INTRODUCTION

Dengue fever is the most rapidly spreading mosquito-borne viral disease worldwide with an estimated 30-fold increase in incidence over last five decades with an unpredictable clinical course and outcome. An estimated 500,000 people with severe dengue infection require hospitalization annually and 90% of them are children <5 years of age. Without proper treatment, the case fatality rate in severe dengue is more than 20% and with timely intervention, it can be reduced to <1%. The recent epidemics has seen changing pattern of presentation of dengue fever in children especially in the presence of coinfections such as enteric fever and malaria thereby making the clinical decision...
more difficult and often mislead the physician’s initial impression.

Puducherry has not experienced any major epidemics of dengue fever in the past, but since 2012 has seen unprecedented rise in dengue fever cases due to rapid urbanization, population migration and increased number of construction sites, open drains and lack of proper public health measures. This study was undertaken to evaluate clinical profile and outcome of dengue fever in children at a tertiary care hospital in Puducherry.

**MATERIALS AND METHODS**

This study was conducted on confirmed cases of dengue fever admitted in the department of pediatrics at a tertiary care hospital in Puducherry from August 2012 to January 2015. All the data from hospital case records were entered in a structured clinical proforma, and the data were retrospectively analyzed. The case definition, diagnosis, and management for dengue fever were as per the revised World Health Organization (WHO) guidelines 2011. Children were classified as dengue fever without warning signs (D), dengue fever with warning signs (DW) and severe dengue infection. Children with severe plasma leakage, severe organ involvement, and severe thrombocytopenia were categorized as severe dengue infection. The diagnosis was confirmed by NS1 antigen-based ELISA test (J. Mitra Kit, India) or dengue serology for IgM and IgG antibodies (Kit from National vector born disease control program Puducherry and National Institute of virology Pune, India) during the acute phase and convalescent phase of illness. Blood samples were collected from children with a provisional diagnosis of dengue fever and were screened by both NS1 Ag assay and MAC-ELISA. All children suspected of dengue fever in the 1st week of illness who were NS1 Ag negative were retested during the 2nd week of illness. All other relevant and other additional investigations were done as per the clinical course of illness. The study protocol was approved by the Institute Ethics committee of Indira Gandhi Medical College and Research Institute at Puducherry, India.

**Statistical analysis**

The data were analyzed using SPSS for window version 16.0 (SPSS, IL, 233 South Wacker drive, 11th Floor, Chicago, 60606-3412, USA) using descriptive statistics. Categorical variables were expressed as frequencies and percentages and then analyzed using the Chi-square test or Fishers exact test, where appropriate. Continuous data expressed as mean ± SD, or median (range), were analyzed using Students t-test, analysis of variance-one-way, or Mann-Whitney U-test. Significance was taken at $P < 0.05$.

**RESULTS**

Out of 398 children admitted with dengue fever the diagnosis was confirmed in 261 cases (65.5%). Non-severe dengue was seen in 159 cases (60.9%) and severe dengue infection was seen in 102 cases (39.1%). The mean age of presentation was 6.9 years and 6-12 years was the most commonly affected age group (62.0%). Infants were the least affected sub-group with 7 cases (2.7%). The youngest was a neonate, and the oldest was 12 years of age. The male: female ratio was 1.2:1. The children that were admitted with dengue fever were from Puducherry (76.3%) and Tamil Nadu (23.7%) [Figure 1]. The mean duration of hospital stay was 6.5 (2.7) days.

The common clinical presentations included fever (94.6%), conjunctival congestion (89.6%), myalgia (81.9%), coryza (79.7%), headache (75.1%), palmar erythema (62.8%), retro-orbital pain (51.3%), joint pain (28.7%), and rash (17.2%). The common atypical manifestations of dengue fever at admission were lymphadenopathy (52.3%), splenomegaly (20.7%), epigastric tenderness (16.4%) biphasic fever (15.7%), right hypochondriac pain (8.4%), seizures (6.5%), and febrile diarrhea (6.5%), [Table 1]. The mean duration of fever was 4.8 (1.8) days at admission.

