



Original Article

Effects of minimal processing on the respiration rate and quality of rambutan cv. 'Rong-Rien'

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Abstract

Respiration rate at 4°C and minimal processing of rambutan cv. 'Rong-Rien' were investigated. Rambutan was harvested from Amphur Ban Na San, Surat Thani Province, at the stage when its skin was turning into a combination of red, green and yellow. After harvesting, the fruits were size-graded to 27-30 fruits/kg, hydrocooled to 14°C, packed with ice in Styrofoam boxes and transported to the laboratory at Prince of Songkla University within 6 h. The respiration rate of fresh rambutan fruits was monitored. For minimal processing, the fruits were soaked in warm solution (55°C) of 100 ppm sodium hypochlorite for one min and immediately cooled in cold water until their internal temperature reached 14°C. The minimal process included peeling, with and without coring. The peeled and peeled and cored rambutan samples were immersed in a solution of 0.5% citric acid + 0.5% CaCl₂ at 4°C for 2 min. The average respiration rates (within 6 h) at 4°C of whole fruit, peeled, and peeled and cored rambutan samples were measured and found to be 122, 134 and 143 mg CO₂/kg/h, respectively. These findings indicated that a preparation style as peeled rambutan without coring, nylon/LLDPE bag, storage temperature of 4.0±1°C, were suitably applied for processed rambutans. To obtain a longer extended shelf life (>12 days) of minimally processed peeled rambutans, further study on food additives, including acidulants and preservative used and gas composition in modified atmosphere packaging (MAP) is needed.

Keywords: respiration rate, rambutan, rambutan cv.Rong-Rien, minimally processed rambutan, modified atmosphere packaging

1. Introduction

Rambutan (*Nephelium lappaeum* Linn.) cv. 'Rong-Rien' is one of Thailand's economic crops with a sizeable export market. It has been classified as nonclimacteric fruit (O'Hare, 1995). The fruit will not ripen if removed from the tree before full maturity, the state of which can be determined principally by appearance (O'Hare, 1995). Rambutan shelf

life is limited by dehydration of the fruit and browning of spinterns (long soft hair) and skin (Ketsa, 1985; Wanichkul and Kosiyachinda, 1982). The darkening of spinterns and skin renders rambutan unmarketable, even though its pulp is still of acceptable eating quality.

The deterioration of rambutan skin can occur rapidly within three days under ambient conditions (O'Hare, 1995). There have been many reports on shelf life extension of rambutan with modified atmosphere packaging (MAP) combined with low storage temperature (Mendoza *et al.*, 1972; Klaewkasetkorn, 1993; Srilaong *et al.*, 2002; Patika-

