Effect of pre-transplant anaemia on perioperative cardiovascular morbidity and graft function in renal transplant patients

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Abstract

Introduction: Anaemia is a known complication of end stage renal disease (ESRD). The Kidney Disease: Improving Global Outcome (KDIGO) guidelines recommend that haemoglobin (Hb) level in patients with ESRD should be maintained at 11-11.5 g/dl. However, it is difficult to achieve these figures in under resourced countries with limited setup where patients with ESRD frequently run low haemoglobin concentrations.

The aims of this study were to review the Hb levels and prevalence of anaemia in ESRD patients undergoing kidney transplantation in a transplant centre in Sudan and to explore the correlation of pre-operative anaemia with post-operative adverse cardiovascular events, delayed graft function (DGF), and frequency of blood transfusion.

Methods: Records of all patients who underwent kidney transplantation during the period Jan 2012 to Dec 2012, were retrospectively reviewed. The variables analysed were pre-operative Hb, frequency of perioperative blood transfusion, cardiovascular events during the perioperative period and frequency of DGF.

Results: Forty two patients were included in this study. The mean pre-operative Hb level was 10.26 g/dl (SD: 2.05). Twenty six patients (62%) had moderate or severe anaemia, as defined by the World Health Organisation, at the time of transplantation. None of the patients developed cardiovascular events during the perioperative period. There was no statistically significant association between pre-operative anaemia and DGF and frequency of blood transfusion.

Conclusion: Offering renal transplant to anaemic ESRD patients was not associated with adverse cardiovascular events. The frequency of DGF in anaemic patients was equivalent to those with normal Hb.
assess the frequency of pre-transplant anaemia in a cohort of Sudanese renal transplant patients and to assess the association of anaemia with post-operative cardiovascular events, DGF and frequency of allogenic blood transfusion (ABT).

Methods

All patients who underwent living-related renal transplantation at Sharg El-Neel hospital during the period 1/1/2012 till 31/12/2012 and satisfied the set inclusion and exclusion criteria, were included in this study. Inclusion criteria included those patients who were ≥ 18 years of age at the time of transplantation, patients with Hb level ≥ 7 g/dl (the cut-off Hb level for accepting patients for transplantation in our centre) and those undergoing their first kidney transplantation.

Anaemia was defined according to the WHO classification of anaemia [12]. A patient is considered to have moderate anaemia if the Hb level is 8.0-10.9 g/dl or severe anaemia if the Hb level is < 8.0 g/dl. However, due to the small sample size, patients were categorized as either anaemic if the Hb level was ≤ 10.9 g/dl or as normal Hb if the level was ≥ 11.0 g/dl.

Due to the lack of a universal definition, DGF was defined in this study as the need for dialysis during the first seven days post-transplantation [13,14].

Perioperative cardiac event was defined as the occurrence of symptoms compatible with acute myocardial ischemia, arrhythmia or congestive heart failure within 30 days post-transplantation. Symptoms compatible with acute myocardial ischemia included myocardial infarction (ST-segment elevated or non-ST-segment elevated) or unstable angina. Diagnosis of myocardial infarction (MI) was made based on the clinical picture, recent ECG changes and elevation of the serum biomarkers, specifically Troponin T, analysed at different time intervals. Diagnosis of unstable angina was made based on clinical picture and recent ECG changes [15].

In our center, as part of the assessment of the potential renal transplant recipients, Hb level was checked during the initial assessment and at admission for the operation. Generally, the cut-off Hb level for accepting patients for transplantation in our center was 7 g/dl; however, some exceptions were made on case by case basis. Pre-transplant assessment included a detailed cardiac history and clinical examination to detect any significant cardiac conditions including Acute Coronary Syndrome (ACS), heart failure, arrhythmias and valvular disease. Assessment for the functional class was also made (ability to climb two flights of stairs) [16]. Baseline ECG and echocardiogram were done for all cases. Any patient who showed features of cardiac compromise from history, clinical examination or investigations was referred to the cardiology service for further assessment and optimization.

Our immunosuppression protocol consisted of triple therapy. Tacrolimus was started 2 days pre-operatively at a dose of 0.15 mg/kg body weight/day given in 2 divided doses and continued post-operatively, with the dose adjusted to keep the trough level at 10-15 ng/ml. Methylprednisolone was given intra-operatively at a dose of 500 mg before release of clamps and continued for 2 days post-operatively at a dose of 250 mg/day. Oral prednisolone was started at the third postoperative day at a dose of 20 mg/day. Azathioprine was started at the first post-operative day at a dose of 5mg/kg/day. Tapering of immunosuppression was started six weeks post-operatively. Induction therapy was not used during the study due to the financial limitations, although it was introduced later.

Post-operatively, kidney function was monitored by measuring hourly urine output, daily measurement of serum urea, creatinine and electrolytes level and calculating creatinine clearance using Cock-Croft Gault formula. Laboratory assessment of full blood count was carried out on daily basis till the time of discharge, usually within 5 to 7 days post-operatively. Decision to give blood transfusion was made jointly between the surgeon and anaesthetist intra-operatively, if it was felt that there was excessive bleeding. Post-operatively, decision to give blood transfusion was made jointly between the surgeon and nephrologist if it was felt that there was excessive output from the surgical drains or if the Hb level dropped to below 7g/dl.

According to our protocol, ECG and cardiac enzymes were requested if the patient complained of any symptoms suggestive of a cardiac event. These were repeated after 12 hours or as deemed necessary.

The variables included in this study were age at time of transplantation, sex, history of blood transfusion before transplantation, Hb level immediately before transplantation, need for transfusion during the perioperative period and number of units transfused, occurrence of cardiovascular events and episodes of DGF.

Ethical approval was granted, from the hospital’s medical directorate, before the commencement of the study.

