Post encephalitic parkinsonism following dengue viral infection

B. V. K. M. Bopeththa* and U. Ralapanawa

Abstract

Background: Incidence of dengue fever as well as dengue hemorrhagic fever is increasing in Sri Lanka especially among elderly population. As the number of cases is rising, rare complications of dengue illness also can be seen in clinical practice when compared to the past few years. Prompt identification and treatment of such complications is challenging due to lack of awareness and unavailability of standard treatment.

Case presentation: 69 years old man presented with acute onset fever and was diagnosed as having dengue illness. On the 4th day of illness, the disease was progressed into dengue haemorrhagic fever and recovered uneventfully. Although he recovered from primary illness, his general condition continued to deteriorate due to new onset of features of parkinsonism. Cerebrospinal fluid analysis and electro encephalogram showed evidence of encephalitis. Cerebrospinal fluid analysis also revealed positive IgM antibodies against dengue virus. Then the diagnosis of post encephalitic parkinsonism following dengue viral infection was made and started on. He was started on SINEMET (carbidopa 10 mg and levodopa 100 mg) half tablet 6 hourly started. After 1 week of treatment he showed marked improvement and then patient was discharged with further follow up plan.

Conclusion: Although the management of dengue illness and dengue haemorrhagic fever is straightforward, recognition and treatment of rare complications like post encephalitic parkinsonism following dengue viral infection is difficult without great clinical suspicion.

Keywords: Dengue haemorrhagic fever, Viral encephalitis, Parkinsonism

Background

Currently dengue illness is the most common public health concern in Sri Lanka. Up to now (from 01/01/2017 to 24/07/2017) 105,153 cases have been reported [1]. In history, the first serologically confirmed case has been reported in 1962 and first outbreak occurred in 1965. First major epidemic occurred in 1989 and became endemic since then. Cyclical epidemics occurred from 2002 onwards [2]. Actually dengue infection has multi system involvement. Because of multisystem involvement, dengue infection can have variety of atypical presentations including neurological manifestations. In this case report we present a case with post encephalitic parkinsonism following dengue infection.

Case presentation

69 years old retired teacher presented to us with 3 days history of acute onset febrile illness with headache, myalgia, joint pain and loss of appetite. He is a known patient with Non Hodgkin lymphoma since 2013 and treatment has been completed. He was regularly followed up in oncology clinic at Teaching hospital Kandy, Sri Lanka. He is a non smoker as well as never used alcohol. He was not on medications in the recent past. He was well and active at home prior to this admission. On admission he was conscious, rational walked to the admission room in a normal way. Body temperature was 101 °F and mild dehydration was present. Peripheries were warm with good capillary refill. Pulse rate was 80 beats per min with blood pressure of 130/80 mmHg. Chest, abdomen and neurological examination were unremarkable. Full blood count on admission showed white cell count of 8.9/μL with 80% of neutrophils and 13% of lymphocytes. Hemoglobin level
was 10.2 g/dL. His platelet count was 200 cells/μL. Serum sodium level was 133 mmol/L and potassium level was 4.3 mmol/L. Serum creatinine was 84 μmol/L. C reactive protein level was within normal range. Liver enzyme results revealed aspartate aminotransferase (AST) level of 55 U/L and alanine aminotransferase (ALT) level of 26 U/L. Serum albumin and non fasting cholesterol levels were within normal range. Immediate ultrasound examination abdomen revealed no free fluid in the abdomen or pelvis. Dengue NS 1 antigen (non structural protein 1) was negative on admission.

Because of ongoing outbreak with high clinical suspicion, diagnosis of uncomplicated dengue fever was made and management started with intravenous normal saline, strict input and output chart maintenance and 6 hourly packed cell volume (pcv) monitoring according to national Dengue management guidelines.

On the 2nd day of admission (4th day of illness) he complained of postural symptoms, vomiting and increasing tiredness. On examination apart from mild dehydration other vital parameters were stable. With the suspicion of onset of critical phase ultrasound abdomen was performed and revealed thin layer of free fluid in hepatorenal pouch. The patient was taken into the high dependency unit and management was continued according to national guidelines. Throughout the critical phase his vital parameters, hourly urine output and hourly packed cell volume measurement remained stable. There was no indication for blood or blood products transfusion during this critical period. Table 1 demonstrates the laboratory results according to the hours spent in the critical phase. In addition to following, non fasting serum cholesterol was low and corrected serum calcium was within the normal range. Arterial blood gas analysis was normal.

