Research Article



Influence of Harvesting Stages and Packaging on Storage Life of Apricot (*Prunus armeniaca* L.) Cv. Trevett

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Abstract | Supply of quality fruits to consumer is a major challenge due to storage and packing issues in developing countries. The current study was designed to evaluate postharvest quality of apricot at different harvesting stages i.e., M1 (greenish), M2 (light yellow) and M3 (orange-yellow) while, storage conditions i.e., packed in plastic bags with F_0 (closed), F_5 (5 pores), F_{10} (10 pores), F_{15} (15 pores), F_{20} (20 pores) and F_{25} (25 pores) at 5±2°C. The fruits were then analyzed for different quality parameters at 0 and 25 days storage. The harvesting stages (HS), perforated plastics (PP), storage duration (SD) as well as HS×SD and PP×SD significantly influenced various quality attributes of apricot fruits. The fruits harvested at the greenish stage had the lowest physiological weight loss and decay index that shows an increase in fruits harvested at orange yellow stage respectively after 25 days of storage. Similarly, after 25 days of storage, fruits stored in plastic bags having 10 pores retained fruit weight and attained minimum physiological weight loss with low decay index. Similarly, the fruits harvested at the greenish stage and stored for 25 days attained the highest titratable acidity, fruit firmness, and ascorbic acid in comparison with other harvesting stages. The highest titratable acidity, fruit firmness, and ascorbic acid were recorded in 10 pores plastic bags, while the lowest titratable acidity, fruit firmness and ascorbic acid were recorded in closed plastic bags after 25 days of storage. The lowest value of TSS and TSS: acid was recorded at a greenish stage, which increased at later stages of harvesting. In case of Pp ×SD, the range of TSS and TSS: Acid at day 0 increased after 25 days of storage with the lowest TSS and TSS: Acid was observed in plastic bags having 10 perforations. Apricot fruits harvested at the greenish stage (10.31 ° Brix) packed in plastic bags having 10 perforations shows the best in retaining most of the quality parameters at storage.

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Keywords | Apricot shelf life, Fruit quality parameter, Harvesting stages, Perforated plastics, Stages and Packaging



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open Oaccess Introduction

pricot (Prunus armeniaca L.) is a deciduous $\boldsymbol{\Gamma}$ fruit belongs to the family Rosaceae and is mostly grown in temperate climate of all continents of the world. Apricot is referred to stone fruits, with outer fleshy exocarp and mesocarp (skin and flesh) surrounds a stone of hardened endocarp having seed inside. Stone fruits also include peach, plum, nectarine, almond and cherry, which are closely related to each other, because of large and hard stone inside. All these belong to the same genus: Prunus, family Rosaceae (Penn state Extension, 2015). Among all stone fruits apricot has unique importance because of its great market value as of fresh as well as dried food commodity and has good market share of agriculture income. It has enough amount sugar, protein crude fibre, crude fate, Vitamin A, C, K, B complex and organic acid (Fatima et al., 2018). Turkey is the world largest producer of apricot with total production of 750,000 tonnes, followed by Uzbekistan and Iran with production of 493842 and 342479 tonnes respectively. Pakistan ranked seventh globally with total production of 128382 tonnes (FAO, 2018). This production is much lower as compared to advance countries and there is still a huge potential to boost the apricot production in the country. Post apricot is a climacteric fruit, which makes it susceptible to spoilage and post-harvest decay after being picked and became susceptible to more shrinkage and dehydration due to non-waxy nature of the skin (Ali et al., 2013). To keep these commodities fresh for as long as possible, their respiration rate should be decreasing without harming the quality of the fruit commodities. Generally, the respiration process is reduced by low temperature and modified atmosphere (Hussain, 1986). For this purpose, polymeric films have been used for packaging of fresh fruits and vegetables for a long time to control water loss, protect it from skin abrasion and reduce several other metabolic activities. It also has the ability to minimize the spread of disease from one fruit to another (Kader, 1980). The reduced metabolic activities depend on the relative permeability of the films, which allow the regulation of respiratory gases. This makes a modified atmospheric condition, which allows low level of oxygen and high level of carbon dioxide within the package and ultimately reduces the respiration rate of the fruits (Scetar et al., 2010). A single film in modified atmosphere packaging cannot provide all the properties needed for a modified atmosphere package (Alique et al., 2003).