Statistical Analysis

Statistical analysis was performed using JMP software version 12. Continuous and categorical variables were displayed as means ± standard deviation (SD) and frequencies and percentages, respectively. Student’s t-test was used to assess the differences between continuous variables. Differences between categorical variables were analyzed using Fisher’s exact test. P values < 0.05 were considered to be statistically significant.

Results

Forty three patients, with ESRD and on haemodialysis, underwent living related kidney transplantation during the study period. One patient was excluded because he was undergoing his second kidney transplantation. The remaining 42 patients satisfied the inclusion and exclusion criteria and were suitable for analysis. None of these patients was undergoing a pre-emptive transplant. Table 1 displays the characteristics of the study group and the distribution of variables included in the study. The patients’ age ranged from 19 to 55 year with a mean (SD) of 32.8 (9.5) years. Twenty two of the sample (66.7%) were males. Twenty six patients (62%) were anaemic at the time of transplantation.

Table 1. Renal transplant patient characteristics. DGF: Delayed graft function

<table>
<thead>
<tr>
<th>Subject characteristics</th>
<th>N (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>Mean (SD) 32.8 (9.5)</td>
</tr>
<tr>
<td>Sex</td>
<td>male 28 (66.7) female 14 (33.3)</td>
</tr>
<tr>
<td>Pre-operative Hb</td>
<td>≤10.9 g/dl 26 (61.9) ≥11.0 g/dl 16 (38.1)</td>
</tr>
<tr>
<td>Pre-transplant blood transfusion</td>
<td>Yes 12 (28.6) No 30 (71.4)</td>
</tr>
<tr>
<td>DGF*</td>
<td>Yes 5 (11.9) No 37 (88.1)</td>
</tr>
<tr>
<td>Acute rejection</td>
<td>Yes 9 (21.4) No 33 (78.6)</td>
</tr>
</tbody>
</table>
The mean Hb level for the whole group was 10.2 g/dl (range: 6.1-14). Six patients had a Hb level less than 8 g/dl (severe anaemia according to the WHO classification). Two patient were operated on with a Hb level below the set threshold of 7.0 g/dl as both patient were running out of access sites and so were offered transplantation in spite of the very low Hb. It was noticeable that males had a higher pre-operative Hb level compared to females, although that did not reach statistical significance ($p=0.076$). Twelve patients (28.6%) gave a past history of transfusion at some stage before coming for transplantation. The mean (SD) Hb on discharge for the studied group was 9.0 g/dl (2.08). The mean (SD) postoperative creatinine level was 1.40 mg/dl (0.66).

There was neither perioperative mortality nor graft loss in the studied group. None of the patients included in the study developed any acute cardiovascular event during the perioperative period.

Of the 42 patients studied, five patients (12%) developed DGF. Two of these patients were haemodialysed in the immediate post-operative period due to high serum potassium following intra-operative blood transfusion. Of these five patients with DGF, 4 patients were anaemic pre-operatively and one patient had normal Hb level. Table 2 summarizes the association of the examined parameters with pre-transplant anaemia. There was no statistically significant association between pre-operative anaemia and DGF ($p=0.38$). Also, there was no association between pre-operative Hb level and DGF ($p=0.9$).

Eleven patients (25.6%) received blood transfusion during the perioperative period. The mean number of units (SD) transfused was 2.72 units (1.48). One patient needed ten units of blood due to excessive intra-operative bleeding. Eight of the patients who needed blood transfusion were anaemic pre-operatively. There was no significant association between pre-operative anaemia and the need for transfusion ($p=0.48$). Three patients who received blood transfusion developed acute rejection. There was no statistically significant association between blood transfusion and episodes of acute rejection ($p=0.1$).

**Discussion**

Anaemia is a common complication of chronic kidney disease, the pathophysiology of which is related to reduced erythropoietin production, iron and vitamin deficiency, reduced erythrocytes life span, chronic inflammation and ureamic milieu [17-19]. Anaemia in CKD significantly increases the risks of cardiovascular morbidity and mortality as well as risk of stroke [20-22]. The prevalence of anaemia in Sudanese renal transplant population and its association with adverse cardiac events and DGF. To the best of our knowledge, there are no similar reports from sub-saharan Africa. There are, however, some limitations to this study. The main limitation is the small sample size which may have resulted in type II error. There are plans to reproduce this study with a larger sample size to be recruited from our center as well as other local centers. The other drawback is that most of the patients in this study group belonged to young age groups. It must be stressed that in our practice, patients are not denied transplantation because of old age (there was no upper age limit included in our protocol). Another limitation is that this was a retrospective study and thus causality cannot be determined. Future studies using prospective cohort design may provide greater control for bias and confounding factors.

In conclusion, offering renal transplantation to moderately anaemic patients was neither associated with adverse cardiac events nor with DGF. Patients should be carefully selected to ensure a favorable outcome. Further investigation with larger samples is required to determine the safety of this practice.

**Table 2. Association of the parameters tested with pre-transplant anaemia.** DGF: Delayed graft function

<table>
<thead>
<tr>
<th>Parameters tested</th>
<th>Association with pre-transplant anaemia</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex</td>
<td>Mean Hb (SD)</td>
<td>0.076</td>
</tr>
<tr>
<td>Male</td>
<td>10.66 mg/dl (1.95)</td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>9.47 mg/dl (2.1)</td>
<td></td>
</tr>
<tr>
<td>DGF (n=5)</td>
<td></td>
<td>0.38</td>
</tr>
<tr>
<td>Anaemic</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>Normal</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Patients requiring transfusion (n=11)</td>
<td>4</td>
<td>0.48</td>
</tr>
<tr>
<td>Anaemic</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>normal Hb</td>
<td>3</td>
<td></td>
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</table>

**References**


