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Prevalence and Patterns of Hyponatremia in Patients with Chronic Liver Disease

Kiranya B¹, Chaudhari Saumya², Saloni Singh³, Dr Priyam Singh⁴, Charishma Parla⁵

¹MBBS, Coimbatore Medical College, Coimbatore

²Intern, Pacific Institute of Medical Sciences (Sai Tirupati University)

³Doctor of Medicine, University of Perpetual Help System Dalta Jonelta Foundation School of Medicine

⁴Lumbini Medical College and Teaching Hospital Palpa, Kathmandu University Nepal

⁵MBBS Graduate, Bukhara State Medical Institute Named

Abstract

Background: Hyponatremia is the most frequent electrolyte disturbance in chronic liver disease (CLD), especially cirrhosis, and is associated with poor prognosis.¹,²

Objectives:

- 1. To determine the prevalence of hyponatremia among CLD patients.
- 2. To describe clinical and biochemical patterns of hyponatremia.
- 3. To correlate severity of hyponatremia with disease stage and complications.

Methods: A cross-sectional study was conducted in the Department of Medicine, from **July 2023** – **June 2024**. A total of 120 CLD patients were enrolled. Serum sodium was categorized as mild (130–134 mEq/L), moderate (125–129 mEq/L), and severe (<125 mEq/L).³ Disease severity was assessed using Child-Pugh and MELD scores.^{10–12} Complications including ascites, hepatic encephalopathy, and spontaneous bacterial peritonitis (SBP) were documented.

Results: Hyponatremia prevalence was 45% (54/120). Distribution: mild 22.5%, moderate 13.3%, severe 9.2%. Hyponatremia was significantly associated with **Child-Pugh C** (**64%**) and higher MELD scores (23.4 vs. 16.7; p < 0.01). Ascites (80% vs. 52%), hepatic encephalopathy (42% vs. 21%), and SBP (18% vs. 7%) were more common in hyponatremic patients.

Conclusion: Hyponatremia is highly prevalent in CLD patients at hospital and correlates with advanced disease and complications. Regular sodium monitoring and early correction are essential.

Keywords: Hyponatremia, CLD, Cirrhosis, Child-Pugh, MELD



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1. Introduction

Chronic liver disease (CLD) is a major cause of morbidity and mortality worldwide. In India, alcohol, viral hepatitis, and non-alcoholic steatohepatitis are leading etiologies. Hyponatremia is the most common electrolyte disorder in cirrhosis and has important prognostic significance.¹,²

Pathogenesis involves **non-osmotic vasopressin release**, systemic vasodilation, and impaired renal perfusion, leading to dilutional hyponatremia.⁷,¹⁰ It predisposes patients to refractory ascites, hepatic encephalopathy, hepatorenal syndrome, and worsens outcomes after liver transplantation.⁶,⁸,⁹,¹³,¹⁵

Reported prevalence varies between **30–50%** depending on study setting.¹,³,⁷ Limited data exist from tertiary centers in North India. This study was therefore conducted at UPUMS, Saifai to determine prevalence and patterns of hyponatremia in CLD patients.

Methods

Study Design

A **cross-sectional observational study** was carried out. This design was chosen as it allows estimation of the prevalence of hyponatremia and description of its patterns at a single point of time among patients with chronic liver disease (CLD).

Study Period

The study was undertaken over a period of 12 months, from July 2023 to June 2024.

Sample Size and Sampling

A total of **120 consecutive patients** diagnosed with chronic liver disease (CLD) and admitted to the Department of Medicine during the study period were included. A consecutive sampling technique was used to avoid selection bias.

Inclusion Criteria

- Adult patients aged ≥18 years.
- Patients with clinically, biochemically, and radiologically confirmed chronic liver disease, irrespective of etiology.

Exclusion Criteria

- Patients with chronic kidney disease (CKD).
- Patients with **congestive heart failure** (CHF).
- Patients with **thyroid or parathyroid disorders** that could alter serum sodium.
- Patients on long-term diuretics for non-hepatic causes.



