

EASL Clinical Practice Guidelines for the management of patients with decompensated cirrhosis[☆]

European Association for the Study of the Liver^{*}

Summary

The natural history of cirrhosis is characterised by an asymptomatic compensated phase followed by a decompensated phase, marked by the development of overt clinical signs, the most frequent of which are ascites, bleeding, encephalopathy, and jaundice. The following Clinical Practice Guidelines (CPGs) represent the first CPGs on the management of decompensated cirrhosis. In this context, the panel of experts, having emphasised the importance of initiating aetiologic treatment for any degree of hepatic disease at the earliest possible stage, extended its work to all the complications of cirrhosis, which had not been covered by the European Association for the Study of the Liver guidelines, namely: ascites, refractory ascites, hyponatremia, gastrointestinal bleeding, bacterial infections, acute kidney injury, hepatorenal syndrome, acute-on-chronic liver failure, relative adrenal failure, cirrhotic cardiomyopathy, hepatopulmonary syndrome, and porto-pulmonary hypertension. The panel of experts, produced these GPGs using evidence from PubMed and Cochrane database searches providing up to date guidance on the management of decompensated cirrhosis with the only purpose of improving clinical practice.

© 2018 European Association for the Study of the Liver. Published by Elsevier B.V. All rights reserved.

Introduction

When the panel of experts nominated by the European Association for the Study of the Liver (EASL) governing board began work to update the Clinical Practice Guidelines (CPGs) on ascites, spontaneous bacterial peritonitis (SBP), and hepatorenal syndrome (HRS),¹ it became obvious that all other complications of decompensated cirrhosis had to be covered. Within this framework, a formal definition of decompensated cirrhosis was sought. The natural history of cirrhosis is characterised by a silent, asymptomatic course until increasing portal pressure and worsening liver function produce a clinical phenotype. In the asymptomatic phase of the disease, usually referred to as compensated cirrhosis, patients may have a good quality of life, and the disease may progress undetected for several years. Decompensation is marked by the development of overt clinical

signs, the most frequent of which are ascites, bleeding, encephalopathy, and jaundice. Following the first appearance of any of these, the disease usually progresses more rapidly towards death or liver transplantation (LT). This phase of the disease has been designated “decompensated cirrhosis”.² Progression of the decompensated disease may be further accelerated by the development of other complications such as rebleeding, acute kidney injury (AKI), with or without the features of HRS, hepato-pulmonary syndrome (HPS), portopulmonary hypertension (PPHT), cirrhotic cardiomyopathy (CCM), and bacterial infections. Indeed, the development of bacterial infections as well as hepatocellular carcinoma may accelerate the course of the disease at any stage, but especially in decompensated cirrhosis.³ Having defined the potential field of action, and having emphasised the importance of initiating aetiologic treatment for any degree of hepatic disease at the earliest possible stage, the panel decided to extend the work to all those complications of cirrhosis which have not yet been covered by EASL guidelines, namely: gastrointestinal (GI) bleeding, bacterial infections other than SBP, acute-on-chronic liver failure (ACLF), adrenal failure, HPS, PPHT and CCM. In doing so, we have had to deal with the recommendations regularly proposed by very well recognised international expert groups who have worked in the field of GI bleeding or ascites and ascites-related complications for many years. Given their extreme importance in clinical practice, only specific aspects of their recommendations were further developed in an attempt to give a more integrated view of the pathophysiology and management of patients with decompensated cirrhosis. Thus, this document can no longer be considered an update of earlier guidelines, but rather the first CPG on the management of decompensated cirrhosis with the sole purpose of improving clinical practice.

Guidelines development process

A panel of hepatologists with a great interest in decompensated cirrhosis, approved by the EASL Governing Board, wrote and discussed this CPG between March 2017 and February 2018. The guidelines were independently peer reviewed, and all contributors to the CPG disclosed their conflicts of interest by means of a disclosure form provided by the EASL Office prior to work commencing. The EASL Ethics Committee reviewed the composition of the panel to eliminate the potential for real or perceived bias. The CPG panel conflict of interests are declared in this submission. These guidelines have been produced using evidence from PubMed and Cochrane database searches before 27 March 2018. Tables describing

[☆] **Clinical Practice Guideline Panel:** Paolo Angeli (Chair), Mauro Bernardi (Governing Board representative), Cándid Villanueva, Claire Francoz, Rajeshwar P. Mookerjee, Jonel Trebicka, Aleksander Krag, Wim Laleman, Pere Gines

^{*} Corresponding author. Address: European Association for the Study of the Liver (EASL), The EASL Building – Home of Hepatology, 7 rue Daubin, CH 1203 Geneva, Switzerland. Tel.: +41 (0) 22 807 03 60; fax: +41 (0) 22 328 07 24.

E-mail address: easloffice@easloffice.eu.



ELSEVIER

Journal of Hepatology 2018 vol. xxx | xxx–xxx

Please cite this article in press as: The European Association for the Study of the Liver. EASL Clinical Practice Guidelines for the management of patients with decompensated cirrhosis. J Hepatol (2018), <https://doi.org/10.1016/j.jhep.2018.03.024>

Table 1. Level of Evidence and Grade of Recommendations.

Level of evidence	
I	Randomised, controlled trials
II-1	Controlled trials without randomisation
II-2	Cohort and case-control analytical studies
II-3	Multiple time series, dramatic uncontrolled experiments
III	Opinions of respected authorities, descriptive epidemiology
Grade of recommendations	
1	Strong recommendation: Factors influencing the strength of the recommendation included the quality of the evidence, presumed patient-important outcomes, and cost
2	Weaker recommendation: Variability in preferences and values, or more uncertainty: more likely a weak recommendation is warranted. Recommendation is made with less certainty: higher cost or resource consumption

the rationale behind the levels of evidence and of recommendations are provided (Table 1).

Pathophysiology of decompensated cirrhosis

The transition from compensated asymptomatic cirrhosis to decompensated cirrhosis occurs at a rate of about 5% to 7% per year.⁴ Once decompensation has occurred, cirrhosis becomes a systemic disease, with multi-organ/system dysfunction.⁵ At this stage, patients become highly susceptible to bacterial infections because of complex cirrhosis-associated immune dysfunction, which involves both innate and acquired immunity.⁶ In turn, patients with bacterial infections are burdened by severe morbidity, up to ACLF, and high mortality.^{6,7} Because of these events, decompensation represents a prognostic watershed, as the median survival drops from more than 12 years for compensated cirrhosis to about two years for decompensated cirrhosis.⁴ For decades the clinical manifestations of decompensated cirrhosis have been seen as the consequence of a haemodynamic disturbance, the hyperdynamic circulatory syndrome, ascribable to peripheral arterial vasodilation that mainly occurs in the splanchnic circulatory area. The extent of such vasodilation is to endanger effective volaemia, ultimately leading to peripheral organ hypoperfusion, the kidney being most affected.⁸ Indeed, reduced effective volaemia brings about the activation of vasoconstrictor and water and sodium retaining mechanisms, such as the renin-angiotensin-aldosterone (RAAS), sympathetic nervous system and arginine-vasopressin secretion. This explains some of the cardinal features of decompensated cirrhosis, such as renal retention of sodium and water leading to ascites formation and HRS. Other manifestations attributable to haemodynamic abnormalities include HPS, increased susceptibility to shock, and a reduced cardiovascular responsiveness to physiological and pharmacological vasoconstrictor stimuli. Subsequent studies have highlighted that a cardiac dysfunction, due to CCM,⁹ is also involved in the pathogenesis of effective hypovolaemia.¹⁰ This occurs particularly in the most advanced stages of decompensation, when such an abnormality prevents cardiac output from increasing enough to comply with the needs of systemic circulation. Although the molecular mechanisms responsible for arterial vasodilation, consisting of an enhanced endothelial production of vasodilating substances, such as nitric oxide, carbon monoxide, prostacyclin and endocannabinoids have been convincingly demonstrated,¹¹ the primary causes of such abnormalities remained somewhat obscure until it became clear that patients with advanced cirrhosis present a state of chronic inflammation, as witnessed by increased circulating levels of pro-inflammatory cytokines and chemokines.¹² This is likely caused by the systemic spread of bacteria and bacterial products, called pathogen associated molecular patterns (PAMPs), as a

result of an abnormal bacterial translocation (BT). Changes in the microbiome and increased intestinal permeability account for this phenomenon. A similar role is likely played by other molecules, called danger associated molecular patterns (DAMPs), released by the diseased liver because of local inflammation and cell apoptosis and necrosis. Both PAMPs and DAMPs bind with innate recognition receptors of immune cells that, once activated, produce and release pro-inflammatory molecules, along with reactive oxygen and nitrogen species. This cascade of events contributes to the development of circulatory dysfunction and, along with it, directly favours the development of multi-organ dysfunction and failure (Fig. 1).⁵ Current strategies for prophylaxis and treatment of decompensation and organ failure in cirrhosis rely on measures aimed to prevent or improve the outcome of each complication, that is renal sodium retention leading to ascites formation, ammonia production in hepatic encephalopathy, effective hypovolaemia after large-volume paracentesis (LVP) or during HRS, renal dysfunction induced by SBP, and intestinal dysbiosis or bacterial overgrowth in patients predisposed to develop infections. All these strategies will be discussed in these CPGs. However, the improved knowledge of the pathophysiological background of decompensated cirrhosis now offers the opportunity for more comprehensive therapeutic and prophylactic approaches to disease management. Indeed, besides treating the underlying aetiologic factor(s), whenever possible, mechanistic approaches to counteract key pathophysiological mechanisms may prevent or delay disease progression and the incidence of complications and multi-organ dysfunction, thus improving patient survival and quality of life, as well as reducing the economic burden of the disease.

Management of decompensated cirrhosis

Ideally, the strategy of management of patients with decompensated cirrhosis should be based on preventing cirrhosis progression (*i.e.* further decompensation) rather than treating complications as they occur. The ultimate treatment for decompensated cirrhosis would be one that targets primarily the pathological alterations within the liver with the aim of restoring the integrity of liver architecture by suppressing inflammation, causing fibrosis regression, regularising the portal and arterial circulation, and normalising cell number and function. Unfortunately, such a treatment does not exist at present. Several antifibrotic or anti-inflammatory drugs have shown promise in experimental models of chronic liver diseases, but no treatment has yet been translated into clinical practice.¹³ Meanwhile, the overall management of decompensated cirrhosis can be addressed using two approaches. The first approach is the suppression of the aetiological factor(s) that has caused liver inflammation and cirrhosis development, whereas the second

approach is based on targeting key factors of pathogenesis of cirrhosis decompensation and progression.

Effects of suppression of aetiological factor on outcome of decompensated cirrhosis

Removal of the aetiological factor(s) causing liver injury is an important cornerstone in the management of cirrhosis. This approach is clearly effective in preventing decompensation and improving outcome in patients with compensated cirrhosis. However, results in patients with decompensated cirrhosis are less efficacious and probably depend, among other factors, on the actual status of liver disease at the time of removing the aetiological factor of liver injury. For example, although in some patients with decompensated alcoholic cirrhosis suppression of alcohol consumption is associated with progressive “re-compensation” of cirrhosis and excellent long-term outcome, in other patients alcoholic cirrhosis progresses despite stopping alcohol intake.^{14,15} Likewise, in patients with cirrhosis due to hepatitis B virus (HBV) infection, treatment with antiviral agents is associated with improved outcome in some, but not all patients.¹⁶ Moreover, treatment of patients with decompensated cirrhosis due to hepatitis C virus infection with direct antiviral agents is associated with beneficial effects in liver function and portal hypertension and likely improves outcome, but these effects are unfortunately not generalisable to all patients treated.^{17,18} The beneficial effects of removing responsible factors in other aetiologies of decompensated cirrhosis are less clear, perhaps with the exception of autoimmune hepatitis.

Effects of targeting key pathogenic events in prevention of cirrhosis progression

Several strategies have been evaluated to prevent disease progression in patients with decompensated cirrhosis, including i) targeting microbiome abnormalities and BT, to improve gut-liver axis; ii) improving the disturbed circulatory function; iii) treating the inflammatory state; and iv) targeting portal hypertension.

Administration of rifaximin has been shown to reduce the risk of development of several complications of cirrhosis besides hepatic encephalopathy in retrospective studies and small case series.¹⁹ Nonetheless, data from prospective randomised double-blind studies are lacking. In patients with decompensated cirrhosis, treatment with norfloxacin reduces the risk of SBP and HRS,^{20,21} but its use is hampered by the possibility of increased risk of infection by resistant bacteria. The potential effectiveness of improving circulatory and kidney function by long-term administration of albumin to patients with decompensated cirrhosis has been explored in two recent randomised controlled trials (RCTs), both published in abstract form, with contradictory findings.^{22,23} The discrepant findings may be related to different doses of albumin used and/or heterogeneity in the study population. Further studies are needed to find out whether long-term albumin administration is efficacious in decompensated cirrhosis. Interestingly, treatment with statins, through their pleiotropic effects, has been shown to reduce portal hypertension and improve survival in patients with advanced cirrhosis.^{24,25} These remarkable effects require validation in future studies. Another potential therapeutic strategy in the prevention of decompensation may be anticoagulation. Indeed, in a small RCT, a 12-month course of enoxaparin was safe and effective in preventing portal vein thrombosis (PVT) in patients

with cirrhosis and a Child-Pugh scores of 7–10. In addition, enoxaparin appeared to delay the occurrence of hepatic decompensation and to improve survival suggesting that both PVT and decompensation may be related to a worsening of portal hypertension and the consequent progressive damage of the intestinal mucosal barrier.²⁶ From the same perspective, two other strategies should be considered. In 2010, it was shown that pentoxifylline treatment significantly reduced the risk of liver-related complications compared to placebo in an RCT of patients with advanced cirrhosis. The prevention of these complications, which included bacterial infections, renal failure, and hepatic encephalopathy was probably related to the fact that pentoxifylline prevents intestinal BT and the consequent development of systemic inflammation.²⁷ Finally, some investigations have shown that treatment with propranolol is not only effective in reducing portal hypertension and the consequent the risk of variceal bleeding but also in decreasing the risk of other complications of cirrhosis related to portal hypertension, such as ascites, HRS, SBP, and hepatic encephalopathy.²⁸ These effects occur specifically in patients who respond to propranolol treatment by markedly decreasing portal pressure, emphasising the strong relationship between pressure and complications of cirrhosis. Nevertheless, in these studies most of patients had compensated cirrhosis. Therefore, studies should be performed in the group of patients with decompensated cirrhosis with the objective of assessing these beneficial effects in cirrhosis progression.

Recommendations

- In patients with decompensated cirrhosis, the aetiological factor, should be removed, particularly alcohol consumption and hepatitis B or C virus infection as this strategy is associated with decreased risk of decompensation and increased survival (**II-2,1**).
- Strategies based on targeting abnormalities in gut-liver axis by antibiotic administration (*i.e.* rifaximin), improving the disturbed systemic circulatory function (*i.e.* long-term albumin administration), decreasing the inflammatory state (*i.e.* statins), and reducing portal hypertension (*i.e.* beta-blockers) have shown potential benefit to decrease cirrhosis progression in patients with decompensated cirrhosis. However, further clinical research is needed with these strategies to confirm their safety and potential benefits as therapeutic approaches with the aim of preventing cirrhosis progression in decompensated patients.

Management of specific complications of decompensated cirrhosis

Ascites

Ascites is the most common cause of decompensation in cirrhosis, as 5% to 10% of patients with compensated cirrhosis per year develop this complication.²⁹ The mainstay of ascites formation is renal sodium retention due to the activation of sodium retaining systems, such as the renin-angiotensin-aldosterone system (RAAS) and sympathetic nervous system. The resulting positive fluid balance ultimately leads to extracellular fluid volume expansion. Reduced effective volemia secondary to splanchnic arterial vasodilation is a main determinant of these alterations,⁸ but renal function abnormalities induced by systemic inflammation

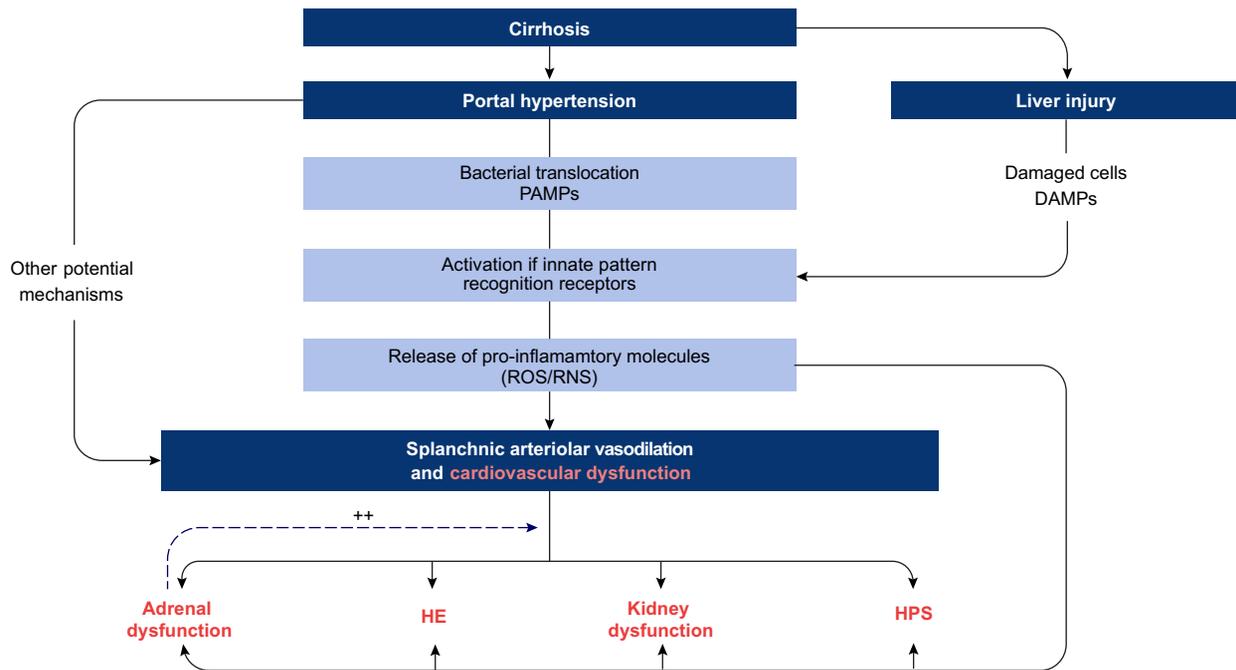


Fig. 1. The new theory on the development of complications and organ failure/s in patients with cirrhosis (adapted from Ref. 5). DAMP, damage-associated molecular pattern; HE, hepatic encephalopathy; HPS, hepatopulmonary syndrome; PAMP, pathogen-associated molecular pattern; RNS, reactive nitrogen species; ROS, reactive oxygen species.

also play a role, especially in the most advanced stages of cirrhosis.⁵ Portal hypertension also contributes³⁰ by acting as a compartmentalising factor of the expanded extracellular fluid volume.

The occurrence of ascites impairs patient working and social life, often leads to hospitalisation, requires chronic treatment and is a direct cause of further complications, such as SBP, restrictive ventilatory dysfunction, or abdominal hernias. The appearance of ascites heralds a poor prognosis, as the five-year survival drops from about 80% in compensated patients to about 30% in patients with decompensated cirrhosis and ascites.⁴

Uncomplicated ascites

Evaluation of patients with ascites

Cirrhosis is the main cause of ascites in the Western world, being responsible for about 80% of cases. Malignancy, heart failure, tuberculosis, pancreatic disease, or other rarer diseases account for the remaining cases. Initial patient evaluation should include history, physical examination, abdominal ultrasound, and laboratory assessment of liver and renal functions, serum and urine electrolytes, as well as an analysis of the ascitic fluid.

Diagnosis of ascites

Ascites can be graded from 1 to 3 according to the amount of fluid in the abdominal cavity³¹ (Table 2). The ascites that recurs at least on three occasions within a 12-month period despite dietary sodium restriction and adequate diuretic dosage is defined as recidivant.³²

Diagnostic paracentesis is indicated in all patients with new onset of grade 2 or 3 ascites and in those admitted to the hospital for any complication of cirrhosis.^{31,32} Manual or automated neutrophil count, total protein and albumin concentration, and culture should be always assessed. A neutrophil count above 250 cells/ μ l denotes SBP.³³ A total protein concentration <1.5 g/dl is generally considered a risk factor for SBP, although there are conflicting data.^{33,34} Ascitic fluid culture requires the

bedside inoculation of at least 10 ml into blood culture bottles to enhance its sensitivity.³⁵ The calculation of serum-ascites albumin gradient (SAAG) may be useful when the cause of ascites is not immediately evident, as SAAG \geq 1.1 g/dl indicates that portal hypertension is involved in ascites formation with an accuracy of about 97%.³⁶ Other tests, such as amylase, cytology, or culture for mycobacteria should be guided by clinical presentation. Ascitic cholesterol determination followed by cytology and carcinoembryonic antigen (CEA) determination in samples where cholesterol concentration exceeds 45 mg/dl appears to be a cost-effective method for the differential diagnosis between malignancy-related and non-malignant ascites.³⁷

Recommendations

- A diagnostic paracentesis is recommended in all patients with new onset grade 2 or 3 ascites, or in those hospitalised for worsening of ascites or any complication of cirrhosis (II-2;1).
- Neutrophil count and culture of ascitic fluid culture (bedside inoculation blood culture bottles with 10 ml fluid each) should be performed to exclude bacterial peritonitis. A neutrophil count above 250 cells/ μ l is required to diagnose SBP (II-2;1).
- Ascitic total protein concentration should be performed to identify patients at higher risk of developing SBP (II-2;1).
- The SAAG should be calculated when the cause of ascites is not immediately evident, and/or when conditions other than cirrhosis are suspected (II-2;1).
- Cytology should be performed to differentiate malignancy-related from non-malignant ascites (II-2;1).

Prognosis of patients with ascites

The development of ascites in patients with cirrhosis is associated with a poor prognosis, as their one and two-year mortality is about 40 and 50%, respectively.¹ Thus, patients with ascites should generally be considered for referral for LT. Hyponatraemia, low arterial pressure, glomerular filtration rate (GFR) and low renal sodium excretion are independent predictors of mortality in cirrhosis with ascites.³⁸ As these parameters are not included in the Child-Pugh score, and only serum creatinine (SCr), which overestimates GFR in cirrhosis,³⁹ is included in the model for end-stage liver disease (MELD) score, the most commonly used prognostic scores can underestimate the mortality risk in patients with ascites. Modifications of the MELD score, such as the MELD-Na and MELD-Ascites scores have only partially overcome this limitation.⁴⁰ Thus, patients with ascites may not receive adequate priority in transplant lists, and improved methods to assess prognosis in these patients are needed. A prognostic score able to identify patients with low MELD score (<18) at high risk of 12-month adverse outcome has recently been proposed, but it still has limited application.⁴¹

Recommendation

- Since the development of grade 2 or 3 ascites in patients with cirrhosis is associated with reduced survival, LT should be considered as a potential treatment option (II-2;1).

Management of uncomplicated ascites

Ascites is uncomplicated when it is not infected, refractory or associated with HRS.^{31,32}

Grade 1 or mild ascites. No data on the evolution of grade 1 ascites are available, nor it is known whether its treatment modifies its natural history.

Grade 2 or moderate ascites. Patients who develop grade 2 ascites do not require hospitalisation, unless other complications are present. They have a positive sodium balance, which can be corrected by reducing the dietary sodium intake and increasing renal sodium excretion with diuretics. Although upright posture favours renal sodium reabsorption⁴² and attenuates the response to diuretics,⁴³ there is no evidence that a prolonged maintenance of the supine position eases the treatment of ascites.

Sodium restriction. The prophylactic use of salt restriction in patients who never had ascites is not supported by evidence. Dietary sodium restriction can lead to the resolution of ascites in about 10% of patients,⁴⁴ especially in those with the first episode of ascites. A clear advantage from the use of low-sodium diets associated with diuretics has not emerged from clinical trials comparing different dietary regimens.^{44,45} Extreme sodium restriction favours the development of diuretic-induced hyponatraemia and renal failure.⁴⁶ Moreover, even moderate sodium restriction, when not prescribed with an adequate educational programme, is often associated with reduced calorie intake,⁴⁷ and may impair nutritional status. The current opinion is that dietary sodium should only be moderately restricted (80–120 mmol/day), mainly to avoid excess salt intake.

Recommendations

- A moderate restriction of sodium intake (80–120 mmol/day, corresponding to 4.6–6.9 g of salt) is recommended in patients with moderate, uncomplicated ascites (I;1). This is generally equivalent to a no added salt diet with avoidance of pre-prepared meals. Adequate nutritional education of patients on how to manage dietary sodium is also recommended (II-2;1).
- Diets with a very low sodium content (<40 mmol/day) should be avoided, as they favour diuretic-induced complications and can endanger a patient's nutritional status (II-2;1).
- Prolonged bed rest cannot be recommended because there is insufficient evidence that it is beneficial in the treatment of ascites (III;1).

Diuretics. Neither diuretics nor LVP are associated with a survival benefit because they act downstream of the pathophysiological cascade, being symptomatic therapies. The negative fluid balance induced by diuretics should not lead to a body weight loss exceeding 0.5 kg/day in patients without peripheral oedema and 1 kg/day in the presence of peripheral oedema to avoid plasma volume contraction, ultimately leading to renal failure and hyponatraemia.⁴⁸ Since secondary hyperaldosteronism plays a pivotal role in the renal sodium retention in patients with cirrhosis,^{49,50} anti-mineralocorticoid drugs (spironolactone, canrenone or K-canrenoate) represent a mainstay in the medical treatment of ascites.⁵⁰ Four hundred mg/day represents the maximal dosage usually recommended.^{31,32} The mechanism of action of anti-mineralocorticoids explains their slow effect. In fact, the activated aldosterone pathway, which involves interaction with a cytosolic receptor and, then, a nuclear receptor, needs to be exhausted before their natriuretic effect arises. Therefore, the dosage of these drugs should not be increased earlier than 72 h. Amiloride, a diuretic acting in the collecting duct, is less effective than anti-mineralocorticoids, and should only be used in patients who develop severe side effects with aldosterone antagonists.⁵¹

Proximal tubular sodium reabsorption promotes renal sodium retention through various mechanisms, such as increased angiotensin II production, sympatho-adrenergic hyperactivity and reduced renal perfusion.⁴⁹ As proximal tubular sodium reabsorption can become relatively prevalent in patients with long-standing ascites,^{52,53} loop diuretics are indicated in this setting. However, they should be combined with but not substituted for anti-mineralocorticoids. Indeed, despite their potent activity, the natriuretic effect of loop diuretics can be completely blunted by unopposed hyperaldosteronism.⁵⁴ Whether diuretic treatment should be initiated with anti-mineralocorticoids alone or should also include a loop diuretic has long been debated. Two studies have addressed this matter providing apparently conflicting results because of differences in patient populations.^{55,56} In both studies, the effects of a diuretic regimen initially consisting of spironolactone or K-canrenoate alone at stepwise increasing

Table 2. Grading of ascites.

Grade 1.	Mild ascites: it is only detectable by ultrasound examination
Grade 2.	Moderate ascites: it is manifest by moderate symmetrical distension of abdomen
Grade 3.	Large or gross ascites: it provokes marked abdominal distension

Clinical Practice Guidelines

dosages (from 100/200 to 400 mg/day), with furosemide added in non-responder patients, were compared with those of the combination of anti-mineralocorticoids with furosemide (from 40 to 160 mg/day) from the beginning of treatment. In one study,⁵⁶ the response rate, the rapidity of ascites mobilisation and the incidence of diuretic-induced complications were similar in both regimens. However, as the sequential treatment required less dose adjustments, it appeared to be more suitable for treating ascites on an outpatient basis. In the other study,⁵⁵ the combined regimen achieved the resolution of ascites in a shorter time, with a lower incidence of side effects, mainly hyperkalemia. Such divergent results likely arose from differences in the patient populations. In one study,⁵⁶ patients with ascites at the first appearance and well preserved renal function prevailed, while, in the other,⁵⁵ most patients had recurrent ascites and many showed a substantial reduction of GFR. Thus, patients with ascites at the first appearance can confidently be treated with anti-mineralocorticoids alone, as they will likely develop a satisfactory response with few side effects. Patients with long-standing, recurrent ascites should receive the combination therapy, which likely shortens the time to achieve natriuresis and lowers the incidence of hyperkalemia.¹ In a randomised double-blind crossover trial torasemide induced greater cumulative 24 h diuresis than furosemide, suggesting that torasemide might be more advantageous in patients exhibiting a weak response to furosemide.⁵⁷

Following mobilisation of ascites, diuretics should be tapered to the lowest dosages able to maintain patients with minimal or no ascites, to minimise side effects. Whenever possible, an aetiological treatment of the underlying cirrhosis should be instituted, as this eases the control of ascites in many cases.

Complications of diuretic therapy. The haemodynamic status of patients with cirrhosis and ascites⁸ makes them highly susceptible to rapid reductions in extracellular fluid volume, which mostly occur with loop diuretics. Thus, renal failure is frequent in this setting,⁴⁸ as is hepatic encephalopathy, also favoured by increased renal ammonia production. Loop diuretics can also lead to potassium and magnesium depletion. Hyponatraemia is another common diuretic-induced side effect in cirrhosis. It mostly, but not exclusively, occurs with loop diuretics, as they inhibit Na-K-Cl transporter and, therefore, solute-free water generation. Plasma volume contraction can also enhance arginine-vasopressin release. Thus, hyponatraemia can also ensue with anti-mineralocorticoid administration, albeit infrequently. Most experts agree on at least temporarily withdrawing diuretics when serum sodium concentration decreases below 120–125 mmol/L. Hyperkalemia, especially in patients with reduced renal perfusion, and painful gynecomastia are the most common side effects induced by anti-mineralocorticoids.

Muscle cramps can impair quality of life in patients receiving diuretics. Albumin infusion can relieve cramps,⁵⁸ as well as baclofen (10 mg/day, with a weekly increase of 10 mg/day up to 30 mg/day), which was safely used in a recent RCT.⁵⁹ One RCT investigated the use of quinidine at the dose of 400 mg/day for four weeks in patients with cirrhosis with painful muscle cramps. Although more effective than placebo, quinidine was associated with diarrhoea in about one-third of cases requiring treatment withdrawal.⁶⁰ Because of the frequency of diuretic-induced side effects, especially during the first month of treatment,⁵⁵ serial measurements of SCr, sodium, and potassium are warranted. The assessment of urine sodium excretion can be limited to non-responders, to unveil excessive sodium intake.

Recommendations

- Patients with the first episode of grade 2 (moderate) ascites should receive an anti-mineralocorticoid drug alone, starting at 100 mg/day with stepwise increases every 72 h (in 100 mg steps) to a maximum of 400 mg/day if there is no response to lower doses (**I;1**).
- In patients who do not respond to anti-mineralocorticoids, as defined by a body weight reduction of less than 2 kg/week, or in patients who develop hyperkalemia, furosemide should be added at an increasing stepwise dose from 40 mg/day to a maximum of 160 mg/day (in 40 mg steps) (**I;1**).
- Patients with long-standing or recurrent ascites should be treated with a combination of an anti-mineralocorticoid drug and furosemide, the dose of which should be increased sequentially according to the response, as explained (**I;1**).
- Torasemide can be given in patients exhibiting a weak response to furosemide (**I;2**).
- During diuretic therapy a maximum weight loss of 0.5 kg/day in patients without oedema and 1 kg/day in patients with oedema is recommended (**II-2;1**).
- Once ascites has largely resolved, the dose of diuretics should be reduced to the lowest effective dose (**III;1**).
- During the first weeks of treatment patients should undergo frequent clinical and biochemical monitoring particularly on first presentation (**I;1**).
- In patients presenting with GI haemorrhage, renal impairment, hepatic encephalopathy, hyponatraemia, or alterations in serum potassium concentration, these abnormalities should be corrected before starting diuretic therapy (**III;1**). In these patients, cautious initiation of diuretic therapy and frequent clinical and biochemical assessments should be performed (**III;1**). Diuretic therapy is generally not recommended in patients with persistent overt hepatic encephalopathy (**III;1**).
- Diuretics should be discontinued if severe hyponatraemia (serum sodium concentration <125 mmol/L), AKI, worsening hepatic encephalopathy, or incapacitating muscle cramps develop (**III;1**).
- Furosemide should be stopped if severe hypokalemia occurs (<3 mmol/L). Anti-mineralocorticoids should be stopped if severe hyperkalemia occurs (>6 mmol/L) (**III;1**).
- Albumin infusion or baclofen administration (10 mg/day, with a weekly increase of 10 mg/day up to 30 mg/day) are recommended in patients with muscle cramps (**I;1**).

Grade 3 or large ascites. The treatment of choice for the management of patients with grade 3 ascites is represented by LVP. Paracentesis should be performed under strict sterile conditions using disposable sterile materials. The procedure is associated with a very low risk of local complications, particularly bleeding^{61,62} even in patients with international normalized ratio

(INR) > 1.5 and platelet count < 50,000/ μ l, minor bleeding from puncture site occurred in two out of 142 paracentesis.⁶¹ Thus, there are no data supporting the prophylactic use of fresh frozen plasma of pooled platelets, even though these are employed in many centres when prothrombin activity is below 40% and platelet count < 40,000/ μ l. LVP should be avoided in the presence of disseminated intravascular coagulation. Other contraindications to LVP are reported (Table 3).

The removal of large volumes of ascitic fluid is potentially associated with a further reduction of effective blood volume, a condition known as post-paracentesis circulatory dysfunction (PPCD).⁶³ The clinical manifestations of PPCD are renal failure, dilutional hyponatraemia, hepatic encephalopathy and decreased survival.⁶³ Plasma volume expansion should be performed at the completion of LVP to prevent this complication. Artificial plasma expanders, such as dextran-70 (8 g/L of ascites removed)⁶⁴ or polygeline (150 ml/L),⁶⁴ saline solution (170 ml/L),⁶⁵ only show a similar efficacy to 20% albumin (8 g/L)⁶⁴ when less than 5 L of ascites are removed. However, polygeline is no longer used in many countries because of the potential risk of transmission of prions and dextran carries the risk of severe allergic reaction and renal failure. A meta-analysis of randomised trials showed that albumin is superior to any other plasma expander or vasoconstrictor not only in preventing PPCD, but also its clinical consequences, such as hyponatraemia and mortality.⁶⁶ Moreover, albumin infusion after LVP appears to be more cost-effective than a cheaper plasma volume expander, such as polygeline, because of the lower number of liver-related complications and hospital costs for a 30-day period.⁶⁷ LVP combined with infusion of albumin in patients with grade 3 ascites is more effective and safer than diuretics.^{68,69} However, LVP does not modify the underlying pathophysiological abnormalities leading to ascites formation. Thus, patients treated with LVP require diuretic therapy to prevent the re-accumulation of ascites.⁷⁰

Recommendations

- LVP is the first-line therapy in patients with large ascites (grade 3 ascites), which should be completely removed in a single session (**I;1**).
- LVP should be followed with plasma volume expansion to prevent PPCD (**I;1**).
- In patients undergoing LVP of greater than 5 L of ascites, plasma volume expansion should be performed by infusing albumin (8 g/L of ascites removed), as it is more effective than other plasma expanders, which are not recommended for this setting (**I;1**).
- In patients undergoing LVP of less than 5 L of ascites, the risk of developing PPCD is low. However, it is generally agreed that these patients should still be treated with albumin because of concerns about use of alternative plasma expanders (**III;1**).
- After LVP, patients should receive the minimum dose of diuretics necessary to prevent re-accumulation of ascites (**I;1**).
- When needed, LVP should also be performed in patients with AKI or SBP (**III;1**).

Table 3. Contraindications to paracentesis.

• Uncooperative patient
• Abdominal skin infection at the proposed puncture sites
• Pregnancy
• Severe coagulopathy (accelerated fibrinolysis or disseminated intravascular coagulation)
• Severe bowel distension

Drugs contraindicated in patients with ascites

As non-steroidal anti-inflammatory drugs inhibit renal prostaglandin synthesis, they should not be used in patients with cirrhosis and ascites, where an increased vasodilating prostaglandin synthesis counteracts the renal vasoconstrictor effects of angiotensin II. Indeed, their administration can lead to acute renal failure, hyponatraemia, and diuretic resistance.⁷¹ It would appear that selective inhibitors of cyclooxygenase-2 do not impair renal function and response to diuretics in patients with ascites.⁷² However, it is not known whether these drugs can be safely used in clinical practice when analgesia is needed. Patients with ascites are also particularly sensitive to the renal vasoconstrictor effect of endogenous adenosine, and dipyridamole can induce a marked reduction in renal perfusion.⁷³

The maintenance of an adequate arterial pressure in cirrhosis with ascites is assured by the activation of endogenous vasoconstrictor systems. Thus, angiotensin-converting enzyme inhibitors,⁷⁴ angiotensin II receptor antagonists, and α 1-adrenergic blockers⁷⁵ should be avoided, as they can induce arterial hypotension and renal function impairment. Aminoglycosides should be avoided in the treatment of bacterial infections, except in specific cases (discussed later), because they are associated with high incidence of nephrotoxicity.⁷⁶ Although cirrhosis with ascites and preserved renal function does not appear to be a risk factor for renal failure induced by contrast media,⁷⁷ this cannot be excluded in patients with impaired renal function. In these cases, preventive measures such as plasma volume expansion with saline may be employed.⁷⁸

Recommendations

- Non-steroidal anti-inflammatory drugs should not be used in patients with ascites because of the high risk of developing further sodium retention, hyponatraemia, and AKI (**II-2;1**).
- Angiotensin-converting-enzyme inhibitors, angiotensin II antagonists, or α 1-adrenergic receptor blockers should not generally be used in patients with ascites because of increased risk of renal impairment (**II-2;1**).
- The use of aminoglycosides is discouraged, as they are associated with an increased risk of AKI. Their use should be reserved for patients with severe bacterial infections that cannot be treated with other antibiotics (**II-2;1**).
- In patients with ascites and preserved renal function, the use of contrast media does not appear to be associated with an increased risk of renal impairment (**II-2**). There are insufficient data in patients with renal failure. Nevertheless, a cautious use of contrast media and the use of preventive measures for renal impairment are recommended (**III;1**).

Refractory ascites*Evaluation of patients with refractory ascites*

According to the criteria of the International Ascites Club, refractory ascites is defined as “ascites that cannot be mobilised or the early recurrence of which (*i.e.*, after LVP) cannot be satisfactorily prevented by medical therapy”.^{31,32} The diagnostic criteria of refractory ascites are shown in Table 4. Refractoriness of ascites is associated with a poor prognosis, with a median survival of about six months.⁷⁹ Therefore, if a patient with refractory ascites has not yet been considered for LT, he/she should be immediately referred to a liver transplant center. The potential underestimation of the mortality risk by commonly used prognostic scores, as discussed earlier also applies to patients with refractory ascites.⁸⁰

Recommendations

- The diagnosis of refractory ascites relies on the assessment of the response of ascites to diuretic therapy and salt restriction. Such an evaluation should be done in stable patients without associated complications, such as bleeding or infection, after ascertaining patient compliance to treatment (III;1).
- Patients with refractory ascites should be evaluated for LT (III;1).

Management of refractory ascites

Large-volume paracentesis. There is general agreement that LVP is an effective and safe treatment of refractory ascites,^{31,35} which should be associated with albumin administration to prevent PPCD.

Diuretics in patients with refractory ascites. Once refractoriness of ascites has been ascertained, diuretics should be discontinued. Only when renal sodium excretion on diuretics exceeds 30 mmol/day, maintenance of diuretic therapy can be considered, when tolerated.³¹

Non-selective beta-blockers in patients with refractory ascites. The controversial issue on the use of non-selective beta-blockers (NSBBs) in patients with ascites and, particularly, in those with

refractory ascites will be developed in the section dedicated to GI bleeding.

Transjugular intrahepatic portosystemic shunts. Transjugular intrahepatic portosystemic shunts (TIPS) decompresses the portal system by shunting an intrahepatic portal branch into a hepatic vein. Its insertion accentuates peripheral arterial vasodilation in the short term. However, within 4–6 weeks its result is an improvement in effective volemia and renal function, ultimately leading to an increase in renal sodium excretion.^{81–85} TIPS-induced natriuresis can be delayed by advanced age and reduced pre-TIPS GFR,⁸⁴ and prevented by intrinsic kidney disease.⁸⁶ TIPS may also exert beneficial effects on nitrogen balance and nutrition⁸⁷ and quality of life.⁸⁸ A major complication after TIPS insertion using bare stent grafts is the development of hepatic encephalopathy, which can occur in up to 50% of patients.^{89,90} The incidence of this complication can be significantly reduced to about 18% with the use of polytetrafluoroethylene (PTFE)-covered stent grafts of 8 mm,⁹¹ a result confirmed by a recent randomised trial comparing 8 mm and 10 mm stent grafts.⁹² Notably, this favourable effect is better than with larger stent grafts underdilated to 8 mm. Indeed, it has been shown that underdilated 10 mm stent grafts passively expand to almost the full diameter within 1–6 weeks.⁹³ It must be underlined that the indication for TIPS insertion in these studies was the prevention or treatment of recurrent bleeding, which may restrict the relevance of these results in patients with refractory ascites. Dysfunction of TIPS with bare stent grafts because of stent thrombosis and stenosis can develop in up to 80% of cases.⁸⁹ This complication has been significantly reduced with the use of PTFE-covered stents.⁹⁴

Controlled studies and meta-analysis. The clinical effects of TIPS with bare stents in patients with refractory or recurrent ascites have been assessed in six prospective RCTs,^{95–100} whose main features are reported (Table 5). Based on these RCTs, seven meta-analyses were performed.^{101–107} The final messages can be summarised as follow: i) TIPS controlled ascites better than LVP, and ii) TIPS is followed by a greater incidence of hepatic encephalopathy. However, discrepant results were obtained with respect to survival. A better survival with LVP, mainly because of a detrimental effect of TIPS in Child-Pugh class C patients, was reported by one study,⁹⁶ while no difference was reported by two.^{95,100} A better survival with TIPS was reported

Table 4. Definition and diagnostic criteria for refractory ascites in cirrhosis.

Definition	
Diuretic-resistant ascites	Ascites that cannot be mobilized or the early recurrence of which cannot be prevented because of a lack of response to sodium restriction and diuretic treatment
Diuretic-intractable ascites	Ascites that cannot be mobilized or the early recurrence of which cannot be prevented because of the development of diuretic-induced complications that preclude the use of an effective diuretic dosage
Diagnostic criteria	
Treatment duration	Patients must be on intensive diuretic therapy (spironolactone 400 mg/day and furosemide 160 mg/day) for at least one week and on a salt-restricted diet of less than 90 mmol/day
Lack of response	Mean weight loss of <0.8 kg over four days and urinary sodium output less than the sodium intake
Early ascites recurrence	Reappearance of grade 2 or 3 ascites within four weeks of initial mobilisation
Diuretic-induced complications	Diuretic-induced hepatic encephalopathy is the development of encephalopathy in the absence of any other precipitating factor Diuretic-induced renal impairment is an increase of serum creatinine by >100% to a value >2 mg/dl (177 μmol/L) in patients with ascites responding to treatment Diuretic-induced hyponatremia is defined as a decrease of serum sodium by >10 mmol/L to a serum sodium of <125 mmol/L Diuretic-induced hypo- or hyperkalemia is defined as a change in serum potassium to <3 mmol/L or >6 mmol/L despite appropriate measures Invalidating muscle cramps

Table 5. Characteristics and results of six randomised controlled trials comparing bared TIPS and LVP in patients with cirrhosis and refractory or recidivant ascites.

Refs.	Refractory/ recidivant ascites (%)	Exclusion criteria	Enrolled patients (N)		Ascites improved (%)		Encephalopathy (%)		Survival (%)	
			TIPS	LVP	TIPS	LVP	TIPS	LVP	TIPS	LVP
Lebrec <i>et al.</i> 1995	100/0	Age >70 yr; severe extra-hepatic diseases; HCC; pulmonary hypertension; HE, bacterial infection; severe alcoholic hepatitis; portal or hepatic vein obstruction or thrombosis; obstruction of biliary tract; obstruction of hepatic artery; serum creatinine >1.7 mg/dl	13	12	38	0	15	6	29	60
Rössle <i>et al.</i> 2000	55/45	HE ≥grade 2; serum bilirubin >5 mg/dl, serum creatinine >3 mg/dl; portal-vein thrombosis, hepatic hydrothorax; advanced cancer; failure of LVP (ascites persisting after LVP or need for LVP >once per week)	29	31	84	43*	23	13	58	32
Ginés <i>et al.</i> 2002 ⁹⁰	100/0	Age >18 or >75 yr; serum bilirubin >10 mg/dl; prothrombin time <40% (INR 2.5); platelet count <than 40,000/mm ³ ; serum creatinine >3 mg/dl, HCC, complete portal vein thrombosis; cardiac or respiratory failure; organic renal failure; bacterial infection; chronic HE	35	35	51	17*	60	34	26	30
Sanyal <i>et al.</i> 2003 ¹⁰⁰	100/0	Causes of ascites other than cirrhosis; advanced liver failure (serum bilirubin >5 mg/dl, PT INR >2); incurable cancers or nonhepatic diseases that were likely to limit life expectancy to 1 yr; congestive heart failure; acute renal failure; parenchymal renal disease; portal vein thrombosis; bacterial infections; HE ≥grade II; florid alcoholic hepatitis, HCC; gastrointestinal hemorrhage within 6 w of randomisation.	52	57	58	16*	38	21	35	33
Salerno <i>et al.</i> 2004 ⁹⁹	68/32	Age > 72 yr; recurrent HE ≥grade 2; serum bilirubin >6 mg/dl; serum creatinine >3 mg/dl; Child-Pugh score >11; complete portal vein thrombosis; HCC; gastrointestinal bleeding within 15 d of randomisation; serious cardiac or pulmonary dysfunctions; bacterial infection; SAAG gradient <11 g/L.	33	33	79	42*	61	39	59	29*
Narahara <i>et al.</i> 2011 ⁹⁷	100/0	Age >70 yr, chronic HE, HCC and other malignancies, complete portal vein thrombosis with cavernomatous transformation, bacterial infection, severe cardiac or pulmonary disease, organic renal disease.	30	30	87	30*	20	5	20	5*

HCC, hepatocellular carcinoma; HE, hepatic encephalopathy; INR, international normalized ratio; LVP, large volume paracentesis; PT, prothrombin time; SAAG, serum-ascites albumin gradient; TIPS, transjugular intrahepatic portosystemic shunt.

* Significantly lower than TIPS.

in another two studies,^{97,99} while, in the remaining one,⁹⁸ although a survival advantage was not found, TIPS was independently associated with transplant-free survival at multivariate analysis. In four meta-analyses including the five studies available at that time no survival advantage with TIPS emerged. However, a trend towards reduced mortality with TIPS was seen¹⁰⁴ after the exclusion of an outlier trial.⁹⁶ The latter was also excluded in the only meta-analysis on individual patient data, and an increased transplant-free survival was found.¹⁰⁷ Finally, the two meta-analyses that included all six trials^{102,103} provided contrasting results, as an improved transplant-free survival was found in one,¹⁰⁷ while a survival advantage with TIPS was limited to patients with recurrent ascites in the other.¹⁰²

Fewer studies assessing the effects of TIPS with PTFE-covered stent grafts are available. Two retrospective studies^{108,109} reported better control of ascites and one-year¹⁰⁸ or two-year¹⁰⁹ survival with covered stent grafts than bare stent grafts in patients with refractory ascites. A survival benefit of TIPS vs. serial paracentesis in patients with refractory ascites has been reported in a single-centre case-control propensity score analysis.¹¹⁰ In a recent RCT comparing covered TIPS vs. LVP in patients with recurrent ascites, a better one-year transplant-free survival was seen in patients treated with covered stents, without any significant increase in occurrence of hepatic encephalopathy.¹¹¹ Thus, currently available data suggest that TIPS improves survival compared to LVP in patients with recurrent ascites, but it does not in those with refractory ascites.

