Introduction

Hypophosphatemia is a metabolic disturbance with potential serious complications and is often unrecognized in critically ill children (CIC) [1]. In a review of clinical studies done on hypophosphatemia in pediatric intensive care unit (PICU) patients, its prevalence exceeded 50% [2]. Phosphate ions are critical for normal bone mineralization, and phosphate plays a vital role in a number of other biological processes such as signal transduction, nucleotide metabolism, adenosine triphosphate (ATP) production and enzyme regulation [3].

Symptoms of hypophosphatemia tend to be nonspecific in the majority of cases and include fatigue and irritability. However, severe hypophosphatemia (less than 1.0 mg/dl) may lead to more serious problems [4] such as reduced diaphragmatic contractility [5], cardiac arrhythmias [6], myocardial reduction and severe congestive cardiac insufficiency in the postoperative period of cardiac surgery [7], leukocyte dysfunction [6] and neuromuscular disturbances [8]. Potential risk factors in most patients with hypophosphatemia in the literature include refeeding syndrome [9], malnutrition [10], starvation for over 3 days, sepsis, Pediatric Index of Mortality (PIM) [2,11], catecholamine’s and antacids [12], trauma, diuretic, steroid therapy [2], excessive parenteral glucose administration and respiratory alkalosis [13].

The aim of this study was to estimate the prevalence of hypophosphatemia and to identify risk factors and outcome associated with this disturbance in children admitted to our PICU.

Materials and Methods

Study design and Population

All infants and children admitted to Cairo University Children Hospital Medical PICU in the period from July 2010 through December 2010 were consecutively and prospectively enrolled excluding those with renal insufficiency. Sample size was determined using power analysis. Power was computed through the statistical package used and the value was 0.896 with alpha level = 0.05, power = 80%. Sample size was 68 and we have chosen 72 cases. The study was approved by Cairo University Ethics Committee and was conducted in accordance with the bylaws for human research. The study was explained and consents were obtained from all parents or legal guardians before enrollment. We had 4 drop out patients due to laboratory test availability (absent serum phosphorus level).
Drop outs were included in the study and were considered as normophosphatemic patients during the statistical analysis.

Patients were subjected to

Initial evaluation:

a. History and clinical examination.

b. Routine laboratory investigations including: complete blood picture, erythrocyte sedimentation rate (ESR), C-reactive protein (CRP), serum electrolytes, serum aminotransferases, blood urea nitrogen (BUN), serum creatinine as well as the appropriate bacterial cultures.

c. Assessment of severity of illness on admission using the PIM. Variables are: Elective admission, specific diagnosis, pupils fixed more than 3mm in bright light, mechanical ventilation, systolic blood pressure, Base excess, FiO2, and PCO2.

d. The following variables were collected: age, gender, diagnosis, malnutrition (± - 2 SD weight for height), phosphorus intake if any, clinical severity score at admission (PIM score), sepsis (whether clinically suspected (multisystem organ failure, bad general condition) or laboratory proved (+ve culture, +ve CRP, leucocytosis), use of dopamine, frusemide, steroids and starvation period (the period without enteral feeding at the PICU).

e. Special emphasis was out on monitoring serum phosphorus levels during the first 7 days of admission (within first 48 hrs of admission and at day 7).

Intervention

As normal serum phosphorus level varies according to age, we used levels in (Table 1) [14] as a guide to start phosphorous supplementation to our patients. Since we didn’t have patients younger than 2 mo, we started to supplement any patient with serum phosphorus level if less than 3 mg/dl.

Statistical Analysis

Nominal data were expressed as frequency and percentage and were compared using Chi square test. Numerical data were expressed as mean, standard deviation and range and compared using T test. Non-parametric data were compared using Mann Whitney test. Pearson’s correlations were used to compare the numerical variables. A p value < 0.05 was considered statistically significant. Logistic regression model was constructed using (hypophosphatemia or normal) as the dependent variable while other data were the independent (predictor) variables. Forward stepwise regression method was used. In this method, the independent variables with the highest significant scores were entered in the analysis. Finally, the model tells us which independent (predictor) variables had the most significant effect on the dependent variable. For each variable in the model at each step, the loss attributed to removing that variable is computed. The more a variable contributes to the model, the larger the change in -2log-likelihood.

