

Original Research Article

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Efficiency of Sweet Potato (*Ipomoea batatas* L.) Genotypes in Retention of Processing Qualities under Ambient Conditions

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ABSTRACT

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Sweet potato (*Ipomoea batatas* L.) is the sixth most important crop grown worldwide after wheat, rice, maize, potato and cassava and is considered as staple food in many developing countries. The largest quantity of sweet potato production is noticed in Asia and the Pacific Islands (93% of global production). However, sweet potatoes face the problem of weevils, shrinkage and loss of nutrients in storage. Therefore, present research on shelf-life and physicochemical parameters of 12 different sweet potato genotypes were studied in order to determine the varieties for better storage. Among those, the genotype BSP₁ was found to be good and exhibited superior qualities like high dry matter (61.7%), low values for TSS (4.4 °Brix), reducing and non reducing sugars (0.67 and 0.21%), starch (0.60%) and physiological loss in weight (13.5%) throughout the storage period. The genotypes Sree Bhadra and BSP₂₃ were the next best genotypes to maintain all the above mentioned characters.

Introduction

The sweet potato (*Ipomoea batatas* L.) is a semi-perishable commodity. Appropriate and efficient post harvest technology and marketing are critical to the entire production-consumption system of sweet potato because of its bulkiness and perishability. The purpose of storage is to maintain tubers in marketable condition and to provide a uniform flow of tubers to market and processing plants throughout the year. It must be realized that storage losses cannot be avoided even by optimal storage (Maldegem, 1999). Storage losses are mainly caused by the processes like respiration, weevil incidence, sprouting,

evaporation of water from the tubers, spread of diseases, changes in the chemical composition and physical properties of the tuber and damage by extreme temperatures.

The main objective of present study was to test efficiency of different sweet potato genotypes for retention of processing qualities under ambient storage conditions.

Materials and Methods

The present investigation was carried out in northern dry zone (Zone-3) of Karnataka state at 16°15' north latitude, 74°45' east longitudes and at an altitude of 612.05 m

above the mean sea level, in the laboratory Rashtriya Krishi Vikas Yojana (RKVY) research unit of the Department of Postharvest Technology, Kittur Rani Channamma College of Horticulture (University of Horticultural Sciences, Bagalkot), Arabhavi, Gokak Taluk and Belgaum district of Karnataka state during the period from 2016-17.

Selection and preparation of sweet potatoes for experiment

Representative even sized fresh sweet potatoes tuber of different varieties were procured from the research field of AICRP - Tuber crops, operating at Regional Horticulture Research and Extension Centre, Dharwad of Karnataka state.

Tubers were well matured and free from damage of pest and disease infestation. Procured sweet potatoes were washed under running tap water to remove adhered soil; damaged and infected tubers were discarded and good tubers were dried under shade.

Total soluble salts (°Brix)

The total soluble solids (TSS) of the sweet potato genotypes were estimated using the hand held refractometer on the different days of observation. A small amount of the flesh of tuber was crushed using mortar-pestle and its juice was obtained by filtering it using multiple layers of muslin cloth. The obtained clear juice was applied in drops on the prism of the calibrated refractometer and the values were read.

Beta-carotene content (mg/100g)

Beta-carotene present in sweet potato tubers was estimated by using petroleum ether method.

Dry matter content (%)

Dry matter content of sweet potato tubers was determined by drying the finely sliced piece of tuber in microwave oven (Onida Power Barbecue-28, MIRC Electronics Ltd., Mumbai) at 40 and 60 power intensity until the constant weight was achieved. The dry weight was calculated using the following formula.

$$\text{Dry matter content (\%)} = \frac{W_2}{W_1} \times 100$$

Where,

W_1 = Fresh weight of the tubers

W_2 = Dry weight of the tubers

Reducing and Non-reducing sugars (%)

The reducing and non reducing sugars present in sweet potato tuber genotypes were estimated using 3, 5-Dinitro Salicylic Acid (DNSA) method (Miller, 1972).

Starch (%)

Starch content in sweet potato tubers was estimated by anthrone reagent method. The sample was treated with 80 per cent alcohol to remove sugars and then starch was extracted with perchloric acid (52%).

