Physicochemical characterization and antimicrobial activity in novel systems containing buriti oil and structured lipids nanoemulsions

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\textbf{A B S T R A C T}

Buriti oil nanoemulsions were prepared using non-interesterified buriti oil or buriti oil interesterified for 6 or 24 h (NBO, NBO6h, and NBO24h), respectively. The aim was to investigate the effects of interesterified oils on the physicochemical and biological properties of nanoemulsions. Samples were stored at 4 and 25 °C for 30 days, and their physicochemical properties and biological activities were evaluated. The mean droplet diameter of nanoemulsions ranged from 196 to 270 nm. NBO24h had the smallest droplet size and was the most stable during the storage period. Furthermore, NBO24h demonstrating the good oxidative stability, had a high antioxidant capacity, and was less susceptible to droplet aggregation. NBO and NBO24 h had similar biological activity against Gram-negative bacteria (Escherichia coli O157: H7); bacterial growth was inhibited by at least 60% at 3.12 mg mL\textsuperscript{-1}. The nanoemulsions have interesting properties for the production of pharmaceutical, cosmetic, and food formulations with antimicrobial activity.

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1. Introduction

The buriti palm (Mauritia flexuosa L.f.) occupies extensive areas in Brazil, especially in the central part of the country and in the southern Amazon lowlands. Buriti oil, extracted from the fruit of the buriti palm, is traditionally used in cooking and for medicinal purposes. The unprocessed oil has considerable levels of minor compounds, such as \( \beta \)-carotene and tocopherols, which help reduce total and low-density lipoprotein cholesterol and lower the risk of coronary artery disease [1,2]. The functional properties of this unconventional vegetable oil have been extensively investigated: buriti oil is rich in oleic acid and other compounds that exert positive health effects, shows antimicrobial activity, and has potential applications in the cosmetic, pharmaceutical, and food industries [3].

Chemical or enzymatic interesterification reactions can be used for the synthesis of structured lipids. Structured lipids are triacylglycerols (TAGs) that have been modified in their fatty acid composition or restructured in their positional distribution of fatty acids in the glycerol backbone [4] to impart desirable physicochemical, nutritional, and functional properties for food applications or other purposes. Structured lipids are also referred to as the “new generation of nutraceutical fats” [5]. Previous work by our research group has shown that enzymatic interesterification can improve physicochemical and biological properties of buriti oil. Enzymatic interesterification is widely used to synthesize oils and fats, as it has potential health and nutrition benefits [6].

Oils and fats find many commercial applications in the form of emulsions. Nanoemulsions are dispersion or delivery systems that are composed of at least two immiscible liquids, one of which is present as nanosized droplets [7,8]. These systems are regarded as valuable alternatives for encapsulating bioactive lipophilic compounds of hydrophobic nature. Oil-in-water (O/W) colloidal dispersions are often used for this purpose. Nanoemulsions show advantages over conventional emulsions because of their smaller droplet size and greater stability against gravitational separation. Previous studies confirmed the advantages of nanoemulsions in the food industry, showing that these systems can be used to improve the bioavailability of complex food matrices and extend product shelflife [9,10]. Nanoemulsions are prepared by either low- or high-energy methods [11]. Examples of the latter include high-pressure homogenization, microfluidization, and ultrasonication [12].

The choice of the emulsifier is of critical importance for producing nanoemulsions because this component facilitates emulsification and promotes physical stability [13]. There are
various types of food-grade emulsifiers that are used to produce nanoemulsions based products in the food industry, such as gum arabic and small-molecule surfactants (e.g., Tweens) [14]. Tween 80 is one of the most used synthetic emulsifiers for the preparation of nano and conventional emulsions because of its good emulsifying properties [15,16]. The amphiphilic character of emulsifiers allows the entrapment of hydrophilic or hydrophobic bioactive compounds, improving their dissolution in the emulsion system.

