



Trabajo Original

Metabolic, inflammatory and oxidative stress markers in the nitric oxide variation of hemodialysis subjects

Marcadores de estrés metabólico, inflamatorio y oxidativo en la variación del óxido nítrico de los individuos de hemodiálisis

Andreza P. Santos Epifânio¹, Karka P. Balbino¹, Mônica de P. Jorge¹, Sônia. M. R. Ribeiro¹, Ana Vládia B. Moreira¹, Jerusa M. Oliveira², Leandro L. Oliveira² and Helen Hermana M. Hermsdorff¹

¹Department of Nutrition and Health. Universidade Federal de Viçosa. Viçosa, Minas Gerais. Brazil. ²Department of General Biology. Universidade Federal de Viçosa. Viçosa, Minas Gerais. Brazil

Abstract

Introduction: Oxidative stress markers such as nitric oxide (NO) have been investigated in hemodialysis (HD).

Objective: Evaluate the association of NO variation with adiposity indicators, metabolic, inflammatory and oxidative stress markers in individuals to HD.

Methods: Cross-sectional study with 85 subjects on HD treatment (≥ 18 years). The clinical-nutritional status was evaluated through subjective global assessment modified (SGAm), anthropometric measurements and body composition. Dietary intake was evaluated using a food frequency questionnaire. Metabolic markers were obtained from medical records. Inflammatory markers (IL-6 and IL-10) and oxidative stress, (TACs), (SOD), (GST), (MDA) and NO were determined using standardized protocols.

Results: Those individuals with a high concentration of NO ($> 4.32 \mu\text{mol/L}$) had lower values for SGAm score ($p = 0.012$) and higher iron values ($p = 0.050$), Fe saturation ($p = 0.037$) and triacylglycerol ($p = 0.003$). The same subjects still had lower consumption of copper ($p = 0.026$), manganese ($p = 0.035$), vitamin E ($p = 0.050$), $\omega 3$ ($p = 0.021$) and $\omega 6$ ($p = 0.020$). In a multiple regression model, concentrations of ferritin, triacylglycerol, IL6 and SOD contributed to a 54.8% increase in NO concentrations, whereas triacylglycerol and SOD concentrations were independent factors for NO variation ($p < 0.001$).

Conclusions: The clinical and nutritional status as well as intake of nutrients with antioxidant properties (Cu, Zn, Mn, vitamin C and $\omega 3$) appears to modulate the variation of NO in this population.

Key words:

Reactive nitrogen species. Inflammation. Superoxide dismutase. Food intake. End-stage renal disease.

Resumen

Introducción: se han investigado marcadores de estrés oxidativo como el óxido nítrico (NO) en hemodiálisis (HD).

Objetivo: evaluar la asociación de la variación del NO con los indicadores de adiposidad, los marcadores metabólicos, inflamatorios y de estrés oxidativo en individuos a HD.

Métodos: estudio transversal con 85 sujetos en tratamiento HD (≥ 18 años). El estado clínico-nutricional se evaluó a través de la evaluación global subjetiva modificada (SGAm), medidas antropométricas y composición corporal. La ingesta dietética se evaluó mediante un cuestionario de frecuencia alimentaria. Marcadores metabólicos se obtuvieron de los registros médicos. Se determinaron marcadores inflamatorios (IL-6 e IL-10) y estrés oxidativo (TAC), (SOD), (GST), (MDA) y NO mediante protocolos estandarizados.

Resultados: los individuos con una alta concentración de NO ($> 4,32 \mu\text{mol/L}$) tuvieron valores más bajos de puntuación de SGAm ($p = 0,012$) y mayores valores de hierro ($p = 0,050$), saturación de Fe ($p = 0,037$) y triacilglicerol ($p = 0,003$). Los mismos sujetos tuvieron un menor consumo de cobre ($p = 0,026$), manganeso ($p = 0,035$), vitamina E ($p = 0,050$), $\omega 3$ ($p = 0,021$) y $\omega 6$ ($p = 0,020$). En un modelo de regresión múltiple, las concentraciones de ferritina, triacilglicerol, IL6 y SOD contribuyeron a un aumento de 54,8% en las concentraciones de NO, mientras que las concentraciones de triacilglicerol y SOD fueron factores independientes para la variación del NO ($p < 0,001$).

Conclusiones: el estado clínico y nutricional así como la ingesta de nutrientes con propiedades antioxidantes (Cu, Zn, Mn, vitamina C y $\omega 3$) parecen modular la variación del NO en esta población.

Palabras clave:

Especies nitrogenadas reactivas. Inflamación. Superóxido dismutasa. Ingesta de alimentos. Enfermedad renal terminal.

Author's contribution: A. P. Santos Epifânio: Design study, field work, data collection, analysis, and writing of the manuscript. K. P. Balbino and M. P. Jorge: Field work, and data collection. S. M. R. Ribeiro; A. V. B. Moreira and L. L. Oliveira: design, and data interpretation. J. M. Oliveira: Data analysis. H. H. M. Hermsdorff: project leader in the Universidade Federal de Viçosa, general coordination, design, and data interpretation. All authors critically reviewed the manuscript and approved the final version submitted for publication.

Received: 01/06/2017 • Accepted: 28/08/2017

Santos Epifânio AP, Balbino KP, Jorge MP, Ribeiro SMR, Moreira AVB, Oliveira JM, Oliveira LL, Hermsdorff HHM. Metabolic, inflammatory and oxidative stress markers in the nitric oxide variation of hemodialysis subjects. Nutr Hosp 2018;35:176-184

DOI: <http://dx.doi.org/10.20960/nh.1319>

Correspondence:

Helen Hermana M. Hermsdorff. Department of Nutrition and Health. Universidade Federal de Viçosa. Avenue PH Rolfs, s/n. Viçosa, Minas Gerais, 36570-900 Brazil
e-mail: helenhermana@ufv.br

INTRODUCTION

Chronic kidney disease (CKD) is a worldwide health problem due to its high incidence (1). High morbidity and mortality in CKD is associated with its own progression to end-stage renal disease (ESRD) and the development of other metabolic disorders that increase the risk for cardiovascular diseases (2).

In turn, hemodialysis (HD), an essential treatment in ESRD, contributes to the increase of oxidative stress due to a diminished antioxidant system, and subsequently to the manifestation of inflammation and endothelial dysfunction, all risk factors for atherosclerosis in this population (3). Thus, oxidative stress markers have gained interest as non-traditional cardiometabolic risk factors in ESRD (4).

