Effect of a late evening meal on nitrogen balance in patients with cirrhosis of the liver

G R Swart, M C Zillikens, J K van Vuure, J W O van den Berg

Abstract

Objective—To assess whether a late evening meal would improve nitrogen balance in patients with cirrhosis of the liver.

Design—Randomised crossover study of meal schedules comparing three meals a day with four or six meals a day, the four and six meal schedules both including a late evening meal (2300).

Setting—Metabolic ward.

Patients—Seven men and two women aged 34-66 with cirrhosis of the liver (Child's grade B).

Interventions—Patients spent two seven day periods in the ward. For five days of each period they received, in random order, isonitrogenous isocaloric diets supplied in three meals a day and in four or six meals a day.

Main outcome measure—Nitrogen balance, calculated as the difference between dietary intake and the total of urinary, faecal, and integumental nitrogen loss.

Results—Faecal nitrogen loss was no different between three meals a day and four or six meals a day. On both four and six meals a day, however, patients had nitrogen balances that were more positive (or less negative) than on three meals a day (1-26 (SD 2-1) g/24 h v 0-26 (2-2) g/24 h, p<0-01). Six meals a day did not produce significantly better improvements in nitrogen balance than four meals a day.

Conclusions—A late evening meal seemed to improve the efficiency of nitrogen metabolism, but longer term studies are needed to assess whether this leads to a better nutritional state.

Introduction

Patients with cirrhosis of the liver are often malnourished. They need more protein to maintain nitrogen equilibrium than healthy people, and their rates of protein turnover—both synthesis and breakdown—are increased. Nevertheless, protein intolerance can lead to portosystemic encephalopathy, and dietary protein may have to be reduced to levels below those needed to maintain nitrogen balance.

Increasing the efficiency of nitrogen metabolism would help to overcome the therapeutic dilemma of choosing between protein need and protein tolerance; it might also improve the nutritional state. We hypothesized that the high protein requirements could be explained by reduced liver glycogen stores, the early onset of gluconeogenesis from amino-acids at night leading to an additional amino acid loss. We recently showed that nocturnal glucose supplementation does indeed reduce raised protein turnover rates and leads to a better nitrogen balance in patients with cirrhosis. As nocturnal glucose feeding is not easy to continue in practice we studied the effects of meal frequency on nitrogen balance, on the assumption that a late evening meal would delay the otherwise early onset of nocturnal amino acid breakdown for gluconeogenesis; it would thus reduce amino acid loss and improve nitrogen balance.

Patients and methods

Nine patients with Child grade B cirrhosis of the liver confirmed at biopsy were studied (table I). During the study they stayed in a metabolic ward after admission to a medical ward for clinical stabilisation. They had no fever, no signs of infection, and no ascites or portosystemic encephalopathy; they had not recently drunk alcohol. Their mean age was 54 (range 34-68) years. The body mass index (Quetelet index) was 26·4 (range 20·2-34·3) kg/m².

The patients were admitted to a metabolic ward for two consecutive periods of seven days. In each period two days were spent on metabolic equilibration and nitrogen balance was measured over the following five days. During the two periods the patients consumed isonitrogenous isocaloric diets in two of three schedules. All the patients received three meals a day in one period; in the other they received either four meals a day or six meals a day. The order of the meal schedules was random. Meals were given at 0800, 1200, and 1700 (three meal periods); 0800, 1200, 1700, and 2300 (four meal periods); or 0800, 1000, 1200, 1500, 1700, and 2300 (six meal periods). Nitrogen balance was calculated as the difference between the measured dietary nitrogen intake and the total of the measured daily urinary nitrogen excretion, the measured faecal nitrogen loss (five day pools), and the integumental nitrogen loss, estimated to be 0·5 g/24 h. To correct for incomplete collection of faeces we gave the patients 1200 mg of polyethylene glycol daily and determined faecal polyethylene glycol excretion.

Diets—The total caloric content of the diets was chosen according to each patient's appetite as found by dietary questioning (table II). A mean of 114 kJ/kg/day (range 80-139) was chosen. The protein content of the diet was decided on by the physician: a mean of 67·4 (range 59·6-99·0) g/day was given. The late evening meal was scheduled to contain about 20% of the total daily energy and protein intake (table II); on average it supplied 17% of the energy and 20% of the protein.