Results obtained by our research group in a previous study provided the fundamental perspective that enzymatic interesterification of Amazonian oils yields structured lipids with interesting characteristics and antimicrobial potential [3]. Thus, the aim of this study was to produce O/W nanoemulsions of non-interesterified and interesterified buriti oil and compare their physicochemical characteristics and biological potential.

2. Materials and methods

2.1. Materials

Buriti oil was purchased from Beraca (São Paulo, Brazil). Commercially purified and immobilized lipase (Lipozyme R⃝ TL IM) was purchased from Novozymes (São Paulo, Brazil). Tween 80 (polyoxyethylene sorbitan monooleate) was purchased from Dinâmica (São Paulo, Brazil). Nile red was purchased from Sigma–Aldrich (St. Louis, USA). Escherichia coli O157:H7 ATCC 11,775 was obtained from the Chemical, Biological, and Agricultural Pluridisciplinary Research Center (CPQBA) of the University of Campinas, Campinas, Brazil.

2.2. Methods

2.2.1. Preparation of nanoemulsions

Buriti oil was interesterified for 6 and 24 h, following the method of Speranza et al. [17]. O/W nanoemulsions with 10% oil phase were prepared according to Ozturk et al. [18]. Three different formulations were manufactured by varying the oil phase: nanoemulsions of buriti oil (NBO), nanoemulsions of buriti oil interesterified for 6 h (NBO6h), and nanoemulsions of buriti oil interesterified for 24 h (NBO24h). Coarse emulsions were homogenized in a high-pressure homogenizer (GEA Niro Soavi, Parma, Italy) for 3 cycles at 800 bar. After homogenization, 100 mL of each sample was stored in amber glass bottles and kept in the dark at 4 or 25 °C for 30 days.

2.2.2. Mean droplet size determination

The mean droplet diameter (d32) of nanoemulsions was measured using a static light-scattering system (Mastersizer 2000, Malvern Instruments Ltd., Malvern, UK). Aliquots of nanoemulsions were diluted with distilled water under constant stirring (1750 rpm) until an obscuration of 5% was achieved. Nanoemulsions were analyzed after 1, 15, and 30 days of storage at refrigeration (4 °C) or room temperature (25 °C). Each sample was analyzed in triplicate.

2.2.3. Zeta-potential measurements

The surface charge of nanoemulsions during the storage period (days 1, 15, and 30 at 4 and 25 °C) was determined using Instrument Zetasizer Nano-Z (Malvern Instruments Ltda., Malvern, UK). Measurements were carried out in triplicate.

2.2.4. Confocal laser scanning microscopy

Confocal laser scanning microscopy (CLSM) was carried out using a 63× objective lens (TCS SPS II C1 Digital Eclipse, Leica, Tokyo, Japan). Samples were stained with Nile red, a fat-soluble fluorescent dye, by mixing 100 mL of sample with 20 μL of 1 mg mL−1 Nile red in propylene glycol. Excitation and emission wavelengths were respectively 488 nm and 515 nm. Images were processed using LAS lite (Zeiss Inc., Toronto, Canada).

2.2.5. Oxidative stability

The thiobarbituric acid reactive species (TBARS) assay is based on the spectrophotometric measurement of the pink pigment produced by the reaction of thiobarbituric acid (TBA) with malondialdehyde (MDA) and other secondary lipid peroxidation products. The assay was performed by adding 100 μL of nanoemulsion and 1 mL of 2-TBA into screw cap tubes, stirring the mixture, and reading the absorbance at 532 nm [19]. TEP (1,1,3,3-tetraethoxypropane) was used to construct a standard curve.

