

**QUALITY PARAMETERS, PROBIOTIC VIABILITY AND SENSORY
PROPERTIES OF PROBIOTIC STIRRED SESAME YOGURT**

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Abstract

The beneficial effects of sesame seeds are of great interest for the conception of healthy dairy products such as probiotic stirred yogurt. We investigated the effects of adding raw or roasted sesame seeds on the probiotic viability, quality parameters and consumers acceptability of stirred yogurt during 28 days cold storage (4 °C). All yogurts were analyzed for microbial counts (starter culture and probiotic), pH, titratable acidity, proteolytic activity, syneresis and antioxidant activity. Yogurts containing sesame seeds showed the highest probiotic counts, proteolytic activity, radical scavenging activity high titratable acidity, and low pH. Raw and roasted sesame can selectively impact probiotic growth with limited effect on yogurt starter culture especially at long cold storage (14-28 days). Yogurts enriched with roasted sesame had higher sensory acceptability compared to control and probiotic yogurts. Roasted sesame can be successfully incorporated to improve probiotic viability and sensory properties of stirred yoghurt, as well as to improve the antioxidant properties.

Keywords: probiotic, sesame yogurt, sensory evaluation, fermentation, proteolytic activity, antioxidant capacity

Introduction

In recent years, consumers around the world are very interested in foods that provide positive health benefits. This social phenomenon is certainly linked to increased awareness of the link between food quality and health. Yogurt is the most known and consumed fermented milk with high nutritional value and health

promoting ingredients (Allgeyer *et al.*, 2010). It is coagulated milk made by lactic fermentation without drainage with two symbiotic and homofermentative cultures: *Streptococcus thermophilus* and *Lactobacillus delbrueckii* subsp. *Bulgaricus* (Shah, 2007; Sodini *et al.*, 2004). Consumption of yogurt helps improve diet quality because it is an excellent source of protein, vitamin B2 and B12, folate, niacin, calcium, magnesium, phosphorus and zinc (Mckinley, 2005). Yogurt is also a rich source of bioactive peptides released during lactic fermentation by lactic acid bacteria (Akalm *et al.*, 2012). In addition, it can be consumed by people with lactose intolerance, due to the modification of allergenic properties of milk (Bernat *et al.*, 2015; Viljoen, 2001). The increase in per capita annual consumption of yogurt can be attributed to the diversification (probiotic or symbiotic yogurt, light or defatted yogurt, fruit yogurt and liquid yogurt drinks) and availability of the low priced product in the market. Probiotics have experienced a renewed interest actually, marked by an explosion of scientific publications/papers underpinned by the partnership with the agri-food industry. The publications are based on Élie Metchnikoff theory, who proposed that the acid-producing organisms in fermented dairy products can prevent “fouling” in the large intestine and thus prolong the consumer’s life span (Anukam and Reid, 2007; Heller, 2001). Probiotics exert several health benefits, including improving lactose digestion, immune and gastrointestinal system (Hatcher and Lambrecht, 1993; Plessas *et al.*, 2012), restoring microbial balance in the host gut flora (Gibson and Roberfroid, 1995), preventing cancers (Reddy *et al.*, 1983), lowering cholesterol (Gilliland and Walker, 1990) and enhancing protection against pathogenic bacteria using three types of biological processes: immunomodulation; direct antimicrobial activity and reinforcement of the epithelium barrier function (Alexandre *et al.*, 2014). Among the probiotic strains, *B. animalis* subsp. *lactis* (Bb12) is the most studied and documented in the literature with over 200 scientific publications (Garrigues *et al.*, 2010). It can withstand harsh conditions compared to other strains and hence its common use in probiotic foods. It can survive passage through the gastrointestinal tract with high capacity to adhere to enterocytes (Haschke *et al.*, 1998), it is also a technologically suitable strain because without adversely affecting taste and texture of food products (Moller and De Vrese, 2004). The viable count of each probiotic bacteria is very important to provide health benefits to the host. Consumption of probiotic yogurt over 100 g per day is generally recommended (Rybka and Kailasapathy, 1995) and the minimum viable counts of each probiotic strain must be 10⁶cfu/mL (Tamime *et al.*, 2005).

Sesame (*Sesamum indicum* L.) is the first cultivated oilseeds by humans and one of the world’s most important crops (Uzun *et al.*, 2007). It has long been considered a healthy food providing energy and preventing aging in the orient (Halvorsen *et al.*, 2002). Sesame seeds are a potential source of nutrients, containing 35-57% oil, 20-25% protein, 20-25% carbohydrate and 5-6% ash (Salunke *et al.*, 1992). Seeds are a rich source of beneficial bioactive compounds and endogenous antioxidants mainly phenolic lignans (sesamol, sesamolol, sesamin and sesamino-lglucosides) (Bedigian, 2004; Uzun *et al.*, 2007), tocopherols and phytosterols (Bae *et al.*, 2016). Regular consumption of sesame seeds helps to low cholesterol (maintain

HDL and lower LDL) and blood pressure and protects the liver from oxidative damage due to the presence of sesamin and sesamol (Anilakumar *et al.*, 2010). Actually, much attention has been directed toward the utilization of oilseed for dairy manufacture including dairy drinks and vegetable milks (soy, almond and sesame milk). The major uses of sesame in human food are: oil production, Tahina and Halawa manufacture and garnish of bakery products (Kinsella *et al.*, 1985).

