



TIPS – All that Non-interventional Radiologists Should Know

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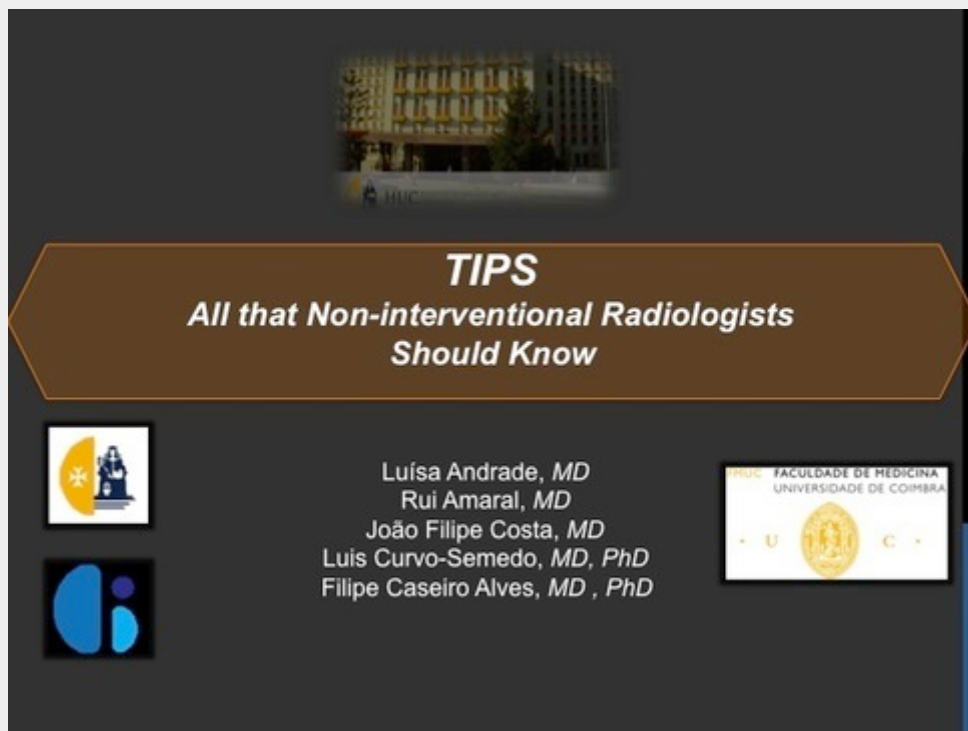
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1. Learning Objectives

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Learning Objectives

LEARNING OBJECTIVES

- To present a review on TIPS (Transjugular Intrahepatic Portosystemic Shunt): indications, contraindications, technique and complications.
- To describe and explain the pre and post-procedure imaging evaluation using mainly Doppler ultrasound and the signs for direct and indirect evidence of TIPS failure.
- To discuss the technique, focusing on tips and tricks.

2. Background

Background

BACKGROUND

- TIPS is a percutaneous imaging-guided procedure in which a channel is constructed within the liver with the intent of reducing portal pressure by diverting blood from the portal to the systemic circulation.
- The parenchymal tract created between the portal vein and hepatic veins is reinforced with placement of a stent graft.
- This procedure is used to treat severe complications of portal hypertension such as refractory ascites and variceal bleeding and can act as a bridge to liver transplantation.

3. Imaging Findings/Procedure Details


Imaging Findings

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LIVER DOPPLER ULTRASOUND – Normal Findings

> LIVER VASCULAR ANATOMY

- 1. PORTAL VEIN** – Supplies 70% of incoming blood to the liver (relatively desoxygenated blood but rich in nutrients from the stomach, intestine, spleen and pancreas)
 - Formed by the confluence of the splenic and superior mesenteric veins.
 - Accompanied by the hepatic artery and common bile duct to form the portal triad.
 - The main portal vein (MPV) enters the liver at the porta hepatis and divides into the right and left portal veins.
 - The MPV diameter does not exceed 13mm in quiet respiration (measured at the site where the portal vein crosses the IVC). Should be normally seen an increased in this diameter, with deep inspiration.



Two B-mode ultrasound images of the portal vein. The top image is labeled 'Quiet respiration' and shows a portal vein with a diameter of 10.4 mm. The bottom image is labeled 'Normal elevation of the diameter of the portal vein during deep inspiration' and shows a portal vein with a diameter of 12.3 mm.

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LIVER DOPPLER ULTRASOUND – Normal Findings

➤ LIVER VASCULAR ANATOMY

2. HEPATIC ARTERY – Supplies 30% of incoming blood to the liver (oxygenated blood)

- The common hepatic artery originates in the majority of patients from the celiac artery.
- After the origin of the gastroduodenal artery it is called the proper hepatic artery.
- It enters the liver alongside and anterior to the portal vein, where it divides into left and right hepatic arteries.
- There are numerous variants of hepatic artery anatomy.

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LIVER DOPPLER ULTRASOUND – Normal Findings

➤ LIVER VASCULAR ANATOMY

3. HEPATIC VEINS – hepatic venous drainage into inferior vena cava (IVC)

- They converge into IVC approximately 1cm below its confluence with the right atrium.
- In most people the right, middle and left hepatic veins enter the IVC in a "crows foot" configuration (transverse plane).
- 30% of individuals have additional hepatic veins.
- The walls of the hepatic veins are relatively hypoechoic, which helps to differentiate them from the portal veins in the more echogenic portal triads.



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LIVER DOPPLER ULTRASOUND – Normal Findings

- The right hepatic vein (RHV) is the largest of hepatic veins and courses between the anterior and posterior segments of the right lobe and drains into the right postero-lateral aspect of the IVC.
- The middle hepatic vein (MHV) courses along the major lobar fissure and drains the anterior right lobe segments (V, VIII) and segment IVb. In 85% of people the MHV forms a common trunk with the left hepatic vein (LHV), which then drains into the left antero-lateral aspect of the IVC.
- The LHV is the smallest of the hepatic veins and drains the lateral segments of the left lobe (II & III) and segment IVa.
- The venous drainage from the central portion of the liver (including caudate lobe) empties directly into the IVC (and cannot be perceived by color Doppler).

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LIVER DOPPLER ULTRASOUND – Normal Findings

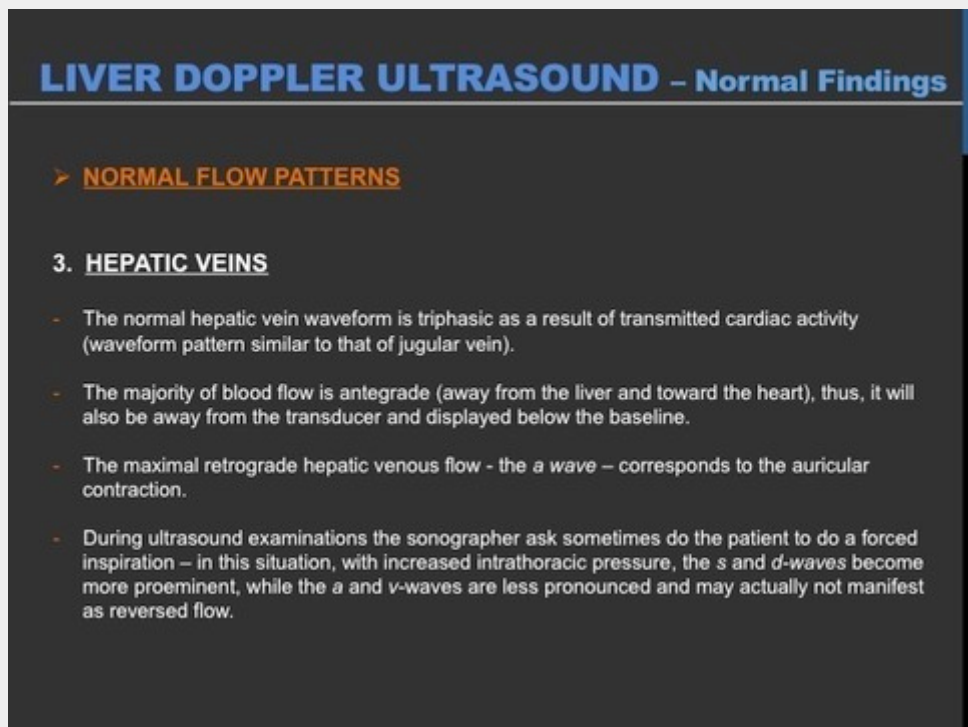
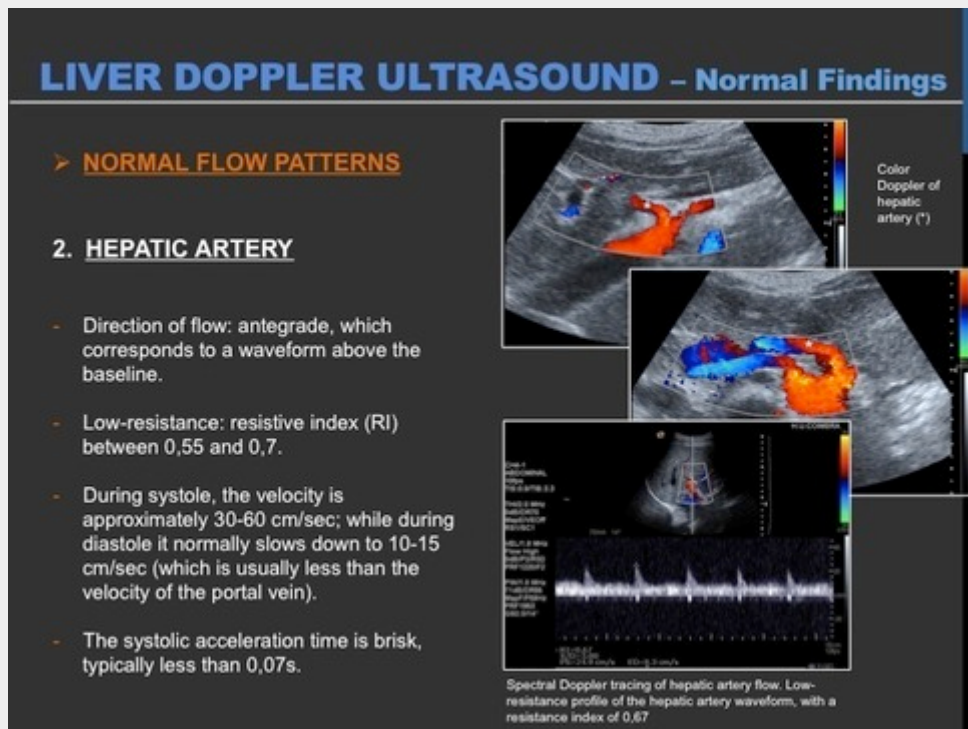
➤ NORMAL FLOW PATTERNS

