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Microbiological stability of chemically preserved apricot pulp

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Key words: Apricot pulp, Sodium Benzoate, Potassium Metabisulphite, Total fungal count, Total bacterial count

Abstract

Background: There exist different methods to preserve the nutrition, color and taste of the fruit pulp for prolonged availability. Bacterial and fungal growth greatly affect the texture and taste of the pulp, if stored for longer period of time. Evaluation of different chemical preservatives to check efficacy and effects on microbial culture growth holds prime importance. The efforts are made in the present investigation to analyze the effect of various concentrations of Sodium Benzoate and Potassium Metabisulphite as preservative on microbial quality of apricot pulp during storage.

Methods: The uniformly ripened Halman Apricot pulp was extracted and preserved by chemical preservatives such as Sodium Benzoate (SB), and Potassium Meta-Bisulphite (PMS) at different concentrations. The pulp was investigated for Microbiological parameters i.e. total bacterial count (TBC) and total fungal count (TFC). The inhibitory activity of chemical preservatives was tested periodically by simulating the industrial storage conditions for apricot pulp in the lab (30-42°C in the dark), for a duration of 60 days.

Results: Significant inhibition in total bacterial count (TBC) was observed in chemically preserved samples. Potassium Metabisulphite was found to be more effective and the highest inhibitory effects on bacterial growth in apricot samples were observed at a concentration of 250mg/250g and 125mg/250g. These were followed by Sodium Benzoate at concentrations of 250mg/250g and 125mg/250g.

Conclusion: This study confirms that the preservatives significantly reduced bacterial and fungal growth in apricot pulp during storage and the pulp was safe for two months without spoilage.

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153 | Advancements in Life Sciences Vol. 1, Issue 2

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Introduction

Apricot (Prunusarmeniaca), a member of Rosaceae family, is a stone fruit with an enlarged mesocarp of the ovary wall being the edible part. The pit or stone consists of endocarp whereas the exocarp constitutes the skin of the fruit. The true seed is found enclosed within the endocarp. The fruit is nearly smooth and s of yellow or orange color. The fruit is eaten raw as well as in dried and canned form [1]. World annual production of apricot is approximated to be 579,000 tones [2]. Apricot is cultivated almost all over the world but Turkey, Iran are the world's largest producers of apricot, accounting for 20% and 10% of world production. Other producers include Italy (5%), Pakistan (5%), Spain (4%), Syria (4%), and United States (3%) France and Morocco (3%) (ERS, 2004). Pakistan is the 4th largest apricot producer of the world with productions reaching 1, 29,652 tones and over an area of 13,758 hectares. Apricot grown in Pakistan is soft and juicy but it has a very short shelf life. Even cold storage cannot lengthen it much. In Pakistan apricot pulp is utilized for manufacturing different value added products.

Apricot fruit is not only consumed in fresh, frozen or dried form but also used for manufacturing pulp, juice, nectar, marmalade, jelly, extrusion products etc. Apricot kernels are also used in aroma perfumes, benzaldehyde cosmetics, oils, and active carbon [3].

Apricot is a perishable fruit having 3-5 days of storage life at optimum conditions and 2-4 weeks at cold storage. The storage life varies with variety. The short storage life of this fruit is due to short time period from commercial ripening to the degradation process characteristic like senescence [4,5].

Different methods are used to preserve apricot pulp but chemical preservation is considered to be the cheapest among numerous methods and is quite common. It is widely used in Pakistan as well. Chemical preservatives are used to prevent microbe caused food spoilage. Apricot pulp is most commonly spoiled by Bacterial and fungal attack which can be controlled by chemical preservatives.

Studies have shown that no single preservative is completely effective against all microorganisms [6]. Calcium (such as Calcium Chloride) is known to conserve the quality of fruits, prevent physiological disorders, reduce the rate of respiration, lessen the solubilization of pectic substance, and maintain the firmness and also to slow down the process of ripening [7]. However, Sodium benzoate (SB) and Potassium metabisulphite (PMS) are more commonly used for long term fruit storage as they have shown better antimicrobial activity [8-10]. SB has been found quite effective against food poisoning, growth and survival of some yeast strains and other spoilage organisms [11,12]. Growth of microorganisms can be inhibited using different concentrations of SB as higher concentrations of SB showed greater antimicrobial effect on different species of Aspergillus [13-15].

