Case Report



Funneling venoplasty for anomalous graft left hepatic vein in living donor liver transplantation using a split left lateral section graft for an infant patient

Jung-Man Namgoong¹, Shin Hwang¹, Tae-Yong Ha¹, Young-In Yoon¹, Yong Jae Kwon¹, Hyunhee Kwon¹, Kyung Mo Kim², Seak Hee Oh²

¹Department of Surgery, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea, ²Department of Pediatrics, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea

The left lateral section (LLS) can have an unusual variant left hepatic vein (LHV) anatomy. We present a case of customized funneling venoplasty of the graft LHV in a 22-month-old girl diagnosed with ornithine transcarbamylase deficiency undergoing deceased donor liver transplantation (LT) using a split LLS graft. The split LLS graft weighed 350 g, yielding a graft-to-recipient weight ratio of 3.2%. Notably, the graft LHV opening was located at the graft liver cut surface, which was only 1 cm in size and 2 cm away from the cepha-lad apex of the LLS graft. Since such a variant location of the small LHV opening was unsuitable for direct anastomosis, we performed a funneling venoplasty using an inferior vena cava fragment homograft obtained from the same donor. The graft implantation was performed according to standard procedures of infant split LT. Follow-up imaging studies showed no vascular complications. The patient recovered uneventfully from the LT operation. She had normal blood test findings, including normal ammonia level. She has been doing well for 6 months after the transplantation. In conclusion, our surgical technique using a funneling venoplasty enabled successful reconstruction of the anomalous graft LHV. Our results suggest that individualized reconstruction techniques should be applied to infant patients undergoing LT using a LLS graft with variant types of graft LHV anatomy.

Key Words: Left hepatic vein; Anatomical variation; Venoplasty; Interposition; Left lateral section graft

INTRODUCTION

A left lateral section (LLS) graft is often used for liver transplantation (LT) in infant patients in the form of living donor liver transplantation (LDLT) or split LT. Although the anatomy of the left hepatic vein (LHV) is diverse [1-4], the majority of graft LHVs are suitable for direct anastomosis to the recipient hepatic vein stumps or directly to the inferior vena cava (IVC). However, in rare instances, the LLS outflow drains through both the LHV and the middle hepatic vein (MHV) or the

Received: January 15, 2021, Revised: January 21, 2021, Accepted: January 23, 2021

Corresponding author: Shin Hwang

Department of Surgery, Asan Medical Center, University of Ulsan College of Medicine, 88 Olympic-ro 43-gil, Songpa-gu, Seoul 05505, Korea Tel: +82-2-3010-3930, Fax: +82-2-3010-6701, E-mail: shwang@amc.seoul.kr ORCID: https://orcid.org/0000-0002-9045-2531

Copyright © The Korean Association of Hepato-Biliary-Pancreatic Surgery This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/by-nc/4.0) which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited. LHV drains directly through the MHV trunk [1]. In donors with such unusual LHV anatomy, it is necessary to preserve the MHV trunk for the safety of the living donor or for the security of the split extended right liver graft. If the graft LHV opening is located at the cut surface of the LLS graft instead of at the cephalic apex, a customized venoplasty technique is necessary to make it suitable for graft hepatic vein reconstruction, especially for an infant recipient with a small IVC. We present a case of customized venoplasty of the graft LHV in an infant patient undergoing deceased donor LT using a split LLS graft.

CASE

The recipient was a 22-month-old girl who was diagnosed with ornithine transcarbamylase (OTC) deficiency. The patient was born through a full-term cesarean-section delivery. She showed irritability and decreased activity from one month after birth. At that time, laboratory studies showed hyperammonemia and metabolic acidosis with high levels of liver enzymes. Gene studies revealed OTC NM_000531.5:c.626C>T. p.Ala209V. Het. The patient was placed on the waiting list of the Korean Network for Organ Sharing (KONOS) with a Pediatric End-stage Liver Disease score of 4 (total bilirubin level of 0.6 mg/dL, albumin level of 4.4 g/dL, prothrombin international normalized ratio of 1.37, and growth failure).

