Intensive enteral feeding in advanced cirrhosis: reversal of malnutrition without precipitation of hepatic encephalopathy

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Abstract
Ten children with advanced cirrhosis and malnutrition (<90% weight for height) were fed for eight weeks with a nasogastric feed comprising whey protein (enriched with branched chain amino acids), fat as 34% medium chain and 66% long chain triglycerides, and glucose polymer. Six of the children were studied for an eight week control period before feeding. Weight, triceps skinfold thickness, mid-arm circumference, mid-arm muscle area, and fasting plasma ammonia and amino acid concentrations were measured before and after the control period and after the consequent feed period. Results showed that despite high energy and protein intakes the children remained malnourished over the control period. All anthropometric indices improved significantly during the feed period, and no child developed clinical encephalopathy. The feed period was associated with a small, and not clinically significant, increase in the plasma ammonia concentration, but no consistent trend in the plasma amino acid concentrations. Thus, in children with advanced hepatobiliary disease awaiting liver transplantation, enteral feeding improved nutritional status without adverse clinical or biochemical effects.

Malnutrition is common in children with hepatobiliary disease. 1 2 It may be associated with a number of adverse effects such as decreased brain growth, 3 4 impairment of mental development, 5 and decreased immunocompetence 6 with an increased susceptibility to infection. 6 The pathogenesis of malnutrition in liver disease is poorly understood. Anorexia, 7 increased metabolic rate, 8 and fat malabsorption 8 are all potentially important contributory factors and raise the possibility that reversal of malnutrition may occur with appropriate intensive nutritional support. We have addressed this question with a study of intensive enteral nutrition in a group of children with severe chronic liver disease. Such an intervention in patients with advanced liver disease might be expected to exacerbate already impaired nitrogen tolerance 9 and precipitate hepatic encephalopathy. 10 Moreover, fat malabsorption is well recognised in children with cholestatic liver disease, 11 12 and an increase in dietary lipid might be expected to precipitate diarrhoea. However, there is some evidence that a nitrogen load high in branched chain amino acids may be less likely than a simple high protein diet to exacerbate such abnormalities or to precipitate hepatic coma. 13 Similarly, medium chain triglycerides may be better tolerated than long chain triglycerides in patients with chronic liver disease. 9 We therefore used a modular feed in which the nitrogen source was enriched with branched chain amino acids and the lipid source was a mixture of long and medium chain triglycerides. We performed an open controlled study with two aims. First, to determine whether malnutrition complicating advanced cirrhosis could be reversed by short term intensive enteral feeding lasting eight weeks. Second, to establish whether such an intervention was associated with dangerous clinical or biochemical sequelae.

Subjects and methods
Ten children (median age 9 months, range 4 months to 8 years; six boys and four girls) were studied. All 10 had cirrhosis that had been proved at biopsy. The plasma bilirubin and albumin concentrations at the start of the study and the aetiology are shown in the table. All were less than 90% of their expected weight for height (range 62–88%, median 78%). 13 16 Three had ascites at the start of the study and five had oesophageal varices.

Six of the children were observed over an eight week control period on their own diet before the eight week period on the trial diet. Such a control period was not possible in four children, who presented urgently to hospital and started immediately on enteral feeding.

The study was approved by the local research ethical committee and all parents gave their informed consent.

Dietary assessment
Nutrient intake before enteral feeding was assessed using seven day weighed records by parents and subsequent computer analysis of the dietary records. 17 Recall dietary assessment at home before feeding and dietetic advice given were noted for the other three children.
ENTERAL FEED
A modular feed was used. The nitrogen source (Generaid, Scientific Hospital Supplies) was whey, enriched with free branched chain amino acids (whey protein 73%, and free branched chain amino acids 27%, in protein gram equivalents). The total branched chain amino acid content of the protein was 31% (including branched chain amino acid content of whey). Each child received 4 g/kg/day of protein. Fat, comprising 34% medium chain triglyceride and 66% long chain triglyceride, and carbohydrate (as glucose polymer) were given as Duocal (Scientific Hospital Supplies). Energy content of the feed was 8% protein, 40% fat, and 52% carbohydrate. Vitamin and mineral requirements were provided for children under 1 year old as Paediatric Seravit (Scientific Hospital Supplies) which contains no sodium or potassium, and for those over 1 year old as Code 544 (Scientific Hospital Supplies) which contains no sodium, potassium, calcium, or phosphate. Sodium and potassium were provided as molar solutions and calcium and phosphate as Calcium-Sandoz and Phosphate-Sandoz (Sandoz). All subjects received additional supplements of vitamins A, D, and E. Initial feed volumes were calculated to provide either 140% of the previous energy intake or 140% of recommended energy requirements per kg for age, whichever was greater. The energy density of the feed ranged between 1 and 1.5 kcal/ml depending on the fluid allowance for the child, with a mean osmolality of 400 mosmol/kg.