Modification in these films alters the gas permeability and rate of respiration at specific temperature (Scetar *et al.*, 2010). Similarly, optimization of pre harvest management practices like application of GA3 and harvesting the fruit at proper physiological maturity stage increases the storage life of stone fruits (Vangdal *et al.*, 2012). The critical factor that influences the fruit quality and storage performance of fruit is to harvest them at proper maturity stage (Crisosto *et al.*, 2004).

Pakistan has massive potential for apricot production and is among the main producers in the world. However, limited literature is available on the storage life of apricot fruit from this region. There is increasing cultivation of apricots year by year in Pakistan and the fruit produced here has good quality and high market value but unfortunately the local producer is totally unaware about its storage under the prevailing environmental conditions which results in huge postharvest losses. Keeping in view the above facts, this study was planned to determine the post-harvest life of apricot fruits in cold storage enclosed in perforated plastics harvested at different harvesting stages.

Materials and Methods

This experiment was planned to retain postharvest quality attributes of fruit for a longer period during cold storage. For this purpose, the fruits of uniform size (20-22gm) were harvested at three different stages of maturity according to the skin color, representing M_1 (greenish stage with total soluble solids 10.31 °Brix), M₂ (light yellow stage with total soluble solids 12.32 °Brix) and M₂ (orange-yellow stage with total soluble solids 13.62 °Brix) from the healthy trees located at Horticulture Research farm, The University of Agriculture Peshawar, Pakistan. Fruits were harvested at each stage and divided into two lots. Lot no 1 fresh fruits of different stages were analyzed for various qualities attributes i.e. (Physiological weight loss, decay index, Fruit firmness, Fruit juice pH, Total soluble solid, Titratable acidity, TSS: acid ratio and Ascorbic acid), while the fruits of lot no 2 (54 no of fruits of each harvesting stage with 09 number of fruits in each packaging) were enclosed in plastic bags (20 x 30 cm, 30 microns) with different perforation (diameter of each pore was 2mm and these perforations were made through 2mm hole puncher) representing F_0 (0 pores), F_5 (5 pores), F_{10} $(10 \text{ pores}), F_{15}(15 \text{ pores}), F_{20}(20 \text{ pores}), F_{25}(25 \text{ pores})$ and shifted to cold storage and stored for 25 days at

 5 ± 2 °C. These perforations were equally distributed in each plastic bag and the plastic bags were placed in such a position in the cold storage so that all the perforations remained open.

The experiment was laid out in Completely Randomized Design (CRD) with three factors replication. The analysis of variance (ANOVA) for different parameters was determined using Statistix 8.1. The data of different parameters were analyzed through ANOVA techniques to observe the differences among different parameters as well as their interactions. In cases where the differences were significant at P≤0.05 the means were separated by Least Significant Difference (LSD) test (Steel and Torrie, 1997). Data on the following parameters were recorded during the experiments.

Physiological weight loss

Physiological weight loss was determined using an electrical balance by the difference between the initial and final weight of each treatment. Each fruit was weighed individually and then averaged. The following equation was used to determine the physiological weight loss, which is expressed in percent (%) (Amayogi and Alloli, 2007).

Decay index (%)

It was determined by dividing the percentage of fruits showing disease or decay incidence visually by total fruits stored in each treatment of all replication.

$$Decay incidence (\%) = \frac{decayed fruits}{Total fruits} \times 100$$

Fruit firmness

Fruit firmness was determined using hand held penetrometer of the randomly selected fruits for each treatment in each replication (Effigi, FT-11) (Pocharski *et al.*, 2000).

