TREATMENT OF HEPATIC ENCEPHALOPATHY: TARGETING THE GUT-LIVER-BRAIN AXIS

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Chairperson
Rajiv Jalan

Speakers
Agustin Albillos,2 Flemming Bendtsen3

1. UCL Institute for Liver and Digestive Health, Royal Free Hospital, London, UK
2. Dept. of Gastroenterology and Hepatology, Hospital Universitario Ramón y Cajal, Universidad de Alcalá, CIBERehd, Madrid, Spain
3. Dept. of Gastroenterology, Hvidovre Hospital, University of Copenhagen, Denmark

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Introduction

Professor Rajiv Jalan

The pathogenesis of hepatic encephalopathy (HE) involves the interaction between two pathophysiological mechanisms: ammonia detoxification and inflammation. The neuropathology of HE is characterised by astrocyte dysfunction and swelling due to ammonia detoxification. The enzyme glutamine synthetase, which is located mainly in astrocytes, protects neurons by absorbing excess ammonia and glutamate, converting it to glutamine. The concentration of ammonia is increased in patients with hepatic failure when compared with healthy individuals, clearly indicating the significance of ammonia in the pathogenesis of HE, and as an important target for treatment.

Bacterial translocation is a significant factor in driving the progression of cirrhosis, hepatic fibrosis, compensated and decompensated cirrhosis and the recurrence of HE (Figure 1). Bacterial translocation appears to be a key event in the transition from well-compensated to decompensated cirrhosis (or acute on chronic liver failure). This transition manifests in severe levels of HE, and probably contributes directly to the ‘second hit’ which is inflammation.

Gut Bacterial Translocation in Cirrhosis

Professor Agustin Albillos

Intestinal microflora and gut bacterial translocation (GBT) have been implicated in the pathogenesis of spontaneous bacterial infections and in the progression of cirrhosis. Bowel decontamination with quinolones or fluoroquinolones has been shown to improve survival in patients with decompensated cirrhosis. Non-absorbable antibiotics such as rifaximin reduce the rate of spontaneous bacterial infection, the rate of portal hypertension related complications, and improve survival.1,2

The basic mechanism that underlies most episodes of spontaneous bacterial infection in cirrhosis is GBT. GBT is defined as the growth of viable bacteria in a mesenteric lymph node culture. GBT is increased in experimental models of cirrhosis and in patients with cirrhosis and ascites.3,4
A positive mesenteric lymph node culture suggests increased passage of enteric bacteria due to increased intestinal permeability, intestinal bacterial overgrowth, or both. More importantly, it also indicates an inability of the immune system to destroy the translocated bacteria. According to the previous definition, GBT was present in about 30-40% of patients with cirrhosis and ascites and in cirrhotic rats with ascites. This demonstrates that bacterial translocation is present in advanced decompensated cirrhosis when severe liver insufficiency has already developed.

Most of the translocated bacteria belong to the common intestinal microbiota, which signifies that there is a disruption of the intestinal barrier in cirrhosis. The intestinal barrier is composed of three interrelated layers, the external composed by mucous and bacterial microflora, the epithelial cells and the sub epithelial where interactions take place between the bacteria and the immune system. The integrity of the epithelial cells is the most important layer of defence against microbiota. Abnormalities in any of these levels of defence have been advocated to explain the high rate of GBT of cirrhosis. Intestinal bacterial overgrowth, increased intestinal permeability and impaired immune system response indicate changes in each of the three layers of the gut barrier in cirrhosis. In cirrhosis, there are abnormalities in the function and structure of the intestinal mucosa, involving tight junction proteins, which lead to increased permeability of macromolecules. There are also qualitative (dysbiosis) and quantitative changes in gut microbiota that indicate intestinal overgrowth, which is associated with most episodes of bacterial translocation. This is mainly attributed to the presence of intestinal hypomotility in cirrhosis, although impaired immunity can also contribute. However, other elements are necessary for GBT to develop; these are predominantly linked to the hepatic insufficiency that is found in cirrhosis. The exact mechanism is unclear, but it is possibly related to impaired immune function considering the role of the liver in innate immune function. There might also be contribution of the neuroendocrine abnormalities present in liver cirrhosis, specifically sympathetic nervous system hyperactivity and changes in bile flow and composition. Therefore, GBT in liver cirrhosis is
the result of damage at different levels of the intestinal barrier, i.e. changes in microbiota, changes in the integrity of the epithelium and impaired immunity (Figure 2).

