Coconut Water-Based Probiotic Drink Proposal: Evaluation of Microbiological Stability and Lactic Acid Estimation

Ana Beatriz Praia*, Gilson Celso Albuquerque Chagas Júnior2, Adalgisa Gabriela dos Santos Guimarães1, Flávia Lopes Rodrigues1 and Nelson Rosa Ferreira1,3*

1Faculty of Nutrition / Federal University of Pará, Brazil
2Graduate Program in Food Science and Technology / Federal University of Pará, Brazil
3Faculty of Food Engineering / Federal University of Pará, Brazil

Abstract

Probiotics are in high demand for their role as a health promoter, with lactose-fermented foods being the main source of getting them. However, the options for probiotics are lower for a group of consumers who are allergic or lactose intolerant. As an alternative to a non-dairy product, this work aims to propose the formulation of a low-cost drink with probiotic characteristics based on coconut water, free from lactose and fermented by Lactobacillus casei shirota. Two products were made: one with packaged coconut water and the other with fresh coconut water. The inoculum was obtained from a commercial fermented drink (Yakult®). In the first stage (fermentation), the total cultivation time was 48h; however, the most suitable time was 12h at 36ºC for both cultures, with monitoring of pH, total acidity and cell concentration. Cultivations were carried out in duplicates with a repetition of the process for each product. After the first stage, the second stage (microbiological stability) at refrigeration temperature (5°C to 8°C) was started. The total refrigeration time was 120h; however, the most suitable time was 72h for both drinks. The estimate of lactic acid production was investigated using infrared spectrometry with Attenuated Total Reflectance (FTIR-ATR). It was possible to observe specific bands of carboxylic acids. The results obtained were promising and show potential to produce probiotic non-dairy drinks with inoculum and low-cost substrate.

Keywords: Coconut water; Fermentation; Functional foods; Lactobacillus; Probiotics

Introduction

Chronic-degenerative diseases have been the main cause of death in the world and controlling the risk factors for these diseases is a way to increase life expectancy. Aspects such as the practice of physical activities and a balanced diet are essential for the treatment and prevention of such diseases, contributing to the maintenance of health [1]. In this context, there is an increase in interest in functional foods, to get the beneficial, metabolic and physiological effects that these can cause.

Probiotics are gram-positive bacteria, defined internationally as “live microorganisms that, when administered in adequate amounts, confer a health benefit on the host” [2]. Probiotics work to maintain healthy intestinal function in the prevention and treatment of diarrhea and food allergies, relief of lactose intolerance and normalization of intestinal transit [3]; decreased serum cholesterol, reduced plasma LDL levels and glucose homeostasis and in the immune function, regulating the defense system [3-5].

Allergies and intolerances related to milk are the main causes of restricting the consumption of dairy products. Allergies and intolerances are manifested through the biochemical incapacity of the organism to digest, absorb or metabolize a specific component. Lactose intolerance is characterized by the inability to digest lactose due to the lack or decrease of the enzyme lactase [6].

Bearing in mind that the most common form of access to probiotics is through fermented milk and yogurts, people who are lactose intolerant cannot consume this food and enjoy its benefits. It is predicted that about 75% of the population, when they reach adulthood, develop this intolerance, because of the loss of the ability to digest lactose [7].

Probiotic foods must meet safety and functionality criteria, which include having no adverse effects and not being associated with infectious diseases, besides containing a microorganism that can survive, maintain metabolic activity and grow at the destination site [5].

Coconut water is the liquid part of the coconut fruit, is low in calories, has a sweet taste and its composition contains mineral salts, vitamins, amino acids and carbohydrates. Coconut water acts as an excellent natural isotonic and among the functional properties attributed is rehydration and electrolyte replacement [8-10].

Some studies show the development of probiotic drinks with coconut water [11-15]. None of these studies show the use of a low-cost...
inoculum and using two types of coconut water (industrialized and fresh). The results got were satisfactory and showed the probiotic potential of the planned drinks.

**Materials and Methods**

**Probiotic material and coconut water**

The probiotic bacteria used, *Lactobacillus casei shirota*, were obtained from a commercial fermented milk purchased under conditions suitable for consumption and kept under refrigeration (4°C) until the moment of use. The counting of lactic bacteria was performed on Man Rogosa Shape (MRS) Agar (Sigma®) and incubated at 36°C for 72h. The result was expressed in Colony Forming Units per mL of sample (CFU/mL).

