Antioxidants extraction from Pinhão (Araucaria angustifolia (Bertol.) Kuntze) coats and application to zein films

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A R T I C L E   I N F O

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Phenolic compounds
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Microstructure
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A B S T R A C T

Seeds from Araucaria angustifolia (Bertol.) Kuntze are consumed after cooking and their coats discarded. Both coats and the cooking water present phenolic compounds, which may be used to improve mechanical properties and provide antioxidant characteristics to films. The objective of this work was to obtain and pinhão coat extracts and to apply these polyphenolic-rich extracts in zein films. Phenolic compounds composition, extraction yield and antioxidant activity (DPPH, ABTS and FRAP) of the extracts were determined. The most abundant molecules present in the hydroethanolic extract were ( + )-catechin and an (epi)catechin dimer, whereas protocatechuic acid were predominant in the both cooking water and ethanolic extracts. Glass transition temperature of zein was not found in the extract-loaded films. Morphological changes were also caused by the presence of the extracts yielding smoother surfaces. The extracts added to zein films led to a three-fold increase in tensile strength (from 5.80 MPa to 17.65 MPa) and two-fold increase in the elongation at break (from 1.60% to 3.18%).

1. Introduction

Seeds produced by Araucaria angustifolia (Bert.) O. Kuntze (Paraná tree), referred to as “pinhão”, are formed by a resistant external tegument (shell or coat) composed of lignocellulosic material rich in tannins and an internal edible pulp (Bello-Pérez et al., 2006; Conforti & Lupano, 2008; Lima et al., 2007; Santos et al., 2013). Seeds are produced seasonally and typically consumed after boiled in water. Although pinhão production is not an organized culture (Conab, 2014), there is an increasing interest in the industrial development of pinhão based products, such as pickled pinhão (Riele Conservas, 2016; WWF, 2016) and beverages such as beer (Cervejaria Campos do Jordão, 2016). In 2007, Brazil generated about 10 ton of pinhão wastes (Lima et al., 2007).

Residual pinhão coats, which represent about 20% of the weight (Conforti & Lupano, 2008; Cordenunsi et al., 2004), has also been attracting interest due to its antioxidant properties and potential biological activity, da Mota et al. (2014) obtained methanolic extracts from the shell and pulp of A. angustifolia seeds and identified in both extracts the presence of molecules with free radicals-trapping ability (e.g. polyphenolic compounds, flavonoids and proanthocyanins). The biological activity of pinhão extracts was reported by da Silva et al. (2014) who investigated the efficacy of a pinhão coat extract (70% ethanol in water) to decrease the postprandial glycemic levels in rats after starch administration. Authors claimed that the extract may be potentially used to suppress postprandial hyperglycemia in diabetic patients due to its inhibitory properties. In the study developed by Oliveira et al. (2015), a pinhão coat extract proved to be an effective inhibitor of pancreatic lipase and to effectively decrease plasma triglyceride levels in mice after a load of olive oil. Also, Branco et al. (2015) demonstrated that an aqueous extract of A. angustifolia bracts presented selective cytotoxicity and pro-apoptotic activity in laryngeal carcinoma HEP-2 cells.

The application of phenolic compounds in film formulations to obtain antioxidant properties, or active packaging materials, has been widely studied. For instance, studies have been published on gelatin-
based films added with curcuma ethanol extract (Bintencourt, Fávaro-Trindade, Sobral, & Carvalho, 2014), low density polyethylene incorporated with rosemary extract and natural obtained from a brewery residual waste (Barbosa-Pereira, Aurrekoetxea, Angulo, Paseiro-Losada, & Cruz, 2014), starch-based films with propolis extract (De Araújo et al., 2015), and zein and chitosan matrices supplemented with phenolic compounds (ferulic and gallic acids, respectively) (Cheng, Wang, & Weng, 2015).

