Are potassium levels in children with hemolytic uremic syndrome predictive of outcome?*

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Received 21 June 2012; revised 17 September 2012; accepted 3 October 2012

ABSTRACT

Objective: To ascertain the role of serum potassium levels in predicting clinical outcomes in diarrhea-associated hemolytic uremic syndrome (HUS D+). Methods: We reviewed clinical and laboratory data from HUS D+ patients at our tertiary care institution from 2001 to 2008. Serum potassium concentration at presentation and during the acute phase of acute renal failure were recorded and related to laboratory parameters and clinical outcomes. Results: 15 HUS D+ cases were identified. E. coli 0157:H7 was found in 9/15 cases (70%). Potassium levels were not predictive of clinical outcomes. Normal serum potassium levels were found in the majority of patients. Potassium levels <3.6 mmol/L were evident at presentation in 3/15 patients (23%), and no patient manifested hyperkalemia even when creatinine levels were concurrently increase. Conclusions: This study suggests the presence of vigorous compensatory mechanisms in the homoeostasis of serum potassium levels during HUS D+ disease since neither the increase stool volumes associated with diarrhea nor the presence of renal failure resulted in clinically significant changes in serum potassium levels.

Keywords: Diarrhea; Hemolytic Uremic Syndrome; Hypokalemia; Potassium Metabolism; Acute Renal Failure

1. BACKGROUND

Hemolytic-Uremic Syndrome (HUS) is defined by the triad of microangiopathic hemolytic anemia, thrombocytopenia and acute renal failure. Annual incidence approaches 1 case per 100,000 children with peak prevalence in the summer [1], and most cases occur in children under five years of age [2]. HUS is a heterogeneous group of similar entities characterized by a variety of prothrombotic host abnormalities [3], with variable clinical expression and severity [4,5]. It is often associated with diarrhea (HUS D+) caused by gastrointestinal infections with Shiga toxin-producing E. coli, which likely produce their effect through the systemic spread of bacterial derived toxins [6,7]. The disease has been linked to consumption of hamburger meat contaminated with toxin-producing E. coli 0157:H7 [1,8].

HUS is the most frequent cause of non-pre-renal acute renal failure in pediatrics. HUS D+ accounts for >80% of pediatric cases. The majority of children with HUS D+ develop some degree of renal insufficiency, and approximately two thirds of children with HUS will require dialysis therapy, while about one third will have milder renal involvement and no need for dialysis. Electrolyte abnormalities encountered in HUS D+ include hypotension and both hypo- and hyper-kalemia [9] although no large studies have examined changes in potassium levels in children with HUS D+. We therefore sought to examine potassium levels in patients with HUS D+ at our institution in an attempt to do understand the pathophysiology of potassium metabolism in HUS D+.
2. METHODS
A retrospective chart review was performed of all patients aged <16 years with HUS D+ seen from April 1st 2001-November 2008. Twenty five patients had HUS, and the medical records of the 15 patients who were HUS D+ were reviewed. Potassium levels at presentation and during hospitalization were recorded. Serum potassium concentrations of >5 mmol/l were used to define hyperkalemia and <3.6 mmol/l to define hypokalemia. Serum sodium levels <135 mmol/l being defined as hyponatremia. No medication that could affect potassium homeostasis was used by any patient prior to presentation.

Clinical outcomes were judged by the extent and duration of renal dysfunction, the duration of dialysis, hypertension, and the length of hospital stay, modeled after the classification of Trompeter, et al. [10]. Adapting these criteria, patients were classified into one of three categories:

Group 1: Patients who had a self-limited clinical course with mild thrombocytopenia and mildly decreased renal function, not requiring dialysis. The hospital stay was typically less than seven days with normal renal function and only minor abnormalities in urinary sediment present at discharge.

Group 2: Patients with more severe and prolonged thrombocytopenia with renal failure requiring 1 - 21 days of peritoneal dialysis. No hypertension developed, and patients were discharged within 2 - 3 weeks. Mild proteinuria with otherwise normal renal function was present at discharge.