1. PORTAL VEIN

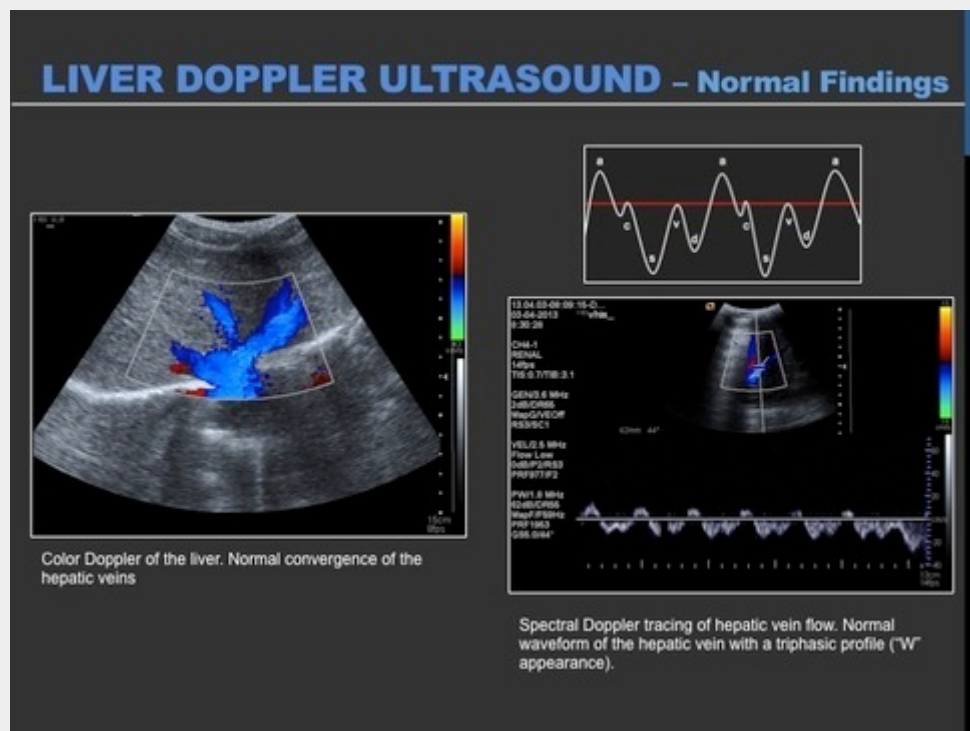
- Antegrade flow (toward the transducer), therefore with a waveform that is above the baseline.
- Hepatopetal flow (towards the liver) throughout the entire cardiac cycle.
- Portal flow velocity varies with cardiac activity and respiration, giving the portal waveform an undulating appearance.
- Mean flow velocity relatively low (but >16cm/s)



Spectral Doppler tracing of portal vein flow. Hepatopetal portal vein flow, with an undulating appearance and a mean velocity > 16 cm/sec.



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PORTAL HYPERTENSION

- Portal hypertension accompanies cirrhosis of the liver in over 90% of the cases.
- Cirrhosis disrupts the normal hepatic sinusoidal pattern, impairing venous drainage and leading to increase in portal vein pressure.
- Portal hypertension is the increased in porto-systemic pressure gradient in any portion of the portal venous system.

Porto-Systemic Pressure Gradient:

- This gradient is assessed by measuring the wedge hepatic venous pressure (a measure of sinusoidal hepatic pressure) and subtracting the free hepatic venous pressure (systemic pressure) thus obtaining the hepatic venous pressure gradient (HVPG).
- Normal: HVPG = 2-5mmHg.
- Sub-clinically Portal Hypertension: HVPG > 6mmHg
- Clinically significant Portal Hypertension: **HVPG > 10-12 mmHg** (predicts clinical course in patients with cirrhosis including development of varices, variceal hemorrhage and encephalopathy, decompensation or death after liver resection, and hepatocellular carcinoma).

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PORTAL HYPERTENSION

Pathophysiology:

Two important factors - vascular resistance and blood flow - exist in the development of portal hypertension:

$$P = FR$$

Where P = pressure gradient through the portal venous system; F = volume of blood flowing through the portal venous system; R = resistance to the flow.

Changes in either F or R affect the pressure. In most types of portal hypertension, both the blood flow and the resistance are altered.

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PORTAL HYPERTENSION - Etiology			
INCREASED RESISTANCE TO FLOW	PREHEPATIC	Portal or splenic vein thrombosis or extrinsic compression (tumors) Splanchnic arteriovenous fistula	
	HEPATIC	Presinusoidal	Schistosomiasis Primary biliary cirrhosis Idiopathic portal hypertension Granulomatous diseases Myeloproliferative diseases Polycystic disease Hepatic metastases
		Sinusoidal	<u>Cirrhosis</u> of various causes (alcoholic and related to hepatitis virus B and C; Wilson disease; Hemochromatosis) Acute and fulminant hepatitis Schistosomiasis, Primary biliary cirrhosis and Idiopathic portal hypertension in advanced stages Peliosis hepatitis Budd-Chiari syndrome Sclerosing cholangitis
		Postsinusoidal	Veno-occlusive disease
INCREASED VASCULAR FLOW	POSTHEPATIC	Budd-Chiari Syndrome and thrombosis of the IVC, Cardiac disease (right sided heart failure, constrictive pericarditis and severe tricuspid regurgitation), Trauma	
	Arteriportal shunts / fistulas (congenital, traumatic, iatrogenic, tumor) Increased portal blood flow Increased splenic flow		

Doppler Ultrasound – Cirrhosis and Portal Hypertension

In patients with cirrhosis and indication to TIPS the radiologist should evaluate the following parameters:

1. Hepatic size and morphology:

- In patients with advanced cirrhosis the liver is small with relative enlargement of caudate lobe and left lobe relatively to the right lobe (in early stages of cirrhosis the liver is enlarged or could appear absolutely normal).



US demonstrates a dysmorphic liver with coarse appearance and irregular borders. This patient has cirrhosis and portal hypertension complicated with ascites.

2. Liver echogenicity and texture:

- Echogenicity could be increased in cirrhotic livers with fat infiltration.
- Irregular borders and nodular surface (in relation to the presence of regenerating nodules and fibrosis).
- Exclude the presence of definable nodules (dysplastic nodules – portal venous supply vs. HCC – arterial hepatic venous supply).
- Exclude the presence of multiple cysts (polycystic liver disease precludes the TIPS technique).



Cirrhotic liver. The hyperechoic nodule proved to be a small hepatocarcinoma.

Doppler Ultrasound – Cirrhosis and Portal Hypertension

In cirrhosis, most of the normal liver architecture is replaced by distorted vascular channels that provide increased resistance to portal venous blood flow and obstruction to hepatic venous outflow.

3. Portal vein:

- Diameter:

- Although the caliber of the portal vein initially may be increased (>13mm) in PH, with the development of porto-systemic shunts, the portal vein caliber will decrease.
- Evaluate the response to respiratory manoeuvres: if the diameter of portal vein does not increase at least 20% from quiet respiration to deep inspiration the patient has probably PH.



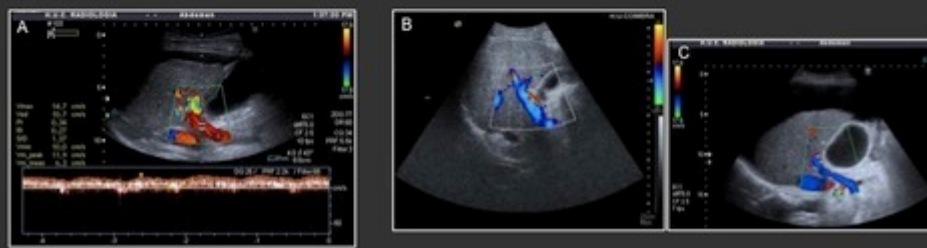
B mode US shows dilatation of the portal vein (A) and normal portal vein diameter (B) after the development of varices.

Doppler Ultrasound – Cirrhosis and Portal Hypertension

Permeability, flow direction and velocity:

- In severe cirrhosis portal venous flow may become continuous (without undulating appearance) or pulsatile due to arteriovenous shunting;
- Portal flow velocity decreases ($< 16\text{cm/s}$) (A);
- Flow may become to and fro (bidirectional) or hepatofugal (retrograde) (B and C). As with slow flow, this finding is diagnostic for portal hypertension from whatever cause.
- A narrow portal vein and a prominent hepatic artery are commonly associated findings when flow is hepatofugal in the main portal vein.

However, we should always kept in mind that, the flow direction in the portal vein is a variable finding in PH and depends on the development of collaterals.



Doppler Ultrasound – Cirrhosis and Portal Hypertension

- Absent (aphasic) portal venous flow may be due to stagnant flow (portal hypertension) or occlusive disease, usually caused by bland or malignant thrombosis.
- In the setting of subacute/chronic occlusive portal vein thrombosis could be development of collateral vessels in or around the occluded portal vein (*cavernous transformation of portal vein*).



(A). Absent flow in the portal vein (RPV) that is markedly dilated due to malignant portal vein thrombosis.



A normal portal vein could not be visualised. Instead there is a plexus of small vessels - anechoic serpiginous structures (arrow). After portal vein thrombosis this web of small vessels reconstitutes portal flow to the liver and is known as cavernomatous transformation of the portal vein.

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Doppler Ultrasound – Cirrhosis and Portal Hypertension

4. Hepatic Artery: (A)

- In patients with PH and decreased portal flow velocity, the hepatic artery appears enlarged with increased flow to compensate for the diminished portal vein flow.
- In patients with cirrhosis the resistive index in the hepatic artery is elevated (>0.7); with development of portal hypertension and arteriovenous/arterioportal shunts the RI decreases (<0.55).