A careful selection of patients is also crucial to maximise the beneficial effects of TIPS, as TIPS can even be detrimental in patients with the most advanced stages of cirrhosis, such as

those belonging to Child-Pugh class C.⁹⁶ The main exclusion criteria for TIPS insertion in the seven RCTs are reported in Table 5. A score system based on SCr, INR, serum bilirubin and aetiology of cirrhosis has been proposed to predict survival after TIPS insertion for refractory ascites.¹¹² Another simple predictor of survival suggested for patients receiving TIPS for refractory ascites consists of the combination of serum bilirubin concentration and platelet count.¹¹³ Another factor that seems to influence mortality is the number of TIPS procedures performed in a centre, as the risk of inpatient mortality is lower in hospitals performing ≥20 TIPS per year.¹¹⁴

Other treatments. Based on the exclusion criteria reported (Table 5), a substantial portion of patients with refractory ascites are not candidates for TIPS insertion. Thus, the search for alternative treatments is warranted.

Medical treatments. Therapies aimed at improving circulatory and renal function have been proposed. The α1-adrenergic agonist midodrine has been shown to improve systemic and renal haemodynamics in patients with cirrhosis and uncomplicated ascites.¹¹⁵ In a small RCT comparing the addition of midodrine (7.5 mg t.i.d) to diuretic treatment with diuretic treatment alone in patients with refractory or recurrent ascites for six months, only a transient beneficial effect on the control of ascites was seen at the third month.¹¹⁶ The use of terlipressin, an analogue of vasopressin with a predominant vasoconstrictor effect in the splanchnic circulatory area in patients with refractory ascites has only been assessed in acute studies. In one,¹¹⁷ terlipressin administration (1 to

Clinical Practice Guidelines

2 mg intravenous [i.v., according to body weight) only increased renal sodium excretion when associated with exogenous atrial natriuretic factor. In another,¹¹⁸ 2 mg of terlipressin led to an increase in GFR, renal plasma flow and renal sodium excretion. However, in this study only eight patients with refractory ascites were included. Whether a prolonged treatment with terlipressin may lead to a clinically relevant improvement of renal function and sodium excretion in refractory ascites is not known.

The α 2-adrenoceptor agonist clonidine, a sympatholytic drug, which suppresses RAAS activity and improves the response to diuretics in patients with cirrhosis and ascites was tested in a large prospective RCT. It was shown that clonidine administration on top of diuretics for three months led to an overall response to diuretics in 60% of cases, while no response was seen with diuretics alone. This effect was associated with significant reductions of RAAS and sympathetic nervous system activity. Interestingly, the favourable effects of clonidine were predicted by the variant genotype of G protein (GNB3 C825T) and adrenergic receptor (ADRA2C Del 322–325) polymorphisms, and the baseline norepinephrine level.¹¹⁹ Small scale or pilot studies evaluated the effects of various combinations of midodrine with either clonidine,¹²⁰ the antagonist of vasopressin V2-receptors tolvaptan,¹²¹ or octreotide and albumin¹²² in patients with refractory and recurrent ascites. Some promising results were obtained, but they need to be confirmed by sufficiently powered RCTs. A recent RCT¹²³ compared the effects of the combined administration of midodrine (5 mg t.i.d) and rifaximin (550 mg b.i.d) on top of diuretics for 12 weeks with diuretics alone. After 12 weeks, 80% of patients in the active arm were complete responders with a significant improvement in survival in the midodrine/rifaximin arm. Due to weakness in the study design, these results are not definitive, but they certainly warrant further investigation.

Alfapump®. The automated low-flow ascites pump (Alfapump®) system consists of a subcutaneously implanted battery-powered programmable pump. It is connected to catheters that transfer ascites from the peritoneal cavity to the bladder, from which it is eliminated with urine. The device has internal sensors that monitor pump function. In two multicentre safety and efficacy studies,^{124,125} Alfapump® ensured a significant reduction of the number and volume of paracentesis in patients with advanced cirrhosis and refractory ascites. However, adverse effects directly related to the device occurred in about one-third¹²⁴ to half¹²⁵ of cases. In a multicentre RCT in patients with refractory ascites, Alfapump® reduced the median number of paracentesis per month by 85% with respect to LVP, and significantly improved quality of life and nutritional parameters, as assessed by hand-grip strength and body mass index. Alfapump® had no effect on survival and was associated with a significantly higher incidence of serious adverse events (85.2 vs. 45.2%), mainly represented by AKI.¹²⁶ Thus, even though Alfapump® is effective in reducing the need for paracentesis in patients with refractory ascites, its frequent side effects require close monitoring of patients. Indeed, in addition to device-related adverse event, it should be noted that the evaluation of kidney and circulatory function in 10 patients with cirrhosis and refractory ascites

carrying Alfapump® has shown a significant GFR decline within six months, which was associated with a marked increase in plasma renin activity and norepinephrine concentration.¹²⁷ This likely represented the pathophysiological background of 18 episodes of AKI experienced by seven patients.

Recommendations

- Repeated LVP plus albumin (8 g/L of ascites removed) are recommended as first line treatment for refractory ascites (**I;1**).
- Diuretics should be discontinued in patients with refractory ascites who do not excrete >30 mmol/day of sodium under diuretic treatment (**III;1**).
- Although controversial data exist on the use of NSBBs in refractory ascites, caution should be exercised in cases of severe or refractory ascites. High doses of NSBB should be avoided (*i.e.* propranolol >80 mg/day) (**II-2;1**). The use of carvedilol can not be recommended at present (**I;2**).
- Patients with refractory or recurrent ascites (**I;1**), or those for whom paracentesis is ineffective (*e.g.* due to the presence of loculated ascites) should be evaluated for TIPS insertion (**III;1**).
- TIPS insertion is recommended in patients with recurrent ascites (**I;1**) as it improves survival (**I;1**) and in patients with refractory ascites as it improve the control of ascites (**I;1**).
- The use of small-diameter PTFE-covered stents in patients is recommended to reduce the risk of TIPS dysfunction and hepatic encephalopathy with a high risk of hepatic encephalopathy is recommended (**I;1**).
- Diuretics and salt restriction should be continued after TIPS insertion up to the resolution of ascites (**II-2;1**), as well as close clinical follow-up (**III,1**).
- Careful selection of patients for elective TIPS insertion is crucial, as is the experience of the centre performing this procedure. TIPS is not recommended in patients with serum bilirubin > 3 mg/dl and a platelet count lower than $75 \times 10^9/L$, current hepatic encephalopathy grade ≥ 2 or chronic hepatic encephalopathy, concomitant active infection, progressive renal failure, severe systolic or diastolic dysfunction, or pulmonary hypertension (**III;1**).
- At present the addition of clonidine or midodrine to diuretic treatment cannot be recommended (**III;1**).
- Alfapump® implantation in patients with refractory ascites not amenable to TIPS insertion is suggested in experienced centres. However, close patient monitoring is warranted because of the high risk of adverse events including renal dysfunction and technical difficulties (**I;2**).

Hepatic hydrothorax

Hepatic hydrothorax describes the accumulation of transudate in the pleural space of patients with decompensated cirrhosis in the absence of cardiac, pulmonary or pleural disease. Its formation is secondary to small diaphragmatic defects, more often located in the right side, through which ascites moves into the pleural space because of the negative intrathoracic pressure induced by inspiration. Hepatic hydrothorax can lead to respiratory failure and can be complicated by spontaneous bacterial infections (empyema). Its appearance is associated with poor prognosis, as the median survival of patients with hepatic hydrothorax ranges from 8–12 months.^{128,129} Notably, the most common prognostic scores, such as Child-Pugh and MELD, seem to underestimate such an adverse outcome.¹²⁸

Diagnosis of hepatic hydrothorax

Once pleural effusion has been ascertained, cardiopulmonary and primary pleural diseases should be excluded by standard clinical approaches. Diagnostic thoracentesis is required to rule out bacterial infection, whose diagnosis relies on the same criteria described for ascites. The protein content of pleural effusion in uncomplicated hepatic hydrothorax is low and the serum to pleural fluid albumin gradient is greater than 1.1 g/dl.¹²⁸

The presence and extent of diaphragmatic defects can be assessed indirectly, by radioisotope techniques, or directly by magnetic resonance imaging or colour-Doppler ultrasonography.^{130,131}

Treatment of hepatic hydrothorax

The first-line management relies on the treatment of ascites with diuretics and/or LVP as discussed earlier. However, it is not rare for pleural effusion to persist despite successful treatment of ascites (refractory hydrothorax). Therapeutic thoracentesis is required to relieve dyspnoea. Its efficacy in refractory hepatic hydrothorax is transient and repeated thoracentesis are required, which increase the risk of complications such as pneumothorax, pleural or soft tissue infection, and bleeding.¹³² The frequent occurrence of these complications discourages the use of chronic pleural drainage, which can also be followed by renal dysfunction from fluid loss.¹³³

Whenever indicated and possible, LT represents the best option for patients with refractory hepatic hydrothorax, which does not seem to adversely affect the outcome of transplantation.^{134,135} TIPS has been effectively employed as definitive treatment or bridge to transplantation in patients with refractory hepatic hydrothorax, whose general outcome seems to be related to the severity of the underlying cirrhosis.^{136,137} These results have been confirmed by a more recent meta-analysis.¹³⁸

Pleurodesis induced by various agents, such as talc, tetracycline, doxycycline, bleomycin and povidone-iodine, can be offered to patients who are not candidates for TIPS or LT. A recent meta-analysis showed that the pooled rate of complete response after pleurodesis was 72%. However, the pooled rate of complications related to this procedure was as high as 82%.¹³⁹ Finally, thoracoscopic repair with mersilene mesh appears to be effective in patients with well-defined diaphragmatic defects.¹⁴⁰ Advanced liver disease, as assessed by MELD score, and preoperative renal dysfunction appear to adversely affect three-month survival. Unfortunately, clear cut-off values cannot be retrieved from that study.

Recommendations

- Patients with hydrothorax should be evaluated for LT (III;1).
- Cardiopulmonary and primary pleural disease should be ruled out before diagnosing hepatic hydrothorax (III;1). Diagnostic thoracentesis should be performed especially when infection of the pleural effusion is suspected (III;1).
- Diuretics and thoracentesis are recommended as the first-line management of hepatic hydrothorax (III;1).
- Therapeutic thoracentesis is indicated in patients with dyspnoea (III;1). Chronic pleural should not be performed because of the frequent occurrence of complications (II-2;1).
- In selected patients, TIPS insertion for recurrent symptomatic hepatic hydrothorax is recommended (II-2;1).
- Pleurodesis can be suggested to patients with refractory hepatic hydrothorax not amenable to LT or TIPS insertion. However, the frequent occurrence of side effects related to this technique restricts its use to selected patients (I;2).
- Mesh repair of diaphragmatic defects is suggested for the management of hepatic hydrothorax in very selected patients. The best results can be achieved in patients with non-advanced cirrhosis without renal dysfunction (II-2;2).

Hyponatremia

Definition and pathophysiology

Hyponatremia is common in patients with advanced cirrhosis, and has been arbitrarily defined as serum sodium concentration lower than 130 mmol/L.^{141,142} However, according to guidelines on hyponatremia in the general patient population,¹⁴³ reductions below 135 mmol/L should also be considered. Patients with hyponatremia have a poor prognosis, as it is associated with increased mortality^{144,145} and morbidity, particularly neurological complications,^{146,147} and reduced survival after LT.¹⁴⁸ Incorporating serum sodium concentration into the MELD score, a new score (MELD-Na) was generated that provides more accurate survival predictions than MELD alone,¹⁴⁹ especially in patients with ascites and hyponatremia with intermediate MELD score values.¹⁵⁰ Both hypovolaemic and hypervolaemic hyponatremia can occur in patients with cirrhosis. The second, most common, is characterised by an expansion of the extracellular fluid volume, with ascites and oedema. It may occur spontaneously, or because of excessive hypotonic fluids (*i.e.*, 5% dextrose), or secondary to complications of cirrhosis leading to an abrupt worsening of effective volemia. The main drivers are non-osmotic hypersecretion of vasopressin and enhanced proximal nephron sodium reabsorption, which impair free water generation and are both caused by effective hypovolaemia. As opposed to hypervolaemic hyponatremia, hypovolaemic hyponatremia is characterised by the frequent absence of ascites and oedema.

Clinical Practice Guidelines

It is caused by a prolonged negative sodium balance with marked loss of extracellular fluid often due to excessive diuretic therapy.

Management of hyponatremia

It is generally considered that hyponatremia should be treated when serum sodium is lower than 130 mmol/L, although there is no good evidence regarding the level of serum sodium at which treatment should be initiated. Hypovolaemic hyponatremia requires plasma volume expansion with saline solution and the correction of the causative factor. The management of hypervolemic hyponatremia requires attainment of a negative water balance. Non-osmotic fluid restriction is helpful in preventing a further decrease in serum sodium levels, but it is seldom effective in improving natremia. Hypertonic sodium chloride administration to patients with decompensated cirrhosis may improve natremia but enhances volume overload and worsens the amount of ascites and oedema. Therefore, it should be limited to severely symptomatic hyponatremia, as defined by life-threatening manifestations, cardio-respiratory distress, abnormal and deep somnolence, seizures and coma, which do not frequently occur in patients with cirrhosis. Furthermore, hypertonic sodium chloride administration can be considered in patients with severe hyponatremia who are expected to get a liver transplant within a few days. In these cases, hyponatremia must not be corrected completely and rapidly to avoid the risk of central pontine myelinolysis that is increased in advanced cirrhosis.¹⁴³ In practice, after an initial rapid correction aimed at attenuating clinical symptoms (5 mmol/L in the first hour), serum sodium concentration should not increase more than 8 mmol/L per day.¹⁴³ Albumin infusion appears to improve serum sodium concentration, but more information is needed.¹⁵¹

Vaptans

Vaptans are selective antagonists of the V2-receptors of arginine-vasopressin in the principal cells of the collecting ducts that enhance solute-free water excretion.¹⁵² Indeed, these drugs are effective in improving serum sodium concentration in conditions associated with high vasopressin levels, such as the syndrome of inappropriate antidiuretic hormone secretion (SIADH) and heart failure.¹⁵² The effects of the administration of vaptans to hyponatremic patients with cirrhosis and ascites have been assessed in several studies. Namely, tolvaptan, satavaptan and lixivaptan lead to an increased urine volume, a solute-free water excretion, and an improvement of hyponatremia in 45–82% of cases.^{153–155} In another study, a short-term intravenous infusion of conivaptan for one to four days in patients with end stage liver disease awaiting OLT was also effective in increasing serum sodium concentration.¹⁵⁶ However, the safety of vaptans has only been established for short-term treatments lasting from one week to one month. When satavaptan was used long term, in addition to diuretics, despite improving both serum sodium concentration and control of ascites, a higher all-cause mortality rate, mostly associated with known complications of cirrhosis, was reported compared to standard medical treatment.^{157,158} Moreover, a recent study cast doubt on the efficacy of tolvaptan in patients with cirrhosis and severe hypervolemic hyponatremia (serum sodium ≤ 125 mEq/L) in a real-life setting.¹⁵⁹ At present, both conivaptan and tolvaptan have been approved in the US by the FDA, while only tolvaptan in Europe has been approved

by the EMA for management of severe hypervolemic hyponatremia (<125 mmol/L). The unique indication given for tolvaptan by the EMA is SIADH, while the FDA also included heart failure and liver cirrhosis. However, the occurrence of serious hepatic injury in three patients with autosomal dominant polycystic kidney disease treated with tolvaptan in a double-blind placebo-controlled trial¹⁶⁰ led the FDA to conclude that this drug should not be used in patients with underlying liver disease.

Recommendations

- The development of hyponatremia (serum sodium concentration <130 mmol/L) in patients with cirrhosis carries an ominous prognosis, as it is associated with increased mortality and morbidity. These patients should be evaluated for LT (**II-2;1**).
- The removal of the cause and administration of normal saline are recommended in the management of hypovolemic hyponatremia (**III;1**).
- Fluid restriction to 1,000 ml/day is recommended in the management of hypervolemic hyponatremia since it may prevent a further reduction in serum sodium levels (**III;1**).
- The use of hypertonic saline in the management of hypervolemic hyponatremia should be limited to the rare cases presenting with life threatening complications. It could also be considered in patients with severe hyponatremia who are expected to get LT within a few days. The correction of serum sodium concentration, once an attenuation of symptoms has been obtained, should be slow (≤ 8 mmol/L per day) to avoid irreversible neurological sequelae, such as osmotic demyelination (**II-3;1**).
- Albumin administration can be suggested in hypervolemic hyponatremia, but data are very limited to support its use (**II-3;2**).
- At present, the use of vaptans should be limited to controlled clinical studies (**III;1**).

Gastrointestinal bleeding

Pathophysiology

Variceal haemorrhage (VH) occurs because of the rupture of the variceal wall due to excessive wall tension. Variceal wall tension is an intrinsic property of the vessel wall that opposes the expansive force determined by variceal transmural pressure, which depends on portal pressure and vessel size. Tissue support surrounding the varix may counteract the increase in variceal pressure and size, protecting the wall from rupture.¹⁶¹ Once variceal wall rupture occurs, the amount of bleeding is related to transmural pressure (which mainly depends on portal pressure), to the area of rupture in the vessel wall and to blood viscosity and/or alterations of haemostasis.¹⁶¹ All these factors can be influenced by therapy. Drug therapy and portal-systemic derivative procedures, reduce portal (and variceal) pressure. Endoscopic therapies and other physical methods, such as balloon tamponade or expandable prosthesis, act merely by both interrupting the blood flow into the varix

and sealing the vascular wall. Portal pressure is a key factor determining both variceal rupture and the severity of the bleeding episode.¹⁶¹ During acute bleeding, portal pressure may increase because of different factors such as over-transfusion or absorption of blood from the gut, which may have a role in failure to control bleeding and/or precipitating rebleeding. Portal pressure is usually assessed by the hepatic venous pressure gradient (HVPG).

Natural history of gastro-oesophageal varices and variceal haemorrhage

Variceal haemorrhage, causative of 70% of all upper GI bleeding events in patients with portal hypertension, remains one of the most severe and immediate life-threatening complications in patients with cirrhosis and constitutes the second most frequent decompensating event after ascites.^{162,163} Decompensated patients have 'clinically significant portal hypertension' (CSPH) per definition and, consequently, a high risk of having gastro-oesophageal varices. In fact, while only 42% of Child A patients have gastro-oesophageal varices, 72% of Child B/C patients do.¹⁶⁴ When decompensation develops, patients without varices on a previous endoscopy should have a repeat endoscopy performed given the risk of developing varices due to worsening of portal hypertension and liver dysfunction. In those without varices at screening, "de novo" varices develop at a rate of 7–8%/year,^{165,166} which could be higher in patients decompensated due to worsening of portal hypertension and liver dysfunction. The progression rate from small to large varices runs up to 22% at one year and 51% at three years in patients with Child B/C cirrhosis, especially when alcoholic in origin and/or when red wale marks are present at first endoscopy, compared to 2% and 16%, respectively, in compensated patients without those risk factors.^{165,166} Prospective studies have consistently demonstrated that the risk of VH, estimated overall at 5–15% per year, is related to variceal size.^{166–170} This risk is further amplified by the severity of liver dysfunction (Child B/C) and/or the presence of red wale marks on varices. Thus, not only medium/large varices (i.e. varices that do not collapse with insufflation at endoscopy), but also small varices with red signs or in Child C should be considered 'high-risk' varices. Despite improvement in therapy, overall mortality with each episode of VH remains around 15% to 25% at six weeks. Such risk is much higher in patients who develop VH in addition to other decompensation (over 80% at five-years) than in those presenting with VH as an isolated decompensating event (20% at five-years).^{170,171} Mortality risk is particularly high when VH is associated with AKI and/or concomitant bacterial infections.¹⁷² Without secondary prophylaxis, rebleeding occurs in approximately 60% to 70% of patients, usually within one to two years of the index haemorrhagic event.¹⁷³ Although increasing efforts are performed to test non-invasively for the presence of gastro-oesophageal varices, these efforts largely remain restricted to compensated cirrhosis.¹⁶⁷ Given the high prevalence of 'high-risk varices' in decompensated cirrhosis, oesophago-gastroduodenoscopy (EGD) should be performed to detect the presence, size of varices and presence of red wale marks.^{168,169}

Recommendations

- Because they are deemed high risk, patients in whom decompensation develops should have EGD performed to screen for gastro-oesophageal varices, unless previously diagnosed and treated (II-2;1).
- If EGD is performed, the presence, size and presence of red wale marks should be reported (II-2;1).
- In patients without varices on screening EGD in whom the aetiological factor persists and/or the state of decompensation continues, screening EGD should be repeated every year. In the remaining patients the screening could be prolonged, but the exact interval is unclear and more data is required (III;2).

Prevention and treatment of variceal haemorrhage

Considering the high-risk of death when VH occurs in patients with decompensated cirrhosis, implementation of strategies to adequately treat VH and to prevent (re)bleeding and death should be actively pursued in patients with decompensated cirrhosis. It should be noted that the current recommendations will concentrate on decompensated patients given the focus of these CPGs.

Primary and secondary prophylaxis of VH in decompensated patients

The Baveno VI¹⁶⁸ and American Association for the Study of Liver Diseases (AASLD)¹⁶⁹ guidelines primarily recommend NSBBs for primary prophylaxis of VH in patients with cirrhosis who have high-risk varices and also, combined with endoscopic band ligation (EBL), for secondary prophylaxis of VH. Both NSBBs and EBL have shown to be equally effective in preventing first bleeding in patients with high-risk varices. The choice between options depends on factors such as patient preference, contraindications or adverse events. Although numerically EBL induces less side effects, it has been associated with more severe and potentially life-threatening complications, resulting from bleeding EBL-ulcers. Moreover, EBL does not impact on portal hypertension. Thus, it does not reduce/prevent other complications and surveillance endoscopies are required to detect variceal recurrence, supporting overall primary preference for NSBBs.^{174,175} For prevention of rebleeding (secondary prophylaxis), combined therapy with NSBBs plus EBL is recommended because combination therapy significantly decreases the probability of rebleeding compared to monotherapy using either EBL or drug therapy. NSBBs are the cornerstone of combined therapy because a meta-analysis shows an improvement in survival with the addition of NSBBs (\pm nitrates) to EBL, while the addition of EBL to NSBBs (\pm nitrates) has no effect on mortality.¹⁷⁶ Recent RCTs indicate that guiding therapy according to the HVPG response to NSBBs can be valuable in this high-risk setting.^{91,177} HVPG-guided therapy may improve the outcomes achieved with current first-line therapy combining NSBBs and EBL,⁹¹ and may achieve a similar survival as covered TIPS, which is the most effective therapy in terms of preventing bleeding.¹⁷⁷

Clinical Practice Guidelines

Accordingly, HVPG-guided therapy can be used when available. However, this approach has relevant drawbacks such as invasiveness and limited availability and, therefore, cannot be widely recommended. NSBBs, such as propranolol and nadolol, act on portal hypertension because non-selective beta-blockade reduces cardiac output and splanchnic blood flow while the unopposed effect of alpha-1 adrenergic receptors leads to splanchnic vasoconstriction, thus reducing portal pressure and its consequential complications. Nonetheless, haemodynamic response rates to NSBBs are modest: approximately 46% of cases according to meta-analyses,^{178,179} endorsing the overall search for novel therapeutic options. Carvedilol, an NSBB with intrinsic anti-alpha-1 receptor activity, has been associated with a greater reduction in portal pressure than the traditional NSBBs and has therefore become a valuable alternative.¹⁸⁰ Its beneficial action on alpha-1 receptors reduces both porto-collateral and intrahepatic resistance, however, this is at the cost of more profound effects on systemic arterial pressure, particularly in decompensated patients. The problem with all the recommendations mentioned so far is that they are based on high quality RCTs that usually excluded patients with more advanced cirrhosis, while major controversy has arisen in recent years regarding the use and safety of NSBBs in patients with advanced disease, particularly in those with refractory ascites and/or SBP. The discussion was initiated by the Clichy group,¹⁸¹ who reported poor survival and increased risk of PPCD among patients with refractory ascites on NSBB therapy. The mechanism underlying these findings was thought to relate to further induction of systemic arterial hypotension and exhaustion of cardiac reserve, in light of the progressive hyperdynamic circulation typically associated with end-stage disease. As a result, end-organ perfusion becomes critical and sets off a multitude of complications, like HRS. Therefore, “the window hypothesis” was proposed which suggested refractory ascites as a critical juncture where the protective effects of NSBBs may cease and a detrimental impact may begin.¹⁸² However, this hypothesis was challenged by opposing reports suggesting protective effects with NSBBs even in decompensated patients.^{183–186} Illustratively, a recent *post hoc* analysis of three RCTs where vaptans and NSBBs were co-administered to patients with ascites showed that NSBBs did not increase mortality.¹⁸³ On the contrary, during follow-up, 29% of initial NSBB users stopped taking NSBBs, inducing a marked rise in mortality and coinciding with variceal bleeding, bacterial infection and/or development of HRS.¹⁸³ Non-haemodynamic effects of NSBBs, like reduction of intestinal permeability, inflammation and BT, are considered to contribute to the beneficial effect, particularly in this advanced stage.^{187–189} Whether NSBBs are detrimental in some patients with advanced cirrhosis should be clarified by future studies (ideally RCTs), as well as the optimal drug-schedule in such stages. Meanwhile some considerations could be made regarding dosing, type of NSBB and titration.^{168,184,185,190} Firstly, dosing of NSBBs was suggested as a potential determinant according to a Danish study in which low propranolol doses (<160 mg/day) were associated with reduced mortality after experiencing an SBP compared to higher doses.^{184,190} Secondly, not all NSBBs proved equal. Carvedilol, which exhibits additional vasodilatory anti-alfa-1-adrenergic activity, might be deleterious in decompensated patients as it is more likely to cause a systemic

haemodynamic depressive effect and may be best avoided or very closely monitored.¹⁸⁵ Thirdly, the concept of titration of NSBBs to a target heart rate of 50–55 bpm might be challenged in decompensated patients given that, in parallel to the progression of liver disease, the hyperdynamic state evolves similarly, which may lead to treatment of the most vulnerable patients paradoxically with higher, and potentially hazardous, doses. Therefore, the use of NSBBs should be based on a critical risk/benefit evaluation in patients with refractory ascites and signs of systemic circulatory dysfunction.^{168,191} Parameters such as severe hyponatraemia,¹⁹¹ low mean arterial pressure³⁸ or cardiac output,¹⁹² and increasing SCr¹⁹³ identify more vulnerable patients among those with decompensated cirrhosis, in whom a dose reduction or temporal discontinuation of NSBB treatment should be considered. The recent BAVENO VI consensus¹⁶⁸ proposed that in patients with refractory ascites and (i) systolic blood pressure <90 mmHg, or (ii) SCr >1.5 mg/dl, or (iii) hyponatraemia <130 mmol/L, the NSBB dose should be reduced or even temporarily discontinued. Abrupt interruption of beta-blockers for a mean of three to six days was recently found to be associated with neither an apparent increase in the risk of variceal bleeding nor with a haemodynamic rebound.¹⁹⁴ If upon rechallenge, NSBB intolerance occurs, EBL should be considered as an alternative in primary prophylaxis.¹⁶⁸ In the setting of refractory ascites and secondary prophylaxis, covered TIPS placement may be considered if the patient is an appropriate candidate.^{111,168}

Recommendations

- Primary prophylaxis must be initiated upon detection of “high-risk varices” (*i.e.* small varices with red signs, medium or large varices irrespective of Child-Pugh classification or small varices in Child-Pugh C patients) because of increased risk of VH (**I;1**).
- Patients with small varices with red wale marks or Child-Pugh C should be treated with NSBBs (**III;1**).
- Patients with medium-large varices should be treated with either NSBBs or EBL (**I;1**). The choice of treatment can be based on local resources and expertise, patient preference, contraindications and adverse events (**III;2**). NSBBs could be preferred because in addition to lowering portal pressure, they also exert other potential beneficial effects (**II-2;2**).
- Although ascites is not a contraindication for NSBBs, caution should be exercised in cases of severe or refractory ascites (**I;1**). High doses of NSBB should be avoided (**II-2;1**). The use of carvedilol can not be recommended at present (**I;2**).
- In patients with progressive hypotension (systolic BP <90 mmHg), or in patients who develop an acute inter-current conditions such as bleeding, sepsis, SBP or AKI, NSBBs should be discontinued (**III,1**). After recovery, reinstatement of NSBBs can be attempted (**III,2**). When NSBB intolerance or contraindications persist, patients bleeding risk should be managed by expeditious EBL (**III,1**).

Recommendations

- Combination therapy of NSBBs + EBL is recommended since it reduces the risk of rebleeding compared with monotherapy (I,1).
- Similar recommendations as for primary prophylaxis can be made with respect to NSBB usage in patients with ascites or developing an acute inter-current condition (III,2).
- If the patient continues to be intolerant to NSBB, covered TIPS placement is recommended provided that there are no absolute contraindication (cf. criteria in ascites section) (III,1).

Variceal haemorrhage

Acute GI bleeding in cirrhosis, either because of gastro-oesophageal varices or non-variceal lesions, is a medical emergency with a high incidence of complications and high mortality and therefore requires intensive care (Fig. 2). Acute variceal haemorrhage (AVH) must be suspected in any cirrhotic patient presenting with upper acute GI bleeding and treatment should be started as soon as bleeding is clinically confirmed, regardless the lack of confirmation by upper endoscopy.¹⁹⁵ Initial therapy should be directed at restoring volaemia.¹⁹⁶ Vasoactive drug therapy^{197,198} and antibiotic prophylaxis^{195,196} should be initiated as soon as AVH is suspected. Goals of therapy in AVH include the control of bleeding, as well as the prevention of early recurrence and the prevention of six-week mortality,

which is considered the main treatment outcome by consensus.^{168,199} Blood volume restitution should be initiated promptly to restore and maintain haemodynamic stability to ensure tissue perfusion and oxygen delivery. To facilitate resuscitation at least two catheters should be placed, large enough to allow rapid volume expansion, which can usually be done with crystalloids.¹⁹⁶ No benefit has been demonstrated with the use of colloids compared to crystalloids.²⁰⁰ Red blood cells are used to improve oxygen delivery to tissues in case of severe anaemia. A restrictive transfusion strategy is adequate in most patients with acute GI bleeding, with a haemoglobin threshold for transfusion of 7 g/dl and a target range after transfusion of 7 to 9 g/dl.²⁰¹ The threshold for transfusion may be higher in patients with massive haemorrhage or in those with underlying conditions that preclude an adequate physiological response to acute anaemia. Recommendations regarding management of coagulopathy and thrombocytopenia cannot be made based on currently available data.^{168,169,199}

As mentioned above vasoactive drug therapy should be initiated as soon as AVH is suspected. Starting vasoactive drugs before endoscopy decreases the incidence of active bleeding during endoscopy and facilitates endoscopic therapy, improving the control of bleeding, and potentially survival.^{197,198} Either terlipressin, somatostatin or octreotide are accepted drugs with proven efficacy.²⁰² All these drugs require i.v administration. The recommended dose of terlipressin is 2 mg/4 h during the first 48 h, followed by 1 mg/4 h thereafter. The recommended dose of somatostatin is a continuous infusion of 250 µg/h (that can be increased up to 500 µg/h) with an initial bolus of 250 µg. The recommended dose of octreotide is a continuous infusion of 50 µg/h with an initial bolus of 50 µg. A bolus of somatostatin

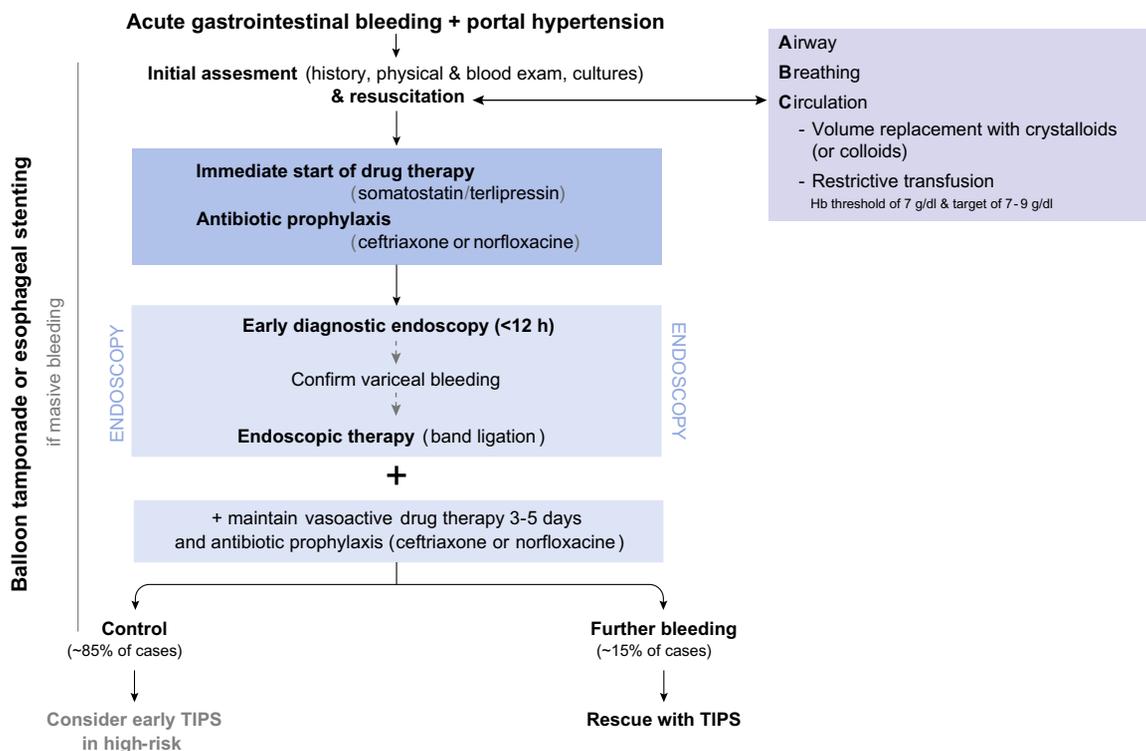


Fig. 2. Algorithm for the management of acute gastrointestinal bleeding in patients with cirrhosis (adapted from Ref. 168). TIPS, transjugular portosystemic shunts.

Clinical Practice Guidelines

or octreotide can be given again if bleeding is ongoing. Once AVH is confirmed, vasoactive drug therapy should be administered for five days to avoid early rebleeding.^{168,169} Shorter administration of vasoactive drugs (48–72 h) can be considered in less severe episodes although more data are required.²⁰³ Once blood volume restitution has been initiated and haemodynamic stability has been achieved, upper endoscopy should be performed, as soon as possible within the first 12 h after admission, to ascertain the cause of haemorrhage (up to 30% of cirrhotic patients bleed from non-variceal causes) and to provide endoscopic therapy if indicated.^{168,169} Erythromycin should be considered before emergency endoscopy (250 mg i.v., 30–120 min before) to facilitate the procedure by improving visibility, in the absence of contraindications (QT prolongation).²⁰⁴ When AVH is confirmed by endoscopy, EBL should be performed within the same procedure. EBL is more effective than sclerotherapy to control bleeding, with fewer adverse effects, and may even improve survival.¹⁹⁶ Sclerotherapy can be used when ligation is not feasible. The combination of endoscopic therapy and vasoactive drugs is more effective than the isolated use of either of these options alone,^{205,206} because it combines the local haemostatic effect on the varices induced by endoscopic treatment and the portal hypotensive effect achieved with drugs. This combination is currently considered the standard of care in AVH.^{168,169} Cyanoacrylate injection and EBL are accepted options for endoscopic therapy in patients bleeding from gastric (cardiofundal) varices as both therapies are equally effective.²⁰⁷ However, EBL should only be performed on small gastric varices in which the complete vessel can be suctioned into the ligation device. Other endoscopic therapies, such as the endoscopic ultrasound-guided insertion of coils and/or cyanoacrylate, are available for fundal varices. Prevention of complications should run simultaneously to haemostatic therapies from admission of patients with cirrhosis and acute GI bleeding. The main complications, whatever the cause of bleeding, include bacterial infections (such as aspiration pneumonia or SBP), hepatic encephalopathy and deterioration of renal function. Bacterial infections are observed in more than 50% of patients and may already be present at the time of bleeding (20%) acting as a precipitating event.¹⁹⁶ Moreover, the presence of bacterial infection is an independent predictor of failure to control bleeding and death.²⁰⁸ Antibiotic prophylaxis is recommended because it reduces the incidence of infections and improves control of bleeding and survival.^{199,208} Ceftriaxone (1 g/24 h) for up to seven days, is the first choice in patients with advanced cirrhosis, in those on quinolone prophylaxis and in hospital settings with high prevalence of quinolone-resistant bacterial infections.^{209,210} Oral quinolones (norfloxacin 400 mg b.i.d) can be used in the remaining patients. These recommendations are however best evaluated and cross-checked from the perspective of local resistance patterns. Renal function should be preserved by the adequate replacement of fluids and electrolytes.²¹¹ Nephrotoxic drugs (such as aminoglycosides and non-steroidal anti-inflammatory drugs [NSAIDs]) as well as LVP, beta-blockers, vasodilators and other hypotensive drugs should be avoided during the course of AVH. Oral non-absorbable disaccharides may be used to prevent the development of hepatic encephalopathy,¹⁶⁹ although more studies are needed. When encephalopathy develops, lactulose or lactitol should be used.^{168,169} Proton pump inhibitors (PPIs) have not shown efficacy for the management of AVH. However, a short course therapy with PPI after EBL may reduce the size of post-banding ulcers.²¹² Despite therapy with vasoactive drugs plus EBL and

prophylactic antibiotics, up to 10–15% of patients with AVH have persistent bleeding or early rebleeding.^{195,199} In such cases, TIPS should be considered as the rescue therapy of choice.^{168,169} When TIPS is not feasible or in case of modest rebleeding, a second endoscopic therapy may be attempted while vasoactive drugs can also be optimised, by doubling the dose of somatostatin and/or changing to terlipressin if not used previously. Balloon tamponade should be used in case of massive bleeding, as a temporary “bridge” until definitive treatment can be instituted and for a maximum of 24 h, preferably under intensive care facilities.^{168,169} Because of the high risk of aspiration pneumonia, tamponade should be preceded by prophylactic orotracheal intubation in comatose or encephalopathic patients. Removable, covered and self-expanding oesophageal stents are an alternative to balloon tamponade, and may have lower rates of serious adverse events.²¹³ RCTs suggest that in high-risk patients, early (preemptive) PTFE-coated TIPS placed within 72 h (ideally in less than 24 h) may result in better permanent control of bleeding and may improve survival.^{214,215} However, these studies had relevant drawbacks such as the inclusion of a highly selected population because of strict exclusion criteria, while observational studies have not confirmed the effect on survival.^{216,217} The use of Child-Pugh class B plus active bleeding at endoscopy as a criterion to select high-risk patients has also been criticised.²¹⁸ It has also been suggested that a recalibrated MELD score may better identify patients at high risk than other scores.²¹⁹ At present, early TIPS should be considered in patients with Child-Pugh class C, with a score <14. However, future studies should clarify which criteria may be preferred to select high-risk patients before a wide implementation of early TIPS. Future studies should also clarify whether an adequate stratification of risk in patients with AVH can optimise therapy.

Recommendations

- Acute GI bleeding, both due to gastro-oesophageal varices or to non-variceal lesions, carries a high incidence of complications and mortality in decompensated cirrhosis and therefore requires close monitoring (**II-2;1**).
- Volume replacement should be initiated promptly to restore and maintain haemodynamic stability (**III;1**). Either colloids and/or crystalloids should be used (**III;1**). Starch should not be used for volume replacement (**I;1**).
- A restrictive transfusion strategy is recommended in most patients with a haemoglobin threshold for transfusion of 7 g/dl and a target range of 7–9 g/dl (**I;1**).
- Antibiotic prophylaxis is recommended in cirrhotic patients with acute GI bleeding because it reduces the incidence of infections and improves control of bleeding and survival. Treatment should be initiated on presentation of bleeding and continued for up to seven days (**I;1**). Ceftriaxone (1 g/24 h) is the first choice in patients with decompensated cirrhosis, those already on quinolone prophylaxis, and in hospital settings with high prevalence of quinolone-resistant bacterial infections. Oral quinolones (norfloxacin 400 mg b.i.d) should be used in the remaining patients (**I;1**).

Recommendations

- Vasoactive drug therapy should be initiated as soon as acute variceal bleeding is suspected, and before endoscopy. Terlipressin, somatostatin or octreotide are accepted options. In patients with acute variceal bleeding drug therapy should be administered for three to five days (**I;1**).
- Gastroscopy should be performed within the first 12 h after admission once haemodynamic stability has been achieved, to ascertain the cause of haemorrhage and to provide endoscopic therapy (**II-2;1**).
- When acute variceal bleeding is confirmed by endoscopy, variceal ligation should be performed within the same procedure (**I;1**).
- In the absence of contraindications (QT prolongation) pre-endoscopy erythromycin (250 mg i.v., 30–120 min before) can be used to facilitate the procedure (**I;2**).
- The combination of vasoactive drugs and ligation is recommended as the first therapeutic option in acute variceal bleeding (**I;1**).
- Early pre-emptive covered TIPS (placed within 24–72 h) can be suggested in selected high-risk patients, such as those with Child class C with score <14 (**I;2**). However, the criteria for high-risk patients, particularly Child B with active bleeding, remains debatable and needs further study.

Recommendations

- Up to 10–15% of patients have persistent bleeding or early rebleeding despite treatment with vasoactive drugs plus variceal ligation, and prophylactic antibiotics. TIPS should be used as the rescue therapy of choice in such cases (**I;1**).
- Balloon tamponade should be used in case of uncontrolled bleeding, but with pre-requisite of expertise and as a temporary “bridge” until definitive treatment can be instituted and for a maximum of 24 h (**III;1**). Removable, covered and self-expanding oesophageal stents can be used as alternative to balloon tamponade (**I;2**).
- In the context of bleeding, where encephalopathy is commonly encountered, prophylactic lactulose may be used to prevent encephalopathy, but further studies are needed (**I;2**).
- Beta-blockers and vasodilators should be avoided during the acute bleeding episode (**III,1**).

Portal hypertension gastropathy and intestinopathy

Portal hypertension gastropathy (PHG) often presents in decompensated patients given that its natural history is significantly influenced by the severity of liver disease and portal hypertension. The presence of oesophageal varices and a Child-Pugh class

B or C at enrollment were found to predict the incidence of PHG, which might range between 30 and 45%.^{220,221} The incidence and severity of PHG may increase following endoscopic treatment for oesophageal varices.²²² Portal hypertension should be distinguished from gastric antral vascular ectasia (GAVE or watermelon stomach), which have different underlying pathophysiologies and different therapeutic implications. The diagnosis of PHG is made by endoscopy and typically shows a snake-skin mosaic pattern (mild subtype), which may have superimposed red signs (severe PHG) and is most commonly located in the proximal stomach (fundus and body) whereas GAVE is characterised by the presence of red spots without a background mosaic pattern, typically located in the gastric antrum.¹⁷⁰ Similar endoscopical lesions, as documented in PHG, may be observed in other areas of the GI tract where they have been termed portal hypertensive duodenopathy, portal hypertensive enteropathy or portal hypertensive colopathy depending on the location of the lesions.²²³ PHG and every form of enteropathy might be clinically important because they are sometimes responsible for insidious blood loss (chronic iron deficient anaemia) and in exceptional cases even overt acute bleeding. When PHG is found as an incidental asymptomatic finding without concomitant oesophageal or gastric varices, its relevance is unclear and endoscopic follow-up or prophylactic treatment is not recommended.¹⁶⁸ First-line therapy for chronic haemorrhage from PHG is an NSBB.^{168,224,225} The same considerations regarding the use of NSBBs in decompensated patients should be made as for gastro-oesophageal varices, except that there is no alternative, endoscopic, standard intervention available for PHG. In addition, iron supplementation should be provided.^{168,226} In patients with medically refractory PHG and compensated cirrhosis, TIPS has shown to improve the endoscopic appearance and decrease the transfusion requirement.²²⁷ In case of acute PHG bleeding, albeit rare, small and uncontrolled studies have suggested pharmacological intervention with somatostatin-analogues or terlipressin because of their portal hypotensive effects and reduction in gastric blood flow.^{226,228} In addition, similar measures are to be taken as for AVH (antibiotic prophylaxis, restrictive transfusion policy). For portal hypertension intestinopathy, there is no established standard of treatment and an approach analogous to that for PHG is suggested. As for any given complication, LT should be considered as part of the management of decompensated patients.

Recommendations

- NSBB and iron supplementation and/or blood transfusion, when indicated, are recommended as first-line therapy for chronic haemorrhage from PHG is an (**I;1**).
- In patients with transfusion-dependent PHG in whom NSBBs fail or are not tolerated, covered TIPS placement may be used provided the patient has no contraindication for TIPS (**II-3;2**).
- Acute PHG bleeding may be treated with somatostatin-analogues or terlipressin but substantiating data are limited (**I;2**).

Gastric varices

The Sarin classification is most commonly used for risk stratification and management of gastric varices (Table 6).²²⁹ Gastric

Clinical Practice Guidelines

Table 6. Classification, prevalence and risk of bleeding of gastric varices.

Type	Definition	Relative frequency	Overall bleeding risk without treatment
GOV 1	OV extending below cardia into lesser curvature	70%	28%
GOV 2	OV extending below cardia into fundus	21%	55%
IGV 1	Isolated varices in the fundus	7%	78%
IGV 2	Isolated varices else in the stomach	2%	9%

GOV, gastro-oesophageal varices; IGV, isolated gastric varices; OV, oesophageal varices.

varices are present in about 20% of patients with cirrhosis. Gastro-oesophageal varices type 1, which are the most common (75% of gastric varices), are oesophageal varices extending below the cardia into the lesser curvature and, in the absence of specific studies, are commonly managed following guidelines for oesophageal varices.¹⁶⁸ Cardiofundal varices (gastro-oesophageal varices type 2 & isolated gastric varices type 1) bleed less frequently. However, haemorrhage from cardiofundal varices is often more severe, more difficult to control and shows a higher risk of recurrent bleeding and mortality (up to 45%) compared to oesophageal varices.²²⁹ Cardiofundal varices are more frequent in patients with splanchnic venous thrombosis, which should be investigated by imaging. The evidence to support recommendations for management of gastric VH is much less robust than that for oesophageal varices. Regarding primary prophylaxis of bleeding from gastric varices, a single randomised trial suggested that cyanoacrylate injection may be more effective than NSBBs in preventing first bleeding in patients with large cardiofundal varices, although survival was similar.²³⁰ Therefore, the last BAVENO consensus concluded that further studies are needed to evaluate the risk/benefit ratio of using cyanoacrylate in this setting before a formal recommendation can be made and meanwhile propose NSBBs as the primary approach.¹⁶⁸ Acute gastric VH is medically treated like bleeding oesophageal varices. However, injection therapy with cyanoacrylate ('glue') may be the preferable option for endoscopic haemostasis.²³¹ Although equally effective as EBL in initial haemostasis, the rebleeding rate is significantly lower.²³² TIPS, with or without additional embolisation of collaterals, is equally effective in gastric and oesophageal VH for control of acute bleeding events and prevention of rebleeding.²³³ In case of massive bleeding, balloon tamponade with the Linton-Nachlas tube may serve as a bridge to other treatments. Regarding secondary prophylaxis, in one RCT repeated cyanoacrylate injection was superior to NSBBs to prevent rebleeding from cardiofundal varices,²³² while the addition of NSBBs to cyanoacrylate did not improve the outcomes achieved with glue alone in another RCT.²³⁴ Another trial comparing TIPS to glue injection showed that TIPS proved more effective in preventing rebleeding from gastric varices, with similar survival and frequency of complications.²³⁵ The option of early TIPS should be strongly considered, particularly in cardiofundal varices given the high rebleeding rate, provided that patient is an appropriate candidate for such a procedure. Alternatively, balloon-occluded retrograde transvenous obliteration (BRTO) can be considered. This interventional radiological procedure enables treatment of fundal varices associated with a large gastro/splenorenal collaterals, which has the theoretical advantage over TIPS of not diverting portal blood inflow from the liver. However, no

randomised trials are available comparing BRTO with other therapies. Several variations of this technique are available, such as balloon-occluded antegrade transvenous obliteration (BATO).²³⁶

Recommendations

- NSBBs are suggested for primary prevention of VH from gastro-oesophageal varices type 2 or isolated gastric varices type 1 (**III;2**).
- Primary prevention for gastro-oesophageal varices type 1 follow the recommendations of oesophageal varices (**III;2**).
- Acute gastric VH should be treated medically, like oesophageal VH (**I;1**). Cyanoacrylate is the recommended endoscopic haemostatic treatment for cardiofundal varices (gastro-oesophageal varices type 2 or isolated gastric varices type 1) (**I;2**).
- TIPS with potential embolisation efficiently controls bleeding and prevents rebleeding in fundal VH (gastro-oesophageal varices type 2 or isolated gastric varices type 1) and should be considered in appropriate candidates (**II-2;1**).
- Selective embolisation (BRTO/BATO) may also be used to treat bleeding from fundal varices associated with large gastro/splenorenal collaterals, although more data is required (**III;2**).

