

Assessment of Bacteriological Quality of Ice-cream During Manufacturing Processes*

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ABSTRACT

To determine the sources of contamination during ice-cream manufacture, fifteen batches of ice-cream were examined bacteriologically. It was observed that total bacterial count varied from 0 to 1,980,000/ml and coliform count 0 to 3000/ml containing in different ingredients of ice-cream mix at the different processed stages. Equipment coming in contact with the product as well as ingredients like colour and essence added after pasteurization of ice-cream mix were found to be the sources of contamination. No pathogenic organisms were detected.

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INTRODUCTION

ICE-CREAM has earned its popularity as dessert and has been judged as finest food nutritionally. Ice-cream provides high caloric value. It has a superior quality of milk proteins which are rich in amino acids and a rich source of calcium and phosphorous along with lactose which favours the assimilation of calcium.

According to Fabian (1945) manufacture of frozen dessert is more complicated than the manufacture of butter and cheese. It is reasonably easy to maintain the chemical composition of ice-cream practically the same throughout the year but difficult to maintain the bacteriological clean conditions of its manufacture and of the equipment involved in manufacturing process. Study on the bacteriological quality of ice-cream mix and ice-cream during different

stages of manufacture in the USSR revealed a heavy contamination with enterococci which did not decrease even after three months storage at -20°C (Mann 1967).

Laxminarayana (1961) stated that high grade ice-cream of acceptable quality can never be produced from raw materials which are initially of poor hygienic quality, irrespective of subsequent methods of treatments and of handling. Similarly the equipment with which the ice-cream mix usually comes in contact at various stages of manufacturing are also important lest they may act as serious sources of contamination for lack of thorough cleaning and sterilization.

Pagaria and Sarswat (1969) observed that pathogenic bacteria can easily gain access to ice-cream mix from fresh or inadequately heat treated milk products.

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or other ingredients and survival in the frozen and hardened finished product unless the mix is properly pasteurized.

Bacterial count of ice-cream is perhaps a better index to judge the conditions prevalent during the production. The present investigation was aimed at collecting the information regarding the bacteriological quality of ice-cream manufactured at the Students' Training Dairy (by a private concern) and to locate the sources of contamination during its manufacture and storage.

MATERIALS AND METHODS

A total number of 15 batches of ice-cream mix and finished ice-cream were examined for the bacterial quality. The bacteriological tests like standard plate count, coliform count, presumptive coliform test were conducted on raw materials such as milk, cream, whole milk powder, sugar, emulsifier, stabilizer, essence and colour when received on the dairy plant as per procedure outlined by Standard methods for examination of Dairy products (1967) except where otherwise stated.

Similarly these ingredients were mixed in proper proportion as per requisite standard and the composite samples of the mixed ingredients were taken at various stages of its manufacture. Such as raw mix (all ingredients mixed), after pasteurization, after aging and freezing and during hardening process also. Along with the standard plate count, coliform count, presumptive coliform test, phosphatase and stormy fermentation tests were also carried out to detect

the sources of contamination. The bacteriological cleanliness of various equipments involved in manufacture of ice-cream like pasteurization vat, mix supply trough: freezing chamber, packaging materials were tested as per ISI, 1963 specifications.

Packaging materials like paper cartons (packages) paper cups were tested with Swab and Rinse method according to ISI specifications.

Necessary precautions were taken while examining the finished ice-cream by first wiping the cup's outer surface with cotton wool dipped in alcohol before making a transfer in the previously sterilized container. The container with ice-cream was then placed in waterbath at 45°C (113°F) for 16 minutes and was thoroughly agitated to ensure proper mixing. The portions were then removed for necessary tests.

Detection of *Salmonella* Spp.

Various media have been suggested for the preparation of differentiation of closely related organisms. The procedure followed for the detection of pathogens (*Salmonella*) was as suggested by Difco manual (1966) and Davis (1963).

Similarly the bottom of the petri dishes with sterile media (S.P.C.A. and MacConkey's Agar) were exposed for the specific period of time i.e. 3, 5 and 7 minutes at different locations in the processing room during ice-cream manufacture to detect bacteriological quality of air as stated by Hedrick and Heldman (1969) circulating in processing room.

RESULTS AND DISCUSSION

The problem of safeguarding ice-cream mix and finished ice-cream against microbial contamination is considerably more difficult than any other dairy products, because the mix ingredients and finished ice-cream have to pass through various stages of preparation while coming in contact with various types of equipment. Frandsen (1938) and Fabian (1942) have listed three agencies which may cause increased microbial count to frozen products. They are (i) ingredients (ii) equipments (iii) workers.

The results obtained from the bacteriological analysis of 15 batches of the mix ingredients have been averaged and are presented in Table 1.