The common early warning signs at the time of admission were persistent vomiting (75.1%), liver enlargement (59.8%), cold and clammy extremities (45.2%), peripheral circulatory failure (40.6%), pain abdomen (31.0%), hypotension (29.5%), restlessness (26.4%), giddiness (23.0%), oliguria (18.4%), ascites (6.9%), lethargy (6.3%),
pleural effusion (5.0%), and impaired consciousness with Glasgow coma scale (GCS) <8 (2.7%) [Table 1].

Hemorrhagic manifestations were present in 52 children (19.9%), which mainly included skin bleeding (39.2%), gum bleeding (34.6%), epistaxis (23.1%), melena (26.9%), hematemesis (9.6%), intracranial bleed (3.8%), and pulmonary bleed (7.7%). Tourniquet test was positive in 33 cases (12.6%). Melena was the most common form of gastrointestinal (GI) bleed.

Shock was present in 102 cases (39.1%) in children with severe dengue infection and among them significant bleeding was present in only 22 cases (21.6%) and decompensated shock in 16 cases (15.7%). There was difficulty in classifying cases as dengue hemorrhagic fever (DHF) as all children did not fulfill the four criteria of fever, thrombocytopenia, bleeding, and plasma leakage. The predominant mode of presentation of severe dengue infection was with features of peripheral circulatory failure without spontaneous bleeding. Six children (2.3%) with decompensated shock had fluid refractory shock and required inotropic support.

The common complications seen with severe dengue infection were liver dysfunction, acute respiratory distress syndrome, pneumonia, impaired consciousness (GCS <8), myocarditis, myositis, hemophagocytic syndrome, acute kidney injury (AKI), and disseminated intravascular coagulopathy (DIC). Co‑infections were seen in 19 cases (7.5%) and among them, enteric fever in 10 cases, malaria in 4 cases, urinary tract infection in 4 cases and one case had acute bacterial meningitis [Table 2].

The hematological parameters showed anemia (29.5%), leukopenia (19.1%) and thrombocytopenia in 215 (82.4%) cases. Hemoconcentration was in 46.4% of cases, and the mean hematocrit was 38.9 (4.4). Seven children (2.7%) had platelet count <10,000/mm³, five children (1.9%) were between 10,000 and 20,000/mm³, 31 children (11.9%) were between 20,000 and 50,000/mm³, 58 (22.2%) children had platelet count between 50,000 and 100,000/mm³, 114 (43.7%) children had platelet count between 1 and 1.5 lakhs/mm³ and 46 children (17.6%) were >1.5 lakhs/mm³). Severe thrombocytopenia (platelet count <50,000/mm³) was seen in 43 (16.5%) cases and among them 36 (83.7%) cases had severe dengue infection. All children who had platelet count <20,000/mm³ had severe dengue infection. NS1 Ag was negative with IgM antibody positive in 44 cases (16.9%). Dengue IgG antibody was positive in 17 cases (6.5%) and among them, severe dengue infection was seen in 16 cases (94.1%). Disordered coagulation (prolongation of the prothrombin and/or activated partial thromboplastin time) was seen in 13 children (5.0%). Altered liver enzymes were seen in 29 cases (11.1%). Ultrasonography of abdomen showed gallbladder wall edema in severe dengue infection in 25 cases (9.6%) [Table 3].

Fifty-two children (19.9%) had spontaneous bleeding and among them, shock was present in 22 children (42.3%). Bleeding manifestations were present in 20 (39.2%), 8 (15.7%), 13 (25.5%) and 11 (21.6%) cases with platelet count between <50,000/mm³, 50,000-100,000/mm³, 1-150,000 and >150,000/mm³, respectively. About 78.8% of children with bleeding had thrombocytopenia, but bleeding manifestations did not always correlate with platelet counts in non-severe dengue infection in comparison to severe dengue infection.

