Hyperglycemia and acute kidney injury in critically ill children

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Background: Hyperglycemia and acute kidney injury (AKI) are common in critically ill children and have been associated with higher morbidity and mortality. The incidence of AKI in children is difficult to estimate because of the lack of a standard definition for AKI. The pediatric RIFLE (Risk, Injury, Failure, Loss of kidney function, and End-stage kidney disease) criteria can be used to define AKI in children. Various biomarkers in urine and blood have been studied to detect AKI in critically ill children. However, it is not clear whether hyperglycemia is associated with AKI. Our objective was to evaluate the effect of hyperglycemia on kidney function and its effect on neutrophil gelatinase-associated lipocalin (NGAL) in children.

Methods: We studied retrospective and prospective cohorts of pediatric critically ill subjects admitted to the pediatric intensive care unit (PICU). We analyzed data from admission that included estimated glomerular filtration rate, plasma and urine NGAL, serum glucose and peak glycemia (highest glycemia during PICU admission), and length of hospital and PICU stay from two different institutions.

Results: We found that the prevalence of hyperglycemia was 89% in the retrospective cohort and 86% in the prospective cohort, P=0.99. AKI was associated with peak glycemia, P=0.03. There was a statistically significant correlation between peak glycemia and hospital and PICU stays, P<0.001 and P<0.001, respectively. Urine NGAL and plasma NGAL were not statistically different in subjects with and without hyperglycemia, P=0.99 and P=0.85, respectively. Subjects on vasopressors had lower estimated glomerular filtration rate and higher glycemia, P=0.01 and P=0.04, respectively.

Conclusion: We conclude that in critically ill children, hyperglycemia is associated with AKI and longer PICU stays.

Keywords: AKI, hyperglycemia, NGAL, peak glycemia, eGFR, PICU

Introduction

Hyperglycemia is common in critically ill patients, and it has been associated with increased mortality and morbidity in adults and children. Srinivasan et al found that blood glucose level >150 mg/dL was associated with a 3.5-fold increased risk of mortality. In critically ill adults, a large prospective randomized clinical study showed that strict control of blood glucose with insulin therapy reduced the incidence of acute kidney injury (AKI). In children, the incidence of AKI is difficult to estimate because of the lack of a standard definition for AKI and lack of large prospective multicenter studies involving general intensive care population, making the AKI incidence prone to definition and center-specific bias. Over 30 definitions of AKI have been published, and pediatric AKI definition has been “extrapolated” from adult studies. AKI in children is associated with significant mortality, as high as 60%. Therefore, prevention of AKI is crucial, and includes hemodynamic support, maintenance of euvoembia, and avoidance of nephrotoxic medications. However, it
is not clear whether hyperglycemia is associated with AKI, and whether it has an impact on AKI outcomes in critically ill children. A number of biomarker studies focused on early detection of AKI, and reference levels of these biomarkers in a healthy pediatric population have since been established. Neutrophil gelatinase-associated lipocalin (NGAL) is a protein that is over-expressed in the kidneys after ischemia, and elevated NGAL serum and urine levels have been used as an early biomarker of AKI. However, it is unclear if NGAL levels are affected by hyperglycemia in critically ill children with AKI.

The aim of our study was to evaluate the association of hyperglycemia with AKI in critically ill children and examine the effect of hyperglycemia on NGAL levels.

Materials and methods

Pediatric subjects admitted to the pediatric intensive care unit (PICU) were recruited in two cohorts in this study; retrospective cohort in University of South Alabama Children’s & Women’s Hospital with recruitment between June 2007 and March 2009, and prospective cohort in Children’s Hospital of Illinois between January 2012 and July 2013. The institutional review board of University of South Alabama Children’s & Women’s Hospital, and the Children’s Hospital of Illinois evaluated and approved this study. Informed consent was obtained from all parents in the prospective arm of the study - the retrospective arm was a chart review and no consent was required.

