

## DEVELOPMENT OF INFANT FORMULAS BASED ON *Lactobacillus helveticus* OR *Bifidobacterium lactis Bb12* BACTERIA AND BARLEY

Yonis, A.A.M. and Hanaa F. El-Meheiry.

Dept. of Home Economics, Fac of Specific Education, Mansoura Univ.

### ABSTRACT

The present work was designed to prepare fermented porridge by (*Lactobacillus helveticus* or *Bifidobacterium lactis Bb12*) as baby food with high nutritional value from malt flour, rice flour, skim milk powder, honey and carrot with different percents. Prepared diet were chemically, microbiologically biologically and sensory evaluated directly after fermentation and after storage period (21 days) at (4-5 °C). The results indicated that the protein, ash, lactic acid, and acetic acid in all treatment increased during storage period, while fiber, carbohydrate, glucose and fructose values were decreased. The pH values varied according to the growth of both strain, *L. helveticus* recorded a higher decreasing effect on pH values (pH 4.12). The viable count of *L. helveticus* reached maximum growth after storage period 7 days in all treatments, then slightly decreased till the end of the storage period, while *Bifidobacterium lactis Bb12* increased during the storage up to 15 days, and then gradually decreased. All samples had highest values of calcium, phosphorus, magnesium, thiamine and riboflavin to meet Recommended Dietary Allowances (R.D.A) requirements. Sensory evaluation showed that the best formula for infants were samples treated with carrot and both strain. The biological study of fermented porridge by rats indicated that rats`body weight gain significantly increased compared to the control with no effect on relative liver and spleen weights. Also, in comparison with the control group, rats fed the fermented porridge resulted in a considerable increase in their intestinal fecal content of *L. helveticus* and *Bifidobacteria*, however the count of staphylococci and coliforms significantly reduced.

### INTRODUCTION

Human milk is the ideal food for infants during the first 4 – 6 months of age, since it contains all breast – fed infant nutritional requirements (Tojo *et al.*, 1995). In trying to simulate the breast – fed infant`s pattern of gut colonization, the addition of lactobacilli and / or bifidobacteria (also known as probiotics) to infant formulas has also been described (Vandenplas, 2002). Probiotics and prebiotics are also incorporated together (known as synbiotics) in foods in order to improve the survival and establishment of beneficial bacteria in the host large intestine, such as addition of probiotics (bacteria) with carbohydrates that promote their growth (prebiotics) to normal infant formula or weaning foods (Edwards and Parrett, 2002).

Cereal can be used as sources of nondigestible carbohydrates that besides promoting several beneficial physiological effects, can also selectively stimulate the growth of lactobacilli and bifidobacteria present in the colon and act as prebiotics. Barley contain water-soluble fiber such as B-glucan and arabinoxylan, oligosaccharides, such as galacto- and fructo-oligosaccharides and resistant starch, which have been suggested to fulfil the prebiotic concept (Charalampopoulos *et al.*, 2002). In addition, barley B-glucan has been reported to selectively support the growth of lactobacilli and bifidobacteria in rat experiments (Ryhanen *et al.*, 1996) and in in-vitro studies (Jaskari *et al.*, 1993).

Fermentation is an effective method of food preservation. The process of fermentation by lactic acid bacteria (LAB) is capable of lowering the pH to below 4 in food products, including barley-based fermenting cereal gruels used as infant foods. This results in growth reduction of pathogenic bacteria such as *B.cereus*, *E.coli*, *Salmonella* spp., *Shigella* spp. and *S.aureus*. (Nout *et al.*, 1989; Simango and Rukure, 1992 and Kingamkono *et al.*, 1995).

Cereals are limited in essential amino acids such as threonine, lysine, and tryptophan, thus making their protein quality poorer compared with animals, and milk (Chavan and Kadam, 1989). Lactic acid fermentation of barley has been found effectively to reduce the amount of phytic acid, tannins and improve protein availability (Chavan and Kadam, 1989). Increased amounts of riboflavin, thiamine, niacin, and lysine due to the action of LAB in fermented blends of cereals were also reported (Hamad and Fields, 1979). On the other hand, Khetarpaul and Chauhan (1990) reported improved minerals availability of cereal fermented with pure cultures of lactobacilli and yeasts.

The present paper aimed to study the effect of using *Bifidobacterium lactis* Bb12, and *Lactobacillus helveticus* (probiotic) on the quality of fermented barley porridge (as weaning food) during the storage period (21 days) at refrigerator temperature (4-5°C).

## **MATERIALS AND METHODS**

-Hull-less barley variety Giza 131 was obtained from the Barley Research Dept. Field Crops Research Insti. A.R.C. Giza, Egypt.

-Skim milk powder was imported from Holland,. It contained 36% protein, 51% lactose, 5% fat, 7.3% Ash and 4% moisture.

-Pure fructose produced and backed by Misr. Scientific Company, Egypt.

-Carrot, natural honey, rice powder were obtained from local markets.

-Starter cultures : *Bifidobacterium lactis* ( Bb 12) and *Lactobacillus helveticus* were supplied by Chr. Hansen Laboratories, Copenhagen, Denmark.

### **- Germination of barely seeds and getting the flour:**

Barly seeds was cleaned ,washed and soaked overnight. After that it incubated for 48 hour at 30°C to germination. Then it were dried in an oven at 50°C for 36 h. After that rotes were removed. The germinated barley was ground into flour by using mill machine and sieved through a 250 µm screen.