Zein films may serve as carriers for antioxidant compound in food packaging applications (Arcan & Yemencioglu, 2011; Forato, Britto, Scramin, Colnago, & Assis, 2013; Park et al., 2012). In zein films, for instance, the addition of gallic acid, p-hydroxy benzoic acid, ferulic acid, flavone, (+)-catechin, and quercetin (Arcan & Yemencioglu, 2011), butylated hydroxyanisole and butylated hydroxytoluene (Kleen, Paseiro-Losada, & Cruz, 2014), green tea extract (Lee, Lee, & Song, 2004), gallic acid (Neo et al., 2013), and ferulic acid and gallic acid (Cheng et al., 2015), have been reported. However, to the best of our knowledge polyphenolic compounds from pinhão extracts have not yet been applied in films formulation as a carrier of antioxidant substances. In such case, the actual impact of the extracts on the films properties must also be determined. Also, applications involving zein films are worth investigating since zein is an important co-product of maize starch production, as well as a by-product of the bioethanol industry.

In this context, the current work aimed to obtain and to characterize pinhão coat extracts as well as to apply these extracts as a source of natural polyphenolic compounds in zein films. Extracts were obtained from the cooking water extract and also by subsequent ethanolic and hydroethanolic extraction of the residual pinhão coats.

2. Materials and methods

2.1. Materials

Pinhão seeds were acquired from the local market in Brazil in June 2015. Zein, Folin–Ciocalteu reagent, gallic acid, 2,2-diphenyl-1-picrylhydrazyl (DPPH), ferric reducing/antioxidant power (FRAP) reagent (0.3 M acetate buffer, pH 3.6 and 10 mM 2,4,6-tris(2-pyridyl)-s-triazine (TPTZ)), Trolox (6-hydroxy-2,5,7,8-tetramethylenon-2-carboxylic acid), ABTS (2,2’-azinobis [3-ethylbenzothiazoline-6-sulfonic acid) diammonium salt) and potassium persulfate (dipotassium peroxydisulfate) were purchased from Sigma–Aldrich. Glycerol, calcium carbonate and ethanol (Vetec) were analytical grade. Acetonitrile (99.9%, Fisher Scientific, HPLC grade) was used in the chromatographic analyses. Formic acid was purchased from Panreac Quimica SLU. (Barcelona, Spain). Phenolic standards were from Extrasynthese (Genay, France). Water was treated in a Milli-Q water purification system (TGI Pure Water Systems, USA).

2.2. Pinhão preparation and antioxidant extraction

Pinhão seeds (526 g) were cooked in water (1 L) for 2 h, using a pressure cooker. Seeds were recovered using a sieve and the extract (cooking water, CW) was frozen (−18 °C).

The cooked pinhão seeds were opened and their coats dried in a forced air convection oven (Cienlab, Brazil) at 40 °C for 24 h. Thereafter, they were crushed in a domestic blender and classified using a vibrating set of sieves (200/+ 400 mesh Tyler). The resultant material was stored in the freezer (−18 °C) until extraction.

The antioxidant ethanolic (EtOH) and hydroethanolic extracts (HA) were prepared using pure ethanol and an 80% (v/v) hydroethanolic mixture, respectively. First, sieved pinhão coats (5 g) and solvent (120 g) were stirred using a magnetic stirrer for 5 h. Extraction yield (% w/w) was determined gravimetrically. All extraction procedures were carried out in triplicate.

2.3. Antioxidant activities

The antioxidant activity of the extracts was assessed by the ABTS and DPPH radical scavenging assays. The ABTS assay was based on the method of Arnao, Cano, and Acosta (2001), as adapted by Thaipong, Boonprakob, Crosby, Cisneros-Zevallos, and Hawkins Byrne (2006). First, a stock solution was prepared from equal volumes of aqueous solutions of 7.4 mM ABTS and 2.6 mM sodium persulfate. The stock solution was kept in the dark for 12 h to allow the formation of the ABTS radical (ABTS+), then diluted with methanol until an absorbance of 1.100 ± 0.010 units at 734 nm was reached (Ocean Optics, Red Tide US650 Fiber Optic Spectrometer, USA) to form the working solution. In test tubes, 2.85 mL of the working solution was added to 150 μL of the extract (or 150 μL ethanol as the control). After homogenization, the solutions were stored in the dark for 2 h and the absorbance at 734 nm was determined. All samples were analyzed in triplicate. Results were calculated using a previously obtained Trolox calibration curve (50–500 μM, y = 1.0595x − 0.0017; R² = 0.9991) and expressed as μmol of Trolox equivalent (TE) per 100 g of pinhão coat (μmolTE.100 gpinhão coat−1).

