

# Altered Doppler flow patterns in cirrhosis patients: an overview

Pooya Iranpour<sup>1</sup>, Chandana Lall<sup>1</sup>, Roozbeh Houshyar<sup>1</sup>, Mohammad Helmy<sup>1</sup>, Albert Yang<sup>1</sup>, Joon-Il Choi<sup>2</sup>, Garrett Ward<sup>1</sup>, Scott C Goodwin<sup>1</sup>

<sup>1</sup>Department of Radiology, University of California Irvine, Orange, CA, USA; <sup>2</sup>Department of Radiology, Seoul St. Mary's Hospital, College of Medicine, The Catholic University of Korea, Seoul, Korea

Doppler ultrasonography of the hepatic vasculature is an integral part of evaluating pre-cirrhotic and cirrhotic patients. While the reversal of the portal venous flow is a well-recognized phenomenon, other flow patterns, although not as easily understood, may play an important role in assessing the disease status. This article discusses the different characteristic flow patterns observed from the portal vein, hepatic artery, and hepatic vein in patients with liver cirrhosis or related complications and procedures. Knowledge of these different flow patterns provides additional information that may reinforce the diagnosis of cirrhosis, help in staging, and offer prognostic information for determining the direction of therapy. Doppler ultrasonography is invaluable when liver transplantation is being considered and aids in the diagnosis of cirrhosis and portal hypertension.

**Keywords:** Liver cirrhosis; Hypertension, portal; Ultrasonography, Doppler, color

## Introduction

Liver cirrhosis is a major cause of morbidity and mortality and is currently the 14th leading cause of death worldwide. Early diagnosis may help prevent the associated detrimental effects, including variceal bleeding, hepatic encephalopathy, and portal vein thrombosis. Staging of liver cirrhosis helps with prognostic information and directs appropriate therapy. Parenchymal changes seen in patients with diffuse liver disease and cirrhosis include surface nodularity, increased echogenicity, coarsened echotexture, and a high caudate-to-right hepatic lobe size ratio. Secondary signs such as splenomegaly and varices represent sequelae of portal hypertension in the case of cirrhosis [1,2]. Spectral and color Doppler ultrasonography can demonstrate many characteristic flow patterns from hepatic vasculature in patients with liver cirrhosis with or without portal hypertension. In this article, we discuss the many characteristic flow patterns identified by Doppler ultrasonography within the portal veins, hepatic veins, and hepatic arteries, as well as the pathophysiology and secondary signs and pitfalls in this patient population.

# ULTRA SONO GRAPHY

## REVIEW ARTICLE

<http://dx.doi.org/10.14366/usg.15020>  
pISSN: 2288-5919 • eISSN: 2288-5943  
Ultrasonography 2016;35:3-12

Received: April 4, 2015  
Revised: May 27, 2015  
Accepted: May 27, 2015

### Correspondence to:

Roozbeh Houshyar, MD, Department of Radiology, University of California Irvine, 101 The City Dr. South, Orange, CA 92868, USA

Tel. +1-714-456-5642  
Fax. +1-714-456-6832  
E-mail: rhoushya@uci.edu

This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (<http://creativecommons.org/licenses/by-nc/3.0/>) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

Copyright © 2016 Korean Society of Ultrasound in Medicine (KSUM)



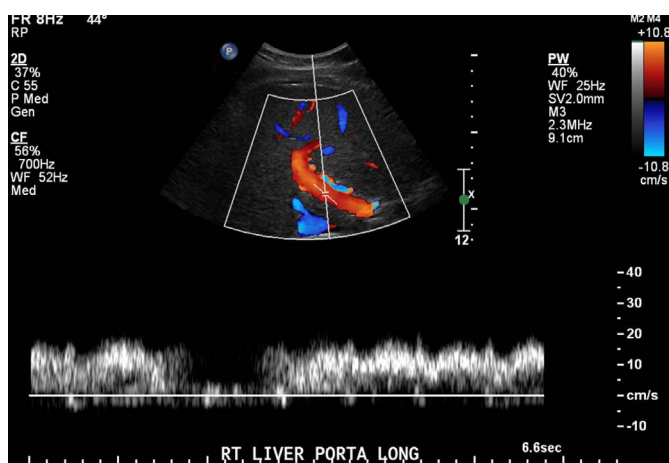
### How to cite this article:

Iranpour P, Lall C, Houshyar R, Helmy M, Yang A, Choi JI, et al. Altered Doppler flow patterns in cirrhosis patients: an overview. Ultrasonography. 2016 Jan;35(1):3-12.

## Portal Venous Flow Patterns in Cirrhosis

### Portal Vein Pulsatility

The normal portal vein waveform normally shows gentle undulations (Fig. 1). In patients with cirrhosis, arterio-portal shunting is a major cause for pulsatility. Increased pulsatility may be measured by calculating the pulsatility index, which is measured as a fraction of the difference between the peak systolic velocity and the end diastolic velocity divided by the average velocity. Increased pulsatility, defined as an increase in the pulsatility index, occurs due to the transmission of pressure across the hepatic sinusoids during the end diastole [3,4]. Alternatively, an increased pulsatility index ( $>0.5$ ) may occur with high right atrial pressure, which can be seen with right heart failure and tricuspid regurgitation. The basis of arterio-portal shunting is the structural distortion that may be seen in patients with cirrhosis in combination with a reversed portal venous flow and increased hepatic venular resistance. Further, hepatocellular carcinoma, which is commonly seen in cirrhotic patients, may cause peri-lesional shunting via draining veins. Liver biopsies performed on cirrhotic patients may also result in arterio-portal fistula formation. Arterio-portal shunting in cirrhosis cases can result in increased portal vein pulsatility (Fig. 2) [5]. Therefore, increased portal vein pulsatility in cirrhotic patients, depending on the clinical history, may prompt an evaluation for the presence of a vascular lesion or arterio-portal fistula. A secondary sign that may help in diagnosing arterio-portal shunts is pulsatility that is in sync with the adjacent hepatic artery waveforms [6].



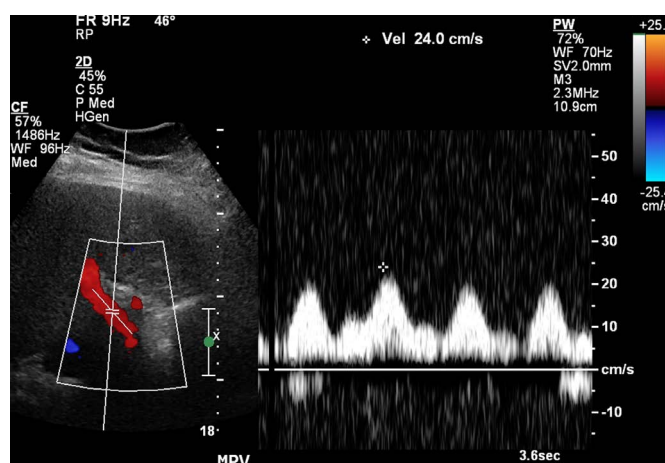
**Fig. 1.** A 35-year-old female with hepatic steatosis. Spectral analysis demonstrates a normal Doppler waveform of the main portal vein. There are gentle waveform undulations, normal peak systolic velocities, and normal hepatopetal flow direction.

