

# Biodegradable Films Based on Poly(lactic acid) Coatings and Natural Olive-Wastewater Extracts for Active Food Packaging

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The goal of this work was to develop innovative, 100% biodegradable films with antioxidant effectiveness, based on poly(lactic acid) (PLA) and a natural olive wastewater extract (OWE). Active PLA coatings were realized by spreading a PLA/OWE coating solution, with an OWE amount up to 20 wt%, on a Poly(lactic acid)/Poly(butylene adipate-co-terephthalate) substrate. The study of active films antioxidant activity and release kinetics in foods with high lipid content was accomplished using 95% ethanol as food simulant. Preliminary shelf-life tests on a sensitive greasy food matrix (i.e. avocado fruits) were also carried out, in order to qualitatively examine the films potential in limiting fruit oxidative/browning phenomena. Finally, the effect of the OWE on the films color and photo-oxidative stability under natural light exposure was also examined. The results pointed out the influence of the morphology and distribution of the active agent on the coatings release rate, and the antioxidant activity and equilibrium time increased by increasing the antioxidant concentration. The shelf-life tests highlighted the promising perspectives for the films in retarding the oxidation/browning reactions of short shelf-life foods, and pave the way to more in-depth investigations on the most appropriate strategies to prevent the transfer of the OWE brown color while keeping intact the benefits of the antioxidant packaging.

## 1. Introduction

In recent years, the growing concern towards ecological issues due to the persistence of plastic waste in ecosystems has pushed scientific and industrial research towards the development of new materials and new processes that represent effective eco-compatible solutions. The flexible packaging field has recently been identified as a key sector to face the challenge of global sustainability, thanks the reduced weights and raw materials employment, yielding reduced waste and disposal issues (Apicella et al., 2018). In terms of raw materials, the use of biodegradable and/or compostable materials aims at minimizing the environmental impact induced by post-consumer synthetic plastic waste (Apicella et al., 2019a). The biodegradable polymers, thanks to the undoubted ecological advantages and their functional characteristics, are ideal for the realization of short life-cycle products such as food packaging. In a circular economy perspective, these materials can be incorporated with natural, bioactive molecules deriving from reevaluation of agri-food waste, obtaining active packaging capable of significantly improving the shelf-life of sensitive foods (Apicella et al., 2019c) and reducing food waste (Guillard et al., 2018).

To this aim, recent innovative research has focused on the addition of active agents from olive wastes and by-products (i.e. olive leaves, pomace and mill wastewaters) within biodegradable films (Martiny et al., 2020; Apicella et al., 2019b). Olive mill wastewaters (OW) represent the liquid fraction of residues in the three phase oil extraction system, which involves the use of a large amount of water (Nunes et al., 2016). Different organic compounds can be found in olive wastewaters as sugars, phenolic compounds, polyalcohols, lipids and pectins. Hydroxytyrosol is the main phenolic compound of OW, demonstrating several benefits such as inhibition of low-density lipo-protein oxidation, free radical scavenging activity and *in-vitro* antimicrobial activity (De Marco et al., 2007). Several extraction and membrane separation processes have been described to obtain olive wastewater extracts (OWEs) yielding to different phenolic composition and antioxidant effectiveness (Sabatini, 2010). For

42 this reason, previous research published by Apicella et al. (2019b) addressed the study of antioxidant activity,  
43 chemical-physical properties, and compatibility with the polymer matrix of several OWEs, obtained from different  
44 separation treatments. The most suitable OWE was selected to preliminarily realize poly(lactic acid) (PLA) coated  
45 antioxidant films. In this study, new biodegradable coatings were produced with higher load of OWE (up to 20  
46 wt%) and a detailed analysis on the release rate and *in vitro* antioxidant activity of the active films was  
47 conducted. The influence of films composition on the mass diffusive transport and polyphenols release in fatty  
48 food simulant was discriminated, evaluating the possibility to modulate films performance according to the food  
49 requirements. Moreover, preliminary shelf-life tests on avocado fruits were carried out, in order to qualitatively  
50 examine the potential of the active packaging in limiting modifications of the visual quality characteristics due to  
51 oxidative phenomena. Finally, the effect of the OWE on the films color and photo-oxidative stability under natural  
52 light exposure was also examined.