**Table 1: Clinical profile of children with dengue fever**

| Clinical variables | n (%) |
|--------------------|-------|
| Fever              | 247 (94.6) |
| Conjunctival congestion | 234 (89.6) |
| Myalgia            | 214 (81.9) |
| Coryza             | 208 (79.7) |
| Headache           | 196 (75.1) |
| Persistent vomiting* | 196 (75.1) |
| Palmar erythema*   | 164 (62.8) |
| Hepatomegaly       | 156 (59.8) |
| Lymphadenopathy**  | 137 (52.3) |
| Retro-orbital pain | 134 (51.3) |
| Cold and clammy extremities* | 118 (45.2) |
| Poor peripheral pulses* | 106 (40.6) |
| Facial flush       | 92 (35.2) |
| Pharyngeal congestion | 85 (32.6) |
| Pain abdomen*      | 81 (31.0) |
| Hypotension*       | 77 (29.5) |
| Joint pain         | 75 (28.7) |
| Restlessness*      | 69 (26.4) |
| Giddiness*         | 60 (23.0) |
| Splenomegaly**     | 54 (20.7) |
| Bleeding*          | 52 (19.9) |
| Oliguria*          | 48 (18.4) |
| Rash               | 45 (17.2) |
| Epigastric tenderness** | 43 (16.4) |
| Biphasic fever**   | 41 (15.7) |
| Tourniquet test    | 33 (12.6) |
| Right hypochondriac pain** | 22 (8.4) |
| Pleural effusion*  | 13 (5.0) |
| Ascites*           | 18 (6.9) |
| Febrile diarrhea** | 17 (6.5) |
| Seizures**         | 17 (6.5) |
| Pedal edema        | 09 (3.6) |
| Lethargy**         | 09 (3.6) |
| Impaired consciousness** | 07 (2.7) |

*Warning signs, **Atypical manifestations, Data as number (%)

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Table 2: Complications in severe dengue infection

| Complications                              | n (%) |
|--------------------------------------------|-------|
| Encephalopathy                             | 7 (2.7) |
| Hepatitis                                  | 29 (11.1) |
| Intracranial hemorrhage                    | 2 (0.8) |
| Fulminant hepatic failure                  | 2 (0.8) |
| Acute kidney injury                         | 6 (2.3) |
| Myocarditis                                | 5 (1.9) |
| Pericardial effusion                        | 3 (1.1) |
| Pneumonia                                  | 31 (11.9) |
| Pulmonary hemorrhage                        | 4 (1.5) |
| Acute respiratory distress syndrome         | 4 (1.5) |
| Hemophagocytic syndrome                    | 2 (0.8) |
| Refractory shock                            | 6 (2.3) |
| Myositis                                    | 3 (1.1) |
| Disseminated intravascular coagulopathy    | 4 (1.5) |
| Co-infections                              | 19 (7.2) |

Data as number (%)

Table 3: Laboratory parameters in children with dengue fever

| Laboratory diagnosis                          | n (%) |
|-----------------------------------------------|-------|
| Thrombocytopenia (PLC <150,000/mm³)           | 215 (82.4) |
| Leukopenia (TLC <4000/mm³)                    | 59 (19.1) |
| Anemia (hemoglobin <10 g/dl)                  | 77 (29.5) |
| Platelet count <50,000/mm³                    | 43 (16.5) |
| Platelet count <20,000/mm³*                  | 12 (4.6) |
| HCT >20% with PLC <50,000/mm³*               | 30 (11.5) |
| Hemconcentration (HCT >40)                    | 121 (46.4) |
| NS1Ag positive                               | 237 (83.1) |
| NS1Ag negative and dengue IgM antibody positive | 44 (16.9) |
| Dengue IgG antibody                           | 17 (6.5) |
| Deranged LFT (SGOT and SGPT >150 IU/L)        | 29 (11.1) |
| Deranged RFT (serum creatinine >3 mg/dl)      | 6 (2.3) |
| Hyponatremia (sodium <135 Meq/L)             | 14 (5.4) |
| Gall bladder wall edema on ultrasound         | 25 (9.6) |
| Prothrombin time (PT/INR >1.0)               | 9 (3.4) |
| APTT (>1.5 times normal)                     | 4 (1.5) |

*Warning signs, Data as number (%). HCT: Hematocrit, PLC: Platelet count, TLC: Total Leukocyte count, LFT: Liver function test, RFT: Renal function test, SGOT: Serum glutamic oxaloacetic transaminase, SGPT: Serum glutamic pyruvic transaminase, INR: International normalized ratio, APTT: Activated partial thromboplastin time, PT: Prothrombin time

