THESIS

SWEET CREAM BUTTER

by

R. E. Caldwell A. H. Wright

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1908.

on

Sweet Cream Butter.

Experiment stations of late years have been working upon the subject of flavor and keeping quality of butter. They have taken up the subject of the bacterial content of milk, ripening of cream by control of temperature, and control by commercial starters, but practically no work has been done with regard to producing sweet cream butter. It is for this purpose, therefore, that we study of intend to make a butter made from sweet cream, as compared with sour cream, with particular reference to the following points: keeping quality, ripening quality, and its bacteriological content.

In order to have a clear conception of what we mean when we speak of sweet or sour cream butter, it would probably be best to define just what we meant by each. By sweet cream butter is meant butter made from cream having less than .2 per cent. acidity; and by sour cream butter we mean butter made from cream having more than .2 per cent. acidity, usually .4 to .5 per cent.

A study of sweet cream butter would naturally be very closely related to the subject of natural and commercial starters. In order, therefore, to have a more definite idea concerning the relation that exists between them, we shall discuss somewhat in detail the different processes and the results sought for.

The ripening of cream has for many years been considered one of the essential factors in the production of a good butter. The object of ripening cream is to produce desirable flavors by means of growing certain species of bacteria, which act upon the milk sugar and form acid. This acid is transmitted to the butter, giving it a very palatable flavor. It also was thought that the acid destroyed the coating surrounding the fat globule, and

in that way cream could be churned more quickly. Until very recently the desirable flavor of butter was a pure acid taste. At present, however, the best trade is calling for a butter that is low in acid and characterized by a sweet nutty taste; or a butter that is characterized by the absence of any objectionable flavor.

As favors in milk and butter are produced almost entirely by the action of bacteria upon its different parts, a control of this factor will also control the flavor. Therefore, it is the OBJECT OF THIS THESIS to make a careful study of sweet cream butter as compared with sour cream butter, in order to determine the relative keeping quality, ripening quality, and bacteriological content of each.

DISCUSSION.

Ripening of cream is a process brought about by a series of chemical changes caused by the action of micro-organisms, but little understood by chemists. A combination of these changes into one general term is called fermentation. It is definitely know that certain organisms are desirable in milk, and that certain other organisms are not desirable. It has been found by analysis that there are over one hundred species of bacteria that produce acid in milk. Obviously, it is not necessary to have all of these species present. In fact only a few of them are desirable, and just what these are is a question which is not yet settled. It also remains a doubt whether or not the same organism produces the desired acid, aroma, and flavor; but one thing is certain, when conditions are favorable for one the other also develops, showing their close relation.

Conn claims that the bacteria which act upon the nitrogenous part of the milk are closely associated with the bacteria that produce the desirable flavor in butter. Weighman claims that the best results are obtained when a variety of species work together in the cream. Eckles cane: to a similar conclusion, and asserted that the flavor and aromatic # Storr's Station, Conn substances produced during the ripening of cream may be produced by a variety of acid producing bacteria. Storch reaches a like conclusion, and states that flavor and aromatic products are the result of a joint action of a great many species of lactic acid producing bacteria.

From the foregoing conclusions of these investigations it seems evident that no single organism produces the desirable flavor in butter and cream. It is evident, however, that the organism or organisms that produce the required flavor belong to the lactic acid group.

What composes the aromatic substances given off from butter is not known, but they undoubtedly are volatile compounds of ether like substances produced by the action and reproduction of bacteria in the milk.

The next question which we must now deal with is the best method of controlling the growth of the bacteria in the cream, not only by inoculating it with the desired organism to begin with, but also to properly control the organisms after inoculation. In order to do this it was planned to inoculate cream while sweet with a culture of desirable lactic acid bacteria, cool and churn at once before any bacterial growth has taken place. The theory is that by churning immediately enough bacteria will be incorporated in the butter to produce any shade of variation desired in the flavor. In other words, by excluding all organism from the cream except those which produce desirable results we have complete control of the bacteriological content of the butter, and can market the butter whenever it is found that the ripening process has reached the most desirable stage.

The method used almost universally at present, and advised by our dairy schools, is what is known as Pasteurization and Artificial Inoculation' This process consists chiefly in two separate parts: First, The cream is taken as soon as delivered and heated to about 175° F.. This kills almost all germ life present. Second, The cream is then cooled to 70° F. and inoculated with a

pure culture of lactic acid bacteria;--usually about 10% is used. The cream is then held at a temperature of about 70° F. until the desired acidity is reached. The acidity is used as an index to determine the degree to which the bacteria have developed. The starter used is kept in stock by **in**oculating pasteurized skim milk with a commercial starter, many kinds of which are now on the market.

The work in this experiment on Sweet Cream Butter is divided into two parts. A good grade of cream was first selected and **pasteurized** to 175° F., then equally divided and to #1 we added 10% starter and churned as soon as cool. No. 2 was carried out by the common creamery method, as above described.

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Sweet Cream Butter

Outline #1.

