Influence of *Aloe vera* L. Based Ascorbic Acid and Lactic Acid Edible Coatings on Quality and Shelf Life of Tomato

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**ABSTRACT**

This study was done to evaluate the effect of edible coatings based on *Aloe Vera* L. (AV) in combination with ascorbic acid and lactic acid on the postharvest life and quality of tomato. Several coatings based on different natural components such as gum Arabic, carnauba, mineral oil, etc. have shown an increase in the shelf life with better retention in postharvest quality. Two edible coatings of natural *Aloe vera* gel (10%) along with Ascorbic Acid (AA; 1%) and Lactic Acid (LA; 1%) were applied to mature tomatoes as an edible coating and stored at room temperature (25-29°C) and 82-84% Relative Humidity (RH) for 30 days. After application, weight loss, total Titratable Acidity (TA), Soluble Solids Content (SSC), ascorbic acid content, pH value, total phenolic content, total antioxidant activity, and decay percentage were measured at 0, 7, 14, 21, and 30 days. Compared with untreated tomato, coated ones exhibited a significant (P≤ 0.05) delay in weight loss and higher retention of SSC, vitamin C, and titratable acidity. Between the two coatings applied to tomato, AV+1% AA+1% LA coating was found to be the most effective in delayed ripening and maintaining the postharvest losses. Results obtained in this study support using AV and AA+LA edible coating as an effective alternative to preserving tomato, delay ripening processes, and extend shelf life.

**Keywords:** *Aloe vera* gel; Antioxidant activity, *Lycopersicon esculentum*, Post-harvest quality.

**INTRODUCTION**

Tomato (*Lycopersicon esculentum*) is one of the most cultivated, consumed and important vegetables worldwide (Dursun *et al*., 2019). Tomatoes are a rich source of fiber, vitamins A, C, lycopene and epidemiological studies indicate that increased consumption of tomato lycopene is related to a lower occurrence of cardiovascular disease and certain types of cancers (Beckles, 2012). There is a growing consumer concern about the eating quality of tomatoes (Jürkenbeck *et al.*., 2019). Tomato is a climacteric fruit and has a relatively short postharvest life since, after harvest, many physical and biological factors including postharvest diseases, increased ripening, and transpiration affect quality loss (Ali *et al*., 2010; Ghaderi *et al*., 2018). In tropical regions where the temperature is high, the main factors that influence the postharvest storage life of tomato are increased respiration and ethylene production rates which result in faster ripening and decline in quality (Ali *et al*., 2010). During mass transpiration of water, vapor moves from the product to the environment, which results in a weight loss of the produce affecting storage quality (de Jesús Salas-Méndez *et al*., 2019). One of the major problems of tomatoes during storage, distribution, and ripening is softening, since it increases vulnerability to damage. These

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factors decrease the postharvest shelf-life of tomato and cause economic loss over time (Vieira et al., 2016; Yousefi Javan and Gharari, 2018). For these constraints, the potentiality of different techniques has been studied and applied to increase the shelf life of fruits and vegetable products. Generally, ripening of tomato is maintained using gas, temperature, and humidity, which are responsible for freshness and storage period as they reduce the rate of respiration and thermal decomposition and ultimately spoilage. The massive production of tomato during the harvest time and lack of efficient post-harvest processing, preservation and storage techniques leads to rapid spoilage (Ameyapoh et al., 2008). However, storage below 12.5°C causes chilling injury and affects the quality of the produce. The storage life of tomatoes can be extended by a controlled atmosphere and hypobaric storage, but these processes are costly (Beckles, 2012). The use of natural constituents as a cheaper alternative for both extending postharvest shelf life and keeping production costs low is investigated and developed in recent times (Ali et al., 2010). The needs for high-quality storage technologies without harmful effects have accentuated the notion of using the natural element as edible coatings to extend the shelf life of fresh products. Edible coatings create an additional wall that modifies the atmosphere, acting against O₂, CO₂, moisture, solute movement, delays rate of respiration, oxidation, and establishes the ability to maintain product quality and extending the shelf life of fresh food (Valverde et al., 2005; Martínez-Romero et al., 2006). These coatings also showed reduced microbiological proliferation in the food surface (Dutta et al., 2009). A model coating is well-defined as an efficient and effective system for the reduction of degradation of quality attributes in the postharvest storage period and decreases rates of loss without causing microbiological spoilage to lengthen shelf-life of fruits and vegetables (Raghai et al., 2016). For better microbial stability, appearance and texture, different functional components such as nutraceuticals, antioxidants, firming agent, preservatives, etc. can be added to coating materials (Cerqueira et al., 2009; Vieira et al., 2016).

