Q: What is the role for terlipressin in hepatorenal syndrome?

A: With careful patient selection, terlipressin, a synthetic analogue of vasopressin, is an effective therapy for hepatorenal syndrome (HRS). The drug has compared favorably with placebo and, based on limited data, other vasoconstrictors.

HEPATORENAL SYNDROME: UPDATED DEFINITIONS

HRS is a serious complication of advanced liver disease defined by kidney dysfunction associated with complex changes in the splanchnic circulation resulting in vasoconstriction and renal hypoperfusion. Traditionally, acute HRS, characterized by a rapid decline in kidney function, has been referred to as type 1 HRS; chronic HRS, characterized by progressively worsening kidney function, has been referred to as type 2 HRS.

The diagnostic criteria for HRS were revised recently based on the International Club of Ascites definition of acute kidney injury (AKI) in patients with cirrhosis. The updated definition addresses the potential overestimation of renal function based on the serum creatinine (Scr) level in patients with cirrhosis, where Scr is reduced because of malnutrition and muscle wasting. The revised definition reclassifies type 1 HRS as HRS-AKI and type 2 HRS as HRS-chronic kidney disease. HRS-AKI is considered a diagnosis of exclusion.

The ultimate therapy for HRS-AKI may be liver transplant in appropriate candidates. However, because AKI is associated with significantly increased mortality risk, therapies are needed that target reversal of HRS-AKI and potentially serve as a bridge to liver transplant. Unfortunately, there are limited treatment options for HRS-AKI. Current therapies include the combination of midodrine and octreotide, norepinephrine, and terlipressin.

TERLIPRESSIN: THE ONLY APPROVED THERAPY FOR HRS-AKI

Terlipressin, available in some parts of the world for several years, was just recently approved by the US Food and Drug Administration (FDA). It is the only drug with an FDA-labeled indication for the treatment of HRS-AKI. The 2021 American Association for the Study of Liver Diseases and the 2018 European Association for the Study of the Liver guidelines both recommend terlipressin in combination with albumin as first-line treatment for patients with HRS-AKI.

EVIDENCE FOR TERLIPRESSIN

Terlipressin has been compared with placebo and with other vasoconstrictors. The efficacy and safety of terlipressin for HRS reversal was demonstrated in CONFIRM (Terlipressin Plus Albumin for the Treatment of Type 1 Hepatorenal Syndrome), a multi-center, randomized, placebo-controlled, double-blind study that led to the FDA approval of terlipressin. It included patients with cirrhosis, ascites, and type 1 HRS, defined as an Scr of at least 2.25 mg/dL without improved renal function within 48 hours of discontinuing diuretic therapy and administration of albumin. CONFIRM was designed before the new International Club of Ascites AKI definition, so enrolled patients generally had higher Scr levels than would qualify for inclusion based on the new HRS-AKI definition (mean Scr 3.5 mg/dL). Further, patients may have been excluded from CONFIRM who would now qualify for treatment based on the updated HRS-AKI definition. CONFIRM exclusion criteria included an Scr greater than 7 mg/dL, shock, large-volume paracentesis.

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(> 4 L) within 48 hours of randomization, and uncontrolled bacterial infection. Concomitant vasopressors were not permitted. Patients were randomized to receive either terlipressin (N = 199) at an initial dose of 1 mg terlipressin acetate (equivalent to 0.85 mg terlipressin) every 6 hours or placebo (N = 100). Doses were adjusted based on clinical response according to prescribing-information guidelines (Figure 1). Both treatment arms received daily albumin replacement (median 335 g including 1 g/kg body weight for 2 treatment arms received daily albumin replacement as the cause of cirrhosis.6

The primary outcome in CONFIRM was verified reversal of HRS, defined as 2 consecutive SCr values of 1.5 mg/dL or less and survival without need for renal replacement therapy for 10 days. This occurred in 32% of patients in the terlipressin group compared with 17% of patients in the placebo group (P = .006). In addition, HRS reversal without need for renal replacement therapy for 30 days occurred in 34% of terlipressin patients compared with 17% of placebo patients (P < .001). However, there was no difference in 90-day mortality rates (51% terlipressin vs 45% placebo, 95% confidence interval [CI] –6 to 18), nor was there a difference in liver transplant rates (23% terlipressin vs 29% placebo).6

Terlipressin was associated with an increased risk of abdominal pain, nausea, diarrhea, and respiratory failure (14% terlipressin vs 5% placebo). In post hoc analyses, populations that were potentially more likely to benefit from terlipressin for HRS reversal included patients with systemic inflammatory response syndrome at baseline, mean arterial pressure less than 70 mm Hg at initiation, and alcohol-associated hepatitis as the cause of cirrhosis.6

Terlipressin compared with other vasoconstrictors

The comparative efficacy of norepinephrine and terlipressin was evaluated along with daily albumin replacement in 120 patients with HRS-AKI in a 2020 randomized, open-label trial. The study included patients who met the recently updated International Club of Ascites criteria for HRS. Consequently, patients in this study had lower SCr levels at initiation than in CONFIRM (mean 1.79 mg/dL in the terlipressin group vs 2.02 mg/dL in the norepinephrine group), so this study may reflect earlier initiation of HRS therapy. The study was conducted in India, where terlipressin was administered as a continuous intravenous infusion, whereas in the United States, FDA-approved dosing is intermittent intravenous administration. Compared with norepinephrine, terlipressin administration was associated with improved reversal of HRS (40% vs 16.7%, P = .004), reduced need for renal replacement therapy (56.6% vs 80%, P = .006), and improved 28-day survival (48.3% vs 20%, P = .001). The rate of adverse effects was significantly higher with terlipressin (23.3% vs 8.3%, P = .02), and effects were mainly gastrointestinal.

