

Active Modified Atmosphere Packaging of Fresh-cut Bell Peppers: Effect on Quality Indices

Helen Manolopoulou¹, Gregory Lambrinos² & George Xanthopoulos²

¹ Technological Educational Institute of Kalamata, Department of Crop Production, Antikalamos, Kalamata, Greece

² Agricultural University of Athens, Department of Natural Resources Management and Agricultural Engineering, Athens, Greece

Correspondence: George Xanthopoulos, Agricultural University of Athens, Department of Natural Resources Management and Agricultural Engineering, 75 Iera Odos Str. 118 55, Athens, Greece. Tel: 30-210-529-4031. E-mail: xanthopoulos@aua.gr

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Abstract

Fresh-cut green bell peppers (*Capsicum annuum L.*) were stored in modified atmosphere packaging (MAP) made of impermeable high-density polyethylene film. Two in-packaging atmospheres and storage temperatures (0 °C and 5 °C) were tested. The respiration rate of the unpackaged produce and the in-package gas concentration, mass loss, firmness, skin colour, ascorbic acid and visual quality of the packaged produce were estimated. Cutting, increased respiration rate of the unpackaged produce by 24% compared to the intact produce for the same storage temperature. After 5 days of storage at 5 °C, significant O₂ depletion of the active modified atmosphere was found. Limited mass loss (0.4-0.5% of the initial mass) and firmness degradation were estimated in both storage temperatures due to the beneficial effect of packaging. The hue angle (h*) reduction was limited in all cases and the initial green colour was preserved. Initial ascorbic acid content was preserved at 0 °C, but significantly increased at 5 °C. The visual quality of the packaged produce was assessed by six trained panelists and found that was not significantly changed at 0 °C storage. In conclusion, the tested active MAP maintained the initial quality indices of fresh-cut peppers (*cv. Twingo FI*) for up to 10 days at 0 °C but not at 5 °C.

Keywords: capsicum annuum, active modified atmosphere packaging, quality indices, cut peppers

Nomenclature:

a*: red/green balance

b*: yellow/blue balance

C*: chroma

df: degrees of freedom

e_i: residuals of Eq. (1)

h*: hue

L*: black/white balance

LSD: least significant difference

P1, P2, P3: gas mixtures (O₂/CO₂: 5% : 10%, 5% : 15%, 21% : 0.03%)

R²_{adj}: coefficient of determination adjusted for the degrees of freedom

SEE: standard error of estimate

σ²: variance of the residuals e_i

1. Introduction

Minimally processed products are very popular with consumers since they are easy in use and healthy (Ohlsson, 1994). However, these products are highly perishable, deteriorate rapidly, and have short shelf-life. Therefore is important to extend their shelf-life with no compromise of their final quality (Huxsoll & Bolin, 1989; Artés, Gómez, Aguayo, Escalona, & Artés-Hernández, F., 2009; Beaulieu, 2011). Ahvenainen (2000) reported that the

sensorial and nutritional shelf-life of the minimally processed products should be at least 4-7 days but generally longer storage times are needed for commercial use. Minimally processed products characterised by high respiration rates due to the wounds caused by the processing while biochemical changes are also observed. The establishment of a beneficial modified atmosphere is difficult to be achieved, since few packaging films are permeable enough to match the respiration rate of the fresh-cut products. Perforation-mediated packaging, are used to increase packaging permeability and reduce potential anaerobiosis (Ahevainainen, 2000).

Bell pepper has been adopted in various diets for their spicy taste and characteristic flavour and are significant source of vitamins A and C, antioxidants and carotenoid zeaxanthin. The latter are important in human diet and sensitive in minimal processing techniques (Raffo, Baimonte & Paoletti, 2008). The recommended controlled atmospheres (CA) vary widely depending on the processed cultivars and the degree of processing. Information about the physiological behaviour of minimally processed bell peppers is limited and is mainly focused to passive modified atmosphere packaging (Conésa, Artés-Hernández, Geysen, Bart, & Artés, 2007). Abe, Yoshimura, Zhou, Abe & Iwata (1992) and Lopéz-Galvéz, El-Bassuoni, Nie & Cantwell (1997) reported that minimally processed bell peppers stored at 0 °C retained their composition and visual quality for 6 to 15 days. Senesi, Prinzivalli, Sala & Gennari (2000) argued with the adoption of low storage temperatures (2 °C) since these can cause pitting due to sensitivity of peppers in chilling injuries. Senesi et al. (2000) suggested storage at 8 °C, which retains satisfactorily the initial quality of the produce. Gorny (2001) suggested as pre-processing storage temperature of pepper, 7-10 °C, and as post-processing, 0-5 °C adding that an atmosphere of 3% O₂ and 5-10% CO₂ can reduce the microbial growth and retain the visual quality of fresh-cut peppers.

In this paper was studied the physiological behaviour of fresh-cut green peppers (*cv. Twingo FI*) packaged in impermeable high density polyethylene (HDPE) film, implementing two initial in-package O₂/CO₂ concentrations and two storage temperatures, 0 °C and 5 °C, for 10 days. A new empirical model was developed to relate mass and firmness loss during the modified atmosphere packaging storage of fresh-cut green bell peppers.

