BRIEF ARTICLES

Percutaneous portal venoplasty and stenting for anastomotic stenosis after liver transplantation

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Abstract

AIM: To review percutaneous transhepatic portal venoplasty and stenting (PTPVS) for portal vein anastomotic stenosis (PVAS) after liver transplantation (LT).

METHODS: From April 2004 to June 2008, 16 of 18 consecutive patients (11 male and 5 female; aged 17-66 years, mean age 40.4 years) underwent PTPVS for PVAS. PVAS occurred 2-10 mo after LT (mean 5.0 mo). Three asymptomatic patients were detected on routine screening color Doppler ultrasonography (CDUS). Fifteen patients who also had typical clinical signs of portal hypertension (PHT) were identified by contrast-enhanced computerized tomography (CT) or magnetic resonance imaging. All procedures were performed under local anesthesia. If there was a PVAS < 75%, the portal pressure was measured. Portal venoplasty was performed with an undersized balloon and slowly inflated. All stents were deployed immediately following the predilation. Follow-ups, including clinical course, stenosis recurrence and stent patency which were evaluated by CDUS and CT, were performed.

RESULTS: Technical success was achieved in all patients. No procedure-related complications occurred. Liver function was normalized gradually and the symptoms of PHT also improved following PTPVS. In 2 of 3 asymptomatic patients, portal venoplasty and stenting were not performed because of pressure gradients < 5 mmHg. They were observed with periodic CDUS or CT. PTPVS was performed in 16 patients. In 2 patients, the mean pressure gradients decreased from 15.5 mmHg to 3.0 mmHg. In the remaining 14 patients, a pressure gradient was not obtained because of > 75% stenosis and typical clinical signs of PHT. In a 51-year-old woman, who suffered from massive ascites and severe bilateral lower limb edema after secondary LT, PVAS complicated hepatic vein stenosis and inferior vena cava (IVC) stenosis. Before PTPVS, a self-expandable and a balloon-expandable metallic stent were deployed in the IVC and right hepatic vein respectively. The ascites and edema resolved gradually after treatment. The portosystemic collateral vessels resulting from PHT were visualized in 14 patients. Gastroesophageal varices became invisible on poststenting portography in 9 patients. In a 28-year-old man with hepatic encephalopathy, a pre-existing meso-caval shunt was detected due to visualization of IVC on portography. After stenting, contrast agents flowed mainly into IVC via the shunt and little flowed into the portal vein. A covered stent was deployed into the superior mesenteric vein to occlude the shunt. Portal hepatopetal flow was restored and the IVC became invisible. The patient recovered from hepatic encephalopathy. A balloon-expandable Palmaz stent was deployed into hepatic artery for anastomotic stenosis before PTPVS. Percutaneous transhepatic internal-external biliary drainage was performed in 2 patients with obstructive jaundice. Portal venous patency was maintained for 3.3-56.6 mo (mean 33.0 mo) and all patients remained asymptomatic.

CONCLUSION: With technical refinements, early detection and prompt treatment of complications, and advances in immunotherapy, excellent results can be achieved in LT.

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Key words: Portal vein; Anastomotic stenosis; Venoplasty; Stent; Liver transplantation

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INTRODUCTION

Liver transplantation is an important option in the management of end-stage liver disease, severe acute liver failure and some metabolic liver disorders. Postoperative vascular complications have been well documented. The incidence of portal venous complications following liver transplantation is considered to be relatively uncommon in comparison with hepatic arterial complications, yet they can be potentially devastating and lead to graft loss. In the past, portal venous complications were managed with surgical treatments such as thrombectomy, anastomosis revision or retransplantation. However, surgical management of these complications has been limited by technical difficulties due to postsurgical fibrosis and limitations in the length of the involved venous structures[1]. Percutaneous interventional procedures have gained worldwide acceptance for alleviating the symptoms of portal hypertension and preserving the graft, due to their minimal invasiveness as well as low complication and high success rates[6]. In this study, we retrospectively reviewed 16 cases that received percutaneous transhepatic portal venoplasty and stenting (PTPVS).

MATERIALS AND METHODS

Patients

From April 2004 to June 2008, 16 of 18 consecutive patients (11 male and 5 female; aged 17-66 years, mean age 40.4 years) underwent PTPVS for portal vein anastomotic stenosis (PVAS) after liver transplantation (LT). Nine patients were from other hospitals. One patient had a living donor LT; another patient received a graft from a right-sided intercostal approach (KMP; COOK, USA or Cobra; Terumo, Japan). If there was an anastomotic stenosis < 75%, the portal pressure was measured at the postanastomotic main portal vein or at the level of the hepatic hilar portal bifurcation. A disposable pressure transducer system (Utah Medical Products, USA) was used for measuring portal pressure. The catheter was guided by the 0.035-inch guidewire through the stenotic segment. Postportography including the splenic vein and superior mesenteric vein was obtained (Figure 1A) and the preanastomotic portal pressure was also measured. The criteria for definite diagnosis of PVAS were as follows: stenosis > 50% of the main portal venous diameter[9] and a pressure gradient across the stenosis > 5 mmHg[3,7,8].

