Clinical history of liver cirrhosis is characterised by two phases: the asymptomatic phase, also termed ‘compensated cirrhosis’, and the phase of complications due to the development of portal hypertension and liver dysfunction, also termed ‘decompensated cirrhosis’, in which patients may develop ascites, the most frequent and clinically relevant complication of liver cirrhosis. Ascites can be classified into uncomplicated and complicated according to the development of refractoriness, spontaneous bacterial peritonitis (SBP) or the association with hepatorenal syndrome (HRS). In this narrative review, we will extensively discuss the optimal pharmacological and non-pharmacological management of cirrhotic ascites with the aim to offer an updated practical guide to Internal Medicine physicians. According to the amount of fluid in the abdominal cavity, uncomplicated ascites is graded from 1 to 3, and the cornerstone of its management consists of restriction of salt intake, diuretics and large-volume paracentesis (LVP); in recent years, long-term administration of human albumin has acquired a new interesting role. Refractory ascites is primarily managed with LVP and transjugular intrahepatic portosystemic shunt (TIPS) placement in selected patients. The occurrence of renal impairment, especially HRS, worsens the prognosis of patients with cirrhotic ascites and deserves a specific treatment. Also, the management of SBP faces the rising and alarming spread of antibiotic resistance. Hepatic hydrothorax may even complicate the course of the disease and its management is a challenge. Last but not least, liver transplantation (LT) is the ultimate and more effective measure to offer to patients with cirrhotic ascites, particularly when complications occur.

Key words: liver cirrhosis, ascites, hepatorenal syndrome, spontaneous bacterial peritonitis, hepatic hydrothorax, human albumin, narrative review

INTRODUCTION

Chronic liver disease may irreversibly lead to cirrhosis. Inflammation, fibrosis and regenerative necrotic events are the main underlying pathogenetic processes. Clinical history is characterised by two phases: the asymptomatic phase, also termed ‘compensated cirrhosis’, and the phase of complications due to the development of portal hypertension and liver dysfunction, also termed ‘decompensated cirrhosis’, in which patients may develop ascites, portal hypertensive gastrointestinal bleeding, encephalopathy, jaundice, hepatorenal syndrome (HRS), hepatopulmonary syndrome, spontaneous bacterial peritonitis (SBP) and/or hepatocellular carcinoma (HCC). Ascites is certainly the most frequent and clinically relevant complication. In this narrative review, we will extensively discuss updated evidence on the optimal pharmacological and non-pharmacological management of cirrhotic ascites.

PATHOPHYSIOLOGY OF CIRRHOTIC ASCITES

Ascites is the presence of fluid in the peritoneal cavity. Almost 60% of patients develop ascites within 10 years from the diagnosis of cirrhosis. Portal hypertension
In recent years, because no treatment has definitely been found effective and poor prognosis. Albumin is exclusively synthesised by hepatocytes and, therefore, liver function impairment leads to hypoalbuminaemia; however, hypoalbuminaemia per se is not the main driver for ascites formation: experimental models demonstrated that in patients with cirrhosis and ascites, the colloid osmotic pressure of plasma remains higher than that of ascitic fluid (19.9 ± 0.5 vs 4.8 ± 1.0 mmHg). According to the peripheral arterial vasodilatation hypothesis, portal hypertension causes an increase of vasodilators in the blood stream and a consequent reduction of effective circulating volume. It results in an activation of sympathetic nervous system (SNS), renin–angiotensin–aldosterone system (RAAS), non-osmotic secretion of vasopressin and a consequent water and sodium retention that provokes a fluid transition to the extravascular space, in particular, to the peritoneal cavity. Another source of vasodilators may be the gut, in which bacterial overgrowth and intestinal dysbiosis may cause intestinal inflammation and breakdown of the intestinal barrier. Bacterial products may stimulate the release of proinflammatory cytokines that increase splanchnic arterial vasodilatation.

**CLINICAL FEATURES AND PROGNOSIS OF CIRRHOTIC ASCITES**

According to the amount of fluid in the abdominal cavity, cirrhotic ascites is graded as follows: mild – grade 1, if detectable only by ultrasound; moderate – grade 2, if it causes symmetrical distension of the abdomen and tense – grade 3, if it causes marked abdominal distension. Moreover, ascites can be classified into uncomplicated or complicated according to the development of refractoriness, SBP or the association with HRS.

Refractory ascites is defined as ascites that cannot be mobilised or the early recurrence because of a lack of response to medical therapy. Refractory ascites includes diuretic-resistant ascites and diuretic-intractable ascites: these terms describe the failure of diuretic therapy due to unsatisfactory efficacy and development of unacceptable side effects, respectively.

Finally, ascites is defined as ‘recidivant’ when it occurs at least three times in 12 months despite the optimisation of therapy. A complete list of definitions is shown in Table 1.

Beyond physical examination, complete blood and urine tests and abdominal ultrasound, physicians should perform diagnostic paracentesis in every patient with a new-onset ascites of grade 2 or 3 or acutely decompensated. Peritoneal fluid analysis includes neutrophil count, total protein and albumin concentration, which allow to distinguish cirrhotic ascites from ascites due to other aetiologies; notably, cirrhosis is the main cause of ascites, being responsible for about 80% of cases. Moreover, peritoneal fluid analysis allows to diagnose SBP and provide important prognostic information.

Ascites is associated with poor quality of life and 2- and 5-year mortality rates at 38% and 78%, respectively, after its occurrence. The median survival for patients with compensated cirrhosis is 12 years and for decompensated cirrhosis 2 years. Main poor prognosis predictors are hyponatraemia, low arterial pressure, increased serum creatinine and low urine sodium.

**MANAGEMENT OF CIRRHOTIC ASCITES**

Table 2 lists a summary of selected recommendations from 2018 European Association for the Study of the Liver (EASL) guidelines and 2012 American Association for the Study of Liver Diseases (AASLD) guidelines.

