

Original Research Article

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Effect of Pretreatments and Drying on the Quality of Oyster Mushrooms

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ABSTRACT

Keywords

Oyster mushrooms, Pretreatments, Cabinet drying, Moisture content, Water activity, Dehydration ratio, Rehydration ratio, Color, Browning index, Microbial count, Sensory evaluation

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Mushroom slices were subjected to different pretreatments (0.1% KMS, 0.2% KMS, 1% CaCl₂, 2% CaCl₂, 0.5% glycerol, 1% glycerol) for 15 minutes and then dried in cabinet tray dryer at 50°C with a purpose to enhance the quality and for improved drying. Mushrooms pretreated with 1% glycerol were best on the basis of lowest moisture content (8.30%), water activity (0.44), browning index (0.04), microbial count (1.32 c.f.u/g) and highest dehydration ratio (9.84), rehydration ratio (3.85), L* value (65.08) and crude fat (1.88%). Also, on the basis of organoleptic acceptability, T₇ (1% Glycerol) was recorded as the best pretreatment for maintaining the quality of mushrooms.

Introduction

Mushrooms assume considerable importance in the human diet as they are rich in non-starchy carbohydrates, dietary fibre, minerals, vitamin-B and low in fat content. In India, there are mainly three species of mushrooms, namely, white button mushroom (*Agaricus bisporus*), oyster mushrooms (*Pleurotus sajor caju*) and paddy straw mushrooms (*Volvariella volvacea*), that are grown commercially. Among these, oyster mushrooms (*Pleurotus sajor caju*) possess unique nutritional and medicinal values, characteristic aroma and taste. Oyster mushrooms are suitable for cultivation in plains and even in hilly areas. They contain

89.8% moisture, 2.9% protein, 0.36% fat, 5.3% carbohydrates, 1% fiber, 1% ash and 33.9 Kilocalories of energy. Oyster mushrooms are perishable, because they have a high respiration and transpiration rates leading to rapid post-harvest deterioration. Browning reactions and dehydration affect the shelf-life of fresh oyster mushrooms within a few days, even if refrigerated. Pre-treatment is necessary to check discolouration during mushroom processing (Mudahar and Bains, 1982 and Pruthi *et al.*, 1984). Pre-treatments of mushrooms before drying in one form or other viz., washing in water, potassium metabisulphite (KMS), sugar, salt either alone or in combination help in checking enzymatic browning, stabilizing colour, enhancing

flavour retention and maintaining textural properties (Singh *et al.*, 2001). Sodium hypochlorite, sodium metabisulphite and glycerol are effective in reducing the browning of dried mushrooms. Addition of glycerol also improves the texture of rehydrated dried mushrooms while CaCl₂ produces a firm textured product. Drying is a method of preservation in which the water activity of the food is reduced. Traditionally mushrooms are dried under open sun, which results in unhygienic and poor quality products (Chua *et al.*, 2001). Due to long drying time and overheating of surface during sun drying, the problems of darkening in colour, loss in flavour and decrease in rehydration ability occur. Solar drying can be considered as an elaboration of sun drying and it is an efficient system of utilizing solar energy (Mulhbauer, 1986 and Bala, 1997).

Conventional hot-air drying is one of the most frequently used methods for mushroom dehydration, which involves thermal or chemical pretreatment and drying at temperatures between 50 and 80°C. In order to avoid darkening of the mushroom surface during hot-air drying, a two phase drying process is employed starting with 30°C or 40°C followed by a final temperature of 60°C. Due to low costs of initial investment, operation and ease of controlling the process, convective drying is the most commonly used method in the food industry (Mundada *et al.*, 2010 and Hiranvarachat *et al.*, 2011). The dehydrated product offers, apart from increased shelf life, the advantage of decreased mass and volume which have the potential for savings in the cost of packaging, handling, storage and transport of the product (Amuthan *et al.*, 1999; Karimi, 2010). Therefore, dehydration combined with some pretreatments appear to be a cost effective method of preservation (Rama and Jacob, 2000) for Indian conditions as dehydrated mushrooms are easy to transport as compared

to canned, pickled and frozen products (Chandra and Samsher, 2002).

Materials and Methods

Mushrooms were purchased from M/S Romesh Chander and Sons, Fresh Vegetable and Mushroom Shop, Parade, Jammu. They were washed with tap water and then kept on blotting paper to remove surface moisture. The research was conducted in the department of Food Science and Technology, SKUAST-J. Mushrooms were cut into slices of 1 cm wide by 1 cm long for the stipe, while 1.5 cm by 3 cm long for the cap and subjected to following pretreatments and then dried in cabinet dryer at temperature 50°C.