### **-Preparation of fermented barley porridge:**

Malted barley flour (140 g) was blended with 500 ml distilled water for 10 min. and cooked at 90°C for 5 min. Then the skim milk powder (120 g) and pure fructose (10 g) were added slowly to the mixture by using blender. The mixture was pasteurized at 90°C for 30 min., and cooled at 37°C. The mixture was inoculated with 2% *Bifidobacterium lactis* Bb 12 or *lactobacillus helveticus* culture under sterilization condition. The initial acidity of porridge was at pH value 6.7 and the fermentation was run to final pH of 4.8. Fermentation pasteurized carrot 10 g or honey 10 g or both (5g+5g) were added to the mixture, which was then stored at refrigerator temp. (5- 7°C) for 21 days.

-Moisture, protein and fiber contents were determined using the methods described by AOAC. (1990).

- Carbohydrate content was calculated as difference between total weight and the sum of moisture, fat, protein, fiber and ash contents,
- Energy value was calculated by the following formula : 3.47 K cal/ g for protein, 8.38 K cal/g for fat and 4.2 K cal / g for total carbohydrate content.
- pH values were measured according to Ling (1963) using digital pH meter, and total solids was determined by using Digital Refractometer
- Sugars and organic acids were analyzed by high performance liquid chromatography (HPLC) according to the method of Black and Bagley, (1978) and Adhikari *et al.*, (2000) .
- Total bacterial counts: Plate count (agar medium) determined according to the method of Lee *et al.*, (1973) plates were incubated for 24 hr at 37 °C.
- L. helveticus* were counted using MRS agar, while, *Bifidobacterium lactis* determined by MRS agar + 0.05% L-cystein- HCl according to the method of Dinaker&Mistry (1994).
- Staphylococcus spp.* were counted using Stph 110 media (Difco, 1974) incubated at 37 °C for 48 h
- Total coliform were estimated by plating suitable dilutions on violet Red Bile Agar medium as recommended by the APHA (1992). Plates were incubated for 25 h at 37 + 1°C.
- Sensory evaluation was carried out by a regular score panel according to Sanni *et al.*, (1998).
- Biological assay: Old male Albino rats were carried out 3-4 weeks (Food Technology Research Institute (ARE) ,Giza, Egypt.) were acclimatized on commercial chow (Protein 8.2%, Fat 2.6%, Ash 8.6%,Carbohydrate 62.2%, Fiber 7.6% and Moisture 10.8%). The animals were arranged in three groups with six rats in each group. A control group was fed on the commercial chow diet and drank tap water. The other groups were fed on the commercial chow diet and drank the following fermented porridge diluted with water by 80:20, (v/v) for six weeks. Diet and drinks were provided *ad libitum* and fresh fermented porridge were supplied twice a day.
- At the end of the experimental period the small intestines of rats were taken off and washed with 20 ml sterilized saline using a sterilized syringe in a sterilized flask and serial dilutions were done. Fecal samples were collected from rectum in a sterilized Petri dish and 1.0 g of feces was transferred to a flask with 99 ml sterilized water. Samples were analyzed immediately using aseptic sterile dilution technique as described by Klaver *et al.*,(1993).

## **RESULTS AND DISCUSSION**

Initial pH values of fermented porridge with *Bif.lactisBb-12* and *L. helveticus* ranged between 4.32 to 4.52 as listed in table (1). Little less acidic pH was observed after 21 days. It was clear that, *L. helveticus* gave the lowest value of pH in all blends compared with *Bif.lactisBb-12*. These results are in agreement with Ghaly *et al.*, (2003) and Zaki, Hala *et al.*, (2004) who used the whey and *L. helveticus* for producing pH 4.5 and 5.5. Sneath (1986) mentioned that the optimum pH for initial growth of *Bifidobacterium* was 6.5-

7.0.Also, Kabeir *et al.*, (2005) studied the growth of *Bif.longum* BB536 in fermented cereal porridge and their survival during storage up to 21 days at (4-5°C). The low pH value as well as presence of high *Bif. longum* BB536 count asserted the safety of the fermented cereals.

**Table (1): Changes of pH value of fermented porridge with *Bif.lactis* Bb12 and *L. heliviticus* during storage up to 21 days at (4-5°C).**

Storage period Treatments	Fermentation by <i>Bifidobacterium lactis</i> Bb12.			
	Zero time	7 day	15 day	21 day
B <sup>1</sup>	4.52	4.46	4.36	4.25
B <sup>1</sup> .H	4.46	4.36	4.34	4.27
B <sup>1</sup> .C	4.43	4.38	4.28	4.20
B <sup>1</sup> .C.H	4.40	4.36	4.29	4.23
	Fermentation by <i>L. heliviticus</i>			
B <sup>2</sup>	4.42	4.33	4.21	4.18
B <sup>2</sup> .H	4.38	4.24	4.19	4.16
B <sup>2</sup> .C	4.32	4.25	4.17	4.12
B <sup>2</sup> .C.H	4.37	4.21	4.14	4.12

B: barley, B.H: barley with honey, B.C barley with carrot, B.C.H: barley with carrot & Honey. 1 : *L. heliviticus*, 2 : *Bif.lactis* Bb-12 .

Data in table (2) show sugars acids profile of fermented carbohydrate using starter cultures during fermentation period. It was found that both glucose and fructose concentration in (B<sup>1</sup>.H & B<sup>2</sup>.H) treatment were higher than all other treatments. This may be due to the high ratio of inverted sugar in honey than carrot . In general glucose and fructose values decreased with increasing time of storage.

