Microbiological Quality of Asadero Cheese Manufactured with a Plant Based Coagulant from Solanum elaeagnifolium

Nina del Rocío Martínez-Ruiz, Sandra Flores Enriquez, Rosa Emma Vázquez-Nájera, José Alberto López-Díaz

Departamento de Ciencias Químico Biológicas, Instituto de Ciencias Biomédicas, Universidad Autónoma de Ciudad Juárez, Ciudad Juárez, México.

Email: joslopez@uacj.mx

Received March 28th, 2013; revised April 29th, 2013; accepted May 7th, 2013

ABSTRACT

In Chihuahua, north Mexico, the berries of Solanum elaeagnifolium (trompillo or silverleaf nightshade) have been used in the manufacture of artisanal filata-type asadero cheese. Solanum elaeagnifolium is a wild plant that possesses proteases in its fruit; those enzymes exhibit general proteolytic activities, which are useful in traditional asadero cheesemaking as a rennet substitute. These type of cheese is softer than those made with rennin or chymosin due to their water content and proteolysis. Thus, the aim of this work was to explore microbiological susceptibility of asadero cheese made with an extract from ripe berries and compared with those elaborated with commercial rennet. Crude extract of such fruit was obtained by salting out with ammonium sulfate (40%). Cheeses were obtained by a standardized process only changing the enzyme, packaged in plastic bag or vacuum bag and storage at 4˚C - 6˚C. The study included the analysis of spoilage bacteria (total aerobic count, yeasts and molds), indicators of hygiene (total coliforms and E. coli) and fecal contamination (fecal coliforms) after 1, 7, 14, 21 and 28 days of production. Although cheese made with S. elaeagnifolium had lower microbial growth, the aerobic count and yeasts and molds count of all cheese samples showing a 28 days shelf life. Total coliforms, in limits beyond the established ones by the Mexican legislation for human consumption with no effect of the enzyme type used for production, as well as for fecal coliforms. Differences were detected just for packaged method, suggesting that good manufacturing practices take place not the plant coagulant.

Keywords: Milk Coagulation; Plant Coagulant; Silverleaf Nightshade; Trompillo; Microbiological Quality

1. Introduction

In Chihuahua Mexico, asadero cheese is produced and consumed along the state. It is a fresh cheese of stretched curd type, pleasantly tangy white cheese, and production is typically a home-based enterprise. The cheeses with these characteristics, especially stretchability, are members of the pasta filata group and include Mozzarella, Provolone, and Kashkaval [1].

The berry of the Solanum elaeagnifolium Cav (silverleaf nightshade) or so called trompillo in Mexico, is used as a rennet substitute in the production of asadero cheese. Chihuahua’s natives use the trompillo as a coagulating enzyme source [2]. S. elaeagnifolium is a deep-rooted, native perennial, which rarely reaches a height of more than 3 feet; leaves are alternate and entire. Leaf margins are normally wavy. Flowers are typical of the nightshade family, potatoes, peppers, and tomatoes [3]. The fruit, initially spherical, green and fleshy, becomes yellow to orange at maturity.

Trompillo is a wild plant that possesses proteases in its berry; those enzymes exhibit general proteolytic activities, which are useful in traditional asadero cheesemaking, reported since about one century [4]. Plant coagulants share biochemical features with chymosin, the enzyme of the rennet. Both, chymosin and plant coagulants cleave the Phe105-Met106 bond of the milk protein κ-casein, but plant coagulants are more proteolytic and have broader specificity on αs1- and β-caseins than chymosin [5-7]. However, it is possible to obtain cheeses with adequate functional properties with S. elaeagni-
folium berries as an enzyme source. Several authors have reported the use of plant’s enzymes (cardosins from Cynara) in different handmade cheeses and showed that the sensory characteristics of those cheeses are similar or better than cheeses manufactured with commercial rennet [8-10]. Similar cheeses to asadero cheese have a shelf life of 6 wk, but usually develops an off-flavor and loses textural integrity by 4 wk, potentially due to the lack of salt and high moisture that allow the outgrowth of undesirable bacteria [11].