Among the markers of oxidative stress, nitric oxide (NO), a recognized vasodilator and cardioprotector, is prominent (5). Many cells are able to synthesize NO by the activity of the enzyme nitric oxide synthase (NOS), which converts the amino acid L-arginine to NO and L-citrulline (6). NO has important functions in renal physiology such as maintenance of homeostasis in blood flow, renal excretion and renin secretion; and tubule glomerular return (7). However, NO may be toxic under conditions of oxidative stress, from the generation of reactive oxygen species (ROS) and deficiency of the antioxidant system (8).

In this context, NO concentrations could be associated with other markers of oxidative stress and inflammation in the ESRD, although there were little studies regarding this topic (9).

Overall, the present cross-sectional study aimed to evaluate the potential association of NO variation with adiposity indicators, as well as metabolic, inflammatory and oxidative stress markers in individuals submitted to HD.

MATERIAL AND METHODS

STUDIED POPULATION

This is a cross-sectional study with 85 subjects on HD treatment (≥ 18 years old), the majority of participants were men (65.9%; $n = 56$) and the average age was 62 ± 13.7 years old attended in a single dialysis center. Patients underwent three weekly sessions of HD with an average duration of 4 hours, blood flow greater than 250 mL/min and dialysate flow of 500 mL/min. Individuals who did not show interest in participating in the study, with a treatment time of less than one month in HD and those with hearing impairment, newly implanted catheters, hemodynamic instability, evaluated by the doctor of the sector, and those unable to stand for anthropometric evaluation were not included in the study.

CLINICAL-NUTRITIONAL STATUS ASSESSMENT

The clinical-nutritional status was assessed using nutritional risk score subjective global assessment modified (SGAm), and anthropometric and body composition measurements. The SGAm

used was based on the model proposed by Kalantar-Zadeh et al. (10) for renal patients on dialysis.

Anthropometry and tetrapolar electrical bioimpedance (BIA) were performed approximately 30 minutes after the end of HD. The anthropometric measures included dried weight (kg), height (cm), and waist circumference (WC), which were performed according to previously standardized procedures (11-13).

Body mass index (BMI) was calculated and individuals were classified according to the cut-off points of the World Health Organization (14) for adults and Lipschitz (15) for the elderly. The individuals were classified according to the cardiometabolic risk, according to the CP values, using the WHO cut-off points, 1995 (14).

DIETARY INTAKE ASSESSMENT

A semi-quantitative food frequency questionnaire was constructed, based on an Australian questionnaire validated for renal patients (16,17). Thus, the food portions of each group were analyzed according to the Food Guide for the Brazilian Population (2006) (18): cereals, tubers and roots, fruits, vegetables and legumes, and other vegetable foods rich in proteins, milk and dairy products, meat and eggs, fats, sugars and salt, water. The oilseeds group was also inserted, according to the Food Guide for Brazilian Population (2014) (19). The reference portion was based on the Family Budget Survey (POF) 2008 table (19). The fruit and vegetable group was divided into high, medium and low potassium.

Nutrient intake was calculated as frequency \times nutrient composition of each portion size for each consumed food item, in spreadsheet in Microsoft Excel 2010, according to the nutritional composition of foods of Brazilian tables (20). Therefore, we evaluated the daily caloric intake (kcal), carbohydrates, protein, lipids and fatty acid profile (in percentage of caloric intake), fiber (g), calcium (g), phosphorus (mg), potassium (mg) and sodium (mg), magnesium (mg), manganese (mg), iron (mg), selenium (μ g), thiamine (mg), niacin (mg), cyanocobalamin (μ g), vitamin E (IU), vitamin C (mg) and folate (mg).

METABOLIC MARKERS

The metabolic markers analyzed in the present study were those obtained from medical records: albumin, urea, urea removal rate (URR), creatinine, potassium, phosphorus, calcium, calcium-phosphorus (Ca-P) product, parathyroid hormone (PTH), hemoglobin (Hb), hematocrit (Ht), ferritin, iron (Fe), transferrin saturation (SatFe), C-reactive protein (CRP), triacylglycerol and total cholesterol. The Kt/V was calculated using the equation proposed by Daugirdas II (21). Values of urea Kt/V > 1.2 were considered indicative of efficiency in HD.

INFLAMMATORY AND OXIDATIVE STRESS MARKERS

Blood samples were collected before the beginning of HD. Blood was collected in (Vacutainer®) tubes containing EDTA as anticoag-

ulant. Serum was separated in a refrigerated centrifuge (15 min, 3000 rpm, 4 °C) and both stored at -80 °C for posterior analysis. Serum IL-2, IL-4, IL-6, IL-10 concentrations were measured by flow cytometry technique with the BD FACVerse Cytometer, using the Human Th1/Th2/Th17 CBA Kit (BD Biosciences, USA) at the Laboratory of General Biology, Department of General Biology, Universidade Federal de Viçosa. The results were obtained using the software FacSuite (BD®). Serum total antioxidant capacity (TAC), enzymes activity superoxide dismutase (SOD), and glutathione S-transferase (GST), the lipid peroxidation product malondialdehyde (MDA) and NO were considered as biomarkers of oxidative stress. TAC was measured by colorimetric assay using the Antioxidant Assay Kit (CS0790, Sigma Aldrich), according to the protocol provided by the manufacturer, and other markers were assessed standardized protocols of the Laboratory of Echophysiology of Chiroptera - Department of Animal Biology, Universidade Federal de Viçosa, as follows.

Activity of antioxidant enzymes

SOD activity was measured in serum in a microplate reader ($\lambda = 570$ nm) (22), based on the ability of this enzyme to catalyze the reaction of superoxide radical (O_2^-) thus decrease the auto-oxidation ratio of pyrogallol. The results were expressed as U SOD/ mg protein.

GST was measured through the formation of GSH conjugate, 2,4-dinitrobenzene (CDNB), and estimated by the change in absorbance at 340 nm for 60s. The molar extinction coefficient of CDNB at 340 nm is $9.6 \text{ mM}^{-1}\text{cm}^{-1}$, which was used for the calculations (23). GST activity was expressed as $\mu\text{mol}/\text{min}/\text{g}$.