### Sluggish or Slow Portal Venous Flow

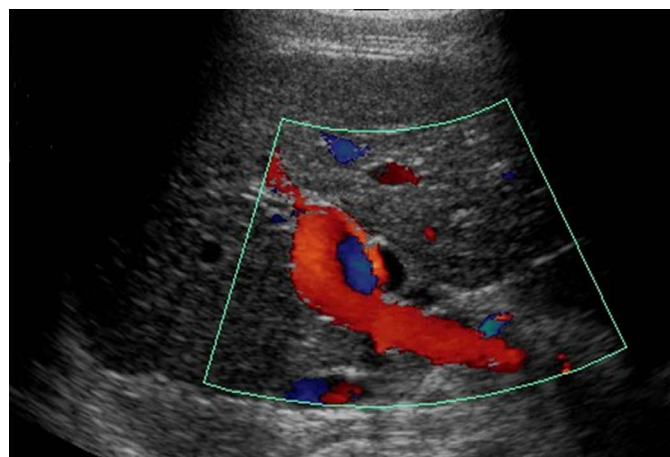
Normal main portal vein (MPV) peak systolic velocities range between 20 cm/sec and 40 cm/sec. A low flow velocity of  $<16$  cm/sec in addition to a caliber increase in the MPV are diagnostic features of portal hypertension [7]. Cirrhosis results in intrahepatic portal hypertension secondary to the increased resistance in hepatic venules caused by the intrahepatic fibrosis [8]. Accurate velocity measurements within the MPV depend on many parameters, including the Doppler/beam-flow angle. False-positive velocities may be secondary to a Doppler angle that is closer to  $90^\circ$  with respect to the flow direction [9]. The choice of Doppler frequency also affects velocity measurements, as higher frequencies produce more accurate velocities while lower frequencies allow for better penetration. Finally, the presence of a turbulent flow and sampling error may result in the under- or overestimation of the flow velocity [10].

### Helical Flow Pattern

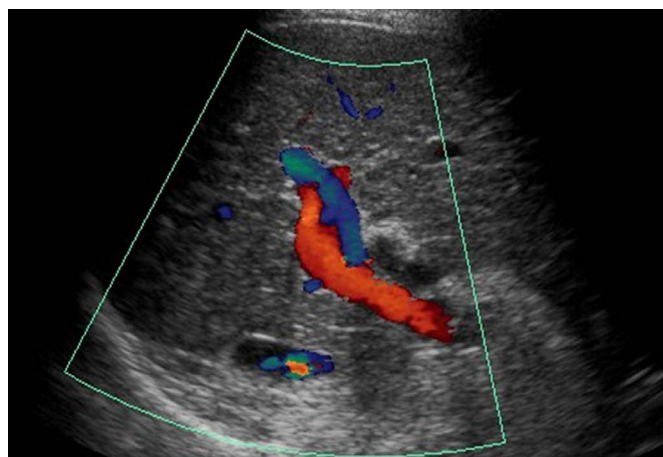
Helical blood flow is a type of secondary flow (defined as a minor flow superimposed on the primary flow pattern). A helical flow is caused by a disturbance in the laminar flow [11,12]. A change in viscosity, vessel geometry, local asymmetries along the vessel wall, and changes in the flow direction and speed can cause accentuated helical patterns [13]. Color Doppler imaging shows alternating red and blue bands in a spiral-like pattern (Fig. 3). Helicity was first characterized within a region of caliber change at the level of the carotid bulb. Studies of the carotid artery indicate that a helical flow may be a marker of exposure to disturbed shear or turbulence [14]. Further, some researchers have hypothesized that this pattern may play a significant role in preventing atherogenesis and fibrointimal



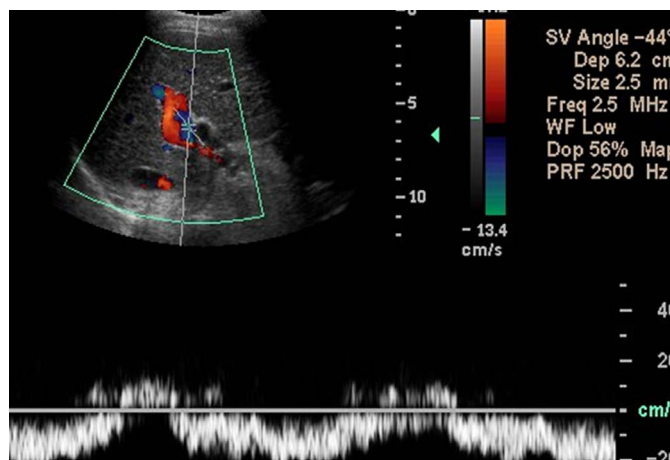
**Fig. 2.** A 79-year-old male with right heart failure. Spectral analysis of the main portal vein (MPV) demonstrates increased pulsatility with an elevated pulsatility index of 1.0 (difference in the peak-to-trough velocities divided by the average velocity [22–9/13]).



A



B



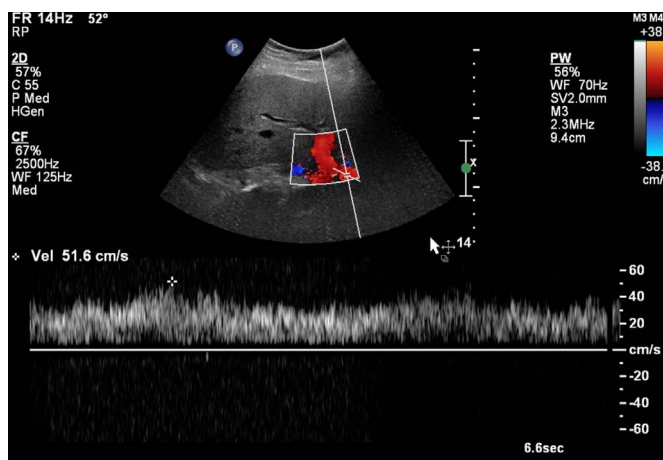
C

**Fig. 3.** A 38-year-old female with liver transplant.

A, B. Color Doppler images demonstrate the helical flow in the main portal vein, with alternating bands of red and blue flows within the main portal vein. C. Spectral analysis shows that there is alternating hepatopetal and hepatofugal flow.

hyperplasia [15].

