SHORT COMMUNICATION

Effect of integration of oxalic acid and hot water treatments on postharvest quality of rambutan (Nephelium lappaceum L. cv. Anak Sekolah) under modified atmosphere packaging

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Abstract The shelf life of rambutan is often limited due to rapid water loss from the spinterns and browning of the pericarp. An integrated approach, which combined hot water treatment (HWT) (56 \degree C for 1 min), oxalic acid (OA) dip (10% for 10 min) and modified atmosphere packaging (MAP), was used to study their effectiveness on the quality of rambutan during storage (10 \degree C, 90–95%) relative humidity). Significant differences were observed in rambutan quality with the combination of $MAP + HW$ $T + OA$ after 20 days of storage. This treatment combination resulted into better retention of firmness and colour (L and a* values) than in the control. Change in the total soluble solid content was significantly delayed however the titratable acidity showed no significant change in comparison to the control at the end of storage.

Keywords Rambutan - Oxalic acid - Firmness - Colour - Weight loss

Introduction

Rambutan (Nephelium lappaceum L.) is a non-climacteric tropical fruit in the family Sapindaceae which includes litchi and longan (Sun et al. [2010](#page-4-0)). It was domesticated in Southeast Asia with West Malaysia and Sumatra thought to be the place of origin. In 2000, approximately 6960 tonnes of fresh rambutan, worth about 4.8 million ringgit, were

 \boxtimes Asgar Ali Asgar.Ali@nottingham.edu.my exported from Malaysia (Sarip [2002\)](#page-4-0). The fruit is well appreciated globally due to the attractive colour of its peel and spinterns as well as the distinctive flavour of the arils.

The marketability of rambutan is often limited due to rapid skin desiccation and browning during storage (O'Hare [1995](#page-4-0)). The spinterns (hair-like protuberances) can facilitate water loss from the peel which makes the fruit highly perishable (Yingsanga et al. [2008](#page-4-0)). Landrigan et al. [\(1996](#page-4-0)) reported that browning of the pericarp and spinterns was positively correlated. They suggested that the marketability of rambutan following storage was unacceptable when 25–40% of water was lost and there was rapid development of browning of the pericarp. Pericarp browning of rambutan postharvest is due to the oxidation of two main phenolic substrates, epicatechin and proanthocyanin, by the enzyme polyphenol oxidase (PPO) to quinones, which further polymerise to melanin (Sun et al. [2010](#page-4-0)).

Fumigation with sulphur dioxide $(SO₂)$ and dipping in hydrochloric acid have been applied commercially to inhibit pericarp browning and extend the shelf life of longan, litchi (Saengnil et al. [2006](#page-4-0)) and rambutan (Mohamed et al. [1988](#page-4-0)). However, Mohamed et al. [\(1988](#page-4-0)) reported that the organoleptic properties of rambutan fumigated with $SO₂$ were adversely affected as the skin was bleached, due to the formation of colourless anthocyanin- $SO₃$ complex, and a strong sulphurous odour was developed in the flesh. Similarly, Latifah et al. ([2009\)](#page-4-0) reported that rambutan packed in polyethylene bags resulted in better retention of red colour, higher relative humidity and significantly less weight loss when compared to that of control fruit which showed accelerated senescence at the end of storage. However, the method applied can only extend the shelf life of rambutan for 6 days. Also, Nurhuda et al. ([2013\)](#page-4-0) reported that water blanching was effective to inhibit the

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enzymatic activity of PPO and polyphenol peroxidase (POD), but the peel colour was bleached after 5 min. To overcome the problems inherent with these treatments, an alternative needs to be developed to deliver safe and high quality fresh commodities to consumers.

Recently, oxalic acid (OA) has received much attention due to its potential application in the pre- or post-harvest preservation of fresh commodities. Being a natural organic acid which has some antioxidant activity, OA may confer significant benefits in relation to environmental stress, systemic resistance and anti-senescence in harvested fruits (Huang et al. [2013;](#page-4-0) Jin et al. [2014\)](#page-4-0). For example, OA delayed ripening and enhanced resistance against pathogen invasion in banana (Huang et al. [2013](#page-4-0)), (Wang et al. [2009](#page-4-0)), plum (Wu et al. [2011](#page-4-0)) and mango (Zheng et al. [2007\)](#page-4-0). The application of OA alone or in combination with other treatments to inhibit pericarp browning has been demonstrated in studies on litchi (Saengnil et al. [2006;](#page-4-0) Zheng and Tian [2006\)](#page-4-0) and longan (Whangchai et al. [2006\)](#page-4-0).

To the best of our knowledge, no study has been conducted on the combination of OA, HWT and MAP on the storage life of rambutan. Therefore, the aim of this study was to investigate the effects of this integrated approach on the physicochemical properties of rambutan postharvest.

