

INAL ARTICLE

MINATION OF BACTERIOLOGICAL AND
SICAL QUALITY OF RAW MILK AND LOCALLY
TLED MILK IN CAFETERIAS OF JIMMA TOWN,
THWEST OF ETHIOPIA

se Sisay¹ (BSc), Esayas Alemayehu² (MSc), Eshetu Gizaw² (BSc)

RACT

GROUND: Food – borne diseases are major public health concern worldwide. people around the world acquired food poisoning due to consumption of raw, inated milk which spread either from infected cows, by handling, or during milk ing. This study aims to examine the bacteriological quality and adulteration of k and locally bottled milk in Cafeterias of Jimma Town.

ODS: Laboratory-based cross-sectional study was conducted in February 2004. six Cafeterias were selected randomly. Hygienic practice and sanitary condition terias were assessed using observational checklist. Raw milk samples were d from Cafeterias bulk (storage) after mixing with sterile plunger, using sterile apped container with 250 ml capacity. One bottle of unopened locally bottled as selected randomly from the shelf and transported using an icebox to mental health laboratory, Jimma University. Nutrient agar, Colombia agar and e water were used to grow total microbes. Louryl sulphate tryptose broth with ent pad and membrane filter were used to determine coliforms. Lactometer and meter were used to measure the specific gravity and temperature of milk ively.

TS: All the milk handlers have no health check up and certificate. Among the ndlers 53.8% wash their hands before and after break and after visiting toilet, .7% only practice hand washing after coughing and sneezing. 80.8% of raw milk were found in poor or grade C quality, while 19.2% were grade B. Among bottled milk samples 9% were grade B while 91% were grade C. Among various ources, majority of them are obtained from individual breeders 5 (84.6%). m count of raw milk was found that 73.1% were grade B – and 23.1% grade C. of locally bottled milk were grade C. 65.4% of raw milk samples were adulterated ater, most of them were collected from individual breeders. Adulteration was cantly associated with microbial count ($\chi^2 = 5.787, P < 0.025$) this indicates ated milk was found with high microbial count.

CONCLUSION: Consumption of locally bottled milk is not advisable. Thus cafeteria owners should avoid preparation of locally bottled milk, unless proper boiling and bottling is practiced with a license of professional. Intensive health education should be given about hygienic practice to the workers and owners. The concerned authorities should strengthen technical supervision and design strong rule for health check up and certification of workers.

KEY WORDS: bacteriological quality, bottled milk, adulteration, and cafeterias.

INTRODUCTION

Milk is an ideal medium for growth of microorganism. It is highly perishable food because it contains nutrients required for microbial growth, such as, protein (3.5%), fat (3.7%), carbohydrate (4.9%), minerals (0.75%) and water (87.2%), (1,2,3). It is also an excellent culture media for many kinds of microbes, being high in moisture content, nearly neutral pH (6.5 – 6.6) and rich in microbial food; which may easily become a source for the spread of pathogens or a good medium for bacteria and easily adulterated with out any color change (4).

For a number of decades, WHO has recognized the wide spread nature of food borne diseases and their impact on communities in both the developing and developed world. The annual incidence of some 1.5 billion episodes of diarrhea in children under 5 years of age and more than 3 million resultant deaths are an indication of the magnitude of the problem, as a significant proportion of the diarrheal disease cases are food borne in origin (5).

In recent year, a number of industrialized countries have even experienced a significant increase of food borne diseases. In several countries surveys have pointed out to an annual incidence of 5 – 10% of the population involved (5, 6). Even the United States has the safest supply of all nations, however, it is estimated 25 million food borne illness cause and 16,000 deaths occur each year. From this approximately 66% of all food

bacterial pathogen (7) while in developing countries like Ethiopia, the reported food borne cases may be as low as 1%, because individual cases or small out breaks of food borne disease generally remain unnoticed or may not be reported (5,6).

The problem in Ethiopian is not an exception. There was a study done on bacteriological quality of raw cows milk obtained from four dairy farms and a milk collection center in and around Addis Ababa by Godefay and Molla (8). This study showed that milk samples from collecting buckets in the dairy farms had a count of 1×10^5 colony forming units (CFU)/ml, those from storage container had count of 1×10^6 CFU/ml and the count reached 1×10^8 c. f. u/ml up on arrival at the processing plant. The mean coliform counts ranged from 1.3×10^4 CFU/ml (storage container before cooling) to 7.1×10^4 CFU/ml can arrival at the processing plants. The hygienic quality of raw milk from the collection center was poor with a mean total bacterial count of 1.3×10^7 c.f. u/ml. Lack of knowledge about clean milk production, use of unclean milking equipment and lack of potable water for cleaning purposes was some of the factors which contributed to the poor hygienic quality of raw milk in the study farms (8).

