NUTRITIVE AND BACTERIOLOGICAL ANALYSIS OF DIFFERENT TYPES OF CHEESE

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ABSTRACT

In the present investigation five cheese samples were collected and their pH and moisture contents were determined which was found to vary from 6.0 to 6.5 and 35.0-40.0% respectively. Major nutritional factors protein (1N NaOH soluble) and fat (chloroform soluble) of the samples were also analyzed and found 57.9 to 100.0 mg/g protein and 15-25% fat. The samples showed higher number of total bacterial count (5.3x10^4 - 2.9x10^7 cfu/g) than the load of Lactobacilli (1.9x10^5 cfu/g) and Streptococci (2.1x10^5 cfu/g). All the isolates were grouped into six on the basis of their morphological and cultural features. Six isolates were selected on the basis of their morphological, cultural, physiological and biochemical characteristics. An attempt was made to identify the isolates. They were found to belong to the three genera viz. Bacillus (S1X3, S1X8, S2Y1, and S2Y2), Staphylococcus (S1X7) and Planococcus (S1X11). They were provisionally identified as Bacillus fastidiosus (S1X3); B. sphaericus (S1X8); B. brevis (S2Y1); B. aminovorans (S2Y2); Staphylococcus saprophyticus (S1X2) and Planococcus citreus (S1X11). The selected isolates were screened for their efficacy in terms of curd and cheese production.

Key words: Cheese, Curd, Microbial load, Nutritional factors.

INTRODUCTION

Milk and milk products including raw milk, pasteurized milk, ice cream, milk powder, yoghurt and cheese are very popular and ideal foods for human consumption throughout the world (Varnam and Sutherland 1994., Dubey and Maheshwari 1999). Cheese can be defined as a solid food product made from the curdled of the milk. The milk is curdled using some combination of rennet and acidification (FDA 1949). Microorganism acidifies the milk and plays a role in texture and flavor of cheese (Weimer et al. 1999). In general, cheese having a great deal of calcium, protein, moisture, salt, fat and many other minerals. All these factors contribute to the microbial stability of cheese (Porter and Hotchkiss 1995). In a variety of ways various microorganisms are associated with cheese which may influence the quality, availability and quantity of cheese (Pelczar et al. 1995).
1993). Two main groups of bacteria are involved in cheese, first groups are starter bacteria and the second groups are adventitious microorganisms from the environment which contaminate the cheese. This group includes *Micrococcus*, *Staphylococcus*, *Bacillus*, *E-coli* and many other organisms (Hoque *et al.* 1999, Cosentino *et al.* 1997, Bareak *et al.* 1993, Robbs *et al.* 1993).

A number of reports are available on the processing, nutritional value, microbial association of cheese and their probiotic activity in different corners of the world. But a few reports are available in Bangladesh. Considering the above facts, the present investigation was undertaken to determine the nutritional factors, microbial load of local and foreign cheese and also the ability of isolates to produce curd and cheese.

**MATERIALS AND METHODS**

*Collection of Samples*

The samples of cheese were collected from different shops of Chittagong and Dhaka (Table-1) and preserve at 4°C in the laboratory.

* Determination of the pH and moisture contents of the samples*

The pH and moisture content of the collected cheese samples were determined by applying traditional laboratory methods.

*Nutritive Analysis of Cheese Samples*

Major nutritional factors of these cheese samples were also analyzed. Protein content of the cheese determined by Lowry method (Lowry *et al.* 1951). For the determination of chloroform soluble fat we used Raghuramulu *et al.* method (1983) with a little modification. One gm of sample was crushed in 10ml of chloroform and allowed to stay for 1 hour. Then the chloroform layer was separated. The process repeated thrice, chloroform extracts poured together in vial and kept open. When the extract was reduced to 1ml, it was transferred in pre-weighed small size screw cap tube and allowed to dry. Again the screw cap tube was weighed and finds out the difference between final and previous weight that indicates the weight of chloroform extract fat of cheese.