Sesame has previously been incorporated in fermented dairy products, but the seed was unroasted. Raw sesame seeds increased titratable acidity, protein and fat content, amino acids especially some essential amino acids and omega fatty acids. Bacterial counts of sesame enriched yogurt increased and the best sensory scores was obtained with 2% sesame supplementation (Aziz and Aboeleinen, 2010).

Due to the huge popularity of yogurt among consumers, producers and manufacturers continue to develop new yogurt assortments with value-added ingredients. Nutraceutical functions and positive effects of sesame seeds were previously reported (Namiki, 2007). It is important to combine the nutritional value of sesame and the health benefits of probiotic culture in yogurt. Our study (i) evaluated the effects of ground raw and roasted sesame on stirred yogurt quality parameters (pH, titratable acidity and syneresis) and probiotic growth, (ii) determined the acceptability of sesame yogurt by consumers comparing between roasted and unroasted sesame yogurts.

Materials and methods

Material and microbial cultures

White sesame seeds (*Sesamum indicum*) imported from India were purchased from a local market in Bejaia (Algeria). Impurities such as dust, sand, stones, spoiled seeds and other extra materials were sieved through a 5mm screen. One part of the seeds was roasted at 180°C for 20 minutes using an electric oven. Roasted and unroasted seeds were ground (GM 200; Retsch GmbH, Germany) to achieve particle size of 2-3 mm then stored in sealed plastic bags (4 °C) until use to avoid oxidation.

The freeze-dried mixture of starter cultures (*Lactobacillus delbrueckii* ssp. *bulgaricus* and *Streptococcus salivarius* ssp. *thermophilus*) and the probiotic strain (*Bifidobacterium animalis* ssp. *lactis*, BB-12) were purchased from CHR Hansen (France).

Determination of optimum sesame incorporation

Pasteurized milk was mixed with skim milk powder to have milk with 18 % total solids and stirred at 85 °C for 20 min. After cooling to 40 °C, the starter culture was incorporated and the milk was divided into conical tubes (50 mL). Variable concentrations of ground sesame seeds (0-10%, w/v) were added in order to determine the best incorporation rate that can be added to milk before fermentation without producing syneresis and disrupting bacterial growth (Agil et al., 2013). Incubation was carried out at 40–42° C until pH 4.5.

Yogurt manufacture

Yogurt was prepared in a laboratory scale using the method described previously (Kailasapathy *et al.*, 2008). Pasteurized milk (3.5% fat) was purchased locally from a commercial source (Candia, Bejaia). Skim milk powder (SMP) was added with speed stirring, to have milk with 18 g/100 g total solids. Then it was heated to 85 °C for 20 min and cooled quickly to 40 °C. Commercial sugar (8%) and 2 g/100 g (w/w) of frozen yogurt starter culture was incorporated and the mixture stirred continuously then divided in 150mL plastic yogurt cups. Freeze-dried culture of *B. animalis ssp. Lactis* (10^8 cfu/g) 2 g/100 g (w/w) was incorporated and sesame seeds (6 %) added to make six yogurts: Y, YP, YUS, YRS, YPUS, and YPRS (Table 1).

Table 1. Formulation of standard or control yogurt (Y), probiotic yogurt (YP), unroasted sesame yogurt (YUS), roasted sesame yogurt (YRS), probiotic unroasted sesame yogurt (YPUS) and probiotic roasted sesame yogurt (YPRS).

Samples	Total solid%	Lactic ferment %	Sugar %	Probiotic %	Sesame seed %
Y	18	2	8	0	0
YP	18	2	8	2	0
YUS	18	2	8	0	6
YRS	18	2	8	0	6
YPUS	18	2	8	2	6
YPRS	18	2	8	2	6

Incubation was carried out at 40–42 °C until a pH of 4.5. After fermentation yogurts were quickly cooled in ice water bath then stirred. Samples were prepared in triplicate and the yogurt cups were sealed then stored in the refrigerator at 4 °C for 28 days for analysis.