An early study conducted by Asenafi and Mogesie (9) on college of dairy farm in Awassa compared the microbial load of milk directly obtained from the udder and fresh milk from utensils. This study revealed that aerobic mesospheric counts for udder milk ranged between 10^3 – 10^4 c.f.u/ml. where as, raw milk from

collecting utensils had counts as high as 10^6 c. f. u/ml. Coliform counts in udder milk was about 10 c.f.u/ml, while this count in fresh milk (raw) collected from utensils was as high as 10^5 c.f.u/ml indicating that major contamination occurred during the milk collection process.

In Ethiopia, raw milk, and milk products are frequently consumed in different establishments and individuals home. But the hygienic status or quality of milk and the prevalence of milk – related out breaks was not well assessed. Different studies were conducted on bacteriological quality of milk on dairy farms of Addis Ababa, Awassa and in Jimma (8-10). However no study was carried out at food and drinking establishments on bacteriological quality of foods that are consumed by the peoples for immediate consumptions or stayed for longer time.

This study assessed the bacteriological quality of raw milk and locally bottled milk and adulteration of these products. It helps to give clue for the quality of milk served in cafeterias and for other researchers to study the bacteriological quality of any food items in different food serving areas. In addition, it may initiate the local government as well as the country at large to give priority attention to modernize or alter the existing hygienic condition of food and drinking establishments, which are the potential sources of food borne diseases.

METHODS AND MATERIALS

Study Design: Laboratory based cross-sectional study was carried out to examine bacteriological quality and adulteration of raw milk and locally bottled milk in 26 cafeterias in February 2004. It was conducted in Oromiya regional state Jimma special zone, Jimma town, which is found 335 km from Addis Ababa, South west of

Ethiopia. The study variables include; total colony count, coliform count, specific gravity of milk, source of milk, ownership of cafeterias, type of container used, educational level of workers, and hygienic practice of milk handlers.

Sampling Technique: 26 cafeterias were selected by applying the population correction formula from municipality documentation and stratified based on ownership status; institutional and private cafeterias, then study samples were selected using simple random sampling technique.

Milk sample: Milk sample was collected from the selected cafeterias.

1. Locally bottled milk – 1 bottle of unopened sample was selected randomly from the stack or shelf.
2. Raw milk – sample was taken from the bulk or tank in sufficient amount.

Raw milk samples from the bulk were collected after thoroughly mixed with a sterile plunger. Sample was collected below the surface with a sterile dipper and aseptically poured into a screw – cupped container of a capacity 250 ml.

The samples were transported using icebox with a suitable temperature ranges (0 – 4°C) to environmental health laboratory, Jimma University and examined within 24 hours (11).

Data were collected using observational checklist. Data regarding laboratory results was collected on daily basis at the end of each laboratory examination of milk.

Colony Count (plate count): Colony count is used for the determination of the quality of raw and pasteurized milk, it assesses the number of viable bacteria in milk.

A sample of 1ml milk was diluted with sterile distilled water in 1: 10, 1: 10^2 , 1: 10^3 and 1: 10^4 ranges and these were inoculated to Nutrient agar and Colombia agar and then incubated at 35°C to 37°C for 48 hours or 72 hours.

Determination of milk quality was made based on standard maximum acceptable level of colony count and *E. coli* count based on most probable number result.

Specific Gravity: Specific gravity of milk was measured using lactometers; to determine adulteration of milk with water. Specific gravity is usually taken at 15.5°C of milk temperature. If possible correct to this temperature (13). This temperature of milk was maintained using refrigerator and evaporative food cooler.

All data collected by observational checklist were checked for the completeness and fulfillment daily. The data were processed manually using scientific calculator and analysis was done using descriptive and analytical methods.

RESULTS

Hygienic Practice: The study revealed that, there was no regular health checkup to milk handlers and no workers found with health certificate. Twenty two (84.6%) of the milk handlers wear protective cloth, of these 20 (76.9%) use outer garment, 2 (7.7%) use hair cover, and the rest 4 (15.4%) never use protective cloth.

Among the milk handlers 18 (69.2%) wear their protective cloth always, 4 (15.4%) wear sometimes and another 4 (15.4%) do not use at all.