Fruit juice pH

pH meter (Model No. INOLAB. pH 720) was used for measuring fruit juice pH of the randomly selected fruit.

Total soluble solids (°Brix)

Total soluble solids (TSS) was recorded with the help of hand refractometer (Ranganna, 1986). A small quantity of crushed fruit pulp of randomly selected fruit was used for extracting few drops of juice that were placed on clean dry prism of the hand refractometer (Kernco Instruments Co Texas) and the reading was recorded. Distilled water and tissue paper were used to clean the prism between consecutive readings.

Titratable acidity (%)

Randomly selected fruits in each replication for all treatments were analyzed for titratable acidity as determined by the recommended method of (AOAC, 2000). 10 ml juice sample of randomly selected apricot fruit was taken in a beaker and their volume was increased up to 100 ml by addition of distilled water. Then the 10 ml diluted sample was titrated against 0.1N NaOH solution using 2-3 drops of phenolphthalein as an indicator. Data was recorded after the appearance of pink color, which persisted for 15 seconds. The data was recorded three times in each treatment and the following formula was used to calculate percent acidity. Average was calculated later.

$$Titratable \ acidity \ (\%) = \frac{T \times F \times N \times 100}{D \times S} \times 100$$

Where; T = Burette solution (ml of NAOH); N= Normality of sodium hydroxide (NAOH); S = Quantity of diluted sample (ml) taken for titration; D = Quantity of sample (gm) taken for dilution; F = Constant acid factor (primary acid in the fruit) = 0.0067 (malic acid in apricot).

TSS: Acid ratio

The TSS: Acid was calculated by using the formula given below:

$$TSS: Acid = \frac{Total Soluble Solid (TSS)}{Titratable Acdity} (Lacey et al., 2009)$$

Ascorbic acid (mg. 100g⁻¹)

Dye method was used for determining ascorbic acid contents in randomly selected fruits in each replication for all the treatments according to the method of (AOAC, 2000). 0.4% oxalic acid solution was used for the dilution of 10 gm of apricot fruit pulp and then its volume was raised to 100 ml. 10 ml of this diluted sample was then titrated against the dye solution till the appearance of light pink color. Ascorbic acid was then calculated using the following formula.

Ascorbic Acid content =
$$\frac{T \times F \times 100}{D \times S} \times 100$$

Where; F= dye factor; T= dye solution (ml) used for titration; D= diluted sample (ml) taken for titration; S= Apricot fruit pulp (gm) taken for titration.



OPEN access Results and Discussion

Physiological weight loss (%)

Perforated plastic (Pp), harvesting stages (HS) and storage duration (SD) had as significant influence on physiological weight loss of apricot fruits (Table 1). The highest physiological weight loss (4.18%) occurred in fruits harvested at orange yellow stage (M_2) , followed by fruit harvested at light yellow stage (M_2) with (3.11%) weight loss after 25 days of storage. However, fruits harvested at greenish stage (M_2) retained their weight with minimum (2.87%) weight loss after 25 days of storage in comparison with fresh fruits. Similarly, maximum weight loss (4.04%) occurred after 25 days of cold storage duration in closed plastic bags (F_0) , while the minimum weight loss (2.82%) occurred in plastic bags having 10 pores (F_{10}), which again increased to (3.69 %) in plastic bags with 25 perforations F_{25}). Respiration and water loss in fruits is a continuous process, which causes fruits deterioration and post-harvest losses. Maximum fruit weight loss in closed plastic bags could be because of O_2 injury and low O_2 levels (anaerobic condition) (Tariq et al., 2001). To avoid such conditions, there must be control flow of gases at optimum rate depending on the nature of the film and type of fruits (Pretel et al., 2000). Similarly, more physiological weight loss occurred in fruits harvested at later stages of maturity could be attributed to higher respiration and transpiration rates (Ngcobo *et* al., 2012). In addition, the Hardenburg et al. (1990) also stated that water loss through transpiration and loss of carbon in respiration process is responsible for weight loss in different fruits, which generally increases with delaying harvesting. The results are also in line with Rashidi et al. (2014) who reported that moisture content of nectarine could be conserved with wrapping and packaging material. Similarly, lower moisture loss was recorded in grapes during storage. Further, Sabir et al. (2011) also reported that water loss in minimally processed table grapes can be prevent through MAP, as a most common wrapping material. Rab et al. (2016) also reported less weight loss in plum fruits harvested at pale green stage and packed in plastic bags having 15 pores after 25 days of cold storage duration.

