A modified cows' milk formula suitable for low birthweight infants

T J FRENCH, M COLBECK, D BURMAN, B D SPEIDEL, AND R A HENDEY

Department of Paediatrics, Bristol Maternity Hospital; Cow and Gate, Trowbridge, Wilts

SUMMARY Low birthweight babies fed standard modified cows' milk formulae are at risk from the high fluid intake needed for adequate nutrition, and very preterm babies often develop late hyponatraemia if the sodium intake fails to match large renal losses. A new cows' milk formula (Cow and Gate Prematalac) provides 120 kcal, 3.6 g protein, 7.5 g fat, and 4 mmol sodium in 150 ml. Ten low birthweight babies were fed the new formula at 150 ml/kg a day and compared with 12 similar babies fed a standard modified cows' milk formula (Wyeth SMA Gold Cap) at 180 ml/kg a day. All the babies grew at intrauterine rates and there was no difference in clinical course. None fed the new formula developed hypernatraemia, oedema, or dehydration and none fed the standard formula developed hyponatraemia. The Prematalac group safely excreted the increased osmotic load and had a higher urinary sodium concentration which should protect less mature preterm infants from late hyponatraemia.

In the UK most preterm infants are fed their mother's milk, banked breast milk, or milk formulae. These formulae contain 67 kcal/100 ml and have a low sodium concentration to suit the needs of term infants.¹ In the 3rd trimester the fetus gains weight at about twice the rate of a term infant in early postnatal life, and major changes take place in body composition as organs develop and stores of energy and minerals accumulate.² It is debatable whether the extrauterine growth rate of the preterm infant needs to match the intrauterine growth rate but such comparison is widely considered a yardstick of the growth and development that are taking place in the central nervous system.

Intrauterine growth rates can be achieved in preterm infants fed breast milk or milk formulae if an amount of 110-150 kcal/kg a day is given,³ but high fluid intakes are required which may lead to a poor tolerance of feed, or to an increased incidence of patent ductus arteriosus⁴ and necrotising enterocolitis.⁵ In addition late hyponatraemia may occur in the very low birthweight baby in whom renal handling of sodium is less developed than in the more mature infant.^{6,7} A sodium intake of 3 mmol/kg a day prevents hyponatraemia in very low birthweight babies who are well⁶ but sick preterm infants have greater sodium losses and require higher intakes.⁸

We have devised a cows' milk formula with a higher energy, protein, and sodium content than in

standard formulae and have fed it at 150 ml/kg a day to low birthweight infants. Growth rates and clinical and metabolic data were compared with those of a similar group fed a standard milk formula at 180 ml/kg a day.

Patients and methods

Newborn babies weighing less than 2500 g entered the trial provided they were not multiple births, large for gestational age, had major congenital abnormalities, or were babies of mothers wishing to breast feed. At birth each baby was placed in one of two groups so that the distribution of birthweights and the proportion of small for gestational age babies in each group were similar. A baby was regarded to be large or small for gestational age if the birthweight was above the 90th centile or less than the 10th centile for gestational age on Gairdner and Pearson growth and development charts.⁹ One group was allocated the new milk (Cow and Gate Prematalac) at a volume of 150 ml/kg a day; the other was given Wyeth SMA Gold Cap at 180 ml/kg a day. The composition of the milks and intakes on full feeding are shown in Table 1. The osmolalities of Cow and Gate Prematalac and Wyeth SMA Gold Cap are 342 and 300 mmol/kg respectively. Feeding began when clinically indicated and infants entered the trial after full oral feeding had been established. None received phototherapy,

