

# Protease and Lipase Activities of *Pseudomonas Spp.* Isolated from Pasteurized Milk

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## ABSTRACT

Raw milk deteriorates in only a few days even when stored under refrigeration temperatures. Moreover, pasteurized and refrigerated milk has a shelf life of 7 to 10 days. Psychrotrophic bacteria have been recognized as a recurring problem in refrigerated storage and distribution of perishable food products. This study was performed to isolate, characterize and evaluate the prevalence of *Pseudomonas spp.* in pasteurized milk. Thirty five pasteurized milk samples were collected aseptically from different dairy shops, supermarkets and groceries in different areas in Sadat city, Menuofiya Governorate, Egypt and analyzed for total viable count and *Pseudomonas spp.* occurrence. *Pseudomonas aeruginosa* found to be the predominant from the *Pseudomonas spp.* followed by *Pseudomonas fluorescens*. The total viable count (cfu per ml) average  $\pm$  SD were  $3.50 \times 10^4$  cfu/ml  $\pm$  0.085. Out of ten *Pseudomonas* species isolates were examined to its proteolytic and lipolytic activity, two isolates had proteolytic activity, while three had lipolytic activity and five isolates had both.

**Key words:** *Pseudomonas aeruginosa*, *Pseudomonas fluorescens*, proteolytic and lipolytic activity.

## INTRODUCTION

Milk has many nutritious qualities that make it an important part of children's diet. To produce the best quality milk and to achieve all the nutritious benefits of it, the highest quality raw milk must be obtained. Pasteurization and shelf stable milk products through Ultra High Temperature (UHT) continuous flow sterilization, are available (Goff and Griffiths, 2006). The conditions of heat treatment used for pasteurization depend on the final product, lower temperatures are used for refrigerated products and higher heat treatments are used for products stored at room temperature (USCFR, 2006). Psychrotrophic bacteria have been recognized as a recurring problem in the refrigerated storage and distribution of fluid milk and perishable dairy products for several decades. So, the psychrotrophic have received increased attention by investigators during recent years, because modern developments in the handling and transportation of milk have resulted in milk being held for longer period at refrigeration temperature before processing, manufacture or consumption.

Some psychrotrophic bacteria may grow at a temperature of 7°C although their optimum temperature is higher. Rapid cooling and cold storage of raw milk favor the growth of psychrotrophic bacteria in milk (Barbano *et al.*, 2006). They become dominant microflora during cold storage of milk and their extracellular enzymes, particularly proteases and lipases, contribute to the spoilage of milk products (Hantis-Zacharov and Halpern, 2007). Microbial growth and metabolism shorten the shelf life of milk by producing undesirable changes in aroma and taste attributes that influence consumer acceptability of the products (Fromm and Boor, 2004). Factors limiting milk stability are well established: bacterial contamination, inadequate packaging system and improper temperature control. Vulnerability of milk's fat and protein to physical-chemical alterations can also lead to deterioration, thus reducing its quality. Cromie (1991) reported the factors that influence the shelf life of pasteurized milk include the quality of the raw material, the binomial temperature/time pasteurization, resistant microorganisms to pasteurization (particularly psychrotrophics), the presence and activity of post pasteurization contaminants, the packaging system and storage temperature post pasteurization which had the greatest impact on the stability of the product. Shelf life of pasteurized milk. Extra cellular lipolytic enzymes produced from psychrotrophics can produce deteriorate effects as rancid flavours and odours in milk and dairy products that make a subsequent reduction of shelf-life and the products become unacceptable to consumers (Downey, 1980). The proteolytic enzymes produced by psychrotrophics in milk are more powerful in its action on milk proteins than that naturally present in milk and that produced by leucocytes even if present by great amount (Grieve and Kitchen, 1985). *Pseudomonas spp.* produces a large number of extracellular toxins, which include phytotoxic factor, pigments, hydrocyanic acid, proteolytic enzymes, phospholipase and enterotoxins. Exotoxins are the most factor responsible for *pseudomonas spp.* pathogenicity because it can produce leucopenia, circulatory collapse, necrosis of liver, pulmonary odema, hemorrhage and kidney tubular necrosis. The enterotoxin produced is responsible for diarrhea disease. Higher temperature for shorter times were less effective

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in destruction of the *Pseudomonas* spp. than that of temperature for longer times because the condition created in the milk at high temperatures may favor of injured cells (Cousin,1982). Generally they are considered incapable of surviving pasteurization and they may be post-processing contamination. Although, they can cause gastroenteritis if ingested in large number(>10<sup>6</sup>), but also the food would be clearly spoiled before such numbers would reached (Johnson,1990). Moreover, its presence in milk and its products was considered as a possible indicator of fecal contamination. Also *Pseudomonas aeruginosa* has been implicated in epidemics of moderate to severe diarrhea in children in the form of enteric fever (Todar, 2002).

