# Concurrent infection of dengue fever and hepatitis A infection: A case report

Sir,

Both dengue fever and hepatitis A infection are endemic in developing countries and are associated with poor sanitation and low socioeconomic status. Their coexistence can present a diagnostic dilemma to the treating physician.<sup>[1]</sup>

A four-year-old girl was admitted with high-grade fever, vomiting, and abdominal pain for 7 days and jaundice for 3 days. On admission she was conscious and hemodynamically stable. Deep icterus was present. The liver was tender with a span of 10 cm and the spleen palpable 2 cm. Other systems were normal. The differential diagnoses considered were malaria, typhoid fever, dengue fever, and leptospirosis. Investigations revealed: hemoglobin 14.8 gm/dL, total leukocyte count 6200/mm<sup>3</sup>, and platelet count 59,000/mm<sup>3</sup>. Peripheral blood smear was negative for malarial parasite. Serum electrolytes, blood culture, urine culture, renal function tests, serum calcium were normal. Widal test and leptospirosis serology were negative. Polymerase chain

Table 1: Liver function test during the course of hospital stay

|  | Day I | Day 5 | Day 8 | Discharge |
|--|-------|-------|-------|-----------|
| Serum bilirubin (mg/dL)                    | 1.8   | 6.6   | 4.7   | 3.8       |
| Direct bilirubin (mg/dL)                   | 1.2   | 3.2   | 2.9   | 1.7       |
| Aspartate aminotransferase (IU/L)<br>(AST) | 4567  | 7654  | 2340  | 450       |
| Alanine aminotransferase (IU/L)            | 1654  | 2476  | 2340  | 879       |
| Lactate dehydrogenase (IÙ/L) (LDH)         | 510   |       | —     | —         |
| AST/LDH                                    | 8.95  |       | _     | _         |
| Prothrombin time (s)                       | 24    | 19    | 19    | 16        |

| Table 2: Mixed infection reported from different regions in the last 15 years |                     |   |  |
|---|---------------------|---|--|
| Mixed infection Region  |                     | Author                                    |  |
| Leptospira, dengue, and hepatitis E   | Delhi (India)       | Behera et al. (2009) <sup>[2]</sup>       |  |
| Dengue fever and malaria  | Karachi (Pakistan)  | Abbasi et al. (2009) <sup>[3]</sup>       |  |
| Dengue fever and hepatitis A and hepatitis E                                  | Karachi (Pakistan)  | Yakoob et al. (2007) <sup>[1]</sup>       |  |
| Dengue fever and typhoid fever  | Bandung (Indonesia) | Sudjana and Jusuf (1998) <sup>[4]</sup>   |  |
| Typhoid fever and viral hepatitis   | Delhi (India)       | Mishra et al. (2008) <sup>[5]</sup>       |  |
| Malaria and leptospirosis   | Bangkok (Thailand)  | Singhsilarak et al. (2006) <sup>[6]</sup> |  |
| Dengue fever and leptospirosis  | Mumbai (India)      | Zaki and Shanbag (2010) <sup>[7]</sup>    |  |
| Hepatitis A and malaria   | Mumbai (India)      | Zaki (2009) <sup>[8]</sup>                |  |

reaction for dengue virus was positive. Ultrasonography revealed pseudothickening of gall bladder, bilateral pleural effusion, ascites, and hepatomegaly with altered echotexture. Liver function tests during the course in hospital are shown in Table 1. Highly elevated liver enzymes and deranged prothrombin time alerted us to the possibility of coexistent viral hepatitis. Serological test for viral hepatitis was positive for HAV-IgM: 1.4 (N # 0.8) and negative for hepatitis B, C, and E viruses. Intravenous fluids and antipyretics were started. As the general condition of the child was improving and all the cultures were negative, supportive treatment was continued. She finally became afebrile on the 11<sup>th</sup> day and was discharged on the 13th day of admission. She is well on follow-up after 2 months with normal liver enzyme tests.