Malondialdehyde

The concentration of MDA was estimated as described by Wallin et al. (24). 200 μl aliquots of each serum sample were separated and added to a 400 μl of heated, vortex homogenized solution TBARS whit trichloroacetic acid (15%) / thiobarbituric acid (0.375%) / hydrochloric acid (0.25 M) for 40 minutes in boiling water (90 °C) and then cooled in an ice bath for 5 minutes. 600 μl of butyl alcohol were added and again homogenized in vortex for ~ 2 minutes. The solutions were centrifuged at 3,000 rpm at room temperature (10 minutes at 900g). 200 μl of the supernatant were separated for quantifying the MDA concentration in microplate reader ($\lambda = 535$ nm). The concentration of MDA was determined by standard curve from known concentrations of 1,1,3,3-tetramethoxypropane (TMPO). The results were expressed as $\mu\text{M}/\text{mg}$ protein.

Nitric oxide

The serum for the NO tests was prepared as described above. The production of nitric oxide was indirectly quantified through nitrite content in the serum sample by the Griess reaction(25),

composed of 1% sulfanilamide and 0.1% naphthyl-ethylene-diamine in 2.5% in 2,5% H_3PO_4 . Thus, 50 μl of the supernatant from the samples were added to microplates with equal volume of the Griess Reactant and incubated at room temperature for 15 minutes, then determined on a microplate reader ($\lambda = 570$ nm). The nitrite concentration of the samples was determined using standard curve with known concentrations of sodium nitrite (NaNO_2) and expressed in Mm/mg protein.

The protein concentration used in the calculations of the activity of antioxidant enzymes, MDA, and NO was measured by the method of Lowry et al. (1951), using bovine serum albumin as previously standardized (26).

STATISTICAL METHOD

Normal distribution of the data was determined using the Kolmogorov-Smirnov test. Data were expressed as mean \pm standard deviation, median (interquartile range). The study population was divided by the median NO concentrations (4.32 $\mu\text{mol}/\text{L}$) in low and high NO concentration. The median cutoff criteria have been previously applied based (27) on a valid and reliable method to assign two groups of risk in epidemiological studies (28).

Comparisons between groups were performed using Student's t-test for parametric variables, or Mann-Whitney, for non-parametric variables. The correlation analysis between variables of interest was performed using Pearson or Spearman correlation coefficient, as appropriate. Multiple regression analysis was used to determine indicators of the variation of NO concentration of the sample studied. For the construction of multiple linear models, the value of $p \leq 0.20$ obtained in the bivariate analysis was used as criterion for inclusion of the variables. In the final model, the backward method was used, for which the variables with less significance (greater p value) were removed one by one from the model.

Statistical analysis was performed using the SPSS 20.0 program (SPSS, Inc., Chicago, IL, USA) and a significance level of less than 5% was applied.

RESULTS

STUDY SAMPLE

The majority of the participants were men (65.9%; $n = 56$) and elderly (61.2%, $n = 52$). The main causes of CKD in the study population were hypertensive nephrosclerosis (41.2%; $n = 35$) and diabetes mellitus (32.9%; $n = 28$). The HD time ranged from 1 to 245 months, with a median of 41.5 months, presenting a statistical tendency when associated with NO ($p = 0.062$). In addition, the sample presented mean of Kt/V (1.52 ± 0.39) and serum albumin (4.08 ± 0.24 g/dL) as expected to HD efficiency and nutritional adequacy. Regarding dietary intake, these individuals have an energy balance for macronutrients, with a high consumption of food sources of potassium 2,548.50 mg/d (935.4-8, 276, 0).

NO AND CLINICAL-NUTRITIONAL STATUS

In relation to weight status evaluated by BMI, 9.1% (n = 3) of adults were classified as underweight, 72.7% (n = 24) as normal weight, 15.2% (n = 5) pre-obese and 3% (n = 1) obesity class I. Among the elderly, 34.6% (n = 18) were classified as underweight, 44.2% (n = 23) normal weight and 21.2% (n = 11) were overweight. There was no statistical difference in relation to NO concentration (p = 0,395), according to weight status.

According to mSGA, nutritional status was adequate in 10.6% (n = 9) of the individuals, while 89.4% (n = 76) were at nutritional risk / mild malnutrition. Interestingly, the mSGA score was statistically lower in subjects with high NO (p = 0.012). By the WC, central adiposity indicator according to WHO (1997) (18), 22.4% of the patients had a high risk and a very high risk

of obesity-related metabolic complications, with a very high risk being greater among women. By total body fat, 20.0% had fat shortage and 23.5%, excess fat. There was no statistical difference with these adiposity indicators in relation to NO concentration (Table I).

NO AND DIETARY INTAKE

Daily intake of total calories, alpha-linolenic fatty acid (ω 3) and linoleic acid (ω 6) were different according to median of NO concentration (Table II). In relation to micronutrients, a higher consumption of manganese, copper, zinc, selenium, vitamin B12 and Vitamin C was observed as well as a statistical tendency for lower consumption of vitamin E and niacin in those individuals with high NO.

Table I. Clinical and metabolic characteristics of the studied sample (n = 85), according to the median of nitric oxide concentrations (4.32 μ mol/L)

Variables	Low NO (n = 43)	High NO (n = 42)	p-value
Age (years)	61.3 \pm 14.0	62.7 \pm 13.4	0.637
Dried body weight (kg)	60.4 \pm 10.2	62.4 \pm 11.8	0.425
HD time (months)	52.0 (0.0-245.0)	33.5 (1.0-147.0)	0.062
BMI (kg/m ²)	23.1 \pm 3.9	23.8 \pm 3.5	0.395
WC (cm)	88.1 \pm 9.6	91.0 \pm 10.35	0.198
Lean mass (kg)	44.5 \pm 9.0	44.8 \pm 8.7	0.877
Visceral fat mass (kg)	8.6 \pm 3.8	9.9 \pm 3.9	0.145
Total body fat (%)	21.9 \pm 10.9	23.6 \pm 7.6	0.145
SGAm	13.0 (9.0-21.0)	11.0 (8.0-19.0)	0.012
Albumin (g/dL)	4.0 (3.8-5.0)	4.0 (4.0-5.0)	0.449
RRU	67.5 \pm 7.4	69.0 \pm 8.3	0.389
Ferritin (ng/mL)	455.5 \pm 333.1	606.1 \pm 449.1	0.084
Iron (μ g/dL)	58.2 \pm 21.8	67.1 \pm 21.1	0.050
SatFe (%)	26.7 \pm 12.0	32.1 \pm 11.4	0.037
Hb (mg/dL)	11.0 (5.0-14.0)	11.0 (7.0-14.0)	0.664
Creatinine (mg/dL)	8.89 \pm 2.4	8.69 \pm 3.7	0.743
Pre-dialysis urea (mg/dL)	123.9 \pm 32.2	121.9 \pm 34.3	0.778
Post-dialysis urea (mg/dL)	37.0 (14.5-83.0)	42.0 (10.0-110.0)	0.283
Kt/V	1.49 \pm 0.31	1.54 \pm 0.47	0.601
PTH (pg/mL)	308.0 (75.0-1771.0)	250.5 (34.9-883.6)	0.044
Calcium (mg/dL)	9.0 (4.5-10.0)	9.0 (7.0-10.0)	0.956
Phosphorus (mg/dL)	4.0 (2.0-8.0)	4.9 (2.0-10.0)	0.618
Ca-P product	39.23 \pm 14.9	39.3 \pm 16.5	0.971
Triacylglycerol (mg/dL)	147.0 (42.0-330.0)	181.0 (65.0-934.0)	0.003
Cholesterol (mg/dL)	183.3 \pm 41.8	195.5 \pm 42.4	0.185