Decay index (%)

Fruits harvested at various stages were observed with varying decaying indexes More fruits were decayed (18.05%) at orange-yellow stage (M_3) , followed

by fruits harvested at light yellow stage (M_2) with (4.86%) decay while minimum increase (4.17%) occurred in fruits harvested at greenish stage (M₁) at the end of 25th day of storage in comparison with fresh fruits. Similarly, after 25 days of storage in cold storage conditions maximum decay index (26.39%) was observed in fruits stored in closed plastic bags (F_{0}) , followed by F_{5} (plastic bags with 5 pores), while minimum decay index (2.77%) occurred in plastic bags with 10 pores (F_{10}), which again increased 6.94%) in plastic bags with 25 perforations (F_{25}). Fruits in closed plastic bags were found more susceptible to deterioration as compared to the perforated ones. Closed plastic bags do not allow the moisture contents to go out, released during continuous transpiration and hence accumulate more humidity. Besides this, high CO₂ injury and humid conditions in closed plastic bags encourage microbial activities and as a result decaying percentage increase (Jawandha et al., 2012). Navjot and Sukhjit (2010) further reported that increase in respiration rate, enzymatic activities and disassociation of cells in the cell wall results in higher spoilage in later harvested fruits, which leads to the softening and ripening of fruits. The Juan *et al.* (1999) observed similar results in apple.

Rab *et al.* (2016) reported least disease incidence in plum fruit harvested at pale green stage, and packed in plastic packaging having 15 perforations after 25 days of cold storage duration. Our results are also in line with Khan *et al.* (2013), who reported that fruit firmness of plum fruit can be retained with different packaging material. Similarly, decrease in plum fruit firmness was recorded by with increasing durations (Rashidi and Bahri, 2014).

Fruit firmness (kg.cm⁻²)

Fruit firmness is one of the important quality attributes. It was observed that fruits showed significant variations in firmness at different stages stored for different durations in cold and normal conditions (Table 1). Initial fruit firmness of greenish stage (M_1), light yellow stage (M_2) and orange yellow stage (M_3) was recorded 3.55 kg.cm⁻², 2.01 kg.cm⁻² and 1.54 kg.cm⁻², respectively which, finally decreased to 0.64 kg.cm⁻², 0.57 kg.cm⁻², and 0.38 kg.cm⁻², respectively. Greenish (M_1) stage showed maximum fruit firmness (0.64 kg.cm⁻²), while minimum fruit firmness was recorded in orange yellow (M_3) after 25 days of cold storage. Similarly, initial firmness of the fruits for F_0 , F_5 , F_{10} , F_{15} , F_{20} and F_{25} was 2.37 kg.cm⁻²,



2.36 kg.cm⁻², 2.38 kg.cm⁻², 2.35 kg.cm⁻², 2.36 kg. cm⁻² and 2.36 kg. cm⁻², respectively, which decreased to 0. 42kg.cm⁻², 0.54 kg.cm⁻², 0.65 kg.cm⁻², 0.61 kg.cm⁻², 0.5 kg.cm⁻² and 0.47 kg.cm⁻², respectively. The maximum fruit firmness (0.64 kg. cm⁻²) was recorded in F_{10} , while minimum fruit firmness (0.42 kg. cm⁻²) was recorded in F_0 after 25 days of storage. Our results are in agreement with (Tariq *et al.*, 2001); they stated that the fruit firmness decreases as result of fruit damages due to the decreased concentration of O_2 and increase CO_2 in modified atmosphere. Such condition leads to an anaerobic condition, which further accelerates the catabolic process fruits became softer and reduced firmness. Our finding also confirms the results of Jan et al. (2012) who concluded that the fruits harvested at early stage of maturity have maximum fruit firmness, while least firmness was recorded in later harvesting stages. The texture of the flesh and changes in primary cell wall during ripening is responsible for the firmness of the fruit. Enzymatic activities, pectin solubilization and disassociation of the primary cell wall and middle lamella structures might be involved in it, which results in the decrease of mechanical strength of cell walls and thus the fruit firmness decreases (Kov and Felf, 2003; Kov et al., 2005).