Results

Of the 72 patients admitted to PICU during the study period, 10 stayed less than 48 hours, 54 had been discharged and 18 died. The mean serum phosphorus level was (3.5 mg/dl for day 1; 3.7 mg/dl for day 7). The mean serum phosphorus level was significantly lower in those less than 3 yrs old when compared to those above 3 yrs old, p=0.0001. As shown in (Table 2), the prevalence of hypophosphatemia on day 1 was 58% (n=42/72). Seven percent of the patients (n=5) developed hypophosphatemia during their PICU stay, and 12.5% (n=9) of children who were hypophosphatemic at day 1 remained hypophosphatemic at day 7, while 8% (n=6) recovered from hypophosphatemia with treatment.

Regarding the admission diagnoses of patients participating in the study, the majority were suffering of respiratory disorders as pneumonias, bronchiolitis, bronchial asthma, pneumothorax, pleural effusion and stridor. Thirty five percent (n=25) of all children were mechanically ventilated. Sixty eight percent of ventilated children were hypophosphatemic (n=17 vs n=8). Hypophosphatemic patients tended to spend more days being ventilated (7.0 vs 2.75 days) (p = 0.0362) (Table 2). Malnutrition (defined by patient’s weight less than 2SD) was not a significant risk factor when compared to patient’s outcome (discharged/died) (p=0.1753); whereas the number of starvation days (4.69 vs 2.5 days; for hypophosphatemia and normo-phosphatemia at day 7 respectively) affected the phosphorus level during the PICU stay and it was highly statistically significant (p <0.0004).

Although 58% of our children were taking ranitidine and omeprazole during their PICU stay, they did not affect the serum phosphorus level. It was similar for all the drugs taken in consideration that they cause hypophosphatemia. Apart from positive culture for methicillin resistant Staphylococcus aureus (MRSA) (p=0.01), none of the other risk factors for infection correlated significantly with hypophosphatemia including CRP and positive cultures. PIM values were significantly higher in hypophosphatemic patients (6.4) compared to normo-phosphatemics (4.1) (p = 0.0237) (Table 3). Comparing the duration of stay at PICU, those with the normal serum phosphorus level were discharged earlier than those with hypophosphatemia (1.6 vs 2.5 days; respectively) p=0.0001.

Patients with normal serum phosphorus level on admission (day 1) had a better outcome (discharged) compared to those with hypophosphatemia (expired), although the difference was not statistically significant (p = 0.2567) (Table 4, 5).

Table 1: Serum Phosphorus levels During Childhood.

<table>
<thead>
<tr>
<th>Age</th>
<th>Phosphorus (mg/dl)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-5 days</td>
<td>4.8-8.2</td>
</tr>
<tr>
<td>1-3 yrs</td>
<td>3.8-6.5</td>
</tr>
<tr>
<td>4-11 yrs</td>
<td>3.7-5.6</td>
</tr>
<tr>
<td>12-15 yrs</td>
<td>2.9-5.4</td>
</tr>
</tbody>
</table>
Discussion

The prevalence of hypophosphatemia in our prospectively studied CIC was 42%; lower than the 76% reported in a retrospective study conducted in CIC by Menezes et al. [15]. Children younger than 3 years old were more affected due to their lower body reserve compared older ones. In the present study, 56% of patients presenting with respiratory disorders were hypophosphatemic. The adding-on effect of hypophosphatemia to their respiratory problems might be attributed to the fact that hypophosphatemia is known to lead to muscle weakness and hypotonia. Fiaccadori et al. [16] found that 25% of adult patients admitted to the ICU with chronic obstructive pulmonary disease were found to be hypophosphatemic.