Bacterial infections

The risk of bacterial infection in cirrhosis is caused by multiple factors that include liver dysfunction, portosystemic shunting, gut dysbiosis, increased BT, cirrhosis-associated immune dysfunction,^{237,238} and genetic factors. This immune defect facilitates BT, induced by increased intestinal permeability and gut bacterial overgrowth observed in cirrhosis.²³⁹ Genetic immune defects can contribute to the high risk of bacterial infections in cirrhosis, particularly SBP. Cirrhotic patients carrying *NOD2* variants associated with impaired recognition of the bacterial product muramyl dipeptide have a higher risk of SBP and a reduced survival time.²⁴⁰

Spontaneous bacterial peritonitis

Definition

Spontaneous bacterial peritonitis has been defined as a bacterial infection of ascitic fluid without any intra-abdominal surgically treatable source of infection. SBP is very common in patients with cirrhosis and ascites.^{241,242} When first described, its mortality exceeded 90% but it has been reduced to approximately 20% with early diagnosis and treatment.²⁴³

Diagnosis

The diagnosis of SBP is based on diagnostic paracentesis.^{33,244} All patients with cirrhosis and ascites are at risk of SBP and the prevalence of SBP in outpatients is 1.5–3.5% and ~10% in hospitalised patients.²⁴⁵ Half the episodes of SBP are present at the time of hospital admission while the rest are acquired during hospitalisation.³³ Patients with SBP may have one of the following:³³ i) local symptoms and/or signs of peritonitis:

abdominal pain, abdominal tenderness, vomiting, diarrhoea, ileus; ii) signs of systemic inflammation: hyper or hypothermia, chills, altered white blood cell count, tachycardia, and/or tachypnoea; iii) worsening of liver function; iv) hepatic encephalopathy; v) shock; vi) renal failure; and, vii) GI bleeding. However, it is important to point out that SBP may be asymptomatic, particularly in outpatients.²⁴⁵ In an observational study in 239 patients with SBP, delayed diagnostic paracentesis (>12 h after admission) was associated with a 2.7-fold increase in mortality.²⁴⁶ Peritoneal infection causes an inflammatory reaction resulting in an increased number of neutrophils in ascitic fluid. Despite the use of sensitive methods, ascites culture is negative in as many as 60% of patients with clinical manifestations suggestive of SBP and increased ascites neutrophil count.³³ The gold standard for ascitic neutrophil count is manual microscopy, but it is labour intensive and associated with interobserver variability, time and costs. In most places this has been substituted with automated counts based on flow cytometry for counting and differentiating cells. This technique has been documented to have high linearity with manual microscopy and thus sensitivity and specificity close to 100%.^{247,248} The greatest sensitivity for the diagnosis of SBP is reached with a cut-off neutrophil count of 250/mm³, although the greatest specificity is reached with a cut-off of 500 neutrophils/mm³.³³ The use of reagent strips cannot be recommended for the rapid diagnosis of SBP.²⁴⁹ Although the presence of bacterial DNA in plasma and/or ascites is associated with an impairment of circulatory function,²⁵⁰ there are not enough data to support its use in clinical practice.²⁵¹ Ascites culture is essential to guide antibiotic therapy. Patients with an ascitic fluid neutrophil count ≥ 250 cells/mm³ and negative culture have culture-negative SBP.²⁵² Their clinical presentation is like that of patients with culture-positive SBP and should be treated in a similar manner. Some patients have 'bacterascites' in which cultures are positive but there is normal ascitic neutrophil count (<250/mm³).³³ In some patients bacterascites is the result of secondary bacterial colonisation of ascites from an extraperitoneal infection. These patients usually have general symptoms and signs of infection. In other patients, bacterascites is due to the spontaneous colonisation of ascites, and can either be clinically asymptomatic or lead to abdominal pain or fever. While in some patients, particularly in those who are asymptomatic, bacterascites represents a transient and spontaneously reversible colonisation of ascites, in other patients, mainly those who are symptomatic, bacterascites may represent the first step in the development of SBP.³³ Spontaneous fungal peritonitis is a rare, less recognised and studied complication, occurring in <5% of cases, but observational data suggest a worse prognosis.²⁵³

Spontaneous bacterial pleural empyema

Infection of a pre-existing hydrothorax, known as spontaneous bacterial pleural empyema, is uncommon. One study followed 3,390 patients with cirrhosis for four years and observed it in 2.4% of the overall population and 16% of patients with pre-existing hydrothorax, with associated mortality of 38%.²⁵⁴ The diagnosis is based on pleural fluid analysis obtained by diagnostic thoracentesis. In the largest observational study reported so far, the diagnosis of spontaneous bacterial empyema was established when the pleural fluid analysis showed a positive culture and more than 250 neutrophils/mm³ or a negative culture and more than 500 neutrophils/mm³, in the absence of lung infection.²⁵⁵ Pleural fluid culture in blood culture bottles was

positive in 75% of cases.²⁵⁵ Spontaneous bacterial pleural empyema was associated with SBP in ~50% of cases.²⁵⁵

Secondary bacterial peritonitis

A small proportion (~5%) of patients with cirrhosis may develop peritonitis due to perforation or inflammation of an intra-abdominal organ, a condition known as secondary bacterial peritonitis.²⁵⁶ The differentiation of this condition from SBP is important. Secondary bacterial peritonitis should be suspected in patients who have localised abdominal symptoms or signs, presence of multiple organisms on ascitic culture, very high ascitic neutrophil count and/or high ascitic protein concentration, or in those patients with an inadequate response to therapy.²⁵⁶ Patients with suspected secondary bacterial peritonitis should undergo prompt computed tomography (CT) scanning and early consideration for surgery.

Recommendations

- A diagnostic paracentesis should be carried out in all patients with cirrhosis and ascites without delay at hospital admission to rule out SBP. A diagnostic paracentesis should also be performed in patients with GI bleeding, shock, fever or other signs of systemic inflammation, GI symptoms, as well as in patients with worsening liver and/or renal function, and hepatic encephalopathy (**II-2;1**).
- The diagnosis of SBP is based on neutrophil count in ascitic fluid of >250/mm³ (**II-2;1**). Neutrophil count is determined by microscopy, but can be substituted with a flow cytometry based automated count. The use of reagent strips has no clear evidence to support it in routine practice (**II-2;1**).
- Although ascitic fluid culture positivity is not a prerequisite for the diagnosis of SBP, culture should be performed in order to guide antibiotic therapy (**II-2;1**).
- Blood cultures should be performed in all patients with suspected SBP before starting antibiotic treatment (**II-2;1**).
- Patients with bacterascites (neutrophil count less than 250/mm³ but positive bacterial culture) exhibiting signs of systemic inflammation or infection should be treated with antibiotics (**II-2;1**). Otherwise, the patient should undergo a second paracentesis. If the culture results come back positive again, regardless of the neutrophil count, the patient should be treated (**III;1**).
- The diagnosis of spontaneous bacterial pleural empyema should be based on positive pleural fluid culture and increased neutrophil count of >250/mm³ or negative pleural fluid culture and a neutrophil count of >500/mm³ in the absence of pneumonia (**II-2;1**).
- Secondary bacterial peritonitis should be suspected in case of multiple organisms on ascitic culture, very high ascitic neutrophil count and/or high ascitic protein concentration, or in those patients with an inadequate response to therapy. Patients with suspected secondary bacterial peritonitis should undergo prompt CT scanning and early considerations for surgery (**III;1**).

Clinical Practice Guidelines

Management of spontaneous bacterial peritonitis

Empirical antibiotic therapy. Empirical antibiotic therapy must be initiated immediately after the diagnosis of SBP.³³ Potentially nephrotoxic antibiotics (*i.e.*, aminoglycosides) should not be used as empirical therapy.⁷⁶ In the 1990 s, cefotaxime, a third-generation cephalosporin, was extensively investigated in patients with SBP because at that time it covered most causative organisms and because of its high ascitic fluid concentrations during therapy.^{1,33} Infection resolution was obtained in 77 to 98% of patients. A dose of 4 g/day is as effective as a dose of 8 g/day.²⁵⁷ A five-day therapy is as effective as a 10-day treatment.²⁵⁸ Alternatively, amoxicillin/clavulanic acid, first given *i.v.* then orally, has similar results with respect to SBP resolution and mortality as cefotaxime²⁵⁹ and at a much lower cost. However, there is only one comparative study with a small sample size and results should be confirmed in larger trials. In addition, some concern exists regarding amoxicillin/clavulanic acid as its use is associated with a high rate of drug induced liver injury (DILI).²⁶⁰ Administration of *i.v.* ciprofloxacin for seven days results in a similar SBP resolution rate and hospital survival as cefotaxime, but at a significantly higher cost.²⁶¹ However, switch therapy (*i.e.*, use of *i.v.* antibiotic initially, followed by oral step-down administration) with ciprofloxacin is more cost-effective than *i.v.* ceftazidime.²⁶² Oral ofloxacin has shown similar results as *i.v.* cefotaxime in uncomplicated SBP, without renal failure, hepatic encephalopathy, GI bleeding, ileus, or shock.²⁶³ However, the spread of resistant bacteria in the healthcare environment during the last two decades has led to an alarming increase in the number of infections caused by multi-drug resistant organisms (MDROs)²⁶⁴ that are defined by an acquired non-susceptibility to at least one agent in three or more antimicrobial categories.²⁶⁵ Patients with advanced cirrhosis are highly susceptible to the development of infections caused by MDROs, because they require repeated hospitalisations, are often submitted to invasive procedures and are frequently exposed to antibiotics, either as prophylaxis or as treatment. All these factors are well known risk factors for the development of infections sustained by MDROs.²⁶⁶ Bacterial resistance increases four fold the risk of mortality of SBP.²⁶⁷ In particular, nosocomial SBP has been associated with multi-drug resistance and poor outcomes.²⁶⁶ The landscape of bacterial resistance is continuously changing and challenging recommendations for antibiotics. Thus, it is crucial to separate commu-

nity-acquired SBP from health care-associated and nosocomial SBP^{6,266–268} and to consider both the severity of infection and the local resistance profile in order to decide the empirical antibiotic treatment of SBP. Piperacillin/tazobactam has been recommended as the primary approach for health care and nosocomial SBP in areas with low prevalence of infections sustained by MDROs. On the contrary meropenem alone or/and combined with glycopeptides or with daptomycin has been suggested as the primary approach for health care-associated SBP when severe, or in areas with high prevalence of MDROs, and for nosocomial SBP in general.^{6,266,268,269} Regarding the severity of infection, it should be highlighted that, recently, the new criteria for the definition of sepsis, namely qSOFA and Sepsis-3²⁷⁰ have been validated in patients with cirrhosis and bacterial infections, proving that they are more accurate than those related to the systemic inflammatory response syndrome in predicting hospital mortality.²⁷¹ Accordingly, a new algorithm has been proposed for the application of qSOFA and Sepsis-3 in the management of cirrhotic patients (Fig. 3). Some more detailed recommendations on the empirical antibiotic treatment of SBP based on the severity and the environment of the infection as well as on local resistance profiles are provided (Fig. 4). A randomised trial with 32 nosocomial episodes of SBP, found meropenem plus daptomycin more effective (86.7%) than ceftazidime (25%) to manage SBP, defined as >25% decrease of neutrophil count at 48 h and to <250/mm³ at day seven.²⁴³ If ascitic fluid neutrophil count fails to decrease to less than 25% of the pretreatment value after two days of

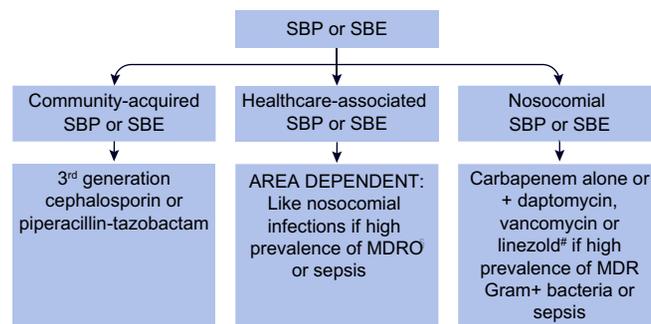


Fig. 4. Recommended empirical antibiotic treatment of SBP or SBE (adapted from Ref. 6). SBE, spontaneous bacterial empyema; SBP, spontaneous bacterial peritonitis; MDRO, multidrug resistant organism.

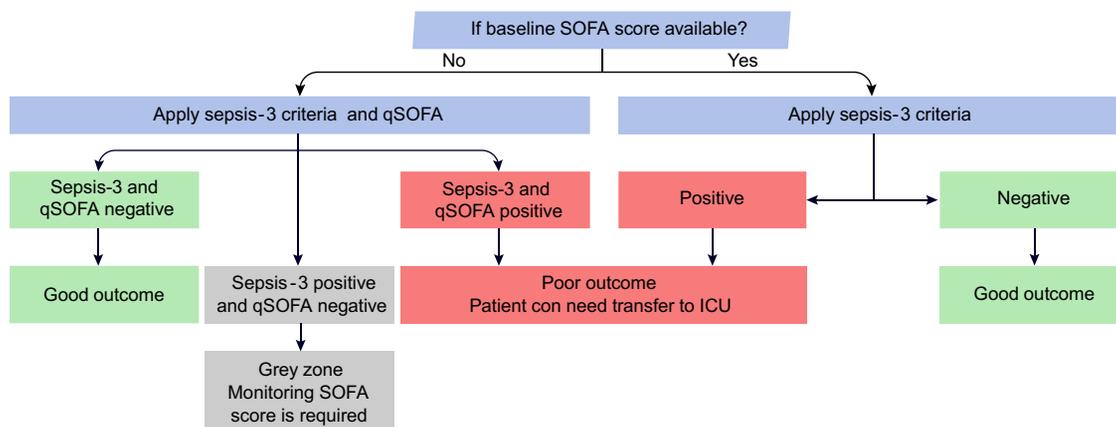


Fig. 3. Algorithm for the application of qSOFA and Sepsis-3 criteria in patients with cirrhosis and bacterial infections (adapted from Ref. 271). ICU, intensive care unit.

antibiotic treatment, there is a high likelihood of failure to respond to therapy.³³ This should raise the suspicion of an infection caused by bacteria resistant to antibiotic therapy, indicating the need for modification of antibiotic treatment according to *in vitro* sensitivity or on an empirical basis, or the presence of 'secondary peritonitis'. In this context, it should be highlighted that the progressive increase of the use of carbapenems because of the worldwide pandemic of extended spectrum beta-lactamases (ESBLs) producing *Enterobacteriaceae* has promoted the emergence of carbapenem-resistant *Enterobacteriaceae*. This implies a potential shift from MDR bacteria to extensively drug resistance (XDR) bacteria defined by a non-susceptibility to at least one agent in all but two or fewer antimicrobial categories or to pandrug resistance (PDR) bacteria defined by a non-susceptibility to all agents in all antimicrobial categories.²⁶⁵ The shift requires an active surveillance in patients at risk, in order to identify patients who are colonised or infected by these clones and prevent their dissemination. The shift may also seriously affect the effectiveness of the broadest spectrum empirical antibiotic treatment among those previously recommended for SBP and infections other than SBP. Carbapenemase-producing and carbapenem-resistant non-carbapenemase-producing *Enterobacteriaceae* can be treated with tigecycline or with the combination of tigecycline at high doses and a carbapenem in continuous infusion. Addition of i.v. colistin could be necessary in severe infections. Severe infections caused by *Pseudomonas aeruginosa* resistant to carbapenems and quinolones usually require the combination of i.v. amikacin/tobramycin or colistin plus a carbapenem/ceftazidime (needed as synergic antibiotics despite antibiotic resistance). Vancomycin resistant *Enterococci* should be treated with linezolid, daptomycin or tigecycline. All this means reintroducing into clinical practice antibiotics known to be highly nephrotoxic in patients with cirrhosis. It follows that serum levels of aminoglycosides and vancomycin must be monitored closely in these patients, to decrease the risk of renal failure. The shift from MDR to XDR bacteria re-emphasises the interest of the pharmaceutical industry for the development of new antibiotics. Several new glycopeptides such as oritavancin, new oxazolidinones such as tedizolid phosphate, new cephalosporins, such as ceftaroline and ceftobiprole and razupenem, a new carbapenem, display extended activity against gram-positive bacteria including vancomycin-resistant *Enterococci*. In contrast, few newly developed antibiotics are active against gram-negative MDROs. Temocillin, a derivative of ticarcillin, is effective against organisms producing ESBLs. Among cephalosporin-beta-lactamase inhibitor combinations, ceftazidime/avibactam and ceftolozane/tazobactam represent further new alternatives to carbapenems for the treatment of patients with infections sustained by ESBL producing, carbapenem-resistant *Enterobacteriaceae* and *Pseudomonas aeruginosa*. However, there are currently no data regarding the clinical use of these drugs in cirrhosis.²⁶⁶

Recommendations

- Empirical i.v. antibiotics should be started immediately following the diagnosis of SBP (II-2;1).
- Environment (nosocomial vs. community acquired), local bacterial resistance profiles and severity of infection should guide empirical antibiotic treatment (I;1).

- Third-generation cephalosporins are recommended as first-line antibiotic treatment for community-acquired SBP in countries with low rates of bacterial resistance (I;1). In countries with high rates of bacterial resistance piperacillin/tazobactam or carbapenem should be considered (II-2;1).
- Healthcare associated and nosocomial SBP is more likely to harbour resistance to antibiotics. Piperacillin/tazobactam should be given in areas with low prevalence of multi-drug resistance while carbapenem should be used in areas with high prevalence of ESBL producing *Enterobacteriaceae*. Carbapenem should be combined with glycopeptides or daptomycin or linezolid in areas with high prevalence of gram positive MDR bacteria (I;1).
- Severe infections sustained by XDR bacteria may require the use of antibiotics known to be highly nephrotoxic in patients with cirrhosis, such as vancomycin or aminoglycosides. In these cases, patients' plasma level should be monitored in accordance with local policy thresholds (III;1).
- De-escalation according to bacterial susceptibility based on positive cultures is recommended to minimise resistance selection pressure (II-2;1).
- The efficacy of antibiotic therapy should be checked with a second paracentesis at 48 h from starting treatment. Failure of first-line antibiotic therapy should be suspected if there is worsening of clinical signs and symptoms and/or increase or no marked reduction in leucocyte count (at least 25%) in 48 h (II-2;1).
- The duration of treatment should be at least 5–7 days (III;1).
- Spontaneous bacterial empyema should be managed similarly to SBP (II-2;2).

Intravenous albumin in patients with spontaneous bacterial peritonitis. SBP without septic shock may precipitate deterioration of circulatory function with severe liver failure, hepatic encephalopathy, and type 1 HRS and has approximately 20% hospital mortality despite infection resolution.²⁷² A randomised, controlled study in patients with SBP treated with cefotaxime showed that albumin (1.5 g/kg body weight at diagnosis, followed by 1 g/kg on day three) significantly decreased the incidence of type 1 HRS (from 30% to 10%) and reduced mortality from 29% to 10% compared with cefotaxime alone. Treatment with albumin was particularly effective in patients with baseline serum bilirubin $\geq 68 \mu\text{mol/L}$ (4 mg/dl) or SCr $\geq 88 \mu\text{mol/L}$ (1 mg/dl). It is unclear whether i.v. albumin is useful in patients with baseline bilirubin $< 68 \mu\text{mol/L}$ and creatinine $< 88 \mu\text{mol/L}$, as the incidence of type 1 HRS in patients meeting these criteria was very low in the two treatment groups (7% without albumin and 0% with albumin).²⁷² The application of the schedule of this therapeutic option should be implemented in clinical practice.²⁷³ Non-randomised studies in patients with SBP also show that the incidence of renal failure and death are very low in patients with moderate liver failure and without renal dysfunction at diagnosis of SBP, so albumin is probably not necessary.²⁷⁴

Recommendation

- The administration of albumin (1.5 g/kg at diagnosis and 1 g/kg on day 3) is recommended in patients with SBP (**I;1**).

Prophylaxis of SBP

Since most episodes of SBP are thought to result from the translocation of enteric gram-negative bacteria, the ideal prophylactic agent should be safe, affordable and effective at decreasing the amounts of these organisms from the gut while preserving the protective anaerobic flora (selective intestinal decontamination).²⁶⁷ Given the high cost and inevitable risk of developing resistant organisms, the use of prophylactic antibiotics must be strictly restricted to patients at high risk of SBP.²⁶⁷ Three high-risk patient populations have been identified: i) patients with acute GI haemorrhage; ii) patients with low total protein content in ascitic fluid and no prior history of SBP (primary prophylaxis), and iii) patients with a previous history of SBP (secondary prophylaxis).²⁷⁵

Primary prophylaxis in patients with low total protein content in ascitic fluid without prior history of SBP.

Cirrhotic patients with low ascitic fluid protein concentration (<10 g/l) and/or high serum bilirubin levels are at high risk of developing a first episode of SBP.²⁶⁷ Several studies have evaluated prophylaxis with norfloxacin in patients without prior history of SBP.²⁶⁷ Fernandez *et al.* randomised 68 patients with cirrhosis and low ascites protein levels (<15 g/l) with advanced liver failure (Child-Pugh score ≥ 9 points with serum bilirubin level ≥ 3 mg/dl) or impaired renal function (SCr level ≥ 1.2 mg/dl, blood urea nitrogen level ≥ 25 mg/dl, or serum sodium level ≤ 130 mEq/L) to receive norfloxacin (400 mg/day for 12 months) or placebo.²⁷⁶ Norfloxacin significantly improved the three-month probability of survival (94% vs. 62%; $p = 0.03$) but at one year the difference in survival was not significant (60% vs. 48%; $p = 0.05$). Norfloxacin administration significantly reduced the one-year probability of developing SBP (7% vs. 61%) and HRS (28% vs. 41%). In a double-blind placebo-controlled trial, 100 patients with ascitic fluid total protein level <15 g/l were randomised to ciprofloxacin (500 mg/day for 12 months) or placebo.²⁷⁷ The probability of survival at one year was higher in patients receiving ciprofloxacin (86% vs. 66%; $p < 0.04$). Meta-analyses of all the trials together or including only pure primary prophylaxis support a significant preventive effect against SBP (RR 0.2; 95% CI 0.07–0.52; $p = 0.001$).^{278,279} The survival benefit is most pronounced at three months (94% vs. 62%, $p = 0.003$) and seems to decrease over time and may be lost after 12 months follow-up (RR 0.65; 95% CI; 0.41–1.02).²⁸⁰

Recommendations

- Primary prophylaxis with norfloxacin (400 mg/day) in patients with Child-Pugh score ≥ 9 and serum bilirubin level ≥ 3 mg/dl, with either impaired renal function or hyponatraemia, and ascitic fluid protein lower than 15 g/L is recommended (**I;1**).
- Norfloxacin prophylaxis should be stopped in patients with long-lasting improvement of their clinical condition and disappearance of ascites (**III;1**).

Patients with prior SBP

In patients who survive an episode of SBP, the cumulative recurrence rate at one year is approximately 70%.³³ Probability of survival at one year after an episode of SBP is 30–50% and falls to 25–30% at two years. Therefore, patients recovering from an episode of SBP should be considered for LT. There is only one randomised, double-blind, placebo-controlled trial of norfloxacin (400 mg/day orally) in patients who had a previous episode of SBP.²⁸¹ Treatment with norfloxacin reduced the probability of recurrence of SBP from 68% to 20%. In an open-label, randomised study comparing norfloxacin 400 mg/day to rifloxacin 400 mg/week in the prevention of SBP recurrence, the one-year probability of SBP recurrence was 26% and 36%, respectively ($p = 0.16$).²⁸² Norfloxacin was more effective in the prevention of SBP recurrence due to *Enterobacteriaceae* (0% vs. 22%, $p = 0.01$). The use of intermittent ciprofloxacin has been associated with a higher rate of quinolone-resistant organisms and should be avoided.^{282,283} It is uncertain whether prophylaxis should be continued without interruption until LT or death in all patients with prior SBP, or if treatment could be discontinued in patients showing an improvement of liver disease. Many patients receive rifaximin to prevent recurrent episodes of HE.²⁸⁴ However, rifaximin may also be effective against recurrent SBP.²⁸⁵ There are no data to guide new indications for primary or secondary prophylaxis of SBP among patients already on rifaximin. More in detail, it is not known whether norfloxacin prophylaxis should be started in patients being treated with rifaximin for prevention of recurrent HE. Likewise, it is not known whether norfloxacin prophylaxis should be stopped in patients who would require rifaximin to prevent HE. Prospective studies are required to investigate the potential benefits and side effects of combined therapy with norfloxacin and rifaximin.

Recommendations

- The administration of prophylactic Norfloxacin (400 mg/day, orally) is recommended in patients who recover from an episode of SBP (**I;1**).
- Despite some promising evidence, at present, rifaximin cannot be recommended as an alternative to norfloxacin for secondary prophylaxis of SBP (**I;2**). Thus, at present, no recommendation can be given to guide primary or secondary prophylaxis of SBP among patients already on rifaximin for the prevention of recurrent HE.
- Patients who recover from SBP have a poor long-term survival and should be considered for LT (**II-2,1**).
- Since it has been suggested that PPI may increase the risk for the development of SBP, its use should be restricted to those with a clear indication (**II-2,1**).

Concomitant medications

Very frequently PPIs are used in patients with cirrhosis, which may increase the risk of SBP. Indications for long-term use should be carefully assessed and PPIs discontinued when possible.^{286,287} NSBBs may be detrimental in end-stage liver disease with haemodynamic derangement, patients should be monitored closely and doses adjusted or drug discontinued if contraindications occur.^{168,190,288} Probiotics have been assessed as

combination therapy with norfloxacin in one randomised trial in a mixed group of patients on primary and secondary prevention of SBP. No additional benefits were demonstrated.²⁸⁹

Infections other than SBP

Prevalence, diagnosis and impact on prognosis

Non-SBP infections are frequent in patients with cirrhosis and present or develop during hospitalisation in 25–30% of patients. The most frequent infections other than SBP are: urinary tract, pneumonia, skin and soft tissue infections, and bacteraemia.^{242,290} They constitute a heterogeneous group regarding clinical course and prognosis. Non-SBP infections increase the odds ratio for death by 3.75 and are associated with a 30% one-month and 63% 12-month mortality.²⁹¹ Endocarditis, secondary peritonitis, pneumonia and bacteraemia have worse prognoses. The combination of data on liver and renal dysfunction and the type of infection enables the identification of patients with poor prognosis.²⁹⁰ In particular, non-SBP infections, as well as SBP, are known as common precipitating factors for ACLF.³ An early diagnosis of all these infections and of SBP is a crucial step in the management of patients with cirrhosis. Since the presentation and the initial course of any bacterial infection may be subtle and not very specific, clinical suspicion is important. Indeed, all inpatients with cirrhosis should be considered as potentially infected until proven otherwise. Therefore, a complete work-up should be carried out at admission and at any time during the hospital stay when clinical deterioration occurs.⁶ In addition, close microbiological surveillance is needed in patients who are at risk of developing infections caused by methicillin resistant organisms. C reactive protein and procalcitonin can be used for detecting infection and to define the severity of the infection,⁶ while their use in the stewardship of antibiotic treatment deserves further investigation.²⁹² To optimise the empirical antibiotic treatment, it is quite important to distinguish among community acquired, health care associated and nosocomial infections. Mortality for nosocomial infections is higher (25–48%) than for community-acquired infection (7–21%) since they are more commonly sustained by MDR bacteria.^{6,266,268} Like in SBP, there is an increasing challenge of resistant bacteria among non-SBP infections. Among 312 patients with cirrhosis and blood stream infections gram-negative bacteria, gram-positive bacteria and *Candida* were the cause of blood stream infections episodes in 53%, 47% and 7% of the cases, and 31% of the infections were caused by MDR bacteria.²⁹³

Management of infections other than SBP

In a randomised trial 94 patients with cirrhosis and infections (most prevalent were urinary tract infections [46%], SBP [22%], and pneumonia [19%]) were randomised to a broad-spectrum antibiotic regimen or a standard regimen. In-hospital mortality was significantly higher in the standard than in the broad-spectrum group (25% vs. 6%; $p = 0.01$).²⁹⁴ Some more specific suggestions on the empirical antibiotic treatment of infections other than SBP based on the type, the severity and the environment of the infection as well as on local resistance profiles are given (Figs. 5–7). In patients who fail to respond to a broad-spectrum antibiotic treatment a fungal infection, including fungal SBP²⁹⁵ should be suspected and investigated.²⁹⁶

Finally, in two randomised trials, concomitant albumin may protect against deterioration in renal and circulatory function.^{297,298} However, albumin did not improve survival and thus

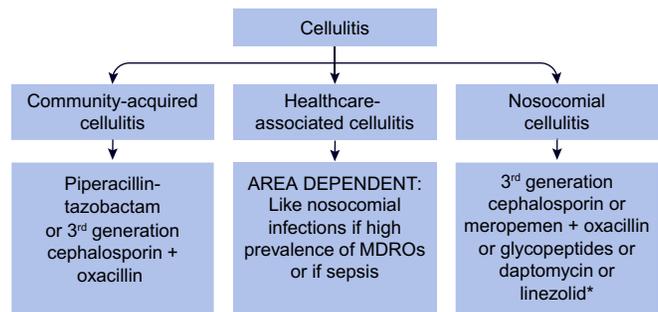


Fig. 5. Recommended empirical antibiotic treatment of soft tissue infections (adapted from Ref. 6). MDRO, multidrug resistant organism.

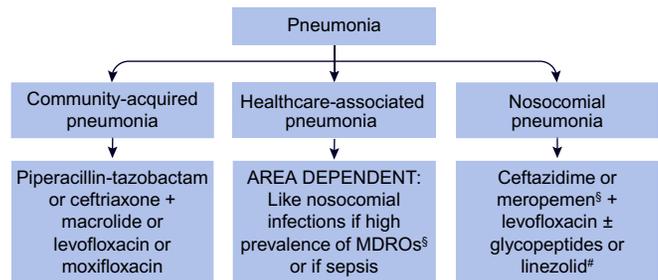


Fig. 6. Recommended empirical antibiotic treatment of pneumonia (adapted from Ref. 6). MDRO, multidrug resistant organism.

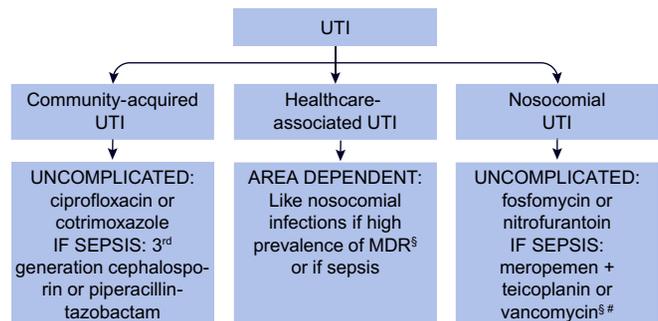


Fig. 7. Recommended empirical antibiotic treatment of UTI (adapted from Ref. 6). MDR, multidrug resistant; UTI, urinary tract infection.

it cannot be recommended. The issue of the management of infections sustained by XDR bacteria has been previously developed.

Recommendations

- Infections other than SBP are frequent and associated with increased mortality. Hospitalised patients with cirrhosis should be assessed and monitored closely for the presence of infections to enable early diagnosis and appropriate treatment (II-1;1).
- Empirical antibiotic therapy should be commenced promptly at suspicion of infection (II-1;1).
- The choice of empirical antibiotic therapy should be based on several factors including: environment (nosocomial vs. health care associated or community acquired), local resistance profiles, severity and type of infection (I;1).

- In the context of high bacterial resistance to antibiotics, carbapenem alone or in combination with other antibiotics proved to be superior to third-generation cephalosporins in healthcare associated infections other than SBP, and therefore, should be preferred (**I;1**).
- Severe infections sustained by XDR bacteria can require the use of antibiotics known to be highly nephrotoxic in patients with cirrhosis such as vancomycin or aminoglycosides. In these cases, patients' plasma level should be monitored in accordance with local policy thresholds (**III;1**).
- Routine use of albumin is not recommended in infections other than SBP (**I;1**).

Prophylaxis of infections other than SBP

There is preliminary evidence that in patients with Child-Pugh class C, norfloxacin administration can reduce the risk of infections and can decrease six-month mortality. However, more data are needed before a recommendation can be made.²¹

Renal impairment

Definition and diagnosis

Renal impairment in patients with cirrhosis was defined more than 30 years ago by an sCr value ≥ 1.5 mg/dl because this value was considered an index of GFR ≤ 40 ml/min.³² The use of sCr in the evaluation of renal function in patients with cirrhosis, has several well-known limitations. However, the diagnosis of renal dysfunction in liver disease is still based on it.^{32,299} The diagnosis should be based on different diagnostic categories including chronic kidney disease (CKD) and acute renal failure (ARF). When only based on reduction of GFR, the diagnosis of CKD in patients with cirrhosis is still challenging, because all the sCr-based equations that have been proposed overestimate GFR in patients with cirrhosis.^{300–304} It can be reasonably assumed that patients with decompensated cirrhosis frequently have CKD caused by certain comorbidities (*i.e.* diabetes, arterial hypertension) and/or specific causes (*i.e.* IgA nephropathy, virus-induced

glomerulopathy),³⁰⁵ however the prevalence of CKD in this population is still unknown. ARF is a common complication in patients with decompensated cirrhosis.³⁰⁶ Historically, the diagnosis was based on an increase in sCr of 50% from baseline to a final value >1.5 mg/dl ($133 \mu\text{mol/L}$).^{1,32,307} Recently, the term ARF was replaced by AKI,^{308–310} irrespectively of its different types. AKI is now defined, as proposed by the Kidney Disease Improving Global Outcomes (KDIGO) group,³¹⁰ as either an absolute increase in sCr of more than or equal to 0.3 mg/dl ($\geq 26.4 \mu\text{mol/L}$) in less than 48 h, or by a percentage increase in sCr of more or equal to 50% (1.5-fold from baseline) in less than seven days. A new staging system was also introduced, mainly based on the percentage increase of sCr from baseline (Table 7), either at the time of the first fulfillment of the KDIGO criteria (initial stage) or at the peak value of sCr during hospitalisation in case of progressive AKI (peak stage).³¹⁰ Based on the staging system and according to the results of several prospective studies,^{311–317} a new algorithm for the management of AKI in patients with cirrhosis has been proposed³¹⁸ (Fig. 8). Recent studies have suggested that in patients with cirrhosis, in AKI stage 1, sCr <1.5 mg/dl is associated with a worse outcome than an sCr ≥ 1.5 mg/dl.^{313,314,317} Thus, in contrast with the KDIGO staging system, it has been proposed to distinguish between a stage 1A (sCr <1.5 mg/dl) and a stage 1B (sCr ≥ 1.5 mg/dl) within AKI stage 1.^{313,314,317} It should be highlighted that the KDIGO criteria also include criteria based on urinary output in the diagnosis of AKI (Fig. 9).³¹⁰ These criteria were not considered by the recent International Club of Ascites (ICA) consensus because (a) these patients are frequently oliguric with avid sodium retention, despite a relatively normal GFR, (b) they may have an increased urine output because of diuretics, and (c) on a regular ward, urine collection is often inaccurate and always untimely.³¹⁸ However, these criteria may also be applied whenever a patient with cirrhosis requires a bladder catheter. The definition of baseline sCr used in the KDIGO criteria is crucial since it has been observed that about 25–30% of episodes of AKI occur before hospitalisation, representing the so-called “community-acquired AKI”. Ideally, “community-acquired AKI” should be diagnosed at the time of hospital admission, requiring, according to the KDIGO criteria, an sCr value dated within

Table 7. International Club of Ascites (ICA-AKI) new definitions for the diagnosis and management of acute kidney injury in patients with cirrhosis.

Subject	Definition		
Baseline sCr	A value of sCr obtained in the previous three months, when available, can be used as baseline sCr. In patients with more than one value within the previous three months, the value closest to the admission time to the hospital should be used. In patients without a previous sCr value, the sCr on admission should be used as baseline.		
Definition of AKI	- Increase in sCr ≥ 0.3 mg/dl ($\geq 26.5 \mu\text{mol/L}$) within 48 h; or, - A percentage increase sCr $\geq 50\%$ which is known, or presumed, to have occurred within the prior seven days		
Staging of AKI	- Stage 1: increase in sCr ≥ 0.3 mg/dl ($\geq 26.5 \mu\text{mol/L}$) or an increase in sCr ≥ 1.5 -fold to 2-fold from baseline; - Stage 2: increase in sCr >2 -fold to 3-fold from baseline; - Stage 3: increase of sCr >3 -fold from baseline or sCr ≥ 4.0 mg/dl ($353.6 \mu\text{mol/L}$) with an acute increase ≥ 0.3 mg/dl ($\geq 26.5 \mu\text{mol/L}$) or initiation of renal replacement therapy		
Progression of AKI	Progression	Regression	
	Progression of AKI to a higher stage and/or need for RRT	Regression of AKI to a lower stage	
Response to treatment	No response	Partial response	Full response
	No regression of AKI	Regression of AKI stage with a reduction of sCr to ≥ 0.3 mg/dl ($\geq 26.5 \mu\text{mol/L}$) above the baseline value	Return of sCr to a value within 0.3 mg/dl ($\geq 26.5 \mu\text{mol/L}$) of the baseline value

AKI, acute kidney injury; sCr, serum creatinine; RRT, renal replacement therapy.

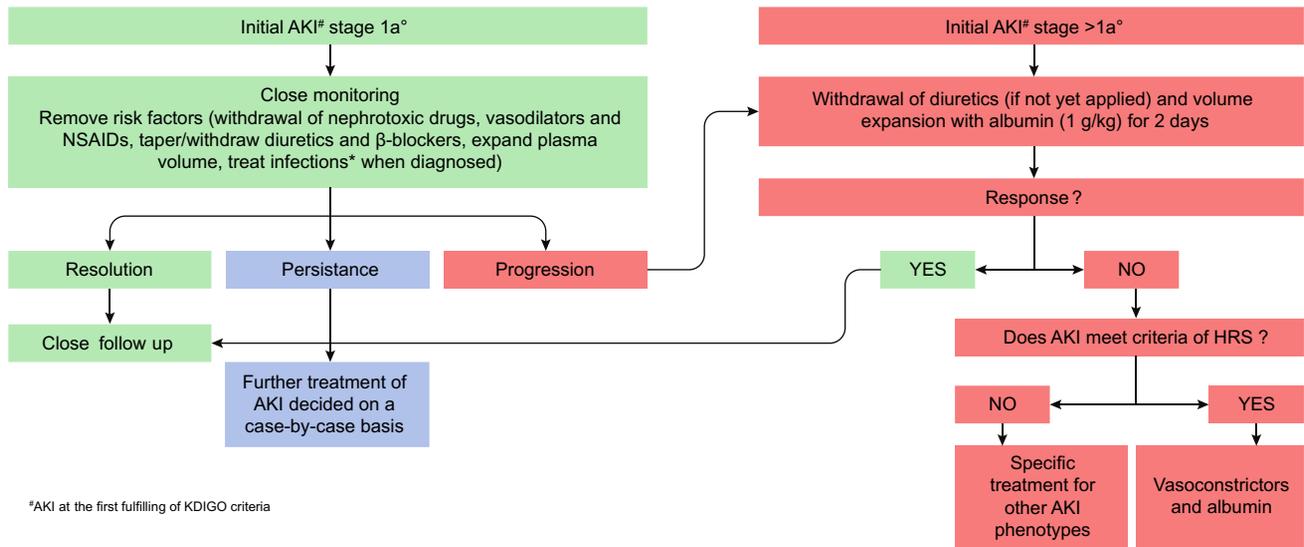


Fig. 8. Algorithm for the management of AKI in patients with cirrhosis (adapted from Ref. 318). AKI, acute kidney injury; HRS, hepatorenal syndrome; NSAID, non-steroidal anti-inflammatory.

KDIGO urine output criteria = an urinary output <0.5 ml/kg B.W./h x 6-12 h

Stage	Serum creatinine criteria
1°	An urinary output <0.5 ml/kg B.W./h x 6-12 h
2°	An urinary output <0.5 ml/kg B.W./h x 12 h
3°	An urinary output <0.5 ml/kg B.W./h x 24 h or anuria per 12 h

Fig. 9. Criteria based on urinary output for the diagnosis of AKI (adapted from Ref. 310). AKI, acute kidney injury; B.W., body weight.

the last week before admission. This point is so crucial for the application of the KDIGO criteria that it has been suggested that an SCr value be calculated when not available the seven days prior to admission. The baseline SCr can be calculated by inversely applying the formulas that are used to calculate the estimated GFR, considering normal values of GFR of 75 ml/min.³¹⁹ Whilst an imputed SCr is accepted in the general population, it can not be used in patients with cirrhosis.³²⁰ Indeed, all SCr-based formulas overestimate the true GFR in these patients leading to an overestimation of the baseline SCr and thus underestimating the prevalence of AKI on admission.³²⁰ Therefore, it has been proposed that not only the value obtained in the last seven days, but also that within the last three months be considered as a baseline value of SCr in patients with cirrhosis (Table 7). In addition, an SCr value obtained within the last three months is the reference to define acute kidney disease (AKD), a third category of renal impairment, along with AKI and CKD, which has been recently proposed in KDIGO recommendations. AKD is clearly a distinct category with different outcome, whether or not it is associated with AKI. AKD is defined by a GFR <60 ml/min/1.73 m² for less than three months, or a decrease in GFR ≥35% for less than three months, or an increase in SCr <50% within the last three months (Table 8). However, no data exist about the prognostic impact of AKD, with or without AKI, in patients with cirrhosis. Thus, waiting for these data, it seems even more justified to make the diagnosis of AKI in patients with cirrhosis on an increase in SCr ≥50% during the last three months. This assumption may also facilitate the diagnosis of AKI overlapping CKD.

Table 8. Definitions of kidney disease.

Definition	Functional criteria	Structural criteria
AKI	Increase in sCr ≥50% within seven days, or increase in sCr ≥0.3 mg/dl within two days	No criteria
AKD	GFR <60 ml/min per 1.73 m ² for <3 months, or decrease in GFR ≥35% for <3 months, or increase in sCr ≥50% for <3 months	Kidney damage for <3 months
CKD	GFR <60 ml/min per 1.73 m ² for ≥3 months	Kidney damage for ≥3 months

AKD, acute kidney disease; AKI, acute kidney injury; CKD, chronic kidney disease; GFR, glomerular filtration rate; sCr, serum creatinine.

Recommendations

- In patients with liver diseases, even a mild increase in SCr should be considered since it may underlie a marked decrease of GFR (II-2;1).
- The first step to be addressed in the diagnostic process is to establish if the patient has a CKD, AKD or AKI as well as an overlap between these diagnostic categories (II-2;1).
- The diagnosis of CKD should be based on a GFR <60 ml/min/1.73 m² estimated by SCr-based formulas, with or without, signs of renal parenchymal damage (proteinuria/haematuria/ultrasonography abnormalities) for at least three months (II-2;1).
- The diagnostic process should be completed by staging CKD, which relies on GFR levels, and by investigating its cause. It should be highlighted that any SCr based formula overestimates GFR in patients with cirrhosis (II-2, 1).

- In patients with cirrhosis the diagnosis of AKI should be based on adapted KDIGO criteria, thus, either on an increase in SCr of >0.3 mg/dl from baseline within 48 h, or an increase of ≥50% from baseline within three months (II-2,1).
- The staging of AKI should be based on an adapted KDIGO staging system, thus distinguishing within AKI stage 1, between AKI stage 1A and AKI stage 1B according to a value of SCr <1.5 or ≥1.5 mg/dl, respectively (II-2,1).

Precipitating factors

Infections, diuretic-induced excessive diuresis, GI bleeding, therapeutic paracentesis without adequate volume expansion, nephrotoxic drugs, and NSAIDs are the other common precipitating factors of AKI in patients with cirrhosis.^{20,242,306} The nephrotoxicity of contrast agents is still debated in patients with cirrhosis³²¹ but contrast imaging should be performed cautiously, particularly in decompensated cirrhosis or in patients with known CKD. Finally, the increase in intra-abdominal pressure associated with tense ascites may lead to AKI, by increasing renal venous pressure.^{322–324}

Management

The cause of AKI should be investigated as soon as possible, to prevent AKI progression. However, even in the absence of a definitive recognised cause of AKI, the management should be immediately started according to the initial stage (Fig. 2). Irrespective of the stage, diuretics should be discontinued. Similarly, even if there are controversial data, beta-blockers should be stopped.¹⁶⁸ Other precipitating factors of AKI should be identified and treated, including screening and treatment of infection, volume expansion when appropriate, and discontinuation of all nephrotoxic drugs, such as vasodilators or NSAIDs.³¹⁸ Volume replacement should be used in accordance with the cause and the severity of fluid loss. Patients with diarrhoea or excessive diuresis should be treated with crystalloids, whilst patients with acute GI bleeding should be given packed red blood cells to maintain haemoglobin level between 7–9 g/dl.³²⁵ In patients with AKI and tense ascites, therapeutic paracentesis should be associated with albumin infusion since it improves renal function.³²⁶ In case of no obvious cause and AKI stage >1A, 20% albumin solution at the dose of 1 g of albumin/kg of body weight (with a maximum of 100 g of albumin) for two consecutive days should be given.³⁰⁷ All other therapeutic options, especially renal replacement therapy (RRT) and kidney transplantation will be discussed in the section dedicated to the management of HRS-AKI.

Recommendations

- When a diagnosis of AKI is made, its cause should be investigated as soon as possible to prevent AKI progression. Even in absence of an obvious cause, the management should be immediately started. Maximal attention in the screening and treatment of infections should be carried out (II-2,1).
- Diuretics and/or beta-blockers as well as other drugs that could be associated with the occurrence of AKI such as vasodilators, NSAIDs and nephrotoxic drugs should be immediately stopped (II-2,1).

- Volume replacement should be used in accordance with the cause and severity of fluid losses (II-2,1).
- In case of no obvious cause of AKI, AKI stage >1A or infection-induced AKI, 20% albumin solution should be used at the dose of 1 g of albumin/kg of body weight (with a maximum of 100 g of albumin) for two consecutive days (III,1).
- In patients with AKI and tense ascites, therapeutic paracentesis should be associated with albumin infusion even when a low volume of ascetic fluid is removed (III,1).

