

PREVALENCE OF MALARIA AMONG THE PATIENTS LIVING IN AREAS OF DISTRICT SBA (SHAHEED BENAZIR BHUTTO) AND MIRPURKHAS.

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Abstract:

Objective: Malaria is a major cause of morbidity in the tropics and about 300 million cases were reported world wide in 2006 among the 100 species of genus plasmodia, the four species such as PL: falciparum, vivax, ovale and malaria causes malaria. The malaria is transmitted by the bite of female anopheles mosquitoes. .

Methodology: This descriptive and experimental study was carried out at department of pathology, People's University of Medical & Health Science (PUMHS), Nawabshah. The cases were collected from paediatrics & Medical outpatients departments of PUMHS Hospital Nawabshah and also from Muhammad Medical College (MMC) Hospital & Civil hospital Mirpurkhas (CHM) from January 2010 to December 2011. A total of 1200 patients were included. The prevalence of malaria on the basis of age, sex, areas of resident, and clinical finding of all patients were recorded and blood tests performed.

Results: Plasmodium Vivax in 70.8% of cases and Plasmodium Falciparum in 29.2% of cases.

Conclusion: In the areas (Nawabshah and Mirpurkhas), Plasmodium Vivax and Plasmodium Falciparum are the cause of Malaria.

Keywords: Malaria, Plasmodium, Pakistan, Nawabshah, Mirpurkhas.

Introduction:

Malaria is a major cause of morbidity in the tropics and about 300 million cases were reported world wide in 2006 among the 100 species of genus plasmodia, the four species such as PL: falciparum, vivax, ovale and malaria causes malaria. The malaria is transmitted by the bite of female anopheles mosquitoes.

The life cycle of malaria parasite is completed in human and female anopheles mosquito. The sporozoites are transmitted into blood by mosquito bite and they first infect the liver cells, then red blood cells by releasing merozoites which mature into the male and female gametocyte. When a mosquito bites a malaria infested human, these gametocytes in the mosquito's stomach unite together to form zygotes that develop into oocysts, which grow and rupture to release sporozoites and cycle starts again. Malaria causes haematological complications such as anemia, leucocytosis and thrombocytopenia, fever, rigors, sweating, body aches, headache, vomiting, pallor and splenomegaly. Death can occur due to complications including cerebral malaria and haematological complications. The microscope examination of peripheral blood for detection of malarial parasite and along with estimation of hemoglobin concentration, ESR and complete blood count are important laboratory investigations for the diagnosis of malaria and its haematological complication.

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The aim of this study was to evaluate prevalence of malaria among the patients living in areas of Districts of Nawabshah and Mirpurkhas. We also studied the haematological complication in these patients.

Patients and Methods:

This descriptive and experimental study was carried out at department of pathology, People's University of Medical & Health Science (PUMHS), Nawabshah. The cases were collected from paediatrics & Medical outpatients departments of PUMHS Hospital Nawabshah and also from Muhammad Medical College (MMC) Hospital & Civil hospital Mirpurkhas (CHM) from January 2010 to December 2011. A total of 1200 patients were included. The prevalence of malaria on the basis of age, sex, areas of resident, and clinical finding of all patients were recorded. To establish the laboratory diagnosis of malaria and its haematological complication, 3ml of venous blood sample was taken from each patient in the tubes containing EDTA and sent to the pathology department. Thick and thin blood smears were made on the clean glass slide and examined under the microscope for detection of various developmental stages of malarial parasites after staining with Giemsa stain. The (CBC) including Hemoglobin Concentration, Total Leucocyte count (TLC), Differential Leucocyte Count (OLC) and Platelet Count were determined by hematology analyzer from the blood sample. The ESR and Malaria Rapid Diagnosis Test were also done from the same blood sample.