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butr, 2001). Luckanatinwong (2005) reported that packaging rambutan in polyethylene bags (70 μm thick) under 5% CO_2 , 5% O_2 and 90% N_2 and stored at 10°C could extend its shelf life for 23 days. Minimally processed fruits and vegetables have recently become popular products due to consumer demand for convenience and fresh or near-fresh quality. It has many advantages over fresh produce, especially for export market. For example, it has lower product volume, thus lower transport cost. It can circumvent problems common to exporting fresh whole produce, e.g. dehydration and rotting, and the needs for quarantine process. Minimally processed fruits and vegetables provide consumer with a new class of products where convenience, quality and relatively long shelf life are desirable (Siriphannich, 1994). However, quality and shelf life of minimally processed fruits and vegetables depend greatly on factors such as respiration rate, environment they are packaged under, processing techniques, storage temperature, etc. (Zagory and Kader, 1988). Respiration is the process by which stored organic materials (carbohydrates, proteins, fats) are broken down into simple end products with a release of energy for cellular functions (Kader, 2002). After the fruits are separated from the tree, the respiratory process must continue to produce energy or the tissues will die leading to cellular senescence. The processing procedures of peeling and cutting cause tissue wounding and induce a high respiration rate, which triggers faster water loss and susceptibility to microbial deterioration compared to intact tissues (Dong *et al.*, 2000; Kader, 2002). Edible coating with many types of additives may be applied on minimally processed fruits to serve several different purposes such as reduction in water loss and microbial decay, and maintaining an acceptable appearance (Olivas and Barbosa-Cánovas, 2005). Fresh-cut mango, similar in minimal process as other various tropical fruits, cutting into 3.0 x 4.0 in., soaking in mixture solution of 2-5% citric acid, 2-5% ascorbic acid and 0.5-2.0% calcium chloride (by volume) at 3-4°C for 1 min, placing 100-150 g each on a plastic tray, packaging in Nylon/PE/LLDPE bags, flushing with a mixture (by volume) of 15-25% CO_2 , 0-6% O_2 (balance N_2), sealed and stored at 10°C, had acceptable qualities for consumption after 14 days storage (Ngarmsak *et al.*, 2003). Agar *et al.* (1999) reported that fresh-cut kiwifruit slices had a shelf life of 9-12 days if treated with 1% CaCl_2 or 2% Ca lactate in combination with cholorine water ($100\mu\text{l}\cdot\text{L}^{-1}$) dip and stored in 1.9-L jars at 0-2°C, RH >90%, with C_2H_4 scrubbing, in 2 to 4 kPa O_2 and/or 5 to 10 kPa CO_2 . Rambutan is a major export fruit in Thailand; however, there have been only a few reports dealing with minimal processing of fresh-cut products (Ngarmsak *et al.*, 2003). The objectives of this research were to study the effects of minimal processing including peeling with and without coring, suitable additives, types of packaging films, modified atmosphere and storage temperatures on the respiration rate and qualities of rambutan.

2. Materials and Methods

2.1 Raw materials

Fresh rambutans cv. 'Rong-Rien', from the orchards in Amphur Ban Na San, Surat Thani Province, were harvested at the stage when their skin was turning into a combination of red, green and yellow (100 days after blooming). After harvesting, the fruits were size-graded to 27-30 fruits/kg, hydrocooled to 14°C, packed with ice in styrofoam boxes and transported to the laboratory at Prince of Songkla University within 6 h.

1) Respiration rate measurement

Respiration rate measurements of fresh fruits and minimally processed rambutan were carried out. Fresh rambutan fruits were washed with tap water, and excess water removed by basket centrifugation. For minimal processing, the fruits were soaked in warm solution (55°C) of 100 ppm sodium hypochlorite for one minute and immediately cooled in ice water until their internal temperature reached at 14°C. The minimal processing was conducted in a temperature-controlled (25°C) room fitted with a UV-lamp (40Wx2, Narva), which was turned on at least 12 h prior to the experiment. The minimal process included peeling, with and without coring. The peeled and peeled and cored rambutan samples were immersed in a solution of 0.5% citric acid + 0.5% CaCl_2 at 4°C for 2 min. Approximately 200 g each of peeled and peeled and cored rambutan samples were placed in 1,200 ml sterile jars. Each set had six replicates. The jar lids had stoppers for gas sampling and rubber tubes were connected to allow airflow through the jars. The inlet tube was inserted down to the bottom of the jars to ensure consistent flushing of the air. The jars were stored at 4°C and a flow-through system was used to allow quantification of the CO_2 production rate over time. Humidified air at 4°C was fed to the jars at a flow rate of 152-154 ml/min. Gas samples of 10 ml were taken at the jar outlet at 1 h interval for 6 h with a 10 ml NIPRO (NIPRO, Thailand) plastic syringe with 26G (0.45x25mm) needles. CO_2 production rate (%) at a given time was determined by a gas chromatograph (Shimadzu Scientific Instruments Inc., Model GC-8A, Kyoto, Japan) operated at 70°C oven temperature, 150°C for the injector port and thermal conductivity detector, and calculated as mg CO_2 /kg/h, according to the method of Watada and Pratt (1980, cited in Postharvest Research Laboratory, Kasetsart University Kamphaeng Saen Campus, 2004). The amounts of N_2 , O_2 , and CO_2 , were determined in the gas chromatograph using a stainless steel column packed with Porapak Q (80/100 mesh, ID 3.0mm) and 1.93 m in length. The respiratory rates (average CO_2 production from six replicates) at 4°C were plotted against time.

