**Case Report**

An 8-year-old boy was admitted to BMH Rinteln with severe left upper quadrant pain. He gave a history of having been kicked in the abdomen by his brother during play 3 weeks previously. The pain had become worse and he then developed anorexia, listlessness and had lost 3 kg in weight. On examination he was pale and had an obvious tender diffuse mass involving the epigastrium and left upper quadrants. Investigations: ESR = 48 in first hour, Hb = 13.2 gram per cent, WBC = 6.0x10⁹/L. An IVU was normal but ultrasound scan showed a transonic collection anterior to the superior pole of the left kidney consistent with the clinical diagnosis of a subcapsular splenic haematoma.

For 5 days his symptoms improved only to deteriorate over the subsequent 2 weeks with anorexia and vomiting. The blood profile was normal except an ESR = 32 and a polymorph leucocytosis of 83%. Repeat ultrasound scan showed one large cyst and several smaller cysts in the splenic region. Barium meal suggested there was a mass behind the stomach displacing it forward. At this stage the plasma amylase = 3089 International units.

Laparotomy confirmed the presence of a large lesser sac cyst (pseudocyst) with multiple smaller cysts of the pancreas each 2-3 cms in diameter. The smaller cysts of the pancreas were marsupialised and the lesser sac was drained through a wide bore tube drain to the exterior through a separate stab incision.

Postoperatively he made slow but steady progress. An external fistula formed in the laparotomy wound 2 weeks following operation but resolved over 3 weeks. Ultrasound scan at this stage showed resolution of the cysts, Hb = 12.2 gm%, WBC = 5.0x10⁹/L and amylase = 257 Somogyi units. The patient has made a complete recovery.

**Discussion**

Pseudocyst of the pancreas is uncommon in childhood; only 113 cases are reported in persons under 18 years of age up to 1981.¹ The mean age of the reported cases is 7.5 years with an age range of 2 months to 17 years.² Boys appear to be affected twice as often as girls and trauma is the initiating cause in well over 50% of cases.³ Kicks, bicycle handlebars and child abuse are the most common forms of trauma implicated. In the remainder, 4% of cases follow mumps and acute pancreatitis, a small percentage are due to gastric duplication, hereditary pancreatitis and malnutrition, but 30% of all cases are of unknown aetiology.⁴

Symptoms most frequently complained of are abdominal pain or a feeling of fullness, nausea and vomiting, anorexia and weight loss. On examination an abdominal mass is present in 64% of cases and 80% of patients have an elevated serum amylase.² Ultrasound scanning is now recognised as the most effective method for diagnosis; cysts as small as 3 cms may be identified. Barium meal may identify a large lesser sac collection.

Management of this rare condition in children has, until recently, been conducted on adult pseudocyst management policies. In adults 50% of pseudocysts may regress spontaneously;⁵ however in children the pancreatitis and cyst formation causes such systemic upset that early surgical intervention is the best method of treatment. The surgical procedure most commonly used is that of internal drainage by cystogastrostomy or less usually by Roux-en Y cystojejunostomy. Cystogastrostomy is technically easy and has the lowest rate of recurrence and operative mortality. External drainage or marsupialisation using firm rubber or silicone drains has been advocated⁶ and was the method used in this case. The argument against this therapy is its high recurrence rate (25%)⁷ and cutaneous fistula formation with the associated electrolyte loss, skin excoriation and enzyme loss. However, in the series reported by Pokorny et al, the fistula resolved spontaneously as occurred in this case.

Recently external drainage in the form of percutaneous fine needle aspiration using ultrasound as a guide has achieved excellent results.⁸ Recurrence of the cyst formation has been lessened by a recent modification whereby the 16 gauge Teflon intravenous catheter used for percutaneous aspiration is left in place to allow complete drainage. There have been no complications and the technique is well tolerated. Complete excision of the cyst wall is not well formed in this age group and dissection is difficult. Non-operative treatment of a pseudocyst is...
reserved for those children whose symptoms are improving in association with resolution of the cyst on ultrasound scan.

Conclusion

Pancreatic pseudocyst in children is uncommon. The case reported highlights the presentation and management with the excellent outlook for cure. The recurrence rate is low after internal drainage, but recently percutaneous fine needle aspiration has revealed equally good results. The disease pathogenesis seems to be non-progressive in children when compared with the adult form due to lack of pancreatic insult from alcohol ingestion or secondary to biliary tract pathology.

Acknowledgement

I wish to thank Colonel W G Johnston for his help and encouragement in writing this article about his patient.

REFERENCES

Pseudocyst of the Pancreas in Childhood

C L Griffiths

*J R Army Med Corps* 1985 131: 70-71
doi: 10.1136/jramc-131-02-04

Updated information and services can be found at:
http://jramc.bmj.com/content/131/2/70.citation

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/