Biochemical and Microbiological Evaluation of Fermented Camel Milk

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Abstract: Camel milk has an important role in human nutrition in the hot regions and arid countries. The general composition of camel milk varies in various part of the world with range of 3.07-5.50% fat, 3.5-4.5% protein, 0.7-0.95% ash and 3.4-5.6 % lactose, 12.1-15% total solid. Camel milk is different from other ruminant milk, having low cholesterol, low sugar, high minerals (sodium, potassium, iron, copper, zinc and magnesium), high vitamin C. Camel milk contains more proteins and whey protein than cow milk. Camel milk is remedy for viruses causing diarrhea as Rota Virus considering it could be important from public health point of view to anticipate the vital hazard in camel raw milk and its produces. In this study, we used rat model of rotavirus infection which causing diarrhea. This investigation was designed to prepare fermented Camel milk with low fat (1.5%) and using it in formulas for feeding diarrhea rats (25%, 50% and 75%) to perform the role of the macro elements (sodium, potassium, phosphorous and calcium) as well as detection of Staphylococcus aureus of the three prepared formulas for diarrhea rats. Results indicated that the fermented camel milk had higher content in sodium and potassium than the raw camel milk and there was stopping diarrhea on formula (3), this is due to the formula (3) can be considered antimicrobial and a strong vital inhibitor to human Rota Virus which represents the main diarrhea-causing agent in infants. It can be concluded that fermented camel milk can be considered as a good food of high nutritive and therapeutic applications. Meanwhile, the high content of antimicrobial agents in camel milk may explain its potential as an activity especially against diarrhea.

Key words: Camel milk- diarrhea-antimicrobial- minerals

Introduction
Camel’s milk in particular is a good source of various vitamins and minerals and is characterized for its low cholesterol and high concentration of insulin Agrawal et al., 2005 & (Badriah., 2012). Camel, which is also known as ship of desert, is used for transportation and a source of milk, meat and wool (Meiloud et al., 2011). It has medicinal properties and antibacterial and antiviral activity (El-Agamy et al., 1992). which may be due to higher concentration of lactoferrin in camel’s milk (Yagil, 1994). Camel milk is used therapeutically against hepatitis, dropsy, problems of spleen, asthma, (Mal et al., 2000. Camel milk is also having low cholesterol, low sugar, high minerals (sodium, potassium, iron, copper, zinc and magnesium), high vitamin C, low protein and large concentration of insulin (Agrawal et al., 2004) & ( Arrowal,.2005). There are no allergens, and it can be consumed by lactase deficient persons and those with weak immune systems. The milk is considered as having medicinal properties. In Sahara, fresh butter is not eaten, but is often used as base for medicines. The products developed also include cosmetics or pharmaceuticals. It claimed that the value of camel milk is to be found in the high concentrations of volatile acids especially, linoleic acid and polyunsaturated acids, which are essential for human nutrition (Agrawal et al., 2004). A series of metabolic and autoimmune diseases are successfully being treated with camel milk. In India, camel milk is used therapeutically against dropsy, jaundice, problems of spleen, asthma, anemia, piles and diabetes role of raw camel milk in chronic pulmonary tuberculosis patients has been observed (Mal et al., 2000 & Agrawal et al., 2004).

Milk plays a significant role in human’s nutrition for conducted to test the effect of camel’s milk in comparison the wonderful reason that they are excellent source of to cow’s milk on blood glucose level, triglycerides, various nutrients. Milk diet has been suggested in the cholesterol and ALT and AST levels in healthy albino management of various diseases (Badriah., 2012).

In several parts of world, camel milk is used as a remedy for some diseases as tuberculosis, juvenile diabetes (Beg et al., 1985), liver cirrhosis, rickets, constipation, asthma (Yagil, 1987). Camel milk was not only contains more nutrients compared to cow milk, but also it has therapeutic and antimicrobial agents (Barbour et al, 1984 & El-Agamy et al., 1992). The antiviral activity in camel milk was also found (El-Agamy et al., 1992). Camel milk is remedy for viruses causing diarrhea as Rota Virus considering it could be important from public health point of view to anticipate
the vital hazard in camel raw milk and its produces. Therefore, Camel milk can be considered a strong vital inhibitor to human Rota Virus which represents the main diarrhea-causing agent in infants (El-Mougi, 1999).

