Pancreatic Panniculitis: A Rare Manifestation of Acute Pancreatitis

Ronak Patel, Ali Safdar Khan, Sami Naveed, Jason Brazleton, Mel Wilcox

University of Alabama at Birmingham, USA

ABSTRACT

Context Pancreatic panniculitis is a very rare complication associated with pancreatic disease and perhaps even a presage to pancreatic pathology. Case report We present a case of pancreatic panniculitis in a 61 year old patient who was treated for sudden onset of abdominal pain associated with nausea and vomiting secondary to acute pancreatitis of unknown etiology. He subsequently developed skin lesions consistent with pancreatic panniculitis which gradually improved after resolution of his acute condition and treatment with topical steroid cream. Conclusion We discuss and review the literature along with highlighting for the readers the important clinical and histopathologic features of acute pancreatitis associated pancreatic panniculitis.

INTRODUCTION

Pancreatic panniculitis (PP), also known as pancreatic fat necrosis and enzymatic panniculitis is a rare, lobular panniculitis that infrequently develops in individuals with pancreatic disease [1]. It is an uncommon manifestation of pancreatic disease associated in 2-3% of pancreatic pathology [2]. The first reports in the English literature are from 1947 [3] although the oldest reports go back as far as 1883 [4]. It is a rare entity with fewer than 150 cases reported in the literature [5]. Due to the rarity of PP, the diagnosis can be missed in its early stages often leading to delay in treatment in certain instances. Its pathogenesis is still not well understood. PP typically presents with tender, edematous and erythematous subcutaneous nodules that spontaneously ulcerate and exude an oily brown substance that results from liquefaction necrosis of adipocytes. The lesions most commonly develop on the lower legs, though other sites like thighs, buttocks, arms, abdomen, chest and scalp have also been reported [6].

CASE REPORT

We present a 61-year-old patient who presented to an outside hospital with sudden onset of acute abdominal pain, nausea, and vomiting lasting for two days. Subsequently the patient was admitted with a diagnosis of acute pancreatitis of unknown etiology. At the time of admission laboratory evaluation revealed serum lipase to be significantly elevated at 5367 units/L. CT scan of the abdomen was consistent with the diagnosis of acute pancreatitis as shown in Figure 1.

Patient returned to the hospital with a relapse of his original symptoms and a fever of 101 degree Fahrenheit three weeks later. Repeat imaging studies showed pancreatic fluid collection (5x3 cm) in the head of the pancreas along with what he reported to be multiple ’spider bites’ on his legs (Figure 2). He was placed on intravenous fluids, antibiotics (topical as well as Intravenous) and analgesics but due to the continuation of symptoms (abdominal pain, nausea) he was transferred to a tertiary care hospital for further management of severe pancreatitis and cellulitis.

Dermatologic exam at that time revealed several scattered, tender erythematous subcutaneous nodules in bilateral lower extremities without ulceration. A punch biopsy of the lesions was performed which showed lobular panniculitis with prominent fat saponification and calcification necrosis. Scattered neutrophils and eosinophils were noted amongst the infiltrate (Figure 3).

An EUS showed hyperechoic foci in the body & tail of the pancreas while the head of pancreas was replaced by a mixed echogenic lesion with anechoic, hyper and hypoechoic areas. Fine needle aspiration revealed amorphous debris with calcification. No malignant cells were seen. He was discharged after four days of conservative inpatient treatment.

Figure 1. CT scan of the abdomen was consistent with the diagnosis of acute pancreatitis.
management and was placed on prednisone for pancreatic panniculitis. He was unable to tolerate the Prednisone or Naproxen due to their side effects and was subsequently placed on topical 1% betamethasone ointment (BID). On recent follow up the lesions were seen to be almost completely resolved (Figure 4) and a repeat EUS showed a 6 cm pancreatic pseudocyst which was asymptomatic.

**DISCUSSION**

Pancreatic panniculitis is a rare complication in the setting of pancreatic disease, in which fat necrosis takes place in the subcutaneous tissue and elsewhere [7]. The exact pathophysiology is becoming clear but not entirely understood. The release of pancreatic enzymes, such as lipase, trypsin, amylase, phosphorilase and trypsin, may be involved in increasing the micro vascular permeability allowing hydrolysis of neutral fat. The resultant glycerol and free fatty acids, leads to fat necrosis and a brisk inflammatory response. The pathogenic role of pancreatic lipase is supported by the finding of the enzyme in the areas of subcutaneous necrosis, and also anti-lipase monoclonal antibodies within the necrotic tissue [8]. Pancreatic disorders are much more common when compared to pancreatic panniculitis which might indicate that there are still unrecognized factors involved in the development of PP. Also there are some PP cases in the literature that have been described with normal levels of pancreatic enzymes (Table 1) further supporting the likely involvement of other unidentified factors in the causality of this entity. An immune complex mechanism has been described in one patient with pancreatic panniculitis [9].

The dermatologic manifestations are independent of the severity of pancreatic pathology [10] and precede the diagnosis of pancreatic pathology about 40% of the times. The lesions are generally located in the lower extremities as tender, erythematous nodules that are red-brown in color, which may ulcerate with drainage of oily fluid [11, 12]. The clinical features of pancreatic panniculitis are similar despite the variety of pancreatic disorders it is associated with. It is important to differentiate PP from other forms of panniculitis such as Erythema nodosum, erythema induratum, lupus panniculitis, infectious, traumatic or alpha 1- antitrypsin deficiency panniculitis [13].

PP is usually confirmed with a subcutaneous biopsy and if the characteristic features of this entity are identified earlier on, an invasive biopsy could be averted [14]. The main histopathologic feature is a mostly lobular panniculitis without vasculitis. But, in the very early stage, a septal pattern has been described, which results from enzymatic damage of the endothelial septa, allowing pancreatic enzymes to cross from blood to fat lobules resulting in coagulative necrosis of the adipocytes, which leads to pathognomonic "ghost cells" (enucleated necrotic cells that have a thick wall with a fine basophilic granular material within their cytoplasm from dystrophic calcification) [15].

Specific subgroups have known to have a higher association (alcoholic men [1], patients with acinar neoplasms [16-19]), and PP may be the harbinger of pancreatic pathology by several days [15] or even months.

The treatment of PP is mainly directed towards the underlying pancreatic pathology, following which a
complete to near complete resolution of symptoms is experienced as illustrated in our report. Corticosteroids may provide some symptomatic relief however NSAID’s or immuno suppressors are usually not effective for the treatment of skin lesions [10]. As more cases of PP are recognized and better characterized, the underlying mechanisms may also be better understood perhaps leading to better vigilance to its incidence, causes and treatment.

**Conflicting Interest**

The authors had no conflicts of interest

**References**


