Research Article

Effects of clarified açaí (Euterpe oleracea Mart) supplementation on oxidative stress markers in hemodialysis patients: A randomized, controlled pilot study

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Abstract

Introduction: Patients with Chronic Kidney Disease (CKD) on Hemodialysis (HD) present a reduction of antioxidant enzymes levels and increased production of free radicals. Nutritional strategies have been used to mitigate the this exacerbated oxidative stress in these patients and supplementation with açai (Amazon fruit Euterpe oleracea) can be a good candidate to exert a protective effect against oxidative stress. The aim of this pilot study was to evaluate the effects of açai supplementation on oxidative stress markers in patients with CKD on HD.

Methods: Eighteen patients were randomized to receive 20 mL of clarified açai pulp juice three times a week for eight weeks (8 patients, 55.5 ± 4.9 years; BMI, 24.8 ± 2.5 Kg/m²) and to be the control group who did not receive any supplementation (10 patients, 56.1 ± 3.4 years, BMI, 25.2 ± 0.7 Kg/m²). Oxidative stress markers plasma levels such as Malondialdehyde (MDA), nitrite, Total Glutathione (TG), Catalase (CAT) and Glutathione Peroxidase (GPx) were evaluated before and after supplementation.

Results: MDA plasma levels were significant reduced from 6.8 ± 1.2 to 5.8 ± 0.9 nmol/mL in açai group (p=0.037), compared to no significant change in control group. The GSH total level presented slight increase after supplementation in açai group (320.7 ± 59.2 to 492.7 ± 84.8 nmol/mL, p=0.085). For Nitrite, CAT and GPx activity after eight weeks of supplementation with clarified açai (Euterpe oleracea Mart) pulp there was no change.

Conclusion: The consumption of clarified açai for eight weeks improved lipid peroxidation in HD patients. This result suggests the promising application of açai as an antioxidant food to patients with CKD on HD.

Introduction

Oxidative stress is a very common complication in Chronic Kidney Disease (CKD) patients and becomes more severe in the late stage of the disease, which is exacerbated by the Haemodialysis (HD) procedure [1-3].

Several nutritional strategies have focused on reducing oxidative stress in CKD patients. Among these interventions are the consumption of antioxidants naturally present in some foods whose range of physiological intake is considered safe and can be achieved through food intake [4,5].

Açai (Euterpe oleracea Mart), a typical fruit from the Eastern Amazon basin in Brazil, where is consumed daily by the general population of the northern region of Brazil, has expanded to other regions of the country, especially in recent years for its export to various countries [6,7].

Studies observed that açai fruit has a high antioxidant capacity and is able to bind to free radicals due the range of bioactive compounds, mainly polyphenols, α-tocopherol, anthocyanins and flavones [8-12]. Additionally, these bioactive compounds interact with various enzymes and transcription factors, resulting in protective effects against oxidative stress-induced damage [13-15].

In previous clinical trials with healthy individuals, açai pulp improved lipid profile, lipid peroxidation and free radicals balance, as well as increased levels of antioxidant enzymes [16,17]. Thus, our hypothesis is that açai supplementation in CKD patients may mitigate the oxidative stress, thus leading to clinical improvement. Therefore, the aim of this study is to evaluate the effects of two months of oral açai supplementation on oxidative stress markers in patients with CKD on Hemodialysis (HD).

Material and methods

Participants

Twenty-three HD patients (açai group: 11 patients and control group: 12 patients) from a hemodialysis Center in Belém, Pará, Brazil were eligible for the randomized controlled trial. Six patients in the açai group and four in the control group withdrew from the study. Therefore, completed the intervention eight patients comprised the açai group and ten the control group (Figure 1).

Patients with access via arteriovenous fistula in an upper extremity; a dialysis duration of 3-4 hours per session three times weekly on a blood flow rate of 350 mL/min to 400 mL/min; and at least a six month history of hemodialysis were included. Patients with autoimmune and infectious diseases, cancer or AIDS; with serum levels of phosphorus and potassium above 5.5 mg/dL; or who used catabolic drugs, antioxidant vitamin supplements or açai were excluded.

The study was performed according to the Ethical Norms for Research with Humans, Resolution 164/96. Approval was granted by the Ethics Committee of Núcleo de Medicina Tropical (No. 2.338.781). The procedures used in this study adhere to the tenets of the Declaration of Helsinki. All patients gave written informed consent to participate in the study and were instructed not to use any supplement during intervention.

