Portal Vein Thrombosis Complicating Islet Transplantation in a Recipient with the Factor V Leiden Mutation

A 31-year-old Native American/white female with Type 1 diabetes mellitus of 26 years duration enrolled in the international multicenter trial of the Immune Tolerance Network (NS01) to replicate the Edmonton Protocol for islet transplantation. She used oral contraceptives. She had undergone two prior islet transplant procedures complicated by sirolimus induced mucositis, diarrhea, and anemia, treated with replacement of sirolimus with mycophenolate mofetil. She received a third transplant of 314,039 islets (4,906 islet equivalents per kg) with a purity of 30% in 9 ml of packed-cell volume in two aliquots through a 5-Fr cholangiogram catheter in the right branch of the portal vein. Each aliquot contained heparin, 35 u/kg of patient body weight. After one hour, she developed abdominal pain, hypotension, and anemia. Enoxaparin was withheld. An abdominal ultrasound on postprocedure day 1 revealed a perihepatic fluid collection consistent with blood and patent hepatic vessels without evidence of thrombosis. She received 4 units of packed red blood cells over 48 hours, and her hemoglobin stabilized.

An abdominal ultrasound on postprocedure day 7 demonstrated a nonocclusive thrombus in the right portal vein, 30 mm in length (Fig. 1). She was anticoagulated with heparin and then warfarin. Serial ultrasounds revealed resolution of the thrombosis by 12 weeks after the event. A thrombophilia evaluation for antithrombin III, protein C and S, anticardiolipin antibodies, homocysteine, prothrombin gene mutation, and Factor V Leiden (FVL) revealed heterozygosity for the FVL mutation, and negative or normal otherwise. Subsequent genetic analysis of her mother revealed heterozygosity for FVL.

Portal vein thrombosis after islet allograft transplantation has been reported previously (1–3). The perceived risks include transplantation of large packed cell volumes of partially purified islets (1), technical difficulties including intraportal injection of Gelfoam (2), and repeated transplant procedures (3). Thrombophilia has not been previously identified as a risk factor.

Our subject had several identified risks for development of portal vein thrombosis: infusion of larger than average packed cell volume (although within protocol guidelines), third transplant, anticoagulation discontinued after infusion of the islets because of bleeding complications, and use of oral contraceptives. Gelfoam was not used with any procedure.

Approximately 5% of whites are heterozygous for the FVL mutation, the most common cause of hereditary thrombophilia (4). For individuals presenting with venous thrombosis, FVL occurs in 11–20% of those in all age groups and 50% of individuals under 50 years of age (5). Heterozygosity for this mutation produces a 7-fold relative risk of venous thrombosis. Our patient used oral contraceptives. The risk of venous thrombosis with both oral contraceptive use and the Factor V Leiden mutation is increased more than 30-fold compared with neither risk factor (6).

Given the relatively high prevalence of this mutation in the general population (5%), we recommend thrombophilia screening for potential islet transplant recipients. If transplantation is performed in such individuals, the procedure should be optimized by using high purity isolations, limiting the packed cell volume to less than 3 ml, limiting bleeding, avoiding oral contraceptives, and using prolonged anticoagulation prophylaxis.

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FIGURE 1. Hepatic ultrasound of the portal vein showing a 30 mm thrombus.
REFERENCES