In hot acidic medium starch was hydrolyzed to glucose and dehydrated to hydroxymethyl furfural, this compound forms a green colored product with anthrone (Bates *et al.*, 1943).

Ascorbic acid content (mg/100g)

Ascorbic acid content of sweet potato tubers was estimated by using the method given by AOAC (1990), which was based on the reduction of 2,6-dichlorophenol indophenols (2,6-DCPIP) by ascorbate.

Total titratable acidity (%)

The total titratable acidity of sweet potato was estimated by titrating the sample against strong base i.e. NaOH.

Physiological loss in weight (%)

The physiological loss in weight (PLW) was

$$\text{Physiological loss in weight (\%)} = \frac{\text{Initial weight} - \text{Final weight}}{\text{Initial weight}} \times 100$$

Results and Discussion

The observations on different parameters of sweet potatoes stored in ambient condition are discussed below.

Total soluble solids (°Brix)

In present investigation, the total sugars of sweet potato tubers increased continuously up to the end of the experiment (Table 1). Accumulation of total sugars could be attributed to the dormancy release and onset of sprouting as thereafter the cultivars showed sprouting, which continued to increase up to the end of experiment (Abdullah and Safraiy, 2015).

Initially the maximum value for TSS (7.1°Brix) was recorded genotype BSP₂₃. On 15 DAS, the genotype BSP₃ had maximum value (7.7°Brix) for TSS and on 30 DAS, maximum amount for TSS (7.8°Brix) was recorded in the genotype BSP₂. The change in sugar content during storage consequently increased the ratio of reducing sugars to non-reducing sugar. Observations made in this experiment are similar to the findings of Kumar *et al.*, (2002) and Huang *et al.*, (2014).

Beta-carotene (mg/100g)

Beta-carotene is sensitive to heat and oxygen as opined by Emmanuel *et al.*, (2010).

estimated at an interval of 15 days during storage. Initial tuber weight was recorded at the beginning of the storage period. The tubers were weighed and the weight was termed as final weight on the particular date of observation. The following formula was employed to calculate the PLW for each date of observation.

Initial weight

Oxidation of carotenoids results in the loss of beta-carotene in the fruits and vegetables as reported by Shekhar *et al.*, (2015). In the present investigation, the highest beta-carotene was recorded in the orange fleshed genotype BSP₂₃ on initial day, 15 DAS and 30 DAS (13.64, 13.00 and 12.46 mg/100g respectively).

Sree Bhadra was found to be next to BSP₂₃ by maintaining second highest content of beta-carotene. This variation occurred due to the difference in flesh colors which is a genotype dependent factor (Blessington *et al.*, 2010, Desai *et al.*, 2013).

Dry matter content (%)

The dry matter in the tubers increases during storage. However the amount of increase varies between genotypes (Elong *et al.*, 2014). In the present investigation, the highest dry matter was observed in the genotype BSP₂₃ on all the days of storage i.e. initially, 15 and 30 DAS (44.0, 46.2 and 48.1% respectively). These results are in conformity with study undertaken by Serge and Tom (1996).

Reducing and non-reducing sugars (%)

In an investigation by Kumar *et al.*, (2002) opined that the reducing sugars is a variety dependent factor. In the present study, the

genotype BSP₂₃ exhibited the maximum amount of reducing sugars (1.44, 1.37, 1.19% respectively) on first, 15th and 30th day. The minimum value for reducing sugars (0.35%) was recorded in BSP₁ at the end of experiment. It also recorded the maximum content of non-reducing sugars (0.42 and 0.59%) on 15 and 30 DAS.

Decrease in reducing sugars and increase in non-reducing sugars was due to utilization of reducing sugars for the metabolic processes in the tubers during storage and accumulation of non-reducing sugars due to sprouting (Ingabire and Hilda, 2011).

Pressey and Shaw (1996) also observed a decline in reducing sugars during storage at higher temperature with sweet potato. Similar results were cited by and Agbemafle *et al.*, (2014) in potato.

