

Nutritional status and overhydration: can bioimpedance spectroscopy be useful in haemodialysis patients?

Cristina Garagarza¹, Patrícia João-Matias², Catarina Sousa-Guerreiro³, Tiago Amaral², Inês Aires², Carina Ferreira², Cristina Jorge², Célia Gil², Aníbal Ferreira⁴

¹ Departamento de Nutrición. Centro de diálisis Dialverca Lisboa (Portugal)

² Servicio de Nefrología. Centro de diálisis Dialverca Lisboa (Portugal)

³ Servicio de Nefrología. Escola Superior de Tecnologia da Saúde de Lisboa (Portugal)

⁴ Servicio de Nefrología. Centro de diálisis Dialverca Lisboa. Facultad de Ciencias Médicas de la Universidad de Lisboa (Portugal)

Nefrologia 2013;33(5):667-74

doi:10.3265/Nefrologia.pre2013.Jun.12008

ABSTRACT

Background: Protein-energy wasting (PEW), associated with inflammation and overhydration, is common in haemodialysis (HD) patients and is associated with high morbidity and mortality. **Objective:** Assess the relationship between nutritional status, markers of inflammation and body composition through bioimpedance spectroscopy (BIS) in HD patients. **Methods:** This observational, cross-sectional, single centre study, carried out in an HD centre in Forte da Casa (Portugal), involved 75 patients on an HD programme. In all participating patients, the following laboratory tests were conducted: haemoglobin, albumin, C-reactive protein (CRP) and 25-hydroxyvitamin D3 [25(OH)D3]. The body mass index of all patients was calculated and a modified version of subjective global assessment (SGA) was produced for patients on dialysis. Intracellular water (ICW) and extracellular water (ECW) were measured by BIS (Body Composition Monitor®, Fresenius Medical Care®) after the HD session. In statistical analysis, Spearman's correlation was used for the univariate analysis and linear regression for the multivariate analysis (SPSS 14.0). A *P* value of <.05 was considered statistically significant. **Results:** PEW, inversely assessed through the ICW/body weight (BW) ratio, was positively related to age (*P*<.001), presence of diabetes (*P*=.004), BMI (*P*=.01) and CRP (*P*=.008) and negatively related to albumin (*p*=.006) and 25(OH)D3 (*P*=.007). Overhydration, assessed directly through the ECW/BW ratio, was positively related with CRP (*P*=.009) and SGA (*P*=.03), and negatively with 25(OH)D3 (*P*=.006) and BMI (*P*=.01). In multivariate analysis, PEW was associated with older age (*P*<.001), the presence of diabetes (*P*=.003), lower 25(OH)D3 (*P*=.008), higher CRP (*P*=.001) and lower albumin levels (*P*=.004). Over-

hydration was associated with higher CRP (*P*=.001) and lower levels of 25(OH)D3 (*P*=.003). **Conclusions:** Taking these results into account, the ICW/BW and ECW/BW ratios, assessed with BIS, have proven to be good markers of the nutritional and inflammatory status of HD patients. BIS may be a useful tool for regularly assessing the nutritional and hydration status in these patients and may allow nutritional advice to be improved and adjusted.

Keywords: Body composition. Haemodialysis. Inflammation. Nutritional status.

Estado nutricional e hiperhidratación: ¿la bioimpedancia espectroscópica es válida en pacientes en hemodiálisis?

RESUMEN

Antecedentes: El desgaste proteico-energético (DPE), asociado a inflamación e hiperhidratación, es común en pacientes en hemodiálisis (HD) y se asocia a mayor morbilidad y mortalidad. **Objetivo:** Evaluar la relación entre el estado nutricional, los marcadores inflamatorios y la composición corporal a través de bioimpedancia espectroscópica (BIS) en pacientes en HD. **Métodos:** En este estudio observacional, transversal, unicéntrico, realizado en un centro de HD en Forte da Casa (Portugal), participaron 75 pacientes en programa de HD. En todos los participantes se hicieron las siguientes determinaciones analíticas: hemoglobina, albúmina, proteína C reactiva (PCR) y 25-hidroxivitamina D3 [25(OH)D3]. Se calculó el índice de masa corporal (IMC) de todos los pacientes y se aplicó una versión modificada de la valoración global subjetiva (VGS) para pacientes en diálisis. El agua intracelular (AIC) y extracelular (AEC) se midió con BIS (Body Composition

Correspondence: Cristina Garagarza

Centro de diálisis Dialverca.

