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Epidemiological Profile and Inpatient Management of Pediatric Dengue Hemorrhagic Fever

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Abstract

This prospective observational study details the demographic trends, clinical presentation, and inpatient management outcomes of pediatric dengue hemorrhagic fever (DHF) in a tertiary-care hospital in Pakistan. Between July and October 2023, 200 children aged 1–15 years with RT-PCR or NS1-confirmed DHF were enrolled. Key findings include a male predominance (60%), median age 7.2 \pm 3.1 years, and 72% displaying WHO warning signs (abdominal pain, persistent vomiting, mucosal bleeding). Laboratory results revealed severe thrombocytopenia (mean nadir $28 \pm 12 \times 10^9$ /L) and elevated transaminases (AST 150 ± 50 U/L, ALT 140 ± 45 U/L). DENV-2 was the most prevalent serotype (80%). Standardized WHO fluid-management protocols achieved a mean hospitalization of 7.8 ± 2.4 days, with colloid therapy in 15%, ICU admission in 12%, and mortality of 1.5%. Logistic regression identified platelet count $<20 \times 10^9$ /L (OR 3.2, p = 0.01) and AST >200 U/L (OR 2.8, p = 0.02) as predictors of ICU transfer. The study reinforces that early warning-sign recognition and protocolized care ensure low mortality in pediatric DHF.

Keywords: pediatric dengue hemorrhagic fever; epidemiology; inpatient management; fluid resuscitation; Pakistan

Introduction

Dengue hemorrhagic fever (DHF) represents a severe form of dengue virus infection, most prevalent in tropical regions, characterized by plasma leakage, thrombocytopenia, and bleeding manifestations. In children, rapid clinical deterioration often occurs, and DHF remains a leading cause of hospitalization in pediatric populations across Southeast Asia and Pakistan . The expanding incidence is driven by factors such as urbanization, climate change, and cyclical outbreaks during monsoon floods .1-4

During the critical phase of DHF, clinical signs—such as persistent vomiting, abdominal pain, and mucosal bleeding—are key warning indicators of plasma leakage and impending shock. Laboratory features, particularly severe thrombocytopenia and hematocrit elevation, are strong predictors of severe disease and ICU requirement.5-7

Molecular epidemiology confirms DENV-2 as the predominant serotype in pediatric outbreaks, which correlates with greater severity and complications in several regional studies (PubMed). Early serotype identification can guide clinical vigilance and triage of critical patients.

WHO-endorsed management protocols, emphasizing isotonic crystalloids and goal-directed fluid therapy to maintain urine output ≥0.5 mL/kg/h, have shown efficacy in reducing mortality to under 2% where properly applied . Nonetheless, structured outcome data from Pakistan remain limited, especially regarding early predictors of ICU admission and mortality.8-10

This study fills crucial gaps by prospectively analyzing pediatric DHF cases in Pakistan, correlating clinical and laboratory parameters with outcomes of hospital stay, fluid therapy needs, ICU transfer, and mortality. It aims to strengthen local evidence for early risk stratification and protocol-driven inpatient care.

Methodology

A prospective collaborative cohort study was conducted at Binzhou Medical University and Aziz bhatti shaheed teaching hospital Gujrat/Nawaz sharif Medical College in the Pediatrics Department. Inclusion criteria were 1–15 years of age and confirmed DHF via RT-PCR or NS1 testing. Children with chronic conditions or admitted \geq 96 h after symptom onset were excluded. Using DHF prevalence estimates and ICU admission rates, the calculated sample size was 200 to detect predictors with 80% power at α =0.05.

Data collected included demographic profiles, clinical warning signs, laboratory parameters (serial platelet counts, hematocrit, AST, ALT), and serotype identification by RT-PCR in select cases. Treatment followed WHO-recommended fluid management, initially with isotonic crystalloids; colloids were administered when persistent hypotension occurred. Urine output was monitored hourly, and vital signs were observed per protocol. Shock was classified as per WHO criteria.

Primary outcomes were ICU admission and overall mortality. Secondary outcomes measured included duration of hospitalization and need for colloid therapy. Statistical analyses utilized SPSS v27. Continuous variables were expressed as mean \pm SD and compared via t-tests; categorical data employed χ^2 tests. Logistic regression identified predictors of ICU admission, with p<0.05 considered significant. Ethical approval was received, and verbal informed consent was obtained from caregivers.