Starch (%)

Amount of starch in different varieties is a character which varies with genotype and different chemical constituents in the tubers as quoted by Zhang *et al.*, (2002). In the present investigation, the maximum amount of starch was recorded in BSP₂₃ (1.30%) initially and same genotype had maximum of 1.23 and 1.07 per cent starch respectively after 15 and 30 days of storage. Starch content of sweet potato tubers, which is indeed a varietal trait, slightly decreased during storage (Table 2). The decline in starch content was correlated with α -amylase activity in storage. There was a stronger positive and significant correlation between starch and reducing sugar content. The reduction in starch was due to the catabolic reactions in storage leading to conversion of complex starch molecules into simpler sugars. Similar reports have also been made by Khayatnezhad *et al.*, (2011).

Table.1 Performance of sweet potato genotypes in retaining TSS (°Brix), Beta-carotene (mg/100g) and Dry matter (%) during ambient storage

Genotypes	Total soluble solids (°Brix)			Beta-carotene (mg/100g)			Dry matter content (%)		
	DAS								
	0	15	30	0	15	30	0	15	30
BSP ₁	4.4	6.1	6.3	1.24	0.93	0.74	38.3	41.5	43.3
BSP ₂	6.1	7.7	7.8	1.65	1.34	1.19	33.4	38.1	41.7
BSP ₃	5.2	5.8	5.5	0.79	0.68	0.63	31.6	33.6	39.5
BSP ₄	4.8	6.1	6.3	0.25	0.21	0.16	29.6	35.7	40.2
BSP ₅	4.9	5.7	5.9	0.61	0.43	0.41	36.5	40.7	46.8
BSP ₆	6.2	7.0	7.3	0.31	0.30	0.25	26.8	34.4	40.2
BSP ₇	6.8	7.4	7.5	0.97	0.69	0.56	32.1	32.5	39.1
BSP ₈	5.9	6.4	6.6	1.49	1.14	1.09	38.2	40.3	42.8
BSP ₉	6.1	7.2	7.4	0.32	0.15	0.09	33.5	39.4	39.2
BSP ₁₀	6.2	6.6	6.9	0.32	0.22	0.09	37.0	40.3	40.1
BSP ₂₃	7.1	7.4	7.7	13.64	13.00	12.46	44.0	46.2	48.1
Sree Bhadra	5.8	6.1	6.6	2.95	2.61	2.40	32.1	34.1	37.9
Mean	5.8	6.6	6.8	2.05	1.81	1.67	34.4	38.1	41.6
S.Em±	0.11	0.09	0.12	0.03	0.04	0.05	0.40	0.33	0.17
C. D. @ 1%	0.42	0.35	0.47	0.11	0.14	0.20	1.57	1.30	0.69

DAS: days after storage

Table.2 Performance of sweet potato genotypes in retaining reducing sugars (%), non-reducing sugars (%) and starch content (%)

Genotypes	Reducing sugars (%)			Non-reducing sugars (%)			Starch (%)		
	DAS								
	0	15	30	0	15	30	0	15	30
BSP ₁	0.67	0.55	0.35	0.21	0.42	0.59	0.60	0.50	0.32
BSP ₂	0.80	0.76	0.72	0.25	0.29	0.35	0.72	0.69	0.65
BSP ₃	0.91	0.86	0.70	0.18	0.24	0.40	0.82	0.78	0.63
BSP ₄	0.68	0.64	0.48	0.13	0.21	0.47	0.61	0.57	0.43
BSP ₅	0.72	0.60	0.54	0.10	0.25	0.34	0.64	0.54	0.49
BSP ₆	0.61	0.57	0.51	0.11	0.22	0.34	0.55	0.51	0.46
BSP ₇	0.61	0.51	0.46	0.08	0.21	0.32	0.55	0.46	0.41
BSP ₈	0.60	0.54	0.52	0.22	0.34	0.40	0.54	0.49	0.47
BSP ₉	0.69	0.65	0.63	0.17	0.23	0.29	0.62	0.59	0.56
BSP ₁₀	0.63	0.62	0.54	0.19	0.28	0.39	0.57	0.55	0.49
BSP ₂₃	1.44	1.37	1.19	0.13	0.23	0.42	1.30	1.23	1.07
Sree Bhadra	0.68	0.65	0.62	0.29	0.35	0.42	0.61	0.59	0.55
Mean	0.75	0.69	0.60	0.17	0.27	0.39	0.68	0.62	0.54
S.Em±	0.01	0.01	0.03	0.03	0.02	0.04	0.01	0.01	0.03
C. D. @ 1%	0.04	0.05	0.13	0.10	0.07	0.14	0.05	0.04	0.11

