



Role of Controlled Atmosphere, Ultra Low Oxygen or Dynamic Controlled Atmosphere Conditions on Quality Characteristics of ‘Scarlet Spur’ Apple Fruit

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ABSTRACT

In this study, the effects of three cold storage technologies, (i) controlled atmosphere-CA (CO₂ 4%, O₂ 3%), (ii) ultra low oxygen-ULO and (iii) dynamic controlled atmosphere-DCA, were investigated on fruit quality of ‘Scarlet Spur’ apples stored during 10 months plus 7 days of shelf life at 20 °C. After harvest, apples were stored at 0 °C and 90±5% relative humidity during 10 months in CA, ULO (CO₂ 3%, O₂ 1%) and DCA (CO₂ 1%, O₂ 0.5%) conditions. HarvestWatch™ sensors were used for assessment of lower oxygen limit (LOL) of fruit during DCA storage.

DCA was the best storage condition suppressing ethylene synthesis and respiration rate during storage. The ULO and DCA conditions showed similar results in the maintenance of firmness and TA amount. Weight loss in these conditions was also lower than CA. No significant difference was observed between storage conditions in terms of SSC. DCA technology gave better results in maintaining color of ‘Scarlet Spur’ than other conditions during cold storage. Result showed that; ULO and DCA conditions were more effective in maintaining quality compared to CA in terms of all quality parameters.

Keywords: Apple, Controlled atmosphere, Postharvest, Cold storage, Chlorophyll fluorescence

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1. Introduction

Controlled atmosphere (CA) storage is a widely used technology for the storage of apple which is one of the most produced and consumed fruit in the world. Reducing the oxygen (O₂) concentration in the storage atmosphere and increasing the concentration of carbon dioxide (CO₂) is the most important factors for prolonging storage period in CA storage technology. Thus fruit quality is kept for longer periods, and postharvest losses during storage are reduced (Both et al. 2014). The possibility to obtain the longest storage period in CA conditions depends on (i) fruit maturity at harvest time, (ii) atmosphere composition during storage and (iii) the cultivar (Thompson 2010).

Storage under proper conditions plays an important role in maintaining product quality especially in climacteric fruit such as apples (Bertone et al. 2012). The main purpose of optimizing the CA storage conditions is to prevent ripening and aging of the fruit by decreasing O₂ level and reducing respiration rate and ethylene production (Veltman et al. 2003). Suppressing respiration rate and ethylene synthesis of fruit are the key postharvest processes throughout cold storage (Wright et al. 2015). O₂ is the most important factor to decrease metabolic activity and reduce biochemical changes after harvest. Therefore, the use of low O₂ levels in storage is an important potential (Tuna Gunes & Horzum 2017).

The equipment developed in CA storage technology has allowed to work at low O₂ or ultra low oxygen (ULO) conditions (Batu & Sen 2014). Standard CA storage involves keeping the oxygen content at 2-3% while the O₂ level in the ULO conditions can be reduced to 1%. ULO storage is more successful than standard CA technology in terms of preventing disease and physiological disorders (Balla & Holb 2007; Mattè et al. 2005). Additionally, it can protect some quality characteristics such as fruit flesh firmness and ground color better than standard CA storage procedure (Thewes et al. 2015). In developed countries, ULO storage has been extensively used in fruit industry in order to maintain fruit quality for a longer period (Watkins 2008).

Dynamic controlled atmosphere (DCA) is the new and popular technology in apple industry (Mditshwa et al. 2018). During DCA storage, O₂ level is reduced to the lowest level that the fruit can tolerate which is just above the so-called critical O₂ concentration (LOL). Quality losses related to anaerobic condition increase when fruit are stored under LOL. Ideally, fruit should be stored at levels just above the critical O₂ concentration (Gasser et al. 2008). It has been reported that fruit kept in these conditions could be stored for a long time without significant losses (Prange et al. 2007; Zanella et al. 2008; Wright et al.