Table 1 Composition of milks and intakes of full feeds

	Cow and Gate Prematalac		SMA Gold Cap	
	Per 100 ml	Per 150 ml/kg a day	Per 100 ml	Per 180 ml/kg a day
Energy (kJ)	335	502	275	495
(kcal)	79	119	65	117
Protein (g)	2.4	3.6	1.5	2.7
Fat (g)	5.0	7.5	3.6	6.5
Lactose (g)	6.6	9.9	7.2	13
Sodium (mmol)	2.60	3.90	0.65	1.17
Potassium (mmol)	2.43	3.65	1.43	2.57
Chloride (mmol)	2.25	3.38	1.04	1.87
Calcium (mmol)	1.67	2.51	1.1	1.98
Magnesium (mmol)	0.47	0.71	0.22	0.40
Phosphorus (mmol)	1.71	2.56	1.1	1.98

assisted ventilation, or xanthine medication during the study. Weight was recorded every other day, occipitofrontal circumference weekly, and length was measured at the beginning and end of the trial. The frequency and nature of stools were recorded. Plasma urea, sodium, potassium, and chloride concentrations together with haemoglobin and white cell counts were estimated twice weekly. Plasma calcium concentration was measured weekly as were urinary osmolality, sodium, and potassium concentration on a random urine sample.

Results

The clinical data for each group are shown in Table 2, and the growth, haematological, and biochemical data in Table 3. Weight gain (Table 3) is expressed as g/kg a day rather than as g/day as weight gain *in utero* in the last trimester is exponential¹⁰ and rate of gain of weight dependent on body weight. There was no significant difference between the proportion of small for gestational age infants in the groups (Fisher's exact test $P = 0.485$). As the data were not normally distributed, medians and ranges are quoted and a nonparametric statistical method—namely the Mann-Whitney U test—was used to determine the significance of differences between the two groups in the trial.

Rates of gain of weight, length, and occipitofrontal circumference of both groups were similar and equalled intrauterine growth.^{11 12} No infant in either group developed diarrhoea, steatorrhoea, oedema, dehydration, patent ductus arteriosus, or necrotising enterocolitis. None became anaemic or leucopenic.

Plasma sodium, potassium, urea, and calcium concentrations were not significantly different and were within the normal range. The plasma chloride concentration of the Prematalac group was higher

Table 2 Clinical data

	Cow and Gate Prematalac group (n = 10)		SMA Gold Cap group (n = 12)	
Birthweight (g)	1780	(1230–2180)	1890	(1240–2220)
Gestation (weeks)	34.0	(27.7–37)	33.6	(30–35.9)
Age (weeks)	1.4	(0.9–6.1)	1.7	(1.0–6.1)
Maturity (weeks)	35.0	(31.6–38.8)	35.7	(33.3–37.7)
Weight at start of trial	1810	(1440–2140)	1885	(1340–2370)

Data expressed as median value and range.

Table 3 Growth, haematological plasma, and urinary biochemistry

	Cow and Gate Prematalac	SMA Gold Cap	P value
Growth			
Weight (g/kg a day)	15.2 (9.7–18.4)	13.6 (8.1–19.4)	NS
Length (cm/day)	0.16 (–0.21–0.53)	0.12 (0.00–0.34)	NS
Occipitofrontal circumference (cm/day)	0.14 (0.11–0.25)	0.18 (0.10–0.25)	NS
Haematology			
Haemoglobin (g/dl)	17.6 (8.9–20.7)	14.6 (9.0–19)	NS
White cell count	11.0 (8.2–15.2)	11.6 (9.3–15.2)	NS
Plasma biochemistry			
Urea (mmol/l)	2.49 (0.9–3.8)	1.52 (0.5–4.1)	NS
Sodium (mmol/l)	140 (134–144)	137 (133–139)	NS
Potassium (mmol/l)	4.7 (4.1–5.9)	4.8 (4.2–6.0)	NS
Chloride (mmol/l)	107 (102–108)	103 (97.5–106)	<0.01
Calcium (mmol/l)	2.44 (1.79–2.60)	2.49 (2.14–2.74)	NS
Urinary biochemistry			
Sodium (mmol/l)	28.0 (13–63)	3.9 (0.5–49)	<0.001
Potassium (mmol/l)	26.8 (8–39)	16.8 (4.5–39)	NS
Osmolality (mmol/kg)	192.0 (63–323)	92.5 (63–195)	<0.005

Data expressed as median value and range.