BMI: body mass index; WC: waist circumference; SGAm: subjective global assessment modified; RRU: rate of reduction of urea; SatFe: transferrin saturation; Hb: Hemoglobin; PTH: parathyroid hormone. Values expressed as mean \pm SD or median and confidence interval according to distribution; p-values by Student t-test or Mann-Whitney test, as appropriated.

Table II. Dietary intake of the studied sample (n = 85), according to the median of nitric oxide concentrations (4.32 µmol/L)

Daily intake	Low NO (n = 43)	High NO (n = 42)	p-value
Caloric intake (Kcal)	2726.5 (821.0-6666.9)	2205.3 (912.4-6241.6)	0.045
Protein (%VCT)	11.6 (7.07-19.37)	12.9 (8.52-24.20)	0.035
Lipid (%VCT)	31.5 (18.51-62.83)	30.1 (17.02-46.50)	0.705
Carbohydrate (%VCT)	56.52 ± 9.28	56.21 ± 6.35	0.857
Fiber (g)	27 (12.2-86.1)	25.5 (9.2-68.1)	0.271
Cholesterol (mg)	242.0 (35.4-759.5)	232.3 (63.6-729.8)	0.806
Saturated fat (%VCT)	7.95 (4.66-23.51)	8.37 (5.49-15.33)	0.429
Monounsaturated fat (%VCT)	9.63 (4.98-26.87)	9.07 (4.46-16.40)	0.660
Polyunsaturated fat (%VCT)	10.93 ± 3.92	9.72 ± 3.38	0.131
Linoleic fatty acid (ω6) (g)	39.9 (5.2-103.1)	21.6 (3.7-61.9)	0.020
α-linoleic fatty acid (ω3) (g)	4.3 ± 2.9	3.2 ± 1.9	0.021
Calcium (g)	684.1 (199.4-2952.7)	645.6 (229.1-2036.8)	0.847
Magnesium (mg)	271.1 (115.3-907.2)	247.9 (102.0-779.9)	0.345
Manganese (mg)	5.0 (1.5-16.7)	3.8 (1.1-21.4)	0.035
Potassium (mg)	2564 (1005-8276)	2439 (935-8268)	0.285
Sodium (mg)	1526.0 (448.6-8346.5)	1517.3 (279.3-5189.1)	0.368
Phosphorus (mg)	417.3 (120.6-1213.0)	350.0 (121.61574.6)	0.277
Iron (mg)	14.4 (4.4-180.2)	12.3 (4.8-337.7)	0.482
Copper (µg)	1.6 (0.5-219.7)	1.2 (0.4-221.0)	0.026
Zinc (mg)	116.3 (3.1-385.8)	133.4 (3.9-559.8)	0.900
Selenium (µg)	100.3 (27.8-302.4)	87.2 (28.1-290.0)	0.552
Thiamine (mg)	1.6 (0.6-5.5)	1.3 (0.5-5.2)	0.084
Niacin (mg)	14.0 (4.2-42.5)	13.0 (4.0-59.5)	0.058
Vitamin B6 (mg)	1.8 (0.4-5.5)	1.6 (0.4-6.8)	0.117
Vitamin B9 (mg)	417.3 (120.6-1213.0)	350 (121.6-1574.4)	0.277
Vitamin B12 (µg)	2.7 (0.4-20.2)	3.5 (1.0-39.9)	0.549
Vitamin E (UI)	8.1 (2.8-19.5)	7.2 (1.5-19.0)	0.050
Vitamin C (mg)	177.1 (19.3-2575.6)	151.7 (11.6-1505.8)	0.241

Values expressed as mean ± SD or median and confidence interval according to distribution; p-values by Student t-test or Mann-Whitney test, as appropriated.

NO AND METABOLIC MARKERS

Regarding metabolic markers, mean values of iron and satFe are in the normal range and are statistically higher in subjects with a high NO concentration (Table I). PTH presented lower values for median in subjects with high NO concentrations ($p = 0.044$), whereas an inverse behavior was observed with serum triglyceride levels ($p = 0.003$) at high NO concentrations. On the other hand, ferritin presented a significant trend ($p = 0.084$) with a mean higher than that recommended for high NO concentrations.

NO, INFLAMMATORY MARKERS AND OXIDATIVE STRESS

Interestingly, there was a positive correlation of NO with the SOD enzyme ($r = 0.616$ $p < 0.001$), and negative correlation with total protein ($r = -0.214$ $p = 0.049$), as shown in figure 1. No significant

correlations were found among others markers such as MDA and GST, or TAC when related to NO. Furthermore, NO concentrations were negatively correlated with IL-6 and IL-10 concentrations, pro- and anti-inflammatory markers, respectively (Fig. 2).

In addition, the possible contribution of clinical and anthropometric variables, as well as metabolic, inflammatory and oxidative stress markers, and dietary intake to the NO variation in HD individuals were evaluated through bivariate regression analysis. Thus, IL-6, SOD, triacylglycerols, iron, transferrin saturation, ferritin and ingestion of ω6 were significantly association with NO concentration (Table III). In relation to inflammatory and oxidative stress markers, MDA, SOD, GST and IL-10 were positively associated, whereas IL-2, IL-4 and IL-6 formed negative predictors of NO (Table IV).

