incidence of breast feeding in all social classes today, however, it would surely be wrong to abandon the use of breast milk, with all its advantages, for a cows' milk formula for low birthweight infants on the mistaken premise that adequate growth cannot be achieved with the former.

We thank the sisters and nurses on special care baby unit for the accurate record keeping which made this study possible.

References

High volume milk feeds for preterm infants

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SUMMARY Two sisters with transferase deficiency galactosaemia presented with hypergonadotropic hypogonadism. In the younger girl galactosaemia was documented first at 9 months of age, although she had never been exposed to exogenous galactose in utero or after birth.

Galactosaemia due to deficiency of galactose-1-phosphate uridyl transferase (transferase deficiency galactosaemia) is characterised by a well known toxicity syndrome. Dietary restriction of galactose may prevent or reverse most of the symptoms. Only recently has ovarian failure been recognised as a complication of this metabolic disorder and we are following two sisters who presented with hypergonadotropic hypogonadism at a very young age.

Case reports

Patient 1. The older of the two girls was born at term by caesarean section to a 23 year old gravida I, para 0. Her birthweight was 3220 g. She was breast fed but started to vomit and when admitted to hospital at age 9 days was moderately dehydrated and jaundiced. Total bilirubin was 262 μmol/l, plasma glucose 3·1 mmol/l, serum glutamic oxaloacetic transaminase 123 IU/l, serum glutamic pyruvic transaminase 136 IU/l, alkaline phosphatase 527 IU/l (normal 22 to 98), prothrombin time 5%, plasma galactose 1·4 mmol/l (25·4 mg/dl), and erythrocyte galactose-1-phosphate 4·8 mmol/l (125 mg/dl). Generalised aminoaciduria was present. She was given intravenous glucose and one day later was started on a lactose free soy based formula. Galactosaemia was confirmed by a practically absent transferase activity of 0·007 μmol/min/g haemoglobin (1·4% of normal). After 8 days of treatment her erythrocyte galactose-1-phosphate concentration had dropped to 1·9 mmol/l but this was still greatly raised five weeks later (542 μmol/l). Subsequent galactose-1-phosphate concentrations, determined three monthly, were between 0 and 235 μmol/l. At 2 years of age her IQ (Brunet-Lézine) was 94. Liver enzymes were normal but her basal follicle stimulating hormone concentration was raised (Table); sex chromatin was normal. At 3½ years of age a luteinising hormone releasing hormone test (100 μg iv) showed particularly high basal and peak follicle stimulating hormone concentrations as well as a raised peak luteinising hormone concentration. She is now 4 years old and her height, weight, and head circumference are within normal limits. There are no cataracts on slit lamp examination but psychological testing has shown an IQ (Kramer) of 75 with substantial deficits in expressive speech and articulation.

Heterozygosity for galactosaemia was proved in both parents by quantitative measurement of trans-
ferase activity and electrophoretic determination of phenotypes. The mother became pregnant again when her first child was 1½ years old. Even before this pregnancy she had observed a galactose restricted diet which she rigorously adhered to until she gave birth again.

**Patient 2.** This girl was also born by caesarean section and weighed 3450 g. She never received lactose and was placed on a soy based formula from the first day of life. Transferase deficiency was confirmed with an activity of 0-009 µmol/min/g haemoglobin. When first seen in our outpatient clinic at 3 months of age she was a healthy and normally developed infant. Her transaminase concentrations were slightly raised (serum glutamic oxaloacetic transaminase 46 and serum glutamic pyruvic transaminase 62 IU/l), but later became normal. Erythrocyte galactose-1-phosphate was 146 µmol/l and has never exceeded that value at regular follow up. Basal and luteinising hormone releasing hormone stimulated gonadotrophin concentrations were raised at 9 months of age and this finding was confirmed 7 months later (Table). Sex chromatin was positive. At present she is 1 year 10 months old and her height and weight are between the 10th and 25th centile. Psychological testing has suggested slight retardation with IQ (Brunet-Lézine) of 87.

**Discussion**

The two siblings with transferase deficiency galactosaemia presented with primary hypogonadism documented by several substantially raised basal serum follicle stimulating hormone concentrations with long lasting and excessive increases in follicle stimulating and luteinising hormone after luteinising hormone releasing hormone stimulation. The older girl was treated with a galactose restricted diet from day 9 of life with rapid clinical improvement. The younger girl has been treated by dietary exclusion of galactose, practically since conception.

Hypergonadotrophic hypogonadism with primary or premature ovarian failure was present in 12 of 18 adult galactosaemic patients described by Kaufman et al.1 No ovarian tissue could be found in three of their patients and although one patient had had normal appearing gonads at age 7 years, streak gonads were found on laparoscopy 10 years later. Dietary treatment had been instituted much later on average in the patients with hypogonadism than in those with normal ovarian function (61 days v 22 days). Steinmann et al2 in a series of 15 galactosaemic patients reported a girl aged 13 years 7 months with normal female karyotype, streak gonads, and primary ovarian failure who had never been exposed to galactose either in utero or after birth. The fact that not all galactosaemic complications may be prevented by early dietary intervention has been recognised for some time. In the earlier study restriction of dietary galactose had been carried out in 11 pregnancies resulting in transferase deficient infants.3 One child had cataracts at birth; all children had noticeably raised erythrocyte galactose-1-phosphate concentrations of 150 to 631 µmol/l.

Certainly compliance with dietary galactose restriction in a heterozygote mother is difficult to monitor, however, even if dietary indiscretion has occurred, plasma galactose concentrations would not be expected to be particularly high. After intravenous administration of galactose to adult heterozygotes, plasma disappearance rate was not different from that of normal subjects (J Bircher, personal communication). Nonetheless, even low maternal plasma galactose concentrations may be detrimental to the fetus. Galactosaemia in utero with an erythrocyte galactose-1-phosphate concentration of 990 µmol/l, high liver concentrations of galactose, galactose-1-phosphate and galactitol,
Familial persistent pulmonary hypertension

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SUMMARY We report three siblings who presented with a clinical picture of persistent pulmonary hypertension of newborn and died between 4 and 15 days of age. Pulmonary artery pressure in all was above systemic values, with a right to left shunt via either the foramen ovale or ductus arteriosus, or both. Histology of the pulmonary vascular bed showed extension of muscle into small arteries which are normally non-muscular.

The term persistent pulmonary hypertension of newborn refers to a condition in which pulmonary artery hypertension prevents the successful transition from the fetal to adult circulation. The neonate continues to have raised pulmonary vascular resistance, with pulmonary artery pressure higher than systemic values. Persistent right to left shunting through the foramen ovale and ductus arteriosus results in hypoxaemia, acidosis, and respiratory distress. Although most cases are associated with well defined clinical entities, primary hypertrophy of the muscular layers of the pulmonary arterioles has been described.

We report three siblings with persistent pulmonary hypertension of newborn associated with abnormal muscularisation of the intra-acinar pulmonary arteries. To our knowledge, this is the first report of familial primary pulmonary hypertension of the newborn, and the occurrence in three siblings suggests a recessively inherited genetic trait.

Case reports

Two boys and one girl from the same family with clinical and pathological features of persistent pulmonary hypertension of newborn are described. The infants were born to healthy parents of Tunisian origin who are first degree cousins. Two additional siblings born after the death of the first two sick infants are normal. No history of unexpected fetal or neonatal deaths in the extended family was recorded. The mother received no medication during...