

## Effect of Different Sanitizers on Microbial, Sensory and Nutritional Quality of Fresh-Cut Jalapeno Peppers

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**Abstract: Problem statement:** Sanitation is a critical step to insure safety of fresh-cut produce. The inadequacies of chlorine, currently used as a sanitizer, have stimulated interest in finding safer, more effective sanitizers, however little is known on the impact of these novel sanitizers on sensory and nutritional quality of the treated products. **Approach:** The effect of four sanitizers: Sodium hypochlorite (OCl), Peroxiacetic Acid (PA), Acidified Sodium Chlorite (ASC) and carvacrol on microbiological, sensorial and nutritional quality (total phenols, vitamin C and antioxidant capacity) of fresh-cut jalapeno peppers stored at 5°C during 27 days was evaluated. **Results:** All sanitizers (except carvacrol) maintained microbiological and overall quality of jalapeno peppers during 27 days. ASC (500 and 250 mg L<sup>-1</sup>) maintained the best microbiological and sensorial properties at the end of the storage period. Carvacrol, active ingredient of oregano essential oil, maintained shelf life for only 17 days. At the end of the storage period, all treatments showed a decrease of 12-43% respect to the initial vitamin C values. Total phenols and antioxidant capacity decreased in a lesser degree. None of the treatments except ACS 500 mg L<sup>-1</sup>, induced higher losses of vitamin C, total phenols or antioxidant capacity compared to control. **Conclusion:** Our results showed that all sanitizers were capable of controlling microbial growth without inducing major loss of antioxidant capacity and photochemical. Carvacrol was the only sanitizer that reduced sensory acceptability of fresh-cut jalapeno peppers, however carvacrol treated samples retained the highest levels of photochemical and antioxidant capacity. ASC was the most effective sanitizer even though it was used at concentrations lower than those currently approved by the FDA.

**Key words:** Jalapeno pepper, sanitizers, antioxidant capacity, microbial growth, overall quality

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### INTRODUCTION

Hot peppers are very important in the Mexican diet; together with corn “tortillas” and beans they are part of the basic diet of a large percentage of the population. Hot and sweet peppers (*Capsicum annum* L.) are good sources of vitamins A, C and antioxidants (Chuah *et al.*, 2008; Deepa *et al.*, 2007; Materska and Perucka, 2005; Topuz and Ozdemir, 2007). Concentration of these compounds depends on cultivar,

maturity, growing conditions and postharvest manipulation. Considering their high concentration of bioactive compounds and widespread use among the Mexican population, there has been an interest in producing fresh-cut peppers (Gonzalez-Aguilar *et al.*, 2008a; Raffo *et al.*, 2008).

Studies have shown that fresh-cut products are particularly susceptible to microbial growth owing to the removal of plant protective tissues and the release of cellular fluids from cutting (Heard, 2002), which

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results in shelf life reduction and food-borne illnesses. Given that fresh-cut products are marketed as pre-washed and ready to eat and not subject to further microbial killing steps, the development and proper application of sanitizing agents to remove microorganisms and control pathogen cross-contamination effectively is critical to ensure the quality and safety of fresh-cut produce (Ruiz-Cruz *et al.*, 2006a). Fresh-cut produce deteriorates faster than intact product; this deterioration affects not only sensorial and microbiological quality, but also nutritional quality, since concentration of bioactive compounds decreases as a consequence of deterioration.

Chlorine (sodium hypochlorite, OCl) has been widely used as a sanitizer during produce washing (Sapers, 2003). However, numerous studies have shown that chlorine used at concentrations permitted by the FDA lacks efficacy in removing human pathogens and spoilage microorganisms (Zhang and Farber, 1996). Additionally, chlorine may react with organic matter in water to form carcinogenic products (Parish *et al.*, 2003). The inadequacies of chlorine as a sanitizer have stimulated interest in finding safer, more effective sanitizers (Sapers, 2001; 2003). Alternatives to chlorine, such as chlorine dioxide, ozone, peroxyacetic acid, acidified sodium chlorite and some essential oils have been already proposed (Beuchat, 1998). Acidified sodium chlorite has shown strong antimicrobial activity against *E. coli* O157:H7, *Salmonella*, *Listeria monocytogenes* and spoilage microorganisms on fresh-cut produce (Gonzalez-Aguilar *et al.*, 2004; Ruiz-Cruz *et al.*, 2007). Peroxyacetic acid has also been shown to be effective against spoilage and pathogenic microorganisms (Rodgers *et al.*, 2004). Among essential oils, oregano essential oil and its active component carvacrol have been widely recognized for its antimicrobial properties (Lambert *et al.*, 2001; Zhou *et al.*, 2007), however they are also known to produce undesirable flavors in treated produce (Rojas-Grau *et al.*, 2009).