Enumeration of probiotic and starter cultures

Bacterial count was carried out weekly for a total of four weeks (days 1, 7, 14, 21, and 28). For each lot at different dilutions (four to five serial dilution of 1/10), 100µL of each of the last three dilutions were spread by the streaks method on Petri dishes. For enumeration of starter cultures, the inoculated media were incubated at 40 °C for 24 h (Espírito Santo *et al.*, 2010). The probiotic was enumerated on MRS agar containing L-cysteine hydrochloride (0.5 g/L) and incubated (40°C, 72h) under anaerobic conditions (Rodrigues *et al.*, 2012). Plates containing 30-300 colonies were enumerated and the results were expressed as log colony forming units per milliliter (log cfu/mL) using the equation 1 (Sun-Waterhouse *et al.*, 2013):

$$\text{cfu} = \text{number of colonies/plate factor} \times \text{dilution factor} \quad (1)$$

where the plate and dilution factors refer to the amount of sample pipetted and the dilution series of the yogurt sample, respectively.

Determination of pH and total titratable acidity (TTA)

pH was measured at 25°C by electrode immersion with a pH meter (211 HANNA) each week until the 28th day. Yogurt (1 mL) was mixed with distilled water (9 mL)

with a few drops (3 to 5) of phenolphthalein 0.1% (w/v). The titration was made with a solution of NaOH (0.1N) until the persistence of a pink color. The volume of NaOH used for titration was noted and the titratable acidity (TTA %) was expressed as a percentage lactic acid equivalent and calculated according to equation 2:

$$\text{TTA (\%)} = V_{\text{NaOH}} \times 0.1\text{N} \times 100\% \times 0.009 \times 10 \quad (2)$$

where: V_{NaOH} - Volume of NaOH in mL used for titration; 0.0090 - coefficient corresponding to lactic acid; 10 - dilution factor.

Proteolytic activity

To measure proteolysis of yogurt bacteria during cold storage, the o-phthalaldehyde (OPA) method was used (Donkor *et al.*, 2007). The concentration of free amino groups and peptides is proportional to the absorbance at 340 nm.

Syneresis

Syneresis was determined using the centrifugation method (Aprianita *et al.*, 2009). Yogurt (10g) from different days storage was centrifuged (Bench-top centrifuge NF 200, Belgium) 700×g at 8 °C for 10 min. The clear supernatant was weighed, and syneresis was expressed as percent weight of supernatant relative to the original yogurt weight using equation 3:

$$\text{Syneresis (\%)} = (\text{weight of collected whey}/\text{weight of yogurt}) \times 100 \quad (3)$$

Antioxidant activity by DPPH inhibition assay

First, water yogurt extracts were prepared using the method of (Amirdivani and Baba, 2011). Yogurt (10 g) was mixed with 2.5 mL of distilled water; the mixture was stirred, and adjusted to pH4 with HCl solution (0.1 M). The mixture was heated in a water bath (45°C, 10 min) and centrifuged (5000g, 10 min, 4°C) to remove precipitated proteins. The supernatant was adjusted to pH 7 with 0.1M NaOH and centrifuged (5000g, 10 min) to remove residual proteins and salts then stored at -20°C for later use. Extractions were carried out in triplicates.

The antioxidant capacity of yogurt extracts was determined as described previously (Behrad *et al.*, 2009). Yogurt water extract (250 µL) was mixed with ethanol solution of 60 µM DPPH (3 mL). The mixture was shaken vigorously and then incubated for 20 min at room temperature in the dark. The absorbance was measured at 517 nm. The readings were compared with the control which contained distilled water instead of yogurt extract. The % inhibition of DPPH was determined using equation 4:

$$\% \text{ inhibition} = (A_{\text{control}} - A_{\text{sample}}) / A_{\text{control}} \times 100 \quad (4).$$

Yogurts sensory evaluation

Ten untrained panelists familiar with yogurt (students in food sciences and technology from the University of Bejaia, Algeria) were selected for sensory evaluation. The sensory analysis was based on a nine-point 1 (poor) to 9 (excellent) hedonic scales for some sensory parameters including color, odor

(flavor), taste, texture, acidity, syneresis and overall acceptability (Haque and Ji, 2003). The stored yogurt samples were evaluated by nonsmoker's panel members on the first day of storage. Water and bread were provided between samples to cleanse the palates.

Data analysis

Three determinations were made for all the assays. Analysis of variance was performed by the general linear models (GLM) procedure, means comparison by Duncan's test, and Pearson correlation according to Statistical Analysis System, SAS 9.1 for Windows.

Results and discussion

Optimum sesame incorporation

Yogurts prepared with an incorporation rate of 8% and 10% have a liquid consistency and texture like a yogurt drink which is due to the phenomenon of syneresis, this will not be accepted by consumers, but yogurts containing 0 to 6% of ground sesame have a consistent texture. Consequently, the best concentration was 6% of ground sesame which can be incorporated into yogurt before fermentation under our experimental conditions without whey separation from gel.

