EFFECT OF CAMEL MILK ON GLYCEMIC CONTROL, RISK FACTORS AND DIABETES QUALITY OF LIFE IN TYPE-1 DIABETES: A RANDOMISED PROSPECTIVE CONTROLLED STUDY

R.P. Agrawal*, S.C. Swami, R. Beniwal, D.K. Kochar, M.S. Sahani, F.C. Tuteja and S.K. Ghouri

S.P. Medical College and National Research Centre on Camel, Bikaner-334001, INDIA * S.P. Medical College, Bikaner-334 001, INDIA

ABSTRACT

The efficacy of camel milk on glycemic control risk factors and diabetes quality of life in patients of type 1 diabetes was evaluated. Twenty four randomly selected patients with type 1 diabetes were enrolled in the study. These patients were divided into two groups. Group 1 (N=12) received usual care (diet, exercise and insulin) and group 2 (N=12) received 500 ml camel milk in addition to usual care for 3 months. Frequent blood sugar monitoring was done to maintain euglycemia by titrating the doses of insulin. HbA $_1$ c, Lipid profile, plasma insulin and c-peptide estimation was done at the beginning and after 3 months. BMI, diabetes quality of life questionnaire were prepared every week. In each visit patient was asked for any untoward effects after starting camel milk.

Baseline data of both the groups were similar in demographic and clinical variables. After 3 months of treatment there was a significant improvement in fasting blood sugar (115.66 \pm 7.17 to 100 \pm 16.2, p<0.002) and HbA₁c levels (9.54 \pm 2.1 to 9.08 \pm 1.77, p<0.002) and significant reduction in insulin requirement (mean doses of insulin 41.16 \pm 10.32 to 30 \pm 12.6, p<0.002) in patients receiving camel milk. Diabetes quality of life score improved significantly in the form of change in satisfaction score from 28 \pm 5.16 to 22.5 \pm 3.96 (p<0.002), impact score from 34 \pm 4.84 to 28.08 \pm 5.26 (p<0.003) and worry score from 15.5 \pm 3.2 to 11.91 \pm 1.24 (p<0.002). There was 30% reduction in doses of insulin in 92% of patients of group 2. However, there was no statistically significant changes in lipid profile, plasma insulin and c-peptide.

Camel milk proved effective supplementation in the management of type 1 diabetes as there was significant reduction in doses of insulin along with betterment in BMI, diabetes quality of life however, there was no change in lipid profile and insulin levels.

Type I diabetes mellitus is an organ specific auto immune disease, characterised by chronic hyperglycemia and disturbances of carbohydrates, fat and protein metabolism associated with insulin deficiency. Cow milk feeding induces primary immunisation to insulin in infants at genetic risk for type 1 diabetes (Vaarala *et al*, 1999). The incidences of diabetes mellitus world wide appear to be increasing (Onkamo *et al*, 1999). Prevention and early treatment is important because diabetes interrupts normal developments in children and carries the threat of severe complication in more active period of life (Dahlquist, 1999). Its primary treatment is insulin replacement, however, at

present, entire physiological insulin replacement can not be achieved in clinical practice and metabolic disturbances can not be normalised. Insulin therapy is still the best treatment but in our country needle phobia and cost of the treatment forces these patients to adopt alternative treatments. In this connection we have heard many folklore stories which describe the use of camel milk in type-1 diabetes mellitus. There is also an account in memories of Emperor Jahangir (1579 - 1627 AD) about usefulness and acceptability of camel milk (Rogers, 1989). It is found that one of the camel milk protein has many characteristics similar to insulin (Beg et

SEND REPRINT REQUEST TO R.P. AGRAWAL

al, 1986b) and it does not form coagulum in acidic enviornment (Wangoh, 1993). This lack of coagulum formation allows the camel milk to pass rapidly through the stomach together with the specific insulin like protein/insulin and remains available for absorption in intestine. Radioimmunoassay of camel milk has revealed high concentration of insulin i.e. 52 units/litre (Singh, 2001). The concentration of insulin in human milk is also significantly higher (60.23 \pm 41.05 micro u/ml) whereas it is very low in cow milk (16.32 \pm 5.98 micro u/ml) (Shehadeh *et al*, 2001) but probably because of coagulation in stomach it is not available for absorption in the intestine.