Definitions

We defined critically ill children as those who required mechanical ventilation within 36 hours of admission to the PICU. AKI was defined by the pediatric RIFLE (Risk, Injury, Failure, Loss of kidney function, and End-stage kidney disease) criteria. Peak glycemia was defined as the highest serum glucose level during PICU admission. Hyperglycemia was defined as serum glucose of 126 mg/dL and higher. Estimated glomerular filtration rate (eGFR) was calculated by Schwartz’s formula.

Study criteria

Children aged 1–18 years admitted to PICU who were critically ill were included in the study. Children with the diagnosis of diabetes mellitus type 1 and 2, children who died within 48 hours of admission, children who had undergone a kidney transplant, children on renal replacement therapy, children with known chronic kidney disease, children with immediate postcardiac surgery, children with tracheostomy, and pregnant children were excluded from the study.

Data collection

Following laboratory data were collected upon admission to PICU: serum glucose, sodium, potassium, chloride, bicarbonate, creatinine, and blood urea nitrogen. Glomerular filtration rate was estimated (eGFR) based on bedside Schwartz equation 8, and urine was collected using a Foley catheter. In the prospective cohort, 1.5 mL aliquots of plasma were centrifuged at 4°C for 15 minutes at 800 g, and then stored at −80°C. Urine was also stored at −80°C before shipping to Cincinnati Children’s Hospital Medical Center to measure NGAL.

Data analysis

Except where otherwise noted, data were expressed as mean ± standard deviation or as percentages. The software used for our analysis included SAS version 9.4 (SAS Institute Inc., Cary, NC, USA) and Microsoft Office Excel 2010 (Microsoft Corporation, Redmond, WA, USA). Statistical tests used included Mann–Whitney test for non-normally distributed variables, Fisher’s exact test to examine the differences in proportions between the groups and Spearman correlation coefficients. All analyses were two sided unless specified otherwise. Statistical significance for all analyses was set at P<0.05.

Results

A total of 67 subjects met inclusion criteria; 37 were recruited retrospectively and 30 prospectively. All the subjects had respiratory failure due to various diagnoses: asthma, septic shock, and pneumonia. The incidence of hyperglycemia was 59/67 (88%) and it was similar in the retrospective and prospective cohorts, 33/37 (89%) and 26/30 (86%), respectively, P=0.99. Table 1 shows all subject characteristics. Peak glycemia was significantly higher in subjects with AKI than in subjects without AKI. The length of hospital and PICU stays were longer in subjects with AKI, but the length of hospital stay did not reach a statistically significant difference. Likewise, serum NGAL and urine NGAL were higher in the AKI group, but did not reach a significant difference (Table 1).

<table>
<thead>
<tr>
<th>Subject demographics</th>
<th>Non-AKI (n=42)</th>
<th>AKI (n=25)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>6.96±0.89</td>
<td>6.2±1.01</td>
<td>0.92**</td>
</tr>
<tr>
<td>Female (%)</td>
<td>47.6 (N=20)</td>
<td>48 (N=12)</td>
<td>0.57**</td>
</tr>
<tr>
<td>Caucasian (%)</td>
<td>52.4 (N=22)</td>
<td>44 (N=11)</td>
<td>0.31**</td>
</tr>
<tr>
<td>Hyperglycemia (%)</td>
<td>85.7 (N=36)</td>
<td>92 (N=23)</td>
<td>0.7**</td>
</tr>
<tr>
<td>Peak glucose (mg/dL)</td>
<td>184.73 (IQR 74)</td>
<td>249.84 (IQR 107)</td>
<td>0.003*</td>
</tr>
<tr>
<td>Length of hospital stay (days)</td>
<td>25 (IQR 14)</td>
<td>40.2 (IQR 31)</td>
<td>0.26*</td>
</tr>
<tr>
<td>Length of PICU stay (days)</td>
<td>16.6 (IQR 11)</td>
<td>30.8 (IQR 18)</td>
<td>0.03*</td>
</tr>
<tr>
<td>Urine NGAL (ng/mL)</td>
<td>21.63 (IQR 15.8)</td>
<td>407.9 (IQR 111.6)</td>
<td>0.47*</td>
</tr>
<tr>
<td>Plasma NGAL (ng/mL)</td>
<td>113.9 (IQR 85.8)</td>
<td>291.4 (IQR 418.6)</td>
<td>0.11*</td>
</tr>
</tbody>
</table>

Note: *Mann–Whitney. **P-value tested by Fisher’s exact test.