Estrada Nacional, 10, 2625-437 Forte da Casa. Lisboa. (Portugal).

cgagarza@hotmail.com

originals

Monitor®, Fresenius Medical Care®) después de la sesión de HD. En el análisis estadístico se utilizó la correlación de Spearman para el análisis univariante y la regresión lineal para el análisis multivariante (SPSS 14.0). Una $p < 0,05$ se consideró estadísticamente significativa. **Resultados:** El DPE, evaluado inversamente a través de la relación AIC/ peso corporal (PC), se relacionó positivamente con la edad ($p < 0,001$), la presencia de diabetes ($p = 0,004$), el IMC ($p = 0,01$) y la PCR ($p = 0,008$) y negativamente con la albúmina ($p = 0,006$) y la 25(OH)D3 ($p = 0,007$). La hiperhidratación, evaluada directamente a través de la relación AEC/PC, se relacionó positivamente con la PCR ($p = 0,009$) y con la VGS ($p = 0,03$), y negativamente con la 25(OH)D3 ($p = 0,006$) y el IMC ($p = 0,01$). En el análisis multivariante, el DPE se asoció a edad más elevada ($p < 0,001$), presencia de diabetes ($p = 0,003$), 25(OH)D3 más baja ($p = 0,008$), PCR más elevada ($p = 0,001$) y niveles de albúmina más bajos ($p = 0,004$). La hiperhidratación se asoció a PCR más elevada ($p = 0,001$) y niveles de 25(OH)D3 más bajos ($p = 0,003$). **Conclusiones:** Teniendo en cuenta estos resultados, las relaciones AIC/PC y AEC/PC, evaluadas con BIS, han demostrado ser buenos marcadores del estado nutricional e inflamatorio de pacientes en programa de HD. La BIS puede ser una herramienta útil para evaluar regularmente el estado nutricional y de hidratación en estos pacientes y puede permitir mejorar y adecuar el asesoramiento nutricional.

Palabras clave: Composición corporal. Hemodiálisis. Inflamación. Estado nutricional. Diálisis.

INTRODUCTION

Protein-energy wasting (PEW), associated with inflammation and overhydration, is common in patients on haemodialysis (HD) and is associated with high morbidity and mortality.¹ C-reactive protein (CRP) is an independent predictor of cardiovascular risk and mortality in advanced chronic kidney disease (ACKD).¹

Subjective global assessment (SGA) is the tool most commonly used for assessing nutritional status and is valid in HD patients.³ Anthropometric data, such as the body mass index (BMI), are indicators of nutritional status in these patients,² but they have their limitations when considered alone. Serum albumin has been widely used as a nutritional marker in HD patients and is one of the most sensitive mortality markers⁴ and a major marker of inflammation.^{1,5} For this reason, a decrease in albumin may reflect an inflammatory state instead of PEW¹ and as albumin decreases with fluid overload, it is less relevant in HD patients.¹ The anti-inflammatory effects of vitamin D are well documented⁶ and several authors, including our research group, have reported a strong correlation between the deficiency of

25-hydroxyvitamin D3 [25(OH)D3] and mortality in both the general population,^{7,8} and in HD patients.^{9,10}

Bioimpedance spectroscopy (BIS) seems to be a valid method for assessing and monitoring hydration and nutritional status in HD patients¹¹ and, apart from determining the individual fluid status (total body water, intra and extracellular water [ICW/ECW]) independently,¹² it analyses body composition (body cell mass, lean mass and fat mass) simply, objectively and non-invasively.¹³ Total body water can be divided into ICW and ECW and fat-free mass (FFM) in ECW and body cell mass (BCM), which includes ICW (Figure 1). The ECW compartment predominantly shows overhydration,^{13,14} which often exists in HD patients and is associated with inflammation and an increased risk of mortality.¹⁵ BCM is the compartment that shows nutritional status¹⁶ and, as it is not greatly affected by changes in hydration, which mainly occur in the compartment that shows overhydration (ECW), it gives us more information than FFM in these patients. The ICW compartment comprises 72% of BCM and is not affected by iso-osmotic changes that occur in the ECW compartment. Furthermore, ICW is often used to estimate BCM, which reflects nutritional status.^{13,14} To assess nutritional status and hydration, two relationships can be used: ICW/body weight (BW), which shows nutritional status (the higher it is, the better nourished the patient is and vice-versa), and ECW/BW, which reflects overhydration.^{14,17}

The main aim of this study was to assess the relationship between nutritional status, inflammation markers and body