Materials and Methods

Subjects: A total of 24 type 1 diabetic patients were randomly recruited from the outpatient diabetic clinic in PBM Hospital, Bikaner, India. Ethical committee of S.P. Medical College, Bikaner approved the protocol and all subjects gave written consent before participation in the study. The patients were advised to follow strict diet, exercise and insulin treatment for 1 month. During this period frequent monitoring of blood sugar was done to maintain euglycemia. After onemonth period these patients were again randomly divided into two groups. Group 1 patients (N=12) received usual care i.e. diet, exercise and insulin and the Group 2 patients (N=12) received 500 ml of camel milk in addition to usual care for 3 months. Patients with any acute metabolic complications like hypoglycemia, ketoacidosis, cardiovascular event, renal or acute infections were not included in the study.

Study Design and Analysis

This was a randomised, open case control, parallel design study. Blood sugar was measured twice in a week before breakfast and before dinner and blood sugar concentration was measured using the glucose oxidase method. Plasma insulin and C-peptide were estimated by fully automated chemi-illuminescence (CLIA test). Anti-insulin antibodies were estimated by radioimmunoassay. HbA1c was measured by high performance liquid chromatography (HPLC). Plasma total cholesterol, triglycerides, VLDL, HDL and LDL were estimated by fully automated biochemistry analyser. Urine microalbumin was tested by

micral test. Body mass index, waist hip ratio, and 'diabetes quality of life' score were also measured every week (Surwit *et al*, 1992; TDCCTRG, 1996).

Statistical Analysis

As the normality of the variables in the study could not be assured, Wilcoxan matched pair test and Mann-Whitney U test were used instead of t tests. The two groups had equal number of participants and they were compared with each other using Mann-Whitney U test. The value of p<0.05 was considered significant for mann-whitney U-test after Satterthwaite correction. The variables were compared at three months to that at the start of the study using Wilcoxan matched pair test with cut off value being decided at p<0.05.

Results

Demographic characteristics are summarised in table 1. The group 1 (control group) and group 2 (camel milk group) were similar in age (20.33 \pm 4.97 Vs 19.5 \pm 8.15), sex (10M, 2F in both groups), body mass index (18.43 \pm 3.59 Vs 20.21 \pm 2.97), fasting blood sugar (117.16 \pm 17.73 Vs 115.66 \pm 7.17), plasma insulin (16.37 \pm 7.57 Vs 16.39 \pm 6.57), c-peptide (1.24 \pm 0.60 Vs 1.26 \pm 0.61) plasma lipids alongwith different clinical, demographical and biochemical variables (table 1).

After three months of treatment there was statistically significant increase in body mass index (20.21 \pm 2.97 to 21.3 \pm 2.95, p <0.05), and improvement in fasting blood sugar (115.66 ± 7.17 to 100 \pm 16.2, p<0.002) and in HbA1c (9.54 \pm 2.1 to 9.08 \pm 1.77, p<0.002), in the camel milk group. These parameters were either unchanged or there was a slight increase in group 1 patients (table 2). Fasting plasma insulin and C-peptide levels did not reveal a significant change in either group and so were the levels of lipid profile, after 3 months of treatment. The diabetes quality of life questionnaire score changed significantly in favour of camel milk (i.e. satisfaction score 26.08 ± 4.11 to 22.5 ± 3.96 , p < 0.05, impact score 32.5 ± 2.71 to 28.08 ± 5.26 , p < 0.05 and worry score 14.66 ± 1.15 to 11.9 ± 1.24 , p<0.05). There was a significant reduction in the mean doses of insulin $(41.16 \pm 10.32 \text{ to } 30 \pm 12.6, p<0.002)$ in patients receiving camel milk (Table 3, Fig 1). The acceptability of camel milk was very good and only 1 patient complained of mild flatulence

Table 1. Base line characteristics of study groups.