ADMINISTRATION OF FEED
Feeding was started in hospital and given by continuous nasogastric infusion over 18–22 hours/day. The initial protein content of the feed was the same as the intake during the control period when this information was available or at 1 g/kg/day where not available. Intake was increased by 1 g per kg per day every third day to a maximum of 4 g/kg/day. The carbohydrate and fat components were also increased stepwise until final energy requirements were reached by five days. The children were encouraged to have additional food during the study period but none took a significant amount. The parents were taught to make up the feed, to care for the nasogastric tube (6FG Silk, E Merck), and to control the infusion pump (Kangaroo Pump, Sherwood Medical) before their child was discharged. The feed was given for eight weeks; blood samples were obtained and anthropometric recordings taken at the start and the end of this period. The children were examined for ascites and signs of hepatic encephalopathy at least every two weeks and the nutrient intake per kg body weight maintained throughout the eight week period.

BIOCHEMISTRY
Venous blood was taken after a four hour fast at the same time as anthropometry. Plasma ammonia concentration was assayed by a continuous flow ion selective electrode system. Plasma for amino acid analysis was deproteinised immediately after collection by the addition of an equal volume of 3% sulphosalicylic acid. The supernatant was stored at minus 20°C until analysis. Amino acids were measured by ion exchange chromatography using a lithium buffer system and ninhydrin detection (Kontron Chromakon 500). The reference standards used were Brodehl and Gelilissen and Scrivenger and Davies.

STATISTICAL METHODS
The Wilcoxon signed rank test was used to compare the paired results. Comparisons between the feed and control periods entered the analysis as a quadratic time component. The significance of this comparison was assessed using a repeated measures analysis of variance. Raw data was used for this analysis apart from ammonia concentrations where logarithmic transformation was performed.

Results
DIETARY ASSESSMENT
Seven children had dietary assessments before enteral feeding. Energy intakes were 90 to 165% (mean 130%) of the recommended intake per kg for age and protein intakes 2.6 to 5.7 g/kg/day (median 3.4 g/kg/day). Six children had intakes above that recommended by the World Health Organisation, and these were fed 140% of their recorded intake for the feeding period. The other four children received 140% of their recommended energy intake. Before the trial four children were on normal diets, four were taking energy supplements (carbohydrate and medium chain triglycerides), and two were on Pregestimil (Bristol-Myers), one of whom was also on energy supplementation. Nine of the...
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children had dietetic advice to maximise energy intake.

TOLERANCE OF FEEDING

Both the nasogastric tube and the enteral feed were well tolerated with no adverse clinical events. In particular, diarrhoea was not a problem. Regurgitation, which had been present before the trial, was exacerbated in two infants but lessened with feed thickening. If the polyurethane nasogastric tube continued to be displaced frequently by the vomiting it was replaced by a stiffer polyvinyl chloride tube (Portex). None of the 10 children developed fluid overload (development of ascites or peripheral oedema). Two of the three with ascites before feeding had lost it by the end of the feed period. The modular nature of the feed allowed precise control of fluid and sodium intake, which aided the control of the ascites. There were no episodes of variceal haemorrhage. All 10 children maintained their target intakes throughout the eight week period.

ANTHROPOMETRY

The weight, mid-arm circumference, and mid-arm muscle area before the eight week control period, where available, and before and after the consequent eight week feeding period are shown in fig 1. Malnutrition was severe before feeding, and nine of the children were more than 2 SDs below the reference mean for mid-arm circumference. All children increased their weight, triceps skinfold thickness, mid-arm circumference, and arm muscle area during the feed period and these changes were statistically significant (p<0.005). Statistically significant improvements over the intervention period compared with the control period were obtained for all measurements (weight p<0.005; mid-arm circumference, triceps skinfold thickness, and arm muscle area p<0.001).

BIOCHEMICAL MEASUREMENTS

Individual plasma ammonia concentrations at the start of the eight week control period and before and after the eight week feed period are shown in fig 2. Prefeeding plasma ammonia concentrations in eight out of the 10 children were above the laboratory reference range for well children. Analysis of the changes in ammonia concentration both during the feed period and between the control and feed period were statistically significant (p<0.05). However, there was much variance in the data and the concentrations never approached those usually associated with clinical problems.27 No patient developed hepatic encephalopathy, and no parent reported increased drowsiness in their child. Indeed, nine out of the 10 children were reported by parents to have become more active and lively during the trial.