Hypophosphatemic patients were more likely to be

### Table 2: Demographic characteristics of the study population.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Total of Patients</th>
<th>Percentage of Patients</th>
<th>Hypo-phosphatemia</th>
<th>Normo-phosphatemia</th>
<th>P Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients Count</td>
<td>72</td>
<td></td>
<td>42</td>
<td>30</td>
<td>0.7885</td>
</tr>
<tr>
<td>Age (mo)</td>
<td></td>
<td>15.08±(34.19)</td>
<td>13.1±(28.38)</td>
<td></td>
<td>0.0653</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.2184</td>
</tr>
<tr>
<td>Female</td>
<td>33</td>
<td>46%</td>
<td>22</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>39</td>
<td>54%</td>
<td>20</td>
<td>19</td>
<td></td>
</tr>
<tr>
<td>Admission Diagnosis</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.1544</td>
</tr>
<tr>
<td>Respiratory Disorders</td>
<td>48</td>
<td>67%</td>
<td>27</td>
<td>21</td>
<td></td>
</tr>
<tr>
<td>Sepsis</td>
<td>6</td>
<td>8%</td>
<td>3</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>Heart Failure</td>
<td>3</td>
<td>4%</td>
<td>1</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Others*</td>
<td>15</td>
<td>21%</td>
<td>11</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>Mechanical Ventilated</td>
<td>25</td>
<td>35%</td>
<td>17</td>
<td>8</td>
<td>0.0362</td>
</tr>
</tbody>
</table>

*Others: Gastroenteritis, Guillain-Barre Syndrome, Encephalitis

### Table 3: Risk factors of hypophosphatemia.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Percentage of Patients</th>
<th>Hypo-phosphatemia</th>
<th>Normo-phosphatemia</th>
<th>P Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Malnutrition</td>
<td>15</td>
<td>21%</td>
<td>7</td>
<td>8</td>
</tr>
<tr>
<td>Drugs taken</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Catecholamines</td>
<td>19</td>
<td>26%</td>
<td>1</td>
<td>18</td>
</tr>
<tr>
<td>Steroids</td>
<td>28</td>
<td>39%</td>
<td>4</td>
<td>24</td>
</tr>
<tr>
<td>Diuretics</td>
<td>8</td>
<td>11%</td>
<td>0</td>
<td>8</td>
</tr>
<tr>
<td>Ranitidine/omeprazole</td>
<td>42</td>
<td>58%</td>
<td>7</td>
<td>35</td>
</tr>
<tr>
<td>Anticonvulsants</td>
<td>14</td>
<td>19%</td>
<td>3</td>
<td>11</td>
</tr>
<tr>
<td>Culture</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Not available</td>
<td>27</td>
<td>38%</td>
<td>13</td>
<td>14</td>
</tr>
<tr>
<td>Negative</td>
<td>16</td>
<td>22%</td>
<td>13</td>
<td>3</td>
</tr>
<tr>
<td>Positive</td>
<td>29</td>
<td>40%</td>
<td>16</td>
<td>13</td>
</tr>
<tr>
<td>CRP</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Not available</td>
<td>7</td>
<td>10%</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Negative</td>
<td>31</td>
<td>43%</td>
<td>19</td>
<td>12</td>
</tr>
<tr>
<td>Positive</td>
<td>34</td>
<td>47%</td>
<td>19</td>
<td>15</td>
</tr>
<tr>
<td>PIM</td>
<td>72</td>
<td>58%</td>
<td>42</td>
<td>30</td>
</tr>
</tbody>
</table>

NB: the not available results were considered a negative result

### Table 4: Predictors of Hypophosphatemia.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Model Log Likelihood</th>
<th>Change in -2 Log Likelihood</th>
<th>df</th>
<th>Sig. of the Change</th>
</tr>
</thead>
<tbody>
<tr>
<td>Step 1 PO4atDay1</td>
<td>-10.974</td>
<td>9.810</td>
<td>1</td>
<td>.002</td>
</tr>
<tr>
<td>Step 2 Ca</td>
<td>-14.209</td>
<td>20.516</td>
<td>1</td>
<td>.000</td>
</tr>
<tr>
<td>Step 2 PO4atDay1</td>
<td>-10.943</td>
<td>13.983</td>
<td>1</td>
<td>.000</td>
</tr>
</tbody>
</table>

### Table 5: Survival to discharge of study population.

<table>
<thead>
<tr>
<th>Number of Patients</th>
<th>Percentage of Patients</th>
<th>Patients with Hypo-phosphatemia</th>
<th>Patients with Normo-phosphatemia</th>
<th>P Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Outcome</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Discharged</td>
<td>54</td>
<td>75%</td>
<td>29</td>
<td>25</td>
</tr>
<tr>
<td>Expired</td>
<td>18</td>
<td>25%</td>
<td>13</td>
<td>5</td>
</tr>
</tbody>
</table>

ventilated and to spend more days on ventilation than normo-
phosphatemic patients. This might be explained by the fact
that hypophosphatemia causes deficiency in the intermediary
compounds for energy production, such as adenosine tri-
phosphate and 2,3-diphosphoglycerate and alterations in energy
metabolism, which may lead to respiratory muscle weakness
and consequent worsening of respiratory insufficiency [5]. The
difficulty of weaning patients from mechanical ventilation is
because of reduced efficiency of respiratory muscular contraction
[17].