Abbreviations: AKI, acute kidney injury; IQR, interquartile range; PICU, pediatric intensive care unit; NGAL, neutrophil gelatinase-associated lipocalin.
There were two deaths, one in the AKI group and the other in the non-AKI group. Subjects with hyperglycemia had significantly longer stays in PICU than subjects without hyperglycemia (Table 2). Subjects with hyperglycemia had also longer hospital stay, but it did not reach statistical significance. The correlation between peak glycemia and length of hospital stay and PICU stay was statistically significant, Spearman’s rho = 0.49, P ≤ 0.001 and Spearman’s rho = 0.53, P ≤ 0.001, respectively. In the prospective cohort, urine NGAL was higher in subjects with hyperglycemia but did not reach a statistically significant difference. Plasma NGAL also did not reach a significant difference between the prospective cohort subjects with and without hyperglycemia.

The subjects who required vasopressors had significantly higher peak glycemia and lower eGFR than the subjects who did not require vasopressors (Table 3). The correlation between peak glycemia and eGFR on admission and the correlation between peak glycemia and urine output during PICU stay were statistically significant, Spearman’s rho = −0.277, P = 0.047 and 0.26, P = 0.039, respectively.

Logistic regression was used to assess the effect of peak glycemia on AKI. Peak glycemia had a statistically significant effect on AKI, with an odds ratio of 1.12 (1.03–1.2), P = 0.0051.

Table 2 Kidney function, NGAL, and length of hospital and PICU stays in subjects with hyperglycemia and normoglycemia

<table>
<thead>
<tr>
<th>Variable</th>
<th>Normoglycemia</th>
<th>Hyperglycemia</th>
<th>P-value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>eGFR on admission (mL/min/1.73 m²)</td>
<td>110.6±37</td>
<td>111.2±56.9</td>
<td>0.716</td>
</tr>
<tr>
<td>Length of hospital stay (days)</td>
<td>14.5</td>
<td>32.9</td>
<td>0.07</td>
</tr>
<tr>
<td>Length of PICU stay (days)</td>
<td>7.6</td>
<td>23.8</td>
<td>0.03</td>
</tr>
<tr>
<td>Urine NGAL (ng/mL) (IQR 25.76)</td>
<td>16.5</td>
<td>119.1</td>
<td>1.00</td>
</tr>
<tr>
<td>Plasma NGAL (ng/mL) (IQR 16.31)</td>
<td>195.8 (IQR 326.1)</td>
<td>159.8 (IQR 88.54)</td>
<td>0.85</td>
</tr>
</tbody>
</table>

Note: *P-value calculated by Mann–Whitney test.
Abbreviations: NGAL, neutrophil gelatinase-associated lipocalin; PICU, pediatric intensive care unit; eGFR, estimated glomerular filtration rate; IQR, interquartile range.

Table 3 Peak glycemia and kidney function in subjects on vasopressors and without vasopressors

<table>
<thead>
<tr>
<th>Variable</th>
<th>Without vasopressors N=50</th>
<th>With vasopressors N=17</th>
<th>P-value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Peak glycemia (mg/dL)</td>
<td>194.52±9.75</td>
<td>251.7±26.03</td>
<td>0.04</td>
</tr>
<tr>
<td>eGFR (mL/min/1.73 m²)</td>
<td>118.88±7.46</td>
<td>88.69±13.76</td>
<td>0.01</td>
</tr>
</tbody>
</table>

Note: *P-value calculated by Mann–Whitney test.
Abbreviation: eGFR, estimated glomerular filtration rate.