*Enumeration of Bacteria*

Ten-fold serial dilution and pour plate technique was followed for isolation and enumeration of bacteria. The total microbial load as well as Lactobacilli and Streptococci load of the samples was determined using Nutrient agar (NA), De Man Rogosa Sharpe (MRS) agar and Yeast Glucose Lemco Agar (YGLA) media respectively.
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Isolation and Identification of the bacteria from samples

The selected bacterial colony were isolated on the basis of their colony morphology and identified on the basis of their morphological, cultural, physiological and biochemical characteristics etc. All the characteristics were compared with the standard description of Bergey’s Manual of Determinative Bacteriology, 8th edition (Buchanan and Gibbons 1974).

Screening of the efficient isolates for cheese production

The identified isolates were screened for their efficacy in terms of the formation of curd and cheese. For the production of cheese, the Foster et al. (1958) method was followed.

RESULTS AND DISCUSSIONS

The pH and moisture content of the cheese samples were determined (Table-1). All the samples were found to be acidic with a pH range of 6.0 to 6.5 and the moisture content of the samples was found within the range of 35.0 to 40.0%. Major nutritional factors protein (1N NaOH soluble) and fat (chloroform soluble) of the samples were analyzed and found 57.9 to 100.0 mg/g & 15-25% respectively. No considerable differences in chloroform soluble fat content of the samples were recorded whereas the protein content was found higher in foreign cheese than the local samples (Table-1).

TABLE 1: DETERMINATION OF pH, MOISTURE AND NUTRITIONAL FACTORS (PROTEIN AND FAT) OF COLLECTED CHEESE SAMPLES.

<table>
<thead>
<tr>
<th>No. of Sample</th>
<th>Description of Sample</th>
<th>Place of Collection</th>
<th>pH</th>
<th>Moisture Content (%)</th>
<th>Protein (mg/g)</th>
<th>Fat (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Local Cottage Cheese</td>
<td>Reazuddin Bazar, Chittagong</td>
<td>6.0</td>
<td>38.5</td>
<td>58.8</td>
<td>25</td>
</tr>
<tr>
<td>2</td>
<td>Kisan Cheese</td>
<td>Narayangunj, Dhaka</td>
<td>6.2</td>
<td>39.2</td>
<td>70.0</td>
<td>17</td>
</tr>
<tr>
<td>3</td>
<td>Milk Vita Cheese</td>
<td>GEC Circle, Chittagong</td>
<td>6.3</td>
<td>40.0</td>
<td>57.9</td>
<td>15</td>
</tr>
<tr>
<td>4</td>
<td>Sliced Anchor Cheese(NZ)</td>
<td>Kamal Store, Chittagong</td>
<td>6.0</td>
<td>36.8</td>
<td>80.4</td>
<td>17</td>
</tr>
<tr>
<td>5</td>
<td>Mozzarella Cheese (Italian)</td>
<td>Kamal Store, Chittagong</td>
<td>6.5</td>
<td>35.0</td>
<td>100.0</td>
<td>22</td>
</tr>
</tbody>
</table>
Frau et al. (1997) proposed that the fat contents of industrially manufactured cheese was significantly different from those in traditionally manufactured cheese (P<0.001) Anonymous 1998, reported that protein content was a little higher in foreign cheese than that of local one. Our results are in concurrence with the above findings. Besides, Taleb et al. (1995) pointed out the chemical changes in protein fraction during maturation of cheese.

In the present study it was found that local cheese showed higher total bacterial count than the foreign samples. Besides, cheese samples showed higher number of total bacterial count (5.3×10⁴- 2.9×10⁷ cfu/g) than the load of Lactobacilli (1.9×10³ cfu/g) and Streptococci (2.1×10³ cfu/g) (Table 2). Lactobacilli and Streptococci are useful for cheese production, cheese moisture retention and also for probiotic activity. (Low et al.1998, Benech et al. 2003, Songisepp et al. 2004.)