This study aimed to explain the occurrence of *Pseudomonas* species contamination and observation of the proteolytic and lipolytic behavior of it in the pasteurized milk sold in Sadat city, Menuofiya Governorate, Egypt.

## MATERIALS AND METHODS

Thirty five samples of pasteurized milk were collected randomly from various local milk supermarket in different areas in Sadat city, Menuofiya Governorate, Egypt. Collected samples were examined for total viable cells and bacteriologically for occurrence and behavior of *Pseudomonas* organisms. Psychrotrophic isolates were obtained from pasteurized milk by pour plating appropriately diluted sample using nutrient agar. Incubation was done at 7°C for 10 days (Cousin *et al.*, 2001). The psychrotrophic isolates were subjected to biochemical characterization after activating the culture in nutrient broth at 37°C for 24 h (Barrow and Feltham,1993).

### Identification of *Pseudomonas* isolates

The representative suspected colonies were purified and then identified according to Bergey's Manual of Systemic Bacteriology (1984) and API 20NE.

### Determination of the proteolytic and lipolytic activities of *Pseudomonas* species

Ten strains belonging to *Pseudomonas* species isolated previously from pasteurized milk samples were investigated for their proteolytic and lipolytic activities as described by Harrigan and McCance (1976).

### Preparation of isolates

*Pseudomonas* isolates were subcultured onto nutrient agar plates and incubated at 30°C for 24 h. Pure cultures were inoculated into nutrient broth and incubated overnight at 30°C prior to testing.

### Proteolytic activity using skim milk agar

Overnight cultures were spot inoculated onto milk agar, standard plate count agar supplemented with 10%

sterile skim milk. The inoculated plates were incubated at 20°C and 4°C for 10 days. The presence of transparent zones around the spots was recorded as positive strains referring to protease production, and subsequently flooded with 10% v/v acetic acid solution. Clear zone around the colonies after one minute exposure were regarded as positive (Harrigan and McCance,1976).

### Lipolytic activity

Lipolytic counts (LP) were determined using nutrient agar (NA) containing tributyrin. The medium was prepared 10 g of tributyrin and 28 g of NA. Plates were incubated at 20°C and 4°C for 72 h determine viable lipolytic counts, and lipolytic activity was determined by measuring clear zone around each colony (Harrigan and McCance,1976).

## RESULTS AND DISCUSSION

### Incidence of *Pseudomonas* spp. among the examined samples:

The samples showed an average  $\pm$  SD of total viable count of  $3.50 \times 10^4 \pm 0.085$  cfu/ml. According to Prevention of Food Adulteration Act (PFA, 2007) total plate count of pasteurized milk should not exceed 30,000 cfu/ml. However it was observed that all the standard plate count plates were characterized by the predominance of typical small circular entire smooth and shining uncolored colonies. Furthermore, only this type of colony was evident in the plates used for taking psychrotrophic count (Table 1). The *Pseudomonas* spp. count (cfu per ml) average  $\pm$  SD was  $1.5 \times 10^2 \pm 0.01$  among different samples, and the isolates were subjected to bacteriological examination (Table 2) such as lactose negative and blue-black and greenish bright dark colonies are lactose positive as a result of incubation during 48 hours at 30°C. Among these, with the lactose negative ones are continued to be studied on and these colonies were applied oxidation-fermentation test. *Pseudomonas* react oxidative. Among the same samples catalase(+), motility(+) and oxidase(+) colonies were studied on growth at 4°C and 41°C have been the first taken into consideration. So, *Pseudomonas*, which incubated into yeast extract medium were left at 4°C for 7-10 days and at 41°C for 24 hours respectively (Mickova *et al.*,1989).

The results showed that 10 samples were positive for occurrence of *Pseudomonas* spp., with a percentage of 26.7%. Identity of the isolates were confirmed by using API 20NE for detection and isolation of *Pseudomonas* spp. as *Pseudomonas aeruginosa* (7 isolates) and *Pseudomonas fluorescens* (3 isolates). Many investigators indicated that some *Pseudomonas* spp. could survive heat treatment used in pasteurization of

**Table 1. Prevalance and count of *Pseudomonas spp.* in examined pasteurized milk samples**

No. of samples	Positive Samples		Min.	Max.	Mean $\pm$ SD*
	No.	%			
35	10	28.5	1.5 $\times$ 10	2.5 $\times$ 10 <sup>3</sup>	1.5 $\times$ 10 <sup>2</sup> $\pm$ 0.01

\*SD= Standard deviation.

**Table 2. The Bacteriological Examination of *Pseudomonas spp.* In examined pasteurized milk samples**

No. of isolates	Catalase	Oxidase	O/F	Lactose Fermentation	Motility	MR	VP	Citrate Utilization	Gelatin Hydrolysis	Indol Production	Nitrate Reduction
1	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
2	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
3	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
4	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
5	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
6	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
7	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
8	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
9	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+
10	+	+	O <sup>+</sup> /F <sup>-</sup>	-	+	-	-	+	+	-	+

milk (Abad *et al.*,1993). Incidence of *Pseudomonas aeruginosa* in pasteurized milk has been reported by Kumaresan and Annal Villi (2008). According to them, post processing contamination led to the predominance of heat sensitive *Pseudomonas spp.* in pasteurized milk.

