disorders such as Marfan's syndrome, Friedreich's ataxia, or muscular dystrophy, the presence of congenital cardiac malformation, weak respiratory muscles, or extreme scoliosis.

The management suggested for any child with spinal curvature is continuing supervision throughout the years of growth by both a children's specialist and an orthopaedic surgeon oriented towards scoliosis, early correction of any congenital heart lesion, very early treatment of respiratory infection by antibiotics, and admission where practicable to an intensive-care unit for any scoliotic child becoming distressed. If these measures are carried out there is a good chance that most cases will survive.

P. T. Bray. Cardiff. 'Newborn screening for cystic fibrosis'. The results of 80,000 screening tests in newborn infants carried out as part of the work of the European Working Group for Cystic Fibrosis were presented and analysed. Techniques employed included analysis of meconium for abnormal protein content by chemical methods, Albusnix, Labstix, and a recently introduced Boehringer 'Test-Strip', as well as immunodiffusion methods. Sweat electrolyte determinations by direct reading ion-specific electrodes were also used, and measurements of the electrical conductivity of the skin, neutrophil activation analysis of nails and hair, and estimation of the sodium content of parotid saliva and electrolytes in the tears.

The work is related to the case for screening in general and screening for cystic fibrosis in particular, with regard to the possibility of earliest diagnosis leading to prevention or mitigation of the severe bronchopulmonary manifestations of the disease. The results obtained so far also enable one to assess the incidence of the disease in Europe with useful implications for genetics.

The data presented have been assembled by the subcommittee on screening of the European Working Group.

C. J. Rolles introduced by Charlotte M. Anderson. Birmingham. 'Usefulness of a modified d-xylose absorption test in the preliminary diagnosis of coeliac disease and its later confirmation'. Though jejunal biopsy remains the definitive procedure in coeliac disease (CD), its use should be selective, and there remains a need for an accurate screening test.

Seventy-one children suspected clinically to have CD had a simple estimation of blood xylose one hour after a 5 g oral dose (given in the fasting state). Later, each had a jejunal biopsy. In all 30 subsequently proven coeliac patients, the xylose level was below 20 mg/100 ml: similar levels were found in 3 noncoelics. Had the xylose result been used to select the patients for jejunal biopsy, only 33 biopsies would have been performed, and no case of CD would have been missed.

The clinical, biochemical, and histological features of CD in young infants may be difficult to differentiate from a postinfective state. In coeliac infants withdrawal of gluten from the diet led to a prompt return to normal of the xylose test, usually within a week, while gluten reintroduction caused xylose absorption to fall within a few days. Children who had been on a strict gluten-free diet for over a year showed no immediate response to a gluten challenge, but did so if gluten was continued for 6 to 8 weeks.

D. N. Challacombe. Birmingham. 'Study of duodenal microflora and bile salts in contaminated small bowel syndrome'. As bacterial overgrowth of the small intestine may occur in infancy in association with chronic diarrhoea and in the absence of anatomical abnormalities of the bowel, the term 'contaminated small bowel syndrome' has been suggested.

This paper reports a qualitative and quantitative study of the aerobic and anaerobic microflora of the duodenum in infants with chronic diarrhoea. The bacteriological results are compared with a group of control infants in hospital with disorders unrelated to the gastrointestinal tract. The duodenal juice has also been examined for the presence of bile salt abnormalities which might be associated with bacterial colonization of the small intestine.

The absence of Esch. coli in the duodenum of control infants and their presence in chronic diarrhoeal disorders suggests that this organism may play a role in the aetiology of chronic diarrhoea. As Esch. coli serotypes isolated from our infants were not among those commonly considered to be enteropathogenic, revision of the present concept of enteropathogenicity to include an increasing number of Esch. coli serotypes is proposed.

Unlike previous reports, deconjugated bile salts were not found in the duodenal juice of infants with secondary monosaccharide intolerance, but were present in one infant with secondary lactose intolerance. Concentrations of taurine conjugated trihydroxy and dihydroxy bile salts in the duodenal juice were significantly lower in patients with chronic diarrhoea than in age-matched controls.

Ann Banister introduced by G. W. Hatcher. Brighton. 'Management of hypernatraemia in infancy'. Controversy still exists over the optimal treatment of hypernatraemic infants. Limited information is available from one previous controlled trial. 38 infants with hypernatraemic dehydration (plasma sodium concentration more than 150 mEq/l) and measured plasma osmolality greater than 350 mOsm/kg water were admitted to a trial of treatment using differing regimens of intravenous rehydration.

The effects of using 0-45% sodium chloride solution with dextrose at two rates of infusion and that of using 0-18% sodium chloride solution with dextrose were compared. The use of the latter solution given at the rate of 100 ml/kg estimated rehydrated weight per 24 hours is recommended, with the early introduction of potassium. This regimen produces a satisfactory rate of fall of osmolality and of effective rehydration, with minimal risk of producing convulsions or over-expansion of the extracellular fluid volume. A plasma expander must be used in the early stages of treatment where circulatory failure is suspected. Sources of continuing excessive fluid losses from the skin and the respiratory tract must be controlled.

Details of mortality and morbidity were given, and the

C J Rolles

Arch Dis Child 1973 48: 825
doi: 10.1136/adc.48.10.825-a

Updated information and services can be found at:
http://adc.bmj.com/content/48/10/825.2.citation

Email alerting service

These include:

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/