Evaluation of clinical and laboratory variables associated with anemia in pediatric patients on hemodialysis

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KEYWORDS
Anemia; Adolescent; Child; Kidney dialysis; Renal failure

Abstract
Objective: To identify the occurrence of anemia in pediatric patients on hemodialysis and the association between hemoglobin levels and anemia in CKD-related variables.
Methods: This was a retrospective study. Patients aged up to 18 years with chronic kidney disease undergoing hemodialysis at this service between January of 2009 and December of 2010 were selected. Clinical and laboratory data were obtained from medical records. Statistical analysis was performed with chi-squared test, Student’s t-test and general estimating equations (GEE) using SPSS 20.0, assuming a significance level of 5%.
Results: A total of 357 medical records depicting the monthly evolution of 29 patients were analyzed. The most common etiology for chronic kidney disease was malformations of the genitourinary tract (28%). Hemoglobin showed a mean (standard deviation) value of 9.20 (1.8) g/dL, with the occurrence of anemia in 65.3% of cases. Anemia was associated with hospitalization; antibiotic use; transfusion; use of intravenous iron hydroxide; low values of creatinine, hemoglobin, and albumin; and high values of ferritin, aluminum, and equilibrated Kt/V (p < 0.05). The odds ratio for anemia with the use of intravenous iron hydroxide was 0.36 (95% CI: 0.25 to 0.89), i.e., a 2.78-fold higher chance of developing anemia without the use of this medication.
Conclusions: Anemia predominated in children and adolescents with chronic kidney disease; intravenous iron hydroxide use was a protective factor.
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Introduction

The prevalence of renal replacement therapy in children aged 0-19 years in Brazil is of 23:1,000,000 in the related age range. The survival of these children is approximately 30-fold lower than that of their healthy peers. The main cause of mortality in this group is cardiovascular disease, accounting for 40% of deaths, and anemia is identified as a factor related to higher rates of mortality.\(^1,2\)

The prevalence of anemia in Brazilian children without a diagnosis of any disease ranges from 25.6% to 63.7%,\(^3,4\) while in children with chronic kidney disease (CKD), chronic hypoprogenetic anemia is common. If untreated, it can lead to growth and development impairment, left ventricular hypertrophy, and tachycardia. When on hemodialysis, blood loss in the dialysis circuit and blood collections due to frequent examinations contribute to further exacerbation of anemia.\(^5\)

Anemia is a limiting survival factor of children on hemodialysis. Thus, this study aimed to identify the occurrence of anemia in pediatric patients on hemodialysis and the association between hemoglobin levels and anemia-related variables in CKD.

Methods

Study design and setting

This was a retrospective cohort study approved by the Ethics Committee in Human and Animal Medical Research, of Hospital das Clínicas da Universidade Federal de Goiás (HC/UFG). Data were collected at the Renal Replacement Therapy Service, Hemodialysis Sector, HC/UFG.

Study participants

The study participants were subjects undergoing hemodialysis in HC/UFG between January of 2009 and December of 2010 who met the inclusion criteria: age < 18 years and diagnosis of CKD. Cases whose medical records were not available were excluded.

Data collection and variables

An expert researcher (a pediatric nephrologist) collected the data on the variables of interest from patients’ records, specifically the records of monthly medical progress, and recorded them in digital form. Record survey was conducted at the Hemodialysis Sector and the Division of Medical Records and Health Information of HC/UFG.

The dependent variable was the occurrence of anemia, measured by hemoglobin levels (mg/dL).

The independent variables were: clinical variables - hospitalizations, antibiotic use in the month (whether the patient used or not), transfusions of packed red blood cells, and main drugs used to treat anemia (folic acid, B complex, vitamin C, erythropoietin, and intravenous iron hydroxide); laboratory variables - serum tests performed as routine at the service, defined by Resolution No. 154 of the Brazilian National Health Surveillance Agency, with relevant adjustments: urea at pre-hemodialysis session, creatinine, hemoglobin, hematocrit, pH, bicarbonate, ferritin, transferrin saturation index (TSI), iron, parathyroid hormone (PTH),...
albumin, and aluminum. Equilibrated Kt/V (EKT/V) was calculated as proposed by Fischbach et al.

### Statistical Analysis

Data were tabulated and analyzed using descriptive and inferential statistics using IBM SPSS Statistics software, release 20 (IBM Corporation, NY, USA).

After exploratory data analysis, continuous variables were categorized based on clinical parameters to better interpret the results, considering the limited number of subjects included in the study.