Types of AKI

All types of AKI can occur in patients with cirrhosis, namely pre-renal AKI, HRS-AKI, intrarenal or intrinsic AKI, and post-renal AKI. The most common cause of AKI in hospitalised patients with decompensated cirrhosis is pre-renal, accounting for approximately 68% of the cases.^{306,327,328} Intrarenal-AKI is mainly represented by acute tubular necrosis (ATN).³⁰⁶ Finally, post-renal AKI is uncommon in decompensated cirrhosis.³²⁸ Considering that most cases of pre-renal AKI are resolved by volume expansion and that post-renal AKI is uncommon, the key point is to differentiate HRS-AKI from ATN. As described in the section “Hepatorenal syndrome”, the concept that HRS is only a functional injury has been challenged during the last decade and, thus, the definition of HRS probably has to be revised. In addition, as kidney biopsy is rarely performed in the setting of AKI in clinical practice, the distinction between HRS-AKI and ATN is difficult. Recently, novel biomarkers have emerged in this setting and urinary neutrophil gelatinase-associated lipocalin (NGAL) is the most promising. Indeed, several studies have shown that urinary NGAL, a marker of tubular damage, could help to determine the type of AKI.^{329–335} However, cut-off values differ greatly according to series, there are overlaps between the different types of AKI and it should be highlighted that no study has confirmed the diagnosis by reference kidney biopsy. Diagnosis based on a combination of multiple biomarkers may be interesting but needs further evaluation.^{329,330,332–334}

Recommendations

- All types of AKI can occur in patients with cirrhosis, namely pre-renal, HRS, intrinsic, particularly ATN, and post-renal. Therefore, it is important to differentiate among them (II-2,1).
- The diagnosis of HRS-AKI is based on revised ICA criteria. As kidney biopsy is rarely performed in the setting of AKI, biomarkers should be implemented. In clinical practice among the different biomarkers to date, urinary NGAL can be used to distinguish between ATN and HRS (II-2;2).

Prognosis

In patients with decompensated cirrhosis, AKI has a negative impact on hospital survival according to either the initial stage,³¹⁴ or the peak stage.^{313,317} Even transient episodes of AKI are associated with a negative impact on mid-term survival.³¹⁵ Nevertheless, a more comprehensive prognostic classification

also considering extra-renal organ failures is much more accurate than the KDIGO criteria for the prognosis in these patients. Finally, looking to the data in the general population, it should be highlighted that the risk for developing CKD is higher in patients with severe or repeated episodes of AKI.³³⁶ Since patients with decompensated cirrhosis are prone to develop frequent episodes of AKI, it can be speculated that they are at higher risk of developing CKD.

Hepatorenal syndrome

Definition, diagnosis and classification

For a long time, HRS has been defined as “a functional renal failure caused by intrarenal vasoconstriction which occurs in patients with end-stage liver disease as well as in patients with acute liver failure or alcoholic hepatitis”.^{32,307} Several data challenge this definition of HRS as well as the classification in type 1 and type 2. Firstly, as described below, pathogenesis of HRS includes both haemodynamic and inflammatory changes. Secondly, the absence of renal parenchymal damage, defining the functional nature, has never been proven by renal biopsies.^{337,338} The absence of significant proteinuria and/or haematuria do not rule out renal lesions, particularly tubular and interstitial lesions.³⁰⁷ In addition, studies assessing novel kidney biomarkers have shown that tubular damage can occur in patients with HRS-AKI when HRS is diagnosed according to the traditional criteria.^{328–330,332} Finally, it should be noted that HRS-AKI can occur in patients with underlying CKD. Type 1 and type 2 were historically defined based on time frame SCr increase.^{32,307} In the recent revised classification, type 1 HRS now corresponds to HRS-AKI.³¹⁸ Consequently, type 2 HRS should now include renal impairment which fulfills the criteria of HRS but not of AKI, namely non-AKI-HRS (NAKI), and only HRS-CKD as previously proposed.³³⁹

Pathophysiology

According to the new theory that has been developed on the pathophysiology of decompensated cirrhosis,⁵ the view on HRS has been changed in recent years, moving from the idea that it was only related to renal hypoperfusion due to macrocirculatory dysfunction (*i.e.* splanchnic arterial vasodilation and reduction of cardiac output).^{192,338} The new theory is that the increased circulating levels of pro-inflammatory cytokines and chemokines^{340,341} may exercise a direct relevant role in the development of HRS. Such cytokines have been associated with renal impairment in patients and in animal models of cirrhosis with infection.^{342–345} Moving from the concept that AKI and HRS-AKI are often precipitated by bacterial infection, the new hypothesis on the pathogenesis of sepsis-induced AKI should also be considered.^{346–348} This theory proposes that a synergic interplay of inflammation and microvascular dysfunction is responsible for the amplification of the signal that PAMPs and DAMPs exert on proximal epithelial tubular cells. The recognition of this signal and its subsequent spread to all the other proximal tubular epithelial cells cause a mitochondria-mediated metabolic downregulation and reprioritisation of cell functions to favour survival processes above all else.³⁴⁹ The sacrificed functions include the absorption on the lumen side of sodium and chloride. The consequent increases of sodium chloride delivery to the macula densa triggers further intrarenal activation of the RAAS and thus lowers GFR. Finally, severe cholestasis may further impair renal function by worsening inflammation and/or macrocirculatory dysfunction, or by

promoting bile salt-related direct tubular damage.^{350,351} All these findings suggest that the pathophysiology of AKI, and particular of HRS-AKI, in patients with decompensated cirrhosis seems more complex than previously hypothesised, supporting the concept that AKI-HRS is not purely functional in nature.

Management

The non-specific management of AKI as been previously described. Thus, in this section, drug therapy, TIPS, RRT, LT and simultaneous liver and kidney transplantation (SLK) will be considered.

Drug therapy. Once the diagnosis of HRS-AKI has been made, patients should promptly receive vasoconstrictive drugs, in association with albumin. The rationale for using vasoconstrictors is to counteract the splanchnic arterial vasodilation, improving renal perfusion.³⁵² Terlipressin, a vasopressin analogue, is the most commonly used. The efficacy of terlipressin plus albumin in the treatment of HRS has been proven in many studies.^{353–360} In the most recent studies, rates of response (complete or partial response) to this treatment range from 64 to 76%, with a complete response, from 46 to 56%.^{358–360} These response rates must now be evaluated according to the new definitions of responses in HRS-AKI recently proposed by the ICA (Table 7). In two meta-analyses terlipressin plus albumin was proven to improve not only renal function but also short-term survival in patients with HRS.^{361,362} Terlipressin was initially proposed to be administered by *i.v.* boluses at a starting dose of 0.5–1 mg every 4–6 h, progressively increased to a maximum of 2 mg every 4–6 h in case of a reduction of baseline SCr < 25%.^{353–358} Adding albumin to terlipressin is more effective than terlipressin alone.³⁵⁴ One possible explanation is that albumin, by increasing volemia, may counteract the decrease in cardiac output associated with HRS¹⁹² but also by terlipressin.³⁶³ In addition, antioxidant and anti-inflammatory properties of albumin may have a beneficial effect.³⁶⁴ The dose of albumin in HRS treatment has not been well established. Studies have suggested adapting the dose according to the level of central venous pressure (CVP), but there is evidence that CVP is inaccurate to manage volume expansion and to assess cardiac output in patients with cirrhosis. In contrast, CVP may be helpful to prevent circulatory overload. Albumin has been used intravenously at the mean dose of 20–40 g/day. Treatment should be maintained until a complete response (SCr below 1.5 mg/dl) or for a maximum of 14 days either in case of partial response (decrease of SCr \geq 50 with a final value still higher than 1.5 mg/dl) or in case of non-response. More recently, continuous *i.v.* infusion of terlipressin at an initial dose of 2 mg/day was proposed,^{359,365} demonstrating a similar rate of response but lower adverse effects than the administration of the drug by *i.v.* boluses.³⁶⁰ Indeed, terlipressin, when administered by continuous *i.v.* infusion, has a more stable lowering effect on portal pressure, even when used at lower doses than those provided by *i.v.* boluses.³⁶⁰ The most common side effects of terlipressin are diarrhoea, abdominal pain, circulatory overload and cardiovascular ischaemic complications which have been reported in up to 45–46% of patients when the drug was delivered by *i.v.* boluses.³⁶⁰ The rate of discontinuation because of side effects, mainly cardiovascular, is around 20%.³⁶⁰ Accordingly, a careful clinical screening including electrocardiogram is recommended in all patients before starting treatment. Patients can be treated on a regular ward but the decision to transfer to a higher level of

Clinical Practice Guidelines

care should be case based. Recurrent HRS in responders, after the end of the treatment, has been reported in up to 20% of cases. Re-treatment is usually effective, however, in some cases, continuous recurrence occurs, thus a long-term treatment with terlipressin plus albumin and a long-term hospitalisation are required.³⁶⁶ The possibility of treating some of these patients outside the hospital has recently been proposed³⁶⁷ but even if promising, further studies are needed. Other vasoconstrictive drugs include i.v. noradrenaline and oral midodrine plus subcutaneous or i.v. octetide, both in combination with albumin. Noradrenaline, given by continuous i.v. infusion at the dose of 0.5–3 mg/h, has been proven to be as effective as terlipressin regarding the increase in mean arterial pressure, the reversal of renal impairment and one-month survival.^{368–371} However, the number of patients treated with noradrenaline remains too small to definitively confirm its efficacy. In addition, in contrast to terlipressin, the use of noradrenaline always requires a central venous line and, in most countries, the transfer of the patient to an intensive care unit (ICU). The combination midodrine plus octetide, used in countries where terlipressin is not yet available,³⁷² has been shown to be much less effective than terlipressin in the treatment of type 1 HRS in a recent RCT.³⁵⁹ Vasoconstrictors, in particular terlipressin, in association with albumin, have also been proposed in the treatment of type 2 HRS. The treatment has been proven to be effective in most cases but, unfortunately, recurrence after the withdrawal of treatment is the norm. In addition, there are controversial data about the impact of this treatment on outcomes, especially in candidates for LT.^{373,374} This may be, at least in part, due to the suboptimal definition of type 2 HRS, as previously discussed. The most relevant factors that may impair the response to vasoconstrictors are: a) the baseline value of SCr, b) the degree of inflammation and c) the degree of cholestasis.^{375–377} The finding that the higher baseline values of SCr, the lower rate of response to terlipressin plus albumin,³⁷⁵ probably reflects the presence of renal parenchymal damage³³⁷ and represents one of the main reasons behind the adoption of the KDIGO criteria for the definition of AKI in patients with cirrhosis, and the introduction of the new algorithm for its management. Regarding inflammation, it has been recently shown that, for the same value of baseline SCr, the rate of response is related to the number of extra-renal organ failures.³⁷⁶

Recommendations

- Vasoconstrictors and albumin are recommended in all patients meeting the current definition of AKI-HRS stage >1A, should be expeditiously treated with vasoconstrictors and albumin (III;1).
- Terlipressin plus albumin should be considered as the first-line therapeutic option for the treatment of HRS-AKI. Terlipressin can be used by i.v. boluses at the initial dose of 1 mg every 4–6 h. However, giving terlipressin by continuous i.v. infusion at initial dose of 2 mg/day makes it possible to reduce the global daily dose of the drug and, thus, the rate of its adverse effects. In case of non-response (decrease in SCr <25% from the peak value), after two days, the dose of terlipressin should be increased in a stepwise manner to a maximum of 12 mg/day (I;1).

- Albumin solution (20%) should be used at the dose 20–40 g/day. Ideally, apart from routinely monitoring patients with HRS-AKI, the serial measurement of CVP or other measures of assessing central blood volume, can help to prevent circulatory overload by optimising the fluid balance and helping to titrate the dose of albumin (II-2;1).
- Noradrenaline can be an alternative to terlipressin. However, limited information is available (I;2).
- In contrast to terlipressin, the use of noradrenaline always requires a central venous line and, in several countries, the transfer of the patient to an ICU. Midodrine plus octetide can be an option only when terlipressin or noradrenaline are unavailable, but its efficacy is much lower than that of terlipressin (I;1).
- According to the new definition of HRS-AKI, complete response to the treatment should be defined by a final SCr within 0.3 mg/dl (26.5 µmol/L) from the baseline value, while partial response should be defined by the regression of AKI stage to a final SCr ≥0.3 mg/dl (26.5 µmol/L) from the baseline value (III;1).
- Adverse events related to terlipressin or noradrenaline include ischaemic and cardiovascular events. Thus, a careful clinical screening including electrocardiogram is recommended before starting the treatment. Patients can be treated on a regular ward, but the decision to transfer to higher dependency care should be case based. For the duration of treatment, it is important to closely monitor the patients. According to the type and severity of side effects, treatment should be modified or discontinued (I;1).
- In cases of recurrence of HRS-AKI upon treatment cessation, a repeat course of therapy should be given (I;1).
- Terlipressin plus albumin is also effective in the treatment of HRS outside the criteria of AKI (HRS-NAKI), formerly known as HRS type II. Unfortunately, recurrence after the withdrawal of the treatment is the norm, and controversial data exists on the impact of the treatment on long-term clinical outcome, particularly from the perspective of LT. As such, vasoconstrictors and albumin are not recommended in this clinical scenario (I;1).

Transjugular intrahepatic portosystemic shunts. The use of TIPS may improve renal function in patients with type 1 HRS.^{378,379} However, the applicability of TIPS in this clinical setting is usually very limited because, in most patients, TIPS is contraindicated because of severe degree of liver failure. TIPS has been studied in patients with type 2 HRS³⁸⁰ and in the management of refractory ascites, frequently associated with type 2 HRS. In these patients, TIPS has been shown to improve renal function.^{95,379}

Renal replacement therapy. Renal replacement therapy should be considered in the management of AKI, whatever the type. As far as HRS-AKI, it should be considered in non-responders to vasoconstrictors. RRT should also be considered in patients with end-stage kidney disease. The indications for RRT are the same in patients with cirrhosis as in the general population including: severe and/or refractory electrolyte or acid-base imbalance,

severe or refractory volume overload, and/or symptomatic azotaemia. However, published data on RRT in patients with cirrhosis are scant, with controversial effects on survival.^{381,382} It has been stated that indications for RRT depend on the perspective of LT. It has been stated that RRT may be considered in patients who are candidates for LT, while, in contrast, the decision to initiate RRT in non-candidates should be individualised to avoid futility.²⁰ However, it has recently been observed that critically ill liver cirrhotic patients requiring RRT have very high mortality independent of LT options. Thus, RRT and treatment at the ICU should not be limited to LT candidates but should be based on the individual severity of illness.³⁸³ Therefore, repeated risk stratification is necessary during the course of treatment, assisted by prognostic scores in addition to clinical judgment and patients' wishes.³⁸³ The ideal timing for RRT initiation has not been defined in patients with cirrhosis. However, data on AKI in patients with acute liver failure as well as in critically ill patients without liver disease suggest that early RRT improves survival.^{384–386} Both haemodialysis or continuous renal replacement therapy (CRRT), have been used in patients with cirrhosis. Despite the available evidence,³⁸⁷ CRRT is probably better tolerated, providing greater cardiovascular stability and allowing a slower correction of severe or refractory hyponatraemia than haemodialysis.

Liver support systems. In two controlled studies, both the so-called artificial liver support systems, either the molecular adsorbents recirculating system (MARS[®]) or Prometheus[®], showed promising beneficial effects in patients with type 1 HRS, but should be further investigated.^{388,389}

Liver transplantation and simultaneous liver-kidney transplantation. The best therapeutic option in patients with HRS is LT.³⁹⁰ However, several studies have shown that SCr after LT is higher in patients transplanted with HRS, compared to those without HRS at the time of LT. In addition, the presence of HRS at the time of LT has a negative impact on survival after LT.³⁹¹ The treatment of type 2 HRS before LT has given conflicting results on the clinical outcome after LT^{373,374} and thus, requires further investigation. SLK can be indicated in patients with cirrhosis and CKD in the following conditions: a) estimated GFR (using MDRD6 equation) ≤ 40 ml/min or measured GFR using iothalamate clearance ≤ 30 ml/min, b) proteinuria ≥ 2 g a day, c) kidney biopsy showing $>30\%$ global glomerulosclerosis or $>30\%$ interstitial fibrosis, or d) inherited metabolic disease. SLK is also indicated in patients with cirrhosis and sustained AKI irrespective of its type, including HRS-AKI when refractory to drug therapy, in the following conditions: a) AKI on RRT for ≥ 4 weeks or b) estimated GFR ≤ 35 ml/min or measured GFR ≤ 25 ml/min ≥ 4 weeks.³⁹² Beyond these two conditions, in a candidate with high priority for LT due to a high MELD score, the option of SLK may be considered in the presence of risk factors for underlying undiagnosed CKD (diabetes, hypertension, abnormal renal imaging and proteinuria >2 g/day).³⁹² The development of new biomarkers of kidney fibrosis, a common and irreversible feature of CKD, is also promising in this context.³⁹³

Regarding the priority allocation of patients with HRS-AKI to the waiting list, some rules should be applied in case of response to drug therapy. In fact, by lowering SCr and increasing serum sodium concentration, the treatment can significantly lower the MELD and MELD-Na score, potentially delaying LT. Considering that the survival rate in responders at three months

is almost 50%, a specific policy of priority allocation is needed for these patients. This can be made either by continuing to consider the baseline MELD and/or MELD-Na score³⁹⁴ rather than those during or after the end of the treatment, or by providing an exception to the MELD score.³⁹⁵

Recommendations

- There is insufficient data to advocate TIPS in HRS-AKI but it could be suggested in selected patients with HRS-NAKI (II-2;2).
- LT is the best therapeutic option for patients with HRS regardless of the response to drug therapy (I;1).
- The decision to initiate RRT should be based on the individual severity of illness (I;2).
- The indication for liver-kidney transplantation remains controversial. This procedure should be considered in patients with significant CKD or with sustained AKI including HRS-AKI with no response to drug therapy (II-2;1).

Prevention of hepatorenal syndrome

The prevention of HRS-AKI, as for other causes of AKI, is based on the use of albumin in patients who develop SBP²⁷² and the prevention of SBP using norfloxacin,²⁷⁶ as discussed before. In addition, the use of pentoxifylline may decrease the incidence of renal failure in patients with cirrhosis²⁷ and of type 1 HRS as well as mortality in patients with severe alcoholic hepatitis.³⁹⁶ However, recent papers do not confirm these results^{397,398} and further studies are needed.

Recommendations

- Albumin (1.5 g/kg at diagnosis and 1 g/kg on day three) should be given in patients with SBP to prevent AKI (I;1).
- Norfloxacin (400mg/day) should be given as prophylaxis of SBP to prevent HRS-AKI (I;1).

Acute-on-chronic liver failure

Definitions and pathophysiology

Since the CANONIC study, the first major international observational study characterising the syndrome of ACLF,³ a large number of publications have described the association of this syndrome with different clinical, diagnostic and therapeutic approaches. ACLF occurs in 30% of admitted patients^{3,399} and in 25% of outpatients,⁴⁰⁰ and is a major cause of death in cirrhosis, with an approximately 50% mortality rate.⁴⁰⁰ Even though there is an ongoing debate regarding the definition of ACLF,^{401–405} the concept of the development of ACLF is similar across different continents and health systems. There is agreement that ACLF is not just decompensation of liver cirrhosis, but a distinct syndrome.⁴⁰⁶ The reason is that ACLF is defined by a multi-organ failure and has a higher short-term mortality than a "simple decompensation" of cirrhosis.^{3,401,406} The risk of developing ACLF is higher in outpatients with advanced liver disease according to the presence of ascites, low mean arterial

Box 1. CLIF-C Acute Decompensation Score (Ref. 407).**CLIF-C Acute Decompensation score**

$$10 \times [0.03 \times \text{Age} + 0.66 \times \text{Ln}(\text{Creatinine}) + 1.71 \times \text{Ln}(\text{INR}) + 0.88 \times \text{Ln}(\text{WBC}) - 0.05 \times \text{Sodium} + 8]$$

Age in years; creatinine in mg/dl; WBC (white blood count) in 10^9 cells/L; sodium in mmol/L

pressure or anaemia and with a high MELD score.⁴⁰⁰ ACLF develops on the background of acute decompensation (AD) of cirrhosis, but a remarkable number of patients (~40%) admitted to hospital developed ACLF on the first episode of AD of their liver disease.³ Thus, the presence of AD is an important clinical feature for the diagnosis of ACLF.^{3,401,406} The EASL-CLIF Consortium has proposed and validated a prognostic score (CLIF-C AD score) for patients with AD who do not develop ACLF.⁴⁰⁷ The CLIF-C AD score (Box 1) was proved to be more accurate for predicting outcome in these patients than the MELD or MELD-Na score.⁴⁰⁷ Once developed, ACLF is characterised by hepatic and extrahepatic organ dysfunction and/or failure, highly activated systemic inflammation, and a high 28-day mortality.^{3,12} The overwhelming and devastating inflammatory response is a key pathogenic mechanism in the development of ACLF, probably explaining why ACLF frequently happens in younger patients.^{3,401,406,408} The trigger of ACLF and this inflammatory response could not be identified in 40–50% of the patients in CANONIC study,³ which might be associated with genetic predisposition, severe portal hypertension or other factors predisposing the patients to development of AD and ACLF.⁴⁰⁹ However, identification of the precipitating events of AD are of great importance to prevent and manage ACLF.^{410,411}

Precipitating events

The precipitating events vary between different populations, geographic areas and aetiologies. While in Western countries (Europe, North and Latin America) bacterial infection, followed by active alcohol intake or binge are major precipitating events,^{3,412,413} in Eastern countries (Asia, Pacific region) the exacerbation of hepatitis B, followed by alcohol or bacterial infections are the major causes of AD and ACLF development.^{414–416} But there are a number of other insults, which might induce ACLF, such as superimposed infection with hepatotropic viruses (especially HAV, HEV), DILI, GI bleeding, circulatory dysfunction upon different situation (e.g. surgery, LVP without albumin). Therefore, in general the precipitating factors might be differentiated into three major categories, hepatotoxic injury (active alcohol intake or binge, DILI), immunological insults (flares of viral or autoimmune hepatitis, bacterial, fungal and viral infections, common cold, subclinical infections, etc.) and haemodynamic derangement following procedures (haemorrhage, surgery, LVP).

Bacterial infections

Overall, the major precipitating factor for ACLF is bacterial infections accounting for 30–57% of cases.^{409,410} The importance of bacterial infections for the development of organ failures and ACLF was also underlined by the studies of North American Consortium for End-stage Liver Disease (NACSELD), who have defined ACLF by the development of two organ failures in presence of bacterial infections.⁴¹² By contrast, bacterial infections were not considered to be precipitating events for ACLF accord-

ing to the definition of Asian Pacific Association for the Study of the Liver (APASL).⁴⁰² However, nowadays there is evidence that bacterial infections are mainly involved in the development of organ failures and thereby of ACLF in Asia as well.^{415,416} In Western countries bacterial infections are the precipitating events in one-third of patients admitted with ACLF and in two-thirds of patients developing ACLF during follow-up.^{3,409,412,413} Based on these data, preventive and early therapeutic interventions for the treatment of infections are of major importance to prevent the development of ACLF. The role of bacterial infections as triggers of AD and development of organ failures has already been discussed.

Active alcohol intake or binge

Alcoholic liver disease was the most prevalent in patients with AD and ACLF in the CANONIC study, as well as in recent reports from India.^{415–417} Interestingly, active alcoholism and alcohol binge were not only a major trigger in these patients, but led to a more severe syndrome than other triggers in alcoholic cirrhosis patients without heavy active alcoholism.³ The role and mechanisms of active alcoholism need further investigation, especially regarding prevention and treatment.

Reactivation and superimposed viral hepatitis

Reactivation of HBV in patients with cirrhosis is the main precipitating event in the non-Caucasian Asian population,^{413,415} occurring mostly in genotypes B and D, and hepatitis B e antigen positive patients. Interestingly, superimposed HAV and HEV can also trigger ACLF in 14–18%.^{414,416} According to the Western experience, these are unusual causes.^{3,413} However, the role of HEV might have been overlooked, and might gain more importance now because of advances in diagnostics and increases in awareness.^{3,418} A timely recognition and treatment of the precipitating event might prevent ACLF and improve outcome in these patients.

Clinical and diagnostic features of ACLF

As discussed previously, organ failures in the presence of AD of cirrhosis are the basis for the diagnosis of ACLF. However, in the CANONIC study the presence and grading of ACLF was based on mortality and the independent association of organ dysfunction/failure with mortality, which was chosen to be $\geq 15\%$ at 28 days.³ Organ failures were defined based on a sequential organ failure assessment (SOFA) score, which was adapted to patients with cirrhosis, the CLIF SOFA score (Table 9). However, two organs received special attention, the kidney and the brain.³ In fact, it has been observed that even mild renal or brain dysfunction in the presence of another organ failure, is associated with a significant short-term mortality and therefore defines the presence of ACLF. Thus, patients with renal failure, defined as creatinine ≥ 2 mg/dl, were classified as ACLF grade Ia while patients with a non-renal and non-cerebral organ failure combined either with mild renal dysfunction (creatinine between 1.5 and 1.9 mg/dl) and/or grade I and II hepatic encephalopathy, as well as those with cerebral failure combined with mild renal dysfunction were classified as ACLF grade Ib (Table 10).³ Thereafter, patients with two organ failures are classified as grade II ACLF, and have a 28-day mortality rate of 32%. Patients with three or more organ failures are classified as grade III ACLF and have an average 28-day mortality of 78% (Table 9). According to this EASL-CLIF definition of ACLF, approximately one-quarter of patients admitted to the hospital

Table 9. CLIF-Sequential Organ Failure Assessment (SOFA) score (adapted from Ref. n° 3).

Organ/system	The CLIF-Sequential Organ Failure Assessment (SOFA) score				
	0	1	2	3	4
Liver (bilirubin mg/dl)	<1.2	≥1.2–<2.0	≥2.0–<6.0	≥6.0–<12.0	≥12.0
Kidney (creatinine, mg/dl)	<1.2	≥1.2–<2.0	≥2.0– <3.5	≥3.5– <5.0	≥5.0
Cerebral (HE grade)	No HE	Grade I	Grade II	Grade III	Grade IV
Coagulation (INR and PLT count)	<1.1	≥1.1–<1.25	≥1.25–<1.5	≥1.5–<2.5	≥2.5 or PLT ≤ 20.000/mm ³
Circulation (MAP, mmHg and vasopressors)	≥70	<70	Dopamine ≤5[†] or dobutamine or terlipressin	Dopamine >5[†] or E ≤0.1 or NE ≤0.1[†]	Dopamine >15[†] or E >0.1 or NE >0.1[†]
Lungs					
PaO ₂ /FiO ₂ , or	>400	>300–≤400	>200–≤300	>100–≤200	≤100
SpO ₂ /FiO ₂	>512	>357–≤512	>214–≤357	>89–≤214	≤89

E, epinephrine; FiO₂, fraction of inspired oxygen; HE, hepatic encephalopathy; NE, norepinephrine; PaO₂, partial pressure of arterial oxygen; SpO₂, pulse oximetric saturation. The bold text indicates the diagnostic criteria for organ failures.

[†] μg/kg/min.

Table 10. Classification and grades of ACLF (adapted from Ref. 3).

Grades of ACLF	Clinical characteristics
No ACLF	No organ failure, or single non-kidney organ failure, creatinine <1.5 mg/dl, no HE
ACLF Ia	Single renal failure
ACLF Ib	Single non-kidney organ failure, creatinine 1.5–1.9 mg/dl and/or HE grade 1–2
ACLF II	Two organ failures
ACLF III	Three or more organ failures

ACLF, acute-on-chronic liver failure; HE, hepatic encephalopathy.

for AD of cirrhosis had ACLF at admission or develop it during the hospitalisation. After having simplified the CLIF SOFA score into the CLIF Organ Failure score (Table 11), the EASL-CLIF Consortium formulated a new score, the CLIF-C ACLF score, which enables the prediction of mortality in patients with ACLF.⁴¹⁹ The CLIF-C ACLF score (Box 2) has been validated by different independent series of patients.^{417,420,421} Other scores were recently proposed by the APASL ACLF Research Consortium and by the NACSELD, but they were not compared specifically with the CLIF-C-ACLF score.^{422,423}

Box 2. CLIF-C Acute Liver Failure (ACLF) score (Ref. 419).**CLIF-C ACLF score**

$$10 \times [0.033 \times \text{Clif OFs} + 0.04 \times \text{Age} + 0.63 \times \text{Ln(WBC)} - 2]$$

Age in years; CLIF OF score as in Table 10; sodium in mmol/L

Recommendations

- The diagnosis of ACLF should be made in a patient with cirrhosis and AD (defined as the acute development or worsening of ascites, overt encephalopathy, GI-haemorrhage, non-obstructive jaundice and/or bacterial infections), when organ failure(s) involving high short-term mortality develop (II-2;1).
- The diagnosis and the grading of ACLF should be based on the assessment of organ function as defined by the CLIF-C Organ Failure score (II-2;1).
- Potential precipitating factor(s), either hepatic (*i.e.* heavy alcohol intake, viral hepatitis, DILI, autoimmune hepatitis) and/or extrahepatic (*i.e.* infections haemodynamic derangements following haemorrhage, surgery) should be investigated. However, in a significant proportion of patients, a precipitant factor may not be identified (II-2;1).

Management of ACLF*General management*

Unfortunately, there is no specific effective treatment for ACLF.⁴²⁴ Therefore, treatment is currently based on organ support and management of associated complications. The cause of liver injury can be specifically treated only in certain situations such as in ACLF secondary to HBV infection, as described

Table 11. Chronic Liver Failure – Organ Failure score system (adapted from Ref. 419).

Organ/system	1 point	2 points	3 points
Liver	Bilirubin <6 mg/dl	6 ≤Bilirubin <12 mg/dl	Bilirubin ≥12 mg/dl
Kidney	Creatinine <2 mg/dl	2 Creatinine <3.5 mg/dl	Creatinine ≥3.5 mg/dl or renal replacement
Brain/HE (West Haven criteria)	Grade 0	Grades 1–2	Grades 3–4^a
Coagulation	INR <2.0	2.0 ≤INR <2.5	INR ≥2.5
Circulation	MAP ≥70 mmHg	MAP <70 mmHg	Use of vasopressors
Lungs	PaO ₂ /FiO ₂ >300, or SpO ₂ /FiO ₂ >357	PaO ₂ /FiO ₂ ≤300–>200, or SpO ₂ /FiO ₂ >214–≤357	PaO₂/FiO₂ ≤200^b or SpO₂/FiO₂ ≤214^b

Note: The bold text denotes criteria for diagnosing organ failures.

FiO₂, fraction of inspired oxygen; HE, hepatic encephalopathy; INR, international normalized ratio; MAP, mean arterial pressure; PaO₂, partial pressure of arterial oxygen; SpO₂, pulse oximetric saturation.

^a Patients submitted to mechanical ventilation due to HE and not to a respiratory failure were considered as presenting a cerebral failure (cerebral score = 3).

^b Other patients enrolled in the study with mechanical ventilation were considered as presenting a respiratory failure (respiratory score = 3).

Clinical Practice Guidelines

later. Patients with ACLF should ideally be admitted to intensive care or intermediate care units, yet this decision should be individualised based on certain factors, particularly patients' age and associated comorbidities. Moreover, patients suitable for LT should be referred to a transplant centre early in the course of ACLF. Late referral may make transplantation impossible due to the rapid evolution of ACLF in most patients.⁴²⁵ In patients in whom ACLF is associated with precipitating factors, such as bacterial infections, GI bleeding, or drug toxicity, early identification and management of these conditions is crucial to patient survival. Nonetheless, it should be emphasised that this early treatment of triggering factors may not prevent the progression of ACLF in all patients. Meanwhile, as already stated, in approximately half of patients with ACLF a precipitating factor cannot be identified.³ Organ support is very important in the management of patients with ACLF.⁴²⁴ Haemodynamic function should be monitored and vasopressor therapy administered in case of marked arterial hypotension. Hepatic encephalopathy should be treated early with standard therapy. Special care should be taken to preserve airway patency to prevent aspiration pneumonia. In patients with coagulation failure, either because of impairment of coagulation factors or low platelet count, substitutive therapy should be given only if there is clinically significant bleeding. If there is respiratory failure, patients should be given oxygen therapy and ventilation, if required. Finally, if there is kidney failure its cause should be identified and managed accordingly. Volume expansion should be given to patients with fluid loss or in the setting of SBP. Excessive volume expansion should be avoided. Patients meeting the criteria of AKI-HRS should be treated with terlipressin and albumin or norepinephrine, if terlipressin is not available. Patients with suspected ATN should be treated with RRT if they meet criteria for this treatment.³⁹²

Specific therapies

Liver support systems. Extracorporeal liver support systems, particularly albumin dialysis (MARS system) and fractionated plasma separation and adsorption (Prometheus system) have been evaluated as therapies for ACLF. These systems remove albumin-bound substances and other substances that accumulate in the context of ACLF and may have deleterious effects on the function of different organs. Both methods have been evaluated in large RCTs in patients with ACLF and no significant effects on survival could be demonstrated.^{388,389} It should be emphasised however, that the definition of ACLF in both trials was different than the current definition of ACLF based on the CANONIC study.³ Moreover, a sub-analysis of the Prometheus study showed a beneficial effect on survival in patients with MELD score higher than 30.³⁸⁹ This finding deserves further investigation. Nonetheless, based on the results of available RCTs, extracorporeal liver support systems do not improve survival of patients with ACLF and should not be recommended in this indication.

Antiviral therapy for chronic hepatitis B. Reactivation of hepatitis B is a very common cause of ACLF in certain areas of the world, particularly in southwest Asia.⁴¹⁴ A number of non-randomised studies and an RCT have shown that treatment with lamivudine, tenofovir or entecavir is associated with inhibition of HBV replication, improvement of liver function, and higher survival in patients with ACLF secondary to hepatitis B infection.^{426–429} The only RCT included 24 patients, 14 treated with tenofovir

and 13 treated with placebo, and showed significant differences in three-month survival (57% vs. 15%, respectively).⁴²⁶ Therefore, it seems evident that the presence of HBV infection should be investigated in all patients with ACLF and antiviral therapy should be started as soon as possible.

Other therapies. A number of therapies have been assessed in patients with ACLF, including dexamethasone, plasma exchange, chinese herbs, caspase inhibitors, mesenchymal stem cells transplantation, and administration of granulocyte-colony stimulating factor (G-CSF).^{430–432} In most cases, the information is still very preliminary and no recommendations can be made regarding their potential use in clinical practice. However, a note on G-CSF seems pertinent because this approach has been assessed in an RCT.⁴³² The rationale behind this treatment seems to be the mobilisation of stem cells from the bone marrow and their engraftment within the liver, although other beneficial effects may also occur. The only RCT evaluating this therapy included 47 patients with ACLF, as defined by the APASL criteria, 23 treated with G-CSF (12 doses of 5 µg/kg subcutaneously) and 24 treated with placebo in a double-blind manner. The main findings were an improvement in 60-day survival in the G-CSF group vs. the placebo group (66% vs. 26%, respectively; $p = 0.001$) along with a reduction in Child-Pugh, SOFA, and MELD scores and a decrease in the occurrence of HRS, hepatic encephalopathy and sepsis in G-CSF treated patients. Although these results are promising, additional studies in a larger number of patients are needed.

Liver transplantation. Liver transplantation is theoretically the definitive treatment for ACLF because it allows the cure of ACLF syndrome as well as the underlying liver disease.⁴²⁵ However, some important issues regarding LT for ACLF deserve a comment, particularly the accessibility of patients to LT, evaluation of candidate subjects, the outcomes of LT on survival, and futility. The accessibility of patients with ACLF to LT is probably decreased compared to that of patients with other indications for LT, because patients with ACLF have a high mortality rate after diagnosis of the condition. Early referral to transplant centres is therefore crucial. Then, because ACLF is a rapidly evolving syndrome, candidate patients need to be submitted to a "fast-track" clinical evaluation of organ function and potential comorbidities that could contraindicate LT. Data on outcome of patients with ACLF treated with liver LT are scarce but nonetheless, patient survival at three-months after LT is about 80%, much higher than what would be anticipated if patients were not transplanted.^{425,433,434} Almost all patients with ACLF-3 developed complications after LT, especially pulmonary, renal and infectious, compared to patients with no ACLF, or ACLF-1 and -2. This emphasises the need for special management when transplanting patients with ACLF-3, with repeated systematic screening for infection and careful monitoring of renal and respiratory parameters.⁴³⁴ Another point is that some patients with ACLF are potentially too sick for LT. In the context of scarcity of donor livers, the potential benefit of LT for patients with ACLF must be balanced with the rationing. Thus, more data is needed to determine medical futility in patients with ACLF.^{425,434} However, if LT is contraindicated or not available for patients with organ failures ≥ 4 or CLIF-C ACLFs > 64 at days 3–7 after diagnosis of ACLF-3, the intensive organ support should be discontinued owing to futility.⁴²⁵

Recommendations

- At present, there is no specific therapy for ACLF aside from antiviral therapy in patients with ACLF due to reactivation of HBV infection. Treatment of ACLF should be based on organ support and management of precipitants (see point below) and associated complications. Patients should be treated in intermediate care or intensive care settings. Organ function, particularly, liver, kidney, brain, lung, coagulation, and circulation should be monitored frequently and carefully throughout hospitalisation, as ACLF is a dynamic condition. However, monitoring and management should be individualised according to specific circumstances, mainly patients' age and comorbidities (**III, 1**).
- Early identification and treatment of precipitating factors of ACLF, particularly bacterial infections, are recommended. However, in some patients ACLF progresses despite treatment of precipitating factors (**III;1**).
- Administration of nucleoside analogues (tenofovir, entecavir) should be instituted as early as possible in patients with ACLF due to HBV infection (**I;1**).
- Early referral of patients with ACLF to liver transplant centres for immediate evaluation is recommended (**II-3;1**).
- Withdrawal of ongoing intensive care support can be suggested in patients, who are not candidates for LT, with four or more organ failures after one week of adequate intensive treatment (**II-2, 2**).
- Despite promising results, the administration of G-CSF can not be recommended at present (**1;2**).

Relative adrenal insufficiency

Definition and pathophysiology

Relative adrenal insufficiency (RAI) is a condition of inadequate cortisol response to stress in the setting of critical illness,⁴³⁵ also named as "Critical Illness Related Corticosteroid Insufficiency" (CIRCI).⁴³⁶ RAI has also been described in patients with cirrhosis and, although it is mainly present in critically ill patients with sepsis or septic shock (68.9%), it also affects non-critically ill cirrhotic patients (41.8%), including those with compensated cirrhosis.⁴³⁷⁻⁴⁴² The pathophysiology of RAI in cirrhosis is not well defined. Suppression of the hypothalamic-pituitary-adrenal axis activity, reduced effective volemia, which may impair adrenal perfusion, and both impaired cholesterol synthesis and enhanced pro-inflammatory cytokine production likely contribute to impair adrenal steroidogenesis.^{443,444} Adrenal dysfunction blunts the vascular effect of angiotensin II, norepinephrine and vasopressin, leading to further sympathetic nervous system activity.⁴⁴⁵ These effects would worsen the cardio-circulatory dysfunction of advanced cirrhosis, and favour gut bacterial overgrowth, and hence BT, by impairing intestinal motility.⁴⁴⁵ This explains why RAI in decompensated cirrhosis is associated with a higher probability of severe sepsis and type-1 HRS, and higher short-term mortality.^{437,446}

Diagnosis

The diagnosis of RAI is influenced by the method employed, as the measurement of serum total cortisol, either at baseline or

after stimulation by the standard dose- or low dose-short Synacthen tests can be utilised.⁴⁴⁷ The consensus statements from the American College of Critical Care Medicine recommend referring to a delta total serum cortisol <250 nmol/L (9 µg/dl) after adrenocorticotrophic hormone administration or a random total cortisol <276 nmol/L (10 µg/dl) in critically ill patients.⁴³⁶ There is no reason for not employing these indications in patients with cirrhosis. However, the diagnosis of RAI based on serum total cortisol concentration, which is measured by standard assays, may be flawed by the reduced serum levels of cortisol binding globulin (CBG) and albumin frequently seen in patients with cirrhosis. This may lead to an overestimation of RAI, as more than 90% of circulating cortisol is bound to these proteins.⁴⁴⁸ The assessment of serum-free cortisol concentration would overcome this limitation. Serum-free cortisol levels <50 nmol/L at baseline, or <86 nmol/L (9 µg/dl) after adrenocorticotrophic hormone suggest the presence of RAI in critically ill patients.⁴⁴⁹ By comparing RAI diagnosis in clinically stable patients with cirrhosis based on either total or free plasma cortisol, a clear discrepancy emerged, as the prevalence of RAI was 58% using total cortisol criteria and 12% using free cortisol with a peak plasma level <33 nmol/L after stimulation.⁴⁵⁰ Unfortunately, the methods for determining free cortisol are complex and expensive, so they are not used in routine clinical practice. The surrogate methods that have been proposed for the calculation of plasma free cortisol^{451,452} do not seem to be fully reliable in patients with cirrhosis.⁴⁵⁰ For these reasons, salivary cortisol has received attention, as it correlates with free cortisol levels irrespective of the concentration of binding proteins.^{453,454} Baseline salivary cortisol <1.8 ng/ml (<0.18 µg/dl) or an increment <3 ng/ml (0.3 µg/dl)⁴⁵³ following a standard-dose short Synacthen test are suggestive of RAI. However, even the evaluation of salivary cortisol is not without shortcomings.⁴⁵⁴

Recommendation

- Diagnosis of RAI should be based on a delta serum total cortisol after 250 µg corticotropin injection of <248 nmol/L (9 µg/dl) or a random total cortisol of <276 nmol/L (<10 µg/dl) (**II-2,1**). As serum free cortisol concentration can be influenced by the reduced serum levels of CBG and albumin frequently seen in patients with cirrhosis, salivary cortisol determination can be preferred (**II-2;2**).

Treatment of relative adrenal insufficiency

It is not known whether cortisol supplementation in clinically stable cirrhosis with RAI is of any value. Two studies have evaluated the effects of treating RAI in critically ill patients with cirrhosis. In one study 17 patients with cirrhosis and sepsis, in whom RAI was diagnosed, received i.v. hydrocortisone (50 mg/6h), and were compared with 50 consecutive patients with cirrhosis and septic shock who had previously been admitted to the same ICU but did not receive steroids. A higher rate of shock resolution, survival in the ICU and hospital survival were seen in the patients treated with hydrocortisone.⁴⁵⁵ In the second study, 57 patients with cirrhosis, septic shock and RAI were randomised to receive either i.v. 50 mg of hydrocortisone or normal saline every 6 h until haemodynamic stability was achieved, followed by steroid tapering over eight days. Lower vasopressor

Clinical Practice Guidelines

doses and higher rates of shock reversal were seen in patients who received hydrocortisone. However, 28-day mortality did not differ between the two groups. Moreover, shock relapse and GI bleeding occurred more often in the hydrocortisone group.⁴⁵⁶

Recommendation

- At present, hydrocortisone treatment (at a dose of 50 mg/6h) of RAI cannot be recommended (I-2).

Cardiopulmonary complications

Cirrhotic cardiomyopathy

Definition and pathophysiology

Cirrhotic cardiomyopathy (CCM) refers to chronic cardiac dysfunction in a patient with established cirrhosis, characterised by a blunted contractile response to stress (pharmacological/surgical or inflammatory) and an altered diastolic relaxation, often associated with electrophysiological abnormalities such as prolongation of the QTc interval. These phenomena occur in the absence of any other cardiac disease.⁴⁵⁷ Systemic inflammation is thought to be key in inducing myocardial dysfunction associated with impaired diastolic relaxation and decreased left ventricular ejection fraction, however, there are few controlled studies.^{193,458,459} Shear stress generated by portal hypertension exhibiting mechanical forces on myocardial fibres, may also play a part.⁴⁶⁰ CCM is largely subclinical but its presence does influence prognosis in advanced disease,⁴⁶¹ and it certainly impacts on the course of interventions such as TIPS and LT.⁴⁶²

Diagnosis

Characterisation of systolic dysfunction in cirrhotic cardiomyopathy

Systolic dysfunction refers to impaired left ventricle contractile responses to stress on echo, translating to a resting left ventricular ejection fraction (LVEF) <55%. For most patients with cirrhosis, the resting systolic function is normal or even increased, due to the hyperdynamic circulation and reduced afterload to maintain cardiac output. To investigate systolic dysfunction in cirrhosis, it is necessary to induce circulatory stress either pharmacologically or through exercise. Systolic dysfunction then manifests as a lack of an appropriate left ventricular contractile response to the applied stress. As disease advances, the progressive reduction in peripheral vascular resistance unmasks systolic dysfunction. Early studies used exercise stress testing to demonstrate a lack of increment in cardiac output or LVEF^{463,464} and this was even shown when noradrenaline levels were increased, suggesting loss of sympathetic responsiveness.⁴⁶⁵ More recent studies used pharmacological stress echo to show a blunted response.⁴⁶⁶ However, other studies using cardiac MRI, have shown normal chronotropic and inotropic responses suggesting the techniques used may give rise to variability.⁴⁶⁷

Myocardial strain imaging for assessing systolic dysfunction

Myocardial strain imaging is a more recent echocardiographic technique evaluating the degree of shortening of myocardial muscle fibres ('strain') influencing cardiac wall motion. The measurement of left ventricular global longitudinal systolic strain (GLS) is believed to be a sensitive marker of left ventric-

ular systolic function and facilitates the assessment of systolic dysfunction at rest,^{468,469} as well as having prognostic importance in heart failure.⁴⁷⁰ Studies of strain imaging in cirrhosis have demonstrated variable results; some showing impaired systolic strain in patients compared with healthy controls, albeit with no correlation to Child-Pugh score.^{471,472} Others demonstrate systolic strain within normal range and not influenced by the presence of ascites.^{473,474} However, interestingly, when patients undergo LT, the systolic strain improves.⁴⁷¹

Characterisation of diastolic dysfunction in cirrhotic cardiomyopathy

Numerous echocardiographic criteria along with transmitral Doppler evaluation have been used to characterise diastolic dysfunction including, early diastolic/atrial filling ratio (E/A), early diastolic filling/mitral annular velocity (E/e') and tricuspid systolic jet velocity. Such measurements are influenced by the pre- and afterload changes of portal hypertension. The latest American Society of Echocardiography (ASE) and European Association of Cardiovascular Imaging guidelines for the evaluation of diastolic dysfunction recommend the following criteria based on a normal LVEF (often the case in cirrhosis):⁴⁷⁵

- Average E/e' > 14
- Septal e' velocity < 7 cm/s OR Lateral e' velocity < 10 cm/s
- Tricuspid velocity > 2.8 m/s
- Left atrial volume index (LAVI) > 34 ml/m²

Diastolic dysfunction translates to impaired relaxation of the left ventricle, abnormal filling of the left atrium, and a higher left atrial volume. Indeed, increased LAVI has been associated with greater risk of heart failure in ischaemic cardiac disease.⁴⁷⁶ Based on these guidelines, diastolic dysfunction is classified as: grade I if one of the three principle criteria (1, 3 and 4 above) are met; and grade II if two or more of the criteria are met.

However, there is heterogeneity in descriptions of the prevalence of diastolic dysfunction in cirrhosis, in part reflecting the different echo techniques and/or diagnostic criteria applied, and the influence of vasoactive agents such as beta-blockers and terlipressin.