2) Minimal processing of rambutan

The minimal process for rambutan in this part of the study, i.e. peeling with and without coring, was similar to the method described above, except different additive solutions were used. The samples were immersed in the solutions for different periods of time before being placed on polystyrene trays pre-sterilized with 70% ethanol. A water absorbent sheet was placed underneath the samples and the trays were sealed in plastic bags under various modified atmospheres

Table 1. Additive solutions and immersion times used for minimally processed rambutan.

Solution code	Additives used, % (w/w)	Immersion time, min
A	0.7% ascorbic acid + 0.75% alginate + 1.0% CaCl ₂	2
B	0.5% citric acid + 0.5% ascorbic acid + 2.0% sorbital + 0.2% CaCl ₂	5
C	0.5% citric acid + 0.5% ascorbic acid + 2.0% sorbital + 0.2% CaCl ₂ + 1.5% Ca-lactate + 0.1% pectin	5
D	0.5% citric acid + 0.5% ascorbic acid + 1.0% CaCl ₂	3
E	0.5% citric acid + 0.5% CaCl ₂	2

Table 2. Properties of three different types of plastic bags (250x250mm, 800 cc.).

Types	Thickness (μm)	Oxygen Transmission Rate (OTR, cc/m ² /d, at 23°C, 0% RH)
PET/LLDPE ¹ (PET)	52	128.18
CPP/LLDPE ² (CPP)	120	932.80
Nylon/LLDPE ³ (nylon)	80	36.60

Notes: ¹Polyethylene Terephthalate/linear low density polyethylene

²Cast-polypropylene/linear low-density polyethylene

³Nylon/ linear low-density polyethylene

Table 3. Experimental design for minimal processing of rambutan.

Experiment/ preparation style	Solution code / immersion time (min)	Type of plastic bag	Packaging condition	Storage temperature (°C)	Storage time (days)
1. Peeled & cored	A/2	nylon	Air	4.0±1 and 8.0±1	14
2. Peeled & cored	B/5	PET nylonCPP	Air	4.0±1	14
3. Peeled & cored	C/5	nylon	Air	8.0±1	7
4. Peeled	D/3	nylon	Air	8.0±1	10
5. Peeled	E/2	nylon	5% CO ₂ , 5% O ₂ , 90% N ₂	4.0±1	12

Note: Each experiment was done in duplicate.

and stored at either 4.0±1.0 or 8.0±1.0°C for up to two weeks. The following factors affecting quality changes in the product were investigated:

Preparation style: peeled vs. peeled and cored

Food additive: various solutions of food additives and immersion times are shown in Table 1.

Plastic bags: types of plastic bags used in this study are shown in Table 2.

Packaging condition: samples were packaged either in air (control) or modified atmosphere (50% flushing mode) with a gas mixture of 5% CO₂, 5% O₂ and 90% N₂ (by volume).

Storage temperature: either 4.0±1.0 or 8.0±1.0°C, with 76% RH.

2.1) Experimental Design

The experiment design for this study is presented in Table 3.

2.2) Quality measurement

Weight loss with respect to initial fresh weight in minimally processed rambutan during storage was determined. Color change was monitored using a colorimeter (Hunterlab, Model Colorflex, USA). pH was measured according to A.O.A.C. (2000) with a Sartorius pH meter (model PB-20, Germany). Total soluble solids of the samples were determined using a hand refractometer (Atago, model N1, Japan).

Microbiological quality of the samples was evaluated by taking duplicate 25 g samples from a sealed plastic bag and homogenizing them in sterile Stomacher bags with 225 ml of sterile buffer peptone water for 1 min at 200 beats per min. Appropriate dilutions of 0.1 ml homogenate were plated out in triplicate on total plate count agar and incubated at 30.0±1.0°C for 48 h (BAM, 2001).

2.3) Statistical analyses

All experiments were conducted in duplicate. Data were analyzed using a completely randomized design. Significant level was established at p≤0.05. Duncan's New Multiple Range

Table 4. Respiration rate (mg CO₂/kg/h) of whole, peeled, peeled and cored rambutan stored at 4°C.