Helicity, although usually transient, has been seen in both liver transplant and transjugular intrahepatic portosystemic shunt (TIPS) recipients. If there is prolonged persistence of the helical flow, it is associated with an increased portal venous velocity and portal vein stenosis (Fig. 4). The helical flow in transplant patients is due to a change in the portal vein diameter between the donor vessel and the native vessel and is accentuated when the discrepancy between the portal veins is greater than 50%. This is in keeping with other authors who have reported that an expansion of the cross-sectional area of a vessel caliber causes flow separation, inducing a circular pattern [16]. Other studies have shown that shortly after liver transplantation, the helical flow within the portal vein resolves due to the decreased flow velocity and local turbulence [17]. The helical flow has been seen in TIPS recipients as well. In a prospective analysis, the prevalence of the helical flow after TIPS placement was reported to be 28% with a variable timeline after placement. The TIPS procedure changes the hemodynamic parameters of the



**Fig. 4.** A 42-year-old male with hepatitis B. Spectral analysis of the main portal vein demonstrates elevated peak systolic velocity. A peak velocity of 51.6 cm/sec is obtained when the Doppler angle is 52°, which utilizing proper technique is likely accurate.

flow, including the portal vein velocity, flow direction, and flow pulsatility [18]. The amount of turbulence created by these dynamic changes is variable among patients and presumably accounts for the unpredictability of whether a helical flow is present postoperatively.

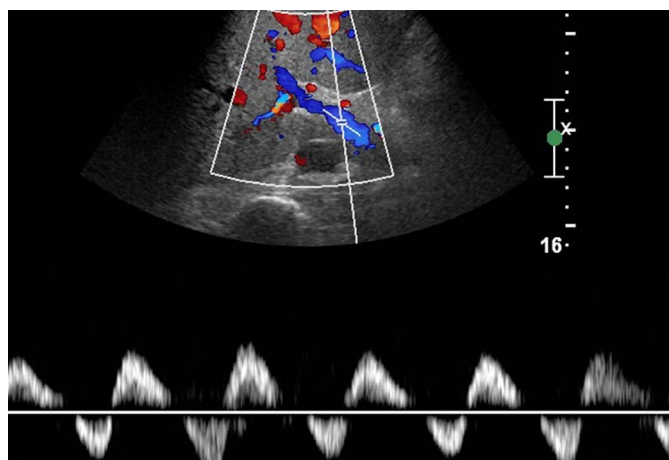
**To-and-Fro Portal Venous Flow Pattern**

With increasing portal venous pressure, there is a progressive decrease in the portal venous flow velocities approaching the level of stagnation. As this occurs, the phenomenon of a to-and-fro flow can be encountered whereby the nearly stagnant blood column in the portal veins is seen to shift into and out of the liver with the respiratory cycle. The effect of transient flow reversal or cessation

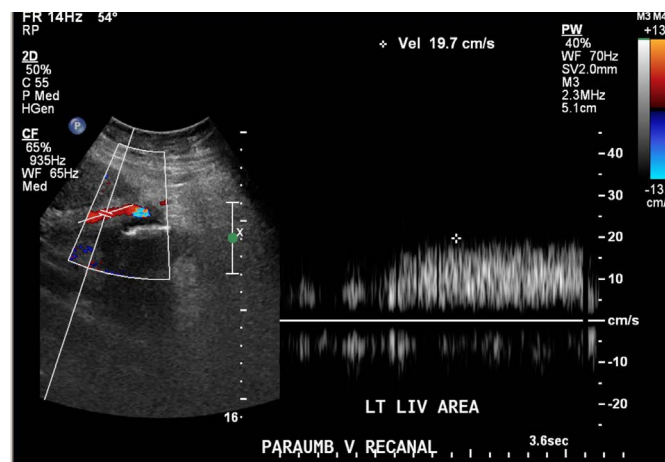
of the forward flow during inspiration can be simulated with the Valsalva maneuver, which also results in a transient hepatofugal flow (Fig. 5). With worsening portal hypertension, stagnation of the blood column can lead to thrombosis or progress to a frank flow reversal.

**Flow Reversal**

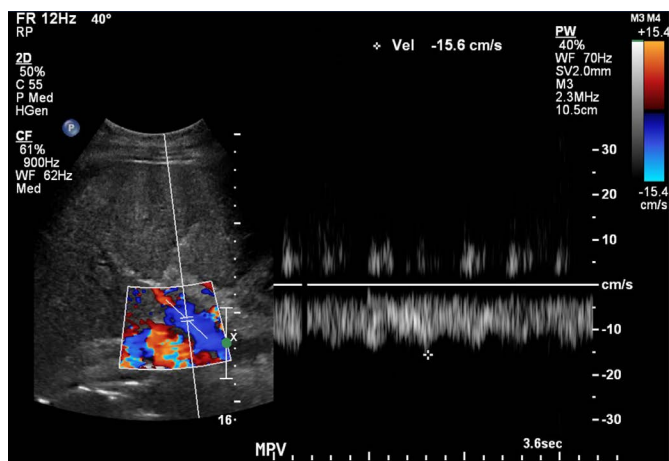
A normal portal venous flow is hepatopetal. A flow reversal (or a hepatofugal flow) is seen in the case of portal hypertension (Fig. 6). In particular, in patients with cirrhosis, obstruction of the hepatic venules and sinusoids by fibrosis, substantiated by arterio-portal and porto-systemic shunting, eventually leads to flow



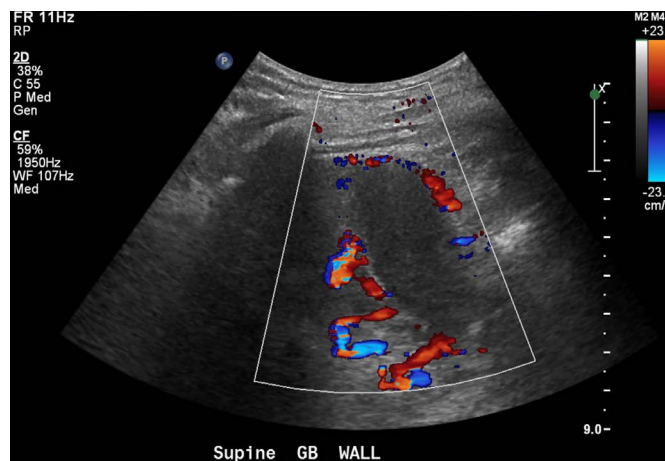
**Fig. 5.** A 52-year-old male with alcoholic cirrhosis. Spectral analysis of the main portal vein demonstrates alternating hepatopetal and hepatofugal flows in a tachypneic patient, compatible with a to-and-fro flow in the main portal vein.



**Fig. 7.** A 47-year-old male with hepatitis C cirrhosis. There is a dilated vessel within the region of the falciform ligament, representing a recanalized paraumbilical vein. Spectral analysis confirms the suspected hepatofugal flow within this vessel.