**Table (2) : Changes of glucose & fructose and the production of lactic acid & acetic acid in fermented porridge with *Bifidobacterium lactis* Bb12 and *L. heliviticus* during storage up to 21 days at (4-5°C).**

Storage period Treatments	Glucose %		Fructose %		Lactic acid %		Acetic acid %	
	Zero time	21 days	Zero time	21 days	Zero time	21 days	Zero time	21 days
	Fermentation by <i>Bifidobacterium lactis</i> Bb12							
B <sup>1</sup>	0.19	Nd	Nd	Nd	0.12	0.13	0.04	0.09
B <sup>1</sup> .H	6.58	5.24	5.12	2.20	0.09	0.10	0.02	0.04
B <sup>1</sup> .C	2.77	1.16	3.34	1.56	0.01	0.08	0.02	0.04
B <sup>1</sup> .C.H	3.89	3.19	2.96	1.19	0.09	0.12	0.02	0.04
	Fermentation by <i>L. heliviticus</i>							
B <sup>2</sup>	1.05	0.47	Nd	Nd	0.09	0.18	0.17	0.23
B <sup>2</sup> .H	6.33	1.76	3.20	1.28	0.12	0.13	0.22	0.34
B <sup>2</sup> .C	1.36	0.58	1.30	0.86	0.14	0.15	0.28	0.37
B <sup>2</sup> .C.H	2.19	0.66	1.26	0.52	0.11	0.12	0.18	0.24

B: barley, B.H: barley with honey, B.C barley with carrot, B.C.H: barley with carrot & Honey. 1 : *L. heliviticus*, 2 : *Bif.lactis* Bb-12 .

Sneath *et al.*, (1986) and Holt *et al.*, (1994) described the complete fermentation of sucrose to the high ability of *Bifidobacterium Bb12* to ferment sucrose to fructose and glucose which fermented to acetate and lactate. In case of *L.heliveticus* the values of glucose and fructose were higher than that obtained by *Bif.lactis Bb-12*. Treatment (B<sup>2</sup>.H)& (B<sup>1</sup>.H) had similar trends and the highest values of glucose and fructose. The primary functional properties for lactic acid starter bacteria used in making fermented food products are their ability to produce organic acid by the fermentation of sugar.

The production of lactate and acetate in fermented porridge are shown also in table (2). It could be seen that lactic acid content increased during storage up to 21 days in all treatments. Meanwhile, (B<sup>1</sup>.C) had the highest value of lactic acid and acetic acid percent. These data agreed with (Kabeir *et al.*, 2005) who reported the low pH (< 4.6) of fermented porridge as results of organic acid such as lactic acid and acetic acid was critical for *Bif.longum*.

Data in table (3) shows the changes in chemical contents of fermented blends at zero time as stored up to 21 days. In case of fermented porridge by *L.heliveticus* the values of moisture, protein, ash, fiber and T.S.S were rather lower but had the same trend as shown in porridge with *Bif.lactis Bb-12*. This may be due to the lower activity of *Bif.lactis Bb-12*. on carbohydrate fermentation. These results agreed with results obtained by (Zaki, Hala 2004).

Ash percentage values, were high in (B<sup>1</sup>) & (B<sup>2</sup>) samples at zero time, and after 21 days of storage, this may be due to the high values of ash in plain barley than carrot or honey; according to FAO (1982), while barley grains, malted barley, carrot and honey (on wet weight basis) had 3.9, 1.8, 0.7, & 0.2% ash respectively.

Fiber percentage values were degraded in all samples after 21 days of storage period. This could be resulted from the fermentation activity of the starter cultures. The results were in harmony with those obtained by Lambo *et al.*, (2005) who indicated that insoluble fiber of barley and oat decreased after fermentation by some strains of L.A.B.

Carbohydrate content values decreased in the following B<sup>1.2</sup>.H, B<sup>1.2</sup>.C.H, B<sup>1.2</sup>.C & B<sup>1.2</sup>.B. The high values of honey treatments may be due to the high ratio of carbohydrate in it. All treatments after 21 days of storage were of lower carbohydrates in comparison with zero time. These data agreed with that obtained by Basyony *et al.*, (2002).

The values of T.S.S in (B<sup>1.2</sup>.C) were lower than (B<sup>1.2</sup>.H & B<sup>1.2</sup>.C.H). These may be due to the culture activity, acid production and degradation of carbohydrate. The same trend had been obtained by Basyony *et al.*, (2002).

Energy in fermented porridge with *Bif.lactis Bb-12* was higher than porridge with *L.heliveticus*. This may be due to differences of carbohydrate content.

The obtained data in table (4) clearly show higher contents of calcium, phosphorus and iron in treatments (B<sup>1</sup> & B<sup>2</sup>) which consisted of plain barley without honey or carrot. The available ingredients levels affect markedly the minerals of final products (Flynn and Cashman,1997). While, the treatments which consist of high ratio of carrot (B<sup>1</sup>.C. & B<sup>2</sup>.C.) showed high content of potassium 232 and 231 mg/100g porridge respectively.

Consumption of 100g/days of these formulas would be sufficient to meet the daily requirements of phosphorus and magnesium, while 180-230 g/days of these porridges would be enough to cover the RDA of calcium and iron. On the other hand, all prepared porridge contain low levels of sodium and potassium comparing with the permitted amounts RDA (1989).

Also, table (4) indicated that the values of minerals in fermented formula using *L.helveticus* were in general following the same trend of *Bif.lactis Bb-12*.

Also, the data in Table (4) indicated that all prepared porridge samples contained low level of vitamin A comparing with the RDA (1989) which recommended the daily vitamin A intake by infant to be 375 RE/day (1 RE = 6 µg of all trans β-carotene). On the other hand treatments (B<sup>1</sup>.C& B<sup>2</sup>.C) contained relatively highest values of vitamin A (38.9& 26.6 µg<sup>RE</sup>). This may be due to the high ratio of vitamin A in carrot. While, it had high values of thiamine and riboflavin. The high amounts of thiamine may be due to using malt flour in preparing each formula but high ratio of riboflavin may be due to using skim milk powder.