Laboratory procedures—Urinary nitrogen was measured in an automatic nitrogen analyser (ANA 1400; Carlo Erba, Milan, Italy) by the Dumas procedure. The coefficient of variation for repeated measurements with this instrument is 1·7%. Faecal samples were destroyed by acid and heat; the remaining clear solution was analysed for nitrogen content in

<table>
<thead>
<tr>
<th>Case No</th>
<th>Sex and age</th>
<th>Cause</th>
<th>Quetelet index (kg/m²)</th>
<th>Bilirubin (µmol/l)</th>
<th>Alanine aminotransferase (U/l)</th>
<th>Preadminin (mg/dl)</th>
<th>Albumin (g/dl)</th>
<th>Ammonia (µmol/l)</th>
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<tr>
<td>1</td>
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<td>32</td>
<td>23</td>
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<td>Alcohol</td>
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<td>59</td>
<td>41</td>
<td>0·38</td>
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<td>93</td>
</tr>
<tr>
<td>3</td>
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<td>27·0</td>
<td>40</td>
<td>35</td>
<td>0·08</td>
<td>33</td>
<td>90</td>
</tr>
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<td>4</td>
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<td>32</td>
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<td>85</td>
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The diets were well tolerated, and the patients showed no preference for either meal schedule. Faecal nitrogen loss was no different between the three meals a day schedule and the four or six meals a day schedule (2.01 (0.62) g/24 h x 2.13 (0.56) g/24 h). On three meals a day both positive and negative nitrogen balances were found. There was no significant correlation between the nitrogen balance and the serum albumin or bilirubin concentration, which was taken as a marker of the severity of the liver disease.

On four meals a day three patients were in positive and two in negative nitrogen balance; on six meals a day all four patients were in positive nitrogen balance. On both four and six meals a day nitrogen balances were consistently more positive or less negative than on three meals a day (1.97 (2.17) g/24 h x 2.06 (2.2) g/24 h; p<0.01 Wilcoxon test; table III); the mean difference was 1.1 (0.3) g/24 h (range 0.24-3.17). Six meals a day did not produce a significantly better improvement in nitrogen balance than four meals a day (1.33 (0.08) g/24 h x 0.73 (0.11) g/24 h).

Discussion
In this study nitrogen balance was always more positive, or less negative, on isonitrogenous and isocaloric diets when a late evening meal was given. The additional nitrogen retention with an extra meal at bedtime was 1.0 g/24 h (range 0.24-3.17) on average. Thus the efficiency of nitrogen metabolism seemed to improve when a late evening meal was given. A retention of 1 g nitrogen, when incorporated into protein, is roughly equivalent to 6.25 g of protein or about 30 g of lean tissue. If such a dietary regimen could be continued over longer periods a better nutritional state and an increase in lean body mass could be expected. Such improved efficiency would be especially useful in patients who need to take protein restricted diets to prevent portosystemic encephalopathy or in those with signs of protein-energy malnutrition. Further clinical studies, including body composition measurements, will, however, be needed to assess whether this effect of meal frequency on nitrogen balance is clinically important. We are not aware of previous reports of studies on the effects of meal frequency on nitrogen balance in cirrhosis. In healthy people meal frequency does not seem to influence nitrogen balance.

Our results support the hypothesis that increased protein requirements in cirrhosis might result from additional amino acid loss caused by the early onset of nocturnal gluconeogenesis from amino acids, in turn caused by small liver glycojen stores. Might the effects on nitrogen balance also be obtained by giving more frequent small meals only during the daytime? Such a schedule, excluding a late evening meal, has not been tested. Although we did not find a statistically significant difference in nitrogen balance between the schedule with four meals a day and that with six meals (in which five of the six meals were taken during the daytime), the numbers of observations were too small to reach a firm conclusion. A greater absorption of nutrients as a result of more frequent presentations of food to the gut probably does not explain the better nitrogen balance since faecal nitrogen loss was the same with three and four and with three and six meals a day.

Additional studies of the long term effects of a higher meal frequency on body composition and clinical complications in patients with cirrhosis are needed. The clinical implication of these findings is that cirrhotic patients should be advised to take an additional meal at bedtime.

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