For the DPPH assay, the procedure described by Mensor et al. (2001) was used, with some modifications. In a test tube, 2.5 mL of the extract was mixed with 1 mL of 0.3 mM DPPH methanol solution (or pure methanol as the control). After 30 min in the dark, the absorbance was determined at 518 nm. All samples were analyzed in triplicate. The antioxidant capacity was calculated using a Trolox standard curve (15 – 75 μM, y = 1.0728x − 0.0168; R² = 0.9977) and the results expressed as μmolTE.100 gpinhão coat−1.

The FRAP assay was performed according to Benzie and Strain (1996), with minor modifications. First, 100 μL of the extract solution (or 100 μL of distilled water, for the blank sample) and 300 μL distilled water were added to a test tube. Then, 3.0 mL of FRAP reagent (10 mM TPTZ in 40 mM HCl, plus 20 mM ferric chloride and 300 mM acetate buffer, pH 3.6, 1:1:10 v/v/v) were added and the solution homogenized and warmed at 37 °C for 30 min in a water bath. Finally, the absorbance of the colored product (ferrous tripyridyltriazine complex) was determined at 593 nm. A Trolox standard curve (50–1000 μM, y = 0.0013x + 0.0111; R² = 0.9991) was prepared and the results expressed as μmolTE.100 gpinhão coat−1.

2.4. Phenolic compound analyses

The lyophilized extracts were re-dissolved in a 80:20 methanol-water (v/v mixture and the phenolic profile was determined by high-performance liquid chromatography-diode array detection-electrospray ionization multi-stage mass spectrometry (HPLC-DAD-ESI)/MSn (Dionex Ultimate 3000 UPLC, Thermo Scientific, San Jose, CA, USA) as previously described by Bessada, Barreira, Barros, Ferreira, & Oliveira (2016). Double online detection was carried by DAD using 280 and 370 nm as the preferred wavelengths and a mass spectrometer connected to the HPLC system via the DAD cell outlet. The MS detection was performed in negative ion mode, using an LTQ XL linear ion trap mass spectrometer (Thermo Finnigan, San Jose, CA, USA) equipped with an ESI source. Positive identification of the phenolic compounds was performed using standard compounds, when available, by comparing their retention times, UV–vis and mass spectra. Also, tentative identification was made based on data reported in the literature. For quantitative analysis, a calibration curve, obtained for each available phenolic standard, was constructed based on the UV signal. For the phenolic compounds identified in the absence of an available commercial standard, quantification was performed based on the calibration curve of the most similar available standard. Results were expressed as mg gextract−1.
2.5. Zein film preparation

Zein films were prepared by film casting according to Phiriyawirut and Maniaw (2012), with some modifications. Table 1 presents the used formulations. For the control formulation (FC), zein (19.2 g) was solubilized in a 80% v/v ethanol in water solution (100 g) by homogenization (Ultra-Turrax T25, IKA, USA) at 3000 rpm during 5 min. Next, glycerol (3.8 g) was added and the final solution degassed in an ultrasonic bath (Cristófoli, Brazil) for 15 min. The solution was then poured into teflon pans (32 × 22 cm) and oven-dried (50 °C for 24 h). For the films containing CW, EtOH and HA extracts (referred herein as FCW, FEtOH and FHA, respectively) the same procedure was adopted with the ethanol in water solution being replaced by the corresponding extracts according to the formulations presented in Table 1.

