Does blood urea nitrogen level predict severity and high-risk endoscopic lesions in patients with nonvariceal upper gastrointestinal bleeding?

Khalid Al-Naamani MD1, Nabil Alzadjali MD2, Alan N Barkun MD MSc (Epidemiology)1,3, Carlo A Fallone MD1

BACKGROUND: Nonvariceal upper gastrointestinal bleeding (UGIB) is a serious medical condition requiring prompt resuscitation and early endoscopic therapy in those with high-risk endoscopic lesions (HRLs). There are little or no data correlating sole blood urea nitrogen (BUN) level with the severity of nonvariceal UGIB or the presence of HRLs in the adult population.

OBJECTIVES: To determine if the BUN level on presentation correlates with parameters of severity of UGIB (need for blood transfusion or intensive care unit [ICU] admission) or to the subsequent finding of HRL, and in so doing identify patients who will require early endoscopic intervention.

METHODS: The Canadian Registry of patients with Upper Gastrointestinal Bleeding undergoing Endoscopy was used to identify patients enrolled from the McGill University Health Centre (Montreal, Quebec) who presented with or developed acute nonvariceal UGIB while admitted. All comparisons were performed using Student’s t test or Wilcoxon’s signed rank test, as appropriate. Logistic regression modelling using a stepwise method was performed to identify independent predictors of severe nonvariceal UGIB and HRL.

RESULTS: Two hundred nine patients were enrolled in the study. The mean age was 67±18 years and 59.8% were male. The mean BUN level was 13.4±9.4 mmol/L. Univariate analysis demonstrated that the BUN level was a significant predictor of ICU admission (BUN 14.7±10.4 mmol/L versus 12.0±8.0 mmol/L, P=0.035). However, when adjusted for systolic blood pressure, BUN level became a weaker predictor of the severity of nonvariceal UGIB or the presence of HRLs in the adult population.

CONCLUSION: In patients with nonvariceal UGIB, the BUN level at initial presentation is a weak predictor of the severity of UGIB as defined by ICU admission, but is not helpful in identifying patients with a HRL.

Key Words: BUN; High-risk lesions (endoscopic); Predictors; Upper gastrointestinal bleed

A acute nonvariceal upper gastrointestinal bleeding (UGIB) is an urgent medical condition that has significant morbidity, mortality and health care resource use (1).

The estimated annual incidence of acute UGIB is between 100 and 200 per 100,000 population (2,3). More than 50% of these cases are due to peptic ulcer disease (4-8). Patients who have a history of acute UGIB present with a wide range of clinical severity, from insignificant bleeds to catastrophic exsanguinations.

Clinical findings such as low blood pressure, postural hypotension and tachycardia can help primary and specialty physicians to differentiate severe from nonsevere UGIB.
hemostatic therapy, because this intervention has been shown to improve patient outcomes, including mortality (18).

The objectives of the present study were to first determine whether the BUN level on presentation correlates with parameters of severity of nonvariceal UGIB, identified by the need for ICU admission and blood transfusions, and second, to assess the ability of the initial BUN level to predict a subsequent finding of HRL, thus facilitating early risk stratification of patients and the need for early endoscopy.

METHODS

The Canadian Registry on nonvariceal Upper Gastrointestinal Bleeding and Endoscopy (RUGBE) (8) was used to identify patients enrolled from the McGill University Health Centre (MUHC) in Montreal, Quebec who presented with or developed acute nonvariceal UGIB while admitted. Previous publications detail the methods with regard to selection criteria and data validation of this registry (8,19).

Data were collected from the RUGBE database; patient charts were also retrieved and reviewed to collect data more specific to the present study that were not available in the RUGBE database. The registry was approved by the institutional ethics review board.

The following variables were assessed: age, sex, comorbidities, symptoms and signs of gastrointestinal bleeding, the American Society of Anesthesiologists score, vital signs, nasogastric aspiration findings, hemoglobin, serum BUN and creatinine levels, the need for blood transfusion, ICU admission and endoscopic findings (Tables 1 and 2). Patients with known renal failure (creatinine level greater than 120 μmol/L for the last three months or on dialysis), liver cirrhosis and pregnancy, and patients with predefined confounding conditions that may alter the BUN level (ie, fever, trauma, infection, on steroids) were excluded. Patients with no initial serum BUN or creatinine levels available were also excluded.

