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Quality Evaluation of Oranges Stored in Evaporative Coolers

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

Large quantities of oranges are wasted during storage due to its highly perishable nature. Research was therefore conducted on quality evaluation of orange fruits stored in aluminum-cladded (ABBEC) and non-cladded burnt-clay-brick evaporative coolers (NBBEC) to prevent postharvest loss. Essentially, the evaporative coolers comprised of double burnt-brick walls (1.29 × 2.55 × 2.56 m) external and (1.13 × 1.27 × 2.08 m) internal, (L × W × H) with wet sand bed in between and a storage space of (1.13 × 0.36 × 1.32 m), (L × W × H). Metabolic rates of oranges were highest at ambient storage, intermediate in NBBEC with the least value in ABBEC. Beta carotene, ascorbic acid and acidity decreased while total soluble solids, pH and microbial loads increased during storage. ABBEC storage resulted in 4.0 to 5.3°C decrease in ambient temperature with corresponding higher relative humidity over the non-cladded burnt-clay-brick coolers.

Keywords: Postharvest loss; evaporative coolers; ambient; relative humidity.

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1. INTRODUCTION

Citrus is produced globally with Brazil leading as the largest producer followed by USA and China [1]. Nigeria ranked 9th among the major producing countries and it produced 3% of the total world citrus output between the year 2000 and 2004 [2]. Sweet orange constitutes the most important proportion among the citrus varieties grown throughout the world, accounting for more than two-thirds of total global production [1]. The overall world production stood at 51.8million metric tons in 2013-14 with Brazil and China leading [3]. In 2018, citrus fruit production for Nigeria was 4.07 million tonnes [4]. According to [5], Benue State is Nigeria's largest producer of orange. However, [6] reported annual losses of about 30 to 60% during peak harvesting seasons due to the highly perishable, non-climacteric nature of the fruits and process technology. Additionally, there are no package and cold storage facilities for fruits and vegetables in Nigeria; transportation from farms to the market is also crude [2]. Wills et al. [7] indicated that fresh fruits and vegetables deteriorate very easily under tropical ambient conditions mainly due to physiological and microbial activities. Jain [8] observed that the prevailing high temperature in the tropics not only hastens physiological activities such as respiration, transpiration and ripening of fresh produce but also affects the physico-chemical composition eventually leading to spoilage. There is need for further research to reduce the postharvest losses encountered by orange farmers in Nigeria [9]. According to [10], evaporative cooling is the process by which the temperature of a substance is reduced due to the cooling effect from the evaporation of water. The concept of evaporative cooling is that the surrounding air serves as a heat sink where sensible heat is exchanged for latent heat of water. Cooling by means of evaporation therefore provides a low cost and effective way of preserving the freshness and prolonging the shelf life of fruits and vegetables as it reduces the temperature and increases the relative humidity inside the storage structure [11]. Vigyan et al. [12] reported that the evaporative cooler is ecofriendly, less energy requiring which improves the quality and productivity of fruits and vegetables by reducing field heat, increasing shelf life and ultimately reducing postharvest losses. Considering acute energy shortage and inadequate cold storage facilities in rural areas. there is tremendous scope for the adoption of low cost evaporative coolers for short-term, onfarm storage of perishable farm produce.

Evaporative coolers could lower temperature range of 10-15°C cooler than the outside temperature and maintain about 95% relative humidity [13].

Many authors such as [14,9,15,16,17]; reported the effectiveness of evaporative coolers for storage of oranges. Adekalu [18] also reported its effectiveness on storage of pre-treated matured sweet oranges. Adekanye et al. [14] evaluated performance of a prototype active evaporative cooling system for fruits and vegetable storage using oranges and tomato. Adekalu and Agboola [9] reported that sensory evaluation, marketability and acceptability of treated oranges were rated good at the end of 35 to 52 days' storage in evaporative coolers. Ubani and Okonkwo [15] reported that oranges stored for six weeks in evaporative cooler. The authors also observed that fruits in the evaporative cooler had weight loss of 4.2% and were in good and acceptable condition after storage.