Several approaches have been addressed in order to minimize deterioration of fresh-cut peppers, such as modified atmosphere packaging, sanitization and cold storage (Raffo *et al.*, 2008; Conesa *et al.*, 2007). However, little information of the effect of these conservation techniques on the nutritional and sensory quality of fresh-cut peppers has been published. For this reason, the aim of the present study was to evaluate the effect of different sanitizer agents on the microbial reduction of fresh-cut Jalapeno Peppers during cold storage and their impact on the sensory quality, antioxidant capacity, total phenols and vitamin C concentration.

## MATERIALS AND METHODS

**Fresh-cut peppers preparation:** Fresh jalapeno peppers fruits (*Capsicum annum* cv. L) cultivated in the area of Delicias, Chihuahua, Mexico and purchased at a local wholesale market in the same city were used for this study. Fruit were sorted to eliminate damaged, poor quality fruit. Jalapeno peppers were washed with tap water to remove residual soil and cut manually into 1 cm width longitudinal strips. Cut peppers were divided into individual 1.6 kg portions contained in nylon mesh bags and washed in the selected sanitizer solutions as described below.

**Sanitization procedure:** Jalapeno strips were submerged in 1:10 ratio (wt/v) solutions containing (i) 200 mL L<sup>-1</sup> sodium hypochlorite (OCl) (Cloralex, NL, Mexico, 6% NaOCl) and pH was adjusted to 6.5 with HCl; (ii) peroxyacetic acid (PA, 80 mL L<sup>-1</sup>, Ecolab, St. Paul, Minn., USA.); (iii) acidified sodium chlorite (ASC, 250 and 500 mg L<sup>-1</sup>) by mixing 0.4 or 0.8 mg L<sup>-1</sup> of NaClO<sub>2</sub> (Sigma-Aldrich, USA) with 2.5 and 5 g L<sup>-1</sup> of citric acid (Sigma-Aldrich, USA), respectively; or iv) 1 or 2.5 mM of carvacrol (Sigma-Aldrich, USA). Tap water washed peppers samples were used as controls. Immersion time was 1 min for ASC and carvacrol treatments and 2 min for OCl, AP and control. After dipping, peppers were drained for 30 sec and spin-dried for 30 s to remove excess of water, using a manually operated commercial salad spinner. Samples of 200 g each, were packaged in polypropylene bags (18×22 cm) with an oxygen transmission rate of 290 mL day<sup>-1</sup> m<sup>-1</sup> O<sub>2</sub><sup>-1</sup> and sealed using a AIE-302 impulse sealer (American IntNI Electric, Taiwan). All samples were stored at 5°C for up to 27 d for subsequent microbiological, sensorial and nutritional analyses at 3 day intervals.

**Sensory evaluation:** The overall visual quality of cut peppers was evaluated following a modified procedure of Mercado-Silva *et al.* (1998) by a six-member trained panel. Before the test, the panel members were trained to recognize and scale the quality attributes of cut peppers. Samples were coded with three-digit random numbers to mask the treatment identity. The acceptability was scored using a nine-point hedonic scale, where 9 = excellent, no defects, 7 = very good, minor defects, 5 = fair, moderate defects, 3 = poor, major defects and 1 = unusable. Minor defects were attributed to dryness of the surface tissue and major defects were usually due to decay. A score of 5 or below was considered to be unsalable.

**Microbial counts:** Cut peppers samples (25 g) were homogenized with sterile phosphate buffers using a stomacher blender (Model 400) for 2 min at 230 rpm. The homogenate was serially diluted by a ten factor. For each dilution, 1 mL was plated on each medium according to Mexican Official Norms (NOM). Plate count agar (Difco) incubated aerobically at 35°C for 24-48 h for total aerobic bacteria; violet red bile agar (Difco) incubated at 35°C for 24-26 h for coliforms and potato dextrose agar supplemented with 10% of tartaric acid for yeast and molds. Results are expressed as log cfu g<sup>-1</sup> of fresh-cut jalapeno pepper.

**Total phenols content:** Jalapeno extracts were obtained according to the methodology proposed by Khkonen *et al.* (1999). Briefly, 0.1 g of powdered dried jalapeno was weighted into a test tube. 5 mL of 80% methanolic solution was added, stirred and sonicated 15 min in the dark. Then the tube was centrifuged (3000 g) for 10 min and the supernatant was collected. Extraction was repeated and total volume was taken to 10 mL. Total phenols were determined as previously reported by Alvarez-Parrilla *et al.* (2007), with the Folin-Ciocalteu reagent, using Gallic acid in methanol (80%) as standard. 50 µL of extract (or standard) was mixed with 250 µL of Folin-Ciocalteu reagent (1:10 dilution with water) in a 96 well microplate and incubated 2 min at room temperature. 200 µL of Na<sub>2</sub>CO<sub>3</sub> (75 mg L<sup>-1</sup>) were added and incubated during 15 min at 50°C. After cooling to room temperature the absorbance at 760 nm was measured with a BioRad BenchmarkPlus microplate reader (BioRad Mexico, Mexico City, Mexico). Results are expressed as mg Gallic Acid Equivalent (GAE) per 100 g FW.