Moisture content

Ten grams of mushroom were dried in hot air oven at 70°C in pre-weighed dishes till constant weight. The dish with dried sample was transferred to desiccators and cooled to room temperature. The dish was then weighed and moisture content in percent was calculated from loss in weight (AOAC, 2002).

$$\text{Percent (\%) moisture} = \frac{\text{Loss in weight (g)}}{\text{Weight of sample (g)}} \times 100$$

Dehydration ratio

Dehydration ratio was calculated by taking the weights of sample before drying and the weight of sample after drying.

$$\text{Dehydration ratio} = \frac{\text{Weight of sample before drying}}{\text{Weight of sample after drying}}$$

Rehydration ratio

The rehydration ratio of dried mushroom flakes was determined by soaking samples

with a defined weight (approx. 5 g) in boiling distilled water at 95°C for 20 minutes. The samples were removed, filtered, dried and weighed. In order to minimize the leaching losses, water bath was used for maintaining the defined temperature (Ranganna, 1986). Rehydration ratio (RR) of the samples was computed as follows:

$$\text{Rehydration ratio} = \frac{M_r}{M_d}$$

Where,

M_r = Mass of rehydrated sample, g;

M_d = Mass of dehydrated sample, g

Water activity

Water activity was estimated using Aqualab water activity meter (Model Series: 3TE).

Color

The color was evaluated by measuring L^* , a^* , b^* parameters by means of Hunter lab colorimeter.

The instrument was standardized against white tile before the measurements. Color was expressed in CIE-Lab parameters as L^* (whiteness / darkness), a^* (redness / greenness), and b^* (yellowness / blueness) (Byrnes and O Beirne, 2008).

Browning index

The degree of non-enzymatic browning of the dried mushrooms was determined following the method of Mudahar and Bains (1982).

The color was extracted from dried mushroom using 60% ethanol, and the absorbance of the filtrate was measured using a spectrophotometer at 440 nm.

Crude fat

Five gram of dried sample was extracted with petroleum ether at 120°C in Soxhlet extraction apparatus for six hours. Ether extract was filtered in pre-weighed beakers. The petroleum ether was completely evaporated from the beakers and the increase in weight of the beaker represented the fat content (AOAC, 2002) and was calculated as below:

$$\text{Per cent fat} = \frac{\text{Weight of fat (g)}}{\text{Weight of sample (g)}} \times 100$$

Microbial count

Spread plate technique described by Palczar and Chan (1997) was used. 1g of each sample was aseptically transferred to 9 ml of sterile water in a separate tube and mixed vigorously. 1 ml of the resulting mixture was transferred to 9 ml of sterile water in a separate tube. The process was continued till 6th dilution (10^{-6}). Nutrient agar (NA) was inoculated with a 0.1 ml of appropriately diluted sample (10^{-6}) by spread plating technique and incubated at 37°C for 24 hours. Colonies were counted and multiplied by the dilution factor.

$$\text{Microbial load (cfu/g)} = \frac{N \times 1 \times D}{V}$$

Where,

N = Numbers of colonies counted

V = Volume of inoculums

D = Dilution factor

Sensory evaluation

Sensory evaluation depends upon the responses given by different sense organs. The samples were evaluated on the basis of appearance, flavour, texture, taste and overall

acceptability by semi-trained panel of 9-10 judges by using 9 point hedonic scale assigning scores from 9 (like extremely) to 1 (dislike extremely). A score of 5.5 and above was considered acceptable (Amerine *et al.*, 1965).

Statistical analysis

The results obtained were statistically analyzed using completely randomized design (CRD) for interpretation of results through analysis of variance.

Results and Discussion

Moisture content and water activity

Moisture content of pretreated and dried mushrooms ranged from 8.30 to 8.83 per cent (Table 1). The percent moisture content was highest in T₁ (Control) with mean value of 8.83 per cent while mushrooms treated with 1% glycerol (T₇) had lowest (8.30 per cent) moisture content. There was significant difference among various treatments. Among various pretreatments, lower moisture content was observed in 1% glycerol pretreated and cabinet dried mushrooms. These results are in agreement with the findings of Manalo and Benedicto (2014) who reported decrease in moisture content with increase in glycerol concentration in RTE dried tocino. The water activity ranged from 0.44 to 0.59 (Table 1). The highest water activity was recorded in T₁ (Control) with mean value of 0.59 followed by T₄ (1% CaCl₂) and T₅ (2% CaCl₂) with mean values of 0.55 and 0.53 while the lowest water activity was recorded in T₇ (1% glycerol) and T₆ (0.5% glycerol) with mean values of 0.44 and 0.46. Lowest water activity was observed in 1% glycerol pretreated and cabinet dried mushrooms. Salim *et al.*, (2016) reported lowest water activity in cabinet dried pear at 60°C and highest in sun dried pear slices. Samples treated with glycerol reached lower