## 53 2. Experimental

### 54 2.1 Materials and preparation of the active coated films

55 Active coated films were produced using as substrate a biodegradable blown film based on a mixture of  
56 Poly(lactic acid) and Poly(butylene adipate-co-terephthalate) (Euromaster, Pistoia, Italy) hereafter referred as  
57 BS (i.e. Biodegradable Substrate). The substrate had a thickness of  $22 \pm 1.0 \mu\text{m}$ . The coating solution, hereafter  
58 referred as PLA, mainly comprised PLA4060 of Natureworks (Minnetonka, USA), having a D-lactide content of  
59 12 wt% which confers an amorphous morphology to the polymer. The PLA coating solution was incorporated  
60 with the olive wastewater extract (named OWE), donated by Fangiano Farming Company (Nocera Terinese,  
61 CZ, Italy). The active agent was added at 0, 5, 10 and 20 wt% based on PLA content. Details on the chemical-  
62 physical properties of the OWE and on the preparation of the PLA active coatings are reported elsewhere  
63 (Apicella et al., 2019b). The PLA dry coating thickness was  $7 \pm 0.8 \mu\text{m}$ , leading to a total structure (BS/PLA)  
64 thickness of  $29 \pm 1.8 \mu\text{m}$ . DPPH (2,2-diphenyl-1-picrylhydrazyl) and Trolox ((±)-6-Hydroxy-2,5,7,8-  
65 tetramethylchromane-2-carboxylic acid) were obtained from Sigma Chemical Co. (St. Louis, Mo., USA). All  
66 organic solvents used were analytical grade.