Cheese is a complex and dynamic microbial ecosystem characterized by the presence of a large variety of bacteria, yeasts, and molds. Some microorganisms, including species of lactobacilli or lactococci, are known to contribute to the organoleptic quality of cheeses, whereas the presence of other microorganisms may lead to spoilage or constitute a health risk [12]. The microbial safety of foods continues to be a major concern to consumers, regulatory agencies and food industries throughout the world. Many food preservation strategies have been used traditionally for the control of microbial spoilage in foods but the contamination of food and spoilage by microorganisms is a problem yet to be controlled adequately. There is no information about effect of S. elaeagnifolium on the microbial safety of asadero cheese. The aim of this work was to explore microbiological susceptibility of asadero cheese made with an extract from ripe berries and compared with those elaborated with commercial rennet.

2. Materials and Methods

2.1. Sample Collection

Wild trompillo’s fruits were obtained from fields in Cd. Juárez, Delicias and Rodrigo M. Quevedo, Chihuahua State, Mexico. Ripe yellow fruits were stored at −20°C. Commercial Chymosin (Rennin Laczyme-20, Enmex, Mexico) was used. Bovine milk was supplied by the Department of veterinarian science, Autonomous University of Ciudad Juarez.

2.2. Enzyme Extract

The berries of S. elaeagnifolium were macerated and homogenized 1:3 with 0.1 M acetate buffer (pH 5.2), 1% Polyvinylpolypyrrolidone (PVPP) was added to prevent oxidation. Supernatant was recovered at 2000 × g for 5 min at 4°C. Crude extract was obtained from the supernatant by salting out with ammonium sulfate (40%), pellet was recovery at 2000 × g for 5 min at 4°C, and solubilized in acetate buffer pH 5.2, to obtain a protein concentrate extract (CE). Extract’s protein was quantified by Bradford method [13] and stored at −20°C until used.

2.3. Preparation of Cheeses

Asadero cheeses were obtained by a standardized process, just changing the enzyme source. Fresh milk pasteurized at 63°C during 30 min was acidified by citric acid (10%) addition until 34°C were reached and warmed to 32°C. Calcium chloride (0.02%), Lactic culture DMS® Delvotec MT-53X/54Y DSL (Latococcus lactis subsp lactis l, Latococcus lactis subsp cremoris and Streptococcus thermophilus) were incorporated according to the manufacturer’s instructions. Commercial rennet (CR) or CE was added at equivalent proportions. Incubation was held at 32°C for 40 min. The coagulum formed was cut with a spatula into small cubes of about 1 cm³, and maintained at 35°C until milk’s whey reached 34°C. Whey was removed and sodium chloride was added (1.4%). Curd was melted at 60°C - 63°C until it formed elastic strands; these were dried at ambient temperature, 15 min each side. The strips were rolled, weighed and divided into samples of 10 g. The samples were packed in both sealable polyethylene bag or in vacuum sealable bag and stored at 4°C - 6°C until analysis.

2.4. Evaluation of Manufacture Practices

Different levels of care in the process were considered because of the implications of the artisanal manufacture of asadero cheese. Cleanup actions were implemented, sanitation and control of pollution sources. Process A, surface cleaning was performed on tables, utensils and equipment, using commercial cleaning agents. The personnel was wearing gown and cap. Process B, the process further A, surfaces were sanitized with a quaternary ammonium solution 200 ppm. Staff was wearing gown, cap and gloves. Process C, as well as the process used in A and B, plus the utensils were immersed in chlorine solution 50 ppm and flamed; aluminum foil was used for draining cheeses. Process D, plus all the practices mentioned above, was denied entry to people to the area of the cheese making, room ventilation was closed, utensils were sterilized during the whole process and staff was wearing gown, cap, gloves and masks [14].

