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This is the author's manuscript

Original Citation:

Availability:

This version is available <http://hdl.handle.net/2318/138839> since 2015-12-01T14:20:15Z

Published version:

DOI:10.1080/19476337.2012.745096

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Effect of different coating in preventing deterioration and preserving the quality of fresh-cut nectarines cv Big Top

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Fresh-cut nectarines are very perishable product, and thus have a short shelf-life. In this study, minimally processed nectarines (cv Big Top) were treated with chitosan (20 g/L), sodium alginate (20 g/L) and Chitoplant[®] (20 g/L), sealed in polypropylene plastic bags and stored at 4°C for 5 days. Effects of the coatings on nectarine slices shelf-life and qualitative attributes were evaluated by investigating changes in colors, browning potential, total soluble solids content and titratable acidity. Changes in quality parameters were lower in coated samples as compared with the control. The alginate coatings were effective on delaying the evolution of the parameters related to postharvest ripening, such as color (Hue, L*) and loss of acidity. The application of sodium alginate coating could be used to reduce deteriorative processes, maintain quality and improve the shelf-life of fresh-cut nectarine stored at 4°C.

Keywords: fresh-cut nectarines; edible coating, quality attributes.

Introduction

Minimal processing has been defined as a combination of procedures, such as washing, sorting, trimming, peeling, and slicing or chopping, that do not affect the fresh-like quality of the food. Ready-to-eat fruits and vegetable market has rapidly grown in recent years due to the health benefits associated with these foods because of busy lifestyles, increasing purchasing power, and

increasingly health-conscious consumers. Nevertheless, because the tissular integrity of fruits is more easily altered during processing, ready-to-use commodities are more perishable than the original materials (Oms-Oliu et al., 2010).

The main factors affecting the loss of consumer acceptability are discoloration, enzymatic browning, dryness and texture loss (Pérez-Gago, González-Aguilar & Olivas, 2005). These parameters determine the visual appearance (Pace, Cefola, Renna & Attolico, 2011). For this reason the fruit processing industry requires the development of techniques able to keep safe shelf-life and preserve the original visual and organoleptic fresh-like characteristics of fresh-cut produces.

The commercial success of fresh-cut peach and nectarine slices has been limited due to their short shelf-life, because of cut surface browning, excessive flesh softening and pit cavity breakdown (Gorny, Hess-Pierce & Kader, 1998). For these reasons makes essential for finding treatments to slow down the browning and ripening process and prolonging their shelf-life. Several treatments have been studied in order to maintain quality and to extend the shelf-life of fresh-cut peach and nectarine like heat treatment (Steiner et al., 2006), chemical treatments (Gorny, Hess-Pierce & Kader, 1999; Costa da Costa, Antunes, Valmor Rombaldi & Arocha, 2011; Zhu, Zhou, Zhu & Guo, 2009), modified atmosphere packaging (Zoffoli, Rodriguez, Aldunce & Crisosto, 1997; Palmer-Wright & Kader, 1997) and edible coatings application (Du, 1997; Ruoyi, Zhifang & Zhaoxin, 2005).

Edible coatings, a new strategy used to extend shelf-life and to improve food quality of whole fruits and fresh-cut fruits, have been applied to many products. Coatings on minimally processed products create a semipermeable barrier to external elements that can reduce moisture loss, solutes migration, respiration and oxidative reaction rates (Barbosa, Dias de Mello Castanho & Rodrigues Monteiro, 2011; Vargas, Pastor, Chiralt, McClements & González- Martínez, 2008). Chitosan has been one of the most promising coating materials for fruits because of its good film-forming property, broad antimicrobial activity, and excellent compatibility with other substances.

Furthermore, chitosan films are tough, highly transparent, long-lasting and flexible (Cé, Cacicano & Brandelli, 2012). Other popular edible coating formulations were polysaccharide-based like alginate or protein-based polymers (Diaz-Mula, Serrano & Valero, 2011). Maintenance of fruit quality has been achieved by using chitosan in peach (Li & Yu, 2001), strawberries (Vu, Hollingsworth, Leroux, Salmieri & Lacroix, 2011) and papaya (Asgar, Tengku, Kamaruzaman & Yasmeen, 2011), pectin coating in melon (Ferrari, Sarantópoulos, Carmello-Guerreiro & Hubinger, 2011), alginate in apple (Rojas-Graü, Tapia, Rodríguez, Carmona & Martin-Belloso, 2007; Olivas, Mattinson & Barbosa-Cánovas, 2007) and in papaya (Tapia et al., 2008).