Variables	Group I (n=12)		Group II (n=12)			n n
	Mean	SD	Mean	SD	t	P
Age (Yrs)	20.33	4.97	19.5	8.15	-1.131	0.257
W/H Ratio	0.75	0.08	0.81	0.05	-1.642	0.109
BMI (Kg/m ²)	18.43	3.59	20.21	2.97	-0.346	0.729
HbA1c (%)	9.51	2.089	9.54	2.10	-1.472	0.140
Dose of Insulin (units/day)	40	8.61	41.16	10.32	-0.028	0.976
Mean Blood Sugar (mg/dl)	117.16	17.73	115.66	7.17	-0.812	0.416
T. Cholesterol (mg/dl)	165.83	19.19	164.58	20.69	0	1
HDL (mg/dl)	61.58	9.1	62.58	13.91	-1.944	0.051
LDL (mg/dl)	89.58	14.7	92	11.62	-1.097	0.272
VLDL (mg/dl)	14.41	4.67	13.5	5	-0.433	0.664
T.G. (mg/dl)	72.39	20.71	66.91	25.6	-0.636	0.524
Micro Albuminuria (mg/dl)	22.54	5.62	22.13	5.10	-0.288	0.772
Plasma Insulin (μIU/ml)	16.37	7.57	16.79	6.57	-0.346	0.729
C. Peptide (ng/ml)	1.24	0.60	1.26	0.61	-0.375	0.707
DQOL Score:						
Satisfaction	26.16	2.58	28	5.16	-1.687	0.091
Impact	29.58	2.60	34	4.84	-1.285	0.198
Worry	13.0	0.05	15.5	3.20	-1.508	0.131

(Values = Mean \pm S.D.) (*p=<0.05);

W/H = Waist/Hip; BMI = Body Mass Index; HbA1c = Glycosylated haemoglobin; HDL = High Density Lipoprotein; LDL = Low Density Lipoprotein; VLDL = Very Low Density Lipoprotein; T.G. = Tri Glyceride; DQOL = Diabetes Quality of Life.

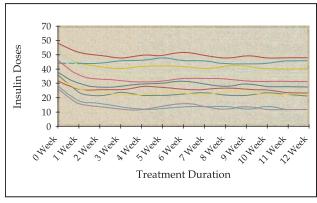


Fig 1. Doses of insulin in individual patient of group II (N-12).

for 3-4 days. Mild diarrohea (2-3 semi-solid) was reported by two patients which also subsided spontaneously.

Discussion

The present study was performed to observe the role of camel milk in achieving glycemic control in type-1 diabetic patients. We observed a significant improvement in mean BMI (20.21 ± 2.97 to 21.3 ± 2.95 , p<0.002) after 3 months of camel milk treatment. The positive effects in weight gain may be because of good nutritional value of camel milk. There was no change in lipid profile and it may be due to lower fat content of camel milk. (i.e. 2.49-3.1 gm% Vs cow milk 3.79 gm%).

We also observed significant reduction in insulin doses to obtain glycemic control along with significant improvement in HbA1c level at the end of three months. No other studies are available for comparison. Improvement in microalbuminuria may be due to good glycemic control or it may be due to direct effect of camel milk. There was marked improvement in diabetes quality of life score after 3 month of camel milk treatment. It may be because of good glycemic control or anabolic effect of camel milk. El-Agamy *et al* (1992) found good amount of lysozyme, lactoferrin, lactoperoxidase, immunoglobulin G and secretory immunoglobulin A in camel milk.

Table 2. Group I vs group II at 3 months.

