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RESEARCH ARTICLE

STUDY OF CLINICAL PROFILE OF P.FALCIPARUM MALARIA IN TERTIARY CARE HOSPITAL

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ARTICLE	INFO
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ABSTRACT

Article History: Received 24th September, 2019 Received in revised form 18th October, 2019 Accepted 07th November, 2019 Published online 30th December, 2019 Malaria being one of the common infections in tropical country like india, this study is based on clinical profie, complications, laboratory investigations as well as prognosis of P. FALCIPARUM malaria.

Key Words:

Malaria, Falciparum, Fever.

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INTRODUCTION

Malaria is probably one of the oldest diseases known to mankind that has profound impact on our history. For centuries it prevented any economic development in vast regions of the earth.In South East Asia (the second most affected region in the world), India has the highest malaria burden (with an estimated 24 million cases per year) [Coatney, 1971], followed by Indonesia and Myanmar. In India, according to NVBDCP 2014 to September 2018 report, there were about 2.9 millions of new cases of plasmodium falciaparum malaria and 1620 deaths were recorded from 2014 to septemmber 2018, C.U.Shah hospital located in Surendranagar which is a rural locality. We get patients from surrounding rural localities as well nearby urban areas. With this background, we decided to conduct a study in this tertiary care center in Surendranagar to note the changing clinical profile of patients admitted with P.falciparum malaria along with its complications and outcome. All through study (in diagnosis and management) guidelines given by WHO were followed.

AIMS AND OBJECTIVES

- To study the different clinical profile in a patient with plasmodium falciparum malaria.
- To determine the hematological abnormalities in patients with plasmodium falciparum malaria.

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• To correlate the hematological and clinical profile with the severity and final outcome.

MATERIALS AND METHODS

This prospective study was conducted in C.U.Shah Medical Hospital during period of 1 year after obtaining approval from Saurastra University Ethical Committee. Total 100 patients were included in this study.

Methodology: Patients who were diagnosed having P.falciparum malaria by conventional microscopy, after obtaining written and informed consent from them, they were all screened for possible inclusion in the study based on inclusion and exclusion criteria.

Inclusion Criteria

- Patients belonging to either sex.
- Age > 18 years.
- Fever (axillary temp ≥ 37.5°C) at the time of examination or history of fever during preceding 48 hours.
- Patients with atypical symptoms
- Peripheral smear positive for Plasmodium falciparum.
- Patients giving informed/written consent.

Exclusion Criteria

- Age < 18 year.
- Presence of P. Vivex on peripheral smear.
- Patients diagnosed having mixed infections (P.vivax and P. falciparum).

Age In Year	Male (n= 66)	Female $(n=24)$	Total n=100(%)
18-30	27(40.90)	12(50%)	39(39%)
31-40	18(27.2)	03(12.5)	21(21%)
41-50	09(13.6)	04(16.6)	13(13%)
51-60	07(10.6)	01(4.16)	08(08%)
61-70	02(3.03)	03(12.5)	05(05%)
71-80	03(4.54)	01(4.16)	04(04%)
	Age In Year 18-30 31-40 41-50 51-60 61-70 71-80	Age In YearMale $(n=66)$ 18-3027(40.90)31-4018(27.2)41-5009(13.6)51-6007(10.6) $61-70$ 02(3.03)71-8003(4.54)	Age In YearMale (n= 66)Female (n= 24) $18-30$ $27(40.90)$ $12(50\%)$ $31-40$ $18(27.2)$ $03(12.5)$ $41-50$ $09(13.6)$ $04(16.6)$ $51-60$ $07(10.6)$ $01(4.16)$ $61-70$ $02(3.03)$ $03(12.5)$ $71-80$ $03(4.54)$ $01(4.16)$

Table 1. Age and Sex distributions

Table 2. Habitat

Age In Year	Male (n= 66)	Female (n= 24)	Total n=100(%)
18-30	27(40.90)	12(50%)	39(39%)
31-40	18(27.2)	03(12.5)	21(21%)
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71-80	03(4.54)	01(4.16)	04(04%)

Habitat	No. of Cases n=100
Urban	24(24%)
Rural	76 (76%)

Table 3. Clinical presentation of P.FALCIPARUM malaria

Type of Malaria	Total n=100(%)
Uncomplicated	65(65%)
Complicated	35(35%)