Harvesting stages	Physiological weight loss (%)		Decay index (%)		Fruit firmness (Kg. cm ⁻²)	
	0 day	25 th day	0 day	25 th day	0 day	25 th day
Greenish stage (M_1)		2.87c		4.17b	3.55a	0.64d
Light yellow stage (M_2)		3.11b		4.86b	2.01b	0.57d
Orange yellow stage (M_3)		4.18a		18.06a	1.54c	0.38e
LSD at 0.05		0.21		2.48		0.04
Perforated plastics						
F ₀		4.04a		26.39a		0.42e
F ₅		3.28cd		8.33b		0.54cd
F ₁₀		2.82e		2.78e		0.65b
F ₁₅		3.05de		4.17cd		0.61bc
F ₂₀		3.44bc		5.55bcd		0.50de
F ₂₅		3.69b		6.94bc		0.47de
LSD at 0.5		0.30		3.52	ns	0.07

Table 1: Effect of harvesting stages × storage duration and perforated plastics × storage duration on physiological weight loss (%), decay index (%), and fruit firmness (kg.cm⁻² of apricot fruit Cv. Trevatt after 25 days of storage (5+2 °C).

Means followed by similar letter(s) in column do not differ significantly from one nother. Ns: Non significant.

Table 2: Effect of $HS \times SD$ and perforated plastics \times storage duration on ascorbic acid (mg. $100g^{-1}$), total soluble solid (° Brix), titratable acidity (%) and total soluble solids: Acid of apricot fruit Cv. Trevatt after 25 days of storage (5±2 °C).

Harvesting stages	Ascorbic acid		Total soluble solid		Titratable acidity		TSS:Acid	
	0 day	25 th day	0 day	25 th day	0 day	25 th day	0 day	25 th day
Greenish stage (M_1)	1.00a	0.33d	10.31f	12.92d	0.94a	0.41d	10.95e	31.71c
Light yellow stage (M_2)	0.75b	0.24e	12.33e	14.09b	0.84c	0.40d	14.73d	35.90b
Orange yellow stage (M_3)	0.63c	0.15f	13.63c	16.23a	0.88b	0.6e	15.47d	49.37a
LSD at 0.05		0.09		0.01		1.21		0.04
Perforated plastics								
F _o	0.80	0.13f	12.09	14.71a	0.88	0.37e	13.76	42.02a
\mathbf{F}_{5}	0.79	0.26cd	12.12	14.33bc	0.89	0.40cd	13.70	37.74c
F ₁₀	0.81	0.36b	12.07	14.17d	0.89	0.42b	13.64	36.22c
F ₁₅	0.79	0.30c	12.09	14.26cd	0.89	0.41bc	13.64	37.26c
F ₂₀	0.79	0.22de	12.10	14.40b	0.88	0.38de	13.79	39.90b
F ₂₅	0.77	0.17ef	12.07	14.60a	0.88	0.38de	13.75	40.80ab
LSD at 0.5	ns	0.13	ns	0.02	ns	1.71	ns	0.06

Means followed by similar letter(s) in column do not differ significantly from one another. Ns: non significant.