In the absence of overt infection, GBT contributes to cirrhosis progression by inducing an activation of the immune system at the systemic level. Systemic inflammation results from the production and release of pathogen-associated molecular patterns that activate specific receptors on the surface of immune cells. This results in the production of proinflammatory lymphokines and monokines, and circulating immune system cells in cirrhosis produce proinflammatory cytokines. This mechanism modulates the clinical expression of cirrhosis, for example the modulation of the activity of astrocytes leading to the neurological changes of cirrhosis or the regulation of vascular tone. Persistent activation of immune cells worsens the immunodeficiency of cirrhosis because it leads to the exhaustion and death of the cells (Figure 3). This situation can be reversed by reducing the enteric bacterial load with non-absorbable or poorly absorbable antibiotics. This was shown in a study where bowel decontamination with antibiotics improved the dendritic cell function of the intestinal lamina propria of cirrhotic rats with bacterial translocation.

Multiple intestinal damage in cirrhosis involves different potential therapeutic targets. These include bile acids, farnesoid X receptor (FXR) agonists for impaired bile flow or composition, beta-blockers for intestinal hypomotility, antioxidants for inflammation oxidative stress and antibiotics, and probiotics for microbiota. The most effective treatment is controlling the microbiota with non-absorbable antibiotics to prevent spontaneous bacterial infection. The use of other specific

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**Figure 2. Mechanisms of gut bacterial translocation in advanced cirrhosis.**
targeted agents (FXR agonists, antioxidants and probiotics) has only been studied in experimental settings; therefore, efficacy data in patients are not available. However, the use of beta-blockers in cirrhotic patients has been shown to be beneficial by reducing portal pressure and GBT. This was demonstrated in cirrhotic rats; propranolol was shown to accelerate intestinal transit and lower intestinal bacterial overgrowth and GBT to mesenteric lymph nodes.

In summary, GBT in cirrhosis is the pathophysiological hallmark of spontaneous bacterial infections. GBT contributes to further decompensation in already decompensated cirrhosis by driving a persistent activation of the inflammatory immune system and exacerbating immunodeficiency. As the pathogenesis of GBT in cirrhosis involves damage at different levels of the intestinal barrier, there are multiple potential targets for the control of GBT. Although most targets have only been tested in an experimental setting, it has been shown that bowel decontamination improves outcomes in patients with decompensated cirrhosis. Furthermore, beta-blockers improve survival by reducing the variceal bleeding risk and bacterial translocation.

Clinical Consequences of Bacterial Translocation – Does Gut Decontamination Improve Outcome?

Professor Flemming Bendtsen

Bacterial translocation is associated with cirrhosis and portal hypertension and contributes to splanchnic vasodilation and systemic vasodilation. Therefore, clinicians should focus on bacterial translocation in patients with ascites and decompensated liver disease. As a consequence of portal hypertension and possibly GBT, patients may develop an increased hepatic venous pressure gradient (HVPG) resulting in the formation of oesophageal varices. Furthermore, systemic inflammatory response leads to disease progression, with an associated increased risk of infection. Structural and functional changes in the gut mucosa, bacterial overgrowth in the small intestine, impairment of defence mechanisms, and decreased gut motility can all lead to bacterial translocation.