Results:

A total of 1200 cases were studied. Among these 700 (58.3%) were children and 500 (41.7%) were adults. The age of these patients ranged between 5 and 65

year, and their mean age was (35+30), while male to female ratio was 1.7:1. Of 1200 patients, 400(32.3%) were residents of Nawabshah city and 800(66.7%) were resident of the rural areas of district SBA and Mirpurkhas. (Table I). The clinical finding in these patients as shown in table II were fever with rigor, sweating or feeling of cold and hot pallor, body ache and splenomegaly. The laboratory finding in these patients as shown in table III showed that the mean value of hemoglobin, RBC and platelet counts were significantly reduced, while WBC count with percentage of neutrophils and ESR were significantly increased. The microscope examination of stained thick and thin blood smears of all these patients showed Plasmodium Vivax in 70.8% of cases and Plasmodium Falciparum in 29.2% of cases. The ICT malaria test was positive for Plasmodium Vivax in 70.8% cases and 29.8% positive for Plasmodium Falciparum.

TABLE 1 Prevalence of malaria among the children and adults on the basis of age, sex and area OF residence of district SBA/Mirpurkhas

N=1200

AGE	SEX	RESIDANCE
Age in years 5-65 years mean age 35+3	Male 770 (64.1%) Female 430 (35.9%) Male to Female ratio 1.7:1	Rural 800 (66.7%) Urban 400 (33.3)
Adult	children	total
500 (41.7%)	700 (58.3)	1200 (100%)

N= Number of patients

TABLE-II Clinical finding in patients with malaria

N=1200

S#	Clinical Finding	No. of Patients	Percentage
1	Fever	1200	100%
2	Associated symptoms with fever like chill, sweating or feeling of coldness and hotness	980	81.7%
3	Baby Ache	750	62.5%
4	Headache	600	50.0%
5	Pallor	800	66.6%
6	Splenomegaly	300	25.0%

N=Total Number of patients

Table III Laboratory finding in patients with malaria and its hematological complication

N=1200

S. #	Laboratory Finding	No. of Patients	Percentage
1	Hemoglobin concentration 5.5 - 11.5/dl(8.5+3)	800	66.6%
2	ESR 40-11mn 37.5+72.5	1200	100.0%
3	Total Leukocytes count 6500 - 25000/cumcn (1625+8750)	900	75.0%
4	Red Cell Count 2.5-4.5 m/cumcn 3.5+1.0	700	58.3%
5	Differential Leukocytes count neutrophils 67-85% (80.5+5.5) Lymphocytes 10- 14%(11+3) Monocytes 10-18%(14+4) Eosinophils 2-4% (3+1)	1000 900 950 1200	83.3% 75.0% 79.1% 100.0%
6	Platelets count 40000-110000/cumcn (75000+35000)	750	62.5%
7	Microscopy P1:vivax P1:Falciparum	850 350	70.8% 29.2%
8	Malaria Diagnosis test Immunochrom atography (IC) Technique +ve for P1:vivx +ve for P1: Falciparum	350 850	70.8% 29.2%

Discussion:

Malaria remain a major cause of morbidity and mortality in Asian as well as African countries of the world and about 300-500 million cases of malaria while 1 million death case per year due to malaria occurs globally². 90% of malaria caused by plasmodium falciparum occurs in Africa³. Prevalence of malaria is common cause of death among the children and pregnant woman^{4,6}. Many Pakistani studies have shown that the ratio of P.vivax to Plasma Falciparum is at least 2 or just above that⁷⁻¹¹. The hematological complications of malaria such as anemia, leukocytosis with neutrophilia and thrombocytopenia have been reported. Anaemia is one of the most common complications in malaria that result from a combination of haemolytic mechanisms and accelerated removal of both parasitized and non-parasitized red blood cells, depressed and ineffective erythropoiesis. Age as a risk factor for thrombocytopenia and anaemia in children treated for acute uncomplicated falciparum malaria¹²⁻¹⁶. In our study, prevalence of P.vivax malaria (70.8%) is more common than the P. Falciparum. (29.2%) among the 700 children (58.3%) and 500 adults (41.7%) out of total 1200 cases in district SBA and Mirpurkhas. The significant clinical finding in these patients were fever with

rigors, pallor, body ache and headache. Hematological complication in these patients were anemia, leukocytosis with neutrophilia, ESR and thrombocytopenia and these were detected by hemoglobin, BSR and complete blood count estimations. In our study. Hemoglobin, platelet count and RBC count were significantly reduced while BSR, leukocyte count and percentage of neutrophils were significantly increased.

