CASE REPORT

Total Pancreatectomy and Islet Auto-Transplantation as Treatment for Ampullary Adenocarcinoma in the Setting of Pancreatic Ductal Disruption Secondary to Acute Necrotizing Pancreatitis. A Case Report

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ABSTRACT

Context Ampullary adenocarcinoma is the third most common periampullary malignancy. Obstruction of the main pancreatic duct is linked with an increased incidence of acute pancreatitis. Acute necrotizing pancreatitis leading to pancreatic duct disruption carries significant morbidity. When these conditions occur in combination, the treatment can be drastically limited as pancreatectoduodenectomy is not a viable option in the setting of friable ductal tissue, which precludes pancreatic ductal anastomosis and can lead to the complications of leak or stricture. Case report Our patient is a 72-year-old woman who developed pancreatic ductal disruption and splenic vein thrombosis as a result of acute necrotizing pancreatitis. Concurrently, she was found to have an ampullary adenoma with high-grade dysplasia. Her treatment options were limited, as she was neither a candidate for pancreatectoduodenectomy given the ductal disruption nor total pancreatectomy, which would render her a brittle diabetic. She was successfully treated with total pancreatectomy and islet auto-transplantation thereby resecting her ampullary lesion while both avoiding a pancreatic anastomosis and preserving pancreatic endocrine beta-cell function. Conclusion We report a case where total pancreatectomy and islet auto-transplantation can be considered as a viable option for treatment of ampullary lesions in a setting where standard surgical options are suboptimal.

INTRODUCTION

Ampullary adenocarcinoma is the third most common periampullary malignancy, with an incidence of 6 new cases per million persons per year [1]. Peak incidence occurs in the sixth or seventh decade of life with a slightly greater propensity among males. Patients typically present with waxing and waning symptoms of biliary obstruction. Due to lower pathologic aggressiveness in the initial stages, the tumors are resected early on in the course of the disease, thereby improving 5-year survival (up to 55%) when compared to pancreatic ductal adenocarcinoma, for example, which has a markedly lower 5-year survival. Most benign ampullary growths can be treated with endoscopic resection or with open surgical resection. Invasive tumors have been historically treated with pancreatectoduodenectomy.

CASE REPORT

Our patient is a 72-year-old woman who initially presented to an outside hospital in January 2011 with severe epigastric abdominal pain and near syncopal episode. Initial diagnostic tests revealed elevated amylase, lipase and transaminase levels. An abdominal computed tomography (CT) scan showed a diffusely edematous pancreas with peripancreatic inflammation. The patient then underwent magnetic retrograde cholangiopancreatography (MRCP), which indicated an intraluminal filling defect in the conjoined portion of the common bile duct and pancreatic duct near the ampulla suspicious for stone, clot or tumor (Figures 1 and 2), in addition to the marked pancreatic edema and peripancreatic inflammation (Figure 3). The patient then underwent endoscopic retrograde cholangiopancreatography (ERCP), which confirmed a lesion in the ampulla of Vater. This lesion was biopsied and thereby only partially resected. Additionally, a pancreatic stone was discovered and removed. The patient received biliary sphincterotomy and stent placement. The final pathology of the biopsy specimen revealed duodenal papillary adenoma with high-grade dysplasia. The patient convalesced from this acute
episode and was discharged with planned follow-up to discuss the treatment strategy for the ampullary lesion. Unfortunately, our patient went on to have quite a complicated course after the index admission. She was readmitted several times with recurrent symptoms of abdominal pain and vomiting over the next several months. She underwent drainage of a peripancreatic fluid collection that was felt to be an abscess and eventually developed necrosis of the mid portion of the pancreas, which resulted in ductal disruption. In addition, she developed thrombosis of the splenic vein. She underwent ERCP and stent placement across the disrupted duct (Figure 4) and was managed over this time period with total parenteral nutrition and bowel rest.

Pancreaticoduodenectomy was considered for management of her papillary adenoma with high-grade dysplasia; however, there was significant concern for performing a pancreatic anastomosis in the setting of recent acute necrotizing pancreatitis. Total pancreatectomy was also considered but rendering the patient a brittle diabetic was felt to be prohibitive. Alternatively, total pancreatectomy with islet auto-transplantation was presented as an option that would adequately treat the ampullary adenoma, avoid an anastomosis in a high-risk field while still preserving islet cell function. As total pancreatectomy with islet auto-transplantation also includes splenectomy, the splenic vein thrombosis would be adequately treated as well. The patient was referred to our institution given our extensive experience with this procedure.

The patient underwent a thorough multi-disciplinary evaluation to optimize her for surgery by endocrinology, internal medicine, gastroenterology and general surgery. All laboratory levels, including baseline insulin levels and liver function tests were well within normal limits. Her nutritional status was satisfactory; as she had returned to an oral diet with

Figure 1. Coronal MRCP (T2 HASTE sequence, thin slice) image demonstrating intraluminal filling defect due to ampullary lesion (white arrow).

Figure 2. Axial MRCP (T2 HASTE sequence, thin slice) image demonstrating intraluminal filling defect due to ampullary lesion (white arrow).

Figure 3. Axial MRCP image demonstrating significant pancreatic edema and peripancreatic inflammation (white arrows).

Figure 4. Coronal CT image demonstrating pancreatic ductal stent (black arrow).
improved appetite while still being maintained on nocturnal total parenteral nutrition. She also received appropriate vaccinations in anticipation of splenectomy two weeks prior to surgery. She successfully underwent total pancreatectomy with islet auto-transplantation in June 2011, which she tolerated well. She received 1,326 islet equivalents per kilogram of body weight. Post-operatively, she required a small dose of daily insulin with meal correction. She was discharged from the hospital two weeks later on supplemental enteral feeds, which were eventually discontinued. Unfortunately, her final pathology indicated a 0.3 cm invasive adenocarcinoma of the ampulla within a background of extensive adenoma with high-grade dysplasia. The margins were widely negative with a final pathologic stage of pT1N0MX G1. On a 3-month follow-up visit, she demonstrated steady and gradual improvement with improved appetite, weight gain and required only moderate serum glucose correction.