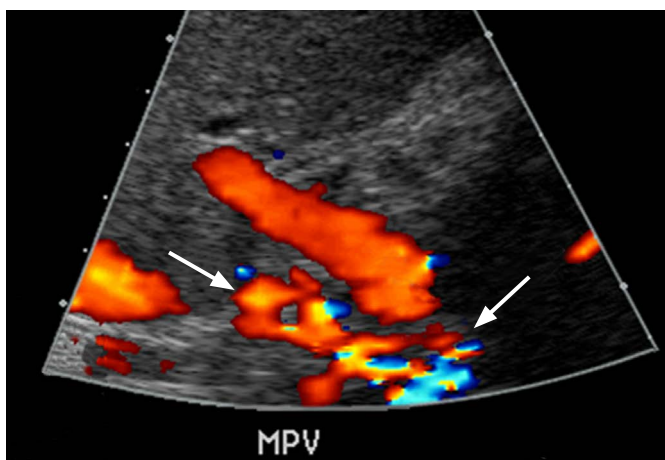
**Fig. 6.** A 43-year-old female with hepatitis C cirrhosis. Spectral analysis of the main portal vein (MPV) demonstrates a reversed hepatofugal (away from the liver) flow.



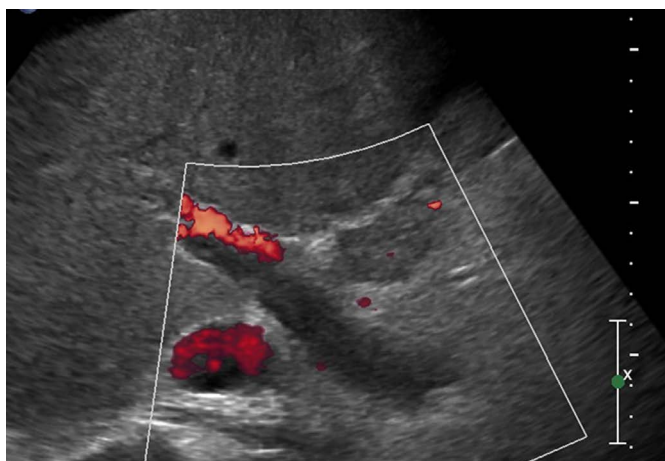
**Fig. 8.** A 39-year-old male with hepatitis B cirrhosis. Color Doppler image demonstrates an increased number of prominent vessels surrounding the gall bladder (GB), representing an example of collateral vasculature in the form of GB wall varices.

reversal. Flow reversal is detected using spectral and color Doppler. Sonographic pitfalls may occur due to the presence of variant portal vein anatomy, resulting in an apparent, false-positive flow reversal depending on the relative location of the ultrasound probe. Atypical intrahepatic arterial collaterals, which are mainly seen after liver transplantation rather than in patients with liver cirrhosis, show a flow from the hepatic capsule to the hilum and may be confused with a hepatofugal flow within the portal venous system [19]. Further, aliasing due to a low pulse repetition frequency setting is a sonographic artifact that may be misinterpreted as a reversed portal venous flow.

Porto-systemic shunting or porto-systemic collateral formation



**Fig. 9.** A 44-year-old male with alcoholic cirrhosis. Color Doppler image of the porta hepatis demonstrates a prominent tangle of vessels deep in the main portal vein (MPV) (arrows), compatible with periportal collaterals.



**A**

**Fig. 10.** A 49-year-old female with hepatitis C cirrhosis.

**A.** Color Doppler image of the main portal vein demonstrates no definite flow within. **B.** However, spectral analysis demonstrates complete stagnation of the portal venous flow.

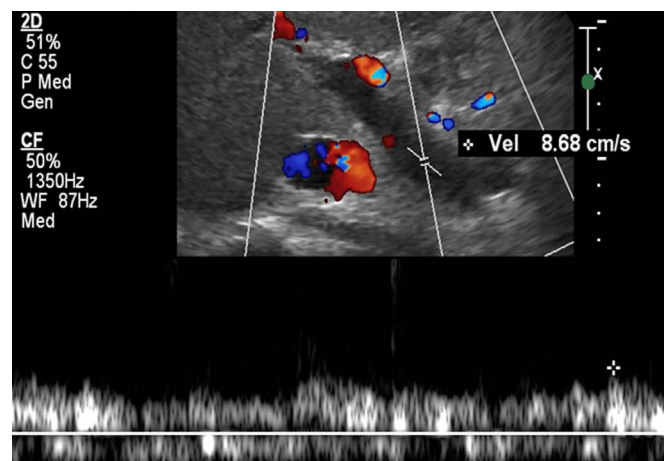
is caused by the opening of normally collapsed porto-systemic anastomoses. Some common porto-systemic collateral vessels include coronary (left gastric) venous collaterals, gastric varices, esophageal and paraesophageal varices, abdominal wall varices, and recanalized paraumbilical veins (Fig. 7), retroperitoneal-paravertebral varices, and pericholecystic (Fig. 8), periportal (Fig. 9), and perisplenic varices. Gastrorenal and splenorenal shunts are other examples of varices [20].

### Stagnant Portal Flow and Thrombosis

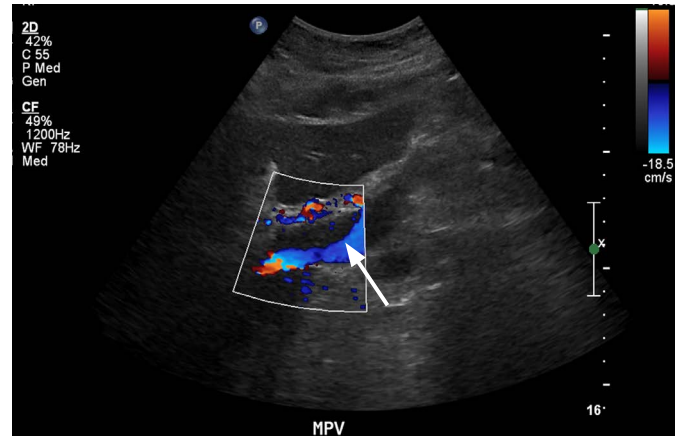
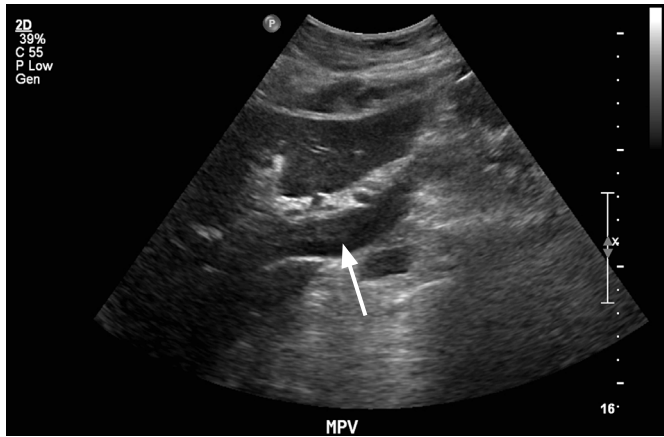
Complete stagnation of portal venous flow is a function of advanced portal hypertension and is associated with severe liver fibrosis such that forward flow is virtually absent (Fig. 10). Further, the portal vein thrombus is usually hypoechoic and can be difficult to discern on grayscale imaging alone. Therefore, evaluation of the MPV on both grayscale and color flow Doppler is necessary (Fig. 11). An apparent absence of flow may also be seen within areas of occlusion secondary to a bland or tumor thrombus. A helpful differentiating factor between a bland thrombus and either a tumor thrombus or recanalization of an occlusive bland thrombus is the identification of whether the blood flow is pulsatile, as in the case of tumor thrombi, or of a waveform similar to that of portal venous flow, as seen in the case of recanalized bland thrombi. The secondary signs of portal vein thrombosis observed by Doppler include the presence of periportal collaterals, representing cavernous transformation, with a flow in the hepatopetal direction (Fig. 12) [21].