Aloe vera (Aloe barbadensis Miller) belongs to the family Liliaceae, and is a rich source of medicinal, antimicrobial, and antioxidant agents such as phenolic compounds, carbohydrate polymers, organic acids, fibers, vitamins, amino acids, and mineral salts. Therefore, it has a wide range of use in food, pharmaceutical and cosmetic industries (Rodríguez et al., 2010; Kahramanoğlu et al., 2019). In the past few years, Aloe vera used as a novel edible coating has captured much attention for safe and environmentally friendly postharvest treatment (Odriozola-Serrano et al., 2008 Valverde et al., 2005). Aloe vera concentrations used in these studies varies depending on the produce. These studies have proven that respiration and ethylene production rate, weight loss, and softening are reduced by coating application. Moreover, aloe gel coating using a total solids content of 1.1–1.2% have been proved to maintain physicochemical factors such as color and firmness in apple slices (Chauhan et al., 2011). The gel treatments as edible coatings have shown potentiality to maintain postharvest quality with several fruit commodities such as sweet or sour cherry (Martínez-Romero et al., 2006), Hayward kiwifruit (Benítez et al., 2013), nectarine (Ahmed et al., 2009), mangoes (Dang et al., 2008), apples (Ergun and Satici, 2012), table grape (Valverde et al., 2005; Serrano et al., 2006), strawberry (Sogvar et al., 2016), blueberry (Vieira et al., 2016), orange fruit (Rasouli et al., 2019), and papaya (Mendy et al., 2019).

Components similar to essential oils, acetic acid, and ascorbic acid were incorporated with A. vera coatings and showed increased activity (Sogvar et al., 2016). As an antioxidant, Ascorbic Acid (AA) reduces vitamin C loss and can be added to the edible coating material. Antimicrobial properties of AA on fresh-cut
fruit such as papaya (Tapia et al., 2008) have been reported.

This work aimed to study the consequence of *A. vera* incorporated with ascorbic acid and lactic acid as an edible coating on the alteration in physicochemical factors linked to quality throughout storage and role in prolonging the shelf life of tomato.

**MATERIALS AND METHODS**

**Plant Material**

Tomatoes (*Lycopersicon esculentum*) harvested at the mature green stage were procured from a native farmer in Sylhet, Bangladesh. At the laboratory, tomatoes were visually categorized based on consistency in size, shape, and color, the absence of injuries, blotches, and malady. Tomatoes were divided into three batches of control, only AV coated, and AV+AA+LA coated batch. A total of 60 tomatoes were taken containing 20 tomatoes in each batch. The procured tomatoes were washed thoroughly with running water and exterior dried before application of the coating.

*Aloe vera* (AV) Coating Preparation, Treatment, and Storing

The *Aloe vera* gel coatings were prepared through a slight modification of the method followed by Brishiti et al. (2013) *Aloe vera* leaves were collected from Natore district, Bangladesh. After a thorough wash with warm (40°C) tap water, the leaves were divided longitudinally, and the colorless tissue was scratched out, liquefied in a food-grade blender (HJ-H176P, Malaysia) at maximum speed for 10 seconds and homogenized. The liquid acquired was *Aloe Gel* (AG; 100%). The liquid was pasteurized at 70°C for 45 minutes, followed by immediate cooling at ambient temperature. To increase coating ability, 1% gelling agent was used for thickening. To enhance the plasticizing effect, 2% glycerol was added. Oleic acid was added to avoid precipitation; for uniform dispersion, an antimicrobial, namely, Cinnamaldehyde was added alongside oleic acid. The solution was then filtered after adding water to form a 10% AV solution.

The solution was then divided into half, and in one solution, 1% LA and 1% AA were added. The coating application was accomplished by dipping the samples in the corresponding liquid solution for 10 minutes. Later, all fruits were air-dried at room temperature for 1 h, then placed in the tray. Four tomatoes from each batch were sampled at 0, 7, 14, 21 and 30 days of storage.

**Physicochemical Analyses of Tomato**

**Weight Loss Percentage**

Tomato samples were weighed using an electronic analytical balance (AY 220, Shimadzu Corporation, Japan) at day 0 and at the end of each storage interval for 30 days. Weight loss was measured by taking initial and final weight differences of samples and expressed as a percentage.

