SPECIAL REPORT

Recommendations for the management of galactosaemia

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There is controversy about certain aspects involving the detection, outcome, and management of galactosaemia.^{1 2} The relative rarity of the disorder and lack of prospective studies have made it difficult for paediatricians to base their advice to parents and their management of patients on good clinical evidence. The UK Galactosaemia Steering Group was established in 1994 to address these problems. It is now maintained by the research division of the Royal College of Paediatrics and Child Health. The primary aim of this steering group is to set up a national register of all cases of galactosaemia and to obtain detailed prospective longitudinal data.

Having established this register it has become apparent that the management of galactosaemia varies widely. Patients are seen in many centres within the UK, both specialist and non-specialist. We believe it would be helpful for clinicians involved in their care to have recommendations for management based on currently available information. The recommendations represent a consensus view of the steering group. Those for the treatment of hypergonadotrophic hypogonadism in galactosaemia have been agreed with the British Society for Paediatric Endocrinology and Diabetes.

Classic galactosaemia

Classic galactosaemia is caused by an inherited deficiency of the enzyme galactose-1-phosphate uridyl transferase.³ Incidence in the UK is approximately 1:45 000 live births; most patients are either homozygous or heterozygous for the Q188R mutation.⁴⁻⁶ Classic galactosaemia most often presents in the neonatal period with life threatening illness. Excluding galactose from the diet combined with supportive care is essential to treat this acute illness and prevent further deterioration. Galactose restriction throughout life is necessary to stop recurrence of severe toxicity. However, early diagnosis and appropriate dietary treatment does not prevent long term complications including cognitive dysfunction, which may worsen with age, and gonadal dysfunction in female patients.⁷ Recent evidence suggests that there is endogenous production of galactose and this may be responsible for these long term effects.⁸ Follow up throughout life is necessary.

Diagnosis

Early diagnosis is essential. Neonatal screening is not routine in England and Wales. Even with screening the results of the assay are likely to be available only after the onset of clinical illness.⁴ Most infants present in the first weeks of life. Rarely patients may present after the newborn period.

MOST COMMON SYMPTOMS AND SIGNS

- poor feeding and poor weight gain
- vomiting and diarrhoea
- jaundice
- cataracts
- lethargy and hypotonia
- hepatomegaly
- encephalopathy
- full fontanelle
- bleeding or excessive bruising
- improvement on intravenous fluids.

ABNORMAL FINDINGS

Liver dysfunction

- unconjugated or combined hyperbilirubinaemia
- abnormal liver function tests
- abnormal clotting
- raised plasma amino acids (particularly phenylalanine, tyrosine and methionine. Raised phenylalanine may result in a false positive neonatal screening test for phenylketonuria).

Renal tubular dysfunction

- metabolic acidosis
- galactosuria (after feeds containing galactose or lactose) and glycosuria
- albuminuria
- aminoaciduria.

Abnormal carbohydrate metabolism

- increased plasma galactose
- increased red blood cell galactose-1-phosphate
- increased urine and blood galactitol

Haemolytic anaemia

Septicaemia (particularly *Escherichia coli*)

DIAGNOSTIC INVESTIGATIONS

Beutler test—A fluorescent spot test for galactose-1-phosphate uridyl transferase activity, now widely used for the diagnosis of galactosaemia. False negative results may occur following recent blood transfusions (within three

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Accepted 8 May 1998

months) and false positive results with glucose-6-phosphate dehydrogenase deficiency.

Red blood cell galactose-1-phosphate uridyl transferase assay—A quantitative assay to confirm the diagnosis. It also identifies variants with residual enzyme activity. False negative results may occur within three months of a blood transfusion. If there are problems establishing a diagnosis, galactose-1-phosphate uridyl transferase can be measured in red cells from both parents to establish their carrier state.

Red blood cell galactose-1-phosphate—Galactose-1-phosphate concentrations are always raised in classic galactosaemia and are not significantly affected by blood transfusions.

DNA analysis—Approximately 60% of UK patients with classic galactosaemia are homozygous for the Q188R mutation.⁵

Urine reducing substances—The presence of reducing substances or galactose in urine is neither sensitive nor specific. Small quantities of galactose are commonly found in the urine of any patient with liver disease and a non-specific glycosuria can occur with galactosaemia caused by tubular damage. Galactose may also disappear rapidly from the urine in patients with galactosaemia.

Prenatal diagnosis—Galactose-1-phosphate uridyl transferase assay in cultured amniotic fluid cells or in chorionic villus biopsies, by galactitol estimation in amniotic fluid supernatant, or by mutation analysis of DNA extracted from chorionic villus biopsy if the genotype of the index case has been characterised.⁹

Initial management

If there is any suspicion of the diagnosis (either biochemical or clinical) galactose must be excluded from the diet (table 1 lists galactose free formulae).