A. Cream,-

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1.	Amount taken,	39
2.	Per cent. acidity	10 01
3.	Bacteriological Content	#1
4.	Temperature of pasteurization	•
5.	Bacteriological Content	#2
6.	Per cent. acidity 14	
7.	Starter, kind, Erricsson Per cent used,	10
	Acidity	
		₿3
		19
	Churn #1, when cooled to 55° F.	
	Wash with water pasteurized to • • • 175° F.	
	Churn #2 when $.35\%$ acidity at $ 54^{\circ}$ F.	
	Wash with water pasteurized to 175° F.	
Butte		

- 1. Bacteriological Content every 10 days:
 - (1) #4, sweet cream butter
 - (2) #5, sour " " " "
- 2. Keep #1 and #2 at 50° F.

__THESIS .---

Sweet Cream Butter

Outline #2.

A. Cream,-

1.	Amount taken 38#
2.	Per cent. acidity 125
3.	Bacteriological Content 1'
4.	Temperature of pasteurization 175° F.
5.	Bacteriological Content #2
6.	Per cent. acidity
7.	Starter, kind Percent used 10
8.	
8. 9.	
8. 9. 10.	Bacteriological Content · · · · · · · #3' Acidity 74% Divide equally: Pounds each 19
8. 9. 10.	Bacteriological Content · · · · · Acidity 74% #3' Divide equally: Pounds each 19 Churn #1 when cooled to · · · · · 52° F.
8. 9. 10. 11.	Bacteriological Content · · · · · · Acidity 74% #3' Divide equally: Pounds each 19 Churn #1 when cooled to · · · · · 52° F. Wash with water pasteurized to · · · 180° F.

B. Butter,-

1. Bacteriological Content every 10 days:

(1) #4', sweet cream butter

(2) #5; sour " "

2. Keep #1 and #2 at 50° F.

---THESIS---

Sweet Cream Butter.

Sub-outline.

Cream,-

1. Take acidity

- 2. Bacteriological Content.
 - (a) 1 1 1 100 500 1000 agar and gelatin, 12 plates.
 - (b) Count.
 - (c) Relation of liquefiers.
 - (d) Take sub-cultures on agar plant.
 - (e) Run sub-cultures thro:--
 - 1'. Glucose agar
 - 2'. Gelatin stab
 - 3'. Plain milk
 - 4'. Litmus milk or litmus gelatin
 - (f) Microscopic examination for germs.
- 3. Pasteurize sample.
- 4. Bacteriological Content -- same as (2).

Methods Used in the Bacteriological Study and Description.

It is thought that a few words here are needed as to the method of obtaining the data herein tabulated. The following bacteriological analyses were taken:--#1, sweet cream before pasteurization,--#2, cream after pasteurization,--#3, starter,--#4, sweet cream butter,--#5, sour, or ripened, cream butter. The history analyses were taken sixteen days from date of making.

The study of bacteria was divided into two heads, --quantitative and qualitative. The quantative was made by using plain agar and litmus gelatine. Four plates were made on each media. The dilutions used were varied according to the estimated content of the sample. In each case check plates were made, and the average of the two were computed, thus dispensing with any undue chance of error. In the qualitative analysis the following methods were used:--The isolation of the bacteria was accomplished by making inoculations from plate cultures, usually about two days old. The inoculations were made from the plate cultures on agar slants, and plain bouillon. When the growth on either of these media was moderately abundant, sub-cultures were made on the various differential media. The following were generally used:--Litmus milk, gelatine stab, glucose agar, and plain milk.

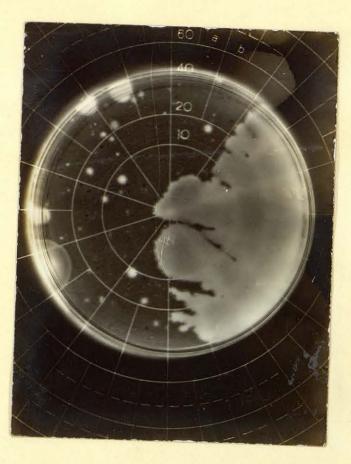
The morphology of the organisms was determined from the bouillon cultures in most cases, althought in some it was secured from fresh agar slants. The organisms were first examined under the microscope in hanging drops in order to determine the motility. No determination of spores or flagella was attempted. After this, stained preparations were made and reserved for future identifications and comparison.

In general, in isolating and determining the species and families of bacteria we have followed the methods used by Conn.



This represents a small regular spreading 1 colony, surface growth. Very dense. A _____ 2,000

dilution. Culture features found, 1 -- B -- 2.



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This represents a very characteristic specimen of a large whitish, spreading colony. Surface growth; bluish floresence.

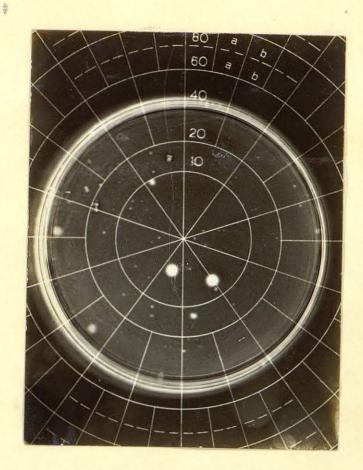
Culture features found in 3 -- B -- 1.