### 67 2.2 Characterization of the biodegradable active films

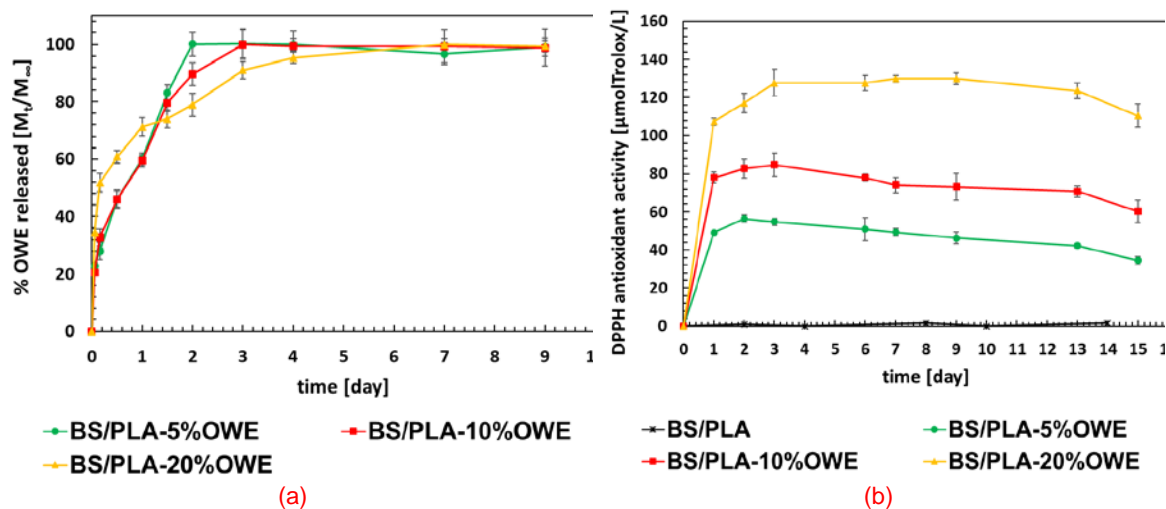
68 The release kinetic of the antioxidant films was evaluated by total immersion method as reported by Apicella et  
69 al., 2019b. 95% v/v Ethanol was selected as fatty foods simulant (D2) according to the Regulation (EU) 10/2011.  
70 Film samples were cut in rectangles of surface area equal to  $1 \text{ dm}^2$  and immersed in 100 mL of release medium.  
71 The flasks were kept in the dark under magnetic stirring, at room conditions up to 15 days. **An aliquot of the  
72 simulant was periodically sampled for measurements and then reinserted.** The concentration of antioxidants  
73 released into the simulant was quantified using a UV-Vis spectrophotometer (Lambda Bio 40, Perkin Elmer,  
74 Waltham, MA, USA) at 280 nm, on the basis of a OWE Concentration vs. Absorbance curve in the range 0-600  
75 ppm). The selected wavelength was the one at which the maximum absorbance of the OWE was found. To  
76 eliminate the influence of other polymer additives, release studies were also conducted on the control films (BS  
77 and BS/PLA) and no significant absorbance at 280 nm was observed. Results were expressed as percent ratio  
78 of  $M_t/M_\infty$  ( $M_t$  is the concentration of antioxidant (mg/mL) diffused at time  $t$ , and  $M_\infty$  represents the concentration  
79 of antioxidant diffused at equilibrium). The antioxidant activity released into the simulant solution was also  
80 measured by DPPH method, as reported by Adiletta et al. 2017 with some modifications. **50  $\mu\text{L}$  of the simulant  
81 was withdrawn from the flask and mixed with 1.95 mL of DPPH methanolic solution ( $6 \times 10^{-5} \text{ M}$ ) in a capped  
82 cuvette.** The mixture was shaken vigorously and allowed to stand at room temperature in the dark for 20 min,  
83 then the absorbance was measured at 517 nm. The blank was conducted using the pure release medium. The  
84 obtained values were expressed as  $\mu\text{molTrolox/L}$ , based on the standard curve of Trolox, and the maximum  
85 antioxidant activity was also expressed per unit volume of the coating, in  $\text{mmolTrolox/dm}^3$ . All analyses were  
86 performed in triplicate. The potential of the active films in preserving vegetables with high lipid content was  
87 investigated by sensory evaluation, in particular analysing colour and texture changes over time, of packaged  
88 fresh-cut avocado (cv. Bacon) fruit. The fruit samples were cut in slices of ca. 0.5 cm thickness, mixed thoroughly  
89 in order to allow a random selection and divided into single bags, each one measuring  $15 \times 10 \text{ cm}^2$  in size and  
90 approximatively of the same weight (ca. 40 g). The bags were sealed and a good contact among the active  
91 coating and the food surface was ensured. **Then, the packages were stored in a refrigerator at  $6^\circ\text{C}$  up to 9 d.  
92 Three bags were prepared for each testing day and were opened to better evaluate the color changes on the  
93 avocado surface.** Finally, colour measurements were carried out on the films by using a colorimeter CIE-Lab  
94 (Chroma Meter II Reflectance CR-300, Minolta, Japan) and the results were expressed according to colour  
95 coordinates  $L^*$  (darkness/lightness),  $a^*$  (greenness/redness),  $b^*$  (blueness/yellowness). The chromatic  
96 parameters were evaluated soon after the coatings production (i.e. at time 0) and at regular time intervals (up  
97 to 73 d) exposing films to natural light, in order to evaluate possible effects due to photo-oxidation phenomena.

98 The colour variation of the films over time was evaluated by " a\* and " b\* parameters, and by means of the  
 99 colour-difference equation CIELAB "  $E^*_{ab} = [(L^*-L_0^*)^2 + (a^*-a_0^*)^2 + (b^*-b_0^*)^2]^{1/2}$ , based on the coordinates L\*, a\*  
 100 and b\*, where the subscript 0 refers to the BS sample.

