THE EFFECTS OF DIFFERENT POSTHARVEST APPLICATIONS ON SOME PHYSICOCHEMICAL PROPERTIES IN ‘RUBYGEM’ AND ‘SABRINA’ STRAWBERRY (FRAGARIA X ANANASSA DUCH.) CULTIVARS

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Abstract. This study aimed to determine the efficacy of two standards strawberry (Fragaria x ananassa Duch.) cultivars ‘Rubygem’ and ‘Sabrina’ grown under protected cultivation system in Mersin province in Turkey. The fruit quality preservation during storage in Modified atmosphere packaging (MAP) at different storage temperatures as well as preservative compatibility under Ultraviolet-C (UV-C), hot water and combined applications were studied. Strawberry fruits kept in cold temperatures (0 °C and 5 °C) for 20 days and biochemical changes (total phenolic content, total antioxidant capacity, vitamin C content and sugar content) were measured after harvest and storage. The ‘cv. Rubygem’ kept its quality for 16 days in all treatments and for 20 days in UV-C and UV-C + hot water applications when kept at 0 °C. Also, the fruits stored at 5 °C kept their quality for 12 days and it was observed that hot water and UV-C + hot water treatments got better results in the same period and at the same storage temperature. In the ‘cv. Sabrina’ results showed that fruit samples at 0 °C were successfully maintained in control as well as the other 3 treatments until the 12th day. The preservation was continued in UV-C, hot water and combined treatments on the 16th day of the storage and the best ones were the hot water and UV-C + hot water at the end of the 20th day. As a result, it was observed that total phenolic contents, total antioxidant activity and sugar were found decrease at the end of storage for both cultivars while fluctuations in vitamin C.

Keywords: biochemical changes, hot water, UV-C, sugar, vitamin C, total phenolic, antioxidant

Introduction

Strawberry (Fragaria x ananassa Duch.) fruit is consumed abundantly by consumers and has many positive effects (Çağlırmak, 2006) such as strengthening defense mechanisms and protecting against cancer and infections, protection against DNA damage in the organism (Vinson et al., 2001), protection against cardiovascular disease and lipid peroxidation, because it is rich in antioxidants (Wang et al., 1996; Araque et al., 2018) including anthocyanins, flavanoids and phenolic materials, as well as its unique color, flavor (Gao et al., 2016), aroma, texture, high-order elagic acid (Aybak, 2005).

It is very important to protect the post-harvest quality of strawberries, because they are located in a group of fruit berries that break down quickly and easily after harvest. The main soluble sugars found in the structure of strawberry berry are glucose, fructose and sucrose, and glucose and fructose are known to dominate sucrose in total sugar content (Cordenunsi et al., 2003).

Strawberry fruit contains as high as 92% water. For this reason, water loss is one of the important issues after harvest. In addition to water loss, a number of insecticides and fungicides are used both on the field and before harvesting to prevent the losses
caused by diseases and pests. Applications that are environmentally friendly to prevent postharvest decay include applications such as hypobaric applications (Romanzzi et al., 2001), modified atmosphere packaging (Silva et al., 1999), Ultraviolet-C (UV-C) and hot water applications (Keskin et al., 2014, 2015).

For the last ten years UV-C (180-280 nm $\lambda = 254$ nm) in horticulture products has been considered as an alternative to fungicides to control post-harvest diseases. Moreover, many investigators have reported that UV-C protects fruits and vegetables against decay after harvesting (Marquenie et al., 2003; Allende and Artes, 2003; Allende et al., 2006; Keskin et al., 2014, 2015). It has also been used to delay the ripening of fruits and vegetables by extending their shelf life (Darvishi et al., 2012). When UV-C is applied in appropriate dose and duration, it increases the accumulation of phytoalexins (Keskin and Kunter, 2017), which is effective on the mechanism against disease and damage plants (Keskin et al., 2014, 2015). However, there is still a lot of work to be done to optimize UV-C applications.

In recent years, hot water applications have been implemented to increase the shelf life of fruits and vegetables and to reduce the use of chemical substances in the fight against diseases and pests (Lurie, 1998). Hot water applications are made by immersion (Barkai-Golan and Phillips, 1991) or by special hot water spraying (Fallik et al., 1996).

Due to short life time of post harvested strawberry fruit, the effect of modified atmosphere packaging (MAP) with different treatments have been widely considered.