Finally, in the multiple regression model, the concentrations of ferritin, triacylglycerols, IL-6 and SOD contributed with a 54.8% of variation in NO concentrations, whereas triacylglycerols and SOD concentrations were independent predictors (Table V).

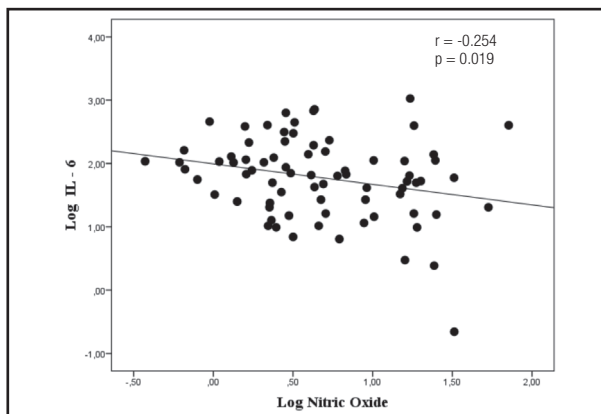


Figure 1.

Spearman correlation between nitric oxide and Interleukin 6 in subjects in HD (n = 85).

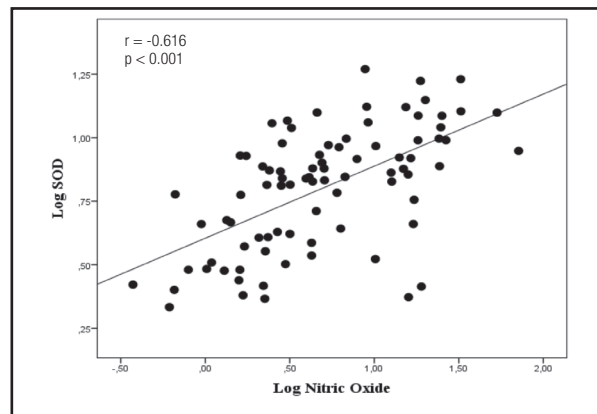


Figure 2.

Spearman correlation between nitric oxide and SOD (superoxide dismutase) in subjects in HD (n = 85).

Table III. Bivariate linear regression to explain the variation of NO concentrations (dependent variable) in subjects in HD (n = 85) in relation to clinical-metabolic variables

Independent variables	Coefficient (β)	CI 95%	p-value	R ²
Gender	-0.360	-0.262-0.191	0.756	0.00
Age (years)	-0.165	-0.382-0.053	0.135	0.02
HD time (months)	-0.001	-0.003-0.001	0.219	0.01
Kt/V	0.128	-0.142-0.398	0.348	0.01
BMI (kg/m ²)	0.018	-0.010-0.047	0.206	0.01
WC (cm)	0.008	-0.003-0.018	0.154	0.01
RCQ	1.171	-0.212-2.555	0.096	0.03
Lean mass (kg)	0.000	-0.013-0.012	0.967	0.00
Visceral fat (kg)	0.019	-0.008-0.046	0.159	0.02
Total corporal fat (%)	0.004	-0.008-0.015	0.536	0.00
Liquid rate	-0.008	-0.021-0.005	0.217	0.01
SGAm	0.032	-0.073-0.010	0.135	0.02
Albumin (g/dL)	-0.046	-0.493-0.400	0.837	0.00
PTH (pg/mL)	0.000	0.000-0.000	0.258	0.01
Calcium (mg/dL)	0.009	-0.113-0.132	0.879	0.00
Triacylglycerol (mg/dL)	0.002	0.001-0.002	< 0.001	0,26
Ferritin (ng/mL)	0.000	0.000-0.001	0.026	0.05
Iron (μ g/dL)	0.004	- 0.001-0.009	0.111	0.03
SatFe (%)	0.009	0.00-0.018	0.049	0.04
Calorie intake (Kcal)	0.005	0.000-0.000	0.096	0.03
Protein consumption (g)	-0.002	-0.005-0.001	0.232	0.01
Lipid consumption (g)	-0.002	-0.004-0.000	0.070	0.03
Carbohydrate consumption (g)	0.000	-0.001-0.000	0.108	0.03
Polyunsaturated fat (g)	-0.005	-0.011-0.000	0.050	0.04
ω 3 (g)	-0.044	-0.091-0.003	0.064	0.04
ω 6 (g)	-0.006	-0.012-0.000	0.049	0.21
Manganese (mg)	-0.022	-0.056-0.011	0.187	0.02
Copper (mg)	-0.002	-0.005-0.001	0.208	0.01
Zinc (mg)	0.000	-0.001-0.001	0.819	0.00
Selenium (μ g)	0.001	-0.002-0.001	0.569	0.00
Vitamin E (mg)	-0.019	-0.047-0.009	0.176	0.02
Vitamin C (mg)	0.000	-0.001-0.000	0.237	0.01
Niacin (mg)	-0.009	-0.021-0.003	0.153	0.02

BMI: body mass index; WC: waist circumference; SGAm: subjective global assessment modified; RRU: rate of reduction of urea; SatFe: saturation transferrin; PTH: parathyroid hormone.

Table IV. Bivariate linear regression to explain the variation of NO concentrations (dependent variable) in subjects in HD (n = 85) in relation to inflammatory markers and oxidative stress

Independent variables	Coefficient (β)	CI 95%	p-value	R ²
Superoxide dismutase	0.532	0.352-0.713	< 0.001	0.293
Malondialdehyde	0.279	0.073-0.485	0.009	0.080
Glutathione-S- transferase	0.140	- 0.073-0.353	0.194	0.020
Total antioxidant capacity	-0.134	- 0.347-0.079	0.216	0.018
IL-2	-0.024	- 0.053-0.005	0.101	0.032
IL-4	- 0.113	- 0.214-0.013	0.028	0.057
IL-6	- 0.253	- 0.461- 0.045	0.018	0.066
IL-10	0.171	- 0.180- 0.024	0.011	0.075
IL-17	- 0.010	- 0.093-0.072	0.807	0.001
C-reactive protein	0.002	- 0.994-0.998	0.997	0.000

Table V. Multiple linear regression to explain the variation of NO concentrations (dependent variable) in subjects in HD (n = 85)

Independent variables	Coefficient (β)	CI 95%	(β) Standardized	p-value
Ferritin (ng/mL)	0.084	-0.100-0.267	0.073	0.366
Triacylglycerols (mg/dL)	0.931	0.598-1.265	0.455	< 0.001
IL-6 (pg/ml)	- 0.55	-0.183-0.072	-0.070	0.388
SOD (U/mg protein)	0.973	0.644-1.302	0.479	< 0.001

Adjusted R²: 0.548. F-test p < 0.0001. IL-6: interleukin 6; SOD: superoxide dismutase.