**Ascorbic acid content:** AA was determined as previously described by Gonzalez *et al.* (2004) 2 g of fresh-cut Jalapeno peppers were homogenized with 15 mL of metaphosphoric acid (4%, wt/v). The mixture was centrifuged (3000 rpm) and the aqueous extract filtered throughout 0.45 µm pore size nylon membrane (Millipore Corp., Bedford, MA). Filtered extracts were manually injected (20 µL) into a Varian ProStar HPLC system (Varian Inc., Walnut Creek, CA) equipped with UV-Vis detector and fitted to a Microsorb-MV C18 reversed-phase column (4.6×250 nm) (Varian Inc., Walnut Creek, CA). 0.2 M KH<sub>2</sub>PO<sub>4</sub> (adjusted at pH 2.4 with phosphoric acid) was used as mobile phase, at a 1 mL min<sup>-1</sup> flow rate. Ascorbic acid was monitored at 254 nm. Results are expressed as mg AA per 100 g FW.

**Antioxidant capacity:** The antioxidant capacity of the different jalapeno samples was determined by both

Ferric Reducing/Antioxidant Power assay (FRAP) and DPPH radical-scavenging assay. FRAP was performed as previously described by Alvarez-Parrilla *et al.* (2007) FRAP reagent was daily prepared and kept at 37°C, by mixing 0.3 M acetate buffer, pH 3.6 with 10 mM TPTZ solution in 40 mM HCl plus 20 mM FeCl<sub>3</sub>•6H<sub>2</sub>O, at a 10:1:1 ratio. Measuring solutions were prepared in a 96 well microplate, by mixing 180 µL of FRAP reagent with 24 µL of diluted jalapeno extract (1:3), distilled water or FeSO<sub>4</sub>•7H<sub>2</sub>O stock solutions. All measurements were carried out at 37°C. Absorbance was measured at 595 nm, every 30 s, during 60 min, using a Biorad microplate reader. Values of antioxidant capacity are reported as µmol Fe<sup>2+</sup> per 100 g FW.

DPPH radical-scavenging assay was performed according to Wojdylo *et al.* (2007), with slight modifications. 5 µL of jalapeno extract (or Trolox standard solution) were mixed with 195 µL of 60 µM DPPH (1,1-Diphenyl-2-picrylhydrazil radical, Sigma Aldrich) in a 96 well microplate, at room temperature. Absorbance was measured at 517 nm, every 60 sec, during 60 min, using a Biorad microplate reader. DPPH values were determined at 30 min and are reported as µmol Trolox Equivalents (TE) per 100 g FW.

**Statistical analysis:** Experiments were repeated twice, each with three replications. Results of the two experiments were combined and values represent the mean ± SD of six replicates. ANOVA and DMS analyses were performed in order to determine differences between sanitizer treatment and storage period (t-test), using the commercial software SPSS 13.0.

## RESULTS

**Sensory evaluation:** Overall visual quality for all treated fresh-cut Jalapeno Peppers decreased continuously when stored at 5°C (Fig. 1). Water loss was the most important detrimental factor that affected fresh-cut jalapeno peppers quality. No statistical difference between control and PA treated jalapeno peppers were observed during the whole experiment. Shelf-life for these samples was 20-22 days. Peppers treated with OCI and both ASC concentrations showed the highest overall quality at the end of the storage (27 day at 5°C), in contrast, Carvacrol treated samples presented the lowest scores during all the storage period. Shelf-life of Carvacrol treated fresh-cut jalapeno peppers was only 14 day at 5°C (overall quality score of 5). The low quality score in these treatments was mainly due to the decayed and slimy appearance of samples.

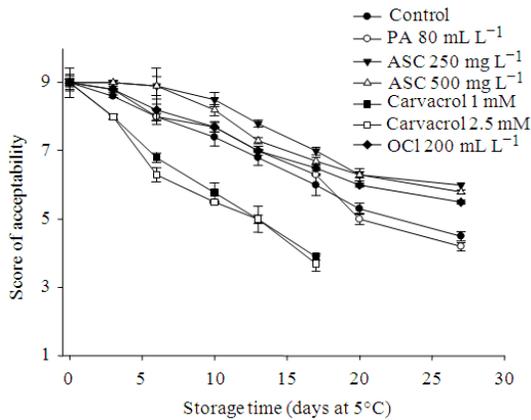


Fig. 1: Effect of carvacrol and sanitizer treatment on overall quality of packaged fresh-cut jalapeno peppers stored at 5°C for up to 27 days. Data points are means  $\pm$  SEM of six replicates

**Microbial enumeration:** Initial populations of total aerobic bacteria (Fig. 2a), total coliforms (Fig. 2b) and molds and yeasts (Fig. 2c) on unwashed peppers were 3.1, 1.4 and 1.0 log cfu g<sup>-1</sup>, respectively. Treatment with all sanitizers decreased initial total aerobic bacterial counts compared with control (water-washed). PA showed the larger initial reduction. Although the aerobic bacteria grew rapidly during cold storage, at the end of the experiment (day 27), all treatments reduced total aerobic counts by 2-3 log cfu g<sup>-1</sup> compared to control. It is important to mention that at day 17, both carvacrol concentrations showed the best aerobic bacteria growth reduction, however, as previously described, at this day the products were not accepted by panelist. ASC treated samples (both concentrations) maintained the lowest bacterial counts throughout the storage.