In this study, effects of different postharvest treatments including UV-C and hot water were studied on biochemical changes of two commercially strawberry varieties stored at different temperatures in in MAP (modified atmosphere packaging) condition.

Materials and methods

Cultivars used in research

In the study, ‘Sabrina’ and ‘Rubygem’ strawberry cultivars, which are grown under greenhouse by a private company named Roseland and harvested in similar maturity in Tarsus district in Mersin province in Turkey, were used as research materials.

Strawberry fruit was first pre-cooled at 0 °C for 1 day. Later, the fruits of the same maturity were divided into 4 different groups. The first group of fruits was separated as controls. The second group of fruits was irradiated with 254 nm UV-C (0.25 kJ m$^{-2}$) with Vilber Lourmat UV-C lamp on both surfaces as 5 min at 20 cm distance. The third group of fruits was kept at 60 °C for 10 seconds in a hot water bath. In the fourth group, both UV-C and hot water applications were applied. Then fruit samples were placed in 250 gr plastic bags.

Fruits were maintained at 0 °C and 5 °C in Van Yüzüncü Yıl University Faculty of Agriculture Department of Horticulture’s cold-air storages with a relative humidity of 90-95% for 20 days.

Total phenolic substance amount and total antioxidant activity

Total phenolic contents in ‘Rubygem’ and ‘Sabrina’ strawberry cultivars were determined by spectrophotometer (Varian Bio 100, Australia) by Folin-Ciocalteau calorimetric method (Swain and Hillis, 1959). Absorbance values of the solutions
were read in spectrophotometer (Thermo Fisher Scientific, G10S UV-Vis) at 725 nm wavelength and the total amount of phenolic substance expressed as gallic acid equivalent (GAE) mg kg\(^{-1}\) fresh weight (FW).

Ferric Reducing Antioxidant Power (FRAP) (Iron (III) Reduction Antioxidant Power) method was used to determine antioxidant activity (Benzie and Strain, 1996). 300 \(\mu\) L freshly prepared FRAP reagent [25 mL acetate buffer (300 mmol/L, pH 3.6), 2.5 mL TPTZ (10 mmol/L) in 40 mmol/L HCl, and 2.5 mL FeCl\(_3\) \(\cdot\) 6H\(_2\)O solution (20 mmol/L) were mixed as required to prepare working FRAP reagent] was warmed to 37 °C, and a reagent blank reading was taken at 593 nm; 10 \(\mu\) L of each fraction extract solution of an appropriate concentration was then added along with 30 \(\mu\) L H\(_2\)O. The 0–4 min absorbance change (\(\Delta A\)\(_{593\,\text{nm}}\)) was calculated for each sample and related to \(\Delta A\)\(_{593\,\text{nm}}\) of a Fe\(^2\+) standard solution tested in parallel to obtain the FRAP value of each sample. Antioxidant activity values were reported as \(\mu\)mol trolox equivalent (TE) mg\(^{-1}\).

**Vitamin C (ascorbic acid)**

The method specified by Cemeroğlu (2007) was applied to determine the content of vitamin C. 3 g of the strawberry fruit were weighed and homogenized in 6 ml of 6% metaphosphoric acid, then centrifuged at 12000 rpm for 15 min at 4 °C. In the HPLC (Agilent 1100 series) analyzes, C vitamins C18 (Phenomenex Luna C18, 250 x 4.60 mm, 5 \(\mu\)) were performed. Readings was carried out in the DAD (DE33225146) detector at a wavelength of 254 nm. L-ascorbic acid (Sigma A5960), prepared at different concentrations, was used to identify vitamin C pick and to determine the amount of vitamin C (Karadogan and Keskin, 2017).

**Sugar content**

After the fruit samples of ‘cv. Rubygem’ and cv. ‘Sabrina’ were homogenized, a 5 g sample from each replicate was added in 10 ml of 6% metaphosphoric acid and then centrifuged at 15.000 rpm for 15 min at 4 °C. Sugar determinations were made on HPLC (Agilent 1100 series) after filtration through filters with a hole diameter of 0.45 mm in the supernatant sample. For this purpose, Phenomenex Rezex RCM monosaccharide column, 80% acetonitrile and refractive index detector were used as carrier phase. The definitions for sucrose, glucose, fructose and maltose were made using external standards, taking into consideration the structural changes of ‘Rubygem’ and ‘Sabrina’ (Özgökçe, 2016).