On the 5th day of admission, Dengue IgM antibodies became positive and IgG was negative. Temperature also came to the base line. Although the patient recovered from dengue hemorrhagic fever, continued to deteriorate his general condition. His speech became slow and there was no energy in his purposeful movements. It took minutes to hours to get up from the bed. His care giver noticed increased stiffness of his arms and limbs. On examination he had expressionless face with cog wheel rigidity. He was severely bradikinetic and there was characteristic stoop posture with broad base gait when walking. Detailed neurological examination revealed Glasgow Coma Scale of 15/15 and there were no cranial nerve abnormalities. Sensory system examination also was normal. He was severely bradikinesic when performing finger nose testing and heel knee shin test.

Further laboratory investigations were performed as follows. Urgent MRI (magnetic resonance imaging) brain was done and was completely normal. EEG (electro encephalogram) showed evidence of encephalitis. Cerebrospinal fluid (CSF) analysis showed white blood cells (WBC) of 45 cells/mm³ (lymphocytes 92%), CSF protein of 58 mg/dL, no red cells and normal sugar level. CSF gram stain and bacterial cultures were negative CSF cytology for malignant cells was negative. CSF was tested for dengue IgM antibodies and it became positive CSF for herpes viruses, Japanese encephalitis virus, polio virus, coxsackie viruses and enteroviruses were negative. Rest of basic investigations including HIV (human immune deficiency virus) antibodies, ESR (erythrocyte sedimentation rate), C reactive protein level, serum electrolytes and full blood count were revealed no abnormality. According to the clinical picture the diagnosis of post encephalitic parkinsonism following dengue viral infection was made. He was started on SINEMET (carbidopa 10 mg and levodopa 100 mg) half tablet 6 hourly started. After 1 week of treatment he showed marked improvement and then patient was discharged with further follow up plan. After 4 weeks of treatment he showed almost complete recovery of his clinical symptoms.

Discussion and conclusion

The dengue is an arbo viral infection caused by four different serotypes (DEN 1, DEN 2, DEN 3, and DEN 4). The dengue virus belongs to the flavivirus group and it is a single stranded enveloped RNA virus. The infection is transmitted by Aedes mosquitoes.

The dengue infection can have a presentation varying from a mild viral fever to dengue hemorrhagic fever and

| WBC | Neu (%) | Lym (%) | PCV | Hb (g/dL) | Platelets | AST | ALT |
|-----|---------|---------|-----|----------|-----------|-----|-----|
| 0 h | 7.8     | 77      | 14  | 33       | 106       | 186 | 57  |
| 12 h| 6.8     | 67      | 32  | 34       | 108       | 160 | 59  |
| 24 h| 7.1     | 70      | 29  | 34       | 11        | 120 | 57  |
| 36 h| 5.59    | 65      | 34  | 33       | 10.4      | 100 | 62  |
| 48 h| 6.24    | 66      | 33  | 32       | 11        | 173 | 60  |
shock. And also it is a multisystem disease with variety of atypical presentations. Neurological manifestations are one such. They include Meningoencephalitis, Acute disseminated encephalomyelitis, Transverse myelitis and Guillain–Barre Syndrome. Encephalitis, the most common neurological manifestation, thought to occur in up to 6.2% of patients with dengue haemorrhagic fever but is rare in uncomplicated dengue fever [3]. Post encephalitic parkinsonism following dengue infection is one such rare presentation and there is very few case reports have been reported in the literature.

The parkinson's disease is named after the English doctor called James Parkinson who published the first detailed description in an essay on the Shaking Palsy in 1817 [4]. Although parkinson's disease has multiple etiologies, idiopathic parkinson's disease is the commonest. Viral etiology is a known fact. It has been shown that numerous viruses can enter nervous system, i.e. they are neurotropic, and induce a number of encephalopathies. One of the secondary consequences of these encephalopathies can be parkinsonism. One of highlighted and controversial cases of viral parkinsonism is that which followed the 1918 influenza outbreak. Other viruses known to induce parkinsonism are coxsackie, Japanese encephalitis, West Nile, HIV, varicella zoster, Herpes Simplex, Epstein Barr and cytomegalovirus [5]. But acute onset parkinsonism following dengue viral infection less commonly reported.