DISCUSSION

The present study evaluated the potential association of NO variation with metabolic, inflammatory and oxidative stress markers in individuals submitted to HD. Our most relevant result was the concentration of SOD, also recognized marker of oxidative stress, as an independent predictor of NO variation.

In this context, NO produced by eNOS under favorable conditions may induce the production of SOD in the muscular layer of the vessel and extracellularly reducing available superoxide radicals (O_2^-) and, consequently, peroxynitrite production (ONOO-) and oxygen reactive species (ROS) expression. In fact, increased production of ROS, such as superoxide, hydrogen peroxide and lipoperoxides, in addition to decreased NO synthesis and concentrations of antioxidants such as vitamin E and SOD, has been observed in hypertension patients when compared to normal individuals. These individuals with hypertension still have decreased concentrations of antioxidants such as vitamin E and SOD (29). It is worth mentioning that vitamin E could also have a pro-oxidant action under special conditions that can be found in HD patients (30). In fact, oral administration of α -tocopherol (500 mg/day) for 1 year for HD patients caused a reduction in SOD activity and total antioxidant status (31). This may be due to the low level of other antioxidants needed to restore the reduced form of vitamin E (e.g., vitamin C) (31). Although vitamin E therapy has been extensively

studied in patients with CKD, there is no consensus on the benefit obtained from its administration (32). The same was found in study by Hambali et al. (33) also found reduced plasma NO in all subjects after HD when compared to controls and consequently reduced SOD, demonstrating a direct relationship between SOD and NO.

Moreover, the cytosolic SOD enzyme is copper and zinc dependent. The decrease of these ions in patients receiving HD may contribute to a decrease in SOD activity and a consequent increase in inflammatory expression (34). In this context, some studies have described interactions between zinc/ copper deficiency and nitrosamine stress with iNOS induction and inflammation, which may contribute to the pathogenesis of diarrheal and cardiovascular diseases (35). Zinc, in turn, has anti-inflammatory properties *in vivo* because of its ability to suppress the induction of cytokines by iNOS, since it is an antioxidant enzyme. Zn supplementation can improve taste and smell and gastrointestinal function, increase food intake, and reduce protein-energy waste (36). Thus, patients with Zn deficiency receiving Zn supplementation have improvements in their antioxidant-antioxidant balance and nutritional status (37), which probably contributes to the increase in plasma SE status. In this sense, consumption above recommendations was observed both in individuals with high and low NO concentrations of minerals (Mn, Cu and Zn), vitamin C and α -linolenic fatty acids (ω 3), all nutrients with antioxidant properties. Thus, we hypothesized that our sample

presents a favorable antioxidant system, since these nutrients act as enzymatic cofactors. Studies highlight them by inhibiting lipoprotein oxidation as an anti-peroxidation agent, and indirectly promote iNOS activation antagonistic action, improving NO vasodilator vascular action, decreasing the available (O_2^-) (38). Evidence also shows that polyunsaturated fatty acids, especially $\omega 3$, promote an increase in the regulation of the NO system by iNOS (39). Thus, the results of dietary intake in relation to NO concentrations reinforce our hypothesis that there is a suitable system for NO activity. However, these benefits were not confirmed by Kooshki et al. (40) who did not observe improvements in F2-isoprostane levels nor in carbonylic proteins after supplementation with 2.08 g/day EPA 1 DHA and 800 mg/day DHA 1 vitamin E, respectively. These findings corroborate the results of the study by Mattos et al. 2017 (41) in which supplementation at physiological doses of n-3 PUFA was not able to alter oxidative stress profiles. However, linear regression analysis showed that n-3 PUFA is associated with improved rates of isoprostane and advanced oxidation protein products (AOPP) in HD patients.

The second relevant result of this study was a negative association between inflammatory markers and NO concentrations. In fact, the increase of NO, through the regulation of iNOS by inflammatory cytokines, such as IL-1, IL-6, and TNF in patients submitted to HD has been demonstrated in the literature (42). In addition, Amore et al. (43) demonstrated that abnormal stimulation of iNOS by cytokines was closely associated with the development of vasculopathy in long-term dialysis patients, consistent with the inverse assumption in our study.

Another interesting result of our study was the higher concentrations of triacylglycerols in those individuals with high NO concentration. The same was observed in the study by Volpe et al. (44). Free fatty acids are stored in the body in the form of triacylglycerols and are released into the tissues by lipolysis. This triacylglycerol increases NO production in the skeletal muscle, through the iNOS, contributing to the initiation of the inflammatory cascade, by activation of transcription factor NF- κ B (45). Taken together, our and previous outcomes suggest that high triacylglycerol may contribute to increase NO production in situations of inadequate antioxidant defense.

In addition, our results showed a positive association between serum iron values and transferrin saturation and a significant trend for ferritin and NO concentration in the individuals studied. Iron supplementation is a common recommendation for patients with renal disease; however, excess iron can act as a pro-oxidant factor, thus contributing to the oxidation of molecules, such as NO. This, produced by eNOS, induces the synthesis of ferritin, which binds to free iron ions and prevents the generation of O_2^- . However, under conditions of vascular endothelium impairment, activated macrophages produce O_2^- , express iNOS and produce NO. In this way, ONOO $^-$ and hydroxyl radical (OH $^\cdot$) are produced, compromising tissue integrity, which favors the activation of coagulation and contributes to vascular lumen obstruction, increasing the response of vasoconstrictors such as Angiotensin II (All) (46). Thus, it appears that iron being free can aggravate oxidative stress in individuals in HD and, consequently, contribute to atherosclerosis and oxidative stress.

Another important finding is that anemia is a common complication observed in renal patients and the administration of recombinant human erythropoietin (RHE) and intravenous iron are recommended by the Kidney Disease Outcomes Quality Initiative (KDOQI) Clinical Practice Guidelines (47). Previous studies have indicated that anemia acts as a contributing factor associated with CKD and oxidative stress, while adjuvant therapies, mainly intravenous (i.v.) iron, seem to further increase this process (48). However, in our study, mean values of ferritin higher than 500 ng/mL were observed in those individuals with high NO concentrations and adequate Fe and SatFe values in the majority of participants, remembering that Fe supplementation in this population is common. Since ferritin with higher values in the group of individuals with high NO concentration and eNOS stimulating the synthesis of ferritin to sequester the free Fe (49) we could suggest that there is an adequate production of NO with vasodilator function in this population.