Regarding hand washing habit 10 (38.5%) wash their hands before and after work, 14 (53.8%) wash their hands before and after break and after visiting toilet, the remaining 2 (7.7%) practice hand washing after sneezing and coughing.

Twenty one (80.7%) of milk handlers keep their fingers clean and trimmed and 5 (19.3%) do not. All the studied cafeterias have access to pipe water, which is from the public distribution system of the town. Equipment-washing activities of cafeteria were found that 4 (15.4%) use warm

water, 20 (76.9%) use soap and water, while the remaining 2 (7.7%) use only cold water. Among the studied cafeterias, 23 (88.5%) use sink and the rest 3 (11.5%) use bowl to wash equipments.

The current condition of milk storage materials found that 15 (57.7%) clean, 9 (34.6%) stained and the rest 2 (7.7%) rusted.

Milk: From a total of 26 cafeterias studied, 22 (84.6%) rented milk from individual breeders, 3 (11.5%) use their own cattle and 1 (3.8%) obtains from market.

The study findings indicated that all cafeterias sale raw milk. Among the cafeterias studied 11 (42.3%) were selling locally bottled milk, while 5 (19.2%) cafeterias were selling Yoghurt and 1 (3.8%) cafeteria was selling powder milk.

These cafeterias use different kinds of storages or raw milk collected from sources. Among the storage containers used bucket were 12 (46.15%), Plastic were 10 (38.46) while Tin cans, which used least, were 4 (15.38%). Methods of drawing milk from the bulk is by pouring (50%) and dipping (50%). Availability of functional refrigerators in cafeterias is shown in Figure 1.

Laboratory Analysis: Examination of microbial quality of raw milk and locally bottled milk were done on 26-raw milk and 11 locally bottled milk samples from 26 cafeterias of Jimma town.

Among the studied samples of raw milk colony count, no one was found in grade A standard, 5 (19.2%) were found with fair quality (grade B) and the rest of the majority 21(80.8%) were in poor quality (grade C), based on American Public Health Service Standard (6). Out of 11 locally bottled milk samples only 9% were found in grade B and the rest 10 (91%) were poor quality (grade C) (Table 1).

In addition all samples were analyzed for coliforms, of which 3.8% were found in

good quality, which is less than 100 C.f.U/ml of milk, 19 (73.%) were found in fair quality and the rest 6 (23.1%) were poor quality (Table 2).

Coliform test of locally bottled milk revealed that 2 (18.18%) of samples have

good quality and the rest 9 (81.81%) have poor quality of milk, which have a colony count of greater than 10 C.f.U/ml of milk (Table 2).

Table 1. Microbial growth in standard plate count of milk sample in cafeterias of Jimma town, February 2004

Type of milk sample	Colony count (Cfu/ml of milk)	Number of cafeterias	Percent (%)
Raw milk	$\leq 2 \times 10^2$	0	0
	$2 \times 10^5 - 1 \times 10^6$	5	19.2
	$\geq 1 \times 10^6$	21	80.8
	Total	26	100
Locally bottled Milk	$\leq 3 \times 10^4$	0	0
	$3 \times 10^4 - 5 \times 10^4$	1	9
	$\geq 5 \times 10^4$	10	91
Total		11	100

Table 2. Coliform count in milk samples of cafeterias in Jimma town, February 2004.

Type of milk Sample	colony count per/ml of milk	Number of cafeterias	Percent (%)
Raw milk	< 100	1	3.8
	101-2000	19	73.1
	≥ 2000	6	23.1
	Total	26	100
Locally bottled Milk	< 10	2	18.2
	> 10	9	81.8
Total		11	100

Score: Colony count $\leq 2 \times 10^5$ c. f. u/ml is Grade A, Good Quality

$2 \times 10^5 - 1 \times 10^6$ c. f. u/ml is Grade B, Fair quality

$\geq 1 \times 10^6$ c. f. u/ml is Grade C, poor quality

The same grading system is used for figures in coliform count table i.e.

< 100 Grade A, good quality

101 - 2000 Grade B, fair quality

≥ 2000 Grad C, poor quality

The American public health service standard for milk and milk products (5, 6, 12).

Specific gravity: A total of 37 raw milk and locally bottled milk samples were collected from 26 cafeterias. Specific gravity of milk was measured using lactometer to determine whether the milk is adulterated with water or not. Out of 26 raw milk samples 17 (65.38%) were found adulterated and the rest 9 (34.6%) were

found in the normal range of milk specific gravity (Table 3).