All treatments of fermented porridge with *L.heliveticus* had high values of thiamine and riboflavin than that fermented by *Bif.lactis Bb-12*

The production of lactic acid bacteria during fermentation period lead to 50% higher in the concentration of thiamine and riboflavin in comparing with the initial concentrate in these vitamins (Alm, 1982). Moreover (Gurr, 1987) found the increase in the vitamin content in the formulated cereals blend at the end of the fermentation.

Microbiological analysis of different treatments in zero time and after storage up to 21 days at 4-5°C, using *L. heliveticus* and *Bif. lactis Bb12* were determined.

Data obtained in fig. (1) show the *L. heliveticus* count in fermented porridge during the storage up to 21 days at 4-5°C. The counts gradually increased during the storage period (7 days). The count of this strain decreased after storage period (21 days).

Also, data shown in fig. (2) indicate the total count of *Bifidobacterium lactis Bb-12* in the porridge. It might be observed from the results that the maximum population count occurred after 15 days of storage at (4-5 °C). On the other hand at the end of storage period (21 days) data showed marked decrease in the population. This trend agreed with that obtained by (Kabeir *et al.*, 2005) and (Kim *et al.*, 2000). It was clear from figures (1 & 2) that the growth of *Bifidobacterium lactis Bb-12* was lower than *L.heliveticus*. These results were in line with that obtained by Zaki, Hala *et al.*, (2004).

Because foods containing probiotic bacteria should contain at least 6 or 7 log cfu live microorganism per gram or per milliliter at the time of consumption, in order to benefit the consumer (Ishibashi and Shimamura 1993), all samples of treatments were fully considered as probiotic food.

Coliforms were not detected in all treatments either when zero time or during the storage period. Kunene *et al.*, (1999) and Kingamacono *et al.*, (1995) explained the reduction in level of coliform bacteria by the production of organic acid which would reduce the proliferation of gram negative bacteria and bacterial spores found in fermented porridge.

T3

***Yonis, A.A.M. and Hanaa F. El-Meheiry.***

t4

**10842**

f1-2

Sensory evaluation scores of fermented porridge with *L.heliveticus* & *Bifidobacterium lactisBb12* during storage up to 21 days was presented in Table (5). Results show that the scores gained for colour, flavour, consistency, taste, and texture. The consumer acceptability of porridges production from the different blends showed that all the samples rated above average provided that B.C& B.C.H had the highest preference among treatments which fermented with *Bifidobacterium lactisBb12* or *L. heliveticus*

**Table (5): Sensory evaluation of fermented barley with *Bif.lactisBb12* and *Lb.heliveticus* after storage up to 21 days at 4-5°C.**

Treatments		Treatments of porridge with <i>Bif.lactisBb12</i> and <i>L. heliveticus</i>							
		B <sup>1</sup>	B <sup>2</sup>	B <sup>1</sup> .H	B <sup>2</sup> .H	B <sup>1</sup> .C	B <sup>2</sup> .C	B <sup>1</sup> .C.H	B <sup>2</sup> .C.H
Colour	10	7.00 <sup>C</sup>	7.00 <sup>C</sup>	6.25 <sup>C</sup>	7.75 <sup>C</sup>	8.00 <sup>B</sup>	9.00 <sup>B</sup>	7.50 <sup>A</sup>	8.00 <sup>A</sup>
Flavor	10	7.25 <sup>B</sup>	7.00 <sup>C</sup>	8.50 <sup>B</sup>	7.50 <sup>B</sup>	8.50 <sup>B</sup>	8.50 <sup>B</sup>	7.50 <sup>B</sup>	8.25 <sup>B</sup>
Consistency	10	6.25 <sup>B</sup>	7.00 <sup>C</sup>	7.00 <sup>B</sup>	6.50 <sup>B</sup>	8.00 <sup>B</sup>	8.50 <sup>B</sup>	7.00 <sup>A</sup>	8.00 <sup>A</sup>
Taste	10	6.00 <sup>B</sup>	6.75 <sup>C</sup>	7.50 <sup>B</sup>	7.75 <sup>C</sup>	8.75 <sup>A</sup>	9.00 <sup>B</sup>	8.50 <sup>A</sup>	7.50 <sup>A</sup>
Texture	10	5.50 <sup>C</sup>	6.00 <sup>C</sup>	7.00 <sup>B</sup>	6.50 <sup>B</sup>	8.00 <sup>B</sup>	8.75 <sup>B</sup>	7.00 <sup>A</sup>	8.25 <sup>A</sup>
Total	50	32.00	33.75	36.75	36.00	41.25	43.75	37.50	40.25

B: barley – B.H: barley with honey. – B.C: barley with carrot. –B.C.H: barley with carrot & Honey. 1 : *Bif.lactis Bb-12* 2 : *L. heliveticus*

Table (6) results show the growth parameters of rats. Rats fed on the fermented porridge (B<sup>1</sup>.C & B<sup>2</sup>.C) exhibited significantly higher final body weight and body weight gain than the control groups. However, rats` relative liver and spleen weights (organ weight / 100g body weight ) were more or less similar among all groups. These results may be due to probiotic bacteria which produces organic acids such as lactic acid and other by-products such as hydrogen peroxide and antibiotics, and breaks down the bile acids, that improve the intestinal flora and create an environment for the efficient utilization of nutrients (Nakazawa and Hosono, 1992).