2. Materials and Methods

2.1 Raw Material

Green bell peppers (*Capsicum annuum L. cv. Twingo FI*) were handpicked at their commercial maturity (firm and bright green) during June, July and August from a local farm in southern Greece. After harvest, peppers were immediately transported to the lab and sorted, removing those of poor quality (blemishes or defects) and non uniform. The sorted produce was disinfected with sodium hypochlorite (100 ppm NaOCl) in water at 5 °C for 2 min and rinsed with tap water at 5 °C for 2 min (Varoquaux & Mazollier, 2002; Cantwell & Suslow, 2002; González-Aguillar et al., 2004; Gil, Selma, López-Gálvez, & Allende, 2009; Gil, Allende, & Selma, 2011). Although in some European countries (Germany, the Netherlands, Switzerland and Belgium) the use of chlorine as disinfectant is not allowed (Oms-Oliu & Soliva-Fortuny, 2011) in other European countries, chlorine is still in use as a 'processing aid' under the Directive 89/107/EEC (Gil et al., 2009). Finally, peduncle was removed from the disinfected peppers and the fruits chopped in uniform rings of 5 mm thickness. Equipment used in the processing, was sanitised before use with a chlorine solution. The pepper rings were sanitised again with chlorinated water at 5 °C for 2 min and rinsed with tap water at 5 °C for 2 min. The remaining water on the pepper rings removed carefully with absorbing paper to prevent damaging the pepper rings. The adopted hygiene rules and low storage temperatures are consistent with the food safety regulations (González-Aguilar, Ayala-Zavala, Ruiz-Cruz, Acedo-Félix, & Diaz-Cinco, 2004; Gil et al., 2011).

2.2 Packaging and Storage Conditions

Pepper rings of 100±2 g were placed in 20 µm thick HDPE (0.96 g cm⁻³) packages, practically impermeable (O₂ permeability, 35.01 mL m⁻² d⁻¹ bar⁻¹ and CO₂ permeability 50.58 mL m⁻² d⁻¹ bar⁻¹, S=P_{CO₂}/P_{O₂}=1.4 at 20 °C, according to data provided by the film manufacturer) since gas flushing was applied. The overall packaging surface area was 1,200 cm². The gas flushing was taken place in vacuum emptied packages, using a gas mixer (MAP Mix 9000, Denmark) and a vacuum compensation chamber (Multivac A. 300/16, Germany). Three initial in-package gas concentrations were tested: (i) 5% O₂ and 10% CO₂ balanced with N₂ [P1], (ii) 5% O₂ and 15% CO₂, balanced with N₂ [P2], (iii) packaging film with one macro-perforation, 3 mm in diameter (the in-packaging atmosphere was near-ambient) [P3], (iv) control samples [C] placed on polystyrene trays (120×215×15 mm) and over wrapped with 13 µm thick PVC film to minimise mass loss.

Gorny (2001) suggested for CA storage of fresh-cut bell peppers O₂ and CO₂ of 3% and 5-10% respectively, noting that injuries due to increased CO₂ levels vary significantly among cultivars. Lopéz-Galvéz et al. (1997) reported that storage conditions, 5 °C and 10% CO₂ can retain product quality and delay decay. Based on the previous suggestions was decided to test two different initial atmospheres, one of which with CO₂>10%, to

examine the physiological behaviour of the specific pepper variety and compare it with the respective under ambient conditions. All the tested packages were stored at 0 °C and 5 °C, and 90% relative humidity in darkness for 10 days as Kang & Lee (1997) suggested. Six packages were prepared per sampling date, treatment (P1, P2, P3 and C) and storage temperature (0 °C and 5 °C). For gas analysis and mass loss estimation, ten bags per treatment and storage temperature were prepared. Every experiment conducted in triplicate.

2.3 Gas Analysis of the In-package Atmosphere

The in-package atmosphere (O₂ and CO₂) was daily measured with a headspace gas analyser (CheckMate 9000, PBI Dansensor Co., Denmark), drawing up 2 mL of air samples. Sampling took place with a hypodermic needle through a septum pasted on the packaging. Ten bags per treatment (P1, P2, P3) and storage temperature were prepared for gas analysis.

2.4 Respiration Rate (RR) of Whole and Fresh-cut Peppers

The CO₂ production rate of whole and fresh-cut unpackaged peppers was measured with a closed respiratory cell system having 3.9-6.3% uncertainty in measurements (Mitropoulos, Lamprinos, & Manolopoulou, 2000). 110±5 g of unpackaged whole peppers and 100±2 g fresh-cut peppers were placed in 962 mL gas tight containers under atmospheric conditions (20.95% O₂, 0.038% CO₂, 79.012% N₂) and left for the rest of the measuring period. Six gas samples per storage temperature (0 °C and 5 °C) were daily taken from the headspace and used for estimation of the respiration rate.

2.5 Mass Loss

Mass loss was calculated as (%) of the initial mass, with an electronic scale of ±0.01 g accuracy. Weighting carried out in controlled air conditions to avoid moisture condensation on the packages. In the packages used for mass loss estimation, was not detected visible water drips from the packaged pepper rings. The mass loss of the 10 packages per treatment (P1, P2, P3, and C) and storage temperature was estimated at the beginning of the experiment and on the 3rd, 6th, 8th and 10th day of the experiment.