Despite being the largest producer of oranges in the country, there is little or no research carried out in Benue State on the use of evaporative cooling in controlling postharvest losses of fruits and vegetables. The aim of this study therefore was to evaluate the storage of fresh oranges in evaporative coolers so as to prolong their shelf life and consequently reduce the postharvest loss of citrus in Nigeria.

2. MATERIALS AND METHODS

2.1 Study Area /Scope of Research

Makurdi is the capital of Benue State, Nigeria. The town is dominated by guinea savannah type of vegetation. The mean annual rainfall is favourable for food production. Makurdi has a sub-humid, semi-arid tropical climate with mean annual precipitation at 1200-1300 mm. About 90% of total annual rainfall occurs in the months of June to September [19]. Temperature rarely falls below 22°C with peaks of 40 and 30°C in February/March. In the wet season, the average temperature is within the range of 23.0-32.7°C. Data generated were the average for 2014 to 2017 for the evaporative coolers located beside the College of Food Technology Complex at the Makurdi University of Agriculture, (Latitude:07.78915° N, Longitude 008.61864° E).

2.2 Design and Construction of Evaporative Coolers

Two almost identical burnt-clay-bricks evaporative coolers were designed and

constructed adjacent and about 1 m apart under two trees. One had two internal aluminum claddings and was designated as aluminum cladded burnt-clay-brick evaporative cooler (ABBEC); the outer aluminum wall was perforated. The other cooler had no internal aluminum cladding and was referred to as native burnt brick evaporative cooler (NBBEC). The pictorial views of the cooling structures are shown in Plate 1. Essentially, the evaporative coolers consist of double jacketed rectangular burnt-clay-brick wall (1.29 × 2.55 × 2.56 m) external and (1.13 × 1.27 × 2.08 m) internal, (L × W × H) with wet sand bed in between and a storage space of (1.13 \times 0.36 \times 1.32 m), (L \times W × H). The cavity between the inner and outer walls of each cooler was filled with river-bed sand. The floors were cemented with mortar (cement, sand and water mixture) to an even 2 cm thickness. The doors to the storage spaces were made of white wood with zinc roofing sheet cladding for protection against rodents and termites. A make-shift thatched roof cover was built above each of the coolers to provide extra protection against direct sunlight in addition to the shade provided by the trees so that the fullest advantage of evaporative cooling could be

harnessed. In order to maintain the sand completely wet during the study, 500litres of water was used to wet the sand twice a day [20].

2.3 Commodity Storage Test

10 kg of ripe orange fruits (Ibadan sweet variety) were purchased from Makurdi Wurukum market and transported to the laboratory in jute bags. They were then washed with tap water to remove adhering sand and other foreign matter.

2.3.1 Weight loss

Weight loss was measured before and after storage using an electronic weighing balance (Model: Mettler P1210). Ten orange fruits were drawn at random on the 1st, 7th and 21st days of storage. Weight loss for each sample of known initial weight was calculated as follows:

$$PWL (\%) = (W_0 - W_t) / W_0 \times 100$$
 (1)

Where, PWL= physiological weight loss; W_o = initial weight of sample and W_t = weight of sample at time, t. The mean for the ten samples were then reported.



Plate 1. Evaporative Coolers 1 & 2

EC1= Non-Cladded Burnt-Clay-Brick Evaporative Cooler (NBBEC). EC2=Aluminum-Cladded Burnt-Clay Brick Evaporative Cooler (ABBEC)

2.3.2 Chemical analyses

Chemical analyses were performed according to the standard official methods described in [19]. Clear orange juice was extracted by pulping 100 g of edible portion in a household electric blender followed by straining using double-layered muslin cloth.