Group 3: Patients with marked and prolonged thrombocytopenia with renal failure, requiring permanent dialysis.

3. RESULTS
The study group consisted of six (40%) females and nine (60%) males with ages ranging from 2.5 - 15 (median 6.6) years. Utilizing the Trompeter classification [10], 9 (60%) of the children had a very good outcome (Group 1) and the other 6 (40%) recovered fully after a more prolonged clinical course (Group 2). No patient developed end stage renal disease (Group 3)—See Table 1.

As per definition diarrhea was recorded in all 15 patients and bloody stools were present in 13 (93%). The diagnostic triad associated with HUS D+ occurred in 13 patients (93%); in two patients the platelet counts remained normal throughout the illness (see Table 1). Stool cultures showed evidence of a bacterial pathogen in 10 cases (76%), E. coli 0157:H7 accounted for nine of these. Non-0157 E. coli, Salmonella non-typhi and Toro-virus, were each found in the stool of one patient. The median duration of hospital stay was 14 days (range 3 - 40 days) and two (14%) patients were admitted to the pediatric in-

Table 1. Children with HUS D+.

<table>
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<tr>
<th>Patient #</th>
<th>Age in Years</th>
<th>Creatinine in mmol/L</th>
<th>Initial Platelet Count × 10^9</th>
<th>Sex</th>
<th>Potassium on Admission in mmol/L</th>
<th>Dialysis</th>
<th>Received Blood</th>
<th>Trompeter Grade</th>
<th>Agent</th>
<th>Sodium on Admission in mmol/L</th>
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<td>No</td>
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<td>57</td>
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tensive care unit. Arterial hypertension was observed in two patients (14%) and neurological defects in three (21%). Renal replacement therapy was instituted in six (46%) patients: five received peritoneal dialysis and one haemodialysis.

Hypokalemia was seen in three patients (21%). No patient developed hyperkalemia, despite acute renal failure with mean serum creatinine levels of 202 mmol/L. Hypokalemia was seen in seven patients (50%), but was not severe (lowest value 129 mmol/L), nor was there any association with clinical outcomes.

4. DISCUSSION

At presentation, serum potassium levels in our patients were low or normal, and no patients manifested hyperkalemia. Potassium levels, even prior to dialysis, were not predictive of clinical outcomes, which were uniformly favorable with no mortality.

Acute renal failure in HUS generally manifests as hypertension, oliguria, metabolic acidosis and hyperkalemia and management consists of appropriate fluid and electrolyte management, antihypertensive therapy, and the initiation of renal replacement therapy when appropriate [11]. In vitro studies have found that there may be local activation of the renin-angiotensin system during the course of HUS, contributing to the thrombotic microangiopathy which is a significant modulator of severity disease in HUS [12]. Renin-angiotensin system activation then contributes to hypertension. Diarrhea in children can cause electrolyte abnormalities such as hyponatremia, hypokalemia and metabolic acidosis with normal anion gap [13]. The concentration of potassium in normal stool is 80 - 90 mmol/L, but because of the low volume of water in normal stool, only about 10 mmol of potassium is normally lost each day [14]. In diarrheal states, although the potassium concentration in stool decreases, large quantities of potassium can nevertheless be lost as the volume of stool increases [15]. Thus any condition, such as infectious diarrhea, that increases stool volume can result in clinically significant potassium depletion and hypokalemia. Profuse watery diarrhea is usually not present at the time of presentation in patients with HUS D+ and that neither initial levels of potassium nor those found during the course of hospitalization are predictive of clinical outcomes. The number of patients in our study is low, limiting the strength of conclusions that can be drawn regarding potassium levels and disease outcomes in HUS D+. Measurement of blood pressure, blood pH, renin and aldosterone levels, stool and urine volumes and the concentration of potassium in each, would have helped clarify potassium homeostasis in our patients. Metabolic acidosis with a random urine potassium-creatinine ratio less than 1.5 may suggest excessive gastrointestinal losses due to diarrhea, or a shift of potassium into cells.

REFERENCES


[10] Trompeter, R.S., Schwartz, R., Chaltiner, C., Dillon, M.J.,