**pH, Total Soluble Solids (TSS) and Total Titratable Acidity (TTA)**

At every 7 days of intervals, except 30th days, three fruits per treatment were analyzed. The samples were grounded in a blender (HJ-H176P, Malaysia) after cutting into small pieces. To determine the pH of each treated sample, a potentiometer (Hanna Instruments Inc., Romania) was used. A digital refractometer (Brix 0–32%, Atago Co Ltd) at 20°C was used to determine TSS according to the AOAC method (932.14) and the results were recorded as %. Then, a filter paper (Whatman No. 1) was used to filter the grounded mixtures. To determine TA, 10 mL of pulp from each sample was taken and phenolphthalein (1%, 2 drops) was added and titrated with 0.1 N NaOH.
(AOAC, 1990). Results were shown in percent of Citric Acid (% CA).

**Vitamin C Assay, Total Phenolic Content and Antioxidant Activity (AA)**

Dye 2, 6-Dichlorophenolindophenol (DCPIP) visual titration method by Ranganna (2004) was used to estimate AA contents as mg per 100 gram of fresh weight. A slightly modified colorimetric method (Singleton et al., 1999) by the Folin-Ciocalteu reagent was used to determine total phenols. Tomato samples were centrifuged at 2,700×g for 15 minutes and filtered through Whatman No 1 filter paper. Afterward, 0.5 mL of the extract was mixed with 0.5 mL of Folin-Ciocalteu reagent and 10 mL of saturated Na₂CO₃ solution was added after 3 minutes. The absorbance was measured at 725 nm after allowing the sample to stand for 1 hour. By comparing the absorbance of the samples with standards, concentrations were determined. Results were expressed as gallic acid equivalents per kilogram of fresh weight (mg kg⁻¹). DPPH free radical scavenging assay was performed by taking 100 µL of tomato extract mixed carefully with methanolic 2 mL of freshly prepared 0.1 mmol L⁻¹ DPPH solution. The absorbance was measured in a UV spectrometer (T60 U, PG Instruments LTD) at 517 nm. Blank was prepared by Methanol (100 µL) without the extract. The scavenging ability of the DPPH radical was calculated by the following formula:

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\text{DPPH-scavenging effect (\%)} = \left[1 - \left(\frac{A_{517, \text{sample}}}{A_{517, \text{blank}}}\right)\right] \times 100
\]

**Decay Percentage**

The decay of treated and untreated samples was determined as the number of the decayed samples divided by the initial number of all sample and the result was multiplied by 100 (El-Anany et al., 2009)

**Statistical Analysis**

Triplicate evaluations were performed and means were expressed as the results. Significance in differences between samples for all parameters was calculated using a two-way ANOVA. Statistical analyses were carried out using SPSS. The chosen p-value was \( P \leq 0.05 \).

**RESULTS AND DISCUSSION**

**Weight Loss**

The results for all tomato samples showed increased weight loss as the time of storage increased (Figure 1-a; \( P \leq 0.05 \)) and significant differences were found. AV+AA+LA-treatments prevented more weight loss than AV alone. Tomatoes not coated had a statistically higher loss in mass compared to tomato coated during the storage time. The natural transpiration process is the main reason for this weight loss tendency during storage. The coating reduces respiration, water loss, and oxidation reaction rates because it creates a semi-permeable barrier against gas and moisture movement (Vieira et al., 2016). Similar results were observed when tomatoes were treated with gum Arabic (Ali et al., 2010), Aloe vera gel treated “Arctic Snow” nectarines (Ahmed et al., 2009) and aloe gel treated sweet cherries (Martínez-Romero et al., 2006), 33% gel treated table grapes (Valverde et al., 2005) and in 10% gel treated ‘Granny Smith’ apples (Ergun and Satici, 2012). The greatest effect observed on the reduction of weight loss was in the combinations of AA+LA with AV.

**SSC, pH and Titrable Acidity (TA)**

A gradual increment in SSC was found during the total storage time Figure 1-b. The soluble solids, however, were significantly higher in control tomato and lowest SSC.
was found in AV+AA+LA coated tomato. The lowest SSC was observed in tomato coated with AV+LA+AA.

This indicates that the bio preservative coatings provided a semi-permeable layer on the samples to reduce O2 production or elevates CO2 and subdues ethylene production by modifying the internal atmosphere. Results similar to this were found in tomato coated with mango kernel and tomato coated with almond gum trees exudate (Nawab et al., 2017).