### 101 3. Results and discussion

#### 102 3.1 Study of films release kinetic, antioxidant activity and inhibition of food oxidation phenomena.

103 The characterization of the release behavior of OWE from PLA coatings and the comparison among the release  
 104 curves for all the films investigated are reported in Figure 1: Figure 1(a) depicts the %OWE released as percent  
 105 ratio of  $M_t/M_\infty$ , where measured  $M_\infty$  was 74, 100 and 154 mg/L for films at 5%, 10% and 20% OWE, respectively;  
 106 Figure 1(b) shows the antioxidant activity in the food simulant during the time, expressed as  $\mu\text{molTrolox/L}$ ; Table  
 107 1 reports the maximum antioxidant activity per unit volume of the coating, expressed as  $\text{mmolTrolox/dm}^3$  and  
 108 the time at which it was reached. As can be observed from Figure 1(a), the films at 5% and 10% OWE follow  
 109 almost the same release kinetic within the first 36 h. The percentage of antioxidant released is equal to ~20%,  
 110 ~30%, 46%, 60% and ~80% after 1, 3, 12, 24 and 36 h, respectively, suggesting for these samples similar  
 111 morphology and mass transport interactions regulating the diffusion mechanisms. The equilibrium was reached  
 112 after 48 h and 72 h, respectively. By contrast, the BS/PLA-20%OWE film shows a different release behavior at  
 113 short ( $t < 24$  h) and long ( $t > 24$  h) times. For  $t < 24$  h, the release rate is the highest and the % OWE released  
 114 is equal to ~34%, 52%, 61% and 71% after 1, 3, 12 and 24 h, respectively. After 24 h, a slower kinetic is  
 115 established, and the system reaches the equilibrium after ca. 7 d of test. This behavior suggests an uneven  
 116 distribution of the active phase within the coating thickness, being more concentrated on the coating surface  
 117 and leading to a faster initial release of the antioxidant. The resistance to diffusive transport increases when the  
 118 diffusion involves the inner side the coating and the concentration of the OWE increases. A less than linear  
 119 increase of antioxidant activity (Figure 1(b)) was observed by increasing OWE content, with a maximum equal  
 120 to  $28.21 \pm 1.93$ ,  $42.31 \pm 2.2$  and  $64.86 \pm 1.2$   $\text{mmolTrolox/dm}^3$  for BS/PLA-5%OWE, BS/PLA-10%OWE and  
 121 BS/PLA-20%OWE samples, respectively (Table 1). These outcomes confirmed the effectiveness of the  
 122 produced films as carriers for antioxidants release, with the possibility to tune the release kinetic with a proper  
 123 design of the systems composition and to tailor the films performance on the preservation requirements of the  
 124 target food.  
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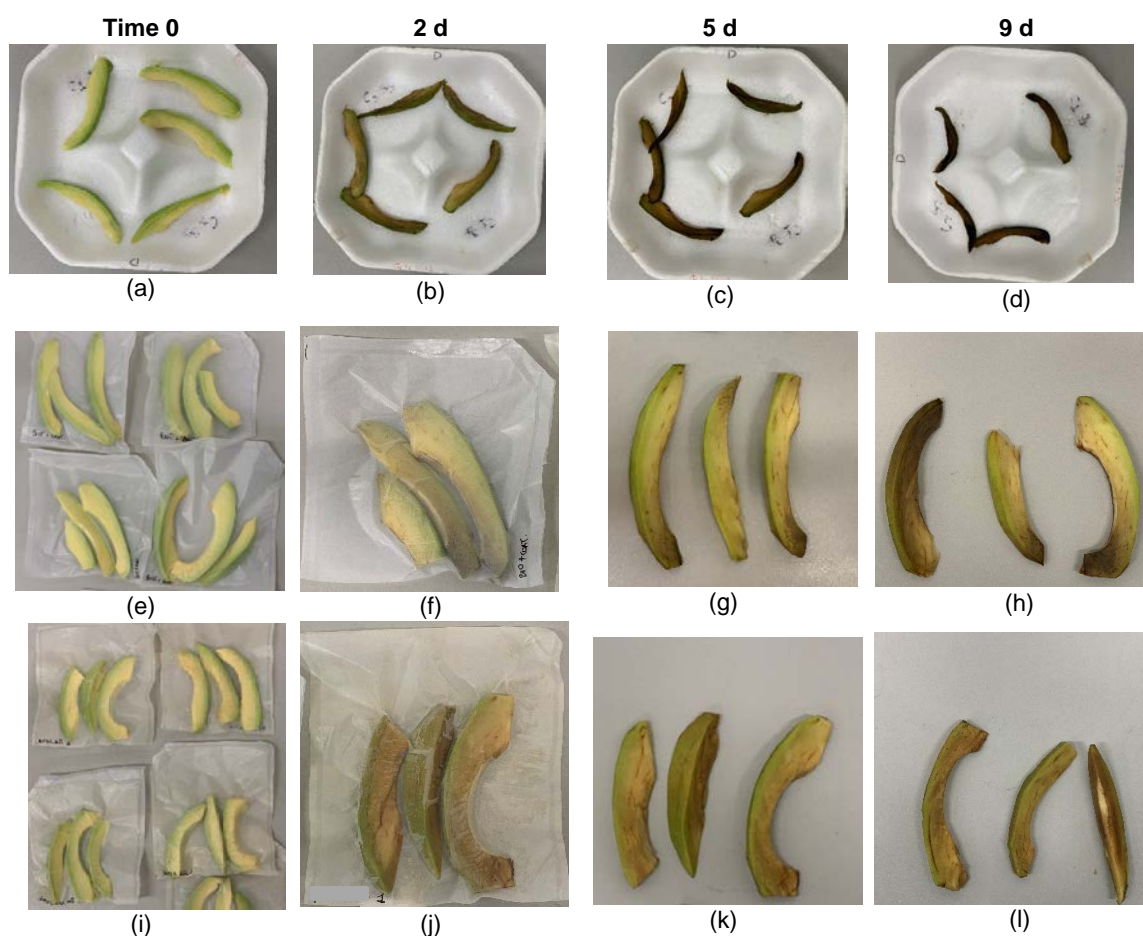