Bacterial counts

Figure 1a shows the variations in bacterial counts of starter culture in the six yogurts. Bacterial count was affected by sesame supplementation ($P<0.05$), probiotic ($P<0.0001$), storage time ($P<0.01$) and their interaction ($P<0.001$). Sesame and probiotic exerted beneficial effects on the control yogurt (Y) culture ($P<0.05$). On the first day of cold storage, *S. thermophilus* and *L. bulgaricus* counts did not differ and varied from 8.12 to 8.39 log cfu/mL; then it declined to 7.45-8.35 log cfu/mL. Reduction rates varied from 0.47 to 8.28% in the following order: Y>YUS>YRS>YPRS>YPUS>YP. In fact, bacterial counts decreased linearly in Y ($Y = -0.18x + 8.27$; $r^2 = 0.93$) and YUS ($YUS = -0.14x + 0.37$; $r^2 = 0.98$).

From 14 to 28 days, the highest count of starter culture was recorded in probiotic yogurt (YP) and the viability was considerably stable and maintained during storage. At day 7, the number of counts increased by 2.8 % in YPUS followed by a sharp decline at the 14th day. The presence of probiotic bacteria supported and helped growth of yogurt starter culture, even in the absence of sesame; this can be explained by a competition of starter culture with the probiotic for the nutritive elements by increase in the number of living cells. Significant enhancement of *L. delbrueckii* ssp. *bulgaricus* counts was reported in yogurt made with a mixture of probiotic strains (*Lactobacillus acidophilus*, *Bifidobacterium lactis* and *L. paracasei*) (Donkor et al., 2006). An important increase (15-20%) was observed in *S. thermophilus* counts during probiotic yogurts storage, but decreased after 21 day from 7 to 33% (Dave and Shah, 1997). *L. bulgaricus* counts also increased initially then declined during the later stage. The population counts were similar for YPUS-YPRS and YUS-YRS, without significant differences particularly at the beginning

and end of storage indicating similar behavior of yogurt bacteria in the presence of raw or roasted ground sesame.

The viable counts of *B. lactis* (log cfu/mL) during the four weeks cold storage (Figure 1b) was significantly ($P<0.001$) affected by sesame supplementation and storage time. Except on day one, where the difference was insignificant. Probiotic supplemented yogurts with roasted or unroasted sesame exhibited higher *B. lactis* ($P<0.05$) counts compared to the control (YP) demonstrating its prebiotic potential. Probiotic count was within the 7.83 to 8.36 log cfu/mL range in unsupplemented yogurt (YP) with optimum at day 1 and 7.73 to 8.44 log cfu/mL in supplemented yogurt with an optimum at day 14 and 21 for YPUS and YPRS, respectively. The number of probiotic colonies varied between the supplemented yogurts (YPUS and YPRS); but difference on day 1 and 28 was not significant between them. Probiotic growth decreased initially for the first week for both supplemented yogurts, then increased in YPUS and remained stable until 21 day in YPRS. From day 1 to 28, in un-supplemented probiotic yogurt (YP) the number of viable count decreased linearly by 1 log cfu/mL ($YP=-0.22x+8.56$; $r^2=0.95$). At 28 days, all sesame yogurts had significantly higher probiotic ($P<0.05$) counts compared to the control by over 0.34 log cfu/mL.

Ground sesame (*Sesamum indicum*) addition promoted total bacterial organisms of *B. animalis ssp lactis* probably due to their improved survival by providing essential nutrients and oligosaccharides. Probiotic strains have nutritional requirement and important proteolytic activity to support their growth. For this, they need free amino acids to increase their viable cells (Vasiljevic and Shah, 2008). Indeed, proteins are the second predominant compounds in sesame seeds (24 g/100 g) that is rich in methionine, cystine, arginine, and leucine (36,25,140, and 75 mg/g protein, respectively) (Namiki, 2007).

Sesame is an excellent source of essential amino acid significantly higher than that of FAO/WHO requirement for human except lysine (Johnson *et al.*, 1979). The high (18-20%) carbohydrate content of *Sesamum indicum* (11% dietary fiber) with low glucose, fructose and starch and the presence of an oligo sugar planteose *O*- α -D-galactopyranosyl-(1,6)- β -D-fructofuranosyl- α -D-glucopyranoside unaffected by the human digestive enzymes maybe responsible for the prebiotic potential of sesame (Namiki, 1995).

During storage, the viability of *B. animalis* in probiotic yogurts was superior to the minimum recommended level (>6 log cfu/g) to provide the beneficial health effects to the gut (Kongo *et al.*, 2006). The viability loss/reduction in probiotic bacteria may be due to post-acidification during refrigerated storage (Sun and Griffiths, 2000). The co-culture of starter culture with bifidobacteria, which are a proteolytic species that is not recommended because they create acidic conditions (Abu-Taraboush *et al.*, 1998). Some probiotic strains are sensitive to antimicrobial substances produced by yogurt bacteria during refrigerated storage (Shah, 2001).

Our results showed that raw and roasted sesame may selectively impact probiotic growth with minimal effect on yogurt starter culture especially at the long cold storage (14-28 days).