2.6. Antioxidant activity of the films

The antioxidant activity (DPPH, ABTS and FRAP) of the films were performed as described in Section 2.3, following an extraction procedure according to the method described by De Araújo et al. (2015). Briefly, ethanol (30 mL) was added to the film sample (1.5 g) homogenized using an Ultra-Turrax at 16,000 rpm for 5 min followed by stirring at 6000 rpm for 15 min. The solution was centrifuged (NT825 Nova Técnica, Brazil) at 6000 rpm for 15 min and the supernatant collected. The extraction was repeated twice, by adding ethanol (30 mL) to the centrifuged precipitate. The three resulting ethanol fractions were stored at −20 °C. Antioxidant results were expressed as μmolTE film−1 sample−1.

2.7. Mechanical properties of the films

The tensile strength tests were performed using a TA-XT Express Enhanced texture analyzer (TA-XT Express Enhanced, Texture Analyzer – Stable Micro Systems, UK) based on the American Society for Testing and Material Standards (ASTM, 2002). Ten samples (50 × 10 mm) of each formulation were pre-conditioned at 23 ± 2 °C and 53% relative humidity for 48 h. Sheet thickness was determined using a digital micrometer (Starrett, 0.001 mm resolution). For each sample, ten random points were measured. Tensile strength (MPa), elongation at break (%), and Young’s modulus (MPa) were evaluated.

2.8. Water solubility of the films

Water solubility is defined as the dry mass content obtained from film samples that were solubilized by immersion in water at 25 °C for 24 h. The adopted procedure was the one described by Pizzoli et al. (2016). Film samples (2 × 2 cm) were weighed (m0, dry basis) and then immersed in water (200 mL, 25 ± 2 °C) for 24 h. Next, the residual film was removed and dried at 70 °C in a forced air oven for 24 h. The samples were then weighed (mF) and the water solubility (SOL%) was calculated using Eq. (1) on a dry basis.

\[
SOL = \left( \frac{(m_0 - m_F) - m_f}{m_0 - m_f} \right) \times 100
\]

2.9. Field emission gun-scanning electron microscopy (FEG-SEM)

The morphology of the zein films was characterized by FEG-SEM (JEOL JSM-6701F, USA) operating at 5 kV. Samples were kept in a desiccator with silica for 2 weeks and then covered with a gold layer, prior to analysis.

3. Results and discussion

3.1. Phenolic compounds in Zein extracts

The peak characteristics (retention time, \( \lambda_{\text{max}} \) in the visible region, mass spectral data) and tentative identification of the phenolic compounds present in the extracts of A. angustifolia (Bertol.) Kuntze. are presented in Table 2. A phenolic profile recorded at 280 nm is exemplified in Fig. 1. The quantification of the phenolic compounds present in the extracts is shown in Table 3. Thirteen phenolic compounds were identified in all samples, comprising ten proanthocyanidins (catechin and epicatechin derivatives), two phenolic acids (protocatechuic and ferulic acid derivatives), a flavonol (quercetin-3-O-glucoside) and a flavanone (eriodictyol-3-O-hexoside).

Compounds 3, 6, 9 and 13 were positively identified as protocatechuic acid, (+)-catechin, (−)-epicatechin and quercetin-3-O-glucoside, respectively, according to their retention time, mass and UV−vis characteristics, by comparison with commercial standards. Catechin, (−)-epicatechin and quercetin have been previously identified in Paraná pine (da Silva et al., 2014), whereas protocatechuic acid and quercetin have been previously reported in the dead bark of

