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Sincerely,

Pr. Fabienne Remize

la science pour la vie, l'humain, la terre

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**Antimicrobial, sealable and biodegradable packaging to maintain the quality of shredded carrots and pineapple juice during storage**

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1 **Antimicrobial, sealable and biodegradable packaging to maintain the quality of**  
2 **shredded carrots and pineapple juice during storage**

3

4

5 **Research highlights**

6 - Biodegradable and sealable packaging films were coated with antimicrobials

7 ~~Films contained potassium sorbate and/or potassium benzoate as antimicrobials in a~~  
8 ~~coating layer~~

9 - Potassium sorbate coated films exerted the strongest antimicrobial effect

10 - Active films limited color modification and microbial development of pineapple juice

11 - Active layer mostly unchanged the sealable film physical-mechanical properties

12

13

## 14 **Abstract**

15 Increasing consumer demand for foods with high nutritional quality, prolonged shelf life and  
16 low environmental impact of the package, is driving innovation towards the development of  
17 new packaging. Multifunctional food packaging films, **biodegradable, heat-sealable and**  
18 **antimicrobial**, were developed. **A PLA coating layer incorporating either sodium benzoate,**  
19 **potassium sorbate, or a combination of them was deposited** onto a poly(lactic)  
20 acid/poly(butylene adipate-co-terephthalate) substrate film. The effectiveness of the  
21 developed systems to preserve the quality of foods was tested in shelf-life experiments  
22 performed on shredded carrots and pineapple juice, selected as model processed raw foods.  
23 **The best performance was observed for the active film containing potassium sorbate:**  
24 **microbial populations increased less rapidly and were 0.7 to 1.8 log CFU/g lower at the end**  
25 **of storage period in this film than in control packs.** Of the two model foods, the pineapple  
26 juice was better preserved: **after 7 days in active packaging, color change and microbial**  
27 **counts of juice were below that of control, observed after one day and after 3 days of storage**  
28 **respectively.** Moreover, the incorporation of the active phases did not significantly affect the  
29 mechanical, barrier and optical properties of the films, opening new ways to prolong shelf-life  
30 of minimally processed foods.

## 32 **Key-words**

33 Active packaging; biodegradable films; preservative; shelf-life; pineapple; carrot; heat-  
34 sealability

## 36 **Abbreviations**

37 **TA:** Titratable Acidity; **PS:** Potassium Sorbate; **SB:** Sodium Benzoate; **PLA:** Poly(Lactic)  
38 **Acid;** **PBAT:** Poly(Butylene Adipate-co-Terephthalate); **TR:** TRansparency; **TPC:** Total Plate  
39 **Counts;** **YMC:** Yeast and Mold Counts; **EC:** Enterobacterium Counts

## 40 Introduction

1  
2 41 Packaged minimally processed fruit and vegetables are of growing popularity for their high  
3  
4 42 nutritional value and convenience. However, these foods are highly susceptible to microbial  
5  
6 43 growth, which represent the main cause of their spoilage (Ragaert et al., 2007). In fact,  
7  
8 44 minimal processing operations, such as cutting, shredding, juicing, lead to the disruption of  
9  
10 45 subcellular compartmentalization and the release of cellular nutrients, which promote the  
11  
12 46 growth of microorganisms (Klaiber et al., 2005) and cause undesirable biochemical and  
13  
14 47 physiological changes. These changes decrease the safety and the nutritional and sensory  
15  
16 48 quality and thus, shorten food shelf life (Wang et al., 2015).

17  
18  
19 49 Among the minimally processed fruit and vegetables, shredded carrots and pineapple juice  
20  
21 50 are among the most widespread and consumed for their sensory and nutritional properties  
22  
23 51 (Klaiber et al., 2005; Leneveu-Jenvrin et al., 2020). Typically, the shelf-life of fresh pineapple  
24  
25 52 juice or shredded carrots stored at 4°C is comprised between three and eight days (Leneveu-  
26  
27 53 Jenvrin et al., 2020; Mahendran, 2015; Piscopo et al., 2019). The decrease in quality of fresh  
28  
29 54 pineapple juice is observed particularly through yeast and mold population increase and  
30  
31 55 modification of color towards browning, leading to sensory descriptors of “fermented”  
32  
33 56 (Leneveu-Jenvrin et al., 2020). On the contrary, quality decrease of shredded carrots during  
34  
35 57 storage is mainly due to gram-negative bacteria development, especially Pseudomonaceae,  
36  
37 58 and surface color change (Klaiber et al., 2005; Xylia et al., 2018). Hence, the limitation of  
38  
39 59 quality loss in the two products has different microbial targets, fungi for pineapple juice and  
40  
41 60 bacteria for shredded carrots.

42  
43  
44 61 The feasibility of a single treatment to limit microbial growth in these two foods requires  
45  
46 62 investigations. Lowering temperature is the most potent way to decrease the microbial  
47  
48 63 growth rate, but it cannot be enough. The addition of antimicrobial agents is another  
49  
50 64 approach to limit the growth of microorganisms (Durango et al., 2006). Among the  
51  
52 65 antimicrobials, sorbates and benzoates hold the major share of the market (Kuplennik et al.,  
53  
54 66 2015). Potassium sorbate (PS) is the salt of sorbic acid. It is classified as a food additive  
55  
56 67 (E202) according to EC 1333/2008 and inhibits yeasts and molds and some bacteria, at pH