**Table (6): Growth parameters of rats fed on fermented porridge (BC) by *L. heliveticus* or *Bif.lactis Bb-12*.**

Diet groups	Initial body weigh (g)	Final body weigh (g)	body weigh gain(g)	Liver %	Spleen %
Control	63.2 + 3.9	106 + 12.6	43.1 + 10.0	4.47 + 0.1	0.45 + 0.05
B <sup>1</sup> .C	74.2 +7.5	161 + 9.5	87.2 + 10.5	4.77 + 0.4	0.64 + 0.05
B <sup>2</sup> .C	67.4 + 3.3	169 + 3.9	102 + 3.9	4.38 + 0.3	0.44 + 0.05

– B.C: barley with carrot. 1 : *Bif.lactis Bb-12* 2 : *L. heliveticus*

The effect of feeding the fermented porridge (B<sup>1</sup>.C & B<sup>2</sup>.C) on small intestinal and feces content of *L. heliveticus* , *Bifidobacteria spp.*, staphylococci and coliform population is shown in table (7). In comparison with the control group, rats fed the fermented porridge resulted in a considerable increase in their intestinal content of *L. heliveticus*, *Bifidobacteria spp.*, however, the count of staphylococci and coliforms significantly decreased. These results are in agreement with that obtained by

Patel *et.al*,(1992) Several studies showed the increase of (LAB) & bifidobacteria population and reduction of pathogens microorganisms in human and rats`intestinal and feces by feeding on fermented products by (LAB) and *Bifidobacteria spp.* ( Silva *et al.*, 1999, Meddah *et al.*, 2001 and Bruno & Shah, 2002).

**Table (7): Effect of fermented porridge (BC) with *L. heliveticus* or *Bif.lactis Bb-12* on small intestinal and feces.**

Parameters (log cfu/ml)	Control	B. <sup>1</sup> C	B. <sup>2</sup> C
<b>Small intestinal</b>			
<i>L.heliveticus</i>	7.1 + 0.09	10.3 + 0.12	10.5 + 0.13
Bifidobacteria	6.8 + 0.14	9.1 + 0.22	9.4 + 0.11
Coliforms	4.2 + 0.10	2.9 + 0.23	2.4 + 0.13
<i>Staphylococci</i>	4.0 + 0.20	3.4 + 0.19	2.9 + 0.20
<b>Staphylococcc Feces</b>			
<i>L. heliveticus</i>	7.5 + 0.04	10.2 + 0.04	10.3 + 0.04
Bifidobacteria	6.6 + 0.07	6.6 + 0.12	9.7 + 0.07
Coliforms	4.8 + 0.11	3.1 + 0.12	3.1 + 0.09
<i>Staphylococci</i>	4.2 + 0.03	4.0 + 0.11	2.7 + 0.08

B.C: barley with carrot. 1 : *Bif.lactis Bb-12* 2 : *L. heliveticus*

## REFERENCES

- Adhikari, L., Mustapha,A., Gru,I.U. and Fernando,L. (2000). Viability of microencapsulated bifidobacteria in set yoghurt during refrigerated storage.J.Dairy Sci. 83:1946-1951.
- APHA (1992). Standard methods for examination of dairy products. American U.S.A public Health. Assoc. 16<sup>th</sup> Ed, Washington D.C.
- Alm,L. (1982). Effect of fermentation on B-vitamin content of milk in Sweden. Journal of Dairy Science, 65:'353-359.
- AOAC (1990). Official Methods of Analysis. Arlington Virginia, of The Association of Official Analytical Chemists, Inc., USA.
- Barnes, G.; Beaton, S. and Goldenberg, N. (1979). Royal Society of health .J.,99 (3): 107. c.f. Robinson, R.K. (1990). Dairy Microbiology: the microbiology of milk product. Vol. 2, 2nd Ed., Elsevier Applied Science., London and New York.
- Basyony,A.E.; Abd-Elrahman. A. H.; and Assem, A.H.(2002). Production of yoghurt like using hull-less barley as a substitution material. Egypt. Journal Agriculture, 82(3).
- Black,L.T.and Bagley,E.M. (1978). Determination of oligosaccharides in soybeans by high pressure liquid chromatography using an internal standard. J. of the American Oil Chemists Society .55:228.
- Bruno,F.A. and Shah,N.P.(2002).Inhibition of pathogenic and putrefactive microorganisms by *Bifidobacterium spp.* Milchwissenschaft,57:612-621.