Quality defects in pasteurized milk are most often the result of microbial contamination, growth and spoilage. Microbial defects usually become evident in the finished product through shelf-life evaluations or consumer complaints. Post-pasteurization contamination with psychrotrophic spoilage bacteria is most detrimental. In most cases, product contamination is the result of insufficient cleaning and sanitation of the processing equipment and plant environment. Practices that are being followed in the dairy processing unit would have singled out psychrotrophics as the major category of spoilage organism (Alatossava and Alatossava, 2007).

### The proteolytic and lipolytic activities of *Pseudomonas* species

The high spoilage potential of *Pseudomonas* species is not only because of its ability to multiply at refrigeration temperatures but also because of their ability to produce thermostable proteases and lipases (Sorhaug and Stepaniak, 1997).The importance of

proteases and lipases, which causes bacterial virulence, were proven in several studies. However, there was less study in the protease and lipase activities of *Pseudomonas spp.*

This study showed that the *Pseudomonas spp.* isolates are not showed proteolytic and lipolytic activities at 4°C, while at 20°C the *Pseudomonas aeruginosa* and *Pseudomonas fluorescens* isolates shown both of these activities which illustrated in (Table 3) .Two isolates were positive for proteolytic ; three lipolytic and five isolates for both proteolytic and lipolytic. This finding agrees with the findings of Dogan and Boor (2003) and Alatossava and Alatossava (2006) .

Psychrotrophics were implicated in many defects in milk and dairy products which are a problem resulting from prolonged refrigeration storage (Swart *et al.* 1989; Garg, 1990) and distribution of perishable food products. *Pseudomonas* species have been implicated in the spoilage of processed milk kept under chilled condition because of their capacity to multiply under refrigeration with the production of thermostable proteases and lipases (Rajmohan *et al.*, 2002) so *Pseudomonas spp.* plays an important role in milk spoilage.

**Table 3. Proteolytic and lipolytic activities of *Pseudomonas spp.* isolated from pasteurized milk samples**

<i>Pseudomonas</i> isolates	No.of tested isolates	Positive proteolytic isolates	Positive lipolytic isolates	Positive proteolytic & lipolytic isolates
<i>Pseudomonas aeruginosa</i>	7	1	2	4
<i>Pseudomonas fluorescens</i>	3	1	1	1
<b>Total</b>	10	2	3	5

### Conclusion

The previous information indicated that psychrotrophs are still inevitable because they are widely distributed in nature, withstand sanitizers and can liberate heat stable enzyme causing spoilage of food and some of them considered as food borne pathogens. This knowledge increasing attention toward the way by which the restriction of these microorganisms must be done. In this study the results cleared that after pasteurized 10 samples from 35 samples were positive for occurrence of *Pseudomonas* spp. , with a percentage of 26.7% that may be due to insufficient pasteurization or the post contamination by this genus and the temperature which milk and dairy products are exposed influences the type of microorganisms that will grow in them. As dairy equipment and utensils constitute the major source of many types of psychrotrophs in milk, so special attention should be considered in their cleaning and sanitation to produce milk of low bacterial count or even completely free of psychrotrophs bacteria.

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## الملخص العربي

### النشاط التحليلي للبروتين و الدهن لبكتريا *Pseudomonas spp.* المعزولة من اللبن المبستر

هدى محروس

ومن النتائج المتحصل عليها تواجد بكتريا *Pseudomonas spp.* في 10 عينات من العينات تحت الدراسة وفيها كانت بكتريا *Pseudomonas aeruginosa* هي السائدة ويليهما بكتريا *Pseudomonas fluorescens* وكان متوسط العد الكلي  $\pm 0.085$  مستعمرة/مل. و بدراسة النشاط التحليلي للبروتين والدهن لهذه العشرة عزلات التابعة لجنس *Pseudomonas spp.* وجد عزلتان موجبة لتحلل البروتين فقط و 3 عزلات محللة للدهن فقط و 5 عزلات محللة للبروتين و الدهن معا.

اللبن الخام يتعرض للفساد خلال أيام قليلة حتى لو حفظ في ظروف حرارة التبريد. أما البسترة والتبريد يعطي اللبن فترة حفظ أطول لمدة تصل إلي 7-10 يوم وتمثل البكتريا المحبة للبرودة المشكلة الأساسية في اللبن المبستر.

تهدف هذه الدراسة إلي عزل و تقدير البكتريا المحبة للبرودة في اللبن المبستر وفيها تم تجميع 35 عينة لبن مبستر من محلات الألبان و السوبر ماركت من مدينة السادات-محافظة المنوفية-مصر و تم تقدير العد الكلي المتواجد فيها وتقدير بكتريا *Pseudomonas spp.*