Portal vein anastomotic venoplasty was performed with a percutaneous transluminal angioplasty balloon dilatation catheter (Powerflex P3; Cordis, Johnson & Johnson Co., USA or Synergy; Boston Scientific Co., USA). The balloon had a smaller diameter than that of the allograft portal vein. Balloon pressure was slowly and gradually elevated using an inflation syringe (Basix Compak; Merit Medical Systems Co., USA) within a pressure limit of 10-15 atm until the balloon’s waist was effaced (Figure 1B). Postvenoplasty portography was repeated to assess the results (Figure 1C). A self-
expandable metallic stent (SMART Control; Cordis, USA) was deployed to cover the stenosis with minimal angulation between the portal vein and the stent. The stent had the same or 1-2 mm larger diameter than that of the allograft portal vein. Poststenting portography (Figure 1D) and the pressure gradient were obtained repeatedly to assess the results. If the deployed stent showed an hourglass deformity of > 50% of its normal diameter, balloon postdilatation was performed. If a satisfactory result had been achieved, the catheter was removed and the puncture tract was embolized with compressed gelfoam bars through the cut vascular sheath.

No intravenous or systemic heparinization was used. Poststenting anticoagulation was achieved by oral administration of aspirin enteric-coated tablets (Bayaspirin, Bayer S.p.A., Italy) 100 mg/d for at least 6 mo.

Technical success, complications, clinical signs and symptoms, laboratory values including the liver function test and the imaging surveillance results after PTPVS were documented. Technical success of the procedure was defined as < 30% residual stenosis being observed on portography with the absence of varices or collateral circulation[3]. Follow-up, including clinical course, stenosis recurrence and stent patency which were evaluated by CDUS and CT, were performed.

RESULTS

Technical success of PTPVS was achieved in all 16 patients. No procedure-related complications occurred. In these patients, liver function was normalized gradually and clinical manifestations related to PHT were improved following PTPVS.

In three asymptomatic patients, a Yashiro type catheter (Terumo, Japan) was introduced into the splenic artery or superior mesenteric artery via a right femoral artery approach and indirect portography was obtained to confirm the diagnosis of PVAS. More than 50% stenosis of portal vein anastomosis was further demonstrated using percutaneous transhepatic portography, but portal venoplasty and stenting were not performed in two patients because pressure gradients across the stenosis were < 5 mmHg. They were observed with periodic CDUS or CT. Portal venoplasty (using 8-10 mm diameter and 40 mm length balloons) and stenting (using 10-12 mm diameter and 40-60 mm length stents) was performed in 16 patients. In 2 patients, the mean initial pressure gradient across the stenosis was 15.5 mmHg and then it decreased to 3.0 mmHg after PTPVS. In the remaining 14 patients, a pressure gradient was not obtained because of > 75% stenosis and typical clinical signs of PHT.

In a 51-year-old woman, who suffered from massive ascites and severe bilateral lower limb edema after secondary LT, CDUS and CT detected PVAS complicated by hepatic vein stenosis and inferior vena cava stenosis. These venous stenoses were identified by indirect portography, percutaneous transhepatic hepatic venography and inferior vena cavaography. First, a self-expandable metallic stent (COOK-Z, GZV-30-75; COOK, USA) was placed in the inferior vena cava via a right femoral vein approach. Second, a balloon-expandable metallic stent (IntraStent, SPM16-26-08-B; ev3 Co., USA) was deployed in the right hepatic vein via an intercostal transhepatic route. Last, PTPVS was performed. The ascites and edema resolved gradually after treatment.
The portosystemic collateral vessels resulting from PHT were visualized by initial portography in 14 patients. Gastroesophageal varices became invisible on poststenting portography in 9 patients. To avoid variceal bleeding, the residual gastroesophageal flow was obstructed with platinum embolization coils (Tornado, MWCE-35-8/4-5-Tornado; COOK, USA) in 2 remaining patients. No further procedure was performed for anorectal varices and pre-existing splenorenal shunt because the abnormal flow became reduced after stenting.