Table 4 (A): Symptomatology of uncomplicated P. falciparum malaria

Presenting Symptoms	Male (n=66)	Female (n=34)	Total n=100
Fever with chills	66(66%)	34(34%)	100(100%)
Headache and Bodyache	36(54.5%)	20(58.8%)	56(56%)
Vomiting	44(66.6%)	30(88.2%)	74(74%)
Pain in abdomen	30(45.4%)	12(35.2%)	42(42%)
Diarrhoea	03(8.8%)	02(3.03%)	05(05%)

Table 4 B. Symptomatology of complicated P. falciparum malaria

Presenting Symptoms	Male (n=66)	Female (n=34)	Total n=100(%)
Jaundice	06(9.09%)	08(23.5)	14(14%)
Altered Sensorium	05(7.5%)	04(11.7%)	09(09%)
Decreased Urine Output	08(12.1%)	04(11.7%)	12(12%)
Cough and breathlessness	06(9.09%)	02(5.8%)	08(08%)
Bleeding Manifestation	03(4.54%)	02(5.88%)	05(05%)

Table 5 (A): Duration of symptoms

< 3 days			– 7 days			>7 days		
Male	Female	Total	Male	Female	Total	Male	Female	Total
09	07	16(16%)	56	19	75(75%)	06	02	08(08%)

Table 5 B. Duration of symptoms and complications

< 3 days			3– 7 days			>7 days		
Uncomp-licated 08	Complic-ated 08	Total 16(16%)	Uncomp-licated 55	Comp-licated 20	Total 75(75%)	Uncomp-licated 02	Comp-licated 06	Total 08(08%)

Table 6. Clinical Signs

Signs	Male (n=66)	Female (n=34)	Total n=100(%)	
Splenomegaly	16(24.2%)	08(23.5%)	24(24%)	
Hepatomegaly	34(51.5%)	24(70.5%)	58(58%)	
Icterus	06(9.09%)	10(29.4%)	16(16%)	
Pallor	10(15.1%)	08(23.5%)	18(18%)	
Altered Higher Function	05(7.5%)	04(11.7%)	09(09%)	
Respiratory Rales	16(24.2%)	06(17.6%)	22(22%)	
Petechiea	03(4.54%)	02(5.88%)	05(05%)	
Oedema	06(9.09%)	03(8.8%)	09(09%)	

Table 7 A. Complication Patterns (According to WHO guidelines)

Criteria	Male (n=66)	Female (n=34)	Total (n=100)	
Hepatits (T. bilirubin>3mg/dl)	10(15.1%)	12(35.2%)	22(22%)	
Anaemia (Hb <5gm %)	14(21.2%)	12(35.2%)	26(26%)	
Cerebral malaria	08(12.1%)	04(11.7%)	12(12%)	
Renal failure (S. creatinine $> 3 \text{ mg \%}$)	12(18.1%)	06(17.6%)	18(18%)	
Spontaneous bleeding, DIC	02(3.03%)	01(2.94%)	03(03%)	
Pulmonary oedema, ARDS	04(6.06%)	06(17.6%)	10(10%)	
Circulatory collapse, Shock. (SBP <70 mmHg)	06(9.09%)	02(5.8%)	08(08%)	
Metabolic acidosis(PH < 7.25, bicarbonate < 15mmol/L)	04(6.06%)	02(5.8%)	06(06%)	
Hypoglycemia(RBS< 40 mg/dl)	02(3.03%)	03(8.8%)	05(%)	
Hemoglobinuria (black urine)	05(7.5%)	04(11.7%)	09(09%s)	

Table 7 B. Number of complications

Number of complications	Total(n=34)
One	08(23.5%)
Two	12(35.29%)
Three or more	14(41.17%)

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	<3 days (n=10)	3-/days (n=6/)	>7 days (n=24)	Total ($n=101$)
Hepatitis	02	18	02	22
Severe anaemia	06	14	06	26
Renal failure	00	10	04	14
Cerebral malaria	00	04	04	08
Spontaneous bleeding	00	01	02	03
ARDS	02	06	02	10
Shock	00	04	04	08
Hypogycemia	00	04	00	04
Acidosis	00	06	00	06

- Presence of concomitant bacterial infection.
- Patients who refused to give the written consent.
- Patients giving history of consumption of antimalarials or antibiotics with antimalarial activities in past 7 days.