Time(h)	Respiration rate (mg CO ₂ /kg/h) ¹		
	Whole rambutan	Peeled rambutan	Peeled and cored rambutan
1	146.90±54.36	156.05±41.43	174.40±59.78
2	110.16±24.61	156.06±41.43	165.25±53.61
3	110.16±31.07	123.95±13.12	142.29±50.53
4	128.52±45.32	142.29±35.20	169.85±67.75
5	119.35±26.58	123.93±41.93	110.15±18.05
6	114.75±34.71	105.57±17.82	96.40±12.20

Note: ¹ average respiration rate ± standard deviation from six replicates.

Test (DMRT) was used to determine significant difference between treatment means. SPSS for Window Version 10.5 was used for all statistical analyses.

3. Results and Discussion

3.1 Respiration rate

Table 4 shows the respiration rate (mg CO₂/kg/h) of whole, peeled, and peeled and cored rambutans over 6 h at 4°C. The average rates for whole intact fruits, peeled, peeled and cored rambutans were 122, 134, 143 mg CO₂/kg/h, respectively. After processing 1 h, the maximum respiration rates of whole, peeled, peeled and cored rambutans were observed, and subsequently decreased throughout the 6 h period. The response of tissues to processing wounds generally increases as the severity of the injury increases (Cantwell and Suslow, 2002; Del Aguila *et al.*, 2006). Damaging rambutan tissues through peeling and coring resulted in higher respiration rates as compared to that of whole fruits. In addition, peeling and coring severely injured the tissues more than peeling alone, resulting in the highest increase in respiration rate. Cantwell and Suslow (2002) reported that whole carrots produced 1.0 µg CO₂/kg/s at 0°C, while carrot slices produced 1.7 µg CO₂/kg/s at 0°C. Sliced carrots showed an increase of 62.2% in respiration rate at 0°C compared to whole carrots. Del Aguila *et al.* (2006) also reported that whole, sliced and shredded radish, after processing and storage at 5°C for 1 and 4 h, produced 4.5 and 2.4, 4.6 and 2.9, and 5.6 and 3.4 µg CO₂/kg/s, respectively. Furthermore, Saltveit (2003) hypothesized that the initial increase in respiration rate was due to the stress caused by cutting or the cellular repair mechanism. The subsequent decrease in respiration rate was probably due to severity of tissue damage from cutting contributing to juice leakage, less in respiratory substrates and faster deterioration (Saltveit, 2003).

3.2 Qualities of minimally processed rambutan

Figure 1(a) shows weight loss (%) of peeled and cored

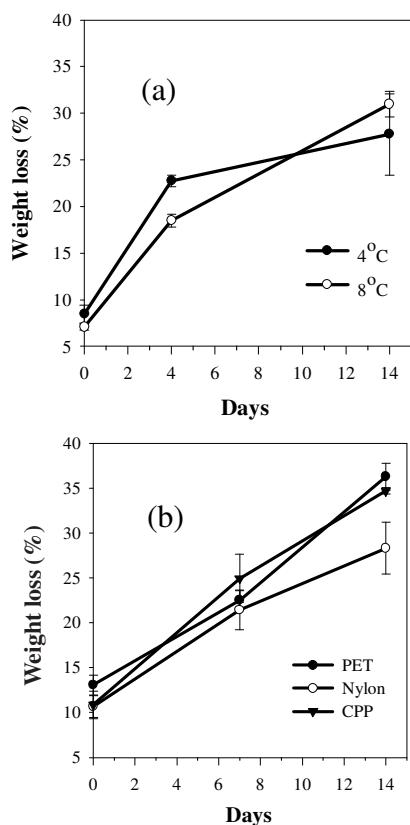


Figure 1. Weight loss (%) of peeled and cored rambutan soaked in solution A, packed in nylon bags and stored at 4°C and 8°C (a), and in solution B, packed in PET, nylon, CPP bags stored at 4°C (b). (Solution A = 0.7% ascorbic acid + 0.75% alginate + 1.0% CaCl₂ for 2 min, Solution B = 0.5% citric acid + 0.5% ascorbic acid + 2.0% sorbital + 0.2% CaCl₂ for 5 min)

rambutan immersed in solution A, packed in nylon bags, stored at 4.0±1.0 and 8.0±1.0°C for up to 14 days. Storage temperature appeared to significantly influence (P<0.05) on weight loss when stored for up to 4 days. Weight loss increased as the storage time increased in both at 4.0 and 8.0°C storage. At 14 days, samples stored at 8.0°C lost more weight than those stored at 4.0°C. Landrigan *et al.* (1996)