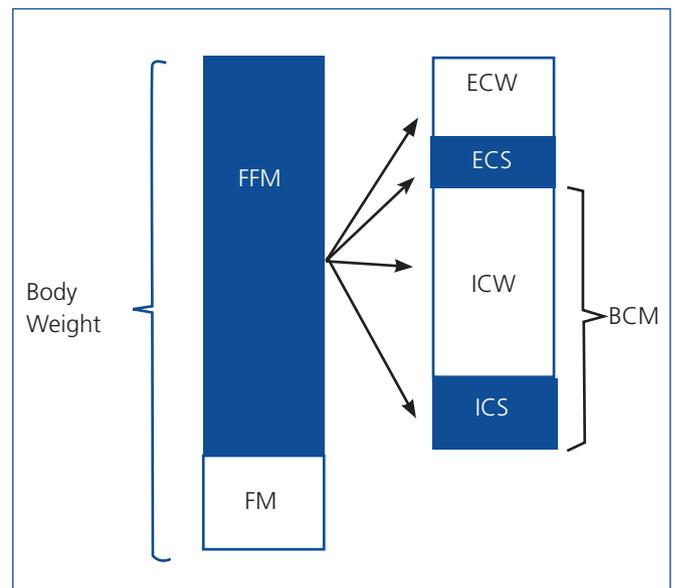


Figure 1. Body composition

BCM: body cell mass; ECW: extracellular water; FFM: fat-free mass; FM: fat mass; ICW: intracellular water; ECS: extracellular solids; ICS: intracellular solids

Adapted from Woodrow, et al.¹³

composition, assessed through BIS, in maintenance HD patients.

METHODS

Study design

An observational, cross-sectional, single centre study of a cohort of maintenance HD patients.

Patients

We included 75 patients on HD three days a week (with an online haemofiltration technique), who had been stable for at least three months, 51% male, 29% diabetic, and all older than 18 years old. The aetiologies of chronic renal failure were: diabetic nephropathy (n=17), hypertensive nephrosclerosis (n=13), chronic glomerulosclerosis (n=9), polycystic kidney disease (n=4), chronic pyelonephritis (n=6), other (n=13), unknown (n=13). The other clinical and nutritional characteristics of the population studied are displayed in Table 1.

All patients were dialysed with high-flux (Helixone®, Fresenius®) membranes and ultrapure water in accordance with the criteria of ISO regulation 13959:2009 - Water for haemodialysis and related therapies. The mean time on HD was 32.3±27.1 months. One of the limitations recognised by the manufacturer of the Body Composition Monitor consists of its use in patients with major amputations of limbs or with pacemakers, since it does not guarantee that the measurements will be accurate in these patients. Therefore, these patients were excluded from the study.

Laboratory tests

The following laboratory tests were carried out: haemoglobin, albumin, CRP and 25(OH)D3. These were calculated before the midweek dialysis session, close to the day on which the BIS was carried out.

Albumin was measured using the colourimetric technique with bromocresol purple (reference value >4.0g/dl) and CRP was obtained by the immunoturbidimetric method. Serum 25(OH)D3 was measured by radioimmunoassay (IDS, Boldon, United Kingdom). The reference value for 25(OH)D3 was 10-60ng/ml.

Analysis of body composition

In all patients BIS was carried out using the Body Composition Monitor (Fresenius Medical Care Deutschland GmbH,

Germany), which takes measurements at 50 frequencies in a range of 5 to 1000KHz. The measurement was performed approximately 30 minutes after the midweek HD session, with four conventional electrodes being placed in the patient, who was lying in the supine position: two in the hand and two in the foot contralateral to the vascular access. Regarding the quality of measurements, all exceeded 95%. The manufacturer of the Body Composition Monitor (Fresenius Medical Care) indicated that 30 minutes after the HD session, the balance between intra-and extracellular fluid was restored and no significant differences in relation to pre-dialysis values¹⁸ were observed. Parameters obtained directly through BIS that were used in this study were ICW and ECW. PEW, represented

Table 1. Patients' characteristics.

n	75
Female % (n)	49 (37)
Male % (n)	51 (38)
Months on haemodialysis ^a	32.3 ± 27.1
Age (years) ^b	67.5 (20-86)
Diabetics % (n)	29 (22)
BMI (kg/m ²) ^a	25.40 ± 4.56
BMI < 19.5 % (n)	4 (3)
19.5 < BMI > 24.9 % (n)	47 (35)
25 < BMI > 29.9 % (n)	32 (24)
30 < BMI > 34.9 % (n)	15 (11)
BMI > 35 % (n)	2 (2)
SGA	
Adequate nutritional status	0
Nutritional risk % (n)	97 (73)
Mild to moderate malnutrition % (n)	3 (2)
Severe malnutrition	0
ICW/BW (L/kg) ^b	0.26 (0.15-0.74)
ECW/BW (L/kg) ^b	0.20 (0.13-0.28)
KtV ^a	1.4 ± 0.23
Albumin (g/dl) ^a	4.12 ± 0.37
25(OH)D3 (ng/ml) ^a	22.3 ± 11.23
CRP (mg/dl) ^b	0.3 (0-12)
Ferritin (ng/dl) ^b	367 (69.3-838)
Haemoglobin (g/dl) ^a	11.83 ± 0.78

25(OH)D3: 25-hydroxyvitamin D3; ECW/BW: extracellular water/body weight; ICW/BW: intracellular water/body weight; BMI: body mass index; CRP: C-reactive protein; SGA: subjective global assessment.