2012). Researches in some apple varieties showed that DCA is more effective than CA for maintaining quality during storage (Veltman et al. 2003; Gasser et al. 2005; Delong et al. 2007; Bessemans et al. 2016; Thewes et al. 2017).

DCA technology involves monitoring of gas concentrations in the storage room via sensors. Up to now, three sensors has been developed in this technology; chlorophyll fluorescence (CF), respiration quotient (RQ) and ethanol (ET) (Thewes et al. 2018). While very little research has been done with RQ and ET sensors, CF is the most common used sensor in the pome fruit industry (Mditshwa et al. 2018).

CF technique measures the stress occurring in fruit during storage period. In this method, while the O₂ level is reduced, the CF signal on the fruit surface is measured by the sensor (Vanoli et al. 2010). Detection technology senses the response from the produce and feeds it back to an analytical software tool (HarvestWatch™) where the output is displayed in graph format (Stephens & Tanner 2005). The increase in the fluorescence signal indicates that the product enters low O₂ stress (Watkins 2008). The O₂ level is maintained over the LOL level by adapting according to fruit metabolism during storage.

In researches on effect of CA, ULO and DCA storage conditions for maintaining significant quality criteria in apples, results has changed based on cultivars (Aubert et al. 2015; Both et al. 2017; Kitemann et al. 2015; Thewes et al. 2015; Tran et al. 2015; Brizzolara et al. 2017). Therefore, in this study, the effects of CA, ULO and DCA on fruit quality of ‘Scarlet Spur’ apple cultivar was evaluated during a storage period of 10 months plus a shelf life period of 7 days at 20°C.

2. Material and Methods

2.1. Plant material

Experimental fruit were obtained from the commercial apple orchard located in Isparta/Eğirdir (38° 17' North, 30° 55' East), in 2012. The uniform trees were 8 years old cv. ‘Scarlet Spur’ apple on MM106 rootstock. Standard cultural practices were applied to the trees during fruit growth and development period.

2.2. Fruit harvest and storage conditions

Fruit were harvested at commercial harvest stage and transported to the postharvest physiology laboratory. Apples were randomly divided into three groups and stored at 0 °C and 90±5% relative humidity (RH) during 10 months in CA (CO₂ 4%, O₂ 3%), ULO (CO₂ 3%, O₂ 1%), or DCA (0.5% O₂ and 1% CO₂) conditions, respectively. Cabinets manufactured with gas tight plastic material and each was 0.5 m³ volume. HarvestWatch™ was used to assess lower O₂ limit (LOL) of fruit during DCA storage. LOL stress in fruit under DCA was assessed by CF sensors placed over a sample of 6 fruit each batch. LOL level in DCA was determined as 0.2%. The samples were stored at 0.5% O₂ level by adding 0.3% safety margin to the determined LOL level under DCA conditions. After cold storage, apples were kept at 20 °C and 60±5 % RH for 7 days to determine the effects of treatments on some quality parameters investigated in this research during shelf life.

2.3. Respiration rate and ethylene production

Fruit (1 kg) were kept in 5 L airtight jars at room condition (20 °C) for determination of ethylene emission and respiration rate. After 3 h, the gas sample was taken from the closed jars by a gastight syringe and injected into loop of gas chromatography (GC) (Agilent 6840). Ethylene emission and respiration rate were measured by GC equipped with flame ionization (FID) and thermal conductivity detectors (TCD), respectively. Measurements were made in split/splitless (S/SL) of inlet in split mode with gas sampling valve with 1-mL gas sample by using fused silica capillar column (GS-GASPRO, 30 m x 0.32 mm I.D., U.S.A). Results were calculated as µL kg⁻¹ h⁻¹ and ml CO₂ kg⁻¹ h⁻¹ for ethylene production and respiration rate, respectively.

2.4. Fruit flesh firmness

Fruit flesh firmness was measured by using a texture analyzer (Güss FTA Type GS14 Fruit-Texture Analyzer Model, Strand, South Africa). The measurements were performed on both side of apple after skin removal using a stainless probe (11.1 mm). Firmness was measured over 10 fruit in each replication and results were presented in Newton (N).

