To Study the Incidence of Papilledema with Relation to Outcome in Children with Cerebral and Non-Cerebral Malaria

Authors
Dr Puneet Agrawal¹, Dr Jyoti Singh², Dr Sujata Lakhtakia³
¹Senior Resident, Department of Pediatrics, S.S. Medical College, Rewa
²Professor and Head, Department of Pediatrics, S.S. Medical College, Rewa
³Assistant Professor, Department of Ophthalmology, S.S. Medical College, Rewa

Corresponding Author
Dr Puneet Agrawal
Room no.8, PG Boys Hostel, S.S. Medical College, Rewa (M.P.), Pin – 486001 India
Phone: 9425721482, Email: dr.puneetagrawal@gmail.com

Abstract
Objective: To study the incidence of Papilledema with relation to outcome in children with Cerebral and Non-cerebral malaria
Design: Prospective, observational, cohort study, comprising of all malaria positive patients.
Setting: Department of Pediatrics, S.S.M.C and associated G.M. Hospital Rewa, Madhya Pradesh during the period of 1st August 2015 to 31st July 2016.
Participants: 100 consecutive patients with cerebral malaria and 100 patients with non-cerebral malaria were included in the study. All 200 cases were malaria positive. All children were evaluated by ophthalmologist for changes of retinopathy
Main Outcome Measure(s): Retinopathic changes and its correlation with mortality and duration of hospitalisation.
Results: In our study papilledema was seen in 25% and 6% of cases in CM and non-CM groups respectively. Mortality was 56% and 17% in patient having papilledema in CM and non-CM groups respectively. Mortality was significantly associated with papilledema in CM group.
Conclusions: In our study mortality was 56% and 17% in patient having papilledema in CM and non-CM groups respectively. This shows that in Cerebral malaria group a very high mortality was present in patients who had papilledema. Mortality was significantly associated with papilledema in CM group suggesting that patients having papilledema are at increased risk of dying. Therefore presence of papilledema in malaria can be used as a marker of serious disease and as an indication for intensive management of these cases.
Keywords: Malaria, Cerebral Malaria, Papilledema, Mortality.

Introduction
Malaria is highly prevalent in India and especially in Madhya Pradesh. It is a major health problem and leading cause of mortality and morbidity in this region. Cerebral malaria is one of the most common non-traumatic encephalopathy in the world[1]. It can be fatal in the absence of, prompt recognition of the disease and its complication,
and non-institution of active appropriate management of patients, especially in young children. Vindhya area of Madhya Pradesh is classified under hyper endemic zone for malaria [2,3]. By this study we aim to recognize cases early and to limit mortality and morbidity related to malaria. The detection of malarial retinopathy can be a good diagnostic and prognostic tool for cerebral malaria. There is a set of retinal abnormalities unique to cerebral malaria. These abnormalities include blurred disc margins, papilledema, retinal haemorrhages, retinal whitening, retinal oedema, vascular changes and soft exudates [4,5,6]. Of these retinal whitening and vascular changes are specific to cerebral malaria [7].

**Objective**

The objective of present study is to study the incidence of Papilledema with relation to outcome in children with Cerebral and Non-cerebral malaria.

**Methodology**

The study was carried out in the Department of Pediatrics, SS Medical College and associated GM Hospital, Rewa, Madhya Pradesh during the period of 1<sup>st</sup> August 2015 to 31<sup>st</sup> July 2016, after clearance from the Institutional Ethics Committee. The study design is prospective, observational, cohort study.

The study group comprised of 100 consecutive children with cerebral malaria presenting with acute febrile encephalopathy with Glasgow Coma Scale ≤10 with or without seizures. Hundred patients with non-cerebral malaria served as control who presented with acute febrile illness with no evidence of encephalopathy. All cases and controls were malaria positive either by peripheral smear examination or by rapid diagnostic kit.

All children were managed as per WHO standard guidelines for treatment of cerebral and non-cerebral malaria [8]. All cases and controls were evaluated by ophthalmologist for any changes of retinopathy within 24 hours of admission.

A detailed clinical evaluation including history and examination was carried out for all study participants at the time of admission. A base line evaluation in the form of blood sugar estimation (glucose strip), complete blood counts, liver function and renal function tests were done at the time of admission in all children. Using aseptic precautions, finger prick sample of blood was collected to prepare thick and thin smears of blood on glass slides, and evaluated for presence of any malarial parasite under oilimmersion, as per standard procedures.

Rapid diagnostic test kits were also used for the diagnosis of malaria. (SD BIOLINE Malaria Ag Pf./Pan kit manufactured by Standard Diagnostics (Alere) limited, Korea.)

Fundus examination was performed by ophthalmologist in all patients, after pupils were fully dilated using mydriatic eye drops. Presence of papilledema, retinal hemorrhages, vessel changes, peripheral whitening, and blurring of disc margins were noted and recorded separately, in addition to any other ophthalmologic abnormality.

All children were followed till discharge or death. The duration of hospitalisation was noted in both eventualities.

**Statistical analysis:** The data of the study were entered and analysed using the software Microsoft Excel 2013 for windows. Appropriate univariate and bivariate analysis were carried out using the Student t test for the continuous variable / proportion test (z test / t test) and two-tailed Fisher exact test or chi-square (χ²) test for categorical variables. The critical levels of significance of the results were considered at 5% i.e. P< 0.05 was considered significant.