- Charalampopoulos,D.; Pandiella, S. S. and Webb, C. (2002). Growth studies of potentially probiotic lactic acid bacteria in cereal based substrates. *Journal of Applied Microbiology.*, (92): 851-859 .
- Difco Manual of Dehydrate Culture Media and Reagents .(1974).Pub. Difco Laboratory Incorporated, Detroit, Michigan, U.S.A.48:201.
- Dinaker, P. and Mistry, V.V. (1994). Growth and viability of *Bifidobacterium bifidum* in cheddar cheese. *J. Dairy Sci*, 77 (10):2854-2864.
- Edwards,C.A. and Parrett, A.M. (2002). Intestinal flora during the first months of life: new perspectives. *British J. of Nutr.*88:11-18.
- FAO (1982). Food Composition Tables for the Near East, Food & Nutrition Paper No. 26, Food & Agriculture Organization of the United Nation, Rome.
- Flynn,A. and Cashman, K. (1997). Nutritional aspects of minerals in bovine and human milk., *Advance Dairy Chemistry*, Vol.3, pp 127 – 154, London, Chapman & Hall.
- Food and Nutrition Board: Recommended Dietary Allowances, ed 10, National Research Council, Washington, DC, 1989 National academy press.
- Ghaly,A.E., Tango,M.S.A. and Adams,M.A. (2003). Enhanced lactic acid production from cheese whey with nutrient supplement addition. *Agricultural Engineering International: the CIGR Journal of Scientific Research and Development Manuscript FP02009.*
- Gurr,M.I. (1987). Nutritional aspects of fermented milk products. *FEMS Microbiology Reviews*, 46:337-342.
- Hamad,A.M. and Fields,M.L. (1979). Evaluation of the protein quality and available lysine of germinated and fermented cereals. *J.Food Sci.* 44:456-459.
- Chavan,J.K. and Kadam,S.S. (1989). Nutritional improvement of cereals by fermentation. *Crit.Rev. Food Sci. Nutr.* 28:349-400.
- Holt,J.G., Krieg,N.R. Sneath, P.H.A., Staley, J.T. and Williams, S.T. (1994). *Bergeys Manual of Determinative Bacteriology*. 8<sup>th</sup>.ed Williams and Wilkins, Awavelry Co.
- Ishibashi,N., and Shimamura, S. (1993). Bifidobacteria: research and development in Japan. *Food Technol.*, 46: 128–135.
- Jaskari,H. Salovaara,T. Mattilla-Sandholm and Putanen,K. (1993). The effect of oat  $\beta$ -glucan on the growth of selected *Lactobacillus* spp. And *Bifidobacterium* spp.In: T.Aalto-Kaarlehto and H. Salovaara,Editors, Proceeding of the 25<sup>th</sup> Nordic Cereal Congress, Univ. Helsinki, Helsinki, pp.242-244.
- Kabeir,B.M., Abd-Aziz,S., Muhammad, K., Shuhaimi, M. and Yazid, M.A. (2005). Growth of *Bifidobacterium Longum* BB536 in medida (fermented cereal porridge) and their survival during refrigerated storage. *Letters In Applied Microbiology.*, 41:125–131.
- Khetarpaul and Chauhan, Khetarpaul,N. and Chauham,B.M. (1990). Effect of fermentation by pure cultures of yeasts and lactobacilli on the available carbohydrate content of pearl millet.*Trop.Sci.*31:131-139.

- Kim,H., Min,J., Lee,J. and Ji,G. (2000). Growth of lactic acid bacteria and bifidobacteria in natural media using vegetable, seaweeds, grains and potatoes. *Food Sci. Biotechnol.*, 9: 322–324.
- Kingamacono, R., Sjogren,E., Svanberg,U. and Kaijser,B. (1995). Inhabitation of deferent strains of entropathogenies in lactic fermenting cereals gruel. *World of Journal Microbial. Biotechnology.*, 11: 299-303.
- Klaver,F.A.M., Kingma,F. and Weerkamp, A.H. (1993). Growth and survival of bifidobacteria in milk. *Netherlands. Milk Dairy J.*,47:151-164.
- Kunene,N.F., Hastings,J.W., and Von Holy,A. (1999). Bacterial populations associated with a sorghum-based fermented weaning cereal., *International Journal of Food Microbiology.*, 49: 75–83 .
- Lambo,M.A. Margareta,O.R. and Nyman,G.E. (2005). Dietary fiber in fermented oat and barley  $\beta$ -glucan rich concentrates. *Food Chemistry.*, 89: 283–293.
- Lee,S.Y., Vedemuthu,E.R. Washam,C.J. and Reinbold,B.W. (1973). An agar medium for the differential enumeration of yoghurt starter bacteria. *J. Milk Food Tech.*, 9(37): 272-275.
- Ling,E.R. (1963). *A Text Book of Dairy Chimistry*. Vol.Π Chapman and Hall Ltd., London , UK 3rd Ed.
- Meddah,A.T., Yazourh,A., Risbourg,B., Verstraete,W. and Romond,M.B. (2001). The regulatory effects of whey retenate from bifidobacteria fermented milk on the microbiota of the simulator of the human intestinal microbial ecosystem (SHIME).*J.Appl. Microbiol.*,91:1110-7.
- Nakazawa,Y. and Hosono,A. (1992). *Functions of Fermented Milk*,Cholleges for Health Sciences. Elsevier Science publishers, London.
- Nout,M.J.R., Rombouts.F.M. and Hautvast,G.J. (1989). Accelerated natural lactic acid fermentation of infant food formulations.*Food Nutr. Bull.* 11:65-73.
- Patel,J.R., Dave,J.M. and Sannabhdti,S.S. (1992).Effect of feeding milk fermented with mixed culture of human stain of Lactobacilli on a faecal Lactobacilli and Coliform counts in human test subjects. *Indian J.Dairy Sci.*,45:379-382.
- Ryhanen,E.L., Mantere-Alhonen,S. and Salvaara,H. (1996). Effect of oat bran and rye bran diet on intestinal Lactobacillus and Bifidobacterium flora on Wistar rats.in:Y.Malkki and J.H.Cummings, Editors, *Dietary Fiber and Fermentation in the Colon*,Office for Official Publications of European Communities, Luxembourg, 55-57.
- Sanni,I.A., Onilude,A.A. and Ibidapo,T.O. (1998). Biochemical composition of infant weaning food fabricated from fermented blends of cereal and soybean. *Food Chemistry.*, 65 : 35-39.
- Silva,A.A., Bambirra,E.A., Oliveir,A.L., Souza,P.P., Gomes,D.A., Vierira,E.C. and Nicoli,J.R. (1999). Protective effect of bifidus milk on the experimental infection with *Salmonella enteritidis* subsp. Typhimurium in conventional and gnotobiotic mic. *J.of Appl. Microbiol.*, 86:331-336.
- Simango,C. and Rukure,G. (1992). Survival of bacterial enteric pathogens in traditional fermented foods. *J.Appl. Bacteriol.* 73:37-40.