DISCUSSION

We have presented a patient with a complex problem that served as a rare indication for total pancreatectomy and islet auto-transplantation. The pancreatic duct disruption secondary to necrotizing pancreatitis had been temporized by ERCP and stent placement; however, the diagnosis of papillary adenoma dictated that appropriate intervention be performed in a timely fashion. Not only did the papillary adenoma represent the likely etiology of her acute necrotizing pancreatitis [2, 3, 4] but it also represented a pre-malignant lesion that, with the presence of high-grade dysplasia, may already harbor invasive carcinoma, as the final pathology report indeed revealed.

The treatment options for this patient were limited and associated with significant pitfalls. Pancreateico-duodenectomy, the standard of care for invasive cancer, would have effectively treated the lesion. However, the reconstruction would have been undermined by the pancreatic necrosis present at the usual pancreateicojenunostomy site as a result of the patient’s recent episode of acute necrotizing pancreatitis. A pancreatic anastomosis in this setting would have had a markedly elevated risk of leakage and subsequent morbidity. Total pancreatectomy, on the other hand, also effectively treats the lesion and yields the benefit of avoiding a pancreatic anastomosis. However, this option would have left the patient a brittle diabetic with its resultant morbidity. Another option for benign lesions that has gained some recent popularity is endoscopic resection. However, this option in the setting of high-grade dysplasia is associated with a high risk of recurrence and is suboptimal treatment for potential coexistent invasive adenocarcinoma that can be found in up to 50% of adenomas with high-grade dysplasia [5]. In addition, it is associated with a risk of post-procedural pancreatitis [6] that may have had devastating consequences in this patient.

Total pancreatectomy with islet auto-transplantation afforded an opportunity to adequately treat the lesion with pancreatic resection while also preserving beta-cell function and minimizing or avoiding brittle diabetes. Initially performed in 1977 at the University of Minnesota [7], total pancreatectomy with islet auto-transplantation is being used with increasing frequency for treatment of chronic pancreatitis [8]. Total pancreatectomy with islet auto-transplantation has been successful in preventing brittle diabetes in the majority of patients undergoing this procedure with up to 71% of patients in the recent era (2001-2007, n=106) demonstrating either full islet function (i.e. insulin independence, 33%) or partial islet function (euglycemia on once daily long-acting insulin, 32%) at one year post-transplant. Moreover, once islet function (full or partial) was established, it persisted, with rates of 85%, 68% and 57% at 2, 5 and 10 years, respectively [9].

While this procedure is performed almost exclusively for chronic pancreatitis, other indications have been reported. Liu et al., reported a case of pancreaticoduodenectomy for pancreatic adenocarcinoma of the head of the pancreas. The procedure was complicated by life-threatening leakage of the pancreaticojejunostomy that was salvaged with completion pancreatectomy and islet auto-transplantation [10]. This patient remained dependent on insulin (16-18 U/day) at one-year follow-up. Unfortunately, he died of local tumor recurrence 2.5 years after transplant without radiographic evidence of tumor metastases in the liver [11]. Similarly, Alsaif et al. published a case of total pancreatectomy with islet auto-transplantation performed for management of uncontrolled pancreatic fistula complicating pancreaticoduodenectomy [12]. This patient was insulin independent at discharge and remained so without evidence of tumor recurrence at one-year follow-up.

Of obvious concern is the potential to inoculate the liver with malignant cells during the portal vein infusion of islet cells. This theoretical risk is not only concerning for the cases above where total pancreatectomy with islet auto-transplantation was performed in the setting of frank pancreatic adenocarcinoma, but also in patients with chronic pancreatitis who are at increased risk for development of malignancy. To date, there are no published reports of liver metastases occurring after total pancreatectomy with islet auto-transplantation. Indeed, even the patient noted above who died of local recurrence of pancreatic adenocarcinoma had no evidence of liver metastases. Lee et al. published their data on total pancreatectomy with islet auto-transplantation in Korea for indications other than chronic pancreatitis [13]. The majority of the 10-patient cohort underwent pancreatic resection for neoplasia: serous cystic adenomas (n=4), pseudo-papillary neoplasms (n=3) and intraductal papillary mucinous neoplasms (IPMN, n=2), which are thought to be both precursor of and markers of malignancy. For
the patients with IPMN, preoperative evaluation as well as intraoperative frozen section confirmed the disease to be focal (as opposed to multicentric) and confined to the body of the pancreas (negative ductal margin). Both lesions had moderate dysplasia and a background of adenoma. At 1-year follow-up, there was no evidence of recurrence or metastatic disease in the liver. In fact, none of their 10 patients had evidence of liver disease following total pancreatectomy with islet auto-transplantation.

The paucity of data regarding this issue makes it extremely difficult to determine the true risk of malignancy. Nonetheless, it remains of serious concern and should certainly be considered when considering total pancreatectomy with islet auto-transplantation in the setting of a neoplastic process. If performed for this indication, the patient should be thoroughly counseled regarding this risk and should receive close follow-up for detection of this complication.

CONCLUSION

We have presented a case of total pancreatectomy and auto-islet transplantation for treatment of ampullary adenocarcinoma in the setting of recent acute necrotizing pancreatitis with mid pancreatic ductal disruption and splenic vein thrombosis. Total pancreatectomy with islet auto-transplantation should be considered if resources are available and standard therapeutic options are suboptimal.

Disclosures None

Conflicts of interest None

References