Liver Transplantation as Ultimate Tool to Treat Portal Hypertension

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Introduction

A variety of liver diseases may cause portal hypertension, an entity that causes complications per se, such as variceal bleeding, ascites, and hepatic encephalopathy. Complications of portal hypertension can be managed by several therapeutic options like drug therapy, interventional endoscopy, transjugular intrahepatic portosystemic shunt (TIPS), or surgical shunt- and non-shunt procedures. In the acute state, the patient has to be treated according to his or her condition, e.g., gastrointestinal bleeding or spontaneous bacterial peritonitis. In advanced disease, liver transplantation is very often the only definitive causal therapy, which cures portal hypertension and most of its complications as well as the underlying liver disease.

Portal hypertension is defined as an increase of 5–10 mm Hg above normal portal vein pressure and may be classified as presinusoidal, sinusoidal, and postsinusoidal (table 1) [1]. It is initiated by the increase in resistance of portal blood flow, and is then sustained by the increase in portal blood flow and hyperdynamic systemic circulation. This cascade is triggered by several neurohumoral and paracrine substances such as nitric oxide (NO), prostacyclin (PGI₂), overactivity of the sympathetic nervous system, β₂-adrenergic receptor agonists, certain neuropeptides, glucagon, and products of tissue hypoxia [2]. These mediators cause peripheral vasodilation and thereby increase the splanchnic blood flow and constitute the ‘forward’ component of portal hypertension [3].

Key Words
Portal hypertension · Liver transplantation · Portal thrombosis · Hepatopulmonary syndrome · Pulmonary hypertension

Abstract

Portal hypertension is a complication of liver cirrhosis that may itself cause complications such as variceal bleeding, ascites and hepatorenal syndrome. There are several options for symptomatic treatment including drug therapy, endoscopy, transjugular intrahepatic portosystemic shunt (TIPS), and various surgical procedures, notably liver transplantation, the only causal treatment. The indication for liver transplantation has to be defined carefully. Progression of the primary disease, evaluation of comorbidity and overall prognosis have to be considered. Conservative symptomatic treatment is used for bridging purposes until liver transplantation can be provided to cure portal hypertension and the underlying primary disease. Careful timing of the transplantation is necessary as well as reorganization of the waiting lists by introducing new priority systems as the Model for End-Stage Liver Disease (MELD) in order to reduce mortality. Furthermore, living donor liver transplantation and split liver transplantation are methods to enlarge the donor pool, and thus accessibility of transplantation to a greater number of patients. This review evaluates the indication of liver transplantation in the treatment of portal hypertension.
Because of this increased blood flow and the increased portal pressure, collaterals are formed such as the short gastro-splenic veins and esophageal varices [1].

**Complications of Portal Hypertension**

Complications of portal hypertension develop independently of the underlying liver disease, and consist mainly of gastrointestinal bleeding, i.e. bleeding of esophageal varices, ascites, hepatic encephalopathy, hepatorenal syndrome, portopulmonary hypertension, and hepatopulmonary syndrome.

In portal hypertension, collaterals are formed to decompensate the portal system. Collateralization is accompanied by an intensified splanchnic vasodilation due to a hyperdynamic circulation, resulting in an increase in portal venous inflow [4]. Esophageal varices, representing the clinically most relevant portosystemic collaterals, affect about 50% of patients with newly diagnosed liver cirrhosis [5]. In the case of portal hypertension >12 mm Hg, the incidence of variceal bleeding is increasing and affects 30% of patients with compensated liver disease and up to 60% of patients with decompensated liver disease [6].

Ascites formation is multifactorial: increased lymphatic flow, a neurogenic hepatorenal reflex, and a decrease in effective arterial blood volume because of splanchnic arterial vasodilation [7] leads to an increase in renal sodium and water retention by activation of the sympathetic nervous system and renin-angiotensin-aldosterone system [8]. Thereby, total body water increases and, in severe cases, dilutional hyponatremia occurs [9]. Water excretion can be further impaired by several factors including diuretics, nonsteroidal anti-inflammatory drugs (NSAIDs), large volume paracentesis performed without plasma expansion and the use of vasopressin analogues during treatment of variceal bleeding [7]. In addition, the filtration rate to the ascending limb of the loop of Henle is reduced, ADH secretion is increased and the renal synthesis of prostaglandins is reduced [9].