Severe UGIB was defined as the need for blood transfusion within 24 h of presentation to the emergency room or the need

### TABLE 1

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>n (%)</th>
<th>95% CI*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mean ± SD, years</td>
<td>67±18</td>
<td></td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>125 (59.8)</td>
<td>53.1–66.5</td>
</tr>
<tr>
<td>Female</td>
<td>84 (40.2)</td>
<td>33.5–47.2</td>
</tr>
<tr>
<td>BUN, mean ± SD, mmol/L</td>
<td>13.4±9.4</td>
<td></td>
</tr>
<tr>
<td>ICU admission</td>
<td>111 (53.1)</td>
<td>46.3–60.0</td>
</tr>
<tr>
<td>Blood transfusion</td>
<td>98 (45.1)</td>
<td>46.8–61.5</td>
</tr>
<tr>
<td>ICU admission and blood transfusion</td>
<td>67 (32.1)</td>
<td>25.7–38.4</td>
</tr>
<tr>
<td>Neither ICU admission nor blood transfusion</td>
<td>45 (21.5)</td>
<td>15.9–27.2</td>
</tr>
<tr>
<td>Endoscopic high-risk lesion</td>
<td>78 (37.3)</td>
<td>30.7–43.9</td>
</tr>
</tbody>
</table>

*95% CIs are reported for the percentage of patients for each characteristic. BUN Blood urea nitrogen; ICU intensive care unit

### TABLE 2

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>n (%)</th>
<th>95% CI*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Systolic blood pressure (&lt;90 mmHg)</td>
<td>190 (90.9)</td>
<td>87.0–94.8</td>
</tr>
<tr>
<td>ASA score†</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1–2</td>
<td>65 (31)</td>
<td>24.8–37.4</td>
</tr>
<tr>
<td>3</td>
<td>93 (44)</td>
<td>37.7–51.3</td>
</tr>
<tr>
<td>4–5</td>
<td>51 (24)</td>
<td>18.5–30.3</td>
</tr>
<tr>
<td>Rectal examination</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bright red blood per rectum</td>
<td>18 (9)</td>
<td>4.8–12.6</td>
</tr>
<tr>
<td>Melena</td>
<td>98 (47)</td>
<td>40.5–54.2</td>
</tr>
<tr>
<td>Occult blood-positive</td>
<td>59 (28)</td>
<td>22.3–34.7</td>
</tr>
<tr>
<td>Occult blood-negative</td>
<td>18 (9)</td>
<td>4.8–12.6</td>
</tr>
<tr>
<td>Not documented</td>
<td>16 (8)</td>
<td>4.0–11.3</td>
</tr>
<tr>
<td>Nasogastric aspirate yield</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bright red blood</td>
<td>45 (22)</td>
<td>15.7–27.4</td>
</tr>
<tr>
<td>Coffee grounds</td>
<td>52 (25)</td>
<td>19.0–30.8</td>
</tr>
<tr>
<td>Bile</td>
<td>10 (5)</td>
<td>1.5–7.1</td>
</tr>
<tr>
<td>Tube not placed or not documented</td>
<td>102 (49)</td>
<td>42.0–55.6</td>
</tr>
</tbody>
</table>

*95% CIs are reported for the percentage of patients for each characteristic; †American Society of Anesthesiologists (ASA) classification of a patient’s severity and acuity of disease

However, these clinical findings are not always reliable, especially in elderly populations with multiple comorbidities and who take many medications that may alter vital signs. In addition, the initial hemoglobin level and, in particular, the hematocrit level may not reflect a patient’s true blood level, because they may be falsely elevated as a result of hemococoncentration (9,10) or may be affected by a chronic anemic state in addition to the acute bleeding event.

Many scoring systems have been established to help clinicians manage patients with UGIB (11). One of these scoring systems, the Blatchford scale, uses blood urea nitrogen (BUN) as one of the variables to prognosticate the outcome of a patient with UGIB. However, the study population used by Blatchford et al (12) included both patients with variceal and those with nonvariceal UGIB. High BUN is mentioned in many medical textbooks as a sign of UGIB, either because of ingested blood proteins or prerenal azotemia (13).