2.3.3 Moisture content

Moisture content was determined by weighing 5g of sample in crucibles whose weight have been determined. The crucibles and the samples were heated at 110°C in a Gallenkamp oven until constant weights were obtained. The dishes and their contents were cooled in a dessicator and then reweighed. The loss in weight was expressed in percentage:

$$\% Moisture = \frac{loss inweight on drying \times 100}{Initial sample weight}$$
 (2)

2.3.4 Ascorbic acid and total carotenoids

Ascorbic acid and carotenoids were determined by [21] methods. Ascorbic acid content was determined by titrimetric method with the titration of filtrate against 2,6-dichlorophenol indophenols and the result expressed as mg/100 g.

2.3.5 Total soluble solids (TSS)

TSS in degree brix was directly measured using Abbe refractometer (Model: Bellingham & Stanley Limited, England) by placing a drop of supernatant on the prism of refractometer.

2.3.6 pH and titratable acidity determination

The digital pH meter (Model pH 211, HI Hanna Instruments, Italy) was used to measure the pHof the orange juice while total titratable acidity (expressed as citric acid %) was determined by titrating 5ml of orange juice with 0.1N sodium hydroxide- using phenolphthalein as an indicator [21].

2.4 Microbiological Analysis

Samples for total plate counts and fungal counts were prepared as described by [22]. Triplicate 2 g portions of orange fruit were squeezed and homogenized in a Warring blender which was previously washed and sterilized with 100 ppm sodium hypochlorite solution and rinsed with sterile deionized water. Serial dilutions of

homogenate ranging from 10⁻¹ to 10⁻⁵ were obtained using sterile saline solution. Total aerobic plate counts and fungal counts were performed on nutrient agar and Saboraud dextrose agar respectively using the pour-plate method described by [23].

2.5 Sensory Evaluation

A consistent panel of 12 semi-trained judge was used to evaluate the appearance, texture and overall acceptability of orange fruit sample using the descriptive sensory profile developed based on perceptions of the judges for quality of fruits and vegetables. Sensory evaluation was conducted under fluorescent light in a special sensory testing room with partitioned booths. The degrees of preference based on the descriptive terms were then converted to scores with 7=very firm and 1=Putrid/mushy for texture, 7=very fresh and 1=extremely mouldy for appearance and 7=highly acceptable and 1=disgusting for overall acceptability [24].

2.6 Statistical Analysis

The results obtained were evaluated using the analysis of variance with the aid of Statisca 6.0 software package (Stafso, Inc. USA). The means of factors showing significant (p=0.5) differences were separated using Tukey's LSD test [25]. For this storage studies with orange fruits, the variables evaluated were influences of 3 storage times (1st, 7th and 21st days) and 3 storage conditions (Atmosphere, NBBEC and ABBEC).

3. RESULTS AND DISCUSSION

3.1 Physiological Loss in Weight

Fig. 1 shows the effect of storage condition on physiological weight changes of orange fruit. Physiological loss in weight is one of the main factors in determining the quality of stored fruits and vegetables. Minimal weight loss was recorded for oranges stored in aluminum burntclay-brick evaporative cooler (5.2%), 7.5% for those stored in NBBEC and maximal weight loss of 48.0% for oranges in ambient storage. These findings showed that oranges under ambient storage conditions shriveled fast and lost weight more quickly than those stored in evaporative coolers. The increase in weight loss with storage period may be due to the effect on reduction in moisture content on respiration. Oranges stored at ambient had wilted and shriveled after only 3 days of storage while those in evaporative

coolers remained fresh and appealing even after 21 days of storage. Ubani and Okonkwo [15] reported that oranges stored at ambient lost weight rapidly resulting in fresh weight loss of 31.1% after one month of storage while oranges stored in evaporative cooler had weight loss of 4.2%; were good and acceptable even after six weeks of storage. Similarly, [9] reported that treated oranges stored for between 35 and 52 days. They reported that weight losses of evaporatively stored pretreated oranges was below 5% unlike the control that was over 7%. According to [1], the reduction in weight could make orange fruits wrinkle and less firm and therefore may be less attractive to consumers if storage conditions are not modified. The authors observed that the decrease in fruit firmness at ambient condition could be attributed to fruit senescence due to poor storage condition. Nunes [26] also reported that reduced humidity during storage results in loss of moisture and orange dehydration, but may also lead to peel damage.