After a waiting period of one year, a 35-year-old female de-

ceased donor was allocated for split LT. At organ allocation, the patient's height and body weight were 85 cm and 11 kg, respectively. The split LLS graft weighed 350 g, yielding a graft-to-recipient weight ratio (GRWR) of 3.2%.

Notably, the graft LHV opening was located at the graft liver cut surface, which was only 1 cm in size and 2 cm away from



Fig. 1. Classification of the hepatic vein anatomy in the left lateral section in terms of patterns of the left lateral section graft hepatic vein openings. Type 1 makes a single opening. Type 2 makes two widely spaced openings. Type 3 makes large and small adjacent openings. Type 4 makes two widely spaced openings. Crossed circles indicate the location of the umbilical portion. Cited from the article of Hwang et al. (Liver Transpl 2013;19:184-190) [1].



Fig. 2. Presumed anatomy of the donor liver. The anatomy of the left hepatic vein appears to be a mixed type of two images with narrow (A) and wide (B) distances between the large and small hepatic vein openings. The small opening indicates the superficial branch of the left hepatic vein (arrows).



Fig. 3. Design of customized funneling venoplasty for the graft left hepatic vein opening. The 1 cm-sized orifice (A) is partially incised to increase the diameter (B). A vein patch is attached at the enlarged graft hepatic vein opening to make a funnel-shaped conduit (C). Arrow indicates a slit incision.

the cephalad apex of the LLS graft. The native anatomy of the donor LHV was presumed to be type 3 (Fig. 1, 2) [1]. Since such a variant location of the small LHV opening was unsuitable for direct anastomosis, we performed a funneling venoplasty using an IVC fragment homograft obtained from the same donor (Fig. 3, 4).

Because there was no anatomical variation in the recipient (Fig. 5), standard procedures of pediatric split LT were performed. After dissection of the recipient native liver was completed, the hepatic parenchyma was incised with a surgical knife, leaving a bulk of hepatic parenchyma around the hepatic vein trunks. The hepatic parenchyma was forcefully pulled out to detach it from the hepatic vein stumps, which made stump walls long and thick. No venoplasty was applied to the recipient IVC orifice. The interposed funnel-shaped orifice of the graft hepatic vein was anastomosed with the recipient IVC orifice by 1 : 1 size matching (Fig. 6). The recipient portal vein was normal-looking, thus it was anastomosed with the graft portal vein stump using a branch patch (Fig. 7). The graft hepatic artery was reconstructed under surgical microscopy. Finally, Roux-en-Y hepaticojejunostomy was performed.

The pathology report of the explant liver revealed enlarged hepatocytes with pale cytoplasm and halo nuclei, and multifocal aggregates of apoptotic cells, consistent with OTC (Fig. 8). Early follow-up computed tomography scan showed no vascular complications (Fig. 9). The patient recovered uneventfully



Fig. 4. Intraoperative photographs of bench work. (A–C) An inferior vena cava fragment homograft is attached to the graft hepatic vein orifice to make a funnel-shaped conduit. (D) The cephalad end of the conduit is much larger than the graft side.



Fig. 5. Preoperative computed tomography images showing no gross abnormality in liver shape (A) or hepatic vasculature (B).

www.ahbps.org

from the LT operation. She had normal blood test findings including normal ammonia level. She has been doing well for six months after the transplantation.