Splanchnic vasodilation and loss of effective arterial blood volume lead to renal hypoperfusion and subsequently to progressive activation of vasoconstrictor systems as the renin-angiotensin-aldosterone system and the sympathetic nervous system [7]. Vasoconstriction of the renal cortex leads to reduced renal blood flow and glomerular filtration rate, causing hepatorenal syndrome [10]. There are two types of hepatorenal syndrome: type 1 is characterized by a rapid progression of renal failure with a twofold increase in initial serum creatinine or a 50% reduction of initial 24-hour creatinine clearance to below 20 ml/min in less than 2 weeks, whereas type 2 develops more moderately without fulfilling the criteria of type 1 [7]. Portopulmonary hypertension is defined as raised pulmonary arterial pressure of greater than 25 mm Hg at rest [11]. High cardiac output and hyperdynamic circulation in patients with portal hypertension increase shear stress on the pulmonary circulation. By this, vasoconstriction leads to increased vascular resistance and to progressive pulmonary vascular remodeling due to proliferation of pulmonary arterial endothelial cells and smooth muscle cells, which result in pulmonary hypertension [12]. Patients mainly suffer from progressive dyspnea, less frequently from fatigue, palpitations, syncope, or chest pain [12].

In hepatopulmonary syndrome, hyperdynamic circulation and portal hypertension lead to altered bowel perfusion and an increased rate of enteral translocation of gram-negative bacteria and endotoxin, which stimulate the release of vasoactive mediators, including TNF-α, hem-oxygenase-derived carbon monoxide, and NO [12]. This causes abnormal intrapulmonary dilation and shunting. Patients eventually suffer from progressive hypoxemia with increasing cyanosis. Some patients may develop clubbing, and cutaneous telangiectasia (spider angiomas) are frequently present [12].

**Options for Treatment prior to Liver Transplantation**

Depending on the severity of liver disease causing portal hypertension and on the main complications caused by portal hypertension, there are several options to treat portal hypertension and its complications prior to liver transplantation.

Drug therapy mainly consists of nonselective β-blockers in order to reduce portal venous pressure [13]. If no reduction of hepatic venous pressure gradient of >20% or <12 mm Hg is achieved, β-blockers can be combined with isosorbide mononitrate [14]. In the case of acute bleeding, terlipressin, a vasopressin analogue, can be used as an effective agent to control variceal bleeding [15]. Vasoactive drugs should be given immediately and maintained for 5 days, and should be combined with initial endoscopic treatment [16]. Recommendations for ascites treatment include both spironolactone and furosemide up to a daily dose of 400 and 160 mg, respectively [10]. In long-term ascites therapy, doses of diuretics have to be limited to
avoid prerenal renal failure. In addition, periodic paracentesis alongside the administration of human albumin (50 ml of 25% albumin solution for every 1.5 liters removed) is advocated [17]. In the case of acute deterioration of kidney function due to acute exacerbation of liver disease or failure of diuretic therapy, dopamine infusions in combination with plasma expanders can be used. Continuous intravenous administration of epoprostenol shows good results for primary therapy of portopulmonary hypertension. In an experimental setting, the endothelin-receptor antagonist bosentan demonstrated beneficial effects on hemodynamics and exercise capacity in patients with portopulmonary hypertension. However, due to its hepatotoxic potential, it has not yet been approved for clinical use in patients with liver diseases [12].

If variceal bleeding occurs, interventional endoscopy is the treatment of choice. This includes sclerotherapy, application of tissue adhesives, banding of varices, and some other methods. Variceal banding is more difficult in acute bleeding due to less visibility, but can control bleeding as effectively as sclerotherapy with less complications and thus seems to improve survival compared to sclerotherapy [6].