Experimental design

The patients were manually allocated with a simple randomization method to the experimental group or the...
control group by a person who was not involved in the research. Patients received clarified açaí (*Euterpe oleracea Mart*) pulp juice that was lyophilized from 250 mL of açaí pulp juice resulting in 20 mL of açaí pulp juice diluted in 20 mL of water with the final volume of 40 mL. The açaí juice was given orally to the patients after hemodialysis sessions (3x/week) for two months, and the control group did not receive any supplementation. Both groups were advised not to consume açaí juice during the period of the research. Measures of biochemical, clinical, anthropometric, oxidative stress data were performed at baseline and after two months.

The routine biochemical parameters, albumin, creatinine, hemoglobin, potassium, phosphorus and Parathyroid Hormone (PTH), were quantified according to standard methods at the hemodialysis center.

**Clarified açaí (*Euterpe oleracea Mart*) pulp juice**

The samples of the açaí pulp juice with fresh fruits were lyophilized and pasteurized. The açaí pulp juice were clarified by a patented process licensed (PI 1003060-3). This process consists of the partial purification of the gross aqueous extract produced by the adsorption technique on macroporous resins that resulted an açaí pulp juice higher purity antioxidant extract with absence of lipids, proteins and fibers [18,19].

The phytochemical composition to evaluate metabolites with antioxidant activity was performed to measure total polyphenols determined by the Folin–Ciocalteu method with antioxidant activity was performed to measure total polyphenols determined by the Folin–Ciocalteu method and total anthocyanins by a pH differential method [18]. Potassium mineral determination was quantified using an atomic absorption spectrophotometer (Thermo, ICE3000) and expressed in mg/L.

**Anthropometric parameters and dietary intake**

For the anthropometric evaluation, the body mass index (BMI) was calculated as weight divided by height squared, and then the nutritional status was classified according to the World Health Organization [20].

Food intake was determined by a 24-hour food recall administered by the researchers on three different days – a hemodialysis day, a day without hemodialysis and a weekend day – throughout the experimental period. The quantitative analysis of energy and macronutrient intake were calculated by the software Avanutri & Nutrição, 4.0 version (Três Rios, Brazil).

**Collection and separation of blood**

The blood was collected by venipuncture in the antecubital region with a vacuum system in tubes containing ethylenediaminetetraacetic acid (EDTA) as an anticoagulant, before the HD session in earlier in the week. Plasma was separated from blood by centrifugation at 3000 x g for 15 minutes at 4 °C, extracted with an automatic pipette, packed in 1.5 mL tubes, and stored at -80°C until analysis.

**Determination of thiobarbituric acid-reactive substances (TBARS)**

This test indicates estimated by the presence of products derived from lipid peroxidation, mainly malonaldehyde (MDA) using the Ohkawa et al (1979) method [21]. The absorbance was measured in a spectrophotometer at 532 nm (BIO-RAD Model 450 Microplate Reader, Hercules, CA, USA).

**Nitrite quantification**

Plasma nitrite, final product of nitric oxide (NO), was evaluated by the Griess Reagent method [22]. The samples were analyzed by in a spectrophotometer at 540 nm (BIO-RAD Model 450 Microplate Reader, Hercules, CA, USA).

**Total Glutathione (TG)**

The detection of total glutathione (GSH and GSSG) [23] was conducted by GSH oxidation by the 5,5-dithio-bis (2-nitrobenzoic acid) reagent, and the GSSG formed was recycled to GSH by glutathione reductase in the presence of NADPH. The absorbance was measured in a spectrophotometer at 412 nm (BIO-RAD Model 450 Microplate Reader, Hercules, CA, USA).

**Catalase activity and glutathione peroxidase**

The evaluation of catalase (CAT) activity was based on the reaction of the enzyme with methanol in the presence of an optimized concentration of H₂O₂ [24]. The analysis was performed by in a spectrophotometer at 540 nm (BIO-RAD Model 450 Microplate Reader, Hercules, CA, USA). Glutathione peroxidase (GPx1) activity in serum was determined by kit, was measured indirectly by a coupled reaction with glutathione reductase. The absorbance was measured with a spectrophotometer at 340 nm (Synergy H1M, Winooski, VT, USA).