Materials and methods

Plant material

Fresh rambutan fruits (N. lappaceum cv. Anak Sekolah) of colour index 4 with light red peel and green spinterns were purchased from Pasar Borong Selayang, Selangor during the commercial harvest in September, 2013. Fruits of uniform size, shape, free from any mechanical damage and insect or pathogenic infection were selected for this study. They were washed with 0.05% (v/v) sodium hypochlorite and air dried at ambient temperature $(25 \degree C)$ for 30 min.

Experimental design

Hot water treatment of fruit was carried out in a water bath (dimensions $590 \times 350 \times 220$ mm) at 56 °C. Polyethylene (PE) bags (thickness 0.08 mm) were purchased from a local company. Based on a preliminary study (results not shown), 10% (w/v) of OA was selected as the optimum concentration for the experiment. Fruits were divided into eight treatments with three replicates of four fruits per treatment. The following treatments were applied:

1. C: Fruits not subjected to any treatment served as control

- 2. MAP: Fruits packaged in PE bag
- 3. OA: Fruits dipped in 10% OA solution for 10 min
- 4. HWT: Fruits immersed in hot water bath at 56° C for 1 min
- 5. HWT $+$ OA: Fruits immersed in hot water bath at 56 °C for 1 min and dipped in 10% OA solution for 10 min
- 6. MAP $+$ OA: Fruits dipped in 10% OA solution for 10 min and packaged in PE bag
- 7. MAP $+$ HWT: Fruits immersed in hot water bath at 56 \degree C for 1 min and packaged in PE bag
- 8. MAP $+$ HWT $+$ OA: Fruits immersed in hot water bath at 56 \degree C for 1 min, dipped in 10% OA solution for 10 min and packaged in PE bag.

After treatment, fruits were stored at 10 $^{\circ}$ C and 90–95% relative humidity (RH). Data were collected on day 0 and at intervals of 5 for 20 days.

Firmness

Fruit firmness was measured using an Instron Universal Testing Machine (Model 5540, USA) in the compression mode. Four fruits in each replication were assessed using a 2 mm diameter probe at a speed of 300 mm/min and a load range of 100 N. The maximum force (N) to penetrate the fruit was expressed as the mean of readings taken at three positions selected randomly on the equator of the fruit.

Colour

Peel colour was determined using a HunterLab MiniScan Xe Plus (Minolta Corp., Japan). Two specific points along the equator of the fruit were used to determine the colour. The data were expressed as mean \pm SE for L and a* values.

Weight loss

Four fruits in each replicate for each treatment were marked before storage and weighed on a digital balance at the beginning of the experiment and at the end of each storage interval. The results were expressed as the percentage loss of initial weight.

Total soluble solid content (TSS) and titratable acidity (TA)

The TSS (\textdegree Brix) and TA were done by method given by Ali et al. (2011) (2011) . Results were expressed as $(^{\circ}Brix)$ reading and percentage of citric acid per 100 g fresh weight, respectively.

Statistical analysis

The experiment was arranged in a completely randomised design (CRD) with three replicates in each treatment. The data were subjected to analysis of variance (ANOVA) using SAS 9.1 software and means were separated using Duncan's Multiple Range Test at $P < 0.05$.

Results and discussion

Firmness

Fruit treated with $MAP + HWT + OA$ showed significantly ($P < 0.05$) better retention of firmness when compared with the control (Fig. 1). Regardless of treatment, there were no differences in the firmness of treated and untreated rambutan on days 0, 5 and 10. On day 20, MAP alone was insufficient to retain firmness of the fruit. The highest firmness (11.48 N) was recorded at the end of the storage in fruit treated with $MAP + HWT + OA$, but was not significantly different from that of HWT or OA treated fruit. Significantly higher retention of firmness was observed in fruits treated with $MAP + OA$ and OA on day 20 when compared to the control.

In this study, MAP alone was insufficient to retain the firmness of rambutan at the end of storage suggesting that the retention of firmness of $MAP + HWT + OA$ treated rambutan is mainly due to the synergistic effect of OA and HWT. Martínez-Esplá et al. [\(2014](#page-4-0)) suggested that less softening in OA treated fruits may be due to the formation of oxalate insoluble pectin which slows down the polymerization of pectin. It has been shown that optimizing the concentration of OA and the conditions for HWT can

Fig. 1 Effect of various treatments on firmness of rambutan during 20 days of storage at 10 °C and 90–95% relative humidity (RH). Values are mean \pm SE

inhibit the activity of cell wall degrading enzymes, such as polygalacturonase (PG) and pectin methyl esterase (PME), which in turn maintains the firmness of fruits (Amnuaysin et al. [2012](#page-4-0); Wu et al. [2011](#page-4-0)). Similar results were also obtained in OA treated fruits of banana (Huang et al. [2013](#page-4-0)), mango (Li et al. 2014) and sweet cherry (Martinez-Esplá et al. [2014\)](#page-4-0).