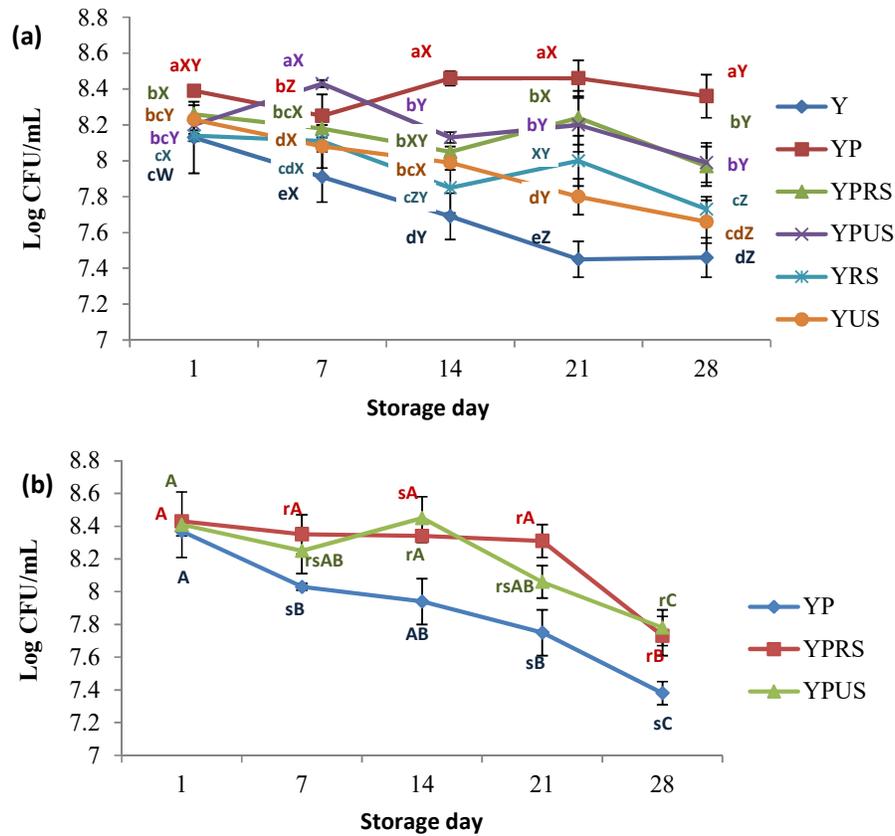


Figure 1. Bacterial viability of starter culture (a) and probiotic (b) in yogurts during 28 days storage at 4°C. Means with different lower and upper case letters are significantly different ($P < 0.05$).

Post-acidification (pH) and titratable acidity

Yogurt pH were significantly affected by sesame supplementation ($P < 0.0001$) and storage time ($P < 0.01$). Yogurt bacteria metabolize carbohydrates for growth and energy producing various organic acids such as lactic, butyric, propionic, acetic and citric acids (Fernandez-Garcia and McGregor, 1994). pH is a measurement of hydrogen (H^+) concentration contributed by the released organic acids during fermentation and storage. At day 1, the pH among all yogurt treatments ranged from 4.41 to 4.52 (Figure 2a), then it decreased linearly until 3.68 to 3.91. There was no significant difference ($P > 0.05$) between unsupplemented yogurts (Y and YP) during storage except on day 14. From day 1 to 14 there were no differences

between all supplemented yogurts with or without probiotic, indicating that probiotic had no effect on pH ($P > 0.05$).