2.5. Soluble solids content (SSC) and titratable acidity (TA)

The fruit juice from 10 apples in each replication was extracted with the help of a juicer for analysis. The soluble solids content (SSC) of apple juice (%) was determined with a refractometer (Digital-Atago Pocket PAL-1). The titratable acidity (TA) in apple juice was measured by titration of 10 mL of juice with NaOH solution (0.1 mol L⁻¹) to an end-point pH of 8.1 by a pH meter (Hanna pH 330 model, WTW, Germany). The results were expressed as % malic acid.

2.6. Fruit skin color

Fruit skin color of apples was measured with a colorimeter (Minolta CR 400, USA). Color measurements were made on both sides of 10 fruit in each replication along the equatorial axes. The calibration of color measurement apparatus was performed using an original calibration plate (white). The fruit colors were evaluated as CIE L*, a* and b*.

2.7. Weight loss

Weight loss of fruit was measured based on the initial weight and calculated as percent (weight loss % = [(first weight - last weight) / first weight × 100]) during cold storage. In order to measure the weight loss during the shelf life period, weight measurements were made at the beginning and at the end of the shelf life. Weight loss of apples was measured over 10 fruit in each replicate.

2.8. Statistical analysis of results

The completely randomized design (with three replications) was chosen for this experiment. Using software package (JMP7), the general linear model was used for statistical analyses. The differences among means (at a significance level of 0.05) were analyzed using LSD test.

3. Results and Discussion

3.1. Respiration rate and ethylene production

During storage and shelf life, respiration rate increased in all storage conditions (Figure 1). The differences between conditions and periods and their interactions were statistically significant in both cold storage and room conditions ($P < 0.001$, 0.0001). The highest respiration rate during storage was determined in samples in CA (mean $9.87 \text{ mL.CO}_2 \text{ kg}^{-1}\text{h}^{-1}$) while the lowest respiration rate was observed in DCA (mean $7.22 \text{ mL.CO}_2 \text{ kg}^{-1}\text{h}^{-1}$). DCA was the best storage condition to suppress respiration rate. In room conditions, respiration rate values obtained from samples stored in ULO ($11.86 \text{ mL.CO}_2 \text{ kg}^{-1}\text{h}^{-1}$) and DCA ($11.39 \text{ mL.CO}_2 \text{ kg}^{-1}\text{h}^{-1}$) gave similar results. CA conditions were again resulted the highest ($14.93 \text{ mL.CO}_2 \text{ kg}^{-1}\text{h}^{-1}$) respiration rate. It was determined that DCA storage of ‘Granny Smith’ apple suppressed respiratory rate better than CA storage (Eren et al. 2015). Similarly, previous studies have showed that limiting O_2 levels, significantly reduces respiration rate (Gasser et al. 2008; Wright et al. 2012; Thewes et al. 2015). Respiration is the breakdown of complex molecules (starch, sugar and organic acids) to simple molecules (CO_2 and H_2O) in the cell (Kader 2002). In the final stage of the respiratory reaction; as O_2 acts as the ultimate electron acceptor in the mitochondrial electron transport chain, the metabolism of the fruit can be slowed down by lowering the O_2 concentration in the storage (Bekele et al. 2016).