In the present study, we prepared fermented camel milk with low fat (1.5%) in formulas for feeding diarrhea rats and perform the role of the macro elements (sodium, potassium, phosphorous and calcium) as well as detection of Staphylococcus aureus of the three prepared formulas for diarrhea rats.

Material & Methods
I- Materials:
(A). Camel milk used in the current study obtained from local market at GIZA, Egypt freeze dried direct vet set (FD-DVS) MYE 96 yoghurt culture was obtained from Rodia Food France. The culture was intended for direct inoculation to the process milk at 0.1%.

(B). Animals: Twenty four Sraye-Dawley White Albino rats, of an average weight 90 ± 0.2 gm were used in this study. The rats were obtained from experimental animal house in Food Technology – Research Institute, Agriculture Research Centre, Giza - Egypt.

Animal inoculation: Rats were inoculated by oral gavage with 0.5 ml phosphate-buffered saline (PBS), 6.75 x 108 PFU of RRV, or 1.45 x 108 PFU of HAL1166 rotavirus. Control animals were always handled prior to virus-inoculated animals. To determine the serum 50% antigenemia dose (AD50) and the 50% diarrhea dose (DD50) of RRV, rats were inoculated with 0.5 ml of serial 10-fold dilutions of RRV. For a subset of experiments, the rats were examined daily for evidence of rotavirus-induced diarrhea by gentle abdominal palpation. Diarrhea was noted and scored from 0 to 4 based on stool color, amount, and consistency. A score of 2 or greater was considered diarrhea ( Ciarlet, 2002).

II-Methods:
Biochemical analysis
Milk sample were analyzed for Moisture, pH, titratable acidity, specific gravity, total solids, and fat according to Ling (1963).

Determination of minerals:
Calcium, Sodium, Phosphorus, Potassium, Iron, Zinc concentration were determined by atomic absorption (Thermo-Tarrell, Ash, Smith-Hiefje (1000) in their digested solutions according to A.O.A.C (2000).

Protein determination: Protein concentration was measured spectrophotometrically according to AOAC (1980).

Prepared fermented Camel milk
Fermented Camel milk was prepared with low fat (1.5%). Camel milk was heated after decreased fat to 1.5% at 90°C for 10 min, immediately cooled to 42°C and inoculated with 0.1% DVS yoghurt culture. The inoculated milk was dispensed into plastic cups, fitted with press on lids. Cups were incubated at 42°C until titratable acidity of 0.8% was reached. Control diet (Casein-based diet) was composed of 11.6% case in (equal 10% protein), 5% corn oil, 4% mineral mixture, 1%vitamin mixture, 5% cellulose and corn starch up to 100%.

Microbiological analysis
Fermented camel milk samples (11 ml) were homogenized for one minute in 99 ml (1/10) of a sterile solution of 0.1% (w/v) peptone water (Oxoid CM9) using a Stomacher Lab blender (Model400, Seward Laboratory, London). From these samples serial decimal dilutions were prepared in sterile 0.1% peptone water. The microorganism's counts were carried out by the pour-plate method with duplicate plating on selective agar media (Parrow, 1978). The coliforms were estimated in duplicate pour plates of Violet Red Bile Agar (VRBA, Oxoid CM107) medium and the plates were overlaid after solidification with 3 to 4 ml of additional Violet Red Bile Agar. The plates were incubated in an inverted position at 30°C ± 2 for 18 – 24 h (Mehlman, 1984). The yeasts and molds were counted on acidified Potato Dextrose Agar, (Oxoid CM139) which was acidified by the addition of the proper amount of sterile 10% tartaric acid (Fluka-Ag-Buchs.SG), then the plates were incubated at 25°C ± 1 for 3 - 7 days (Koburger and Marth, 1984). The laetic acid bacteria were enumerated in pour plates of de Man, Rogosa and Sharp (MRS) medium (Oxoid CM359). The plates were incubated at 37°C for 48 h under microaerobic conditions using Gas Pak (H2+CO2) (BBL, Microbiology Systems, Div. Becton Dickinson and Co., Cockeysville. Med.) anaerobic systems (Gilliland et al., 1984). The detection of Staphylococcus aureus, was applied according to the methods described in the FDA (1998).