water activity values than those treated with glucose and sucrose. This can be due to glycerol, a polyhydric alcohol which decreased the a_w by means of hydrogen binding with it. Panwar *et al.*, (2013) reported the similar results in aonla segments. Manalo and Benedicto (2014) also reported decrease in water activity with increase in glycerol concentration in RTE dried tocino.

Dehydration and rehydration ratio

The dehydration ratio ranged from 9.38 to 9.84 and there was significant difference among various treatments (Table 1).

The highest dehydration ratio was recorded in mushrooms treated with 1% glycerol (T₇) with mean value of 9.84 and lowest (9.38) in T₁ (Control) whereas the highest rehydration ratio was recorded in mushrooms treated with 1% glycerol (T₇) with mean value of 3.85 and lowest in T₁ (Control) with mean value of 2.91. Dehydration and rehydration ratio of pretreated and dried oyster mushrooms were more as compared to untreated mushrooms (control). Similar findings were reported by Doymaz (2014) regarding drying kinetics and rehydration characteristics of convective hot-air dried white button mushroom slices.

Color and browning index

Color is an important quality parameter of mushroom. The L* value was highest in T₇ (1% glycerol) with mean value of 65.08 while it was lowest in T₁ (Control) with mean value of 45.81 (Table 2). The highest a* (6.42) and b* (26.78) values were recorded in T₁ (Control) whereas the lowest a* (3.86) and b* (19.36) values were recorded in T₇ (1% glycerol). Oyster mushrooms pretreated with 1% glycerol showed best color as there was highest L* value and lowest a* and b* values as compared to control. Similar results were observed by (Mohamed and Hoo, 1994).

Table.1 Effect of pretreatments on moisture content (%), water activity (a_w), dehydration ratio and rehydration ratio of dried oyster mushrooms

| Pre-treatments | Moisture content (%) | Water activity | Dehydration ratio | Rehydration ratio |
|--|----------------------|----------------|-------------------|-------------------|
| T ₁ (Control) | 8.83 | 0.59 | 9.38 | 2.91 |
| T ₂ (0.1% KMS) | 8.65 | 0.51 | 9.60 | 3.73 |
| T ₃ (0.2% KMS) | 8.50 | 0.48 | 9.62 | 3.79 |
| T ₄ (1% CaCl ₂) | 8.72 | 0.55 | 9.44 | 3.66 |
| T ₅ (2% CaCl ₂) | 8.70 | 0.53 | 9.47 | 3.70 |
| T ₆ (0.5% Glycerol) | 8.40 | 0.46 | 9.81 | 3.78 |
| T ₇ (1% Glycerol) | 8.30 | 0.44 | 9.84 | 3.85 |
| Mean | 8.58 | 0.51 | 9.59 | 3.63 |
| CD _(0.05) | 0.09 | 0.08 | 0.08 | 0.08 |
| ±S.E.(m) | 0.03 | 0.02 | 0.02 | 0.02 |

Table.2 Effect of pretreatments on color and browning index of dried oyster mushrooms

| Pre-treatments | Color | | | Browning index |
|--|--------------|-------------|--------------|----------------|
| | L* | a* | b* | |
| T ₁ (Control) | 45.81 | 6.42 | 26.78 | 0.98 |
| T ₂ (0.1% KMS) | 58.20 | 5.24 | 24.39 | 0.14 |
| T ₃ (0.2% KMS) | 58.36 | 5.18 | 24.18 | 0.13 |
| T ₄ (1% CaCl ₂) | 62.29 | 4.28 | 20.42 | 0.05 |
| T ₅ (2% CaCl ₂) | 63.11 | 4.15 | 20.19 | 0.04 |
| T ₆ (0.5% Glycerol) | 64.24 | 3.98 | 19.62 | 0.06 |
| T ₇ (1% Glycerol) | 65.08 | 3.86 | 19.36 | 0.04 |
| Mean | 59.58 | 4.73 | 22.13 | 0.20 |
| CD _(0.05) | 0.12 | 0.11 | 0.12 | 0.08 |
| ±S.E.(m) | 0.04 | 0.04 | 0.04 | 0.03 |