CONCLUSION AND DISCUSSION

- This prospective study included 100 patients (66 males and 24 females). Thus male: female ratio was 2.8:1, reflecting male preponderance.
- Age ranged between 18 to 80 years. Age distribution patterns showing majority of patients 65% were below age 40 and prevalence declined with increasing age and fall sharply beyond 60 year.
- Majority of patients 76 % were from rural area.
- 65% patients presented with uncomplicated malaria. However 35% presented with severe and complicated malaria as per WHO guidelines.
- Fever with chills was found in all (100) patients, & out of them 74 percent patents had vomiting. Followed by headache, bodyache, abdominal pain and diarrohea.
- 34% presented with symptoms suggestive of complicated or severe forms of malaria which included jaundice (14%), altered sensorium (9%), decreased urine output(12%), cough & breathlessness(8%) and bleeding manifestations(18%) like hemoptysis and hematuria.
- Hepatomegaly was most common clinical sign found in 58% patients followed by Splenomegaly(24%), icterus(16%), pallor (18%), altered higher functions (9%), respiratory rales (22%).

- 16 % patients presented within 3 days starting of symptoms, of these 8 patients had complication. Number of Patients present in between 3-7 days of symptoms was 75%, of which 20 patients had complications. However 8 % who patients presented with duration of symptoms more than 7 days, 6 percent patient had one or more complications of malaria. Distribution pattern revealed that patients presenting with longer duration had highest rate of complication. These differences were statistically highly significant (P value < 0.001).
- Among 34% complicated P. Falciparum malaria patients most common complication observed was anaemia in 26 percent patient followed by severe Hepatitis (22%), renal failure (18%), spontaneous bleeding (14%), cerebral malaria(12%), ARDS in 10 percent patient, Hemoglobinuria in 9 percent patient and Hypoglycaemia in 5 percent patient.
- Majority of patient 23.5 % had 1 complication, 35.29 % patients had 2 complications and 41.17% patient developed more than 3 complications.(Out of 34 complicated)
- Majority of patients 91 % with one and two complications recovered completely. However 9 percent patient with 3 or more complications died due to multi organ failure.
- Thrombocytopenia (<1.5 lakh) was found in 94% patients. So it is most significant finding.
- Mean duration of hospital stay in 65 uncomplicated P. falciparum malaria patients had mean 4.18 ± 1.50SD. Among 35 complicated P.falciparum malaria patients duration of hospital stay were mean7.59 ± 2.91SD days.

- Mean of haemoglobin level in 65 uncomplicated P. falciparum malaria patients had mean 11.95 ± 1.98 SD. Among 35 complicated P.falciparum malaria patients level of haemoglobin were mean 6.59 ± 3.77 SD.
- Mean of platelet count in 65 uncomplicated P. falciparum malaria patients had mean 82151.52 ±56713.11SD. Among 35 complicated P.falciparum malaria patients had mean 40941.18 ± 46983.52 SD.

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REFERENCES

- Coatney GR., Collins WE., Warren M., Contacos PG. 1971."22 Plasmodium falciparum (Welch, 1897)". The primate malarias. Division of Parasitic Disease, CDC. p. 263.
- Rich, S. M., Leendertz, F. H., Xu, G., Lebreton, M., Djoko, C. F., Aminake, M. N., Takang, E. E., Diffo, J. L. D., Pike, B. L., Rosenthal, B. M., Formenty, P., Boesch, C., Ayala, F. J., Wolfe, N. D. 2009. "The origin of malignant malaria". Proceedings of the National Academy of Sciences. 106 (35): 14902–14907. Bibcode:2009PNAS..10614902R. doi:10.1073/ pnas.0907740106. PMC 2720412. PMID 19666593.Roberts &Janovy 2005.
- Perkins, D. J., Were, T., Davenport, G. C., Kempaiah, P., Hittner, J. B., Ong'Echa, J. M. 2011. "Severe malarial anemia: Innate immunity and pathogenesis". International Journal of Biological Sciences. 7 (9): 1427–1442. doi:10.7150/ijbs.7.1427. PMC 3221949. PMID 22110393.
- Perlmann, P; Troye-Blomberg, M (2000). "Malaria blood-stage infection and its control by the immune system". Folia biologica. 46 (6): 210–8. PMID 11140853.
- Vaughan, Ashley M., Aly, Ahmed S.I., Kappe, Stefan H.I. 2008. "Malaria Parasite Pre-Erythrocytic Stage Infection: Gliding and Hiding". Cell Host & Microbe. 4 (3): 209–218. doi:10.1016/j.chom.2008.08.010. PMC 2610487. PMID 18779047.
- Loy, Dorothy E., Liu, Weimin; Li, Yingying; Learn, Gerald H., Plenderleith, Lindsey J., Sundararaman, Sesh A., Sharp, Paul M., Hahn, Beatrice H. 2017. "Out of Africa: origins and evolution of the human malaria parasites Plasmodium falciparum and Plasmodium vivax". International Journal for Parasitology. 47 (2–3): 87–97. doi:10.1016/ j.ijpara.2016.05.008. PMC 5205579. PMID 27381764. WHO. Retrieved 17 March 2018.
- World Malaria Report 2008" (PDF). World Health Organisation. 2008. p. 10. Retrieved 2009-08-17.