DISCUSSION

LLS grafts rarely have LHV variations of sufficient significance to preclude direct reconstruction because a variant LHV can join with the MHV trunk in the left medial section. Because the left medial section parenchyma is sacrificed during LLS graft procurement [5,6], the LHV branches in the left medial section can be harvested to obtain a single graft hepatic vein orifice. However, infant recipients usually require small LLS grafts to adequately match the GRWR. Thus, LLS grafts usually only have the LHV trunk. If the graft LHV orifice is

unsuitable for direct anastomosis, it is necessary to use customized venoplasty of the graft LHV to prevent graft hepatic

We have previously classified the LHV anatomy of 300 potential LLS graft donors into four types according to the

number and location of graft LHV openings: single opening

(type 1; n = 218, 72.7%), two large adjacent openings (type 2; n

= 29, 9.7%), one large and one small adjacent opening (type 3;

n = 34, 11.3%), and two widely spaced openings (type 4; n = 19,

Fig. 6. Intraoperative photographs of recipient hepatic vein reconstruction. (A) The three hepatic vein openings are widely opened to make an enlarged orifice. (B) The hepatic vein openings at the recipient inferior vena cava and the graft are well matched in size. (C) The posterior wall of the hepatic vein reconstruction is visible. (D) The anterior wall of the hepatic vein reconstruction is visible.

Fig. 7. Intraoperative photographs of left lateral section graft implantation. (A) The hepatic vein reconstruction is located at the orthodox position. (B) Portal vein reconstruction is performed using a branch patch of the recipient portal vein.

Fig. 7. Intraoperative lateral section graft in hepatic vein reconst at the orthodox positi reconstruction is perfo

vein outflow obstruction (HVOO) [1,7].



6.3%) (Fig. 1). LHV types 2 and 3 require wedged unification venoplasty and type 4 requires additional vein interposition [1]. The LHV anatomy of the present LLS graft can be classified as a type 3, in which one large LHV branch is drained directly through the MHV trunk with the presence of a small superficial LHV branch. We had thought that this superficial LHV branch could be used for unification venoplasty of the LHV



Fig. 8. Gross photograph of the explant liver.

opening [7]. However, we had to sacrifice it in the present case because it was widely apart from the LHV trunk with a size less than 2 mm.

In our previous LDLT case using a LLS graft with type 4 LHV, customized interposition-wedged unification venoplasty was used and the source of the interposition vein conduit was an ilio-femoral vein homograft [1]. In the present case, because the required length of the vein conduit was 2 cm and the diameter of the graft LHV opening was only 1 cm, we thought that direct anastomosis might have a high risk of graft HVOO. To cope with the anatomical variation, we made a funneling venoplasty using an IVC fragment homograft patch to connect the recipient IVC orifice and the graft LHV orifice. The final configuration of the graft hepatic vein reconstruction appeared natural and stream-lined, as in an LLS graft with type 1 LLV anatomy.

LDLT or split LT for infant recipients is vulnerable to vascular complications because the graft and recipient vessels are much smaller than those in adult-to-adult LT. Anastomotic stenosis following hepatic vein reconstruction using an LLS graft is usually attributed to the small size of vascular anastomosis. Once HVOO develops, it is often difficult to treat it through radiologic angioplasty [8-11]. Insertion of a wall stent to treat HVOO is considered a life-saving procedure, with a high likelihood of the need for retransplantation later because such a vascular wall stent may not expand sufficiently during physical growth of the recipient from infancy to adolescence [12,13].



Fig. 9. Follow-up computed tomography images taken 4 days after liver transplantation. The hepatic vein reconstruction (A, C) appears smooth and streamlined without stenosis. The portal vein reconstruction (B, D) shows a size discrepancy without noticeable stenosis.

Therefore, graft hepatic vein reconstruction requires a secure surgical design, particularly in infant recipients.

In conclusion, our surgical technique using funneling venoplasty enabled successful reconstruction of the anomalous graft LHV. Our results suggest that individualized reconstruction techniques should be applied for infant patients undergoing LT using a LLS graft with variant types of graft LHV.