2.2.6. Antioxidant assays

The ferric reducing antioxidant power (FRAP) assay was carried out according to Benzie and Strain [20] with modifications. A solution of TPTZ (2,4,6-tripyridyl-s-triazine) was prepared in 40 mM HCl, and a solution of 20 mM FeCl3·6H2O was prepared in distilled water. Nanoemulsions were diluted to 1:2, 1:3, and 1:5. Absorbance was measured at 595 nm using a UV–vis spectrophotometer. Antioxidant activity was expressed as μmol Trolox equivalents (TE) mL−1 nanoemulsion according to a standard curve. The modified oxygen radical absorbance capacity-fluorescein (ORAC-fl) assay was carried out following the method of Dávalos et al. [21]. ORAC values were calculated by the difference between the area under the fluorescence decay curve of the sample and that of the blank. A standard curve was constructed using Trolox, and results were expressed as μmol TE mL−1 nanoemulsion.

2.2.7. Polymorphism

The polymorphic form of nanoemulsion crystals was investigated by X-ray diffraction (XRD) according to the AOCS method Cj 2–95 [22]. The measurements were obtained with steps of 0.02° in 2θ acquisition time of 2 s, with scans of 15–40°, at 25 °C using a Philips PW 1710 diffractometer with Bragg–Brentano geometry (θ/2θ) and Cu Ka radiation (λ = 1.54178 Å; 40 kV; 30 mA). XRD. Polymorphic forms were identified by analyzing the short spacings of each crystal. Relative proportions of the different crystal types were estimated from the relative intensity of the short spacings. Data were analyzed using Origin (Origin Lab Corporation, Northampton, USA).

2.2.8. Antimicrobial activity

The minimum inhibitory concentration (MIC) of NBO and NBO24h against E. coli O157:H7 ATCC 11,775 was determined using the broth microdilution method NCCLS [23]. Briefly, 50 μL aliquots of each nanoemulsion (NBO and NBO24h) were sequentially diluted (25.0 - 0.39 mg mL−1) in a 96-well plate containing 100 μL of Mueller–Hinton Broth (Difco®). The final concentration of bacterial suspension was adjusted to 1.5 × 106 colony forming units (CFU) mL−1. Plates were incubated at 37 °C for 24 h. Chloramphenicol was used as positive control.

2.3. Statistical analysis

Data were submitted to analysis of variance (ANOVA) followed by Tukey’s test using Minitab v. 16 (Minitab Inc., State College, USA) to evaluate significant differences (p < 0.05) between mean values.

3. Results and discussion

3.1. Size distribution and zeta potential

Overall, nanoemulsions were relatively stable at 4 °C and had little or no variation in droplet size during storage. The results show that temperature had no significant effect on droplet size.
Data also suggest that structured nanoemulsions were more resistant to coalescence and gravitational separation, especially NBO24h, as the mean droplet size was smaller than 200 nm and no visible changes in phase separation were observed during storage [24]. Similar droplet diameters were observed in systems containing vitamin E, vitamin D, or lime oil [25]. Physicochemical changes in TAGs caused by the interesterification result in a predominance of unsaturated fatty acids at the sn-1 and sn-3 positions, leading to the formation of tri-unsaturated TAGs that are rich in oleic acid and thereby reduce the melting range of buriti oil. Furthermore, the substitution of palmitic acid for oleic acid at the sn-1 and sn-3 positions of TAG in structured buriti oil [26] might have contributed to the reduction in droplet size. The lower the melting range of a structured lipid, the lower its viscosity, which directly affects droplet size. Oleic acid has a certain degree of mobility provided by the presence of a double bond, and this may alter the crystalline arrangement and the density at which chains can be stacked [27] (Table 1).

Another important factor associated with the stability of droplet size during storage is the efficiency of homogenization, which depends on the physicochemical properties of the lipids that form the emulsion, as smaller droplets are formed when the lipid phase is less viscous. When oil droplets are more viscous, they are more difficult to be dissolled by the homogenizer because they may exit the break zone before deforming and rupturing [12]. Emulsifiers play an important role in the preparation of nanoemulsions because they reduce interfacial tension and facilitate the formation of small droplets during homogenization through adsorption of the oil–water interface. However, if the hydrophilic–lipophilic balance is too high, then the interfacial tension of the emulsifier may be too high to form fine oil droplets [28,29]. The high-energy homogenization process combined with the interesterification reaction yielded nanoemulsions that were more stable. Although no studies have investigated the effect of fatty acid position in TAGs on nanoemulsion stability, some authors evaluated the effect of oil type on stability and the behavior of different nanoemulsions during in vitro digestion [30,31]. After droplet formation, the stability of nanoemulsions depends on the oil phase composition, as oils differ in polarity and solubility [32]. A previous study showed that nanoemulsions containing small droplets (d<sub>32</sub> < 200 nm) may find practical application in pharmaceutical, food, and cosmetic industries because of their structure and easy manipulation [28].