68 lower than 6.5 (Davidson et al., 2005). Sodium benzoate (SB) is the salt of benzoic acid: it  
69 has been classified as a food additive (E211) in EC 1333/2008 and inhibit yeasts, molds and  
70 bacteria at pH lower than 4-4.5 (Musyoka et al., 2018). Synergistic effects between these  
71 compounds have also been reported (Stanojevic et al., 2009). Usually, these antimicrobial  
72 agents are directly added into the food. However, the incorporation of the antimicrobials into  
73 a package could be more efficient than their direct addition into the food, because they may  
74 gradually migrate from the package onto the food surface. **Release rate of antimicrobials**  
75 **depends on temperature, pH, and on the food surface for diffusion, as reported by many**  
76 **authors on several foods and packaging systems (Glicerina et al., 2021; T. Z. Jin, 2017; Uz &**  
77 **Altinkaya, 2011; Vasile & Baican, 2021). Pineapple juice and shredded carrots differ by their**  
78 **liquid or solid nature, which modifies their interaction with packaging film. For that reason and**  
79 **because of different microbial targets to inhibit, these two minimally-processed products were**  
80 **chosen as model foods for this study.**

81 Different authors reported that the incorporation of PS or SB or their combination at  
82 concentration ranging from 10 to 15 wt% into biodegradable and non-biodegradable  
83 polymeric systems was effective to delay the growth of microorganisms (Careli-Gondim et  
84 al., 2020; T. Jin et al., 2010; Shen et al., 2010). However, most of the existing investigations  
85 focused on the antimicrobial activity of the films *in vitro*, while a few studies are available on  
86 *in vivo* tests that demonstrated the effectiveness of these biodegradable systems on real  
87 food such as fresh noodles, strawberry puree, berries and avocado (Careli-Gondim et al.,  
88 2020; T. Jin et al., 2010; Junqueira-Gonçalves et al., 2016; Wangprasertkul et al., 2021).  
89 In this work, we intended to develop innovative, multifunctional and environmentally friendly  
90 food packaging films able to prolong the shelf-life of minimally processed fruits and  
91 vegetables.

92 To this aim, biodegradable systems consisting of a PLA/PBAT substrate film, coated with an  
93 amorphous PLA layer incorporating either PS, SB or a combination of them, were prepared.

94 **The coating layer constituents were selected to ensure a biodegradable film structure,**



95 combined with heat-sealing ability and antimicrobial activity. The ability to increase food  
96 shelf-life and the technical suitability of the developed films were determined.

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## 98 **Materials and Methods**

### 99 *Fruit and vegetable sampling and processing*

100 Carrots (*Daucus carota* cv. Maestro) and pineapples (*Ananas comosus* cv. Queen Victoria)  
101 were collected from local markets in Reunion island (France). They were transferred to the  
102 laboratory and stored at 10°C until processing, within 24h. All the equipment used was  
103 previously rinsed with sodium hypochlorite solution.

104 Carrots were rinsed into chlorinated water 200 ppm for 3 min, washed with distilled water and  
105 then allowed to dry naturally. Then, they were subjected to shredding and put into a tray.  
106 Fresh lemon juice ( $100 \pm 20$  mL for 1 kg of carrots) was added in order to obtain a pH equal  
107 to 4.

108 Pineapple fruit at a similar stage of ripeness were manually peeled and cut and prepared into  
109 juice (Extractor Wismer EW-01, CAPAVENIR, France).

### 110 *Active films*

#### 111 *Film composition and production*

112 Active films were developed through a coating technique. The support surface used for the  
113 coatings was a commercial biodegradable film named as Biopolymer (Euromaster, Italy).  
114 This film is made by a blend of PLA and PBAT, has a medium thickness of  $22.0 \pm 1.0$   $\mu\text{m}$ , an  
115 oxygen and water vapor permeability of  $5.15 \pm 0.35$   $(\text{cm}^3 \times \text{cm}) / (\text{m}^2 \times \text{d} \times \text{bar})$  and  $9.19 \pm 0.37$   
116  $(\text{g} \times \text{m}) / (\text{m}^2 \times \text{Pa} \times \text{s})$  respectively (Apicella et al., 2019). The material selected as matrix for the  
117 coating layer was PLA4060D supplied by NatureWorks™ (Minnetonka, USA). The active  
118 phases added in the coating layer were potassium sorbate (PS) and sodium benzoate (SB).  
119 Acetone and a surfactant (Tween 85) were also used for the coating production. They were  
120 all supplied by Sigma Aldrich Co. (Missouri, USA). All organic solvents were analytical grade.  
121 A polymeric solution of consisting of acetone and PLA, in a mass ratio of 80:20, was mixed  
122 with an aqueous solution containing the active phases and the surfactant (at a concentration  
123 of 1 wt% of the total solid content) in a volumetric ratio of 100:10. Then, the obtained  
124 emulsion was applied to the surface of the support, which was the biopolymer film, by means  
125 of a threaded hand coater. The active phases were PS, SB and a combination of them in a

126 50% mass ratio and were added at a concentration of 0 wt%, 5 wt%, 10 wt% or 15 wt% of  
127 the total solid content of the coating layer. A level of 15 wt% was the highest limit for the  
128 complete solubilization of SB in water.

### 129 *Film characterization*

130 Mechanical tests were performed on rectangular specimens of the produced films (width =  
131 12.7 mm, length = 30 mm) using a SANS dynamometer equipped with a 100 N load cell. The  
132 testing speed was set according to the ASTM 882. A minimum of five specimens for each  
133 film were tested.

134 Oxygen permeability measurement on the films was carried out using a permeabilimeter  
135 (GDP - C 165 of Brugger), with a manometric operation, connected to a thermo-controlled  
136 bath (ThermoHaake). Before the test, an evacuation was performed on both the upper and  
137 lower half-cells to remove the moisture and other residual gases. The test temperature was  
138 set at 23°C and the oxygen flow to 80 mL/min, following ISO 15105-1. Sample area was 16  
139 cm<sup>2</sup>. The obtained oxygen transmission rate (OTR) was multiplied by the respective  
140 thickness of the film to calculate the permeability coefficient (PO<sub>2</sub>). All tests were made in  
141 triplicate and the averaged values are reported (standard deviation < 8%).