Experimental diets:
The composition of experimental diets used in this work for feeding diarrhea rats is shown in table (1):
Table (1): The experimental rat groups and their diets

<table>
<thead>
<tr>
<th>Items</th>
<th>Basel diet</th>
<th>25%</th>
<th>50%</th>
<th>75%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1 (control)</td>
<td>100</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Group 2</td>
<td>75</td>
<td>25</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Group 3</td>
<td>50</td>
<td>-</td>
<td>50</td>
<td>-</td>
</tr>
<tr>
<td>Group 4</td>
<td>25</td>
<td>-</td>
<td>-</td>
<td>75</td>
</tr>
</tbody>
</table>

Biological experiments

Twenty four rats were fed the control diet for 7 consecutive days, rats were infected by rotavirus to induce diarrhea. After that, the rats were divided into four groups; each of it consists of six rats. The first group was the control one that fed on Casein-based diet. The rats of other groups (groups 2, 3, and 4) were fed on fermented camel milk. The total feeding period was 28 days. The rats were weighted after the 7 and 14 days at the beginning of the experiment and once a week. The gain weight was calculated by differences between the final body weight (gm) and the initial body weight (gm). The protein efficiency ratio (PER) was calculated according to A.O.A.C (2000) as follow:

\[
\text{PER} = \frac{\text{Gain in body weight (gm)}}{\text{Protein consumed (gm)}}
\]

The corrected PER was calculated according to Jansen et al., 1978 as follow:

\[
\text{Correct PER} = \text{PER} \times 2.5
\]

Blood Sampling:

In all previously mentioned animal groups, blood samples were collected after 12 hours fasting at the start of the experiment and after 2 and 4 weeks, from orbital venous by means of micro capillary glass tubes. The blood of the six rats of each group were placed in a dry clean centrifuge tubes and left to clot in a water bath (37ºC) for half an hour . The blood was centrifuged for 10 minutes at 4000 r.p.m to separate the serum. Serum was carefully aspirated and transferred into clean curvet quartz tubes and kept frozen at -20ºC till analysis.

Statistical analysis

Statistical analysis was performed using the SPSS software package for Windows [SPSS (UK) Ltd., Surrey, United Kingdom]. ANOVA was used to determine the difference between the means of the groups. Further analysis was carried out using a t-test for comparing two variables. P value considered significant when it was < 0.05

RESULTS AND DISCUSSION

Camel milk used in this study has Total Solids (T.S), Fat, Total Nitrogen (T.N) and Titratable Acidity (T.A), with contents of 11.92%, 3.1%, 0.49% and 0.18% respectively. This data is similar with Badran.,2004, While fermented camel milk was analysis T.S, Fat, P.N, N.P.N and acidity were 12.3%, 1.5%, 0.432%, 0.054 and 0.88% respectively. In this study we used separation fat partly from camel milk to (1.5%) before manufacture of fermented camel milk. we prepared quantity different formulas from it (25%, 50% and 75%) as shown as in table (1). Three prepared formulas from fermented camel milk used for therapeutics diarrhea.