All treatments (except carvacrol 1 mM) caused significant reductions in total coliforms count on day 0 compared with the control (Fig. 2b). This reduction was maintained throughout the storage. Reduction of 1.4-3.2 log cfu g<sup>-1</sup>, were observed for the different treatments. Fresh-cut peppers washed with ASC presented the lowest microbial growth, while control peppers showed the highest coliforms growth.

The yeast and mold counts on fresh-cut peppers during 27 days of storage are shown in Fig. 2c. No microorganisms were detected during the first 3 days of storage, starting at day 6, all treatments showed similar growth of yeasts and molds. At 17 day no statistical difference was observed between control and Carvacrol treated samples. At the end of the experiment (27 day), a significant decrease in yeast and mold growth was observed between all treatments (except OCI) and control.

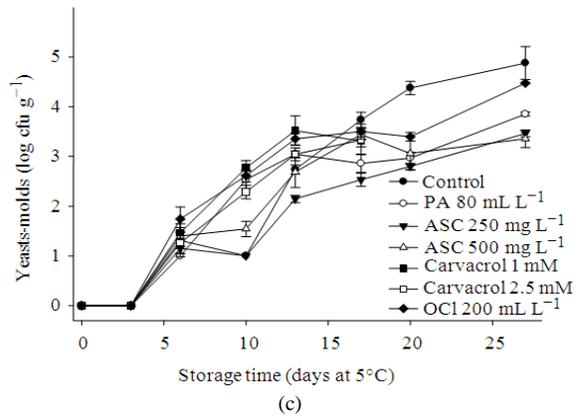
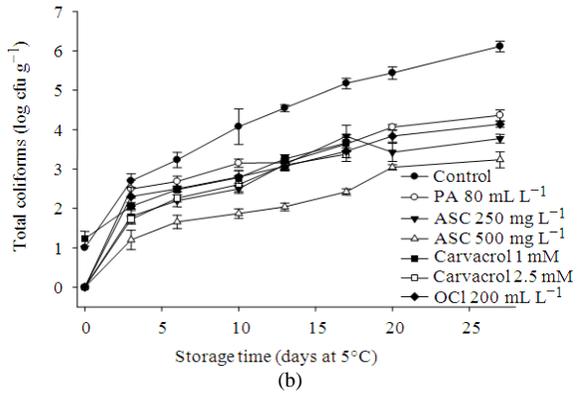
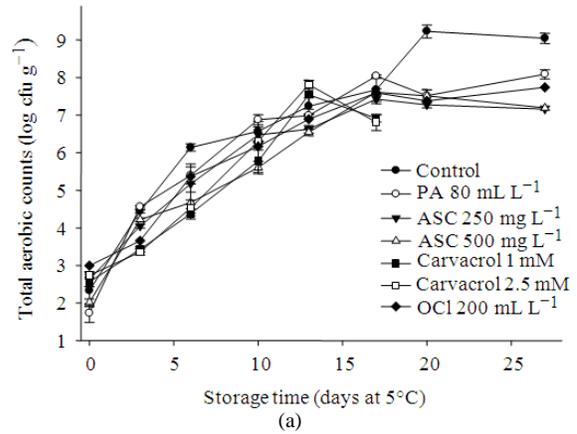


Fig. 2: Effect of carvacrol and sanitizers treatment on (a) total aerobic counts; (b) total coliforms and (c) yeasts and molds in fresh-cut jalapeno peppers stored at 5°C for up to 27 days. Data points are means  $\pm$  SEM of six replicates

**Total phenols content:** Figure 3a shows the modification of total phenol concentrations over time, for fresh-cut jalapeno peppers treated with different sanitizers.