^a Mean ± standard deviation.

^b Median (interquartile range).

originals

by the ICW/BW ratio, and overhydration, represented by the ECW/BW ratio were analysed.

Subjective global assessment

We used the modified version of the SGA for dialysis patients.¹⁹ The data were obtained taking into account changes in BW, eating habits, the presence of gastrointestinal symptoms, functional activity and the presence of comorbidities. We performed a brief physical test to assess the loss of muscle mass and fat mass and the presence of oedema. Each component of the SGA was assigned a score of 1 (normal) to 5 (very severe). A final rating was attributed to each patient after adding up all the scores of the different components of the SGA; the higher the score, the greater the risk of PEW.

Anthropometric parameters

The following data were collected: height (m) measured with a precision telescopic stadiometer (Seca 222®), dry weight (kg) measured with a calibrated scale (Soehnle® S20) and BMI (kg/m²).

Statistical analysis

Categorical variables are expressed as frequencies, normally distributed variables are expressed as mean \pm standard deviation and variables whose distribution is not normal are expressed as a median (interquartile range). We used Spearman's correlation for univariate analysis and performed a multivariate linear regression analysis (confidence interval 95%).

For statistical calculations, we used the SPSS 14.0 software (SPSS Inc., Chicago, IL, USA). A *P* value of $<.05$ was considered to be statistically significant.

RESULTS

In our study, BMI was not significantly related to nutritional or inflammatory parameters.

Overall, 16 patients (22%) had albumin below 4.0g/dl. The mean value of serum albumin was above 4.0g/dl and the median CRP was below 1mg/dl, in accordance with the K/DOQI guidelines.³ Serum 25(OH)D3 was below the desirable values (<30 ng/ml) in 59 (80%) patients.

In univariate regression analysis, PEW, assessed inversely by the ICW/BW ratio, was positively associated with age, diabetes, BMI, CRP and ferritin. PEW was negatively related

to albumin and 25(OH)D3 (Table 2). Moreover, a negative correlation was found between serum albumin and CRP ($r=-.49$, $P<.001$) and lower concentrations of 25(OH)D3 were associated with higher levels of CRP ($r=-.29$, $P=.03$).

The multivariate analysis shows that PEW is associated with older age, presence of diabetes, lower levels of 25(OH)D3 and albumin and higher CRP levels (Table 3).

In the univariate analysis, overhydration, assessed by the ECW/BW ratio, was positively related to CRP and SGA, and negatively to serum 25(OH)D3 and BMI (Table 2). In the multivariate analysis, overhydration was associated with higher levels of CRP and lower 25(OH)D3 (Table 4).

DISCUSSION

In this study, a worse nutritional status (lower ICW/BW) was associated with older age, diabetes and inflammatory markers. Diabetes mellitus has been identified as a risk factor for PEW in HD patients.²⁰ The value of albumin as a marker of nutritional status has been questioned because its concentration is affected by non-nutritional parameters such as hydration status, loss of albumin in the urine and acidosis,²¹ infection or inflammation, with the latter factor

Table 2. Spearman's correlation coefficients between intracellular water/body weight and extracellular water/body weight with nutritional and inflammatory markers (univariate analysis)

ICW/BW	r	P
Albumin	0.44	0.006
25(OH)D3	0.42	0.007
Age	-0.57	<0.001
Diabetes	-0.45	0.004
BMI	-0.39	0.01
CRP	-0.41	0.008
Ferritin	-0.37	0.01
AEC/PC		
CRP	0.34	0.009
SGA	0.25	0.03
25(OH)D3	-0.40	0.006
BMI	-0.37	0.01

25(OH)D3: 25-hydroxyvitamin D3; ECW/BW: extracellular water/body weight; ICW/BW: intracellular water/body weight; BMI: body mass index; CRP: C-reactive protein; SGA: subjective global assessment.

Table 3. Association between intracellular water/body weight and different parameters, including markers of inflammation and nutrition (multivariate analysis)

Dependent variable	Independent variables	OR (CI 95 %)	P	r ²
ICW/BW	Age	0.22 (0.16. 0.04)	< 0.001	0.411
	Diabetes	0.27 (0.12. 0.07)	0.003	
	CRP	0.25 (0.13. 0.03)	0.001	
	Albumin	2.13 (1.32. 4.80)	0.004	
	25(OH)D3	1.18 (1.12. 3.14)	0.008	

25(OH)D3: 25-hydroxyvitamin D3; ECW/BW: extracellular water/body weight; ICW/BW: intracellular water/body weight; BMI: body mass index; CRP: C-reactive protein; SGA: subjective global assessment.