**Results**

In our study 59% and 61% children were males in study and control group respectively. The incidence of malaria was minimum in below 6 months age in both groups while highest incidence was observed between 6 years-12 years (Table no 1).
Papilledema was seen in 25% & 6% of cases in CM and non-CM groups respectively. Mortality was 56% and 17% in patient having papilledema in CM and non-CM groups respectively. Mortality was significantly associated with papilledema in CM group.(Table no 2)

### Table 1 Sample characteristics

| 1. Gender | Cerebral malaria (CM) (n=100) | Non cerebral malaria (Non CM) (n=100) |
|-----------|-------------------------------|--------------------------------------|
| Male (M)  | 59 (59%)                      | 61 (61%)                             |
| 2. Age    |                               |                                      |
| <6 month  | 5 (5%)                        | 2 (2%)                               |
| 6 –60 mon | 27 (27%)                      | 29 (29%)                             |
| 6 yrs–12yrs | 47 (47%)                    | 51 (51%)                             |
| Above 12yrs | 21 (21%)                    | 18 (18%)                             |

### Table no. 2 Papilledema in relation to Outcome in Cerebral and Non cerebral malaria

| Fundus – Papilledema | Cerebral malaria (CM) (n=100) | Non cerebral malaria (Non CM) (n=100) |
|----------------------|-------------------------------|--------------------------------------|
|                      | Mortality | Alive | Total | Mortality | Alive | Total |
| Normal               | 11 (15%) | 64 | 75 (100%) | 6 (6%) | 88 | 94 (100%) |
| Present              | 14 (56%) | 11 | 25 (100%) | 1 (17%) | 5 | 6 (100%) |
| Total                | 25 (25%) | 75 | 100 (100%) | 7 (7%) | 93 | 100 (100%) |
| P value              | 0.0001 | 0.89 |

### Discussion

In our study papilledema was seen in 25% and 6% of cases in CM and non-CM groups respectively. Mortality was 56% and 17% in patient having papilledema in CM and non-CM groups respectively. Mortality was significantly associated with papilledema in CM group suggesting that patients having papilledema are at increased risk of dying (p=0.0001) but mortality was not significantly associated with papilledema in non CM group. Beare, et al. found papilledema prevalence of 15% in cerebral malaria[9]. Retinopathy was associated with subsequent death (relative risk, 3.7) and papilledema conferred the highest risk (RR=4.5) [10]. Lewallen found the relative risk of death in patients with papilledema was 6.7 times that in patients without papilledema and also found that patients with papilledema had poor outcome 5.2 times greater than those without this finding[11]. According to Hirneiss, papilledema was not commonly seen (8%) but was a poor prognostic sign[12].

### Conclusions

In our study papilledema was seen in 25% and 6% of cases in CM and non-CM groups respectively. In CM group mortality was observed in 56% of patients who had papilledema, compared to 15% who did not have papilledema. In Non CM group mortality was observed in 17% of patients who had papilledema, compared to 6% who did not have it. This shows that in Cerebral malaria group a very high mortality was present in patients who had papilledema. Mortality was significantly associated with papilledema in CM group suggesting that patients having papilledema are at increased risk of dying. Therefore presence of papilledema in malaria can be used as a marker of serious disease and as an indication for intensive management of these cases.

### References

1. Snow RW, Newton CRJC, Craig MH &Steketee RW. The public health burden of plasmodium falciparum malaria in Africa: deriving the numbers. Disease
2. Jain V, Avinash C, Pradeep K, Manmohan S, Mrigendra P, Rasik B, et al. Burden of cerebral malaria in Central India (2004–2007). Am J Trop Med Hyg. 2008;79:63642.
3. Singh J, Soni D, Mishra D, Singh HP, Bijesh S. Placental and neonatal outcome in maternal malaria: A prospective cohort study from Central India. Indian Pediatr. 2014;51:2858.
4. Kochhar DK, Shubhakaran, Kumawat BL, Thanvi I, Joshi A, Vyas SP. Ophthalmoscopic abnormalities in adults withfalciparum malaria. QJ Med. 1998;91:84552.
5. Poncet 1878. De La Retino-Coroidiste Palustre. Ann D'oculistique. 1878; 79:201–218.
6. Hero M, Harding SP, Riva CE, Winstanley PA, Peshu N, Marsh K. Photographic and angiographic characterization of the retina of Kenyan children with severe malaria. Arch Ophthalmol 1997;115: 997–1003
7. Beare NAV, Taylor TE, Harding SP, Lewallen S, Molyneux ME. Malarial retinopathy: a newly established diagnostic sign in severe malaria. Am J Trop Med Hyg 2006 75 790-797.
8. www.who.int/malaria
9. Beare NAV, Taylor TE, Harding SP, Lewallen S, Molyneux ME. Malarial retinopathy: a newly established diagnostic sign in severe malaria. Am J TropMedHyg2006 75 790-797.
10. Beare NA, Southern C, Chalira C, Taylor TE, Molyneux ME, Harding SP; Prognostic significance and course of retinopathy in children with severe malaria. Arch Ophthalmol. 2004 Aug;122(8):1141-7.