Milk and Cream:

For formulating the ice-cream mix, pasteurized milk and cream were purchased. Standard plate count showed bacterial count from 10, 250 per ml to 1,11,000 per ml in milk and 17,500 bacteria per ml to 1,65,000 per ml in cream (Table 1). About 40% of the total milk samples studied from 15 batches of the ice-cream and about 53.3% of cream showed higher bacterial count than the standard of pasteurized milk and cream. This can be reasonably attributed to improper care during transportation as well as unsatisfactory conditions of containers used.

Whole Milk Powder:

The bacteriological characteristics of whole milk powder used as one of the

ingredients in manufacture of ice-cream can be seen in Table 1. The standard plate count varied from 3200 to 28,500 count/ml while surprisingly no sample was found to contain coli. The count was less than the one suggested by ISI (1963) as 50,000 SPC/ml and 90 coli/ml for the whole milk powder. Herbert *et al* (1967) recorded that 85 per cent of the samples from different dryers were below 30,000 per g. and only 0.9% samples exceeded the 90 coli per g.

Sugar:

The maximum and minimum standard plate count in sugar recorded 29,500 and 2000 bacteria per ml respectively against the standard of 200 bacteria/ml recommended by Foster *et al* (1958) (Table 1). Except for one sample which showed presence of coli less than 10/ml, the rest of the samples were all negative for coli. Foster *et al* (1958) stated that the sugar is well protected there are less chances of the microbial contamination.

Emulsifiers and Stabilizers:

No bacterial standard have been laid down for the emulsifiers however these can be judged on the line of stabilizers. Total bacterial count in emulsifiers has been recorded from 150 to 37,450/ml (Table 1) and coli were present only in three samples out of 15, counting 30, 50 and 100 coli per ml.

Sixty-six per cent of total (stabilizers) samples examined showed SPC count below 10,000/ml while the rest showed higher, some of this upto 28,000/ml. Except one sample rest of the samples

TABLE 1
Maximum and minimum bacterial counts in mix ingredients.

Sr. No.	Ingredients	No. of samples	SPC		Coli.		Below standard			Above standard			Standard recommended count
			Max.	Min.	Max.	Min.	No. of SAM.	SPC	%	No. of SAM	SPC	%	
1.	Milk	15	111000	10250	3000	20	6	40	0	Nil	9	60	15 100 30,000; 10
2.	Cream	15	165000	17500	3000	20	8	53.3	0		7	46.6	15 100 60,000; SPC 10 coli.
3.	W.M.P.	15	28500	2300	0	0	15	100	15	100	Nil	0	Nil 0 50,000 SPC 90 coli.
4.	Sugar	15	29500	2000	10	0	Nil	0	—	—	15	100	— 200 SPC no report coli.
5.	Emulsifier	15	37450	150	100	30	8	53.3			7	46.6	no St. (SPC 10,000 SPC) Coli.
6.	Stabilizer	15	28000	0	90	0	10	66.6			5	33.3	10,000 or less no report coli.
7.	Essence	15	21050	100	0	0	1	6.6			14	93.4	100 SPC
8.	Colour	15	165000	3250	2670	0	6	40			9	60	10,000 SPC

showed absence of coli (Table 1). The quality of the stabilizer used can be said to be of satisfactory standard.

Foster *et al* (1958), Gayer *et al* (1966) both have suggested that stabilizers should not exceed the count more than 10,000/ml.

Colours:

It can be seen from Table 1 that the average standard plate count for colour varied from 3,250 to 1,65,000 count/ml and coliforms from 0 to 2,670 count/ml; 60 per cent of the samples are above the standard plate count of 10,000/ml and 73 per cent of the samples showed more than 10 coliforms/ml indicating improper preparation, preservation and handling.

Flavouring Materials:

The results obtained in Table 1 showed 100/ml to 21,050/ml bacterial

count of which 93.4% samples failed to attain the standard of 100 bacteria per ml. However, all the samples were free from coli count.

Raw Mix (all ingredients mixed).

All the ingredients were mixed together in pasteurizing vat and the bacterial count was taken which varied from 20,000 to 1,65,000 bacteria/ml and coliforms were 70 to 3,000/ml. Shitskov (1954) recorded 1,25,000 to 1,80,000 count per ml from the samples obtained from two different factories for the raw ice-cream mix.

Pasteurization Process:

The mix was pasteurized at 72°C for 20 mts. Table 2 depicts the bacterial count of the pasteurized mix. The efficiency of the pasteurization process was 98% indicating reasonably low bacterial count in pasteurized mix. The

TABLE 2

Total bacterial counts at various stages.