### Portal Vein Aneurysm

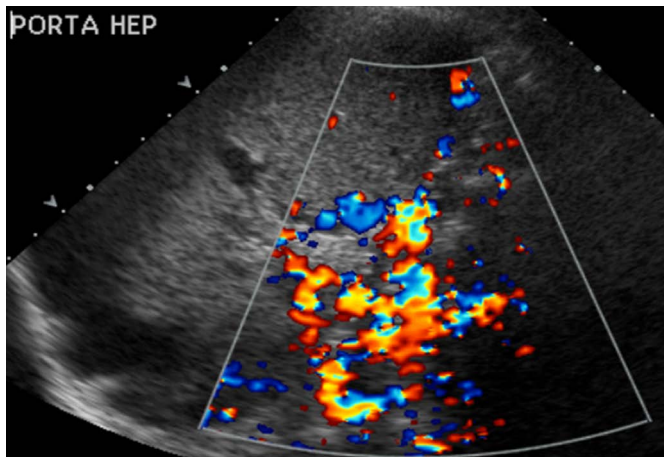
The normal MPV has a caliber of up to 13 mm, and a portal vein



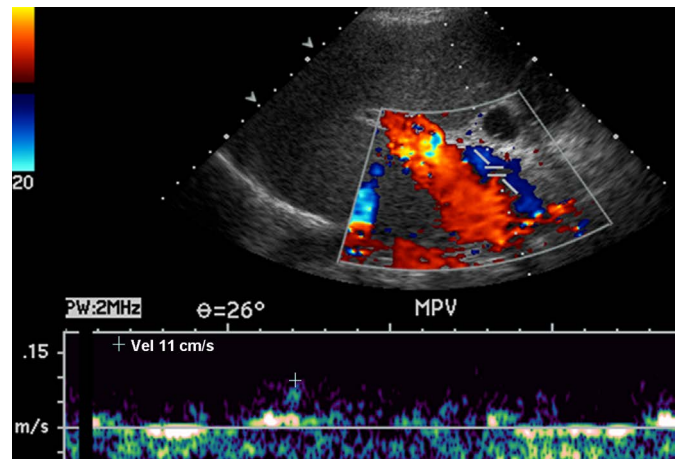
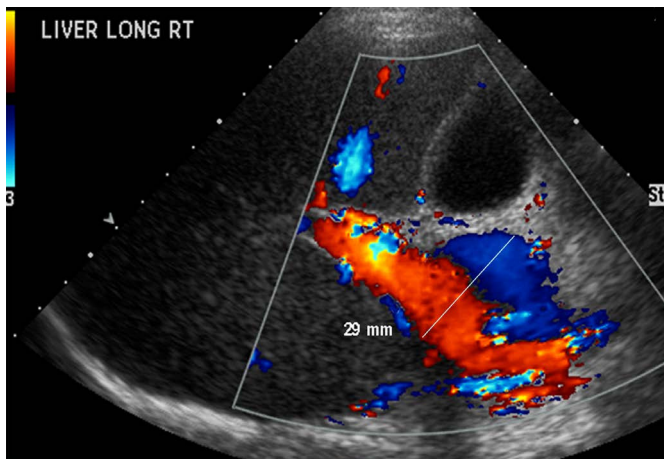
**B**



**A**  
**Fig. 11.** A 53-year-old male with hepatitis C.  
**A.** Grayscale imaging of the porta hepatis demonstrates a barely perceptible hypoechoic thrombus within the main portal vein (MPV) (arrow).  
**B.** Color Doppler imaging clearly demonstrates the partial thrombus within the MPV (arrow).



**Fig. 12.** A 49-year-old male with alcoholic cirrhosis. Color Doppler image of the porta hepatis demonstrates an absence of a normal main portal vein with a tangle of periportal collaterals, compatible with a cavernous transformation in the case of chronic portal vein thrombosis.



**A**  
**Fig. 13.** A 42-year-old male with hepatitis B.  
**A.** Color Doppler image of the porta hepatis demonstrates the focal saccular dilatation of the main portal vein (MPV), measuring 29 mm in diameter with a turbulent or yin-yang flow pattern within the aneurysm. **B.** Spectral analysis demonstrates the to-and-fro flow within the aneurysm.

aneurysm is a focal saccular or fusiform dilatation of the portal venous system. A portal vein aneurysm can be congenital or acquired and occurs due to vein wall weakening with a caliber of at least 20 mm. The acquired causes include portal hypertension, pancreatitis, trauma, and post-surgical etiologies [22]. Grayscale images typically show a focally enlarged MPV, while Doppler imaging shows a turbulent or yin-yang flow within the aneurysm, a stagnant flow, or a thrombus associated with the aneurysm (Fig. 13).

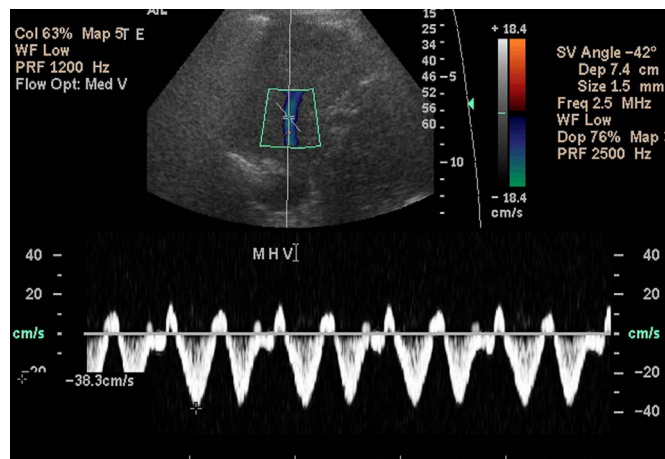
### Pseud thrombosis

An apparent absence of flow may actually be artifactual, secondary to the Doppler settings. For example, a high velocity scale setting may miss velocities below the threshold value, resulting in an apparent absent flow. The transducer probe type (i.e., frequency setting), depth of the vessel, Doppler angle/probe position, or the presence of any overlying, obstructing structure may prevent an accurate velocity measurement, resulting in a false-positive area of absent flow [23]. Therefore, attention to Doppler gain settings, wall filter, frequency of probe used, and Doppler angle of insonation is critical to avoid this issue (Fig. 14).

