Successful use of vancomycin hydrochloride in the treatment of lactulose resistant chronic hepatic encephalopathy

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Abstract
Vancomycin hydrochloride (2 g daily) was administered to 12 patients with cirrhosis and lactulose resistant portal systemic encephalopathy in a double blind crossover trial. All 12 patients showed a remarkable clinical improvement after vancomycin treatment. The mean (SE) electroencephalographic (EEG) frequency changed from 6-3 (0·2) to 8·5 (0·2) cps (p<0·001) and the mean arterial ammonia concentration from 152 (4) μg/ml to 97 (8) μg/ml (p<0·001). Their clinical condition deteriorated when treatment was switched to lactulose, returning to the previous slower EEG frequency and high arterial ammonia concentrations. Vancomycin seems to be effective in chronic portal systemic encephalopathy in patients who are not helped by lactulose alone.

In recent years, chronic recurrent portal systemic encephalopathy in patients with cirrhosis has generally been treated with restriction of dietary protein and oral lactulose. There are, however, occasional patients with portal systemic encephalopathy who do not respond to oral lactulose treatment and develop hepatic encephalopathy despite this drug. Although a specific amino acid solution, Fischer’s solution, has also been advocated in severe hepatic encephalopathy in patients with cirrhosis, it is difficult, in practice, to administer over a long period of time.

We have recently shown that vancomycin hydrochloride, a non-absorbable antibiotic which is effective against anaerobic Gram negative rods but ineffective against aerobic ones, is useful in controlling portal systemic encephalopathy in patients with cirrhosis. It has also been shown that changes in blood ammonia concentrations correspond very well with changes in the number of anaerobic Gram negative rods in faeces.

We therefore gave vancomycin hydrochloride to cirrhotic patients with portal systemic encephalopathy that was resistant to oral lactulose. We examined changes in their faecal bacterial flora and report the effectiveness of this antibiotic in these subjects.

Methods

Patients
Fourteen of 29 consecutive patients with cirrhosis and portal systemic encephalopathy admitted to Kanagawa Cancer Center Hospital and Kawasaki Kyodo Hospital between 7 April 1986 and 10 March 1988, whose portal systemic encephalopathy was resistant to continuous lactulose administration, were admitted to the study. There were 13 men and one woman. The proportion of lactulose resistant patients was high because many had been transferred to these two centres from outlying hospitals. The diagnosis of liver cirrhosis was made by laparoscopy and examination of liver biopsy specimens. Rupture of oesophageal varices occurred in one man soon after vancomycin hydrochloride administration, and he was excluded from the study. One woman was unable to continue taking vancomycin because of severe nausea and she was also excluded. Twelve patients finished the study.

None of the 12 patients had abused alcohol for more than two years and the liver lesion was biochemically and histologically inactive in all but one, who showed only mild activity.

All patients had given informed consent and the research was carried out according to the declaration of Helsinki.

Study design
Before the trial the patients were admitted to hospital for two weeks (the control period) for assessment of their clinical condition. During this period lactulose was given to all patients and the amount was adjusted individually to induce two to four bowel movements a day. Protein intake was restricted to about 50 g daily and there were no changes in diet during the trial period. At the end of the two week assessment period, the mental status of the patients was evaluated and graded according to the criteria suggested by Parsons-Smith et al., modified by Conn et al. The mean electroencephalographic (EEG) frequency was estimated, the fasting arterial ammonia concentration was measured by a modified method of Okuda and Fujii, and a bacteriological study of stools was done.

The trial then started. During the first eight weeks all patients were given 1000 mg of vancomycin hydrochloride (VCM, Shinogi Pharmacological Co, Tokyo, Japan) orally twice daily – at 6 am and at 6 pm. The vancomycin was diluted in distilled water to a volume of 30 ml. At this dose vancomycin has a mildly laxative property and produces one to three semi-formed stools per day. Once the vancomycin administration had been established, the patients were discharged and followed for eight weeks as outpatients. During this period, clinical and EEG assessments were made twice weekly. After
Vancomycin and hepatic encephalopathy

Figure 1: Design of the vancomycin hydrochloride trials. During the first eight weeks all 12 patients received vancomycin hydrochloride in a single blind trial. The patients were then randomised to a double blind crossover trial of vancomycin hydrochloride v lactulose.

During the eight weeks the patients were readmitted to hospital for a further two weeks and the same assessments were performed as during the control period.