The substrate used for fermentation had as raw material only coconut water and for preparing the drink, two formulations were initially used: one with industrialized coconut water and the other with fresh coconut water.

The industrialized coconut water showed sucrose (less than 1%) and antioxidant sodium metabisulfite (manufacturer’s data). The commercial condition was sterile (Ultra High Temperature-UHT) in 200mL Tetra Pak® packages. The entire contents were removed from the original packaging and distributed aseptically in sterile glass containers.

Fresh coconut water was obtained from mature coconuts (*Cocos nucifera*) purchased at a fair in Belém-PA (Brazil). The coconuts were washed in running water and punctured. The water was removed, also removing suspended particles. The distribution was carried out in duplicate in glass containers (200mL/each) and autoclaved for 15 minutes at 121°C.

**Inoculum addition**

The inoculation temperature in fresh coconut water (*W*) and industrialized coconut water (*Wf*) was stabilized at 38°C. The initial pH was 4.6 and 4.5 for *W* and *Wf* respectively. The pH was adjusted to 6.0 and subsequent inoculation (3% v/v) of fermented commercial milk was performed concerning the volume of coconut water. Cultivation occurred in static mode for 48h at 36°C.

**Determination of fermentation time**

During the 48h period of cultivation, microbial growth and physical-chemical characteristics (pH and total acidity) were evaluated, according to the method of IAL [16]. The quantification of lactic acid bacteria was performed as described in APHA [17]. The serial dilution technique with peptone water (0.1% w/v) was used to count cells in Man Rogosa Sharpe (MRS) Agar (Sigma®) and incubated at 36°C at 72h. The result was expressed in Colony Forming Units per mL of sample (CFU/mL). From these results, the best fermentation times for each substrate were defined.

**Evaluation of microbiological stability**

In the assessment of microbiological stability, the same conditions were repeated for *W* and *Wf* and now the fermentation time was established as a function of the minimum cell concentration values, as previously described. After the fermentation step, the material was stored under refrigeration (5°C to 10°C) for 120 hours. During this period, analyzes of pH, acidity and microbiological count were performed at 24-hour intervals.

**Microbiological analysis**

Analysis of total coliforms, thermotolerant coliforms (45°C), *Salmonella* sp and mold and yeasts count were carried out before fermentation, after fermentation and after refrigeration [17].

**Estimation of lactic acid**

The presence of lactic acid was estimated by spectroscopy in the infrared region with the Fourier transform and Attenuated Total Reflectance (FTIR-ATR) (Agilent Carry 360) in the spectral region from 4000 to 600cm⁻¹ with a measurement of 4cm⁻¹ and 32 scans. A liquid sample (fermented medium) was used without previous treatment. The spectrometric baseline fitting was performed by subtracting the effects of the water bands.

**Results and Discussion**

**Bacterial growth and fermentation time**

Quantification of lactic acid bacteria in commercial fermented milk was performed using quantification on MRS agar. The result obtained was in the order of 10⁹ CFU/mL. An inoculum of 3% (v/v) was added, corresponding to 6mL of fermented milk concerning coconut waters (200mL), where a theoretical concentration of 10⁹ CFU/mL was obtained. Figure 1 shows an enlargement of the lactobacillus (1000 X) present in fermented milk.
values in the order of $10^8$ of cell concentration were reached, which shows the probiotic potential of the product in both types of substrates (Table 1).

After analyzing cell growth, it was observed that the fermentation time for probiotic viability in $W_f$ was 12h, while in $W_i$ it was 6h. In the time after 12h, only $W_i$ did not reach the value of $10^9$ CFU/mL until 48h of culture. Thermal treatment was carried out in fresh coconut water to eliminate fermentative micro-organisms, thus guaranteeing a better evaluation of growth with the inoculum, despite the possibility of nutritional loss because of the heating of the raw material. Even with the heat treatment, a better adaptation of the lactobacillus in $W_i$ was observed in principle. The response of microbial growth in $W_f$ was considered satisfactory, considering that $W_f$ undergoes heat treatment at high temperatures ($130^\circ$C to $150^\circ$C) for a short time (2s to 4s) and rapid cooling (below $32^\circ$C). It is believed that preservatives may have caused some inhibitory effect concerning the growth of lactic acid bacteria. The beverages planned preliminarily showed similar sensory characteristics. Both had an odor like fermented milk and a whitish color (Figure 2).