Bacterial decontamination is believed to modify the risk of complications of portal hypertension. There are no large clinical trials that evaluate clinical endpoints such as death, development of variceal bleeding, hepatorenal syndrome, and
other complications of cirrhosis. However, there are trials that evaluate surrogate markers such as the effect on haemodynamics and inflammation, vasoactive hormones, and inflammatory markers. Albillos et al.\(^2\) evaluated lipopolysaccharide binding protein (LBP) levels in cirrhotic patients with marked immune and haemodynamic derangement. Patients were randomised to either norfloxacin (which targets most of the Gram-negative bacteria in the intestine) or to placebo. A further sub-division of patients was made into those that had signs of inflammation with an increased LBP at baseline, and those with normal LBP. The patients with increased LBP had more severe signs of derangement in their haemodynamic evaluations; these included decreased blood pressure (BP) and higher pulse rate leading to a hyperdynamic circulation with increased cardiac output. Hepatic venous catheterisation was performed to measure HVPG. In patients with a high LBP randomised to norfloxacin, 4 weeks’ treatment had a beneficial effect on BP and systemic vascular resistance when compared with baseline measurements. However, no effect was seen on HVPG in this study. Norfloxacin demonstrated a clear effect on LBP (a marker of bacterial translocation) whereas no effect was seen with placebo (Figure 4).

The results of this study concur with those of a randomised crossover trial\(^8\) in which patients with cirrhosis were randomised to placebo or norfloxacin for 28 days, and then crossed over to the treatment not previously received. A clear effect was seen on systemic vascular resistance, but no effect was seen on HVPG in patients receiving norfloxacin. Furthermore, the study evaluated blood flow and found that BP increased and systemic vascular resistance increased in patients receiving norfloxacin. These studies demonstrate that treatment with norfloxacin generates a significant beneficial change in systemic haemodynamic parameters, but not in splanchnic haemodynamic parameters.

Although the non-absorbable antibiotic norfloxacin does not appear to have an effect on splanchnic haemodynamics, recent studies indicate that rifaximin might reduce HVPG.\(^9\) Patients were given rifaximin 1200 mg daily for 29 days, HVPG was measured at baseline and at day 29. The results showed a decrease in HVPG. However, this study was uncontrolled and it is unclear whether this limitation skewed the results, or whether there is a true treatment effect. Unlike norfloxacin, rifaximin has a broader microbial spectrum and has an effect on Gram-positive bacteria.
In an uncontrolled study, rifaximin has been shown to improve systemic haemodynamics and renal function in patients with alcohol-related cirrhosis and ascites. In an open label study, 15 patients with ascites and Child-Pugh B or C cirrhosis received 1200 mg of rifaximin daily for 4 weeks. Haemodynamic parameters, renal function tests and measurement of inflammation markers were measured at baseline and at day 28. The results showed that rifaximin had a positive effect on the systemic haemodynamic parameters (Table 1). This study indicates that rifaximin has an effect on surrogate markers by decreases in cytokine levels, (which are the signals of bacterial translocation), and on haemodynamics.

None of the studies of the effects of gut decontamination in cirrhosis are double-blind or placebo controlled. However, it appears from the evidence available that gut decontamination prevents relapse of HE, improves systemic haemodynamics by an increase in BP and a decrease in cardiac output, decreases immune activation, and may improve renal function. The effect of gut decontamination on splanchnic haemodynamics is yet to be proven.

A randomised, double-blind, placebo controlled study is in progress. This study is investigating intestinal decontamination with rifaximin in cirrhotic patients with ascites, and assessing the effects on haemodynamic and inflammatory factors. The aim of the study is to stop progression of liver disease by inhibiting bacterial translocation via bowel decontamination. The outcome measures are to reduce portal hypertension, diminish vasodilation, increase glomerular filtration rate, normalise inflammation and decrease risk of infection (Figure 5).

There is a need for a large scale, randomised, double-blind, placebo controlled study, addressing gut decontamination in decompensated cirrhosis. The clinical endpoints should include mortality, the risks of variceal bleeding, infections (especially spontaneous bacterial peritonitis), and hepatorenal syndrome.

### Table 1. Systemic haemodynamics, endogenous vasoactive systems, renal function, body weight, and inflammatory markers pre and post treatment with rifaximin.