Table 4. Associations between extracellular water/body weight and markers of inflammation and nutrition (multivariate analysis)

Dependent variable	Independent variables	OR (CI 95 %)	P	r ²
ECW/BW	PCR	2.34 (1.38. 5.23)	0.001	0.437
	25(OH)D3	0.24 (0.12. 0.05)	0.003	

25(OH)D3: 25-hydroxyvitamin D3; ECW/BW: extracellular water/body weight; CI: confidence interval; OR: odds ratio; CRP: C-reactive protein.

being the most important cause leading to the decrease in albumin.^{1,22} through BIS analysis, we demonstrated that a worse nutritional status was associated with lower levels of albumin and increased inflammation (higher levels of CRP and ferritin). Like other authors,²³ our results also indicated a negative correlation between serum albumin and CRP, which suggests that lower albumin levels that are observed in patients with PEW may be secondary to inflammatory processes. According to the data obtained, lower levels of 25(OH)D3 were associated with poorer nutritional status and its deficiency contributes to the development of chronic inflammation because of its association with higher CRP concentrations.²⁴ In our study, lower levels of 25(OH)D3 were associated with higher CRP levels. In this regard, it has been shown that this vitamin supplement leads to reduced CRP levels.²⁵

In accordance with these results, the ICW/BW ratio analysed through BIS appears to be adequate for monitoring the nutritional status of maintenance HD patients, since a worse nutritional status is significantly associated with common conditions in HD patients such as: advanced age, diabetes, inflammation and deficiency of 25(OH)D3.

Our data on BMI are in line with recent studies,^{26,27} which have shown a progressive increase in BMI in patients with ACKD, as well as a greater number of overweight and obese patients.²⁸ In this study, we found a negative association between ICW/BW and BMI, which means that better nourished patients were those with lower BMI values. It should be noted that for patients in this study, in accordance with BMI, only 4% were underweight. In maintenance HD patients, there is an unusual relationship (called a “reverse

epidemiology") between BMI and survival, with increased mortality in patients with low BMI (BMI <20kg/m² [29]) and increased survival in overweight patients (BMI ≥27.5kg/m² [30], BMI >25kg/m² [29,31]) and obese patients (BMI >30kg/m² [31]).^{29,30,32} It should be noted that BMI does not discriminate between lean mass, fat and water, it only determines a ratio between weight and height and not all patients who have a high BMI have it at the expense of fat.³³ However, the use of BMI alone has limitations in assessing the nutritional status of these patients³⁴ because it can hide PEW in overhydrated patients.²

With regard to the SGA, most patients had risk of malnutrition or mild malnutrition, very few had moderate and none severe. It is important to highlight that in the modified version of SGA for dialysis patients, all patients who had been on HD for more than a year, even with normal clinical history parameters and physical examination, obtained a score of 9, which classified them as having nutritional risk or being at risk of mild malnutrition. This situation explains the high prevalence of malnutrition (assessed by SGA) observed in our population, as the mean time on dialysis was greater than 2 years (68% of patients were on HD for more than a year). Similar results have been reported by other authors.³⁵ In our study, the SGA was not related to the ICW/BW ratio, which may have been because there were only two patients with moderate and none with severe malnutrition. Although this nutritional score tool is useful for differentiating severely malnourished patients from those with an adequate nutritional status, it may not be a reliable indicator of the degree of malnutrition.^{1,19,36}

Nonadherence to fluid restrictions can lead to fluid overload³⁷ and this is one of the most difficult aspects of the HD treatment regimen,¹⁵ with a prevalence of perceived failure by patients ranging from 30% to 74%.^{38,39} In our study, overhydration was associated with higher CRP concentrations. This association has previously been described in dialysis patients.^{40,41} The link between inflammation and expansion of extracellular volume is probably caused by intestinal oedema that facilitates the translocation of bacterial toxins.^{42,43} Furthermore, our results show that patients with overhydration had poorer nutritional status (higher SGA and lower 25(OH)D3). It has been reported that the prevalence of PEW, evaluated through the SGA, decreased in patients in whom hydration improved (ECW decreased and ICW increased), and vice-versa.⁴⁴ The same association was found using a different nutritional score (the Bilbrey nutritional index).⁴⁵ Deficiency in 25(OH)D3 has been associated with inhibition of the renin-angiotensin⁴⁶ pathway, which may be related to overhydration. The negative correlation observed between overhydration and BMI suggests that higher BMI values were associated with greater accumulation of body fat. This supports the fact that patients with better nutritional status were those with normal weight, according to the BMI classification. Therefore, hydration status, assessed through regular BIS analysis, can allow detecting

changes in both nutritional and inflammatory status. As has already been described by other authors,¹¹ BIS is an important technique to assess nutritional and hydration status and has proved to be as accurate as the reference methods considered as the gold standard.⁴⁷ Some authors believe that the spectroscopic measurement technique has high reproducibility and specificity.^{47,48} In patients on HD, it has been used to assess dry weight,^{49,50} but it is also valid for assessing body composition, nutritional status,^{14,51} and overhydration.^{52,53} Moreover, it has been shown that BIS, through measuring hydration status, may be suitable for assessing the prognosis of HD patients.^{54,55}