5. Hepatic Veins:

- These veins become compressed and narrowed in cirrhotic patients (B).
- With PH they lose the classic pulsatility and turn into a biphasic or monophasic waveform (C).




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Doppler Ultrasound – Cirrhosis and Portal Hypertension

6. Spleen

- Access spleen greatest dimension – in the setting of PH there could be mild to moderate splenomegaly ($>13\text{cm}$).
- Reversed flow in the splenic vein (towards the spleen). Indirect sign of splenogastric or splenorenal collaterals, and thus PH.

7. Presence / absence of ascites and volume quantification

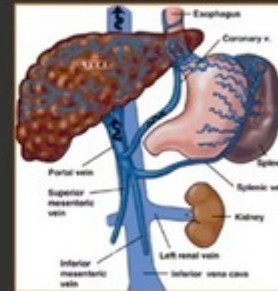


Doppler Ultrasound – Cirrhosis and Portal Hypertension

8. Porto-systemic venous collaterals – shunting of blood from the portal to the systemic circulation.

- **Short gastric varices:** courses between the spleen and the greater curvature of the stomach.
- **Left gastric varices (coronary varices):** courses from the splenic or portal veins to the lesser curvature of the stomach. Dilatation of the coronary vein ($>7\text{mm}$) is associated with severe PH (HVPG $>10\text{mmHg}$).

These veins communicate with systemic esophageal veins and are known as gastroesophageal varices.



- **Paraumbilical veins:** originates from the left portal vein and runs inferiorly through the falciform ligament towards the anterior abdominal wall where it connects with the systemic epigastric veins or with the subcutaneous veins originating the *caput medusae*.
- **Splenorenal-mesenteric collaterals:** courses between the splenic hilum and the left renal vein.

Doppler Ultrasound – Cirrhosis and Portal Hypertension



(A) US scan of the spleen. Tangle of vessels are seen at the upper pole of the spleen, corresponding to short gastric varices.



(C) Color Doppler image shows a recanalized paraumbilical vein that carries blood towards the umbilicus.



(B) Longitudinal US image shows a tortuous left gastric varix coursing from the region of celiac axis towards the gastro-esophageal junction.




(D) US scan of the spleen revealed anechoic tubular structures adjacent to the lower pole of the spleen, communicating to the left renal vein (not shown).

(E) A large varix occupies the space between the splenic hilum (s) and the left kidney (k).



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Doppler Ultrasound – Cirrhosis and Portal Hypertension



Pre-TIPS ultrasound Doppler evaluation:

Detecting *stigmas* of cirrhosis and portal hypertension in the liver and hepatic vasculature

Exclude hepatic cysts

Detecting complications of portal hypertension:

- Neoplastic processes – HCC
- Portal vein thrombosis ± cavernous transformation
- Splenomegaly
- Ascites
- Collateral pathways

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TRANSJUGULAR INTRAHEPATIC PORTOSYSTEMIC SHUNT - TIPS

DEFINITION

- Percutaneous imaging-guided method in which a channel is constructed within the liver with the intent of reducing portal pressure by diverting blood from the portal to the systemic circulation.
- Intrinsically functions as a side-to-side porta-caval shunt, designed to function as a partial shunt that preserves a portion of portal flow to the liver.

OBJECTIVE

- Reduce portal hypertension in patients with complications related to portal hypertension, such as refractory ascites and refractory variceal bleeding.


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TIPS - Indications

- **Refractory ascites** (the most frequent indication for TIPS)
- **Secondary prevention of variceal bleeding** (second-line therapy after failure of medical and endoscopic therapy)
- Refractory acute variceal bleeding
- Portal hypertensive gastropathy
- Hepatorrenal syndromes
- Bleeding gastric / ectopic varices
- Refractory hepatic hydrothorax
- Budd-Chiari syndrome
- Hepatic veno-occlusive disease
- Hepatopulmonary syndrome
- In centers with extensive experience TIPS should be considered in patients with portal vein thrombosis with or without cavernomatous transformation, when intrahepatic portal venous branches are patent.

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TIPS – Patient Selection



- Patients with a Child-Pugh score of >12 are in high risk of post-procedural death.
- MELD scores of $>15-18$ or a bilirubin level of $>3.5\text{mg/dl}$ have poor prognosis and TIPS should only be performed in the absence of any other treatment possibilities.

TIPS – Contra-Indications

Absolute

- Unproved portal hypertension
- Primary prevention of variceal bleeding
- Congestive heart failure
- Polycystic liver disease
- Uncontrolled systemic infection or sepsis
- Unrelieved biliary obstruction
- Severe pulmonary hypertension
- Severe tricuspid regurgitation

Relative:

- Hepatocarcinoma large and central
- Hepatic venous obstruction
- Portal vein thrombosis
- Severe coagulopathy (INR>5)
- Thrombocytopenia (<20 000)
- Moderate pulmonary hypertension
- Hepatic encephalopathy poorly controlled

TIPS – Technique

1. Jugular access (right internal jugular vein) – Seldinger technique

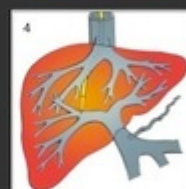
2. A catheter is placed and guided through the superior vena cava, right atrium and inferior vena cava into a hepatic vein. (Because of large size and consistent and predictable relation to the portal veins, the right hepatic vein is the preferred hepatic vein branch for the creation of a TIPS shunt).

3. Selective catheterization of a suitable branch of the RHV

(Pressure measurements are performed at this point, including central venous pressures, at the level of right atrium and IVC and free and wedged hepatic venous pressures).

4. A needle inserted through the catheter is then used to puncture the liver from a central portion of the hepatic vein and enter the main portal branch (usually the right portal vein).

The puncture could be navigated with ultrasound.




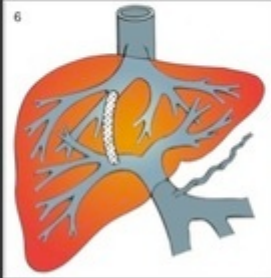

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TIPS – Technique

5. Dilatation of the needle tract by a balloon catheter – establishing of a connection between the portal and systemic circulation directly inside the liver parenchyma.


6. Insertion of a TIPS stent graft

7. The final step of the TIPS procedure is placement of a pigtail catheter over the portal vein guide wire for follow up portography and pressure measurements.




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TIPS – Technique



Suprahepatic flebography

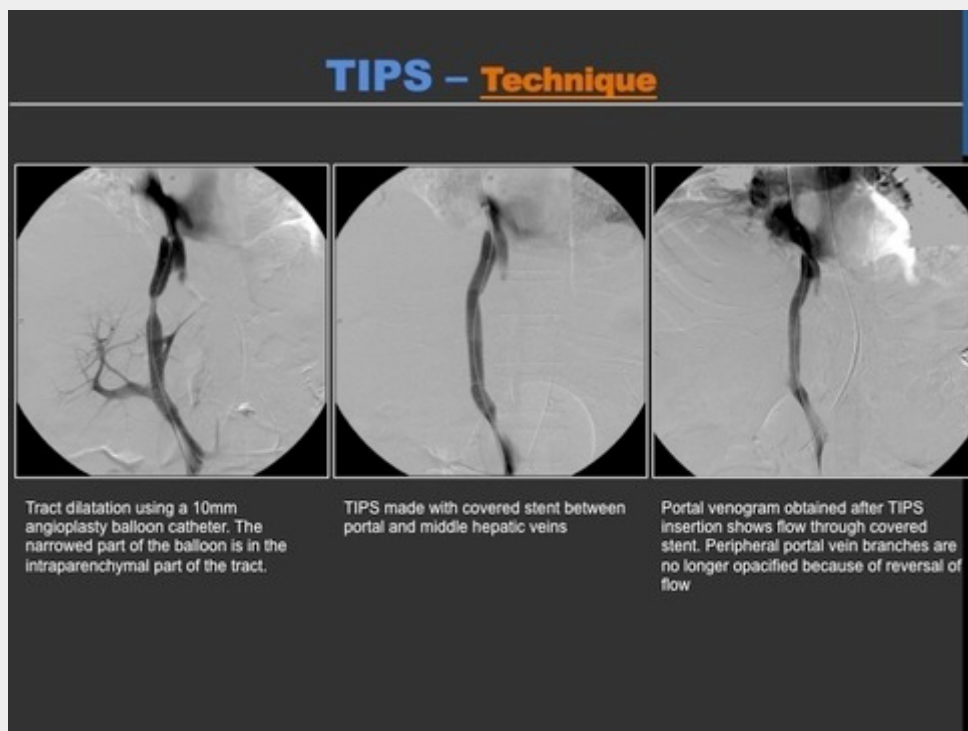


Advance of needle into the portal vein



Transhepatic portogram shows collateral circulation – left gastric vein varices

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TIPS – Complications

Complications	Frequency (%)
<u>TIPS Dysfunction</u> ^{*1}	
- Occlusion / Stenosis	18-78
- Thrombosis	10-15
Encephalopathy	
- New/worse	10-44
- Chronic	5-20
Transcapsular puncture	33
Stent migration or non-optimal placement	10-20
Hemolysis	10-15
Sepsis	2-10
Hemobilia	<5
Intraperitoneal bleeding	1-2
Hepatic infarction	1
Fistulae and endotipsitis	rare

^{*1} When TIPS were created with bare metal stents, the loss of primary patency was around 50% at one year follow-up. In this era, with use of expanded polytetrafluoro-ethylene (ePTFE)- covered stent-graft, the patency has vastly improved and stenosis occurs in just 8-20% of TIPS.

TIPS – TIPS Function / DYSFUNCTION

- Technical success = decreased of the HVPG to < 12mmHg or a reduction of at least 20%
- Clinical success = cessation of variceal bleeding, decrease of ascites, and conversion into diuretic-sensitive ascites, as well as improvement of liver function.
- The significance of TIPS dysfunction is that it can lead to recurrent portal hypertension and put patients at risk for reaccumulation of ascites or further variceal bleeding.
- Stenosis or occlusion (reduced caliber >50%) of the TIPS endograft may lead to loss of decompression of portal venous pressure. It can occur either by the result of thrombosis or intimal hyperplasia.
- An increased in HVPG > 12mmHg or a recurrence of the complication may indicate dysfunction of the TIPS.
- Thrombosis is an early complication but was more often seen (10-15% of patients) when bare metal stents were used.
- With the development of covered stents the frequency of TIPS dysfunction has been reduced and some controlled trials revealed primary patency rates of the covered stent > 80%.