142 The transparency of the films was measured following the ASTM D1746 – 03 through a UV-  
143 VIS spectrophotometer (Lambda 800, USA). Squared samples of the films (length = 5 cm)  
144 were placed on the internal side of the spectrophotometer cell and the transmittance was  
145 measured at 560 nm. Three replicates of each film were tested. The percentage of  
146 transparency (TR) was calculated according to:

$$147 \quad TR = T_r/T_0 \times 100 \quad (1)$$

148 where T<sub>r</sub> was the transmittance with the specimen in the beam and T<sub>0</sub> was the transmittance  
149 with no specimen in the beam.

150 Hot seal strength (hot-tack) tests were performed with a heat seal tester model HSG-C  
151 (Brugger), according to standard ASTM F1921-98, Method B. Samples were cut into strips  
152 (width = 1.5 cm and length = 30 cm) and then were hot pressed at a temperature of 85°C

153 under a pressure of 15 N/cm<sup>2</sup> with a welding time of 0.5 s. The heat-sealing strength was  
154 measured right after the sealing. A minimum of five replicates was tested for each sample.

#### 155 *Food packaging*

156 Films were cut into squared shapes of 11×11 cm<sup>2</sup> and sealed in pairs on three sides using a  
157 Multivac sealing machine. The obtained bags were filled either with 25g of shredded carrots  
158 or with 10 ml of fresh pineapple juice, sealed on the remaining side and stored at 4 °C. One  
159 pack was prepared for each date of analysis. All the experiments were performed in triplicate

#### 160 *Food quality determination*

##### 161 *Microbiological counting*

162 For the microbiological analysis, 4.5 ± 0.5 g of shredded carrots or of pineapple juice were  
163 removed aseptically from each bag and were transferred into a sterile stomacher bag. Then,  
164 the same weight of saline peptone water was added (SPW, Condalab, Torrejón de Ardoz,  
165 Madrid, Spain). Carrots and SPW were blended for 60 s by using a stomacher.

166 Total aerobic plate counts (TPC), and yeast and mold counts (YMC), were determined after  
167 plating on Plate Count Agar (PCA, Biokar Diagnostics Solabia, Beauvais, France) incubated  
168 at 30°C for 72h and Sabouraud glucose agar with 100 mg/L chloramphenicol (SGA, Biokar  
169 diagnostic, Solabia, Allonne, France) incubated at 30°C for 5 days, respectively.

170 Enterobacterium plate counts (EC) were determined on VRBG agar  
171 (Biokar diagnostic) after 48h of incubation at 37°C.

##### 172 *pH and titratable acidity*

173 Shredded carrots (1.5 g) were mixed with 10 mL of distilled water prior to pH and titratable  
174 acidity (TA) determination. The pH value was determined by a pH meter (5231 and GLP22,  
175 Crison Instruments S.A. Barcelona, Spain), and TA was determined by titration with 0.05 M  
176 NaOH (TitroLine easy, Schott, Mainz, Germany). TA was expressed in citric acid equivalents  
177 in g/100 mL.

##### 178 *Visual appearance and color determination*

179 Pictures of 3.0 ± 0.5 g of shredded carrots were taken using a viewing booth Just Normlicht  
180 in order to have a constant light source.

181 The color of samples (mixed carrots or juice) was assessed with a spectrophotometer CM  
182 3500d (Minolta®, Carrières-sur-Seine, France). The coordinates  $L^*$ ,  $a^*$  and  $b^*$  of the CIELAB  
183 space were measured. The total colour variation was calculated as follow:

$$\Delta E = \sqrt{(L^*_a - L^*_0)^2 + (a^*_a - a^*_0)^2 + (b^*_a - b^*_0)^2} \quad (2)$$

185 in which  $L^*_a$ ,  $a^*_a$  and  $b^*_a$  refer to the assay condition and  $L^*_0$ ,  $a^*_0$  and  $b^*_0$  to the initial  
186 condition used as a control.

### 187 *Statistical analysis*

188 The statistical analysis of the data was performed with XLSTAT software (Addinsoft, Paris,  
189 France). A confidence interval of 95% was used for all analyses. The Fisher (LSD) test was  
190 applied for ANOVA for carrots. The Bonferroni test was applied for pairwise comparisons for  
191 pineapple juice with a p-value of 0.0001.

192

## 193 Results and Discussion

1  
2 194 The effectiveness of the developed systems was tested on two different foods, shredded  
3  
4 195 carrots and pineapple juice. The foods were packed in active bags produced from films of  
5  
6 196 different compositions and stored in refrigerated conditions for 10 days, during which they  
7  
8 197 were periodically tested for appearance and nutritional quality. Films having a content of  
9  
10 198 active phase of 5 wt% or 10 wt% were not effective to delay the microbial growth of the  
11  
12 199 selected foods (data not shown). This is in accordance with literature (Shen et al., 2010), in  
13  
14 200 which a minimum content of 15 wt% PS in starch films was required to exhibit *in vitro*  
15  
16 201 antimicrobial function, due to the hydrogen bonding interaction between the hydroxyl group  
17  
18 202 of the polymer matrix and the carboxyl group of potassium sorbate. Therefore, only the  
19  
20 203 results corresponding to films with an active phase content of 15wt% are presented.

### 204 *Effect of antimicrobial films on shredded carrot quality*

25 205 ~~The physicochemical and color parameters and the microbiological quality of shredded~~  
26  
27 206 ~~carrots packed in the control and in the antimicrobial films were monitored during refrigerated~~  
28  
29 207 ~~storage at 4°C.~~

30  
31  
32  
33 208 The pH values and titratable acidity of shredded carrots during 10 days of storage are  
34  
35 209 reported in **Table 1**. Since the antimicrobial activity of both PS and SB is pH-dependent  
36  
37 210 (Davidson et al., 2005; Musyoka et al., 2018), lemon juice was added during processing of  
38  
39 211 carrots in order to reduce their initial pH. The lemon juice addition led to a decrease in the  
40  
41 212 initial pH value of shredded carrots from  $6.2 \pm 0.5$  to  $4.1 \pm 0.1$ , which is consistent with the  
42  
43 213 antimicrobial activity of both PS and SB.

