

Inverted Donor Inferior Vena Cava Conduit to Bypass an Inferior Vena Cava Obstruction in a Kidney Transplant Recipient

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Background	Ileocaval venous obstruction in kidney transplant recipients may preclude a conventional external iliac venous anastomosis if adequate allograft venous outflow cannot be achieved. We share the case of a 69-year-old male with a history of inferior vena cava (IVC) ligation who presented for kidney transplantation. A unique surgical approach was utilized for venous drainage of an iliac fossa deceased-donor allograft.
Summary	Our patient presented for a deceased-donor renal transplant. He had a history of deep vein thrombosis (DVT) requiring remote infrarenal IVC ligation after developing intraabdominal bleeding while on anticoagulation. After inspection of the patient's computed tomography (CT) imaging and the deceased-donor right kidney, which had been procured with a substantial length of proximal IVC, he was deemed a candidate for iliac fossa allograft implantation using an inverted donor IVC conduit. The donor renal vein was divided from the donor IVC, then rotated 180°. The donor renal vein was re-implanted at its original orifice, creating a conduit from the renal hilum running superiorly. An end-to-side anastomosis between the inverted donor IVC and recipient IVC was created above the IVC obstruction. He experienced immediate and sustained allograft function at his latest follow-up three months post transplantation.
Conclusion	Iliac vein and IVC anomalies represent a surgical challenge for renal transplantation. We present a case where an inverted donor IVC conduit was created to bypass an infrarenal IVC obstruction, resulting in successful iliac fossa allograft implantation. This case highlights how preoperative planning, procedural creativity, and knowledge regarding alternative venous drainage options and implantation sites for renal allografts can assist in identifying solutions to these technical challenges.
Key Words	kidney transplantation; surgical technique; ileocaval obstruction
Abbreviations	COPD: chronic obstructive pulmonary disorder DOAC: direct-acting oral anticoagulant DVT: deep vein thrombosis GSV: greater saphenous vein IVC: inferior vena cava PE: pulmonary embolism

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Case Description

Ileocaval venous obstruction can be caused by a variety of etiologies, including congenital atresia, thrombosis, or iatrogenic. When present in renal transplant recipients, these anomalies may preclude a conventional external iliac venous anastomosis, as a low-pressure venous outflow from the allograft is required to minimize the risk of graft failure. The patient should therefore be evaluated for an alternative site of venous drainage. The anatomical characteristics of the procured organ may also influence the surgical approach. We report a unique case of surgical management of a kidney transplant recipient with an inferior vena cava (IVC) obstruction.

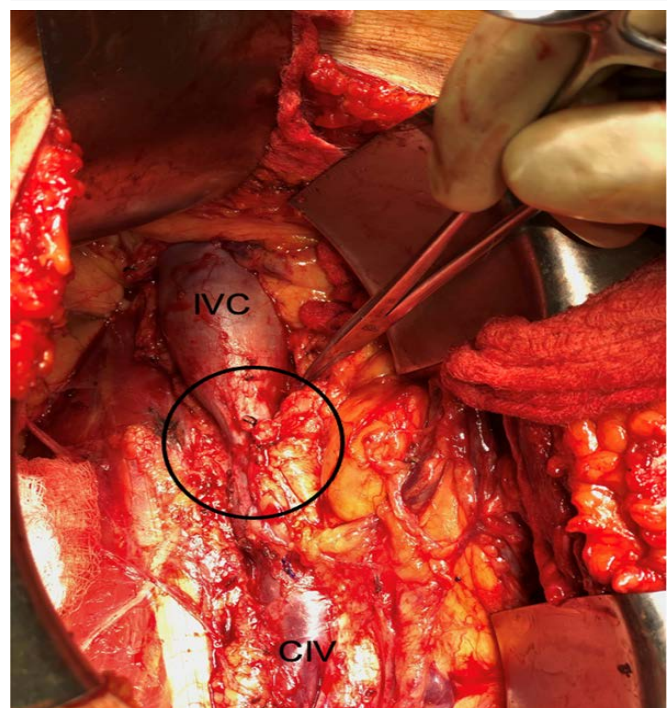
The patient is a 69-year-old male with a history of severe chronic obstructive pulmonary disorder (COPD) requiring bilateral lung transplants and end-stage renal disease secondary to resulting calcineurin toxicity, who presented to our institution for deceased-donor kidney transplantation. He also had a history of deep vein thrombosis and pulmonary embolism (DVT/PE) requiring remote infrarenal IVC ligation to prevent thrombosis propagation after developing intraabdominal bleeding while on anticoagulation. A computed tomography (CT) scan of the abdomen and pelvis was obtained on his arrival at our facility to delineate his abdominal venous outflow better and assist with surgical planning (Figure 1).