**Statistical analysis**

Experiments were conducted with three replication. The descriptive statistics for the traits studied are expressed as mean and standard error. In terms of these characteristics, factorial (Four Factors) Variance Analysis was performed to determine whether there was a difference between storage period, applications, temperature and cultivars. Following the variance analysis, Duncan test was used to identify the different groups. The statistical significance level was taken as p < 0.05 in the calculations and the SPSS statistical package program was used for the calculations.
Results

Total phenolic content

During storage, it was observed that total phenolic contents of the ‘cv. Rubygem’ and ‘cv. Sabrina’ strawberry cultivars decreased at both storage temperatures. In the ‘cv. Rubygem’ strawberry cultivar; the highest total phenolic content at 0 °C was found to be 498.486 mg kg⁻¹ with UV-C on day 16 and the lowest value was found in the control group as 247.946 mg kg⁻¹ on day 20. At 5 °C, the highest total phenolic content was determined in hot water application on day 8 as 496.054 mg kg⁻¹, while the lowest value was found in the control group with 206.189 mg kg⁻¹ on the 12th day.

The highest total phenolic content in ‘cv. Sabrina’ was found to be in hot water application with 515.919 mg kg⁻¹ on 8th day and the lowest value was found in UV-C application on the 20th day with 117.405 mg kg⁻¹. At 5 °C, it has been identified that the highest value application was 484.297 mg kg⁻¹ with UV-C + hot water application on day 0 and the lowest total phenolic compound value was with UV-C application on day 8 with 385.784 mg kg⁻¹ (Fig. 1).

Figure 1. Changes in total phenolic content as a result of storage of ‘cv. Rubygem’ and ‘cv. Sabrina’ strawberry cultivars at 0 and 5 °C. (a, b, c, d: The difference between the applications of the same kind, same storage period and different lower case letters at the same storage temperature is significant (p < 0.05) [Comparison of applications]. A, B, C, D: The difference between storage times of the same kind, same application and different capital letters at the same storage temperature is significant (p < 0.05) [Comparison of storage periods])

Total antioxidant activity

Total antioxidant activity was found to decrease at the end of storage at both types and at the storage temperature. Therefore, in the ‘cv. Rubygem’, the highest value in total antioxidant activity at 0 °C was found to be 73.650 mg 100 g⁻¹ in the hot water application and the lowest value in the control group with 9.600 mg 100 g⁻¹ in the 20th day. The highest total antioxidant capacity at 5 °C was found to be 66.300 mg 100 g⁻¹ in hot water at 0th day, and the lowest total antioxidant value in 12th day at 3.250 mg 100 g⁻¹ in UV-C + hot water application. The ‘cv. Sabrina’ has the highest antioxidant value at 0 °C, Hot water application with 47.250 mg 100 g⁻¹ on the 8th day and hot water application on the 20th day with the lowest value of 6.600 mg 100 g⁻¹. At 5 °C, the
highest antioxidant amount was 40.100 mg 100 g$^{-1}$ UV-C + hot water application on day 0, while the lowest was 23.050 mg 100 g$^{-1}$ UV-C on day 8 (Fig. 2).

**Vitamin C (ascorbic acid)**

During storage, fluctuations in vitamin C value are observed. In the ‘cv. Rubygem’, the highest vitamin C concentration at 0 °C was on 0th day with 343.723 mg 100 g$^{-1}$ and the lowest vitamin C concentration was 16th day with 78.041 mg 100 g$^{-1}$ in UV-C + hot water application. According to analysis results, at 5 °C, the highest value of vitamin C was detected in the application of UV-C + hot water with 343.723 mg 100 g$^{-1}$ on day 0, while the lowest value of vitamin C was in the control group with 32.492 mg 100 g$^{-1}$ on day 8. When the ‘cv. Sabrina’ was examined, it was determined that the highest value at 0 °C was in hot water application with 382.854 mg 100 g$^{-1}$ on day 0 (Fig. 3).
382.854 mg 100 g\(^{-1}\) on day 0. The lowest value was determined in UV-C application on day 8 with 22.906 mg 100 g\(^{-1}\).