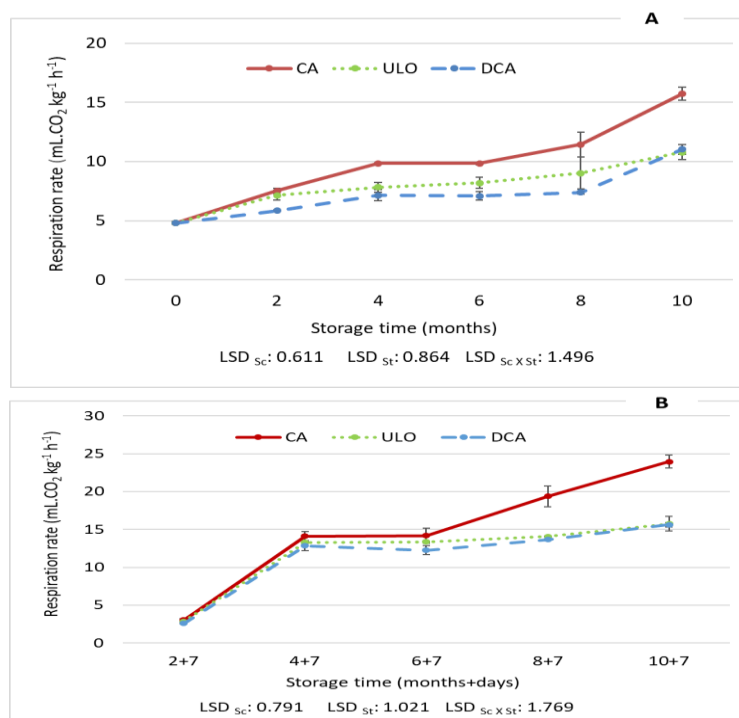


Figure 1- Respiration rate of ‘Scarlet Spur’ apples stored under different atmosphere conditions during 10 months (A) and plus 7 days for shelf life (B). Vertical bars represent standard error (n=3)

The effects of storage conditions and periods on ethylene production were statistically significant. The interaction between time and condition was also significant ($P < 0.05$, 0.0001). During cold storage and shelf life, the amount of ethylene production increased in all three storage conditions (Figure 2). The highest increase was observed in CA storage. In the 8th month of storage, the ethylene production in the CA ($7.46 \mu\text{L.C}_2\text{H}_4 \text{ kg}^{-1} \text{ h}^{-1}$) showed a noteworthy increase compared to other conditions. This rapid increase in shelf life began to be observed since the 4th month ($41.32 \mu\text{L.C}_2\text{H}_4 \text{ kg}^{-1} \text{ h}^{-1}$). The highest average ethylene production during storage was obtained under CA storage. ULO and DCA caused in similar results in terms of ethylene production. For some apple cultivars, higher ethylene production was found in fruit stored in CA conditions compared to fruit stored in ULO and DCA conditions (Mattheis et al. 1998; Hennecke et al. 2008; Çalhan et al. 2012; Thewes et al. 2015). Since ethylene initiates the ripening process in fruit, its production is reduced to the lowest possible level, resulting in higher fruit quality after storage. (Watkins 2006).