L* (Whiteness/darkness)

a* (redness/greenness)

b* (yellowness/blueness)

Table.3 Effect of pretreatments on crude fat (%) and microbial count (cfu/g) of dried oyster mushrooms

| Pre-treatments | Crude fat (%) | Microbial count ($\times 10^2$ cfu/g) |
|--|---------------|--|
| T ₁ (Control) | 1.31 | 1.93 |
| T ₂ (0.1% KMS) | 1.77 | 1.47 |
| T ₃ (0.2% KMS) | 1.79 | 1.42 |
| T ₄ (1% CaCl ₂) | 1.73 | 1.54 |
| T ₅ (2% CaCl ₂) | 1.76 | 1.51 |
| T ₆ (0.5% Glycerol) | 1.82 | 1.36 |
| T ₇ (1% Glycerol) | 1.88 | 1.32 |
| Mean | 1.72 | 1.51 |
| CD _(0.05) | 0.08 | 0.09 |
| ±S.E.(m) | 0.03 | 0.03 |

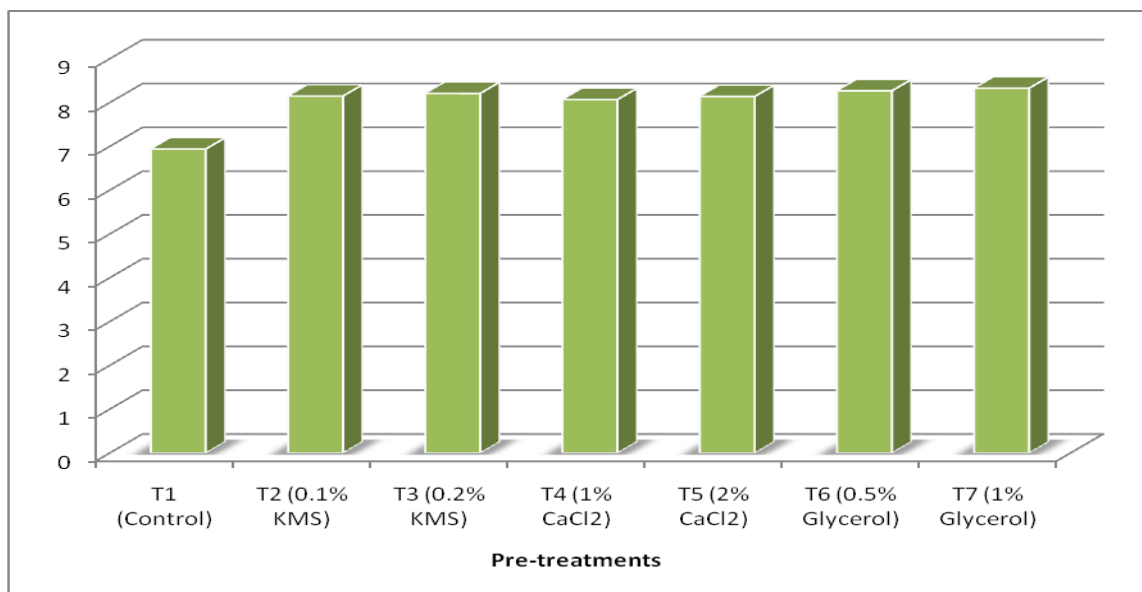
Table.4 Effect of pretreatments on appearance, flavour, texture, taste and overall acceptability of dried oyster mushrooms

| Pre-treatments | Appearance | Flavour | Texture | Taste | Overall acceptability |
|--|-------------|-------------|-------------|-------------|-----------------------|
| T ₁ (Control) | 7.08 | 6.84 | 6.96 | 6.81 | 6.94 |
| T ₂ (0.1% KMS) | 8.43 | 8.05 | 8.15 | 8.03 | 8.15 |
| T ₃ (0.2% KMS) | 8.47 | 8.10 | 8.18 | 8.10 | 8.21 |
| T ₄ (1% CaCl ₂) | 8.40 | 7.90 | 8.22 | 7.91 | 8.07 |
| T ₅ (2% CaCl ₂) | 8.44 | 7.97 | 8.28 | 7.96 | 8.14 |
| T ₆ (0.5% Glycerol) | 8.56 | 8.19 | 8.35 | 8.21 | 8.27 |
| T ₇ (1% Glycerol) | 8.62 | 8.25 | 8.44 | 8.27 | 8.33 |
| Mean | 8.28 | 7.90 | 8.08 | 7.89 | 8.01 |
| CD _(0.05) | 0.07 | 0.07 | 0.08 | 0.07 | 0.07 |
| ±S.E.(m) | 0.02 | 0.02 | 0.03 | 0.02 | 0.02 |