This represents the growth of an irregular

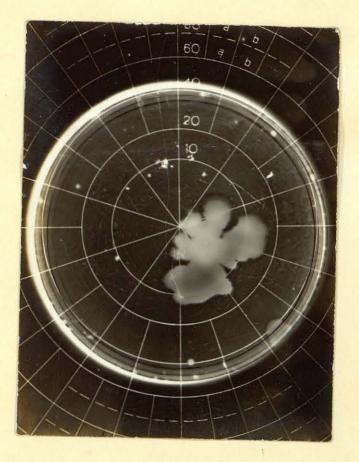
shaped, surface growing colony. Cream, white.
1
A ----- dilution.
10 000

Culture features found on 4 -- B -- 2.



1

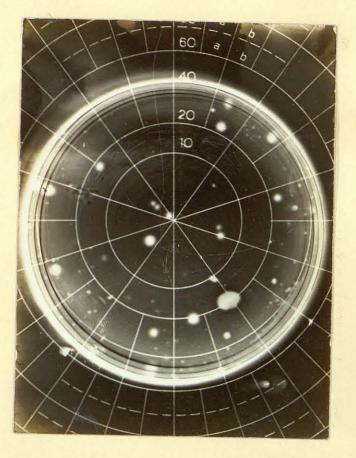
This represents a typical lactic acid colony. Small, oblong, grayish color. Deep and surface l growth. A ----- dilution. 10 000 Culture features found on 3 -- B -- 2.



This represents the growth of a yellow spreading colony. Very irregular, cloudy, surface growth.

1 A ----- dilution. 1 000

Culture features found on 1 -- B -- 4.



This represents a very small, regular colony.

Both surface and deep growth. Grayish white. 1 a ----- dilution. 2,000

Culture features found on 1 -- B -- 3.



This represents a very characteristic growth of mold. The radial edge is its truest characteristic.

Qualitative Analysis.

Content #1'

No. B-1

(a) Description: ... Large white spreading colony.

Surface growth.

(b) Relative per cent .: ... Trace.

No. B-2

(a) Description: Small distinct surface colony, round

and regular.

(b) Relative per cent .: 5%.

No. B-3

(a) Description: Nut shaped organism, deep and surface

growth.

(b) Relative per cent .: 70.

No. B-4

- (a) Description: Yellow spreading colony, surface growth.
- (b) Relative per cent., 5.

No. B-5

(a) Description: Medium size, radial spreading colong,

surface.

(b) Relative per cent., 1.

Quantitative Analysis.

Content #1'.

Agar Plates. -A- 24 hours. 1 --- solution 10,080,000 per c.c. 200 1 " " 10,650,000 per c.c. 1000 1 " <u>10,120,000</u> per c.c. 2000 -Average 10,283,000 per c.c. Agar Plates. -B-1 --- solution per c.c. 200 1 "" " 4,380,000 per c.c. 1000 1 """ " 3,920,000 per c.c. 2000 Average, 4,150,000 per c.c. Final Average on Agar, 7,216,500 per c.c. Gelatin Plates. -A- 48 hours. 1 --- solution, No growth per c.c. 200 per c.c.

Qualitative Analysis.

Content #1.

No. B-1 (a) Description: ... Large white spreading colony. Faintly regular, surface growth. 7 (b) Relative per cent. 100,000 No. B=2 (a) Description: ... Small, distinctly white, surface colony. Very round and regular (b) Relative per cent .: 20. No. B-3 (a) Description: ... Very small, regular colony. Both surface and deep growth. Grayish white. (b) Relative per cent .: 20. No. B-4. (a) Description: ... Yellow spreading colony. Very irregular, cloudy, surface growth. (b) Relative per cent .: About 1 to 1,000,000. No. B-5 (a) Description: ... Large spreading colony. Distinctly

radial. Yellowish white.

(b) Relative per cent .: About 3 or 4 to 1,000,000

Quantitative Analysis,

Content #1.

Agar Plates. -A-- 24 hours. 4 1 ----solution, to numerous per c.c. 100 1 " " 3,460,000 per c.c. -----500 1 " " 5,760,000 per c.c. -----1,000 Average 4,610,000 per c.c. Agar Plates. -B-1 solution, to numerous, per c.c. 100 1 " " 2,540,000 per c.c. -----500 1 " " 5,800,000 per c.c. -----1,000 Average 4,176,000 per c.c. Final Average on Agar, 4,390,000 per c.c. Gelatin Plates. -A-- 48 hours. solution, no growth per c.c. 100 1 " " 2,340,000 per c.c. ------500 1 " " 2,720,000 -----per c.c. 1,000 Average 2,530,000 per c.c. Gelatin Plates. -B-