The limitations of our study are its sample size and its cross-sectional, single centre design. However, successive measurements with BIS over time may be very useful for assessing the trend of nutritional status, inflammation and hydration in HD patients, which would allow therapy to be optimised and nutritional advice to be adjusted.

Conflicts of interest

The authors declare that they have no conflicts of interest related to the contents of this article.

REFERENCES

1. Mitch WE, Ikizler TA. Handbook of nutrition and the kidney. 6th ed. Philadelphia, Pa.; London: Lippincott Williams & Wilkins; 2010.
2. Byham-Gray L, Burrowes JD, Chertow GM. Nutrition in kidney disease. Totowa, N.J.: Humana; London: Springer [distributor]; 2008.
3. Clinical practice guidelines for nutrition in chronic renal failure. K/DOQI, National Kidney Foundation. Am J Kidney Dis 2000;35(6 Suppl 2):S1-140.
4. Kalantar-Zadeh K, Kilpatrick RD, Kuwae N, McAllister CJ, Alcorn H Jr, Kopple JD, et al. Revisiting mortality predictability of serum albumin in the dialysis population: time dependency, longitudinal changes and population-attributable fraction. Nephrol Dial Transplant 2005;20(9):1880-8.
5. Friedman AN, Fadem SZ. Reassessment of albumin as a nutritional marker in kidney disease. J Am Soc Nephrol 2010;21(2):223-30.
6. Helming L, Bose J, Ehrchen J, Schiebe S, Frahm T, Geffers R, et al. 1alpha,25-Dihydroxyvitamin D3 is a potent suppressor of interferon gamma-mediated macrophage activation. Blood 2005;106(13):4351-8.
7. Heaney RP. Vitamin D in health and disease. Clin J Am Soc Nephrol 2008;3(5):1535-41.
8. Holick MF. Vitamin D: importance in the prevention of cancers, type 1 diabetes, heart disease, and osteoporosis. Am J Clin Nutr 2004;79(3):362-71.
9. Wolf M, Shah A, Gutierrez O, Ankers E, Monroy M, Tamez H, et al. Vitamin D levels and early mortality among incident hemodialysis patients. Kidney Int 2007;72(8):1004-13.