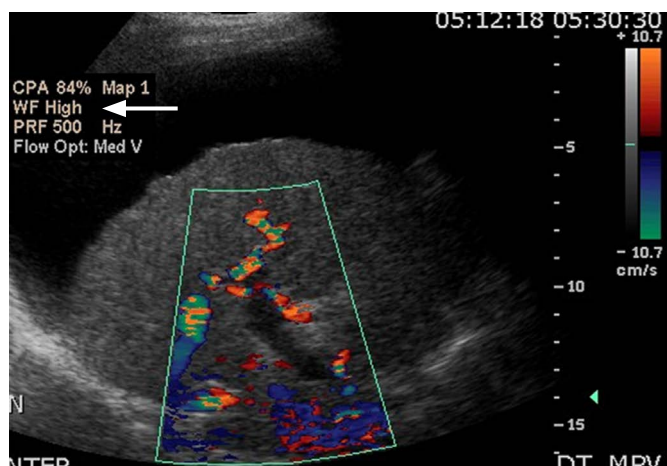
## Hepatic Vein Flow Patterns in Cirrhosis Patients

The normal waveform within the hepatic veins is triphasic with two hepatofugal phases related to the atrial and ventricular diastole (Fig. 15). Fibrotic or inflammatory changes as well as fat deposition

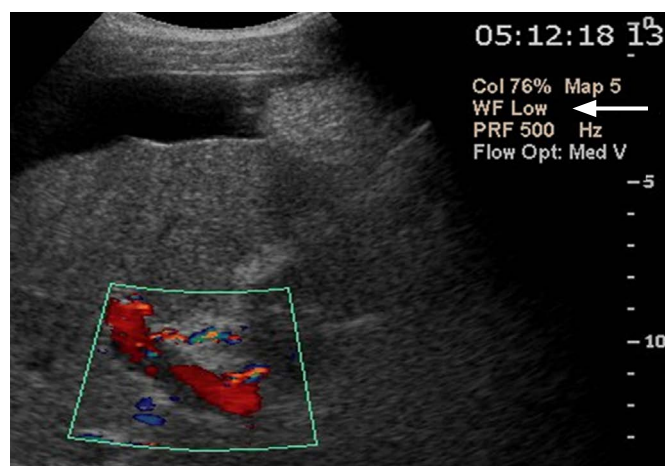
in the liver may create a monophasic flow pattern [24]. In the case of end-stage cirrhosis, distorted architecture with changes in the underlying liver architecture can cause a striking reduction in the caliber or absence of the visualization of the hepatic veins [25]. Early waveform changes in cirrhosis patients include spectral broadening and dampening of the normal, retrograde, pre-systolic wave of the hepatic vein waveform [26]. Later, the normal triphasic waveform pattern may be diminished or replaced with a monophasic pattern



**Fig. 15.** A 33-year-old male with hepatitis C. Spectral analysis of the main hepatic vein (MHV) demonstrates normal triphasic waveforms. The “a” wave corresponds to the atrial contraction, which occurs at the end diastole followed by the “S” wave caused by the motion of the atrioventricular septum during midsystole. The “v” wave corresponds to the opening of the tricuspid valve followed by the “D” wave of the early diastolic right ventricular filling.



**A**  
**Fig. 14.** A 52-year-old female with alcoholic cirrhosis.



**B**  
A. Initial color Doppler imaging of the main portal vein demonstrates no definite flow within the main portal vein; however, note that the wall filter was set to high (arrow). B. Repeat color Doppler with the wall filter set to low (arrow) now demonstrates some degree of flow within the main portal vein. This is an example of pseud thrombosis of the main portal vein.

(Fig. 16). Therefore, the monophasic hepatic vein waveform indicates relatively high portal pressures.

The changes in collaterals may also affect hepatic vein pulsatility. The patency of the paraumbilical vein is a rather frequent finding in cirrhosis patients, which alters the hepatic hemodynamics. The portal venous flow has also been reported to be affected by the patency of the paraumbilical vein, but hepatic arterial resistance and flow are not affected by such patency [27]. A decrease in the phasicity of the hepatic vein is also associated with hepatic vein

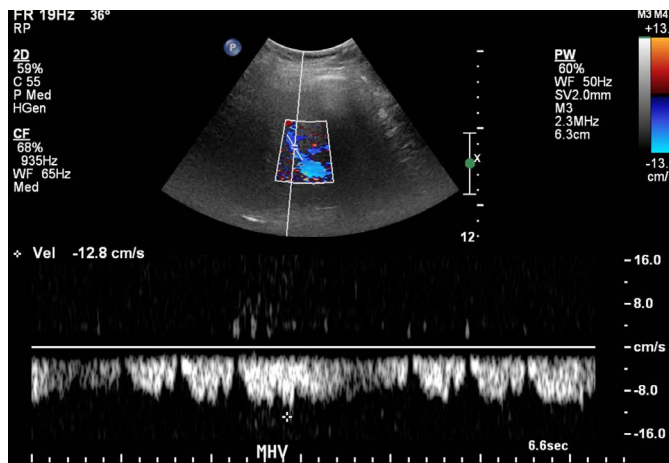
stenosis or thrombosis. There are additional physiologic factors such as high intra-abdominal pressure due to the Valsalva maneuver or suspended respiration in the case of end expiration, which may result in monophasic waveforms.

### Hepatic Arterial Changes

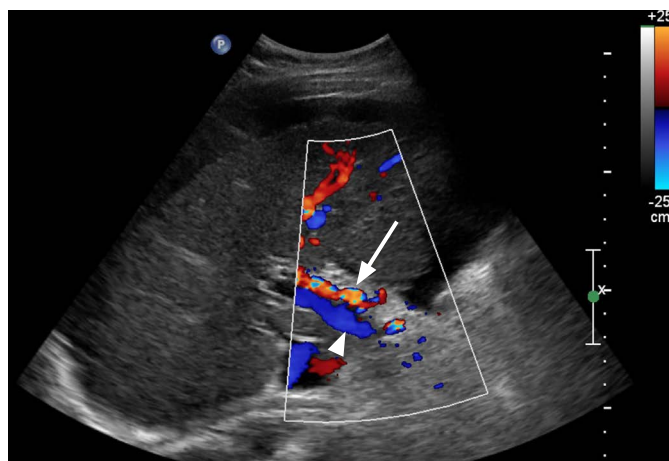
Unlike the portal vein flow patterns in cirrhosis patients, neither the hepatic artery nor the superior mesenteric artery resistive indices correlate with the cirrhosis stage. However, in patients with cirrhosis, hepatic artery compensation occurs when the portal venous flow is compromised [28]. Further, the hepatic arterial flow regulation in cirrhosis patients is maintained via a mechanism called the hepatic arterial buffer response, which increases the flow in this case [29].