Follow up Doppler Ultrasound

Baseline Examination : 1 week after the procedure

Note: In the first 48-72 hours after TIPS placement there are trapped air microbubbles between the layers of encapsulated PTFE hindering the evaluation of TIPS and making it impermeable to insonation with ultrasound.

In our practice, when early shunt thrombosis is suspected, assessment of the shunt is performed with multidetector CT or venography.

Another approach is to simply wait for symptoms of portal hypertension to recur. We discourage this approach, specially for patients with a history of variceal bleeding, because the first symptom suggesting a problem with the TIPS might be a fatal hemorrhage.

Protocol of Follow up Doppler examinations : 1 week after the procedure (or before patient discharge) and then at 1-3 months, 6 and 12 months after TIPS creation and every 6-12 months thereafter.

Whenever clinical status justifies, we made evaluations at other time intervals to check patency of the TIPS.

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Follow up Doppler Ultrasound – What evaluate ?

- Despite reports of variable success rates, ultrasound has remained an integral part of TIPS surveillance because it is non-invasive, inexpensive, and widely available.

1. Liver morphology and focal liver lesions

2. Vascular evaluation:

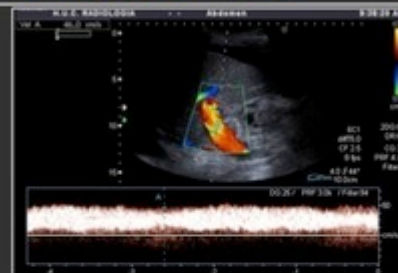
- Assessment and study of the permeability and flow direction in:
 - *Portal vein and its branches*
 - *Hepatic veins* and any hepatic vein segment between the cephalic portion of the TIPS and the IVC _The hepatic vein with the shunt, usually the right, is checked in both proximal to and just beyond its junction with the stent.
 - Inferior vena cava
 - Splenic vein

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Follow up Doppler Ultrasound

Portal Vein

- Main portal vein velocity after TIPS is typically higher than its pre-TIPS value
- MPV velocity: $> 30\text{cm/s}$
- With shunt dysfunction, the flow in the portal vein leading up to the TIPS is diminished and the MPV velocity drops often below 30cm/s .
- Flow direction in the portal system is towards the portal vein end of the stent:
- MPV and splenic vein = hepatopetal flow
- Right and left portal veins = hepatofugal flow (towards the inflow of the shunt). If the patient has patent paraumbilical vein collaterals, the flow within the left portal vein will continued to be hepatopetal.



Doppler tracing obtained at portal vein shows flow direction towards the portal vein end of the TIPS (hepatopetal) and velocity of 46cm/s.



Left portal vein with hepatofugal flow - flowing out of the diseased liver and towards the inflow of the shunt

Follow up Doppler Ultrasound

3. Evaluation of TIPS

- Position of the stent: the cephalic end of the shunt is most commonly located immediate to the connection of the right hepatic vein with the IVC, and the caudal end is located in the right portal vein.
- Lumen of TIPS : anechoic lumen without wall thickening



Two cases of baseline examination following TIPS placement. Grey-scale image of the right lobe of the liver. The TIPS is seen coursing from the porta hepatis region to the junction of the right hepatic vein with inferior vena cava. The mesh like appearance of the shunt is due to the echo reflection of the individual wire elements of the stent

Follow up Doppler Ultrasound

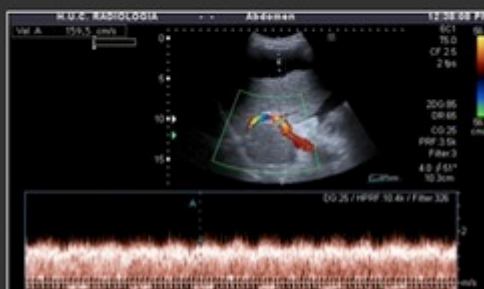
- Check permeability and flow direction : the lumen should be completely filled by the turbulent, periodic flow and the direction of flow should be from the portal vein to the hepatic vein.



Color Doppler image of the TIPS. The entire lumen is saturated with color indicating patency. Note the non-uniformity of color encoding.

Follow up Doppler Ultrasound

- Measure the flow velocity, with angle correction, in proximal, middle and distal portion of the TIPS.
- Thresholds velocities that are too high ($> 200\text{--}250\text{ cm/s}$ – jet-effect zone beyond the stenosis) or too low ($< 50\text{ cm/s}$) are associated with high specificity and sensitivity for shunt dysfunction.
- **Normal flow velocity:** $90\text{--}190\text{ cm/s}$
- Changes in the peak flow velocity $\pm 50\text{ cm/s}$ within TIPS, in relation with the baseline examination also suggests TIPS dysfunction and should also prompt shunt venography and portosystemic pressure measurements.
- Velocities can also be variable through the shunt itself in a normal situation – usually velocities increased from the portal venous end to the hepatic venous end of the shunt.
- The flow is also more turbulent when multiple stents components are used and when overriding stents cause a relative narrowing of the shunt lumen.



Spectral Doppler of the TIPS show an appropriately functioning TIPS.

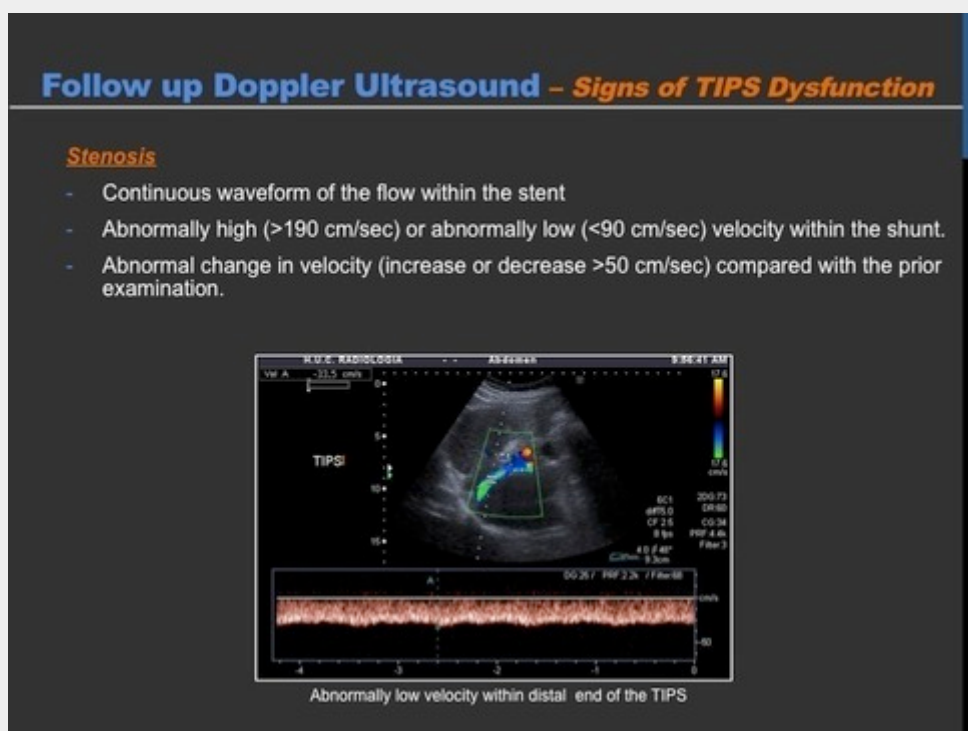
Follow up Doppler Ultrasound – Signs of TIPS Dysfunction

- Shunt malfunction is the result of narrowing or occlusion caused by intimal hyperplasia or in situ thrombosis.
- Can occur anywhere within the stent but is most commonly in the cephalic portion of the TIPS or in the hepatic vein between the stent and the IVC.

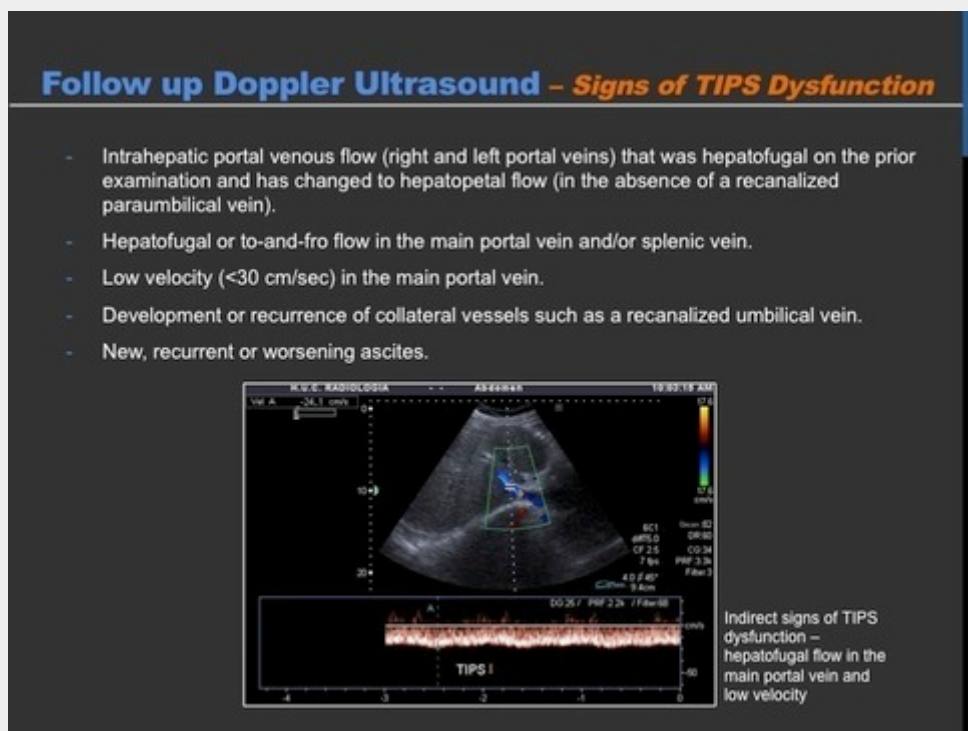
Occlusion

- Absent flow at color Doppler US (you can use *Power Doppler* and low filters to identify low flow conditions and differentiate them from true occlusion).
- Aphasic spectral waveform.
- When flow velocity and direction in the portal veins is the same as it was prior to TIPS placement.