3.2 Chemical Analysis

3.2.1 Moisture content

Moisture content is an important quality feature that directly influences storability of fruits and vegetables. It is an index of water activity of many foods. The high moisture content of oranges (84.69%) in this study as shown in Table 1 implies that these fresh produce might be hiahlv perishable because spoilage microorganisms thrive in high moisture foods and is also indicative of low total solids. Similarly, the high moisture content provides for greater activity of water soluble enzymes and co-enzymes needed for metabolic activities of fresh produce [27].

Table 1. Moisture content of orange

Fruit	Moisture content (%)
Orange	84.69

3.2.2 Ascorbic acid and total carotenoids

The effect of storage condition on ascorbic acid content of orange fruit is shown in Fig. 2. The ascorbic acid values for orange juice decreased significantly during storage. The initial ascorbic acid content of fresh orange fruit was 37.72mg/100g which decreased to the lowest value of 22.20 mg/100 g in ambient storage, 23.35 mg/100 g in NBBEC storage and 26.83

mg/100 g in ABBEC storage. The differences in ascorbic acid content of orange fruits observed for all the storage duration and different storage conditions were quite significant (p<0.05). These values were slightly lower than those obtained by [28] which ranged from 23.01 to 42.50 mg/100 g and [29] who reported a similar value of 43.78 mg/100 g. According to Holcombe [30], ascorbic acid content of citrus fruit is never constant but varies with factors which include climatic/environmental conditions, maturity stage, handling and storage, ripening stage, species, and variety of the citrus fruit as well as temperature.

Beta carotene content of oranges decreased significantly from 2385.20 to 1158.50µg/100 g. After 21 days of storage, higher carotenoid content was recorded for oranges stored in ABBEC (2147.40 µg/100 g), intermediate in NBBEC (1167.20 μ g/100 g) while the minimum was recorded for oranges stored at room temperature (1158.50 µg/100 g). [31] reported a much lower value of 355 µg/100g for oranges. Rodriquez-Amaya [32] observed concentrations of beta carotene vary with the species and variety of orange and with growing, harvesting and storage conditions.

3.2.3 Total soluble solids of oranges

Fig. 3 presents the effect of storage condition on the total soluble solids of orange fruit in this study. The total soluble solids of fresh oranges before storage was 3.47°Brix which increased significantly to the highest value of 14.18°Brix in ambient, 12.70°Brix in NBBEC and the lowest value of 10.40°Brix in ABBEC storage condition. The increasing TSS could be due to increase in sugar content as acidity level in the fruit decreased in holding. Fruit palatability might increase as a result of high TSS to acidity ratio. The values reported by [28] were slightly lower and within the range of 5.50 to 11.80% while that of [29] was a high value of 33.88% as a result of the addition of sugar to the orange juice. According to AOAC [33], about 15% of the soluble constituents of orange juice are other than sugars and organic acids. This fraction consists of inorganic compounds, amino acids, ascorbic acids and small amounts of pectins, essential oils, esters, glucosides and other organic compounds. These compounds are commercially important in that the occurrence of off-flavours is either natural or processed orange juice is due to the oxidation and decomposition of these substances.

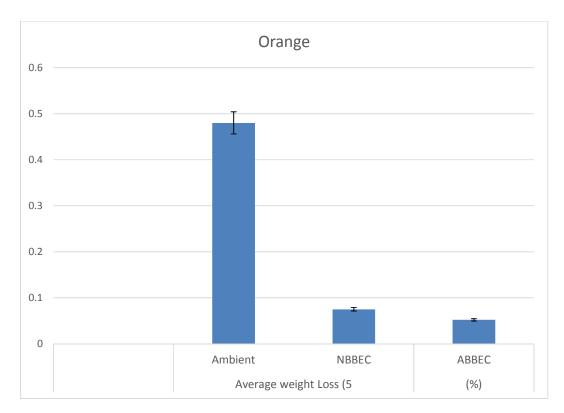


Fig. 1. Effect of storage condition on physiological weight changes of orange fruit