Discussion

In summary, our study demonstrates that hyperglycemia is common in critically ill children and is associated with AKI. We showed that odds of having AKI increase 12% for an every increase in peak glycemia by 10 mg/dL. We also showed that hyperglycemia has an impact on PICU length of stay. Other studies have reported that the incidence of hyperglycemia in critically ill children is between 14% and over 90%, we found an incidence of 88%. A better glycemic control in these children might improve outcomes.

It has been reported that oxidative stress plays an important role in the complications of diabetes, and hyperglycemia causes oxidative stress in different ways. Vanhorebeek et al reported a rabbit mode of prolonged critical illness and found elevated plasma creatinine, severe morphological kidney damage correlated with elevated glucose levels. Renal cortical perfusion and oxygen delivery were lower in the rabbits with hyperglycemia. Mitochondrial respiratory chain activities were severely reduced in rabbits with hyperglycemia. Langouche et al showed a protective effect of intensive insulin therapy on the vascular endothelium and hepatocyte mitochondria of critically ill subjects. We can speculate that a mitochondrial injury also occurs in critically ill children with hyperglycemia. The proximal tubule is sensitive to mitochondrial injury; this might be part of the AKI pathogenesis. In rabbits, it has been shown that hyperglycemia causes a morphological kidney damage that correlates with diabetes level and plasma creatinine, and this is associated with mitochondrial dysfunction. Mitochondrial injury plays an important role in AKI. In mice, it has been shown that hyperglycemia increases oxidative and nitrosative stress and accelerates renal injury. We can speculate that subjects with AKI and hyperglycemia may have kidney dysfunction at the mitochondrial level. This can also explain our finding of glycemia having inverse correlation with eGFR.

Our subjects with hyperglycemia stayed longer in the PICU than subjects with normoglycemia. It has been shown that hyperglycemia occurs more frequently in sicker patients, and this might explain the longer stays in hospital and PICU. As sicker patients may require cardiovascular support, it may explain the increased use of pressors in this population. We did find a significant correlation between peak glycemia and PICU/hospital stay and this is in agreement with a study of pediatric subjects.

Subjects with higher glycemia were more likely to be on vasopressors; this is in agreement with other studies. Preissig and Rigby found that with increasing support measures (mechanical ventilation, vasopressors), the risk...
of hyperglycemia increases. As mentioned before, sicker subjects have higher glycemia and these subjects usually require vasopressor support. Our data suggest that subjects that were on vasopressors were sicker (lower eGFR) and had higher glycemia; this is in agreement with other studies.

Our study found no significant difference in urine and plasma NGAL between the subjects with and without hyperglycemia, and subjects with and without AKI. It is possible that the injury caused by hyperglycemia in the mitochondria does not cause overexpression of NGAL. Another explanation could be the small number of subjects in this study.

Limitations
This study was limited by the small sample size. None of our subjects had kidney biopsy as they are rarely performed in patients with AKI. We cannot therefore, make a statement about structural/mitochondrial damage, as this is an observational study, cause–effect relationship cannot be ascertained. Another limitation is that we have no data on steroid and intravenous dextrose use in our subjects. It has been shown that hyperglycemia is more common in critically ill children and also with the use of steroids for various indications, such as shock and asthma, and all of our subjects were ventilated. The use of dextrose in intravenous fluids might explain the high incidence of hyperglycemia in these children. However, both of our groups were ventilated and we do not suspect that steroid/dextrose was used preferentially in one group over the other creating systematic bias.

Conclusion
Despite those limitations, to the best of our knowledge, this is the first study that describes relationship of AKI and hyperglycemia in critically ill children. We conclude that critically ill children with hyperglycemia have higher incidence of AKI and longer PICU stays. Peak glycemia correlates with lower eGFR on admission. We acknowledge that several variables might affect kidney function in these subjects; however, confirmation of these findings in larger prospective studies may provide clinical evidence for serum glucose as a biomarker of risk for AKI in critically ill children and lay foundation to clinical intervention trials to tightly regulate glycemia in selected critically ill children.

Author contribution
All authors contributed in data analysis and revised the paper.

Disclosure
The authors report no conflicts of interest in this work.

References