Finally, in the present study, mGSA presented significantly higher scores, corresponding to malnutrition status, being statistically significant at low NO concentrations ($p = 0.012$). In this sense, the pro-inflammatory state, oxidative stress, endothelial dysfunction and malnutrition resulting from these pathological processes are common to CKD. mGSA is a reliable tool for assessing early malnutrition (50). The presence of malnutrition may contribute to the reduction of NO synthesis, release and activity by eNOS, activating several components of the atherogenic process, such as vasoconstriction (51). The study by Silva et al. corroborates with these findings, since they observed a blockade in the transport of L-arginine and synthesis of nitric oxide, being this one associated with the increase of the platelet aggregation in individuals with malnourished DRT (52). In our study, there was no association between the markers of body composition and NO. One explanation for this result is that the majority of study participants presented normal albumin values. In this sense, Danielski et al. (53) demonstrated that inflammatory and oxidative stress markers were increased in patients with hypoalbuminemia when compared to normoalbuminemic patients. In addition, our samples presented in majority normal weight and adequate body fat mass. Thus, the lack of association between oxidative stress and body fat composition markers may be influenced by the low prevalence of malnutrition or by the antioxidant effect of albumin.

In conclusion, the present cross-sectional study showed a significant association of NO with markers of lipid and iron metabolism, as well as with inflammatory markers (IL6 and IL10) and oxidative stress (SOD) in HD patients, indicating its important risk mediator of this population. In addition, the clinical-nutritional status and nutrient consumption with antioxidant properties (Cu, Zn, Mn, vitamin C and $\omega 3$) seem to modulate this relationship.

ACKNOWLEDGEMENTS

The authors declare that they have no conflicts of interest. We thank the volunteers and the master's and doctoral scholarships from CAPES and CNPq. We also thank the FAPEMIG (Minas Gerais/Brazil) and CNPq by financial support. H.H.M. Hermsdorff is CNPq fellow.