The objective of this work were to evaluate the effectiveness of different coatings in order to prolong shelf-life of nectarine slices, and to study the effect of the best treatment on the quality of fresh-cut nectarine slices stored in modified atmosphere packaging, as the commercial practice.

Materials and methods

Material

Nectarines (*Prunus persica* (L.) Batsch., cv Big Top) were harvested at commercial maturity, and stored (4°C and 95% RH) before processing. The fruits were peeled, cored and cut into 5-mm-thick slices using a hand-operated slicer.

Methods

Sample preparation

Three different coating solutions were prepared: (1) 20 g/L chitosan (90–95% of deacetylated degree and 90–100 mPa.s of viscosity, Sigma-Aldrich Co., Steinheim, Germany), (2) 20 g/L sodium alginate (Sigma-Aldrich Co., Steinheim, Germany), (3) 20 g/L Chitoplant[®], containing 95% chitosan, 2.5% boron and 2.5 % zinc (Agritalia, Italy), a commercial chitosan formulation. Nectarines slices used as controls were not treated with coating.

(1) Film-forming solutions were prepared by dissolving chitosan (20 g/L) powders in an aqueous solution of 20 g/L citric acid. Then, the solution was added with 500 ml/L glycerol and 1,5 ml/L Tween 20 to improve the wettability (Duan, Wu, Strik & Zhao, 2011). The mixture was homogenized for 1 min and stirred in a 60 °C water bath for 30 min, followed by cooling to room temperature.

(2) Film-forming solutions were prepared by dissolving sodium alginate (20 g/L) powders in deionized water while heating on a stirring hot plate for 10 min at 70 °C until the mixture became clear (Chiumarelli, Ferrari, Sarantópoulos & Hubinger, 2011). Coating solutions also contained glycerol (15 ml/L) as an emulsifier and calcium chloride (20 g/L) (Sigma-Aldrich Co., Steinheim, Germany). Calcium chloride was used to induce crosslinking reaction (Rojas-Graü, Tapia, Rodríguez, Carmona & Martin-Belloso, 2007; Tapia et al., 2008).

(3) Film-forming solutions were prepared by dissolving Chitoplant[®] (20 g/L) powders in deionized water with continuous shaking until the solution became clear. No plasticizer was added.

Slices were immersed in the coating solutions for 1 min at 20°C and were then allowed to drip off. Fruits dipped in sodium alginate based coating, in addition, were immersed in calcium chloride for 30 s. Control fruits were dipped in distilled water.

25 slices were randomly selected and packaged in polypropylene plastic bags (20cm x 30 cm size and 90 µm thickness) with 50 cm³ O₂/m²/bar/day, 150 cm³ CO₂/m²/bar/day and 2.8 g/m²/bar/day water vapor transmission rate (I.Plast, Italy). The packages were completely sealed (UNIMEC packaging systems, Italy) and then were stored at 4°C and 95% RH in darkness for 5 days.

Analysis

Gases composition

The concentrations of oxygen and carbon dioxide inside the packages were monitored daily by sampling (0.5 ml) the headspace using a CANAL 121 (Vizag, Gas Analysis, France). A syringe was inserted into the package through a rubber seal placed on the film. Gases were analysed with an electrochemical sensor for O₂ level and an infrared sensor for CO₂ level. The instrument was calibrated towards air. Results were expressed as kPa of O₂ and CO₂ inside the bags.

Quality measurements

Color analysis were performed at 0, 1, 2 and 5 days of cold storage at 4°C.

L*, a*, and b* values were determined at two points along each side of the cut surface using a Minolta chroma-meter (CR400; Minolta, Ramsey, NJ, USA). The flesh color was also expressed as whiteness index ($WI = 100 - [(100 - L^*)^2 + (a^*)^2 + (b^*)^2]^{1/2}$) according to Bolin and Huxsoll (1991).