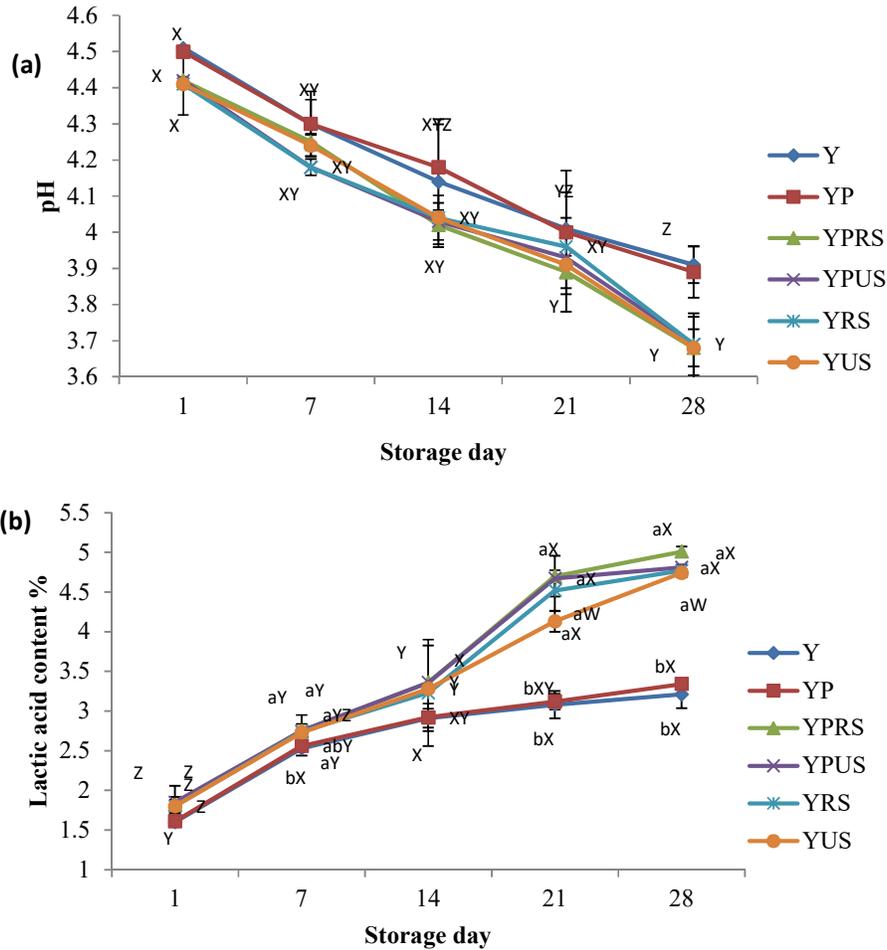


Figure 2. Post-acidification (a) and titratable acidity (b) in yogurts during 28 days of storage at 4°C. Means with different lower and upper case letter are significantly different ($P < 0.05$).

TTA measurement is an indicator of bacterial metabolic activity in fermented dairy product. At day 1 the TTA values ranged between 1.6 and 1.85 then increased until 3.21 and 5 (Figure 2b). TTA for the four sesame yogurts was higher than that of control and probiotic yogurts (Y and YP) during storage; the evolution of titratable acidity in those yogurts (YUS, YRS, YPUS, YPRS) may be described in two distinct phases. Between day 1 and 14, there was a gradual significant increase and between day 14 and 28 the rate of increase was greater and rapid

(YUS=0.27x+1.15; $r^2=0.99$; YPRS=0.83x+1.02; $r^2=0.97$). The addition of both raw and roasted ground sesame was helpful in increasing total titratable acidity ($P < 0.01$) during storing indicating that microorganisms were more active in the presence of sesame seeds. Sesame contains various carbohydrates including D-glucose (3.63%), D-fructose (3.43%), D-galactose (0.40%), sucrose (0.17%), raffinose (0.59%), stachyose (0.38%), planteose (0.23%), and sesamose (0.14%) which may easily be available and metabolized by yogurt bacteria (Wankhede and Tharanathan, 1976). The reduction of pH and liberation of lactic acids reflects the high metabolic activity of yogurt bacteria. Lactic acid bacteria (LAB) used milk proteins (nitrogen source) and carbohydrate from sesame seeds (carbon source) to produce organic acids and other volatile compounds including acetaldehyde and diacetyl. Organic acids play an important role as natural preservatives and the volatile compounds contribute to the organoleptic properties and yogurt flavor (Fernandez-Garcia and McGregor, 1994).

Proteolysis by o-phthalaldehyde (OPA) assay

Proteolytic activity improved significantly ($P < 0.05$) in the presence of probiotic and sesame seeds (Table 2). All yogurts had higher OPA values (0.9-1.04) than that of the control yogurt (0.62) at the beginning and during refrigerated storage. The highest proteolytic activity (1.26) was recorded in both supplemented yogurts (YRS and YUS) at days 14 and 21, respectively. Proteolytic activity of bacteria was similar in probiotic sesame containing yogurt (YPUS and YPRS) indicating that seed roasting had no effect on yogurt proteolysis. Proteolysis increased in all yogurts until the optimum absorbance recorded on day 14 or 21 depending on yogurt sample then decreased at the end of storage.

Table 2. Proteolysis (OPA) of yogurts during 28 days storage at 4°C

Sample	Storage time (Days)				
	1	7	14	21	28
Y	0.62±0.01 ^{eZ}	0.78±0.02 ^{eXY}	0.85±0.02 ^{dXY}	0.87±0.01 ^{dX}	0.74±0.03 ^{cY}
YP	0.90±0.01 ^{bY}	0.99±0.06 ^{bXY}	1.12±0.02 ^{cX}	1.04±0.03 ^{cXY}	0.99±0.04 ^{bXY}
YPRS	1.01±0.02 ^{abZ}	1.13ab±0.04 ^{XYZ}	1.25±0.06 ^{abX}	1.07±0.08 ^{bcYZ}	1.14±0.02 ^{aXY}
YPUS	1.04±0.04 ^{aZ}	1.17±0.06 ^{aXY}	1.22±0.01 ^{abX}	1.08b±0.03 ^{cYZ}	1.08±0.08 ^{abYZ}
YRS	0.93±0.01 ^{abY}	1.19±0.02 ^{aX}	1.27±0.05 ^{aX}	1.17±0.03 ^{abX}	1.16±0.07 ^{aX}
YUS	0.98±0.03 ^{abZ}	1.06±0.02 ^{abYZ}	1.18±0.06 ^{bcXY}	1.26±0.02 ^{aX}	1.14±0.01 ^{aXY}

Results are expressed as absorbance at 340 nm; means with different lower and upper case letters in the same column or row are significantly different ($P < 0.05$).