REFERENCES

1. Balasubramanian S. Progression of chronic kidney disease: mechanisms and intervention in retardation. *J Ap Med* 2010;10:19-28.
2. Shah S, Baliga R, Rajapurkar M, et al. Oxidants in chronic kidney disease. *J Am Soc Nephrol* 2007;18:16-28.
3. Cachofeiro V, Goicochea M, Vinuesa S. Oxidative stress and inflammation a link between chronic kidney disease and cardiovascular disease. *Kidney Int* 2008;74:4-9.
4. Kuo K, Tarng D. Oxidative stress in chronic kidney disease. *Adaptive Medicine*; 2010;2:87-94.
5. Snyder SH, Bredt DS. Biological role of nitric oxide. *Science Am* 1992;266:68-77.
6. Förstermann U, Sessa W. Nitric oxide synthases: regulation and function. *European Heart Journal*, Oxford 2011;31:829-37.
7. Baylis C. Nitric oxide deficiency in chronic kidney disease. *American Journal of Physiology - Renal Physiology*, Bethesda 2008;294:F1-F9.
8. Modlinger PS, Wilcox CS, Aslam S. Nitric oxide, oxidative stress, and progression of chronic renal failure. *Seminars in Nephrology* 2004;24:354-65.
9. Harrison DG. The mosaic theory revisited: common molecular mechanisms coordinating diverse organ and cellular events in hypertension. *Journal of the American Society of Hypertension* 2013;7:68-74.
10. Kalantar-Zadeh K, Kleiner, M, Dunne E, et al. A modified quantitative subjective global assessment of nutrition for dialysis patients. *Nephrol Dial Transplant* 1999;14:1732-8.
11. Jeliffe DB. Evaluation del estado de nutrición de la comunidad com especial referencia a las encuestas em las regions in desarrollo. *Genebra: Organización Mundial de la Salud*; 1968.
12. World Health Organization – WHO Physical status: the use and interpretation of anthropometry. *Geneva: Technical Report Series*, 854; 1995.
13. Coelho MASC, Amorim RB. Avaliação nutricional em geriatria. In: Duarte ACG. *Avaliação nutricional: Aspectos clínicos e laboratoriais*. São Paulo: Atheneu; 2007. pp. 155-76.
14. World Health Organization Obesity: preventing and managing the global epidemic. *Genebra*; 1997.
15. Lipschitz DA. Screening for nutritional status in the elderly. *Prim Care* 1994;21:55-67.
16. Kalantar-Zadeh K, Kopple JD, Deepak S et al. Features food consumption G. Block hemodialysis patients as obtained by food frequency questionnaire. *J Nutr Ren* 2002;12:17-31.
17. Molina MDB, Bensenor I, Cardoso LO, et al. Reprodutibilidade e validade relativa do Questionário de Frequência Alimentar do ELSA-Brasil. *Cad. Saúde Pública* 2013;29:79-389.
18. Brasil, Ministério da Saúde: Guia Alimentar para População Brasileira promovendo a alimentação saudável. Normas e manuais técnicos: Brasília; 2006.
19. Brasil, Ministério da Saúde (2014): Guia Alimentar para População Brasileira promovendo a alimentação saudável. 2ªed. Normas e manuais técnicos: Brasília. IBGE – Instituto Brasileiro de Geografia e Estatística. Pesquisa de Orçamentos Familiares 2002-2003 – POF. Rio de Janeiro, 2004. Pesquisa de Orçamentos Familiares 2008-2009 – POF. Rio de Janeiro; 2010.
20. NEPA – Núcleo de estudos e Pesquisas em Alimentação. Tabela Brasileira de Composição de Alimentos (TACO). 1ª ed. Campinas: NEPA – UNICAMP 2004;42.
21. Hemodialysis Adequacy 2006 Work Group. Clinical practice guidelines for hemodialysis adequacy, update 2006. *Am J Kidney Dis* 2006;48(1):S2-S90.
22. Dieterich S, Bieligg U, Beulich K. Gene expression of antioxidative enzymes in the human heart: increased expression of catalase in the end-stage failing heart. *Circulation* 2000;101:33-9.
23. Habig W, Jakoby WB. Assays for determination of glutathione S-transferase. *Methods Enzymol* 1981;77:398-405.
24. Wallin B, Rosengren B, HG Shertzer, et al. Lipoprotein oxidation and measurements of thiobarbituric acid reacting substances formation in single microtiter plate: its use of evaluation of antioxidants. *Analytical Biochemistry* 1993;208:10-5.
25. Green LC, Wagner DA, Glogowski J. Analysis of nitrate, nitrite and [¹⁵N] nitrate in biological fluids. *Anal Biochem* 1982;126:131-8.
26. Lowry OH, Rosebrough NJ, Farr AL, et al. Protein measurement with the folin phenol reagent. *J Biol Chem* 1951;193:265-75.
27. Hermsdorff HHM, Puchau B, Bressan J, et al. Association of body Fat distribution with proinflammatory gene expression in peripheral blood mononuclear cells from young adult subjects. *OMICS: J Integr Biol* 2010;14:297-307.
28. Willett WC. *Nutritional epidemiology*. 2nd ed. New York: Oxford University Press; 1998.
29. Kumar KV, Das UN. Are free radicals involved in the pathobiology of human essential hypertension? *Free Rad Res Comms* 1993;19:59-66.
30. Antoniadì G, Eleftheriadis T, Liakopoulos V, et al. Effect of one-year oral α -tocopherol administration on the antioxidant defense system in hemodialysis patients. *Ther Apher Dial* 2008;12:237-42.
31. Sesso HD, Buring JE, Christen WG, et al. Vitamins E and C in the prevention of cardiovascular disease in men: The Physicians' Health Study II randomized controlled trial. *JAMA* 2008;300:2123-33.
32. Chen S-C, Lee M-Y, Huang J-C, et al. Association of diabetes mellitus with decline in ankle-brachial index among patients on hemodialysis: A 6-year follow-up study. *Cheng X, editor. PLoS ONE* 2017;12(4).
33. Hambali Z. Oxidative Stress and Its Association with Cardiovascular Disease in Chronic Renal Failure Patients. *Indian Journal of Nephrology* 2011;21(1):21-5.
34. Gheddouchi S, Mokhtari-Soulimane N, Merzouk H, et al. Low SOD activity is associated with overproduction of peroxynitrite and nitric oxide in patients with acute coronary syndrome. *Nitric Oxide* 2015;49:40-6.
35. Torregrossa A, Berkson, Moncrief J. *Nitric Oxide* 2011;24:S21-S22.
36. Guo CH, Wang CL, Chen PC. Micronutrient Metabolism in Hemodialysis Patients. In: Maria GP, editor. *Hemodialysis - Different Aspects*. Croatia: InTech: Rijeka; 2011. pp. 173-204.
37. Mazani M, Argani H, Rashtchizadeh N, et al. Effects of zinc supplementation on antioxidant status and lipid peroxidation in hemodialysis patients. *J Ren Nutr J Ren Nutr* 2013;23(3):180-4.
38. Thomas SR, Stocker R. Molecular action of vitamin E in lipoprotein oxidation: implications for atherosclerosis. *Free Radic Biol Med* 2000;28:1795.
39. Kooshki A, Taleban FA, Tabibi H, et al. Effects of omega-3 fatty acids on serum lipids, lipoprotein (a), and hematologic factors in hemodialysis patients. *Ren Fail* 2011;33:892-8.
40. Kooshki A, Taleban FA, Tabibi H, et al. Effects of Marine Omega-3 Fatty Acids on Serum Systemic and Vascular Inflammation Markers and Oxidative Stress in Hemodialysis Patients. *Ann Nutr Metab* 2011;58(3):197-202.
41. de Mattos AM, da Costa JAC, Jordão Júnior AA, et al. Omega-3 fatty acid supplementation is associated with oxidative stress and dyslipidemia, but does not contribute to better lipid and oxidative status on hemodialysis patients. *J Ren Nutr* 2017;17:S1051-2276.
42. Madore F, Prud'homme L, Austin JS, et al. Impact of nitric oxide on blood pressure in hemodialysis patients. *Am J Kidney Dis* 1997;30:665-671.
43. Amore A, Bonaudo R, Ghigo D, et al. Enhanced production of nitric oxide by blood-dialysis membrane interaction. *J Am Soc Nephrol* 1995;6:1278-83.
44. Volpe, CMO et al. The Production of Nitric Oxide, IL-6, and TNF-Alpha in Palmitate-Stimulated PBMCs Is Enhanced through Hyperglycemia in Diabetes. *Oxidative Medicine and Cellular Longevity* 2014. 479587. PMC. Web. 30 Sept. 2016.
45. Suganami T, Tanimoto-Koyama K, Nishida J, et al. Role of the Toll-like receptor 4/NF-kappaB pathway in saturated fatty acid-induced inflammatory changes in the interaction between adipocytes and macrophages. *Arterioscler Thromb Vasc Biol* 2007;27:84-91.
46. Dusse LMS, Vieira LM, Carvalho MG. Revisão sobre óxido nítrico. *Jornal Brasileiro de Patologia e Medicina Laboratorial*, Rio de Janeiro 2003;39:343-50.
47. National Kidney Foundation (KDOQI). Clinical Practice Guidelines and Clinical Practice Recommendations for Anemia in Chronic Kidney Disease. *Am J Kidney Dis* 2006;47:S11-145.
48. Capusa C, Mircescu G. Oxidative stress, renal anemia, and its therapies: Is there a link? *J Ren Nutr* 2010;20:S71-S76.
49. Davis DW, Lewis ML, Hart DA. Age dependent expression and distribution of nitric oxide (NO) synthase isoforms in the ovine kidney. *Open Journal of Molecular and Integrative Physiology* 2013;3:61-70.
50. Jones CH, Newstead CG, Will EJ, et al. Assessment of nutritional status in CAPD patients: serum albumin is not a useful measure. *Nephrol Dial Transplant* 1997;12:1406-13.
51. Noris M, Benigni A, Boccardo P, et al. Enhanced nitric oxide synthesis in uremia: Implications for platelet dysfunction and dialysis hypotension *Kidney Int* 1993;44:445-50.
52. Silva CDD, Brunini MCT, Reis MB, et al. Effects of nutritional status on the L-arginine-nitric oxide pathway in platelets from hemodialysis patients. *Kidney International* 2005;68:2173-9.
53. Danielski M, Ikizler TA, McMonagle E, et al. Linkage of hypoalbuminemia, inflammation, and oxidative stress in patients receiving maintenance hemodialysis therapy. *Am J Kidney Dis* 2003;42:286-94.