Quality analysis were performed at 0 on the nectarines whitout coating and after 5 days of cold storage at 4°C in all the treatments. Total soluble solids content (TSS) was determined in the juice from 25 slices with a digital refractometer Atago PR-101 (Atago, Japan) at 20 °C and results expressed as °Brix. Titratable acidity (TA) was determined by titration with 0.1 N NaOH up to pH 8.1, using 10 ml of diluted juice in distilled H₂O and results were expressed as meq/L.

Browning potential

Browning potential was determined according to the method of Arias, Gonzalez, Lopez-Buesa and Oria (2008) at day 0, and after 1 and 5 days of storage. The extract was obtained as followed: nectarine slices (25g) from each treatment were homogenized for 2 minutes at 13,500 rpm with an Ultra-Turrax T25 (IKAs WERKE, Germany), the homogenates were centrifuged (Centrifuge AVANTITM J-25, Beckman Instruments Inc., Fullerton, Calif., U.S.A.) at 4000 rpm for 10 min and the supernatant was filtered through Whatman 4 filter paper (Whatman Intl., U.K.). The absorbance

of the clear juice was then measured spectrophotometrically (Hitachi, U-5100, Japan) at 440 nm to determine browning potential (BP). This measurement was replicated three times.

Statistical analysis

The basic experimental design consisted of three coating treatments, each having three replicates. In sliced nectarine measures, a package contained 25 slices, which was equivalent to two nectarines, was considered a replicate. Data were analyzed by analysis of variance using statistical procedures of the STATISTICA ver. 6.0 (Statsoft Inc., Tulsa, OK, USA). The sources of variance being coating treatments. Tukey's test HSP (honestly significant differences) was used to determine significant differences among treatment means. Means values were considered significantly different at $P \leq 0.05$.

Results and discussion

Gases composition

Significant differences were observed between coated and uncoated nectarine slices regarding the composition of O_2 and CO_2 in the head space along the evaluated period (Fig. 1-2). Higher O_2 levels were found in alginate coating slices, lower in chitosan coating slices. This result, probably depend on the different barrier effect and mechanical properties of the coatings solutions used (Chiumarelli & Hubinger, 2012). Moreover, the application of coatings in fresh-cut products can create a modified atmosphere around each piece, reducing the respiration rate and the metabolic processes (Rojas-Graü, Tapia & Martín-Belloso, 2008).

CO_2 production was lower in alginate-coated slices with respect to control ones, especially in the first 2 days of storage (Fig. 2), in agreement with previous reports in peaches coated with chitosan, methylcellulose or alginate (Li & Yu 2001; Maftoonazad, Ramaswamy & Marcotte, 2008). Alginate coating acts as barrier to gas exchange, so there is less oxygen available for plant

tissues respiration, resulting in lower release of carbon dioxide by the product, and, consequently, reducing the metabolic processes (Rojas-Graü, Tapia, Rodríguez, Carmona & Martín-Belloso, 2007). On the contrary, CO₂ production was significantly higher in Chitoplant[®] treated slices.

Quality measurements

Cut surface browning is a major problem for several minimal processed fruits. On non-treated samples a darkening effect on the color of cut slices was noted, when compared to treated samples, in according with instrumental color evaluation (Hue angle).

During storage, a reduction in Hue angle was observed in all the treatments (Fig. 3) and in control slices. The highest values were obtained in slices treated with sodium alginate, with final value of 95.12, showing a reduction of the ripening process.

No significant differences were observed in those slices treated with 2% chitosan, 2% chitoplant and control respect to color evolution during cold storage. A sharp decrease in Hue angle would indicate an over-ripe and senescence process of fruit, which is considered as detrimental (Rojas-Graü, Tapia & Martín-Belloso, 2008). On the contrary, the alginate coated fruits (with higher Hue angle) maintained the typical bright color of recently processed fruits.

Fruit color is crucial in purchase decisions, especially if the product is packaged and cannot be touched or smelled. During storage the L* value decreased for all the control and treated fruits (Fig. 4). However, a smaller decrease was observed for the fresh-cut fruit treated with sodium alginate and control. Lightness was better preserved by the treatment with sodium alginate coating. On the contrary, chitosan coating slices showed the lowest L*. The lower L* level could be also correlated with the reduction of O₂ and increase of CO₂ inside the packages and with a greater deterioration of the fruit (Fig. 1). A rapid deterioration of the fruit involved an increase in respiration rate and enzymatic metabolic processes that led to a loss of quality of fresh-cut produce (Gonzalez-Aguilar et al., 2009).