CONFLICT OF INTEREST

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

ORCID

Jung-Man Namgoong, https://orcid.org/0000-0002-9237-7440 Shin Hwang, https://orcid.org/0000-0002-9045-2531 Tae-Yong Ha, https://orcid.org/0000-0001-9932-0212 Young-In Yoon, https://orcid.org/0000-0002-9308-0366 Yong Jae Kwon, https://orcid.org/0000-0001-9490-1229 Hyunhee Kwon, https://orcid.org/0000-0001-6647-9155 Kyung Mo Kim, https://orcid.org/0000-0001-7896-6751 Seak Hee Oh, https://orcid.org/0000-0002-9672-8877

AUTHOR CONTRIBUTIONS

Conceptualization: SH. Data curation: TYH, YIY, YJK, HK, KMK, SHO. Methodology: JMN, SH. Visualization: SH. Writing - original draft: JMN, SH. Writing - review & editing: All authors.

REFERENCES

- Hwang S, Kim KH, Kim DY, Kim KM, Ahn CS, Moon DB, et al. Anomalous hepatic vein anatomy of left lateral section grafts and customized unification venoplasty for pediatric living donor liver transplantation. Liver Transpl 2013;19:184-190.
- 2. Hwang S, Lee SG, Choi ST, Moon DB, Ha TY, Lee YJ, et al. Hepatic vein anatomy of the medial segment for living donor liver transplantation using extended right lobe graft. Liver Transpl 2005;11:449-455.
- 3. Radtke A, Sotiropoulos GC, Sgourakis G, Molmenti EP, Schroeder T, Saner FH, et al. Hepatic venous drainage: how much can we learn

from imaging studies? Anatomic-functional classification derived from three-dimensional computed tomography reconstructions. Transplantation 2010;89:1518-1525.

- Mochizuki K, Takatsuki M, Soyama A, Hidaka M, Obatake M, Eguchi S. The usefulness of a high-speed 3D-image analysis system in pediatric living donor liver transplantation. Ann Transplant 2012;17:31-34.
- Hwang S, Lee SG, Lee YJ, Park KM, Ahn CS, Kim KH. Postoperative changes in remnant medial segment parenchyma of living donor livers after procurement of left lateral segment graft. Hepatogastroenterology 2006;53:773-777.
- 6. Seda-Neto J, Godoy AL, Carone E, Pugliese V, Fonseca EA, Porta G, et al. Left lateral segmentectomy for pediatric live-donor liver transplantation: special attention to segment IV complications. Transplantation 2008;86:697-701.
- Namgoong JM, Hwang S, Park GC, Ahn CS, Kim KH, Kim KM, et al. Outflow vein venoplasty of left lateral section graft for living donor liver transplantation in infant recipients. Pediatr Transplant 2021;25:e13970.
- Galloux A, Pace E, Franchi-Abella S, Branchereau S, Gonzales E, Pariente D. Diagnosis, treatment and outcome of hepatic venous outflow obstruction in paediatric liver transplantation: 24-year experience at a single centre. Pediatr Radiol 2018;48:667-679.
- Katano T, Sanada Y, Hirata Y, Yamada N, Okada N, Onishi Y, et al. Endovascular stent placement for venous complications following pediatric liver transplantation: outcomes and indications. Pediatr Surg Int 2019;35:1185-1195.
- Zhang ZY, Jin L, Chen G, Su TH, Zhu ZJ, Sun LY, et al. Balloon dilatation for treatment of hepatic venous outflow obstruction following pediatric liver transplantation. World J Gastroenterol 2017;23:8227-8234.
- Lu KT, Cheng YF, Chen TY, Tsang LC, Ou HY, Yu CY, et al. Efficiency of transluminal angioplasty of hepatic venous outflow obstruction in pediatric liver transplantation. Transplant Proc 2018;50:2715-2717.
- Yeh YT, Chen CY, Tseng HS, Wang HK, Tsai HL, Lin NC, et al. Enlarging vascular stents after pediatric liver transplantation. J Pediatr Surg 2017;52:1934-1939.
- 13. Namgoong JM, Hwang S, Yoon YI, Cho YP, Kang WH, Kwon YJ, et al. Third retransplantation using a whole liver graft for late graft failure from hepatic vein stent stenosis in a pediatric patient who underwent split liver retransplantation. Ann Hepatobiliary Pancreat Surg 2021;25:299-306.