Serum hemoglobin values were classified as adequate or inadequate (anemia) as recommended by the National Institute for Health and Clinical Excellence (NICE) criteria of 2011, which considers both age groups (0-2 years and over 2 years). Student’s t-test and the chi-squared test were then used to test associations between the described variables and the occurrence of anemia.

This longitudinal observational study included repeated measurements for the same variable in the same subject. As the dependent variable was categorical (having or not anemia), the analysis of variance for repeated measures could not be used. Moreover, multiple linear regression would be contraindicated because the assumption of independence of random variables could not be met, as the data had an interdependent association (repeated measures). Thus, the statistical analysis that would better determine predictive variables for anemia were the generalized estimating equations (GEEs).

The GEE approach, which is an extension of generalized linear models, was developed to produce more efficient and less biased regression estimates for use with correlated data, as repeated longitudinal measures. Thus, GEE analyzes data depending on exposure over successive periods of time. For the analysis of this study, a model for the dependent variable “anemia” was created. The independent variables were “hospitalization”, “infection of double-lumen catheter (DLC)”, “antibiotic use,” “use of intravenous iron hydroxide.” Missing data were treated by the GEE mechanism that uses all available data to include evasion data. The odds ratio (OR) and confidence intervals (95% CI) were calculated to reflect the possible associations between predictive factors and dependent variable.

All statistical tests were two-tailed, with a significance level of 0.05.

### Results

A total of 31 patients were included, but two were excluded due to inaccessible files. The final sample consisted of 29 patients, corresponding to 357 files of monthly clinical evolution throughout 24 months. Each patient was followed-up for one to 23 consultations. The mean age of patients at baseline was 10 years and 3 months (standard deviation [SD] 34.6 months). Of the 29 patients included, 21 were males (72.4%). The etiologies of CKD with their respective percentages are shown in Fig. 1.

All patients but one had anemia at some point during their follow-up, which corresponded to 65.3% of consultations (n = 233). The mean hemoglobin (g/dL) for the categorized groups were: anemic - 8.2 g/dL (SD 1.2); non-anemic - 11.2 g / dL (SD 1.0) (p < 0.001). The mean hemoglobin value for the total sample was 9.2 (SD 1.8).

The bivariate analysis showed that anemia was associated with several clinical situations (Table 1): need for hospitalization, antibiotic use, transfusion, and less frequent use of intravenous iron hydroxide (p < 0.05). As for the laboratory findings, anemia was related to lower levels of creatinine, hematocrit, and albumin, as well as to higher values of ferritin and aluminum, and EKT/V (Table 1).

Based on the results of the chi-squared and Student’s t-test, the authors sought to determine the predictive clinical factors for anemia in this group of patients, considering the clinical variables that reached p < 0.2 in the bivariate analysis. Among these, the variable “need for transfusion” was excluded, as it is not logical to evaluate this variable as a predictor of anemia. Thus, through GEE, it was