The b^* value was used to estimate changes in the yellow color during the storage time (Gonzalez-Aguilar et al., 2009). Control fruit showed the lowest b^* values, high values of coating fruits indicated that chitosan and alginate treatments were able to maintain the highest b^* values during the storage time. The severity of surface whitening for the coated and uncoated nectarine slices was quantitatively expressed by WI scores at different treatments (Fig. 6). Higher WI scores indicated greater development of surface whiteness (Gounga, Xu, Wang & Yang, 2008). For uncoated slices, the WI score range from 49.4 to 47.1 while the chitosan coating showed the lowest WI score of 39.11, at the end of storage period. This result is related with L^* values and with the result of previous work (Raybaudi-Massilia, Mosqueda-Melgar & Martín-Belloso, 2008; Vargas et al., 2009) and could be due to the film forming effect of chitosan on surface pores of the samples, which limits water losses. Changes in WI was also reported by Aguayo, Allende and Artés (2003) in “Piel de sapo” melon and others varieties, indicating that WI decreased when translucency injury increased on fresh-processed melon, as a consequence of a physiological disorder characterized by dark and glassy flesh.

TSS at harvest was 8.2° Brix with significant effect ($P \leq 0.05$) of coating treatments during storage on nectarine slices. The Chitoplant[®] coating induced a significant ($P \leq 0.05$) decrease in TSS content and titratable acidity values compared to chitosan treated slices. The faster decrease in acidity gave rise to a faster senescence (Asgar, Tengku, Kamaruzaman & Yasmeen, 2011). On the contrary, the chitosan and alginate coatings were significantly ($P \leq 0.05$) effective in delaying the loss of acidity, which occurred during cold storage (titratable acidity at harvest was 97.98 meq/l). The levels of titratable acidity were correlated with the antibrowning efficacy of the treatments. Han, Zhao, Leonard and Traber (2004) reported that in raspberry and strawberry, the chitosan coatings slowed down the changes in titratable acidity, effectively delaying fruit ripening. The effects of edible coating on decreasing acidity losses have been also found in chitosan and alginate coated peaches (Li & Yu 2001; Maftoonazad, Ramaswamy & Marcotte, 2008). The effect of

coating on acidity retention could be a result of the lower respiration rate found in coated fruits, since organic acids are substrates for many reactions during aerobic respiration in plant cell. In chitosan coating significant ($P \leq 0.05$) higher acidity values are also due to the effect of citric acid utilized for film-forming solution.

Table 1 shows pH variations during storage for control and coated samples. As expected, since coated fruits showed less variation in titrable acidity, the associated variation in their pH was also relatively lower and no significant (pH at harvest was 5.48). At the end of storage, control nectarines had significant higher pH than coated fruits, confirming some previous results (Togrul & Arslan, 2004).

Browning Potential

Browning potential was monitored in all the samples. An increase in browning potential was found throughout storage (Fig.7) in according with other works (Martin-Diana et al., 2007; Castaner, Gil, Ruíz & Artes 1999). Control samples presented the highest and Chitoplant[®] the lowest values (Fig. 7). The application of Chitoplant[®] coating proved to be the treatment that was the most effective in reducing browning index of nectarine slices. This could be due to the inactivation of the enzymes by the use of coating and to the lower viscosity of Chitoplant[®] (Martin-Diana et al., 2007, Romanazzi et al., 2009).

Chitosan coatings were able to decrease browning of fruit by reducing polyphenol oxidase and peroxidase activities, these changes were directly related to color changes of the flesh of different fruits and vegetables. A chitosan coating form a protective barrier on the surface of the fruit and reduce the supply of oxygen for enzymatic oxidation of phenolics (Zhang & Quantick, 1998).

Conclusion

The use of edible coatings can help maintain desirable quality characteristics of fresh-cut Big Top nectarine slices. The coated slices maintained their initial color during the refrigerated storage, in particular, the samples treated with alginate coating. Moreover, coatings reduced browning potential and decay of fresh-cut nectarine, in particular with the use of Chitoplant[®]. The alginate coatings were effective on delaying the evolution of the parameters related to postharvest ripening, such as color and loss of acidity, which could be explained by the gas barrier provided by the coatings creating a modified atmosphere in the fruit. On the contrary, chitosan coating was not effective on delaying color browning, in particular the lightness and the whiteness. According to the results obtained in this study, sodium alginate (20 g/L) and Chitoplant[®] appears to be a promising conservation alternative for fresh-cut nectarine.