1 solution, to numerous per c.c. 100 1 " 2,920,000 per c.c. 500 1 " 2,800,000 per c.c. 1,000 Average 2,860,000 per c.c. Final Average of Gelatin 2,695,000 per cc. Grand Average per c.c. 3,542,500

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2.81

	Qualitative Analysis No. I.																
No. of Plate																	
No. per cc																	
1083	Diameter over 100 Chains Spores Motility																
Lod	C	hains	+	+	+		+	+	+	+	+						
rp	S	pores											-				
Mo	M	otility	+	+	+		+	+	+	1	I						
	G	raham Stain				1											
	th	Turbidity	+	+	+		*	+	÷	÷	÷						
	Broth	Scum	1	ł	1	-	1	1	1	1	1						
	E A	Sediment	1	+	1		I	+	1	1	+						
20	н	Dull	1	1	I		1	1	1	1	4.						
ILO	Agar	Wrinkled	+	T	1		+	1	1	I	1						
Features	-	Chromogenesis	1	1	1		1	1	1	ł	1						
0 El		Round-compact	*	+	1		+	+	1								
Н	я.	Proteus-like	1														
Cultural	atin	Rhizoid	1	Ŧ	+		1	1	- +1	ŧ							
1 tr	Gele	Filamentous	1	1	E		1	1	1								
Cu	60	Curled	1	ŧ	1		1	1	1								
	Gel. Stab	Surface growth	+	1	1		+	1	1	1	1						
	00 0 0 0	Needle growth	+	+	+		+	+	+	+	+						
	*	Gelatin	I	ŀ	ł		1	1	1	.1	1						
61		Coagulated	+	+	+	63	÷	+	+	+	+						
nic		Acid	+	+	+	No	+	+	+	+	+						
Bio-Chemical	R	Alkaline	1	1	11	102	1	1	1	I	1						
01	Milk	Saponified	1	1	1	ys	1	1	1	1	1						
Bic		Peptonized		- 6		Apsl											
	+	Gas	1	1	i	A	1	1	1	ł	1						
	Organism name																
*	=	Liquefaction +	= G	1	Su	gai											

Qualitative Analysis,

Content #2'.

No. B-1

(a) Description: ... Large white surface, colony

regular.

(b) Relative per cent .: Trace.

No. B=2

(a) Description: ... Single distinct surface.

Colony regular.

(b) Relative per cent .: 1.

No. B-3

(a) Description: ... Small regular. Deep surface

growth.

(b) Relative per cent.

No. B=4

(a) Description: ... Large dense white spreading

colony. Regular.

(b) Relative per cent .: 2.

No. B-5

(a) Description: ... Small regular yellow colony,

dense.

(b) Relative per cent .: Trace

Quantitative Analysis,

Content #2'.

Agar Plates. -A- 24 hours.

solution, plates neutralized.

Final Average on Agar 192,000 per c.c.

Gelatin Plates. -A- 48 hours.

No growth on gelatin.

Qualitative Analysis.

Content #2.

No. B-1

- (a) Description: .. Large white surface growth.
- (b) Relative per cent. 1.

No. B-2

(a) Description: .. Small distinct surface colony.

Regular.

(b) Relative per cent .: 3.

No. B-3

(a) Description: .. Small regular, deep surface

growth. Lactic acid.

(b) Relative per cent. 40.

No. B-4

(a) Description: ... Large white, dense surface colony.

Regular.

(b) Relative per cent. 5.

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Quantitative Analysis,

Content #2.

Agar P	lates.	-A-						
	1	solution,	22,500	per o	c.c.			
	1	н н	31,000	per o	c.c.			
	500 1		30,000	per o	G.C.			
	1,000	Average	27,800					
Agar P	lates.	-B-						
	1	Solution	36,000		C • C •			
	1	11 11	32,000	per c	c.c.			
	500	Average	34,000	per c	C . C .			
		Fina	1 Average	on Age	ar,	30,900 pe	r c.	с.
Gelati	n Plate	esA-						
	1							
	100	solution	6,000	per c	C • C •			
	1 500	и п.	7,500	per c	C.C.			
	1	n n	5,000	per c				
	1000	Average	6,166	per c				
Gelati	n Plate	esB						
	1							
	100	solution	6,000	per c	·			
	500	17 11	2,000	per c)•C•			
	1	н н	6,000	per c	• • • •			
		Average	4,666 Final Ave Grand Ave	erage o	on Gel	atin 4600 17950		
			ur and Ave	area		11750	per	0 0.