<table>
<thead>
<tr>
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<th>Baseline</th>
<th>Rifaximin</th>
<th>P</th>
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<tr>
<td><strong>Systemic haemodynamics and endogenous vasoactive systems</strong></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Mean arterial pressure, mm Hg</td>
<td>90 (81.6–101.6)</td>
<td>92 (86.3–106.6)</td>
<td>0.05</td>
</tr>
<tr>
<td>Cardiac output, L/min</td>
<td>6.01 (4.77–8.76)</td>
<td>5.76 (4.1–8.04)</td>
<td>0.02</td>
</tr>
<tr>
<td>Systemic vascular resistance, dynes * sec * cm⁻⁵</td>
<td>1500 (960–1815)</td>
<td>1619 (1163–2243)</td>
<td>0.01</td>
</tr>
<tr>
<td>Plasma renin activity, ng/mL/h</td>
<td>6.85 (117–16.98)</td>
<td>5.59 (106–14.6)</td>
<td>0.02</td>
</tr>
<tr>
<td>Plasma aldosterone level, pg/mL</td>
<td>637.2 (311–1323)</td>
<td>495.2 (231.6–1245)</td>
<td>0.06</td>
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<tr>
<td><strong>Renal function and body weight</strong></td>
<td></td>
<td></td>
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<tr>
<td>Serum creatinine level, mg/dL</td>
<td>0.9 (0.8–1.2)</td>
<td>0.9 (0.8–1.1)</td>
<td>0.6</td>
</tr>
<tr>
<td>Serum urea level, mg/dL</td>
<td>34 (18–50)</td>
<td>33 (20–46)</td>
<td>0.2</td>
</tr>
<tr>
<td>Glomerular filtration rate, mL/min</td>
<td>65 (40–84)</td>
<td>66 (39–91)</td>
<td>0.006</td>
</tr>
<tr>
<td>Urinary sodium level, mmol/d</td>
<td>47.5 (22–100)</td>
<td>48.6 (25.4–106)</td>
<td>0.03</td>
</tr>
<tr>
<td>Weight, kg</td>
<td>72.8 (60.2–81.5)</td>
<td>72 (59.6–81)</td>
<td>0.1</td>
</tr>
<tr>
<td><strong>Circulating endotoxin and cytokine concentrations</strong></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Plasma endotoxin, EU/mL</td>
<td>1.07 (0.65–10)</td>
<td>0.56 (0.37–5.23)</td>
<td>0.005</td>
</tr>
<tr>
<td>Serum interleukin-6, pg/mL</td>
<td>18.82 (3.97–54.85)</td>
<td>12.49 (1.87–30.11)</td>
<td>0.01</td>
</tr>
<tr>
<td>Serum tumour necrosis factor-, pg/mL</td>
<td>5.51 (3.84–10.47)</td>
<td>3.78 (2.34–7.26)</td>
<td>0.02</td>
</tr>
</tbody>
</table>

Note: Data are expressed as the median and range. Kalambokis GN et al.¹⁰
Treatment of Hepatic Encephalopathy: Targeting the Gut-Liver-Brain Axis. Gut Decontamination and HE. The More Things Change, the More They Remain the Same

Professor Rajiv Jalan

There have been significant advances in the treatment of HE, predominantly involving targeting the gut-liver-brain axis and gut decontamination. The current treatment concepts are illustrated by the following case study.

A 53 year-old male Afro-Caribbean lawyer was transferred from a neurology hospital where he had been an inpatient for 2 months. The patient had a history of haematemesis and melaena, spastic paraparesis, and a previous history of alcohol abuse and mildly abnormal liver function tests. On admission to the Intensive Care Unit (ICU) the patient had fluctuating levels of consciousness with a Glasgow Coma Scale (GCS) of 4-5. Immediate management included resuscitation, transfusion and an upper gastrointestinal endoscopy. This showed Grade 3 oesophageal varices which were treated with banding and achieved good control of the bleeding. The patient was treated with terlipressin for 3 days, antibiotics, thiamine and lactulose. The initial diagnosis was cirrhosis and oesophageal varices (a complication of the disease). The patient had been previously fit and well, and had a flourishing law practice until 2006. He retired prematurely in July 2009 due to progressive leg weakness that started in 2006, and became wheelchair bound in 2011. The patient had undergone extensive investigations at the specialist neurology hospital and a working diagnosis of spastic paraplegia of unknown origin was made.