- Sneath, P.H.A., Mair, N.S., Elisabeth Sharpe, M. and Holet, J.G. (1986). Bergey's manual of systematic bacteriology. Vol.2. Williams and Wilkins, London.
- Tojo, R., Leis, R., Pavon, P. and Moran, J. (1995). Leche humanay formulas infantiles: comparacion nutricional,- in New Perspectives- in Infant Nutrition. Ergon, Madrid, pp.23-28.
- Vandenplas, Y. (2002). Oligosaccharides in infant formula. Br J Nutr. 87:293-296.
- Zaki, M. Hala., Saleh, F.A. and Ahmed, A.I. (2004). Production of functional food using bacterial fermentation. Egypt. J. Agric. Res. 82:1-12.

## **Lactobacillus helveticus or Bifidobacterium lactis Bb12**

علاء الدين أحمد مرسى يونس و هناء فاروق المهيري  
قسم الإقتصاد المنزلي - كلية التربية النوعية- جامعة المنصورة

تم في هذه الدراسة تحضير أغذية متخمرة باضافة *Lactobacillus helveticus*

### **(1) or Bifidobacterium lactis Bb12 (2)**

للأطفال ذات قيمة غذائية عالية من دقيق الشعير العارى المنبت, و دقيق الأرز, و اللبن الفرز المجفف, و العسل, و الجزر بنسب مختلفة . حيث تم تقييم هذه الوجبات المحضرة كيميائيا, و ميكروبيولوجيا, و حسيا, و بيولوجيا بعد تجهيزها مباشرة و خلال فترة التخزين لمدة 21 يوم على درجة حرارة التلاجة (4-5م) .  
و قد أوضحت النتائج زيادة كل من البروتين و الرماد و حمض اللاكتيك و حمض الخليك بعد فترة التخزين, بينما أنخفضت قيم كل من الألياف و الكربوهيدرات و الجلوكوز و الفركتوز.

أختلفت قيم الأس الهيدروجيني نتيجة لأختلاف نمو السلالتين فقد سجلت السلالة (1) أقل القيم و هي (4.12) . كما أظهر العد الكلي لبكتريا السلالة (1) أعلى معدل نمو و حتى اليوم السابع في كل المعاملات ثم بدأ في الانخفاض ببطئ حتى نهاية فترة التخزين. بينما كان أعلى معدل نمو لبكتريا السلالة (2) عند اليوم 15 ثم حدث انخفاض واضح في العدد حتى نهاية التخزين .

كما أوضحت النتائج ارتفاع محتوى العينات من أملاح الكالسيوم و المغنسيوم و فيتاميني ب1 و ب2 لتصل الى حدود الاحتياجات اليومية حسب التوصيات العالمية. أما عن النتائج البيولوجية فقد أوضحت أن المنتجات المتخمرة أدت الى زيادة معنوية في وزن جسم الفئران مقارنة بمجموعة المقارنة بدون أى تأثير على الوزن النسبي (الوزن / 100 جم من وزن الجسم ) للكبد و الطحال . و بالمقارنة مع مجموعة المقارنة وجد أن الفئران التي غذيت على المنتجات المتخمرة تميزت بزيادة أعداد بكتريا السلالة (1 و 2) في أمعائها و برازها في حين حدث انخفاض كبير في أعداد بكتريا القولون و البكتريا العنقودية.

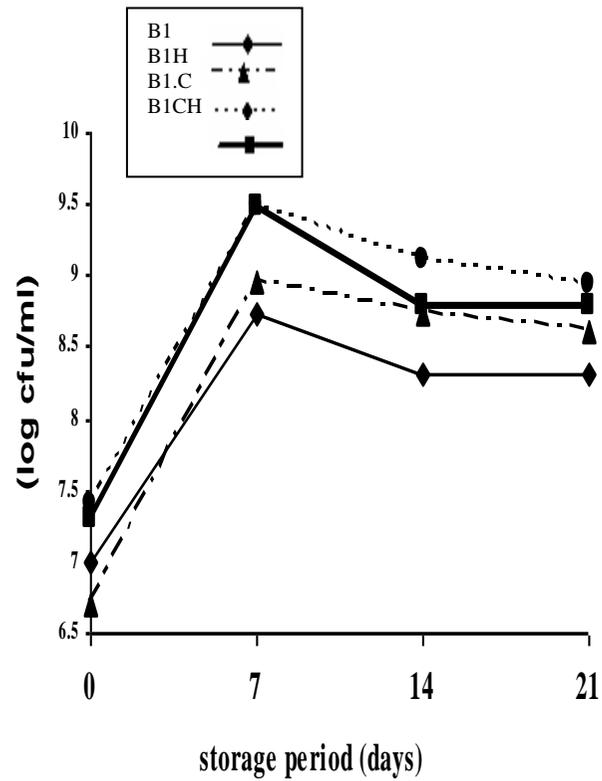


Fig (1): Viability of *L. helveticus* count (log cfu/g) during storage up to 21 days at (4-5°C).

