Study of Serum Lactate Dehydrogenase Levels in Critically Ill Dengue Patients Admitted in PICU

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ABSTRACT

Background: Certain biochemical markers have been found useful in monitoring the progress and predicting the severity of dengue. Serum lactate dehydrogenase (LDH) is one such potential biomarker.

Aim and objective: To study the serum LDH levels in critically ill patients of dengue with warning signs and severe dengue and its relationship with severity of illness.

Materials and methods: Retrospective analytical study.

Results: Sixty patients with severe dengue and dengue with warning signs were included. Mean duration of fever at admission was 4 ± 1.6 days and the mean age was 9.6 ± 5.06 years. Male:female ratio was 1:1. Half the children were above 10 years. The median LDH value was 1,133.5 IU (interquartile range (IQR) 640–1,732). The mean LDH value was significantly higher in patients with severe dengue (2,986.65 ± 3,638.54) as compared to dengue with warning signs (1,209.87 ± 1,370.20) (p=0.047). The majority (70%) of patients with severe dengue had LDH >1,000 IU and complications like severe bleeding, pleural effusion, ARDS, and shock were higher in this group. Mean hospital stay in patients with LDH >1,000 was 14.685 ± 5.993 days and in those with LDH <1,000 was 8.732 ± 3.312 days (p=0.000). Mean platelet count was significantly lower in severe dengue (56,405.00 ± 49,918.74) as compared to dengue with warning signs (922,257.50 ± 71,235.44) (p value 0.028) and there was a weak negative correlation between LDH and platelet count which was non-significant (r = Karl Pearson coefficient −0.055; p value 0.676). The case fatality rate was 9%. The mean LDH (4,783 ± 5,131) in non-survivors was much higher than survivors (1,531.1 ± 1,986) though this was not statistically significant.

Conclusion: Serum LDH level at admission was significantly raised in severe dengue as compared to dengue with warning signs. Similarly, mean LDH values were higher in survivors as compared to non-survivors. There was a weak negative correlation between LDH and platelet count.

Clinical significance: Lactate dehydrogenase can be used early in the disease to identify those who may progress to severe dengue and predict mortality. This will help optimize resource allocation and more effective care in a disease-endemic country like India.

Keywords: Dengue with warning signs, Lactate dehydrogenase, Severe dengue.

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INTRODUCTION

Dengue fever is endemic in tropical countries. The clinical spectrum of dengue infection varies from no symptoms to severe dengue with shock. Nearly 100 million cases of dengue fever and between 250,000 and 500,000 cases of severe dengue are annually reported to the World Health Organization (WHO). Severe dengue is characterized by thrombocytopenia, spontaneous hemorrhages, and plasma leakage that can lead to shock.¹

The name dengue was derived from the Swahili word for “bone-breaking fever” or the Spanish word for “the walk of a Dandie”.² It is a self-limiting arboviral infection characterized by fever, rash, joint pains, nausea, vomiting, headache, and retro-orbital pain. As per the revised WHO criteria in 2009, the cases are classified as dengue without warning signs, dengue with warning signs, and severe dengue.

The disease severity may vary from mild fever which does not require hospitalization to severe disease with features of dengue hemorrhagic fever (DHF)/dengue shock syndrome (DSS) requiring intensive care management.² The disease goes through three clinical stages like febrile phase, critical phase, and recovery phase. The febrile phase generally lasts for 2–10 days after which the patient may either enter the recovery phase or progress to a critical phase marked by defervescence and appearance of plasma leakage. Most complications of dengue occur in the above phase.

Despite the severity, the acute phase of dengue starts with a fever that is indistinguishable from other acute febrile infections. Certain biochemical alterations in dengue may help in early diagnosis and predicting the severity. Studies have reported that patients with DHF have elevated levels of transaminases [aspartate aminotransferase (AST) and alanine aminotransferase (ALT)], amylase, lactate dehydrogenase (LDH), and creatine kinase (CK).³ The serum LDH levels have been reported to be increased in DF and still higher in DHF and DSS cases.³ It has been considered to be a diagnostic marker of dengue infection as well as an independent predictor of DHF.⁶ Sirikutt and Kalayanarooj observed mean LDH
level to be >500 IU in dengue patients and <500 IU in non-dengue patients. Increasing levels of LDH were also seen toward the end of a febrile phase in DHF and DSS cases. Lactate dehydrogenase is an intracellular enzyme abundantly found in body tissues, e.g., muscles, liver, placenta, RBCs, and reticuloendothelial system. Its serum levels increase after cell injury. It has been evaluated as a prognostic marker of various inflammatory states like sepsis, infections, myocardial infarction, malignancies, and cardiopulmonary compromise. It is thought to be a marker of vascular permeability in immune-mediated lung injury. An early increase in LDH (three times the normal value) was found to be an independent predictor of DHF.