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Data Collection

Data were collected using a structured proforma. The following parameters were recorded:

- **Demographic details**: age, sex, residence (rural/urban).
- **Etiology of CLD**: alcoholic liver disease, viral hepatitis, non-alcoholic steatohepatitis (NASH), others.
- Laboratory parameters: serum sodium levels, liver function tests, renal function tests.
- Classification of hyponatremia:
 - o Mild: 130–134 mEq/L
 - o Moderate: 125–129 mEq/L
 - o Severe: <125 mEq/L
- Disease severity indices:
 - o **Child-Pugh score** (grading into A, B, and C).
 - o Model for End-Stage Liver Disease (MELD) score.
- **Complications documented**: ascites, hepatic encephalopathy, spontaneous bacterial peritonitis (SBP).

Statistical Analysis

Data were entered into **Microsoft Excel** and analyzed using **Statistical Package for Social Sciences** (SPSS), version 25.0 (IBM, USA).

- Continuous variables were expressed as mean ± standard deviation (SD) and compared using Student's t-test.
- Categorical variables were expressed as **frequency and percentage** and compared using the **Chi-square test** (χ^2).
- A p-value <0.05 was considered statistically significant.

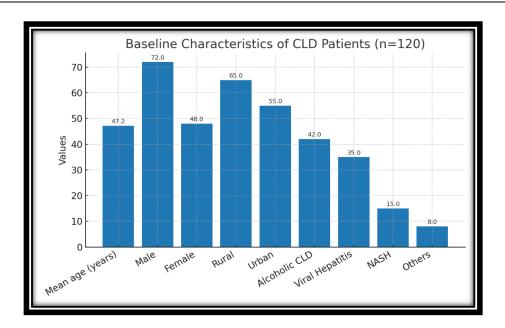
Results

Table 1. Baseline Characteristics (n = 120)

| Characteristic | Value | |
|-------------------------|---|--|
| Mean age (years) ± SD | 47.2 ± 11.8 | |
| Male : Female ratio | 72 (60%) : 48 (40%) | |
| Residence (Rural/Urban) | 65 (54%) / 55 (46%) | |
| Etiology of CLD | Alcoholic – 42%; Viral Hepatitis – 35%; NASH – 15%; Others – 8% | |



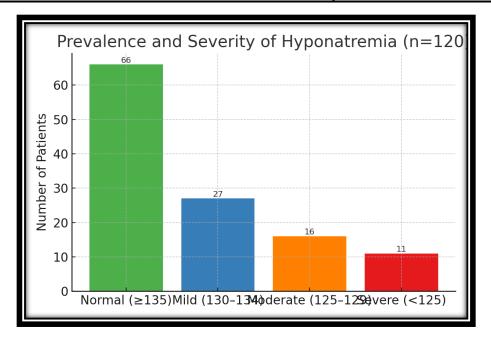
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This table shows the demographic and clinical profile of the 120 CLD patients included. The mean age was 47.2 years, with a male predominance (60%). Slightly more patients belonged to rural areas (54%). Alcoholic liver disease (42%) and viral hepatitis (35%) were the leading etiologies.

Table 2. Prevalence and Severity of Hyponatremia

| Serum Sodium (mEq/L) | Patients (%) |
|----------------------|--------------|
| Normal (≥135) | 66 (55%) |
| Mild (130–134) | 27 (22.5%) |
| Moderate (125–129) | 16 (13.3%) |
| Severe (<125) | 11 (9.2%) |
| Total Hyponatremia | 54 (45%) |



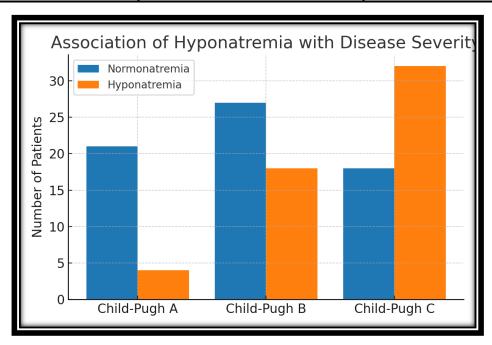


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Out of 120 CLD patients, 54 (45%) had hyponatremia. Most cases were of mild type (22.5%), followed by moderate (13.3%) and severe (9.2%). More than half of the patients (55%) maintained normal sodium levels. This highlights that nearly one in two patients with CLD develops hyponatremia.