<table>
<thead>
<tr>
<th>Peak</th>
<th>Rt (min)</th>
<th>Imax (nm)</th>
<th>[M−H]− (m/z)</th>
<th>MS2 (m/z)</th>
<th>Tentative identification</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>5.01</td>
<td>279</td>
<td>593</td>
<td>575(28),549(3)467(36),441(57),425(100),423(31),407(11),305(18),289(8)</td>
<td>Prolphenidin B3</td>
</tr>
<tr>
<td>2</td>
<td>5.47</td>
<td>280</td>
<td>577</td>
<td>451(23),425(100),407(22),289(12),287(10)</td>
<td>B-type (epi)catechin dimer</td>
</tr>
<tr>
<td>3</td>
<td>5.74</td>
<td>260/292</td>
<td>153</td>
<td>109(100)</td>
<td>Protocatechuic acid</td>
</tr>
<tr>
<td>4</td>
<td>6.31</td>
<td>280</td>
<td>577</td>
<td>451(16),425(100),407(19),289(8),287(7)</td>
<td>B-type (epi)catechin dimer</td>
</tr>
<tr>
<td>5</td>
<td>6.67</td>
<td>314</td>
<td>355</td>
<td>193(100)</td>
<td>Ferulic acid hexoside</td>
</tr>
<tr>
<td>6</td>
<td>7.21</td>
<td>280</td>
<td>289</td>
<td>245(100),203(50),187(10),161(9),137(3)</td>
<td>(+)−Catechin</td>
</tr>
<tr>
<td>7</td>
<td>7.78</td>
<td>280</td>
<td>577</td>
<td>451(24),425(100),407(22),289(11),287(10)</td>
<td>B-type (epi)catechin dimer</td>
</tr>
<tr>
<td>8</td>
<td>7.9</td>
<td>280</td>
<td>577</td>
<td>451(18),425(100),407(22),289(9),287(7)</td>
<td>B-type (epi)catechin dimer</td>
</tr>
<tr>
<td>9</td>
<td>9.89</td>
<td>280</td>
<td>289</td>
<td>245(100),203(35),187(6),161(8),137(3)</td>
<td>(−)−Epicatechin</td>
</tr>
<tr>
<td>10</td>
<td>11.35</td>
<td>279</td>
<td>865</td>
<td>739(78),71(47),695(100),577(62),575(42),425(12),407(9),289(6),287(11)</td>
<td>B-type (epi)catechin trimer</td>
</tr>
<tr>
<td>11</td>
<td>12.02</td>
<td>281</td>
<td>865</td>
<td>739(78),71(56),695(100),577(46),575(82),425(11),407(9),289(6),287(9)</td>
<td>B-type (epi)catechin trimer</td>
</tr>
<tr>
<td>12</td>
<td>16.49</td>
<td>283/324</td>
<td>449</td>
<td>287(100)</td>
<td>Eriodictyol-3-O-hexoside</td>
</tr>
<tr>
<td>13</td>
<td>19.31</td>
<td>350</td>
<td>463</td>
<td>301(100)</td>
<td>Quercetin-3-O-glucoside</td>
</tr>
</tbody>
</table>
Additional fragments from the alternative cleavages of different interflavan bonds. Compound 1 presented a pseudomolecular ion [M - H]⁻ at m/z 593, which was tentatively identified as prodelphinidin B3, while compound 5 ([M - H]⁻ at m/z 355) was identified as ferulic acid hexoside. Compound 12 ([M - H]⁻ at m/z 449) was an eriodictyol-O-hexoside.

Proanthocyanidins and biflavonoids have been previously described as the most abundant compounds in A. angustifolia (Freitas et al., 2009; Peralta et al., 2016; da Silva et al., 2014; Yamaguchi, Kato, & Di, 2009; Yamaguchi, Vassa, Kato, & Mascio, 2005). Nevertheless, to the best of our knowledge, a detailed characterization of the proanthocyanidin fraction in A. angustifolia seeds has not been previously described.

Similar phenolic profiles were observed among all the samples, differing only in the amount of the identified compounds. The most abundant molecule present in samples CW and EtOH was protocatechuic acid, whereas, for sample HA, (+)-catechin (compound 6) and an (epi)catechin dimer (compound 4) predominated.

3.2. Antioxidant activity of pinhão extracts

The extraction yield and antioxidant activity (DPPH, ABTS and FRAP) of the pinhão extracts are presented in Table 4. The initial extraction applied to pinhão, simulated its preparation for consumption, being the residual cooking water (CW) collected. As observed in Table 2, this extract presented the highest solids content. The subsequent EtOH and HA extracts were obtained from the coats of the cooked pinhão, hence, lower extraction yields were obtained. The extraction yields, between EtOH and HA, were significantly different (P < 0.05), revealing that water is a good solvent for pinhão coat extraction.