Several studies using an E/A ratio of ≤ 1 criteria have demonstrated left atrial enlargement in patients with ascites and advanced disease.^{464,477} Therapeutic paracentesis improves E/A ratio and importantly, in all studies, there is no relation to aetiology.⁴⁷⁸ In patients treated with TIPS there was no relation to aetiology, but diastolic dysfunction does show a positive correlation with higher MELD scores.^{462,479} A further study using E/e' ratio to define diastolic dysfunction in patients with ascites and elevated plasma renin, demonstrated that an increased E/e' was an independent predictor of development of HRS type 1 and one-year mortality.⁴⁸⁰ By contrast, other studies fail to show a clear relationship with disease severity or survival,^{460,472,481} albeit in two studies echo criteria are not specified.^{460,472} Studies using the LAVI criteria suggest a closer association between left atrial enlargement and Child-Pugh C disease.⁴⁸²

Newer techniques such as cardiac MRI with 'T1 mapping' and Late Gadolinium enhancement are being deployed to assess whether fibrosis or oedema modulates myocardial function in conditions such as amyloid and Fabry disease.⁴⁸³ Literature on using such techniques in liver disease are very limited. A recent study in patients with chronic hepatitis C showed no significant differences in echo parameters to controls, but demonstrated

lower post-contrast myocardial T1 time and higher partition coefficients, indicative of diffuse myocardial fibrosis.⁴⁸⁴

Natural history

Impact of disease related physiological stress: sepsis, decompensation and/or GI bleeding

There are limited studies of cardiac assessment during acute cirrhosis decompensation and associated haemodynamic instability. In a seminal study of acute cirrhosis decompensation with SBP, a subgroup with HRS were shown to have lower cardiac outputs at diagnosis and this correlated inversely with high inflammatory indices.¹⁹³ A follow-on study by the same group identified that patients with HRS had cardiac outputs that were further reduced at follow-up, compared to patients who do not develop HRS after SBP, and these patients had higher plasma noradrenaline and renin.⁴⁸⁵ Other studies recapitulate this with a demonstration of reduced kidney blood flow and importantly, suggest those with low cardiac index, also have increased mortality.^{192,485}

In relation to systemic inflammation and sepsis, one study showed lipopolysaccharide binding protein (LBP) levels (a surrogate for BT and lipopolysaccharide) in patients with ascites to be associated with significant diastolic dysfunction and left atrial enlargement. The E/e' ratio in these patients correlated with LBP levels. This data supports findings from experimental studies, which have shown a role for inflammation, signalling through the inflammasome and macrophage activation, as key pathological processes related to myocardial dysfunction.^{486–488}

Acute GI bleeding in cirrhosis is understandably associated with significant haemodynamic disturbances and has not been studied systematically in relation to cardiac function. Data assessing chronotropic function suggest the QTc interval is increased in cirrhotic patients during an acute bleeding episode compared to non-cirrhotic patients, and that this is associated with higher MELD scores and independently predicts survival.⁴⁸⁹ This contrasts with a more recent study that fails to demonstrate a clear link between QTc prolongation and mortality.⁴⁹⁰ Possible reasons for this heterogeneity in outcomes are the variable nature of vasoactive agents and their respective doses required for the control of bleeding in these studies. For example, one study showed terlipressin decreased cardiac output by 17% and the reduction in wall motion after terlipressin correlated with the Child-Pugh score.⁴⁹¹

Impact of interventions on cirrhotic cardiomyopathy

TIPS. Cardiac reserve is a major clinical consideration for elective TIPS placement and a 2D echo to assess LVEF is standard practice. Despite this, some patients do have cardiac decompensation post TIPS insertion. Several studies show an association between presence of diastolic dysfunction at the time of TIPS and poor survival.^{462,479} By contrast, others have not shown any difference in survival between patients with and without diastolic dysfunction at the time of TIPS.⁴⁹² However, uniformly, studies suggest an increase in left ventricular and atrial volume over time, implying that such patients may be at greater risk of future heart failure, based on literature for ischaemic heart disease and dilated cardiomyopathy.^{476,493}

Liver transplantation. Just as data on the effects of cirrhosis complications on cardiac function are variable, data on the impact of

the physiological stress of LT on patients with pre-existing cardiac dysfunction is heterogenous, largely because of the different echo criteria and thresholds applied.

One study in 173 transplant recipients assessed systolic (resting ejection fraction <55%) and diastolic (E/A ratio <1 or a deceleration time >200 ms) dysfunction and reported it occurring in 2% and 43% of patients, respectively. Whilst patients with diastolic dysfunction were older, interestingly, outcomes were not influenced by the presence of diastolic dysfunction.⁴⁹⁴ By contrast, another study used echo and brain natriuretic peptide (BNP) levels to grade severity of cardiac dysfunction. Those patients with higher BNP levels (>391) on day one tended to have higher mortality and longer dialysis requirements post transplantation. Of these, a subset with BNP levels >567 had ejection fractions <50%, and some of these died of cardiogenic shock within 72 h post-transplant. Autopsy in these patients showed diffuse myocardial fibrosis. In the main, BNP levels tended to decrease towards normal values over a week.⁴⁹⁵

A further study performed a detailed echo assessment, including myocardial strain assessment with speckle tracking, in patients undergoing LT compared to non-transplanted patients over a median follow-up of 18 months. Whilst patients pre-transplant had increased left ventricular mass and diastolic dysfunction, following transplantation, there was a significant improvement in systolic strain and reduced left ventricular mass. Conversely, cirrhosis patients who were not transplanted had an increase in left ventricular mass, albeit systolic strain did not change significantly.⁴⁷¹ This implies that some of the pathophysiological changes in CCM, such as increased left ventricular mass and size, are reversible with resolution of the disease. However, studies with comprehensive characterisation of cardiac function post transplantation are limited.

Prognosis for cirrhotic cardiomyopathy

Data relating cardiac dysfunction (especially diastolic dysfunction) with survival is variable. Some prospective studies, despite detailed evaluation of patients, including those with ascites and using speckle tracking, show no relation between cardiac dysfunction and survival, even among more decompensated patients.^{473,496} Many of the patients in these studies have evidence of diastolic dysfunction and some with even advanced grade II diastolic dysfunction albeit the GLS values in these studies are within the normal range.

Conversely, other studies suggest an association between presence of diastolic dysfunction and higher two-year mortality, with diastolic dysfunction ranging from 38–67%, especially in patients with severe ascites.^{461,497} Indeed, in one such study, a multivariate analysis showed left ventricular diastolic dysfunction was an independent predictor of mortality.⁴⁶¹ Another study followed 80 patients to assess one-year mortality, finding 46% had diastolic dysfunction on echo criteria and about half of these had grade II dysfunction, in whom mean arterial blood pressure was lower and MELD score higher than grade I patients. The presence of diastolic dysfunction was associated with a higher degree of ascites and plasma renin levels and 38% of these patients developed criteria for HRS type I. Survival was 95% in those without diastolic dysfunction, compared to 79% in those with grade I dysfunction and 39% with grade II diastolic dysfunction. E/e' ratio was an independent predictor of survival.⁴⁸⁰

Recommendations

- Evaluation of cirrhosis patients with echocardiography should be performed with dynamic stress testing either pharmacologically, or through exercise, given that systolic dysfunction may be masked by the hyperdynamic circulation and reduced afterload. Failure to increment cardiac output after physiological/pharmacological stress (and in the absence of influence of beta-blockade) indicates systolic dysfunction (II-1;1).
- Myocardial strain imaging and assessment of GLS may serve as a sensitive marker of left ventricular systolic function and facilitate its assessment at rest and in decompensated patients (II-2;2). Cardiac MRI may also identify structural changes. However, with all these techniques, there is the need for more controlled studies and correlation with clinical endpoints (III;2).
- Diastolic dysfunction may occur as an early sign of cardiomyopathy in the setting of normal systolic function, and should be diagnosed using the recent ASE guidelines, namely: Average $E/e' > 14$; Tricuspid velocity > 2.8 m/s and LAVI > 34 ml/m² (II-1;1).
- In patients with AD of cirrhosis, reduced cardiac output (as a manifestation of CCM) is of prognostic significance as it is associated with the development of AKI (specifically hepatorenal dysfunction) after infections such as SBP (II-1;1).
- Prolongation of the QTc interval is common in cirrhosis and can be evaluated since it may indicate a poor outcome. Agents that can prolong the QT interval should be used cautiously (II-2;2).
- Detailed functional cardiac characterisation should be part of the assessment for TIPS insertion (II-2;2) or LT (II-1;1).
- Standardized criteria and protocols for the assessment of systolic and diastolic function in cirrhosis are needed (II-2;2).

Hepato-pulmonary syndrome

Definitions and clinical manifestations

The association of chronic liver disease with respiratory symptoms and hypoxia is well recognised. Four main pulmonary

complications may occur in patients with chronic liver disease: pneumonia, hepatic hydrothorax, HPS and PPHT. HPS is defined as a disorder in pulmonary oxygenation, caused by intrapulmonary vasodilatation and, less commonly, by pleural and pulmonary arteriovenous communications occurring in the clinical setting of portal hypertension.^{498,499} It is most commonly diagnosed in patients with cirrhosis^{498,499} and portal hypertension⁵⁰⁰ but, it has also been described in patients with pre-hepatic portal hypertension,⁵⁰¹ with venous obstruction but without cirrhosis, and even in patients with acute or chronic hepatitis⁵⁰⁰ (Table 12). A severe impairment of liver function and a specific aetiology of liver disease are not needed for the development of HPS,⁴⁹⁸ based on the profiles of the patients studied. In terms of prevalence, HPS has been reported in 10% of patients with chronic viral hepatitis in 15–23% of those with cirrhosis and in 28% of those with Budd-Chiari syndrome.^{502–504} However, the prevalence of HPS reported in patients with cirrhosis undergoing LT evaluation ranges from 5–32%,^{504–508} while intrapulmonary vascular dilatation (IPVD) can be detected by echocardiography in 50–60% of cirrhosis patients undergoing LT evaluation. No relationship seems to exist between HPS and CCM.⁵⁰⁴ The clinical manifestations of HPS in patients with chronic liver disease primarily involve dyspnoea and platypnoea.^{498,502,506} Dyspnoea is the most common respiratory complaint in patients with HPS, but it is non-specific. Its onset is insidious, usually occurring on exertion. Platypnoea, which is a shortness of breath exacerbated by sitting up and improved by lying supine, is a less sensitive but a more specific finding in these patients. Hypoxemia with exertion or at rest is common and it is exacerbated in the upright position (orthodeoxia). There are no signs or hallmarks of HPS on physical examination. However, tachypnoea and polypnoea, digital clubbing and/or cyanosis in patients with the hallmarks of chronic liver disease suggest the presence of HPS.^{498,502,506}

Pathophysiology

The pathophysiology of HPS is characterised by an IPVD occurring within the pulmonary arterial circulation. This vascular abnormality consists of diffuse or localised abnormal dilated pulmonary capillaries and, less commonly, pleural and pulmonary arteriovenous communications,⁵⁰⁹ which result in impaired oxygenation of venous blood as it passes through the pulmonary circulation. IPVD impairs ventilatory/perfusion (V/Q) ratio and may result in anatomic and functional shunt leading to hypoxaemia (Fig. 10). In patients with advanced liver cirrhosis this leads to a subtle increase in intrapulmonary blood shunting, which is more pronounced in patients with HPS. The consequent increase of shunting and V/Q mismatch in the

Table 12. Diagnostic criteria of hepatopulmonary syndrome.

Hypoxia with partial pressure of oxygen < 80 mmHg or alveolar–arterial oxygen gradient ≥ 15 mmHg in ambient air (≥ 20 mmHg in patients older than 65 years).

Pulmonary vascular defect with positive findings on contrast-enhanced echocardiography or abnormal uptake in the brain ($> 6\%$) with radioactive lung-perfusion scanning

Commonly in presence of portal hypertension, and in particular:

- hepatic portal hypertension with underlying cirrhosis
- pre-hepatic or hepatic portal hypertension in patients without underlying cirrhosis

Less commonly in presence of:

- acute liver failure, chronic hepatitis

All criteria were determined by means of positive contrast-enhanced echocardiography (*i.e.*, microbubble opacification of the left heart chambers three to six cycles after right atrial passage). The abbreviated formula for the alveolar–arterial gradient is as follows: $PaO_2 - PaO_2 = (FIO_2 [Patm - PH_2O] - [PaCO_2/0.8]) - PaO_2$, where PaO_2 denotes partial pressure of alveolar oxygen, PaO_2 partial pressure of arterial oxygen, FIO_2 fraction of inspired oxygen, $Patm$ atmospheric pressure, PH_2O partial pressure of water vapor at body temperature, and $PaCO_2$ partial pressure of arterial carbon dioxide (0.8 corresponds to the standard gas-exchange respiratory ratio at rest); the normal range is 4 to 8 mmHg.

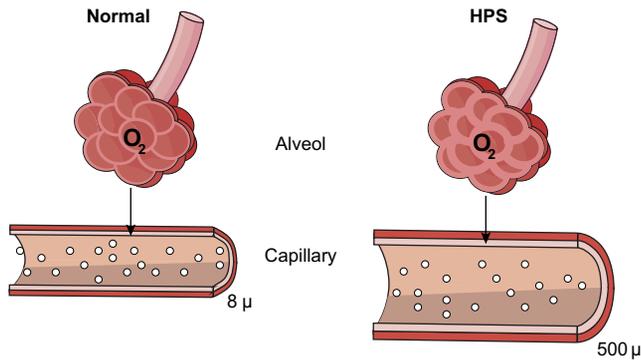


Fig. 10. The pathophysiology of hepatopulmonary syndrome (adapted from Ref. 498). HPS, hepatopulmonary syndrome.

upright position is the cause of the orthodeoxia.⁵¹⁰ The pathogenesis of IPVD is probably multifactorial (Fig. 11). The release of nitric oxide, which is a potent vasodilator, plays a critical role in the development of HPS. The increased release of nitric oxide in the pulmonary circulation is related to an increased expression and activity of two isoforms of nitric oxide synthase (NOS), the endothelial NOS (eNOS) and the inducible NOS (iNOS).^{511–516} Meanwhile, BT and the BT-related endotoxaemia and pro-inflammatory response also contribute to the accumulation of macrophages in the pulmonary microvasculature.⁵¹⁷ Endothelial activation of fractalkine (CX3CL1), a chemokine, in the lung may favour the adherence of monocytes in the pulmonary microcirculation.⁵¹⁸ Monocytes express iNOS and produce heme oxygenase-1, leading to increased carbon monoxide production, further enhancing vasodilatation.⁵¹⁹

CX3CL1 and vascular endothelial growth factor (VEGF) A, produced by circulating monocytes, also contribute to angiogenesis, recently recognised as a further pathogenetic factor of pulmonary IPVD in experimental HPS.^{520–522} A downregulation of miRNA-199 a-5p has recently been described as a contributory mechanism of pulmonary microvascular endothelial cell proliferation and thus pathogenesis of HPS.⁵²³ Polymorphisms in genes involved in the regulation of angiogenesis have also been associated with the risk of HPS in patients with cirrhosis⁵²⁴ (Fig. 11). Finally, it has recently been observed that rosuvastatin, by down-regulating protein expression of nuclear factor kappa B and VEGF-1,2 and Rho-associated A kinase, may improve the intrapulmonary angiogenesis and the alveolar-arterial oxygen pressure gradient in common bile duct ligation rats.⁵²⁵

Diagnosis

In patients with portal hypertension and the clinical suspicion of HPS partial pressure of oxygen (PaO₂) in arterial blood gas (ABG) should be assessed. A PaO₂ lower than 80 mmHg and or an alveolar-arterial oxygen gradient (P[A-a]O₂) ≥15 mmHg while breathing ambient air at sea level should lead to further investigations (Table 12). For adults ≥65 years a P[A-a]O₂ ≥20 mmHg cut-off is used.⁵²⁶ However, it should be highlighted that although these criteria are well established, enabling one to unify the diagnostic methods and thus to better understand the disease, they are based on a consensus of experts. Pulse oximetry indirectly measures oxygen saturation (SpO₂), it is non-invasive and may be useful in the diagnosis of HPS in adults since a SpO₂ <96% was found to be highly sensitive (100%) and specific (88%) for detecting HPS in patients with a PaO₂ <70

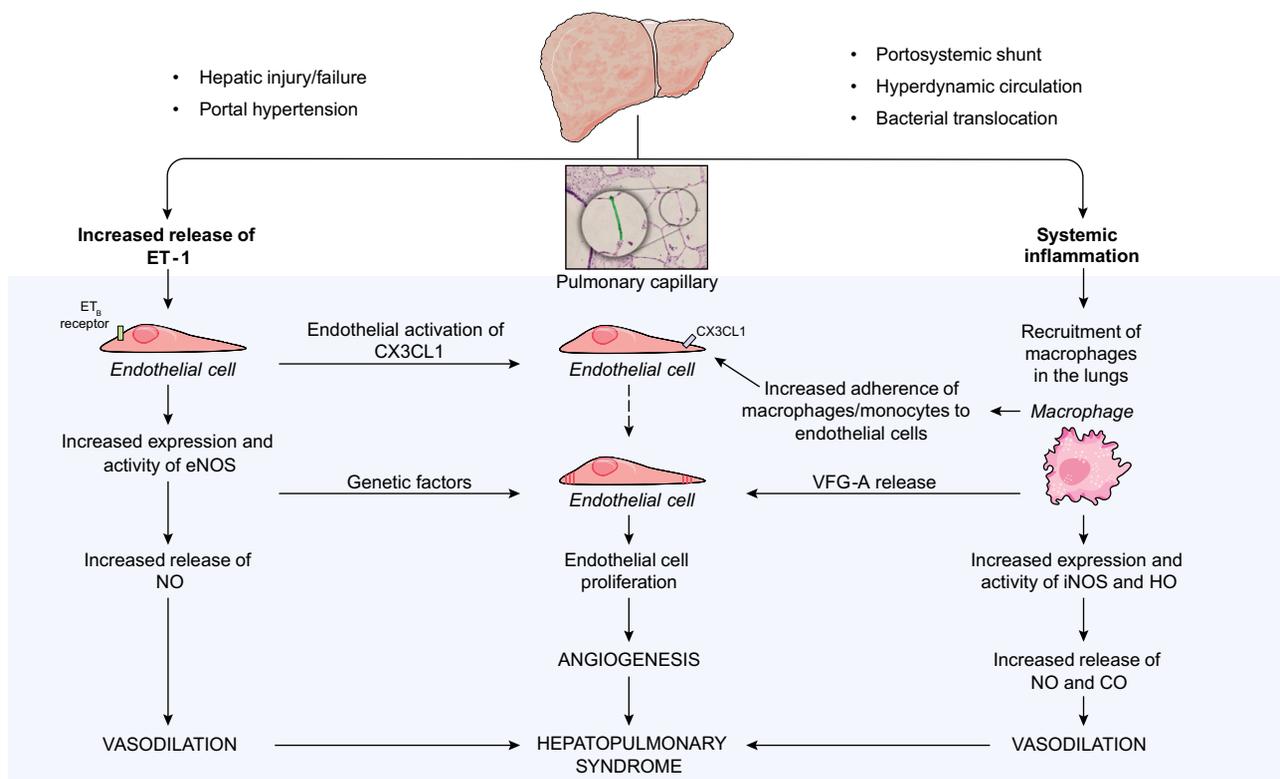


Fig. 11. The pathogenesis of hepatopulmonary syndrome. ET, endothelin; eNOS, endothelial nitric oxide synthase; iNOS, inducible nitric oxide synthase; NO, nitric oxide; HO, Heme oxygenase-1; CO, carbon monoxide; CX3CL1, fractalkine; VEGF-A, vascular endothelial growth factor A.

Clinical Practice Guidelines

mmHg, limiting ABG testing to only 14% of patients.⁵²⁷ The validity of this non-invasive approach was not confirmed, recently, in paediatric patients with HPS.⁵²⁸ Serial SpO₂ measurements may be useful to monitor impaired oxygenation over time in patients with HPS. The ABG is essential for the staging of the severity of HPS. HPS can be categorised as mild (PaO₂ ≥80 mmHg), moderate (PaO₂ 60–79 mmHg), severe (PaO₂ 50–59 mmHg), and very severe (PaO₂ <50 mmHg).^{498,500,501,503} Recently, it has been observed that HPS is associated with elevated von Willenbrand factor antigen (vWF-Ag) levels. Thus, vWF-Ag has been proposed as a potentially useful screening tool for early detection of HPS, but further studies are needed to validate it.⁵²⁹ The chest X ray is usually non-specific, nevertheless, it can be used to effectively rule out other concomitant pulmonary diseases since only a mild interstitial pattern in the lower part of the lungs may be found, because of pulmonary vasodilatation.^{498,500,501,503} A decrease in the single-breath diffusing capacity for carbon monoxide is the only alteration of the routine pulmonary function test that is frequently and consistently abnormal in patients with HPS. However, it is not specific and it may not normalise after LT.^{498,500,501,503} All the other respiratory function tests are non-specific, showing normal or reduced forced vital capacity or maximum forced expiratory volume during the first second (FEV1). Thus, they can only be used to rule out other concomitant pulmonary diseases. Thoracic CT scans have also been proposed as a complementary technique to rule out another underlying pulmonary pathology,^{498,499} although there is little information regarding their specific role in the diagnosis of HPS. It has been suggested that thoracic CT scans can be useful to measure the calibre of the peripheral arteries and the bronchial/arterial relationship.^{530,531} Furthermore, CT scanning makes it possible to define the vascular pattern of HPS in a similar manner to arteriography by detecting pleural and pulmonary arteriovenous communications. Contrast-enhanced transthoracic echocardiography with saline (shaken to produce microbubbles >10 μm in diameter) is the most useful method to detect pulmonary vascular dilatation. After the administration of agitated saline in a peripheral vein, microbubble opacification of the left atrium within three to six cardiac cycles after right-atrial opacification indicates microbubble passage through an abnormally dilated vascular bed, since microbubbles do not pass through normal capillaries.⁵³² The injection of technetium-99 m-labeled macro-aggregated albumin (MAA) in the peripheral vein for lung scanning (MAA scan) is a potential alternative diagnostic procedure although it is more invasive and less sensitive. Particles, with a 20–50 μm size, escape through the abnormal pulmonary capillaries and stay in downstream capillary beds supplied by systemic arteries, such as the brain, kidneys, and spleen. Quantitative imaging of the MAA scan in the brain and lung enables calculation of the degree of shunting.^{533,534} The measurement of shunting with MAA scans may be useful as a complementary diagnostic tool in patients with HPS in two clinical situations. Firstly, in patients with a severe hypoxaemia and a coexistent HPS and intrinsic lung disease since a shunting >6% at MAA scan proves the major contribution of HPS to hypoxaemia. Secondly, in patients with HPS and very severe hypoxaemia (PaO₂ <50 mmHg), since the presence of shunting >20% is associated with a poor outcome after LT.⁵³⁵ Despite the potential role of lung perfusion scintigraphy for prognostic use in patients with cirrhosis and IPVD, its diagnostic accuracy for HPS remains to be established.⁵³⁶ Finally, neither contrast eco-

cardiography nor MAA scan can differentiate discrete arteriovenous communications from diffuse precapillary and capillary dilatations or intracardiac shunt. The former distinction can be made by means of pulmonary angiography. The latter distinction can be made by means of transoesophageal contrast-enhanced echocardiography that directly reveals the intra-atrial septum. Pulmonary angiography should not be performed in all patients with suspected HPS, but only in: a) patients with the severe hypoxaemia (PaO₂ <60 mmHg) poorly responsive to administration of 100% oxygen, and b) patients strongly suspected (by means of a CT chest scan) of having arteriovenous communications that would be amenable to embolisation.

Recommendations

- In presence of tachypnoea and polypnoea, digital clubbing and/or cyanosis in a patient with the hallmarks of chronic liver disease, HPS should be suspected and investigated (**II-2;1**).
- Pulse oximetry is the screening tool for HPS in adult patients, but not in paediatric patients. For patients with SpO₂ <96%, ABG analysis should be performed. A PaO₂ lower than 80 mmHg and or an alveolar-arterial oxygen gradient (P[A-a]O₂) ≥15 mmHg while breathing ambient air, should lead to further investigations. For adults ≥65 years a P[A-a]O₂ ≥20 mmHg cut-off should be used (**II-2;1**).
- The use of contrast (microbubble) echocardiography to characterise HPS is recommended (**II-2;1**).
- Trans-oesophageal contrast-enhanced echocardiography can be performed to exclude definitively intra-cardiac shunts, albeit this technique is not devoid of risks (**II-2;2**).
- An MAA scan should be performed as a complementary tool to quantify the degree of shunting in patients with severe hypoxaemia and coexistent intrinsic lung disease, or to assess the prognosis in patients with HPS and very severe hypoxaemia (PaO₂ <50 mmHg) (**II-2;1**).
- Neither contrast echocardiography nor MAA scan can definitively differentiate discrete arteriovenous communications from diffuse precapillary and capillary dilatations or cardiac shunts. Pulmonary angiography should be performed only in patients with the severe hypoxaemia (PaO₂ <60 mmHg), poorly responsive to administration of 100% oxygen, and in whom there is a strong suspicion of arteriovenous communications that are amenable to embolisation (**II-2;1**).

Natural history

The natural history of IPVD as well as of HPS is still unclear. Most patients with IPVD maintain a normal gas exchange over time, and it is not clear the reason why a subset of patients with IPVD develops HPS.⁵³⁷ A diagnosis of HPS is associated with a poor outcome in terms of both survival and quality of life.^{505,507,508} Regarding survival, it should be highlighted that in patients undergoing evaluation for LT, the mortality rate was almost double in patients with HPS compared to patients with cirrhosis without HPS, independent of other potential predictors of mortality such as age, MELD score and comorbidities.⁵⁰⁵ In patients with cirrhosis and HPS, who were not

evaluated for LT, the five-year survival rate was 23% while it was 63% in patients with cirrhosis without HPS who were matched for the aetiology and severity of cirrhosis according to the Child-Pugh classification, age, and MELD score.⁵⁰³ Survival was significantly worse among patients with HPS and a PaO₂ of less than 50 mmHg at the time of diagnosis.^{506,507}

Management

Medical treatment

Spontaneous resolution of HPS is uncommon. There is no established medical therapy currently available for HPS. Several drugs have been applied for the treatment of HPS with conflicting results. However, no large randomised trial has been conducted, probably because of the low number of patients. Data from several uncontrolled clinical studies and anecdotal evidence indicate that treatment with beta-blockers, cyclooxygenase inhibitors, systemic glucocorticoids and cyclophosphamide, almitrine bismesylate, inhaled nitric oxide, nitric oxide inhibitors, and antimicrobial agents has been uniformly unsuccessful.⁵⁰⁵ Pentoxifylline has also been tried in the treatment of HPS in adults and children in two small pilot studies with contradictory results in terms of improvements in oxygenation and frequent GI side effects.^{538,539} Administration of garlic was found to be associated with an improvement in the PaO₂ in a small randomised study.⁵⁴⁰ However, a case of moderate hepatotoxicity associated with short-term, high-dose garlicin therapy in an LT recipient with persistent HPS was recently reported.⁵⁴¹ The use of TIPS has been proposed to reduce portal pressure in patients with HPS. However, data are insufficient even when a systemic analysis review is considered.⁵⁴² In addition, there is some concern that TIPS can enhance pulmonary vasodilation by exacerbating the hyperkinetic circulation. Thus, no recommendation for the use TIPS to treat HPS can be given.^{498,505} Finally, coil embolisation (embolotherapy) has been shown to improve arterial oxygenation temporarily in the context of angiographic arteriovenous communications.^{531,543} Endothelin-1 receptor antagonists or angiogenesis inhibitors have not been tested up to now in patients with HPS. Thus, long-term oxygen therapy remains the most frequently recommended therapy for symptoms in patients with severe hypoxaemia. However, some aspects of this treatment such as efficacy, costs, and compliance, remain to be evaluated.

Recommendations

- Long-term oxygen therapy is recommended in patients with HPS and severe hypoxaemia. Nevertheless, there is no available data concerning effectiveness, tolerance, cost-effectiveness, compliance and effects on survival rates of this therapy (**II-2;1**).
- No recommendation can be proposed regarding the use of drugs or the placement of TIPS for the treatment of HPS (**I;1**).

Liver transplantation

The most common and the only successful treatment for HPS is LT. LT results in a complete reversal or in a significant improvement of HPS in more than 85% of patients with severe hypoxaemia.⁵⁴⁴ In a prospective clinical study performed in the pre-MELD era, a pre-LT severe hypoxaemia, in particular when it

was associated with a large shunting at MAA scan, was found to be a very strong predictor of mortality after LT.⁵³⁵ In 2007, five years after the introduction of MELD in US, the United Network for Organ Sharing (UNOS) recommended assigning a MELD score of 22 for the initial application of patients with severe HPS (PaO₂ <60 mmHg), with further increases every three months, to balance pre- and post-LT outcomes between HPS and non-HPS candidates.⁵⁴⁵ In the largest retrospective study comparing the results of LT between the pre-MELD era and the MELD era in patients with HPS, the five-year survival rate after LT was found to improve from 67% during the pre-MELD era to 88% in the MELD era.⁵⁴⁶ Other data showed that in post-MELD era, there was no association between pre-LT oxygenation and waitlist survival in patients with HPS. These findings reflect not only the results of the introduction of HPS as a MELD exception, but also of an improved perioperative management in patients with HPS. The regular assessment of the severity of hypoxaemia may facilitate LT prior to the occurrence of very severe hypoxaemia. In fact, hypoxaemia can worsen in patients with HPS who are on the active transplantation list, with a median decrease in pO₂ of 5.2 mmHg per year,⁵⁰⁸ and it has been recently confirmed that a pre-LT room-air PaO₂ ≤44.0 mmHg is still associated with increased post-LT mortality.⁵⁴⁷ Thus, it has been suggested that an ABG analysis should be carried out every six months, but no study has clarified which is the best method for conducting this (ABG analysis vs. pulse oximetry) nor how frequently it should be performed. Despite the increased survival rate in patients with HPS after LT in the MELD era, it has recently been observed that HPS MELD exception patients had lower overall mortality compared to others awaiting LT, suggesting that the appropriateness of the HPS MELD exception policy should be reassessed.⁵⁴⁸ There are very few and small studies on the impact of HPS on anaesthetic procedures, as well as in the post-LT management in the ICU. Nevertheless, it seems that inhaled nitric oxide, methylene blue, extracorporeal membrane oxygenation and non-invasive ventilation may improve oxygenation immediately post-LT.⁵⁴⁹⁻⁵⁵¹

Recommendations

- Patients with HPS and PaO₂ <60 mmHg should be evaluated for LT since it is the only treatment for HPS that has been proven to be effective to date (**II-2;1**).
- Since a severe hypoxaemia (PaO₂ <45–50 mmHg) is associated with increased post-LT mortality, an ABG analysis should be carried out every six months in order to facilitate prioritisation to LT (**II-2;1**).

Portopulmonary hypertension

Definition and diagnosis

A diagnosis of PPHT should be considered in a patient with established portal hypertension in the absence of other causes of pulmonary artery or venous hypertension, namely: chronic thromboembolism, chronic lung disease/hypoxia; chronic left heart disease.

Patients may be asymptomatic but often present with exertional dyspnoea and they may have clinical signs of right heart failure when moderate to severe disease develops.⁵⁵² Classification of severity is based on mean pulmonary arterial pressure

Clinical Practice Guidelines

(mPAP) and assumes there is high pulmonary vascular resistance (PVR). PPHT is graded as mild (mPAP ≥ 25 and < 35 mmHg); moderate (mPAP ≥ 35 and < 45 mmHg), and severe (mPAP ≥ 45 mmHg).⁴⁹⁸ The diagnosis also requires there to be normal pulmonary occlusion pressures, to exclude elevation of pulmonary pressure resulting from elevated left ventricular filling pressure. Transthoracic Doppler echocardiography (TDE) is the main screening tool for evaluating the presence of PPHT when screening high-risk patients, such as those being considered for TIPS or LT.^{553–555} As a screening test, some studies suggest a pulmonary artery systolic pressure of > 30 mmHg on TDE has a negative predictive value of 100%, but a positive predictive value of only 59%.⁵⁵⁴ However, when assessing patients for LT, the threshold for right heart catheterisation is less clear, with a right ventricular systolic pressure > 50 mmHg and/or significant right ventricular hypertrophy seen as the trigger for this investigation to rule out significant PPHT.⁵⁵⁵

Pathophysiology

In patients with portal hypertension, PPHT is thought to arise from limited blood flow in the pulmonary arterial circulation because of vasoconstriction. Numerous factors are thought to be responsible for this including: Changes in endogenous vaso-regulators; increased endothelin 1 and reduced prostacyclin synthase from pulmonary endothelial cells; proliferation of smooth muscle cells/endothelial activation and platelet aggregation.

Natural history and prognosis

From studies in patients evaluated for LT, the incidence is thought to be between 3–10% based on haemodynamic criteria. Furthermore, female sex and pre-existing autoimmune liver disease are thought to be independent risk factors.⁵⁵⁶ Genetic variation in oestradiol levels may increase the predisposition to pulmonary artery vasoconstriction. Indeed, women are at three times greater risk than men.⁵⁵⁷ There is also an association between patients who have moderate to severe PPHT and the presence of large portosystemic shunts.⁵⁵⁸ However, there is no clear association between the severity of liver disease or portal hypertension and the development of severe PPHT.^{556,559} Studies quote survival rates at one year of between 35–46% without specific treatment.^{560,561} Mortality is often associated with other complications of liver disease such as hepatocellular cancer, sepsis and GI bleeding and right ventricular failure. Increased rates of mortality are related to higher right atrial pressure and lower cardiac index.^{559,562} In a multicentre registry study, patients with PPHT were shown to have worse outcomes than patients with idiopathic pulmonary hypertension, with a five-year survival of 40% vs. 64%.⁵⁶³ However, a retrospective French study challenges this, whilst reporting increased mortality in those with a lower cardiac index, likely reflecting failed compensation to increased right ventricular dysfunction, and patients with more advanced liver disease.⁵⁶⁴

Medical treatment

The evidence base for pharmacological therapies in PPHT is limited with most data extrapolated from studies in pulmonary arterial hypertension not related to liver disease.^{565,566} Drugs to promote acute vasodilatation during right heart catheterisation assessment, theoretically may be deleterious as they run the risk of further reducing cardiac index. There is a lack of data to clarify this.⁵⁶⁷ Conversely, whilst patients with advanced

portal hypertension may be on treatment with beta-blockers, withdrawing beta-blocker therapy may help to increase cardiac output and thereby help exertional dyspnoea, in patients with advanced PPHT.⁵⁶⁸

Endothelin receptor antagonists. Bosentan has been shown to improve pulmonary artery haemodynamics and exercise tolerance in patients with PPHT, independently of liver disease severity.^{569–572} One retrospective study reports survival rates of up to 89% at three years.⁵⁷³ Others have shown improvements in cardiac index up to 39%, albeit in a small number of patients, but an increase in aminotransferases, which responded to dose reduction or discontinuation.⁵⁷¹ The FDA places a caution on this class of drug in patients with advanced liver dysfunction. There is limited data on the use of other members in this family of agent, including ambrisentan and macitentan, for PPHT.^{574,575}

Phosphodiesterase subtype-5 inhibitors. Blockade of phosphodiesterase-5 inhibitors facilitate the vasodilatory effects of nitric oxide, through reduced metabolism of cGMP. Small case series suggest that sildenafil improves functional capacity and increases cardiac output.^{576–578} It should be noted that sildenafil can precipitate variceal bleeding and as such, caution should be exercised.⁵⁷⁹

Prostacyclin analogues. Prostacyclin analogues have many potential benefits including vasodilatory, reduced vascular smooth muscle proliferation and anti-thrombotic. Case series suggest improved pulmonary haemodynamics with i.v. epoprostenol and the potential for improved five-year survival compared to registry data in pulmonary artery hypertension (70 vs. 40%).^{580–583} However, lower doses than those used in idiopathic pulmonary hypertension are suggested to reduce the development of thrombocytopenia and splenomegaly. Other studies have also looked at use of inhaled iloprost and reported short-term haemodynamic benefit.⁵⁸⁴

Impact of the management of other complications of cirrhosis

Caution should be exercised when considering TIPS placement for the treatment of other complications of cirrhosis in patients with proven PPHT. The anticipated increase in right ventricular filling pressures and cardiac output may precipitate marked increases in PVR and right-sided pressure overload.^{585,586} Moderate PPHT (mPAP > 35 and < 45 mmHg) is a relative contraindication for TIPS placement, and severe PPHT is an absolute contraindication.⁵⁸⁶

Recommendations

- Screening for PPHT should be via TDE in patients deemed potential recipients for TIPS or LT; in those with a positive screening test, right heart catheterisation should be performed (**II-1;1**).
- In patients with PPHT who are listed for transplantation, echocardiography should be repeated on the waitlist, albeit, the specific interval is unclear (**III;1**).
- Beta-blockers should be stopped and varices managed by endoscopic therapy in cases of proven PPHT (**II-3;1**).

- Therapies that have been approved for primary pulmonary arterial hypertension may have benefit in PPHT to improve exercise tolerance and haemodynamics. However, endothelin antagonists should be used with caution because of concerns over hepatic impairment (**II-2;1**).
- TIPS should not be used in patients with PPHT (**II-3;1**).

Liver transplantation

Historically, severe PPHT has been a relative contraindication for LT because of very poor outcomes. However, with the advent of improved haemodynamic control with agents such as i.v. prostacyclin, there are case series showing normal pulmonary haemodynamics almost two years post LT.^{587,588}

Stratifying risk for LT. In patients with an mPAP ≥ 45 –50 mmHg, most centres would deem this an absolute contraindication to transplantation irrespective of therapy applied.^{562,587,589} Patients with an mPAP > 35 mmHg have increased risk post LT, associated with increased hospital stay and longer ventilator requirements.^{562,590,591} If LT is considered in such patients, it is suggested that their PPHT is treated aggressively to lower mPAP and improve right ventricular function.^{588,592,593}

To facilitate access to LT before there is further progression of PPHT to a point where transplantation risks are deemed too high, MELD exception (MELD 22 points) has been granted to patients with PPHT (mPAP > 25 mmHg and PVR > 240 dynes/s per cm^{-5}) with at least moderate disease severity (baseline mPAP > 35 mmHg).⁵⁹⁴ Patients are considered surgical candidates if, after targeted therapy to lower PAP, they have improved mPAP (< 35 mmHg) and PVR (< 400 dyne/s per cm^{-5}) and/or normalise their PVR. Applying this exception has been noted to reduce waitlist mortality.⁵⁹⁵

Per-operative considerations. All patients should be monitored with a pulmonary artery catheter. Therapy to lower mPAP should be continued throughout the operative period, given that there is often a rise in cardiac output post re-perfusion and this may add more stress on any pre-existing impaired right ventricle function.^{595–597} Indeed, in some cases, a severe acute rise in PAP may lead to graft failure because of hepatic congestion through a failing right ventricle. The management of such adverse haemodynamics, in addition to i.v. prostacyclin or inhaled nitric oxide includes the use of extracorporeal membrane oxygen therapy (ECMO).^{598,599}

Postoperative considerations. Monitoring PAP response to therapy is via serial transthoracic echo with tissue Doppler at 4–6-month intervals and consideration of tapering pulmonary artery targeted therapy, though no controlled data exists to provide guidance on this.^{581,600,601} Case reports and series suggest that 29–64% of patients with moderate to severe PPHT under long-term follow-up post-transplant have been able to discontinue therapy over time.^{599–602} Indeed, some suggest a return to normal right ventricle function following therapy for PPHT in the pre-transplant period and then after transplant surgery.^{581,601} PPHT MELD exception patients have worse one-year mortality or graft failure than patients without PPHT.⁶⁰³

Recommendations

- If mPAP < 35 mmHg and right ventricular function is preserved, LT should be considered (**II-2,1**). A mPAP of ≥ 45 mmHg should be considered an absolute contraindication to LT irrespective of therapy applied (**III,1**).
- Therapy to lower mPAP and improve right ventricular function should be commenced in patients with mPAP ≥ 35 mmHg. Right ventricular function should be periodically evaluated (**II-2,1**).
- MELD exception can be considered in patients with proven PPHT in whom targeted therapy fails to decrease mPAP < 35 mmHg but does facilitate normalisation of PVR to < 240 dynes/s cm^{-5} and right ventricular function (**II-3;2**).
- MELD exception should be advocated in patients with proven PPHT of moderate severity (assessment mPAP ≥ 35 mmHg) in whom targeted treatment lowers mPAP < 35 mmHg and PVR < 400 dynes/s cm^{-5} (**II-2;1**).

Conclusions

These guidelines on the management of patients with decompensated cirrhosis were developed based on a new pathophysiological background that offers the opportunity for more comprehensive therapeutic or prophylactic approaches to manage the disease. The knowledge of the key pathophysiological mechanisms makes it possible nowadays to counteract the progression of cirrhosis and so to prevent its complications. This represents a step forward, shifting our approach from treating the complications of decompensated cirrhosis to preventing their occurrence. However, to make this possible it is crucial to think about new models of specialist care for patients with cirrhosis. A care coordination programme, has been proven to improve survival and to reduce emergent readmission to the hospital in these patients.⁶⁰⁴ Care coordinators can facilitate the development of educational programmes for patients and caregivers optimising their adherence to guideline recommendations. In addition, they can plan invasive procedures in a day hospital, allowing transfer of real-time information to primary care physicians to improve quality and coordination of care. In so doing, it is possible to prevent unnecessary visits to the emergency department and/or emergent readmission to the hospital. These measures will progressively reduce the burden of cirrhosis.

Conflict of interest

Paolo Angeli: Consultancy fee from Sequana Medical AG, Gilead Italy and Biovie; Patent inventor from Biovie; Research grant from Gilead; Speaker's fee from Bhering, Kedrion 2016. Mauro Bernardi: Consultancy fee from CLS Behring GmbH, Baxter Healthcare SA, Grifols SA; Speaker's fee from CLS Behring GmbH, Baxter Healthcare SA, PPTA Europe, Octapharma AG, Gilead Sciences, AbbVie Italia. Wim Laleman: Speaker's fee for Gore, Norgine, 4C, Abbvie, Sirtex; Consultancy fee for AbbVie, Gilead, MSD, Intercept; Research grant from Gilead. Jonel Trebicka: Speaker's fee or Consultancy fee from Gore & associates (TIPS), Sequana medical (alpha-pump), Alexion (PNH), Versantis (liposomes). Aleksander Krag: None. Claire Francoz: None. Pere Gines: Advisory/Consultancy fee for Sequana Grifols, Mallinck-

Clinical Practice Guidelines

rodt, Ferring Pharmaceuticals; Research Funding from Sequana, Grifols, Ferring Pharmaceuticals.

Please refer to the accompanying ICMJE disclosure forms for further details.

Acknowledgments

We would like to thank Alessandra Brocca Dr, Marta Tonon MD, Husain-Syed Faeq MD for the great editorial work. We acknowledge ICREA for the ACADEMIA AWARD given to Pere Ginès. We would like to thank the reviewers of this Clinical Practice Guideline for their time and critical reviewing: EASL Governing Board, Alexander Gerbes, Thierry Gustot, Guadalupe Garcia-Tsao.

References

- [1] EASL clinical practice guidelines on the management of ascites, spontaneous bacterial peritonitis, and hepatorenal syndrome in cirrhosis. *J Hepatol* 2010;53:397–417.
- [2] D'Amico G. The clinical course of cirrhosis. Population based studies and the need of personalized medicine. *J Hepatol* 2014;60:241–242.
- [3] Moreau R, Jalan R, Gines P, Pavesi M, Angeli P, Cordoba J, et al. Acute-on-chronic liver failure is a distinct syndrome that develops in patients with AD of cirrhosis. *Gastroenterology* 2013;144:1426–1437.
- [4] D'Amico G, Garcia-Tsao G, Pagliaro L. Natural history and prognostic indicators of survival in cirrhosis: a systematic review of 118 studies. *J Hepatol* 2006;44:217–231.
- [5] Bernardi M, Moreau R, Angeli P, Schnabl B, Arroyo V. Mechanisms of decompensation and organ failure in cirrhosis: From peripheral arterial vasodilation to systemic inflammation hypothesis. *J Hepatol* 2015;63:1272–1284.
- [6] Jalan R, Fernandez J, Wiest R, Schnabl B, Moreau R, Angeli P, et al. Bacterial infections in cirrhosis: a position statement based on the EASL Special Conference. *J Hepatol* 2013;60:1310–1324.
- [7] Arvaniti V, D'Amico G, Fede G, Manousou P, Tsochatzis E, Pleguezuelo M, et al. Infections in patients with cirrhosis increase mortality four-fold and should be used in determining prognosis. *Gastroenterology* 2010;139:1246–1256.
- [8] Schrier RW, Arroyo V, Bernardi M, Epstein M, Henriksen JH, Rodes J. Peripheral arterial vasodilation hypothesis: a proposal for the initiation of renal sodium and water retention in cirrhosis. *Hepatology* 1988;8:1151–1157.
- [9] Wiese S, Hove JD, Bendtsen F, Moller S. Cirrhotic cardiomyopathy: pathogenesis and clinical relevance. *Nat Rev Gastroenterol Hepatol* 2014;11:177–186.
- [10] Arroyo V, Terra C, Gines P. Advances in the pathogenesis and treatment of type-1 and type-2 hepatorenal syndrome. *J Hepatol* 2007;46:935–946.
- [11] Iwakiri Y, Groszmann RJ. The hyperdynamic circulation of chronic liver diseases: from the patient to the molecule. *Hepatology* 2006;43:S121–S131.
- [12] Claria J, Stauber RE, Coenraad MJ, Moreau R, Jalan R, Pavesi M, et al. Systemic inflammation in decompensated cirrhosis: Characterization and role in acute-on-chronic liver failure. *Hepatology* 2016;64:1249–1264.
- [13] Trautwein C, Friedman SL, Schuppan D, Pinzani M. Hepatic fibrosis: Concept to treatment. *J Hepatol* 2015;62:S15–S24.
- [14] Alvarez MA, Cirera I, Sola R, Bargallo A, Morillas RM, Planas R. Long-term clinical course of decompensated alcoholic cirrhosis: a prospective study of 165 patients. *J Clin Gastroenterol* 2011;45:906–911.
- [15] Powell Jr WJ, Klatskin G. Duration of survival in patients with Laennec's cirrhosis. Influence of alcohol withdrawal, and possible effects of recent changes in general management of the disease. *Am J Med* 1968;44:406–420.
- [16] Shim JH, Lee HC, Kim KM, Lim YS, Chung YH, Lee YS, et al. Efficacy of entecavir in treatment-naive patients with hepatitis B virus-related decompensated cirrhosis. *J Hepatol* 2010;52:176–182.
- [17] Cheung MC, Walker AJ, Hudson BE, Verma S, McLauchlan J, Mutimer DJ, et al. Outcomes after successful direct-acting antiviral therapy for patients with chronic hepatitis C and decompensated cirrhosis. *J Hepatol* 2016;65:741–747.
- [18] Lens S, Alvarado E, Mariño Z, et al. Effects of all-oral antiviral therapy on HVPG and systemic hemodynamics in patients with hepatitis C virus-associated cirrhosis. *Gastroenterology* 2017;153:1273–1283.
- [19] Kang SH, Lee YB, Lee JH, Nam JY, Chang Y, Cho H, et al. Rifaximin treatment is associated with reduced risk of cirrhotic complications and prolonged overall survival in patients experiencing hepatic encephalopathy. *Aliment Pharmacol Ther* 2017;46:845–855.
- [20] Ginès P, Schrier RW. Renal failure in cirrhosis. *N Engl J Med* 2009;361:1279–1290.
- [21] Moreau R, Elkrief L, Bureau C, Pararnau JM, Thavenot T, Saliba F, et al. A randomized trial of 6-month norfloxacin therapy in patients with Child-Pugh class C cirrhosis. *J Hepatol* 2017;66:S1.
- [22] Caraceni P, Riggio O, Angeli P, Alessandria C, Neri S, Foschi FG, et al. Long-term albumin administration in decompensated cirrhosis: an open label randomized trial. *Lancet* 2018. [In press].
- [23] Sola E, Sola C, Simon-Talero M, Martin-Llahi M, Castellote J, Garcia-Martinez R, et al. Midodrine and albumin for prevention of complications of cirrhosis in patients in the waiting list for liver transplantation. A randomized, multicenter, double-blind, placebo-controlled trial. *J Hepatol* 2017;66:S11.
- [24] Abraldes JG, Albillos A, Banares R, Turnes J, Gonzalez R, Garcia-Pagan JC, et al. Simvastatin lowers portal pressure in patients with cirrhosis and portal hypertension: a randomized controlled trial. *Gastroenterology* 2009;136:1651–1658.
- [25] Abraldes JG, Villanueva C, Aracil C, Turnes J, Hernandez-Guerra M, Genesca J, et al. Addition of simvastatin to standard therapy for the prevention of variceal rebleeding does not reduce rebleeding but increases survival in patients with cirrhosis. *Gastroenterology* 2016;150:1160–1170.
- [26] Villa E, Cammà C, Marietta M, Luongo M, Critelli R, Colopi S, et al. Enoxaparin prevents portal vein thrombosis and liver decompensation in patients with advanced cirrhosis. *Gastroenterology* 2012;143:1253–12609.
- [27] Lebec D, Thabut D, Oberti F, Perarnau JM, Condat B, Barraud H, et al. Pentoxifylline does not decrease short-term mortality but does reduce complications in patients with advanced cirrhosis. *Gastroenterology* 2010;138:1755–1762.
- [28] Abraldes JG, Tarantino I, Turnes J, Garcia-Pagan JC, Rodes J, Bosch J. Hemodynamic response to pharmacological treatment of portal hypertension and long-term prognosis of cirrhosis. *Hepatology* 2003;37:902–908.
- [29] Gines P, Quintero E, Arroyo V, Teres J, Bruguera M, Rimola A, et al. Compensated cirrhosis: natural history and prognostic factors. *Hepatology* 1987;7:122–128.
- [30] Ripoll C, Groszmann R, Garcia-Tsao G, Grace N, Burroughs A, Planas R, et al. Hepatic venous pressure gradient predicts clinical decompensation in patients with compensated cirrhosis. *Gastroenterology* 2007;133:481–488.
- [31] Moore KP, Wong F, Gines P, Bernardi M, Ochs A, Salerno F, et al. The management of ascites in cirrhosis: report on the consensus conference of the International Ascites Club. *Hepatology* 2003;38:258–266.
- [32] Arroyo V, Gines P, Gerbes AL, Dudley FJ, Gentilini P, Laffi G, et al. Definition and diagnostic criteria of refractory ascites and hepatorenal syndrome in cirrhosis. *International Ascites Club. Hepatology* 1996;23:164–176.
- [33] Rimola A, Garcia-Tsao G, Navasa M, Piddock LJ, Planas R, Bernard B, et al. Diagnosis, treatment and prophylaxis of spontaneous bacterial peritonitis: a consensus document. *International Ascites Club. J Hepatol* 2000;32:142–153.
- [34] Bruns T, Lutz P, Stallmach A, Nischalke HD. Low ascitic fluid protein does not indicate an increased risk for spontaneous bacterial peritonitis in current cohorts. *J Hepatol* 2015;63:527–528.
- [35] Runyon BA, Canawati HN, Akriviadis EA. Optimization of ascitic fluid culture technique. *Gastroenterology* 1988;95:1351–1355.
- [36] Runyon BA, Montano AA, Akriviadis EA, Antillon MR, Irving MA, McHutchison JG. The serum-ascites albumin gradient is superior to the exudate-transudate concept in the differential diagnosis of ascites. *Ann Intern Med* 1992;117:215–220.
- [37] Gerbes AL, Jünger D, Xie YN, Permanetter W, Paumgartner G. Ascitic fluid analysis for the differentiation of malignancy-related and nonmalignant ascites. Proposal of a diagnostic sequence. *Cancer* 1991;68:1808–1814.
- [38] Llach J, Gines P, Arroyo V, Rimola A, Tito L, Badalamenti S, et al. Prognostic value of arterial pressure, endogenous vasoactive systems, and renal function in cirrhotic patients admitted to the hospital for the treatment of ascites. *Gastroenterology* 1988;94:482–487.