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Doppler Ultrasound – POST-TIPS EVALUATION



Post-TIPS ultrasound Doppler evaluation:

- Detecting **complications** of TIPS procedure:
Exclude hematomas (intra-hepatic, subcapsular, peri-hepatic); biliary obstruction; hemobilia (echogenic content in the common bile duct or gallbladder); hemoperitoneu (increased echogenicity of ascites)
- Assessment of **TIPS function**. Evaluation of direct and indirect signs of **TIPS dysfunction**:
 - Absent flow
 - Shunt velocity abnormally high (>190 cm/sec) or abnormally low (<90 cm/sec)
 - Increase or decrease in shunt velocity of >50cm/s compared with the baseline examination
 - Focal region of increased velocity in the shunt or hepatic vein
 - Hepatopetal flow in left and right portal vein branches
 - Low velocity and hepatofugal flow in the main portal vein
 - Recurrence of colateralls or ascites

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Evaluating TIPS function

- Currently, venography and pressure measurements are the standards for determining shunt dysfunction, but are used only as secondary tests in our institution, because they are more invasive than ultrasound evaluation with Doppler.
- If the Doppler ultrasound findings are suggestive of a occlusion / stenosis or if there is a recurrence of patient symptoms then we performed a venography study and a TIPS revision.

4. Conclusion

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CONCLUSION

- TIPS is a technique developed to treat complications of portal hypertension.
- Doppler ultrasound evaluation is an invaluable non-invasive technique that permits an early diagnosis of shunt dysfunction.

5. References

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6. Author Information

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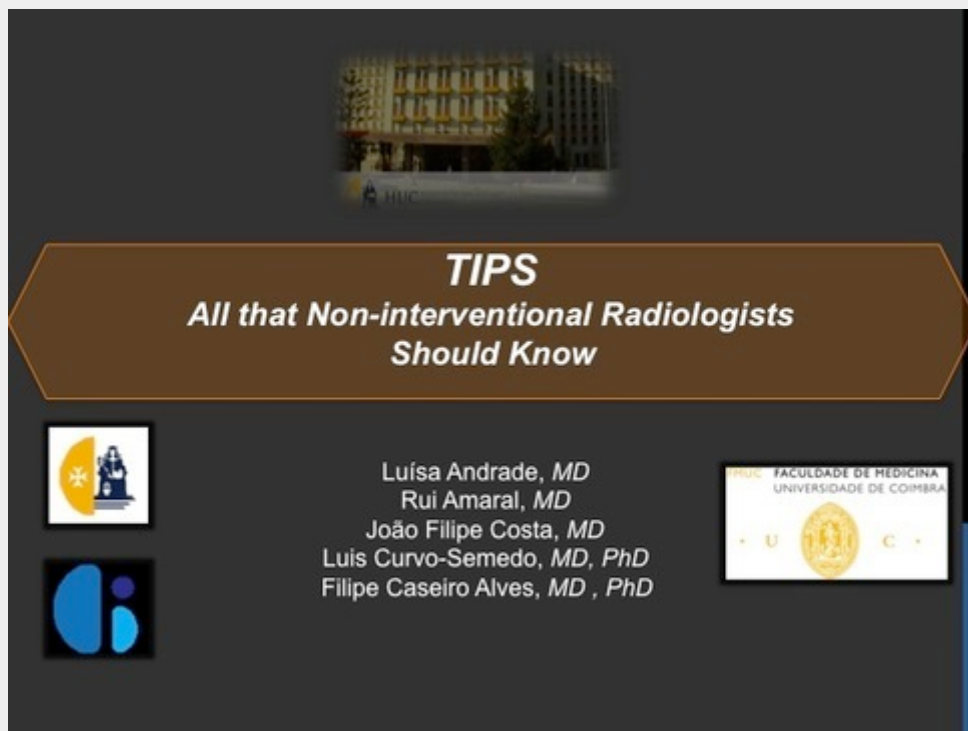
Centro Hospitalar e Universitário de Coimbra

Portugal



7. Mediafiles

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LEARNING OBJECTIVES

- To present a review on TIPS (Transjugular Intrahepatic Portosystemic Shunt): indications, contraindications, technique and complications.
- To describe and explain the pre and post-procedure imaging evaluation using mainly Doppler ultrasound and the signs for direct and indirect evidence of TIPS failure.
- To discuss the technique, focusing on tips and tricks.

BACKGROUND

- TIPS is a percutaneous imaging-guided procedure in which a channel is constructed within the liver with the intent of reducing portal pressure by diverting blood from the portal to the systemic circulation.
- The parenchymal tract created between the portal vein and hepatic veins is reinforced with placement of a stent graft.
- This procedure is used to treat severe complications of portal hypertension such as refractory ascites and variceal bleeding and can act as a bridge to liver transplantation.

LIVER DOPPLER ULTRASOUND – Normal Findings

> LIVER VASCULAR ANATOMY

1. **PORTAL VEIN** – Supplies 70% of incoming blood to the liver (relatively desoxygenated blood but rich in nutrients from the stomach, intestine, spleen and pancreas)

- Formed by the confluence of the splenic and superior mesenteric veins.
- Accompanied by the hepatic artery and common bile duct to form the portal triad.
- The main portal vein (MPV) enters the liver at the porta hepatis and divides into the right and left portal veins.
- The MPV diameter does not exceed 13mm in quiet respiration (measured at the site where the portal vein crosses the IVC). Should be normally seen an increased in this diameter, with deep inspiration.



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➤ LIVER VASCULAR ANATOMY

2. HEPATIC ARTERY – Supplies 30% of incoming blood to the liver (oxygenated blood)

- The common hepatic artery originates in the majority of patients from the celiac artery.
- After the origin of the gastroduodenal artery it is called the proper hepatic artery.
- It enters the liver alongside and anterior to the portal vein, where it divides into left and right hepatic arteries.
- There are numerous variants of hepatic artery anatomy.

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➤ LIVER VASCULAR ANATOMY

3. HEPATIC VEINS – hepatic venous drainage into inferior vena cava (IVC)

- They converge into IVC approximately 1cm below its confluence with the right atrium.
- In most people the right, middle and left hepatic veins enter the IVC in a "crows foot" configuration (transverse plane).
- 30% of individuals have additional hepatic veins.
- The walls of the hepatic veins are relatively hypoechoic, which helps to differentiate them from the portal veins in the more echogenic portal triads.



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LIVER DOPPLER ULTRASOUND – Normal Findings

- The right hepatic vein (RHV) is the largest of hepatic veins and courses between the anterior and posterior segments of the right lobe and drains into the right postero-lateral aspect of the IVC.
- The middle hepatic vein (MHV) courses along the major lobar fissure and drains the anterior right lobe segments (V, VIII) and segment IVb. In 85% of people the MHV forms a common trunk with the left hepatic vein (LHV), which then drains into the left antero-lateral aspect of the IVC.
- The LHV is the smallest of the hepatic veins and drains the lateral segments of the left lobe (II e III) and segment IVa.
- The venous drainage from the central portion of the liver (including caudate lobe) empties directly into the IVC (and cannot be perceived by color Doppler).

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LIVER DOPPLER ULTRASOUND – Normal Findings

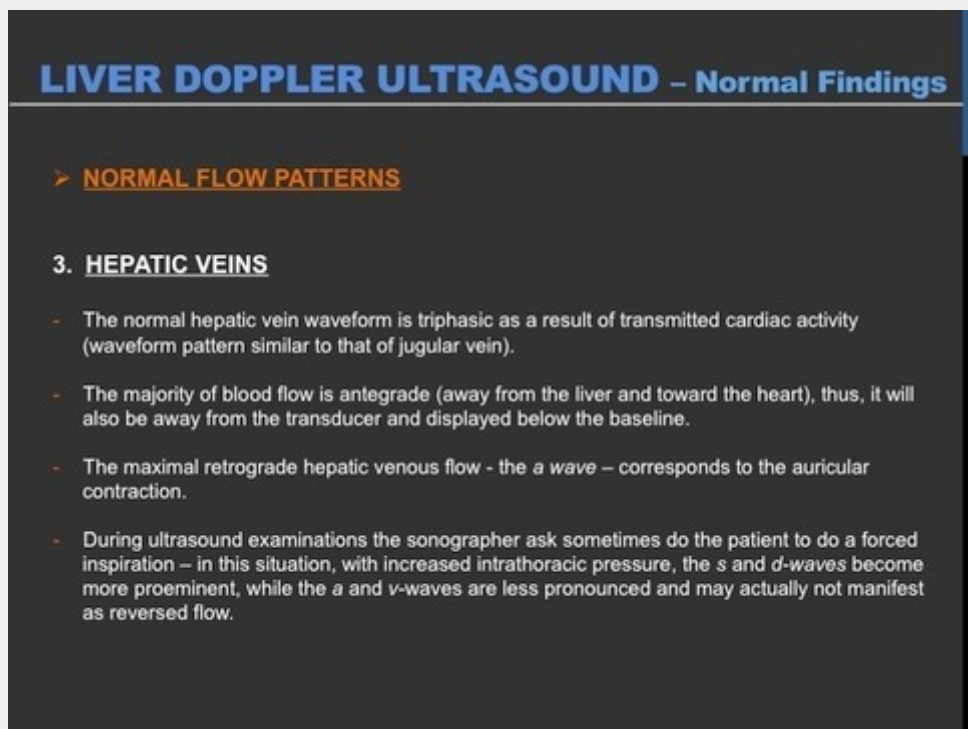
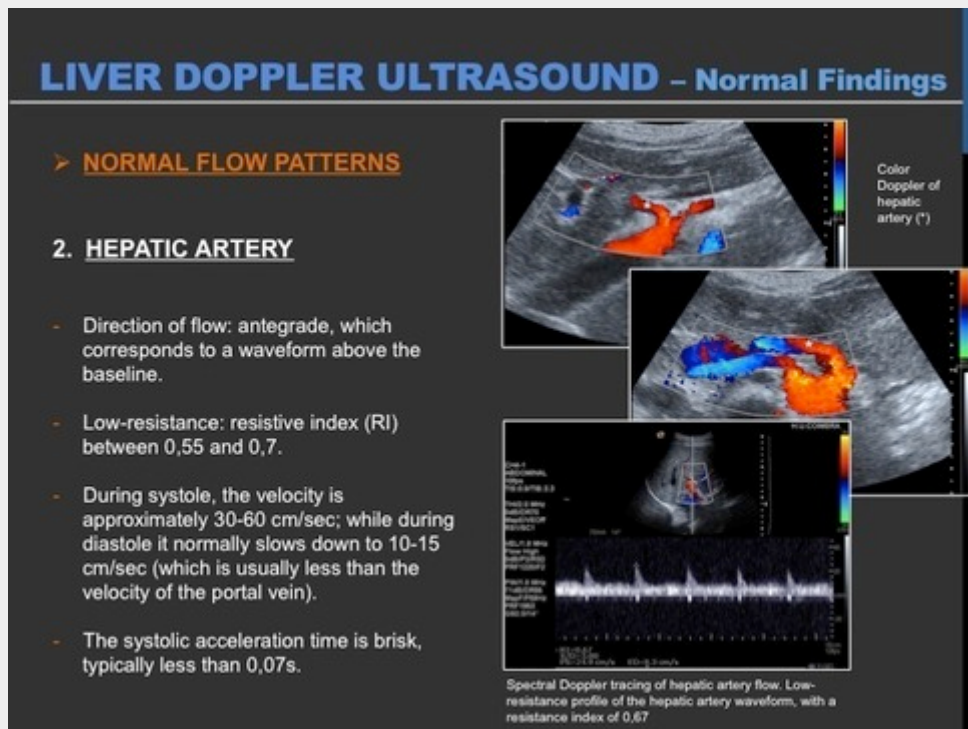
➤ NORMAL FLOW PATTERNS