Conclusion:

The following conclusion has been made from the above study.

1. The prevalence rate of the malaria caused by *P. Vivax* is 2.5 times greater than the malaria caused by *P. Falciparum* among the children and adults in District Shaheed Benazeirabad.
2. The hematological complication such as anemia, leukocytosis with neutrophilia and thrombocytopenia among the children and adult were assessed by hemoglobin RBC estimation and complete blood count. It has been observed that haemoglobin, RBC count and platelet count were decreased while ESR and total leukocyte count with percentage of neutrophils in these Patients were increased.
3. Further studied are needed to determine the cold agglutination test, platelet aggregation test and serum interleukin level in the malaria.

References:

1. World Health Organization: World Malaria Report. 2010.
http://www.who.int/malaria/world_malaria_report_2010/en/index.html
2. Cotter C, Sturrock HI, Hsiang MS, Liu J, Phillips AA, Hwang J, Gueye CS, Fullman N, Gosling RD, Feachem RG: The changing epidemiology of malaria elimination: new strategies for new challenges. *Lancet* 2013,382:900.
3. WHO: Policy recommendation: Seasonal Malaria Chemoprevention (SMC) for *Plasmodium falciparum* malaria control in highly seasonal transmission areas of the Sahel sub-regions in Africa. WHO Global Malaria Program; 2012.
4. O'Meara WP, Beeman JG, McKenzie FE: The promise and potential challenges of intermittent preventive treatment for malaria in infants (IPTi) *Malar J* 2005, 4:33.
5. Cairns M, Roca-Feltrer A, Garske T, Wilson AL, Diallo D, Milligan PJ, Ghani AC, Greenwood BM: Estimating the potential public health impact of seasonal malaria chemoprevention in African children. *Nat Commun* 2012, 3:881
6. Beg MA, Sani N, Mehraj V, Jam W, Khan MA, Malik A, Menezes E, Hussain R, Smego R Jr: Comparative features and outcomes of malaria at a tertiary care hospital in Karachi, Pakistan *Jnt J Infect Dis* 2008,12:37-42.
7. Durrani AB, Durrani ill, Abbas N, Jabeen M: Epidemiology of cerebral malaria and its mortality *J Pak Med Assoc* 1997,47:213-215.
8. WHO: World malaria report. Geneva: World Health Organization; 2011.
9. Nizamani MA, Kalar NA, Khushk IA: Burden of malaria in Sindh, Pakistan: a two years surveillance report. *J Liaquat University of Med Health Sci* 2006, 5:762-83.
10. Ghanchi NK, Ursing J, Beg MA, Veiga MI, Jam S, Martensson A: Prevalence of resistance associated polymorphisms in *Plasmodium falciparum* field isolates from southern Pakistan *Malar J* 2011,10:18.
11. Rana MS, Tanveer A: Chloroquine resistance and *Plasmodium falciparum* in Punjab, Pakistan during 2~2001. *Southeast Asian J Trop Med Public Health* 2004, 35:288-291.
12. Lathia TB, Joshi R: Can hematological parameters discriminate malaria from non malarious acute febrile illness in the tropics? *Indian J Med Sci* 2004, 58:239-244.
13. Reyburn H, Mbakilwa H, Mwangi R, Mwerinde O, Olomi R, Drakeley C, Whitty CJ: Rapid diagnostic tests compared with malaria microscopy for guiding outpatient treatment of febrile illness in Tanzania randomized trial. *BMJ* 2007,334:403.
14. Wever PC, Henskens YM, Kager PA, Dankert J, Tom van Goo1: Detectetion of imported malaria with the cell -Dyn 4000 Hem.atology analyzer. *J Clin Microbiol* 2002, 40:47294731.
15. Price RN, Simpson JA, Nosten F, Luxemburger C, Hkirjaroen, terKuile F, Chongsuphajaisiddhi T, White NJ: Factors contributing to anemia after uncomplicated falciparum malaria. *Am J Trop Med Hyg* 2001, 65:614-22.
16. Wickramasinghe SN, Abdalla SH: Blood and bone marrow changes in malaria. In *Bailliere's Clin Hematol*. Volume 13. Harcourt Pub Ud; 2000::277-299.