44  
45  
46 214 During the storage of shredded carrots, a pH increase, though not significant for all  
47  
48 215 conditions, was observed during the first three days, followed by a slight decrease until day  
49  
50 216 10. As expected, TA followed the opposite trend. The changes of pH and TA during the first  
51  
52 217 days can be explained by spatial repartition at the bottom of the pack of the acidic liquid  
53  
54 218 (lemon juice), whereas the further decrease could result from microbial acidification, as  
55  
56 219 previously reported (Alegria et al., 2010; Piscopo et al., 2019; Pushkala et al., 2012).

220 Color measurement of shredded carrots were also performed, since color variation of food  
1  
2 221 during the time is generally perceived by the consumers as a loss of freshness (Piscopo et  
3  
4 222 al., 2019). For shredded carrots, surface dehydration and production of lignin could result in  
5  
6 223 the discoloration of the vegetable during the storage (Alegria et al., 2010; Pushkala et al.,  
7  
8 224 2012). At the initial day, shredded carrots exhibited a low brightness ( $L^*$ ) and an intense red  
9  
10 225 ( $a^*$ ) and yellow ( $b^*$ ) color (**Online Resource 1**). Fai et al. (Fai et al., 2016) reported  
11  
12 226 comparable values for shredded carrots of  $a^*$  and  $b^*$ , but also a higher value of brightness.  
13  
14 227 **Poorly significant color differences to day 0 were observed during storage for the control**  
15  
16 228 **packaging film (Table 1)**. This could be owed by the presence of the film that acted as a  
17  
18 229 barrier to water, reducing dehydration and thus discoloration of the packed food. Also  
19  
20 230 Piscopo *et al.* (Piscopo et al., 2019) did not observe any relevant discoloration of shredded  
21  
22 231 carrots packaged in polypropylene (PP) pouches up to 10 days of storage at 4°C, while  
23  
24 232 Alegria et al. (Alegria et al., 2010) found a fading/whitening of the characteristic color of  
25  
26 233 shredded carrots packed in PP bags, significant after 7 days. **A large variability of  $L^*$ ,  $a^*$  and**  
27  
28 234  **$b^*$ , and thus color difference was observed (Table 1 and Online Resource 1), which can**  
29  
30 235 **explain the observation**. Moreover, as reported in **Table 1**, no relevant differences in color of  
31  
32 236 carrots packed in the films compared to day 0 were observed, except for the 15 wt% SB + 15  
33  
34 237 wt% PS film after 10 days, sign that the active phase did not modified the global color of the  
35  
36 238 packed food. To support that data, the visual appearance of shredded carrots stored in the  
37  
38 239 different films for 7 days is reported in **Figure 1**. No relevant browning or widespread change  
39  
40 240 was observed during the time, but localized white and black tiny spots appeared on carrot  
41  
42 241 surface whatever the packaging film.  
43  
44 242 Microbial counts of shredded carrots are reported in **Figure 2a** and **Figure 2b**. The TPC of  
45  
46 243 shredded carrots packed in the control film significantly increased during the storage, being  
47  
48 244 significantly different to day 0 from 3 days of storage, and reaching  $11.3 \pm 0.1$  log CFU/g  
49  
50 245 after 10 days. During the same time, the increase of YMC was significant only after 10 days  
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52 246 of storage compared to the initial count. YMC of control conditions reached  $4.7 \pm 0.1$  log  
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54 247 CFU/g.  
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248 The European Regulation (EUR-Lex - 32005R2073 - EN - EUR-Lex, 2005) sets a threshold  
1  
2 249 for the total microbial load of minimally processed vegetables equal to 7 log CFU/g.

3  
4 250 According to this limit, the shelf life of the packed carrots was comprised between 3 and 7  
5  
6 251 days, because of bacterial development. Different results are reported in literature with a  
7  
8 252 range of shelf-life between 6 and 10 days. Corbo *et al.* (Corbo *et al.*, 2004) found a shelf life  
9  
10 253 of shredded carrots treated with chlorinated water equal to 6 days considering a threshold of  
11  
12 254 7.7 log CFU/g. Alegria *et al.* (Alegria *et al.*, 2010) found that shredded carrots treated with  
13  
14 255 chlorinated water reached the European threshold value in 7 days, whereas Piscopo *et al.*  
15  
16 256 (Piscopo *et al.*, 2019) reported that shredded carrots stored in PP bags reached the limit  
17  
18 257 after 10 days of storage.

19  
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22 258 The comparison of the microbial counts of shredded carrots according to the different  
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24 259 packaging films showed that the presence of the active phases slightly reduced the TPC  
25  
26 260 (**Figure 2a**). The strongest inhibitory effect was observed for the film 15 wt% PS after 7 and  
27  
28 261 10 days. The TPC counts exceeded 7 log CFU/g between 3 and 7 days, as for the control,  
29  
30 262 but a difference of 0.8 log CFU/g was observed between the control and the carrots packed  
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32 263 in 15 wt% PS after 7 days. A similar tendency was observed for YMC of carrots packed  
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34 264 antimicrobial films, though with lower populations (**Figure 2b**). Only the film 15 wt% SB was  
35  
36 265 able to maintain YMC at a level not different from the control at day 0, whatever the storage  
37  
38 266 duration, and the fungal population after 7 days of storage was 3.4 log CFU/g, i.e. 1.3 log  
39  
40 267 CFU/g less than the control.