Recently, a few studies have suggested a role for the BUN to creatinine ratio in localizing the site of gastrointestinal bleeding (14-16). Ernst et al (15) showed that the BUN to creatinine ratio correlates significantly with UGIB; a ratio higher than 36 exhibited a sensitivity of 90%, but a specificity of only 27%. Chalasani et al (16) concluded that this ratio is not specific due to a high degree of overlap between UGIB and lower gastrointestinal bleeding, especially in patients without hematemesis. Therefore, this ratio may simply reflect intravascular volume depletion and resultant prerenal azotemia. The same authors also found that the degree of elevation in the BUN to creatinine ratio correlates best with transfusion requirements rather than admission vital signs.

Endoscopic findings, in their own right, also guide the decision of whether to admit a patient, and if so, the need for an intensive care unit (ICU) admission. High-risk endoscopic lesions (HRLs), defined as the presence of active bleeding (spurting or oozing), a nonbleeding visible vessel or an adherent clot are associated with an increased risk for persistent and recurrent UGIB. Laine and Peterson (17) reported that the rate of further bleeding was less than 5% in patients with a clean ulcer base and increased to 10% in patients with a flat spot, 22% in those with an adherent clot, 43% in those with a nonbleeding visible vessel and 55% in those with active bleeding (oozing and spurting). Patients with HRLs benefit from early endoscopy within the first 24 h, as well as aggressive endoscopic
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for ICU admission because of UGIB. According to an established critical path in place at the MUHC, a blood transfusion at the MUHC is considered for patients with a hemoglobin level lower than 70 g/L or a drop in hemoglobin level by 20 g/L or greater that is not due to fluid resuscitation. Preset indications for ICU admissions due to an episode of UGIB included hypotension (systolic blood pressure [SBP] lower than 90 mmHg) refractory to fluid resuscitation. Preset indications for ICU admissions due to an established critical path in place at the MUHC at the time of delivery of care, a blood transfusion at the MUHC is considered for patients with a hemoglobin level lower than 70 g/L or a drop in hemoglobin level by 20 g/L or greater that is not due to fluid resuscitation. Preset indications for ICU admissions due to an episode of UGIB included hypotension (systolic blood pressure [SBP] lower than 90 mmHg) refractory to fluid resuscitation in the emergency room or findings of HRLs (active bleeding, spurting or oozing, nonbleeding visible vessel or adherent clot) on emergency endoscopy (requiring endoscopic hemostasis).

Statistical analysis

Standard descriptive statistics, including means, medians and SDs were used to characterize the cohort. BUN was used as a continuous variable, while blood transfusion was used as a dichotomous variable (ie, yes or no).

Comparisons were made between those with and without severe nonvariceal UGIB, as well as between those exhibiting HRLs, using Wilcoxon's signed rank test. Multivariate analysis using logistic regression modelling (stepwise method) was performed to predict ICU admission, severe UGIB and HRLs, while adjusting for possible covariates (ie, SBP, American Society for Anesthesiologists score, rectal examination and nasogastric aspirate findings) (Table 2). The 95% CIs are reported for ORs of the predictors included in the final model, with a significance threshold of P<0.05. All analyses were performed using SAS 9.1 (SAS Institute Inc, USA).

RESULTS

The RUGBE registry included 1869 patients, of which 239 were treated at the MUHC. Patients with known renal failure (n=17), confounding conditions that could alter BUN level (ie, fever, trauma, infection, on steroids) (n=8) and those with no recorded BUN or creatinine levels (n=5) were excluded, leaving 209 patients for the analysis. (Figure 1).

The study population had a mean (± SD) age of 67±18 years, with a mean BUN level of 13.4±9.4 mmol/L (Table 1). Nearly one-half of the patients (98 patients, 45.1%) required blood transfusions, and just over one-half (111 patients, 53.1%) required an ICU admission for nonvariceal UGIB. Forty-five patients (21.5%) required neither ICU admission nor blood transfusion. HRLs were found in 78 patients (37.3%).