1. PORTAL VEIN

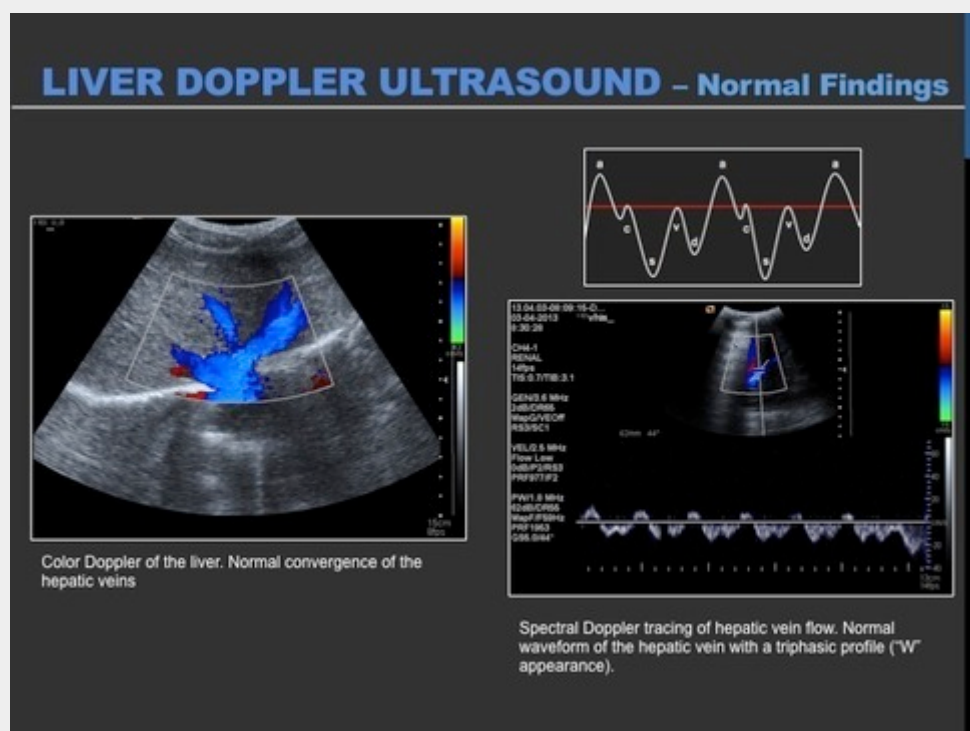
- Antegrade flow (toward the transducer), therefore with a waveform that is above the baseline.
- Hepatopetal flow (towards the liver) throughout the entire cardiac cycle.
- Portal flow velocity varies with cardiac activity and respiration, giving the portal waveform an undulating appearance.
- Mean flow velocity relatively low (but >16cm/s)



Spectral Doppler tracing of portal vein flow. Hepatopetal portal vein flow, with an undulating appearance and a mean velocity > 16 cm/sec.



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PORTAL HYPERTENSION

- Portal hypertension accompanies cirrhosis of the liver in over 90% of the cases.
- Cirrhosis disrupts the normal hepatic sinusoidal pattern, impairing venous drainage and leading to increase in portal vein pressure.
- Portal hypertension is the increased in porto-systemic pressure gradient in any portion of the portal venous system.

Porto-Systemic Pressure Gradient:

- This gradient is assessed by measuring the wedge hepatic venous pressure (a measure of sinusoidal hepatic pressure) and subtracting the free hepatic venous pressure (systemic pressure) thus obtaining the hepatic venous pressure gradient (HVPG).
- Normal: HVPG = 2-5mmHg.
- Sub-clinically Portal Hypertension: HVPG > 6mmHg
- Clinically significant Portal Hypertension: **HVPG > 10-12 mmHg** (predicts clinical course in patients with cirrhosis including development of varices, variceal hemorrhage and encephalopathy, decompensation or death after liver resection, and hepatocellular carcinoma).

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PORTAL HYPERTENSION

Pathofisiology:

Two important factors - vascular resistance and blood flow - exist in the development of portal hypertension:

$P = FR$

Where P = pressure gradient through the portal venous system; F = volume of blood flowing through the portal venous system; R = resistance to the flow.

Changes in either F or R affect the pressure. In most types of portal hypertension, both the blood flow and the resistance are altered.

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PORTAL HYPERTENSION - Etiology

INCREASED RESISTANCE TO FLOW	PREHEPATIC	Portal or splenic vein thrombosis or extrinsic compression (tumors) Splanchnic arteriovenous fistula	
	HEPATIC	Presinusoidal	Schistosomiasis Primary biliary cirrhosis Idiopathic portal hypertension Granulomatous diseases Myeloproliferative diseases Polycystic disease Hepatic metastases
		Sinusoidal	Cirrhosis of various causes (alcoholic and related to hepatitis virus B and C; Wilson disease; Hemochromatosis) Acute and fulminant hepatitis Schistosomiasis, Primary biliary cirrhosis and Idiopathic portal hypertension in advanced stages Peliosis hepatitis Budd-Chiari syndrome Sclerosing cholangitis
		Postsinusoidal	Veno-occlusive disease
INCREASED VASCULAR FLOW	POSTHEPATIC	Budd-Chiari Syndrome and thrombosis of the IVC, Cardiac disease (right sided heart failure, constrictive pericarditis and severe tricuspid regurgitation), Trauma	
		Arterioportal shunts / fistulas (congenital, traumatic, iatrogenic, tumor) Increased portal blood flow Increased splenic flow	

Doppler Ultrasound – Cirrhosis and Portal Hypertension

In patients with cirrhosis and indication to TIPS the radiologist should evaluate the following parameters:

1. Hepatic size and morphology:

- In patients with advanced cirrhosis the liver is small with relative enlargement of caudate lobe and left lobe relatively to the right lobe (in early stages of cirrhosis the liver is enlarged or could appear absolutely normal).



US demonstrates a dysmorphic liver with coarse appearance and irregular borders. This patient has cirrhosis and portal hypertension complicated with ascites.

2. Liver echogenicity and texture:

- Echogenicity could be increased in cirrhotic livers with fat infiltration.
- Irregular borders and nodular surface (in relation to the presence of regenerating nodules and fibrosis).
- Exclude the presence of definable nodules (dysplastic nodules – portal venous supply vs. HCC – arterial hepatic venous supply).
- Exclude the presence of multiple cysts (polycystic liver disease precludes the TIPS technique).



Cirrhotic liver. The hyperechoic nodule proved to be a small hepatocarcinoma.

Doppler Ultrasound – Cirrhosis and Portal Hypertension

In cirrhosis, most of the normal liver architecture is replaced by distorted vascular channels that provide increased resistance to portal venous blood flow and obstruction to hepatic venous outflow.

3. Portal vein:

- **Diameter:**
 - Although the caliber of the portal vein initially may be increased (>13mm) in PH, with the development of porto-systemic shunts, the portal vein caliber will decrease.
 - Evaluate the response to respiratory maneuvers: if the diameter of portal vein does not increase at least 20% from quiet respiration to deep inspiration the patient has probably PH.



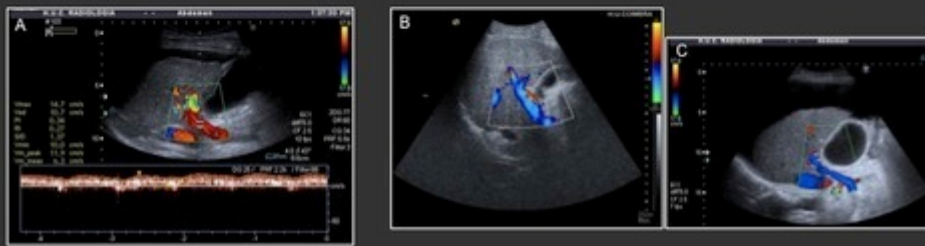
B mode US shows dilatation of the portal vein (A) and normal portal vein diameter (B) after the development of varices.

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Doppler Ultrasound – Cirrhosis and Portal Hypertension

- Permeability, flow direction and velocity:
- In severe cirrhosis portal venous flow may become continuous (without undulating appearance) or pulsatile due to arteriovenous shunting;
- Portal flow velocity decreases ($< 16\text{cm/s}$) (A);
- Flow may become to and fro (bidirectional) or hepatofugal (retrograde) (B and C). As with slow flow, this finding is diagnostic for portal hypertension from whatever cause.
- A narrow portal vein and a prominent hepatic artery are commonly associated findings when flow is hepatofugal in the main portal vein.

However, we should always kept in mind that, the flow direction in the portal vein is a variable finding in PH and depends on the development of collaterals.