In our study, none of the drugs known to deplete serum
phosphorus levels as a side effect to their use (catecholamines,
antacids, anticonvulsants, steroids, diuretics), showed
association with hypophosphatemia. Souza de Menezes et al.
[15] reported same results concerning diuretics, steroids and
sepsis. On the contrary, Santana e Menezes et al. [1] in their study
in PICU in 2009 found that the use of dopamine was associated
with hypophosphatemia and attributed this to increased urinary
phosphorus excretion. Also, more than one study reported the
association between hypophosphatemia and diuretics [5],
steroids [13] and sepsis [18]. The absence of this association in
our study can be due to the heterogeneity of the studied group,
which makes each risk factor of a small sample size. Also, this
can be explained by the dose and the duration of these drugs which
can be not long enough to do this effect.

In contrast to the data reported in the literature, our study
found no significant association with previously described
factors associated with hypophosphatemia, such as malnutrition,
diuretics and sepsis. Only positive cultures for MRSA showed
significant negative correlation with serum phosphorus level. We
hypothesized that patients with MRSA are more severely ill being
hospital acquired and not covered by our first line antibiotics in
the PICU which is amikacin and ampicillin/sulbactum. On the other
hand, Antachopoulos et al. [19] in studying acute infectious
disease in pediatrics, not including CIC, demonstrated significant
negative correlation between serum level of phosphate and CRP.
Barak et al. [20] also demonstrated that infections and sepsis
were correlated with hypophosphatemia.

Although the lack of association observed in our study between
serum phosphorus level and malnutrition, hypophosphatemia
was significantly affected with increased starvation days. We
explain this by the fact that serum phosphorus level is affected
by recent insult (starvation) rather than long standing one
(malnutrition) due to rapid renal excretion of phosphorus.
Santana e Menezes et al. [1] failed to prove this association, and
explained this by having a small sample size to detect statistical
association among the two variables. Our study showed that PIM
values were significantly higher in hypophosphatemic children.
However, Souza de Menezes et al. [15] found no significant
association between PIM and hypophosphatemia.

Our results agreed with the fact that hypophosphatemia
affects the length of PICU stay. This might be explained by the
effect of hypophosphatemia that can trigger myocardial
dysfunction, low ATP for proper respiratory muscles contraction,
leading to an increased need for ventilatory support [17]. In
the context of the hematologic system, the decline in levels
of 2,3-diphosphoglycerate triggered by hypophosphatemia
increases hemoglobin’s affinity for oxygen, thereby causing
tissue hypoxia and leading to changes in erythrocytes and
leukocyte functions, hemolytic anemia, platelet dysfunction, and
thrombocytopenia [21]. Moreover, this study demonstrated that
patients with normal serum phosphorus level on admission had
better outcome (discharged rather than died). This confirmed
the results of the study done by Manary et al. [6] and the
recommendations noted by the review done by Souza de Menezes
et al. [2].

Conclusion

Hypophosphatemia is frequent in children admitted to PICU.
It was more prevalent in those with respiratory failure, those
with more starvation days and higher PIM. More length of stay
and worse outcome were associated with hypophosphatemia.
The association of hypophosphatemia and severity of disease
needs further investigations. Our study highlights the importance
of serum phosphorus level as predictive for the course of illness
and outcome. More studies are needed to delineate the risk
factors of admission hypophosphatemia and development of
hypophosphatemia in the PICU, separately.

References

Hypophosphatemia in critically ill children: prevalence and associated
2. De Menezes FS, Leite HP, Fernandez J, Benzercy SG, de Carvalho WB
regulation of phosphorus homeostasis. Am J Physiol Renal Physiol
Pathophysiologic implications, clinical presentations, and treatment.
Refeeding syndrome with enteral nutrition in children, kwashiorkor is associated with increased mortality. J
cardiac failure after cardiac surgery. Anaesthesia 61(12): 1211-1213.
Refeeding syndrome with enteral nutrition in children: a case report,