The zeta potential is an indirect measure of the surface charge of oil droplets, which provides an indication of stability during storage. The stability of NBO, NBO6h, and NBO24h stored at 4 and 25 °C was analyzed for 30 days (Table 2). Surface charges ranged from −23.5 to −30.8 mV, which indicates that nanoemulsions would be stable against aggregation and coalescence during storage.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Storage period (days)</th>
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<tbody>
<tr>
<td>1</td>
<td>15</td>
</tr>
<tr>
<td>25 °C</td>
<td>25 °C</td>
</tr>
<tr>
<td>NBO</td>
<td>4.0 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>NBO6h</td>
<td>4.0 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>NBO24h</td>
<td>4.0 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
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<sup>a</sup>Means within a row followed by different letters differ significantly at p < 0.05.

### Table 2

<table>
<thead>
<tr>
<th>Sample</th>
<th>Storage period (days)</th>
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<tr>
<td>1</td>
<td>15</td>
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<tr>
<td>25 °C</td>
<td>25 °C</td>
</tr>
<tr>
<td>NBO</td>
<td>4.0 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>NBO6h</td>
<td>4.0 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>NBO24h</td>
<td>4.0 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
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</table>

<sup>a</sup>Means within a row followed by different letters differ significantly at p < 0.05.

### 3.2. Confocal laser scanning microscopy (CLSM) analysis

Confocal laser scanning microscopy observations allowed us to gain a better understanding of the microstructure of nanoemulsions and examine nanoscale characteristics that could not be inferred from droplet size results. CLSM images (Fig. 1) revealed that lipids were present as dispersed nanoparticles in suspension. NBO, NBO6h, and NBO24h (Fig. IB, ID, and IF, respectively) stored at room temperature (25 °C) displayed a very similar distribution pattern, with lipid droplets of less than 1 μm. However, it was difficult to distinguish individual droplets because of the resolution limit of the microscope. The number of relatively large droplets increased as the storage time increased, which suggests droplet aggregation and emulsion instability.

NBO, NBO6h, and NBO24h (Fig. IA, IC, and IE, respectively) stored at 4 °C showed greater susceptibility to droplet aggregation than nanoemulsions stored at room temperature, according to CLSM images. This phenomenon was more intense after 30 days of storage. Although we did not find a significant increase in droplet size, microstructural observations provided evidence that coalescence took place at refrigeration temperature. In contrast, nanoemulsions stored at 25 °C did not exhibit larger agglomerations with time, especially NBO6h and NBO24h, which were composed of structured lipids. These results can be explained by the fact that structured buriti oil has a higher content of unsaturated TAGs than its unstructured equivalent and thus has lower melting point and viscosity. These physicochemical properties influence droplet formation and stability [26].
Interestingly, nanoemulsions exhibited a specific aggregation behavior that may be related to polymorphic changes during storage. Such changes affect nanoemulsions properties, including crystal shape and particle morphology, as will be discussed in detail later. No studies reporting similar behaviors in structured lipid nanoemulsions were found in the literature. Some studies investigated the behavior of conventional emulsions or nanoemulsions by confocal microscopy using gastric digestion models [9,30].

As nanoemulsions produced with lipids interesterified for 6 or 24 h afforded similar results, NBO6h was excluded from the study and only NBO and NBO24h were subjected to further investigation.