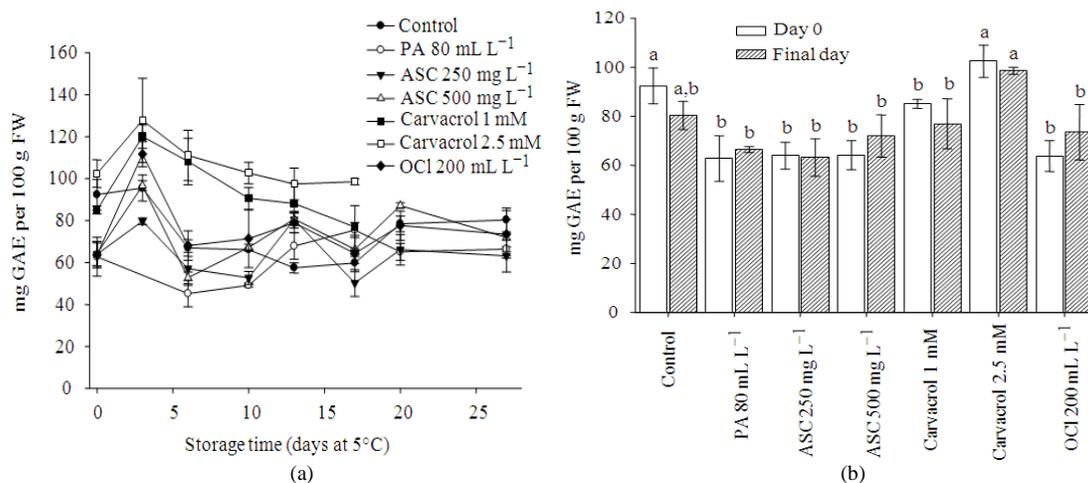


Fig. 3: Phenolic concentration of fresh-cut jalapeno peppers treated with carvacrol and sanitizers stored at 5°C for up to 27 days (a) Time dependent evolution of phenolic compounds (b) Initial and final concentrations. Data points are means  $\pm$  SEM of six replicates. Different letters in bars indicate statistical significant difference between treatments in either initial or final values (DMS test,  $p < 0.05$ ). \*: Means statistical difference between initial and final values (t-test,  $p < 0.05$ )

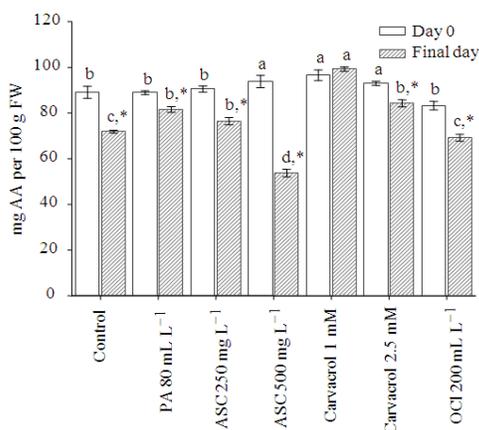


Fig. 4: Initial and final vitamin C concentration of fresh-cut jalapeno peppers treated with carvacrol and sanitizers stored at 5°C for up to 27 days. Data points are means  $\pm$  SEM of six replicates. Different letters in bars indicate statistical significant difference between treatments in either initial or final values (DMS test,  $p < 0.05$ ). \*Means statistical difference between initial and final values (t-test,  $p < 0.05$ )

Initial phenolic concentrations ranged from 62.8 mg GAE per 100 g FW for PA treated peppers to 102.4 mg GAE per 100 g FW for carvacrol 2.5 mM treated peppers. When the initial phenolic concentrations were compared among treatments, a decrease of the initial phenolic concentration, respect to control, after the

sanitization process was observed, due to a bleaching process; this reduction was not observed in carvacrol treated peppers. These results can be explained considering that carvacrol itself is a phenolic compound and consequently it is absorbed into the tissue during the sanitation process (Fig. 3b). In Fig. 3a it is possible to observe that for all treatments there is an increase on total phenols at day 3 and thereafter a slight decrease is observed over time. Figure 3b shows initial and final phenolic concentrations of fresh-cut jalapeno peppers treated with different sanitizers, in which a small, not significant, decrease in the phenolic concentration at the end of the shelf-life was observed. Figure 3a indicates that neither the sanitizers nor the storage period had any effect on the phenolic concentration of fresh-cut jalapeno peppers.

**Ascorbic Acid content:** Effect of sanitization on vitamin C concentration over the shelf-life period is presented in Fig. 4. Initial vitamin C values ranged from 83.2 mg AA per 100 g FW for OCI treated peppers to 96.6 mg AA per 100 g FW for 1 mM carvacrol treated peppers, these values were in the range of other reported for jalapeno peppers (Navarro *et al.*, 2006; Chuah *et al.*, 2008; Marin *et al.*, 2004; Simonne *et al.*, 1997; Howard *et al.*, 2000). Carvacrol treated peppers had statistically higher vitamin C concentration. Vitamin C concentration decreased at the end of the storage period, except for those peppers treated with carvacrol, where the concentration remained practically unchanged, or even a slight increase was observed.

