Quality of mini tomatoes harvested at two maturity stages and kept chilled in three packages

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ABSTRACT

The economic importance of small tomato fruits has been increased considerably due to the significant levels of lycopene and sugars. However, to achieve these compounds, the harvest is limited to a ripening stage demanded by consumers, which could reduce this marked period. Harvesting at an early stage and use of conservation techniques are desirable to allow the marketing period. The aim of this study was to evaluate the quality of small tomato fruits, Sweet Grape cultivar, harvested at two stages of ripening (light red and red), kept in three types of package (perforated PET, PEBD 30 µ and Vegetal Pack 18 µ) at a temperature of 20±1ºC and a relative humidity (RH) of 85±5% during 20 days. During this period, gas composition inside the packaging (O₂, CO₂ and ethylene), peel color, fresh mass loss, soluble solids content (SS), titratable acidity (TA), SS/TA and ascorbic acid levels were evaluated every 5 days. The lycopene content was determined at the beginning and at the end of conservation. The experimental design was completely randomized in a split plot with four replications. Mass loss, peel color and gas composition were affected by both stage of harvest and package. The total SS/TA increased 4.74 during the conservation for two stages, regardless of package. The light red fruits reached after 20 days of conservation, mean levels of lycopene of 16.47 mg 100 g⁻¹ and SS/TA of 18.43; on the other hand the red fruits achieved 15.30 mg 100 g⁻¹ of lycopene and 16.80 of SS/TA. There was a mean increase of 8.4 mg 100 g⁻¹ of lycopene in light red fruits, while in red fruits, the increase was 0.9 mg 100 g⁻¹. The PEDB film was the best to maintain the fresh mass and peel color, therefore it was indicated for the maintenance of these attributes. The Vegetal Pack film showed no advantage in relation to the PET perforated packaging.

Keywords: Solanum lycopersicum, maturity stage, conservation, modified atmosphere.

RESUMO

Qualidade de mini tomates colhidos em dois estádios de maturação e conservados refrigerados em três embalagens

A importância econômica do mini tomate tem crescido consideravelmente, sendo um dos motivos os teores significativos de licopeno e de açúcares redutores. Para alcançar estes atributos, a colheita é realizada no ponto de consumo, o que reduz sua vida útil. Colheita em estádio precoce e uso de técnicas de conservação são desejáveis para ampliar o período de comercialização. Dentro deste contexto, o objetivo deste trabalho foi avaliar a qualidade de mini tomates, cultivar Sweet Grape, colhidos em dois estádios de maturação (vermelho claro e vermelho), mantidos em três embalagens (PET perfurada, PEBD 30 µ e Vegetal Pack 18 µ) a 20±1ºC e 85±5% de UR por 20 dias. Durante este período foram avaliadas a cada 5 dias a composição gasosa no interior das embalagens (O₂, CO₂ e etileno), a coloração da casca, a perda de massa fresca, o teor de sólidos solúveis (SS), a acidez total (AT) e o teor de ácido ascórbico. O teor de licopeno foi determinado no início e no final da conservação. O delineamento experimental foi inteiramente casualizado, em parcela subdividida, com quatro repetições. A perda de massa, a coloração da casca e a composição gasosa foram afetadas tanto pelos estádios de maturação quanto pelas embalagens. A relação SS/AT aumentou 4.74 durante a conservação, independente do tipo de embalagem. Os frutos em estádio vermelho claro alcançaram, aos 20 dias de conservação, teores médios de licopeno de 16,47 mg 100 g⁻¹ e relação SS/AT de 18.43, enquanto os frutos em estádio vermelho apresentaram 15,30 mg 100 g⁻¹ de licopeno e 16,80 de SS/AT. Houve incremento médio de 8.4 mg 100 g⁻¹ dos teores de licopeno nos frutos vermelho claro, enquanto que nos vermelhos, o aumento foi de 0,9 mg 100 g⁻¹. O filme PEBD foi o que melhor manteve a massa fresca e a coloração da casca, por isso foi o indicado para a manutenção destes atributos. O filme Vegetal Pack não apresentou vantagem em relação à embalagem PET perfurada.

