Acute disseminated encephalomyelitis after Plasmodium vivax infection: case report and review of literature

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Abstract

Acute demyelinating encephalomyelitis (ADEM) usually occurs after viral infections or vaccination. Its occurrence after Plasmodium vivax infection is extremely uncommon. We report the case of an 8-year-old girl who had choreo-athetoid movements and ataxia after recovery from P. vivax infection. Diagnosis of ADEM was made on the basis of magnetic resonance imaging findings. The child responded to corticosteroids with complete neurological recovery.

Introduction

Acute disseminated encephalomyelitis (ADEM) is a monophasic, immune mediated, inflammatory and demyelinating disease of the central nervous system (CNS) with diffuse neurological signs and multifocal white matter lesions on neuroimaging.1,2 It usually occurs within 2 to 4 weeks following a viral infection or vaccination.3,4 In some patients, no antecedent trigger can be identified. An autoimmune reaction of T-cells against myelin basic protein, triggered by viral infection or vaccination, has been suggested as a possible pathogenesis.5 Post-malarial ADEM has been previously reported as a complication of Plasmodium falciparum. To the best of our knowledge, only a single case of ADEM following treatment of Plasmodium vivax malaria has been reported in pediatric literature.6

Case Report

An 8-year old female child presented with complaints of fever and vomiting for 2 days and an episode of generalized tonic-clonic seizure on the day of admission. Her mother and sibling were also admitted to our hospital with diagnosis of severe malaria. Vitals revealed tachycardia and hypotension. On general physical examination, the child had mild pallor with no sign of icterus, cyanosis, edema or lymphadenopathy. Respiratory, cardiovascular and abdominal examinations were normal. Neurological examination showed that the child was drowsy with hypertonia, hyperreflexia and positive Babinski response. There were no meningeal or cerebellar signs.

Investigations showed hemoglobin 9.8 g/dL, white blood cell count 9.8×10^9/L and platelets 0.37×10^11/L. Peripheral smear examination showed multiple trophozoites of P. vivax along with thrombocytopenia. Differential leucocyte count was 60% neutrophils, 30% lymphocytes, 3% eosinophils and 7% monocytes. Rapid malaria antigen tests (RMAT) was also positive for P. vivax. Serum biochemistry including blood sugar, renal and liver function tests were normal. Cerebrospinal fluid (CSF) examination was normal.

Child was managed in intensive care unit with supportive care for hypotension with crystalloids and inotropes. In view of peripheral smear and RMAT suggestive of P. vivax, the child was started on intravenous artesunate (as per the guidelines of National Vector Borne Disease Control Programme, India, for severe malaria).7

The child responded well and was shifted to pediatric ward after one week of admission. Patient did not have any neurological deficit and repeat peripheral smears showed parasitological clearance. On day 15 of illness, the child presented with abnormal movements in form of ataxia and choreo-athetoid movements of limbs. There were no meningeal signs or cranial nerve palsies. Deep tendon reflexes were exaggerated and Babinski sign was present. Repeat peripheral blood smear examination did not show any malarial parasite. Magnetic resonance imaging (MRI) of brain with contrast showed high intensity signals in subcortical, cortical, left parietal, periventricular regions and pons (Figure 1). In view of temporal relationship and latency of symptoms and MRI findings, diagnosis of post malaria ADEM (secondary to P. vivax) was made. Patient was started on prednisolone at 2 mg/kg/day for 7 days and tapered over one week. Chorea-athetoid movements decreased and ataxia improved. After 2 weeks of discharge, patient had no neurological sequelae.

Discussion

Severe malaria is known to be classically caused by Plasmodium species, but in recent years, P. vivax is being increasingly recognized as a cause of severe malaria in children.8 ADEM has been reported in literature as a complication of falciparum malaria (Table 1).4,5,8-12 Only a single case of P. vivax malaria followed by ADEM has been reported in pediatric age group.9 This case reiterates the importance of neurological complications occurring even after recovery from P. vivax in concordance with the changing epidemiology of malaria.

Post-malaria neurological syndrome (PMNS) was first described by Nguyen et al. as symptomatic malaria infection (initial blood smear positive for asexual forms of parasite), whose parasites have cleared from the peripheral blood and, in cerebral cases, had recovered consciousness fully, who developed neurological or psychiatric symptoms within two months of acute illness.10 Schnorf, et al. classified PMNS into 3 subtypes according to severity: i) mild and localized encephalopathy, characterized by isolated cerebellar ataxia or postural tremor; ii) diffuse, but relatively mild encephalopathic form, characterized by acute confusion or epileptic seizures; and iii) severe, corticosteroid-responsive encephalopathy, characterized by motor aphasia, generalized myoclonus, postural tremor, and cerebellar ataxia, resembling ADEM.11 PMNS and ADEM have striking similarities e.g. period of latency, multifocal neurological deficits, response to steroids and good prognosis. Increase in levels of cytokines in cerebrospinal fluid, tumor necrosis factor-α and interleukins 2 and 6 have been noted in severe malaria and PMNS.15 The latency to neurological involvement after eradication of parasite and response to corticosteroids in our patient support an immunological mechanism.

References

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Conclusions

Recurrent or new appearance of neurological complications in a known patient of severe malaria should arouse suspicion of ADEM. Cranial MRI should be ordered for confirmation of the diagnosis. Definitive treatment should be instituted as the disease has a good outcome with negligible sequelae.

References

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Table 1. List of cases of acute demyelinating encephalomyelitis reported after recovery from malaria.

| First author   | Species          | Age of patient | Time of neurological complication after recovery from malaria | MRI finding (T2-DWI)                                                                 | Treatment                              | Outcome                      |
|----------------|------------------|----------------|-------------------------------------------------------------|--------------------------------------------------------------------------------------|----------------------------------------|-----------------------------|
| Mohsen⁴        | *P. falciparum*  | 30y            | 8 weeks                                                     | Hyperintensities in subcortical white matter of right frontal and temporal lobes and left cerebellar hemisphere | No steroids                            | Complete neurological recovery |
| Sharma⁹        | *P. falciparum*  | 20y            | 2 weeks                                                     | Hyperintensities of subcortical white matter, corpus callosum and midbrain           | I/V methylprednisolone for 3 days      | Complete neurological recovery |
| Rachita¹⁰      | *P. falciparum*  | 4y             | 1 week                                                      | Hyperintensities in cerebral hemisphere, subcortical white matter and midbrain       | I/V methylprednisolone for 3 days, then oral steroids | Complete neurological recovery |
| Agrawal¹¹      | *P. falciparum*  | 12y            | 16 days                                                     | Hyperintensities in periventricular white matter, centrum semiorale and genou of corpus callosum | I/V methylprednisolone for 5 days, then oral steroids | Complete neurological recovery |
| Koibuchi¹²     | *P. vivax*       | 24y            | 2 weeks                                                     | Hyperintensities in left cerebral cortex and subcortex                              | I/V methylprednisolone for 3 days, then oral steroids for 2 weeks | Complete neurological recovery |
| Goyal⁶         | *P. vivax*       | 18mo           | 1 week                                                      | Diffuse white matter hyperintensities of subcortical deep and periventricular white matter and both external capsules | I/V methylprednisolone for 3 days, then oral steroids for 2 weeks | Complete neurological recovery |

y, years; mo, months; MRI, magnetic resonance imaging.
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