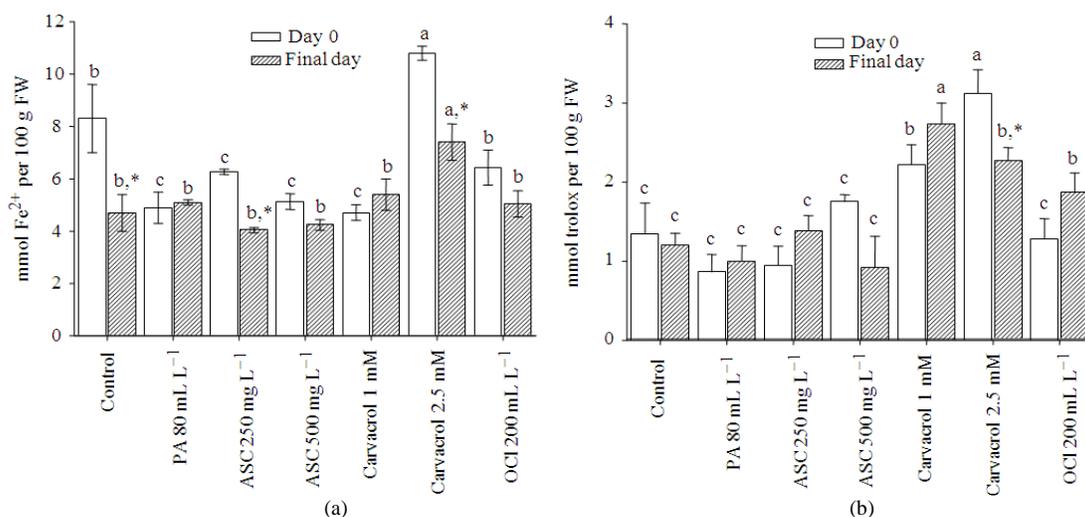


Fig. 5: Initial and final antioxidant capacity of fresh-cut jalapeno peppers treated with carvacrol and sanitizers stored at 5°C for up to 27 days, determined by (a) FRAP and (b) DPPH assays. Data points are means  $\pm$  SEM of six replicates. Different letters in bars indicate statistical significant difference between treatments in either initial of final values (DMS test,  $p < 0.05$ ). \*: Means statistical difference between initial and final values (t-test,  $p < 0.05$ )

However, it should be considered that the storage period for these samples was only 17 days (Fig. 1). The largest vitamin C concentration decrease (42%) was observed for ASC 500 mg L<sup>-1</sup> treated peppers, followed by control. A 16.8% reduction was observed for the OCI treated jalapeno peppers. The lowest vitamin C loss was found in PA treated peppers (8%). These results suggest that elaboration of fresh-cut jalapeno peppers reduces their nutrimental quality due to ascorbic acid losses; however, sanitation processes may help reduce this loss.

**Antioxidant capacity:** In the present study, antioxidant activity was measured by both FRAP and DPPH scavenging assays, two of the most used antioxidant capacity methods for vegetable samples. Initial FRAP antioxidant capacity value for the control was 8.32 mmol Fe<sup>2+</sup> per 100 g FW (Fig. 5a) and the sanitation treatment reduced the initial antioxidant capacity of all samples (values ranged from 4.71 mmol Fe<sup>2+</sup> per 100 g FW for carvacrol 1 mM treated peppers to 10.8 mmol Fe<sup>2+</sup> per 100 g FW for carvacrol 2.5 mM). These results showed that during sanitation, a loss of phenolic compounds (Fig. 3b) and antioxidant capacity is observed for all treatments due to a bleaching process. The only treatment were no reduction (and even a higher FRAP values) was observed was in the carvacrol 2.5 mM treated peppers. These results are in agreement with those observed for phenolic compounds.

When the initial and final FRAP values were compared, it is possible to observe that almost all treatments reduced the antioxidant capacity value of fresh-cut peppers during shelf-life. This reduction in the FRAP value was more evident for control (43%) than for the other treatments. These results are similar to those obtained for vitamin C.

Figure 5b shows the initial and final DPPH values for all treated fresh-cut jalapeno peppers. Initial DPPH value for control was 1.35 mmol TE per 100 g FW. Once again, when the initial antioxidant capacity values of the treated peppers were compared, it is possible to observe that both carvacrol treatments increased DPPH value, however the other sanitizers had no effect on DPPH value. When initial and final values were compared (Fig. 5b), no effects were observed due to storage for all treatments except carvacrol 2.5 mM, where a significant reduction of DPPH antioxidant capacity was observed.

In order to determine which of the two compounds: The phenolic compounds or vitamin C was the main contributor to the antioxidant capacity of jalapeno peppers, a correlation study between the initial and final values of this two compounds with antioxidant capacity was carried out. Results are presented in Table 1. From the analysis of this Table 1, it is possible to conclude that phenolic compounds are the main contributors to antioxidant capacity in jalapeno peppers and that better correlation is observed with the FRAP method. No significant correlation was observed for vitamin C.

Table 1: Correlation coefficient between phenolic compounds, vitamin C and antioxidant capacity assays

Correlation	Correlation coefficient (R)	Pearson probability
FRAP-Folin	0.7172	0.0039
DPPH-Folin	0.6743	0.0082
FRAP-DPPH	0.6209	0.0235
AA-FRAP	0.3906	0.1676
AA-DPPH	0.4902	0.0752
AA + Folin-FRAP	0.7269	0.0032
AA C+ Folin-DPPH	0.7587	0.0016

A good correlation was observed when both antioxidant capacity methods were compared ( $R = 0.62$ ,  $p < 0.05$ ). Finally, it is possible to observe that when both phenols and vitamin C are added, the correlation with both antioxidant capacity assays increased, indicating that in the case of jalapeno pepper, both compounds are responsible (in more or less degree) for antioxidant capacity.