In a 2015 randomized trial conducted in Italy, terlipressin was administered by continuous infusion to 27 patients, and 22 patients received midodrine and octreotide; both groups also received albumin. Patients who received terlipressin had higher rates of HRS reversal (19 of 27, or 70.4%) compared with those who received the combination of octreotide and midodrine (6 of 21, or 28.6%, P = .01).

Finally, despite limited prospective data comparing terlipressin with other vasoconstrictors, a recent meta-analysis of 26 HRS trials concluded that terlipressin is associated with greater HRS reversal compared with midodrine-octreotide (72.5 more reversals per 1,000; 95% CI > 198 to < 12) and norepinephrine (30.4 more reversals per 1,000; 95% CI > 83 to < 14.6). Based on these data, terlipressin may be more effective than norepinephrine and is likely more effective than the combination of midodrine and octreotide, a conclusion supported by consensus guideline recommendations.

### TABLE 1

**Diagnostic criteria for hepatorenal syndrome with acute kidney injury**

| Cirrhosis; acute liver failure; ACLF |
| Increase in serum creatinine: |
| • ≥ 0.3 mg/dL within 48 hours or |
| • ≥ 50% from baseline according to ICA criteria and/or |
| • Urinary output ≤ 0.5 mL/kg ≥ 6 hours. |

No full or partial response after at least 2 days of diuretic withdrawal and volume expansion with albumin (1 g/kg body weight per day to a maximum 100 g/day)

Absence of shock

No current or recent treatment with nephrotoxic drugs

Absence of parenchymal disease as indicated by proteinuria > 500 mg/day, microhematuria and/or abnormal renal ultrasonography

ACLF = acute-on-chronic liver failure


Angeli P, Garcia-Tsao G, Nadim MK, Parikh CR.

*News in pathophysiology, definition and classification of hepatorenal syndrome: a step beyond the International Club of Ascites (ICA) consensus document, with permission from Elsevier.*
**TERLIPRESSIN**

**PATIENT SELECTION CONSIDERATIONS: POPULATIONS AT RISK OF ADVERSE EFFECTS**

Terlipressin is recommended as a first-line treatment of HRS-AKI and has demonstrated efficacy for HRS reversal over other therapies, but its use is not without hazard. It is associated with respiratory failure, especially in the setting of albumin administration. Other comorbidities can increase the risk of respiratory failure and should be addressed before using terlipressin.6,11,12

**Risk of respiratory failure**

The risk of respiratory failure with terlipressin in the CONFIRM study was 14% overall vs 5% with placebo, and death from respiratory failure occurred in 11% of terlipressin patients vs 2% of placebo patients.6 A similar trend emerged in the pooled data from all phase 3 studies of terlipressin compared with placebo: among 598 patients in placebo-controlled studies, respiratory failure occurred in 11.2% of terlipressin patients vs 4.4% of placebo patients.11

Respiratory failure with terlipressin is hypothesized to result from increased systemic vascular resistance,12 which may lead to pulmonary edema in patients with cardiac dysfunction or volume overload. CONFIRM excluded patients with severe cardiovascular disease, including unstable angina, known pulmonary edema, heart failure, and symptomatic peripheral vascular disease.6 These patients are unlikely to be good candidates for terlipressin therapy. Further, all patients in CONFIRM received daily albumin replacement of 1 g/kg on day 1, followed by 20 to 40 g per day thereafter. The median total albumin administered during terlipressin administration was 199 g (± 147 g).6 Volume overload associated with intravenous albumin dosing has been proposed as a risk factor for respiratory failure in patients receiving terlipressin.11 Consequently, excess volume administration should be avoided, particularly excessive albumin, which is associated with pulmonary edema in vulnerable patients.

Additional factors associated with increased risk of respiratory failure include acute-on-chronic liver failure (Aclf) grade 3, grade 3 to 4 hepatic encephalopathy, aspiration pneumonia, and recent upper gastrointestinal bleeding.11 It is recommended that these patients receive optimized therapy for hepatic

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**Figure 1.** Guide to terlipressin dosing. Based on information in reference 7.

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Days 1–3
- Initial dose: 0.85 mg terlipressin every 6 hours
- Record baseline serum creatinine on day 1

Day 4
Assess serum creatinine level compared with baseline

If serum creatinine has decreased by 30% or more from baseline, continue terlipressin 0.85 mg every 6 hours

If serum creatinine has decreased by less than 30% from baseline, increase terlipressin dosage to 1.7 mg (2 vials) every 6 hours

If serum creatinine is at or above baseline value, discontinue terlipressin

Continue until 24 hours after patient achieves a second consecutive serum creatinine value of ≤ 1.5 mg/dL at least 2 hours apart, or for a maximum of 14 days
The increased utilization of healthcare resources asso-

The cost of terlipressin is substantial, with an average wholesale cost more than $1,000 per 0.85-mg vial. Further, although terlipressin may be admin-

## REFERENCES


4. European Association for the Study of the Liver. EASL clinical practice guidelines for the management of patients with decompen-