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43  
44 268 The results of carrot shelf-life tests suggested that the developed films were able to release  
45  
46 269 the antimicrobial agents, as they exerted their antimicrobial function on TPC reduction for the  
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48 270 film containing PS as active agent and on YMC for the film containing SB. No synergistic  
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50 271 effect between PS and SB was observed.

#### 51 272 *Effect of antimicrobial films on pineapple juice quality*

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53  
54 273 The active films were assessed in order to evaluate their impact on the quality of fresh  
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56 274 pineapple juice during storage up to 7 days at 4°C. Fresh pineapple juice is naturally acidic,  
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58 275 with a pH value comprised between 3.1 and 4.2 and its main organic acids are citric and  
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276 malic (Leneveu-Jenvrin et al., 2020). Hence, juice pH was suitable for PS and SB  
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2 277 antimicrobial effect. Until now, beverage active packaging has been mostly investigated for  
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4 278 shelf-stable beverages which require high barrier properties regarding oxygen (Palomero et  
5  
6 279 al., 2016; Ramos et al., 2015).

8  
9 280 **Table 2** shows that the pH value of pineapple juice in the control film did not significantly  
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11 281 change during storage, whereas a slight decrease of TA was noticed after 7 days. After 7  
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13 282 days of juice storage, pH values were not different according to the packaging film, but  
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15 283 differences in TA were observed. In fresh pineapple juice, TA variability is in the range 0.66-  
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17 284 1.35 g/100 mL (Leneveu-Jenvrin et al., 2020) and the values obtained in this study were  
18  
19 285 consistent with these data, except for juice stored for 7 days in 15 wt% SB film.

21  
22 286 The color of the juice stored in the control film changed during storage: a\* and b\* significantly  
23  
24 287 decreased (**Online Resource 2**), resulting in a color difference of  $23.8 \pm 1.2$  after 7 days  
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26 288 compared to the initial color (**Table 2**). This difference is high, as a difference above 5 can be  
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28 289 assigned to two different colors (Mokrzycki & Tatol, 2011). It is also consistent with previous  
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30 290 reports indicating color modification and browning during storage of pineapple juice  
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32 291 (Leneveu-Jenvrin et al., 2020). For the juice stored in the antimicrobial films, the decrease in  
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34 292 a\* and b\* was less pronounced than for the control film, leading to a color variation compared  
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36 293 to the initial juice in the range  $4.4 \pm 0.1 - 7.4 \pm 0.1$ . The color change was even less marked  
37  
38 294 for the juice stored in films with 15 wt% SB or 15wt% PS as active phase, compared to the  
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40 295 one in films with both the antimicrobial compounds, which behave differently with an increase  
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42 296 in L\*.

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44 297 **Figure 2c** and **Figure 2d** shows the development of enterobacteria and yeasts and molds  
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46 298 respectively in pineapple juice stored in control or antimicrobial films. For the control film, a  
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48 299 significant increase is observed after two days for EC and YMC. Both EC and YMC of the  
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50 300 control condition increased by 2 log CFU/g within the 7 days of storage. Initial counts and  
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52 301 population increases were consistent with previous observations (Leneveu-Jenvrin et al.,  
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54 302 2020). The comparison of microbial counts after two days of storage of juice whatever the  
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56 303 antimicrobial packaging film showed lower counts for the two microbial groups compared to  
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304 **the control one.** After 7 days of storage, the difference of counts between the control  
305 **condition** and juice packed into antimicrobial films represented 1.4-1.5 log CFU/g and 1.0-1.6  
306 log CFU/g, respectively for EC and YMC. **Each antimicrobial** proved to be of the same  
307 efficacy than their combination for the inhibition of yeast and molds and **of** enterobacteria.  
308 All together, these results showed that the use of biodegradable films containing  
309 antimicrobial compounds at a concentration of 15 wt% in the coating layer is strongly  
310 effective to limit microbial development in pineapple juice, but also to limit color modification,  
311 extending the juice's shelf life.  
312 The impact of the active films is much more pronounced on pineapple juice than on shredded  
313 carrots, likely because of the lower pH of pineapple compared to carrots that allowed to the  
314 active phases to better **exert** their antimicrobial action. Moreover, the diffusion of the  
315 antimicrobials, **which are non-volatile but water soluble, was probably favored by the liquid**  
316 **nature of the pineapple juice** (Wu et al., 2018). In addition, the ratio food quantity / film  
317 surface was more favorable for pineapple juice than for carrots. Among the tested  
318 antimicrobial compounds, films containing PS were more effective than SB or their  
319 combination to delay the microbial growth in both the analyzed food, in accordance with data  
320 literature reported for fresh noodles (Wangprasertkul et al., 2021).

### 321 *Film properties*

322 The main functional properties of the films and their heat-sealing ability were evaluated in  
323 order to analyze the effect of the incorporation of the active phases on the film's performance  
324 and thus on their suitability as food packaging materials. **Table 3** reports the mechanical,  
325 barrier and optical properties of the films with the highest levels of active phase. All the  
326 coated films showed values of the main mechanical parameters in the same range of  
327 PLA/PBAT based blends (Pietrosanto, Scarfato, Di Maio, & Incarnato, 2020; Pietrosanto,  
328 Scarfato, Di Maio, Nobile, et al., 2020) and comparable to those of polyethylene (PE)  
329 (Mangaraj et al., 2009), **one of the most used conventional polymers for flexible packaging**  
330 **applications.** The presence of the active phases did not significantly affect the stiffness of the  
331 films, while it led to a slight increase of the yield stress. Moreover, it caused a slight reduction