### 3.3. Lipid peroxidation during storage (TBARS assay)

The oxidative stability of NBO and NBO24h was monitored during 30 days of storage by the TBARS method, used to measure the formation of secondary oxidation products. Results are shown in Table 3. A significant increase in TBARS was observed during storage for both NBO and NBO24h, indicating the occurrence of lipid oxidation. NBO and NBO24h had very similar ($p > 0.05$) TBARS levels on the first day of storage at room temperature. TBARS levels increased in NBO as the storage time increased, so that at the end of storage at 25 °C, NBO showed significantly higher TBARS than NBO24h. At the end of storage at 4 °C, NBO24h also had significantly lower TBARS than NBO. Droplet size and interfacial area influence lipid oxidation in O/W emulsions, as do various structural parameters and components [37]. Nanoemulsions have a large specific surface area because they are composed of small droplets [38]. Tween 80 is also known to play a role in the oxidation of emulsified lipids through the formation of hydroperoxides [39].

On the basis of the results presented above, we believe that enzymatic interesterification helps prevent lipid peroxidation in buriti oil nanoemulsions. In general, significant amounts of secondary oxidation compounds are formed when fatty acid chains containing three or more double bonds, different from linoleic acid or oleic acid, are involved [40].

Similar results were reported by Nejadmansouri et al. [37], who observed that the oxidation rate of nanoemulsions was greater at refrigeration temperature than at ambient temperature. Wang et al. [41] concluded that interesterification contributed positively to the stability of oils, retarding or preventing oxidation.

This study has important implications for the design and use of nanoemulsions as delivery systems. The present results show that, because of its smaller droplet size, NBO24h is less susceptible to the formation of secondary oxidation products throughout the storage period at both studied temperatures. It is known that the mechanism of lipid oxidation in O/W emulsions is very complex

### Table 3

Oxidative stability (TBARS levels in mg MDA mL$^{-1}$) of nanoemulsions of buriti oil (NBO) and structured buriti oil (NBO24 h) at days 1, 15, and 30 of storage at 4 or 25 °C.

<table>
<thead>
<tr>
<th>Storage temperature</th>
<th>NBO</th>
<th>NBO24h</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>day 1 (TBARS)</td>
<td>day 15 (TBARS)</td>
</tr>
<tr>
<td>4 °C</td>
<td>11.90 ± 0.24$^a$</td>
<td>5.67 ± 0.04$^a$</td>
</tr>
<tr>
<td>25 °C</td>
<td>10.29 ± 0.62$^a$</td>
<td>6.12 ± 1.13$^a$</td>
</tr>
</tbody>
</table>

$^a$, $^b$ Means within a row followed by different letters differ significantly at $p < 0.05$. 

Fig. 1. Confocal laser scanning micrographs of nanoemulsions of buriti oil (NBO) stored at 4 and 25 °C (IA and IB), nanoemulsions of buriti oil interesterified for 6 h (NBO6h) stored at 4 and 25 °C (IC and ID), and nanoemulsions of buriti oil interesterified for 24 h (NBO24 h) stored at 4 and 25 °C (IE and IF).
and differs from that which occurs in bulk lipid systems [42]. Because emulsified lipids are highly susceptible to oxidation, combined antioxidant strategies are required to delay oxidation reactions and improve shelf-life. This knowledge served as a basis for developing structured nanoemulsions of buriti oil: it has excellent antioxidant, functional, and biological properties [26,43]. Subsequent experiments were performed using NBO and NBO<sub>24h</sub> stored at room temperature, as samples were shown to be less stable at 4°C.