Table 3. Association with Disease Severity

| Disease Severity | Normonatremia (%) | Hyponatremia (%) |
|---------------------|-------------------|------------------|
| Child-Pugh A (n=25) | 21 (84%) | 4 (16%) |
| Child-Pugh B (n=45) | 27 (60%) | 18 (40%) |
| Child-Pugh C (n=50) | 18 (36%) | 32 (64%) |
| Mean MELD Score | 16.7 ± 4.5 | 23.4 ± 5.2 |



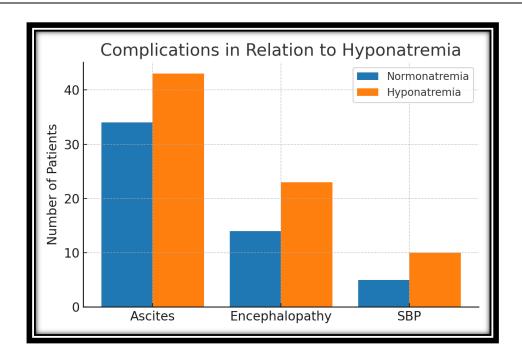
The prevalence of hyponatremia rose markedly with worsening Child-Pugh class. Only 16% of Child-Pugh A patients had hyponatremia, compared to 64% in Child-Pugh C. The mean MELD score was also significantly higher in the hyponatremic group (23.4 vs. 16.7), establishing hyponatremia as a marker of advanced disease.

Table 4. Complications in Relation to Hyponatremia

| Complication | Normonatremia (n=66) | Hyponatremia (n=54) |
|------------------------|----------------------|---------------------|
| Ascites | 34 (52%) | 43 (80%) |
| Hepatic Encephalopathy | 14 (21%) | 23 (42%) |
| SBP | 5 (7%) | 10 (18%) |



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Complications were significantly more frequent among hyponatremic patients. Ascites was present in 80% of the hyponatremia group compared to 52% in normonatremics. Similarly, hepatic encephalopathy (42% vs. 21%) and SBP (18% vs. 7%) showed higher prevalence. This indicates that hyponatremia correlates with increased risk of decompensating events.

Discussion

This study found a **45% prevalence** of hyponatremia in CLD patients at UPUMS, similar to prior Indian and international data.¹,³,⁷ Severe hyponatremia was strongly associated with **Child-Pugh C** and higher **MELD scores**, confirming its role as a poor prognostic marker.⁵,⁶,^{10–12}

Complications such as ascites, encephalopathy, and SBP were significantly more common in hyponatremic patients, consistent with Sindhura et al.¹ and Thuluvath et al.⁷ Hyponatremia results from vasopressin-mediated water retention and impaired solute clearance,⁷,¹⁰ which worsens cerebral edema⁹ and aggravates portal hypertension.⁸,¹³

Clinically, hyponatremia is an independent risk factor for mortality and is incorporated in transplant prognostic models.¹⁴,¹⁵ Early detection and timely management (fluid restriction, vaptans, careful correction) are vital.

Limitations: Single-center, cross-sectional design, limited sample size. Multicenter prospective studies are needed.

Conclusion

Hyponatremia is common (45%) in CLD patients. It correlates with advanced disease (Child-Pugh C, high MELD) and with complications such as ascites and encephalopathy. Routine sodium monitoring and appropriate management are essential for better outcomes.



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