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Figure 1  Etiology of chronic kidney disease.
HUS, hemolytic uremic syndrome; ARPKD, autosomal recessive polycystic kidney disease; FSGS, focal and segmental glomerulosclerosis; GUT, genitourinary tract.
Total = 100%.
```
Table 1  Association between clinical and laboratory variables and the occurrence of anemia in children with chronic kidney disease on hemodialysis (357 files of 29 patients).

<table>
<thead>
<tr>
<th>Independent variables</th>
<th>n</th>
<th>Total sample</th>
<th>Anemia</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Yes (n = 233)</td>
<td>No (n = 124)</td>
<td></td>
</tr>
<tr>
<td><strong>Medical history, n (%)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Need for hospitalization</td>
<td>316</td>
<td>52 (14.6%)</td>
<td>43 (20.7%)</td>
<td>9 (8.3%)</td>
</tr>
<tr>
<td>Antibiotic use</td>
<td>316</td>
<td>60 (16.8%)</td>
<td>48 (23.1%)</td>
<td>12 (11.1%)</td>
</tr>
<tr>
<td>Need for transfusion</td>
<td>316</td>
<td>11 (3.1%)</td>
<td>56 (26.9%)</td>
<td>3 (2.8%)</td>
</tr>
<tr>
<td>Double-lumen catheter infection</td>
<td>316</td>
<td>59 (16.5%)</td>
<td>10 (4.8%)</td>
<td>1 (0.9%)</td>
</tr>
<tr>
<td><strong>Main medications used in the treatment of anemia, n (%)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Folic acid</td>
<td>339</td>
<td>333 (93.3%)</td>
<td>222 (98.7%)</td>
<td>111 (97.4%)</td>
</tr>
<tr>
<td>Vitamin B complex</td>
<td>339</td>
<td>332 (93.0%)</td>
<td>221 (98.2%)</td>
<td>111 (97.4%)</td>
</tr>
<tr>
<td>Vitamin C</td>
<td>339</td>
<td>327 (91.6%)</td>
<td>219 (97.3%)</td>
<td>108 (94.7%)</td>
</tr>
<tr>
<td>Erythropoietin</td>
<td>321</td>
<td>303 (84.9%)</td>
<td>201 (94.4%)</td>
<td>102 (94.4%)</td>
</tr>
<tr>
<td>Intravenous iron hydroxide</td>
<td>318</td>
<td>160 (44.8%)</td>
<td>89 (42.4%)</td>
<td>71 (65.7%)</td>
</tr>
<tr>
<td><strong>Laboratory exams, mean (standard deviation)</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pre-hemodialysis session urea (mg/dL)</td>
<td>353</td>
<td>140.5 (42.3)</td>
<td>143.2 (42.9)</td>
<td>135.5 (40.8)</td>
</tr>
<tr>
<td>Creatinine (mg/dL)</td>
<td>352</td>
<td>6.2 (2.1)</td>
<td>5.9 (2.1)</td>
<td>6.6 (2.1)</td>
</tr>
<tr>
<td>Hematocrit (%)</td>
<td>356</td>
<td>28.3 (5.7)</td>
<td>25.1 (4.0)</td>
<td>34.3 (3.0)</td>
</tr>
<tr>
<td>Power of hydrogen (pH) (AV)</td>
<td>119</td>
<td>7.4 (0.6)</td>
<td>7.4 (0.1)</td>
<td>7.4 (0.1)</td>
</tr>
<tr>
<td>Bicarbonate (mmol/L)</td>
<td>120</td>
<td>20.2 (3.8)</td>
<td>19.8 (3.4)</td>
<td>20.7 (4.1)</td>
</tr>
<tr>
<td>Ferritin (ng/L)</td>
<td>351</td>
<td>709.3 (460.0)</td>
<td>754.0 (479.1)</td>
<td>625.1 (413.1)</td>
</tr>
<tr>
<td>Transferrin saturation index (TSI) (%)</td>
<td>357</td>
<td>35.5 (19.4)</td>
<td>36.0 (20.2)</td>
<td>34.5 (17.9)</td>
</tr>
<tr>
<td>Iron (µg/dL)</td>
<td>340</td>
<td>75.7 (93.5)</td>
<td>74.7 (95.9)</td>
<td>78.0 (89.8)</td>
</tr>
<tr>
<td>Parathormone (PTH) (pg/mL)</td>
<td>347</td>
<td>520.3 (495.1)</td>
<td>526.2 (519.4)</td>
<td>513.2 (450.7)</td>
</tr>
<tr>
<td>Albumin (g/dL)</td>
<td>343</td>
<td>3.9 (0.5)</td>
<td>3.8 (0.6)</td>
<td>4.1 (0.2)</td>
</tr>
<tr>
<td>Aluminum (µg/dL)</td>
<td>167</td>
<td>24.1 (28.6)</td>
<td>28.0 (30.8)</td>
<td>16.0 (21.6)</td>
</tr>
<tr>
<td>Equilibrated Kt/V (eKt/V) (AV)</td>
<td>294</td>
<td>1.9 (0.4)</td>
<td>2.0 (0.4)</td>
<td>1.8 (0.4)</td>
</tr>
</tbody>
</table>

Values in bold show statistically significant associations (p < 0.05).  
AV, absolute value.  
a Pearson’s chi-squared test.  
b Student’s t-test.

Table 2  Clinical factors predictive of anemia in children and adolescents on hemodialysis.

<table>
<thead>
<tr>
<th>Independent variables</th>
<th>Odds ratio</th>
<th>95% Confidence interval</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Minimum</td>
<td>Maximum</td>
<td></td>
</tr>
<tr>
<td>Need for hospitalization</td>
<td>1.00</td>
<td>0.52</td>
<td>1.93</td>
</tr>
<tr>
<td>Antibiotic use</td>
<td>1.97</td>
<td>0.89</td>
<td>4.35</td>
</tr>
<tr>
<td>Double-lumen catheter infection</td>
<td>4.13</td>
<td>0.49</td>
<td>34.60</td>
</tr>
<tr>
<td>Use of intravenous iron hydroxide</td>
<td>0.36</td>
<td>0.25</td>
<td>0.89</td>
</tr>
</tbody>
</table>

demonstrated that the use of intravenous iron hydroxide was a protective factor for anemia (Table 2): the odds the children on hemodialysis to have anemia when using this medication was 0.36 times the odds of those who did not use it. Calculating the inverse of 0.36, it is observed that the chance of children who did not use intravenous iron hydroxide to have anemia was 2.78 times the chance of those who used it. The other variables did not show such significant predictors in the GEE.

Discussion