Ascorbic acid (mg. 100g-1)

Apricot fruits harvested at various stages were observed with different concentrations of ascorbic acid. Similarly, perforated plastic and storage duration also significantly influenced the ascorbic acid of apricot fruits (Table 2). The initial data recorded for different maturity stages was 1.0, 0.75 and 0.63 mg. $100g^{-1}$ for M₁, M₂ and M₃ respectively, which declined to 0.33, 0.24 and 0.15 mg. 100g⁻¹, respectively. The study also showed maximum ascorbic acid (0.33 mg. 100g⁻¹) in fruits harvested at greenish stage (M₁), while the minimum ascorbic acid $(0.15 \text{ mg}, 100\text{g}^{-1})$ was seen in fruits harvested at orange-yellow stage (M_{2}) . Similarly, he initials ascorbic acid contents of the fruits stored in various perforated plastic bags was 0.80, 0.79, 0.81, 0.79, 0.79 and 0.77 mg. 100g⁻¹ for F_0 , F_5 , F_{10} , F_{15} , F_{20} and F_{25} , respectively which, finally decreased to 0.13, 0.26, 0.36, 0.30, 0.22 and 0.17 mg. 100g⁻¹ respectively after 25 days in cold storage. However, maximum ascorbic acid (0.36 mg. 100g⁻¹) was reported in plastic bags with 10 perforations (F_{10}) while the minimum ascorbic acid $(0.13 \text{ mg}, 100\text{g}^{-1})$ was recorded in plastic bags having no pores (F_0) after 25 days of storage. Our result was confirmed by Chambroy et al. (2013), stated that the level of CO_2 and O_2 concentration is dependent on film permeability at a defined temperature. Findings are in line with Shafique et al. (2006) and reported similar decease in ascorbic contents. Nasrin et al. (2008) also recorded similar results and reported slight decline in vitamin content in tomato by packing it in perforated polythene bags during refrigeration storage. Lee and Kader (2000) stated that the regulation of oxidative process might be responsible for the reduction of ascorbic acid in fruit. They also concluded that many other factors like temperature, gaseous composition of storage and post-harvest storage might also induce heavy losses to ascorbic acid.

Total soluble solids (°Brix)

Total soluble solids determine the proper maturity time of the fruits and have close relation with harvesting stages, packaging materials and storage duration. Initially, data showed that there were 10.31, 12.33, and 13.63 (°Brix) for greenish stage M_1), light yellow stage M_2) and orange yellow stage (M_3), which were increased to 12.92, 14.09, and 16.23°Brix), respectively. On the other hand, total soluble solids of the fruits, stored in perforated plastic bags for 25 days were12.09, 12.12, 12.07, 12.09, 12.10 and 12.07 °Brix, for F_0 , F_5 , F_{10} , F_{15} , F_{20} and F_{25} , respectively for

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fresh fruits, which increased to 14.71, 14.33, 14.17, 14.26, 14.40, 14.60 °Brix respectively after 25 days of storage. The conversion of starches and pectins into simple sugar during ripening leads to increase in total soluble solids of the fruits. The results of the current study were in line with (Chambroy *et al.*, 2013; Nasrin *et al.*, 2008; Rajkumar and Mitali, 2009). Juan *et al.* (1999) who stated that more starch and pectin content is present at early mature stage as compared to mid and late mature stage due to which total soluble solid increases with delaying harvesting.

Titratable acidity (%)

Titratable acidity of apricot fruits is significantly influenced by perforated plastic, harvesting stages and storage duration (Table 2). Freshly harvested fruits at various stages such as M_1 , M_2 and M_3 were recorded with 0.94%, 0.84% and 0.88% respectively, which finally decreased to 0.41%, 0.40% and 0.36%, respectively after 25 days. However, fruits stored in plastic bags having 10 holes (F_{10}) . retained the titratable acidity (0.42%) after 25 days in cold storage conditions However, fruit in closed plastic bags (F_0) were observed with minimum (0.37%) titratable acidity after 25 days of storage. In the current study, acidity was decreased in closed plastic bags that might be because of anaerobic conditions Tariq *et al.* (2001) also stated that acidity of citrus fruits decreased in both conditions i.e., 0% oxygen:10% CO₂ and 21% oxygen: 0% CO₂. They also stated that, it is possible that anaerobic condition which is created by lower level of oxygen might be responsible for decreasing fruit acidity, during which the organic acid might be used as a reserve source for energy production. On the other hand, high rate of respiration at higher level of oxygen might decrease acidity.