Palavras-chave: Solanum lycopersicum, estádio de maturação, conservação, atmosfera modificada.
properties are intensified when the fruits are at the most advanced ripeness stage, ideal for consumption (Kader, 2008). For this, the ripeness stage of the tomato at the time of harvesting and also the control of pre-and post-harvest factors are essential to ensure the quality demanded by the consumers (Moura et al., 1999; Beckles, 2012). The ripeness stage recommended for harvesting is when the peel of the mini tomato presents more than 90% red coloring (USDA, 1991). However, as the tomato is a climacteric fruit, it is probable that harvesting at the light red stage (60 to 90% red colored peel) would give the productive and commercial sectors greater flexibility for its management.

Storing tomatoes post-harvest depends on various pre-harvest factors and using procedures that maintain their nutritional and organoleptic quality (Kader, 2008). One of the most successful techniques is using modified atmosphere, with plastic films, that represents a barrier to the exit of water vapor and gas exchanges and reduces the respiratory activity of the fruit (Kader et al., 1989). The permeability of the films to gases allows the carbon dioxide concentration resulting from the fruit respiration increase while the oxygen concentration decreases from its consumption in the respiratory process. Thus the fruit metabolism goes through biochemical and physiological alterations, leading to a considerable increase in its life after harvest (Day, 2002). The atmospheric composition inside the package depends on the permeability characteristics of the plastic film material, on the number of fruits packed and the storage temperature (Gorris & Peppelenbos, 1992; Mosca et al., 1999). Using packaging can favor fruit conservation, ensuring the maintenance of characteristics such as color, aroma, flavor and firmness.

Several studies have been carried out using plastic films associated or not with other post-harvest storage technologies in tomatoes, where use of films based on polyethylene and polypropylene has been emphasized to maintain the quality attributes related to the fresh mass and peel coloring (Vanndy et al., 2008; Sabir & Agar, 2011; Akbudak et al., 2012). Currently the perforated PET type of packaging is used for mini tomato that has no barrier to the carbon dioxide and oxygen gases. Other films, such as low-density polyethylene (PEBD) and Vegetable Pack are barriers against gases and thus alter the internal composition of the packages. The 30 µ thick PEBD film has a permeability rate to O2 of 7,489 mL (CNT) m-2day-1 and to CO2 of 27,900 mL (CNTP) m-2day-1 (Caron, 2009). Vegetable Pack, defined as a modifying self-control atmosphere membrane with zeolite technology (gas absorber), at 18 µ thickness, permits a permeability to oxygen of 9,185 mL (CNTP) m-2 day-1 and to CO2 of 41,100 mL (CNT) m-2 day-1 (Caron, 2009).

Thus the objective of the present study was to assess the quality of mini tomato harvested at two ripeness stages and stored in three types of packaging.

**MATERIAL AND METHODS**

Sweet Grape cultivar mini tomatoes were harvested in a commercial production area located in Mogi Mirim (22º26’0”S; 46º57’0”W), São Paulo state, Brazil, at two ripeness stages called light red and red (USDA, 1991). The fruits were transported in a non-refrigerated vehicle to the post-harvest laboratory of the Escola Superior de Agricultura Luiz de Queiroz. Later they were selected for presence of damage and cleaned in a sodium hypochlorite solution (200 ppm). Fruits with 60 to 90% of the peel with red coloring were considered for the light red ripeness stage. For the red ripeness stage, fruits were considered with more than 90% of the peel with red coloring, according to the USDA classification norms for tomato coloring (USDA, 1991).

The treatments consisted of three types of packaging (PEBD 30 µ, perforated PET and Vegetable Pack 18 µ) and two ripeness stages (light red and red). The fruits were packed on 180 g capacity 16x16x4 cm perforated PET trays. The Vegetable Pack plastic film was wrapped round the whole tray and sealed using an Everest® “Porta Película”. The PEBD film plastic bag covered the whole tray and was sealed using an Everest® Termo Plástico welder the same size as the tray. Mini tomatoes placed only on a tray with a perforated PET lid were considered the control.

Physical and chemical analyses were carried out (soluble solid content, ascorbic acid, titratable total acidity, peel coloring, mass loss and lycopene contents). Shortly after closing the packaging the initial mass and the gas composition (O2, CO2 and ethylene) were determined inside the packaging. The treatments were kept at 20±2°C for 20 days. Physical and chemical assessments were made every five days, except for the lycopene content that was measured on the first and last days of storage. The gas composition was assessed on the first and second day and then every five days.