There are several overlapping clinical features of dengue, leptospirosis, malaria, and viral hepatitis, which can cause substantial misdiagnosis. Existence of simultaneous, multiple infections in an individual has been reported in the literature [Table 2].<sup>[1-8]</sup> Both dengue fever and viral hepatitis can present with fever and jaundice. Liver involvement in dengue can occur due to direct effect of the virus or host immune response on liver cells, circulatory compromise caused by hypotension or localized vascular leakage inside the liver capsule and tissue tropism of particular viral serotypes or genotypes.<sup>[9]</sup> Although hepatic involvement is commonly seen with dengue fever, severe hepatic derangement is rare. Presentation of hepatitis A infection is similar, but with a few differences: Fever usually subsides with the appearance of jaundice and the period between onset of fever and jaundice is 1-7 days.<sup>[5]</sup> Serum aminotransferase levels are markedly elevated in viral hepatitis (8-10 times normal) as compared with those in dengue fever in which they are elevated 2-3 times the normal value and the ratio of AST/LDH (aspartate aminotransferase/ lactate dehydrogenase) is more than 4 in viral hepatitis.<sup>[5]</sup> In dengue fever, aspartate aminotransferase

has been found to increase more quickly and peaking at a higher level and then reverting to normal sooner than alanine aminotransferase.<sup>[9]</sup> This pattern is different from that commonly seen during acute hepatitis caused by hepatitis viruses. Other differentiating features of dengue fever include hemoconcentration, thrombocytopenia, and third space fluid losses.<sup>[7]</sup> The coagulation profile is usually normal in patients with dengue fever.<sup>[9]</sup> Hence an abnormal coagulation profile should alert one to an underlying infection with a hepatotropic virus or disseminated intravascular coagulation associated with sepsis. Usually in dengue fever without complications the fever spikes comes down by day 4-5 of illness. However, fever may be prolonged in patients having coexisting other infections as seen in our case. Highly elevated liver enzymes, deranged prothrombin time, and prolonged fever in the patient alerted us to the possibility of coexistent viral hepatitis.

This case illustrates the importance of physician awareness of mixed infections in endemic areas that can pose diagnostic dilemmas, complications, and prolonged course.

## Syed Ahmed Zaki, Vijay Lad

Department of Pediatrics, Lokmanya Tilak Municipal General Hospital and Medical College Sion, Mumbai- 400 022, Maharashtra, India

#### Correspondence:

Dr. Syed Ahmed Zaki, Room no.509, new RMO guarters, Sion, Mumbai- 400 022, Maharashtra, India. E-mail: drzakisyed@gmail.com

### References

- Yakoob J, Jafri W, Siddiqui S, Riaz M. Dengue fever with hepatitis E and hepatitis A infection. J Pak Med Assoc 2009;59:176-7.
- 2.Behera B, Chaudhry R, Pandey A, Mohan A, Dar L, Premlatha MM, et al. Co-infections due to leptospira, dengue and hepatitis E: a diagnostic challenge. J Infect Dev Ctries 2009;4:48-50.
- 3 Abbasi A, Butt N, Sheikh QH, Bhutto AR, Munir SM, Ahmed SM. Clinical features, diagnostic techniques and management of dual dengue and malaria infection. J Coll Physicians Surg Pak 2009;19:25-9.
- 4. Sudjana P, Jusuf H. Concurrent dengue hemorrhagic fever and typhoid fever infection in adult: case report. Southeast Asian J Trop Med Public Health 1998;29:370-2.
- 5. Mishra D, Chaturvedi D, Mantan M. Typhoid fever and viral hepatitis.

Indian J Pediatr 2008;75:509-10.

- Singhsilarak T, Phongtananant S, Jenjittikul M, Watt G, Tangpakdee N, Popak N, et al. Possible acute coinfections in Thai malaria patients. Southeast Asian J Trop Med Public Health 2006;37:1-4.
- Zaki SA, Shanbag P. Clinical manifestations of dengue and leptospirosis in children in Mumbai: an observational study. Infection 2010;38:285-91.
- 8. Zaki SA, Asif S, Dadge D, Shanbag P. Co-existence of viral hepatitis with malaria. J Postgrad Med 2009;55:233.
- Trung DT, Thao le TT, Hien TT, Hung NT, Vinh NN, Hien PT, et al. Liver involvement associated with dengue infection in adults in Vietnam. Am J Trop Med Hyg 2010;83:774-80.

| Access this article online |  |  |  |
|----------------------------|--|--|--|
| Quick Response Code:       |  |  |  |
|                            | Website:<br>www.ijccm.org              |  |  |
|                            | <b>DOI:</b><br>10.4103/0972-5229.92073 |  |  |