Results indicate a pH increase in all the samples, though the untreated tomato showed a higher increase than treated samples. The reason behind pH value increment is that, during the time of storage, the acid disintegration happens as respiration goes on. Fruit taste quality is greatly influenced by the acid change. While tomato ripened, the major amounts of acids found were citric acid and malic acid. The increase in the pH of tomato with maturation reported here was in the range reported by other studies (Fernández-Ruiz et al., 2004)

The values of titratable acidity of tomato samples decreased with increased storage period (Figure 1-d), and the value for AV+AA+LA treated tomato compared to the control was significantly different, with a maximum reduction for uncoated fruit. The pH value was inversely proportional to the TA value. The reduced TA value in the control tomato in comparison with treated tomato indicates delayed ripening effect of the coating by creating a barrier and reducing the rate of respiration. Climacteric agricultural products use organic acids as primary substrates for the respiration process, and coating prevents the utilization of these acids while reduction is expected for respiring produces (El-Anany et al., 2009). This retains the firmness and retards metabolic processes that increase the storage period of tomato. Modified Atmosphere Packaging (MAP) shows similar results
where the modification behaves like a gas barrier, which slows down the respiration rate by reducing oxygen uptake. (Arowora et al., 2013). Aloe gel-treated nectarines showed similar results (Ahmed et al., 2009; Athmaselvi et al., 2013). It has been found that quality loss is influenced by decreased amounts of acidity (Patanè et al., 2019).

**Vitamin C**

There was an increase in the ascorbic acid or vitamin C content of treated and untreated tomato, which peaked after 7 days and declined afterward (Figure 2-e). AV+AA+LA coated tomato showed a maximum amount of ascorbic acid and there was no significant difference between AV alone and AV+AA+LA coatings. With maturity, ascorbic acid content increased in tomato but once it reached the maximum ripe stage, concentration declined. A spontaneous process, namely, autooxidation occurs when ascorbic acid and oxygen combine, which is a probable reason for the loss in vitamin C (Sogvar et al., 2016). The coating treatment slows down ascorbic acid increment, indicating that the coating layer reduced but did not stop the production of vitamin C. The effective retention of ascorbic acid levels is due to the lower level of gas permeability that retards the physiochemical processes during storage. A similar kind of increase and gradual decrease after maximum value were reported by other studies (Ali et al., 2010; Athmaselvi et al., 2013).

**Total Phenolic Content and Antioxidant Activity**

Figure 2-f shows the changes in the phenolic content of tomato during storage. The contents of the TP of tomato fruits in different maturity stages studied are within the range observed by others (Odriozola-Serrano et al., 2008; Valdivia-Nájar et al., 2018). The mature red stage of tomato showed the maximum polyphenols content; however, the lowest polyphenols content was observed in the mature green phase of tomato. Total phenolic content showed a significant reduction in the over-ripening stage. The increase in physiological activities due to ripening results in an increase in TPC. Similar results were observed in other studies, indicating that the content of total phenols decreases after the maturity stage of tomato fruits (Odriozola-Serrano et al., 2008). However, coated tomato maintained lower values during storage and delayed ripening (Mirdehghan and Valero, 2017). One of the significant biological properties of tomato is the antioxidant capacity of phenols and lycopene. Therefore, maintaining the natural balance of these compounds during storage is very crucial. Results suggest that AV+AA+LA treated tomato showed the lowest amount of TP content.

There was a decrease in antioxidant capacity in all the samples, either treated or untreated Figure 2-g. The antioxidant capacity of aloe extract was reported in many studies. Different studies confirm that Aloe vera gel shows antioxidant capacity because of antioxidant polyphenols, indoles, and alkaloids presence, as confirmed by different analyses (Nejatzadeh-Barandozi, 2013). It is effective in delaying browning, dehydration, and maintains the visual quality of berry without causing any harmful effect on other sensory attributes (Mirdehghan and Valero, 2017). Aloe vera may increase its free radical scavenging capacity to resist tissue damage. Antioxidant activity decreases because of senescence at the time of storage. Decreases in antioxidant activity of the samples may be influenced by the capability of retaining fruit quality characteristics by suppressing enzyme activity that destroys antioxidant components. Vitamin C and phenolic compounds are antioxidant compounds that impart antioxidant activity (Tulipani et al., 2008; Sogvar et al., 2016). The delayed ripening process during storage due to treatment of Aloe vera edible coating was
parallel to the previous studies for fruits including strawberry, papaya, nectarine, etc. (Martínez-Romero et al., 2006; Ahmed et al., 2009).