Further, Beaudry *et al.* (1992) stated that during respiratory metabolism organic acid are consumed. Package perforations generated modified atmosphere, which helps to retain high level of organic acid and thus delaying ripening process. However, Tariq *et al.* (2001) reported that loss of organic acid may induce due to high rate of respiration in sealed packages due to low oxygen. It was reported that acidity of the fruit decline during ripening and storage. Findings are also in line with Rab *et al.* (2016), who recorded highest titratable acidity after 25 days of cold storage in plum fruits harvested at pale green stage and packed in plastic bags having 15 perforations. Khan *et al.* (2007) and Pretel *et al.* (2000) also reported less titratable



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acidity in apricot and persimmon with storage durations.

Total soluble solids: Acid ratio

TSS and Acid dependent on each other and TSS/ acid ratio contributes toward giving many fruits their characteristic flavor. At the start of ripening process the TSS/acid ratio remain low, because of low sugar content and high acidity, which makes the fruit taste sour, during ripening process the sugar content increases and fruit acid are degraded, thus TSS/acid ratio achieves a high value. Previously, it has been observed that both TSS and acidity of the fruits were considerably varied. Similarly, the TSS: Acid was also found responsive to harvesting stages, plastic bags perforation and storage duration (Table 2). Initially, TSS: Acid for the fruits harvested stages remained at lower levels such as 10.95, 14.73, and 15.47 for M_1 , M₂, and M₂ that increased to 31.71, 35.90, and 49.37, respectively after 25 days of cold storage. Similarly, TSS: Acid was increased from $F_0(13.76)$, $F_5(13.70)$, F_{10} (13.64), F_{15} (13.64), F_{20} (13.79) and F_{25} (13.75) of freshly harvested fruits to F_0 (42.02), F_5 (37.74), F_{10} (36.22), F_{15} (37.26), F_{20} (39.90) and F_{25} (40.80), respectively after 25 days of storage. These results are in agreement with (Tariq et al., 2001). The increasing of total soluble solid and decreasing of acidity of fruits with storage and delaying harvesting is responsible for the increases of total soluble solids: Acid. The results of the present study were also in agreement with the earlier conclusion of (Juan et al., 1999; Dhillon and Cheema, 1991) in apple.

Conclusions and Recommendations

This study concluded that harvesting of apricot fruits at greenish stage (10.31 °Brix) attained minimum physiological weight loss and decay index as compared to other harvesting stages after 25 days of storage. Similarly, fruits packed in plastic bags contained 10 pores retained fruit weight and had maximum titratable acidity, fruit firmness and ascorbic acid with minimum physiological weight loss and low decay index. Similarly, the highest value of titratable acidity, fruit firmness and ascorbic acid and lower content of TSS and TSS: Acid was observed in green stage harvested fruit during 25 days of storage. In case of interaction of packaging materials and storage durations, fruits packed in plastic bags having 10 perforation showed increase in TSS and TSS: Acid value with prolonging storage duration up to 25 days. Therefore, it is concluded that apricot fruits should be harvested at green stage (10.31 °Brix) and should be packed in plastic bags having 10 perforations.

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Novelty Statement

To enhance the storage life of apricot, fruits should be harvested at greenish stage (10.31 0 Brix) and should be packed in plastic bag having 10 perforations of 2mm each equally distributed.

Author's Contribution

Rashid Khan: Conducted the research.

Muhammad Sajid: Major supervisor, provided guidelines and idea of the research.

Saud Khan: Prepared the manuscript.

Jehanzeb: Made the layout of research and determined the factors.

Aslam Noor: Finalized the manuscript.

Conflict of interest

The authors have declared no conflict of interest.

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