Enhanced proteolytic activity suggests that enzymatic activity of protease from starter culture and probiotic was markedly improved in the presence of ground sesame seeds that provided divalent ions including Ca^{2+} , Fe^{2+} and Mg^{2+} . Those minerals were reported to be necessary for protease activity (Llorente-Bousquets *et al.*, 2008). Indeed, sesame seed was found to be rich in various minerals. Among them, calcium and iron, which are often deficient in modern diets, and present in high concentrations (1200 and 9.6 mg/100 g, respectively) (Namiki, 2007).

Syneresis

Syneresis is an undesirable feature in yogurt resulting from the separation of liquid phase (serum) from the protein gel (Shah, 2003). Syneresis of investigated yogurts was significantly affected by sesame supplementation ($P<0.001$) and storage time ($P<0.001$) but there was no difference between roasted and unroasted sesame supplementation ($P>0.05$). At day 1, syneresis varied between 41.63 and 42.26 % (Table 3) in all yogurts without significant difference ($P>0.05$). Whey separation of sesame containing yogurt except YPUS followed parallel biphasic trends during storage with inflection at 14 days. From day 1 to 14, differences in level of syneresis were insignificant among the three supplemented yogurts (YUS, YRS, YPRS) and between control (Y) and probiotic yogurt (YP). Storage linearly increased whey separation (14-28 days) especially in supplemented yogurts (YUS= $2.63x+41.8$; $r^2=0.96$; YRS= $3.33x+40.6$; $r^2=0.98$, YPRS = $2.5x+41.33$; $r^2=0.99$).

Syneresis increased with sesame addition compared to control and probiotic yogurts (Table 3). Many studies have reported higher syneresis accompanied with low viscosity for enriched yogurts with various plant ingredients. Therefore, addition of fruits decreased water-holding capacity of protein and viscosity resulting in increased syneresis (Akyüz and Coflkun, 1995; Mohamed *et al.*, 2014; Zainoldin and Baba, 2009). At the end of cold storage, control and probiotic yogurts (Y and YP) displayed the lowest syneresis (45.7 and 44.03 %, respectively) whereas the highest mean value (50.36 %) was recorded in roasted sesame containing yogurt (YRS). The syneresis trend of these yogurts paralleled changes in their proteolysis (OPA) and TTA, thereby indicating their strong association. Therefore, increased syneresis in sesame containing yogurts can be attributed to lactic acid bacteria degradation of milk proteins that are responsible for water retention.

Table 3. Syneresis of yogurts during 28 days storage at 4°C

Sample	Storage time (Days)				
	1	7	14	21	28
Y	41.63±3.12 ^Z	40.93±1.16 ^{cZ}	42±2.39 ^{bZ}	43.30± 1.18 ^{cY}	45.70± 3.31 ^{bcX}
YP	42.03± 2.31 ^Y	41.60± 0.4 ^{bcY}	42.63±4.41 ^{abY}	44.00± 1.31 ^{bcX}	44.03±3.34 ^{cX}
YPRS	41.90±1.86 ^Z	43.33± 1.13 ^{aYZ}	43.87±0.64 ^{abWZ}	46.27±2.91 ^{abXY}	48.87±0.56 ^{abX}
YPUS	41.7±2.45 ^Y	41.53±3.81 ^{bcY}	42.60±0.88 ^{abY}	43.20±2.81 ^{cY}	48.17±2.76 ^{abX}
YRS	41.83±3.78 ^Z	42.90±2.56 ^{abZ}	43.70±2.98 ^{abZ}	47.77±3.14 ^{aY}	50.37±0.97 ^{aX}
YUS	42.27± 1.75 ^Z	42.97±2.84 ^{abYZ}	44.30± 1.16 ^{aY}	46.10a± 1.32 ^{bX}	49.57±4.54 ^{aW}

Results are expressed in % of collected whey; means with different lower and upper case letters in the same column or row are significantly different ($P< 0.05$).

Antioxidant activity

Yogurts containing raw or roasted sesame displayed strong scavenging capacity, whereas control and probiotic yogurts (Y and YP) manifested lower scavenging

capacity (Figure3). Our results agreed with those of Hashish *et al.* (2014) where the antioxidant activity of yogurt increased with white and red sesame tahina (1 to 6%) supplementation. Many studies reported higher antioxidant capacity of yogurt supplemented with various ingredients from plant material (fruits, leaves and seeds) compared to their control (Muniandy *et al.*, 2016; Sah *et al.*, 2016; Van Nieuwenhove *et al.*, 2019). This potential was attributed to the content of phenolic compounds (Baba *et al.*, 2014). In our study, proteolysis was more marked in yogurts enriched with sesame seeds. In addition, bioactive peptides released by lactic acid bacteria during proteolysis may play an important role as antioxidant compound.