2.6 Firmness

Firmness was measured with a Texture Analyser TA-XT2i (SMS, England) using a Kramer shear cell, containing 10 g of sample. The speed of the probe was set to 1 mm s⁻¹. Firmness was evaluated at the maximum rupture force (N), defined from a typical force-distance diagram. Measurements of the firmness were carried out at the beginning of the experiment and on the 3rd, 6th and 10th day in 6 packages per treatment (P1, P2, P3, and C) and storage temperature.

2.7 Colour Measurement

Colour was measured on the CIE L*a*b* chromatic space with a Minolta CR-300 Chromameter (Minolta Corp., Japan). The instrument was initially calibrated using a white ceramic tile (Y=92.6 X=0.313 y=0.319). The L* chromatic variable, ranges from 0 (black) to 100 (white) and is an indicator of the lightening or darkening due to the physicochemical changes taking place during storage. The a* measures the degree of redness (+a*) and greenness (-a*) while the b* the degree of yellowness (+b*) and blueness (-b*). The measured a* and b* values were used to estimate chroma values, $C^*=(a^{*2}+b^{*2})^{0.5}$ and hue angle degrees $h^*=\arctan^{-1}(b^*/a^*)$. In fact, hue angle values greater than 90° correspond to intense green colour while values close to 90° to yellow. Chroma defines the colour intensity or purity of the hue. Values close to 0 correspond to neutral colours and values close to 60 to bright colours (McGuire 1992). Depending on the sampling scheme and accuracy, the L*, a*, b*, h* or C* values can provide a satisfactory colour description (Manolopoulou, Xanthopoulos, Douros & Lamprinos, 2010). Colour was evaluated at the beginning and on the 3rd, 6th and 10th day, in 6 packages per treatment (P1, P2, P3, and C) and storage temperatures.

2.8 Ascorbic Acid

Ascorbic acid was determined using the 2,6 dichlorophenolindophenol method (AOAC 1990) and expressed in mg per 100 g of the initial fruit mass. Measurements were taken at the beginning and at the end of the storage (10th day). Measurements of the ascorbic acid were carried out in 6 packages per treatment (P1, P2, P3, and C) and storage temperature.

2.9 Quality Evaluation

The visual quality of the packaged produce was assessed by a panel of six trained judges adopting a hedonistic scale from 1 to 9, rating with 9=excellent, no visual defects, 7=good minor defects, 5=fair moderated defects, 3=poor moderately severe defects, 1=unusable. In this study, score of 6 was considered as the marketable limit for a fresh-cut produce (López-Gálvez et al., 1997). The assessment was carried out at the beginning of the experiment and on the 3rd, 6th and 10th day, in 6 packages per treatment (P1, P2, P3, and C) and storage

temperature.

2.10 Experimental Design and Statistical Analysis

The experiment followed a full factorial design (storage temperature×treatment×storage time) and subjected to the analysis of variance (ANOVA) using Statgraphics Plus 5.1 (Statpoint Technologies, Inc, USA). Mean values were subjected to Fisher's Least Significant Difference test (LSD) at confidence level $p \leq 0.05$. The adopted test is liberal with respect to the comparison wise error rate, but is powerful for detecting true differences among means (Gacula & Sign 1984). In the figures and tables are presented the mean values from three experimental series, since no significant differences were found among the three experimental series ($p \leq 0.05$).

3. Results and Discussion

3.1 Respiration Rate

Respiration rate (RR) of the unpackaged fresh-cut and whole peppers was not significantly different at 0 °C but was significantly different at 5 °C (cf. Figure 1). At 5 °C the RR of the fresh-cut pepper increased from 0.62 mL CO₂ h⁻¹ 100 g⁻¹ (1st day) up to 0.98 mL CO₂ h⁻¹ 100 g⁻¹ (9th day) and then sharply decreased at 0.47 mL CO₂ h⁻¹ 100 g⁻¹ (10th day) probably due to the exhaustion of the necessary substrates for retaining high RR. The RR of the whole peppers at 5 °C reduced from 0.5 mL CO₂ h⁻¹ 100 g⁻¹ (1st day) to 0.27 mL CO₂ h⁻¹ 100 g⁻¹ and remained constant at this level for the last three days of the storage. At 5 °C, the initial RR of the fresh-cut peppers was 23.5% higher than the RR of whole produce. The measured RR values were similar to those reported by Kang & Lee (1997) who observed an increase of 38% in RR of fresh-cut peppers during a six-day storage at 5 °C. The mean RR values for the fresh-cut pepper at 0 °C was 0.27±0.13 mL CO₂ h⁻¹ 100 g⁻¹ and at 5 °C was 0.69±0.17 mL CO₂ h⁻¹ 100 g⁻¹. At 0 °C the difference of the mean RR values between whole and fresh-cut produce was <1% whereas at 5 °C was 47%. The statistical analysis highlighted that temperature control is important in reducing the injury induced metabolic activity in whole and fresh-cut produce although ANOVA pointed out that the interaction of storage time and treatments was also significant in both storage temperatures.