**TABLE 2: THE TOTAL BACTERIAL LOAD OF CHEESE SAMPLES.**

<table>
<thead>
<tr>
<th>No. of Sample</th>
<th>Description of Sample</th>
<th>Total bacterial count (cfu/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>NA</td>
</tr>
<tr>
<td>1</td>
<td>Local Cottage Cheese</td>
<td>9.3×10⁵</td>
</tr>
<tr>
<td>2</td>
<td>Kisan Cheese</td>
<td>2.9×10⁷</td>
</tr>
<tr>
<td>3</td>
<td>Milk Vita Cheese</td>
<td>3.1×10⁵</td>
</tr>
<tr>
<td>4</td>
<td>Sliced Anchor Cheese</td>
<td>5.3×10⁴</td>
</tr>
<tr>
<td>5</td>
<td>Mozzarella Cheese</td>
<td>2.5×10⁶</td>
</tr>
</tbody>
</table>

During the period of study, six isolates were selected on the basis of their morphological, cultural, physiological and biochemical characteristics. An attempt was made to identify the isolates by conventional method. They were found to belong to the three genera viz. Bacillus (S₁X₃, S₁X₈, S₂Y₁, S₂Y₂), Staphylococcus (S₁X₇) and Planococcus (S₁X₁₁). They were provisionally identified as the following species: Bacillus. fastidiosus (S₁X₃); B. sphaericus (S₁X₈); B. brevis (S₂Y₁); B. aminovorans (S₂Y₂); Staphylococcus saprophyticus (S₁X₇) and Planococcus citreus (S₁X₁₁) compared with the standard description of Bergey’Manual of Determinative Bacteriology, 8th edition (Buchanan and Gibbons 1974).
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In this study the efficacy of curd formation and cheese production by the selected isolates are shown in Table-3.

<table>
<thead>
<tr>
<th>Isolates No.</th>
<th>Curd Formation</th>
<th>Time of Curdling</th>
<th>Amount of Whey</th>
<th>Colors of Cheese</th>
<th>Dry weight of Cheese(gm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>S₁ X₁₁</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>S₁ X₃</td>
<td>+</td>
<td>2-3days</td>
<td>++</td>
<td>Cream</td>
<td>1.3</td>
</tr>
<tr>
<td>S₁ X₇</td>
<td>+</td>
<td>2-3days</td>
<td>++</td>
<td>Cream</td>
<td>2.0</td>
</tr>
<tr>
<td>S₁ X₈</td>
<td>+</td>
<td>3-4days</td>
<td>++</td>
<td>Cream</td>
<td>1.5</td>
</tr>
<tr>
<td>S₂ Y₁</td>
<td>+</td>
<td>4days</td>
<td>+</td>
<td>Cream</td>
<td>0.9</td>
</tr>
<tr>
<td>S₂ Y₂</td>
<td>+</td>
<td>3-4days</td>
<td>++</td>
<td>Cream</td>
<td>1.0</td>
</tr>
</tbody>
</table>

Note: ‘+’= Positive (‘+’ as scanty, ‘++’ as moderate, ‘+++’ as heavy) and ‘-’= Negative.

Among the isolates Planococcus citreus (S₁X₁₁) was unable to produce any curd and cheese. However Bacillus fastidiosus (S₁X₃), B. sphaericus (S₁X₇), B. brevis (S₂Y₁), B. aminovorans (S₂Y₂) and Staphylococcus saprophyticus (S₁X₇) were able to produce very little amount of curd and cheese which are not suitable for commercial use.

In the early part of this century, scientists and commercial companies began to investigate cheese starters (natural) and examine the organisms they contained. As one would accept, in addition to the lactic acid bacteria it was found that there were a number of undesirable contaminants. These undesirable bacteria might arise in cheese as a result of contamination during handling and processing. Robbs et al. (1993) reported the behavior of Staphylococcus aureus in parmesan cheese during production and storage. Raw milk, plant floor, processing water was the main source of bacterial contamination in cheese (Sabeha and Rashid 1988). So, in the present study, the members of the genus Bacillus, Planococcus and Staphylococcus that were recorded in cheese may be the contaminating microbes. Cosentino et al. (1997) reported the frequency and level of Bacillus sp. contamination in dairy products. Recently Hoque et al. (1999) studied the occurrence of Staphylococcus aureus in milk products. The results of present investigation are in concurrence with the results of Cosentino et al. (1997) and Hoque et al. (1999).

REFERENCES


FDA (Food and Drug Administration) 1949. Department of Agriculture. U.S.A.


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