In a 28-year-old man with hepatic encephalopathy, a pre-existing meso-caval shunt was detected due to visualization of the IVC on initial portography (Figure 2A and B). A meso-caval shunt was performed due to refractory massive ascites and recurrent variceal bleeding before LT. After stenting, it was seen that the contrast agents flowed mainly into the IVC via the shunt and little flowed into the portal vein (Figure 2C). To maintain adequate hepatopetal perfusion pressure and to avoid thrombosis in the portal stent or liver failure, a covered stent (Wallgraft Endoprosthesis; Boston Scientific, USA) with a 10 mm diameter and a 50 mm length was deployed into the superior mesenteric vein to occlude the shunt. Once more portography revealed that portal hepatopetal flow was restored and the IVC became invisible (Figure 2D). The patient felt abdominal pain after the procedure, but the symptom subsided a week later. The patient recovered from hepatic encephalopathy.

A balloon-expandable Palmaz stent (Genesis PG1840PMW; Cordis, USA) was deployed into the hepatic artery via a right femoral artery route for anastomotic stenosis 2 mo before PTPVS. Percutaneous transhepatic internal-external biliary drainages were performed with a biliary drainage catheter (Ultrathane MAC-LOC, ULT8.5-38-40-P-32S-CLB-RH; COOK, USA) in 2 patients complicated with obstructive jaundice. The follow-up results of CDUS in all 16 patients and CT scan in 6 patients revealed portal venous patency was maintained for 3.3-56.6 mo (mean 33.0 mo).

DISCUSSION

Portal venoplasty and stenting

The rate of portal venous complications after LT, which include primary portal vein anastomotic stenosis or portal vein thrombosis, has been reported to be below 3% [9-11]. However, in children with reduced-size LT and living donor LT, the incidence of portal venous stenosis or thrombosis is higher than in adults with deceased donor transplantation, because the donor portal segment is relatively short, and due to interposition grafts and size mismatch of the portal vein diameter between donors and recipients. Factors that increased the risk of portal venous complications were pre-existing vein thrombosis or hypoplasia and large portocaval collaterals [12].

Since the first report of portal venous angioplasty and stent placement through a transhepatic approach after LT by Olcott et al [13] in 1990, percutaneous transhepatic interventional procedures have gained worldwide acceptance for treatment of these complications following LT, due to their minimal invasiveness as well as low complication and high success rates [2,14]. Nonetheless, the reported recurrence rate has been relatively high, i.e. 28.6%-36.8%, following balloon angioplasty alone [3,6,7].

Stents have usually been used to treat recurrent and elastic portal venous stenoses following balloon angioplasty, as this procedure has several potential complications [3,6,7]. Nevertheless, Ko et al [8] preferred to perform primary stent placement rather than balloon angioplasty in the early posttransplantation period (less than 1 mo). In our study, all stents were deployed immediately following balloon angioplasty for two reasons. First, balloon predilation can reduce the incidence of stent displacement (most jump forward) during stent deployment, especially when the stenosis is severe. Second, direct venoplasty and stenting can recover the normal portal flow once and for all, because repeat percutaneous transhepatic portal venoplasty may lead to puncture injuries in the transplanted
liver and increase the incidence of procedure-related complications, such as intrahepatic pseudoaneurysm, arteriovenous fistulas, subcapsular hematoma, bleeding through the puncture tract of the graft, or portal venous thrombosis.

The luminal area is proportional to the square of the radius; flow is proportional to the fourth power of the radius. Thus, any small improvement is magnified. Intimal damage can lead to platelet aggregation that can in turn lead to short-term occlusion or long-term restenosis. Therefore, any intimal tears should be avoided at all costs, even to the extent of accepting a less visually appealing angiographic result. Slow and gradual inflation of the balloon during angioplasty results in fewer large intimal tears, flaps, and dissections than the more commonly used method of “blow it up, let it down”[5]. In our study, portal venoplasty was performed with an undersized balloon and the balloon was slowly and gradually inflated. Although portal vein thrombosis or stent-edge stenosis may occur, the follow-up results revealed portal venous patency was maintained for a mean 33.0 mo.

Although the etiology of anastomotic stenosis was unclear in our patients, it suggested fibrosis or intimal hyperplasia, which retained an hourglass deformity on the deployed stent.

Role of imaging examination

It is important to detect vascular complications after liver transplantation, because most stenoses or thromboses are frequently treatable with interventional procedures; but, if left untreated, many vascular complications may progress to graft failure. However, compared with biliary obstructions, early portal venous stenosis is difficult to detect from clinical signs and symptoms alone. Furthermore, sometimes the portal venous anastomotic site cannot be seen with ultrasonography because of intestinal artifacts. In such a case, other noninvasive cross-sectional imaging modalities, CT and MRI, may be valuable.