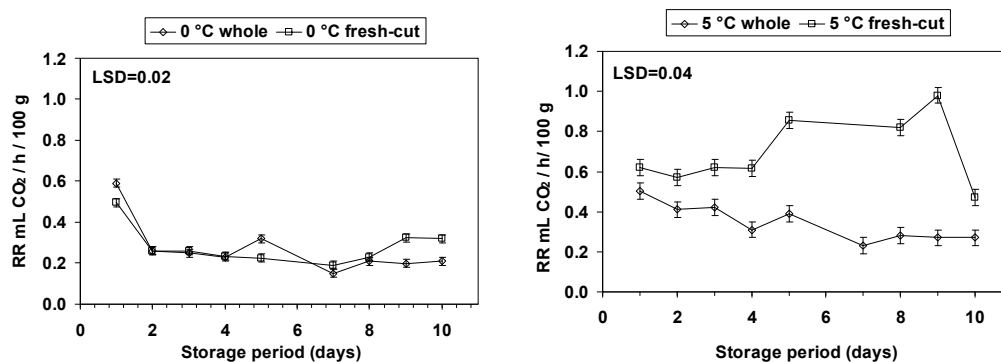


Figure 1. Respiration rates of unpackaged fresh-cut and whole peppers stored at 0 and 5 °C. Data points are means of 18 replicates \pm LSD

3.2 Gas Composition

Evolution of O₂ and CO₂ during the storage of fresh-cut peppers at 0 °C and 5 °C is showed in Figure 2. As it was expected, the CO₂ accumulation in the fresh-cut produce packaging was rapid at 5 °C. The ANOVA showed that O₂ and CO₂ evolution was storage temperature, storage time and treatment dependent. Significant difference noted in the CO₂ levels among the P1, P2 and P3 treatments. Similar O₂/CO₂ evolution observed for P1 and P2 treatments at both tested temperatures. At 0 °C, the O₂ decreased from the initial 5% to the final 0.5 % and 0.2 % for P1 and P2 respectively, whereas in P3 the O₂ concentration stayed above 19%. At 5 °C, the respiratory activity consumed O₂ up to the 5th day of storage although at the end of the storage no tissue injuries or off flavours were detected. Further investigation is needed on the effect of the headspace at low O₂ or high CO₂ concentrations, as well as the tolerance of the packaged peppers at the formed O₂/CO₂ in-package atmospheres. The O₂ levels between P1 and P2 for the two tested temperatures were not significant different. At 0 °C, the CO₂ concentration increased by 28%, from 10% to 12.8% at P1, by 8%, from 15% to 16.2% at P2 and at P3 increased up to 2.28%. At 5 °C, the CO₂ increased up to 6.03% at P3 while at P1 and P2 increased by 58%, from 10% to

15.8% and by 31%, from 15% to 19.6% respectively.

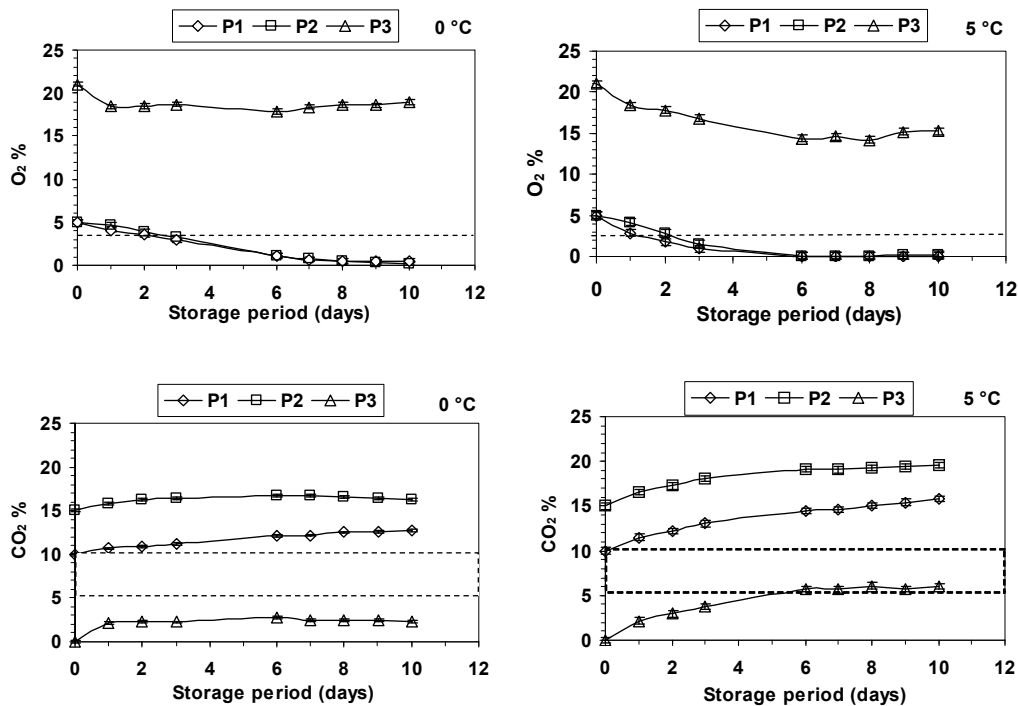


Figure 2. Changes in O₂ and CO₂ of packaged fresh-cut peppers stored at 0 and 5 °C. P1, P2, P3 are O₂:CO₂ mixtures respectively, 5%:10%, 5%:15%, 21%:0.03%. Data points are means of 30 replicates±LSD; [0 °C: LSD_{O₂}=0.25; LSD_{CO₂}=0.14; 5 °C: LSD_{O₂}=0.40; LSD_{CO₂}=0.33]. Horizontal dotted lines demarcate the suggested O₂ to CO₂ levels for fresh-cut green bell peppers

3.3 Mass Loss

Packaging reduced mass loss of *Twingo* F1 whole peppers (Manolopoulou et al., 2010). Similarly, the packaging film, used in P1, P2 and P3 reduced significantly mass loss of the fresh-cut produce (cf. Figure 3) with no visible formation of water drips in the packages. At the end of the storage (10th day) at 0 °C and 5 °C, packaged produce lost <0.5% of its initial weight, whereas control samples lost 1.4% and 2.6% respectively which lies below the acceptable limit of 5% (Manolopoulou et al., 2010). The ANOVA showed that the interaction of the storage temperature, storage time and treatment, affected significantly mass loss ($p \leq 0.001$).