332 in the ductility of the films, more pronounced for the SB phase. It may be hypothesized that  
333 SB, having aromatic nature, and thus lower chemical affinity than the aliphatic PS towards  
334 the PLA matrix of the coating layer, could tend to aggregate into the polymer, creating small  
335 particles that act as stress concentration points and reduce the toughness. However, in all  
336 cases the change **was not large enough to compromise the ductile failure mode of the films**  
337 **(Table 4).**

338 Since the oxygen is one of the main factors that lead to food spoilage, the developed films  
339 were also tested for their oxygen permeability. All coated films showed comparable values of  
340 permeability coefficients (approx.  $40 \text{ (cm}^3 \times \text{mm)} / (\text{m}^2 \times \text{d} \times \text{bar})$ ), independently on the  
341 formulation of the active layer. These values are in the typical range **for** food packaging films,  
342 far above those of polyolefins, but lower than polyethylene terephthalate (PET) (Piergiovanni  
343 and Limbo, 2010). **Compared** to the uncoated substrate film, which permeability coefficient is  
344 equal to  $51.5 \text{ (cm}^3 \times \text{mm)} / (\text{m}^2 \times \text{d} \times \text{bar})$ , **a significant improvement of the oxygen barrier**  
345 **performance of the coated films was observed.** This finding can be **explained by** the lower  
346 molecular mobility of the polymer chains of the coating layer (PLA, at glassy state at 23°C)  
347 with respect to those of the substrate (PLA/PBAT blend, where the PBAT constituent is in the  
348 rubbery state at 23°C) (Apicella et al., 2019; Pietrosanto, Scarfato, Di Maio, Nobile, et al.,  
349 2020). **On the opposite**, the addition of both the preservatives did not lead to significant  
350 changes **of oxygen permeability of coated films.** Different results were reported in literature  
351 on other polymer substrates (Wangprasertkul et al. 2021), where the incorporation of PS and  
352 SB, which are polar molecules, prevented the permeability to oxygen molecules.

353 Films optical properties in terms of “see through” possibility were investigated by measuring  
354 the transparency, *i.e.* the transmission of visible light at 560 nm ( $\text{TR}_{560}$ ). The transparency of  
355 the developed films was lower than the conventional used polymers (e.g. PE and PET)  
356 (Moreno-Vásquez et al., 2017; Nogi et al., 2013), due to the presence of **PBAT, which is**  
357 **white and opaque, in the substrate** (Wang et al., 2016). Moreover, the addition of the  
358 preservatives led to a further reduction **of** the transparency value, which was more significant  
359 for SB than **for** PS. **The** transparency of a polymeric system increases with the increase of

360 the dispersed composite size (Schulz et al., 2007). Thus, the lower transparency of 15SB film  
361 can be attributable to the lower compatibility of SB active agent with PLA compared to PS,  
362 that caused a worse dispersion in the polymer matrix. Since the antimicrobial activity is  
363 affected by the dispersion of the active phase in the polymer matrix (Kuplennik et al., 2015),  
364 this result can also explain the lower antimicrobial effect on both the shredded carrots and  
365 pineapple juice that was observed for films containing SB comparatively to those containing  
366 PS.

367 Finally, to investigate the capability of the coating layer to provide the heat-sealing ability of  
368 the films, hot-tack tests were carried out. Packaging systems able to self-adhere at a  
369 convenient processing condition and provide a flawless hermetic seal are of fundamental  
370 importance for food packaging applications, because it ensures the tightness of the package  
371 and thus the protection of the food. Proper heat-sealing ability also makes the packaging  
372 suitable for applications in aseptic processing and packaging where special technologies as  
373 hot-filling or steam sterilization are used. In the case of the films developed in this work, heat-  
374 sealing ability is made possible by the use of an amorphous PLA coating layer able to self-  
375 adhere. Table 4 reports the results of the hot tack tests. As it can be seen, the addition of the  
376 PLA coating layers to the substrate, which is not sealable, allowed to impart to all of them  
377 heat-sealing ability. In the case of the control film sample, the hot seal strength is 383 g/15  
378 mm and the failure happens mainly through a delamination mechanism, which means that  
379 the adhesion of the PLA coating layer to the substrate was lower than its seal strength.

380 The incorporation of the preservatives in the coating layer affected both the seal strength and  
381 the failure type. The seal strength of the active films was lower than that of the control one:  
382 the presence of active agent salts in the polymer matrix hindered the interactions between  
383 the two melted surfaces of the PLA layer during the heat-sealing process. The seal strength  
384 was so much reduced that it became lower than the adhesion of the PLA coating to the  
385 substrate, resulting in an adhesive failure mechanism of the active films. In addition, films  
386 containing PS had a lower seal strength than those containing SB, because the better

387 compatibility and dispersion of PS into the polymer matrix might have exerted a greater  
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2 388 hindering effect on the interaction between the PLA layers (Voon et al., 2012).  
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389 **Conclusion**

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2 390 Multifunctional, eco-friendly active packaging films were developed by coating a  
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4 391 biodegradable PLA/PBAT substrate with a heat-sealable layer loaded with antimicrobial  
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6 392 agents. The incorporation of the active phases did not relevantly affect the mechanical and  
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8 393 barrier properties of the films, while it led to a slight reduction of transparency and hot-tack  
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10 394 strength. Only the films at highest level of active agents showed effective antimicrobial  
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12 395 activity in shelf-life test performed on shredded carrots and fresh pineapple juice.  
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14 396 The film with PS was very efficient to reduce microbial development and color modification of  
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16 397 pineapple juice. A better dispersion of PS in the polymer matrix probably explains its better  
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18 398 performance compared to films containing SB or a combination of both antimicrobials.  
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20 399 The better impact on shelf-life of pineapple juice compared to shredded carrots, probably  
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22 400 owed to the liquid state and the lower pH of pineapple juice, that may have favored both the  
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24 401 diffusion of the antimicrobial into the food and the antimicrobial activity of the active  
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26 402 compounds. In conclusion, these results open new possibilities for the development of  
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28 403 sustainable active packaging for beverages with a short shelf-life.  
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**Table 1.** pH value, titratable acidity (TA, g/100g) and color difference of shredded carrots packed in the different films during refrigerated storage up to 10 days. Control: no active phase, 15SB: sodium benzoate at 15 wt% in the coating layer, 15PS: potassium sorbate at 15 wt% in the coating layer, 15SB+15PS: sodium benzoate and potassium sorbate in a 50/50 mass ratio at 15 wt% in the coating layer. Means and standard deviations of triplicates are indicated. Values in the same column with different letters show significant differences at  $p < 0.05$ .