Regarding the DPPH antioxidant activity, there were no significant differences (P > 0.05) among the treatments. Previously, Sant’Anna, Sfoglia, Mercali, Corrêa and Brandelli (2016) evaluated the DPPH inhibition capacity of an aqueous pinhão coat extract and obtained up to 87% inhibition.

In contrast to the DPPH results, ABTS antioxidant activity were significantly different among the extraction treatments (P < 0.05). The CW extract was the most effective against the ABTS radical, followed by the HA extract, with the EtOH extract possessing the least ABTS⁺ scavenging activity. Koehnlein et al. (2012) analyzed the ABTS scavenging activity of pinhão coat extracts (70% ethanol in water) and found 50% of the radical was scavenged with 52 μg/mL of extract. In the current study, extraction with ethanol led to significantly different FRAP (P < 0.05), compared to extractions using water (CW and HA extracts). Similarly, Bursal and Köksal (2011) also observed that aqueous extracts from sumac (Rhus cotaria L.) presented higher FRAP results than ethanol extracts.

3.3. Mechanical, thermal and microstructural properties of zein films with pinhão extracts

Table 5 summarizes the mechanical properties of the zein films containing the pinhão extracts. Film with HA extract (FHA) presented...
Mechanical properties, water solubility and antioxidant activity (DPPH, ABTS and FRAP; μmol TE/g film) of zein films with pinhão coat extracts: FC - control formulation (zein); FHA - zein with HA extract; FEtOH - zein with EtOH extract; FCW- zein with CW extract.

<table>
<thead>
<tr>
<th>Film</th>
<th>Young Modulus (MPa)</th>
<th>Tensile strength (MPa)</th>
<th>Elongation at break (%)</th>
<th>Water solubility (%)</th>
<th>DPPH (μmol TE g⁻¹)</th>
<th>ABTS (μmol TE g⁻¹)</th>
<th>FRAP (μmol TE g⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>FC</td>
<td>24.62 ± 11.01</td>
<td>5.80 ± 3.70</td>
<td>1.60 ± 0.38</td>
<td>54.71 ± 1.69</td>
<td>3.10 ± 0.12</td>
<td>36.71 ± 0.16</td>
<td>18.60 ± 1.42</td>
</tr>
<tr>
<td>FHA</td>
<td>6.16 ± 2.25</td>
<td>3.09 ± 1.89</td>
<td>1.92 ± 0.89</td>
<td>53.77 ± 6.08</td>
<td>3.36 ± 0.11</td>
<td>22.52 ± 0.73</td>
<td>50.84 ± 1.15</td>
</tr>
<tr>
<td>FEtOH</td>
<td>33.18 ± 15.81</td>
<td>7.88 ± 6.61</td>
<td>0.98 ± 0.53</td>
<td>55.38 ± 1.68</td>
<td>3.88 ± 0.14</td>
<td>33.33 ± 0.58</td>
<td>15.96 ± 1.10</td>
</tr>
<tr>
<td>FCW</td>
<td>23.47 ± 2.32</td>
<td>17.65 ± 3.67</td>
<td>3.18 ± 0.65</td>
<td>73.76 ± 1.60</td>
<td>4.88 ± 1.90</td>
<td>43.62 ± 0.28</td>
<td>13.53 ± 0.04</td>
</tr>
</tbody>
</table>

a,b Mean ± standard deviation (n = 3), means from the same column followed by different letters present a significant difference in the extraction treatment (P < 0.05) based on Tukey’s test.