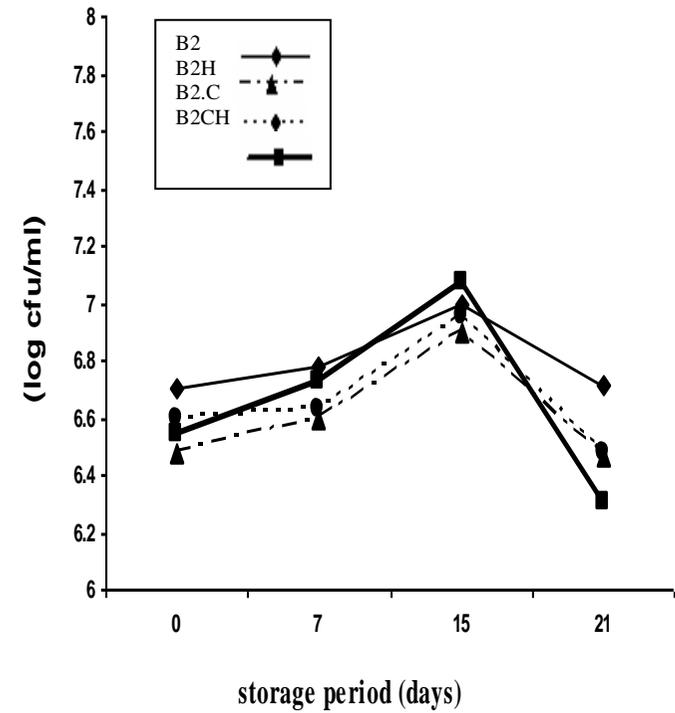


Fig (2): Viability of *Bifidobacterium lactis* Bb-12 count (log cfu/g) in fermented porridge during storage up to 21 days at (4-5°C).

**Table (3): Changes of chemical composition in fermented porridge with *Bif.lactis Bb-12* and *L. helveticus* after storage up to 21 days at (4-5°C) (on wet weight basis).**

Composition %	Treatment of fermented porridge with <i>Bifibobacterium lactis Bb12</i>								Treatment of fermented porridge with <i>Lactobacillus helveticus</i>							
	B <sup>1</sup>		B <sup>1</sup> .H		B <sup>1</sup> .C		B <sup>1</sup> .C.H		B <sup>2</sup>		B <sup>2</sup> .H		B <sup>2</sup> .C		B <sup>2</sup> .C.H	
	0 time	21 days	0 Time	21 days	0 Time	21 days	0 Time	21 days	0 time	21 days	0 time	21 days	0 time	21 days	0 time	21 days
Moisture	78.37	77.79	78.12	77.76	79.88	79.85	77.14	76.92	76.86	75.36	75.88	74.69	75.55	74.64	75.79	74.80
Protein	5.70	6.80	4.35	5.45	5.00	6.25	5.20	6.10	4.90	6.20	4.10	5.14	4.51	5.85	5.10	5.50
Ash	1.20	1.30	0.83	1.16	0.95	1.27	0.91	1.21	1.15	1.30	0.70	0.80	0.90	1.10	0.80	1.08
Fiber	1.00	0.80	0.80	0.68	1.90	1.70	0.70	0.50	1.07	0.91	0.90	0.81	2.10	1.60	1.10	0.90
Carbohydrate	13.73	13.31	15.90	15.04	12.27	10.93	15.89	15.27	16.73	16.73	18.42	18.56	16.94	15.41	17.21	17.02
T.S.S	12.00	14.60	25.00	26.50	13.60	15.60	18.50	19.60	11.20	12.80	22.50	20.90	12.80	11.40	17.80	16.20
Energy (K.cal)	77.72	80.44	81.00	81.96	69.08	68.72	84.36	85.48	86.52	90.52	90.08	95.16	85.80	85.04	89.24	90.08

**B: barley – B.H: barley with honey– B.C: barley with carrot –B.C.H: barley with carrot & Honey  
1: *Bif.lactis Bb-12* 2: *L.helveticus***

**Table (4) Minerals and vitamins contents of fermented porridge mg/100g porridge (on wet weight basis).**

Treatments	Fermented porridge with <i>Bifibobacterium lactis Bb12</i>												
	Ca	P	Mg	Fe	Zn	Cu	Na	K	Vitamin A		Vitamin C (mg/ 100g)	Thiamine (mg/ 100g)	Riboflavin (mg/ 100g)
									β-carotene (µg /100g)	Vitamin A (µg <sup>RE*</sup> )			
B <sup>1</sup>	223	323	47	3.07	1.77	0.24	195	223	16.0	2.6	1.9	17.4	3.0
B <sup>1</sup> .H	198	321	46	2.82	1.45	0.12	222	216	27.0	4.5	2.4	17.6	3.0
B <sup>1</sup> .C	178	315	46	2.40	1.25	0.24	221	232	133.7	38.9	4.5	17.6	4.4
B <sup>1</sup> .C.H	190	320	47	2.67	1.17	0.17	243	225	91.9	18.9	3.3	17.9	2.0
	Fermented porridge with <i>Lactobcillus helveticus</i>												
B <sup>2</sup>	225	328	48	3.13	1.72	0.27	201	222	19.0	1.5	1.8	23.8	3.6
B <sup>2</sup> .H	197	324	45	2.74	1.39	0.10	220	209	25.0	2.8	4.1	23.2	3.4
B <sup>2</sup> .C	176	313	45	2.35	1.29	0.26	217	231	159.0	26.6	6.0	24.8	4.8
B <sup>2</sup> .C.H	188	317	49	2.59	1.24	0.19	244	223	99.0	16.5	5.6	22.8	3.6
R.D.A (mg) Infants (0- 5)months)	400	300	40	6	5	0.4	120**	500**	375 (µg <sup>RE</sup> )		30 (mg)	0.4 (mg)	0.5 – 0.8 (mg)

B: barley – B.H: barley with honey – B.C: barley with carrot –B.C.H: barley with carrot & Honey. 1: *Bif.lactisBb-12* 2: *Lb.helveticus* RE\*: retinol equivalents. R.D.A: Recommended Dietary Allowances (1989).

\*\* : Minimum requirements.