The right-sided deceased donor kidney was then inspected. The donor renal vein had been procured along with 2 cm proximal and 8 cm distal donor IVC. The patient was taken to the operating room for a planned renal transplant in the right iliac fossa, using recipient IVC just proximal to his obstruction as the site for venous drainage of the allograft. An inverted donor IVC conduit was created to maximize the proximity of the allograft to the right iliac artery and bladder. The donor renal vein was divided from the donor IVC, which was then rotated 180°. The donor renal vein was re-implanted at its original orifice, now at the inferior aspect of the IVC, creating a conduit from the renal hilum running superiorly. The inferior IVC cuff was then oversewn. A right lower quadrant curvilinear incision and a right retroperitoneal dissection were performed (Figure 2).

Figure 1. Location of Infrarenal IVC Ligation (arrow). Published with Permission



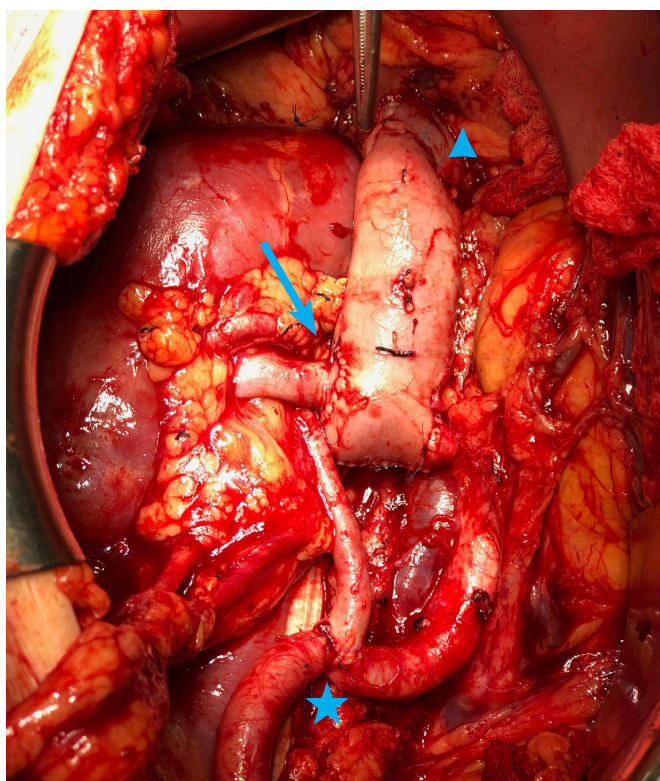
Figure 2. Intraoperative Photo. Published with Permission



Retroperitoneal exposure of the infrarenal IVC obstruction (circled) and common iliac vein (CIV)

An end-to-side anastomosis between the inverted donor IVC and recipient IVC was created above the obstruction and below the native renal veins, and an end-to-side anastomosis between the donor renal artery and recipient external iliac artery was created (Figure 3). A Lich ureterocystostomy was then performed.