**Management of uncomplicated ascites**

**GRADE 1**

No specific pharmacological treatment is suggested in grade 1 ascites because no treatment has definitely been shown to modify natural history. The first step is to treat the underlying liver disease, in particular, stopping alcohol consumption when present; abstinence dramatically improves the reversible component of alcoholic liver disease. Also, viral hepatitis and autoimmune hepatitis can have a dramatic response; conversely, other liver diseases are less reversible.

**GRADE 2**

**Bed rest**

Bed rest was previously recommended based on the assumption that upright position further increases plasma renin levels. There is currently insufficient published evidence to routinely recommend bed rest to all patients.

**Sodium restriction and nutrition**

Although there is no clear evidence, current guidelines suggest a moderate restriction of dietary salt (88 mmol of sodium/day, equivalent to 2 g of salt/day). Moreover, fluid intake should be restricted only in patients with dilutional hyponatraemia (sodium <130 mEq/L). However, published evidence shows a relationship between lower salt intake (in particular, sodium <50 mmol/day) and malnutrition (due to a lower caloric intake), diuretic-induced hyponatraemia and renal failure. Indeed, malnutrition is associated with infection, recurrence of ascites, hepatic encephalopathy and poor prognosis. In recent years, malnutrition has also been found to be associated with obesity and sarcopenic obesity. Therefore, guidelines
suggest to examine all patients with advanced chronic liver disease with a rapid nutritional screen and to ensure an optimal daily energy and protein intake of 35 kcal/kg actual body weight/day and 1.2–1.5 g/kg actual body weight/day, respectively, in order to avoid malnutrition.\[30\]

Diuretics

Diuretics are a cornerstone of ascites management; they are used for symptomatic treatment to induce negative fluid balance. The activation of RAAS and the consequent increase of aldosterone lead to the retention of water and sodium by proximal and distal renal tubules.\[31\] For this reason, aldosterone antagonists (e.g., spironolactone) are more effective than loop diuretics or other potassium-sparing diuretics.\[32\] The initial dose should be 100–200 mg/day and can be increased up to 400 mg/day.\[13\]

Furosemide (from 40 to 160 mg) can be combined with aldosterone antagonists\[13\] to increase the amount of sodium that reaches the distal tubule, and therefore, it indirectly increases the effectiveness of spironolactone. The combination of aldosterone antagonists and loop diuretics is effective in a shorter period of time and is safer than sequential diuretic therapy.\[33,34\]

Table 1: Definitions

| Grade 1 ascites | Ascites detectable only by ultrasound |
|----------------|--------------------------------------|
| Grade 2 ascites| Ascites causes symmetrical distension of the abdomen |
| Grade 3 ascites| Ascites causes marked abdominal distension |
| Complicated ascites | Development of refractoriness, SBP, HRS |
| Recidivant ascites | Ascites occurs at least three times in 12 months despite the optimisation of therapy |
| Refractory ascites | Ascites that cannot be mobilised or it occurs early because of a lack of response to medical therapy: 1. diuretic-resistant ascites: diuretic therapy failure due to unsatisfactory efficacy: intensive diuretic therapy for at least 1 week and salt-restricted diet (<90 mmol/day) with weight loss <0.8 kg over 4 days and urinary sodium output less than the sodium intake or reappearance of grade 2 or 3 ascites within 4 weeks of initial mobilisation 2. diuretic-intractable ascites: development of unacceptable side effects due to diuretic therapy |
| Hepatorenal syndrome | AKI: increase in sCr ≥0.3 mg/dl within 48 h or a percentage increase (≥50%) within 7 days |
| 1. Stage 1: increase in sCr ≥0.3 mg/dl or an increase in sCr ≥1.5-fold to twofold from baseline; a) stage 1A: sCr <1.5 mg/dL b) stage 1B: sCr ≥1.5 mg/dL |
| 2. Stage 2: increase in sCr > twofold to threefold from baseline; |
| 3. Stage 3: increase in sCr >threefold from baseline or sCr ≥4.0 mg/dl with an acute increase ≥0.3 mg/dl or initiation of renal replacement therapy |
| HRS-AKI criteria: 1. Cirrhosis, acute liver failure, acute-on-chronic liver failure 2. Increase in sCr ≥0.3 mg/dL within 48 h or ≥50% from baseline value according to the ICA consensus document and/or urinary output ≤0.5 mL/kg ≥6 h 3. No full or partial response, according to the ICA consensus document, after at least 2 days of diuretic withdrawal and volume expansion with albumin. The recommended dose of albumin is 1 g/kg of body weight per day to a maximum of 100 g/day 4. Absence of shock 5. No current or recent treatment with nephrotoxic drugs 6. Absence of parenchymal disease as indicated by proteinuria >500 mg/day, microhaematuria (>50 red blood cells per high-power field), urinary injury biomarkers (if available) and/or abnormal renal ultrasonography. Suggestion of renal vasoconstriction with FENa of <0.2% (with levels <0.1% being highly predictive) |
| HRS-NAKI 1. HRS-AKD a) eGFR < 60 mL/min per 1.73 m² for <3 months in the absence of other (structural) causes b) Percent increase in sCr <50% using the last available value of outpatient sCr within 3 months as the baseline value 2. HRS-CKD eGFR < 60 mL/min per 1.73 m² for ≥3 months in the absence of other (structural) causes |
| Spontaneous bacterial peritonitis | Bacterial infection of ascites without any intra-abdominal source of infection Neutrophil count in ascitic fluid of >250/mm³ determined by microscopy or flow cytometry-based automated count |