This article emphasizes the high prevalence of anemia in children undergoing hemodialysis, in spite of the administration of recombinant human erythropoietin, and reaffirms the importance of intravenous iron supplementation in this population.

In relation to the clinical aspects of this study, it is noteworthy that, among the causes of CKD, congenital malformations of the urinary tract were the most prevalent, consistent with data obtained in developed and developing countries. However, neurogenic bladder was the second most prevalent cause observed in the present study and by researchers in Turkey, while glomerulonephritis and hereditary nephropathies were observed in developed countries. This may reflect a delay in the diagnosis and treatment of urological problems in developing countries.1,11-13
It is worth mentioning the use of medications for the treatment of CKD-associated anemia: 94.4% used erythropoietin; 50.5% used intravenous iron hydroxide in the present study. In the Brazilian dialysis census of 2011, prescriptions of these medications were 80.0% and 53.1%, respectively.14

The target-hemoglobin values for children and adolescents with CKD in the literature are divergent and have been revised over the past years.12,15,16 It is known that, in children and adolescents, age and gender should be taken into account in order to define target hemoglobin (Hgb) values.17 The recommendation of the last NICE criteria6 to define Hgb in patients as adequate or inadequate was adopted; therefore, the present study showed that Hgb values below the expected (anemia) were found in most monthly records of patients (65.3%), albeit still above those found in the last Brazilian dialysis census (39.3%).14

The authors believe that this difference is due to the study methodology, in which the same patient with anemia was reviewed during successive months, overestimating the frequency of anemia. In other studies, the hemoglobin value is usually considered in a single annual measurement (only one month of the year); the authors believe that this, conversely, may underestimate the prevalence of anemia. The mean Hgb observed was 9.2 (SD 1.8) g/dL. Comparatively, American data show a mean Hgb of 11.5 (SD 1.6) g/dL, with 68% of patients showing the target Hgb values.2

Data from the United Kingdom showed that most children on renal replacement therapy are anemic (47% of children on hemodialysis).18 A Polish study evidenced a mean Hgb of 10.91 (SD 1.2) g/dL.19

It is known that serum Hgb varies over the months, and withdrawal of erythropoietin in the last 60 days and hospitalization are related to Hgb < 11 g/dL. The longer the hospital length of stay, the greater the chance of more significant decreases in Hgb values.20 A 14.6% rate of hospitalization was observed in the present study, which may also have contributed to the anemia in the present population. Appropriate values of serum albumin are related to adequate Hgb values.21 Adequate mean serum albumin levels were observed in the present study, in agreement with other authors.15,19

Good markers of ferritin and transferrin saturation index (TSI) were identified, as the iron stores are replenished intravenously, as recommended in the literature for over a decade for hemodialysis patients receiving human erythropoietin.22,23

In 91.6% of the files, patients were receiving vitamin C, which is important for release of iron stored in the body, ensuring its availability for erythropoiesis.24,25

When the GEE was performed, intravenous iron hydroxide received during that month showed to be protective against anemia. The variable "transfused" was removed from the GEE analysis, as it ultimately reflects the event throughout the month. A blood sample was collected for the monthly tests on the first Thursday of each month. Those who met the criteria for receiving packed red blood cells (including the low Hgb values) received the blood product in subsequent sessions. Thus, the answer "yes" to the variable "transfused" meant that, during that month, hemoglobin levels were very low and the patient required blood transfusion. Therefore, the transfusion event could be erroneously indicated as a risk factor for anemia in the GEE.

The main limitation of this study was its retrospective design, with analysis of data found in medical records. The lack of control in data collection and information loss constitute biases of retrospective studies. These were observed in the present study, compromising the sample and decreasing the power of data representation. The longitudinal repeated measures were treated by GEE.

In conclusion, the study population consisted of children and adolescents with end-stage renal disease on hemodialysis. Anemia was prevalent and the use of intravenous iron hydroxide was a protective factor.

Conflicts of interest

The authors declare to have no conflicts of interest.

References