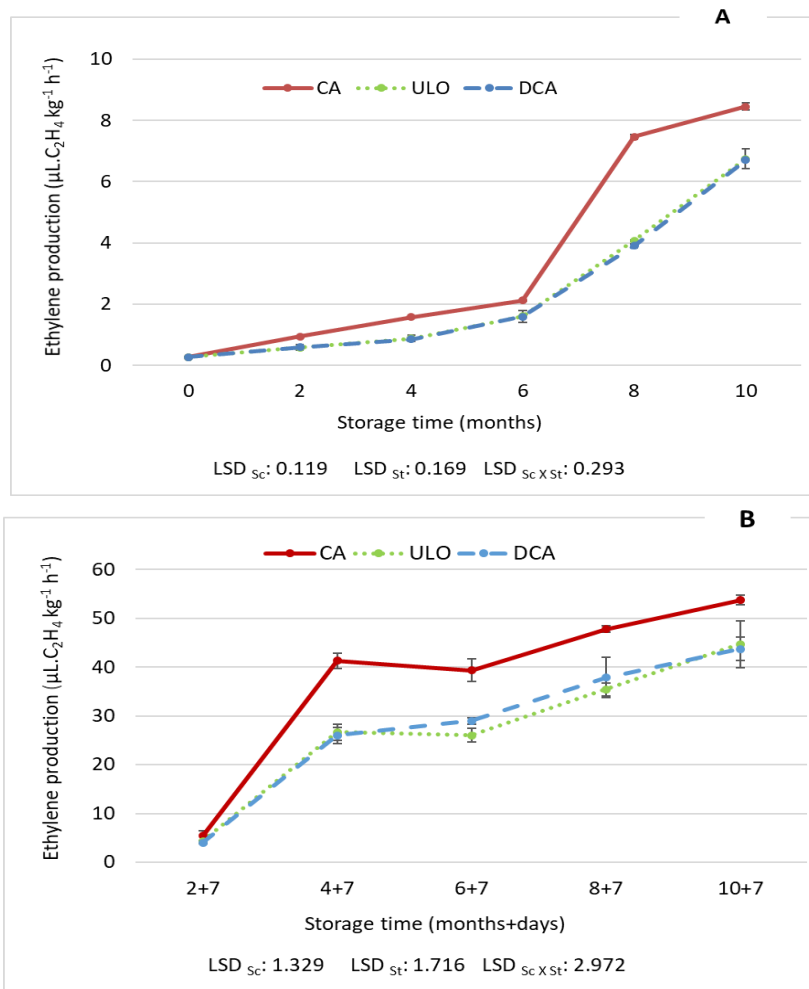


Figure 2- Ethylene production of ‘Scarlet Spur’ apples stored under different atmosphere conditions during 10 months (A) and plus 7 days for shelf life (B). Vertical bars represent standard error (n=3)

Reducing O_2 level in the storage atmosphere decreases the ethylene production of fruit (Gorny & Kader 1996). Storage of apples under DCA conditions significantly reduces ethylene synthesis and maintains long-term quality parameters (Watkins, 2008). DCA storage is effective in suppressing the activity of 1-Aminocyclopropane-1-carboxylate (ACC) oxidase enzyme that plays a key role in ethylene synthesis and production (Thewes et al. 2015; Weber et al. 2015; Thewes et al. 2017). ACC formed during ethylene synthesis pathway is oxidized by ACC oxidase enzyme and convert to ethylene (Nath et al. 2006). Therefore, reduction of O_2 in the storage room inhibits the activity of ACC oxidase enzyme and the conversion of ACC to ethylene.

3.2. Firmness (N), SSC (%) and TA (%)

According to the changes in fruit firmness values, the differences between the storage conditions and the storage period were statistically significant ($P < 0.0001$). With the prolonged storage period, the firmness values of the samples decreased in all conditions. Firmness, which is one of the most important factors affecting apple quality, decreases in relation to water loss during long term cold storage (Mditshwa et al. 2017b). This decrease in the value of firmness during storage has been shown similarly in previous studies (Koyuncu & Bayındır 2013). Gwanpua et al. (2014) reported that the loss of sugar in the

'Jonagold' apples during ripening, increased pectin solubility, and the decrease in the water-soluble pectin molar mass were caused by softening. The highest average fruit flesh firmness values were obtained from the samples stored in DCA and ULO conditions during the storage (68.11 N-67.56 N) and shelf life (54.50 N -52.28 N) period. (Table 1, 2). Fruit flesh firmness was better protected in low O₂ conditions (De Castro et al. 2007). During the maturation, some enzymes cause the polymerization of pectin polymers and loosening the cohesion between the cells (Brummell & Harpster, 2001; Goulao & Oliveira 2008). This loss in cohesion of the pectin network is responsible for softening (Fischer & Bennett 1991). Many enzymes play a role in cell wall modifications during maturation of apples. The activities of these enzymes are related to ethylene production (Gwanpua et al. 2014). Ethylene signals the cell wall degrading enzymes and triggers their activity (Payasi et al. 2009). The lower firmness loss in DCA is related to the low amount of ethylene produced in this condition. It is reported that DCA storage suppresses the enzymes responsible for softening (Mditshwa et al. 2018). Studies on apples have shown that the firmness of fruit flesh in DCA conditions is better protected than CA conditions (Mattheis et al. 1998; Zanella et al. 2005; DeLong et al. 2007; Zanella et al. 2008; Tran et al. 2015; Thewes et al. 2015 Bessemans et al. 2016; Mditshwa et al. 2017a). The ULO conditions also yielded better results than CA storage in maintaining firmness. Similar findings were obtained from previous studies on 'Royal Gala' apple variety (Thewes et al. 2015; Weber et al. 2015; Both et al. 2017).

