Recurrent pancreatoblastoma with inappropriate adrenocorticotropic hormone secretion

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SUMMARY We describe a boy with recurrent pancreatoblastoma who developed Cushing's syndrome due to inappropriate adrenocorticotropic hormone secretion.

Primary epithelial tumours of the pancreas are rare in childhood. Although islet cell tumours have previously been shown to produce inappropriate secretion of peptide hormones, we believe that this is the first report of a pancreatoblastoma producing adrenocorticotropic hormone and an ectopic hormone syndrome.

Case report
A 22 month old boy presented with a firm, non-
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tender, asymptomatic abdominal mass of 12 cm diameter. He was normotensive with a blood pressure of 112/75 mm Hg, normokalaemic with a potassium concentration of 3-6 mmol/l, had an appreciably raised α fetoprotein concentration of 3670 μg/l (normal 2 μg/l), and a normal serum amylase of <300 IU/l (Phadebas method). An ultrasound scan showed a mass of mixed echogenicity crossing the midline and separate from the kidneys. Plain abdominal radiography showed the mass to be focally calcified.

At operation the tumour was found to be arising from the pancreas, and a subtotal pancreatectomy was performed. Histological examination showed a malignant epithelial tumour with areas showing acinar differentiation, areas of poorly differentiated cells with frequent mitoses, and squamoid corporules. Immunoperoxidase methods showed strong staining for α1 antitrypsin and α fetoprotein, but were negative for adrenocorticotropic hormone, neurone specific enolase, and chromogranin. Electron microscopy confirmed acinar differentiation; no neurosecretory granules were seen. The appearance was of a pleomorphic pancreatoblastoma. Tumour was present at the line of resection indicating incomplete removal.

There were severe postoperative hypokalaemia (potassium concentration 2-1 mmol/l) despite intravenous supplements. The blood glucose remained stable. Treatment was commenced with actinomycin D and vincristine in cycles of three weeks, and the serum α fetoprotein concentration fell to 18 μg/l. After three months the concentration of serum α fetoprotein rose; computed tomography and a radionucleide scan using labelled antibody to α fetoprotein showed a tumour deposit in the posterior mediastinum. The chemotherapy was therefore changed to cisplatin/etoposide, ifosfamide/etoposide, and doxorubicin in cycles of three weeks. The serum α fetoprotein then fell to normal concentrations, and treatment was continued for one year. He remained well off treatment for 18 months.

Three and a half years from first presentation he developed increasing fatigue, weight gain, and pigmentation. On examination he had a Cushingoid appearance, a palpable abdominal mass, and his blood pressure was 130/80 mm Hg. He was hypokalaemic (potassium concentration 2-5 mmol/l), and had a raised serum α fetoprotein concentration (590 μg/l). The plasma adrenocorticotropic hormone concentration was 56 pmol/l (normal 2–18 pmol/l) and the urinary cortisol 2000 nmol/l (normal 55–740 nmol/l). A 24 hour serum cortisol profile showed loss of diurnal variation. Abdominal ultrasound examination confirmed the mass arising from the pancreatic remnant. Chest radiographs showed several metastases. A computed tomogram of brain showed no abnormality of the pituitary fossa and this together with the appreciably raised serum adrenocorticotropic hormone, skin pigmentation, hypokalaemia, and lack of response to either dexamethasone or metyrapone, confirmed the diagnosis of inappropriate secretion of adrenocorticotropic hormone. This was recurrent and metastatic pancreatoblastoma, as shown by the recurrent pancreatic mass seen on ultrasound examination and associated with a raised serum α fetoprotein concentration, although histological confirmation was not attempted.

Treatment was resumed with a combination of vinblastine, carboplatin, etoposide, and bleomycin. Unfortunately the tumour showed no response to two courses of chemotherapy, which was therefore withdrawn. The child died three and a half years after diagnosis.