How viruses actually cause parkinsonism still we do not know. Although efforts have been made to detect viral particles, in the brain, or antibodies in the serum or CSF of patient with parkinson's disease, they have been generally unsuccessful. But recent immunohistochemical work has revealed the presence of complement proteins and the interferon induced MxA in association with Lewy bodies and swollen neuronal process. So we propose a hypothesis that neuroviral influenza A and other potent viruses may be responsible for the formation of Lewy bodies and later cell death in nigral region [6]. So routine anti Parkinson medications like carbidopa and levodopa combination may be effective as illustrated by this case study.

Post encephalitic parkinsonism following dengue infection is a very rare presentation and only very few case reports are available. Although viral etiology may be responsible for the formation of Lewy bodies and later cell death in nigral region [6]. So routine anti Parkinson medications like carbidopa and levodopa combination may be effective as illustrated by this case study.

As mentioned according to latest immunohistochemical studies, there is a possibility that potent viruses may be responsible for the formation of Lewy bodies and later cell death in nigral region [6]. So routine anti Parkinson medications like carbidopa and levodopa combination may be effective as illustrated by this case study.

Post encephalitic parkinsonism following dengue infection is a very rare presentation and only very few case reports are available. Although viral etiology may be responsible for the formation of Lewy bodies and later cell death in nigral region [6]. So routine anti Parkinson medications like carbidopa and levodopa combination may be effective as illustrated by this case study.

**Abbreviations**

SNEMET: carbidopa 10 mg and levodopa 100 mg; AST: aspartate aminotransferase; ALT: alanine aminotransferase; Dengue NS 1 Ag: Dengue non structural protein 1 antigen; PCV: packed cell volume; MRI: magnetic resonance imaging; EEG: electroencephalogram; CSF: cerebro spinal fluid; WBC: white blood cell; HIV: human immunodeficiency virus; ESR: erythrocyte sedimentation rate; ELISA: enzyme-linked immunosorbent assay.

**Authors’ contributions**

BVKMB and UR made equal contribution to the identification of the patient, conception of the work, diagnosis and treatment, data collection and analysis, drafting the manuscript, writing references, revising it critically for important intellectual content, final development of case report. Both authors read and approved the final manuscript.

**Acknowledgements**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

**Availability of data and materials**

The date set used and or analyzed during the current study is available from the corresponding author on reasonable request.

**Consent to publish**

Written informed consent was obtained from the patient for publication of this case report.

**Ethics approval and consent to participate**

Not applicable.

**Funding**

None received.
Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Received: 31 August 2017   Accepted: 21 November 2017
Published online: 29 November 2017

References
1. National Dengue Control Unit. Sri Lanka, information on dengue. http://www.dengue.health.gov.lk. Accessed 3 July 2017.
2. Weekly epidemiological report, epidemiology unit. Sri Lanka. vol. 35, issue no 29 (12th to 18th July 2008). 2008.
3. Varatharaj A. Encephalitis in the clinical spectrum of dengue infection. Neurol India. 2010;58(4):S85–91.
4. Parkinson’s disease. https://en.wikipedia.org/wiki/Parkinsonism. Accessed 3 July 2017.
5. Jang Haeman, Boltz David A, Smeyne RJ. Viral parkinsonism. Biochim Biophys Acta. 2009;1792(7):714–21.
6. Takahashi M, Yamada T. Viral etiology for Parkinson’s disease a possible role of inf A virus infection. Jpn J Infect Dis. 1999;52(3):89–98.
7. Laura L, Hermann, Thaisomboonsuk B, Fernandez S. Evaluation of dengue NS-1 Ag detection assay sensitivity and specificity for the diagnosis of acute dengue viral infection. Southeast Asian J Trop Med Public Health. 2011;42(3):570–8.
8. Lam SK, Devine PL. Evaluation of capture ELISA and rapid immunochromatographic test for the determination of IgM and IgG antibodies produced during dengue infection. Clin Diagn Virol. 1998;10(1):75–81.
9. Thomas M, Badachi S, Sarma GRK, Nadig R. "Dot Sign" in dengue encephalitis. Ann Indian Acad Neurol. 2015;18(1):77–9.
10. Ashraf W, Abdul R, Suresh Kumar EX. MRI findings in dengue encephalitis. Ann Indian Acad Neurol. http://www.annalsofian.org. Accessed 3 July 2017.
11. Brijesh KS, Debashree SRD, George RA, Aggarwal, Sivasankar R. MRI findings in dengue encephalitis. Indian J Radiol Imaging. http://www.ijri.org. Accessed 5 July 2017.
12. Sheikh Z, Gaillard AF, et al. Parkinson disease. http://www.radiopaedia.org. Accessed 8 July 2017.