SBP: spontaneous bacterial peritonitis; HRS: hepatorenal syndrome; AKI: acute kidney injury; sCr: serum creatinine; ICA: International Club of Ascites; FENa: fractional excretion of sodium; NAKI: non-acute kidney injury; AKD: acute kidney disease; eGFR: estimated glomerular filtration rate; CKD: chronic kidney disease
| Table 2: Summary of selected recommendations about treatment of cirrhotic ascites and its complication from 2018 EASL guidelines and 2012 AASLD guidelines (details in the main text) |
| --- |
| **Rational behind grading system for recommendations** | **EASL 2018** | **AASLD 2012** |
| Level of evidence | I Randomised, controlled trials | Class of recommendation: |
| | II-1 Controlled trials without randomisation | **Class I** Conditions for which there is evidence and/or general agreement that a given diagnostic evaluation, procedure or treatment is beneficial, useful and effective |
| | II-2 Cohort and case–control analytical studies | **Class II** Conditions for which there is conflicting evidence and/or a divergence of opinion about the usefulness/efficacy of a diagnostic evaluation, procedure or treatment |
| | II-3 Multiple time series, dramatic uncontrolled experiments | **Class IIa** Weight of evidence/opinion is in favour of usefulness/efficacy |
| | III Opinions of respected authorities, descriptive epidemiology | **Class IIb** Usefulness/efficacy is less well established by evidence/opinion |
| Grade of recommendations | 1 Strong recommendations: Factors influencing the strength of the recommendation included the quality of the evidence, presumed patient-important outcomes and cost |
| | 2 Weaker recommendations: Variability in preferences and values, or more uncertainty: a weak recommendation is warranted more likely. Recommendation is made with less certainty: higher cost or resource consumption | **Class III** Conditions for which there is evidence and/or general agreement that a diagnostic evaluation/procedure/treatment is not useful/efficacious and, in some cases, may be harmful |
| **Treatment of the underlying disease** | The aetiological factors should be removed, particularly alcohol consumption and hepatitis B or C virus infections (II-2;1) | Level of evidence |
| | Prolonged bed rest cannot be recommended (III;1) | **Level A** Data derived from multiple randomised clinical trials or meta-analyses |
| | A moderate restriction of sodium intake is recommended in patients with moderate, uncomplicated ascites (I;1) | **Level B** Data derived from a single randomised trial or nonrandomised studies |
| | Diets with a very low sodium content (<40 mmol/day) should be avoided (II-2;1) | **Level C** Only consensus opinion of experts, case studies or standard of care |
| | Patients with the first episode of grade 2 (moderate) ascites should receive an anti-mineralocorticoid drug alone (I;1) | First-line treatment of patients with cirrhosis and ascites consists of sodium restriction (88 mmol/day [2000 mg/day], diet education) and diuretics (oral spironolactone with or without oral furosemide) (Class IIa, Level A) |
| | In patients who do not respond to anti-mineralocorticoids, furosemide should be added (I;1) | Fluid restriction is not necessary unless serum sodium is less than 125 mmol/L (Class III, Level C) |
| | 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) | |
| | Diuretics should be discontinued in patients with refractory ascites who do not excrete >30 mmol/day of sodium under diuretic treatment (III;1) | To be continued... |
### Table 2: Summary of selected recommendations about treatment of cirrhotic ascites and its complication from 2018 EASL guidelines and 2012 AASLD guidelines (details in the main text)

| EASL 2018 | AASLD 2012 |
|-----------|------------|
| **LVP** | **Paracentesis should be performed in patients with tense ascites. Sodium restriction and oral diuretics should then be initiated (Class IIa, Level C)** |
| 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) | For LVP, an albumin infusion of 6–8 g/L of fluid removed appears to improve survival and is recommended (Class IIa, Level A) |
| In patients undergoing LVP of >5 L of ascites, plasma volume expansion should be performed by infusing albumin, as it is more effective than other plasma expanders (I;1) | Post-paracentesis albumin infusion may not be necessary for a single paracentesis of less than 4 to 5 L (Class I, Level C) |
| In patients undergoing LVP of <5 L of ascites, it is generally agreed that these patients should still be treated with albumin (III;1) | Diuretic-sensitive patients should preferably be treated with sodium restriction and oral diuretics rather than with serial paracentesis (Class IIa, Level C) |
| After LVP, patients should receive the minimum dose of diuretics necessary to prevent re-accumulation of ascites (I;1) | Serial therapeutic paracentesis is a treatment option for patients with refractory ascites (Class I, Level C) |
| Repeated LVP plus albumin are recommended as the first-line treatment for refractory ascites (I;1) | Routine prophylactic use of fresh frozen plasma or platelets is not recommended before paracentesis (Class III, Level C) |

**NSBBs**

| **NSBBs** | **The risks versus benefits of beta blockers must be carefully weighed in each patient with refractory ascites. Consideration should be given to discontinuing or not initiating these drugs in this setting (Class III, Level B)** |
| 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 cannot be recommended at present (I;2) |
| The use of carvedilol cannot be recommended at present (I;2) | The use of carvedilol cannot be recommended at present (I;2) |

**TIPS**

| **TIPS** | **TIPS may be considered in appropriately selected patients who meet the criteria similar to those of published randomised trials (Class I, Level A)** |
| Patients with refractory or recurrent ascites (I;1) or those for whom paracentesis is ineffective 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 improves the control of ascites (I;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 improves the control of ascites (I;1) | The use of small-diameter PTFE-covered stents is recommended to reduce the risk of TIPS dysfunction and hepatic encephalopathy is recommended (I;1) |
| The use of small-diameter PTFE-covered stents is recommended to reduce the risk of TIPS dysfunction and hepatic encephalopathy is recommended (I;1) | Careful selection of patients for elective TIPS insertion is crucial (III;1) |

**Other medical treatments**

| **Other medical treatments** | **Oral midodrine should be considered in patients with refractory ascites (Class IIa, Level B)** |
| At present, the addition of clonidine or midodrine to diuretic treatment cannot be recommended (III;1) | Vaptans’ use does not currently appear justified (Class III, Level A) |