Sr. No.	Raw mix		Pasteurized mix		Aging		Freezing		Hardening	
	SPC	Coli.	SPC	Coli.	SPC	Coli.	SPC	Coli.	SPC	Coli.
1.	1,10,000	180	8,000	0	65,000	120	80,000	200	1,65,000	460
2.	1,65,000	720	3,000	0	45,000	20	72,000	30	48,750	150
3.	75,000	70	1,500	0	25,550	20	36,000	300	1,62,500	500
4.	54,850	70	1,600	0	33,000	50	42,000	90	64,050	350
5.	42,550	80	1,400	0	28,150	30	37,000	70	64,500	680
6.	20,200	780	800	0	39,500	210	24,000	190	57,250	260
7.	1,55,000	1250	1,700	0	45,450	990	57,350	480	52,300	320
8.	56,450	200	1,000	0	32,600	120	29,900	170	92,550	3000
9.	1,29,500	900	1,850	0	17,000	0	26,500	100	1,58,550	520
10.	1,40,000	3000	1,800	10	58,500	2000	24,600	3000	21,000	230
11.	35,500	610	700	0	48,000	120	50,050	80	50,100	180
12.	76,700	110	900	0	38,850	90	60,250	30	67,200	1110
13.	58,500	270	950	0	86,500	1110	77,500	650	2,07,500	2300
14.	50,200	150	1,830	0	58,000	140	75,000	230	1,21,000	1600
15.	72,750	2000	2,350	0	1,78,500	110	1,98,500	500	1,69,600	1800

coliforms were absent in all the pasteurized samples. To reduce the protective effect of sugar and stabilizer higher temperature are usually recommended.

Foster *et al* (1958) reported that ice-cream mix if pasteurized properly, will give total count of 10,000 or less bacteria per g.

Aging and Freezing:

It can be seen from Table 2 that during the aging period there was an increase in bacterial count as well as coli count.

The increase in bacterial count during the process of aging and freezing can be attributed to improper handling of mix by the workers involved in manufacturing of ice-cream. It can be due to prolonged storage and exposing the pasteurized mix to atmospheric temp. by often opening and closing the aging vat.

Similarly there was an increase in bacterial count after freezing the mix. Ellikar (1949), Foster *et al* (1958) noted the increase after freezing on account of breaking up of clumps of bacteria and contamination from the freezer.

But Llyod (1969) observed reduction after freezing and concluded that the effect of freezing on various types of bacteria results in physical destruction of the cells by agitation in the presence of ice-crystal. Alexander and Rothwell (1970) reported that the reduction of bacterial count on freezing is mainly at the temperature of maximum ice-crystal formation.

Results recorded in Table 3 indicate that the freezing chamber, mix supply trough and environmental air are reasonably responsible for the increase in the total bacterial count after the freezing process.

The ingredients, colour and essence do not receive any heat treatment and are directly added to the pasteurized mix at the time of freezing which might act as the source of contamination resulting in increased load of bacterial count after the freezing process.

Bacterial count after hardening period in cold storage:

The increase in total bacterial count during storage reveal that due to uncontrolled practice of handling. On and often entry in the hardening room, or unmindfully keeping the doors open result into the fluctuation of the temperature which encourages the increased bacterial count.

The final product of ice-cream taken out from the storage after 48 hours proved to be positive to presumptive coliform test in all samples. Patel and Vyas (1969) reported 90% of the cases positive to presumptive coliform test conducted on the ice-cream sold in Anand market (Gujarat). However, according to Delay (1947) the presence of coliforms in the samples cannot be correlated to the presence of pathogens in the samples examined.

Patel and Vyas (1969) reported a very small incidence of Salmonella organisms in the icecream sold in Anand market (Guj.).

TABLE 3

Microbial condition of the equipments, air and packages used in manufacturing of ice-cream.

Sr. No.	Pasteurization		Mix supply trough		Freezing chamber		Air		Packages		S.P.C.		Coli.	
	SPC	Coli.	SPC	Coli.	SPC	Coli.	SPC	Coli.	SPC	Coli.	Max.	Min.	Max.	Min.
1.	9000	0	25000	0	18500	0	128-28	0-37	13600	0	25500	4250	250	0
2.	5500	0	10750	0	6250	0	30-68	38-68	16000	0				
3.	18750	0	30750	0	15000	0	18-18	0	18200	400	46500	5000	1000	0
4.	19000	0	27500	500	20000	0	6-23	0	2400	200				
5.	5500	0	14250	0	30000	250	31-97	0	14800	800				
6.	11000	0	46500	0	3000	0	5-51	0-29	9400	800	75000	3000	1000	0
7.	25000	0	14500	0	22250	0	67-81	0	5200	0				
8.	4250	0	6000	0	9000	0	53-53	1-78	3600	0	128-28	5-51	38-68	0
9.	11000	0	19500	1000	11500	0	20-55	0-95	6400	0				
10.	5500	0	37500	0	7500	1000	13-48	0-83	9400	0	45000	2400	1800	0
11.	19750	0	34000	0	23500	0	68-80	1-49	7400	200				
12.	13500	0	22250	0	11500	0	30-72	0-83	5400	0				
13.	17500	0	8000	0	2000	0	24-14	0	6400	0				
14.	17500	250	5000	0	8250	0	33-56	1-25	35400	1800				
15.	11250	0	10500	0	3750	0	34-96	.59	45000					

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