Case Report

An unusual presentation of pancreatic serous cystadenoma

Alexandra O. Stathis*, Samuel C. Kuo

Department of Surgery, Hornsby Kur-ring-gai Hospital, Palmerston Road Hornsby, NSW 2077, Australia

Received: 24 February 2020
Revised: 03 March 2020
Accepted: 05 March 2020

*Correspondence:
Dr. Alexandra O Stathis,
E-mail: Alexandra.stathis@health.nsw.gov.au

Copyright: © the author(s), publisher and licensee Medip Academy. This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ABSTRACT

Pancreatic cystic neoplasms (PCNs) are predominantly benign entities which represent almost 50 percent of all cystic lesions of the pancreas. PCNs are often an incidental finding on abdominal imaging and are not indicated for surgical resection unless they show evidence of malignant transformation or become symptomatic due to mass effect. This report examines an unusual presentation of a PCN, in a 70 years old female with sudden onset abdominal pain, who was found to have spontaneous intraabdominal haemorrhage secondary to a benign PCN. Emergency laparotomy was performed and a distal pancreatectomy or splenectomy were required to achieve haemostasis. Incidence of spontaneous haemorrhage in a benign PCN is a rare but serious complication.

Keywords: Pancreatic cystadenoma, Pancreatic tumour, Abdominal surgery

INTRODUCTION

Pancreatic cystic neoplasms are predominantly benign entities which represent almost 50 percent of all cystic lesions of the pancreas.1,2 PCNs are often an incidental finding on abdominal imaging, and patients are usually asymptomatic. Indeed, benign PCNs are not indicated for resection unless they show evidence of malignant transformation or become symptomatic, usually due to a slowly evolving mass effect.3,4

Here we outline the unusual case of a female in her 70s who presented with massive intraabdominal haemorrhage resulting from the rupture of a previously undiagnosed benign pancreatic serous cystadenoma.

CASE REPORT

A female in her 70’s presented to an outer metropolitan emergency department with sudden onset, severe abdominal pain. The patient had a pertinent history of paroxysmal atrial fibrillation, mitral valve replacement and previous open appendicectomy. Medications included warfarin, ivabradine and flecainide.

On arrival to the ED, the patient was found to be hypotensive with a blood pressure of 50/38 mmHg which improved to 130/60 mmHg after a 1500 ml fluid bolus. In the context of flecainide, there was no cardiac compensation and her heart rate remained steady at 65bpm. The patient’s abdomen was distended and guarded though not rigid. Initial bloods showed a haemoglobin of 74 and INR of 2.1.

Mobile chest X-ray demonstrated no free gas, whilst bedside focused assessment with sonography for trauma scan showed significant free fluid in bilateral upper quadrants and pelvis.

A CT mesenteric angiogram demonstrated a partly necrotic lesion involving the distal body and tail of pancreas with an extension to the splenic hilum (Figure 1 and 2). CT evidence of substantial free fluid within the...
abdomen further raised concern of haemorrhage. Emergency laparotomy was immediately pursued.

Intraoperative findings were that of a large multi cystic tumour involving the body and tail of pancreas, densely adherent to the splenic hilum, left kidney and mesocolic vessels. A distal pancreatectomy and splenectomy were undertaken and haemostasis achieved. Postoperatively the patient was admitted to the intensive care unit where she remained intubated for 3 days, required four units of packed red blood cells and inotropic support. Following an extended admission for a pancreatic leak, the patient was discharged home on day 17.

Histology confirmed a serous cystadenoma without malignant features. On recent review, three years post, there was no evidence of recurrence.

**DISCUSSION**

Pancreatic cystic neoplasms are true cystic lesions of the pancreas that exist in an estimated 2–45% of the general population and account for nearly 50% of all pancreatic cysts.\(^1\)\(^2\)

The increasing use of cross-sectional imaging has seen a rise in the incidental diagnosis of PCNs in patients undergoing CT or MR imaging of the abdomen for unrelated indications.

These incidental lesions are largely benign, though malignant potential does exist in some subtypes. PCNs are grouped into four subtypes according to the world health organisation histologic classification.\(^3\)\(^4\) They are: serous cystic neoplasms (SCN), mucinous cystic neoplasms, intraductal papillary mucinous neoplasms (IPMNs): main-duct or branch-duct, and solid pseudo papillary neoplasms.

Age of presentation for PCNs is typically between the 5th and 7th decade, with the exception of the solid pseudo papillary subtype which presents earlier, between the 2\(^{nd}\) and 3\(^{rd}\) decade of life. Intraductal mucinous neoplasms occur equally in men and women, whilst all other subtypes have a female predominance.

Diagnosis of new PCNs is almost always incidental. Once suspected on initial imaging, of which MRI is superior, tissue diagnoses may be sought. Endoscopic ultrasound with fine needle aspiration is the method of choice.\(^5\)\(^6\)

A patient with a pancreatic serous cystadenoma is most likely to be asymptomatic. Symptoms, if present, usually manifest secondary to obstruction or compression of surrounding structures.

Haemorrhagic complication of PCNs is very rare, with only a handful of cases reported in the literature.\(^7\)\(^9\)

Pancreatic serous cystic neoplasms are a wholly benign entity. Early studies suggested that serous cyst adenomas had low malignant potential however there is a growing body of evidence to support that serous cystadenomas are indeed the only PCNs that are benign.\(^10\)\(^11\) It is thought that early previous reports of malignant lesions did not truly satisfy the WHO definition of serous cyst adenomas.\(^12\)\(^13\)

The 2018 European evidence-based guidelines on pancreatic cystic neoplasms recommend just one year of follow-up for newly diagnosed serous cystadenomas.\(^4\) The rationale for this advice is the evidence that confirmed serous cystadenomas are benign and need only be operatively managed if causing compressive symptoms to adjacent organs. The rate of growth of these lesions is considered so slow that new onset of symptoms is exceedingly rare. Indeed, approximately 60% of SCNs

![Figure 1: Serous cystadenoma of the distal pancreas.](image1)

![Figure 2: Serous cystadenoma of the distal pancreas.](image2)
remain of stable size. There is evidence to support that SCNIs of the pancreatic head tend to exhibit more aggressive behaviour including faster rate of growth.  

CONCLUSION

PCNs are common. Most are serous cystadenomas, which are benign. Serous cystadenomas may require elective surgery for mass effect. We describe a rare case where emergency surgery was required. Incidence of spontaneous haemorrhage in a benign PCN is a rare but serious complication.

ACKNOWLEDGEMENTS

The abovenamed authors received no external funding for the preparation of this report, and have nothing to declare.

Funding: No funding sources
Conflict of interest: None declared
Ethical approval: Not required

REFERENCES


Cite this article as: Stathis AO, Kuo SC. An unusual presentation of pancreatic serous cystadenoma. Int Surg J 2020;7:1280-2.