NS = not significant, $P > 0.05$.

Conversion: SI to traditional units—plasma urea: 1 mmol/l $\approx$ 6.02 mg/100 ml; plasma calcium: 1 mmol/l $\approx$ 4.0 mg/100 ml.

than the SMA group but both were within the normal range.

There was a marked difference in the composition of urine formed on the different milks (Table 3). Urinary sodium concentration of the majority of infants fed SMA Gold Cap was lower than 10 mmol/l and in many cases lower than 5 mmol/l; those fed Prematalac had a higher urinary sodium concentration, generally between 15 and 35 mmol/l. Most infants fed SMA Gold Cap had a urinary osmolality less than 120 mmol/kg while the majority of the Prematalac group had a value between 120 and 240 mmol/kg. In only one case did the urinary

osmolality of an infant fed Prematalac exceed 300 mmol/kg. The urinary potassium concentration was similar in each group.

Discussion

The new formula (Cow and Gate Prematalac) differs in three important ways from formulae designed for use in term infants. Firstly, the higher energy density (80 kcal/100 ml; 335 kJ/dl) provides sufficient energy for preterm infants to grow at intrauterine rates at a volume intake of 150 ml/kg a day.³ Lower volume intakes are associated with an improved tolerance of feed and a lower incidence of patent ductus arteriosus⁴ and necrotising enterocolitis.⁵ The main energy source in the new formula is a mixture of butter fat and vegetable oil in a higher concentration than in standard formulae (Table 1). Secondly, the protein content is also higher. An intake of 150 ml/kg a day provides 3.6 g protein with a casein: whey ratio of 40:60. The protein concentration and composition are based on the recommendations of Senterre.¹³ He showed that an oral intake of 3.5–4 g/kg a day with the above casein: whey ratio was optimal for growth of preterm infants provided sufficient energy was available.¹³ Protein intakes within this range produce normal plasma urea levels, give a plasma amino acid profile similar to that of cord blood, and keep the dietary solute and acid loads well within the renal capacity for excretion.¹³ Thirdly, the sodium concentration has been increased to avoid late hyponatraemia (plasma sodium <130 mmol/l) which occurs in one-third to two-thirds of very low birthweight babies (<1300 g) fed low sodium milks.^{6,7} This can be slowly corrected if the dietary sodium is increased to 3 mmol/kg a day.⁶ Late hyponatraemia occurs if sodium intake is below that required for growth (1.2 mmol/kg a day¹⁰) and for loss in the urine. Immature renal function in very low birthweight babies leads to a high urinary sodium loss even in the presence of hyponatraemia.⁷ Leake⁸ has estimated the urinary sodium loss in these infants to be 2.5 mmol/kg a day. Sick preterm infants and those on xanthine derivatives for apnoea have even higher urinary sodium losses.⁸ The new formula provides 4 mmol/kg a day sodium to meet the normal requirement of very low birthweight babies and to allow extra sodium for those who are unwell or being treated with xanthine derivatives.

The clinical course of infants fed the new formula at 150 ml/kg a day was identical with that of infants fed the standard formula at 180 ml/kg a day and no infant had any complication. Both groups gained weight, length, and head circumference at intrauterine rates. The main advantage of the new

formula was growth at these rates with an intake of 30 ml/kg a day less than required with the standard formula. This has several advantages when milk is being introduced gradually to the preterm infant; firstly, a full energy intake is achieved earlier because of the lower volume intake required; secondly, infants who can tolerate only low volume feeding because of intercurrent illness will be better nourished.