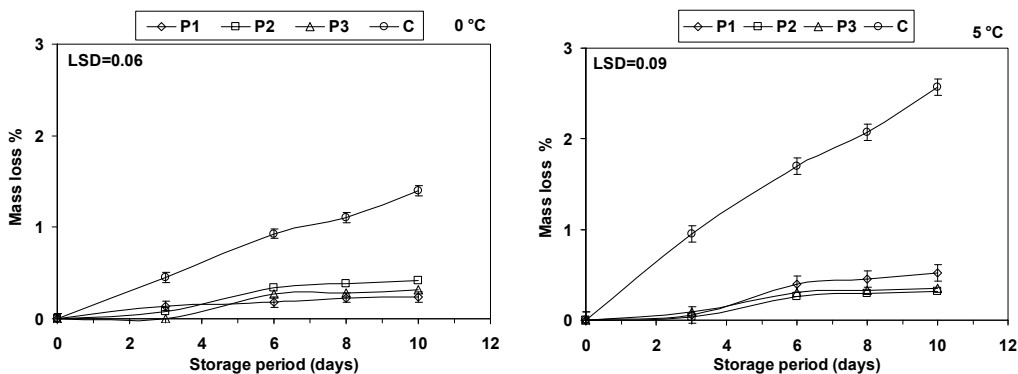


Figure 3. Mass loss of the packaged fresh-cut peppers stored at 0 and 5 °C. P1, P2, P3 are O₂:CO₂ mixtures respectively, 5%:10%, 5%:15%, 21%:0.03%. C denotes control samples. Data points are the means of 30 replicates±LSD

3.4 Firmness

The ANOVA showed that only the interaction of treatment with storage time affected significantly firmness ($p=0.0087<0.05$). At 0 °C, the firmness of the fresh-cut produce, (cf. Table 1) increased by 5-18% at the end of the storage (10th day), which was significant only for P1 and P2. At 5 °C, firmness increased at the end of the storage by 3-11%, which was significant only for P1. The employed firmness test is catastrophic and introduces an experimental error due to inevitable variability among the tested samples, which should be considered in the statistical inference.

Table 1. Effect of storage temperature, treatment and storage time on the mean rupture force (N) of packaged fresh-cut green bell peppers stored at 0 and 5 °C

Storage days	Storage Temperature	P1	P2	P3	C	F ANOVA
0	0 °C	337.4±12.8 A	337.4±12.8 A	337.4±12.8 A	337.4±12.8 A	
3		389.9±10.5 aBC	349.5±16.7 bAB	386.2±16.9 abC	357.9±10.7 abAB	2.31*
6		351.7±11.9 aAB	347.6±13.4 aAB	383.2±17.8 aBC	370.2±16.7 aAB	0.91 ^{NS}
10		399.8±11.7 aC	377.0±11.8 abB	339.5±14.5 bAB	355.8±12.9 bAB	2.92*
<i>F ANOVA</i>		5.52*	2.01 ^{NS}	4.07*	1.61 ^{NS}	
0	5 °C	337.4±12.8 A	337.4±12.8 A	337.4±12.8 A	337.4±12.8 A	
3		371.2±15.0 abBC	349.0±11.7 aA	386.9±13.5 bB	369.9±18.0 abB	1.43*
6		407.0±15.2 cC	327.3±15.8 aA	356.0±16.9 abAB	376.9±18.7 bcB	9.35*
10		373.7±15.2 aBC	356.3±16.5 aA	348.5±18.7 aAB	373.4±16.2 aB	0.68 ^{NS}
<i>F ANOVA</i>		7.42*	0.94 ^{NS}	2.85 *	3.49*	

Values followed by different lowercase letters in the same row, show significant difference among treatments, $p\leq 0.05$; Values followed by different uppercase letters in the same column, show significant difference among storage days, $p\leq 0.05$; Values denote means±std; NS=not significant; *=significant at $p\leq 0.05$.

Nunes & Emond (2003) discussed that increase of firmness is water loss induced and results in toughening of the epidermis in fleshy tissues, fact which was visible in control samples and less visible in P1, P2 and P3 treatments (cf. Figure 4a) where mass loss was limited (<0.3%) for both storage temperatures. Smith, Waldron, Maness & Perkins-Veazie (2003) reported that mass loss and firmness are related. Therefore, regression analysis of mass loss with respect to treatment and firmness, resulted in Eq. (1), where $R^2_{adj}=0.633$ and $SEE=0.279$. Therefore, if firmness is known, then mass loss can be evaluated without prior knowledge of the initial weight.

$$\begin{aligned} \text{Mass loss (\%)} = & 0.076 + 4.656 \times 10^{-4} \times \text{firmness} \times P1 - 1.463 \times 10^{-4} \times \text{firmness} \times P2 \\ & - 2.189 \times 10^{-4} \times \text{firmness} \times P3 + 23.395 \times 10^{-4} \times \text{firmness} \times C \end{aligned} \quad (1)$$