**Alfapump® system**

| **Alfapump® system** | **LT should be considered in patients with cirrhosis and ascites (Class I, Level B)** |
| Alfapump implantation in patients with refractory ascites not amenable to TIPS insertion is suggested in experienced centres (I;2) | LT should be considered in patients with cirrhosis and ascites (Class I, Level B) |

**LT**

| **LT** | **Referral for LT should be expedited in patients with refractory ascites, if the patient is otherwise a candidate for transplantation (Class IIa, Level C)** |
| 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) | Gold standard management procedures are mainly used in LT management in patients with grade 3 ascites. LT should be considered as a potential treatment option (Class IIa, Level C) |
| Patients with refractory ascites should be evaluated for LT (III;1) | Patients with refractory ascites should be evaluated for LT (III;1) |

To be continued...
Third-generation cephalosporins are recommended as the 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)

For healthcare-associated and nosocomial SBP, piperacillin/tazobactam should be given in areas with low prevalence of MDRs, 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)

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 (II-2;1)

The duration of treatment should be at least 5–7 days (III;1)

Administration of albumin (1.5 g/kg at diagnosis and 1 g/kg on day 3) is recommended in patients with SBP (I;1)

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)

Administration of prophylactic norfloxacin (400 mg/day, orally) is recommended in patients who recover from an episode of SBP (I;1)

Patients who recover from SBP have a poor long-term survival and should be considered for LT (II-2,1)

To be continued...
The aim is to achieve a body weight reduction of more than 1 kg/week until ascites is controlled. Weight loss should be less than 500 g/day in patients without peripheral oedema in order to avoid side effects such as renal impairment, hepatic encephalopathy, hyponatraemia, hypokalaemia, hypomagnesaemia and muscle cramps; notably, the last one can be successfully treated with human albumin (HA) infusion. In a small trial, albumin was found to reduce the frequency of cramps compared with placebo by 2.5 ± 2.9 episodes/week. Moreover, baclofen also may be effective in mitigating muscle cramps.

Once ascites disappears, diuretic therapy should be administered at the minimum effective dose to avoid recurrences.

**GRADE 3**

**Large-volume paracentesis**

Large-volume paracentesis (LVP) is the first-line therapy in patients with grade 3 ascites which should be completely removed in a single session.

Physicians should not perform paracentesis in case of uncooperative patient, abdominal skin infection at the proposed puncture sites, pregnancy or severe bowel distension. Despite the fact that underlying coagulopathy is common, LVP is safe; overall, the frequency of bleeding complication is estimated to be 1%–3%, and therefore, international guidelines suggest not to perform paracentesis just in case of severe coagulopathy such as accelerated fibrinolysis or disseminated intravascular coagulation.

The removal of a large volume of ascitic fluid is potentially associated with further reduction of effective blood volume. This condition is known as post-paracentesis circulatory dysfunction (PPCD), which is associated with rapid recurrence of ascites, high incidence of HRS, hyponatraemia and death. Plasma volume expansion should be performed at the completion of LVP greater than 5 L of ascites by infusing albumin (8 g for any litre of ascites removed), as it is more effective than other plasma expanders. European guidelines suggest to administer albumin also after removal of less than 5 L of ascites, while American guidelines do not. However, LVP does not modify the underlying pathophysiological abnormalities leading to ascites formation.

**Diuretics**

After LVP, patients should receive the minimum dose of diuretics necessary to prevent re-accumulation of ascites.

**Albumin**

Albumin is the most abundant protein in serum and extracellular fluids. Albumin plays a central role in maintaining plasma oncotic pressure; moreover, it has other important properties, that is, as an antioxidant and...
Liver cirrhosis does not only reduce the quantity of albumin, but also its quality. The proinflammatory and pro-oxidant state of decompensated cirrhosis affects the function and structure of albumin. The amount of proteins with intact structure and function is called ‘effective albumin concentration’. Therefore, the SA level at baseline and during treatment may predict patient outcomes, and the latter value may be influenced by HA administration.

The MATCH study enrolled 173 patients with cirrhotic ascites awaiting liver transplantation (LT). Patients in the treatment arm received midodrine 15–30 mg/day and HA 40 g every 15 days, while patients in the control arm received matching placebo for 1 year or until LT. The study treatment was not significantly different from placebo in improving survival (during the follow-up period, 7% and 5% of patients died in the treatment group and in the placebo group, respectively; \( P = 0.527 \)) or in the prevention of complications of cirrhosis (37% and 43% of patients developed complications in the treatment group and in the placebo group, respectively; \( P = 0.402 \)); there was not a reduction either in the dose of diuretic treatment or in the requirement of LVP with HA and midodrine. However, the episodes of hyponatraemia and renal failure were more severe in the placebo group.

Notably, the treatment duration was shorter than expected: a median of 80 days in the entire population and 63 days in the midodrine and HA groups; 68% from the treatment group and 55% from the placebo group were transplanted at a median of 42 days and only nine patients received the treatment for 1 year as prevented.

Moreover, compared with the ANSWER study, in the MATCH study, the total amount of HA administered was lower and the increase in SA level was not significantly different between the two groups. Patients enrolled in the MATCH study were sicker than in the ANSWER trial as demonstrated by a higher median MELD score (17 vs. 13) which is connected with a lower ‘effective albumin concentration’.

Long-term HA administration requires an effort by healthcare services and patient compliance. However, it is associated with a better quality of life because of fewer hospital admissions and lesser need for medical intervention, contributing to a favourable cost-effectiveness ratio.

Overall, HA may be defined as a disease-modifying treatment in patients with decompensated cirrhosis and, despite many clinical issues that still need to be clarified and investigated, available evidence supports long-term HA administration in patients with uncomplicated ascites.