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Table

Table 1. Changes in total soluble solid content (°Brix), titratable acidity (meq/l) and pH of nectarines coated with Chitoplant® (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L) after cold storage (4°C and 95% RH).

days of storage	treatment	quality parameters		
		total soluble solids	titratable acidity	pH
day 0	control	8.2 ± 0.2	47.98 ± 0.3	5.48 ± 0.3
day 5	control	7.8 ± 0.2 b	27.73 ± 2.5 c	5.79 ± 0.5 a
	chitosan	8.7 ± 0.1 a	44.47 ± 3.2 a	5.56 ± 0.3 b
	chitoplant	6.8 ± 0.1 bc	22.34 ± 2.7 c	5.63 ± 0.6 b
	alginate	7.4 ± 0.2 b	36.79 ± 2.4 b	5.54 ± 0.5 b

Data are the mean ± SE (n=25). Mean values followed by the same letter in column are not significantly different at $P \leq 0.05$ level.

Fig. 1. Evolution of O₂ (KPa) during cold storage on nectarines coated with Chitoplant® (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L).

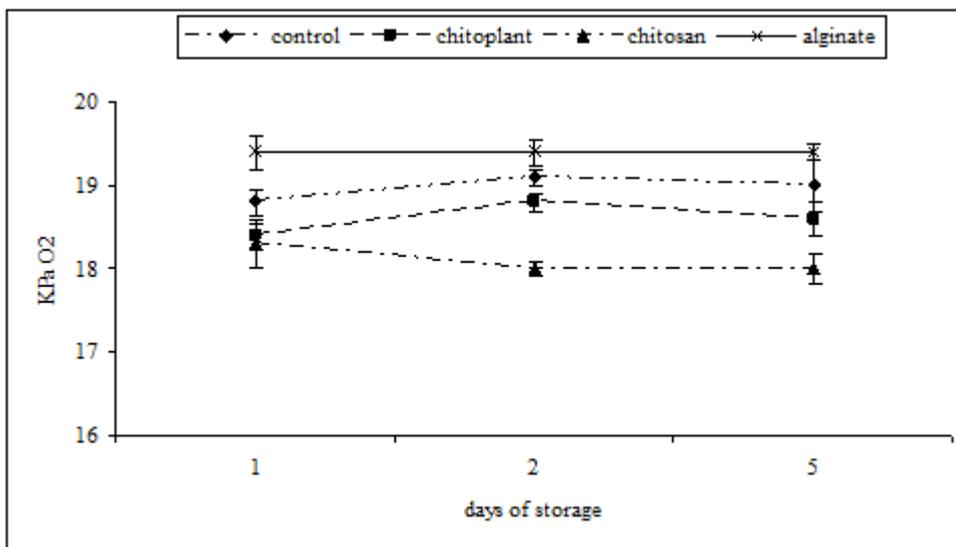


Fig. 2. Evolution of CO₂ concentration (KPa) during cold storage on nectarines coated with Chitoplant[®] (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L).