The P1, P2 and P3 terms are indicator variables taking the value 1 if true and 0 if false. Therefore Eq. (1) yielded four separate lines (cf. Figure 4a), one for each treatment. For P1 treatment, the model becomes:

$$\text{Mass loss (\%)} = 0.076 + 4.656 \times 10^{-4} \times \text{firmness} \quad (2)$$

and for P2 treatment:

$$\text{Mass loss (\%)} = 0.076 - 1.463 \times 10^{-4} \times \text{firmness} \quad (3)$$

Further statistical analysis of Eq. 1 showed that statistically significant relationship exists between the correlated variables and statistical significant difference exists among the slopes for C, P1, P2 and P3 ($p\leq 0.05$). Finally, the efficiency of the model (cf. Eq. 1) was further tested, plotting the residual error graph (cf. Figure 4b) to examine the *homoscedasticity* hypothesis. From this plot (cf. Figure 4b) was found that the *homoscedasticity* hypothesis is fulfilled since the plotted studentised residuals of the predicted mass loss (%) show no systematic patterns and are allocated around zero in a value zone between 3.0 and -3.0.

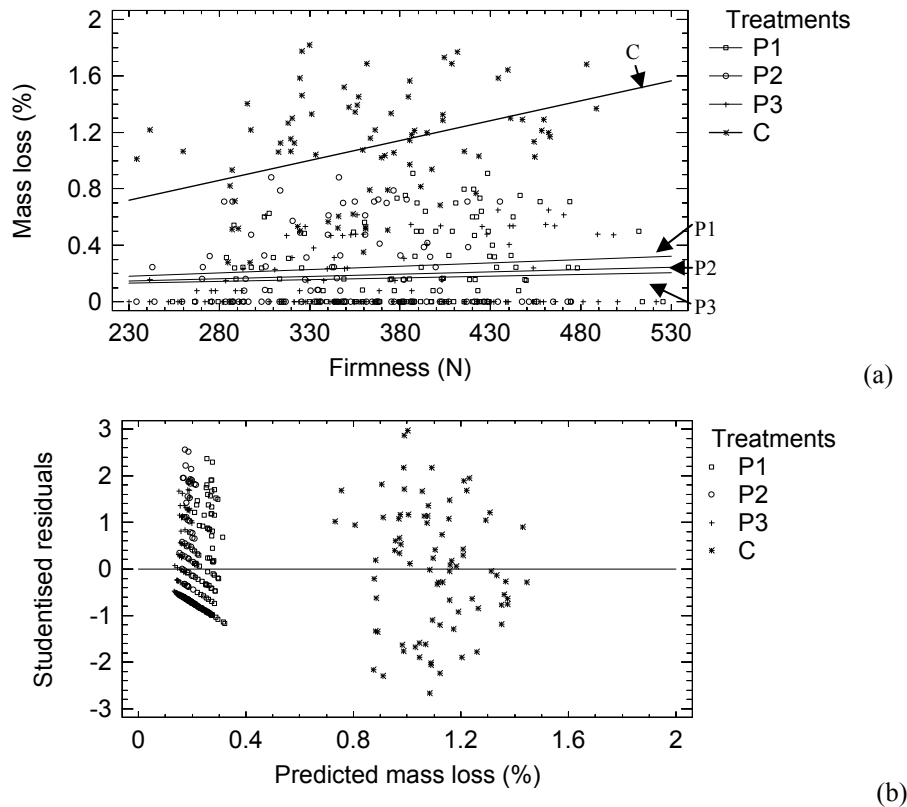


Figure 4. (a) Mass loss (%) change with firmness (N) of fresh-cut packaged peppers for P1, P2, P3 and C; (b) Residual error plot of the predicted mass loss (%). P1, P2, P3 are O₂:CO₂ mixtures respectively, 5%:10%, 5%:15%, 21%:0.03%. C denotes control samples

3.5 Colour Evaluation

The ANOVA of treatment, storage temperature and, storage time and the way these parameters affect colour parameters (L^* , C^* and h^*) are tabulated in Tables 2 and 3. The lightness (L^*) of the fresh-cut produce on the initial day was 32.5 ± 3.4 . At the end of the storage (10th day), the L^* values at 0 °C, were significantly reduced at P1 and P2 and were significantly different from the L^* values at P3. Only P2 had significantly different L^* values from the control L^* values. At 5 °C, the L^* values were significantly reduced at P1 and did not changed significantly at P2, P3 and C. The initial chroma (C^*) value was 25.8 ± 2.6 . At 0 °C, the C^* values were significantly reduced in all treatments at the end of the storage (10th day). At 5 °C, the P1, P2, P3, and C had significant in-between differences while C and P3 samples exhibited the highest, chroma reduction which was 21%.

The initial hue angle (h^*) was 117.5 ± 1.8 and presented not significant variations throughout the storage. The 10th day of the experiment, at 0 °C, the P1 and P2 were significant different from P3 and control samples. At 5 °C, P1, P2 and C did not have any in-between significant differences although they were significant different from the P3. In conclusion, low L^* or high h^* angle values at the end of the storage, indicated retention of the initial green colour which may be due to limited dehydration and/or limited chlorophyll degradation which is responsible for the green colour. Retention of the green colour due to limited chlorophyll degradation in fruits and vegetables can also initiated from the elevated CO₂ and/or low O₂ in-package concentrations according to Weichmann (1986). In all cases, the small L^* reduction (<14%), and h^* retention, throughout the MAP storage indicated retention of the initial green colour of the fresh-cut produce.