126 Figure 1: (a) OWE release rate ( $M_t/M_\infty$ ) from active coated films in 95% Ethanol, at 23°C and (b) DPPH  
 127 antioxidant activity (expressed in  $\mu\text{molTrolox/L}$ ) released from active coated films in Ethanol 95%, at 23°C.

128 Table 1: Time at which the maximum antioxidant activity is reached, and average maximum antioxidant activity  
 129 (expressed as  $\text{mmol Trolox/dm}^3$ ) for the active coated films. Values followed by different letters within the same  
 130 column were significantly different according to Duncan's test ( $P < 0.05$ ).

Sample Film	Time at max antioxidant activity [d]	Average max. antioxidant activity [ $\text{mmolTrolox/dm}^3$ ]
BS/PLA-5%OWE	2	$28.21 \pm 1.93^a$
BS/PLA-10%OWE	3	$42.31 \pm 2.2^b$
BS/PLA-20%OWE	7	$64.86 \pm 1.2^c$

131 The results, coupled with the measured low barrier performance of the films (Apicella et al., 2019b) suggested  
 132 their possible application to preserve sensitive foods with short shelf life as fresh-cut horticultural products. To  
 133 this aim, the efficacy of the active films in inhibiting oxidation/browning phenomena of sensitive foods was  
 134 evaluated by preliminary shelf life tests on fresh-cut avocado slices. Figure 2 shows the comparison among the  
 135 pictures of the samples stored at 6°C up to 9 d without package (first row), in BS/PLA film (second row) and  
 136 BS/PLA-20%OWE film (third row), taken as example. Avocado is a fruit of unusually high oil content (15% to  
 137 30% depending on the variety) and its shelf-life is severely determined by oxidative processes, which affect both  
 138 lipidic and aqueous fractions. The decay is produced by enzyme mediated oxidative reactions, with the formation  
 139 of dark compounds, as well as changes in lipids due to auto-oxidation (Elez-Martinez et al., 2005). **In order to**  
 140 **better evaluate the colour changes undergone by the packaged fruits, different bags were opened at fixed times.**  
 141 As observable in Figure 2 (b), (c) and (d), unpackaged avocado displayed the fastest and most severe  
 142 dehydration and blackening of the tissues. Fruit samples packaged in BS/PLA film (Figure 2 (f), (g) and (h)) also  
 143 showed a progressive darkening attributable to oxidation phenomena, which was mainly concentrated on the  
 144 fruit edges and became more consistent by increasing the storage time. The pulp also exhibited an ongoing loss  
 145 of firmness which was qualitatively evaluated to the touch.  
 146



147 *Figure 2 Pictures of avocado slices unpackaged (first row), packaged in BS/PLA (second row) and BS/PLA-*  
 148 *20%OWE (third row) films. From left to right: slices after 0, 2, 5 and 9 storage days at 6°C.*