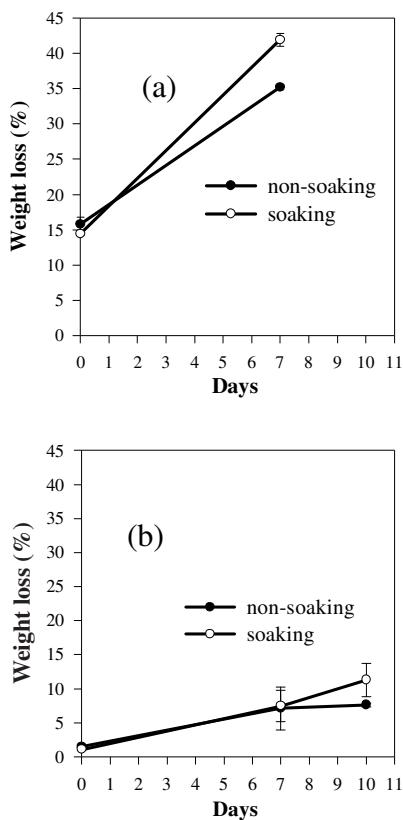


Figure 2. Weight loss (%) of peeled and cored rambutan with and without soaking in solution C packed in nylon bags and stored at 8°C (a), of peeled rambutan with and without soaking in solution D packed in nylon bags and stored at 8°C (b). (Solution C = 0.5% citric acid + 0.5% ascorbic acid + 2.0% sorbital + 0.2% CaCl₂ + 1.5% Ca-lactate + 0.1% pectin for 5 min, Solution D = 0.5% citric acid + 0.5% ascorbic acid + 1.0% CaCl₂ for 3 min)

studied weight loss in components of rambutan fruit stored under 20°C and 60%RH and reported that the flesh lost approximately 20% weight after 7 days of storage. Boonyaritthongchai (1999) also reported that rambutan fruits stored at 20°C lost more weight than those at 13°C. In the current research the storage temperature of 4°C was, therefore, selected for subsequent experiments.

Figure 1(b) shows weight loss (%) of peeled and cored rambutan immersed in solution B. After 14 days of storage, weight loss of samples packed in nylon bags was significantly less ($P<0.05$) than those packed in PET and CPP. Results showed that types of packaging bags with differences in OTR (cc/m²/d, at 23°C, 0%RH) values significantly influenced on weight loss of samples, especially at 14 days storage. The present study implied that under severe test conditions consisting of high temperature at 8°C and preparation style of peeling and coring, samples packed in the nylon bag with the OTR value of 36.60 cc/m²/d exhibited maintaining desirable quality in term of water loss compared to the other two bags. In addition, Kim et al. (2004) reported that film oxygen transmission rates significantly effected quality of fresh-cut salad savoy under the test conditions. Packages with optimum OTR films attained the desired O₂ and CO₂ levels throughout 25 days storage period at 5°C contributing to salad savoy remaining in freshness and acceptable quality. Salad savoy from packages with high in OTR value developed discoloration on the cut-ends due to high O₂ in the package. While the packages with low in OTR value exhibited a rapid depletion of O₂ and accumulated CO₂ in the packages, resulting in off-odor, decay and unacceptable quality.

Figure 2(a) shows weight loss (%) in peeled and cored rambutan with and without soaking in solution C. Results showed that after 7 days of storage, samples soaking in solution C had significant ($P<0.05$) greater in weight loss (41.9%).

Table 5. Quality changes of minimally processed peeled rambutan soaking in solution E¹, packed in nylon bags with and without modified atmosphere and stored at 4°C.