**Sugar content**

At the end of storage of the ‘cv. Rubygem’ at 0 °C, the application with the highest fructose value was found in the control group on day 0 with 30.404 mg g\(^{-1}\), while the lowest value was found in the control group on day 20 with 4.180 mg g\(^{-1}\). At 5 °C, the application with the highest fructose content was 30.404 mg g\(^{-1}\), while the lowest value application was with hot water at 3.876 mg g\(^{-1}\) on the 12\(^{th}\) day. When the variations in the ‘cv. Sabrina’ at 0 °C were examined, it was found that the highest fructose value was 22.793 mg g\(^{-1}\) on 8\(^{th}\) day in UV-C application and the lowest value was 1.539 mg g\(^{-1}\) on 20\(^{th}\) day in UV-C application were determined according to the analysis results. At 5 °C, the highest fructose value was found to be 22.556 mg g\(^{-1}\) with UV-C application on day 0 and the lowest value was found with UV-C application on day 8 with 11.372 mg g\(^{-1}\) (Fig. 4).

![Graphs showing fructose values for 'Rubygem' and 'Sabrina' strawberry cultivars at 0 and 5 °C](image)

**Figure 4. Changes in fructose values resulting from storage of ‘Rubygem’ and ‘Sabrina’ strawberry cultivars at 0 and 5 °C**

Glucose has decreased regularly at both cultivars and storage temperatures. In this context, it has been determined that the lowest values in both cultivars and storage temperatures are in hot water application. As a result of storage of the ‘cv. Rubygem’ at 0 °C, the highest glucose value was found to be in the control group with 26.539 mg g\(^{-1}\) on day 0, while the lowest value was found to be in hot water application with 0.577 mg g\(^{-1}\) on the 20\(^{th}\) day. The highest glucose value for 5 °C was the control group with 26.539 mg g\(^{-1}\) on day 0, while the lowest glucose value was found to be 1.086 mg g\(^{-1}\) on the 12\(^{th}\) day with hot water application. In the ‘cv. Sabrina’, it was determined that the maximum value at 0 °C was 16.945 mg g\(^{-1}\) in UV-C application on 8\(^{th}\) day, while the lowest value was 0.762 mg g\(^{-1}\) in hot water on 20\(^{th}\) day. The highest glucose value at 5 °C was found to be 16.284 mg g\(^{-1}\) in UV-C application on day 0, while the lowest value was found in hot water application on day 8 with 5.676 mg g\(^{-1}\) (Fig. 5).
When the changes in the sucrose value were examined, it was determined that there were fluctuations in both strawberry cultivars and storage temperatures. In the ‘cv. Rubygem’, it was determined that the highest value at 0 °C was 0.304 mg g⁻¹, the UV-C + hot water application on 16th day, and the lowest value was UV-C application on day 0 with 0.027 mg g⁻¹. At 5 °C, the highest sucrose level was detected in fruit samples treated with 0.300 mg g⁻¹ of UV-C + hot water on day 8, while the lowest value was found to be in UV-C application on day 0 with 0.027 mg g⁻¹. According to the results of the analysis, in the ‘cv. Sabrina’, the highest sucrose value at 0 °C was found in the control group at 0.466 mg g⁻¹ in the hot water application on 0th day and the lowest value was in the control group on the 16th day with 0.033 mg g⁻¹. The highest sucrose value at 5 °C was determined to be 0.466 mg g⁻¹ in hot water application on day 0, while the lowest value was found to be in the control group with 0.094 mg g⁻¹ on the 8th day (Fig. 6).

Figure 5. Changes in glucose values resulting from storage of ‘Rubygem’ and ‘Sabrina’ strawberry cultivars at 0 and 5 °C

Figure 6. Changes in sucrose value resulting from storage of ‘Rubygem’ and ‘Sabrina’ strawberry cultivars at 0 and 5 °C
Looking at the changes in maltose value, it was determined that the variations for the ‘cv. Rubygem’ were parallel to each other, while the ‘cv. Sabrina’ was found to be fluctuated. In this case, it was determined that the lowest value of the ‘cv. Rubygem’ was at 0 °C, the highest maltose value was in the control group with 29.497 mg g⁻¹ on day 0. The lowest value was found in the control group with 4.20 mg g⁻¹ on the 16th day. The highest maltose concentration at 5 °C was determined in the control group with 29.497 mg g⁻¹ on 0th day, whereas the lowest value application was found to be the control group with 1.178 mg g⁻¹ on 12th day. In the ‘cv. Sabrina’, the highest maltose value at 0 °C was found to be 20.094 mg g⁻¹ in UV-C application on day 0, while the lowest value was found to be 2.592 mg g⁻¹ in UV-C + hot water application on day 8. At 5 °C, the application of the highest maltose value was found to be 20.095 mg g⁻¹ on day 0 with UV-C application, while the lowest value was found to be in the control group with 3.285 mg g⁻¹ on day 8 (Fig. 7).