**Statistical analysis**

Data were expressed as the mean and Standard Deviation (SD). The normal distribution of the data was verified by the Shapiro–Wilks test. The majority of the variables were normally distributed, except for the kinetic index of dialysis adequacy (Kt/V), parathyroid hormone and CAT activity.

The F test was used on independent parametric variables, and since all tests showed similar variations among the groups, the nonmatched Student’s t-test was applied. The Mann–Whitney test was used for independent nonparametric and ordinal variables, and the chi–squared test was applied to independent and dichotomous variables. The tests were performed using the statistical package Stata with the confidence values set to 95% (p<0.05).

**Results**

The clarified açaí pulp juice offered had total polyphenol content was 1013.2 mg equivalent gallic acid (GAE)/100 mL and total anthocyanins were quantified at 272.1 mg/100 mL, while potassium was quantified at 1174.69 mg/L.
Table 1 shows demographic and biochemical parameters in Açai group (3 males) and Control group (9 males) at baseline moment and Table 2 shows the food intake profile in both groups.

Table 3 shows the effects of supplementation with clarified açai (Euterpe oleracea Mart.) pulp juice on the MDA, nitrite and GSH total level, CAT and GPx activity before and after 2 months in HD patients. The results showed a significant reduction in plasma MDA levels and a slight increase in the total GSH levels after supplementation.

Discussion

During the last decade, açai has been studied because of its richness in bioactive compounds that exert effects on the balance of oxidative stress\(^1\). In hemodialysis patients, antioxidant–rich foods seem to have a protective effect on oxidative stress and cardiovascular damage, which is responsible for the high mortality of this group [25–27]. We found with the supplementation of clarified açai pulp juice for two months, a decrease in plasma MDA levels in HD patients, which may have benefits in improving oxidative stress. This study is the first work describing the effects of intervention with clarified açai (ideal model to evaluate the impact of isolated phenolic components and with high antioxidant capacity detected by the DPPH method) in patients with CKD on HD\(^2\). Compared to other foods with antioxidant capacity, açai has the highest content of total polyphenols (279.3 mg 100g\(^{-1}\)) and antioxidant capacity of 26μM Trolox g\(^{-1}\) than apple, blueberry, raspberry, blackberry and cranberry [28–30]. A pilot clinical study randomized, double-blind, placebo-controlled of phase I in healthy individuals, showed that the ingestion of 120 mL juice containing a mixture of fruits and berries including predominantly açai, increased significantly the serum antioxidant capacity at one hour and two hours after consumption as well as reduced the lipid peroxidation [31].

To evaluate oxidative stress, MDA was used as a lipid peroxidation marker, whose reduced plasma levels is associated with progression of CKD stages [32]. Our pioneering study showed, even as a pilot study, results similar to those found with supplementation of red fruit juice with a slightly larger sample size of 21 patients on HD and shorter intervention time for two months [33]. Similar result was found with supplementation pomegranate juice intake in 101 HD patients for one year [34]. In animal model of renovascular hypertension, the extract of açai showed the highest content of total polyphenols (279.3 mg 100g\(^{-1}\)) and antioxidant capacity detected by the DPPH method) in patients with CKD on HD\(^1\). Phenolic components and with high antioxidant capacity [35]. The mechanism involved appears to be related to the anthocyanins that have an important role on protection in the cells of oxidative damage, this probably involves its action as a reduction in ROS produced in polymorphonuclear cells, and reduced migration of pro-inflammatories [31]. However, it is still uncertain if lipid peroxidation may not only be the cause, but also the result of cell damage, since progression of CKD stages is associated with the value of MDA and increased consumption of antioxidant enzymes [36,37].

