En-Bloc Liver–Pancreas Transplant in Iran

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Abstract
Liver transplantation can be challenging in cirrhotic patients with diabetes mellitus. In chronic liver disease, the glucose metabolism is altered; uncontrolled diabetes negatively influences the outcome of liver transplantation and poses difficulty in the management of immediate post transplantation period. Simultaneous liver-pancreas transplantation is an option to prevent early complications due to diabetes and also to improve the quality of life after transplantation in patients with Insulin-Dependent Diabetes Mellitus (IDDM) and chronic liver disease. We report the first en-bloc liver-pancreas transplant done in the transplant history of Iran. We describe the technical details of the procedure as well as the short term outcome after transplantation. In this case report, we also discuss in some details, the surgical, medical and immunological advantages of combined liver–pancreas transplantation as opposed to separate implantation of both organs.

Keywords: Cirrhosis, diabetes mellitus, en-bloc, liver-pancreas, transplantation


Introduction
Combined liver-pancreas transplantation was formerly used as a salvage procedure for non-resectable upper abdominal malignancies, but the procedure was almost abandoned due to poor outcome.1-3 Recently, the indication for combined liver-pancreas transplantation has been changed and transplant surgeons have started using this technique for patients with IDDM who are candidates for liver transplantation at the same time. There are several advantages to using simultaneous liver-pancreas transplantation such as insulin independence after transplantation, improving patient management and decreasing the risk of post transplant cardiovascular diseases.4 Diabetic patients who receive only liver graft are not only at increased risk of developing cardiovascular diseases but also remain diabetic after transplantation which may itself negatively affect the long-term graft survival.4-6 Though there are so many advantages to combined liver-pancreas transplantation, surprisingly only few cases have been reported in the literature.7

Case Report
A 25-year old male (64 Kg in weight) with a history of IDDM since age 11, presented with intractable pruritis and jaundice at 19 years of age. His laboratory findings revealed elevated cholestatic enzymes and ERCP showed multiple bile duct strictures consistent with primary sclerosing cholangitis (P.S.C) which was later confirmed on liver biopsy.

In June 2012, the patient referred with advanced liver disease. He had intractable and refractory pruritis, was deeply jaundiced and anemic. He was on insulin therapy (90 units/day) and had a history of recurrent hypoglycemic episodes recently. His examination revealed no signs of diabetic neuropathy or retinopathy. At the time of admission, the laboratory parameters were as follows: total bilirubin 18.2 mg/dL, AST 106 U/L, ALT 73 U/L, Alkaline Phosphatase 3919 U/L, BUN 19 mg/dL, Serum creatinine 0.8 mg/dL and Hemoglobin 7.7 g/dL. Bleeding profile was within normal limits.

He received en-bloc liver-pancreas graft from an ABO identical deceased donor aged 15 years. Liver biopsy of the graft showed no macro- or microsteatosis.

Operative procedure started with a classic Mercedes incision. Hepatectomy was performed in standard fashion with caval cross clamping. The liver, along with the en-bloc duodenopancreatic graft, was then transplanted orthotopically. First, supra and infra hepatic caval anastomoses were done. Inflow was established by anastomosing the recipient’s portal vein to the infrapancreatic superior mesenteric vein of the graft. Arterial anastomosis was performed between inferior orifice of the donor’s aorta (including both celiac and the superior mesenteric arteries) to infra renal aorta of the recipient in end-to-side fashion. Finally, a Roux-en-y enterenteric anastomosis was performed between the recipient’s jejunum and graft duodenum for exocrine pancreatic and biliary drainage. Total duration of the operation was 360 minutes and 3 units of packed cell were transfused. Total cold ischemia time was 9 hours while warm ischemia time was 80 minutes. Post operative recovery was uneventful and the patient was discharged on the 15th post operative day with normal liver function tests and free of insulin therapy. Initial immunosuppression included induction with Alemutuzumab (Campath 1H) and maintenance therapy with Tacrolimus (target trough level 12 – 15 ng/mL) and mycophenolate mofetil (MMF) 2 g/day. Steroids were tapered down and completely withdrawn within 6 months post transplantation. The
protect these organs from severe rejection episodes. Despite organ transplants such as pancreas or multiple organs, the liver can evidence. but at the same time allows the patient to have insulin independence. This approach not only corrects liver disease but at the same time allows the patient to have insulin independence.

Certain liver diseases have strong association with diabetes mellitus; P.S.C is one of the diseases which has been described in association with diabetes mellitus type 1. Other diseases of the liver which have direct association with diabetes mellitus are NASH, and cystic fibrosis. In selected cases, these diseases justify combined liver-pancreas transplantation.

As well as those previously mentioned benefits of combined liver-pancreas transplantation, it may have an immunologic advantage. Unlike liver, pancreas is considered a highly immunogenic organ and when liver transplant combines with other organ transplants such as pancreas or multiple organs, the liver can protect these organs from severe rejection episodes. Despite the advantages of combined liver-pancreas transplantation, only few centers have reported this kind of transplantation and only in limited numbers.

In conclusion, en-bloc liver-pancreas transplantation may have some advantages over separate implantation of both organs in patients having IDDM and end-stage liver disease, including better long-term survival and better quality of life after transplantation. Moreover, combined transplant can prevent more serious rejection episodes due to the immunotolerant effect of liver allograft.

References