**Figure 7. Changes in the maltose value resulting from storage of ’Rubygem’ and ’Sabrina’ strawberry cultivars at 0 and 5 °C**

**Discussion**

Statistical analysis among different treatments on total phenolic contents in ‘cv. Rubygem’ and ‘cv. Sabrina’ cultivars showed that there was no significant difference in ‘cv. Rubygem’ stored at 0 °C in 0, 8 and 12 days. On the other hand, the difference between UV-C and hot water applications on the 16th day and also difference between control and UV-C + hot water applications on the 20th day was found statistically significant. The difference between the applications of hot water, UV-C + hot water, and control application on the 8th day was found to be significant, whereas the applications on the 0th and 12th day were not statistically significant when considering the differences between applications at the other storage temperature of 5 °C (Fig. 1). In the ‘cv. Sabrina’; the differences between the applications on days 8, 12 and 20 were not significant. The difference between UV-C + hot water application on day 0 and UV-C application and hot water application on day 16 were significant. Differences between applications at the ‘cv. Sabrina’ at 5 °C, another storage temperature; UV-C + hot water application was statistically significant at day 0, whereas it was not significant at day 8 applications (Fig. 1). In both types and temperatures there was a decrease in the total 

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amount of phenol in almost all applications. These findings are similar to those of Pan et al. (2004). Results of this study showed that the highest amounts of phenolic substances in ‘cv. Rubygem’ could be observed at 0 °C storage temperature by using hot water and at 5 °C by using UV-C and hot water treatments. In this study that we have conducted, it has been determined that the application that protects the phenolic materials, which are so important in terms of human health, is hot water application at 0 °C, and UV-C and hot water applications at 5 °C in the ‘cv. Rubygem’. In the ‘cv. Sabrina’, the result is that the hot water application is the best at both storage temperatures. Previous studies have shown that the total amount of phenolics in strawberries depends on the storage temperature and the composition of the atmosphere (Wang and Zheng, 2001; Ayala-Zavala et al., 2004; Cordenunsi et al., 2005). This high phenolic material in strawberries has increased the shelf life of fruit.

Differences in application with respect to total antioxidant capacity: the difference between the ‘cv. Rubygem’ and the ‘cv. Sabrina’ stored at 0 °C was statistically significant. The difference between the applications in the ‘cv. Rubygem’ type stored at 5 °C is significant, while the ‘cv. Sabrina’ is not significant (Fig. 2). Antioxidant substances naturally occur in plants. These plant antioxidants include anthocyanins and phenolic compounds. It was determined that the amount of antioxidant decreased regularly in both cultivars and 4 different applications. When the applications are evaluated on the basis of the amount of antioxidant in the ‘cv. Rubygem’, UV-C and hot water application is the application that best protects the amount of antioxidant compound at both storage temperatures. In the ‘cv. Sabrina’, it is determined that UV-C and hot water applications are the best application for strawberry fruit stored at 0 °C and control and UV-C + hot water applications for the fruits stored at 5 °C. The successful results have been obtained in terms of protection of antioxidants, especially in applications that we have made in the ‘cv. Rubygem’. The study we conducted is similar to that of Vicente et al. (2004) with a study of grape fruits, indicating that the amount of antioxidants decreased both at the end of storage and that hot water and UV-C treatments were effective.

The comparison of different treatments on amounts of vitamin C showed that in ‘cv. Rubygem’ at 0 °C, the difference between UV-C + hot water application with UV-C and hot water applications on day 0 was statistically significant. Also, there were no significant differences among storage days (8, 12, 16 and 20 days). At 5 °C, the difference between applications on day 0 and 12th day of storage were statistically significant, whereas on 8th day was not meaningful (Fig. 3). Difference between applications in the ‘cv. Sabrina’: at 0 °C, the difference between control and hot water applications of UV-C application was statistically significant at day 0, whereas the difference between applications at 8, 12, 16 and 20 days was not significant. At 5 °C, the difference between UV-C and hot water was significant at day 0 and hot water at day 8 and other applications (Fig. 3). Vitamin C values during storage: In the ‘cv. Sabrina’, except for UV-C applications, there was a decrease in vitamin C value at both temperatures and applications. Both types and at both storage temperatures (except for the ‘cv. Sabrina’ stored at 5 °C) have been the control group that best protects Vitamin C. With this study, both UV-C applications and hot water applications are thought to reduce the amount of vitamin C.