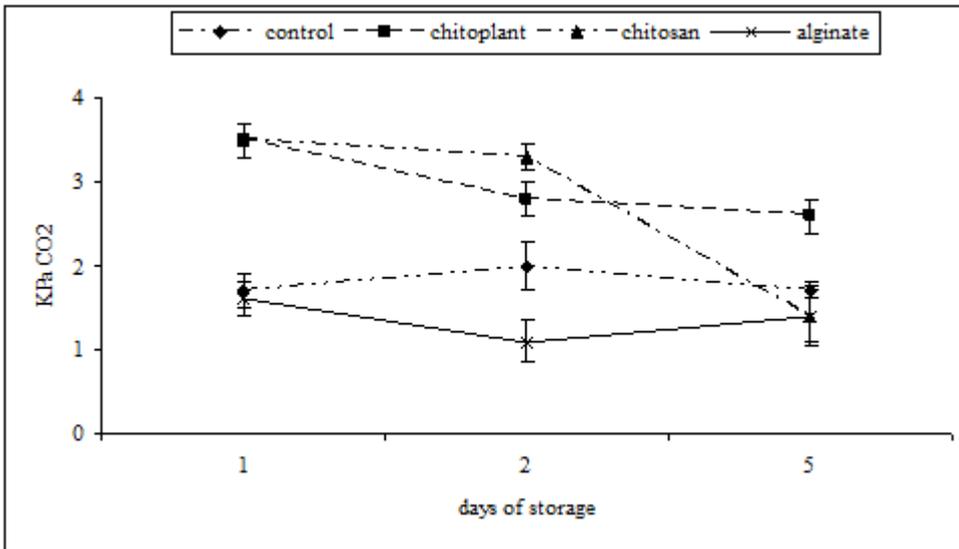


Fig. 3. Evolution of color (Hue angle) during cold storage on nectarines coated with Chitoplant[®] (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L). Data are the mean \pm SE (n=25).

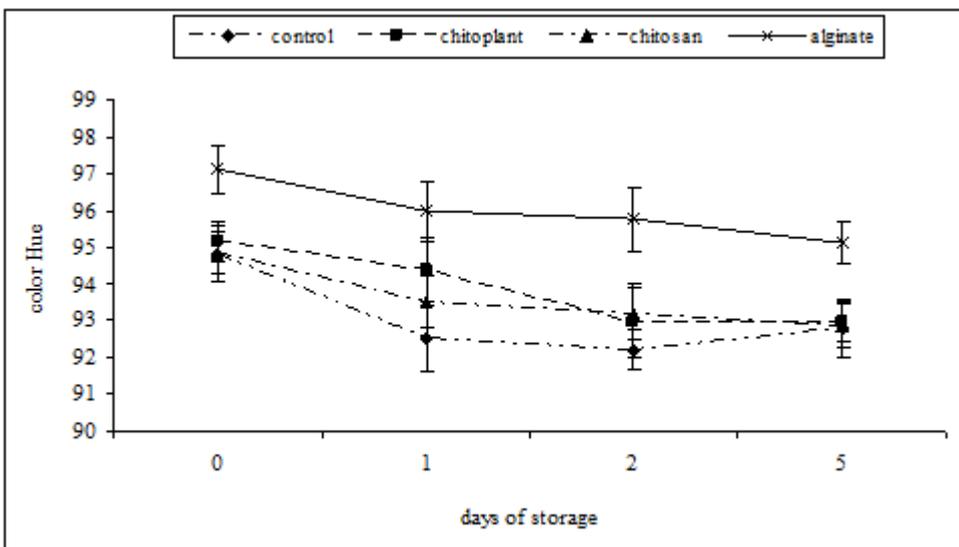


Fig. 4. Evolution of color (L^*) during cold storage on nectarines coated with Chitoplant[®] (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L). Data are the mean \pm SE (n=25).

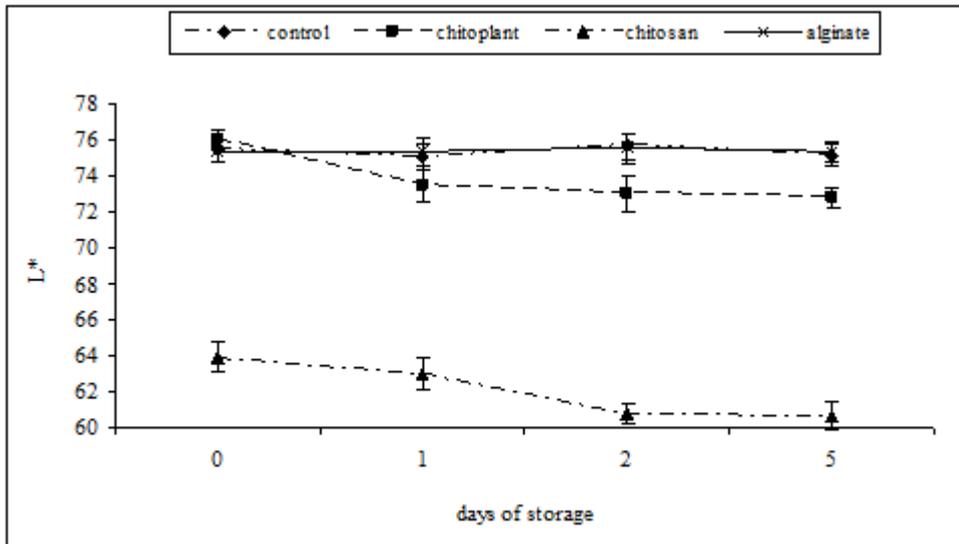


Fig. 5. Evolution of color (b^*) during cold storage on nectarines coated with Chitoplant[®] (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L). Data are the mean \pm SE (n=25).