Since the end of the 1980s, TIPS has become a standard option to reduce portal pressure when conservative treatment has failed [18]. Morbidity and mortality have decreased in contrast to surgery and the reduction in portal hypertension and thereby the incidence of side effects, i.e. hepatic encephalopathy and liver failure, can be adjusted more precisely by varying stent diameter, ranging from 7 to 12 mm [19]. Furthermore, in the case of inappropriate shunt flow, TIPS function can be optimized by a repetitive procedure. TIPS can be used either as an emergency therapy in case of uncontrollable variceal bleeding, or as an elective procedure in patients with complications due to portal hypertension such as refractory ascites and hepatorenal syndrome [19]. Also, there are several reports of transjugular portosystemic shunting that led to the correction of hypoxemia in case of hepatopulmonary syndrome [20]. The direct impact of TIPS on liver transplantation, perioperative outcome, and rate of complications has not been fully evaluated yet. There are some reports of technical difficulty due to incorrect positioning of the stent and damage to the portal vein intima, dislocation of the stent into the portal vein [21], and injury to the bile duct [22]. However, a TIPS placed correctly does not influence the liver transplantation procedure and may serve to bridge the interval to liver transplantation as well as a solution for patients not awaiting liver transplantation [23].

If the insertion of a TIPS is technically not possible, a surgical shunt would be the next line of treatment. Total portosystemic shunts include any shunt with a diameter greater than 10 mm, and control variceal bleeding and ascites in up to 90% of patients [24, 25]. Because of the large shunt diameter, portal flow is diverted through the shunt leading to decompression of sinusesoids with a relatively high risk of liver failure and encephalopathy, however [24, 25]. Partial portosystemic shunts reduce the size of the anastomosis of a side-to-side shunt to 8 mm. Up to this size, portal pressure is reduced to 12 mm Hg and portal flow is maintained in 80% of patients [26]. Selective shunts provide decompression of gastroesophageal varices to control bleeding, while, at the same time, portal hypertension and thus portal flow to the cirrhotic liver are maintained. The distal splenorenal shunt [27], which is most commonly used, combines the anastomosis of the splenic vein to the left renal vein and ligature of the left gastric vein [28]. At present, surgical shunts may be of some value in patients with recurrent variceal bleeding in early or absent liver disease, such as portal vein thrombosis [29]. A previous portosystemic shunt operation is no contraindication to subsequent liver transplantation, although peritransplant morbidity is increased due to the increased complexity of the procedure, portal vein abnormalities, and adhesions [30]. When a surgical shunt is indicated, a distal splenorenal shunt seems to be superior to other shunts to control variceal bleeding and does not significantly compromise future liver transplantation. However, the incidence of postoperative portal vein thrombosis of up to 6% should be taken into consideration [31].

When shunt procedures, either interventional or surgical, are technically impossible, non-shunt operations may be indicated. They comprise devascularization operations including gastric and esophageal devascularization, mostly splenectomy, to reduce a major inflow path of gastroesophageal varices, and in some instances esophageal transection [32]. The incidence of liver failure and hepatic encephalopathy is low following devascularization procedures due to a better maintenance of portal flow, but portal vein thrombosis occurs in up to 20% of patients after splenectomy [28].

Liver Transplantation

All therapeutic options listed above, with some restrictions regarding surgical non-shunt procedures, may be used prior to liver transplantation. However, in most
Portal hypertension is the only causal treatment for liver transplantation and also cures the underlying liver disease. While treating the patient conservatively, the optimal time point for placing the patient on the waiting list should not be missed, in particular with regard to increasing waiting time and higher perioperative morbidity and mortality after liver transplantation in advanced liver failure and hepatorenal syndrome [8, 33]. Furthermore, some complications of portal hypertension, such as severe, fixed pulmonary hypertension, are contraindications to liver transplantation [34].

Without the option of liver transplantation, 5-year survival of patients with Child C cirrhosis and variceal bleeding is approximately 25% whereas after liver transplantation it is 70–75% [35]. Even in patients with a Child-Turcotte-Pugh (CTP) score below 10, 5-year survival after liver transplantation is better than without (85 vs. 73%, respectively) (table 1) [31]. However, due to the shortage of donor organs, not every patient with portal hypertension can receive a liver graft in time, and mortality on the waiting list is increasing [28].