Out of 11 locally bottled samples 5 (45.45%) were found adulterated and 6 (54.54%) were in the normal range.

Specific gravity less than 1.027 is adulterated-normal range of milk specific gravity is between 1.027 and 1.035 (5,6,12).

As shown in table 4, there is a statistical significant association between adulteration and colony count.

Table 3. Specific gravity of milk samples in cafeterias of Jimma town February, 2004

Sample Milk	Specific gravity	Number of cafeterias	Percent
Raw milk	< 1.027	17	65.4
	1.027-1.035	9	34.6
	>1.035	-	-
	Total	26	100
Locally bottled Milk	<1.027	5	45.5
	1.027-1.035	6	54.5
	>1.035	-	-
Total		11	100

Table 4. Association of milk Specific gravity (adulteration) and colony Count in cafeterias of Jimma town, February 2004.

Specific gravity of milk	Colony count/ml		Number of cafeteria	χ^2
	$2 \times 10^5 - 1 \times 10^6$ c.f.u/ml	$\geq 1 \times 10^6$		
<1.027	1(3.26)	16(13.7)	17	$\chi^2 = 5.787$ $P < 0.025$ $df = 1$
1.027-1.035	4(1.7)	5(7.27)	9	
Total	5	21	26	

χ^2 Calculated = 5.787
 χ^2 Tabulated = 3.841

df = 1
P < 0.025

DISCUSSION

This study was made in cafeterias of Jimma town, and the quality of raw and locally bottled milk is unsatisfactory. According to this study, the workers who wear protective cloth were 84.62% and the rest 15.38% do not have protective cloth. This may indicate a better awareness of food and drink establishment owners to provide protective cloth to their workers. Among the milk-handlers only 7.7% was wash their hands after coughing and sneezing. This may affect the bacteriological quality of milk, because the quality of milk was found unsatisfactory. The study identified that; there was no regular health check up to milk handlers, and no continuous follow up towards the quality of milk by the responsible authorities. Probably, this is the main contributing factor for milk quality deterioration. Most of the time milk is contaminated with external source at different processes by micro organisms directly from the milk handlers, who have direct or indirect contact with milk, especially if the milk handlers are in the process of shedding pathogenic organisms, during sneezing, coughing, scratching and from body surface in contact with milk (7). Personal cleanliness is necessary particularly during milking, processing and distribution.

According to Ehlers (13), when milk leaves the cow's udder it contains some bacteria, ordinarily, harmless, but if the animal is diseased some may be pathogens. The other bacteria enter the milk from the air, dust, the milkier hands and surface of milk storage container (3). Similarly from milk storage containers 42.3% were not clean. Probably the unhygienic condition may be the cause to poor quality of milk in cafeterias.

The bacteriological standards of dairy products established or developed by united states public health service in 1978,

indicated that, the plate count of grade A raw milk should be less than 2×10^5 c.f.u/ml, grade B is between 2×10^5 c.f.u/ml to 1×10^6 c.f.u/ml and grade C is greater or equal to 1×10^6 c.f.u/ml of milk (3, 14). Based on this standard, all samples of raw milk were not found in grade A, while 19.2% were grade B and almost all in all 80.8% were in grade C. A similar study done by Godefay and Molla in and around Addis Ababa on storage containers was found 4×10^6 c.f.u/ml (18). Another study by Tadesse *et al.*, shows that the microbial growth was higher in milk containers and bulk cans than in the freshly drawn milk. This indicates that milk handlers as well as milk storage materials, which have direct physical contacts, are most likely to be a source of contamination. So milk utensils and equipments should be designed and cleaned properly to avoid the favorable conditions for microbial growth.

Regarding the quality of locally bottled milk found that, a considerable number (i.e. 91%) was poor with a colony count of greater than or equal to 5×10^4 c.f.u. /ml. This may be due to improper boiling and bottling procedures during preparation. Coliform test for dairy products are not intended to indicate fecal contamination, but to reflect over all dairy farm and distribution place sanitation. The analysis of the study showed that 3.8% of samples were in good quality (i.e. < 100 c.f.u/ml), 73.1% were fair and the rest 21.1% were poor quality. The study is almost similar to the finding of Godefay and Molla in and around Addis Ababa. Over all, it indicates that, coliform counts are much greater than the standard set by APHS (American Public Health Service Standard) (6). This may be due to poor hygienic practice of milk handlers, absence of health check up and usage of stained and unclean storage materials. In addition, milk source of most cafeterias are individual breeders, because 90.9% of milk

were found high in microbial count and 65.4% were adulterated. Whereas, milk collected from their own cattle were in good quality and normal range of specific gravity.

