contaminated soils most times bare footed and could have exposed them to infections. This finding is in line with [22, 23 and 24]. Females had a higher malaria parasite infection prevalence of (51.2%) than the males (50.0%). This might be as a result of hormones and other intrinsic factors such as menstruation in some of the females, which might affect the body's immune system due to loss of blood and also the involvement of females in more outdoor activities like washing, cleaning and cooking.

This is in contrast to the works of [25] where the prevalence of male to female is higher in a proportion of (59.8%:40.2%).

Age groups 9-10 and 5-6 had same prevalence of (42.9%) in helminthes infection followed by 7-8 (30.0%) and lastly 11-12 with a prevalence of (22.0%). This is in line with [24]. While age group, 9-10 has the highest prevalence of (54.2%) in malaria parasite infection, followed by ages 5-6 with prevalence of (50.9%). Age group of 7-8 has prevalence of (50.0%) while 11-12 had the lowest prevalence of (48.35%) ($P= .87$). These findings are not strange as the age ranges fall within susceptible groups.

Six species of intestinal helminthes was recovered from the study area. These includes; *Strongyloides stecoralis*, *Ascaris lumbricoides*, *Enterobius vermicularis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichura*. The prevalence of *Strongyloides stecoralis* in this study was (8.7%) which is in contrast with [26] who had a higher prevalence of (17.0%) in his study. The low prevalence of *S. stecoralis* might

be as a result of improving hygiene practices in the various schools. *Ascaris lumbricoides* had a prevalence of (14.0%). This is similar to the result obtained from the works of [27] who also encountered *Ascaris lumbricoides* in his research. *Enterobium vermicularis*, *Necator americanus*, *Ancylostoma duodenale* and *Trichuris trichuria* had prevalences of (1.3%), (5.3%), (4.7%), and (9.3%) respectively. This is in line with the reports of [28] who found same parasites and similar prevalence in their study.

Large mammals are prone to co-infection as a result of the possession of large body surface areas and appendages. Thirty- seven 24.7% Co-infection of malaria and intestinal parasites was recorded during the study with 32% occurring in An-nur. Males had a lower co-infection than female 31.3% as this might be as a result of their biology. Co-infection in this study is in line with the study of [29].

4.1 Conclusion

Having observed the high prevalence of gastrointestinal helminthes in the study area and the danger it portends for the pupils most of whom are ignorant and lack access good toilet facilities. Regular de-worming of the pupils by parents, and health education are necessary to keep the prevalence in check. This study may also be used to predict risk for communities under consideration.

Consent

Survey visits was made to identify the school and interact with the management and obtain their consent to participate in the study..

Ethical approval

The study protocol was approved by the research ethics committee of Faculty of science, University of Port Harcourt.

References

1. World Health Organization.(2012) Technical report series on research priorities for helminthes infection. WHO press, Geneva 107:16-171.
2. Greenwood D, Barer M, Slack R, Irving W. (2013). Medical microbiology 18th edition, China. 244-247
3. World Health Organization (2015). Achieving the malaria MDG target: reversing the incidence of malaria 2000-2015. Geneva: world health organization and the United Nations children's fund.
4. World Health Organization (2014). World malaria report. Geneva: world health organization.
5. Lamb TJ. (2012). Immunity to parasitic infection. 1st edition, New Delhi India.91-104.
6. National institute for allergy and infectious diseases (NIH-a) (2010). Malaria parasites.
7. Abah AE, and Temple B. Prevalence of Malaria Parasite among Asymptomatic Primary School Children in Angijama Community, Bayelsa State, Nigeria. *Trop Med Surg.* 2015; 4(1): 203.
8. World Health Organization (2013). Soil transmitted Helminthes fact sheets. 2010 report, WHO press, Geneva.107:16-171.