The mean fasting plasma amino acid concentrations before and after feeding are shown in fig 3. Concentrations are expressed as the number of SDs from the reference mean for the age band. The majority of amino acid concentrations were in the normal range before feeding
Before enteral feeding

After enteral feeding

Figure 2  Fasting plasma ammonia concentrations before and after eight week control period and after consequent enteral feeding period in children with cirrhosis. Upper limit of laboratory reference range (95% confidence limits) is 40 μmol/l. Analysis of the paired results before and after the feed period and the change over control period compared with that over the feed period gave p<0.05 for both.

Figure 3 Changes in mean fasting amino acid concentrations of 10 children with cirrhosis before and after eight weeks of enteral feeding expressed as a SD score of reference standards. The bold circle represents the mean amino acid concentration for the reference population. The dashed concentric circles represent 2 SD steps from the reference mean. Methionine has an extended scale. Individual amino acids values are plotted on the radius of the diagram. Analysis of the individual paired results before and after feeding for the amino acids shown (and also glutamine, aspartate, cystine, serine, alanine, proline, and lysine) were statistically non-significant apart from valine, p<0.05 and glycine, p<0.005.

Figure 4 Fasting plasma methionine concentrations for 10 children with cirrhosis before and after the eight week enteral feeding period with a repeat result for an individual (still on the trial regimen) who had an intercurrent viral infection at the time of the eight week sample. Analysis of the paired results was not significant.

(-2 to +2 SD) and were unchanged afterwards. In particular branched chain amino acids were all near the reference mean both before and after feeding. Lysine, threonine, and phenylalanine concentrations were above the reference range initially, but did not increase after feeding. Mean methionine concentrations were raised both before and after feeding, a finding consistent with severe liver disease. While the mean methionine concentration was higher at the end of feed period than before, there was no statistically significant difference between the individual paired results. This rise was biased by the results in three children in whom methionine concentrations increased greatly. Two of those three had deteriorating liver function during the study as determined by prolongation of prothrombin time. The third child had an intercurrent viral infection in week 8 of the feed period at the time of the second sample, and a repeat methionine concentration after recovery while still on nasogastric feeding was similar to the prefeed value (fig 4). The other seven children had appreciably raised methionine concentrations up to 9 SDs above the reference mean both before and after feeding with little individual change.

Discussion

Intensive enteral feeding appreciably improved the nutritional status of the children, assessed clinically by upper arm measurements and by weight. There were no serious deleterious effects. In particular, no episodes of hepatic
encephalopathy were precipitated. Although the children studied were malnourished, we found no evidence of a poor dietary intake. This differs from the observations of Watkins and Glassman. The majority of children we studied had received dietary advice aimed at increasing energy intake before feeding and had an energy intake higher than that recommended for age and weight. Our sample was small but these observations point to malabsorption, poor utilisation, or increased metabolic requirements as more important mechanisms for protein energy malnutrition than inadequate intake. This is supported by the demonstration of Pierro et al that children with extrahepatic biliary atresia have both an increased resting energy expenditure and large energy losses in the stools.

Fasting plasma ammonia concentrations were above the reference range before feeding and had risen further by the end of the study period. However, this was not associated with any clinical deterioration and the concentrations were much lower than seen for example in Reye's syndrome. This study had small patient numbers and until more data and clinical follow up are available, we would advise clinical observation for adverse effects and measuring plasma ammonia in situations where dietary nitrogen is being increased in children with severe liver disease.

The amino acid profiles before feeding showed similar abnormalities to those documented previously, with particular elevations of methionine, threonine, and phenylalanine. Branched chain amino acid concentrations were in the reference range. This pattern changed little with enteral feeding and there were no increases in branched chain amino acid concentrations. Thus, in the group studied, no gross abnormality was attributable to the high nitrogen load. In this study we have demonstrated that malnutrition in children with cirrhosis occurs despite a normal or high energy intake; it can be reversed without deleterious effects by an intensive period of enteral feeding. However, we cannot conclude that branched chain amino acid enrichment has any advantage over standard protein and this will require further study.

The only chance of long term survival for these children is liver transplantation. We speculate that better nourished children will have improved survival before transplantation during their wait for a donor organ. In one centre, 25% of children died while awaiting transplantation. Poor pretransplant nutrition may also be associated with an adverse result after grafting. Prior correction of severe protein energy malnutrition may therefore improve the long term outlook of children with severe liver disease awaiting transplantation.

We thank Alan Girling, School of Mathematics and Statistics, University of Birmingham, for statistical advice; Scientific Hospital Supplies, Liverpool for supply of the feed components; and 'Remember Rebecca Trust' for provision of computer equipment.

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