

# Prevention of Hepatorenal Syndrome, as a Complication of Liver Cirrhosis

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## Dear Editor,

Hepatorenal syndrome (HRS) is defined as the appearance of a renal injury in patients with chronic liver disease. It is of a functional origin and caused by systemic circulatory dysfunction, which leads to renal vasoconstriction secondary to the effect of increased vasoactive substances intended to compensate for initial splanchnic vasodilation [1]. HRS always develops in the context of circulatory dysfunction, mainly in the splanchnic arterial territory, and is generally associated with ascites and hyponatremia due to the activation of neurohormonal systems [2].

The diagnosis of this pathology is clinical and its definition and classification have been updated according to the criteria for acute kidney injury (AKI) [3]. The HRS diagnostic criteria proposed by ICA-AKI are:

(1) Previous diagnosis of chronic liver disease and ascites; (2) AKI diagnosis; (3) No response after 2 consecutive days of diuretic withdrawal and plasma volume expansion with albumin (1 g/kg body weight); (4) Absence of shock; (5) Current or recent use of nephrotoxic drugs (nonsteroidal anti-inflammatory drugs, aminoglycosides, iodinated contrast media, etc.); and (6) No macroscopic evidence of structural kidney injury, defined as no proteinuria (>500 mg/day), no microhaematuria (>50 red blood cells per high-power field), and normal renal ultrasound findings [2-3].

In terms of treatment, the use of albumin is indicated for the prevention of circulatory dysfunction that occurs in the acute context of the disease based on its beneficial effect as a plasma expander; however, more recent studies also indicate that albumin decreases the systemic inflammatory response and reduces proinflammatory substances [4]. In addition, it can improve autoregulation of renal perfusion, which can lead to a reduction in oxidative stress and endothelial activation [5].

Different methods of prevention and reduction of mortality have been studied, such as the placement of a transjugular intrahepatic portosystemic shunt (TIPS) [4]. This technique has been used successfully in individuals with recurrent ascites with excellent results in terms of morbidity and mortality [5]. However, further trials are needed to consolidate its efficacy in the treatment of hepatorenal dysfunction and its benefit-risk assessment [6-8].

## Conflict of Interest

We declare that we have no conflict of interest related to the content of this article.

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## References

1. Heimbach JK, Kulik LM, Finn RS, Sirlin CB, Abecassis MM, et al. (2018) AASLD guidelines for the treatment of hepatocellular carcinoma. *Hepatology* 67: 358-380.
2. Simbrunner B, Trauner M, Reiberger T, Mandorfer M (2021) Recent advances in the understanding and management of hepatorenal syndrome. *Faculty Reviews* 10.
3. Marrero JA, Kulik LM, Sirlin CB, Zhu AX, Finn RS, et al. (2018) Diagnosis, Staging, and Management of Hepatocellular Carcinoma: 2018 Practice Guidance by the American Association for the Study of Liver Diseases. *Hepatology* 68: 723-750.
4. Gullello R, Alejandro E (2015) Insuficiencia renal en la cirrosis: Evaluación de nuevas clasificaciones pronósticas y tratamiento del síndrome hepatorenal tipo 1 asociado a sepsis. *Universitat de Barcelona*.

5. Salerno F, Navickis RJ, Wilkes MM (2013) Albumin infusion improves outcomes of patients with spontaneous bacterial peritonitis: a meta-analysis of randomized trials. *Clin Gastroenterol Hepatol* 11: 123-130.
6. Plauth M, Bernal W, Dasarathy S, Merli M, Plank LD, et al. (2020) ESPEN Guideline on Clinical Nutrition in Liver Disease. *Clin Nutr* 39: 485-521.
7. Belcher JM, Sanyal AJ, Peixoto AJ, Perazella MA, Lim J, et al. (2014) Kidney biomarkers and differential diagnosis of patients with cirrhosis and acute kidney injury. *Hepatology* 60: 622-632.
8. Caraceni P, Riggio O, Angeli P, Alessandria C, Neri S, et al. (2018) Long-term albumin administration in decompensated cirrhosis (ANSWER): an open-label randomised trial. *Lancet* 391: 2417-2429.