2.5. Microbiological Analysis

The milk was analyzed for the total coliforms content, before and after pasteurization. The asadero cheese samples made with CR or CE, packaged in resealable plastic bags and bags under vacuum were analyzed for spoilage bacteria (total aerobic count, yeasts and molds), indicators of hygiene (total coliforms) and fecal contamination (fecal coliforms), according to standardized procedures of the Bacteriological Analytical Manual [15]. Bacteriological tests were performed on days 1, 7, 14, 21 and 28 storage. Yeast and molds were analyzed until day 28 and
35. In determining fecal coliform Most Probable Number, a confirmatory test was conducted for the presence of *Escherichia coli*. The microbiological evaluation results of the two types of cheese were compared using the Student *t*-test (*p* < 0.05) using the statistical program SPSS 15.0 (Statistical Package for Social Sciences, SPSS Inc. Chicago, Ill.).

3. Results and Discussion

3.1. Analysis of Microorganisms in Milk

The total coliform limit for pasteurized milk should be 10 CFU/mL at the production plant and 20 CFU/mL at the point of sale, as provided by the Mexican legislation [16]. No pasteurized milk from this study was 35,000 CFU/mL and <3 CFU/mL for pasteurized. The counts were conducted in the 1:10 dilution and the value found for total coliforms in pasteurized milk was lower than the limit set in the Mexican standards and are consistent with the results determined by others [17,18]. This result is important because it implies that there is no contamination prior to experimental treatments.

3.2. Manufacture Practices

The effect of Good Manufacturing Practices (GMP) in total coliform counts for cheese made with CE and commercial packaging in sealable plastic bag and under vacuum packaging can be seen in Figure 1. The results reflect a decrease of coliforms based on the process used A > B > C > D. The set of operations carried out in the process D proved to be the best in terms of quality healthcare for Asadero cheese making, which is consistent with the statement by related studies [17,19].

The incubation step is the period in which opportunistic microorganisms can grow and affect the final quality of cheese whether appropriate conditions exist. It is necessary to consider the cleaning and disinfection of all equipment used in the processing of cheese or any other food as well as having a HACCP plan [17]; although this is so difficult in the traditional and small cheese plant. An advantage of asadero cheese making is that one of the last stages is carried out at melt temperatures similar to pasteurization, so that this could be a control point if properly done.

3.3. Total Coliforms Count

The Figure 2 shows the total coliform counts in cheese made with CE and CR packaged in resealable plastic bag. The number of colonies tend to decrease over time in both treatments, without representing a significant difference (*p* < 0.05) between them until day 21. At day 28, treatment with CE showed a decrease in the number of coliform colonies (*p* < 0.05), compared with CR cheese. Similar results have been reported in related cheeses during ripening [11,18,20]. This behavior is caused by the development of starter cultures, as they at pH 4 and 5 in dairy products inhibit the common and pathogenic flora, achieving lengthen the shelf life of product. Moreover, some studies indicate that the starter culture has a competition for colonization sites and substrates, fermentation products (organic acids, hydrogen peroxide, etc.) inhibitory to many pathogens [21].

Furthermore, it has been shown that lactic bacteria as *Lactococcus* sp., *Lactis*, *Streptococcus thermophilus* and *Lactobacillus*, and others, produce bacteriocins capable of
of inhibiting both gram positive and gram negative [11, 22,23]. Therefore, the decreasing growth of total coliforms along Asadero cheese storage may be due to lactic microorganisms development.

### 3.4. Effect of Packaging on Fecal Coliform

The results of cheese with CE packing under vacuum during the different storage days, can be seen in Figure 3. According to the Standard Mexican legislation the maximum limit of fecal coliforms in cheese is 100 CFU/g [24]. This limit was exceeded on day one of storage, as shown in the graph, but tends to decrease with advancing storage time. The result shows a behavior similar to that developed for total coliforms. From the seventh day of storage, the cheese in both types of packaging decreased fecal coliform count over that obtained on day one, representing count below the limits set by the legislation. This behavior has been reported in similar studies in different cheeses, in addition to the discussed in the previous section, this is attributing to inhibition by the starter cultures due to the substrate competition [11,21,25]. This trend is repeated in both packages, presenting a positive effect on vacuum packaging with lower microbial load present. This is because the vacuum packing limits the availability of oxygen. The cheese packed under vacuum was favorable for controlling coliform microorganisms, since lower counts were significantly different (p < 0.05) compared with cheese packed in sealable plastic bag, these results are consistent with microbiological studies conducted in different packaged foods [26].