Almost all, 81.8% of locally bottled milk samples were poor quality i.e. > 10 c.f.u/ml, coliform. The presence of coliform indicates that, either improper boiling and bottling or contamination after processing by human or other warm-blooded animal or both has occurred. Probably, this contamination may be from milk handlers during preparation and storage of the milk after boiling.

From the study samples of raw milk 65.4% was adulterated (i.e specific gravity < 1.027) and the rest 34.6% was found in the normal range of specific gravity (i.e. between 1.027 and 1.035). Adulteration with water may be done intentionally for profit or the cafeteria owners may not be aware about the quality of milk rented from individual breeders. Because, 22 (84.6%) of the milk collected from individual breeders were found adulterated (65.4%).

Accordingly, the study indicated that there was a statistical significant association between adulteration and microbial count ($\chi^2 - 5.787, P < 0.025$). The high increment of colony count is due to adulteration with water. Whether the water is clean or not, and contaminated by milk handlers during adulteration may affect the quality. Whereas milk collected from their own cattle was found that all in normal range of specific gravity and grade B. microbial quality.

CONCLUSION AND RECOMMENDATION

Based on the American Public Health Service Standard accepted in many countries of the world, the microbial quality of milk in Jimma is unsatisfactory. Consumption of raw milk and locally

bottled milk from cafeterias is not advisable because both are found, high in microbial count, coliform count and adulterated with water. Especially milk rented from individual breeders is more adulterated and poor in terms of microbial quality.

Locally bottled milk is not guaranteed in its quality as a bailed and bottled product.

Based on the findings of the study the following recommendations are forwarded.

□ The concerned authorities, municipality and local health authorities need to strengthen technical supervision on cafeterias service and frequent inspection should be practiced.

□ Cafeteria owners should to avoid preparation of locally bottled milk, unless properly boiled and bottled with a license and approval of qualified personal.

□ Intensive health education should be given both to the owners and the workers about good hygienic practice and the effect of unhygienic condition.

□ Ministry of health must establish strong rules for any public eating and drinking establishment workers to have periodical health check up.

ACKNOWLEDGEMENTS

We would like to express our appreciation to Jimma University for funding this project. We also express our gratitude to the Jimma community who participated in the study.

REFERENCES

1. Teka, GE. Food hygiene. Principles and methods of food-borne diseases control with special reference to Ethiopia, 1997, Addis Ababa, Ethiopia.
2. National Research Council. Sanitary milk control and its relation to the sanitary, nutritive and other qualities of milk, USA, 1953.
3. Ehlers MV. , Municipal and rural sanitation, TMH edition, New Delhi, 1982 PP.183 – 224.
4. Fraizer, WC. Food microbiology 3rd ed: Westhoff editors, New York, 1978.
5. WHO. Global estimation of food-borne disease. World Health statistics quarterly report, WHO, Geneva: 1997 Vol. 50 (1) Pp. 14-15.
6. WHO. Food contamination and diarrhea World Health Magazine, the WHO Geneva, 1990 (Jan. – Feb.): 18 – 19.
7. Marriot NG, Principles of food sanitation, 3rd ed. Chapman and hall. New York, 1995.
8. Godefay B, Molla B. Bacteriological quality of raw Cow's milk from four dairy forms and a milk collection center in and around Addis Ababa. Berliner and Mucnchener Tieraerztliche Wachschrift: 2000 113 (7-8): 276 – 8.
9. Mogesie A. Bacteriological quality of milk. *Ethiop J Health Sci.* 2002; 25 (1).
10. Getahun T, Gebre-Selassie S. Assessment of the bacteriological quality of milk at dairy farms and individual breeders in Jimma town, Southwest Ethiopia. *Ethiop J Health Sci.* 2003; 13: 21 – 29.
11. Collee JG. Fraser AG, Marmino BP, Simmons A. Practical Medical Microbiology. 14th ed. Edinburgh, Churchill, living stone: 1999 pp 896.
12. Cheesbraugh M. Microbiology: Medical Laboratory manual for tropical countries: 1984 Vol. 2: pp 219 – 220.
13. Ecklescombs M. Milk and milk products. Mc Grow-Vill Inc. New York, 1982 pp 69- 85.
14. John ML. Public Health and Preventive medicine, 12th ed., USA, 1995; Pp 771–772.