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Qualitative Analyses No. II. 281 2B2 Bl B2 B4 B4 B5 No. of Plate No. per cc Diameter over 100 Morphology Dhains * 1 1 + 1 1 + + Spores + 1 + Motility + 1 + 1 1 Graham Stain + 4 + 1 + + + Turbidity 1 Broth Scum 1 1 1 Ŧ * 1 1 1 + afa + 4 + + + + Sediment ÷ + + ÷ + Features Dull + + + Agar 1 + f Wrinkled 1 + 1 + 1 Chromogenesis + 1 1 + 1 1 1 1 Round-compact Latin Proteus-like + ÷ Cultural Rhizoid + + Filamentous + ÷ Curled Gel Surface growth + + 1 + + 1 + 1 1 -d-4 1 + + 1 + Needle growth 1 T J Gelatin 1 I x 1 1 * Bio-Chemical 1 + + 1 + + 4. + Coagulated MO Acid Ŧ + + 1 + + + + Milk Alkaine + 1 1 02 ÷ I 1 1 1 VSI 1 1 1 Ŧ 1 1 1 1 Saponified n 8. Peptonized 1 1 1 1 1 1 1 1 + Gas. ł 1 1 1 1 I 1 Name of Organism

* = Liquefaction + = Gl. Sugar.

Qualitative Analysis,

Content #3'.

No. B-1 (a) Description: ... Surface spreading colony. Blueish floresence. (b) Relative per cent .: Trace No. B-2 (a) Bescription: ... Nut shaped, deep growing colony. Lactic acid. (b) Relative per cent .: 80. No. B-3 (a) Description: ... Very small white colony. Deep and surface growths. (b) Relative per cent.: 30. No. B-4 (a) Description: ... None present. No. B-5 (a) Description: ... Large radial yellow colony. Surface growth. (b) Relative per cent .: trace. No. B-7 (a) Description: ... Large definitely radial white surface growth. (b) Relative per cent .: Trace. No. B-8 (a) Description: ... Regular dense colony, surface growth. (b) Relative per cent .: 3.

Quantitative Analysis,

Content #3'.

Agar Plates. -A- 24 hours. 1 ----- solution 75,000,000 per c.c. 1,000 Ъ " " 15,600,000 per c.c. 2,0000 1 ----- " " 60,000,000 per c.c. 100,000 Average 50,000,000 per c.c.

Agar Plates. -B-

```
1
----- solution 80,000,000 per c.c.
1,000
1
"" " 98,000,000 per c.c.
20,000
 1
---- " " lost
100,000
     Average 89,000,000 per c.c.
```

Final Average on Agar 69,600,000 per c.c.

Gelatin Plates. -A- 48 hours

No growth

Content #3.

- No. B-1
 - (a) Description: ... Large whitish spreading colony.

Surface growth, blueish floresence.

- (b) Relative per cent.: 1 to 1,000,000
- No. B-2
 - (a) Description: ... Small, oblong, yellow colony.

Characteristically shaped.

Irregular, deep growth.

(b) Relative per cent .: 40.

No. B-3

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(a) Description: ... Very small white colony.
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Round, regular, deep and surface growth

- (b) Relative per cent .: 50.
- No. B-4
 - (a) Description: ... Light blue, round colony. Dark yellow in center, light on edges.
 - (b) Relative per cent.: 1-2 to 1,000,000.
- No. B-5
 - (a) Description: ... Large radial yellow colony.
 Irregular surface growth.
 - (b) Relative per cent.: 1-2 to 1,000,000.
- No. B-6
 - (a) Description: ... Medium, definitely radial,

white colony.

(b) Relative per cent .: trace.

Quantitative Analysis. Content #3. Agar Plates. -A- 24 hours. 1 ----- solution too numerous 5 000 1 •••••• " 84,800,000 per c.c. 10,000 1 "124,000,000 per c.c. 50,000 Average104,000,000 per c.c. Agar Plates. -B-1 ----- solution too numerous 5,000 1 "" " 88,000,000 per c.c. 10,000 1 """ " 126,000,000 per c.c. 50,000 Average 102,000,000 per c.c. Final Average on Agar, 103,000,000 per c.c. Gelatin Plates. -A- 48 hours. 1 ----- solution 1,400,000 per c.c. 5,000 1 "" " 1,690,000 per c.c. 10,000 1 "" " 1,800,000 per c.c. 50,000 Average 1,600,000 per c.c. Gelatin Plates. -B-1 ----- Solution 1,500,000 per c.c. 5,000 1 "" " 1,200,000 per c.c. 10,000 " " 1,600,000 per c.c. Average 1,430,000 Final Average on Gelatin 1,515,000 per c.c. 50,000 Grand Average 52,257,000 per c.c.