Univariate analysis demonstrated an association between high BUN level and markers of severe nonvariceal UGIB (Table 3). The mean BUN level in the group of patients requiring ICU admission was higher than those not requiring ICU admission (14.7±10.4 mmol/L versus 12.0±8.0 mmol/L, respectively; P=0.035). There was no significant association between BUN level and blood transfusion requirements (P=0.508). The BUN levels were no different between those with a HRL and those without a HRL (14.2±10.7 mmol/L versus 12.9±8.6 mmol/L, respectively; P=0.605) (Table 3). Multivariate analysis demonstrated that, after adjustment for systolic blood pressure, BUN became a weak predictor of ICU admission for UGIB (OR 1.03, 95% CI 1.00 to 1.06, P=0.08).

DISCUSSION

A relationship between azotemia and gastrointestinal bleeding has been the main focus of many studies in the past (20-23). Elevated BUN level in UGIB patients can be due to ingested blood protein, secondary to prerenal azotemia or renal failure. It has been shown that blood ingestion results in mild to moderate azotemia (21,23). Cohn et al (24) administered 180 g of protein as whole blood (800 cc) and observed an increase in the BUN level of 21% to 35%. In our study, renal failure as a cause of high BUN level has been excluded, such that the only sources of high BUN level (from a source other than bleeding – although the latter is much less likely) would be UGIB, volume contraction or ingestion of protein. Timing of the BUN test from the onset of bleeding and site of bleeding
could be a confounding factor because ingested blood protein needs time for digestion, absorption and BUN production.

Urashima et al (25) found that severity of bleeding could be determined by the BUN to creatinine ratio and that a decrease in the BUN to creatinine ratio indicates a cessation of bleeding. In their study, a linear relationship was found between the change in hemoglobin and the BUN to creatinine ratio. However, the study was based on only 16 UGIB cases in a pediatric population. Other studies on BUN to creatinine ratio that include adult populations (14-16) were unable to correlate this ratio with severity of nonvariceal UGIB. In addition, cirrhotic patients were included in some of these studies.

Contrary to the previous belief of a strong association between high BUN level and severe UGIB, the present study found that the BUN level did not predict the need for blood transfusion \( (P=0.508) \) (Table 3). The high BUN level in the group of patients not requiring blood transfusion could be secondary to prerenal azotemia or volume contraction from non-significant blood loss in patients with borderline renal function or perfusion that is not reflected by an acute rise in creatinine levels. Another explanation for the lack of correlation between high BUN level and the need for blood transfusion, is the strict criterion for blood transfusion implemented at the MUHC. Interestingly, a modified Blatchford score with predictive value was recently proposed that excluded BUN level as a predictive marker of the severity of UGIB (12).

The present study also demonstrates that an elevated BUN level is a weak predictor of the need for ICU admission, especially when BUN level is adjusted for other covariates such as low SBP \( (OR\; 1.03,\; 95\%\; CI\; 1.00\; to\; 1.06,\; P=0.08) \). This is partly due to collinearity between low blood pressure and high BUN levels. It is also possible that the high BUN level may have been taken into account by the physician when deciding to admit a patient to the ICU. Hence, the BUN level may not be a predictor of severity of nonvariceal UGIB but simply a factor that may have partly determined whether the patient was admitted to the ICU.

Endoscopic findings can guide the decision for discharge, hospital admission or admission to an ICU. Active bleeding (spurting or oozing) or a nonbleeding visible vessel seen on endoscopy have the highest potential for rebleeding, with rates of approximately 55% and 43%, respectively \( (17) \). Predicting these findings, especially in patients without frank hematemesis, will help guide the clinician in deciding the need for an urgent endoscopy, given the protective role of endoscopic therapy in patients with HRLs \( (8) \). Unfortunately, our study found that a high BUN level was not significantly associated with the subsequent finding of HRLs \( (P=0.605) \) (Table 3). To our knowledge, this is the first study to examine the correlation between BUN level and HRLs. The lack of correlation between high BUN level and HRLs, however, is not surprising given that a strong association between high BUN level and parameters of severe nonvariceal UGIB was not found.

CONCLUSIONS

In patients with nonvariceal UGIB, an elevated BUN level at initial presentation is a weak predictor of severity of bleeding when also accounting for other factors such as low SBP. Also, an elevated BUN level was not helpful in identifying patients with HRLs.

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BUN level as a predicro of UGIB severity