Infants fed the new formula and the standard formula had plasma sodium, potassium, chloride, calcium, and urea levels within the normal range. In particular none fed the new formula developed hypernatraemia and none fed the standard formula developed hyponatraemia (Table 3). The latter result is not unexpected since the infants fed standard milk formula were more mature than those in whom late hyponatraemia has been reported^{6,7} (Table 2). Sodium content of the standard milk formula will match the 1.2 mmol/kg a day required for growth¹⁰ provided that it is fed at 180 ml/kg a day (Table 1) and that the infant's renal tubular function is sufficiently mature maximally to reabsorb sodium. The very low sodium concentration in the urine of this group (Table 3) shows that this degree of renal sodium reabsorption has been achieved. The higher urinary sodium found in the Prematalac group means that the new formula provides a margin of safety to prevent late hyponatraemia in very low birthweight babies.

Urinary osmolality of the group fed the new formula was higher than that fed the standard milk formula, no doubt reflecting the higher sodium and protein loads and the lower fluid intake of the new formula group. The range of osmolality found in infants fed the new formula was within the concentrating ability of the very preterm infant⁸ and indicates that adequate fluid was provided to meet the obligatory renal solute loss.

Conclusion

The new formula fed at 150 ml/kg a day leads to growth at intrauterine rates. The increased sodium concentration is sufficient for the needs of very preterm infants^{6,7} and although it is greater than the requirement of more mature infants the excess can safely be excreted in the urine. The new formula appears to be a suitable milk for all babies of low birthweight, but further information is needed for the infant of very low birthweight, particularly if he is sick.

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References

- ¹ Burman D. Nutrition in early childhood. In: McLaren D S, Burman D, eds. *Textbook of paediatric nutrition*. Edinburgh: Churchill Livingstone, 1976: 46–75.
- ² Jonxis J H P. Special problems arising in the nutrition of very small prematures and their subsequent growth. In: Wilkinson A W, ed. *Early nutrition and later development*. Tunbridge Wells: Pitman Medical, 1976: 79–91.
- ³ American Academy of Pediatrics Committee on Nutrition. Nutritional needs of low-birth weight infants. *Pediatrics* 1977; **60**: 519–30.
- ⁴ Bell E F, Warburton D, Stonestreet B S, Oh W. Randomized trial comparing high (HV) and low-volume (LV) maintenance fluid administration in low-birth-weight infants (abstract). *Pediatr Res* 1979; **13**: 489.
- ⁵ Bell E F, Warburton D, Stonestreet B S, Oh W. Letter: High-volume fluid intake predisposes premature infants to necrotising enterocolitis. *Lancet* 1979; **ii**: 90.
- ⁶ Roy R N, Chance G W, Radde I C, Hill D E, Willis D M, Sheepers J. Late hyponatremia in very low birth weight infants (<1.3 kilograms). *Pediatr Res* 1976; **10**: 526–31.
- Day G M, Radde I C, Balfe J W, Chance G W. Electrolyte abnormalities in very low birthweight infants. *Pediatr Res* 1976; **10**: 522–6.
- ⁸ Leake R D. Perinatal nephrobiology: a developmental perspective. *Clin Perinatol* 1977; **4**: 321–49.
- ⁹ Gairdner D, Pearson J. A growth chart for premature and other infants. *Arch Dis Child* 1971; **46**: 783–7.
- ¹⁰ Shaw J C L. Parenteral nutrition in the management of sick low birthweight infants. *Pediatr Clin North Am* 1973; **20**: 333–58.
- ¹¹ Usher R, McLean F. Intrauterine growth of liveborn Caucasian infants at sea level: standards obtained from measurements in 7 dimensions of infants born between 25 and 44 weeks of gestation. *J Pediatr* 1969; **74**: 901–10.
- ¹² Lubchenco L O, Hansman C, Boyd E. Intrauterine growth in length and head circumference as estimated from live births at gestational ages from 26–42 weeks. *Pediatrics* 1966; **37**: 403–8.
- ¹³ Senterre J. Nitrogen balances and protein requirements of preterm infants. In: Visser H K A, ed. *Nutrition and metabolism of the fetus and infant*. The Hague: Nijhoff, 1979: 195–212.

Correspondence to Dr T J French, Department of Paediatrics, Musgrove Park Hospital, Taunton, Somerset PA1 5DA.

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