Besides having an inhibitory effect on microflora in the stored pulp, addition of preservatives (SB & PMS) may adversely influence its physico chemical characteristics and sensory profile [16]. Addition of preservatives has been known to influence physico chemical characteristics of mango pulp as an increase in acidity; TSS (Brix), reducing sugars and decrease in sucrose content was observed [17]. Codex Standards adopted in 2001 and 2006 have defined maximum levels for the use of these chemicals in fruit preparations including pulp, purees and fruit. These are 1000mg/kg SB as benzoic acid and 500mg/kg PMS as residual SO₂. However, careless use of these preservatives is not only a great threat to health and well-being of consumers, but is also well documented to be a primary cause for appearance of resistant the microorganisms thus leading to the occurrence of many emerging food borne diseases [18,19]. Therefore there is a need to optimize dosage levels of these preservatives in apricot pulp. Individual and synergistic effects of these preservatives on various quality attributes under local conditions should also be thoroughly investigated. Therefore a study was conducted in the Department of Agriculture and Food Karakorum international Technology university, Gilgit. Apricot is one of the major fruits in northern areas of Pakistan especially in Gilgit-Baltistan, which is important economically, nutritionally and health point of view. Due to its high moisture content, apricot pulp is highly perishable in nature and has shortest shelf life. So, efforts have been made in the present investigation to improve the shelf life of apricot pulp by using various safe preservatives and to optimize their concentrations as well.

Methods

Procurement of materials

A local apricot variety (Halman) was selected for the study. Ripened fruits were obtained from Shigar valley of Gilgit Baltistan, which were uniform in size, weight and color. The fruit was washed thoroughly with distilled water in a pre-heated tray, to remove unwanted entities like dust, dirt, pesticides residues and surface microflora. Two preservatives sodium benzoate (NaC₆H₅CO₂) (Merck 6290) and potassium metabisulphite (K₂S₂O₅) (Merck 106357) were purchased from dealers of local market.

Pulp extraction, Pasteurization, packaging and storage

After washing, the apricots were dried and processed immediately for pulp extraction. Extraction was done in an electric pulper where pulp was separated from the stones and the obtained pulp was pasteurized in a water bath at a temperature of 82oC for 30 minutes. (At this temperature it is possible to completely kill spore forming bacteria which are sensitive to acidity of apricot pulp, with no changes in physical and chemical attributes) After pasteurization chemical preservatives (SB and PMS) as per treatment combination presented in Table 1 were mixed with the pulp. The treated pulp samples (250g each) were then transferred to sterilized glass bottles and stored under ambient conditions (30-40°C) in the dark for a period of 60 days and assessed for microbial attributes at an interval of 20 days.

Analysis of microbiological parameters of apricot pulp

Total microbiological contamination of pulp samples was performed for total bacterial and fungal count of stored pulp samples. One gram sample was acquired from each treated apricot pulp sample by means of aseptic techniques, and placed onto Nutrient medium in autoclaved petri plates. Plates were later incubated (Memmert 100-Germany) for 24-48 hours at 37°C. The pulp samples were analyzed for microbiological quality by considering the total plate count as evaluation index as described by Diliello, 1982 [19]. PDA (potato dextrose agar) was used as media for TFC (Total Fungal Count) and MHA (Mullar Hinton Agar) was used for TBC (Total Bacterial Count) of chemically preserved apricot pulp samples. The experimentation was repeated three times and reported data represents mean values (CFU/g) of these dimensions.

Statistical analysis:

Data of present study were analyzed statistically, using analysis of variance [20]. Multiple Range test was applied to assess the difference between means [22]. Significance was defined at p \leq 0.05. Values are means of three experiments (SD±).

Treatment	Sodium Benzoate (mg/g)	Potassium Meta bisulfate (mg/g)
Τ0		
T1	500	
Τ2		500
Т3	250	
T4		250
T5	125	
Т6		125
Table 1. Trea	tment combinations	(mg/g) of various

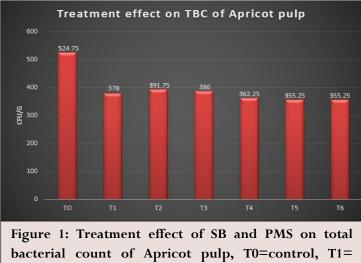
Table 1. Treatment combinations (mg/g) of various chemical

Results

Microbiological analysis of apricot pulp

Total bacterial count (TBC):

The present study was designed to investigate the effect of preservatives in prolonging the storage life of apricot pulp and to find out the optimum concentration required to effectively control fungal and bacterial contamination of stored pulp. For this purpose three different concentrations of each preservative i.e. 125 mg/g, 250 mg/g and 500 mg/g were analyzed.