149 **The different avocado slices stored in the active film at 20%OWE (Figure 2 (j), (k) and (l)), on the other hand,**  
 150 **experienced a rapid color variation from yellow to brown.** However, the homogeneous distribution of the brown  
 151 color over the surface of the fruit, which remained almost unaltered over the time, could be attributable to the  
 152 OWE migration from the film to the avocado pulp. In fact, it is well known that the OWs are characterized by a  
 153 typical dark brown color, due to the polymerization of tannins and to low molecular weight phenolic compounds  
 154 (Apicella et al., 2019b; Otles and Semih, 2012). Moreover, no appearance of oxidation black spots focused on  
 155 the edges was detected, and a good preservation of the pulp texture was observed. These exploratory results  
 156 underlined the potential helpful role of the OWE in retarding the oxidation/browning reactions of sensitive foods,



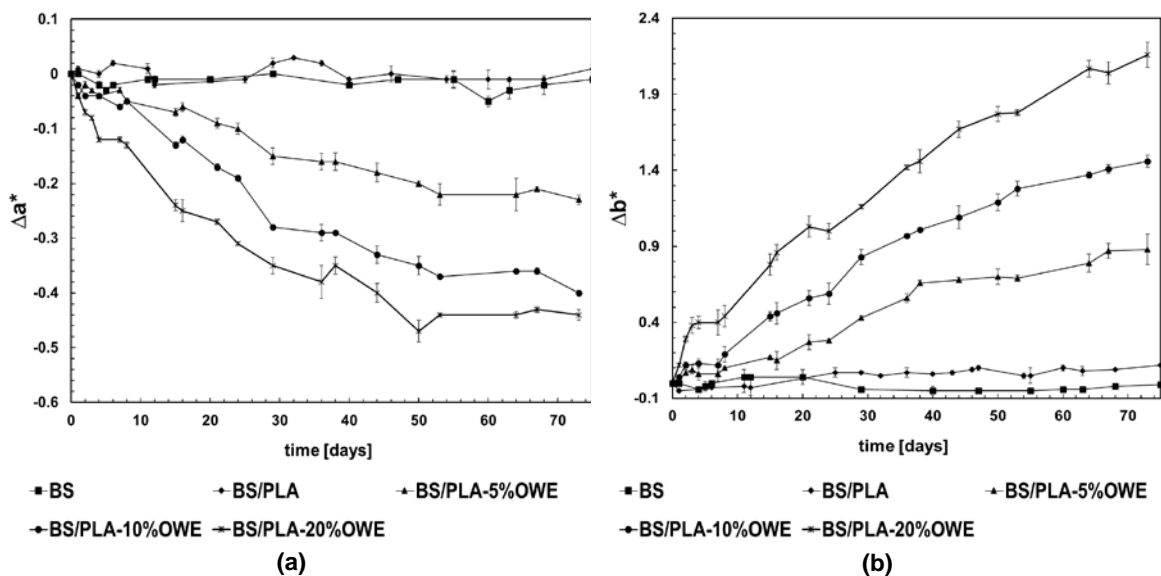
157 and pave the way to more in-depth, quantitative shelf-life studies on the most appropriate strategies to  
 158 prevent/limit the transfer of the OWE brown color while keeping intact the benefits of the antioxidant packaging.

### 159 3.2 Evaluation of optical properties and photo-oxidative stability

160 The effect of the OWE addition on the optical properties and color stability of the PLA coatings was evaluated  
 161 by colorimetric analyses. Table 2 reports the CieLab color coordinates  $L^*$ ,  $a^*$ ,  $b^*$  and the chromatic variation  $E$   
 162 for all the films soon after their production. No significant differences ( $P > 0.05$ ) were measured among BS and  
 163 BS/PLA samples, thanks to PLA excellent optical properties (Apicella et al., 2019a). Instrumental color analyses  
 164 confirmed that the incorporation of the OWE gave a colored brown taint to the films, with an increase in  
 165 greenness ( $a^*$ ) and yellowness ( $b^*$ ) values by increasing the antioxidant concentration.