Following admission to the ICU, neurological examination on day 4 revealed that he was still unresponsive with a GCS of 4-5, had Grade 3-4 HE and had very little power in both legs. Abdominal examination showed mild
hepatosplenomegaly, no ascites and no peripheral stigmata of liver disease. Despite the very large ‘hit’ the patient had experienced, his chest was clear and lung function was normal, heart rate was 78 beats/minute, BP 143/78 mmHg, and kidney function was acceptable (creatinine 78 μmol/L). Liver investigations showed mildly elevated bilirubin (37 μmol/L which equates to 2 mg/dL), and his creatinine was normal (65 μmol/L which equates to between 0.7 and 0.8 mg/dL). The model for end-stage liver disease (MELD) score was 10; indicating very mild early chronic liver disease with portal hypertension. All the viral serology results were negative, except Hep B sAg and Hep B eAg (due to previous exposure to Hepatitis B), but HBV was negative. These findings suggested underlying alcoholic cirrhosis which was well compensated. Liver ultrasound demonstrated a normal liver, splenomegaly (14.5 cm) and a trace of pelvic ascites. MRI of the liver showed no evidence of chronic liver disease; the biliary system was normal and a normal total liver volume of 1,382 mL. Transjugular liver biopsy demonstrated evidence of alcoholic cirrhosis. Consequently, a diagnosis of well compensated alcoholic cirrhosis with portal hypertension was made. The patient had a neurological deficit; he was comatose and had spastic paraplegia which was initially thought to be unrelated. On day 5, the results of an ammonia test were 145 μmol/L (normal is <40), this indicated that the patient’s condition was due to HE. Therefore, lactulose was increased to 15 mL three times a day (tds) and treatment with rifaximin 550 mg twice a day (bd) was commenced. GCS remained at 4–5 for a further 5 days and the patient developed diarrhoea (culture and sensitivity was negative) which led to rifaximin and lactulose being stopped.

A CT scan of the abdomen showed a large spontaneous portacaval shunt emanating from the left renal vein. Brain MRI showed an increased pallidal hyperintensity on T1 imaging, and MR spectroscopy showed an elevated glutamine level. The patient showed evidence of brain dysfunction with hyperammonemia, increased brain glutamine, spastic paraparesis, and pallidal hyperintensity indicating that the syndrome may be a result of brain dysfunction precipitated by cirrhosis. A diagnosis of severe HE was made and embolisation of the patient’s shunt was considered. Embolisation is a useful treatment for HE and a MELD score of ≤11 predicts a good response to therapy.12 However, this patient had portal hypertension and a recent variceal bleed and if the shunt were to be blocked, the portal hypertension would become worse. Consequently, the treatment strategy was to insert a small diameter Transjugular Intrahepatic PortoSystemic Shunt (TIPSS) and through the TIPSS, embolise the shunt and control the shunt. The patient remained in ICU and gradually woke up over the next 6 days.

On the ward between days 18 and 28 it became apparent that the patient had persistent Grade 2 HE and remained significantly hyperammonemic (75–90 μmol/L). Treatment with lactulose 15 mL tds and rifaximin 550 mg bd was restarted. Recommencement of rifaximin was in line with findings by Bass et al.13 that showed that patients discharged from hospital who had been treated with rifaximin had fewer HE relapses and hospital admissions. The study (Figure 6) found a 50% relative risk reduction and a 9% absolute risk reduction in patients treated with rifaximin (hazard ratio 0.50 [95% CI, 0.29–0.87] p=0.01).

The patient was discharged from hospital to a rehabilitation centre on day 42. His ammonia levels had improved (35–42 μmol/L), neurologically the GCS was 15, and he remained mildly encephalopathic (Grade 1). The patient was followed-up closely; the treatment strategy was to continue lactulose and rifaximin, and if no improvement was seen liver transplantation would be considered. Weissenborn et al.14 showed that patients who have large portacaval shunts can develop HE and hepatic myelopathy, which may be reversible with liver transplantation. In this patient, liver transplantation was indicated for hepatic myelopathy because it was thought that the spastic paraparesis may be related to HE.