Table 1- Firmness (N), SSC (%) and TA (% malic acid) of 'Scarlet Spur' apples during cold storage

Storage conditions (Sc)		Storage time (St)						Mean
		0	2	4	6	8	10	
Firmness (N)	CA	76.60	64.84	62.81	60.21	55.25	47.76	61.25B ¹
	ULO	76.60	72.47	68.12	66.35	62.54	59.30	67.56A
	DCA	76.60	71.46	70.18	66.16	65.18	59.08	68.11A
	Mean	76.60a	69.59b	67.03c	64.24d	60.99e	55.38f	
P values		Sc ***	St ***	Sc X St *				
SSC (%)	CA	12.57	14.70	15.60	15.80	15.27	15.93	14.98 ^{NS}
	ULO	12.57	15.30	15.33	15.70	14.97	15.70	14.93
	DCA	12.57	15.00	15.23	15.63	15.73	15.57	14.96
	Mean	12.57 ^{NS}	15.00	15.39	15.71	15.32	15.73	
P values		Sc NS	St NS	Sc X St NS				
TA (% malic acid)	CA	0.36	0.33	0.30	0.29	0.27	0.23	0.30B
	ULO	0.36	0.33	0.31	0.31	0.30	0.26	0.31A
	DCA	0.36	0.33	0.31	0.30	0.29	0.27	0.31A
	Mean	0.36a	0.33b	0.31c	0.30c	0.29cd	0.25e	
P values		Sc ***	St ***	Sc X St NS				

*, P<0.05-0.01; **, P<0.01-0.001; ***, P<0.0001; NS, nonsignificant. ¹Means followed by different letters with in the same row and column are significantly different.

Table 2- Firmness (N), SSC (%) and TA (%) of 'Scarlet Spur' apples during shelf life after cold storage

Storage conditions (Sc)		Storage time (St)					Mean
		2+7	4+7	6+7	8+7	10+7	
Firmness (N)	CA	45.03	52.65	50.13	37.26	24.92	42.00B ¹
	ULO	55.58	63.40	56.30	44.84	41.27	52.28A
	DCA	62.38	56.43	62.41	52.03	39.26	54.50A
	Mean	54.33cd	57.49a	56.28b	44.71d	35.15e	
P values		Sc ***	St ***	Sc X St *			
SSC (%)	CA	16.03	15.47	15.40	16.33	16.73	15.99 ^{NS}
	ULO	15.83	15.67	16.20	15.30	15.80	15.76
	DCA	15.03	15.63	15.93	15.13	15.67	15.48
	Mean	15.63 ^{NS}	15.59	15.84	15.59	16.07	
P values		Sc NS	St NS	Sc X St NS			
TA (%)	CA	0.28	0.24	0.25	0.21	0.17	0.23B
	ULO	0.32	0.28	0.27	0.25	0.23	0.27A
	DCA	0.31	0.31	0.30	0.25	0.22	0.28A
	Mean	0.30a	0.28b	0.27b	0.24c	0.20d	
P values		Sc ***	St ***	Sc X St NS			

*, P<0.05-0.01; **, P<0.01-0.001; ***, P<0.0001; NS, nonsignificant. ¹Means followed by different letters with in the same row and column are significantly different.

In comparison with the DCA and ULO conditions in terms of fruit flesh firmness, it is better protected in the apple cultivars of 'Gloster' (Köpcke 2015), 'Granny Smith' and 'Red Delicious' (Mditshwa et al. 2017a, 2017b; Brizzolara et al. 2017) in the DCA conditions while similar results obtained from the studies regarding apple cultivars 'Golden Delicious' and 'Pinova' (Kitemann et al. 2015), 'Fuji' and 'Gala' (Zanella & Rossi 2015). This is mainly because of different metabolic reactions of apple genotypes (Brizzolara et al. 2017).