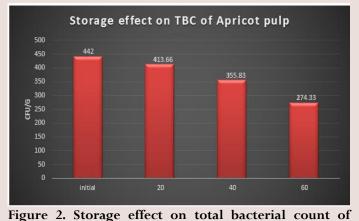


bacterial count of Apricot pulp, T0=control, T1= 500ppm SB, T2=500 ppm PMS, T3=250 ppm SB, T4=250 ppm PMS, T5=125 ppm SB, T6=125ppm PMS.

Controlled samples were also analyzed for comparison. The highest inhibitory effects on bacterial growth in apricot samples were observed by PMS at a concentration of 250mg/250g and 125mg/250g. These were followed by SB at 250mg/250g and 125mg/250g. However, none of the concentrations of the two preservatives used completely inhibited the bacterial growth for period of 60 days of storage. The control

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sample T0 (no preservatives added) showed highest level of contamination in apricot pulp samples after 60 days, whereas, minimum growth was observed in the presence of PMS at various concentrations.



Apricot pulp preserved with SB and PMS at various concentrations. The values are means of three experiments (SD \pm) (α =0.05).

Total fungal count (TFC):

TFC of microbial population is considered as an index of quality of food production. A decrease in fungal growth was observed in the presence of preservatives (PMS and SB).

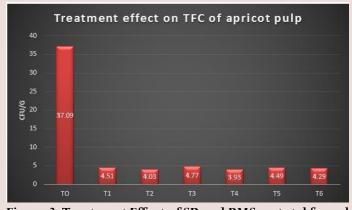
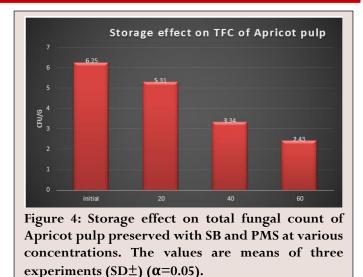


Figure 3. Treatment Effect of SB and PMS on total fungal count of stored Apricot pulp, T0=control, T1= 500ppm SB, T2=500 ppm PMS, T3=250 ppm SB, T4=250 ppm PMS, T5=125 ppm SB, T6=125ppm PMS. The values are means of three experiments (SD \pm) (α =0.05).



The maximum fungal growth was found in Controlled sample (T0) after 60 days of storage, whereas a considerable reduction was observed in chemically preserved samples. Figure 3 and 4 reveals treatment and storage effects of PMS and SB on fungal growth of apricot pulp. From the present study it can be confirmed that PMS and SB both at a concentration of 500mg/250g were most effective in inhibiting fungal growth in apricot pulp.

Discussion

Chemically preserved apricot samples for TBC were periodically analyzed and a progressive decrease in the growth was observed. Figure 1 and 2 reveals treatment and storage effects of PMS and SB on the bacterial growth of apricot pulp. The decrease in bacterial growth was due to the presence of PMS which is considered to be relatively best inhibitor in apricot pulp. These studies were found to be in accord with previous studies of Akhter et al., Hashmi *et al.*, and Hussain *et al.*, on inhibitory effects of PMS [16, 23, 17]. It is more effective in controlling growth of microorganisms in

stored pulp samples. 22 Similar results regarding microbial quality of apricot pulp have also been reported by Sakhale *et al.*, and Hussain *et al.*, [25, 17].

In our study decrease in fungal growth was observed in the presence of preservatives (PMS and SB). A previous study on inhibitory effect of preservatives has already shown the reduction of fungal growth in jam samples due to the presence of preservatives [24]. Awan and Rehman recommended use of 0.1% sodium benzoate in apricot and apple jam for fungal growth. Our results are in complete agreement with Ayub et al., that showed that samples added with 20% sucrose and potassium metabisulphite had overall best results in controlling the microbial population [28]. Brenndor et al., have also reported that addition of sulpher dioxide reduces the microbial population in fruit products [24].

This study reveals the inhibitory effects of Sodium Benzoate (SB) and potassium metabisulphite (PMS) on microbial growth in the apricot pulp stored under ambient temperature. These preservatives may retain quality attributes especially protein and ascorbic acid contents without any spoilage. Addition of the two preservatives significantly helped in controlling microbial and fungal growth of apricot pulp. The Effect of these chemical preservatives on TBC and TFC of stored apricot pulp, as shown by the present research. may facilitate the development of a safer and feasible storage of apricot pulp at industrial level. We found potassium metabisulphite (PMS) to be more effective in controlling growth of microorganisms in apricot pulp during storage. Both SB and PMS at different concentrations inhibit microbial and fungal growth in apricot pulp and prolong the storage life up to two months without any spoilage. The added preservatives may be also helpful to retain physical, chemical and organoleptic attributes of other value added products of apricot pulp like wine, jams, and squashes etc.

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