Variables	Group I n=12		Group II n=12		Mann -Whitney U test	
	Mean	SD	Mean	SD	Z adjusted	p value
Age (Yrs)	20.33	4.097	19.5	8.15	-1.131	0.257
W/H Ratio	0.75	0.08	0.81	0.05	-1.379	0.164
BMI (Kg/m2)	18.41	3.51	21.3	2.95	-1.328	0.184
HbA1c (%)	9.48	1.96	9.08	1.77	-1.905	0.056
Dose of Insulin (units/day)	38.5	8.49	30	12.06	-2.139	0.032*
Mean Blood Sugar (mg/dl)	118.16	7.15	100	16.2	-3.935	8.32E-05*
Total Cholesterol (mg/dl)	168.08	15.61	158.33	21.55	-0.433	0.664
HDL (mg/dl)	58.66	15.61	66.66	11.29	-0.115	0.907
LDL (mg/dl)	89.66	12.26	79.16	17.75	-0.981	0.326
VLDL (mg/dl)	14.25	3.16	12.08	5.08	-1.041	0.297
T.G. (mg/dl)	72.0	14.79	60.16	25.16	-0.520	0.603
Micro Albuminuria (mg/dl)	22.9	5.43	25.17	5.43	-0.230	0.817
Plasma Insulin (IU/ml)	16.31	7.5	16.94	6.54	-0.173	0.862
C.Peptide (ng/ml)	2.28	0.63	2.22	0.5	-0.723	0.469
DQOL Score						
Satisfaction	22.75	2.37	22.5	3.96	-3.034	0.002*
Impact	29.5	2.93	28.08	5.26	-2.175	0.029*
Worry	12.58	1.16	11.91	1.24	-4.073	4.65E-05*

(Values = Mean \pm S.D.) (*p=<0.05)

Requirement of mean doses of insulin/day before treatment in patients of group-2 was 41.16 ± 10.32. It came down very fast initially and then gradually to a mean level of 30±12.06, (p<0.05). Only one patient out of 12 patients required the same doses of insulin and the other 11 patients had lower requirement to maintain euglycemic blood level. Camel milk was found to contain about 52 units/litre insulin (Singh, 2001) and it may be the reason for lesser requirement of insulin in camel milk group. Oral insulin has been known since many years but the important drawback is its coagulum formation in acidic media in stomach thereby neutralising its potency. The lack of coagulum formation of camel milk may act as an effective vehicle to take the insulin present in it unchanged to the intestine and from there it can be absorbed even if some amount is destroyed in the passage. Beg et al (1986a) has found that amino acid sequence of some of the camel milk protein, is rich in half cystine, which

has superficial similarity with insulin family of peptides.

The data of this study shows a significant hypoglycemic effect of camel milk when given as an adjunct therapy. The action is presumed to be due to presence of insulin/insulin like protein in it. Its therapeutic efficacy may be due to lack of coagulum formation of camel milk in acidic media. It has been observed that oral administration of insulin initiated at clinical onset of type 1 diabetes did not prevent the deterioration of beta cell function (Chaillous et al, 2000). Pozzilli et al (2000) in IMDIAB VII study indicates that addition of 5mg of oral insulin does not modify the course of the disease in the first year after diagnosis and probably does not statistically effect the humoral immune response against insulin (Pozzilli et al, 2000). It is important to note that a certain level of scientific testing on camel milk has already been attempted and documented, particularly, insulin levels in

Table 3. Group II before and after treatment.

Variables	0 Month		3 Months		Wilcoxon matched pairs test	
	Mean	SD	Mean	SD	Z	P value
Age (Yrs.)	19.5	8.15	19.8	8.15		
W/H Ratio	0.81	0.05	0.81	0.05	1.01	0.312
BMI (Kg/m2)	20.81	2.97	21.3	2.95	3.06	0.002*
HbA1c (%)	9.54	2.1	9.08	1.77	3.06	0.002*
Dose of Insulin (units/day)	41.16	10.32	30	12.06	3.06	0.002*
Mean Blood Sugar (mg/dl)	115.66	7.17	100	16.2	3.06	0.002*
T. Cholesterol (mg/dl)	164.58	20.59	158.33	21.55	1.29	0.195
HDL (mg/dl)	62.58	13.91	66.66	11.29	0.86	0.388
LDL (mg/dl)	92	11.62	79.16	17.75	2.04	0.040*
VLDL (mg/dl)	13.5	5	12.08	5.08	1.42	0.155
T.G. (mg/dl)	66.91	25.6	60.16	25.16	1.02	0.306
Micro Albuminuria (mg/dl)	22.13	5.1	25.17	5.43	0	1
Plasma Insulin (μIU/ml)	16.79	6.57	16.94	6.54	1.17	0.239
C.Peptide (ng/ml)	2.26	0.61	2.22	0.5	1.45	0.146
DQOL Score		7	•	<u> </u>	•	•
Satisfaction	28	5.16	22.5	3.96	3.06	0.002*
Impact	34	4.84	28.08	5.26	2.93	0.003*
Worry	15.5	3.2	11.91	1.24	3.05	0.002*

(Values = Mean \pm S.D.) (*p=<0.05)

camel milk and this scientific wisdom can be a remarkable achievement for diabetic patients.