- [39] Caregario L, Menon F, Angeli P, Amodio P, Merkel C, Bortoluzzi A, et al. Limitations of serum creatinine level and creatinine clearance as filtration markers in cirrhosis. *Arch Intern Med* 1994;154:201–205.
- [40] Bernardi M, Gitto S, Biselli M. The MELD score in patients awaiting liver transplant: strengths and weaknesses. *J Hepatol* 2011;54:1297–1306.
- [41] Biselli M, Dall'Agata M, Gramenzi A, Gitto S, Liberati C, Brodosi L, et al. A new prognostic model to predict dropout from the waiting list in cirrhotic candidates for liver transplantation with MELD score <18. *Liver Int* 2015;35:184–191.
- [42] Bernardi M, Santini C, Trevisani F, Baraldini M, Ligabue A, Gasbarrini G. Renal function impairment induced by change in posture in patients with cirrhosis and ascites. *Gut* 1985;26:629–635.
- [43] Ring-Larsen H, Henriksen JH, Wilken C, Clausen J, Pals H, Christensen NJ. Diuretic treatment in decompensated cirrhosis and congestive heart failure: effect of posture. *Br Med J* 1986;292:1351–1353.
- [44] Bernardi M, Laffi G, Salvagnini M, Azzena G, Bonato S, Marra F, et al. Efficacy and safety of the stepped care medical treatment of ascites in liver cirrhosis: a randomized controlled clinical trial comparing two diets with different sodium content. *Liver* 1993;13:156–162.
- [45] Gauthier A, Levy VG, Quinton A, Michel H, Rueff B, Descos L, et al. Salt or no salt in the treatment of cirrhotic ascites: a randomised study. *Gut* 1986;27:705–709.
- [46] Reynolds TB, Lieberman FL, Goodman AR. Advantages of treatment of ascites without sodium restriction and without complete removal of excess fluid. *Gut* 1978;19:549–553.
- [47] Morando F, Rosi S, Gola E, Nardi M, Piano S, Fasolato S, et al. Adherence to a moderate sodium restriction diet in outpatients with cirrhosis and ascites: a real-life cross-sectional study. *Liver Int* 2015;35:1508–1515.
- [48] Pockros PJ, Reynolds TB. Rapid diuresis in patients with ascites from chronic liver disease: the importance of peripheral edema. *Gastroenterology* 1986;90:1827–1833.
- [49] Bernardi M, Trevisani F, Gasbarrini A, Gasbarrini G. Hepatorenal disorders: role of the renin-angiotensin-aldosterone system. *Semin Liver Dis* 1994;14:23–34.
- [50] Bernardi M, Servadei D, Trevisani F, Rusticali AG, Gasbarrini G. Importance of plasma aldosterone concentration on the natriuretic effect of spironolactone in patients with liver cirrhosis and ascites. *Digestion* 1985;31:189–193.
- [51] Angeli P, Dalla Pria M, De Bei E, Albino G, Caregario L, Merkel C, et al. Randomized clinical study of the efficacy of amiloride and potassium canrenoate in nonazotemic cirrhotic patients with ascites. *Hepatology* 1994;19:72–79.
- [52] Angeli P, Gatta A, Caregario L, Menon F, Sacerdoti D, Merkel C, et al. Tubular site of renal sodium retention in ascitic liver cirrhosis evaluated by lithium clearance. *Eur J Clin Invest* 1990;20:111–117.
- [53] Gatta A, Angeli P, Caregario L, Menon F, Sacerdoti D, Merkel C. A pathophysiological interpretation of unresponsiveness to spironolactone in a stepped-care approach to the diuretic treatment of ascites in nonazotemic cirrhotic patients. *Hepatology* 1991;14:231–236.
- [54] Perez-Ayuso RM, Arroyo V, Planas R, Gaya J, Bory F, Rimola A, et al. Randomized comparative study of efficacy of furosemide vs. spironolactone in nonazotemic cirrhosis with ascites. Relationship between the diuretic response and the activity of the renin-aldosterone system. *Gastroenterology* 1983;84:961–968.
- [55] Angeli P, Fasolato S, Mazza E, Okolicsanyi L, Maresio G, Velo E, et al. Combined vs. sequential diuretic treatment of ascites in non-azotemic patients with cirrhosis: results of an open randomised clinical trial. *Gut* 2010;59:98–104.
- [56] Santos J, Planas R, Pardo A, Durandez R, Cabre E, Morillas RM, et al. Spironolactone alone or in combination with furosemide in the treatment of moderate ascites in nonazotemic cirrhosis. A randomized comparative study of efficacy and safety. *J Hepatol* 2003;39:187–192.
- [57] Gerbes AL, Bertheau-Reitha U, Falkner C, Jüngst D, Paumgartner G. Advantages of the new loop diuretic torasemide over furosemide in patients with cirrhosis and ascites. A randomized, double blind cross-over trial. *J Hepatol* 1993;17:353–358.
- [58] Angeli P, Albino G, Carraro P, Dalla Pria M, Merkel C, Caregario L, et al. Cirrhosis and muscle cramps: evidence of a causal relationship. *Hepatology* 1996;23:264–273.
- [59] Elfert AA, Abo Ali L, Soliman S, Zakaria S, Shehab El-Din I, Elkhawany W, et al. Randomized placebo-controlled study of baclofen in the treatment of muscle cramps in patients with liver cirrhosis. *Eur J Gastroenterol Hepatol* 2016;28:1280–1284.
- [60] Lee FY, Lee SD, Tsai YT, et al. A randomized controlled trial of quinidine in the treatment of cirrhotic patients with muscle cramps. *J Hepatol* 1991;12:236–240.
- [61] Lin CH, Shih FY, Ma MH, Chiang WC, Yang CW, Ko PC. Should bleeding tendency deter abdominal paracentesis? *Dig Liver Dis* 2005;37:946–951.
- [62] Pache I, Bilodeau M. Severe haemorrhage following abdominal paracentesis for ascites in patients with liver disease. *Aliment Pharmacol Ther* 2005;21:525–529.
- [63] Gines P, Tito L, Arroyo V, Planas R, Panes J, Viver J, et al. Randomized comparative study of therapeutic paracentesis with and without intravenous albumin in cirrhosis. *Gastroenterology* 1988;94:1493–1502.
- [64] Gines A, Fernandez-Esparrach G, Monescillo A, Vila C, Domenech E, Abecasis R, et al. Randomized trial comparing albumin, dextran 70, and polygeline in cirrhotic patients with ascites treated by paracentesis. *Gastroenterology* 1996;111:1002–1010.
- [65] Sola-Vera J, Minana J, Ricart E, Planella M, Gonzalez B, Torras X, et al. Randomized trial comparing albumin and saline in the prevention of paracentesis-induced circulatory dysfunction in cirrhotic patients with ascites. *Hepatology* 2003;37:1147–1153.
- [66] Bernardi M, Caraceni P, Navickis RJ, Wilkes MM. Albumin infusion in patients undergoing large-volume paracentesis: a meta-analysis of randomized trials. *Hepatology* 2012;55:1172–1181.
- [67] Moreau R, Valla DC, Durand-Zaleski I, Bronowicki JP, Durand F, Chaput JC, et al. Comparison of outcome in patients with cirrhosis and ascites following treatment with albumin or a synthetic colloid: a randomised controlled pilot trial. *Liver Int* 2006;26:46–54.
- [68] Gines P, Arroyo V, Quintero E, Planas R, Bory F, Cabrera J, et al. Comparison of paracentesis and diuretics in the treatment of cirrhotics with tense ascites. Results of a randomized study. *Gastroenterology* 1987;93:234–241.
- [69] Salerno F, Badalamenti S, Incerti P, Tempini S, Restelli B, Bruno S, et al. Repeated paracentesis and i.v. albumin infusion to treat 'tense' ascites in cirrhotic patients. A safe alternative therapy. *J Hepatol* 1987;5:102–108.
- [70] Fernandez-Esparrach G, Guevara M, Sort P, Pardo A, Jimenez W, Gines P, et al. Diuretic requirements after therapeutic paracentesis in non-azotemic patients with cirrhosis. A randomized double-blind trial of spironolactone vs. placebo. *J Hepatol* 1997;26:614–620.
- [71] Elia C, Graupera I, Barreto R, Solà E, Moreira R, Huelin P, et al. Severe acute kidney injury associated with non-steroidal antiinflammatory drugs in cirrhosis: a case-control study. *J Hepatol* 2015;63:593–600.
- [72] Claria J, Kent JD, Lopez-Parra M, Escolar G, Ruiz-Del-Arbol L, Gines P, et al. Effects of celecoxib and naproxen on renal function in nonazotemic patients with cirrhosis and ascites. *Hepatology* 2005;41:579–587.
- [73] Llach J, Gines P, Arroyo V, Salmeron JM, Gines A, Jimenez W, et al. Effect of dipyridamole on kidney function in cirrhosis. *Hepatology* 1993;17:59–64.
- [74] Pariente EA, Bataille C, Bercoff E, Lebrec D. Acute effects of captopril on systemic and renal hemodynamics and on renal function in cirrhotic patients with ascites. *Gastroenterology* 1985;88:1255–1259.
- [75] Albillos A, Lledo JL, Rossi I, Perez-Paramo M, Tabuenca MJ, Banares R, et al. Continuous prazosin administration in cirrhotic patients: effects on portal hemodynamics and on liver and renal function. *Gastroenterology* 1995;109:1257–1265.
- [76] Cabrera J, Arroyo V, Ballesta AM, Rimola A, Gual J, Elena M, et al. Aminoglycoside nephrotoxicity in cirrhosis. Value of urinary beta 2-microglobulin to discriminate functional renal failure from acute tubular damage. *Gastroenterology* 1982;82:97–105.
- [77] Guevara M, Fernández-Esparrach G, Alessandria C, Torre A, Terra C, Montañà X, et al. Effects of contrast media on renal function in patients with cirrhosis: a prospective study. *Hepatology* 2004;40:646–651.
- [78] Solomon R, Werner C, Mann D, D'Elia J, Silva P. Comparison of saline, mannitol, and furosemide to prevent acute decreases in renal function induced by radiocontrast agents. *N Engl J Med* 1994;331:1416–1420.
- [79] Salerno F, Borroni G, Moser P, Badalamenti S, Cassara L, Maggi A, et al. Survival and prognostic factors of cirrhotic patients with ascites: a study of 134 outpatients. *Am J Gastroenterol* 1993;88:514–519.
- [80] Guardiola J, Baliellas C, Xiol X, Fernandez Esparrach G, Gines P, Ventura P, et al. External validation of a prognostic model for predicting survival of cirrhotic patients with refractory ascites. *Am J Gastroenterol* 2002;97:2374–2378.
- [81] Huonker M, Schumacher YO, Ochs A, Sorichter S, Keul J, Rossle M. Cardiac function and haemodynamics in alcoholic cirrhosis and effects of the transjugular intrahepatic portosystemic stent shunt. *Gut* 1999;44:743–748.

Clinical Practice Guidelines

- [82] Sanyal AJ, Freedman AM, Luketic VA, Purdum 3rd PP, Shiffman ML, DeMeo J, et al. The natural history of portal hypertension after transjugular intrahepatic portosystemic shunts. *Gastroenterology* 1997;112:889–898.
- [83] Wong F, Sniderman K, Liu P, Allidina Y, Sherman M, Blendis L. Transjugular intrahepatic portosystemic stent shunt: effects on hemodynamics and sodium homeostasis in cirrhosis and refractory ascites. *Ann Intern Med* 1995;122:816–822.
- [84] Wong F, Sniderman K, Liu P, Blendis L. The mechanism of the initial natriuresis after transjugular intrahepatic portosystemic shunt. *Gastroenterology* 1997;112:899–907.
- [85] Gerbes AL, Gülberg V, Waggershauser T, Holl J, Reiser M. Renal effects of transjugular intrahepatic portosystemic shunt in cirrhosis: comparison of patients with ascites, with refractory ascites, or without ascites. *Hepatology* 1998;28:683–688.
- [86] Ochs A, Rossle M, Haag K, Hauenstein KH, Deibert P, Siegerstetter V, et al. The transjugular intrahepatic portosystemic stent-shunt procedure for refractory ascites. *N Engl J Med* 1995;332:1192–1197.
- [87] Plauth M, Schutz T, Buckendahl DP, Kreyman G, Pirlich M, Grungreiff S, et al. Weight gain after transjugular intrahepatic portosystemic shunt is associated with improvement in body composition in malnourished patients with cirrhosis and hypermetabolism. *J Hepatol* 2004;40:228–233.
- [88] Gülberg V, Liss I, Bilzer M, Waggershauser T, Reiser M, Gerbes AL. Improved quality of life in patients with refractory or recidivant ascites after insertion of transjugular intrahepatic portosystemic shunts. *Digestion* 2002;66:127–130.
- [89] Casado M, Bosch J, Garcia-Pagan JC, Bru C, Banares R, Bandi JC, et al. Clinical events after transjugular intrahepatic portosystemic shunt: correlation with hemodynamic findings. *Gastroenterology* 1998;114:1296–1303.
- [90] Riggio O, Angeloni S, Salvatori FM, De Santis A, Cerini F, Farcomeni A, et al. Incidence, natural history, and risk factors of hepatic encephalopathy after transjugular intrahepatic portosystemic shunt with polytetrafluoroethylene-covered stent grafts. *Am J Gastroenterol* 2008;103:2738–2746.
- [91] Sauerbruch T, Mengel M, Dollinger M, Zipprich A, Rossle M, Panther E, et al. Prevention of rebleeding from esophageal varices in patients with cirrhosis receiving small-diameter stents vs. hemodynamically controlled medical therapy. *Gastroenterology* 2015;149:660–668.
- [92] Wang Q, Lv Y, Bai M, Wang Z, Liu H, He C, et al. Eight millimetre covered TIPS does not compromise shunt function but reduces hepatic encephalopathy in preventing variceal rebleeding. *J Hepatol* 2017;67:508–516.
- [93] Pieper CC, Jansen C, Meyer C, Nadal J, Lehmann J, Schild HH, et al. Prospective evaluation of passive expansion of partially dilated transjugular intrahepatic portosystemic shunt stent grafts—a three-dimensional sonography study. *J Vasc Interv Radiol* 2017;28:117–125.
- [94] Bureau C, Garcia-Pagan JC, Otal P, Pomier-Layrargues G, Chabbert V, Cortez C, et al. Improved clinical outcome using polytetrafluoroethylene-coated stents for TIPS: results of a randomized study. *Gastroenterology* 2004;126:469–475.
- [95] Gines P, Uriz J, Calahorra B, Garcia-Tsao G, Kamath PS, Del Arbol LR, et al. Transjugular intrahepatic portosystemic shunting vs. paracentesis plus albumin for refractory ascites in cirrhosis. *Gastroenterology* 2002;123:1839–1847.
- [96] Lebrec D, Guilly N, Hadengue A, Vilgrain V, Moreau R, Poynard T, et al. Transjugular intrahepatic portosystemic shunts: comparison with paracentesis in patients with cirrhosis and refractory ascites: a randomized trial. *French Group of Clinicians and a Group of Biologists. J Hepatol* 1996;25:135–144.
- [97] Narahara Y, Kanazawa H, Fukuda T, Matsushita Y, Harimoto H, Kidokoro H, et al. Transjugular intrahepatic portosystemic shunt vs. paracentesis plus albumin in patients with refractory ascites who have good hepatic and renal function: a prospective randomized trial. *J Gastroenterol* 2011;46:78–85.
- [98] Rossle M, Ochs A, Gulberg V, Siegerstetter V, Holl J, Deibert P, et al. A comparison of paracentesis and transjugular intrahepatic portosystemic shunting in patients with ascites. *N Engl J Med* 2000;342:1701–1707.
- [99] Salerno F, Merli M, Riggio O, Cazzaniga M, Valeriano V, Pozzi M, et al. Randomized controlled study of TIPS vs. paracentesis plus albumin in cirrhosis with severe ascites. *Hepatology* 2004;40:629–635.
- [100] Sanyal AJ, Genning C, Reddy KR, Wong F, Kowdley KV, Benner K, et al. The North American study for the treatment of refractory ascites. *Gastroenterology* 2003;124:634–641.
- [101] Albillos A, Banares R, Gonzalez M, Catalina MV, Molinero LM. A meta-analysis of transjugular intrahepatic portosystemic shunt vs. paracentesis for refractory ascites. *J Hepatol* 2005;43:990–996.
- [102] Bai M, Qi XS, Yang ZP, Yang M, Fan DM, Han GH. TIPS improves liver transplantation-free survival in cirrhotic patients with refractory ascites: an updated meta-analysis. *World J Gastroenterol* 2014;20:2704–2714.
- [103] Chen RP, Zhu Ge XJ, Huang ZM, Ye XH, Hu CY, Lu GR, et al. Prophylactic use of transjugular intrahepatic portosystemic shunt aids in the treatment of refractory ascites: meta-regression and trial sequential meta-analysis. *J Clin Gastroenterol* 2014;48:290–299.
- [104] D'Amico G, Luca A, Morabito A, Miraglia R, D'Amico M. Uncovered transjugular intrahepatic portosystemic shunt for refractory ascites: a meta-analysis. *Gastroenterology* 2005;129:1282–1293.
- [105] Deltenre P, Mathurin P, Dharancy S, Moreau R, Bulois P, Henrion J, et al. Transjugular intrahepatic portosystemic shunt in refractory ascites: a meta-analysis. *Liver Int* 2005;25:349–356.
- [106] Saab S, Nieto JM, Lewis SK, Runyon BA. TIPS vs. paracentesis for cirrhotic patients with refractory ascites. *Cochrane Database Syst Rev* 2006;CD004889.
- [107] Salerno F, Camma C, Enea M, Rossle M, Wong F. Transjugular intrahepatic portosystemic shunt for refractory ascites: a meta-analysis of individual patient data. *Gastroenterology* 2007;133:825–834.
- [108] Maleux G, Perez-Gutierrez NA, Evrard S, Mroue A, Le Moine O, Laleman W, et al. Covered stents are better than uncovered stents for transjugular intrahepatic portosystemic shunts in cirrhotic patients with refractory ascites: a retrospective cohort study. *Acta Gastroenterol Belg* 2010;73:336–341.
- [109] Tan HK, James PD, Sniderman KW, Wong F. Long-term clinical outcome of patients with cirrhosis and refractory ascites treated with transjugular intrahepatic portosystemic shunt insertion. *J Gastroenterol Hepatol* 2015;30:389–395.
- [110] Gaba RC, Parvinian A, Casadaban LC, Couture PM, Zivin SP, Lakhoo J, et al. Survival benefit of TIPS vs. serial paracentesis in patients with refractory ascites: a single institution case-control propensity score analysis. *Clin Radiol* 2015;70:e51–e57.
- [111] Bureau C, Thabut D, Oberti F, Dharancy S, Carbonell N, Bouvier A, et al. Transjugular intrahepatic portosystemic shunts with covered stents increase transplant-free survival of patients with cirrhosis and recurrent ascites. *Gastroenterology* 2017;152:157–163.
- [112] Malinchoc M, Kamath PS, Gordon FD, Peine CJ, Rank J, ter Borg PC. A model to predict poor survival in patients undergoing transjugular intrahepatic portosystemic shunts. *Hepatology* 2000;31:864–871.
- [113] Bureau C, Metivier S, D'Amico M, Peron JM, Otal P, Pagan JC, et al. Serum bilirubin and platelet count: a simple predictive model for survival in patients with refractory ascites treated by TIPS. *J Hepatol* 2011;54:901–907.
- [114] Sarwar A, Zhou L, Novack V, Tapper EB, Curry M, Malik R, et al. Hospital volume and mortality after trans-jugular intrahepatic portosystemic shunt creation in the United States. *Hepatology* 2017.
- [115] Angeli P, Volpin R, Piovano D, Bortoluzzi A, Craighero R, Bottaro S, et al. Acute effects of the oral administration of midodrine, an alpha-adrenergic agonist, on renal hemodynamics and renal function in cirrhotic patients with ascites. *Hepatology* 1998;28:937–943.
- [116] Singh V, Dhungana SP, Singh B, Vijayverghia R, Nain CK, Sharma N, et al. Midodrine in patients with cirrhosis and refractory or recurrent ascites: a randomized pilot study. *J Hepatol* 2012;56:348–354.
- [117] Gadano A, Moreau R, Vachieri F, Soupison T, Yang S, Cailmail S, et al. Natriuretic response to the combination of atrial natriuretic peptide and terlipressin in patients with cirrhosis and refractory ascites. *J Hepatol* 1997;26:1229–1234.
- [118] Krag A, Moller S, Henriksen JH, Holstein-Rathlou NH, Larsen FS, Bendtsen F. Terlipressin improves renal function in patients with cirrhosis and ascites without hepatorenal syndrome. *Hepatology* 2007;46:1863–1871.
- [119] Lenaerts A, Codden T, Meunier JC, Henry JP, Ligny G. Effects of clonidine on diuretic response in ascitic patients with cirrhosis and activation of sympathetic nervous system. *Hepatology* 2006;44:844–849.
- [120] Singh V, Singh A, Singh B, Vijayvergiya R, Sharma N, Ghai A, et al. Midodrine and clonidine in patients with cirrhosis and refractory or recurrent ascites: a randomized pilot study. *Am J Gastroenterol* 2013;108:560–567.
- [121] Rai N, Singh B, Singh A, Vijayvergiya R, Sharma N, Bhalla A, et al. Midodrine and tolvaptan in patients with cirrhosis and refractory or recurrent ascites: a randomised pilot study. *Liver Int* 2017;37:406–414.

- [122] Tandon P, Tsuyuki RT, Mitchell L, Hoskinson M, Ma MM, Wong WW, et al. The effect of 1 month of therapy with midodrine, octreotide-LAR and albumin in refractory ascites: a pilot study. *Liver Int* 2009;29:169–174.
- [123] Hanafy AS, Hassaneen AM. Rifaximin and midodrine improve clinical outcome in refractory ascites including renal function, weight loss, and short-term survival. *Eur J Gastroenterol Hepatol* 2016;28:1455–1461.
- [124] Bellot P, Welker MW, Soriano G, von Schaewen M, Appenrodt B, Wiest R, et al. Automated low flow pump system for the treatment of refractory ascites: a multi-center safety and efficacy study. *J Hepatol* 2013;58:922–927.
- [125] Stirnimann G, Berg T, Spahr L, Zeuzem S, McPherson S, Lammert F, et al. Treatment of refractory ascites with an automated low-flow ascites pump in patients with cirrhosis. *Aliment Pharmacol Ther* 2017;46:981–991.
- [126] Bureau C, Adebayo D, Chalret de Rieu M, Elkrief L, Valla D, et al. Alfapump® system vs. large volume paracentesis for refractory ascites: A multicenter randomized controlled study. *J Hepatol* 2017;67:940–949.
- [127] Sola E, Sanchez-Cabus S, Rodriguez E, Elia C, Cela R, Moreira R, et al. Effects of alfapump system on kidney and circulatory function in patients with cirrhosis and refractory ascites. *Liver Transpl* 2017;23:583–593.
- [128] Badillo R, Rockey DC. Hepatic hydrothorax. Clinical features, management, and outcomes in 77 patients and review of the literature. *Medicine* 2014;93:135–142.
- [129] Garbuzenko DV, Arefyev NO. Hepatic hydrothorax: An update and review of the literature. *World J Hepatol* 2017;1197–1204.
- [130] Zenda T, Miyamoto S, Murata S, Mabuchi H. Detection of diaphragmatic defect as the cause of severe hepatic hydrothorax with magnetic resonance imaging. *Am J Gastroenterol* 1998;9:2288–2289.
- [131] Hewett LJ, Bradshaw ML, Gordon LL, Rockey DC. Diagnosis of isolated hepatic hydrothorax using peritoneal scintigraphy. *Hepatology* 2016;64:1364–1366.
- [132] Alonso JC. Pleural effusion in liver disease. *Semin Respir Crit Care Med* 2010;31:698–705.
- [133] Orman ES, Lok AS. Outcomes of patients with chest tube insertion for hepatic hydrothorax. *Hepatol Int* 2009;3:582–586.
- [134] Xiol X, Tremosa G, Castellote J, Gornals J, Lama C, et al. Liver transplantation in patients with hepatic hydrothorax. *Transpl Int* 2005;18:672–675.
- [135] Sersté T, Moreno C, Francoz C, Razek WA, Paugham C, et al. The impact of preoperative hepatic hydrothorax on the outcome of adult liver transplantation. *Eur J Gastroenterol Hepatol* 2010;22:207–212.
- [136] Gordon FD, Anastopoulos HT, Crenshaw W, Gilchrist B, McEniff N, Falchuk KR, et al. The successful treatment of symptomatic, refractory hepatic hydrothorax with transjugular intrahepatic portosystemic shunt. *Hepatology* 1997;25:1366–1369.
- [137] Siegerstetter V, Deibert P, Ochs A, Olschewski M, Blum HE, Rossle M. Treatment of refractory hepatic hydrothorax with transjugular intrahepatic portosystemic shunt: long-term results in 40 patients. *Eur J Gastroenterol Hepatol* 2001;13:529–534.
- [138] Ditah IC, Al Bawardy BF, Saberi B, Ditah C, Kamath PS. Transjugular intrahepatic portosystemic stent shunt for medically refractory hepatic hydrothorax: A systematic review and cumulative meta-analysis. *World J Hepatol* 2015;7:1797–1806.
- [139] Hou F, Qi X, Guo X. Effectiveness and safety of pleurodesis for hepatic hydrothorax: A systematic review and meta-analysis. *Dig Dis Sci* 2016;61:3321–3334.
- [140] Huang PM, Kuo SW, Chen JS, Lee JM. Thoracoscopic mesh repair of diaphragmatic defects in hepatic hydrothorax: A 10-year experience. *Ann Thorac Surg* 2016;101:1921–1927.
- [141] Angeli P, Wong F, Watson H, Gines P, and the participants to CAPPS. Hyponatremia in cirrhosis: results of a survey. *Hepatology* 2006;44:1535–1542.
- [142] Gines P, Berl T, Bernardi M, Bichet DG, Hamon G, Jimenez W, et al. Hyponatremia in cirrhosis: from pathogenesis to treatment. *Hepatology* 1998;28:851–864.
- [143] Spasovski G, Vanholder R, Allohio B, Annane D, Ball S, Bichet D, et al. Clinical practice guideline on diagnosis and treatment of hyponatremia. *Intensive Care Med* 2014;40:320–331.
- [144] Biggins SW, Rodriguez HJ, Bacchetti P, Bass NM, Roberts JP, Terrault NA. Serum sodium predicts mortality in patients listed for liver transplantation. *Hepatology* 2005;41:32–39.
- [145] Porcel A, Diaz F, Rendon P, Macias M, Martin-Herrera L, Giron-Gonzalez JA. Dilutional hyponatremia in patients with cirrhosis and ascites. *Arch Intern Med* 2002;162:323–328;
- Cordoba J, Garcia-Martinez R, Simon-Talero M. Hyponatremic and hepatic encephalopathies: similarities, differences and coexistence. *Metab Brain Dis* 2010;25:73–80.
- [146] Cordoba J, Garcia-Martinez R, Simon-Talero M. Hyponatremic and hepatic encephalopathies: similarities, differences and coexistence. *Metab Brain Dis* 2010;25:73–80.
- [147] Amodio P, Del Piccolo F, Pettenu E, Mapelli D, Angeli P, Iemmolo R, et al. Prevalence and prognostic value of quantified electroencephalogram (EEG) alterations in cirrhotic patients. *J Hepatol* 2001;35:37–45.
- [148] Londono MC, Guevara M, Rimola A, Navasa M, Taura P, Mas A, et al. Hyponatremia impairs early posttransplantation outcome in patients with cirrhosis undergoing liver transplantation. *Gastroenterology* 2006;130:1135–1143.
- [149] Biggins SW, Kim WR, Terrault NA, Saab S, Balan V, Schiano T, et al. Evidence-based incorporation of serum sodium concentration into MELD. *Gastroenterology* 2006;130:1652–1660.
- [150] Kim WR, Biggins SW, Kremers WK, Wiesner RH, Kamath PS, Benson JT, et al. Hyponatremia and mortality among patients on the liver-transplant waiting list. *N Engl J Med* 2008;359:1018–1026.
- [151] McCormick PA, Mistry P, Kaye G, Burroughs AK, McIntyre N. Intravenous albumin infusion is an effective therapy for hyponatremia in cirrhotic patients with ascites. *Gut* 1990;31:204–207.
- [152] Quittnat F, Gross P. Vaptans and the treatment of water-retaining disorders. *Semin Nephrol* 2006;26:234–243.
- [153] Cardenas A, Gines P, Marotta P, Czerwiec F, Oyuang J, Guevara M, et al. Tolvaptan, an oral vasopressin antagonist, in the treatment of hyponatremia in cirrhosis. *J Hepatol* 2012;56:571–578.
- [154] Gerbes AL, Gulberg V, Gines P, Decaux G, Gross P, Gandjini H, et al. Therapy of hyponatremia in cirrhosis with a vasopressin receptor antagonist: a randomized double-blind multicenter trial. *Gastroenterology* 2003;124:933–939.
- [155] Gines P, Wong F, Watson H, Milutinovic S, del Arbol LR, Olteanu D. Effects of satavaptan, a selective vasopressin V(2) receptor antagonist, on ascites and serum sodium in cirrhosis with hyponatremia: a randomized trial. *Hepatology* 2008;48:204–213.
- [156] O'Leary JG, Davis GL. Conivaptan increases serum sodium in hyponatremic patients with end-stage liver disease. *Liver Transpl* 2009;15:1325–1329.
- [157] Wong F, Gines P, Watson H, Horsmans Y, Angeli P, Gow P, et al. Effects of a selective vasopressin V2 receptor antagonist, satavaptan, on ascites recurrence after paracentesis in patients with cirrhosis. *J Hepatol* 2010;53:283–290.
- [158] Wong F, Watson H, Gerbes A, Vilstrup H, Badalamenti S, Bernardi M, et al. Satavaptan for the management of ascites in cirrhosis: efficacy and safety across the spectrum of ascites severity. *Gut* 2012;61:108–116.
- [159] Pose E, Sola E, Piano S, Gola E, Graupera I, Guevara M, et al. Limited efficacy of tolvaptan in patients with cirrhosis and severe hyponatremia: real-life experience. *Am J Med* 2017;130:372–375.
- [160] Torres VE, Chapman AB, Devuyt O, Gansevoort RT, Grantham JJ, Higashihara E, et al. Tolvaptan in patients with autosomal dominant polycystic kidney disease. *N Engl J Med* 2012;367:2407–2418.
- [161] Bosch J, Groszmann RJ, Shah VH. Evolution in the understanding of the pathophysiological basis of portal hypertension: How changes in paradigm are leading to successful new treatments. *J Hepatol* 2015;62:S121–S130.
- [162] D'Amico G, De Franchis R. Upper digestive bleeding in cirrhosis. Post-therapeutic outcome and prognostic indicators. *Hepatology* 2003;38:599–612.
- [163] Jepsen P, Ott P, Andersen PK, Sorensen HT, Vilstrup H. Clinical course of alcoholic liver cirrhosis: a Danish population-based cohort study. *Hepatology* 2010;51:1675–1682.
- [164] Kovalak M, Lake J, Mattek N, Eisen G, Lieberman D, Zaman A. Endoscopic screening for varices in cirrhotic patients: data from a national endoscopic database. *Gastrointest Endosc* 2007;65:82–88.
- [165] Merli M, Nicolini G, Angeloni S, Rinaldi V, De Santis A, Merkel C, et al. Incidence and natural history of small esophageal varices in cirrhotic patients. *J Hepatol* 2003;38:266–272.
- [166] Groszmann RJ, Garcia-Tsao G, Bosch J, Grace ND, Burroughs AK, Planas R, et al. Beta-blockers to prevent gastroesophageal varices in patients with cirrhosis. *N Engl J Med* 2005;353:2254–2261.
- [167] Abraldes JG, Bureau C, Stefanescu H, Augustin S, Ney M, Blasco H, et al. Non invasive tools and risk of clinically significant portal hypertension and varices in compensated cirrhosis: the "ANTICIPATE" study. *Hepatology* 2016;64:2173–2184.

Clinical Practice Guidelines

- [168] De Franchis RBavero VI faculty. Expanding consensus in portal hypertension: report of the BAVENO VI Consensus Workshop: Stratifying risk and individualizing care for portal hypertension. *J Hepatol* 2015;63:743–752.
- [169] Garcia-Tsao G, Sanyal AJ, Grace ND, Carey W Practice Guidelines Committee of the American Association for the Study of Liver Diseases Practice Parameters Committee of the American College of Gastroenterology. Prevention and management of gastroesophageal varices and variceal hemorrhage in cirrhosis. *Hepatology* 2007;46:922–938.
- [170] North Italian Endoscopic Club for the Study and Treatment of Esophageal Varices. Prediction of the first variceal hemorrhage in patients with cirrhosis of the liver and esophageal varices. A prospective multicenter study. *N Engl J Med* 1988;319:983–989.
- [171] D'Amico G, Pasta L, Morabito A, et al. Competing risks and prognostic stages of cirrhosis: a 25-year inception cohort study of 494 patients. *Aliment Pharmacol Ther* 2014;39:1180–1193.
- [172] Augustin S, Muntaner L, Altamirano JT, Gonzalez A, Saperas E, Dot J, et al. Predicting early mortality after acute variceal hemorrhage based on classification and regression tree analysis. *Clin Gastroenterol Hepatol* 2009;7:1347–1354.
- [173] Bosch J, Garcia-Pagan JC. Prevention of variceal rebleeding. *Lancet* 2003;361:952–954.
- [174] Garcia-Pagan JC, De Gottardi A, Bosch J. Review article: the modern management of portal hypertension—primary and secondary prophylaxis of variceal bleeding in cirrhotic patients. *Aliment Pharmacol Ther* 2008;28:178–186.
- [175] Hernandez-Gea V, Aracil C, Colomo A, Garupera I, Poca M, Torras X, et al. Development of ascites in compensated cirrhosis with severe portal hypertension treated with beta-blockers. *Am J Gastroenterol* 2012;107:418–427.
- [176] Puente A, Hernandez-Gea V, Graupera I, Roque M, Colomo A, Poca M, et al. Drugs plus ligation to prevent rebleeding in cirrhosis: an updated systematic review. *Liver Int* 2014;34:823–833.
- [177] Villanueva C, Graupera I, Aracil C, Alvarado E, Minana J, Puente A, et al. A randomized trial to assess whether portal pressure guided therapy to prevent variceal rebleeding improves survival in cirrhosis. *Hepatology* 2017;65:1693–1707.
- [178] D'Amico G, Pagliaro L, Bosch J. Pharmacological treatment of portal hypertension: an evidence-based approach. *Semin Liver Dis* 1999;19:475–505.
- [179] Albillos A, Banares R, Gonzalez M, Ripoll C, Gonzalez R, Catalina MV, et al. Value of the hepatic venous pressure gradient to monitor drug therapy for portal hypertension: a meta-analysis. *Am J Gastroenterol* 2007;102:1116–1126.
- [180] Li T, Ke W, Sun P, Chen X, Belgaumkar A, Huang Y, et al. Carvedilol for portal hypertension in cirrhosis: systematic review with meta-analysis. *BMJ Open* 2016;6:e010902.
- [181] Serste T, Melot C, Francoz C, Durand F, Rautou PE, Valla D, et al. Deleterious effects of beta-blockers on survival in patients with cirrhosis and refractory ascites. *Hepatology* 2010;52:1017–1022.
- [182] Wiest R, Albillos A, Gluud LL. The window hypothesis: haemodynamic and non-haemodynamic effects of beta-blockers improve survival of patients with cirrhosis during a window in the disease. *Gut* 2012;61:967–969.
- [183] Bossen L, Krag A, Vilstrup H, Watson H, Jepsen P. Non-selective beta-blockers do not affect mortality in cirrhosis patients with ascites: Post hoc analysis of three RCTs with 1198 patients. *Hepatology* 2016;63:1968–1976.
- [184] Bang UC, Benfield T, Hyldstrup L, Jensen JE, Bendtsen F. Effect of propranolol on survival in patients with decompensated cirrhosis: a nationwide study based Danish patient registers. *Liver Int* 2016;36:1304–1312.
- [185] Reiberger T, Mandorfer M. Beta adrenergic blockade and decompensated cirrhosis. *J Hepatol* 2017;66:849–859.
- [186] Leithead JA, Rajoriya N, Tehami N, Hodson J, Gunson BK, Tripathi D, et al. Non-selective β -blockers are associated with improved survival in patients with ascites listed for liver transplantation. *Gut* 2015;64:1111–1119.
- [187] Senzolo M, Cholongitas E, Burra P, Leandro G, Thalheimer U, Patch D, et al. Beta-blockers protect against spontaneous bacterial peritonitis in cirrhotic patients: a meta-analysis. *Liver Int* 2009;29:1189–1193.
- [188] Reiberger T, Ferlitsch A, Payer BA, Mandorfer M, Heinisch BB, Hayden H, et al. Non-selective beta-blocker therapy decreases intestinal permeability and serum levels of LBP and IL-6 in patients with cirrhosis. *J Hepatol* 2013;58:911–921.
- [189] Mookerjee RP, Pavesi M, Thomsen KL, Mehta G, Macnaughtan J, Bendtsen F, et al. Treatment with non-selective beta blockers is associated with reduced severity of systemic inflammation and improved survival of patients with acute-on-chronic liver failure. *J Hepatol* 2016;64:574–582.
- [190] Madsen BS, Nielsen KF, Fiolla AD, Krag A. Keep the sick from harm in spontaneous bacterial peritonitis: Dose of beta blockers matters. *J Hepatol* 2016;64:1455–1456.
- [191] Serste T, Gustot T, Rautou PE, Francoz C, Njimi H, Durand F, et al. Severe hyponatremia is a better predictor of mortality than MELDNa in patients with cirrhosis and refractory ascites. *J Hepatol* 2012;57:274–280.
- [192] Krag A, Bendtsen F, Henriksen JH, Moller S. Low cardiac output predicts development of hepatorenal syndrome and survival in patients with cirrhosis and ascites. *Gut* 2010;59:105–110.
- [193] Ruiz-del-Arbol L, Urman J, Fernandez J, Gonzalez M, Navasa M, Monescillo A, et al. Systemic, renal, and hepatic hemodynamic derangement in cirrhotic patients with spontaneous bacterial peritonitis. *Hepatology* 2003;38:1210–1218.
- [194] Payance A, Bissonnette J, Roux O, Elkrief L, Gault N, Francoz C, et al. Lack of clinical or haemodynamic rebound after abrupt interruption of beta-blockers in patients with cirrhosis. *Aliment Pharmacol Ther* 2016;43:966–973.
- [195] Garcia-Tsao G, Bosch J. Management of varices and variceal hemorrhage in cirrhosis. *N Engl J Med* 2010;362:823–832.
- [196] Villanueva C, Escorsell A. Optimizing general management of acute variceal bleeding in cirrhosis. *Curr Hepatol Rep* 2014;13:198–207.
- [197] Avgerinos A, Nevens F, Raptis S, Fevery J. Early administration of somatostatin and efficacy of sclerotherapy in acute oesophageal variceal bleeds: the European Acute Bleeding Oesophageal Variceal Episodes (ABOVE) randomised trial. *Lancet* 1997;350:1495–1499.
- [198] Levacher S, Letoumelin P, Pateron D, Blaise M, Lapandry C, Pourriat JL. Early administration of terlipressin plus glyceryl trinitrate to control active upper gastrointestinal bleeding in cirrhotic patients. *Lancet* 1995;346:865–868.
- [199] Garcia-Tsao G, Abraldes JG, Berzigotti A, Bosch J. Portal hypertensive bleeding in cirrhosis: Risk stratification, diagnosis, and management: 2016 practice guidance by the American Association for the study of liver diseases. *Hepatology* 2017;65:310–335.
- [200] Myburgh JA. Fluid resuscitation in acute illness—time to reappraise the basics. *N Engl J Med* 2011;364:2543–2544.
- [201] Villanueva C, Colomo A, Bosch A, Concepcion M, Hernandez-Gea V, Aracil C, et al. Transfusion strategies for acute upper gastrointestinal bleeding. *N Engl J Med* 2013;368:11–21.
- [202] Seo YS, Park SY, Kim MY, Kim JH, Park JY, Yim HJ, et al. Lack of difference among terlipressin, somatostatin, and octreotide in the control of acute gastroesophageal variceal hemorrhage. *Hepatology* 2014;60:954–963.
- [203] Azam Z, Hamid S, Jafri W, Salih M, Abbas Z, Abid S, et al. Short course adjuvant terlipressin in acute variceal bleeding: a randomized double blind dummy controlled trial. *J Hepatol* 2012;56:819–824.
- [204] Altraif I, Handoo FA, Aljumah A, Alalwan A, Dafalla M, Saeed AM, et al. Effect of erythromycin before endoscopy in patients presenting with variceal bleeding: a prospective, randomized, double-blind, placebo-controlled trial. *Gastrointest Endosc* 2011;73:245–250.
- [205] Banares R, Albillos A, Rincon D, Alonso S, Gonzalez M, Ruiz-del-Arbol L, et al. Endoscopic treatment vs. endoscopic plus pharmacologic treatment for acute variceal bleeding: a meta-analysis. *Hepatology* 2002;35:609–615.
- [206] Villanueva C, Piqueras M, Aracil C, Gomez C, Lopez-Balaguer JM, Gonzalez B, et al. A randomized controlled trial comparing ligation and sclerotherapy as emergency endoscopic treatment added to somatostatin in acute variceal bleeding. *J Hepatol* 2006;45:560–567.
- [207] Rios Castellanos E, Seron P, Gisbert JP, Bonfill Cosp X. Endoscopic injection of cyanoacrylate glue vs. other endoscopic procedures for acute bleeding gastric varices in people with portal hypertension. *Cochrane Database Syst Rev* 2015;5:CD010180.
- [208] Bernard B, Grange JD, Khac EN, Amiot X, Opolon P, Poynard T. Antibiotic prophylaxis for the prevention of bacterial infections in cirrhotic patients with gastrointestinal bleeding: a meta-analysis. *Hepatology* 1999;29:1655–1661.
- [209] Fernandez J, Ruiz del Arbol L, Gomez C, Durandez R, Serradilla R, Guarner C, et al. Norfloxacin vs ceftriaxone in the prophylaxis of infections in patients with advanced cirrhosis and hemorrhage. *Gastroenterology* 2006;131:1049–1056.