For the destructive analyses, 10 whole fruits from each replication were ground using a Mixer and a homogeneous mass was obtained.

A complete randomized experimental design was used in a factorial scheme, with split plots in time with four replications consisting of 10 fruits each. The factors consisted of five assessment times, three types of packaging and two fruit ripeness stages.

**Monitoring gas composition inside the packaging (O2, CO2 and ethylene)**

**Carbon dioxide and oxygen evolution:** samples of air from the free space in the packages were collected using a PBI Dansensor gas analyzer, model Check Mate that removed approximately 2 mL gas through silicone scepters fixed in the packaging. The results were expressed in percentages of O2, CO2 and ethylene.

**Ethylene evolution:** 1 mL samples of air were removed from the free space in the package with a Hamilton 2.5 mL Gastight syringe and injected in a gas chromatograph (Thermofinnigan, model GC Trace 2000), with a flame ionization detector (FID), using a “Propak N” column. The drag gas was hydrogen with a 30 mL minute-1 flow. The temperatures maintained in the apparatus were 80°C for the column, 100°C for the injector, 250°C for the detector and 350°C for the metanador. To establish the standard curve, aliquots
of 1 mL of the 0.58 and 1.94 ppm ethylene standards were injected in the gas chromatographer. The ethylene evolution was expressed in ppm.

**Peel coloring** - two readings were made on the equatorial region of each fruit with a Minolta colorimeter, model cr-300, using the L*a*b* system. The results were expressed by the \( a^*/b^* \) ratio that designates the development index of the red color in tomatoes.

**Lycopene** - it was determined following methodology described by Sadler et al. (1990). One gram pulp was weighed from each replication in a 250 mL Erlenmeyer already wrapped in aluminum foil, and homogenized in a mixture of 50 mL hexane, 12.5 mL ethanol and 12.5 mL acetone at the proportion of 2:1:1, v:v:v. Shortly after mixing, the samples were centrifuged 30 min and then transferred to a separation funnel to which was added 10 mL distilled water. The solution was separated in polar and apolar fractions. The apolar fraction, with pigment, was measured by reading the hexane solution absorbency at 420 nm, using the hexane as standard. The results were expressed in milligrams of lycopene per 100 g pulp.

**Soluble solid content** - a sample of the pulp from each replication was placed in a digital refractometer (Atago PR-101), with automatic temperature correction to 20°C. The results were expressed in percentage.

**Titratable acidity** - 10 g of pulp from each replication were weighed and placed in 90 mL distilled water. The potentiometric titration was detected with sodium hydroxide 0.1 mol L\(^{-1}\) to pH 8.10. The calculations were made according to Carvalho et al. (1990) and the results expressed in percentage of citric acid in the pulp.

**Soluble solids/acidity ratio:** obtained by the ratio between the percentage of soluble solids and acidity.

**Mass loss** - calculated by the difference between the initial mass and the mass at the time of assessment, using semi-analytical scales, and the

![Figure 1](image.png)

**Figure 1.** Gaseous composition inside the package with small tomatoes during storage at 20±1°C and 85±5% RH. (A) PEBD and red stage; (B) PEBD and light red stage; (C) Vegetal Pack and red stage; (D) Vegetal Pack and light red stage. Vertical bars represent the standard deviation [composição gasosa no interior das embalagens com mini tomates ao longo do armazenamento a 20±1°C e 85±5% de UR. (A) PEBD e estádio vermelho; (B) PEBD e estádio vermelho claro; (C) Vegetal Pack e estádio vermelho; (D) Vegetal Pack e estádio vermelho claro. Barras verticais representam o desvio padrão da média]. Piracicaba, ESALQ, 2010.
results were expressed in percentage of initial mass.

**Ascorbic acid content** - it was determined according to methodology by Carvalho et al. (1990) based on reduction of the 2,6-dichlorophenolindolphenol-sodium (DCFI) indicator by ascorbic acid. Five gram of pulp were weighed from each replication and placed in an Erlenmeyer flask containing 25 mL 1% oxalic acid solution. The titratable acidity was assessed with DCFI until a persistent pink coloring was obtained for 15 seconds. The results were expressed in milligrams of ascorbic acid per hundred grams pulp.