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Doppler Ultrasound – Cirrhosis and Portal Hypertension

- Absent (aphasic) portal venous flow may be due to stagnant flow (portal hypertension) or occlusive disease, usually caused by bland or malignant thrombosis.
- In the setting of subacute/chronic occlusive portal vein thrombosis could be development of collateral vessels in or around the occluded portal vein (*cavernous transformation of portal vein*).



(A). Absent flow in the portal vein (RPV) that is markedly dilated due to malignant portal vein thrombosis.
(B) Spectral Doppler of the "thrombus" show an arterial waveform in a hepatofugal direction.



A normal portal vein could not be visualised. Instead there is a plexus of small vessels - anechoic serpiginous structures (arrow). After portal vein thrombosis this web of small vessels reconstitutes portal flow to the liver and is known as cavernomatous transformation of the portal vein.

slide20.jpg


Doppler Ultrasound – Cirrhosis and Portal Hypertension

4. Hepatic Artery: (A)

- In patients with PH and decreased portal flow velocity, the hepatic artery appears enlarged with increased flow to compensate for the diminished portal vein flow.
- In patients with cirrhosis the resistive index in the hepatic artery is elevated (>0.7); with development of portal hypertension and arteriovenous/arterioportal shunts the RI decreases (<0.55).

5. Hepatic Veins:

- These veins become compressed and narrowed in cirrhotic patients (B).
- With PH they lose the classic pulsatility and turn into a biphasic or monophasic waveform (C).




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Doppler Ultrasound – Cirrhosis and Portal Hypertension

6. Spleen

- Access spleen greatest dimension – in the setting of PH there could be mild to moderate splenomegaly ($>13\text{cm}$).
- Reversed flow in the splenic vein (towards the spleen). Indirect sign of splenogastric or splenorenal collaterals, and thus PH.

7. Presence / absence of ascites and volume quantification



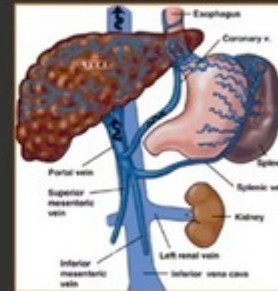
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Doppler Ultrasound – Cirrhosis and Portal Hypertension

8. Porto-systemic venous collaterals – shunting of blood from the portal to the systemic circulation.

- **Short gastric varices:** courses between the spleen and the greater curvature of the stomach.
- **Left gastric varices (coronary varices):** courses from the splenic or portal veins to the lesser curvature of the stomach. Dilatation of the coronary vein ($>7\text{mm}$) is associated with severe PH (HVPG $>10\text{mmHg}$).

These veins communicate with systemic esophageal veins and are known as gastroesophageal varices.



- **Paraumbilical veins:** originates from the left portal vein and runs inferiorly through the falciform ligament towards the anterior abdominal wall where it connects with the systemic epigastric veins or with the subcutaneous veins originating the *caput medusae*.
- **Splenorenal-mesenteric collaterals:** courses between the splenic hilum and the left renal vein.

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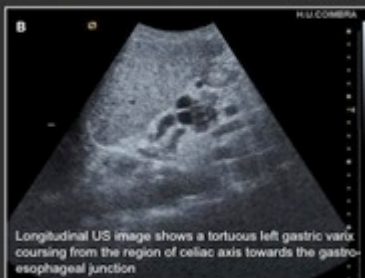
Doppler Ultrasound – Cirrhosis and Portal Hypertension



US scan of the spleen. Tangle of vessels are seen at the upper pole of the spleen, corresponding to short gastric varices.



Color Doppler image shows a recanalized paraumbilical vein that carries blood towards the umbilicus.



Longitudinal US image shows a tortuous left gastric varix coursing from the region of celiac axis towards the gastro-esophageal junction




(D) US scan of the spleen revealed anechoic tubular structures adjacent to the lower pole of the spleen, communicating to the left renal vein (not shown).

(E) A large varix occupies the space between the splenic hilum (s) and the left kidney (k)



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Doppler Ultrasound – Cirrhosis and Portal Hypertension



Pre-TIPS ultrasound Doppler evaluation:

Detecting *stigmas* of cirrhosis and portal hypertension in the liver and hepatic vasculature

Exclude hepatic cysts

Detecting complications of portal hypertension:

- Neoplastic processes – HCC
- Portal vein thrombosis ± cavernous transformation
- Splenomegaly
- Ascites
- Collateral pathways

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TRANSJUGULAR INTRAHEPATIC PORTOSYSTEMIC SHUNT - TIPS

DEFINITION

- Percutaneous imaging-guided method in which a channel is constructed within the liver with the intent of reducing portal pressure by diverting blood from the portal to the systemic circulation.
- Intrinsically functions as a side-to-side porta-caval shunt, designed to function as a partial shunt that preserves a portion of portal flow to the liver.

OBJECTIVE

- Reduce portal hypertension in patients with complications related to portal hypertension, such as refractory ascites and refractory variceal bleeding.


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TIPS - Indications

- **Refractory ascites** (the most frequent indication for TIPS)
- **Secondary prevention of variceal bleeding** (second-line therapy after failure of medical and endoscopic therapy)
- Refractory acute variceal bleeding
- Portal hypertensive gastropathy
- Hepatorrenal syndromes
- Bleeding gastric / ectopic varices
- Refractory hepatic hydrothorax
- Budd-Chiari syndrome
- Hepatic veno-occlusive disease
- Hepatopulmonary syndrome
- In centers with extensive experience TIPS should be considered in patients with portal vein thrombosis with or without cavernomatous transformation, when intrahepatic portal venous branches are patent.

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TIPS – Patient Selection



The illustration shows a line of stylized human figures. The first figure is green and has a large red checkmark next to it, indicating selection. The subsequent figures are grey. Below the figures are several rectangular boxes, some of which are empty, suggesting a process of evaluation or selection.

- Patients with a Child-Pugh score of >12 are in high risk of post-procedural death.
- MELD scores of $>15-18$ or a bilirubin level of $>3.5\text{mg/dl}$ have poor prognosis and TIPS should only be performed in the absence of any other treatment possibilities.

TIPS – Contra-Indications

Absolute

- Unproved portal hypertension
- Primary prevention of variceal bleeding
- Congestive heart failure
- Polycystic liver disease
- Uncontrolled systemic infection or sepsis
- Unrelieved biliary obstruction
- Severe pulmonary hypertension
- Severe tricuspid regurgitation

Relative:

- Hepatocarcinoma large and central
- Hepatic venous obstruction
- Portal vein thrombosis
- Severe coagulopathy (INR>5)
- Thrombocytopenia (<20 000)
- Moderate pulmonary hypertension
- Hepatic encephalopathy poorly controlled

TIPS – Technique

1. Jugular access (right internal jugular vein) – Seldinger technique

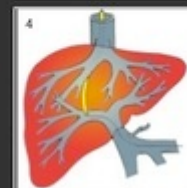
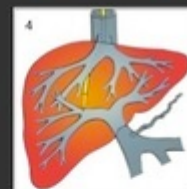
2. A catheter is placed and guided through the superior vena cava, right atrium and inferior vena cava into a hepatic vein. (Because of large size and consistent and predictable relation to the portal veins, the right hepatic vein is the preferred hepatic vein branch for the creation of a TIPS shunt).

3. Selective catheterization of a suitable branch of the RHV

(Pressure measurements are performed at this point, including central venous pressures, at the level of right atrium and IVC and free and wedged hepatic venous pressures).

4. A needle inserted through the catheter is then used to puncture the liver from a central portion of the hepatic vein and enter the main portal branch (usually the right portal vein).

The puncture could be navigated with ultrasound.




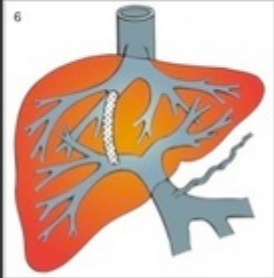

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TIPS – Technique

5. Dilatation of the needle tract by a balloon catheter – establishing of a connection between the portal and systemic circulation directly inside the liver parenchyma.


6. Insertion of a TIPS stent graft

7. The final step of the TIPS procedure is placement of a pigtail catheter over the portal vein guide wire for follow up portography and pressure measurements.




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TIPS – Technique



Suprahepatic flebography

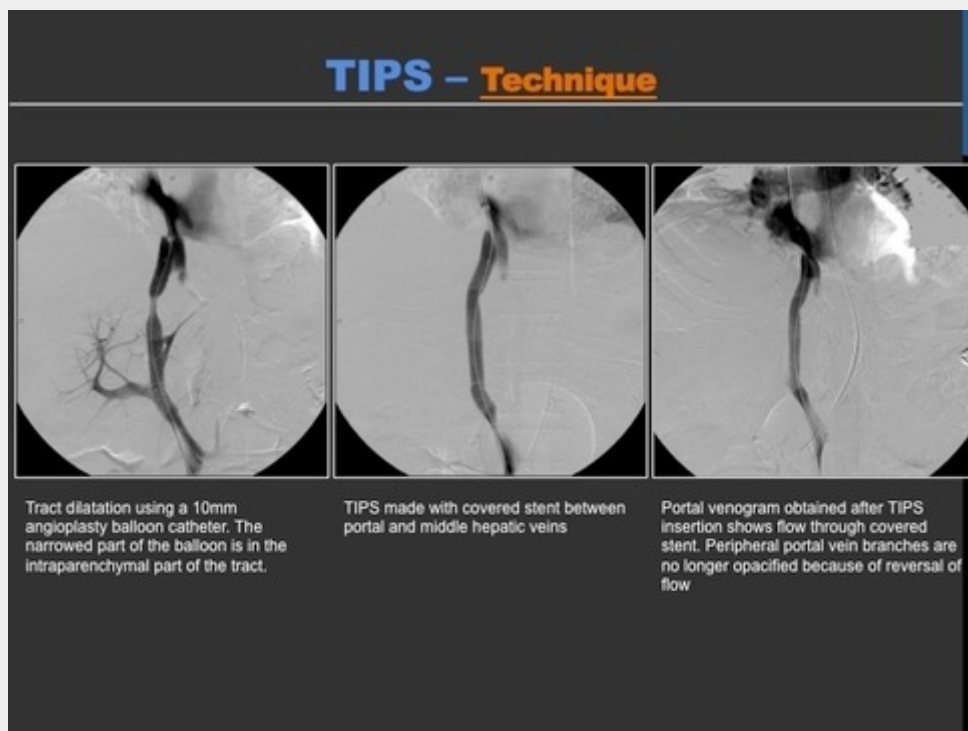


Advance of needle into the portal vein



Transhepatic portogram shows collateral circulation – left gastric vein varices

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slide33.jpg

TIPS – Complications

Complications	Frequency (%)
TIPS Dysfunction ^{*1}	
- Occlusion / Stenosis	18-78
- Thrombosis	10-15
Encephalopathy	
- New/worse	10-44
- Chronic	5-20
Transcapsular puncture	33
Stent migration or non-optimal placement	10-20
Hemolysis	10-15
Sepsis	2-10
Hemobilia	<5
Intraperitoneal bleeding	1-2
Hepatic infarction	1
Fistulae and endotipsitis	rare

^{*1} When TIPS were created with bare metal stents, the loss of primary patency was around 50% at one year follow-up. In this era, with use of expanded polytetrafluoro-ethylene (ePTFE)- covered stent-graft, the patency has vastly improved and stenosis occurs in just 8-20% of TIPS.