9. Alum A, Rubino JR, Ijaz MK. The global war against intestinal parasites should be use in a holistic approach? *Int. J infected disease*, 2010; 14:732-730.
10. Mulu A, Maier M, Liebert UG. Deworming of intestinal helminthes reduces HIV-1 subtype C viremia in chronically co-infected individuals. *International journal of infectious Diseases* 2013; **10**.(6): 02-016.
11. Murray CJ, Rosenfeld LC, Lim SS, Andrews KG, Foreman KJ. Global malaria mortality between 1980 and 2010: a systematic analysis. *Lancet* 2012; 379:413-431.
12. Akpoghomeh, O. (2001). Street map of Port Harcourt and its Environs. Kraft Books Ibadan. 214-312.
13. Ogba CO, (2002). Problem of Urban slum: A case study of Port Harcourt City. Unpublished Thesis, Enugu State University of technology Enugu. 102pp.
14. Cheesbrough M. District Laboratory Practice in Tropical Countries. Part 1 (2ed) Cambridge University Press 2005: 194-202.
15. Sarpong N, Owusu-Dabo E, Kreuels, B. Prevalence of malaria parasitemia in School Children from two districts of Ghana earmarked for indoor residual spraying; a cross-sectional study. *Malar J.* 2015; 14, 260.
16. Eze CN, Owhoeli O, Ganale SS. Assessment of intestinal helminthes in community school children of Khana Local Government Area, Rivers State. Nigeria. *Nigerian Journal of Parasitology* 2016; 37 (1).

17. Odu NN, Okonko IO, Ethi O. Study of neglected tropical diseases (NTD) Gastrointestinal helminthes among school children in Port Harcourt, Rivers State Nigeria. *Report and Opinion* 2011; 3 (9): 6-16.
18. Jonathan W, Abdullah AA, KS. Prevalence and Association of Parasitic Helminths among the Cross Section of Male and Female Gender Groups at University of Guyana, Georgetown, Guyana. *Research Journal of Parasitology* 2015; 10: 50-57.
19. World Health Organization (2002). Prevention and control of schistosomiasis and soil-transmitted helminthiasis: report of a WHO expert committee. World [Health](#) Organization Technical Report Series 912: 1-57.
20. Ukoli FMA. (1992). *Prevention and control of parasitic diseases in tropical Africa: the main issues*. University Press, Ibadan, Nigeria.
21. Aribodor DN, Okechukwu, PA, Eneanya OA, Etaga HO. prevalence and associated risk factors of intestinal infections among 5-14 years olds in Akpor community, Anambra State Nigeria. *Nigeria Journal parasitology* .2013; **34**(2):107-112.
22. Ali I, Mekete G, Wdajo N. Intestinal parasitism and related risk factors among students of Asendabo elementary and junior secondary school, South Western Ethiopia. *Ethiop J [Health](#) Dev*. 2011; 13: 157-161.
23. Adeyeba O. A., (2002). Intestinal parasitic infections among school children in a rural community Southwest Nigeria. *Nig J Parasitol* 23: 11-18.
24. Omotola OA, Ofoezie IE. Prevalence and Intensity of Soil Transmitted Helminths among School Children in Ifetedo, Osun State, Nigeria. *J Bacteriol Parasitol* (2019);10: 352.
25. Lannes, N.S.K.(2013) prevalence of malaria and intestinal helminth co-infection in children presenting with anemia in Freetown, Sierra Leone. Pp10.

26. Rahman M, Islam S, Masduzzaman M, Alam M, Chawdhury MNU, Ferdous J, *et al.* Prevalence and diversity of gastrointestinal helminthes in free-ranging Asian house shrew (*Suncus murinus*) in Bangladesh, *Veterinary World*, 2018; **11**(4): 549-556.
27. Bassey SE, Loko GT. A survey of helminth infections in Amakakala, Ogbio Local Government Area, Bayelsa State, Nigeria. *Biotechnol Res.* 2017; **3**(2):39.
28. Shehu MM, kabiru AU, Abubakar UK, Muhammad NM. Prevalence of Intestinal Helminthes Infections Among school children in Relation to occupation and toilet facilities in Maru Local Government Area, Zamfara State, Nigeria. *Journal of Biology, Agriculture and Health care.* 2017; **3**(19):87.
29. Atting IA, Ukpe IO, Usip LP. The prevalence of excreta-related soil-transmitted helminthiasis and the role of sanitation in its control in primary school children in Uyo metropolis, Akwa Ibom State, Nigeria. *Journal of Agric and Environmental Management.* 2013; **2**(11):341-346.