### Transjugular intrahepatic portosystemic shunt

Transjugular intrahepatic portosystemic shunt (TIPS) is the catheterisation of a hepatic vein by the transjugular approach while the patient is under local anaesthesia,
Gallo et al.: Optimal management of cirrhotic ascites: A review for internal medicine physicians

The beneficial effect of some reports suggested protective. These regarding management of ascites, 1.5 g/kg at diagnosis and 1 g/kg on day 3. Dosage of administration of 25 g/tonce a week for 4 weeks. To prevent further reduction of effective blood volume (post-paracentesis circulatory dysfunction). Human albumin administration improves effective blood volume by attenuating peripheral arterial vasodilation, prevents renal dysfunction, enhances cardiac inotropism and reduces systemic inflammation and endothelial dysfunction, acting as an antioxidant agent. This leads to an improvement of survival and a reduction in the occurrence of spontaneous bacterial peritonitis, sepsis, hepatorenal syndrome type 1, hepatic encephalopathy grade, as well as the evolution rate to refractory ascites and the need of paracentesis.

| Indication                                           | Dosage of administration of albumin 20% | Rational                                                                 |
|------------------------------------------------------|----------------------------------------|---------------------------------------------------------------------------|
| Post-paracentesis                                    | 8 g/L of ascites removed                | To prevent further reduction of effective blood volume (post-paracentesis circulatory dysfunction) |
| Muscle cramps                                         | 25 g once a week for 4 weeks            | To reduce the frequency of muscle cramps by improving effective circulating volume |
| Long-term administration (in particular, in patients with uncomplicated ascites) | 40 g twice a week for 2 weeks and then 40 g weekly | Human albumin administration improves effective blood volume by attenuating peripheral arterial vasodilation, prevents renal dysfunction, enhances cardiac inotropism and reduces systemic inflammation and endothelial dysfunction, acting as an antioxidant agent. This leads to an improvement of survival and a reduction in the occurrence of spontaneous bacterial peritonitis, sepsis, hepatorenal syndrome type 1, hepatic encephalopathy grade, as well as the evolution rate to refractory ascites and the need of paracentesis. |
| Renal impairment (AKI stage >1A without obvious cause) | 1 g/kg body weight for two consecutive days | Human albumin prevents HRS-AKI occurrence |
| HRS-AKI                                              | 20–40 g/day                            | Human albumin reduces systemic inflammation and microvascular dysfunction, besides improving blood volume |
| SBP                                                   | 1.5 g/kg at diagnosis and 1 g/kg on day 3 | Human albumin prevents HRS-AKI occurrence |

AKI: acute kidney injury; HRS: hepatorenal syndrome; SBP: spontaneous bacterial peritonitis

followed by placement of a stent to connect portal and systemic blood systems. Potential complications include encephalopathy, heart and liver failure.

The most important indication to TIPS is the prevention of variceal bleeding. Regarding management of ascites, this treatment option was mostly studied in refractory ascites: among six prospective randomised controlled trials (RCTs) available which compared TIPS versus LVP, only two included patients with recurrent ascites. Interestingly, a meta-analysis of all six studies globally failed to demonstrate a survival advantage with TIPS (OR 0.82; 95% CI 0.46–1.50), while in the subgroups of patients with recurrent ascites, an increased survival with TIPS was found (OR 0.45; 95% CI 0.24–0.81). The beneficial effect of TIPS in patients with recurrent ascites was confirmed by a recent trial performed by Bureau and colleagues. These data suggest that cirrhotic patients with a better clinical status, in terms of liver and renal function, may survive longer than cirrhotic patients with a worse status. However, more data are warranted to investigate if TIPS may be advantageous at an earlier stage of liver dysfunction.

Management of complicated ascites

Refractory ascites

According to the International Club of Ascites (ICA) criteria, a diagnosis of refractory ascites requires intensive diuretic therapy for at least 1 week and a salt-restricted diet of less than 90 mmol/day, mean weight loss of <0.8 kg over 4 days and urinary sodium output less than the sodium intake, reappearance of grade 2 or 3 ascites within 4 weeks of initial mobilisation and development of diuretic-induced complications. Refractory ascites occurs in 5%–10% of patients with cirrhosis and ascites (more than 90% of patients have diuretic-intractable ascites) and is associated with a low probability of survival: about 50% at 6 months.

Management of refractory ascites

Large-volume paracentesis

Repeated LVP plus HA (8 g for a litre of ascites removed) are recommended as the first-line treatment for refractory ascites. This has been shown to be safe and effective with a low incidence of renal injury and electrolyte abnormalities and lesser systemic and haemodynamic disturbances compared with diuretics.

Diuretics

According to EASL guidelines, diuretics should be discontinued in patients with refractory ascites who do not excrete >30 mmol/day of sodium under diuretic treatment.

Nonselective beta-blockers

The administration of nonselective beta-blockers (NSBBs) in patients with refractory ascites is a matter of discussion. NSBBs reduce portal pressure and are currently used for primary and secondary prophylaxis of variceal haemorrhage. Some reports suggested protective effects with NSBBs in patients with decompensated cirrhosis probably mediated by reduction of intestinal permeability and inflammation, particularly in this advanced stage. Illustratively, a recent post hoc analysis of three RCTs explored the co-administration of vaptans.
and NSBBs in patients with ascites: the 52-week cumulative all-cause mortality was similar in the NSBB user and nonuser groups (23.2% vs. 25.3%, adjusted hazard ratio [HR] = 0.92, 95% CI 0.72–1.18). This effect was also confirmed in the subgroup of patients with refractory ascites; however, during follow-up, 29% of initial NSBB users stopped taking NSBBs, entailing an increased risk of mortality, hospitalisation, variceal bleeding, bacterial infection and/or development of HRS.[70]

Conversely, various studies caution the use of NSBBs in situations such as SBP,[71] severe alcoholic hepatitis[72] and refractory ascites. In particular, the Clichy group reported poor survival and increased risk of PPced among patients on NSBB therapy; in one study, the median survival time was 5 months in patients treated with NSBBs compared with 20 months in patients not treated with NSBBs.[73,74]