10. Matias PJ, Ferreira C, Jorge C, Borges M, Aires I, Amaral T, et al. 25-Hydroxyvitamin D3, arterial calcifications and cardiovascular risk markers in haemodialysis patients. *Nephrol Dial Transplant* 2009;24(2):611-8.
11. Zaluska W, Jaroszynski A, Bober E, Malecka T, Kozik J, Ksiazek A. [Measurement of fluid compartments using electrical bioimpedance for assessment of target weight in hemodialysis patients]. *Przegl Lek* 2000;57(12):707-10.
12. Matthie JR. Bioimpedance measurements of human body composition: critical analysis and outlook. *Expert Rev Med Devices* 2008;5(2):239-61.
13. Woodrow G, Devine Y, Cullen M, Lindley E. Application of bioelectrical impedance to clinical assessment of body composition in peritoneal dialysis. *Perit Dial Int* 2007;27(5):496-502.
14. Jacobs LH, van de Kerkhof JJ, Mingels AM, Passos VL, Kleijnen VW, Mazairac AH, et al. Inflammation, overhydration and cardiac biomarkers in haemodialysis patients: a longitudinal study. *Nephrol Dial Transplant* 2010;25(1):243-8.
15. Kalantar-Zadeh K, Regidor DL, Kovesdy CP, Van Wyck D, Bunnapradist S, Horwich TB, et al. Fluid retention is associated with cardiovascular mortality in patients undergoing long-term hemodialysis. *Circulation* 2009;119(5):671-9.
16. Cohn SH, Vaswani AN, Yasumura S, Yuen K, Ellis KJ. Assessment of cellular mass and lean body mass by noninvasive nuclear techniques. *J Lab Clin Med* 1985;105(3):305-11.
17. Earthman C, Traughber D, Dobrzt J, Howell W. Bioimpedance spectroscopy for clinical assessment of fluid distribution and body cell mass. *Nutr Clin Pract* 2007;22(4):389-405.
18. Mamat R, Kong NC, Ba'in A, Shah SA, Cader R, Wong V, et al. Assessment of body fluid status in hemodialysis patients using the body composition monitor measurement technique. *J Clin Nurs* 2012;21(19-20):2879-85.
19. Kalantar-Zadeh K, Kleiner M, Dunne E, Lee GH, Luft FC. A modified quantitative subjective global assessment of nutrition for dialysis patients. *Nephrol Dial Transplant* 1999;14(7):1732-8.
20. Pupim LB, Flakoll PJ, Majchrzak KM, Aftab Guy DL, Stenvinkel P, Ikizler TA. Increased muscle protein breakdown in chronic hemodialysis patients with type 2 diabetes mellitus. *Kidney Int* 2005;68(4):1857-65.
21. Kaysen GA, Rathore V, Shearer GC, Depner TA. Mechanisms of hypoalbuminemia in hemodialysis patients. *Kidney Int* 1995;48(2):510-6.
22. Don BR, Kaysen G. Serum albumin: relationship to inflammation and nutrition. *Semin Dial* 2004;17(6):432-7.
23. Kaysen GA, Stevenson FT, Depner TA. Determinants of albumin concentration in hemodialysis patients. *Am J Kidney Dis* 1997;29(5):658-68.
24. Covic A, Voroneanu L, Goldsmith D. The effects of vitamin D therapy on left ventricular structure and function - are these the underlying explanations for improved CKD patient survival? *Nephron Clin Pract* 2010;116(3):c187-95.
25. Matias PJ, Jorge C, Ferreira C, Borges M, Aires I, Amaral T, et al. Cholecalciferol supplementation in hemodialysis patients: effects on mineral metabolism, inflammation, and cardiac dimension parameters. *Clin J Am Soc Nephrol* 2010;5(5):905-11.
26. Kramer HJ, Saranathan A, Luke A, Durazo-Arvizu RA, Guichan C, Hou S, et al. Increasing body mass index and obesity in the incident ESRD population. *J Am Soc Nephrol* 2006;17(5):1453-9.
27. Barros A, da Costa BE, Poli-de-Figueiredo CE, Antonello IC, d'Avila DO. Nutritional status evaluated by multi-frequency bioimpedance is not associated with quality of life or depressive symptoms in hemodialysis patients. *Ther Apher Dial* 2011;15(1):58-65.
28. Gallar-Ruiz P, Digioia C, Lacalle C, Rodríguez-Villareal I, Laso-Laso N, Hinojosa-Yanahuaya J, et al. Body composition in patients on haemodialysis: relationship between the type of haemodialysis and inflammatory and nutritional parameters. *Nefrologia* 2012;32(4):467-76.
29. Leavey SF, McCullough K, Hecking E, Goodkin D, Port FK, Young EW. Body mass index and mortality in «healthier» as compared with «sicker» haemodialysis patients: results from the Dialysis Outcomes and Practice Patterns Study (DOPPS). *Nephrol Dial Transplant* 2001;16(12):2386-94.
30. Fleischmann E, Teal N, Dudley J, May W, Bower JD, Salahudeen AK. Influence of excess weight on mortality and hospital stay in 1346 hemodialysis patients. *Kidney Int* 1999;55(4):1560-7.
31. Kalantar-Zadeh K, Kopple JD, Kilpatrick RD, McAllister CJ, Shinarberger CS, Gjertson DW, et al. Association of morbid obesity and weight change over time with cardiovascular survival in hemodialysis population. *Am J Kidney Dis* 2005;46(3):489-500.
32. Chazot C, Gassia JP, Di Benedetto A, Cesare S, Ponce P, Marcelli D. Is there any survival advantage of obesity in Southern European haemodialysis patients? *Nephrol Dial Transplant* 2009;24(9):2871-6.
33. Beddhu S. The body mass index paradox and an obesity, inflammation, and atherosclerosis syndrome in chronic kidney disease. *Semin Dial* 2004;17(3):229-32.
34. Yuste C, Abad S, Vega A, Barraca D, Bucalo L, Pérez-de José A, et al. Assessment of nutritional status in haemodialysis patients. *Nefrologia* 2013;33(2):243-9.
35. Oliveira CM, Kubrusly M, Mota RS, Silva CA, Oliveira VN. [Malnutrition in chronic kidney failure: what is the best diagnostic method to assess?]. *J Bras Nefrol* 2010;32(1):55-68.
36. Cooper BA, Bartlett LH, Aslani A, Allen BJ, Ibels LS, Pollock CA. Validity of subjective global assessment as a nutritional marker in end-stage renal disease. *Am J Kidney Dis* 2002;40(1):126-32.
37. Denhaerynck K, Manhaeve D, Dobbels F, Garzoni D, Nolte C, De Geest S. Prevalence and consequences of nonadherence to hemodialysis regimens. *Am J Crit Care* 2007;16(3):222-35; quiz 36.
38. Vlamincck H, Maes B, Jacobs A, Reyntjens S, Evers G. The dialysis diet and fluid non-adherence questionnaire: validity testing of a self-report instrument for clinical practice. *J Clin Nurs* 2001;10(5):707-15.
39. Kugler C, Vlamincck H, Haverich A, Maes B. Nonadherence with diet and fluid restrictions among adults having hemodialysis. *J Nurs Scholarsh* 2005;37(1):25-9.
40. Demirci MS, Demirci C, Ozdogan O, Kircelli F, Akcicek F, Basci A, et al. Relations between malnutrition-inflammation-atherosclerosis and volume status. The usefulness of bioimpedance analysis in peritoneal dialysis patients. *Nephrol Dial Transplant* 2011;26(5):1708-16.
41. Paniagua R, Ventura MD, Avila-Díaz M, Hinojosa-Heredia H, Méndez-Durán A, Cueto-Manzano A, et al. NT-proBNP, fluid volume overload and dialysis modality are independent predictors of mortality in ESRD patients. *Nephrol Dial Transplant* 2010;25(2):551-7.
42. Goncalves S, Pecoits-Filho R, Perreto S, Barberato SH, Stinghen AE, Nefrologia 2013;33(5):667-74