The data were submitted to the Hartley test (p<0.05) to verify the homogeneity of variance among the treatments. Then analysis of variance was carried out by the F test (p<0.05 and p<0.01). According to the significance, the means of the qualitative data were compared by the Tukey test (p<0.05) and the means of the quantitative data were submitted to polynomial regression analysis (p<0.05).

**RESULTS AND DISCUSSION**

**Gas composition** - The oxygen concentration decreased inside the PEBD film and Vegetable Pack packaging during the 20 days of storage at 20°C. The PEBD film packaging had a more accentuated reduction of O₂ from 21 to less than 10%, and the carbon dioxide concentrations increased more, from 0.03 to 0.80%, than the other packaging (Figure 1A and B).

The 30 µ PEBD film resulted in lower oxygen concentrations inside the packaging after two days at 20°C with fruits at the light red stage compared to those with fruits at the red stage (Figure 1A and B). The Vegetable Pack film resulted in few alterations in the gas composition inside the packaging regardless of the fruit ripeness stage (Figures 1C and D) because oxygen decreased from 21 to 17% only at 10 days storage. The gas composition inside the perforated PET packaging (control) was the same as the environment (21% oxygen and 0.03% carbon dioxide) during the 20 days at 20°C (data not shown).

The gas concentration resulting from using PEBD film (10% oxygen and 0.8% carbon dioxide) for 20 days was justifiable because it presented lower permeability rates to carbon dioxide and oxygen than the other packaging types (Vegetable Pack and Perforated PET). This composition was similar to that reported by Akbudak et al. (2007) when they worked with heat treatment and used this film with 50 µ in two cultivars of mini tomato. These authors observed that after 20 days the gas concentration inside the packaging was 10 to 15% oxygen and 1 to 2% carbon dioxide.

The highest ethylene values were obtained on the fifth day of storage and then decreased (Figures 2A and B). Carbon dioxide accumulation and reduced oxygen concentration inside the packaging probably inhibited ethylene production after this date. This performance was similar in all the treatments, except for the red tomatoes packed in PEBD film (Figure 2B). The greatest ethylene accumulation in the PEBD film packaging regardless of the ripeness stage was probably due to the lower permeability to gases of this film. The Vegetable Pack, however, presented smaller ethylene accumulation because it may have permitted greater permeability of the gases and further because the zeolite mineral, that has adsorption capability, is part of its composition. Studies using plastic films impregnated with zeolite have shown that the gas composition varies in function of
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Figure 3. Evolution of peel color ($a^*/b^*$) of small tomatoes of red stage (A) and of light red stage (B) packaged in plastic films during 20 days 20±1ºC and 85±5% RH [evolução da coloração da casca ($a^*/b^*$) dos mini tomates de estádio vermelho (A) e de estádio vermelho claro (B) acondicionados em filmes plásticos durante 20 dias a 20±1ºC e 85±5% de UR]. Piracicaba, ESALQ, 2010.

this mineral inside the packaging and therefore the respiratory rate of the fruit (Huang et al., 2013). The type and quantity of this mineral in the plastic film has also been shown to interfere in ethylene absorption (Patdhanagula et al., 2010).

The difference in ethylene concentration among the ripeness stages in the PEBD film packaging may have been related to the different physiology of each stage. The fruit harvesting process generates stress and mechanical damage that results in cell signaling with the response of greater ethylene production (Ayub et al., 1995) that is accentuated when the fruits are not yet fully ripe, that is, they have not gone through the climacteric peak. When they reach the peak, ethylene production increases, completing the development of their physical and chemical characteristics. Chaves et al. (1998) studied transgenic tomato physiology at two ripeness stages for ethylene production and obtained higher concentrations of this gas in fruits ripened after harvest than in fruits ripened on the plant.

**Peel coloring** - The ratio between ripeness in post-harvest and change in the quality attributes of mini tomato was observed in the evolution of the peel coloring expressed by the $a^*/b^*$ ratio (Figure 3).

The red stage fruits were influenced by the type of packaging. When stored in Vegetable Pack, the $a^*/b^*$ ratio increased for peel coloring until the eighth day, but then decreased. Differently, fruits in the Perforated PET and PEBD film packaging expressed gradual increase in the $a^*/b^*$ ratio over the 20 days storage (Figure 3 A).