Film	Days of storage	pH	TA	$\Delta E$
Control	0	4.1 ± 0.1 b	3.5 ± 0.3 a	0 ± 0 c
Control	3	5.0 ± 0.4 a	2.0 ± 0.8 ab	8.5 ± 5.3 ab
15SB	3	4.8 ± 0.1 ab	2.6 ± 0.4 ab	10.3 ± 11.4 ab
15PS	3	4.7 ± 0.3 ab	2.4 ± 0.6 ab	6.9 ± 3.7 abc
15PS+SB	3	4.8 ± 0.3 ab	2.3 ± 0.6 ab	7.1 ± 6.7 abc
Control	7	5.1 ± 0.2 a	2.2 ± 0.1 ab	3.8 ± 4.5 ab
15SB	7	4.8 ± 0.2 ab	2.9 ± 0.2 ab	4.8 ± 2.2 abc
15PS	7	4.8 ± 0.3 ab	2.4 ± 0.1 ab	6.8 ± 8.3 abc
15PS+SB	7	4.7 ± 0.1 ab	2.8 ± 0.1 ab	10.3 ± 6.8 ab
Control	10	5.1 ± 0.1 a	1.7 ± 0.4 b	6.3 ± 4.9 abc
15SB	10	4.6 ± 0.1 ab	3.0 ± 0.5 ab	4.5 ± 1.7 abc
15PS	10	4.7 ± 0.1 ab	2.9 ± 0.1 ab	8.3 ± 4.0 abc
15PS+SB	10	4.5 ± 0.2 ab	2.9 ± 0.9 ab	14.8 ± 4.9 a

**Table 2.** pH value, titratable acidity (TA, g/100g) and color difference of fresh pineapple juice packed in control film or in antifungal films and stored up to 7 days at 4°C. Means and standard deviations of triplicates are shown. Different letters in the same column indicate significant differences (p-value < 0.0001). Control: no active phase, 15SB: sodium benzoate at 15 wt% in the coating layer, 15PS: potassium sorbate at 15 wt% in the coating layer, 15SB+15PS: sodium benzoate and potassium sorbate in a 50/50 mass ratio at 15 wt% in the coating layer.

<b>Film</b>	<b>Days of storage</b>	<b>pH</b>	<b>TA</b>	<b>ΔE</b>
Control	0	3.7 ± 0.1 ab	0.92 ± 0.10 a	0 ± 0 i
Control	1	3.6 ± 0.1 ab	0.92 ± 0.10 a	9.1 ± 0.5 c
15SB	1	3.6 ± 0.1 ab	0.90 ± 0.10 a	7.5 ± 1.7 cde
15PS	1	3.6 ± 0.1 ab	0.91 ± 0.10 a	3.2 ± 1.5 h
15PS+SB	1	3.7 ± 0.1 ab	0.92 ± 0.10 a	6.4 ± 1.4 gh
Control	3	3.6 ± 0.1 ab	0.92 ± 0.10 a	14.4 ± 1.1 b
15SB	3	3.7 ± 0.1 ab	0.90 ± 0.10 a	8.9 ± 0.2 cd
15PS	3	3.8 ± 0.1 a	0.91 ± 0.10 a	6.3 ± 0.2 defg
15PS+SB	3	3.5 ± 0.1 b	0.91 ± 0.10 a	5.8 ± 0.2 efgh
Control	4	3.7 ± 0.1 ab	0.92 ± 0.10 a	22.6 ± 1.0 a
15SB	4	3.5 ± 0.1 b	0.90 ± 0.10 a	5.7 ± 0.9 efgh
15PS	4	3.6 ± 0.1 ab	0.90 ± 0.10 a	3.7 ± 0.2 gh
15PS+SB	4	3.6 ± 0.1 ab	0.91 ± 0.10 a	5.6 ± 0.2 efgh
Control	7	3.5 ± 0.1 b	0.85 ± 0.10 b	23.8 ± 1.2 a
15SB	7	3.7 ± 0.1 ab	0.45 ± 0.10 d	4.5 ± 0.1 fgh
15PS	7	3.7 ± 0.1 ab	0.76 ± 0.10 c	4.4 ± 0.1 fgh
15PS+SB	7	3.6 ± 0.1 ab	0.84 ± 0.10 b	7.4 ± 0.1 cde

**Table 3.** Film composition, thickness, and functional properties of the films: Elastic modulus (E), yield stress ( $\sigma_y$ ), elongation at break ( $\epsilon_b$ ), oxygen permeability coefficient ( $PO_2$ ), and transparency at 560 nm ( $TR_{560}$ ). Means and standard deviations of triplicates are shown. Different letters in the same column indicate significant differences (p-value < 0.0001). Control: no active phase, 15SB: sodium benzoate at 15 wt% in the coating layer, 15PS: potassium sorbate at 15 wt% in the coating layer, 15SB+15PS: sodium benzoate and potassium sorbate in a 50/50 mass ratio at 15 wt% in the coating layer.