When the changes in sugar content were examined, the difference between the applications of fructose in both types of storage (except the ‘cv. Sabrina’ at 5 °C) was found to be significant (Fig. 4). The difference between the applications at the end of
the storage of the ‘cv. Rubygem’ strawberry in terms of glucose was found statistically significant. In the ‘cv. Sabrina’, the difference between applications was not statistically significant at 0 °C. At 5 °C, the difference between treatments was found to be insignificant on day 0, whereas on day 8, control was statistically significant (Fig. 5). When the difference between the applications for sucrose value was examined, the difference between UV-C + hot water application and control and UV-C applications on day 0 in the strawberry cultivars stored at 0 °C was statistically significant, while the difference between the practices on the day was not statistically significant on days 8, 12, 16 and 20. When the difference between the applications is examined in the ‘cv. Rubygem’, the difference between UV-C + hot water application and control and UV-C applications was statistically significant on day 0 at 5 °C, whereas the difference between the applications on days 8 and 12 found to be insignificant (Fig. 6). The difference between the application of UV-C and UV-C + hot water at 0 °C on the 0th day was found to be statistically significant when looking at the difference between applications for the changes in the value of sucrose in the ‘cv. Sabrina’. On the 8th day, the difference between control and UV-C applications was not statistically significant, while UV-C + hot water application was statistically significant with hot water, control and UV-C applications. The difference between the applications of storage on the 12th, 16th and 20th day was not statistically significant. When the difference between the applications was observed at 5 °C storage temperature of the ‘cv. Sabrina’ fruits, the difference between the applications of hot water and UV-C + hot water on the 0th day was significant, but the difference between the applications on the 12th day was not statistically significant (Fig. 6). At 0 °C, maltose content of the ‘cv. Rubygem’ was statistically significant at day 0, whereas applications on 8, 12, 16 and 20 days were not statistically significant. At 5 °C, the control application on 0th day and the difference between control and UV-C applications on 12th day was found to be statistically significant in the ‘cv. Rubygem’ (Fig. 7). The difference between the applications in the ‘cv. Sabrina’: While the difference between hot water and UV-C + hot water applications at 0 °C on the 8th day was not statistically significant, the difference between control and UV-C applications of these applications was found statistically significant. The difference between the applications on 0, 12, 16 and 20 days was not statistically significant. At 5 °C, the difference between control and hot water applications with UV-C + hot water application on the 8th day of the ‘cv. Sabrina’ was found to be statistically significant (Fig. 7). When fructose, glucose, sucrose and maltose sugar compositions were examined in strawberry fruit, these values decreased in almost all applications at the end of storage. This is why the fruits need energy to perform various reactions inside the cell so that they can survive even when they are picked up. This energy is obtained by using sugars, organic acids and oils present in the fruit. In other words, sugar is used during respiration.

Conclusion

In this study, the ‘cv. Rubygem’ and the ‘cv. Sabrina’ strawberry cultivars were placed in the plastic bags with control, UV-C, hot water and UV-C + hot water applications and stored at 0 and 5 °C, the fruits of the ‘cv. Rubygem’ were successfully preserved in 4 groups of applications until the 16th day and at the end of 20th day UV-C, hot water and UV-C + hot water applied fruit samples were successfully preserved. Changes in the same fruit samples at 5 °C were observed for 12 days and during this
period it was determined that hot water and UV-C + hot water treatments could last 12 days at this storage temperature. When the changes occurred at the end of preservation of the ‘cv. Sabrina’ are examined, it is thought that fruit samples at 0 °C can be successfully preserved in all applications up to 12th day. It has been determined that UV-C application in this cultivar is preserved for 16 days, hot water and UV-C + hot water applications are preserved up to 20 days. It was determined that the ‘cv. Sabrina’ fruit samples kept at 5 °C lasted for only 8 days. As a result, it was observed that total phenolic contents, total antioxidant activity and sugar were found decrease at the end of storage for both cultivars while fluctuations in vitamin C.

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Tekin – Çavuşoğlu: Effects of hot water and UV-C applications in some strawberry cultivars stored in modified atmosphere condition