- et al. Associations between renal function, volume status and endotoxaemia in chronic kidney disease patients. *Nephrol Dial Transplant* 2006;21(10):2788-94.
43. Ortega O, Gallar P, Muñoz M, Rodríguez I, Carreño A, Ortiz M, et al. Association between C-reactive protein levels and N-terminal pro-B-type natriuretic peptide in pre-dialysis patients. *Nephron Clin Pract* 2004;97(4):c125-30.
44. Cheng LT, Tang W, Wang T. Strong association between volume status and nutritional status in peritoneal dialysis patients. *Am J Kidney Dis* 2005;45(5):891-902.
45. Espinosa Cuevas MA, Navarrete Rodriguez G, Villeda Martinez ME, Atilano Carsi X, Miranda Alatrister P, Tostado Gutiérrez T, et al. Body fluid volume and nutritional status in hemodialysis: vector bioelectric impedance analysis. *Clin Nephrol* 2010;73(4):300-8.
46. Li YC, Qiao G, Uskokovic M, Xiang W, Zheng W, Kong J. Vitamin D: a negative endocrine regulator of the renin-angiotensin system and blood pressure. *J Steroid Biochem Mol Biol* 2004;89-90(1-5):387-92.
47. Wabel P, Chamney P, Moissl U, Jirka T. Importance of whole-body bioimpedance spectroscopy for the management of fluid balance. *Blood Purif* 2009;27(1):75-80.
48. Moon JR, Stout JR, Smith AE, Tobkin SE, Lockwood CM, Kendall KL, et al. Reproducibility and validity of bioimpedance spectroscopy for tracking changes in total body water: implications for repeated measurements. *Br J Nutr* 2010;104(9):1384-94.
49. Macheck P, Jirka T, Moissl U, Chamney P, Wabel P. Guided optimization of fluid status in haemodialysis patients. *Nephrol Dial Transplant* 2010;25(2):538-44.
50. Passauer J, Petrov H, Schleser A, Leicht J, Pucalka K. Evaluation of clinical dry weight assessment in haemodialysis patients using bioimpedance spectroscopy: a cross-sectional study. *Nephrol Dial Transplant* 2010;25(2):545-51.
51. Donadio C, Consani C, Ardini M, Bernabini G, Caprio F, Grassi G, et al. Estimate of body water compartments and of body composition in maintenance hemodialysis patients: comparison of single and multifrequency bioimpedance analysis. *J Ren Nutr* 2005;15(3):332-44.
52. van de Kerkhof J, Hermans M, Beerenhout C, Konings C, van der Sande FM, Kooman JP. Reference values for multifrequency bioimpedance analysis in dialysis patients. *Blood Purif* 2004;22(3):301-6.
53. Park J, Chung HC, Kim MS, Kim SJ, Chang JW, Lee JS. Relationship between extracellular water fraction of total body water estimated by bioimpedance spectroscopy and cardiac troponin T in chronic haemodialysis patients. *Blood Purif* 2009;28(1):61-8.
54. Wizemann V, Wabel P, Chamney P, Zaluska W, Moissl U, Rode C, et al. The mortality risk of overhydration in haemodialysis patients. *Nephrol Dial Transplant* 2009;24(5):1574-9.
55. Chazot C, Wabel P, Chamney P, Moissl U, Wieskotten S, Wizemann V. Importance of normohydration for the long-term survival of haemodialysis patients. *Nephrol Dial Transplant* 2012;27(6):2404-10.