Table 2. ANOVA of the factors and their interactions affecting the L*, C* and h* colour parameters of the packaged fresh-cut peppers stored at 0 and 5 °C

		df	L*	C*	h*
<i>Factors</i>	A: treatment	3	1.95*	2.43*	1.64*
	B: storage temperature	1	0.14 ^{NS}	0.82*	2.01*
	C: storage time	3	3.10 ^{NS}	5.18*	2.70 ^{NS}
<i>Interactions</i>	A×B	3	0.15 ^{NS}	0.48*	1.38*
	A×C	9	1.12*	2.08*	0.46 ^{NS}
	B×C	3	2.10 ^{NS}	1.15 ^{NS}	1.48*
	A×B×C	9	1.56 ^{NS}	0.93 ^{NS}	1.20*

NS=not significant; *=significant at $p \leq 0.05$; SST=total sum of squares; df=degree of freedom.

Table 3. Effect of storage temperature, treatment and storage time on the L*, C* and h* colour parameters of the packaged fresh-cut peppers stored at 0 and 5 °C

Colour parameter	0 th day		3 rd day		6 th day		10 th day	
	0 °C	5 °C	0 °C	5 °C	0 °C	5 °C	0 °C	5 °C
<i>L*</i>								
P1	32.5±3.4	32.5±3.4	29.1±2.5	32.4±2.8	28.9±2.9	30.7±2.7	29.2±1.9	30.4±2.2
P2	32.5±3.4	32.5±3.4	33.6±2.6	31.3±2.1	29.8±3.3	30.4±3.0	27.9±2.5	32.8±2.5
P3	32.5±3.4	32.5±3.4	33.2±2.7	32.6±2.8	31.5±4.9	33.9±4.0	32.9±4.0	31.3±4.5
Control	32.5±3.4	32.5±3.4	30.9±3.9	31.5±4.4	31.5±4.7	31.4±4.9	30.8±4.8	32.1±4.5
<i>C*</i>								
P1	25.8±2.6	25.8±2.6	21.5±2.4	23.7±2.6	23.2±2.5	22.7±2.2	23.9±2.1	23.2±2.6
P2	25.8±2.6	25.8±2.6	25.5±2.6	24.1±2.6	24.1±2.6	23.9±2.2	23.6±2.4	25.9±2.2
P3	25.8±2.6	25.8±2.6	24.4±2.7	21.9±2.5	22.9±2.9	21.4±2.1	22.9±3.9	20.3±2.8
Control	25.8±2.6	25.8±2.6	22.4±3.3	20.8±2.5	22.4±2.5	20.5±3.2	21.4±2.6	20.4±3.5
<i>h*</i>								
P1	117.5±1.8	117.5±1.8	118.9±2.1	116.5±2.2	118.6±2.1	118.2±1.9	117.3±3.0	117.9±2.0
P2	117.5±1.8	117.5±1.8	118.1±2.0	116.0±2.0	117.7±2.2	117.4±2.4	117.5±2.4	117.3±1.9
P3	117.5±1.8	117.5±1.8	118.1±2.0	116.0±2.6	117.4±1.8	117.4±2.6	118.4±2.0	115.8±3.0
Control	117.5±1.8	117.5±1.8	118.0±1.8	117.1±2.7	118.1±2.0	116.7±3.2	118.5±1.7	117.3±3.2

0 °C: LSDL*=0.98; LSDC*=0.47; LSDh*=0.36; 5 °C: LSDL*=0.99; LSDC*=0.64; LSDh*=0.30; means±std.

3.6 Ascorbic Acid

The initial ascorbic acid (128.5 ± 2.0 mg 100 g⁻¹ of initial mass) retained in all treatments throughout the storage at 0 °C (cf. Table 4). However, at 5 °C a significant increase in the ascorbic acid (P1: 32%, P2: 25%, P3: 29% and C: 52%) was observed at the end of the storage, highlighting the temperature dependence of the ascorbic acid. These results agree with Senesi et al. (2000) and González-Aguilar et al. (2004) experiments on shelf-life of fresh-cut peppers under passive MAP. The combined effect of low storage temperatures (0 and 5 °C) and MAP was beneficial in retaining the initial ascorbic acid, although Klein (1987) referred that minimal processing result in loss of the ascorbic acid in fresh-cut than in intact produce as the oxidative process is accelerated. In contradictory conclusions (retention or increase of the ascorbic acid) were driven also Barth, Kerbel, Perry & Schmidt (1993), Paradis et al. (1996) for broccoli, and Howard & Hernandez-Brenes (1998) for jalapeno pepper storage.

Table 4. Effect of storage temperature, treatment and storage time on the mean ascorbic acid (mg 100 g⁻¹) of packaged fresh-cut peppers stored at 0 and 5 °C

Storage temperature		Ascorbic acid (mg/100 g)			
		P1	P2	P3	Control
0 °C	0 th day	128.5±2.0 aA	128.5±2.0 aA	128.5±2.0 aA	128.5±2.0 aA
	10 th day	124.0±2.8 Aa	119.5±1.2 aA	127.1±2.4 aA	126.2±1.5 aA
5 °C	10 th day	170.2±1.4 aB	160.2±2.1 aB	166.3±3.2 aB	195.2±4.1 bB

Values in the same column followed by different uppercase letters show significant differences between temperatures, $p \leq 0.05$; Values in the same row followed by different lowercase letters show significant differences among treatments, $p \leq 0.05$; Means±std.