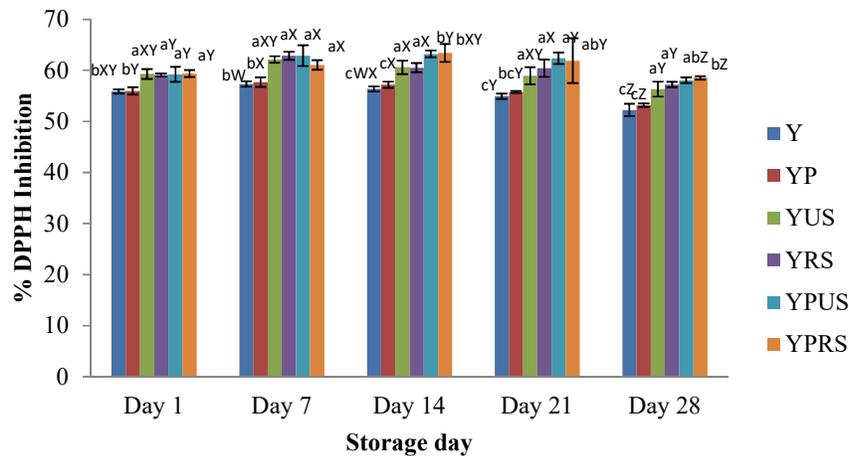


Figure 3. DPPH radical scavenging activity of yogurts during 28 days storage at 4°C. Means with different lower and upper case letters are significantly different ($P < 0.05$).

Sensory analysis

The results of the sensory evaluation after 1 day of refrigerated storage performed in all yogurt samples are reported in Figure 4. Roasted sesame containing yogurts (YRS and YPRS) were preferred by the panelists in color, flavor, taste and sweetness. Sweetness was enhanced with roasted seeds that contained more sugar; whereas taste and flavor was improved with roasting that enhances the pleasant characteristic flavor of hazelnut (Namiki, 2007).

Color is the first characteristic perceived by consumer and the major attribute in food industry. Color of roasted sesame containing yogurts (YRS and YPRS) was preferred due to the pigmentation caused by browning substances released from the Maillard reaction that gave better visual appearance to the product.

Control yogurt (Y) showed the better score for both texture and acidity. This implies that the high acidity of the supplemented yogurt was perceived by the panelists. Because no differences were observed, probiotic roasted sesame yogurt

(YPRS) and roasted sesame yogurt (YRS) were equally accepted ($P>0.05$) at first view indicating that probiotic had no effect on sensory attributes of yogurt. The same results were observed when comparing some sensory characteristics (appearance, flavor, texture) between standard and probiotic yogurt made with *L. rhamnosus* GR-1 and *L. reuteri* RC-14 (Hekmat and Reid, 2006).

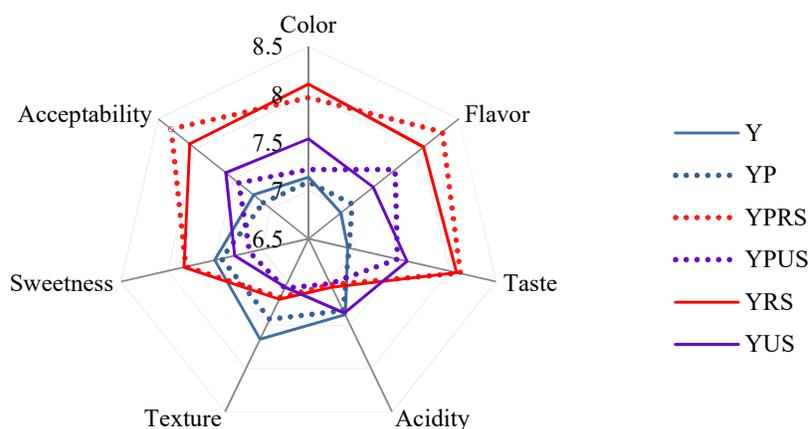


Figure 4. Diagram of sample scores of the sensory evaluation of probiotic stirred sesame yogurt.

Conclusion

In this study, ground (raw or roasted) sesame yogurt supplementation (6%) selectively stimulated bifidogenic microbial growth, increased syneresis, acid production and proteolysis and reduced pH. Antioxidant activity of yogurts also increased in the presence of sesame during storage. The enhanced antioxidant potential and sensory properties of roasted sesame supplemented yogurts can be beneficial to improve human health and consumer's acceptability; this implies that addition of roasted sesame in yogurt manufacture can be recommended without adverse effect on the properties of this product knowing that it provides selective conditions for probiotic growth.

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