Supportive care should be provided as required, dependent on the severity of liver, renal, and central nervous system disease. Antibiotics, intravenous fluids, plasma, and vitamin K are often necessary.

Continue a galactose free formula after confirming the diagnosis. Refer to regional centre for specialist paediatric, dietetic, and biochemical support.

Table 1 Milk substitutes suitable for infants with galactosaemia (and prescribable in the UK on FP10 designated ACBS)

Formula	Manufacturer	Comments
Infasoy	Cow & Gate	Lactose free soya preparation
Wysoy	SMA Nutrition	Lactose free soya preparation
Prosobee	Mead Johnson	Lactose free soya preparation
Farleys Soya Formula	Farleys	Lactose free soya preparation
Isomil	Ross Products Division	Lactose free soya preparation
MCT Pepdite 0-2	SHS	For infants with severe liver disease
		MCT containing preparation, lactose free
Pregestimil	Mead Johnson	For infants with severe liver disease
		MCT containing preparation. Contains up to 0.8 mg lactose/100 ml
Enfamil	Mead Johnson	Contains up to 3 mg lactose/100ml, therefore not recommended as first choice
AL 110	Nestlé	Contains up to 10 mg lactose/100ml, therefore not recommended as first choice
Galactomin 17	Cow & Gate	Contains up to 13 mg lactose/100ml, therefore not recommended as first choice

MCT, medium chain triglyceride.

Table 2 Lactose/galactose free calcium supplements suitable for use in galactosaemia

Product	Manufacturer
Tritalac tablets	3 M
Calcichew	Shire
Calcium—500	Renacare
Sandocal 400 (10 mmol Ca ²⁺ /tablet)	Sandoz
Sandocal 1000 (25 mmol Ca ²⁺ /tablet)	Sandoz
Calcium gluconate effervescent 1 g tabs	Cox
Calcium lactate 300 mg tabs	APS
Calcium lactate 300 mg tabs	Sussex Pharm

Calcium—Sandoz is NOT suitable.

Long term management

Patients should be followed up throughout childhood and adult life. A recommended scheme for outpatient review is: every three months to 1 year of age; every four months to 2 years; every six months to 14 years; and annually thereafter. More frequent monitoring for girls in late childhood and adolescence is necessary to monitor pubertal development.

DIET

Galactose must be excluded from the diet throughout life. In essence this means no milk or dairy products. Breast feeding and cows' milk formulae are contraindicated. Soya based formulae are given as an alternative (table 1) unless there is significant liver disease, when a medium chain triglyceride containing casein based protein hydrolysate such as Pregestimil (Mead Johnson) should be used until the liver disease has resolved.

Foods such as offal, fruits, legumes, and pulses are insignificant sources of galactose compared with endogenous production and there is no evidence to support their restriction.^{8 10}

Soya infant formulae normally provide adequate calcium in infancy; however, by 1 year of age the volume of soya formula taken begins to fall. It is then necessary to change to a commercially available calcium enriched soya milk or to give adequate calcium supplements (table 2). A poor calcium intake, as well as other factors, may lead to a decrease in bone density.¹¹ The nutritional adequacy of the diet should be checked by a dietitian at least annually.

Alcohol—There is no evidence to support the hypothesis that alcohol is more harmful to patients with galactosaemia than to the normal population.

Pregnancy—Galactose ingestion by heterozygous pregnant women has not been shown to have any adverse effect on the fetus.¹² There is no evidence for milk restriction in such women during pregnancy.

Many medications, particularly tablets, contain lactose, and this should be checked before prescribing. However, in many cases, the amounts of galactose (compared with endogenous production) are insignificant, particularly if given for a short period.¹³

Biochemical findings

RED CELL GALACTOSE-1-PHOSPHATE

Concentrations of red cell galactose-1-phosphate have been widely used to monitor dietary compliance. Concentration of this metabolite is often extremely high at diagnosis and it

is essential to demonstrate its decline to acceptable values after introducing the diet. This usually occurs within a few weeks but, for unknown reasons, can occasionally take as long as a year. The units used to express galactose-1-phosphate concentrations vary between different laboratories. Target concentrations differ, to some extent, according to methodology but generally the upper limit of the acceptable range quoted in different units are as follows:

- $\mu\text{mol/l}$ red cells, 150
- $\mu\text{g/ml}$ packed cells, 50
- $\text{mg}/100\text{ ml}$, 5
- $\mu\text{mol/g}$ haemoglobin, 0.5

In patients with classic galactosaemia, despite dietary adherence, galactose-1-phosphate concentrations never decline to those found in healthy individuals (close to 0).