Most of the studies on LDH in dengue infection have been in adults. This study was aimed to determine the admission serum LDH levels in patients of dengue with warning signs and severe dengue admitted to PICU and its relationship with severity.

Materials and Methods
This retrospective study was conducted in a tertiary care teaching hospital in Western Maharashtra from July 2019 to December 2019. The study was approved by the Institutional Ethical Committee.

Case records of all patients with a diagnosis of dengue fever based on a positive NS1 antigen and/or IgM ELISA test were screened. The cases with a discharge diagnosis of severe dengue or dengue with warning signs where serum LDH was done at admission were included for analysis. The details related to history, examination, and laboratory findings were noted. The complications, severity, and mortality were documented.

The WHO criteria for dengue with warning signs and severe dengue were used in the study. Accordingly, dengue with warning signs included the presence of the following:

- Abdominal pain or tenderness.
- Persistent vomiting.
- Clinical fluid accumulation.
- Mucosal bleed.
- Lethargy, restlessness.
- Liver enlargement >2 cm.
- Hematocrit increase with a rapid decrease in platelet count.

Severe dengue was defined as the presence of any of the following:

- Severe plasma leakage leading to (a) shock, (b) fluid accumulation with respiratory distress.
- Severe bleeding.
- Severe organ involvement (a) AST or ALT ≥1,000, impaired consciousness, heart, and other organ involvement.

Statistical Analysis
Data were analyzed using descriptive statistics [mean and standard deviation, median and interquartile range (IQR) or frequency and percentage as deemed appropriate]. Categorical variables were analyzed using the Chi-square test. Continuous variables were analyzed using Student’s t-test and the r value was calculated by Pearson correlation coefficient.

Results
During the study period, 227 patients were admitted with dengue fever which comprised 94 cases of dengue with warning signs and severe dengue. Sixty cases admitted to PICU on whom serum LDH was done at admission were included in the study.

The mean duration of fever at admission was 4 ± 1.6 days and the mean age was 9.6 ± 5.06 years. Male:female ratio was 1:1. Half the children were above 10 years.

The median LDH value was 1,133.5 IU (IQR 640–1,732). The LDH values in the study cases ranged from 190 to 13,668 IU (mean—1,802). The majority (70%) of severe dengue had LDH >1,000 IU as against 52% in dengue with warning signs (p = 0.012). The mean LDH value was significantly higher in patients with severe dengue (2,986.65 ± 3,638.54) as compared to dengue with warning signs (1,209.87 ± 1,370.20) (p = 0.047).

Shock and pleural effusion with respiratory distress were significantly higher in patients with LDH >1,000. Other complications like severe bleeding, transaminitis, and ARDS were also more common in patients with LDH >1,000 IU, though this was not statistically significant. On checking the correlation between LDH values and platelet count, there was a negative but weak correlation between LDH and platelet count which was not significant (r = Karl Pearson coefficient —0.055; p = 0.676).

Mean hospital stay (14.685 ± 5.993 days vs 8.732 ± 3.312 days) was significantly higher in patients with LDH >1,000 as compared to those with LDH <1,000 (p = 0.000).

The case fatality rate was 9%. The mean LDH (4,783 ± 5,131) in non-survivors was much higher than survivors (1,531 ± 1,986) though this was not statistically significant. The causes of death included ARDS, refractory shock, DIC, and multi-organ dysfunction.

Discussion
Biochemical alterations detected after 48–96 hours of fever can predict a more severe form of dengue infection. This suggests that early pathogenic changes occur before complications develop. These potential biochemical markers may be used for monitoring illness and predicting severity. Serum LDH levels have been reported to be increased in dengue fever. However, there are limited studies in children evaluating its relationship to the severity of the disease.

Sixty cases admitted to PICU on whom serum LDH was done at admission were included in the study. Half (51%) of the patients were above 10 years of age and the male:female ratio was 1:1. A similar finding was reported by Shankar and Prarthana.6 Out of 60 patients, 20 (33%) had severe dengue and 40 (66%) had dengue with warning signs (Tables 1 and 2).