Fruits at the light red stage placed in PEBD film and Vegetable Pack presented a linear increase in the red coloring over time, but when they were placed in the Perforated PET packaging (control) the $a^*/b^*$ ratio decreased after 10 days at 21ºC (Figure 3B).

It can be inferred that the fruits stored in the PEBD and Vegetable Pack film were better preserved because the modified atmosphere resulted in reduced net metabolic activity of the fruits and consequently in the attributes that characterize their ripeness. In studies using modified atmosphere for tomato storage, PEBD film also delayed the development of the red coloring longer than the biodegradable packaging (Kantola & Hélen, 2001; Suparlan & Itoh, 2003).

**Lycopene content** - There was interaction among the types of packaging, fruit ripeness stages and storage time for the lycopene content. At 20 days storage at 20-21ºC the lycopene contents were higher in the light red fruit stored in the three types of packaging compared to the contents determined shortly after harvest. Furthermore, the fruits of this stage, when placed in the Perforated PET packaging, presented greater increase in the lycopene contents than the other treatments ($p<0.05$), ending at 20 days with 18.58 mg 100 g$^{-1}$ (Table 1). This fact did not occur for the red fruits whose contents were maintained without significant difference throughout the storage period.

The lycopene content of the tomatoes at the red stage was greater in the PEBD film than in the Vegetable Pack, at 20 days storage. The lycopene content was greatest in the light red tomatoes in the Perforated PET packaging (Table 1).

Lycopene content in tomatoes
depends on the ripeness stage and the development conditions of the fruits (Javanmardi & Kubota, 2006). Furthermore, modification in the atmosphere can interfere in the synthesis of this pigment because it reduces the fruit metabolic activities resulting in lower ethylene production and decreased physiological changes (Fonseca et al., 2002). As the tomato ripening process is associated with increase in lycopene content (Javanmardi & Kubota, 2006), delay in this process can result in lower contents of this compound, as reported by Siripatrawan & Assatarakul (2009) when they used 32 μ commercial film associated with methyl jasmonate to store unripe tomatoes.

In the mini tomatoes studied 8 to 18 mg 100 g⁻¹ lycopene were found during storage, equivalent to the contents determined by Pernice et al. (2010), Siripatrawan & Assatarakul (2009) in cherry tomatoes, with contents from 9 to 14 mg 100 g⁻¹ after 20 days storage.

**Soluble solids and acidity ratio** - The red fruit presented second-degree polynomial performance during storage with the highest soluble solids and acidity ratio at 17 days. The light red fruits resulted in linear regression with increase in this ratio up to 20 days storage (Figure 4C). These results corroborate Guillén et al. (2006), who demonstrated differences among the tomato ripeness stages during storage for the soluble solids and acidity ratio. However, in a study carried out with “Cronos” tomatoes, the soluble solids and acidity contents did not differ significantly for storage time and nor for the partially ripe and ripe ripeness stages (Brackman et al., 2007).

As the acidity and soluble solids contents directly influence fruit flavor, these results showed that tomatoes harvested at the light red stage, regardless of the packaging, showed increase in terms of flavor, while flavor started to decrease on the 17th day of storage at 22°C those harvested at the red stage.

**Mass loss (%)** - The fruits stored in the three types of packaging, regardless of the ripeness stage, presented mass loss over time. However, when in Perforated PET, the fruits lost about 3% fresh matter by the end of storage. When placed in Vegetable Pack and PEBD film, however, they lost 1% and 0.24%, respectively, by the end of the 20 days (Figure 4D).
The table below presents the effect of packaging and storage time (0 and 20 days) in the lycopene content (mg 100 g\(^{-1}\) in small tomatoes stored at 20±1°C and 85±5% RH [efeto das embalagens e do período de armazenamento (0 e 20 dias) no teor de lycopeno (mg 100 g\(^{-1}\)) em mini tomates armazenados a 20±1°C e 85±5% de UR]. Piracicaba, ESALQ, 2010).