The lowest Young’s modulus, and this treatment differed statistically from the others (P < 0.05). Associated with this, a low tensile strength was also observed, however, the sample prepared with EtOH extract (FETOH), as well as the control zein formulation (FC) did not differ statistically from the FHA sample (P > 0.05). In contrast, the film formulated with CW (FCW) presented a significant increase (P < 0.05) in its tensile strength. Likewise, elongation at break significantly increased (P < 0.05) in the film with CW. Zein films are known for their brittleness (Lawton, 2004); thus, the addition of CW extract, which improved the plastic behavior of the zein films, provided a strong and tough material, even when compared to the other treatments. This difference may be attributed to the CW extract composition, which was rich in phenolic acids and flavonoids (Table 3). According to Arcan and Yemenicioglu (2011), the increased flexibility of zein films by the addition of phenolic compounds probably results from the binding of phenolic compounds to the surface of the zein proteins, which leads to an increase in the free volume of the film matrix. Also, a decrease in the hydrophobic interactions among zein molecules may occur due to the action of the hydrophilic groups of the phenolic compounds that contribute to an increase in polymeric chain mobility, eliminating film brittleness.

The CW extract is mainly composed of water-soluble compounds and, proportionally, its solids content is higher than the one present in the other extracts (Table 4, extraction yield results). Accordingly, the FCW film presented a significant increase (P < 0.05) in water solubility when compared with the other samples.

In Fig. 2, the DSC thermograms of the zein films are presented. The FC curve revealed a glass transition temperature (Tg) located at 65 °C. Also, two endothermic peaks, around 100 and 160 °C, could be observed for all samples. These peaks are associated with the evaporation of residual water and glycerol, respectively (Kashiri et al., 2016). According to Ghanbarzadeh and Oromiachi (2009), the Tg value of pure zein occurs at 170–180 °C and decreases to approximately 60–80 °C in the zein films plasticized by polyols (e.g., glycerol, sorbitol and polyethylene glycol), in agreement with the result shown in Fig. 2. For thermograms of the zein films with extracts (FETOH, FHA and FCW, respectively), Tg was not observed, suggesting that it is probably below the evaluated temperature range.

The FEG-SEM images in Fig. 3 shows the morphology of the zein films, both from a surface and fracture perspective. In the fracture images, it is possible to observe that FC, FHA and FETOH samples present irregular structures that can be associated with their brittle mechanical behavior and low Young’s modulus. Also, a porous structure was formed, which has been associated with a fragile behavior (Pizzoli et al., 2016). A rough surface could be observed for all samples, however, the FCW sample was the smoother among the tested treatments.

![Fig. 2. DSC thermograms of zein films: FC - control formulation (zein); FHA - zein with HA extract; FEtOH - zein with EtOH extract; FCW - zein with CW extract.](image1)

![Fig. 3. FEG-SEM images of zein films: FC - control formulation (zein); FHA - zein with HA extract; FEtOH - zein with EtOH extract; FCW - zein with CW extract. Sample name-F: fracture images (450× magnification); Sample name-S: surface image (50,000× magnification).](image2)
Antioxidant activity of zein films with pinhão extracts

The DPPH, ABTS and FRAP antioxidant activity of the produced films are presented in Table 5. The zein control film (CF) presented antioxidant activity in all performed tests since amino acid residues present anti-
oxidant activity (Zhang, Luo, & Wang, 2011). In agreement with the anti-
oxidative evaluation of the pinhão coat extracts alone (Table 4), the DPPH results of the corresponding extracted-loaded films were similar (P > 0.05). Likewise, ABTS results showed that FCW had the highest ABTS- scavenging capacity among the used treatments (P < 0.05), corroborating the observed behavior of the extracts (Table 4). Also, the FHA film presented the highest FRAP, comparatively with the other treatments (P < 0.05) which is in agreement with results in Table 4.

4. Conclusions

Different pinhão coat extracts (cooking water (CW), ethanolic (EtOH) and hydroethanolic (HA)) were obtained and characterized presenting similar phenolic compounds profiles, being protocatechuic acid the most abundant compounds identified in samples CW and EtOH, while (+)-catechin and an (epi)catechin dimer predominated in HA sample. Regarding to extracts antioxidant activity, DPPH presented si-

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