Fever (100%) was the most common presenting symptom followed by pleural effusion (41%) and severe bleeding like severe gastrointestinal bleeding, hematuria in 11% of cases. Shankar and Prarthana reported fever in 100% cases and the bleeding tendency in 2% of cases.6

The mean LDH in the study population was 1,802 IU. Our finding was similar to Sirikutt and Kalayanarooj7 where the mean LDH was 1,873 IU. Other studies by Perveen et al.8 and Liao et al.9

Table 1: Age-wise distribution

| Age (years) | Frequency | Percentage |
|------------|-----------|------------|
| <1         | 8         | 13         |
| 1–5        | 7         | 11         |
| 6–10       | 14        | 23         |
| 11–17      | 31        | 51         |
| Total      | 60        | 100        |
reported much lower mean LDH values (618.38 and 448.17) as compared to the present study. The mean LDH levels at admission were significantly higher in patients with a discharge diagnosis of severe dengue (2,986.65 ± 3,638.54) as compared to dengue with warning signs (1,209.87 ± 1,370.20) (p = 0.047). Several authors have reported higher serum LDH in severe dengue as compared to non-severe dengue.1,6,8

A study by Sirikutt and Kalayanarooj7 from Thailand compared LDH levels in adult patients with dengue fever and other febrile illnesses. They did serial LDH estimations on enrolment day (day 0), day of leakage, and discharge day and found that LDH levels were higher in dengue patients and the LDH levels correlated with disease severity They reported a mean LDH level of 709.2 in DF, 1,873 in DHF, 654.5 in DSS, and 434 IU in non-dengue patients. The mean LDH levels were elevated in all groups of patients, but with different levels. They concluded that LDH >500 can be used to differentiate patients with dengue from non-dengue patients in the early febrile phase and if the level is 1,000 or more on day 0 it may be a predictor of severe dengue infection (Sirikutt). We also observed LDH levels >1,000 to be significantly associated with severe dengue (p = 0.047). The rise in LDH is attributed to skeletal muscle damage and/or liver damage.

Mean platelet count was significantly lower in severe dengue (56,405.00 ± 49,918.74) as compared to dengue with warning signs (922,257.50 ± 71,235.44) (p = 0.028). The association between the severity of thrombocytopenia and the clinical presentation has been described in many earlier studies.10–13 However, there are no published reports on the correlation of LDH and platelet count. On checking the correlation between LDH values and platelet count, we observed a negative but weak correlation between LDH and platelet level which was not significant (r = Karl Pearson coefficient −0.055; p = 0.676). Studies on a larger number of patients are needed to evaluate whether LDH levels have a significant negative correlation with platelet count (Tables 3 and 4).

Mean hospital stay (14.685 ± 5.993 vs 8.732 ± 3.312 days) was significantly higher in patients with LDH >1,000 as compared to those with LDH <1,000 (p = 0.000). In a recent study, Mittal et al. evaluated LDH as an estimate of the duration of hospital stay in dengue cases and observed a correlation between LDH and time of discharge from the hospital. The authors calculated an estimate of the duration of hospital stay based on platelet counts and LDH where LDH levels on the day of least platelet count were used.

The discharge criteria in their study were platelet count showing a significant increase in three consecutive samples.3 Complications like shock and pleural effusion with respiratory distress were significantly higher in patients with LDH >1,000. Mortality in the study was 9%. Kamath et al.14 reported case fatality of 8.3% in severe dengue. The relatively high mortality in the present study was attributed to the late referral of cases. The mean LDH levels were higher in non-survivors although not statistically significant.

**LIMITATIONS**
The study is limited by a small sample size and its retrospective nature. We also did not have dynamic values of LDH in the different phases of illness which would have added valuable information.

**CONCLUSION**
Serum LDH at admission was significantly raised in severe dengue as compared to dengue with warning signs. It may be used early in the disease to identify those who may progress to severe dengue and help optimize resource allocation in a disease-endemic country like India. Lactate dehydrogenase had a weak negative but non-significant correlation with platelet count.

Serum LDH values should be prospectively evaluated in larger studies to assess their usefulness as an early predictor of severity and to determine the cut-off levels so that they can be included in the management guidelines.

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**Table 2:** Distribution according to clinical manifestations

|          | n = 60 | Percentage |
|----------|--------|------------|
| Fever    | 100    | 100        |
| Severe bleeding | 7     | 11         |
| Impaired consciousness | 4    | 6          |
| Pleural effusion   | 25    | 41         |
| Shock      | 13     | 21         |
| ARDS       | 6      | 1          |

**Table 3:** Distribution according to complications and LDH level

| complications                  | LDH <1,000 (n = 25) | LDH >1,000 (n = 35) | p value |
|--------------------------------|---------------------|---------------------|---------|
| Severe bleeding                | 1 (4)               | 6 (17)              | 0.118   |
| AST or ALT >1,000              | 1 (4)               | 5 (14)              | 0.19    |
| Impaired consciousness         | 1 (4)               | 3 (8)               | 0.483   |
| Pleural effusion with respiratory distress | 6 (24) | 19 (54) | 0.018   |
| ARDS                           | 1 (4)               | 5 (14)              | 0.19    |
| Shock                          | 4 (16)              | 14 (40)             | 0.045   |
| Death                          | 1 (4)               | 4 (11)              | 0.303   |

The study evaluated whether LDH levels have a significant negative correlation with platelet count.
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