Long term monitoring of classic galactosaemia by frequent measurements of red cell galactose-1-phosphate is unjustified. The laboratory costs are high; a given value reflects only galactose intake in the previous 24 hours; and no correlation with long term clinical outcome has been shown. Raised concentrations may indicate continuous and serious deviations from the diet but some patients have unusually high values despite meticulous dietary compliance. Galactose-1-phosphate concentrations do not vary greatly in individual patients providing dietary compliance is reasonable. A suggested scheme for the frequency of monitoring galactose-1-phosphate levels is as follows:

- < 1 year, every three months
- 1–14 years, every six months
- > 14 years, annually

Urinary galactitol

Urinary galactitol has been investigated as a means of monitoring diet but, although it may appear to have some practical and theoretical advantages compared with galactose-1-phosphate, its value remains to be validated. It may prove useful in patients in whom cataracts fail to resolve or recur. The assay is only available in a few centres.

Development

Neurodevelopmental problems are common in children with galactosaemia.^{14 15} In preschool years speech problems are frequent and severe. Development may be generally delayed but it is more common for cognitive problems to become apparent during school years. Regular assessment of development and cognitive function are indicated using standardised tests—for example, Griffiths scales, Bailey scales, British ability scales. In particular, assessment should be directed towards early detection of speech impairment. Many children require speech therapy and additional help at school. Ataxia and intention tremor may occur in older children and adults.¹⁶

Ophthalmological findings

Slit lamp examination for cataract assessment should be made at the time of diagnosis, every six months until 3 years of age, and then annually.

Growth

Growth in infancy and childhood should be normal providing the diet is adequately supervised.

Osteoporosis

A satisfactory calcium intake should be maintained to protect patients from osteoporosis. Some units advocate regular assessments of bone density.

Pubertal development in girls with galactosaemia

Hypergonadotrophic hypogonadism occurs in most girls over 14 years of age with galactosaemia.¹⁵ Most women are infertile. Pubertal development and fertility are normal in boys.

Endocrine investigations and treatment in girls

Follicle stimulating hormone, luteinising hormone, and oestradiol should be measured at 6 months and then at 10 and 12 years old (and if necessary yearly thereafter). Referral to a paediatric endocrinologist should be made by the time the patient is 10 years old.

We recommend hormone treatment from 12 years of age (some recommend 13 years if the girl is short) with raised basal gonadotrophin concentrations and low oestradiol. One recommended regimen is as follows:

- Ethinyl oestradiol 2 μg daily for the first year—if pubertal advance is particularly slow then the time on this dose should be reduced to six months
- Ethinyl oestradiol 5 μg daily for the second year
- Ethinyl oestradiol 10 μg daily for the third year—the patient and parents should be informed that vaginal spotting or withdrawal bleeds may occur during this year; if this is the case, a progestogen should be added at the start of each month
- Loestrin (an oral contraceptive preparation containing 20 μg ethinyl oestradiol and a progestogen) given in the usual way (daily for 21 days followed by 7 days abstinence). This preparation should be adequate for most young women. If there is breakthrough bleeding or any other symptoms of oestrogen deficiency then a preparation containing 30 μg of ethinyl oestradiol can be used. (Natural oestrogens would be an acceptable alternative to synthetic oestrogen.)

NB—All tablet preparations contain small quantities of lactose but the total dose is very small compared with endogenous production.

Outpatient review should be every four months during the first two years of hormone treatment and every six months thereafter. The following should be monitored:

- physical development and growth throughout these years of pubertal induction
- bone age yearly from the start of treatment
- pelvic ultrasound (for uterine dimensions) at the start of treatment and after two and four years
- blood pressure.

Special care and advice should be given to patients with a family history or past history of venous thrombosis and to those who are overweight or smoke.

Conclusions

These recommendations are based on our current understanding of galactosaemia. It is evident that there is still much to learn about this disorder. We would welcome comments from others involved in the care of children and adults with galactosaemia. As more data become available from the galactosaemia register and elsewhere, these recommendations will be updated.

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Editors' note

In general we are reluctant to publish consensus statements or guidelines determined by a committee or working party. Such papers are not readily amenable to change following the normal editorial peer review process. We prefer authors to use systematic reviews or a similar evidence-based method when producing guidelines for publication in the journal. In this way readers can decide to what extent they can rely on the recommendations made.

We publish this paper because we realise that many patients with galactosaemia are cared for outside specialist units and it will benefit children and their doctors to have the views of the steering group.

They are recommendations—not rules—and we remind readers that our general policy when deciding whether to accept for publication guidelines and consensus statements remains the same.

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