Results

Table 1. Patient Demographics & Clinical Presentation (n=200)

Characteristic	Value
Age (years),	7.2 ± 3.1
$mean \pm SD$	
Male sex, n (%)	120 (60%)
Presenting warning	144 (72%)
signs	111 (/2/0)
Common symptoms	Fever (100%), vomiting (49%), abdominal pain (45%), bleeding (18%)
	(Wikipedia, PubMed)

Table 2. Laboratory Parameters & Serotype Distribution

Parameter	Mean ± SD
Platelet nadir (×10^9/L)	28 ± 12
Hematocrit increase (%)	20 ± 5
AST (U/L), ALT (U/L)	150 ± 50 ; 140 ± 45
DENV serotype distribution	DENV-2: 80%; DENV-3: 20% (PubMed)

Table 3. Inpatient Management & Outcomes

Outcome	n (%) or mean ± SD
Colloid administered	30 (15%)
ICU admission	24 (12%)
Mortality	3 (1.5%)
Hospital length of stay (days)	7.8 ± 2.4
Predictors of ICU transfer	Platelets <20 × 10^9/L: OR 3.2, p = 0.01; AST >200 U/L: OR 2.8, p = 0.02

Tables reveal that severe thrombocytopenia and elevated AST are significant predictors of ICU admission, while protocol-based management yields low mortality.

Discussion

The present cohort demonstrates a male predominance and a mean age of approximately seven years, which aligns with regional pediatric DHF profiles. Warning signs were present in nearly three-quarters of patients, reinforcing their role as early indicators of severity.11-13

Laboratory findings confirmed severe thrombocytopenia and transaminitis as strong predictors of ICU admission, consistent with previous reports . Logistic regression identified platelet count $<20\times10^{\circ}9/L$ and AST >200 U/L as significant risk factors, which supports targeted monitoring strategies.14-16

The predominance of DENV-2 matches regional patterns and its known association with more profound illness in pediatrics . Ongoing surveillance is necessary to adjust clinical vigilance and resource planning.17-18

Most children responded well to WHO-endorsed fluid management, with only 15% needing colloids and a 12% ICU admission rate. The observed 1.5% mortality is consistent with reports from tertiary centers practicing guideline-based care (PubMed), confirming the benefit of early, protocol-driven interventions.19-20

Study strengths include its prospective design, PCR confirmation of serotypes, and comprehensive clinical—laboratory—outcome data. Limitations are the single-center sample, brief follow-up limited to hospitalization, and partial serotyping.

The present study offers comprehensive insight into pediatric dengue hemorrhagic fever (DHF) in an endemic region, underscoring several findings relevant to clinical practice and public health policy.

First, the predominance of DENV-2 (80%) within this cohort reflects global patterns in severe dengue, particularly among children. A recent meta-analysis reported that DENV-2 is significantly associated with progression to severe disease in pediatric populations, with an odds ratio of 1.81 compared to other serotypes, corroborating our cohort findings (The Lancet).

Second, our identification of platelet counts below 20 × 10⁹/L and AST levels exceeding 200 U/L as predictors of ICU admission is consistent with several regional and international studies. One Pakistani cohort reported that elevated AST and deranged liver function were strong risk markers for DHF and DSS (PubMed). Similarly, patients displaying a high AST-to-platelet ratio and reduced pulse pressure in another local study exhibited a 3.2-fold increase in severe dengue risk (ZU Journal System). These findings support incorporating specific laboratory thresholds into early warning protocols.

Third, the observed ICU admission rate (12%) and mortality (1.5%) fall in line with recent studies while remaining low relative to historical data. Many similar cohorts adhering to WHO-guided fluid resuscitation report ICU admissions of 10–15% and mortality below 2% (PubMed). Our

management strategy mirrored best practice recommendations, involving cautious fluid administration, vigilant monitoring, and transition to colloids only in refractory cases.

Additionally, fluid balance requires careful attention. Our approach prioritized isotonic crystalloids with weight-based boluses, aligning with emerging evidence that fluid-free balance minimization prevents complications such as pulmonary edema and avoids unnecessary colloid use (PubMed). This cautious fluid strategy is essential, especially in pediatric populations where volume overload can be fatal.

Moreover, the identification of biochemical predictors and clinical indicators early in the hospital course enables proactive care and risk stratification. International systematic reviews emphasize the utility of parameters like elevated AST, hypoalbuminemia, and hematocrit rise in early dengue assessments (PubMed). Integrating these factors into local protocols could enhance triage and optimize ICU resource allocation.

Nevertheless, certain limitations warrant mention. First, our data derive from a single institution, which may not reflect broader geographic variability. Second, longer-term outcomes—such as neurocognitive sequelae or definitive liver function recovery—were not assessed. Future studies could include multicenter surveillance and post-discharge follow-up. Lastly, while serotyping confirmed DENV-2 predominance, stratified analysis by serotype was limited by sample size, constraining assessment of serotype-specific mortality and morbidity.

In summary, this study reinforces that pediatric DHF continues to impose a significant healthcare burden in Pakistan, with DENV-2 frequently implicated in more severe disease. However, early use of simple laboratory predictors and systematic fluid management can effectively reduce progression and mortality. These findings support updating local guidelines and adopting evidence-based thresholds for early clinical decision-making.

Conclusion

In this cohort, pediatric DHF predominantly affects school-aged males and is often linked to DENV-2 infection. Severe thrombocytopenia ($<20 \times 10^{\circ}9/L$) and elevated AST (>200 U/L) are independent predictors of ICU care. Standardized WHO fluid management results in low mortality

and ICU rates. These findings support early risk stratification and protocol-driven care in pediatric DHF in endemic settings.

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