Film	Film	Active phase	Concentration of active phase (wt%)	Coating layer thickness ( $\mu\text{m}$ )	Total film thickness ( $\mu\text{m}$ )	E (MPa)	$\sigma_y$ (MPa)	$\epsilon_b$ (%)	$PO_2$ ( $\text{cm}^2 \times \text{mm}) / (\text{m}^2 \times \text{d} \times \text{bar})$	$TR_{560}$ (%)
Control	Control	None	0	12.6 ± 3.1	35.6 ± 3.2	290 ± 18 a	6.4 ± 1.5 a	305 ± 19 a	38.5	11.0 ± 1.2
15SB	15SB	SB	15	14.0 ± 3.0	37.0 ± 3.1	295 ± 31 a	7.3 ± 1.5 a	263 ± 26 b	39.1	6.4 ± 0.7 b
15PS	15PS	PS	15	13.3 ± 3.6	36.3 ± 3.7	292 ± 20 a	7.2 ± 1.4 a	290 ± 25 ab	40.3	10.9 ± 0.5
15PS+SB	15PS+SB	PS+SB (50/50 w/w)	15	13.9 ± 2.3	36.9 ± 2.4	293 ± 26 a	7.1 ± 1.6 a	270 ± 34 b	40.9	7.3 ± 0.9 b

**Table 4.** Hot tack strength (according to ASTM F 1921 – 98) of the films.

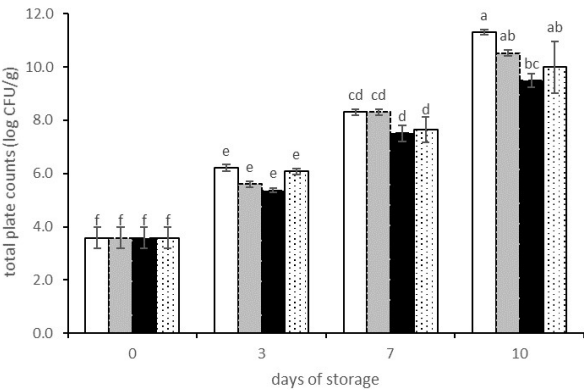
<b>Film</b>	<b>Hot tack strength (g/15 mm)</b>	<b>Failure mode</b>
Uncoated substrate	Not sealable	-
Control	383 ± 14	Delamination
15SB	283 ± 29	Adhesive
15PS	245 ± 25	Adhesive
15PS+SB	267 ± 29	Adhesive

**Figure 1**

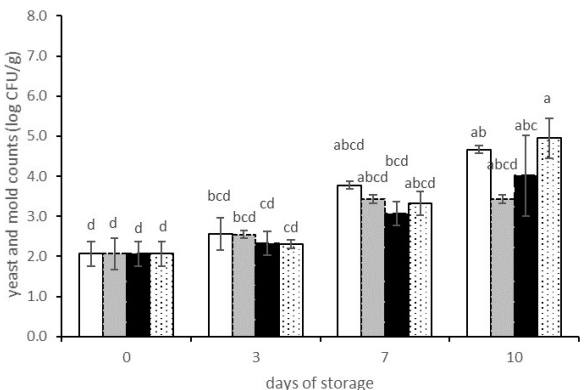


**Figure 2**

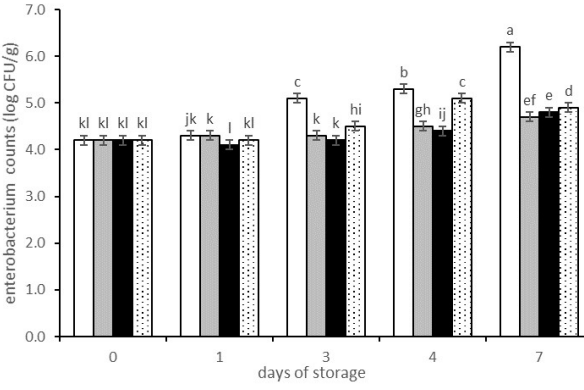
(a)



(b)



(c)



(d)

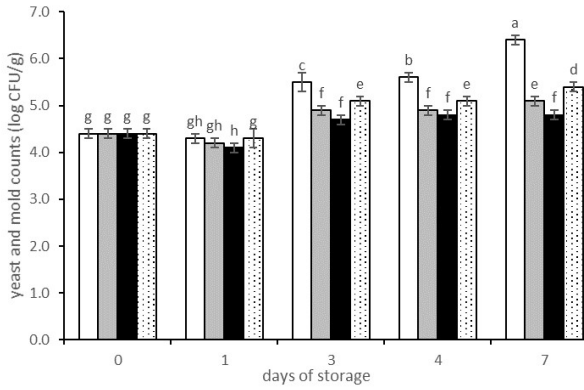




Figure captions

**Fig. 1** Visual appearance of shredded carrots packed into the films containing antimicrobials after 7 days of storage. Arrows indicate black or white spots on the carrot surface. Film names: Control, no active phase; 15SB, sodium benzoate at 15 wt% in the coating layer; 15PS, potassium sorbate at 15 wt% in the coating layer; 15SB+15PS, sodium benzoate and potassium sorbate in a 50/50 mass ratio at 15 wt% in the coating layer.

**Fig. 2** Total aerobic plate counts of shredded carrots (a), yeast and mold counts of shredded carrots (b), enterobacterium counts of pineapple juice (c) and yeast and mold counts of pineapple juice (d), packed in the control film (white bars), in 15SB (grey bars), in 15PS (black bars), in 15SB+15PS (spotted bars) during refrigerated storage. Film names: Control, no active phase; 15SB, sodium benzoate at 15 wt% in the coating layer; 15PS, potassium sorbate at 15 wt% in the coating layer; 15SB+15PS, sodium benzoate and potassium sorbate in a 50/50 mass ratio at 15 wt% in the coating layer. Means and standard errors are indicated. **Same letter in a graph indicates no significant difference** at  $p < 0.05$ .