Both cheeses made with CE and CR showed no growth of *E. coli* during the storage period. The presence of *E. coli* in foods such as cheese, involves a direct or indirect contamination, reflecting poor hygiene and inadequate process, so that the product can be considered safe.

### 3.5. Total Aerobic Count

The aerobic microorganisms count for cheeses made with the two types of coagulant and packaged in resealable plastic bags are shown on Figure 4. The trend for both treatments was increasing the UFC during storage. Both CE and CR cheeses had a higher count, especially at 21 and 28 days. These results are considered as a normal process, some authors have found that most of the lactic acid bacteria in cheese, after 15 days of ripening, occupied the most aerobic count and that after a month have dominated the bacterial community [27,28]. This correlate with the lower coliforms count, since has been discussed that lactic acid bacteria growing in raw milk or cheese and tend to give a protective action against a large number of undesirable microorganisms. It should be noted that the aerobic mesophilic count is not considered at this point as an indicator of spoilage, this microbial flora is part of the starter culture added (*Lactococcus lactis subsp lactis*, *Lactococcus lactis subsp cremoris* and *Streptococcus thermophilus*). This bacteria show an increasing growth throughout the asadero cheese shelf life and this is desirable for the cheese formation and to add acid flavor and sensory characteristics of the final product [29,30].

### 3.6. Yeast and Molds

The trend of growth of yeast and molds in cheeses made with CE or CR is shown in Figure 5. Although the greatest growth observed was on CR cheese, it is within the safe limits allowed by Mexican law, which is 500 CFU/g [24]. Cheeses of the two treatments were within
the limits until 21 and 28 days of storage. CE Cheese still remained within the safe limit established by Mexican law at the day 35, different to the CR treatment that have increased molds, resulting different at 35 days ($p < 0.05$), which probably means that \textit{S. elaeagnifolium} extract may have an anti-mold ability or produce cheese characteristics inappropriate for mold growth at the long term. In some reports with cheese similar to this study it was found that the cheese made from goat milk stored at refrigeration ($4 ^\circ C$) remains unchanged from fresh and microorganism deterioration after 35 days of storage [31, 32]. Yeast and molds in foods often cause impairment, function as health indicators and behave as pathogens through the production of toxins, however in the case of ripened products made with molds, their presence has no a meaning health [12]. The mold on the surface of fresh cheeses suggests early deterioration and lack of freshness. Its microscopic development in food poses a risk in the production of toxic substances to the health of consumers [33]. This kind of microorganisms are most often found in soft cheeses, usually comprising the genera \textit{Alternaria}, \textit{Aspergillus}, \textit{Cladosporidium}, \textit{Monilia}, \textit{Mucor} and \textit{Penicillium} and yeasts \textit{Geotrichum candidum} and \textit{Candida} spp. [34,35].

4. Conclusions

There were no differences between the two types of coagulants in most microbiological parameters; there are no effects of the plant coagulant \textit{S. elaeagnifolium} on the microbiological quality of asadero cheese during 28 days of shelf life. Coagulant plant tends to inhibit the mold growth until 35 days, however further investigation is recommended. Manipulation of pasteurized milk at room temperature is an important point of post-pasteurization contamination. The contamination with undesirable microorganisms in the incubation step can result in a safety hazard for asadero cheese. The terms of packaging and storage temperature of finished product are also important. Good manufacturing practices remain crucial for obtaining an optimal hygienic quality product. Considering that \textit{S. elaeagnifolium} (Trompillo) is a weed with substantial distribution in northern Mexico, could be considered as an industrial source of milk coagulant in the manufacture of different cheeses. Before a further application of \textit{S. elaeagnifolium} extract in cheese, the sensory evaluation is recommended. Development of cost effective isolation procedures that yield standardized extracts as well as safety and toxicology evaluation of this coagulant requires a deeper investigation.

REFERENCES