The mechanism underlying these findings was supposed to be related to the induction of systemic arterial hypotension and exhaustion of cardiac reserve. Furthermore, patients with advanced stages of decompensation often receive higher doses of NSBBs, thus exaggerating the detrimental effects on systemic haemodynamics. Accordingly, a retrospective nationwide study of 3719 Danish patients with cirrhosis found a reduction in mortality for propranolol doses <160 mg/day (HR 0.4; 95% CI 0.2–0.8), but an increase in mortality for doses >160 mg/day (HR 2.5; 95% CI 0.9–6.8).[75] This finding led to the ‘window hypothesis’, which postulates that NSBBs are associated with higher rates of survival in selected patients with moderate-to-large oesophageal varices, but without refractory ascites, hypotension, SBP, HRS, sepsis or severe alcoholic hepatitis.[76,77]

In conclusion, while NSBBs continue to occupy a pivotal role in the treatment of portal hypertension, recent evidence has not only outlined additional, haemodynamically independent beneficial effects in cirrhosis, but also described potentially debilitating effects in advanced stages. According to AASLD and EASL guidelines, physicians should use NSBBs with caution in cirrhotic patients with refractory ascites and discontinue usage if haemodynamic or renal impairment arises. Moreover, carvedilol and high-dose propanolol should be avoided.[73,74]

**Albumin**

Evidence concerning long-term administration of HA in patients with refractory ascites is scant compared to that on uncomplicated ascites. However, an observational study by Di Pascoli and colleagues suggested that mortality may be significantly reduced with the administration of 40 g twice a week than without (41.6% vs. 65.5% over a period of 2 years).[78]

**Transjugular intrahepatic portosystemic shunt**

Evidence supporting the use of TIPS in patients with refractory ascites is controversial. Among four RCTs exclusively concerning patients with refractory ascites, one showed that TIPS with bare-metal stents worsens mortality compared with standard treatment, mainly because of the detrimental effect of TIPS in Child–Pugh class C patients.[79] Two RCTs did not find any difference.[80,81] A better survival with TIPS was reported in just one study.[82] Recent evidence assessed the effects of TIPS with polytetrafluoroethylene (PTFE)-covered stent grafts. Two retrospective studies reported better control with covered stent grafts than with bare stent grafts in patients with refractory ascites.[83,84] It seems that the development of hepatic encephalopathy and stent thrombosis can be reduced with the use of covered stent grafts.[85,86] Other potential side effects are liver and heart failure.[87,88]

A careful selection of patients is crucial to maximise the beneficial effects of TIPS. Score systems have been developed to help clinicians’ choice.[89] Indeed, 2012 AASLD guidelines recommend to reserve TIPS for patients who meet the criteria similar to those of published randomised trials.[14]

**Other treatments**

A substantial portion of patients with refractory ascites are not candidates for TIPS insertion. Several medical treatments such as midodrine, clonidine, terlipressin and tolvaptan have been studied in this scenario, but at present, EASL guidelines do not recommend any of them.[11,13] Conversely, AASLD guidelines suggest the use of midodrine in this setting.[14]

The automated low-flow ascites pump (Alfapump®) system is a subcutaneously implanted pump connected to catheters that transfer ascites from the peritoneal cavity to the bladder, allowing elimination with urine. Alfapump showed efficacy in two RCTs in patients with advanced cirrhosis and refractory ascites, reducing the need for paracentesis by 90%.[89,90] Conversely, its main side effect and contraindication is renal failure.[91] Alfapump is considered in EASL guidelines, but not in AASLD guidelines.[13,14]

**Liver transplantation**

EASL guidelines suggest that LT should be considered not only in all patients with refractory ascites, but also in patients with grade 2 or 3 ascites because of their poor prognosis[13] which can greatly improve after LT: 71% of patients are alive at 10 years.[92]

The Child–Pugh classification and, since 2002, MELD score have been used for patient selection and priority.
RENAL IMPAIRMENT AND HRS

Renal impairment in patients with decompensated cirrhosis consists of chronic kidney disease (CKD) and acute renal failure; the latter term was replaced by the term acute kidney injury (AKI) and is defined by the Kidney Disease Improving Global Outcomes (KDIGO) group as an increase in serum creatinine ≥0.3 mg/dL within 48 h or a percentage increase ≥50% within 7 days. AKI has three severity stages: in the first stage, serum creatinine increases up to twofold from baseline; stage 1 includes stage 1A with serum creatinine <1.5 mg/dL and stage 1B with serum creatinine ≥1.5 mg/dL; in the second stage, it increases up to threefold from baseline and in the third stage, it increases more than threefold from baseline. From an aetiologic point of view, AKI is differentiated into prerenal AKI, HRS-AKI, intrarenal or intrinsic AKI, which is represented by acute tubular necrosis (ATN), and post-renal AKI.

Overall, HRS is just one of the possible reasons of renal impairment in patients with liver disease; a study observed that it is less common than prerenal or infection-associated kidney injury representing just 13% of the underlying causes of renal deterioration in this setting. However, the prevalence of HRS in cirrhotic ascites rises to 39% at 5 years and clearly worsens the prognosis.

HRS was historically defined as a functional renal failure caused by intrarenal vasoconstriction in patients with liver disease; moreover, based on the time of development and prognosis, HRS is classified as type 1 and type 2. Absence of renal parenchymal damage and its potential reversibility are considered as its main characteristics.

In recent years, the ‘functional’ paradigm of HRS has been questioned. The idea that HRS has an exclusive functional nature is now unsustainable. Vasodilatation and inadequate cardiac output, which leads to intrarenal vasoconstriction and renal impairment, was considered the epicentre of pathogenesis, but systemic inflammation and microvascular dysfunction, as well as some degree of parenchymal damage are also considered determinants.