Decay Percentage

The treated and untreated samples showed no sign of decay until day 7, as presented in Figure 2-h. After that, there was a significant reduction in decay in AV+AA+LA treated tomato compared to the other samples, and after 30 days storage, AV+AA+LA treated samples remained free of significant decay. After 21 days of storage, 66% of the control samples showed considerable amount of spoilage. Treated samples decreased decay percentage, perhaps by delaying senescence. Similar results were found in the case of gum Arabic treatment on tomato (Ali et al., 2010). The visual evaluation confirmed that the uncoated tomatoes spoiled extensively after 21 days of storage (Figure 3). The coating solution of A. vera with or without LA+AA appears to have inhibited the ripening when compared with uncoated tomato samples. Figure 3 shows three batches of tomato samples (coated, AV, and AV+LA+AA).

This inhibition in ripening may be due to the effect of the coating layer of Aloe vera and AV+LA+AA on the surface. Less surface color change was observed in AV+AA+LA treated samples. In other studies, Aloe vera has been reported to have an antifungal effect, which is important for the reduced damage of tomato (Vieira et al., 2016).

CONCLUSIONS

Among all the studies that have been conducted with Aloe vera, in this study, for the first time, Aloe vera was used in
combination with AA+LA in tomato. The results suggested that tomato samples coated with AV+AA+LA retarded postharvest ripening and maintained sample quality attributes most efficiently during storage at 25°C up to 30th day as compared to the untreated control tomato that reached its maximum maturity in 14 days. This study indicated that AV with AA application could be used as a biochemical means for maintaining tomato quality and increasing its shelf life. Additional studies need to be conducted to understand the gaseous exchange of AV+LA+AA coating layer and develop new formulations to apply to different climacteric fruit and vegetables at low temperatures. Further research is needed to understand how coating influences microbial growth and thus affects the ripening process. Addition of A. vera with ascorbic acid in the edible coating as bio preservative, which contains antifungal and antioxidant compounds, provides a novel means to enhance safety and increase the postharvest storage life of tomato. It also ensures health and environmental protection by preventing the use of artificial harmful chemical preservative compounds in various agricultural products.

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تاثیر پوشش خوراکی اسکوربیک اسید و لاکتیک اسید بر پایه آلوه و روی کیفیت و ماندگاری گوجه فرنگی

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چکیده

هدف از انجام این پژوهش ارزیابی تاثیر پوشش خوراکی اسکوربیک اسید و لاکتیک اسید بر پایه آلوه و (AV) روی ماندگاری و کیفیت گوجه فرنگی بعد از برداشت محصول بود. چندین پوشش مبتنی بر مواد طبیعی مختلف مانند صمغ عربی، روغن معدنی، و...) به اندازه ماندگاری و نگهداری بهتری کیفیت بعد از برداشت شده اند. در این پژوهش، دو پوشش خوراکی از ول آلوده ورای طبیعی (10%) همراه با اسکوربیک اسید (AA) 1% و لاکتیک اسید (LA) 1% بعنوان پوشش خوراکی مورد استفاده قرار گرفت. پوشش خوراکی به گوجه فرنگی های رسیده که در حرارت اطاق (29-25 درجه سانتی گراد و نسبت RH 84-82%) به مدت 30 روز از شده بود افزوده شد. سپس، مقدار تلفات وزن، اسیدیته کل قابل تیتر (TA)، محتوای جامدات محلول (SSC)، تیتر اسکوربیک اسید، اسیدیته کل، فعالیت آنتی اکسیدانی کل، و درصد پوستیگی در روزهای 14، 7، 21، و 30 اندازه گیری شد. گوجه فرنگی هایی که پوشش داده شده بود در مقایسه با آنها که نامه شده بود ناخیر معانار (P≤0.05) را در تلفات وزن نشان داد و مقدار بیشتری SCC، و اسیدیته کل قابل تیر داشت. در آزمون ورش آزمون شده، موتورترین پوشش برای تاخیر در رسیدن میوه و نگهداری از تلفات بعد از برداشت، پوشش با AV+ 1% AA+1% LA موثرتر بود. نتایج به دست آمده در این بررسی نشان داد که می‌توان از AV+ 1% AA+1% LA به عنوان گزینه‌ای موتور برای حفظ و نگهداری گوجه فرنگی، تأخیر در فرآیند رسیدن میوه، و طولانی کردن ماندگاری استفاده کرد.