The double blind randomised crossover study of vancomycin v lactulose was performed as follows: six patients (group A) selected at random were changed to lactulose alone immediately after the initial eight weeks on vancomycin, while the remaining six (group B) remained on vancomycin for another eight weeks. After this period, the medication was reversed and the study was continued for a further eight weeks (Fig 1). The amount of lactulose given was again adjusted individually as described previously and the procedure for giving vancomycin was the same as it had been at the beginning of the study.

All the analyses of EEG and bacterial flora were conducted on a double blind (ie coded) basis.

During the subsequent study, these 12 patients remained clinically stable with no gastrointestinal bleeding, electrolyte abnormalities, constipation, or infection and they all completed the study.

The patients’ characteristics in each group at the beginning of the study were as follows: group A comprised six men and group B five men and one woman. There was no significant difference in the mean (SE) age between group A (58·3 (4·2) years) and group B (61·0 (4·3) years). The type of cirrhosis in group A was alcoholic in four patients, posthepatitic in one, and cryptogenic in one. In group B it was alcoholic in three, posthepatitic in one, and cryptogenic in two patients.

With regard to liver histology, the ratio of micronodular: macronodular: mixed nodular cirrhosis was 1:3:2 in group A and 1:4:1 in group B. The average duration of encephalopathy was 2·0 (0·6) years in group A and 1·7 (0·4) years in group B.

The biochemical data for the two groups at the beginning of the study were as follows: serum albumin concentration (normal range, 4·0-5·0 g/l) was 3·4 (0·2) in group A and 3·1 (0·2) in group B, serum aspartate transaminase activity (normal range <40 IU/l) was 66 (8) in group A and 45 (8) in group B, serum alkaline phosphatase activity (normal range 5-17 umol/l) was 9·8 (1·0) in group A and 10·3 (1·7) in group B, total bilirubin concentration (normal range 0-10%) was 45·7 (5·5)% in group A and 41·9 (3·6)% in group B. There were no significant differences between group A and group B in any of these parameters.

Bacteriological study

Bacteriological studies of faecal flora were carried out as shown in Figure 2. A 1 g specimen of each fresh stool was subjected to serial 10-fold dilution immediately after defecation. The anaerobic diluent consisted of 4 g KH2PO4, 6 g Na2HPO4, 1 g L-cystein, 1 g agar, and 1000 ml of distilled water. The dilution was carried out in the glove box anaerobic chamber. A standard loopful of the diluent was inoculated onto the specially designed culture media shown in Figure 2. The aerobic plates were incubated for two to four days. The anaerobic plates were incubated for four to seven days by the gas pack method. Isolated strains of both aerobes and anaerobes were identified according to Bergey’s classification.

Statistical analysis

Statistical comparisons of the data were made using the paired Student’s t test.

Results

As shown in Table I the grade of hepatic encephalopathy improved in all 12 patients when vancomycin was given. The improvement was

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Mean (SE): 7·0 (0·3) p<0·001 0·2 (0·2)
usually evident after two to three days of treatment. In 10 patients the encephalopathy resolved completely and in the remaining two only grade 1 encephalopathy persisted. The grade of encephalopathy after vancomycin (0-2 (0-2)) was significantly (p<0-001) less than that beforehand (2-0 (0-3)). The EEG frequency after vancomycin (8-5 (0-2) cps) was significantly (p<0-001) larger than that beforehand (6-3 (0-2) cps) (Table II).

In the crossover double blind study, the tendency was the same. In group A, the mental status deteriorated when patients were crossed to lactulose, and improved again when they crossed back to vancomycin. In group B, the improvement in mental status persisted for about two months, and thereafter it deteriorated in five of six patients when they were crossed to lactulose.

**ARTERIAL AMMONIA CONCENTRATION**

Table III shows the changes in the arterial ammonia concentration in the 12 patients throughout the vancomycin trial. In the control period the mean (SE) ammonia concentration was 152 (4) μg/ml and it fell significantly (p<0-001) to 97 (8) μg/ml after vancomycin therapy. The change to lactulose resulted in a return to the previous high values.

**FAECAL BACTERIAL FLORA**

Table IV shows the changes that occurred in faecal bacterial flora during the study.