Platelet transfusion was given in 17 children (6.5%) with severe dengue infection and out of them 12 children (70.6%) had a platelet count <20,000/mm³; whereas 5 children (29.4%) had platelet count in the range of 20,000-50,000/mm³. Only those children with significant spontaneous bleeding, shock and with severe thrombocytopenia necessitated platelet transfusion. Apart from platelet transfusion blood transfusion was given in 10 (3.9%) cases, fresh frozen plasma in 6 (2.3%) cases, colloids in 3 (1.2%) cases, intravenous fluids in 96 (37.7%) cases, and inotropes in 5 (1.9%) cases. There were six deaths (2.3%) and out of them four presented with impaired consciousness (66.6%) at admission with GCS <8. The common causes for poor outcome (deaths) were multiorgan failure, encephalopathy, DIC, and refractory shock.

DISCUSSION

Dengue fever is a major public health problem with high morbidity and mortality, and the recent epidemic has shown variable clinical presentations with unpredictable clinical evolution and outcome.[10] In our study, children >6 years were the most commonly affected age group and were more at risk to develop severe dengue infection and similar to the previous studies by Faridi et al. and Wichmann et al.[2, 3] However Aggarwal et al., and Gurdeep et al., in their studies showed <6 years of age to be the most common affected age group.[4, 5]

In our study, shock was the most common form of presentation in severe dengue infection and was present in 39.1% of cases in comparison to the previous studies by Ratageri et al., Aggarwal et al., where it was present in 22% and 33%, respectively.[4, 6] The most common mode of presentation of severe dengue infection was a peripheral circulatory failure without bleeding, and there was difficulty in classifying them as DHF as per 1997 WHO classification as most cases failed to fulfill all the four criteria of fever, hemorrhagic phenomenon, thrombocytopenia, and hemoconcentration. They were sub-classified as dengue fever with bleeding without shock and dengue fever associated with peripheral circulatory failure without bleeding based on the revised 2011 WHO guidelines for dengue fever.[1, 7]

Bleeding manifestations were seen in 19.9% of cases and much lower in comparison to the previous studies.[6-10] The most common hemorrhagic manifestations in our study were petechiae and GI bleeding and similar to the previous studies by Ratageri et al., and Rachel et al.[6] Melena constituted the most common form of internal bleeding in our study and also in the study by Shah et al.[8] Hematemesis was reported as the most common manifestations in the study by Narayanan et al., whereas epistaxis was most common in the study by Faridi et al.[10] The tourniquet test in our study was positive in 12.6% of cases and was much lower compared to the previous studies.[10, 14] The tourniquet test did not correlate well with bleeding manifestations or with thrombocytopenia, similar to the finding reported by Wali et al., and Narayanan et al.[10, 15] Bleeding manifestations were highly variable and did not always correlate with the platelet counts as it occurred in 21.2% of cases with normal platelet counts.
During the epidemics at Puducherry, primary infection was more common than secondary infection, but 94.1% of cases with secondary infection had severe dengue infection. During secondary infection T-cells become activated due to interactions with infected monocytes which induce plasma leakage by the release of a cascade of cytokines such as interferon-gamma, IL-2, and tumor necrosis factor-alpha (TNF-α) thereby predisposing to DHF and dengue shock syndrome. Wichmann et al., in their study showed that secondary infection was significantly associated with the development of severe dengue infection in children.

The most pathognomic feature of dengue is an increase in vascular permeability leading to loss of plasma from blood vessels, which causes hemoconcentration, low blood pressure and shock. This may also be accompanied by a combination of hemostatic abnormalities such as thrombocytopenia, vascular changes, and coagulopathy. Hemoconcentration was difficult to interpret in our study due to lack of the availability of preillness hematocrit and with a high prevalence of anemia in the community. However, the change in hematocrit of more than 20% and standard hematocrit cut offs from the previous studies by Gomber et al., and Balasubramanian et al., was used as indicators of hemoconcentration for diagnosing and monitoring severe dengue infection. There was a poor correlation between thrombocytopenia, bleeding, and plasma leakage in the study as 21.2% of children with bleeding had normal platelet counts and only 24.2% of children with shock had bleeding manifestations. Coagulation profile was deranged in 13 cases (5.1%) signifying the fact that factors other than thrombocytopenia such as platelet dysfunction, consumption coagulopathy, and endothelial dysfunction are responsible for bleeding in dengue fever.