The macro elements (sodium, potassium, phosphorous and calcium) as well as micro elements (iron and zinc) of the three prepared formulas for diarrhea rats were determined and the results were presented in table (2):

Table (2): Minerals content of formulas for diarrhea rats (mg/100gm)

<table>
<thead>
<tr>
<th>Formulas</th>
<th>macro elements</th>
<th>micro elements</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Na</td>
<td>K</td>
</tr>
<tr>
<td>Formula 1</td>
<td>70</td>
<td>180</td>
</tr>
<tr>
<td>Formula 2</td>
<td>75</td>
<td>185</td>
</tr>
<tr>
<td>Formula 3</td>
<td>80</td>
<td>195.5</td>
</tr>
</tbody>
</table>

Minerals content noticed in fermented camel milk were distinguished by the higher content of the sodium and potassium with the values 70.0 and 180.0 mg/100g than the row camel milk which recorded 52.0 and 145.0 mg/100g for sodium and potassium, respectively. These obtained results were agree with these obtained by (Watt and Merrill, 1963), (Abu-Lehia, 1989), (Aderson, 1991), (El-Agamy et al., 1998) and (Shamsia, 2009).

The data indicated that formula (3) characterized by the highest levels of Na, K, P and Ca, they were 80.0 and 195.5, 120, and 145 mg/100g, respectively. While formula 1 was the least in its content of these elements, with values of 70.0, 180.0, 113, and 135 mg/100g.
respectively. Also, the values of microelements (Iron and Zinc) in formula 3 were 0.15 and 1.2 mg/100gm respectively compared to formula 1, 2. In general, it was clearly noticed that most formulas have nearly the same content of the studied minerals.

Sodium deficiency was seen when severe diarrhea or vomiting occur. Partially, all the body sodium was found in the extra-cellular fluid which bathes the tissues. It does not cross the cell membrane to any great extent. Sodium functions are important in the regulation of acid-base equilibrium maintenance of osmotic pressure and of water balance. Potassium deficiency occur when there was prolonged failure treat and in pathologic conditions such as severe diarrhea (Robinson and Lowler, 1989).

The results of protein efficiency (PER), protein, sodium and potassium contents in serum of diarrhea rats are showed in table (3) and fig (1):

Table (3): Means of protein efficiency (PER), value of protein, sodium and potassium contents in serum of diarrhea rats fed on control and experimental diets

<table>
<thead>
<tr>
<th>Groups</th>
<th>Serum parameters after periods in days</th>
<th>Protein (mg/l)</th>
<th>Sodium (m mol/l)</th>
<th>Potassium (m mol/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>PER</td>
<td>0   7  21 21</td>
<td>0    7  21 21</td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td>2.8</td>
<td>2.9 3.0 6.5 6.9</td>
<td>7.2 150.0 151.9 151.5</td>
</tr>
<tr>
<td>Group 1</td>
<td></td>
<td>2.4</td>
<td>2.5 2.5 6.8 7.0</td>
<td>7.2 150.2 151.3 151.2</td>
</tr>
<tr>
<td>Group 2</td>
<td></td>
<td>2.6</td>
<td>2.7 2.7 6.9 7.1</td>
<td>7.2 150.7 151.5 151.4</td>
</tr>
<tr>
<td>Group 3</td>
<td></td>
<td>2.8</td>
<td>2.9 3.0 7.0 7.2</td>
<td>7.4 151.0 151.8 151.6</td>
</tr>
<tr>
<td>L.S.D</td>
<td></td>
<td>0.2</td>
<td>0.3 0.1 0.5 0.8</td>
<td>0.9 0.6 0.7 0.8</td>
</tr>
</tbody>
</table>

L.S.D: (Less Significance Difference at = 0.05)

From table (3), we noticed that all prepared diets were of good PER with ranged between (2.4 to 3.0). The highest PER was detected for rats fed on the diet prepared from formula 3; it was the same with rats fed on the control casein diet. Also, this table shows the means of total protein, sodium and potassium contents in the serum of diarrhea rats. These parameters were significantly equal for all rats at all the detected periods. Serum protein levels of rats fed control and tested diets were higher than the normal range this is in agreement with (El-Agamy et al., 1998) who found higher fat, protein (especially casein) and ash contents but lower whey protein and lactose contents in camel milk, Also with (Shamsia., 2009) who revealed that camel milk proteins contained satisfactory balance of essential amino acids. So, camel milk can be considered as good source of protein and can meet part of the daily needs of humans from these nutrients.