### Table 1: Demographic and biochemical profile in Açai and Control Groups.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Açai Group (n=8)</th>
<th>Control Group (n=10)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y), means ± SD</td>
<td>55.5± 4.9</td>
<td>56.1± 3.4</td>
<td>0.919</td>
</tr>
<tr>
<td>Diabetic, n (%)</td>
<td>6 (75)</td>
<td>2 (20)</td>
<td>0.054</td>
</tr>
<tr>
<td>Hypertensive, n (%)</td>
<td>8 (100)</td>
<td>7 (70)</td>
<td>0.216</td>
</tr>
<tr>
<td>Dialysis vintage (mo)</td>
<td>50.3 ± 11.3</td>
<td>53.8± 10.1</td>
<td>0.824</td>
</tr>
<tr>
<td>Kt/V</td>
<td>1.3 (1.3-1.4)</td>
<td>1.2 (1.1-1.8)</td>
<td>0.082</td>
</tr>
<tr>
<td>BMI (kg/m(^2))</td>
<td>24.8± 2.5</td>
<td>25.2± 20.7</td>
<td>0.901</td>
</tr>
<tr>
<td>Albumin (g/dL)</td>
<td>4.2 ± 0.08</td>
<td>4.3 ± 0.07</td>
<td>0.114</td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>9.7 ± 0.8</td>
<td>11.3± 0.5</td>
<td>0.124</td>
</tr>
<tr>
<td>Potassium (mg/dL)</td>
<td>5.0 ± 0.4</td>
<td>4.8 ± 0.4</td>
<td>0.390</td>
</tr>
<tr>
<td>Phosphorus (mg/dL)</td>
<td>4.0 ± 0.3</td>
<td>4.9± 0.2</td>
<td>0.044</td>
</tr>
<tr>
<td><strong>Biochemical parameters</strong></td>
<td></td>
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<tr>
<td><strong>Albumin (g/dL)</strong></td>
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<td><strong>Hemoglobin (g/dL)</strong></td>
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<td><strong>Potassium (mg/dL)</strong></td>
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<td><strong>Phosphorus (mg/dL)</strong></td>
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</tbody>
</table>

BMI: body mass index. Kt/V: kinetic index of dialysis adequacy.

The p value is representative of the intergroup analysis. Data are presented as the mean ± SD or as median and confidence interval.

### Table 2: Dietary intake baseline analysis in Açai and Control Groups.

<table>
<thead>
<tr>
<th>Estimated dietary intake</th>
<th>Açai Group (n=8)</th>
<th>Control Group (n=10)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total energy (kcal/day)</td>
<td>1511 ± 473</td>
<td>1495 ± 662</td>
<td>0.955</td>
</tr>
<tr>
<td>Carbohydrate (g/day)</td>
<td>155 ± 51</td>
<td>176 ± 89</td>
<td>0.859</td>
</tr>
<tr>
<td>Total lipids (g/day)</td>
<td>48 ± 24</td>
<td>63 ± 25</td>
<td>0.220</td>
</tr>
<tr>
<td>Protein (g/day)</td>
<td>64 ± 22</td>
<td>73 ± 27</td>
<td>0.423</td>
</tr>
</tbody>
</table>

The p value is representative of the intergroup analysis. Data are presented as the mean ± SD.

### Table 3: Oxidative stress profile before and after 8 weeks of supplementation with clarified açai (Euterpe oleracea) juice in HD patients.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Açai Group (n=8)</th>
<th>Control Group (n=10)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>MDA (nmol/mL)</strong></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td><strong>Nitrite (μmol/L)</strong></td>
<td></td>
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<tr>
<td><strong>TG (nmol/mL)</strong></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td><strong>CAT (mmol/min/mL)</strong></td>
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<tr>
<td><strong>GPx activity (nmol/min/mL)</strong></td>
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</tbody>
</table>

Data are presented as the mean ± SD or as median and confidence interval.

The p value is representative of the intragroup analysis; values were tested with paired sample T-tests.

MDA: Malondialdehyde; TG: Total glutathione; CAT: Catalase; GPx: glutathione peroxidase

Furthermore, it is inaccurate to assume a dose–time effect due to the low bioavailability of anthocyanins in organisms and the associated absorption factors that lead to synergistic açai components and metabolite effects because they induce intestinal microbiota modification [38].

Our work suggests that supplementation with açai may be beneficial in protecting against damage caused by oxidative stress. Moreover, quantifying açai recommendations is still needed because dietary and pharmacological interaction factors are unknown [38].

As limiting factor of this work, the small size of the sample, the absence of the evaluation of the antioxidant capacity and the total phenolic compounds plasma levels. However, this was the first randomized pilot trial that obtained time–response association between clarified açai juice and reduction in lipid peroxidation in HD patients.

Conclusion

In conclusion, despite our study having a small sample size, the consumption of clarified açai for eight weeks it seems improved lipid peroxidation in HD patients. This result indicates the promising application of açai as an antioxidant action in patients with CKD. Thus, it is desirable outcomes that polyphenol–rich interventions may inhibit or slow down oxidative stress in hemodialysis patients. From this finding, future studies in larger populations are needed to verify how much the açai is able to assist in combating the sequela of free radicals.

References