The SSC of fruit increased at the end of the storage period compared to initial values with fluctuation during cold storage. The amount of SSC was 12.57% at harvest. At the end of the cold storage period, SSC amount was determined as 15.93% in CA, 15.70% in ULO and 15.57% in DCA condition. No significant difference was observed between technologies in terms of SSC (Table 3, 4). TA values decreased significantly during cold storage and shelf life (Table 3, 4). The decrease of TA during storage is due to the consumption of malic acid as a metabolite substrate in fruit respiration (Ackerman et al. 1992). TA was significantly lower in samples stored in CA compared to other conditions ($P < 0.0001$). The ULO and DCA conditions showed similar results in the maintenance of TA amount. Similarly, ‘Granny Smith’ (Eren et al. 2015), ‘Cortland’ (DeLong et al. 2007), ‘Royal Gala’ (Weber et al. 2015), and ‘Red Delicious’ (Brizzolara et al. 2017) apple cultivars have been reported to maintain better TA levels under the DCA and ULO than CA conditions. The CA conditions with low O_2 are advantageous in maintaining TA values (Özer 2002). Generally, a decrease in the concentration of O_2 in atmosphere causes a decrease in consumption rates of citrate and malate in the formation of organic acids in the tricarboxylic acid cycle (Mir & Beaudry 2002).

Table 3- Fruit skin color changes of ‘Scarlet Spur’ apples during storage and shelf life

Storage conditions (Sc)		Storage time (St)(months)						Mean
		0	2	4	6	8	10	
L*	CA	30.53	30.66	29.64	29.59	27.02	27.21	29.11A ¹
	ULO	29.71	29.74	27.89	28.60	25.07	26.50	27.92B
	DCA	29.93	30.43	28.88	28.56	25.49	25.38	28.11B
	Mean	30.06a	30.28a	28.80b	28.92b	25.86c	26.36c	
P values	Sc***	St***	Sc X St	NS				
a*	CA	19.14	20.80	23.31	23.21	24.66	23.52	22.44A
	ULO	19.10	20.12	22.81	22.53	24.81	24.68	22.34A
	DCA	18.44	19.63	21.72	21.68	23.53	23.40	21.40B
	Mean	18.89e	20.18d	22.62bc	22.47c	24.33a	23.87ab	
P values	Sc*	St***	Sc X St	NS				
b*	CA	8.98	9.90	11.06	10.94	11.54	11.63	10.68A
	ULO	8.44	9.13	10.22	10.20	11.21	11.69	10.15AB
	DCA	8.33	9.12	9.99	9.80	10.51	10.65	9.73B
	Mean	8.58c	9.38c	10.42ab	10.32b	11.08ab	11.32a	
P values	Sc*	St***	Sc X St	NS				
Storage conditions (Sc)		Storage time (St)(months+days)					Mean	
		2+7	4+7	6+7	8+7	10+7		
L*	CA		33.97	30.72	33.27	34.78	33.02	33.15 ^{NS}
	ULO		32.37	30.63	31.70	33.07	32.70	32.10
	DCA		32.66	29.15	32.39	34.41	31.03	31.93
	Mean		33.00b	30.17d	32.46bc	34.08a	32.25c	
P values	Sc NS	St**	Sc X St	NS				
a*	CA		25.01	23.65	23.10	25.33	22.52	23.92 ^{NS}
	ULO		24.35	23.26	24.01	21.99	22.05	23.13
	DCA		24.22	23.86	23.09	24.33	22.16	23.53
	Mean		24.53 ^{NS}	23.59	23.40	23.88	22.24	
P values	Sc NS	St NS	Sc X St	NS				
b*	CA		13.61	12.33	11.56	12.46	10.81	12.16 ^{NS}
	ULO		12.23	11.83	11.78	11.23	10.79	11.57
	DCA		11.51	11.26	11.41	10.77	10.57	11.10
	Mean		12.45 ^{NS}	11.81	11.58	11.49	10.72	
P values	Sc NS	St NS	Sc X St	NS				

*, $P < 0.05-0.01$; **, $P < 0.01-0.001$; ***, $P < 0.0001$; NS, nonsignificant. ¹Means followed by different letters with in the same row and column are significantly different.