## DISCUSSION

Gonzalez-Aguilar *et al.* (2004) showed that fresh-cut bell pepper maintained their quality for up to 25 days at 5°C. Therefore, this temperature was chosen in the present study, observing no visual signs of chilling injury, in fresh-cut jalapeno peppers for up to 27 days, contrasting with the study by Kang and Lee (1997) who reported that fresh-cut peppers are sensitive to chilling injury during storage at 5°C.

Several studies have shown that sanitation is essential in preserving quality of fresh-cut fruits and vegetables, since deteriorative microorganisms are ubiquitous in fruits and, during processing, pathogenic microorganisms can infect produce (Gonzalez *et al.*, 2004; Kang and Lee, 1997). The use of different sanitizers, such as chlorine dioxide, Peroxyacetic Acid (PA), Acidified Sodium Chlorite (ASC) and some essential oils has been proposed as an alternative to chlorine (OCI) due to reported lack of efficacy and other disadvantages of the later (Beuchat, 1998). Although effectiveness of these sanitizers in reducing microbial growth has been demonstrated in some fresh-cut products, studies addressing their effect on the total quality of the products are lacking.

Initial populations of total aerobic bacteria, total coliforms and molds and yeasts on unwashed jalapeno peppers were similar to those reported for fresh-cut bell peppers (Gonzalez-Aguilar *et al.*, 2004). Treatment of fresh-cut peppers with ASC at concentrations of 250 and 500 mg L<sup>-1</sup> showed the best results in reducing growth of all microorganisms and, at the same time maintained the highest visual quality of the produce as evaluated by a trained panel. Similar results have been

obtained with ASC for the reduction of microbial growth in shredded carrots (Ruiz-Cruz *et al.*, 2006a; 2006b; 2007; Gonzalez *et al.*, 2004). The antibacterial activity capacity of ASC is attributed to the low pH of ASC solutions (2.5) that probably affect the cell's ability to maintain pH homeostasis, disrupting substrate transport and inhibiting metabolic pathways (Simonne *et al.*, 1997). ASC is currently approved by the FDA to be used in the range 500-1200 mg L<sup>-1</sup> (VLex, 2000). In this concentration range, ASC exhibited strong efficacy over pathogen inactivation on various food products (Gonzalez-Aguilar *et al.*, 2004). However, our results show that even lower ASC concentrations (250 and 500 mg L<sup>-1</sup>) were highly effective at controlling microbial growth, better than other treatments including OCI; ASC was also better at maintaining sensorial quality of fresh-cut jalapeno peppers. Moreover, in preliminary tests, we used ASC at higher concentrations (in the FDA-approved range) on fresh-cut peppers, lettuce, cilantro and spinach and found that it had a negative impact on visual quality, whether or not it was followed by a potable water rinse (data not shown). Similar findings were also reported by Bosilevac *et al.* (2004) and Roller and Seedhar (2002) who found that red meat products treated with 300 mg L<sup>-1</sup> of ASC were superior to those treated with 600 mg L<sup>-1</sup>. Therefore, our results suggest approval for the use of ASC below 500 mg L<sup>-1</sup> should be analyzed since it may better preserve sensory attributes without compromising food safety. FDA approval at these low concentrations is needed before its commercial application.

After ASC, OCI (200 mL L<sup>-1</sup>) treatment was the most effective in maintaining sensory and microbiological quality of fresh-cut jalapeno peppers, followed closely by PA (80 mL L<sup>-1</sup>) treatment. Carvacrol treatment was, on average, as effective as the others in reducing the growth of total aerobic and coliform bacteria; however it showed a strong negative impact on visual sensory attributes of fresh-cut jalapeno peppers and a low effect in reducing growth of yeasts and molds. Carvacrol is a phenolic terpenoid, major component and active ingredient of oregano essential oil and its antimicrobial properties are well documented (Lambert *et al.*, 2001; Zhou *et al.*, 2007), however studies regarding carvacrol's organoleptic impact on fresh-cut produce are limited. Roller and Seedhar (2002) reported that no adverse sensory change was observed on kiwifruit and honeydew melon treated with 1 mM carvacrol. On the contrary and in agreement with the results shown in the present study, Rojas-Grau *et al.* (2009) observed that, in spite of the low concentration of oregano oil used in the treatment of edible-coated

fresh-cut apples, consumers detected a residual herbal taste, which diminished overall preference of these samples. It is clear that, carvacrol and/or oregano essential oil have the potential to be used as sanitizers in fresh-cut products, however its effects on sensory quality must be closely evaluated in each specific product.