![Figure 2](image)

Figure 2: Probiotic drink based on coconut water after 12 hours of fermentation at $36^\circ$C.

**Analysis of total coliforms, Salmonella sp. and molds and yeasts**

Microbiological analysis carried out in coconut waters before the fermentation showed the absence of total and thermotolerant coliforms in MPN/mL. Microbiological analysis performed after fermentation showed the absence of total coliforms and thermotolerant coliforms and the absence of Salmonella sp. Regarding the product after refrigeration (storage), the absence of molds and yeasts was observed. These results showed that the entire process remained within microbiological standards.

**Total acidity and pH analysis**

The initial pH of coconut waters was corrected to 6.0. This procedure aimed to promote the most suitable pH range for the development of lactic acid bacteria, which can preferably grow at slightly acidic to neutral pH (5.5 to 6.2) and at temperatures between $30^\circ$C to $40^\circ$C [19,20].

The results of pH in $W_f$ showed a variation between 5.4 and 5.7 in the 48h period. The acidity value decreased approximately 25% after 6 hours of cultivation and remained stable until 36 hours. After 48 hours, a return to initial acidity was observed. The pH and acidity values show important information about hydrogen concentration and organic acids in the medium. In this case, it is possible to deduce that the reduction in pH and the increase in acidity may be mostly related to the presence of lactic acid. Ramos et al. [21], formulated three fermented milk drinks with cajá flavor (with 40% whey and 15%, 20% and 25% of fruit pulp) with an inoculum of Lactobacillus acidophilus, Bifidobacterium and Streptococcus thermophilus; fermented at $42^\circ$C for 4h. Thus, as in this work, practically constant results of pH and acidity were observed (Figure 3).

![Figure 3](image)

Figure 3: Profile of pH and total acidity with industrialized coconut water ($W_i$).

Otherwise, $W_i$ showed different results. A marked decrease in pH was observed after the initial 24h of fermentation and a more expressive increase in acidity. The total acidity value at the end of the fermentation was 3.2 times higher when compared to the beginning. Based on this observation, it is possible to infer that cultivation in $W_i$ has a greater potential for lactic acid production when compared to $W_f$ (Figure 4). Giri et al. [11], showed similar results in preparing a coconut water drink fermented with Lactobacillus casei L4. In this study, a decrease in pH from 6.5 to 3.32 was observed after 48 hours of fermentation, with a consequent increase in acidity.

The two coconut water substrates had different profiles in terms of pH and acidity. The increase in acidity and consequent decrease in pH observed in the $W_i$ suggests a better adaptation of lactic acid bacteria in this condition. It is believed that this occurs because of the fresh coconut water presenting nutritional characteristics more suitable for microbiological growth.

<table>
<thead>
<tr>
<th>Formulation</th>
<th>Incubation Time (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
</tr>
<tr>
<td>$W_i$</td>
<td>$4.15\times 10^7 \pm 1.63\times 10^7$</td>
</tr>
<tr>
<td>$W_f$</td>
<td>$5.4\times 10^5 \pm 2.25\times 10^9$</td>
</tr>
</tbody>
</table>

Table 1: Concentration of cells in CFU/mL by incubation time.

Note: $W_i$: Industrialized coconut water; $W_f$: Fresh coconut water.
The inverse relationship of pH and total acidity with bacterial growth, suggests a proliferation of lactic acid bacteria, which consume sugars of the production of lactic acid [22]. It is known that *Lactobacillus casei shirota* can ferment different sugars, including galactose, lactose, D-lactose, D-fructose, sorbitol, and others [23].

The bacteria use nutrients from the substrate for fermentation and multiplication. These metabolic activities release more lactic acid, this decreases the pH and increases the acidity of the final product [23]. About industrialized coconut water, bacterial growth occurred, however over a longer time. It is possible that industrial processing has altered the nutritional profile of this substrate. The addition of antioxidants in industrialized coconut water may also have influenced pH stability and acidity.

Although the two formulations showed adequate results regarding the probiotic potential (bacterial growth greater than 10⁸), the fermentation with *Wf* presented the desired response more efficiently. The fermentation with *Wf* reached the appropriate microbial concentration in pH and practically constant acidity. Otherwise, in *Wi* reached the range of 10⁵-10⁹ CFU/mL (Table 1) and showed increased values of acidity during the 48h of analysis, with a consequent decrease in pH.