Weight loss

Weight loss of rambutan increased progressively with storage (Fig. 2). Minimum weight loss (0.35%) was recorded in fruit treated with $MAP + HWT + OA$, but there was no significant difference when compared to fruit treated with MAP, MAP $+$ OA and MAP $+$ HWT. The highest weight loss was recorded in control (24.6%), followed by HWT $+$ OA (23.1%), HWT (22.1%) and OA (20.3%) treated rambutan, respectively.

It should be noted that water loss from rambutan occurs mainly through the spinterns in which the stomatal density is at least five times higher than in the main body of the fruit (Latifah et al. [2009](#page-4-0)). Consequently, the highest percentage weight loss was observed in the control fruit. MAP can control the respiration and transpiration of fruit as well as the relative humidity of the packaged fruit and hence result in lower weight loss when compared to the control.

Colour

Regardless of treatment applied, the L and a* values of rambutan decreased gradually throughout the storage period (Fig. [3](#page-3-0)a, b). After 20 days, the L value of control fruit was reduced from 42.70 to 3.97, i.e. retained only 9.3% of its initial value. The highest retention of the L

Fig. 2 Effect of various treatments on weight loss rambutan during 20 days of storage at 10 $^{\circ}$ C and 90–95% relative humidity (RH). Values are mean \pm SE

50

A

value (59.4%) was observed in fruit treated with $MAP + HWT + OA$. Similarly, a* value of rambutan treated with $MAP + HWT + OA$ was significantly $(P<0.05)$ higher at the end of storage when compared with the control and other treatments. The results showed that the combination of $MAP + HWT + OA$ can retain the bright red colour of rambutan throughout storage. In contrast, untreated rambutan became unacceptable at the end of storage as the pericarp colour changed progressively from red to dark brown. Fruits treated with MAP, HWT or OA alone showed no differences with the control at the end of storage suggesting that the treatments applied alone are ineffective to maintain the colour of rambutan.

It should be noted that the chelating property of oxalic may promote the binding of PPO with copper and POD with iron to form inactive complexes which in turn inhibit the activity of the enzymes (Saengnil et al. [2006;](#page-4-0) Zheng and Tian [2006](#page-4-0)). Saengnil et al. ([2006\)](#page-4-0) reported that HWT

> C MAP OA HWT $HWT + OA$ MAP + OA

followed by oxalic acid inhibited pericarp browning in litchi fruit and suggested that HWT can reduce the high surface tension of the epidermis of the pericarp and hence enhance the permeability of oxalic acid into the pericarp of fruits. Whangchai et al. ([2006\)](#page-4-0) also reported that the combination of ozone and oxalic acid resulted in better colour retention of longan fruit.

TSS and TA

TSS increased more $(P < 0.05)$ in the control as compared to other treatments (Fig. 4) with the highest SSC (18.4 Brix) after storage for 20 days. In contrast, TSS in fruit treated with $MAP + HWT + OA$ was 78.4% lower than that of the control. Regardless of treatment applied, TA of rambutan decreased throughout storage (Fig. [5\)](#page-4-0). There were no differences ($P > 0.05$) between control and treated samples at the end of storage.

The rate of increase in TSS or decrease in TA has been reported to be dependent on the metabolic activity of fruits (Mustafa et al. [2014\)](#page-4-0). The accumulation of reducing sugars and utilization of organic acids was slower in MAP treated rambutan possibly due to the reduced oxygen content in the MAP system which slows down the senescence of rambutan. Similarly, Latifah et al. [\(2009](#page-4-0)) reported that the increase of TSS was delayed in PE packed rambutan when compared with control fruit after 6 days of storage.

Fig. 3 Effect of various treatments on L (a) and a^* (b) value of rambutan during 20 days of storage at 10 $^{\circ}$ C and 90–95% relative humidity (RH). Values are mean \pm SE

Storage Time (Days)

Fig. 4 Effect of various treatments on TSS of rambutan during 20 days of storage at 10 °C and 90–95% relative humidity (RH). Values are mean \pm SE

Fig. 5 Effect of various treatments on TA of rambutan during 20 days of storage at 10 °C and 90–95% relative humidity (RH). Values are mean ± SE

Conclusion

All the treatments tested alone or in combination for different physiochemical parameters under study showed significant differences when compared to control, with different efficacies. However, treatments when used alone (MAP, OA or HWT) were insufficient to maintain firmness, water retention, colour and TSS of rambutan fruits during storage. Amongst all the combination of $MAP + HWT + OA$ was effective in maintaining the most of the physicochemical properties of rambutan for up to 20 days in storage. The combination treatment effect is most likely due to synergistic effect. However, more in depth studies, especially on the anthocyanin content, browning enzymes such as phenylalanine ammonia-lyase (PAL), phenol peroxidase (PPO) and peroxidase (POD) and respiration of treated and untreated rambutan, should be carried out.

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