Figure 3. Intraoperative Photo. Published with Permission



End-to-side donor renal vein to donor IVC conduit anastomosis (arrow). End-to-side donor IVC to recipient IVC anastomosis (arrowhead). Conventional end-to-side donor renal artery to recipient external iliac anastomosis (star).

There were no surgical complications, and the patient experienced immediate allograft function. He was discharged on postoperative day 7. Graft function has remained excellent at his latest follow-up three months post-transplantation.

Discussion

Ileocaval venous anomalies in kidney transplant recipients present a technical challenge to renal transplantation. Venous anomalies may be identified by patient history, physical exam, preoperative imaging, or incidentally intraoperatively. They can be due to various etiologies,

including thrombosis, congenital agenesis, extrinsic compression, or intrinsic stenosis due to intraluminal devices. An ileocaval anomaly impeding venous outflow is not an absolute contraindication to conventional external iliac venous anastomosis; there are case reports of successful renal transplantation when patients are maintained on life-long anticoagulation, and collateral drainage was deemed adequate on venography.^{1,2} However, high-pressure venous outflow can lead to acute renal vein thrombosis, graft failure, delayed thrombosis,³ chronic graft dysfunction,⁴ and or abdominal wall phlebitis.⁵ Therefore, an iliac vein or IVC anomaly in a potential kidney transplant recipient should prompt evaluation of ileocaval venous outflow and consider alternative venous drainage options and allograft implantation sites, taking into account both patient and allograft anatomy.

Successful renal transplantation using nontraditional venous drainage sites has been widely reported and can be broadly categorized into systemic or portal venous anastomoses. Systemic options amenable to iliac fossa graft placement include the IVC,^{1,4,6-10} large collateral vein,¹⁰⁻¹² or gonadal vein.¹³⁻¹⁵ An orthotopic placement using the native renal vein,^{4,16} is another systemic option, though a native nephrectomy is also often required to create space for the graft. Portal venous drainage options require intraabdominal graft placement and are, therefore, at a greater risk of vascular torsion compared to retroperitoneal graft placement. Portal options include the splenic vein,^{7,17-20} IMV,^{21,22} SMV,^{16,23,24} portal vein,^{16,25,26} and hepatic vein.²⁷

Our patient's preoperative imaging demonstrated a patent portion of his infrarenal IVC, which could be easily accessed through an elongated paramedian incision allowing for iliac fossa placement, which was preferred due to its proximity to the iliac artery and bladder. However, elongation of the donor renal vein was required to reach the patient IVC. There are two case reports where an extension of the donor renal vein was used to reach its destination; one case used a donor iliac vein interposition graft to bridge the donor renal vein to the recipient portal vein,²⁵ while the other involved procurement of the ovarian vein and greater saphenous vein (GSV) along with the kidney from a living donor.⁶ In the latter case, the GSV was ultimately used to bridge the donor renal vein to the infrahepatic IVC. Our case demonstrates that the donor IVC procured with the deceased-donor allograft is another valuable option for renal vein elongation and may be inverted to bridge the donor renal vein and the recipient infrarenal IVC.

Conclusion

Iliac vein and IVC anomalies in kidney transplant recipients preclude a conventional external iliac venous anastomosis if there is potential for allograft venous outflow hypertension. These anomalies should prompt consideration of alternative systemic or portal venous drainage options, which may require orthotopic or intraabdominal graft placement. Both patient and allograft anatomy can influence the surgical approach. We report a case of a successful kidney transplant in a patient with an infrarenal IVC obstruction, where the patient and allograft anatomy allowed for an iliac fossa placement of a graft using an inverted donor IVC conduit to bypass his IVC obstruction.

Lessons Learned

Ileocaval venous anomalies in kidney transplant recipients should prompt consideration of unconventional venous drainage options. Both patient and allograft anatomy inform surgical decision-making. An infrarenal IVC obstruction can be successfully bypassed using an inverted donor IVC conduit for venous drainage of an iliac fossa deceased-donor allograft.

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