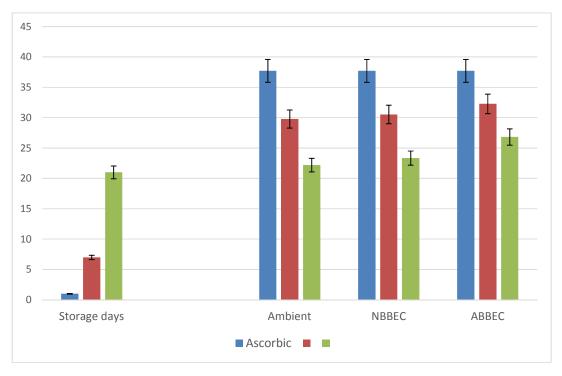


Fig. 2. Effect of storage condition on the ascorbic acid content of orange

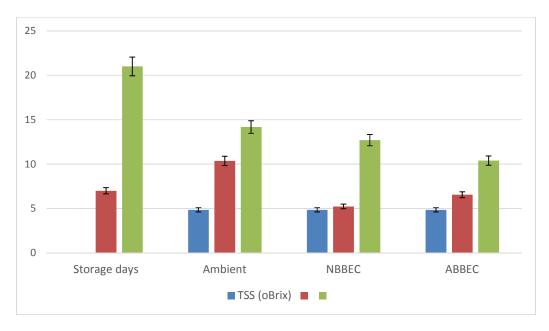


Fig. 3. Effect of storage conditions on the TSS of orange

3.2.4 pH and total titratable acidity

The pH of orange juice in this study showed a gradual increase from 3.36 to 3.80 (Table 2). Significant differences (p<0.05) existed in the level of pH during storage duration from day 1 to day 21. However, the differences in pH was not significant on day 1 and day 7 in NBBEC and ABBEC storage. This could be due to the relatively lower temperature and higher relative humidity exhibited by the evaporative coolers. This result was similar to that of [28] who reported pH values of 3.23 to 4.08. Other researchers such as [34,29] and [35] reported similar values of 3.4 to 4.6; 3.50 and 3.30 respectively. In fruits, the palate acidity depends on the hydrogen ion concentration which is affected by the degree of the acid. pH levels increased as sweet orange fruits were stored over a period of time which enhanced sweetness as the acidity level of fruits decreased due to increase in sugar content.

Changes in total titratable acidity were significantly affected by the rate of metabolism especially respiration which consumed organic acid (Table 2). However, the total titratable acidity of oranges were not significantly (p>0.05) different (0.97-1.25%). Ndife et al. [28] reported values of 0.4 to 1.06% while [29] and [35] reported 0.1% and 0.13% respectively. The gradual decline in TTA levels during storage

could be attributed to increased sugar substrates in the fruit which could be due to increased respiratory activity in orange fruit. Lower acid content is also known to improve fruit flavour.

3.3 Microbiological Analysis of Oranges

The results of microbial analysis of orange fruit samples are presented in Table 3. There were significant differences within the three storage conditions. There was no fungal load in evaporative coolers but ambient storage recorded 0.25Log₁₀cfu/g after 21st day of storage. However, significant differences existed in the fungal counts on days 7 and 21. [34] Recorded higher total plate count of 5 Log₁₀cfu/g and total fungal count of 2 to 2.85Log₁₀cfu/g while [36] reported highest volumes of contaminants ranging from 7.88 to 7.95Log₁₀cfu/g in their orange juice samples. Low temperatures in the evaporative coolers slowed down the plant's metabolic processes such as respiration, ethylene production and enzyme activity. This explains the lower microbial counts recorded in ABBEC storage due to the lower heat sinks of aluminum.