Ultrasonography is often used to screen for vascular abnormalities, including hepatic arterial stenosis and thrombosis, and the less common stenoses or thromboses of the portal vein, hepatic veins, and inferior vena cava. Precise anatomy of the vascular abnormalities is often better determined on CT or MRI, especially when a focal stenosis occurs in the distal IVC, or in the hepatic artery proximal to the porta hepatis where it is difficult to image directly by ultrasonography. CT and MRI give detailed imaging, while ultrasonography tends to give more physiologic data. CT and MRI can provide a more comprehensive evaluation of the transplanted liver; reveal abnormalities of vascular structure; and depict bile ducts, liver parenchyma, and extrahepatic tissues. Moreover, CT angiography and MR angiography can be used to evaluate the extent and degree of the portosystemic collateral vessels resulting from PHT. Magnetic resonance cholangiopancreatography can be valuable to detect focal biliary abnormalities; however, percutaneous transhepatic cholangiography remains the gold standard for biliary complications[6].

When a patient is asymptomatic, indirect portography is a recommended option to identify a portal venous stenosis, as this procedure has a relatively lower incidence of procedure-related complications and no puncture injuries to the graft. When there is a requirement for measurement of portal pressure gradients across a stenosis or further interventional procedures, percutaneous transhepatic portography is necessary.

Portal venography with measurement of pressure gradient across the stenosis remains the most reliable examination[4], but, the procedure is an invasive one. Although some reports have considered a transstenotic pressure gradient of > 5 mmHg as abnormal[7,8], no standard guidelines for a significant pressure gradient have yet been established. Park et al[9] believed that the pressure gradient is not directly correlated with the clinical results, and mentioned that portal venoplasty might not be so helpful for patients whose clinical symptoms are possibly related with graft dysfunction and not with the stenosis. The treatment is valuable if patients have symptoms related to portal venous inflow abnormality or PHT, even though the pressure gradient is not significant[4]. In patients who do not have evidence of PHT, and have normal hepatic function, stenoses may be observed for progression with periodic ultrasound. Moreover, in patients with PHT, the potential contribution of underlying hepatic parenchymal disease (rejection or recurrent hepatitis) must be considered. However, if portal venous stenosis is suspected as being a significant contributor to PHT, therapeutic intervention is necessary.

Negative findings on serial CDUS and the absence of clinical symptoms during the follow-up period might prompt us to regard these patients as not having any hemodynamically significant vascular abnormalities.

In conclusion, percutaneous transhepatic portal venoplasty and stenting for anastomotic stenosis after liver transplantation is a safe and effective procedure for alleviating the signs and symptoms of portal hypertension and preserving the graft. With technical refinements, early detection and prompt treatment of complications, and advances in immunotherapy, excellent results can be achieved in liver transplantation.
recurrent and elastic portal venous stenoses following balloon angioplasty, as this procedure has several potential complications. Nevertheless, Ko et al preferred to perform primary stent placement rather than balloon angioplasty in the early posttransplantation period (< 1 mo).

**Innovations and breakthroughs**

In this study, all stents were deployed immediately following balloon angioplasty for two reasons. First, balloon predilation can reduce the incidence of stent displacement during its deployment. Second, direct venoplasty and stenting can recover normal portal flow once and for all, because repeat percutaneous transhepatic portal venoplasty may lead to puncture injuries to the transplanted liver and increase the incidence of procedure-related complications. Intimal damage can lead to platelet aggregation that can in turn lead to short-term occlusion or long-term restenosis. Slow and gradual inflation of the balloon during angioplasty results in fewer large intimal tears, flaps, and dissections than the more commonly used method of “blow it up, let it down.” In this study, portal venoplasty was performed with an undersized balloon and the balloon was slowly and gradually inflated. Although portal vein thrombosis or stent-edge stenosis may occur, the follow-up results revealed portal venous patency was maintained for a mean of 33.0 mo. The treatment is valuable if patients have symptoms related to portal venous inflow abnormality or portal hypertension even though the pressure gradient is not significant.

**Applications**

Percutaneous transhepatic portal venoplasty and stenting (PTPVS) could be used for portal anastomotic stenosis after liver transplantation to alleviate the signs and symptoms of portal hypertension and preserve the graft.

**Peer review**

PTPVS for anastomotic stenosis after liver transplantation is a safe and effective procedure for alleviating the signs and symptoms of portal hypertension and preserving the graft. The results are encouraging and suggest that the method of undersized balloon and slow and gradual inflation can reduce intimal damage and keep the portal venous patency for a mean of 33.0 mo after stenting.

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