### Increased Resistive Index

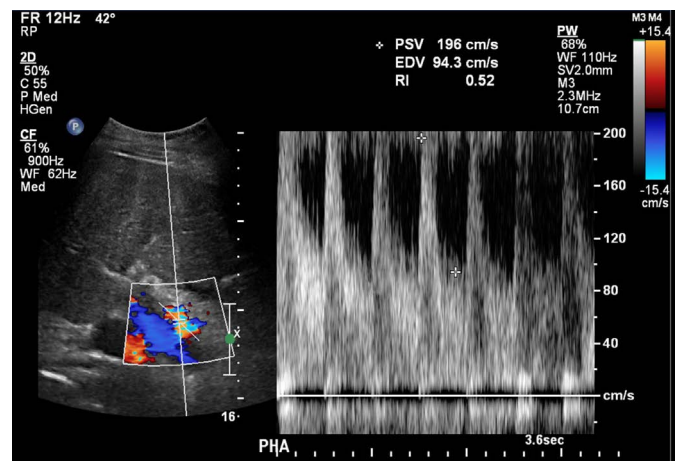
The hepatic arterial resistance changes with increasing portal pressure values [30]. Unfortunately, it has been determined that hepatic arterial resistive indices do not correlate with the actual severity of cirrhosis. It is unclear whether there is a predictable chronological order between changes in the hepatic artery and changes in the portal venous flow [31,32]. The observed specificity and sensitivity of utilizing a hepatic arterial resistive index (RI) cutoff of >0.77 to predict cirrhosis were low at 70% and 68%, respectively [33]. A unique circumstance of the utility of RI is found in the case of a fulminant hepatic failure with acute viral hepatitis. It was found that an RI of >0.74 had an 84% sensitivity and 94% specificity



**Fig. 16.** A 44-year-old male with hepatitis C cirrhosis. Spectral analysis of the middle hepatic vein (MHV) demonstrates dampened monophasic waveforms that are most likely related to portal hypertension in a case of cirrhosis.



**A**  
**Fig. 17.** A 53-year-old male with alcoholic cirrhosis. **A.** Color Doppler of the porta hepatis demonstrates a prominent vessel anterior to the main portal vein, which represents a dilated proper hepatic artery (arrow). There is color aliasing within the hepatic artery, indicating elevated velocities/turbulent flow. A dilated, increased flow within the hepatic artery represents a compensatory mechanism secondary to the lack of hepatic perfusion either by a slow/stagnant flow or by the flow reversal within the main portal vein. In this case, there is a reversed hepatofugal flow within the main portal vein (arrowhead). **B.** Spectral analysis of the proper hepatic artery (PHA) demonstrates markedly elevated velocities.



**B**



for a fulminant hepatic failure in this case [34]. Unfortunately, no predictive values were found in the case of chronic cirrhosis. We know that there is significant variability among RI sampling, which may contribute to the lack of predictability and association with the degree of cirrhosis [35]. RI evaluation in the case of liver transplant rejection was also found to have a poor predictive value of worsening rejection versus improvement [36].

### Increased Hepatic Artery Velocity

In normal patients, the hepatic artery is a low-resistance systemic artery with low-resistance monophasic waveforms. Peak systolic velocities approximate 100 cm/sec. Although not an entirely reliable finding, in the case of portal hypertension or portal vein thrombosis, both of which decrease the portal venous flow within the liver, the arterial flow rate and velocity may increase as a compensatory response (Fig. 17) [3]. Patients with portal vein thrombosis have been shown to have decreased hepatic arterial resistive indices, in some cases less than 0.50. Decreased portal vein flow velocities with the resultant hepatic arterial flow and velocity increase cause a decrease in the liver vascular index, which has been shown to have a high sensitivity and specificity in the diagnosis of cirrhosis and portal hypertension [37].

### Summary

Knowledge of vascular flow patterns in cirrhosis patients provides invaluable information for patients with cirrhosis and portal hypertension. Aberrant portal venous flow patterns have the most significant impact and associative value with cirrhosis given that most of the blood supply is from the portal system. The hepatic arterial flow and the portal venous flow show a correlation in spite of the fact that they are independently regulated. Flow pattern changes within the portal side do not seem to develop in a chronological order. However, in a cirrhotic patient undergoing a routine imaging follow-up, the radiologist is in a perfect position to monitor pattern changes, which may show hemodynamic alterations, indicative of worsening or improving liver function.

ORCID: Pooya Iranpour: <http://orcid.org/0000-0002-4436-6974>; Roozbeh Houshyar: <http://orcid.org/0000-0003-4263-5533>; Joon-Il Choi: <http://orcid.org/0000-0003-0018-8712>

### Conflict of Interest

No potential conflicts of interest relevant to this article was reported.