Indication and time point for liver transplantation have to be evaluated carefully, considering not only etiology, severity, and activity of underlying liver disease, but also comorbidity and possible contraindications [36]. A widely used clinical scoring system is the Child score or one of its modifications, e.g. CTP classification (table 2).

Table 1. Classification of portal hypertension

<table>
<thead>
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<th>Presinusoidal</th>
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<tr>
<td>Extrahepatic</td>
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<tr>
<td>Portal vein thrombosis</td>
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<td>Intrahepatic</td>
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<td>Schistosomiasis</td>
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<td>Idiopathic portal hypertension</td>
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<td>Congenital hepatic fibrosis</td>
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<td>Sarcoidosis</td>
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<td>Felty’s syndrome</td>
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<td>Primary biliary cirrhosis</td>
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| Sinusoidal |  |
| Alcoholic cirrhosis |  |
| Nonalcoholic cirrhosis |  |
| Nodular regenerative hyperplasia |  |

| Postsinusoidal |  |
| Extrahepatic |  |
| Budd Chiari syndrome |  |
| Intrahepatic |  |
| Veno-occlusive disease |  |
| Central hyaline sclerosis (alcohol hepatitis) |  |

Since the prognostic value of the CTP score has never been proven [37] and because this score contains subjective values, such as degree of encephalopathy and ascites, the Model for End-Stage Liver Disease (MELD) has been developed [38]. The United Network for Organ Sharing (UNOS) has been using this score to organize waiting list priority since February 27, 2002. Originally, the MELD score was developed to determine the short-term prognosis of patients undergoing a TIPS procedure based on four variables: serum creatinine, serum bilirubin, international normalized ratio (INR), and the etiology of liver disease. It was demonstrated that implication of the etiology of liver disease into the calculation of the score contributes very little to the ability of MELD score to predict 3-month survival. Therefore, the Organ Procurement and Transplantation Network (OPTN) calculates the MELD score using the following equation:

\[
\text{MELD} = 0.957 \times \ln(\text{creatinine}) + 0.378 \times \ln(\text{bilirubin}) + 1.12 \times \ln(\text{INR}) + 0.643.
\]

The score is rounded and truncated for values of 40 and higher. This new system does not categorize patients into groups but utilizes a continuous score. The predictive value for 3-month survival is excellent, and superior to that of the CTP score. Patients with an MELD score of <9 have a 3-month mortality rate of 1.9 versus 71.3% for patients with an MELD score >40 [39].

Indications to liver transplantation for some complications of portal hypertension as hepatorenal syndrome and pulmonary hypertension have to be discussed separately. Pulmonary hypertension might be reversible after liver transplantation, but if severe, fixed pulmonary hypertension (MPAP >35 mm Hg) is present, mortality is unacceptably high and thus liver transplantation is con-
venous flow exceeds 250 ml/min/100 g liver tissue significantly lower survival prognosis. Sufficient venous outflow of the sinusoidal vascular bed and hyperdynamic circulation results in good long-term outcome. Additionally, this procedure can be used as rescue therapy if portal flow cannot be achieved [54].

LDLT in patients with PVT remains controversial. However, when performed, combination of thrombendvenectomy and a venous jump-graft is required, e.g. recipient saphenous vein or internal iliac vein. Alternatively, cryoconserved grafts can be used. Other techniques already applied in patients with extensive portomesenteric thrombosis include cavo-portal hemitranposition, characterized by an anastomosis of the donor portal vein to the inferior vena cava, which is than ligated cranial to the anastomosis [55].
Conclusions

Liver transplantation is the only curative treatment for patients with portal hypertension in end-stage liver disease. Patients with good liver function despite portal hypertension may be managed satisfactorily without liver transplantation. However, patients with end-stage liver disease should be considered timely for liver transplantation. Patients on the waiting list need symptomatic ‘bridging therapy’, such as drug therapy, endoscopic therapy, and a TIPS procedure, until a liver transplant is available. Improved waiting list management and alternatives to deceased donor transplantation such as split liver transplantation and LDLT can expand the donor pool and help to decrease the waiting time and thereby the mortality of patients on the waiting list.

References

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