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	Qualitative Analysis No. III.																			
	No.	. of	Plate	3B1	332	5B3	334	3B5		Bl	20 20 20	B3	B4	B5	B7	B8.				
	No.	pe	rcc																	
Ĩ	5y	D	iameter over 100																	1
	Log	Chains Chains Spores Motility Crehem Stein				4.	4-	+		1	+	+		+	+	+				
	ho	S	pores																	
	d'r'	M	otility	+	+	1	*	+		+	+	1		+	+	1				
	MC	G	raham Stain																	
		q	Turbidity	+	+	+	+	+		+	*	+		+	+	I				Ī
		Broth	Soum	I	1	1	1	1		1	1	1		1	1	1				1
		B	Sediment	+	+	+	+	+		+	+	4		+	+	.+			2	1
	Features		Dull	+	1	I		1		+	I	1		1	1	+				1
4114	tu	Agar	Wrinkled	1	1	1	1	1		1	1	1		1	1	1				1
	68	Ag	Chromogenesis	1	1	1	j	1		J	1	1		1	i	1				1
			Round-compact																	1
r	ral	In																		1
	0ultural	stin.	Rhizoid																	1
	Cul	Gel	Filamentous							1										1
			Curled	-																1
		ap ap	Surface growth	+	+	+	4			+	+	+		+	ł	+				1
		Gel	Needle growth	1	+	+	+			+	+	+		+	*	1				1
		*	Gelatin	Ł	1	T	1			1	1	1		1	1	1				
	Ц		Coagulated	+	+	+	+	4		+	+	+		+	+	+				
	108		Acid	+	+	+	+	+	MO	+		+	ent	+	+	+				
	Bie-Chemical	M	Alkaline	1	1	1	I	1	is	1	1	1	000	1	1	1				
	-CI	Milk	Saponified	1	1	1	1	1	200	1	1	1	pr	1	1	1		-		
	910	-	Peptonized	1	i	I	1	1	nal	1	1	1	ne	1	1	T				1
	щ	+	Gas	+	+	+	+	+	A	+	+	+	NO	+	1	1				1
	Name of Organism																			

* = Liquefaction + = Gl Sugar.

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Quantitative Analysis,

Content #4'.

Agar Plates. -A- 24 hours.

1 1 ----- slution, 3,840,000 per c.c. 1,000 "" " 4,640,000 per c.c. 2,000 1 " 3,960,000 per c.c. 10,000 Average, 4,133,300 per c.c. Agar Plates. -B-

1,000	soluti	on,	6,000,000	per	c.c.
2,000	11	n	2,880,000	per	C . C .
10,000	Ħ	**	2,640,000	per	c.c.
	Avera	ge	3,840,000	per	C . C .

Final Average on Agar, 3,986,600 per c.c.

Gelatin Plates. -A-

No plates made on gelatin.

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Qualitative Analysis.

Content #4.

History Analysis.

No.

- (a) Description: .. Lactic acid
- (b) Relative per cent .: 90.

No. B-1

(a) Description: ... Cream white, round colony.

Surface growth.

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(b) Relative per cent .: 9.

No. B-2

(a) Description: ... Moulds and irregular

colonies.

(b) Relative per cent .: 1.

No. B-3

(a) Description: ... No spreading colonies.

One mould.

Quantitative Analysis,

Content #4.

History Analysis

Agar PlatesA-	24 hours.
1	71.000
solution 100	31,000 per c.c.
1 2,000 1	200,000 per c.c.
10,000	610,000 per c.c.
	347,000 per c.c.
Agar PlatesB- l	
1,000	320,000 per c.c.
1 2,000 1	216,000 per c.c.
10,000	600,000 per c.c.
	378,600 per c.c.
Fina	1 Average on Agar 362,800 per c.c.
Gelatin PlatesA-	48 hours.
	720 000 non a a

1,000	solution	720,000	per	c.c.	
1 2,000	" "	772,000	per	c.c.	
1 0000	" "1:	iquefied			
1,0000	Average	746,000	per	C.C.	

Gelatin Plates. -B-

No growth on gelatin. -B-

Finel Average on Gelatin 746,000 per c.c. Grand Average 554,400 per c.c. 295

D

Qualitative Analysis.

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Content #4.
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No. B-1

(a) Description: ... Cream white round colony.

Surface growth.

(b) Relative per cent .: Very small.

No. B-2

(a) Description: ... Light yellow surface colony. Star

growth in center very characteristic.

(b) Relative per cent .: Very small.

No. B-3

(a) Description: ... Very small, --filimentous.(Proved to be a mold)

(b) Relative per cent. Comparatively large.

No. B-4

(a) Description: ... Large white surface colony.

Irregular growth.

(b) Relative per cent.: 45.

No. B-5

(a) Description: ... Numerous molds present. Few

non-acidifiers.

(b) Relative per cent of acidifiers, 90.

296

D

Quantitative Analysis.

Content #4.

Agar Plates. -A- 24 hours. 1 solution 210,000 per c.c. 1,500 1 ----- " " 750,000 per c.c. 3,000 1 " " 1,785,000 per c.c. 15,000 Average 915,000 per c.c. Agar Plates. -B-1 solution 550,500 per c.c. 1,500 1 " " 288,000 per c.c. 3,000 1 " 765,000 per c.c. 15,000 Average 534,500 per c.c. Final Average on Agar 724,750 per c.c.

Gelatin Plates. -A- 48 hours

1 500	solution	60,000	per c.c.
1,000		68, 0 00	per c.c.
1 5,000		9,0,000	per c.c.
,,	Average	72,666	per c.c.