### References

1. Tsochatzis EA, Bosch J, Burroughs AK. Liver cirrhosis. *Lancet* 2014;383:1749-1761.
2. Tchelepi H, Ralls PW, Radin R, Grant E. Sonography of diffuse liver disease. *J Ultrasound Med* 2002;21:1023-1032.
3. McNaughton DA, Abu-Yousef MM. Doppler US of the liver made simple. *Radiographics* 2011;31:161-188.
4. Abu-Yousef MM, Milam SG, Farnier RM. Pulsatile portal vein flow: a sign of tricuspid regurgitation on duplex Doppler sonography. *AJR Am J Roentgenol* 1990;155:785-788.
5. Mori K, Yoshioka H, Itai Y, Okamoto Y, Mori H, Takahashi N, et al. Arterioportal shunts in cirrhotic patients: evaluation of the difference between tumorous and nontumorous arterioportal shunts on MR imaging with superparamagnetic iron oxide. *AJR Am J Roentgenol* 2000;175:1659-1664.
6. Bhargava P, Vaidya S, Kolokythas O, Katz DS, Dighe M. Pictorial review: hepatic vascular shunts: embryology and imaging appearances. *Br J Radiol* 2011;84:1142-1152.
7. Al-Nakshabandi NA. The role of ultrasonography in portal hypertension. *Saudi J Gastroenterol* 2006;12:111-117.
8. Berzigotti A, Seijo S, Reverter E, Bosch J. Assessing portal hypertension in liver diseases. *Expert Rev Gastroenterol Hepatol* 2013;7:141-155.
9. Park MY, Jung SE, Byun JY, Kim JH, Joo GE. Effect of beam-flow angle on velocity measurements in modern Doppler ultrasound systems. *AJR Am J Roentgenol* 2012;198:1139-1143.
10. Tola M, Yurdakul M. Effect of Doppler angle in diagnosis of internal carotid artery stenosis. *J Ultrasound Med* 2006;25:1187-1192.
11. Bathurst JC, Thorne CR, Hey RD. Secondary flow and shear stress at river bends. *J Hydraul Div* 1979;105:1277-1295.
12. Gijsen FJ, van de Vosse FN, Janssen JD. The influence of the non-Newtonian properties of blood on the flow in large arteries: steady flow in a carotid bifurcation model. *J Biomech* 1999;32:601-608.
13. Morbiducci U, Ponzini R, Rizzo G, Cadioli M, Esposito A, Montevicchi FM, et al. Mechanistic insight into the physiological relevance of helical blood flow in the human aorta: an in vivo study. *Biomech Model Mechanobiol* 2011;10:339-355.
14. Gallo D, Steinman DA, Bijari PB, Morbiducci U. Helical flow in carotid bifurcation as surrogate marker of exposure to disturbed shear. *J Biomech* 2012;45:2398-2404.
15. Morbiducci U, Ponzini R, Grigioni M, Redaelli A. Helical flow as fluid dynamic signature for atherogenesis risk in aortocoronary bypass: a numeric study. *J Biomech* 2007;40:519-534.
16. White SS, Zarins CK, Giddens DP, Bassiouny H, Loth F, Jones SA, et al. Hemodynamic patterns in two models of end-to-side vascular graft anastomoses: effects of pulsatility, flow division, Reynolds number, and hood length. *J Biomech Eng* 1993;115:104-111.
17. Rosenthal SJ, Harrison LA, Baxter KG, Wetzel LH, Cox GG, Batnitzky

- S. Doppler US of helical flow in the portal vein. *Radiographics* 1995;15:1103-1111.
18. De Gaetano AM, Rinaldi P, Barbaro B, Mirk P, Di Stasi C, Gui B, et al. Intrahepatic portosystemic venous shunts: color Doppler sonography. *Abdom Imaging* 2007;32:463-469.
  19. Herrmann J, Herden U, Ganschow R, Petersen KU, Schmid F, Derlin T, et al. Transcapsular arterial neovascularization of liver transplants increases the risk of intraoperative bleeding during retransplantation. *Transpl Int* 2013;26:419-427.
  20. Hwang HJ, Kim KW, Choi EK, Jeong WK, Kim PN, Kim SA, et al. Hepatofugal portal flow on Doppler sonography in various pathological conditions: a pictorial essay. *J Clin Ultrasound* 2009;37:511-524.
  21. Piscaglia F, Gianstefani A, Ravaioli M, Golfieri R, Cappelli A, Giampalma E, et al. Criteria for diagnosing benign portal vein thrombosis in the assessment of patients with cirrhosis and hepatocellular carcinoma for liver transplantation. *Liver Transpl* 2010;16:658-667.
  22. Koc Z, Oguzkurt L, Ulasan S. Portal venous system aneurysms: imaging, clinical findings, and a possible new etiologic factor. *AJR Am J Roentgenol* 2007;189:1023-1030.
  23. Kruskal JB, Newman PA, Sammons LG, Kane RA. Optimizing Doppler and color flow US: application to hepatic sonography. *Radiographics* 2004;24:657-675.
  24. Dietrich CF, Lee JH, Gottschalk R, Herrmann G, Sarrazin C, Caspary WF, et al. Hepatic and portal vein flow pattern in correlation with intrahepatic fat deposition and liver histology in patients with chronic hepatitis C. *AJR Am J Roentgenol* 1998;171:437-443.
  25. Stark DD, Hahn PF, Trey C, Clouse ME, Ferrucci JT Jr. MRI of the Budd-Chiari syndrome. *AJR Am J Roentgenol* 1986;146:1141-1148.
  26. Barakat M. Non-pulsatile hepatic and portal vein waveforms in patients with liver cirrhosis: concordant and discordant relationships. *Br J Radiol* 2004;77:547-550.
  27. Sacerdoti D, Bolognesi M, Bombonato G, Gatta A. Paraumbilical vein patency in cirrhosis: effects on hepatic hemodynamics evaluated by Doppler sonography. *Hepatology* 1995;22:1689-1694.
  28. Le Couteur DG, Hickey H, Harvey PJ, Greedy J, McLean AJ. Hepatic artery flow and propranolol metabolism in perfused cirrhotic rat liver. *J Pharmacol Exp Ther* 1999;289:1553-1558.
  29. Gulberg V, Haag K, Rossle M, Gerbes AL. Hepatic arterial buffer response in patients with advanced cirrhosis. *Hepatology* 2002;35:630-634.
  30. Schneider AW, Kalk JF, Klein CP. Hepatic arterial pulsatility index in cirrhosis: correlation with portal pressure. *J Hepatol* 1999;30:876-881.
  31. Sacerdoti D, Merkel C, Bolognesi M, Amodio P, Angeli P, Gatta A. Hepatic arterial resistance in cirrhosis with and without portal vein thrombosis: relationships with portal hemodynamics. *Gastroenterology* 1995;108:1152-1158.
  32. Vassiliades VG, Ostrow TD, Chezmar JL, Hertzler GL, Nelson RC. Hepatic arterial resistive indices: correlation with the severity of cirrhosis. *Abdom Imaging* 1993;18:61-65.
  33. Pierce ME, Sewell R. Identification of hepatic cirrhosis by duplex doppler ultrasound value of the hepatic artery resistive index. *Australas Radiol* 1990;34:331-333.
  34. Tanaka K, Numata K, Morimoto M, Shirato K, Saito S, Imada T, et al. Elevated resistive index in the hepatic artery as a predictor of fulminant hepatic failure in patients with acute viral hepatitis: a prospective study using Doppler ultrasound. *Dig Dis Sci* 2004;49:833-842.
  35. Paulson EK, Kliewer MA, Frederick MG, Keogan MT, DeLong DM, Nelson RC. Hepatic artery: variability in measurement of resistive index and systolic acceleration time in healthy volunteers. *Radiology* 1996;200:725-729.
  36. Marder DM, DeMarino GB, Sumkin JH, Sheahan DG. Liver transplant rejection: value of the resistive index in Doppler US of hepatic arteries. *Radiology* 1989;173:127-129.
  37. Iwao T, Toyonaga A, Oho K, Tayama C, Masumoto H, Sakai T, et al. Value of Doppler ultrasound parameters of portal vein and hepatic artery in the diagnosis of cirrhosis and portal hypertension. *Am J Gastroenterol* 1997;92:1012-1017.