166 Table 2: CieLab color coordinates  $L^*$ ,  $a^*$  and  $b^*$  for the neat biodegradable substrate (BS) and for the PLA  
 167 coated films at 0, 5, 10 and 20% OWE concentration. Chromatic variation  $E$  is also reported. Values followed  
 168 by different letters within the same column were significantly different according to Duncan's test ( $P < 0.05$ ).

Sample Film	$L^*$	$a^*$	$b^*$	$E$
BS	$96.9 \pm 0.2^b$	$-0.74 \pm 0.12^c$	$2.28 \pm 0.20^a$	-
BS/PLA	$97.3 \pm 0.8^b$	$-0.80 \pm 0.25^c$	$2.27 \pm 0.26^a$	0.40
BS/PLA-5%OWE	$96.7 \pm 0.3^b$	$-1.04 \pm 0.30^{b,c}$	$3.98 \pm 0.11^b$	1.73
BS/PLA-10%OWE	$96.2 \pm 1.5^b$	$-1.51 \pm 0.33^{a,b}$	$6.02 \pm 0.44^c$	3.80
BS/PLA-20%OWE	$93.9 \pm 0.8^a$	$-1.80 \pm 0.56^a$	$11.9 \pm 0.63^d$	10.1



169 Figure 3: Change in redness ( $\Delta a^* = a^*_t - a^*_{t=0}$ , (a)) and yellowness ( $\Delta b^* = b^*_t - b^*_{t=0}$ , (b)) during the time, for the BS  
 170 substrate and BS/PLA coated films at 0, 5, 10 and 20% OWE concentration.

171 The total color change of the active films ( $E^*$ ) with respect to the neat BS and BS/PLA films was also  
 172 highlighted. Similar outcomes were also found by other authors (Marcos et al., 2014; Manzanarez-López et al.,  
 173 2011) reporting increased yellowness in PLA films containing olive leaves extract or  $\pm$ -tocopherol. In order to  
 174 analyze the photo-oxidative stability of the films on long storage time, the changes of  $a^*$  and  $b^*$  values with  
 175 respect to time 0 was measured over 73 d.  $\Delta a^*$  and  $\Delta b^*$  trends are shown in Figure 3 (a) and (b), respectively.  
 176 No significant color change for the BS substrate and the BS/PLA films were observed, while the BS/PLA-OWE  
 177 coatings undergo an increase both in redness and yellowness of the samples, which became more significant  
 178 by increasing the antioxidant content. In particular, the total increase in  $\Delta a^*$  and  $\Delta b^*$  parameters, after 73 d, was  
 179 equal to -0.23, -0.4, -0.44 and 0.88, 1.46, 2.16 for the films at 5%, 10% and 20% OWE, respectively. The  
 180 progressive browning is attributable to the oxidation of the active phase over the time. Polyphenols are very  
 181 unstable and susceptible to degradation because of high temperatures, light, oxygen, solvents, enzymes,

182 metallic ions, or association with other food constituents (Volf et al., 2014). However, only minor changes in the  
183 photo-oxidative stability of the films occurred considering short shelf-life packaging application times.

#### 184 4. Conclusions

185 In this work, innovative biodegradable antioxidant films based on olive mill wastewaters were developed through  
186 a conventional technique commonly applied in packaging industry. The analysis on the release rate and *in vitro*  
187 antioxidant activity proved that the films are able to be used as carriers for the controlled release of the  
188 antioxidant agent, with the possibility to modulate the release performance by properly designing the films  
189 composition. An increasing antioxidant activity (from 28.21 to 64.86 mmolTrolox/dm<sup>2</sup>) and release time (from 2  
190 to 7 d) was found by increasing OWE concentration up to 20%. Preliminary shelf-life tests endorsed the  
191 perspectives to use films as 100% green alternative for preserving O<sub>2</sub>-sensitive foods with high respiration rates,  
192 such as fresh-cut avocado. However, appropriate strategies to prevent/limit the transfer of the natural OWE  
193 brown color to the food have to be implemented for films application. Finally, only minor changes in the photo-  
194 oxidative stability of the films occurred if considering short shelf-life packaging applications times.

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