<table>
<thead>
<tr>
<th>Package</th>
<th>Day 0</th>
<th>Day 20</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Light red stage</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PET (perforated)</td>
<td>8.04 ± 1.20 B a</td>
<td>18.58 ± 0.56 A a</td>
</tr>
<tr>
<td>PEBD</td>
<td>8.04 ± 1.20 B a</td>
<td>14.72 ± 2.21 A b</td>
</tr>
<tr>
<td>Vegetable Pack</td>
<td>8.04 ± 1.20 B a</td>
<td>16.12 ± 2.15 A b</td>
</tr>
<tr>
<td><strong>Red stage</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PET (perforated)</td>
<td>14.32 ± 1.89 A a</td>
<td>15.66 ± 1.15 A ab</td>
</tr>
<tr>
<td>PEBD</td>
<td>14.32 ± 1.89 A a</td>
<td>16.42 ± 1.39 A a</td>
</tr>
<tr>
<td>Vegetable Pack</td>
<td>14.32 ± 1.89 A a</td>
<td>13.82 ± 2.20 A b</td>
</tr>
</tbody>
</table>

Means followed by the same upper-case letters in the row and lower-case letters in the column did not differ from each other, Tukey test, 5% (médias seguidas de mesmas letras maiúsculas na linha e minúsculas na coluna não diferem entre si pelo teste de Tukey, ao nível de 5% de probabilidade).

Of the two films the PEBD film presented greater fruit mass conservation over time. In other studies, the greater conservation of mass was also obtained in 'Chiranjeevi' cultivar tomatoes when stored in PEBD and non-perforated PET packaging (Yadav et al., 2009). The same occurred for cherry tomatoes when stored in 50 µ PEBD film (Akbudak et al., 2007). This fact was due to the greater accumulation of water vapor inside the PEBD film packaging that helps maintain fruit turgidity (Akbudak et al., 2007).

**Ascorbic acid content** - There was interaction between the fruit ripeness stages, packaging type and storage time for ascorbic acid content (Figure 4A and B). For the red fruits, stored in PEBD film packaging, the ascorbic acid content presented third-degree polynomial performance (Figure 4A). However, for the fruits in Perforated PET and Vegetable Pack packaging, the ascorbic acid content remained constant throughout storage, with a general mean of 28.87 and 29.18 mg 100 g\(^{-1}\), respectively (Figure 4A).

The ascorbic acid content in light red fruits in Vegetable Pack decreased after the 10\(^{th}\) day of storage. However, when packed in Perforated PET they remained with contents of 29.77 mg 100 g\(^{-1}\) throughout the storage period and in PEBD film they presented contents ranging from 25 to 30 mg 100 g\(^{-1}\) (Figure 4B).

Ascorbic acid loss was greater (p<0.05) for the fruits at the light red stage, around 5 mg 100 g\(^{-1}\) at the end of storage, when placed in the PEBD film and Vegetable Pack packaging. This may have occurred as a result of a more accelerated metabolism in these fruits due to stress from early harvest and the lower oxygen concentrations resulting from the modified atmosphere. Studies on the performance of ascorbic acid synthesis and oxidation in tomatoes have shown that stress due to low oxygen levels can reduce synthesis and also oxidation, because the specific genes related to these two processes are deactivated or inhibited for a certain period (Loannidi et al., 2009). This also explains the constant ascorbic acid contents during storage for the fruits packed only in Perforated PET.

The ascorbic acid content in the tomato fruit varies according to the ripening period, cultivar, lightness fertilization and substrate (Sampaio & Fontes, 1998). Holcman (2009) reported ascorbic acid contents from 19 to 24 mg 100 g\(^{-1}\) for the Sweet Grape and Sweet Million tomato cultivars, respectively, cultivated in a protected environment. These contents are similar to those obtained for the mini tomato in the present study (23 and 35 mg 100 g\(^{-1}\), minimum and maximum values).

The Sweet Grape tomatoes harvested at two ripeness stages performed differently in the post-harvest. The fruits harvested at the light red stage imply an increase in the lycopene content, the red index and soluble solids/acidity ratio during the 20 days. This showed that tomatoes can be harvested before they reach the red stage that allows a longer commercialization period.

The PEBD film altered the gas composition inside the packaging with mini tomato, so that the peel color was maintained. Furthermore, the barrier to water vapor resulted in a smaller fresh mass loss, keeping the fruits turgid for longer so that this film is the most indicated for maintaining these attributes. The Vegetable Pack film, however, presented no advantage compared to the Perforated PET packaging (control).

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