Green, yellow and red peppers are known to be excellent sources of vitamin C, polyphenols and carotenoids (Chuah *et al.*, 2008; Deepa *et al.*, 2007; Materska and Perucka, 2005; Topuz and Ozdemir, 2007). These phytochemicals show high antioxidant capacity and a diet rich in such compounds has been linked to a decreased risk of developing chronic and degenerative diseases (Gonzalez-Aguilar *et al.*, 2008a). However, as far as we know, there are no studies that show the effect of sanitizers and conservation period on the phytochemical degradation and antioxidant capacity of fresh-cut peppers. Initial concentrations of total phenols and vitamin C in control fresh-cut jalapeno peppers were similar to those reported by several authors for jalapeno and other sweet and hot peppers (Chuah *et al.*, 2008; Deepa *et al.*, 2007; Howard *et al.*, 1994; Navarro *et al.*, 2006; Oboh *et al.*, 2007; Oboh and Rocha, 2008; Ornelas-Paz *et al.*, 2010). Sanitation process reduced initial total phenols but not vitamin C concentrations; however, storage reduced vitamin C (11-42%) but had no impact on total phenols concentration. These results are in agreement with several authors. Gonzalez-Aguilar *et al.* (2008b) observed a 20-60% vitamin C decrease in fresh-cut mangoes stored at 5°C; Howard and Hernandez-Brenes (1998) reported that ascorbic acid concentration of peppers decreased 17% when jalapeno peppers were maintained at 4.4°C under MAP atmosphere during 15 days; Raffo *et al.* (2008) observed an 11.6% reduction on vitamin C of fresh-cut red sweet peppers treated with OCl (165 mL L<sup>-1</sup>), kept 9 days at 4°C. In fact, it is well known that vitamin C is the most labile phytochemical, which is easily degraded even by mild treatments and during storage (Gonzalez-Aguilar *et al.*, 2010). Nevertheless it should be mentioned that all sanitizers except ASC 500 mg L<sup>-1</sup> prevented vitamin C loss, compared to control.

As mentioned before, neither treatment nor storage significantly affected total phenols concentration; however, in Fig. 3a, it is possible to observe that for all treatments there is an increase on total phenols at day 3 and thereafter a slight decrease over time. Even though no enzymatic studies were carried out, this increase on phenolic content could be due to a ripening process that increases PAL activity, which consequently may increase phenolic content of peppers (Simonne *et al.*, 1997).

The antioxidant capacity of green, yellow and red peppers has been determined by several methods, such as 2,2-diphenyl-1-picrylhydrazyl radical, DPPH (Chuah *et al.*, 2008; Materska and Perucka, 2005; Sun *et al.*, 2007), TEAC (Navarro *et al.*, 2006; Pellegrini *et al.*, 2003), β-carotene bleaching assay (Materska and Perucka, 2005), Fe<sup>2+</sup>-induced lipid peroxidation assay (Oboh and Rocha, 2008) and ferric reducing/antioxidant power assay, FRAP (Deepa *et al.*, 2007; Marquez *et al.*, 2006). Antioxidant capacity of fruits and vegetables is owed to phytochemicals such as phenolic compounds, vitamins C and E, carotenoids, among others. In the present study, antioxidant activity was measured by FRAP and DPPH assays. Similar to total phenols, initial FRAP values of antioxidant capacity were reduced in all treatments (except carvacrol) due to a bleaching process. FRAP values were reduced after storage; however this reduction was highest in control samples (43%). ASC 250 mg L<sup>-1</sup> and carvacrol 2.5 mM treated samples showed a significant reduction in FRAP values (35 and 31% respectively), all other treatments prevented the loss of antioxidant capacity; in fact AP treated samples had higher antioxidant capacity at the end of the storage. When the antioxidant capacity of fresh-cut jalapeno peppers was evaluated with DPPH method, no effect on antioxidant capacity was observed due to sanitation or storage.

The case of carvacrol effect on phytochemicals and antioxidant activity deserves special attention, since it increased initial total phenols and antioxidant capacity and prevented loss of vitamin C and antioxidant capacity in a high degree; therefore carvacrol treated samples showed the highest final values of phenols, vitamin C and antioxidant capacity. However, it has been mentioned that carvacrol rapidly reduced acceptability of fresh-cut jalapeno peppers and carvacrol treated samples had the lowest shelf life.

Finally, the correlation studies between antioxidant capacity and concentration of phytochemicals show that phenols are the main contributors to antioxidant capacity in jalapeno peppers, but vitamin C is also important. These results are in agreement with other authors (Alvarez-Parrilla *et al.*, 2007; Gorinstein *et al.*, 2004).

## CONCLUSION

The following conclusions can be made from the results presented above: (i) ACS at 500 and 250 mg L<sup>-1</sup> was the most effective sanitizer in fresh-cut jalapeno peppers, since it reduced microbial growth without significantly affecting sensory quality nor phytochemical content or antioxidant capacity; (ii)

fresh-cut jalapeno peppers are good sources of phenolic compounds and vitamin C and sanitizer treatment does not significantly affect the concentration of these phytochemicals, in fact it may prevent their loss during storage; (iii) even though carvacrol treatment was effective in preventing microbiological spoilage and phytochemical loss, it had a strong negative effect on sensory acceptability, further studies should be carried out with carvacrol or oregano essential oil in order to overcome this lack of acceptability and take advantage of its beneficial properties.

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