Platelet transfusion was indicated only in children with severe dengue infection with shock with severe thrombocytopenia (platelet count <50,000/mm³) with significant bleeding and prophyactic platelet transfusion was not given similar to the previous studies. Skin bleeds, single episode of epistaxis or mucosal bleed were not given empirical platelet transfusion based on platelet counts.

Children who had features of shock with rising hematocrit, not responding to crystalloids received plasma or colloids whereas with falling hematocrit received a whole blood transfusion.

The classical presentation of dengue fever was not always present in our study. Coryza was one of the common manifestations in our study which is otherwise unusual in dengue fever with very few published reports. Splenomegalgy was in 20.7% of cases in our study and is an unusual manifestation of dengue fever. Faridi et al., in their study similarly showed a high percentage (32.4%) of splenomegalgy in children with dengue. Hemophagocytic syndrome is an unusual manifestation of dengue fever which was present in two cases with very cases reported in the past. Myositis an unusual manifestation was found in two cases in our study. The probable mechanism for myositis is the release of myotoxic cytokines, particularly TNF-α thereby injuring the affected muscle.

Encephalopathy was the most common neurological manifestations in severe dengue infection in our study and 66% of cases with impaired consciousness at admission had a poor outcome. The neurological manifestations in dengue include seizures, encephalopathy, acute disseminated encephalomyelitis, and rarely Guillain-Barre syndrome. The neurological manifestations are secondary to cerebral hypoperfusion, cerebral edema, direct neurotrophic effect, secondary to hepatic dysfunction and metabolic derangements such as hypoglycemia and hyponatremia as reported in the previous studies.

Hepatic dysfunction was seen in 29 children (11.1%) and two children had a fulminating hepatic failure. The etiology of hepatic dysfunction in dengue fever is usually due to direct cytopathic injury, unregulated host immune response, active viral replication, and hypoxia and tissue ischemia due to prolonged shock, hemorrhage, and metabolic acidosis. The other unusual GI complications in the study were pancreatitis, cholecystitis, appendicitis, febrile diarrhea, and parotitis with very few cases reported in the past.

Co-infections were seen in 7.2% of cases, and it is important that they be promptly recognized as they can modify the clinical presentation of dengue fever and can result in missed or delayed diagnosis and treatment. Coexistence of malaria and dengue have been reported to be in the range of 20% to as high as 80%. The complications associated with poor outcome were acute respiratory distress syndrome, AKI, fluid refractory shock, myocarditis and DIC and similar to the previous studies. The mortality in our study was 2.3% was much lower compared to the previous studies. The common factors could be increased public awareness, better health seeking behavior of parents, early recognition and timely intervention in the study, and change in the epidemic pattern of presentation than in the previous years.
The present study highlights the recent increase in the incidence of dengue fever cases at Puducherry because of lack of proper public health measures. It also highlights the change in the epidemic pattern of presentation with more number of atypical manifestations and lack of classical mode of presentation. Clinical vigilance, awareness and timely intervention are vital to reducing the morbidity and mortality in dengue fever.

There are several limitations to this study. The study is a retrospective analysis of dengue fever cases from a single center and included only those cases that were admitted to the hospital. The diagnosis was confirmed by either dengue NS1 antigen test or dengue serology. Viral isolation and serotype identification and serotype identification was not done in the present study. A large multicentric prospective study including a larger sample size would be ideal.

**CONCLUSION**

The clinical awareness of the changing pattern of presentation is lacking among health care personnel, especially at primary health centers from where these cases are often referred. Health education regarding the unusual manifestations, changing epidemic trends, early recognition and mortality due to severe dengue infection.

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**Conflicts of interest**

There are no conflicts of interest.

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