3.7 Overall Visual Quality

The appearance of the fresh-cut produce was evaluated visually. The ANOVA (cf. Table 5) showed that the interaction of storage temperature with storage time affected significantly the visual quality of the fresh-cut produce. The previous inference is showed in Figure 5, where at 5 °C was found higher visual degradation than at 0 °C which agrees with López-Galvez, El-Bassuoni, Nie, & Cantwell (1997) who studied the visual degradation of red and green fresh-cut peppers under CA storage. Although Kang & Lee (1997) referred that storage of fresh-cut pepper at 5 °C can initiate chilling injuries, this was not confirmed in this study where chilling injuries did not found in any of the tested cases. The combined effect of minimal processing and MAP, according to Forney & Lipton (1990) and Gorny (2001), permit the use of low storage temperatures (-5 °C - 0 °C), with no danger of chilling injury occurrence. The low O₂ conditions at 0 °C and 5 °C did not produce off-flavours in the fresh-cut produce according to the panelists, which agree with González-Aguilar et al. (2004) who reported that traces of ethanol and acetaldehyde, produced during storage of fresh-cut green pepper at 8 °C were not sufficient to affect the sensory susceptibility of the end-product.

Table 5. Analysis of variance of storage temperature, treatment and storage time and their interactions as these affect the visual quality of the packaged fresh-cut peppers stored at 0 and 5 °C

		df	Visual Quality
Factors	A: Storage temperature	1	10.57%*
	B: Treatment	3	39.74%*
	C: Storage time	2	11.24%*
Interactions	A×B	3	0.12% ^{NS}
	A×C	2	5.20%*
	B×C	6	3.11% ^{NS}
	A×B×C	6	1.53% ^{NS}

NS=not significant; *=significant to $p \leq 0.05$; SST=total sum of squares; df=degree of freedom.

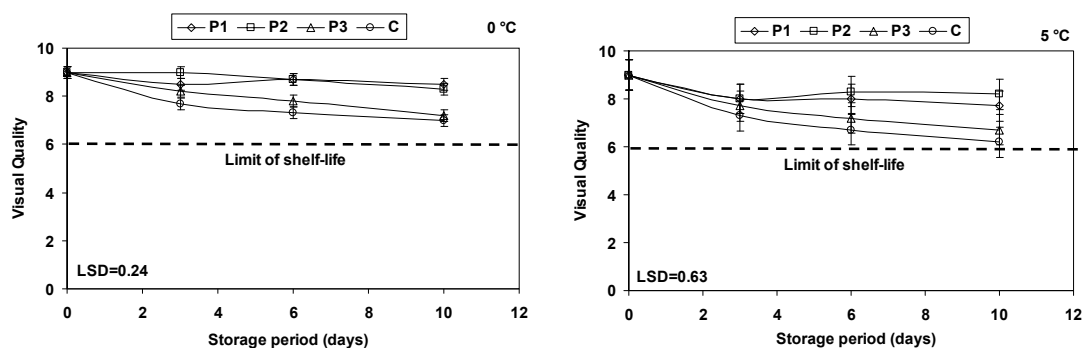


Figure 5. Visual quality of packaged fresh-cut peppers stored at 0 and 5 °C. P1, P2 and P3 are O₂:CO₂ mixtures respectively 5%:10%, 5%:15%, 21%:0.03%. C denotes control samples. Data points are the means of 108 replicates±LSD

4. Conclusions

The RR of the unpackaged fresh-cut produce was 24% higher than of the whole produce at 5 °C. The analysis showed that temperature control in minimally processed products is crucial for the related metabolic activity control. The in-package formed atmospheres found to be temperature, storage time and treatment dependent. At 5 °C the O₂ reduced significantly until the 5th day, although no tissue injuries or off flavours were detected at the end-product. No significant differences were found in the O₂ levels at 0 °C and 5 °C for the tested initial O₂ and CO₂ concentrations (P1 and P2). As the CO₂ level concerns, significant differences detected among the P1, P2 and P3 treatments. Mass loss was significantly affected by the storage temperature, storage time and treatment in P1, P2 and P3 treatments. Only treatment and storage time affected significantly firmness. At 0 °C the increase of the firmness at the end of the storage was significant in P1 and P2 treatments whereas at 5 °C the increase was significant only in P1 treatment. A mathematical correlation was developed relating mass loss, firmness and the tested treatments with satisfactory accuracy ($R_{adj}^2=0.633$). In all treatments the initial green colour was retained throughout the storage based on the final h* value which was not significantly different from the initial h* value. A significant increase of the ascorbic acid was observed at the end of the storage at 5 °C. The visual quality significantly affected from the storage temperature and storage time. Significant visual degradation observed at 5 °C although the scored values were above the acceptable limit. From the previous findings, can be concluded that fresh-cut bell peppers (cv *Twingo F1*) stored at 0 °C under the tested active MAP conditions can maintain a qualitative shelf-life up to 10 days.

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