Table 4- Weight loss (%) of ‘Scarlet Spur’ apples stored in different conditions

Storage conditions (Sc)		Storage time (St)					Mean
		2	4	6	8	10	
Cold Storage (0 °C)	CA	0.44	0.85	1.21	1.51	2.08	1.22A ¹
	ULO	0.36	0.61	0.87	1.08	1.57	0.91B
	DCA	0.36	0.63	0.88	1.07	1.61	0.90B
	Mean	0.38e	0.70d	0.99c	1.22b	1.75a	
Shelf Life (+7 days at 20 °C)	CA	1.71	2.17	2.06	2.76	3.76	2.49A
	ULO	1.57	1.83	1.69	2.16	2.82	2.02B
	DCA	1.58	1.95	1.57	2.18	2.91	2.03B
	Mean	1.62e	1.98c	1.77d	2.36b	3.16a	
P values		Sc	***	Shelf	Sc	*	
	Cold Storage	St	***	Life	St	***	
		Sc X St	***		Sc X St	NS	

*, $P < 0.05-0.01$; ***, $P < 0.0001$; NS, nonsignificant. ¹Means followed by different letters with in the same row and column are significantly different.

3.3. Fruit skin color

L* value, which expresses brightness during storage and shelf life, generally decreased according to initial value. Whereas red color (a*) and yellow ground color (b*) increased due to maturation (Table 3). This increase is caused by the decomposition of chlorophyll forming the green color in the fruit during the storage and turning the color of the green in the fruit to yellow (Çalhan et al. 2012). The effect of different storage conditions on color values during storage was statistically significant ($P < 0.001$). The interaction between the storage conditions and storage time was insignificant in both cold storage and shelf-life conditions. The lowest average a* and b* values (21.40-9.73) were obtained from the samples stored in DCA conditions during the storage period. DCA preserves the quality of fruit better, by contributing to the preservation of the fruit color during storage and shelf life (Zanella et al. 2008; Veltman et al. 2003). Previous studies showed that DCA technology gave better results in maintaining color of ‘Granny Smith’ than CA (Bessemans et al. 2016) and ULO in ‘Elstar’ (Veltman et al. 2003). Additionally, it was reported that DCA delays chlorophyll degradation (Tran et al. 2015).

3.4. Weight Loss

The weight loss during storage and shelf life of ‘Scarlet Spur’ apple samples kept under different atmospheric composition was increased continuously as shown in Table 4. This change was statistically significant ($P < 0.0001$). The maximum weight loss occurred in CA for both storage (1.22%) and shelf life (2.49%) period while the weight loss observed in ULO (0.91%-2.02%) and DCA (0.90%-2.03%) conditions were similar. Weight loss is associated with the respiratory rate of the product. Increases in weight loss are due to the removal of water from the tissues along with the CO₂ released as a result of the respiration of the product during storage (Erbaş et al. 2014). The gas composition in the ULO and DCA storage conditions suppressed the respiration rate better compared to the CA storage and thus the weight loss in these conditions was also lower. The interaction between storage time and storage conditions was statistically significant ($P < 0.0001$) during storage and insignificant during shelf life.

4. Conclusions

Result of this study conducted with ‘Scarlet Spur’ apple showed that ULO and DCA conditions were more effective in maintaining quality compared to CA in terms of all quality parameters. DCA was the best storage condition suppressing respiration rate and ethylene production that expressed maturation during storage. Additionally, DCA was found to be more effective than other conditions to preserve important quality parameters in apple fruit such as color and firmness. As a result, the storage of ‘Scarlet Spur’ apples under DCA was found to be more successful than ULO and CA conditions in terms of preservation of quality criteria in long-term cold storage and during shelf life.

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