- [210] Tandon P, Abraldes JG, Keough A, Bastiampillai R, Jayakumar S, Carboneau M, et al. Risk of bacterial infection in patients with cirrhosis and acute variceal hemorrhage, based on child-pugh class, and effects of antibiotics. *Clin Gastroenterol Hepatol* 2015;13:1189–1196.
- [211] Cardenas A, Gines P, Uriz J, Bessa X, Salmeron JM, Mas A, et al. Renal failure after upper gastrointestinal bleeding in cirrhosis: incidence, clinical course, predictive factors, and short-term prognosis. *Hepatology* 2001;34:671–676.
- [212] Shaheen NJ, Stuart E, Schmitz SM, Mitchell KL, Fried MW, Zacks S, et al. Pantoprazole reduces the size of postbanding ulcers after variceal band ligation: a randomized, controlled trial. *Hepatology* 2005;41:588–594.
- [213] Escorsell A, Pavel O, Cardenas A, Morillas R, Llop E, Villanueva C, et al. Esophageal balloon tamponade vs. esophageal stent in controlling acute refractory variceal bleeding: A multicenter randomized, controlled trial. *Hepatology* 2016;63:1957–1996.
- [214] Monesillo A, Martínez-Lagares F, Ruiz-del-Arbol L, et al. Influence of portal hypertension and its early decompression by TIPS placement on the outcome of variceal bleeding. *Hepatology* 2004;40:793–801.
- [215] García-Pagán JC, Caca K, Bureau C, et al. Early use of TIPS in patients with cirrhosis and variceal bleeding. *N Engl J Med* 2010;362:2370–2379.
- [216] Garcia-Pagan JC, Di Pascoli M, Caca K, Laleman W, Bureau C, Appenrodt B, et al. Use of early-TIPS for high-risk variceal bleeding: results of a post-RCT surveillance study. *J Hepatol* 2013;58:45–50.
- [217] Rudler M, Cluzel P, Corvec TL, Benosman H, Rousseau G, Poynard T, et al. Early-TIPSS placement prevents rebleeding in high-risk patients with variceal bleeding, without improving survival. *Aliment Pharmacol Ther* 2014;40:1074–1080.
- [218] Augustin S, Altamirano J, Gonzalez A, Dot J, Abu-Suboh M, Armengol JR, et al. Effectiveness of combined pharmacologic and ligation therapy in high-risk patients with acute esophageal variceal bleeding. *Am J Gastroenterol* 2011;106:1787–1795.
- [219] Reverter E, Tandon P, Augustin S, Turon F, Casu S, Bastiampillai R, et al. A MELD-based model to determine risk of mortality among patients with acute variceal bleeding. *Gastroenterology* 2014;146:412–419.
- [220] Merli M, Nicolini G, Angeloni S, Gentili F, Attili AF, Riggio O. The natural history of portal hypertensive gastropathy in patients with liver cirrhosis and mild portal hypertension. *Am J Gastroenterol* 2004;99:1959–1965.
- [221] Thuluvath PJ, Yoo HY. Portal Hypertensive gastropathy. *Am J Gastroenterol* 2002;97:2973–2978.
- [222] Yoshikawa I, Murata I, Nakano S, Otsuki M. Effects of endoscopic variceal ligation on portal hypertensive gastropathy and gastric mucosal blood flow. *Am J Gastroenterol* 1998;93:71–74.
- [223] Urrunaga NH, Rockey DC. Portal hypertensive gastropathy and colopathy. *Clin Liver Dis* 2014;18:389–406.
- [224] Hosking SW, Kennedy HJ, Seddon I, Triger DR. The role of propranolol in congestive gastropathy of portal hypertension. *Hepatology* 1987;7:437–441.
- [225] Perez-Ayuso RM, Pique JM, Bosch J, Panes J, Gonzalez A, Perez R, et al. Propranolol in prevention of recurrent bleeding from severe portal hypertensive gastropathy in cirrhosis. *Lancet* 1991;337:1431–1434.
- [226] Ripoll C, Garcia-Tsao G. Treatment of gastropathy and gastric antral vascular ectasia in patients with portal hypertension. *Curr Treat Options Gastroenterol* 2007;10:483–494.
- [227] Kamath P, Lacerda M, Ahlquist D, McKusick MA, Andrews JC, Nagorney DA. Gastric Mucosal Responses to intrahepatic portosystemic shunting in patients with cirrhosis. *Gastroenterology* 2000;118:905–911.
- [228] Zhou Y, Qiao L, Wu J, Hu H, Xu C. Comparison of the efficacy of octreotide, vasopressin, and omeprazole in the control of acute bleeding in patients with portal hypertensive gastropathy: a controlled study. *J Gastroenterol Hepatol* 2002;17:973–979.
- [229] Sarin SK, Lahoti D, Saxena SP, Murthy NS, Makwana UK. Prevalence, classification and natural history of gastric varices: a long-term follow-up study in 568 portal hypertension patients. *Hepatology* 1992;16:1343–1349.
- [230] Mishra SR, Sharma BC, Kumar A, Sarin SK. Primary prophylaxis of gastric variceal bleeding comparing cyanoacrylate injection and beta-blockers: a randomized controlled trial. *J Hepatol* 2011;54:1161–1167.
- [231] Rios CE, Seron P, Gisbert JP, Bonfill CX. Endoscopic injection of cyanoacrylate glue vs. other endoscopic procedures for acute bleeding gastric varices in people with portal hypertension. *Cochrane Database Syst Rev* 2015;5:CD010180.
- [232] Mishra SR, Chander Sharma B, Kumar A, Sarin SK. Endoscopic cyanoacrylate injection vs. beta-blocker for secondary prophylaxis of gastric variceal bleed: a randomised controlled trial. *Gut* 2010;59:729–735.
- [233] Chau TN, Patch D, Chan YW, Nagral A, Dick R, Burroughs AK. “Salvage” transjugular intrahepatic portosystemic shunts: gastric fundal compared with esophageal variceal bleeding. *Gastroenterology* 1998;114:981–998.
- [234] Hung HH, Chang CJ, Hou MC, Liao WC, Chan CC, Huang HC, et al. Efficacy of non-selective beta-blockers as adjunct to endoscopic prophylactic treatment for gastric variceal bleeding: a randomized controlled trial. *J Hepatol* 2012;56:1025–1032.
- [235] Lo GH, Liang HL, Chen WC, Chen MH, Lai KH, Hsu PI, et al. A prospective, randomized controlled trial of transjugular intrahepatic portosystemic shunt vs. cyanoacrylate injection in the prevention of gastric variceal rebleeding. *Endoscopy* 2007;39:679–685.
- [236] Saad WE. Endovascular management of gastric varices. *Clin Liver Dis* 2014;18:829–885.
- [237] Mookerjee RP, Stadlbauer V, Lidder S, Wright GA, Hodges SJ, Davies NA, et al. Neutrophil dysfunction in alcoholic hepatitis superimposed on cirrhosis is reversible and predicts the outcome. *Hepatology* 2007;46:831–840.
- [238] Wasmuth HE, Kunz D, Yagmur E, Timmer-Stranghoner A, Vidacek D, Siewert E, et al. Patients with acute on chronic liver failure display “sepsis-like” immune paralysis. *J Hepatol* 2005;42:195–201.
- [239] Wiest R, Garcia-Tsao G. Bacterial translocation (BT) in cirrhosis. *Hepatology* 2005;41:422–433.
- [240] Appenrodt B, Grunhage F, Gentemann MG, Thyssen L, Sauerbruch T, Lammert F. Nucleotide-binding oligomerization domain containing 2 (NOD2) variants are genetic risk factors for death and spontaneous bacterial peritonitis in liver cirrhosis. *Hepatology* 2010;51:1327–1333.
- [241] Fernandez J, Navasa M, Gomez J, et al. Bacterial infections in cirrhosis: epidemiological changes with invasive procedures and norfloxacin prophylaxis. *Hepatology* 2002;35:140–148.
- [242] Fasolato S, Angeli P, Dallagnese L, et al. Renal failure and bacterial infections in patients with cirrhosis: epidemiology and clinical features. *Hepatology* 2007;45:223–229.
- [243] Piano S, Fasolato S, Salinas F, Romano A, Tonon M, Morando F, et al. The empirical antibiotic treatment of nosocomial spontaneous bacterial peritonitis: Results of a randomized, controlled clinical trial. *Hepatology* 2016;63:1299–1309.
- [244] Runyon BA. The evolution of ascitic fluid analysis in the diagnosis of spontaneous bacterial peritonitis. *Runyon BA. Am J Gastroenterol* 2003;98:1675–1677.
- [245] Evans LT, Kim WR, Poterucha JJ, Kamath PS. Spontaneous bacterial peritonitis in asymptomatic outpatients with cirrhotic ascites. *Hepatology* 2003;37:897–901.
- [246] Kim JJ, Tsukamoto MM, Mathur AK, Ghomri YM, Hou LA, Sheibani S, et al. Delayed paracentesis is associated with increased in-hospital mortality in patients with spontaneous bacterial peritonitis. *Am J Gastroenterol* 2014;109:1436–1442.
- [247] Van de Geijn GM, van Gent M, van Pul-Bom N, Beunis MH, van Tilburg AJ, Njo TL. A new flow cytometric method for differential cell counting in ascitic fluid. *Cytometry B Clin Cytom* 2016;90:506–511.
- [248] Fleming C, Brouwer R, van Alphen A, Lindemans J, de Jonge R. UF-1000i: validation of the body fluid mode for counting cells in body fluids. *Clin Chem Lab Med* 2014;52:1781–1790.
- [249] Gülberg V, Gerbes AL, Sauerbruch T, Appenrodt B. Insufficient sensitivity of reagent strips for spontaneous bacterial peritonitis. *Hepatology* 2007;46:1669.
- [250] Bellot P, García-Pagán JC, Francés R, Abraldes JG, Navasa M, Pérez-Mateo M, et al. Bacterial DNA translocation is associated with systemic circulatory abnormalities and intrahepatic endothelial dysfunction in patients with cirrhosis. *Hepatology* 2010;52:2044–2052.
- [251] Bruns T, Reuken PA, Stengel S, Gerber L, Appenrodt B, Schade JH, et al. The prognostic significance of bacterial DNA in patients with decompensated cirrhosis and suspected infection. *Liver Int* 2016;36:1133–1142.
- [252] Runyon BA, Hoefs JC. Culture-negative neutrocytic ascites: a variant of spontaneous bacterial peritonitis. *Hepatology* 1984;4:1209–1211.
- [253] Gravito-Soares M, Gravito-Soares E, Lopes S, Ribeiro G, Figueiredo P. Spontaneous fungal peritonitis: a rare but severe complication of liver cirrhosis. *Eur J Gastroenterol Hepatol* 2017;29:1010–1016.
- [254] Chen CH, Shih CM, Chou JW, Liu YH, Hang LW, Hsia TC, et al. Outcome predictors of cirrhotic patients with spontaneous bacterial empyema. *Liver Int* 2011;31:417–424.

Clinical Practice Guidelines

- [255] Xiol X, Castellvi JM, Guardiola J, Sese E, Castellote J, Perello A, et al. Spontaneous bacterial empyema in cirrhotic patients: a prospective study. *Hepatology* 1996;23:719–723.
- [256] Soriano G, Castellote J, Alvarez C, Girbau A, Gordillo J, Baliellas C, et al. Secondary bacterial peritonitis in cirrhosis: a retrospective study of clinical and analytical characteristics, diagnosis and management. *J Hepatol* 2010;52:39–44.
- [257] Rimola A, Salmeron JM, Clemente G, Rodrigo L, Obrador A, Miranda ML, et al. Two different dosages of cefotaxime in the treatment of spontaneous bacterial peritonitis in cirrhosis: results of a prospective, randomized, multicenter study. *Hepatology* 1995;21:674–679.
- [258] Runyon Runyon BA, McHutchison JG, Antillon MR, Akriviadis EA, Montano AA. Short-course vs. long-course antibiotic treatment of spontaneous bacterial peritonitis. A randomized controlled study of 100 patients. *Gastroenterology* 1991;100:1737–1742.
- [259] Ricart E, Soriano G, Novella MT, Ortiz J, Sabat M, Kolle L, et al. Amoxicillin-clavulanic acid vs. cefotaxime in the therapy of bacterial infections in cirrhotic patients. *J Hepatol* 2000;32:596–602.
- [260] deLemos AS, Ghabril M, Rockey DC, Gu J, Barnhart HX, Fontana RJ, et al. Drug-induced liver injury network (DILIN). Amoxicillin-clavulanate-induced liver injury. *Dig Dis Sci* 2016;61:2406–2416.
- [261] Terg R, Cobas S, Fassio E, Landeira G, Rios B, Vasen W, et al. Oral ciprofloxacin after a short course of intravenous ciprofloxacin in the treatment of spontaneous bacterial peritonitis: results of a multicenter, randomized study. *J Hepatol* 2000;33:564–569.
- [262] Angeli P, Guarda S, Fasolato S, Miola E, Craighero R, Piccolo F, et al. Switch therapy with ciprofloxacin vs. intravenous ceftazidime in the treatment of spontaneous bacterial peritonitis in patients with cirrhosis: similar efficacy at lower cost. *Aliment Pharmacol Ther* 2006;23:75–84.
- [263] Navasa M, Follo A, Llovet JM, Clemente G, Vargas V, Rimola A, et al. Randomized, comparative study of oral ofloxacin vs. intravenous cefotaxime in spontaneous bacterial peritonitis. *Gastroenterology* 1996;111:1011–1017.
- [264] Fernandez J, Bert F, Nicolas-Chanoine MH. The challenges of multi-drug-resistance in hepatology. *J Hepatol* 2016;65:1043–1054.
- [265] Magiorakos AP, Srinivasan A, Carey RB, Carmeli Y, Falagas ME, Giske CG, et al. Multidrug-resistant, extensively drug-resistant and pandrug-resistant bacteria: an international expert proposal for interim standard definitions for acquired resistance. *Clin Microbiol Infect* 2012;18:268–281.
- [266] Fernandez J, Acevedo J, Castro M, Garcia O, de Lope CR, Roca D, et al. Prevalence and risk factors of infections by multiresistant bacteria in cirrhosis: a prospective study. *Hepatology* 2012;55:1551–1561.
- [267] Wiest R, Krag A, Gerbes A. Spontaneous bacterial peritonitis: recent guidelines and beyond. *Gut* 2012;61:297–310.
- [268] Piano S, Brocca A, Mareso S, Angeli P. Infections complicating cirrhosis. *Liver Int* 2018;38:126–133.
- [269] Lutz P, Nischalke HD, Kramer B, Goeser F, Kaczmarek DJ, Schlabe S, et al. Antibiotic resistance in healthcare-related and nosocomial spontaneous bacterial peritonitis. *Eur J Clin Invest* 2017;47:44–52.
- [270] Singer M, Deutschman CS, Seymour CW, Shankar-Hari M, Annane D, Bauer M, et al. The third international consensus definitions for sepsis and septic shock (Sepsis-3). *JAMA* 2016;315:801–810.
- [271] Piano S, Bartoletti M, Tonon M, Baldassarre M, Chies G, Romano A, et al. Assessment of Sepsis-3 criteria and quick SOFA in patients with cirrhosis and bacterial infections. *Gut* 2017.
- [272] Sort P, Navasa M, Arroyo V, Aldeguer X, Planas R, Ruiz-del-Arbol L, et al. Effect of intravenous albumin on renal impairment and mortality in patients with cirrhosis and spontaneous bacterial peritonitis. *N Engl J Med* 1999;341:403–409.
- [273] Garioud A, Cadranet JF, Pauwels A, Nousbaum JB, Thévenot T, Dao T, et al. Association Nationale des Hépatogastroentérologues des Hôpitaux Généraux de France, Association Française pour l'Etude du Foie, Club de Réflexion des Cabinets et Groupes d'Hépatogastroentérologie. Albumin Use in Patients With Cirrhosis in France: Results of the "ALBU-LIVE" Survey: A Case for Better EASL Guidelines Diffusion and/or Revision. *J Clin Gastroenterol* 2017;51:831–838.
- [274] Poca M, Concepcion M, Casas M, Alvarez-Urturi C, Gordillo J, Hernandez-Gea V, et al. Role of albumin treatment in patients with spontaneous bacterial peritonitis. *Clin Gastroenterol Hepatol* 2012;10:309–315.
- [275] Fernandez J, Tandon P, Mensa J, Garcia-Tsao G. Antibiotic prophylaxis in cirrhosis: Good and bad. *Hepatology* 2016;63:2019–2031.
- [276] Fernandez J, Navasa M, Planas R, Montoliu S, Monfort D, Soriano G, et al. Primary prophylaxis of spontaneous bacterial peritonitis delays hepatorenal syndrome and improves survival in cirrhosis. *Gastroenterology* 2007;133:818–824.
- [277] Terg R, Fassio E, Guevara M, Cartier M, Longo C, Lucero R, et al. Ciprofloxacin in primary prophylaxis of spontaneous bacterial peritonitis: A randomized, placebo-controlled study. *J Hepatol* 2008;48:774–779.
- [278] Loomba R, Wesley R, Bain A, et al. Role of fluoroquinolones in the primary prophylaxis of spontaneous bacterial peritonitis: meta-analysis. *Clin Gastroenterol Hepatol* 2009;7:487–493.
- [279] Saab S, Hernandez JC, Chi AC, et al. Oral antibiotic prophylaxis reduces spontaneous bacterial peritonitis occurrence and improves short-term survival in cirrhosis: a meta-analysis. *Am J Gastroenterol* 2009;104:993–1001.
- [280] Krag A, Wiest R, Gluud LL. Fluoroquinolones in the primary prophylaxis of spontaneous bacterial peritonitis. *Am J Gastroenterol* 2010;105:1444–1445.
- [281] Gines P, Rimola A, Planas R, Vargas V, Marco F, Almela M, et al. Norfloxacin prevents spontaneous bacterial peritonitis recurrence in cirrhosis: results of a double-blind, placebo-controlled trial. *Hepatology* 1990;12:716–724.
- [282] Bauer TM, Follo A, Navasa M, Vila J, Planas R, Clemente G, et al. Daily norfloxacin is more effective than weekly rifloxacin in prevention of spontaneous bacterial peritonitis recurrence. *Dig Dis Sci* 2002;47:1356–1361.
- [283] Terg R, Llano K, Cobas SM, Brotto C, Barrios A, Levi D, et al. Effects of oral ciprofloxacin on aerobic gram-negative fecal flora in patients with cirrhosis: results of short- and long-term administration, with daily and weekly dosages. *J Hepatol* 1998;29:437–442.
- [284] Bass NM, Mullen KD, Sanyal A, Poordad F, Neff G, Leevy CB, et al. Rifaximin treatment in hepatic encephalopathy. *N Engl J Med* 2010;362:071–1081.
- [285] Elfert A, Abo Ali L, Soliman S, Ibrahim S, Abd-Elsalam S. Randomized-controlled trial of rifaximin vs. norfloxacin for secondary prophylaxis of spontaneous bacterial peritonitis. *Eur J Gastroenterol Hepatol* 2016;28:1450–1454.
- [286] Min YW, Lim KS, Min BH, Gwak GY, Paik YH, Choi MS, et al. Proton pump inhibitor use significantly increases the risk of spontaneous bacterial peritonitis in 1965 patients with cirrhosis and ascites: a propensity score matched cohort study. *Aliment Pharmacol Ther* 2014;40:695–704.
- [287] Dam G, Vilstrup H, Watson H, Jepsen P. Proton pump inhibitors as a risk factor for hepatic encephalopathy and spontaneous bacterial peritonitis in patients with cirrhosis with ascites. *Hepatology* 2016;64:1265–1272.
- [288] Mandorfer M, Bota S, Schwabl P, Bucsics T, Pfisterer N, Kruzik M, et al. Nonselective beta blockers increase risk for hepatorenal syndrome and death in patients with cirrhosis and spontaneous bacterial peritonitis. *Gastroenterology* 2014;146:1680–1690.
- [289] Pande C, Kumar A, Sarin SK. Addition of probiotics to norfloxacin does not improve efficacy in the prevention of spontaneous bacterial peritonitis: a double-blind placebo-controlled randomized-controlled trial. *Eur J Gastroenterol Hepatol* 2012;24:831–839.
- [290] Fernandez J, Acevedo J, Arroyo V. Response to the clinical course and short-term mortality of cirrhotic patients with non-spontaneous bacterial peritonitis infections. *Liver Int* 2017;37:623.
- [291] Arvaniti V, D'Amico G, Fede G, Manousou P, Tsochatzis E, Pleguezuelo M, et al. Infections in patients with cirrhosis increase mortality four-fold and should be used in determining prognosis. *Gastroenterology* 2010;139:1256.
- [292] Piano S, Morando F, Carretta G, Tonon M, Vettore E, Rosi S, et al. Predictors of early readmission in patients with cirrhosis after the resolution of bacterial infections. *Am J Gastroenterol* 2017;112:1575–1583.
- [293] Bartoletti M, Giannella M, Lewis R, Caraceni P, Tedeschi S, Paul M, et al. A prospective multicentre study of the epidemiology and outcomes of bloodstream infection in cirrhotic patients. *Clin Microbiol Infect* 2017.
- [294] Merli M, Lucidi C, Di Gregorio V, Lattanzi B, Giannelli V, Giusto M, et al. An empirical broad spectrum antibiotic therapy in health-care-associated infections improves survival in patients with cirrhosis: A randomized trial. *Hepatology* 2016;63:1632–1639.
- [295] Fiore M, Chiodini P, Pota V, Sansone P, Passavanti MB, Leone S, et al. Risk of spontaneous fungal peritonitis in hospitalized cirrhotic patients with ascites: a systematic review of observational studies and meta-analysis. *Minerva Anestesiol* 2017;83:1309–1316.
- [296] Piano S, Angeli P. Reply letter. *Hepatology* 2016;64:998–999.

- [297] Guevara M, Terra C, Nazar A, Sola E, Fernandez J, Pavesi M, et al. Albumin for bacterial infections other than spontaneous bacterial peritonitis in cirrhosis. A randomized, controlled study. *J Hepatol* 2012;57:759–765.
- [298] Thevenot T, Bureau C, Oberti F, Anty R, Louvet A, Plessier A, et al. Effect of albumin in cirrhotic patients with infection other than spontaneous bacterial peritonitis. A randomized trial. *J Hepatol* 2015;62:822–830.
- [299] Piano S, Romano A, Di Pascoli M, Angeli P. Why and how to measure renal function in patients with liver disease. *Liver Int* 2017;37:116–122.
- [300] Gonwa TA, Jennings L, Mai ML, Stark PC, Levey AS, Klintmalm GB. Estimation of glomerular filtration rates before and after orthotopic liver transplantation: evaluation of current equations. *Liver Transpl* 2004;10:301–309.
- [301] Hoek FJ, Kemperman FA, Krediet RT. A comparison between cystatin C, plasma creatinine and the Cockcroft and Gault formula for the estimation of glomerular filtration rate. *Nephrol Dial Transplant* 2003;18:2024–2031.
- [302] Inker LA, Schmid CH, Tighiouart H, Eckfeldt JH, Feldman HI, Greene T, et al. Estimating glomerular filtration rate from serum creatinine and cystatin C. *N Engl J Med* 2012;367:20–29.
- [303] Roy L, Legault L, Pomier-Layrargues G. Glomerular filtration rate measurement in cirrhotic patients with renal failure. *Clin Nephrol* 1998;50:342–346.
- [304] Francoz C, Nadim MK, Baron A, Prie D, Antoine C, Belghiti J, et al. Glomerular filtration rate equations for liver-kidney transplantation in patients with cirrhosis: validation of current recommendations. *Hepatology* 2014;59:1514–1521.
- [305] Francoz C, Glotz D, Moreau R, Durand F. The evaluation of renal function and disease in patients with cirrhosis. *J Hepatol* 2010;52:605–613.
- [306] Garcia-Tsao G, Parikh CR, Viola A. Acute kidney injury in cirrhosis. *Hepatology* 2008;48:2064–2077.
- [307] Salerno F, Gerbes A, Gines P, Wong F, Arroyo V. Diagnosis, prevention and treatment of hepatorenal syndrome in cirrhosis. *Gut* 2007;56:310–318.
- [308] Mehta RL, Kellum JA, Shah SV, Molitoris BA, Ronco C, Warnock DG, et al. Acute Kidney Injury Network: report of an initiative to improve outcomes in acute kidney injury. *Crit Care* 2007;11:R31.
- [309] Nadim MK, Kellum JA, Davenport A, Wong F, Davis C, Pannu N, et al. Hepatorenal syndrome: the 8th International Consensus Conference of the Acute Dialysis Quality Initiative (ADQI) Group. *Crit Care* 2012;16:R23.
- [310] Khwaja A. KDIGO clinical practice guidelines for acute kidney injury. *Nephron Clin Pract* 2012;120:c179–c184.
- [311] Belcher JM, Garcia-Tsao G, Sanyal AJ, Bhogal H, Lim JK, Ansari N, et al. Association of AKI with mortality and complications in hospitalized patients with cirrhosis. *Hepatology* 2013;57:753–762.
- [312] de Carvalho JR, Villela-Nogueira CA, Luiz RR, Guzzo PL, da Silva Rosa JM, Rocha E, et al. Acute kidney injury network criteria as a predictor of hospital mortality in cirrhotic patients with ascites. *J Clin Gastroenterol* 2012;46:21–26.
- [313] Fagundes C, Barreto R, Guevara M, Garcia E, Solà E, Rodríguez E, et al. A modified acute kidney injury classification for diagnosis and risk stratification of impairment of kidney function in cirrhosis. *J Hepatol* 2013;59:474–481.
- [314] Piano S, Rosi S, Maresio G, Fasolato S, Cavallin M, Romano A, et al. Evaluation of the Acute Kidney Injury Network criteria in hospitalized patients with cirrhosis and ascites. *J Hepatol* 2013;59:482–489.
- [315] Tsiens CD, Rabie R, Wong F. Acute kidney injury in decompensated cirrhosis. *Gut* 2013;62:131–137.
- [316] Wong F, O'Leary JG, Reddy KR, Patton H, Kamath PS, Fallon MB, et al. New consensus definition of acute kidney injury accurately predicts 30-day mortality in patients with cirrhosis and infection. *Gastroenterology* 2013;145:1280–1288.
- [317] Huelin P, Piano S, Solà E, Stanco M, Solé C, Moreira R, et al. Validation of a staging system for acute kidney injury in patients with cirrhosis and association with acute-on-chronic liver failure. *Clin Gastroenterol Hepatol* 2017;15:438–445.
- [318] Angeli P, Gines P, Wong F, Bernardi M, Boyer TD, Gerbes A, et al. Diagnosis and management of acute kidney injury in patients with cirrhosis: revised consensus recommendations of the International Club of Ascites. *J Hepatol* 2015;62:968–974.
- [319] Kidney Disease: Improving Global Outcomes (KDIGO) CKD Work Group. KDIGO. Clinical practice guideline for the evaluation and management of chronic kidney disease. *Kidney Int* 2012;2013:1–150.
- [320] Rosi S, Piano S, Frigo AC, Morando F, Fasolato S, Cavallin M, et al. New ICA criteria for the diagnosis of acute kidney injury in cirrhotic patients: can we use an imputed value of serum creatinine? *Liver Int* 2015;35:2108–2114.
- [321] Guevara M, Fernandez-Esparrach G, Alessandria C, Torre A, Terra C, Montana X, et al. Effects of contrast media on renal function in patients with cirrhosis: a prospective study. *Hepatology* 2004;40:646–651.
- [322] Umgelter A, Reindl W, Franzen M, Lenhardt C, Huber W, Schmid RM. Renal resistive index and renal function before and after paracentesis in patients with hepatorenal syndrome and tense ascites. *Intensive Care Med* 2009;35:152–156.
- [323] Cabrera J, Falcón L, Gorriç E, Pardo MD, Granados R, Quinones A, Maynar M. Abdominal decompression plays a major role in early postparacentesis haemodynamic changes in cirrhotic patients with tense ascites. *Gut* 2001;48:384–389.
- [324] de Cleve R, Silva FP, Zilberstein B, Machado DJ. Acute renal failure due to abdominal compartment syndrome: report on four cases and literature review. *Rev Hospital das Clin* 2001;56:123–130.
- [325] Nadim MK, Durand F, Kellum JA, Levitsky J, O'Leary JG, Karvellas CJ, et al. Management of the critically ill patient with cirrhosis: A multidisciplinary perspective. *J Hepatol* 2016;64:717–735.
- [326] Umgelter A, Reindl W, Wagner KS, Franzen M, Stock K, Schmid RM, et al. Effects of plasma expansion with albumin and paracentesis on haemodynamics and kidney function in critically ill cirrhotic patients with tense ascites and hepatorenal syndrome: a prospective uncontrolled trial. *Critical care* 2008;12:R4.
- [327] Moreau R, Lebrec D. Diagnosis and treatment of acute renal failure in patients with cirrhosis. *Best Pract Res Clin Gastroenterol* 2007;21:111–123.
- [328] Moreau R, Lebrec D. Acute renal failure in patients with cirrhosis: perspectives in the age of MELD. *Hepatology* 2003;37:233–243.
- [329] Fagundes C, Pepin MN, Guevara M, Barreto R, Casals G, Sola E, et al. Urinary neutrophil gelatinase-associated lipocalin as biomarker in the differential diagnosis of impairment of kidney function in cirrhosis. *J Hepatol* 2012;57:267–273.
- [330] Belcher JM, Sanyal AJ, Peixoto AJ, Perazella MA, Lim J, Thiessen-Philbrook H, et al. Kidney biomarkers and differential diagnosis of patients with cirrhosis and acute kidney injury. *Hepatology* 2014;60:622–632.
- [331] Qasem AA, Farag SE, Hamed E, Emar M, Bihery A, Pasha H. Urinary biomarkers of acute kidney injury in patients with liver cirrhosis. *ISRN Nephrol* 2014;2014:376795.
- [332] Barreto R, Elia C, Sola E, Moreira R, Ariza X, Rodriguez E, et al. Urinary neutrophil gelatinase-associated lipocalin predicts kidney outcome and death in patients with cirrhosis and bacterial infections. *J Hepatol* 2014;61:35–42.
- [333] Ariza X, Sola E, Elia C, Barreto R, Moreira R, Morales-Ruiz M, et al. Analysis of a urinary biomarker panel for clinical outcomes assessment in cirrhosis. *PLoS One* 2015;10:e0128145.
- [334] Puthumana J, Ariza X, Belcher JM, Graupera I, Ginès P, Parikh CR. Urine interleukin 18 and lipocalin 2 are biomarkers of acute tubular necrosis in patients with cirrhosis: a systematic review and meta-analysis. *Clin Gastroenterol Hepatol* 2017;15:1003–1013.
- [335] Huelin P, Elia C, Solà E, Solé C, Moreira R, Carol M. New diagnostic algorithm of acute kidney injury in cirrhosis that includes categorization of stage 1 and assessment of urine NGAL. Relevance for the differential diagnosis and clinical outcomes. *J Hepatol* 2017;66:S11.
- [336] Chawla LS, Eggers PW, Star RA, Kimmel PL. Acute kidney injury and chronic kidney disease as interconnected syndromes. *N Engl J Med* 2014;371:58–66.
- [337] Trawale JM, Paradis V, Rautou PE, Francoz C, Escolano S, Sallee M, et al. The spectrum of renal lesions in patients with cirrhosis: a clinicopathological study. *Liver Int* 2010;30:725–732.
- [338] Wade HM, Geiger XJ, Cortese C, Mai ML, Kramer DJ, Rosser BG, et al. Kidney allocation to liver transplant candidates with renal failure of undetermined etiology: role of percutaneous renal biopsy. *Am J Transplant* 2008;8:2618–2626.
- [339] Wong F, Nadim MK, Kellum JA, Salerno F, Bellomo R, Gerbes A, et al. Working Party proposal for a revised classification system of renal dysfunction in patients with cirrhosis. *Gut* 2011;60:702–709.
- [340] Stadlbauer V, Wright GA, Banaji M, Mukhopadhyaya A, Mookerjee RP, Moore K, et al. Relationship between activation of the sympathetic nervous system and renal blood flow autoregulation in cirrhosis. *Gastroenterology* 2008;134:111–119.
- [341] Wiest R, Lawson M, Geuking M. Pathological bacterial translocation in liver cirrhosis. *J Hepatol* 2014;60:197–209.

- [342] Albillos A, de la Hera A, Gonzalez M, Moya JL, Calleja JL, Monserrat J, et al. Increased lipopolysaccharide binding protein in cirrhotic patients with marked immune and hemodynamic derangement. *Hepatology* 2003;37:208–217.
- [343] Navasa M, Follo A, Filella X, Jimenez W, Francitorra A, Planas R, et al. Tumor necrosis factor and interleukin-6 in spontaneous bacterial peritonitis in cirrhosis: relationship with the development of renal impairment and mortality. *Hepatology* 1998;27:1227–1232.
- [344] Shah N, Dhar D, El Zahraa Mohammed F, Habtesion A, Davies NA, Jover-Cobos M, et al. Prevention of acute kidney injury in a rodent model of cirrhosis following selective gut decontamination is associated with reduced renal TLR4 expression. *J Hepatol* 2012;56:1047–1053.
- [345] Shah N, Mohamed FE, Jover-Cobos M, Macnaughtan J, Davies N, Moreau R, et al. Increased renal expression and urinary excretion of TLR4 in acute kidney injury associated with cirrhosis. *Liver Int* 2013;33:398–409.
- [346] Alobaidi R, Basu RK, Goldstein SL, Bagshaw SM. Sepsis-associated acute kidney injury. *Semin Nephrol* 2015;35:2–11.
- [347] Emet DR, Shaw AD, Kellum JA. Sepsis-associated AKI: epithelial cell dysfunction. *Semin Nephrol* 2015;35:85–95.
- [348] Prowle JR, Bellomo R. Sepsis-associated acute kidney injury: macrohemodynamic and microhemodynamic alterations in the renal circulation. *Semin Nephrol* 2015;35:64–74.
- [349] de Seigneux S, Martin PY. Preventing the progression of AKI to CKD: the role of mitochondria. *J Am Soc Nephrol* 2017;28:1327–1329.
- [350] Bairaktari E, Liamis G, Tsolas O, Elisaf M. Partially reversible renal tubular damage in patients with obstructive jaundice. *Hepatology* 2001;33:1365–1369.
- [351] van Slambrouck CM, Salem F, Meehan SM, Chang A. Bile cast nephropathy is a common pathologic finding fo kidney injury associated with severe liver dysfunction. *Kidney Int* 2013;84:192–197.
- [352] Durand F, Graupera I, Gines P, Olson JC, Nadim MK. Pathogenesis of hepatorenal syndrome: implications for therapy. *Am J Kidney Dis* 2016;67:318–328.
- [353] Moreau R, Durand F, Poynard T, Duhamel C, Cervoni JP, Ichai P, et al. Terlipressin in patients with cirrhosis and type 1 hepatorenal syndrome: a retrospective multicenter study. *Gastroenterology* 2002;122:923–930.
- [354] Ortega R, Gines P, Uriz J, et al. Terlipressin therapy with and without albumin for patients with hepatorenal syndrome: results of a prospective, nonrandomized study. *Hepatology* 2002;36:941–948.
- [355] Sanyal AJ, Boyer T, Garcia-Tsao G, Regenstein F, Rossaro L, Appenrodt B, et al. A randomized prospective double blind, placebo controlled study of terlipressin for type 1 hepatorenal syndrome. *Gastroenterology* 2008;134:1360–1368.
- [356] Martín-Llahi M, Pepin MN, Guevara M, Diaz F, Torre A, Monescillo A, et al. Terlipressin and albumin vs albumin in patients with cirrhosis and hepatorenal syndrome: a randomized study. *Gastroenterology* 2008;134:1352–1359.
- [357] Boyer TD, Sanyal AJ, Wong F, Frederick RT, Lake JR, O'Leary JG, et al. REVERSE study investigators terlipressin plus albumin is more effective than albumin alone in improving renal function in patients with cirrhosis and hepatorenal syndrome type 1. *Gastroenterology* 2016;150:1579–1589.
- [358] Rodriguez E, Elia C, Sola E, Barreto R, Graupera I, Andrealli A, et al. Terlipressin and albumin for type-1 hepatorenal syndrome associated with sepsis. *J Hepatol* 2014;60:955–961.
- [359] Cavallin M, Kamath PS, Merli M, Fasolato S, Toniutto P, Salerno F, et al. Terlipressin plus albumin vs. midodrine and octreotide plus albumin in the treatment of hepatorenal syndrome: A randomized trial. *Hepatology* 2015;62:567–574.
- [360] Cavallin M, Piano S, Romano A, Fasolato S, Frigo AC, Benetti G, et al. Terlipressin given by continuous intravenous infusion vs. intravenous boluses in the treatment of hepatorenal syndrome: A randomized controlled study. *Hepatology* 2016;63:983–992.
- [361] Gluud LL, Christensen K, Christensen E, Krag A. Terlipressin for hepatorenal syndrome. *Cochrane Database Syst Rev* 2012;CD005162.
- [362] Facciorusso A, Chandar AK, Murad MH, Prokop LJ, Muscatiello N, Kamath PS. Singh S Comparative efficacy of pharmacological strategies for management of type 1 hepatorenal syndrome: a systematic review and network meta-analysis. *Lancet Gastroenterol Hepatol* 2017;2:94–102.
- [363] Narahara Y, Kanazawa H, Taki Y, Kimura Y, Atsukawa M, Katakura T, et al. Effects of terlipressin on systemic, hepatic and renal hemodynamics in patients with cirrhosis. *J Gastroenterol Hepatol* 2009;24:1791–1797.
- [364] Garcia-Martinez R, Caraceni P, Bernardi M, Gines P, Arroyo V, Jalan R. Albumin: pathophysiological basis of its role in the treatment of cirrhosis and its complications. *Hepatology* 2013;58:1836–1846.
- [365] Gerbes AL, Huber E, Gulberg V. Terlipressin for hepatorenal syndrome: continuous infusion as an alternative to i.v. bolus administration. *Gastroenterology* 2009;137:1179–1181.
- [366] Piano S, Morando F, Fasolato S, Cavallin M, Boscato N, Boccagni P, et al. Continuous recurrence of type 1 hepatorenal syndrome and long-term treatment with terlipressin and albumin: a new exception to MELD score in the allocation system to liver transplantation? *J Hepatol* 2011;55:491–496.
- [367] Gow PJ, Ardalán ZS, Vasudevan A, Testro AG, Ye B, Angus PW. Outpatient terlipressin infusion for the treatment of refractory ascites. *Am J Gastroenterol* 2016;111:1041–1042.
- [368] Alessandria C, Ottobrelli A, Debernardi-Venon W, Todros L, Cerenzia MT, Martini S, et al. Noradrenalin vs terlipressin in patients with hepatorenal syndrome: a prospective, randomized, unblinded, pilot study. *J Hepatol* 2007;47:499–505.
- [369] Duvoux C, Zanditenas D, Hezode C, Chauvat A, Monin JL, Roudot-Thoraval F, et al. Effects of noradrenalin and albumin in patients with type I hepatorenal syndrome: a pilot study. *Hepatology* 2002;36:374–380.
- [370] Singh V, Ghosh S, Singh B, Kumar P, Sharma N, Bhalla A, et al. Noradrenaline vs. terlipressin in the treatment of hepatorenal syndrome: a randomized study. *J Hepatol* 2012;56:1293–1298.
- [371] Sharma P, Kumar A, Shrama BC, Sarin SK. An open label, pilot, randomized controlled trial of noradrenaline vs. terlipressin in the treatment of type 1 hepatorenal syndrome and predictors of response. *Am J Gastroenterol* 2008;103:1689–1697.
- [372] Esraillian E, Pantangco ER, Kyulo NL, Hu KQ, Runyon BA. Octreotide/Midodrine therapy significantly improves renal function and 30-day survival in patients with type 1 hepatorenal syndrome. *Dig Dis Sci* 2007;52:742–748.
- [373] Restuccia T, Ortega R, Guevara M, Gines P, Alessandria C, Ozdogan O, et al. Effects of treatment of hepatorenal syndrome before transplantation on posttransplantation outcome. A case-control study. *J Hepatol* 2004;40:140–146.
- [374] Rodriguez E, Henrique Pereira G, Sola E, Elia C, Barreto R, Pose E, et al. Treatment of type 2 hepatorenal syndrome in patients awaiting transplantation: Effects on kidney function and transplantation outcomes. *Liver Transpl* 2015;21:1347–1354.
- [375] Boyer TD, Sanyal AJ, Garcia-Tsao G, Blei A, Carl D, Bexon AS, et al. Predictors of response to terlipressin plus albumin in hepatorenal syndrome (HRS) type 1: relationship of serum creatinine to hemodynamics. *J Hepatol* 2011;55:315–321.
- [376] Piano S, Schmidt HH, Ariza X, Amoros A, Romano A, Solà E, et al. Impact of Acute-on-Chronic Liver Failure on response to treatment with terlipressin and albumin in patients with type 1 hepatorenal syndrome. *J Hepatol* 2017;66:5572.
- [377] Nazar A, Pereira GH, Guevara M, Martín-Llahi M, Pepin MN, Marinelli M, et al. Predictors of response to therapy with terlipressin and albumin in patients with cirrhosis and type 1 hepatorenal syndrome. *Hepatology* 2010;51:219–226.
- [378] Brensing KA, Textor J, Perz J, Schiedermaier P, Raab P, Strunk H, et al. Long term outcome after transjugular intrahepatic portosystemic stent-shunt in non-transplant cirrhotics with hepatorenal syndrome: a phase II study. *Gut* 2000;47:288–295.
- [379] Guevara M, Gines P, Bandi JC, Gilabert R, Sort P, Jimenez W, et al. Transjugular intrahepatic portosystemic shunt in hepatorenal syndrome: effects on renal function and vasoactive systems. *Hepatology* 1998;28:416–422.
- [380] Testino G, Ferro C, Sumberaz A, Messa P, Morelli N, Guadagni B, et al. Type-2 hepatorenal syndrome and refractory ascites: role of transjugular intrahepatic portosystemic stent-shunt in eighteen patients with advanced cirrhosis awaiting orthotopic liver transplantation. *Hepato-gastroenterology* 2003;50:1753–1755.
- [381] Keller F, Heinze H, Jochimsen F, Passfall J, Schuppan D, Buttner P. Risk factors and outcome of 107 patients with decompensated liver disease and acute renal failure (including 26 patients with hepatorenal syndrome): the role of hemodialysis. *Ren Fail* 1995;17:135–146.
- [382] Sourinayanayane A, Raina R, Garg G, McCullough AJ, O'Shea RS. Management and outcome in hepatorenal syndrome: need for renal replacement therapy in non-transplanted patients. *Int Urol Nephrol* 2014;46:793–800.

- [383] Stauffer K, Roedel K, Kivaranovic D, Drolz A, Horvatits T, Rasoul-Rockenschaub S, et al. Renal replacement therapy in critically ill liver cirrhotic patients-outcome and clinical implications. *Liver Int* 2017;37:843–850.
- [384] Wu VC, Ko WJ, Chang HW, Chen YS, Chen YW, Chen YM, et al. Early renal replacement therapy in patients with postoperative acute liver failure associated with acute renal failure: effect on postoperative outcomes. *J Am Coll Surg* 2007;205:266–276.
- [385] Gaudry S, Hajage D, Schortgen F, Martin-Lefevre L, Pons B, Boulet E, et al. Initiation strategies for renal-replacement therapy in the intensive care unit. *N Engl J Med* 2016;375:122–133.
- [386] Zarbock A, Kellum JA, Schmidt C, Van Aken H, Wempe C, Pavenstadt H, et al. Effect of early vs delayed initiation of renal replacement therapy on mortality in critically ill patients with acute kidney injury: the elain randomized clinical trial. *JAMA* 2016;315:2190–2199.
- [387] Wong LP, Blackley MP, Andreoni KA, Chin H, Falk RJ, Klemmer PJ. Survival of liver transplant candidates with acute renal failure receiving renal replacement therapy. *Kidney Int* 2005;68:362–370.
- [388] Banares R, Nevens F, Larsen FS, Jalan R, Albillos A, Dollinger M, et al. Extracorporeal albumin dialysis with the molecular adsorbent recirculating system in acute-on-chronic liver failure: the RELIEF trial. *Hepatology* 2013;57:1153–1162.
- [389] Kribben A, Gerken G, Haag S, Herget-Rosenthal S, Treichel U, Betz C, et al. Effects of fractionated plasma separation and adsorption on survival in patients with acute-on-chronic liver failure. *Gastroenterology* 2012;142:782–789.
- [390] Boyer TD, Sanyal AJ, Garcia-Tsao G, Regenstein F, Rossaro L, Appenrodt B, et al. Impact of liver transplantation on the survival of patients treated for hepatorenal syndrome type 1. *Liver Transpl* 2011;17:1328–1332.
- [391] Gonwa TA, Klintmalm GB, Levy M, Jennings LS, Goldstein RM, Husberg BS. Impact of pretransplant renal function on survival after liver transplantation. *Transplantation* 1995;59:361–365.
- [392] Nadim MK, Sung RS, Davis CL, Andreoni KA, Biggins SW, Danovitch GM, et al. Simultaneous liver-kidney transplantation summit: current state and future directions. *Am J Transplant* 2012;12:2901–2908.
- [393] Francoz C, Nadim MK, Durand F. Kidney biomarkers in cirrhosis. *J Hepatol* 2016;65:809–824.
- [394] Angeli P, Gines P. Hepatorenal syndrome, MELD score and liver transplantation: an evolving issue with relevant implications for clinical practice. *J Hepatol* 2012;57:1135–1140.
- [395] Cillo U, Burra P, Mazzaferro V, Belli L, Pinna AD, Spada M, et al. A multistep, consensus-based approach to organ allocation in liver transplantation: toward a “blended principle model”. *Am J Transplant* 2015;15:2552–2561.
- [396] Akriviadis E, Botla R, Briggs W, Han S, Reynolds T, Shakil O. Pentoxifylline improves short-term survival in severe acute alcoholic hepatitis: a double-blind, placebo-controlled trial. *Gastroenterology* 2000;119:1637–1648.
- [397] Mathurin P, Louvet A, Duhamel A, Nahon P, Carbonell N, Boursier J, et al. Prednisolone with vs without pentoxifylline and survival of patients with severe alcoholic hepatitis: a randomized clinical trial. *JAMA* 2013;310:1033–1041.
- [398] Parker R, Armstrong MJ, Corbett C, Rowe IA, Houlihan DD. Systematic review: pentoxifylline for the treatment of severe alcoholic hepatitis. *Aliment Pharmacol Ther* 2013;37:845–854.
- [399] Allen AM, Kim WR. Epidemiology and healthcare burden of acute-on-chronic liver failure. *Semin Liver Dis* 2016;36:123–126.
- [400] Piano S, Tonon M, Vettore E, Stanco M, Pilutti C, Romano A, et al. Incidence, predictors and outcomes of acute-on-chronic liver failure in outpatients with cirrhosis. *J Hepatol* 2017;67:1177–1184.
- [401] Arroyo V, Jalan R. Acute-on-chronic liver failure: definition, diagnosis, and clinical characteristics. *Semin Liver Dis* 2016;36:109–116.
- [402] Sarin SK, Kedarisetty CK, Abbas Z, Amarapurkar D, Bihari C, Chan AC, et al. Acute-on-chronic liver failure: consensus recommendations of the Asian Pacific Association for the Study of the Liver (APASL) 2014. *Hepatol Int* 2014;8:453–471.
- [403] Jalan R, Yurdaydin C, Bajaj JS, Acharya SK, Arroyo V, Lin HC, et al. Toward an improved definition of acute on chronic liver failure. *Gastroenterology* 2014;147:4–10.
- [404] Bajaj JS. Defining acute-on-chronic liver failure: will East and West ever meet? *Gastroenterology* 2013;144:1337–1339.
- [405] Kim TY, Song DS, Kim HY, Sinn DH, Yoon EL, Kim CW, et al. Characteristics and discrepancies in acute-on-chronic liver failure: need for a unified definition. *PLoS One* 2016;11:e0146745.
- [406] Arroyo V, Moreau R, Jalan R, Gines P. Acute-on-chronic liver failure: A new syndrome that will re-classify cirrhosis. *J Hepatol* 2015;62: S131–S143.
- [407] Jalan R, Pavesi M, Saliba F, Amoros A, Fernandez J, Holland-Fischer P, et al. The CLIF Consortium Acute Decompensation score (CLIF-C ADs) for prognosis of hospitalised cirrhotic patients without acute-on-chronic liver failure. *J Hepatol* 2015;62:831–840.
- [408] Arroyo V, Moreau R, Kamath PS, Jalan R, Gines P, Nevens F, et al. Acute-on-chronic liver failure in cirrhosis. *Nat Rev Dis Primers* 2016;2:16041.
- [409] Trebicka J. Predisposing factors in acute-on-chronic liver failure. *Semin Liver Dis* 2016;36:167–173.
- [410] Sarin SK, Choudhury A. Acute-on-chronic liver failure. *Curr Gastroenterol Rep* 2016;18:61.
- [411] Bajaj JS, O’Leary JG, Reddy KR, Wong F, Biggins SW, Patton H, et al. Survival in infection-related acute-on-chronic liver failure is defined by extrahepatic organ failures. *Hepatology* 2014;60:250–256.
- [412] Silva PE, Fayad L, Lazzarotto C, Ronsoni MF, Bazzo ML, Colombo BS, et al. Single-centre validation of the EASL-CLIF consortium definition of acute-on-chronic liver failure and CLIF-SOFA for prediction of mortality in cirrhosis. *Liver Int* 2015;35:1516–1523.
- [413] Li H, Chen LY, Zhang NN, Li ST, Zeng B, Pavesi M, et al. Characteristics, diagnosis and prognosis of acute-on-chronic liver failure in cirrhosis associated to hepatitis B. *Sci Rep* 2016;6:25487.
- [414] Shalimar, Saraswat V, Singh SP, Duseja A, Shukla A, Eapen CE, et al. Acute-on-chronic liver failure in India: The Indian National Association for Study of the Liver consortium experience. *J Gastroenterol Hepatol* 2016;31:1742–1749.
- [415] Shi Y, Yang Y, Hu Y, Wu W, Yang Q, Zheng M, et al. Acute-on-chronic liver failure precipitated by hepatic injury is distinct from that precipitated by extrahepatic insults. *Hepatology* 2015;62:232–242.
- [416] Shalimar, Kumar D, Vadiraja PK, Nayak B, Thakur B, Das P, et al. Acute on chronic liver failure because of acute hepatic insults: Etiologies, course, extrahepatic organ failure and predictors of mortality. *J Gastroenterol Hepatol* 2016;31:856–864.
- [417] Dhiman RK, Agrawal S, Gupta T, Duseja A, Chawla Y. Chronic liver failure-sequential organ failure assessment is better than the asia-pacific association for the study of liver criteria for defining acute-on-chronic liver failure and predicting outcome. *World J Gastroenterol* 2014;20:14934–14941.
- [418] Pischke S, Suneetha PV, Baechlein C, Barg-Hock H, Heim A, Kamar N, et al. Hepatitis E virus infection as a cause of graft hepatitis in liver transplant recipients. *Liver Transpl* 2010;16:74–82.
- [419] Jalan R, Saliba F, Pavesi M, Amoros A, Moreau R, Gines P, et al. Development and validation of a prognostic score to predict mortality in patients with acute-on-chronic liver failure. *J Hepatol* 2014;61: 1038–1047.
- [420] McPhail MJ, Shawcross DL, Abeles RD, Chang A, Patel V, Lee GH, et al. Increased survival for patients with cirrhosis and organ failure in liver intensive care and validation of the chronic liver failure-sequential organ failure scoring system. *Clin Gastroenterol Hepatol* 2015;13: 1353–1360.
- [421] Lee M, Lee JH, Oh S, Jang Y, Lee W, Lee HJ, et al. CLIF-SOFA scoring system accurately predicts short-term mortality in acutely decompensated patients with alcoholic cirrhosis: a retrospective analysis. *Liver Int* 2015;35:46–57.
- [422] O’Leary JG, Reddy KR, Garcia-Tsao G, Biggins SW, Wong F, Fallon MB, et al. NACSELD acute-on-chronic liver failure (NACSELD-ACLF) score predicts 30-day survival in hospitalized patients with cirrhosis. *Hepatology* 2018.
- [423] Choudhury A, Jindal A, Maiwall R, Sharma MK, Sharma BC, Pamecha V, et al. APASL ACLF Working Party Liver failure determines the outcome in patients of acute-on-chronic liver failure (ACLF): comparison of APASL ACLF research consortium (AARC) and CLIF-SOFA models. *Hepatol Int* 2017;11:461–471.
- [424] Hernaez R, Sola E, Moreau R, Gines P. Acute-on-chronic liver failure: an update. *Gut* 2017;66:541–553.
- [425] Gustot T, Fernandez J, Garcia E, Morando F, Caraceni P, Alessandria C, et al. Clinical Course of acute-on-chronic liver failure syndrome and effects on prognosis. *Hepatology* 2015;62:243–252.
- [426] Garg H, Sarin SK, Kumar M, Garg V, Sharma BC, Kumar A. Tenofovir improves the outcome in patients with spontaneous reactivation of hepatitis B presenting as acute-on-chronic liver failure. *Hepatology* 2011;53:774–780.
- [427] Lin B, Pan CQ, Xie D, Xie J, Xie S, Zhang X, et al. Entecavir improves the outcome of acute-on-chronic liver failure due to the acute exacerbation of chronic hepatitis B. *Hepatol Int* 2013;7:460–467.