### 3.4. Antioxidant capacity of nanoemulsions

The FRAP assay, based on the electron transfer reaction between antioxidants and radicals [44], has been widely used to determine the antioxidant capacity of O/W emulsions and nanoemulsions. But because antioxidant capacity varies according to the radicals that participate in the reaction, it must be examined by more than one method. We analyzed the antioxidant activity of structured and non-structured buriti oil nanoemulsions by the FRAP and ORAC assays. Results are presented in Table 4. By the FRAP method, NBO had an antioxidant activity of 419.83 μmol mL<sup>-1</sup> and NBO<sub>24h</sub> of 611.38 μmol mL<sup>-1</sup>, that is, NBO<sub>24h</sub> exhibited an antioxidant capacity approximately 31% higher than that of NBO. Our data show that structured lipids in NBO<sub>24h</sub> were effective in donating electrons to stabilize free radicals, contributing to the antioxidant effect of the system.

Minor components of oils, such as tocopherols, carotenoids, phenolics, exert antioxidant effects through various chemical mechanisms, including the elimination of free radicals by electron transfer or hydrogen donation and chelation of transition metals [40]. Thus, the antioxidant activity of nanoemulsions of buriti oil and structured buriti oil is probably also due to the presence of carotenoids, mainly β-carotene [26]. In contrast, the presence of free fatty acids in emulsified oils is known to increase lipid oxidation, which is related to their charge in the oil–water interface, and to increase the negative charge of oil droplets, which may attract cationic transition metals such as Fe<sup>2+</sup> or Fe<sup>3+</sup> to the oil droplet surface [45,46]. Transition metals can originate from various sources, including process water, processing equipment, and oils and fats [37]. When present in the emulsion system, transition metals can catalyze oxidation reactions; in these cases, lipid oxidation cannot be prevented by a radical-scavenging antioxidant alone and a metal chelator is needed [37].

The ORAC assay measures the oxidative degradation of the molecule fluorescein, which when inhibited indicates antioxidant activity [47]. The ORAC value of NBO<sub>24h</sub> was 3640.56 μmol mL<sup>-1</sup> and of NBO was 3089.66 μmol mL<sup>-1</sup> (Table 4). NBO<sub>24h</sub> showed greater antioxidant activity than NBO by both FRAP and ORAC methods, which can be confirmed by further oxidative stability assays. To date, no other study has evaluated ORAC in nanoemulsions of structured lipids.

Previous studies have shown that the levels of phenolics of high antioxidant potential, can vary in buriti oil [48,49]. Thus, buriti oil emulsion systems may differ in composition and concentration of phytochemicals and may even contain compounds that interact in an antagonistic manner [43,50].

### 3.5. Crystal polymorphism in nanoemulsions

Crystallization processes occur according to the predominant form of crystals and have a great impact on functionality and application [51]. In this study, evidence was collected to show the influence of droplet size on the crystalline structural properties of nanoemulsions, especially in relation to the presence of TAG molecules, which can crystallize into different forms depending on processing methods. Polymorphism analysis of NBO and NBO<sub>24h</sub> by X-ray diffraction after 30 and 120 days of storage revealed the coexistence of β' and β polymorphs in samples. The short spacings and polymorphic forms of buriti oil, NBO, and NBO<sub>24h</sub> are shown in Table 5. The diffractogram of bulk buriti oil stored for 30 days at room temperature showed peaks at 4.4, 4.2, and 3.9 Å, attributed to the polymorphic forms β' and β. The diffractograms of NBO and NBO<sub>24h</sub> also showed a combination of the polymorphic forms β' and β, as evidenced by the peaks at 4.2 and 3.9 Å, corresponding to the polymorph β, and very strong peaks at 4.4 Å, attributed to β'.

Polymorphic behaviors, such as melting point, may be affected by the chain length of fatty acids in the TAG structure [52]. To avoid interaction with lipid droplets and facilitate crystallization, we performed these experiments using a low concentration of surfactant.

