Correlation between Inflammatory Biomarkers and Red Blood Cell Life Span in Chronic Hemodialysis Patients

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Key Words
Red blood cell life span · Inflammation · Interleukins · Uric acid · Hemodialysis · Anemia

Abstract
Background and Objectives: The pathogenesis of anemia in hemodialysis (HD) patients is dependent on multiple factors, with decreased red blood cell life span (RBCLS) being a significant contributor. Although the impact of reduced RBCLS on anemia is recognized, it is still a subject that is not well researched. The objective of this study was to investigate the relationship between RBCLS and inflammatory biomarkers in chronic HD patients.

Design, Setting, Participants, and Measurements: RBCLS was calculated from alveolar carbon monoxide concentrations measured by gas chromatography. Interleukins (IL) IL-6, IL-18, IL-10, and high sensitivity C-reactive protein were measured using bead-based multiplex assay. Measurements were carried out at baseline and during follow-up. The associations between RBCLS and inflammatory biomarkers were evaluated using linear mixed effects models.

Results: RBCLS measurements were available for 54 HD patients. Their average age was 58.5 ± 14.4 years, 68.5% were males, 48.1% were diabetics, and the HD vintage was 51 ± 48 months. In 4 patients, RBCLS was measured once, while in 50 patients, up to 5 repeated RBCLS measurements were available. RBCLS was 73.2 ± 17.8 days (range 37.7–115.8 days). No association was found between RBCLS and any of the inflammatory biomarkers. Of note, RBCLS was positively correlated with levels of uric acid (p = 0.02) and blood urea nitrogen (BUN; p = 0.01), respectively.

Conclusion: Our study suggests that inflammation pathways reported by these biomarkers only have a limited role in causing premature RBC death. The positive correlation with uric acid and BUN warrants further studies.

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J.M. and Y.D. contributed equally to this work.
Introduction

Anemia is a typical complication of chronic kidney disease. The pathogenesis of anemia in hemodialysis (HD) patients is dependent on multiple factors, including insufficient production of endogenous erythropoietin (EPO), functional iron deficiency, acute and chronic inflammation, hyperparathyroidism, aluminum toxicity, deficiency of nutrients such as folate and others, inhibition of erythropoiesis [1], increased apoptosis of erythroblasts [2], and shortened red blood cell life span (RBCLS). Previous research has indicated that RBCLS in chronic HD patients was reduced to approximately 60–90 days [3]. While exogenous erythropoiesis stimulating agents (ESA) are used successfully in the treatment of anemia in a majority of HD patients, some patients require very high ESA doses, a clinical phenotype called ESA resistance. While ESA resistance is frequently attributed to inflammation, functional iron deficiency and circulating EPO inhibitors, a shortened RBCLS has hardly been recognized as a contributing factor [3, 4]. Inflammation and oxidative stress frequently co-exist in the maintenance HD patients [5] and their interconnection induces anemia complicated with EPO hyporesponsiveness [6]. HD patients present frequently with high levels of inflammatory markers such as high sensitivity C-reactive protein (hs-CRP), interleukin (IL)-6, tumor necrosis factor-α, neutrophil-to-lymphocyte ratio (NLR) [7], and low levels of serum albumin [8]. Increased levels of pro-inflammatory cytokines reduce the production of erythroid progenitor cells and this contributes to hyporesponsiveness to ESAs and poor treatment outcomes [9, 10]. Elevated levels of inflammatory cytokines also contribute to functional iron deficiency. While the relationship between inflammation and renal anemia is well explored [11, 12], the impact of reduced RBCLS although recognized is not yet well researched.

The objective of this study was to further investigate the relationship between RBCLS and inflammatory biomarkers in chronic HD patients.