DAS: Days after storage

Table.3 Performance of sweet potato genotypes in retaining ascorbic acid content (mg/100g), total titratable acidity (%) and physiological loss in weight (%)

Genotypes	Ascorbic acid (mg/100g)			Total titratable acidity (%)			Physiological loss in weight (%)		
	DAS								
	0	15	30	0	15	30	15	30	
BSP ₁	16.30	10.87	5.43	0.19	0.27	0.15	13.5	22.9	
BSP ₂	14.49	10.87	5.43	0.31	0.35	0.27	15.1	31.9	
BSP ₃	21.74	16.30	10.87	0.27	0.35	0.19	12.4	28.5	
BSP ₄	25.36	19.93	12.68	0.31	0.39	0.27	30.3	42.3	
BSP ₅	23.55	18.12	12.68	0.27	0.31	0.23	13.4	31.5	
BSP ₆	12.68	10.87	5.43	0.31	0.31	0.27	15.7	37.7	
BSP ₇	14.49	9.06	5.43	0.23	0.31	0.19	17.8	35.0	
BSP ₈	21.74	14.49	5.43	0.31	0.35	0.23	15.0	31.4	
BSP ₉	18.12	9.06	5.43	0.35	0.39	0.23	14.0	35.7	
BSP ₁₀	18.12	12.68	7.25	0.31	0.31	0.23	11.4	31.1	
BSP ₂₃	21.74	14.49	9.06	0.39	0.46	0.31	10.9	23.6	
Sree Bhadra	18.12	12.68	7.25	0.35	0.39	0.31	14.6	28.5	
Mean	18.87	13.28	7.70	0.30	0.35	0.24	15.3	31.7	
S.Em±	1.73	1.48	1.17	0.06	0.06	0.05	0.88	1.10	
C. D. @ 1%	6.86	5.85	4.63	NS	NS	NS	3.48	4.34	

DAS: Days after storage

Ascorbic acid (mg/100g) and total titratable acidity (%)

The genotypes differ in the amount of ascorbic acid content as stated by Singh *et al.*, (2005) and it decrease on storage in ambient condition. However in the present investigation, the ascorbic acid content in the tubers decreased in all the genotypes as the storage time progressed. The maximum amount of ascorbic acid was recorded in BSP₄ on initial and 15 DAS respectively (25.36 and 19.93 mg/100g). Further on 30 DAS, two genotypes BSP₄ and BSP₅ recorded maximum amount of ascorbic acid (12.68 mg/100g each). A significant decrease in ascorbic acid noticed, could be due to enzymatic loss of L-ascorbic acid where it is converted to 2-3-dioxy-L-gluconic acid (Mapson, 1970). These results were found to be in line with the investigations of Cruz-rus *et al.*, (2011) in strawberry and Brar *et al.*, (2013) in potato. The total titratable was found to vary non-significantly (Table 3).

Physiological loss in weight (%)

The PLW increases with progress in storage period however the values of PLW are a genotype dependent factor (Amoah *et al.*, 2011). In the present study, minimum PLW (10.9%) was recorded in BSP₂₃ on 15th day while on 30th day, minimum PLW among all genotypes was recorded in BSP₁ (22.9%). This difference in PLW might be due to the difference in respiration and transpiration rates in different genotypes and also increase in respiration rate as the storage prolonged (Mehta and Singh, 2002). These reports were found to be in line with the study by Huang *et al.*, (2014) in sweet potato.

As evident from the overall assessment on the results obtained, the genotype BSP₁ was found to perform best among all the genotypes in ambient storage condition with

its desirable characters like low TSS, low amount of reducing sugars, low PLW, low moisture content, high dry matter and so on. It was also able to retain its qualities throughout the storage period.

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