3.4 Sensory Evaluation of Oranges

The effect of storage condition on sensory scores of orange fruit is presented in Table 4. Panelists scored fresh orange sample 6.71 out of total

Table 2. Effect of Storage Conditions on pH and TTA of Orange

Parameter	Storage Time(Days)	Ambient	NBBEC	ABBEC	LSD
рН	0	3.36 ^a	3.36 ^a	3.36 ^a	
-	7	3.53 ^{cd}	3.39 ^a	3.36 ^a	
	21	3.80 ^d	3.43 ^c	3.37 ^c	0.43
TTA (%)	0	1.25 ^a	1.25 ^a	1.25 ^a	
` ,	7	1.08 ^c	1.15 ^{ab}	1.21 ^a	
	21	0.97 ^c	1.05 ^c	1.18 ^{ab}	0.74

Table 3. Effect of storage conditions on microbial load of orange

Microbial Parameter	Storage Time (Days)	Storage Conditions Ambient	Storage Conditions NBBEC	Storage Conditions ABBEC
Total Plate	0	0.31 ^b	0.31 ^b	0.31 ^b
Count(Log ₁₀ cfu/g)	7	0.33 ^a	0.33 ^a	0.32 ^{ab}
, J.,	21	1.35 ^a	2.18 ^d	0.32 ^{ab}
Yeast &	0	0.00^{c}	0.00^{c}	0.00^{c}
MouldCount(Log ₁₀ cfu/g)	7	0.23 ^b	0.22 ^b	0.21 ^b
(310 - 37	21	0.25 ^a	0.24 ^a	0.23 ^a

NBBEC= Non-cladded burnt-clay-brick evaporative cooler. ABBEC= Aluminum-cladded burnt-clay-brick evaporative cooler. Values for each parameter with common superscripts are not significantly (p>0.05) different

Table 4. Effect of storage conditions on sensory scores of orange

Sensory Attribute	Storage Time(Days)	Storage Conditions Ambient	Storage Conditions NBBEC	Storage Conditions ABBEC
Appearance	1	6.62 ^a	6.62 ^a	6.62 ^a
	7	5.15 ^b	6.20 ^{ab}	6.11 ^a
	21	3.67 ^c	5.69 ^b	5.82 ^c
Texture	1	6.71 ^a	6.71 ^a	6.71 ^a
	7	5.19 ^b	6.30 ^b	6.33 ^b
	21	3.68 ^d	5.69 ^c	5.82 ^{bc}
Overall	1	6.69 ^a	6.69 ^a	6.69 ^a
Acceptability	7	4.76 ^{bc}	5.79 ^b	6.16 ^a
	21	3.74 ^e	5.66 ^b	5.71 ^b

Values for each attribute with common superscripts are not significantly (p>0.05) different. Each result is the mean of 12 panelists responses on a scale with 7=excellent and 1=very poor. ABBEC=Aluminum-cladded burnt-clay-brick evaporative cooler. NBBEC= Non-cladded burnt-clay-brick evaporative cooler

value of 7.0 for texture which decreased significantly (p<0.05) to 3.68 in ambient storage, 5.69 and 5.82 in NBBEC and ABBEC storage respectively (Table 3).

Oranges stored in ABBEC were preferred because samples were judged superior organoleptically in quality than those stored in ambient, retaining more of its colour, flavour and viscosity. Idah et al. [37] in agreement with this result inferred that for consumers to obtain optimum nutrients from their products, oranges should not be stored beyond two weeks as the pot-in pot evaporative cooler storage system is concerned. Shrinkage rate for sweet orange in

ABBEC was very slow compared to those stored at ambient conditions.

4. CONCLUSION AND RECOMMENDA-TIONS

The evaporative coolers maintained a better quality in the physicochemical characteristics of orange fruit. In general, percentage weight loss, total soluble solids and pH increased with storage while beta carotene, total titratable acidity, and ascorbic acid decreased. The evaporative coolers gave an alternative approach to mechanical refrigeration for increasing the shelf life and maintaining the quality of oranges.

The ABBEC stored oranges exhibited lower biochemical and physiological reaction rates hence tissue breakdown, colour changes were lower in ABBEC than in NBBEC and ambient. The authors therefore recommend aluminum-cladded burnt-clay-brick evaporative coolers for on- farm storage of oranges.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

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