Acknowledgement

We are thankful to Dr. M.S. Sahani, Director, National Research Centre on Camel, Bikaner for providing camel milk for our study. We are also thankful to Dr. Raghvendra Singh, Senior Scientist, National Research Centre on Camel, Bikaner for chemical analysis of camel milk.

References

Beg OU, Bahr-Lind Ström H Von, Zaidi ZH and Jornvall H (1986a). A camel milk protein rich in half cystine. Primary structure assessment of variations, internal repeat patterns and relationship with neurophysin and other active polypeptides. European Journal of Biochemistry 15(1):195-201.

Beg OU, Bahr-Lind Ström H Von, Zaidid ZH and Jörnvall H (1986b). Characteristics of camel milk protein rich in proline identifies a new beta casein fragment. Regulatory Peptides 15(1):55-61.

Chaillous L, Lefevre H, Thivolet C, Boitard C, Lahlou N, Atlan-Gepner C, Bouhanick B, Mogenet A, Nicolino M, Carel JC, Lecomte P, Marechaud R, Bougneres P, Charbonnel B and Sai P (2000). Oral insulin administration and residual beta-cell function in recent-onset type 1 diabetes: a multicentre randomised controlled trial. Diabetes Insuline Orale group. Lancet 356:545-549.

Dahlquist GG (1999). Primary and secondary prevention strategies of pre-type 1 diabetes: potentials and pitfall. Diabetes Care 22 (Suppl. 2):B4-B6.

El-Agamy EI, Ruppanner R and Ismail A (1992). Antibacterial and antiviral activity of camel milk protective proteins. Journal of Dairy Research 59(2):169-175.

Onkamo P, Vaananen S, Karvonen M and Tuomilehto J (1999). World wide increase in incidence of type 1 diabetes: analysis of the data on published incidence trends. Diabetologia 42:1395-1403.

Pozzilli P, Pitocco D and Visalli N (2000). No effect of oral insulin on residual beta-cell function in recent-onset Type 1 diabetes (the IMDIAB VII). Diabetologia 43:1000-1004.

Rogers A (1989). Memories of Jahangir. Allantic Publishers and Distributor, New Delhi. p 315.

- Shehadeh N, Gelertner L, Blazer S, Perlman R, Solovachik L and Etzioni A (2001). Importance of insulin content in infant diet: suggestion for a new infant formula. Acta Paediatr. 90(1):93-95.
- Singh R (2001). Senior Scientist. National Research Centre on Camel, Bikaner, India. Personal Communication.
- Surwit RS, Schneider MS and Feinglos MN (1992). Stress and diabetes mellitus. Diabetes care 15:1413-1422.
- The Diabetes Control and Complications Trial Research Group (1996). Influence of intensive diabetes treatment on

- quality of life outcomes in the diabetes control and complications trial. Diabetes Care 19(3): 195-202.
- Vaarala O, Knip M, Paronen J, Hamalainen AM, Muona P, Vaatainen M, Ilonen J, Simell O and Akerblom HK (1999). Cow's milk formula feeding induces primary immunisation to insulin in infants at genetic risk for type 1 diabetes. Diabetes 48(7):1389-1394.
- Wangoh J (1993). What steps towards camel milk technology? International Journal of Animal Science 8:9-11

CORRIGENDUM

The legend of figure number 3 should be read for figure 4 and vice-versa in the manuscript entitled, "Mastitis caused by *Pasteurella hemolytica* and *Streptococcus agalactiae*" in December, 2002 issue of JCPR on page no.122.

Camel Publishing House apologises the error.