Fermented milk products are known for their taste, nutritive value and therapeutic properties. According to the International Dairy Federation (1969), fermented milks are “products prepared from milk” whole or fully skimmed, concentrated or milk substituted from partially or full skimmed milk, either homogenized or un-homogenized, pasteurized or sterilized and fermented by mean of specific microorganisms. Milk from camel has been used to make traditional fermented milk products throughout the world. The people who have domesticated these milk animal usual accepted fermented milk by necessity (Kroger et al., 1989), and (Abdel Moneim et al., 2009).

As shown in table (4), the infectious of rats produced by a Rotavirus to induce diarrhea caused a great loss in the initial body weight of all rats. When the diarrhea rats were fed on formula 3, diarrhea was stopped at the second day compared to formula 1, 2 and control casein diet. Formula (2) was followed by formula 3 in stopping diarrhea after three days which contained 50% fermented camel milk, 50% control diet, while formula (1) was stopping diarrhea after four days, its similar with the control casein diet which contained 25% fermented camel milk. These observations could be considered that formula (3) was the adequate one for stopping diarrhea compared to other prepared formula and the control, followed by formula (2) and formula (1) as shown in fig (2).
Table (4): Means of body weigh (B.W), gain weight (G.W) and diarrhea stopping day of diarrhea rats fed on control and experimental diets

<table>
<thead>
<tr>
<th>Groups</th>
<th>Initial body weigh (g)</th>
<th>B.W and G.W (g) after periods in days</th>
<th>Diarrhea stopping day</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>4 days</td>
<td>7 days</td>
</tr>
<tr>
<td></td>
<td></td>
<td>B.W. W. loss</td>
<td>B.W. G.W.</td>
</tr>
<tr>
<td>Group 1 (control)</td>
<td>90.2</td>
<td>80.2</td>
<td>10.0</td>
</tr>
<tr>
<td>Group 2</td>
<td>85.0</td>
<td>70.0</td>
<td>15.0</td>
</tr>
<tr>
<td>Group 3</td>
<td>88.2</td>
<td>75.0</td>
<td>13.2</td>
</tr>
<tr>
<td>Group 4</td>
<td>90.1</td>
<td>80.0</td>
<td>10.1</td>
</tr>
<tr>
<td>L.S.D</td>
<td>0.9</td>
<td>0.8</td>
<td>0.4</td>
</tr>
</tbody>
</table>

Results were agreement with previously reported (Abu-Lehia, 1987, Aderson, 1991 and El-Agamy et al., 1998).

The results of this work showed that final products of fermented camel milk prepared in the lab by using cultures had no Staphylococcus aureus in formula (3).

The stopping diarrhea at the second day on feeding formulas (3) may be attributed to its higher content of fermented camel milk (75%), besides this formula contained the highest level of protein, Na and K. The stopping diarrhea may be due to the interaction effect between all these factors in formula (3) causing diarrhea.

The stopping diarrhea at the second day on feeding formulas (3) may be attributed to its higher content of fermented camel milk (75%), besides this formula contained the highest level of protein, Na and K. The stopping diarrhea may be due to the interaction effect between all these factors in formula (3) causing diarrhea.

The concentrations of macro elements (Na, K, P, and Ca) were higher in formula 3 than in formula 1 and 2. Therefore, it considered that the formula 3 as a good source of these minerals. Also, formula 3 contained higher values of microelements (Fe and Zn).

Besides, the fermented camel milk can stopping diarrhea because it contained specific antibodies to Rotavirus which cause diarrhea and in addition, the fermented camel had antivirus activity against human Rotavirus, so, it used as a remedy for virus (Shamsia, 2009).

It can be concluded that fermented camel milk can be considered as a good food of high nutritive and therapeutic applications. Meanwhile, the high content of antimicrobial agents in camel milk may explain its potential as an activity especially against diarrhea.

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