Second, the historic classification in type 1 and type 2 was recently revised. Type 1 is now called HRS-AKI and fulfills the characteristics of AKI. It occurs in patients with cirrhosis and ascites as well as in patients with acute liver failure in the presence of precipitating factors: not only bacterial infections from any sources (in particular, bacterial translocation from the gut) seem to play a major role in this process, but also LVP without adequate albumin administration and excessive administration of diuretics could provoke HRS-AKI.

Nowadays, the diagnosis of HRS-AKI is based on the revised ICA criteria: 1) presence of cirrhosis and ascites, 2) presence of AKI, 3) no response to diuretic withdrawal and volume expansion with albumin, 4) absence of shock, 5) no nephrotoxic drugs, 6) no macroscopic signs of structural kidney injury (proteinuria, microhaematuria and/or abnormal renal ultrasonography).

Since prerenal-AKI can be recognised by the response to plasma volume expansion, the differential diagnosis between HRS-AKI and ATN-AKI is a challenge. ATN-AKI is characterised by signs of parenchymal damage such as microhaematuria and proteinuria, which cannot be present. Novel biochemical markers are under study to solve this problem, and the most promising is neutrophil gelatinase-associated lipocalin (NGAL).

Nevertheless, the presence of increased levels of some tubular biomarkers in patients with HRS-AKI may suggest that there is a continuum from HRS to ATN. Type 2 is now called HRS-NAKI (non-AKI) because its main clinical consequence is not acute renal failure, but refractory ascites. It can be considered the extreme expression of the impairment in circulatory function during the course of cirrhosis. Angeli and colleagues proposed to refer HRS-NAKI either to CKD, defined by a reduction of the glomerular filtration rate (GFR) under 60 mL/min developed in more than 3 months (HRS-CKD), or to acute kidney disease, if the reduction developed in less than 3 months without the criteria for AKI (HRS-AKD). HRS-AKI and HRS-NAKI should not be seen as a continuum because they are clinically and pathophysiologically independent.

Management of renal impairment and HRS

When a patient with liver disease suffers from AKI, any potentially leading drug should be stopped, volume replacement therapy should be provided and, in the absence of an evident underlying cause, 1 g of 20% HA/kg of body weight has to be administered for two consecutive days.

Based on EASL guidelines, HRS-AKI should be treated with HA associated with vasoconstrictive drugs. The drug that is the most recognised for this purpose is terlipressin, administered at a dose of 0.5 mg every 4–6 h or by continuous i.v. infusion at an initial dose of 2 mg/day. HA...
should be administered at the dose 20–40 g/day, higher doses considered in the treatment of AKI as discussed above have never been studied in patients with HRS-AKI, in particular, in combination with terlipressin. Bearing in mind new theories about the role of systemic inflammation in decompensated cirrhosis and HRS, and the physiologic role of albumin, we may understand the central role of HA administration in this scenario.[11] In one prospective non-randomised trial, reversal of HRS, defined as a serum creatinine level of <1.5 mg/dL, was significantly higher with terlipressin plus HA than with terlipressin alone (77% vs 25%).[52]

An alternative vasoconstrictor is noradrenaline, given by continuous i.v. infusion at the dose of 0.5–3 mg/h. This drug is less studied and requires a central line and an intensive care unit setting.[114] Seven RCTs compared terlipressin with noradrenaline, but only one study found that terlipressin had better survival than noradrenaline.[112] Terlipressin has not been approved for use in the United States, so it is not mentioned in AASLD guidelines, which suggest the use of octreotide plus midodrine as vasoactive drugs.[14] However, this suggestion is based on the results of small studies,[11,114] and the combination octreotide plus midodrine has been shown to be much less effective than terlipressin in the treatment of HRS in a recent RCT.[115]

TIPS has a minor role in HRS-AKI because it is often contraindicated due to disease severity that limits its application;[113] moreover, a recent meta-analysis showed high incidence of hepatic encephalopathy with TIPS.[116] Non-responders to the combination of HA and vasoconstrictors should be considered for renal replacement therapy following the same criteria as in general population.[10] LT is the best therapeutic option for patients with HRS, regardless of the response to drug therapy;[117] moreover, simultaneous liver–kidney transplantation should be considered for HRS-AKI with no response to drug therapy.[113,114]

Based on EASL guidelines, in patients with HRS-NAKI, including HRS-CKD, vasoconstrictors and HA are not recommended despite their efficacy because of the high rate of recurrence possibly due to the underlying renal parenchymal damage.[113,118] TIPS may have a role in the management of HRS-NAKI because the frequent association with of refractory ascites, for which the efficacy of TIPS has been described above.[10]

Renal replacement therapy and LT, in particular, simultaneous liver–kidney transplantation, can be indicated in patients with cirrhosis and significant CKD.[113,114] Despite continuous progress in the treatment of HRS, its prognosis remains severe and more therapeutic options are warranted. A recent systematic review and meta-analysis showed that the combination of Traditional Chinese Medicine (TCM) with conventional therapy seems to lower serum creatinine, blood urea nitrogen, bilirubin, plasma ammonia and abdominal circumference; however, the study was invalidated by significant heterogeneity and methodological issues. At present, evidence is scant and TCM cannot be officially suggested as an adjunctive treatment.[119]

**SPONTANEOUS BACTERIAL PERITONITIS**

SBP is defined as a bacterial infection of ascites without any intra-abdominal source of infection.[128] Any patient with cirrhotic ascites is at risk of SBP, and its prevalence in outpatients is 1.5%–3.5%, increasing sensibly in hospitalised patients; moreover, it makes the prognosis of ascites worse (mortality due to SBP is now estimated to be around 20%).[121] SBP is highly common in cirrhotic ascites, justifying diagnostic paracentesis in all patients. SBP is diagnosed when the neutrophil count is greater than 250/mm³ in ascitic fluid. Conversely, the positivity of bacterial culture is not essential for diagnosis, considering the high rate of false-negative culture; however, cultures should always be performed because any isolated bacteria can guide antibiotic therapy. Bacterascites is instead defined as a neutrophil count less than 250/mm³, but positive bacterial culture.[113,114]