Materials and Methods

Study Participants and Study Design

We enrolled non-smoking HD patients from an urban HD center, and measured their RBCLS up to 5 times during a maximum follow-up of 18 months in a prospective observational study. ESA dosages were adjusted by the attending nephrologist to maintain hemoglobin (Hgb) levels within the range 10.0–11.9 g/dl. We excluded patients with hemorrhagic disorders, chronic lung disease, infection or cancer.

This study was conducted in accordance with the ethical principles set forth in the Helsinki Declaration, and approval was obtained from the New England institutional review board. Written informed consent was obtained from all individuals prior to their participation in this study.

Data Collection

Clinical data, including age, gender, HD vintage, comorbidities, vascular access type, dialysis adequacy (Kt/V), routine laboratory data, and weekly EPO dose were extracted from the study database.

Measurements

Whole blood samples were drawn before the first HD session of the week (Monday or Tuesday), centrifuged when appropriate and analyzed at Spectra Laboratories, Rockleigh NJ, to assess a complete blood count, hs-CRP, blood urea nitrogen (BUN), albumin, iron status (iron, ferritin, transferrin saturation), uric acid, carnitine, as well as total and direct bilirubin. In addition, EDTA blood samples were centrifuged immediately after collection, and plasma was aliquoted and frozen at −70°C for further batch analysis of the pro-inflammatory cytokines IL-6 and IL-18 and the anti-inflammatory cytokine IL-10. Plasma levels of IL-6 and IL-10 were measured in duplicates using the MILLIPLEX MAP High Sensitivity Human Cytokine Panel kit (Millipore Corp., Billerica, MA, USA) with Luminex® 100TM System (Luminex Corporation, Austin, Tex., USA). Quantitative analysis of plasma IL-18 was made in duplicates by high-sensitivity ELISA (MLB, Nagoya, Japan) that uses 2 monoclonal antibodies against 2 different epitopes of human IL-18 according to the instructions in the supplier manual. Body weight was measured before and after each HD session. The doses of EPO (epoetin alpha) are expressed as IU/kg body weight per week. We used the EPO resistance index as the metric of EPO resistance, defined as the weekly weight-adjusted EPO dose (U/kg/week) divided by Hgb level (g/dl) [13, 14].

Measurement of Alveolar Carbon Monoxide Concentrations and Calculation of RBCLS

Breakdown of heme results in the production of carbon monoxide (CO), and the alveolar CO concentration can be used to estimate RBCLS [15]. Alveolar breath samples were obtained twice using the GaSampler™ device (Quintron Instruments Milwaukee, WI, USA) before the first blood sample was drawn. A sample of ambient air from the patient’s home was obtained with an open plastic bottle that was kept overnight in the patient’s home to allow for equilibration of gases. CO concentration was measured by gas chromatography (Model 910, Buck Scientific, East Norwalk, CT, USA). Patient alveolar air, ambient air from the patient’s home, and standard gas (20 parts per million) used for calibration were measured and the area under the curves (AUC) representing the CO concentration were reported. All gas samples were measured in triplicates, and the respective average AUC of each air sample were used to calculate the CO concentrations. The endogenous CO is the difference between the average alveolar and home ambient air CO concentrations. The endogenous CO concentration and RBCLS were computed as previously reported [15–17].

Statistical Analyses

Continuous variables are presented as mean ± SD if normally distributed and as median (25th, 75th percentile) otherwise. Categorical variables are presented as percentages of the respective group.

Linear mixed effects (LME) models were used to explore the associations between RBCLS and independent variables. We used
RBCLS measurements were available in 54 HD patients. Their average age was 58.5 ± 14.4 years, 68.5% were males, 48.1% were diabetics, and the HD vintage was 51 ± 48 months (table 1). In 4 patients, RBCLS was measured once, while in 50 patients up to 5 repeated RBCLS measurements were available. Among these 50 patients, 7 patients had 2 measurements, 9 patients had 3 measurements, 12 patients had 4 measurements, and 22 had 5 measurements.