A study on emulsified blends of interesterified fats reported that crystallization (−10°C) in the β' form was probably due to changes in the configuration of TAGs, which showed an increase in unsaturated fatty acids in the sn−2 position [53]. In the present study, TAGs in buriti oil and interesterified buriti oil contained a high percentage of tri-unsaturated fatty acids, which cannot crystallize in a metastable form [26]. According to some researchers [51,54], the presence of minor lipid components, monoacylglycerol and diacylglycerol in the oil at concentrations above 5% (data not shown), causes an increase in the energy barrier for nucleation, leading to delayed crystallization and the formation of unstable polymers. A better understanding of polymorphic transformations can be gained by analyzing the X-ray diffraction results of nanoemulsions after 120 days of storage. Transition from the polymorphic form α to β occurs more rapidly during storage when the system is emulsified because of the small size of crystals in emulsified systems. The polymorphic form of a lipid was shown to influence the biological activity of hydrophobic drugs in emulsion systems [55], because the molecular packing of lipid crystals can expel the molecules from the crystalline matrix. Although uncharacteristic of emulsified crystalline material, some crystalline nanostructure have previously been found to improve emulsion stability.

### 3.6. Antimicrobial assay

Antimicrobial agents from natural sources are being extensively investigated for their potential application in the food and other industries. In this study, structured and unstructured buriti oil was used to produce nanoemulsions with a mean droplet diameter with approximately 200 nm. It was measured by reading the absorbance in the plate cavities with different concentrations of nanoemulsions and response expressed in (Fig. 2).

<table>
<thead>
<tr>
<th>Sample</th>
<th>Concentration range (mg mL&lt;sup&gt;-1&lt;/sup&gt;)</th>
<th>ORAC (μmol TE&lt;sup&gt;-&lt;/sup&gt; mL&lt;sup&gt;-1&lt;/sup&gt;)</th>
<th>FRAP (μmol TE&lt;sup&gt;-&lt;/sup&gt; mL&lt;sup&gt;-1&lt;/sup&gt;)</th>
</tr>
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<tbody>
<tr>
<td>NBO</td>
<td>0.20−0.05</td>
<td>3089.7 ± 360.40&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;</td>
<td>419.8 ± 7.30&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;</td>
</tr>
<tr>
<td>NBO&lt;sub&gt;24h&lt;/sub&gt;</td>
<td>0.20−0.05</td>
<td>3640.6 ± 235.25&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;</td>
<td>611.4 ± 47.53&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;&lt;/sup&gt;</td>
</tr>
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</table>

<sup>TE, Trolox equivalents. <sup>a</sup>b</sup> Means followed by different letters within a column differ significantly at p < 0.05 by Tukey test.
NBO inhibited bacterial growth by approximately 61% at 3.12 mg mL⁻¹ demonstrating the bacteriostatic effect. Complete inhibition of bacterial growth was not achieved (Fig. 2). The low concentration of unsaturated fatty acids, which have antimicrobial activity, might explain the low antimicrobial effect of the samples [50]. Nevertheless, the results suggest that NBO and NBO₂₄h have physicochemical characteristics that contribute to their biological activity. According to recent studies, droplet size can affect the antimicrobial activity of emulsions. Essential oil nanoemulsions induced faster inactivation of E. coli than conventional emulsions [55]. The antimicrobial activity is not only to the chemical activity of individual components but also to the structure of the O/W nanoemulsion system. Some studies have correlated droplet size with biological activity [56,57]. However, a conventional emulsion of buriti oil showed bactericidal and bacteriostatic activity against E. coli [3]. The different explanations proposed for the experimental observations were considered, but we were not able to compare our results, as no previous data on the antimicrobial activity of buriti oil nanoemulsions are available in the literature. The study provided relevant contributions to the development of value-added products from natural ingredients, such as vegetable oils. Interesterification positively influenced mean droplet size, and oxidative stability of nanoemulsions, exhibited good physical stability, with little evidence of phase separation or droplet growth during 30 days of storage at 25°C. Furthermore, interesterification enhanced the antioxidant activity and, to a lesser extent, the antimicrobial action of buriti oil nanoemulsions. Several challenges remain to be overcome in the development of structured nanoemulsions with industrial applications that provide controlled-release of bioactive compounds, good stability thermodynamically and antimicrobial effects.

Declaration of Competing Interest

None.

Acknowledgments

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