Considering that at 12h the two formulations showed probiotic potential, it was decided to use this time as a fermentation standard for both formulations. The establishment of the standard time was important to test the microbiological stability in refrigeration. Thus, there was no need to carry out a long time for the desired microbial growth during the fermentation phase.

**Microbiological stability under refrigeration**

The quantification of viable cells in *Wf* shower less variation in the refrigeration period (5°C to 8°C) between times 24 to 96h. In this condition, there was the maintenance in the ideal bacterial growth range for probiotics (10⁵-10⁹) for a longer time (96h) during storage under refrigeration, in comparison with *Wi* (72h) (Figure 5).

The drink with *Wf* maintained an average pH equal to 6.2 ± 0.3 up to 72 hours of refrigeration. After this period, there was a decrease to 5.1 ± 0.2. The *Wf* drink showed pH stability of up to 24 hours. Between 24 hours and 120 hours, there was a progressive decrease to pH equal to 4.5. It is believed that the pH decrease in both drinks must be related to the accumulation of lactic acid produced during fermentation and later during refrigeration, as has been described in other studies [13,18] (Figure 6).

**Lactic acid production**

The samples of both drinks were analyzed in the middle infrared region, observing the wavenumbers in 1720 cm⁻¹ (C=O stretch of carboxylic acids), 1230 cm⁻¹ (C-O stretch of carboxylic acids) and 1128 cm⁻¹ (C-O stretch of esters). Figure 7 shows two infrared spectrum profiles for drinks made in 120 hours of storage. For the microbiological characteristic of the inoculum used in this study (lactobacillus), it is possible to infer that the decrease in pH and increase in acidity is because of the production of lactic acid. Spectroscopy in the infrared region with Attenuated Reflectance (FTIR-ATR) is an excellent technique for organic analysis, because it is fast, does not require sample preparation and analytical standard. This mainly facilitates routine analysis in industries with an emphasis on a specific substance as in this study. Figure 7 shows the results of the infrared spectroscopy analysis, resulting from the spectral subtraction of the water molecule in the drinks. This procedure was necessary because of the intensity of the water bands that overlapped the characteristic
bands of the carboxylic acids in the samples of the drinks. Lactic acid is classified as a carboxylic acid, nominated by IUPAC as 2-hydroxypropanoic acid. In this way, it was possible to identify two characteristic bands of carboxylic acids (1720cm⁻¹ and 1230cm⁻¹) [24]. A lower transmittance value of 1720cm⁻¹ for the Wf drink may show a higher concentration of lactic acid. This result is corroborated by figure 6, which shows a lower pH value in 120h when compared to Wf. In 1128cm⁻¹, it is possible to observe characteristic ester bands, this is because lactic acid undergoes a condensation reaction even at low concentrations [25]. The lower transmittance value for the drink with Wf shows a higher concentration of condensed lactic acid molecules. This is believed to be due to the greater stability of lactic acid polymers at less acidic pH [26].

Figure 7: Infrared spectrum: Drink with industrialized coconut water (red) and drink with fresh coconut water (blue).

### Conclusion

When we compare the two formulations, fresh coconut water substrate with industrial coconut water substrate, we noticed that the two formulations showed adequate results about the probiotic potential after 12 hours of fermentation. Complementary, we can consider that the adequate cooling time is 72h for both formulations at temperatures between 5°C to 8°C.

Coconut water proved to be an excellent substrate, both for its adequate nutritional profile and its ease of acquisition, besides allowing the growth of *L. casei shirota*. Despite being initial data, there is a good prospect of future applicability in the development of a low-cost product that can also be safely prepared in homes, as it is with the production of homemade yogurts.

The drinks proposed in this study can serve an increasingly growing market niche of people who have lactose intolerance or allergy, or even those who do not consume products of animal origin.

### Declaration of Conflicting Interests

The authors declare that there is no conflict of interest.

### Acknowledgment

The authors would like to thank professors Alessandra Santos Lopes (FEA/UFPA) and Luiza Helena Meller da Silva (FEA/UFPA) and also the Federal University of Pará (UFPA) for providing the infrastructure.

### References