Gelatin Plates. -B-

P. G. & G

1 500	solution	59,000	per c.c.	¢
1,000	n n	70,000	per c.c.	
# # # # # #	n n	150,000	per c.c.	
5,000	Average Final Ave Grand Ave		alatin 828,333 403,742	3 per c.c 2 per c.c

TY"

297

298

D

Qualitative Analysis No. IIII. B2 B3 B4 B2. B4 Bl BS B1 B4 B5 B6 No. of plate 37 No. per c. c. Diameter over 100 Morphology ł -4-+ ł + Chains + + 43 Spores + õ 1 1 + Motility + 1 + Graham Stain 07 Turbidity E 1 1 th + + * ÷ Soum Bro. 1 HO 1 + 1 1 + 1 54 Sediment 1 + + ÷ 1 + + I Dull + + 1 δΩ ÷ + Agar 1 Features Wrinkled 1 I 1 1 1 I 1 TR Chromogenesis £. 1 1 1 1 1 ŧ 0 Round-compact Gelatin Xt Cultural Proteus-like S.t. Khizoid Filamentous ri. 3 No. Curled 1 1 + + rent Gel Stab 1 + + F + Surface growth + ÷ 40 + 4 -H - 1 Needle growth 1 4 i 07 00 1 1 1 + 1 1 * Gelatin ÷ 20 NO N + Coagulated 1 1 21 4 i. H + ÷ 45 anese' esent 0.0 esent Bio-Chemic 44 Acid 1 1 1 Ē ÷ En. ÷ + MIJK 0 1 1 1 th 00 1 Alkaline 1 I E P 10 IL OL Id 31 Id Id T 1 1 2 1 0 1 Saponified Mone None E.d 1 so Mone None + 1 En Peptonized 1 1 1 1 + E P.O. 11 01 1 1 + 1 1 1 1 Gas. 1 Organism name * = Liquefaction

Qualitative Analysis. Content #5.

No. B-1

(a) Description: ... Large white spreading colony.

Regular edges, surface growth.

(b) Relative per cent .: .000001.

No. B-2

(a) Description: ... Medium large yellow colony.

Very regular, dense surface growth.

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(b) Relative per cent .: 2.

No. B-3

(a) Description: ... Small white deep growth colony.

Very regular, open, rather pale.

(b) Relative per cent .: 25.

No. B-4

(a) Description: ... Deep colony; very small; dense;

Nut like characteristic oblong shape.

(b) Relative per cent .: 70.

Note:-

General: No liquefaction. Very few molds. At least 90 per cent. acidifiers.

Quantitative Analysis.

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Content #5.
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Agar Plates. -A- 24 hours. 1 ----- solution 1,400,000 per c.c. 2,000 1 " " 1,144,000 per c.c. 4,000 1 " " 2,400,000 per c.c. 20,000 Average 1,648,000 per c.c. Agar Plates. -B-1 ----- solution 900,000 per c.c. 2,000 1 -----11 per c.c. 4,000 1 "" 1,500,000 per c.c. 20,000 Average 1,200,000 per c.c. Final Average on Agar 1,424,000 per c.c. Gelatin Plates. -A- 48 hours. 1 ----- solution 1,360,000 per c.c. 2,000 1 ---- " " 3,120,000 per c.c. 4,000 1 ----- " " 1,120,000 per c.c. 20,000 Average 1,833,333 per c.c. Gelatin Plates. -B-1 ----- solution 1,560,000 per c.c. 2,000 1 "" " 1,000,000 per c.c. 4,000 1 " 3,320,000 per c.c. 20,000 Average 1,960,000 per c.c. Final Average on Gelatin 1,896,666 per c.c.

Grand Average

1,660,333 per c.c.

300

Qualitative Analysis.

0

301

Content #5'.

No.	B-1	None	present
No.	B-2	None	present
No.	B-3	None	present

No. B-4

- (a) Description: ... Lactic acid.
- (b) Relative per cent .: 75.

No. B-5

(a) Description: ... Very large distinctly green

surface growth.

(b) Relative per cent .: 4.

No. B-6

- (a) Description: ... Radial surface growth.
- (b) Trace.

Quantitative Analysis.

Content #5'.

Agar	Plates.	-A-	24 hours.								
	1	solution	, too numer	0 us							
	1 2,000	11 II	9,480,000	per c.c.							
	1	11 11	10,800,000	per c.c.							
	10,000	Average	10,320,000	per c.c.							
Agar	Plates.	-B-									
	1 1,000 1	Solution, too numerous									
	2,000	n n	8,760,000	per c.c.							
	10,000	81 11	12,000,000	per c.c.							
		Arronaceo	10 380 000	202 0 0							

Final Average on Agar, 10,350,000 per c.c.

Gelatin Plates.

No Plates on Gelatin.

Average 10,380,000 per c.c.

0

Qualitative Analysis.

Content #5.

History Analysis.

No. B-1

(a) Description: ... White, round colony,

surface growth.

(b) Relative per cent .: 15 to 20.

No. B-2 None

No. B-3

(a) Description: ... Several molds, causing liquefaction

of gelatin.

(b) Relative per cent .: 1.

No. B-4

(a) Description: ... Small nut shaped colonies. Very

numerous and characteristic.

Lactic acid.

(b) Relative per cent .: 75.