- Baron, Christopher; Hamlin, Christopher 2015. "Malaria and the Decline of Ancient Greece: Revisiting the Jones Hypothesis in an Era of Interdisciplinarity". Minerva. 53 (4): 327–358. doi:10.1007/s11024-015-9280-7.
- Hempelmann, Ernst; Krafts, Kristine 2013. "Bad air, amulets and mosquitoes: 2,000?years of changing perspectives on malaria". Malaria Journal. 12 (1): 232. doi:10.1186/1475-2875-12-232. PMC 3723432. PMID 23835014.
- Nerlich, A. 2016. "Paleopathology and Paleomicrobiology of Malaria". Microbiology Spectrum. 4(6). doi:10.1128/microbiolspec.PoH-0006-2015. PMID 27837743.
- Lalchhandama, K. 2014. "The making of modern malariology: from miasma to mosquito- malaria theory" (PDF). Science Vision. 14 (1): 3–17. Archived from the original (PDF) on 2014-04-27.
- Cox, Francis EG. 2010. "History of the discovery of the malaria parasites and their vectors". Parasites & Vectors. 3 (1): 5. doi:10.1186/1756-3305-3-5. PMC 2825508. PMID 20205846.
- Baccetti, B 2008. "History of the early dipteran systematics in Italy: from Lyncei to Battista Grassi". Parassitologia. 50 (3–4): 167–172. PMID 20055226.
- Capanna, E 2006. "Grassi versus Ross: who solved the riddle of malaria?". International Microbiology. 9 (1): 69–74. PMID 16636993.
- Bruce-Chwatt, L.J. 1987. "Falciparum nomenclature". Parasitology Today. 3 (8): 252. doi:10.1016/0169-4758(87)90153-0.
- Christophers, R; Sinton, JA 1938. "Correct Name of Malignant Tertian Parasite". British Medical Journal. 2 (4065): 1130– 1134. doi:10.1136/bmj.2.4065.1130. PMC 2211005. PMID 20781927.
- Liu, W; Li, Y; Learn, GH; Rudicell, RS; Robertson, JD; Keele, BF; Ndjango, JB; Sanz, CM; et al. 2010. "Origin of the human malaria parasite Plasmodium falciparum in gorillas". Nature. 467 (7314): 420–5. Bibcode:2010Natur.467..420L. doi:10.1038/nature09442. PMC 2997044. PMID 20864995.
- Holmes, Edward C. 2010. "Malaria: The gorilla connection". Nature. 467 (7314): 404–405. Bibcode: 2010Natur.467..404H. doi:10.1038/467404a. PMID 20864986.
- Liu, W; Y Li, GH Learn, RS Rudicell, JD Robertson, BF Keele, JN Ndjango, CM Sanz, DB Morgan, S Locatelli, MK Gonder, PJ Kranzusch, PD Walsh, E Delaporte, E Mpoudi-Ngole, AV Georgiev, MN Muller, GM Shaw, M Peeters, PM Sharp, JC Rayner, BH Hahn, 2010. "Origin of the human malaria parasite Plasmodium falciparum in gorillas". Nature. 467 (7314): 420-5. Bibcode:2010Natur.467..420L. doi:10.1038/nature09442. PMC 2997044. PMID 20864995. https://en.wikipedia. org/wiki/Category:CS1 maint: Multiple names: authors 1 ist