Treatment (4°C)	Storage time (days)	Quality changes ²				Total viable count (CFU/g)
		Weight loss (%)	°Brix	pH	L*	
Control	0	0.65±0.06d	16.67±1.20a	5.40±0.40a	47.08±2.16ab	< 250
	3	3.83±0.11bc	15.53±0.23a	4.94±0.05c	45.35±1.19c	
	6	4.92±0.50abc	14.53±0.50a	4.99±0.17bc	44.48±1.94c	
	9	5.90±0.89ab	15.60±0.28a	4.73±0.02d	46.77±0.55bc	
	12	6.95±0.11a	15.40±0.28a	4.80±0.16d	46.32±0.33c	4.5x10 ⁶
5% CO ₂ , 5% O ₂ , 90% N ₂	0	0.60±0.08d	16.10±1.01a	5.40±0.48a	48.20±2.59a	< 250
	3	2.85±0.88cd	15.80±0.53a	5.08±0.02bc	46.96±4.15a	
	6	5.96±2.72a	15.47±0.14a	5.07±0.19b	45.49±2.41bc	
	9	5.95±0.46ab	14.70±0.14a	4.99±0.02c	45.20±1.41c	
	12	7.54±0.43a	15.20±0.85a	4.96±0.06c	45.25±2.12c	6.4x10 ⁴

Note: ¹solution E = 0.5% citric acid + 0.5% CaCl₂ for 2 min.

²Mean±standard deviation of two replicates. Means within each column not sharing a common letter are statistically different ($P<0.05$)

than that of non-soaked ones (35.2%). Figure 2(b) shows weight loss in peeled rambutan with and without soaking in solution D. Similarly results showed that after 10 days of storage, sample soaking in solution D had significant ($P < 0.05$) greater in weight loss (11.3%) than that of non-soaked ones (7.6%). Poovaiah (1986) hypothesized that the major effect of calcium was maintaining the structure integrity of membranes and cell walls by forming cross-links between free carboxyl groups of the pectin chains. However, both peeled, peeled and cored samples exhibited severe leakage of water at the end of storage period, as a consequence of water loss, adverse effects on appearance changes remarkably occurred in soaked samples compared to non-soaked ones. As a result, extension of shelf life of peeled rambutan beyond 10 days appeared possible, as compared to that of peeled and cored samples. However, the use of suitable food additives and lowering the storage temperature may possibly aid longer extension of fresh-cut shelf life (Garcia and Barrett, 2002).

Table 5 shows quality changes in peeled rambutan, immersed in solution E, packed in nylon bags with and without modified atmosphere. The results demonstrated that MAP greatly influenced microbial counts of samples. After 12 days storage at $4.0 \pm 1.0^\circ\text{C}$, samples packed in the atmosphere of 5% CO_2 , 5% O_2 and 90% N_2 had lower microbial count (6.4×10^4 CFU/g) than that of control (4.5×10^6 CFU/g). Modified atmosphere did not significantly affect weight loss, total soluble solids content ($^\circ\text{Brix}$) and L^* values of the samples. pH of MAP and control samples after 12 days of storage, was significantly different ($P < 0.05$), being in the range of 4.73-5.40. Kader (2002) reported that the combination effects of gas composition in MAP (low O_2 level and high CO_2 level) and low storage temperature could delay aerobic microbial spoilage of fresh-cut products. Low level of O_2 suppressed the growth of aerobic microorganisms. In addition, Hu *et al.* (2007) reported that the effects of initial low 5% O_2 of fresh-cut cabbages packed in perforated film with stored at 5°C not only suppressing surface microbial growth but also respiration rates and the oxidation of ascorbic acid leading to obtaining better color and quality of samples. These also suggest that the use of chemicals such as acidulants and preservative may prolong better shelf life of peeled rambutan.

4. Conclusions

The respiration rate at 4°C of whole fruit, peeled, and peeled and cored rambutans can be used to indicate potential quality changes in rambutan tissues as a consequence of minimal processing. Processing method, packaging film, and storage temperature were important factors influencing the quality of minimally processed rambutans. To obtain longer shelf life of peeled rambutans, further study on food additives used and gas composition in MAP is needed.

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