Treatment details

| Pre-treatments | Detail |
|----------------|----------------------|
| T ₁ | Control |
| T ₂ | 0.1% KMS |
| T ₃ | 0.2% KMS |
| T ₄ | 1% CaCl ₂ |
| T ₅ | 2% CaCl ₂ |
| T ₆ | 0.5% Glycerol |
| T ₇ | 1.0% Glycerol |

Fig.1 Effect of pretreatments on overall acceptability of dried oyster mushrooms



Farahnaky *et al.*, (2013) reported that L value increased, a and b values decreased with increase in glycerol concentration as compared to control in wheat starch edible films. The browning index of mushrooms is related to change in color of mushrooms. As the L* value decreased and a* and b* value increased, the browning index increased (Table 2). The highest browning index was observed in T₁ (Control) with mean value of 0.98. The browning index was recorded to be lowest (0.04) in T₅ (2% CaCl₂) and it was at par with T₇ (1% glycerol) with mean value of 0.04. Chakraborty *et al.*, (2014) also observed lower browning index in glycerol monostearate treated samples as compared to samples not containing glycerol monostearate regarding the study of functional properties based statistical optimization of foam mat drying parameters for potato (Kufri Chandramukhi).

Crude fat and microbial count

The crude fat of various pretreatments ranged from 1.31 to 1.88 per cent (Table 3). The crude fat was highest (1.88 per cent) in mushrooms treated with 1% glycerol (T₇) and it was at par with T₆ (0.5% glycerol) with mean value of 1.82 per cent while it was lowest (1.31 per cent) in T₁ (Control). Crude fat contents were more in pretreated oyster mushrooms as compared to untreated mushrooms. Similar findings were reported by Dunkwal *et al.*, (2007) regarding physico-chemical properties and sensory evaluation of *Pleurotus sajor caju* powder as influenced by pre-treatments and drying methods and also Ibrahim *et al.*, (2017) in oyster mushrooms. The highest (1.93×10^2 cfu/g) microbial count (Table 3) was recorded in T₁ (Control) while it was lowest (1.32×10^2 cfu/g) in T₇ (1% glycerol) and it was at par with T₆ (0.5% glycerol) with mean value of (1.36×10^2 cfu/g). Microbial count in pretreated and dried oyster mushrooms was low as compared to untreated

mushrooms (control). Similar results were observed by Ibrahim *et al.*, (2017) regarding effect of pre-treatments and drying methods on the chemical quality and microbial density of wild edible oyster mushroom.

Sensory evaluation

Sensory evaluation of pretreated and dried mushrooms was done using 9-point hedonic rating scale. Mushrooms treated with various chemicals attained more score for overall acceptability as compared to control (Table 4). The highest overall acceptability rating was recorded in T₇ (1% glycerol) with mean value of 8.33 while it was lowest in T₁ (Control) with mean value of 6.94. Pretreated dried oyster mushrooms scored higher sensory scores than untreated mushrooms in terms of color, flavour, texture, taste and overall acceptability (Fig. 1).

Among various pretreatments, mushroom samples pretreated with 1% glycerol and cabinet dried were adjudged best based on overall sensory score. Glycerol treated samples had a better color as they showed higher L* value but relatively lower a* and b* values. Glycerol pretreatment improved the texture score of dried mushrooms. Glycerol treated mushroom samples showed a softer texture than CaCl₂ and KMS pretreated mushrooms because it function as a solvent which replaces water in dried product. Glycerol has also been reported to reduce shrinkage during drying, maintain texture and improve the ability to rehydrate to a level of fresh mushrooms. Similar findings were observed by (Mohamed and Hoo, 1994). These results are in agreement with the findings of Srivastava and Bala (2016) also reported that 1% gum-arabic associated with 1% carboxymethyl cellulose and 1% glycerol coating was effective in increasing the shelf life of button mushrooms. Kim *et al.*, (2013) also reported that organoleptic scores were

higher in glycerol pretreated and hot air dried jujube fruits regarding effect of pretreatment and drying methods on quality and antioxidant activities of dried jujube fruits (*Zizyphus jujuba*).

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