363

Quantitative Analysis.

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Content #5.
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History Analysis

Agar Plates. -A- 16 days from last count. 1 ----- solution 180,000 per c.c. 1,000 1 11 ** ------176,000 per c.c. 2,000 1 530,000 per c.c. 10,000 Average 295,300 per c.c. Agar Plates. -B-1 ---- solution 280,000 per c.c. 1,000 1 -----2 18,000 per c.c. 2,000 1 11 ------170,000 per c.c. 10,000 156,000 Average per c.c. Final Average on Agar, 225,650 per c.c. Gelatin Plates. -A-1 ----- solution 249,000 per c.c. 1,000 1 200,000 per c.c. 2,000 ----1 " " contaminated 10,000 Average 224,500 per c.c. Gelatin Plates. .-B-1 ----- solution, liquefied 1,000 1 88 11 466,000 -----per c.c. 2,000 1 per c.c. 10,000 Average 553,000 per c.c. Final Average on Gelatin, 388,750 per c.c. 307,200 per c.c. Grand Average

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Popper Popper<							-	-	-	_					-	-	_		-	_	-	502
No. per C. C. Dismeter over 100 + <t< td=""><td colspan="13"></td><td></td></t<>																						
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Diameter over 100 * * * * * * * * * * * * * * * * * * *	No.	, pe	er c. c.																			
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Organism name.																						

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<u>SUMMARY</u>.

In making a summary of this work we find it necessary to make several statements concernings its inadequacy. In the first place the work was not as extensive as it should have been in order to give conclusive results. The material and conveniences were somewhat unsatisfactory. The starters used were poor; both containing plenty of acidifiers, but they were gas producers. In spite of these undesirable features the results obtained were good, and as far as they went, fairly conclusive.

Summing up the results of the experiment we shall take up each part separately. First, in the bacteriological analysis it was found that numerous species of bacteria were present in the original dream, but after pasteurizing the cream there was an enormous decrease. In each case, however, there was a larger decrease in the lactic acid groups than in other groups, showing that the lactic acid organisms were more readily killed by heat than are some of the other species. In each case the sweet cream butter contained a smaller percentage of organisms, especially those of the lactic acid group. After the butter had been kept in cold storage (50° F.) for sixteen days it was found that the sweet cream butter had increased over 150,000 bacteria per cubic centimeter, andthat the sour cream butter had decreased over 1,000,000 bacteria per cubic centimeter. In the case of the sweet cream butter the percentage of bacteria produced, other than lactic acid, was relatively large. In the sour cream butter practically all were lactic acid organisms. From the bacteriological standpoint, therefore, the results are greatly in favor of butter churned from ripened cream.

Second, after all bacteriological analyses of the different batches were made, a careful examination of the physical features was also made. The relative conditons of each batch of butter was noted and scored by Professor Kendall, head of the Dairy Department of the College, and by Professor Wilson, State Dairy Commissioner. ,10

The scores are as follows :---

Bat	ch	#1,	Sweet	cream	but	ter,	86%.
11	11	#2,	Sour	11	11	n	89.75%
11	ŧt	<i>#</i> 3,	Sweet	11	11	n	90%
11	11	#4	Sour	11	11	11	95.5%

The results of the scoring correspond very closely with the results obtained from the bacteriological content, as given above. From this it is evident that the butter made from churning cream ripened to .35% acidity gave much better results than that made from the cream which was not ripened at all.

As stated in the beginning, it was thought that if the cream was churned immediately after adding the starter, and before any ripening had taken place, a more complete control of the butter with reference to the flavor could be maintained. The results obtained, however, seem to point in the opposite direction. From the bacteriological standpoint, as has been formerly stated, the evidence is very conclusively in favor of thesour cream butter. The undesirable organisms decreased rapidly in the sour cream butter. In two weeks the total content of the sour cream butter decreased over 1,000,000 bacteria per cubic centimeter, while the sweet cream butter increased in total content. Not only this, but the results also show that the relative per cent. of undesirable organisms, gas produces, etc, in the sweet cream butter increased in relative per cent., while the undesirable organisms in the sour cream butter nearly disappeared, and the lactic acid organisms comprised nearly the entire per cent. It is thought from this, since the object in allowing organisms to grow in the butter is to impart to it a flavor that results from the presence of lactic acid, it is advisable to allow the organism to grow in the cream in order that the desirable acid producing bacteria will have abundance of milk sugar in which to develop; whereas, in the butter there is comparatively no milk sugar present in which the organism can develop. When there is an abundance of milk sugar the lactic acid organisms grow very rapidly and soon destroy all other organisms and ifinally produce so much acid

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that they themselves are destroyed. This, then, gives the ideal butter, i.e., butter with the desired flavor and without any destructive or undesirable bacteria present.

It is our conclusion, therefore, that the best method that can be followed in making butter is to ripen the cream to a relative low per cent. acidity, viz. .35%. This amount of ripening willgive the lactic acid organisms such a start that they can soon destroy the large majority of other bacteria, and produce the desired flavor.