The mean RBCLS was 73.2 ± 17.8 days (range 37.7–115.8 days; fig. 1). We did not observe any significant correlation between RBCLS and white blood cells (WBC), NLR, hs-CRP, IL-6, IL-18, and IL-10, respectively (fig. 2a–f).

Of note, RBCLS was positively correlated with uric acid (p = 0.02) and BUN (p = 0.01), respectively (fig. 3a, b).

**Discussion**

The key finding of this study is that there is lack of correlation between RBCLS and inflammatory indicators. To the best of our knowledge, this is the first study in chronic HD patients to test the hypothesis of a relationship between RBCLS and these inflammatory biomarkers.
Fig. 2. Relationship between inflammatory biomarkers and RBC life span. Each data point is an individual measurement; there are multiple measurements per patient. The solid lines represent the fixed effect estimated by LME models. None of these associations was statistically significant. a WBC (beta = -0.16, p = 0.89); b NLR (beta = 1.18, p = 0.35); c hs-CRP (beta = -0.27, p = 0.23); d IL-6 (beta = -0.09, p = 0.51); e IL-10 (beta = -0.21, p = 0.31); f IL-18 (beta = -0.05, p = 0.13).

Fig. 3. Positive correlations between RBCLS and (a) uric acid (beta = 2.60, p = 0.02) and (b) BUN (beta = 0.27, p = 0.01) levels. Each data point is an individual measurement there are multiple measurements per patient. The solid lines represent the fixed effect estimated by LME models.
The relationship between inflammation and anemia in dialysis patients is well described and it has been shown that inflammation exerts both direct and indirect effects on erythropoiesis [18–20]. However, little is known about an association between inflammatory cytokines and RBCLS. In the present research, we studied about cytokines with known involvement in the genesis of anemia. IL-6 impairs late stages of erythroid development, after cells have been primed for Hgb synthesis [21]. In vitro data show a direct negative effect of IL-10 on erythroid progenitor cells proliferation and indicate that IL-10 could inhibit the growth of these cells [22]. A recent study shows that IL-10 induces the expression of heme oxygenase-1 [23], which may enhance heme degradation, resulting in an increase in the amount of iron within monocytes, and subsequent incorporation within the reticuloendothelial system and the development of anemia. IL-18, a pro-inflammatory cytokine with diverse pleiotropic effects, induces interferon gamma production by natural killer cells, T cells, and activated macrophages. IL-18 also regulates both T helper 1 (Th1) and Th2 responses, thus playing an important role in the inflammatory process [24].

While not being part of the primary data analysis, we found that uric acid and BUN were correlated with RBCLS. It is tempting to speculate that the positive correlation between uric acid levels and RBCLS may be related to its antioxidant actions. Our research also corroborates a recently reported positive relationship between BUN and RBCLS [3], and currently, the underlying biology of that finding remains elusive. The correlation of RBCLS with uric acid and BUN warrants further studies.

We used the CO method developed by Strocchi et al. [15] to measure RBCLS. The main advantages of this method are its noninvasive nature, the avoidance of radioactive label, and its practical simplicity. However, the underlying principle of that method deemed it necessary to exclude smokers from this research, and this certainly was a limitation of this study.

Whenever the results of a study comprise negative findings, considerations of statistical power and sample size are important. While being one of the largest studies with repeated measurements of RBCLS in chronic HD patients, its statistical power is still low. With estimated parameters as the true parameters, our study has only 20% power at the significance level of 5% for NLR as a marker. To have 80% of power, one would need a likely prohibitively large number of 500 patients. The nonsignificant associations may be a result of small sample size and large variation between patients.

In conclusion, our study did not find an association between RBCLS and inflammatory biomarkers in chronic HD patients. This suggests that inflammation only has a limited role in the genesis of premature RBC death.

**Disclosure Statement**

P.K. holds stock in Fresenius Medical Care. M.M.Y.W. is a research consultant with Arbor Research Collaborative for Health.

**References**