Clinical Practice Guidelines

- [428] Zhang Y, Hu XY, Zhong S, Yang F, Zhou TY, Chen G, et al. Entecavir vs lamivudine therapy for naive patients with spontaneous reactivation of hepatitis B presenting as acute-on-chronic liver failure. *World J Gastroenterol* 2014;20:4745–4752.
- [429] Xiang-Hui Y, Lang X, Yan Z, Li Z, Xiao-Feng S, Hong R. Prediction of prognosis to lamivudine in patients with spontaneous reactivation of hepatitis B virus-related acute-on-chronic liver failure: using virologic response at week 4. *Eur J Intern Med* 2014;25:860–864.
- [430] Chen JF, Wang KW, Zhang SQ, Lei ZY, Xie JQ, Zhu JY, et al. Dexamethasone in outcome of patients with hepatitis B virus-related acute-on-chronic liver failure. *J Gastroenterol Hepatol* 2014;29:800–806.
- [431] Guo YM, Li FY, Gong M, Zhang L, Wang JB, Xiao XH, et al. Short-term efficacy of treating hepatitis B virus-related acute-on-chronic liver failure based on cold pattern differentiation with hot herbs: A randomized controlled trial. *Chin J Integr Med* 2016;22:573–580.
- [432] Garg V, Garg H, Khan A, Trehanpati N, Kumar A, Sharma BC, et al. Granulocyte colony-stimulating factor mobilizes CD34(+) cells and improves survival of patients with acute-on-chronic liver failure. *Gastroenterology* 2012;142:505–512.
- [433] Finkenstedt A, Nachbaur K, Zollner H, Joannidis M, Pratschke J, Graziadei IW, et al. Acute-on-chronic liver failure: excellent outcome after liver transplantation but high mortality on the wait list. *Liver Transpl* 2013;19:879–886.
- [434] Artru F, Louvet A, Ruiz I, Levesque E, Labreuche J, Ursic-Bedoya J, et al. Liver transplantation in the most severely ill cirrhotic patients: A multicenter study in acute-on-chronic liver failure grade 3. *J Hepatol* 2017;67:708–715.
- [435] Bouachour G, Tiro P, Gouello JP, Mathieu E, Vincent JF, Alquier P. Adrenocortical function during septic shock. *Intensive Care Med* 1995;21:57–62.
- [436] Marik PE, Pastores SM, Annane D, Meduri GU, Sprung CL, Arlt W, et al. Recommendations for the diagnosis and management of corticosteroid insufficiency in critically ill adult patients: consensus statements from an international task force by the American College of Critical Care Medicine. *Crit Care Med* 2008;36:1937–1949.
- [437] Tsai MH, Peng YS, Chen YC, Liu NJ, Ho YP, Fang JT, et al. Adrenal insufficiency in patients with cirrhosis, severe sepsis and septic shock. *Hepatology* 2006;43:673–681.
- [438] Acevedo J, Fernandez J, Prado V, Silva A, Castro M, Pavesi M, et al. Relative adrenal insufficiency in decompensated cirrhosis: Relationship to short-term risk of severe sepsis, hepatorenal syndrome, and death. *Hepatology* 2013;58:1757–1765.
- [439] Jang JY, Kim TY, Sohn JH, Lee TH, Jeong SW, Park EJ, et al. Relative adrenal insufficiency in chronic liver disease: its prevalence and effects on long-term mortality. *Aliment Pharmacol Ther* 2014;40:819–826.
- [440] Tsai MH, Huang HC, Peng YS, Chen YC, Tian YC, Yang CW, et al. Critical illness-related corticosteroid insufficiency in cirrhotic patients with acute gastroesophageal variceal bleeding: risk factors and association with outcome. *Crit Care Med* 2014;42:2546–2555.
- [441] Fede G, Spadaro L, Tomaselli T, Privitera G, Scicali R, Vasiopoulou P, et al. Comparison of total cortisol, free cortisol, and surrogate markers of free cortisol in diagnosis of adrenal insufficiency in patients with stable cirrhosis. *Clin Gastroenterol Hepatol* 2014;12:504–512.
- [442] Kim G, Huh JH, Lee KJ, Kim MY, Shim KY, Baik SK. Relative adrenal insufficiency in patients with cirrhosis: a systematic review and meta-analysis. *Dig Dis Sci* 2017;62:1067–1079.
- [443] Fede G, Spadaro L, Tomaselli T, Privitera G, Germani G, Tsochatzis E, et al. Adrenocortical dysfunction in liver disease: a systematic review. *Hepatology* 2012;55:1282–1291.
- [444] McNeilly AD, Macfarlane DP, O'Flaherty E, Livingstone DE, Mitic T, McConnell KM, et al. Bile acids modulate glucocorticoid metabolism and the hypothalamic–pituitary–adrenal axis in obstructive jaundice. *J Hepatol* 2010;52:705–711.
- [445] Worlicek M, Knebel K, Linde HJ, Moleda L, Scholmerich J, Straub RH, et al. Splanchnic sympathectomy prevents translocation and spreading of *E coli* but not *S aureus* in liver cirrhosis. *Gut* 2010;59:1127–1134.
- [446] Cholongitas E, Goulis I, Pagkalidou E, Haidich AB, Karagiannis AKA, Nakouti T, et al. Relative adrenal insufficiency is associated with the clinical outcome in patients with stable decompensated cirrhosis. *Ann Hepatol* 2017;16:584–590.
- [447] Trifan A, Chiriac S, Stancu C. Update on adrenal insufficiency in patients with liver cirrhosis. *World J Gastroenterol* 2013;19:445–456.
- [448] Galbois A, Rudler M, Massard J, Fulla Y, Bennani A, Bonnefont-Rousselot D, et al. Assessment of adrenal function in cirrhotic patients: salivary cortisol should be preferred. *J Hepatol* 2010;52:839–845.
- [449] Hamrahian AH, Oseni TS, Arafah BM. Measurements of serum-free cortisol in critically ill patients. *N Engl J Med* 2004;350:1629–1638.
- [450] Tan T, Chang L, Woodward A, McWhinney B, Galligan J, Macdonald GA, et al. Characterising adrenal function using directly measured plasma free cortisol in stable severe liver disease. *J Hepatol* 2010;53:841–848.
- [451] Coolens JL, Van Baelen H, Heyns W. Clinical use of unbound plasma cortisol as calculated from total cortisol and corticosteroid-binding globulin. *J Steroid Biochem* 1987;26:197–202.
- [452] le Roux CW, Sivakumaran S, Alagband-Zadeh J, Dhillon W, Kong WM, Wheeler MJ. Free cortisol index as a surrogate marker for serum free cortisol. *Ann Clin Biochem* 2002;39:406–408.
- [453] Arafah BM, Nishiyama FJ, Tlaygeh H, Hejal R. Measurement of salivary cortisol concentration in the assessment of adrenal function in critically ill subjects: a surrogate marker of the circulating free cortisol. *J Clin Endocrinol Metab* 2007;92:2965–2971.
- [454] Thevenot T, Borot S, Remy-Martin A, Sapin R, Cervoni JP, Richou C, et al. Assessment of adrenal function in cirrhotic patients using concentration of serum-free and salivary cortisol. *Liver Int* 2011;31:425–433.
- [455] Fernandez J, Escorsell A, Zabalza M, Felipe V, Navasa M, Mas A, et al. Adrenal insufficiency in patients with cirrhosis and septic shock: Effect of treatment with hydrocortisone on survival. *Hepatology* 2006;44:1288–1295.
- [456] Arabi YM, Aljumah A, Dabbagh O, Tamim HM, Rishu AH, Al-Abdulka-reem A, et al. Low-dose hydrocortisone in patients with cirrhosis and septic shock: a randomized controlled trial. *CMAJ* 2010;182:1971–1977.
- [457] Moller S, Henriksen JH. Cirrhotic cardiomyopathy: a pathophysiological review of circulatory dysfunction in liver disease. *Heart* 2002;87:9–15.
- [458] Hunter JD, Doddi M. Sepsis and the heart. *Br J Anaesth* 2010;104:3–11.
- [459] Mehta G, Gustot T, Mookerjee RP, Garcia-Pagan JC, Fallon MB, Shah VH, et al. Inflammation and portal hypertension - the undiscovered country. *J Hepatol* 2014;61:155–163.
- [460] De BK, Majumdar D, Das D, Biswas PK, Mandal SK, Ray S, et al. Cardiac dysfunction in portal hypertension among patients with cirrhosis and non-cirrhotic portal fibrosis. *J Hepatol* 2003;39:315–319.
- [461] Karagiannakis DS, Vlachogiannakos J, Anastasiadis G, Vafiadis-Zouboulis I, Ladas SD. Diastolic cardiac dysfunction is a predictor of dismal prognosis in patients with liver cirrhosis. *Hepatol Int* 2014;8:588–594.
- [462] Rabie RN, Cazzaniga M, Salerno F, Wong F. The use of E/A ratio as a predictor of outcome in cirrhotic patients treated with transjugular intrahepatic portosystemic shunt. *Am J Gastroenterol* 2009;104:2458–2466.
- [463] Grose RD, Nolan J, Dillon JF, Errington M, Hannan WJ, Bouchier IA, et al. Exercise-induced left ventricular dysfunction in alcoholic and non-alcoholic cirrhosis. *J Hepatol* 1995;22:326–332.
- [464] Wong F, Girgrah N, Graba J, Allidina Y, Liu P, Blendis L. The cardiac response to exercise in cirrhosis. *Gut* 2001;49:268–275.
- [465] Bernardi M, Rubboli A, Trevisani F, Cancellieri C, Ligabue A, Baraldini M, et al. Reduced cardiovascular responsiveness to exercise-induced sympathoadrenergic stimulation in patients with cirrhosis. *J Hepatol* 1991;12:207–216.
- [466] Kim MY, Baik SK, Won CS, Park HJ, Jeon HK, Hong HI, et al. Dobutamine stress echocardiography for evaluating cirrhotic cardiomyopathy in liver cirrhosis. *Korean J Hepatol* 2010;16:376–382.
- [467] Krag A, Bendtsen F, Dahl EK, Kjaer A, Petersen CL, Moller S. Cardiac function in patients with early cirrhosis during maximal beta-adrenergic drive: a dobutamine stress study. *PLoS One* 2014;9:e109179.
- [468] Stanton T, Leano R, Marwick TH. Prediction of all-cause mortality from global longitudinal speckle strain: comparison with ejection fraction and wall motion scoring. *Circ Cardiovasc Imaging* 2009;2:356–364.
- [469] Lang RM, Bierig M, Devereux RB, Flachskampf FA, Foster E, Pellikka PA, et al. Recommendations for chamber quantification: a report from the American Society of Echocardiography's Guidelines and Standards Committee and the Chamber Quantification Writing Group, developed in conjunction with the European Association of Echocardiography, a branch of the European Society of Cardiology. *J Am Soc Echocardiogr* 2005;18:1440–1463.
- [470] Stampehl MR, Mann DL, Nguyen JS, Cota F, Colmenares C, Dokainish H. Speckle strain echocardiography predicts outcome in patients with heart failure with both depressed and preserved left ventricular ejection fraction. *Echocardiography* 2015;32:71–78.
- [471] Chen Y, Chan AC, Chok SH, Sharr W, Fung J, et al. A detailed evaluation of cardiac function in cirrhotic patients and its alteration with or without liver transplantation. *J Cardiol* 2016;67:140–146.
- [472] Sampaio F, Pimenta J, Bettencourt N, Fontes-Carvalho R, Silva AP, Valente J, et al. Systolic and diastolic dysfunction in cirrhosis: a tissue-

- Doppler and speckle tracking echocardiography study. *Liver Int* 2013;33:1158–1165.
- [473] Nazar A, Guevara M, Sitges M, Terra C, Sola E, Guigou C, et al. LEFT ventricular function assessed by echocardiography in cirrhosis: relationship to systemic hemodynamics and renal dysfunction. *J Hepatol* 2013;58:51–57.
- [474] Cesari M, Fasolato S, Rosi S, Angeli P. Cardiac dysfunction in patients with cirrhosis: is the systolic component its main feature? *Eur J Gastroenterol Hepatol* 2015;27:660–666.
- [475] Nagueh SF, Smiseth OA, Appleton CP, Byrd 3rd BF, Dokainish H, Edvardson T, et al. Recommendations for the evaluation of left ventricular diastolic function by echocardiography: an update from the American society of echocardiography and the European association of cardiovascular imaging. *Eur Heart J Cardiovasc Imaging* 2016;17:1321–1360.
- [476] Takemoto Y, Barnes ME, Seward JB, Lester SJ, Appleton CA, Gersh BJ, et al. Usefulness of left atrial volume in predicting first congestive heart failure in patients > or = 65 years of age with well-preserved left ventricular systolic function. *Am J Cardiol* 2005;96:832–836.
- [477] Valeriano V, Funaro S, Lionetti R, Riggio O, Pulcinelli G, Fiore P, et al. Modification of cardiac function in cirrhotic patients with and without ascites. *Am J Gastroenterol* 2000;95:3200–3205.
- [478] Pozzi M, Carugo S, Boari G, Pecci V, de Ceglia S, Maggolini S, et al. Evidence of functional and structural cardiac abnormalities in cirrhotic patients with and without ascites. *Hepatology* 1997;26:1131–1137.
- [479] Cazzaniga M, Salerno F, Pagnozzi G, Dionigi E, Visentin S, Cirello I, et al. Diastolic dysfunction is associated with poor survival in patients with cirrhosis with transjugular intrahepatic portosystemic shunt. *Gut* 2007;56:869–875.
- [480] Ruiz-del-Arbol L, Achecar L, Serradilla R, Rodriguez-Gandia MA, Rivero M, Garrido E, et al. Diastolic dysfunction is a predictor of poor outcomes in patients with cirrhosis, portal hypertension, and a normal creatinine. *Hepatology* 2013;58:1732–1741.
- [481] Cesari M, Frigo AC, Tonon M, Angeli P. Cardiovascular predictors of death in patients with cirrhosis. *Hepatology* 2017.
- [482] Merli M, Torromeo C, Giusto M, Iacovone G, Riggio O, Puddu PE. Survival at 2 years among liver cirrhotic patients is influenced by left atrial volume and left ventricular mass. *Liver Int* 2017;37:700–706.
- [483] Kozor R, Nordin S, Treibel TA, Rosmini S, Castelletti S, Fontana M, et al. Insight into hypertrophied hearts: a cardiovascular magnetic resonance study of papillary muscle mass and T1 mapping. *Eur Heart J Cardiovasc Imaging* 2016;18:1034–1040.
- [484] Ngu PJ, Butler M, Pham A, Roberts SK, Taylor AJ. Cardiac remodelling identified by cardiovascular magnetic resonance in patients with hepatitis C infection and liver disease. *Int J Cardiovasc Imaging* 2016;32:629–636.
- [485] Ruiz-del-Arbol L, Monescillo A, Arocena C, Valer P, Gines P, Moreira V, et al. Circulatory function and hepatorenal syndrome in cirrhosis. *Hepatology* 2005;42:439–447.
- [486] Zhang W, Xu X, Kao R, Mele T, Kvietys P, Martin CM, et al. Cardiac fibroblasts contribute to myocardial dysfunction in mice with sepsis: the role of NLRP3 inflammasome activation. *PLoS One* 2014;9:e107639.
- [487] Gaskari SA, Liu H, D'Mello C, Kunos G, Lee SS. Blunted cardiac response to hemorrhage in cirrhotic rats is mediated by local macrophage-released endocannabinoids. *J Hepatol* 2015;62:1272–1277.
- [488] Liu H, Lee SS. Nuclear factor- κ B inhibition improves myocardial contractility in rats with cirrhotic cardiomyopathy. *Liver Int* 2008;28:640–648.
- [489] Trevisani F, Di Micoli A, Zambruni A, Biselli M, Santi V, Erroi V, et al. QT interval prolongation by acute gastrointestinal bleeding in patients with cirrhosis. *Liver Int* 2012;32:1510–1515.
- [490] Zhao J, Qi X, Hou F, Ning Z, Zhang X, Deng H, et al. Prevalence, risk factors and in-hospital outcomes of QTc interval prolongation in liver cirrhosis. *Am J Med Sci* 2016;352:285–295.
- [491] Krag A, Bendtsen F, Mortensen C, Henriksen JH, Moller S. Effects of a single terlipressin administration on cardiac function and perfusion in cirrhosis. *Eur J Gastroenterol Hepatol* 2010;22:1085–1092.
- [492] Wannhoff A, Hippchen T, Weiss CS, Friedrich K, Rupp C, Neumann-Haefelin C, et al. Cardiac volume overload and pulmonary hypertension in long-term follow-up of patients with a transjugular intrahepatic portosystemic shunt. *Aliment Pharmacol Ther* 2016;43:955–965.
- [493] Busk TM, Bendtsen F, Henriksen JH, Fuglsang S, Clemmesen JO, Larsen FS, et al. Effects of transjugular intrahepatic portosystemic shunt (TIPS) on blood volume distribution in patients with cirrhosis. *Dig Liver Dis* 2017;49:1353–1359.
- [494] Raevens S, De Pauw M, Geerts A, Berrevoet F, Rogiers X, Troisi RI, et al. Prevalence and outcome of diastolic dysfunction in liver transplantation recipients. *Acta Cardiol* 2014;69:273–280.
- [495] Saner FH, Neumann T, Canbay A, Treckmann JW, Hartmann M, Goerlinger K, et al. High brain-natriuretic peptide level predicts cirrhotic cardiomyopathy in liver transplant patients. *Transpl Int* 2011;24:425–432.
- [496] Sampaio F, Pimenta J, Bettencourt N, Fontes-Carvalho R, Silva AP, Valente J, et al. Systolic dysfunction and diastolic dysfunction do not influence medium-term prognosis in patients with cirrhosis. *Eur J Intern Med* 2014;25:241–246.
- [497] Alexopoulou A, Papatheodoridis G, Pouriki S, Chrysohoou C, Raftopoulos L, Stefanadis C, et al. Diastolic myocardial dysfunction does not affect survival in patients with cirrhosis. *Transpl Int* 2012;25:1174–1181.
- [498] Rodriguez-Roisin R, Krowka MJ, Herve P, Fallon MB. ERS task force pulmonary-hepatic vascular disorders (PHD) scientific committee. Pulmonary-hepatic vascular disorders (PHD). *Eur Respir J* 2004;24:861–880.
- [499] Machicao VI, Balakrishnan M, Fallon MB. Pulmonary complications in chronic liver disease. *Hepatology* 2014;59:1627–1637.
- [500] Kaymakoglu S, Kahraman T, Kudat H, Demir K, Cakaloglu Y, Adalet I, et al. Hepatopulmonary syndrome in noncirrhotic portal hypertensive patients. *Dig Dis Sci* 2003;48:556–560.
- [501] Fuhrmann V, Madl C, Mueller C, Holzinger U, Kitzberger R, Funk GC, et al. Hepatopulmonary syndrome in patients with hypoxic hepatitis. *Gastroenterology* 2006;131:69–75.
- [502] Rodríguez-Roisin R, Agustí AG, Roca J. The hepatopulmonary syndrome: new name, old complexities. *Thorax* 1992;47:897–902.
- [503] Fallon MB, Abrams GA. Pulmonary dysfunction in chronic liver disease. *Hepatology* 2000;32:859–865.
- [504] Voiosu AM, Daha IC, Voiosu TA, Mateescu BR, Dan GA, Băicuș CR, et al. Prevalence and impact on survival of hepatopulmonary syndrome and cirrhotic cardiomyopathy in a cohort of cirrhotic patients. *Liver Int* 2015;35:2547–2555.
- [505] Rodríguez-Roisin R, Krowka MJ. Hepatopulmonary syndrome; a liver-induced lung vascular disorder. *N Engl J Med* 2008;358:2378–2387.
- [506] Schenk P, Schoniger-Hekele M, Fuhrmann V, Madl C, Silberhumer G, Muller C. Prognostic significance of the hepatopulmonary syndrome in patients with cirrhosis. *Gastroenterology* 2003;125:1042–1052.
- [507] Swanson KL, Wiesner RH, Krowka MJ. Natural history of hepatopulmonary syndrome: impact of liver transplantation. *Hepatology* 2005;41:1122–1129.
- [508] Fallon MB, Krowka MJ, Brown RS, Trotter JF, Zacks S, Roberts KE, et al. Impact of hepatopulmonary syndrome on quality of life and survival in liver transplant candidates. *Gastroenterology* 2008;135:1168–1175.
- [509] Schraufnagel DE, Kay JM. Structural and pathologic changes in the lung vasculature in chronic liver disease. *Clin Chest Med* 2006;17:1–15.
- [510] Boryczka G, Hartleb M, Rudzki K, Janik MA. Influence of an upright body position on the size of intrapulmonary blood shunts in patients with advanced liver cirrhosis. *J Physiol Pharmacol* 2015;66:855–861.
- [511] Fallon MB, Abrams GA, Luo B, Hou Z, Dai J, Ku DD. The role of endothelial nitric oxide synthase in the pathogenesis of a rat model of hepatopulmonary syndrome. *Gastroenterology* 1997;113:606–614.
- [512] Luo B, Liu L, Tang L, Zhang J, Ling Y, Fallon MB. ET-1 and TNF- α in HPS: analysis in prehepatic portal hypertension and biliary and nonbiliary cirrhosis in rats. *Am J Physiol Gastrointest Liver Physiol* 2004;286:G294–G303.
- [513] Wiest R, Groszmann RJ. The paradox of nitric oxide in cirrhosis and portal hypertension: too much, not enough. *Hepatology* 2002;35:478–491.
- [514] Ling Y, Zhang J, Luo B, Song D, Liu L, Tang L, et al. The role of endothelin-1 and the endothelin B receptor in the pathogenesis of hepatopulmonary syndrome in the rat. *Hepatology* 2004;39:1593–1602.
- [515] Tang L, Luo B, Patel RP, Ling Y, Zhang J, Fallon MB. Modulation of pulmonary endothelial endothelin B receptor expression and signaling: implications for experimental hepatopulmonary syndrome. *Am J Physiol Lung Cell Mol Physiol* 2007;292:L1467–L1472.
- [516] Frossard JL, Schiffer E, Cikirikcioglu B, Bourquin BJ, Morel DR, Pastor MC. Opposite regulation of endothelial NO synthase by HSP90 and caveolin in liver and lungs of rats with hepatopulmonary syndrome. *Am J Physiol Gastrointest Liver Physiol* 2007;293:G864–G870.
- [517] Rabiller A, Nunes H, Lebrec D, Tazi KA, Wartski M, Dulmet E, et al. Prevention of gram-negative translocation reduces the severity of hepatopulmonary syndrome. *Am J Respir Crit Care Med* 2002;166:514–517.

Clinical Practice Guidelines

- [518] Thenappan T, Goel A, Marsboom G, Fang YH, Toth PT, Zhang HJ, et al. A central role for CD68(1) macrophages in hepatopulmonary syndrome: reversal by macrophage depletion. *Am J Respir Crit Care Med* 2011;183:1080–1091.
- [519] Zhang J, Yang W, Luo B, Hu B, Maheshwari A, Fallon MB. The role of CX(3)CL1/CX(3)CR1 in pulmonary angiogenesis and intravascular monocyte accumulation in rat experimental hepatopulmonary syndrome. *J Hepatol* 2012;57:752–758.
- [520] Carter EP, Hartsfield CL, Miyazono M, Jakkula M, Morris Jr KG, McMurtry IF. Regulation of heme oxygenase-1 by nitric oxide during hepatopulmonary syndrome. *Am J Physiol Lung Cell Mol Physiol* 2002;283:L346–L353.
- [521] Zhang J, Luo B, Tang L, Wang Y, Stockard CR, Kadish I, et al. Pulmonary angiogenesis in a rat model of hepatopulmonary syndrome. *Gastroenterology* 2009;136:1070–1080.
- [522] Zhang J, Yang W, Hu B, Wu W, Fallon MB. Endothelin-1 activation of the endothelin B receptor modulates pulmonary endothelial CX3CL1 and contributes to pulmonary angiogenesis in experimental hepatopulmonary syndrome. *Am J Pathol* 2014;184:1706–1714.
- [523] Zeng J, Chen L, Chen B, Lu K, Belguise K, Wang X, et al. MicroRNA-199 a-5p Regulates the Proliferation of Pulmonary Microvascular Endothelial Cells in Hepatopulmonary Syndrome. *Cell Physiol Biochem* 2015;37:1289–1300.
- [524] Roberts KE, Kawut SM, Krowka MJ, Brown Jr RS, Trotter JF, Shah V, et al. Genetic risk factors for hepatopulmonary syndrome in patients with advanced liver disease. *Gastroenterology* 2010;39:130–139.
- [525] Chang CC, Wang SS, Hsieh HG, Lee WS, Chuang CL, Hc Lin, et al. Rosuvastatin improves hepatopulmonary syndrome through inhibition of inflammatory angiogenesis of lung. *Clin Sci* 2015;129:449–460.
- [526] Rodriguez-Roisin R, Krowka MJ, Herve P, Fallon MB. ERS (European Respiratory Society) Task Force/PHD Scientific Committee. Highlights of the ERS task force on pulmonary-hepatic vascular disorders (PHD). *J Hepatol* 2005;42:924–927.
- [527] Arguedas MR, Singh H, Faulk DK, Fallon MB. Utility of pulse oximetry screening for hepatopulmonary syndrome. *Clin Gastroenterol Hepatol* 2007;5:749–754.
- [528] Hoerning A, Raub S, Neudorf U, Muntjes C, Kathemann S, Lainka E, et al. Pulse oximetry is insufficient for timely diagnosis of hepatopulmonary syndrome in children with liver cirrhosis. *J Pediatr* 2014;164:546–552.
- [529] Horvatis T, Drolz A, Roedel K, Herkner H, Ferlitsch A, Perkmann T, et al. Von Willebrand factor antigen for detection of hepatopulmonary syndrome in patients with cirrhosis. *J Hepatol* 2014;61:544–549.
- [530] Köksal D, Kaçar S, Köksal AS, Tüfekçioğlu O, Küçükay F, Okten S, et al. Evaluation of intrapulmonary vascular dilatations with high-resolution computed thorax tomography in patients with hepatopulmonary syndrome. *J Clin Gastroenterol* 2006;40:77–83.
- [531] Lee KN, Lee HJ, Shin WW, Webb WR. Hypoxemia and liver cirrhosis (hepatopulmonary syndrome) in eight patients: comparison of the central and peripheral pulmonary vasculature. *Radiology* 1999;211:549–553.
- [532] Abrams GA, Jaffe CC, Hoffer PB, Binder HJ, Fallon MB. Diagnostic utility of contrast echocardiography and lung perfusion scan in patients with hepatopulmonary syndrome. *Gastroenterology* 1995;109:1283–1288.
- [533] Wolfe JD, Tashkin DP, Holly FE, Brachman MB, Genovesi MG. Hypoxemia of cirrhosis: detection of abnormal small pulmonary vascular channels by a quantitative radionuclide method. *Am J Med* 1977;63:746–754.
- [534] Abrams GA, Nanda NC, Dubovsky EV, Krowka MJ, Fallon MB. Use of macroaggregated albumin lung perfusion scan to diagnose hepatopulmonary syndrome: a new approach. *Gastroenterology* 1998;114:305–310.
- [535] Arguedas MR, Abrams GA, Krowka MJ, Fallon MB. Prospective evaluation of outcomes and predictors of mortality in patients with hepatopulmonary syndrome undergoing liver transplantation. *Hepatology* 2003;37:192–197.
- [536] Kalambokis G, Tsianos EV. Pitfalls in the assessment of intrapulmonary shunt using lung perfusion scintigraphy in patients with cirrhosis. *Liver Int* 2010;31:138–139.
- [537] Kochar R, Tanikella R, Fallon MB. Serial pulse oximetry in hepatopulmonary syndrome. *Dig Dis Sci* 2011;56:1862–1868.
- [538] Tanikella R, Philips GM, Faulk DK, Kawut SM, Fallon MB. Pilot study of pentoxifylline in hepatopulmonary syndrome. *Liver Transpl* 2008;14:1199–1203.
- [539] Gupta LB, Kumar A, Jaiswal AK, Yusuf J, Metha V, Tyagi S, et al. Pentoxifylline therapy for hepatopulmonary syndrome: a pilot study. *Arch Intern Med* 2008;168:1820–1823.
- [540] De BK, Dutta D, Pal SK, Gangopadhyay S, Das Baksi S, Pani A. The role of garlic in hepatopulmonary syndrome: a randomized controlled trial. *Can J Gastroenterol* 2010;24:183–188.
- [541] Shaikh SA, Tischer S, Choi EK, Fontana RJ. Good for the lung but bad for the liver? Garlic-induced hepatotoxicity following liver transplantation. *J Clin Pharm Ther* 2017;42:646–648.
- [542] Tsauo J, Weng N, Ma H, Jiang M, Zhao H, Li X. Role of transjugular intrahepatic portosystemic shunts in the management of hepatopulmonary syndrome: a systemic literature review. *J Vasc Interv Radiol* 2015;26:1266–1271.
- [543] Poterucha JJ, Krowka MJ, Dickson ER, Cortese DA, Stanson AW, Krom RA. Failure of hepatopulmonary syndrome to resolve after liver transplantation and successful treatment with embolotherapy. *Hepatology* 1995;21:96–100.
- [544] Krowka MJ, Mandell MS, Ramsay MA, Kawut SM, Fallon MB, Manzarbeitia C, et al. Hepatopulmonary syndrome and portopulmonary hypertension: a report of the multicenter liver transplant database. *Liver Transpl* 2004;10:174–182.
- [545] Gupta S, Castel H, Rao RV, Picard M, Lilly L, Faughnan ME, et al. Improved survival after liver transplantation in patients with hepatopulmonary syndrome. *Am J Transplant* 2010;10:354–363.
- [546] Fallon MB, Mulligan DC, Gish RG, Krowka MJ. Model for end-stage liver disease (MELD) exception for hepatopulmonary syndrome. *Liver Transpl* 2006;12:S105–S107.
- [547] Iyer VN, Swanson KL, Cartin-Ceba R, Dierkhising RA, Rosen CB, Heimbach JK, et al. Hepatopulmonary syndrome: favorable outcomes in the MELD exception era. *Hepatology* 2013;57:2427–2435.
- [548] Goldberg DS, Krok K, Batra S, Trotter JF, Kawut SM, Fallon MB. Impact of the hepatopulmonary syndrome MELD exception policy on outcomes of patients after liver transplantation: an analysis of the UNOS database. *Gastroenterology* 2014;146:1256–1265.
- [549] Fleming GM, Cornell TT, Welling TH, Magee JC, Annich GM. Hepatopulmonary syndrome use of extracorporeal life support for life-threatening hypoxia following liver transplantation. *Liver Transpl* 2008;14:966–970.
- [550] Monsel A, Mal H, Brisson H, Luo R, Eyraud D, Vénizet C, et al. Extracorporeal membrane oxygenation as a bridge to liver transplantation for acute respiratory distress syndrome-induced life-threatening hypoxemia aggravated by hepatopulmonary syndrome. *Crit Care* 2011;15:R234.
- [551] Chihara Y, Egawa H, Tsuboi T, Oga T, Handa T, Yamamoto K, et al. Immediate noninvasive ventilation may improve mortality in patients with hepatopulmonary syndrome after liver transplantation. *Liver Transpl* 2011;17:44–48.
- [552] McLaughlin VV, Archer SL, Badesch DB, Barst RJ, Farber HW, Lindner JR, et al. ACCF/AHA 2009 expert consensus document on pulmonary hypertension a report of the American College of Cardiology Foundation Task Force on Expert Consensus Documents and the American Heart Association developed in collaboration with the American College of Chest Physicians; American Thoracic Society, Inc.; and the Pulmonary Hypertension Association. *J Am Coll Cardiol* 2009;53:1573–1619.
- [553] Torregrosa M, Genesca J, Gonzalez A, Evangelista A, Mora A, Margarit C, et al. Role of Doppler echocardiography in the assessment of portopulmonary hypertension in liver transplantation candidates. *Transplantation* 2001;71:572–574.
- [554] Raevens S, Colle I, Reyntjens K, Geerts A, Berrevoet F, Rogiers X, et al. Echocardiography for the detection of portopulmonary hypertension in liver transplant candidates: an analysis of cutoff values. *Liver Transpl* 2013;19:602–610.
- [555] Cotton CL, Gandhi S, Vaitkus PT, Massad MG, Benedetti E, Mrtek RG, et al. Role of echocardiography in detecting portopulmonary hypertension in liver transplant candidates. *Liver Transpl* 2002;8:1051–1054.
- [556] Kawut SM, Krowka MJ, Trotter JF, Roberts KE, Benza RL, Badesch DB, et al. Clinical risk factors for portopulmonary hypertension. *Hepatology* 2008;48:196–203.
- [557] Paulus JK, Roberts KE. Oestrogen and sexual Dimorphism of pulmonary arterial hypertension; a transitional challenge. *Eur Respir J* 2013;41:1014–1016.
- [558] Talwalkar JA, Swanson KL, Krowka MJ, Andrews JC, Kamath PS. Prevalence of spontaneous portosystemic shunts in patients with portopulmonary hypertension and effect on treatment. *Gastroenterology* 2011;141:1673–1679.
- [559] Hadengue A, Benhayoun MK, Lebrec D, Benhamou JP. Pulmonary hypertension complicating portal hypertension: prevalence and relation to splanchnic hemodynamics. *Gastroenterology* 1991;100:520–528.

- [560] Robalino BD, Moodie DS. Association between primary pulmonary hypertension and portal hypertension: analysis of its pathophysiology and clinical, laboratory and hemodynamic manifestations. *J Am Coll Cardiol* 1991;17:492–498.
- [561] Swanson KL, Wiesner RH, Nyberg SL, Rosen CB, Krowka MJ. Survival in portopulmonary hypertension: Mayo Clinic experience categorized by treatment subgroups. *Am J Transplant* 2008;8:2445–2453.
- [562] Cartin-Ceba R, Krowka MJ. Portopulmonary hypertension. *Clin Liver Dis* 2014;18:421–438.
- [563] Condino AA, Ivy DD, O'Connor JA, Narkewicz MR, Mengshol S, Whitworth JR, et al. Portopulmonary hypertension in pediatric patients. *J Pediatr* 2005;147:20–26.
- [564] Le Pavec J, Souza R, Herve P, Lebrec D, Savale L, Tcherakian C, et al. Portopulmonary hypertension: survival and prognostic factors. *Am J Respir Crit Care Med* 2008;178:637–643.
- [565] Krowka MJ, Cartin-Ceba R. Portopulmonary hypertension: formidable dual threat vs. hopeful dual therapy. *Liver Transpl* 2014;20:635–636.
- [566] Kawut SM, Horn EM, Berekashvili KK, Garofano RP, Goldsmith RL, Widlitz AC, et al. New predictors of outcome in idiopathic pulmonary arterial hypertension. *Am J Cardiol* 2005;95:199–203.
- [567] Ricci GL, Melgosa MT, Burgos F, Valera JL, Pizarro S, Roca J, et al. Assessment of acute pulmonary vascular reactivity in portopulmonary hypertension. *Liver Transpl* 2007;13:1506–1514.
- [568] Provencher S, Herve P, Jais X, Lebrec D, Humbert M, Simonneau G, et al. Deleterious effects of beta-blockers on exercise capacity and hemodynamics in patients with portopulmonary hypertension. *Gastroenterology* 2006;130:120–126.
- [569] Halank M, Miehlke S, Hoeffken G, Schmeisser A, Schulze M, Strasser RH. Use of oral endothelin-receptor antagonist bosentan in the treatment of portopulmonary hypertension. *Transplantation* 2004;77:1775–1776.
- [570] Hoepfer MM, Halank M, Marx C, Hoeffken G, Seyfarth HJ, Schauer J, et al. Bosentan therapy for portopulmonary hypertension. *Eur Respir J* 2005;25:502–508.
- [571] Savale L, Magnier R, Le Pavec J, Jais X, Montani D, O'Callaghan DS, et al. Efficacy, safety and pharmacokinetics of bosentan in portopulmonary hypertension. *Eur Respir J* 2013;41:96–103.
- [572] Barth F, Gerber PJ, Reichen J, Dufour JF, Nicod LP. Efficiency and safety of bosentan in child C cirrhosis with portopulmonary hypertension and renal insufficiency. *Eur J Gastroenterol Hepatol* 2006;18:1117–1119.
- [573] Hoepfer MM, Seyfarth HJ, Hoeffken G, Wirtz H, Spiekerkoetter E, Pletz MW, et al. Experience with inhaled iloprost and bosentan in portopulmonary hypertension. *Eur Respir J* 2007;30:1096–1102.
- [574] Cartin-Ceba R, Swanson K, Iyer V, Wiesner RH, Krowka MJ. Safety and efficacy of ambrisentan for the treatment of portopulmonary hypertension. *Chest* 2011;139:109–114.
- [575] DuBrock HM, Channick RN, Krowka MJ. What's new in the treatment of portopulmonary hypertension? *Expert Rev Gastroenterol Hepatol* 2015;9:983–992.
- [576] Reichenberger F, Voswinckel R, Steveling E, Enke B, Kreckel A, Olschewski H, et al. Sildenafil treatment for portopulmonary hypertension. *Eur Respir J* 2006;28:563–567.
- [577] Cadden IS, Greanya ED, Erb SR, Scudamore CH, Yoshida EM. The use of sildenafil to treat portopulmonary hypertension prior to liver transplantation. *Ann Hepatol* 2009;8:158–161.
- [578] Fisher JH, Johnson SR, Chau C, Kron AT, Granton JT. Effectiveness of phosphodiesterase-5 inhibitor therapy for portopulmonary hypertension. *Can Respir J* 2015;22:42–46.
- [579] Tzathas C, Christidou A, Ladas SD. Sildenafil (viagra) is a risk factor for acute variceal bleeding. *Am J Gastroenterol* 2002;97:1856.
- [580] Kuo PC, Johnson LB, Plotkin JS, Howell CD, Bartlett ST, Rubin LJ. Continuous intravenous infusion of epoprostenol for the treatment of portopulmonary hypertension. *Transplantation* 1997;63:604–606.
- [581] Sussman N, Kaza V, Barshes N, Stribling R, Goss J, O'Mahony C, et al. Successful liver transplantation following medical management of portopulmonary hypertension: a single-center series. *Am J Transplant* 2006;6:2177–2182.
- [582] Fix OK, Bass NM, De Marco T, Merriman RB. Long-term follow-up of portopulmonary hypertension: effect of treatment with epoprostenol. *Liver Transpl* 2007;13:875–885.
- [583] Awdish RL, Cajigas HR. Early initiation of prostacyclin in portopulmonary hypertension: 10 years of a transplant center's experience. *Lung* 2013;191:593–600.
- [584] Melgosa MT, Ricci GL, Garcia-Pagan JC, Blanco I, Escribano P, Abalde JG, et al. Acute and long-term effects of inhaled iloprost in portopulmonary hypertension. *Liver Transpl* 2010;16:348–356.
- [585] Colombato LA, Spahr L, Martinet JP, Dufresne MP, Lafortune M, Fenyves D, et al. Haemodynamic adaptation two months after transjugular intrahepatic portosystemic shunt (TIPS) in cirrhotic patients. *Gut* 1996;39:600–604.
- [586] Boyer TD, Haskal ZJ. American Association for the Study of Liver D. The Role of Transjugular Intrahepatic Portosystemic Shunt (TIPS) in the Management of Portal Hypertension: update 2009. *Hepatology* 2010;51:306.
- [587] Yoshida EM, Erb SR, Pflugfelder PW, Ostrow DN, Ricci DR, Ghent CN, et al. Single-lung vs. liver transplantation for the treatment of portopulmonary hypertension—a comparison of two patients. *Transplantation* 1993;55:688–690.
- [588] Krowka MJ, Wiesner RH, Heimbach JK. Pulmonary contraindications, indications and MELD exceptions for liver transplantation: a contemporary view and look forward. *J Hepatol* 2013;59:367–374.
- [589] Kuo PC, Plotkin JS, Gaine S, Schroeder RA, Rustgi VK, Rubin LJ, et al. Portopulmonary hypertension and the liver transplant candidate. *Transplantation* 1999;67:1087–1093.
- [590] Krowka MJ, Plevak DJ, Findlay JY, Rosen CB, Wiesner RH, Krom RA. Pulmonary hemodynamics and perioperative cardiopulmonary-related mortality in patients with portopulmonary hypertension undergoing liver transplantation. *Liver Transpl* 2000;6:443–450.
- [591] Mangus RS, Kinsella SB, Marshall GR, Fridell JA, Wilkes KR, Tector AJ. Mild to moderate pulmonary hypertension in liver transplantation. *J Surg Res* 2013;184:1150–1156.
- [592] Raevens S, De Pauw M, Reyntjens K, Geerts A, Verhelst X, Berrevoet F, et al. Oral vasodilator therapy in patients with moderate to severe portopulmonary hypertension as a bridge to liver transplantation. *Eur J Gastroenterol Hepatol* 2013;25:495–502.
- [593] Ramsay MA, Simpson BR, Nguyen AT, Ramsay KJ, East C, Klintmalm GB. Severe pulmonary hypertension in liver transplant candidates. *Liver Transpl Surg* 1997;3:494–500.
- [594] Krowka MJ, Fallon MB, Mulligan DC, Gish RG. Model for end-stage liver disease (MELD) exception for portopulmonary hypertension. *Liver Transpl* 2006;12:S114–S116.
- [595] DuBrock HM, Goldberg DS, Sussman NL, Bartolome SD, Kadry Z, Salgia RJ, et al. Predictors of waitlist mortality in portopulmonary hypertension. *Transplantation* 2017;101:1609–1615.
- [596] Ramsay M. Portopulmonary hypertension and right heart failure in patients with cirrhosis. *Curr Opin Anaesthesiol* 2010;23:145–150.
- [597] Paulsen AW, Whitten CW, Ramsay MA, Klintmalm GB. Considerations for anesthetic management during veno-venous bypass in adult hepatic transplantation. *Anesth Analg* 1989;68:489–496.
- [598] Ramsay MA, Spikes C, East CA, Lynch K, Hein HA, Ramsay KJ, et al. The perioperative management of portopulmonary hypertension with nitric oxide and epoprostenol. *Anesthesiology* 1999;90:299–301.
- [599] Stratta C, Lavezzo B, Ballaris MA, Panio A, Crucitti M, Andruetto P, et al. Extracorporeal membrane oxygenation rescue therapy in a case of portopulmonary hypertension during liver transplantation: a case report. *Transplant Proc* 2013;45:2774–2775.
- [600] Ashfaq M, Chinnakotla S, Rogers L, Ausloos K, Saadeh S, Klintmalm GB, et al. The impact of treatment of portopulmonary hypertension on survival following liver transplantation. *Am J Transplant* 2007;7:1258–1264.
- [601] Hollatz TJ, Musat A, Westphal S, Decker C, D'Alessandro AM, Keevil J, et al. Treatment with sildenafil and treprostinil allows successful liver transplantation of patients with moderate to severe portopulmonary hypertension. *Liver Transpl* 2012;18:686–695.
- [602] Khaderi S, Khan R, Safdar Z, Stribling R, Vierling JM, Goss JA, et al. Long-term follow-up of portopulmonary hypertension patients after liver transplantation. *Liver Transpl* 2014;20:724–727.
- [603] Goldberg DS, Batra S, Sahay S, Kawut SM, Fallon MB. MELD exceptions for portopulmonary hypertension: current policy and future implementation. *Am J Transplant* 2014;14:2081–2087.
- [604] Morando F, Maresio G, Piano S, Fasolato S, Cavallin M, Romano A, et al. How to improve care in outpatients with cirrhosis and ascites: a new model of care coordination by consultant hepatologists. *J Hepatol* 2013;59:257–264.