Discussion

Primary epithelial tumours of the pancreas are rare in childhood, and have a biphasic age distribution. Pancreatoblastoma like other blastomas of childhood affects children in the early years of life, has a slight male predominance,1 and is sometimes associated with Beckwith-Wiedemann syndrome.2 There have been two previous reports of pancreatoblastoma producing α fetoprotein.3 Another tumour, unique to late childhood and early adult life, affects females almost exclusively, and has been given various names including ductuloacinar adenoma and papillary-cystic tumour of the pancreas. It usually follows a benign course.

Pancreatoblastoma can arise in any part of the pancreas and has distinct histological features including undifferentiated areas, acinar differentiation, squamoid corporules, and the absence (by ordinary light microscopy) of islet cells. Originally thought to arise only from the ventral pancreas and therefore not to contain endocrine cells, it is now known that this tumour may arise in the distal (and therefore from dorsal) pancreas. The latter type seems to have a more insidious clinical onset, is non-encapsulated, and is associated with a worse prognosis.

There have been several reports of islet cell tumours producing hormones such as adrenocorticotropic hormone, gastrin, glucagon, and growth hormone releasing hormone,4 but non-islet cell pancreatic tumours are not associated with inappropriate secretion of peptide hormones. Insulin, gastrin, and somatostatin have been shown by immunohistochemistry in pancreatoblastoma,5 6 but we believe that this is the first case of pancreatoblas-
Intestinal disease in cystic fibrosis

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SUMMARY Three children with cystic fibrosis developed steatorrhoea unresponsive to changes in pancreatic supplements. The final diagnoses were chronic giardiasis, stagnant loop syndrome, and Crohn's disease. Refractory intestinal symptoms in cystic fibrosis merit further investigation.

It is still controversial whether the gastrointestinal manifestations of cystic fibrosis are primary or secondary,1 although recent evidence shows that the basic defect in chloride transport is expressed in the intestinal epithelium.2 Steatorrhoea, flatulence, abdominal pain, distal intestinal obstruction syndrome, and rectal prolapse are usually improved by adequate pancreatic supplementation, particularly with microsphere preparations.1 3 Occasionally, however, high doses of these supplements, with or without H2 antagonists, are ineffective. This can be caused by incorrect use of the supplements, or poor compliance, but coeliac disease and cows' milk protein intolerance have also been described in association with cystic fibrosis.1 4 Investigating three children who presented in this way showed further treatable intestinal pathology.

Case reports

Case 1
A 16 year old girl in early puberty complained of varying bowel habit over several months, passing up to five loose stools a day, often containing 'blobs of grease'. She also experienced rectal incontinence with laughter, and frequent abdominal pain after food. Her weight gain was poor, but this could not be explained by her chest disease. Stool microscopy showed fat globules but no parasites. At first poor compliance was suspected. A jejunal biopsy specimen, however, showed Giardia lamblia between the villi, and motile trophozoites were seen in the jejunal juice. After a three day course of high dose metronidazole she has been asymptomatic and her weight velocity has increased sharply.

Case 2
A 7 year old boy had steatorrhoea, 8 g/day, despite a low fat diet (25 g/day) and up to 50 pancreatin microsphere capsules per day. His weight was between the 10th and 25th centiles, and height was on the 10th centile. Attempts to introduce a higher fat diet produced abdominal discomfort, distention, flatulence, and gross steatorrhoea. Cimetidine had no effect. As a neonate he had a Bishop Koop resection and ileostomy for meconium ileus, which was closed by clamping at 1 month. A year previously he had been admitted with subacute obstruction, which responded to conservative management. Concentrations of serum iron, plasma ferritin, and vitamin A were low, and prothrombin time, vitamin D, and plasma bile acid concentrations were normal. Xylose absorption and a jejunal biopsy specimen were also normal, but fasting breath hydrogen was increased. Barium enema

References

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Accepted 5 August 1988
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Arch Dis Child 1988 63: 1494-1496
doi: 10.1136/adc.63.12.1494

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