Generally, the count of total anaerobic Gram negative rods ran as high as 10^10-10^17 organisms per g stool before vancomycin (ie when lactulose was given), and by contrast the count became as low as 10^10-10^12 after vancomycin. There were significant differences between the control period and the period of vancomycin administration with respect to the log bacterial counts in anaerobic Gram negative rods (p<0-01, p<0-02, respectively).

The anaerobic Gram negative rods most commonly found in faecal specimens before vancomycin administration were bacteroides, and *Bacteroides sp* found in that period are shown in Table V.

On the other hand, the counts of aerobic Gram negative rods were 10^-4-10^-8 per g of stool before vancomycin treatment, and these remained nearly the same or slightly increased afterwards.

**Discussion**

The management of chronic recurrent portal systemic encephalopathy is difficult and unsatisfactory. It consisted of restriction of dietary protein and antibiotic suppression of the intestinal bacteria, usually by neomycin until 1967 when lactulose was introduced. This synthetic disaccharide (1,4-galactoside fructose), which is neither absorbed nor hydrolysed in the upper intestinal tract of man, passes unchanged into the lower bowel where it is metabolised by bacteria with the production of lactic, acetic, and formic acids and carbon dioxide. It was initially suggested that acidification favoured growth of lactobacilli and other acidophilic fermentative bacteria and suppression of acidophilic proteolytic bacteria resulting in a decrease in ammonia production in the colon.

It has recently been suggested, however, that a major effect of lactulose is to augment the incorporation of nitrogen (especially ammonia) into faecal bacteria for synthesis of bacterial protein, although nitrogen in the soluble frac-
tion also increases, and that the increased faecal nitrogen excretion, mostly in the bacterial fraction of stool, would reduce urea synthesis.12-16

The cathartic properties of lactulose and acidification of colonic contents are thought to be of secondary importance in this theory. More recently, it has also been shown that at least half of the total ammonia production in the gut is the result of non-bacterially mediated glutamine degradation in the intestinal wall,17 and that lactulose inhibited degradation of glutamine to ammonia in vitro18 and in vivo.19,20

Lactulose has been reported to be effective in the treatment of portal systemic encephalopathy by a number of investigators,12,21 and in recent years a combination of dietary protein restriction and oral lactulose administration has become the most popular treatment for chronic recurrent portal systemic encephalopathy in Japan. There are, however, many patients with cirrhosis and hepatic encephalopathy in whom this combination does not bring about full recovery. A specific amino acid solution like Fischer’s solution has been advocated in severe hepatic encephalopathy in these patients.11 It is, however, difficult to control chronic recurrent portal systemic encephalopathy for a long period with such a method because of the time and expense needed.22

We recently showed that vancomycin, a non-absorbable antibiotic that suppresses anaerobic Gram-negative rods but not aerobic ones,23 is very effective in chronic recurrent portal systemic encephalopathy.24

We therefore administered vancomycin to patients with intractable chronic recurrent portal systemic encephalopathy which developed in spite of the oral lactulose treatment. This study shows that vancomycin is surprisingly effective in these patients. It improved their clinical symptoms, EEG abnormalities, and blood ammonia values.25

In recent years, it has been shown in vitro that an appreciable proportion of the non-sporing anaerobes, especially bacteroides, produce urease,26 and that Gram-negative anaerobic rods (chiefly bacteroides) are more active than aerobic enterobacteria in the production of ammonia from nutrient medium containing peptides and amino acids.24 Indeed the Bacteroides sp isolated in this study belonged to the most active species in ammonia production. Moreover, in 1982 Morgan et al27 showed clinically that in the treatment of hepatic encephalopathy in a series of 11 mildly or moderately affected and seven severely affected patients with histologically confirmed cirrhosis, metronidazole, which is active against bacteroides and other anaerobes, was as effective as neomycin and that metronidazole could also reduce the production of endogenous ammonia by its effect on the anaerobic intestinal flora.

More recently we have shown that bacteroides may be the main producers of ammonia in hepatic encephalopathy in cirrhotic patients, and that suppression of these bacteria by vancomycin brings about recovery from hepatic encephalopathy in many patients.28

One of the reasons for the effect of vancomycin in lactulose-resistant intractable portal systemic encephalopathy could be its ability to decrease

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We thank emeritus Professor Kunio Okuda of Chiba University for his indispensable assistance in the preparation of the manuscript.

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18 Van Leeuwen PAM, Bogaart EJ, Janssen MA, de Boer JG, van Eyk HMA, Soeters PB. Ammonia production and glutamine metabolism in the small and large intestine of the