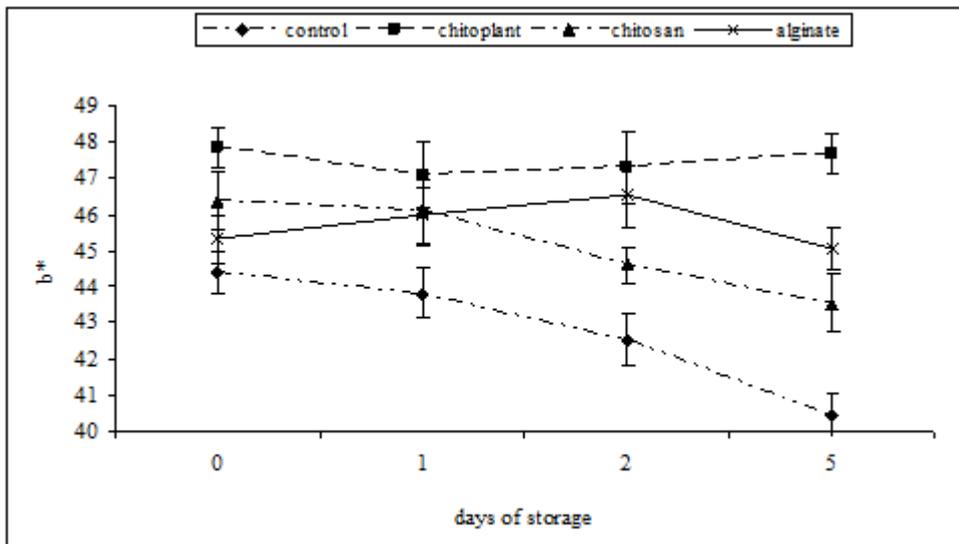


Fig. 6. Evolution of color (Whiteness Index) during cold storage on nectarines coated with Chitoplant® (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L). Data are the mean \pm SE (n=25).

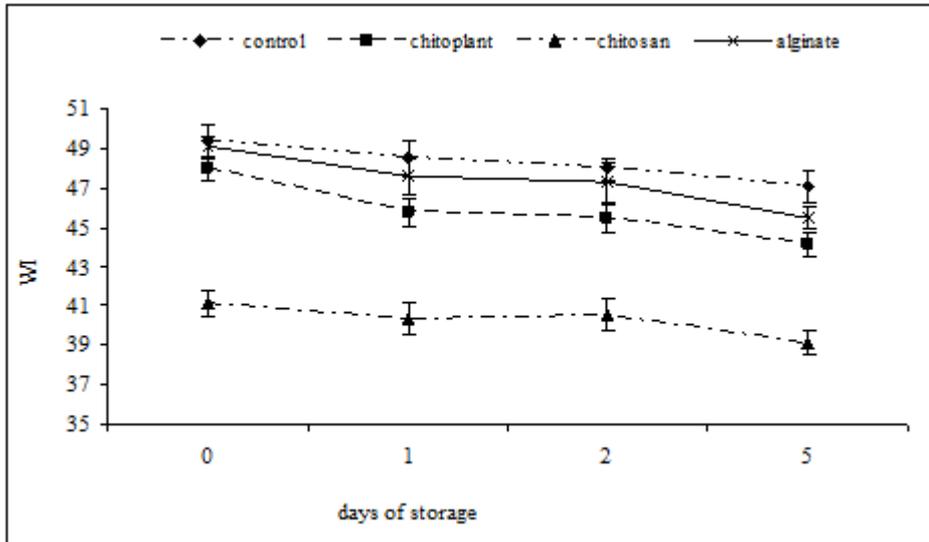


Fig. 7. Browning Potential values during cold storage on nectarines coated with Chitoplant® (20 g/L), chitosan (20 g/L) and sodium alginate (20 g/L). Data are the mean \pm SE (n=25).