On review 8 months later, the patient had no further hospital admissions for HE and continued on treatment with lactulose and rifaximin, which he tolerated quite well. Ammonia levels had varied between 30 and 40 μmol/L, and there had been a gradual improvement in the power of both lower limbs with physiotherapy, nutritional support and drug treatment. The patient had responded to multiple level interventions, not only from a neurological state but from spastic paraparesis as well. The final diagnosis was alcoholic cirrhosis with portal hypertension, HE and hepatic myelopathy.
Knowledge of the pathogenesis of cirrhosis may not have changed. The role of the gut was first described in 1893 by Necki et al.15-17 who made the following observations: ammonia was coming from the gut and portal ammonia was greater than arterial ammonia (the current treatment for this is lactulose and rifaximin); ammonia concentration is increased in the muscle and kidney (the proposed treatment today is ornithine phenylacetate); ammonia concentration is raised in the gastric mucosa even in a fasting state (now there may be a role for H. Pylori eradication therapy); ammonia was raised in the kidney of the portacaval shunted dog, and urinary ammonia excretion increased after a meat-meal or ammonia administration (the current treatments are volume expansion, ornithine phenylacetate, gut lavage, and rifaximin). These observations raise the question of how far we have actually progressed; the gut-liver axis was really described more than 120 years ago.

Panel Discussion: The Role of Non-Absorbed Antibiotics in Gut Decontamination

GBT plays an important role in HE. Key mediators in GBT that drive this include the presence of bacteria in the circulation or tissues, as well as bacterial products that stimulate an inflammatory reaction (e.g. bacterial DNA). Furthermore, it is not the response itself that should be targeted but the location where it originated – the gut-liver axis.

There is a need for more controlled trials on GBT and gut microbiota, with patients with clear signs of bacterial translocation as the focus. Parameters that prove that bacterial translocation is present (e.g. an increased LBP or a high level of cytokines) should be measured. Targeting GBT requires the use of a very broad spectrum, non-absorbable antibiotic, such as rifaximin, which should be tested in this context. Rifaximin has shown efficacy in bowel decontamination with very few side-effects.

A clear target for gut decontamination is a decompensated cirrhotic patient with ascites who has signs of inflammation that are attributable to GBT. In this group of patients, gut decontamination is likely to improve survival. It has been shown that in spontaneous bacterial translocation, norfloxacin would
not be sufficient to eradicate the bacteria and overgrowth in the intestine. Consequently, a drug is required to target both Gram-positive and Gram-negative bacteria: rifaximin has a non-selective, broad antibiotic spectrum and acts on both Gram-positive and Gram-negative bacteria.

The case study demonstrated that the pathology of hepatic myelopathy is a loss of the anterior horn cells; producing a very severe form of HE. There are two types of hepatic myelopathy: a reversible form, and an irreversible form. It is difficult to distinguish the types of hepatic myelopathy in a patient, but both types result from large, long-standing porto-systemic shunt. Liver transplantation is a treatment option for these patients but not all respond and the type of hepatic myelopathy present in responders is unknown.

In the treatment of HE, liver transplantation may not be the first treatment choice – particularly if the chance of recovery is not evident. There are several different treatment options and indicators that can be considered. In comatose patients with liver disease who have clinical signs of HE, ammonia is a useful indicator of the severity of encephalopathy. Accurate measurement can indicate a target for intervention, and in the context of acute liver failure, ammonia provides a measurement to follow as an endpoint to intervention. However, the measurement of ammonia is difficult and must be performed using a validated technique.

The insertion of TIPSS in patients with long-term HE has been suggested to be contraindicated. The case study patient illustrated a treatment strategy when limited options are available. The patient had experienced a variceal bleed and there was a high chance of re-bleeding if the spontaneous large portosystemic shunt was occluded for the treatment of HE. Therefore, a small 8 mm shunt was inserted that could be stretched if the patient bled, providing an opportunity to rescue the patient. Small shunts are at risk of closure therefore a low level of anticoagulation is required.

Faecal transplantation is a therapeutic option that has been suggested in patients with *Clostridium difficile* infection. However, in patients who are immunosuppressed due to cirrhosis, this may not prevent bacterial overgrowth in the small intestine, and as a result is unlikely to benefit many patients.

Clinical experience demonstrates the important role of non-absorbable antibiotics in gut decontamination in patients with HE.

REFERENCES

11. ClinicalTrials.gov Identifier: NCT01769040