**Management of SBP**

Nowadays, major concern about the treatment of SBP gravitates around the alarming increase in the spread of multidrug-resistant (MDR) and extensively drug-resistant (XDR) organisms.[122] Notably, cirrhotic patients are highly susceptible to develop infections by these bacteria due to frequent hospitalisations and antibiotic treatment.[123] The most recent guidelines radically changed the recommendations about the choice of antibiotic treatment, prioritising the probability of infection by MDRs or XDRs; this approach is a major novelty compared with the past.[13]

In the setting of community-acquired SBP, third-generation cephalosporin is the first-line antibiotic treatment,[113,114] in particular, cefotaxime is considered the best option because of its high concentration in the ascitic fluid.[124] The 2012 American guidelines also suggest the use of ofloxacin as an alternative to cefotaxime for patients without prior exposure to quinolones.[14]
In countries with high rates of bacterial resistance, piperacillin/tazobactam or carbapenem should be considered as the first-line option also in a community-acquired setting according to EASL guidelines.\textsuperscript{[13]} For healthcare-associated and nosocomial SBP, EASL guidelines suggest the use of piperacillin/tazobactam in areas with low prevalence of multidrug resistant. Carbapenem should instead be used in areas with high prevalence of extended-spectrum beta-lactamases (ESBLs) producing Enterobacteriaceae, even associated with glycopeptides, daptomycin or linezolid if coexists also high prevalence of gram-positive MDR bacteria.\textsuperscript{[13]} Conversely, AASLD guidelines do not provide specific suggestion to manage empiric antibiotic therapy in nosocomial setting or the presence of risk factors for MDR infections, but just advises to follow local susceptibility testing of bacteria.\textsuperscript{[14]}

HA should be administered in combination with antibiotic therapy in order to prevent HRS-AKI occurrence. In a landmark RCT, mortality was reduced from 29\% to 10\% in patients treated with high-dose HA in combination with cefotaxime than on treatment with cefotaxime alone. Treatment with HA was particularly effective in patients with baseline serum bilirubin ≥4 mg/dL or serum creatinine ≥1 mg/dL, who, however, represented the majority of enrolled patients.\textsuperscript{[5]} Based on these results, EASL guidelines recommend the administration of HA at the dose of 1.5 g/kg at diagnosis and 1 g/kg on day 3 in patients with SBP.\textsuperscript{[13]} Conversely, AASLD guidelines restrict this recommendation to patients with similar characteristics to the subgroup of patients who experienced the maximal benefit in the discussed RCT.\textsuperscript{[14]} Our knowledge about alternative regimens with low-dose HA is scant, since just a few studies investigated this possibility.\textsuperscript{[22,126]}

Prophylaxis of SBP with norfloxacin is another important topic. According to guidelines, primary prophylaxis has to be limited to patients with ascitic fluid protein <1.5 g/ dL along with impaired renal function or liver failure.\textsuperscript{[13,14]} in these categories, 3-month probability of survival was 94\% with norfloxacin and 62\% with placebo in an RCT.\textsuperscript{[27]} Moreover, patients who recover from an episode of SBP should be considered for LT\textsuperscript{[13]} besides receiving secondary prophylaxis, which demonstrated to reduce recurrence from 68\% to 20\% in another RCT.\textsuperscript{[128]}

As for HRS, TCM was investigated as an adjunctive treatment for SBP. In a recent meta-analysis the additional use of Xuebijing injection seems to improve the efficacy of antibiotics for the treatment of SBP in liver cirrhosis; currently, no official suggestion can be made due to a low level of evidence.\textsuperscript{[129]}

**HEPATIC HYDROTHORAX**

Hepatic hydrothorax is the accumulation of ascites in the pleural space (mostly the right) sustained by a pressure gradient through embryologic diaphragmatic defects. Hepatic hydrothorax can lead to respiratory failure and can be complicated by spontaneous bacterial pleural empyema; its appearance worsens the prognosis of decompensated cirrhosis.\textsuperscript{[130]}

Diagnostic approach begins with the exclusion of cardiopulmonary and primary pleural diseases; moreover, diagnostic thoracentesis is required to rule out bacterial infection.\textsuperscript{[13,131]} Diaphragmatic defects can be identified by magnetic resonance imaging or colour Doppler and contrast-enhanced ultrasonography.\textsuperscript{[132,133]}

**Management of hepatic hydrothorax**

The management includes diuretics, LVP, TIPS and LT.\textsuperscript{[13,14]} Moreover, pleurodesis can be suggested to patients who are not candidates for LT or TIPS.\textsuperscript{[13]} Different techniques have been considered, including surgical approach with the possibility of concomitant repair of diaphragmatic defects; however, the frequency of complications is high.\textsuperscript{[134]}

When no other options are available, therapeutic thoracentesis is required to relieve dyspnoea, but its efficacy is transient and multiple repetition is challenging.\textsuperscript{[13,135]} Chest tube insertion is contraindicated because of the high rate of placement-related complications such as bleeding or pneumothorax, infections, protein loss and renal impairment.\textsuperscript{[14,136,137]} Conversely, pigtail tube for drainage seems to have a lower rate of complications.\textsuperscript{[138]} Indwelling pleural catheters (IPC) are tunneled catheters inserted subcutaneously that are often used in the treatment of malignant pleural effusions; although this option is not mentioned by guidelines, some reports showed a favourable cost-effective ratio.\textsuperscript{[139,140]}

**Conflict of Interest**

We have no conflicts of interest to disclose.

**Author Contribution**

A. Gallo and A. Squizzato contributed to the conception and design. A. Gallo, C. Dedionigi and C. Civitelli wrote the manuscript. A. Squizzato, A. Panzeri and C. Corradi contributed to the language and content editing. The manuscript has been read and finally approved by all the authors.
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