TIPS – TIPS Function / DYSFUNCTION

- Technical success = decreased of the HVPG to < 12mmHg or a reduction of at least 20%
- Clinical success = cessation of variceal bleeding, decrease of ascites, and conversion into diuretic-sensitive ascites, as well as improvement of liver function.
- The significance of TIPS dysfunction is that it can lead to recurrent portal hypertension and put patients at risk for reaccumulation of ascites or further variceal bleeding.
- Stenosis or occlusion (reduced caliber >50%) of the TIPS endograft may lead to loss of decompression of portal venous pressure. It can occur either by the result of thrombosis or intimal hyperplasia.
- An increased in HVPG > 12mmHg or a recurrence of the complication may indicate dysfunction of the TIPS.
- Thrombosis is an early complication but was more often seen (10-15% of patients) when bare metal stents were used.
- With the development of covered stents the frequency of TIPS dysfunction has been reduced and some controlled trials revealed primary patency rates of the covered stent > 80%.

Follow up Doppler Ultrasound

Baseline Examination : 1 week after the procedure

Note: In the first 48-72 hours after TIPS placement there are trapped air microbubbles between the layers of encapsulated PTFE hindering the evaluation of TIPS and making it impermeable to insonation with ultrasound.

In our practice, when early shunt thrombosis is suspected, assessment of the shunt is performed with multidetector CT or venography.

Another approach is to simply wait for symptoms of portal hypertension to recur. We discourage this approach, specially for patients with a history of variceal bleeding, because the first symptom suggesting a problem with the TIPS might be a fatal hemorrhage.

Protocol of Follow up Doppler examinations : 1 week after the procedure (or before patient discharge) and then at 1-3 months, 6 and 12 months after TIPS creation and every 6-12 months thereafter.

Whenever clinical status justifies, we made evaluations at other time intervals to check patency of the TIPS.

Follow up Doppler Ultrasound – What evaluate ?

- Despite reports of variable success rates, ultrasound has remained an integral part of TIPS surveillance because it is non-invasive, inexpensive, and widely available.

1. Liver morphology and focal liver lesions

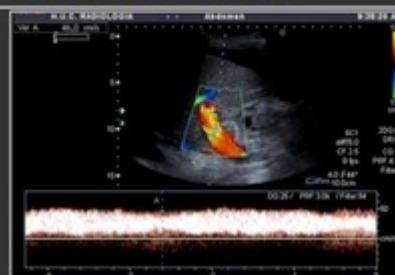
2. Vascular evaluation:

- Assessment and study of the permeability and flow direction in:
 - *Portal vein and its branches*
 - *Hepatic veins* and any hepatic vein segment between the cephalic portion of the TIPS and the IVC _The hepatic vein with the shunt, usually the right, is checked in both proximal to and just beyond its junction with the stent.
 - Inferior vena cava
 - Splenic vein

Follow up Doppler Ultrasound

Portal Vein

- Main portal vein velocity after TIPS is typically higher than its pre-TIPS value
- MPV velocity: > 30cm/s
- With shunt dysfunction, the flow in the portal vein leading up to the TIPS is diminished and the MPV velocity drops often below 30cm/s.
- Flow direction in the portal system is towards the portal vein end of the stent:
- MPV and splenic vein = hepatopetal flow
- Right and left portal veins = hepatofugal flow (towards the inflow of the shunt). If the patient has patent paraumbilical vein collaterals, the flow within the left portal vein will continued to be hepatopetal.



Doppler tracing obtained at portal vein shows flow direction towards the portal vein end of the TIPS (hepatopetal) and velocity of 46cm/s.



Left portal vein with hepatofugal flow - flowing out of the diseased liver and towards the inflow of the shunt

Follow up Doppler Ultrasound

3. Evaluation of TIPS

- Position of the stent: the cephalic end of the shunt is most commonly located immediate to the connection of the right hepatic vein with the IVC, and the caudal end is located in the right portal vein.
- Lumen of TIPS : anechoic lumen without wall thickening



Two cases of baseline examination following TIPS placement. Grey-scale image of the right lobe of the liver. The TIPS is seen coursing from the porta hepatis region to the junction of the right hepatic vein with inferior vena cava. The mesh like appearance of the shunt is due to the echo reflection of the individual wire elements of the stent

Follow up Doppler Ultrasound

- Check permeability and flow direction : the lumen should be completely filled by the turbulent, periodic flow and the direction of flow should be from the portal vein to the hepatic vein.



Color Doppler image of the TIPS. The entire lumen is saturated with color indicating patency. Note the non-uniformity of color encoding.

Follow up Doppler Ultrasound

- Measure the flow velocity, with angle correction, in proximal, middle and distal portion of the TIPS.
- Thresholds velocities that are too high ($> 200\text{--}250\text{ cm/s}$ – jet-effect zone beyond the stenosis) or too low ($< 50\text{ cm/s}$) are associated with high specificity and sensitivity for shunt dysfunction.
- **Normal flow velocity:** $90\text{--}190\text{ cm/s}$
- Changes in the peak flow velocity $\pm 50\text{ cm/s}$ within TIPS, in relation with the baseline examination also suggests TIPS dysfunction and should also prompt shunt venography and portosystemic pressure measurements.
- Velocities can also be variable through the shunt itself in a normal situation – usually velocities increased from the portal venous end to the hepatic venous end of the shunt.
- The flow is also more turbulent when multiple stents components are used and when overriding stents cause a relative narrowing of the shunt lumen.



Spectral Doppler of the TIPS show an appropriately functioning TIPS.

Follow up Doppler Ultrasound – Signs of TIPS Dysfunction

- Shunt malfunction is the result of narrowing or occlusion caused by intimal hyperplasia or in situ thrombosis.
- Can occur anywhere within the stent but is most commonly in the cephalic portion of the TIPS or in the hepatic vein between the stent and the IVC.

Occlusion

- Absent flow at color Doppler US (you can use *Power Doppler* and low filters to identify low flow conditions and differentiate them from true occlusion).
- Aphasic spectral waveform.
- When flow velocity and direction in the portal veins is the same as it was prior to TIPS placement.

slide42.jpg

Follow up Doppler Ultrasound – Signs of TIPS Dysfunction

Stenosis

- Continuous waveform of the flow within the stent
- Abnormally high (>190 cm/sec) or abnormally low (<90 cm/sec) velocity within the shunt.
- Abnormal change in velocity (increase or decrease >50 cm/sec) compared with the prior examination.

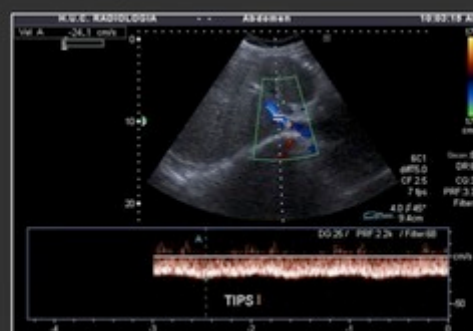


Abnormally low velocity within distal end of the TIPS

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Follow up Doppler Ultrasound – Signs of TIPS Dysfunction


- Intrahepatic portal venous flow (right and left portal veins) that was hepatofugal on the prior examination and has changed to hepatopetal flow (in the absence of a recanalized paraumbilical vein).
- Hepatofugal or to-and-fro flow in the main portal vein and/or splenic vein.
- Low velocity (<30 cm/sec) in the main portal vein.
- Development or recurrence of collateral vessels such as a recanalized umbilical vein.
- New, recurrent or worsening ascites.



Indirect signs of TIPS dysfunction – hepatofugal flow in the main portal vein and low velocity

slide44.jpg

Doppler Ultrasound – POST-TIPS EVALUATION



Post-TIPS ultrasound Doppler evaluation:

- Detecting **complications** of TIPS procedure:
Exclude hematomas (intra-hepatic, subcapsular, peri-hepatic); biliary obstruction; hemobilia (echogenic content in the common bile duct or gallbladder); hemoperitoneu (increased echogenicity of ascites)
- Assessment of **TIPS function**. Evaluation of direct and indirect signs of **TIPS dysfunction**:
 - Absent flow
 - Shunt velocity abnormally high (>190 cm/sec) or abnormally low (<90 cm/sec)
 - Increase or decrease in shunt velocity of >50cm/s compared with the baseline examination
 - Focal region of increased velocity in the shunt or hepatic vein
 - Hepatopetal flow in left and right portal vein branches
 - Low velocity and hepatofugal flow in the main portal vein
 - Recurrence of colateralls or ascites

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Evaluating TIPS function

- Currently, venography and pressure measurements are the standards for determining shunt dysfunction, but are used only as secondary tests in our institution, because they are more invasive than ultrasound evaluation with Doppler.
- If the Doppler ultrasound findings are suggestive of a occlusion / stenosis or if there is a recurrence of patient symptoms then we performed a venography study and a TIPS revision.

CONCLUSION

- TIPS is a technique developed to treat complications of portal hypertension.
- Doppler ultrasound evaluation is an invaluable non-invasive technique that permits an early diagnosis of shunt dysfunction.

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