**CASE REPORT**

**Gastric neuroendocrine carcinoma arising from heterotopic pancreatic tissue**

R Chetty, I Weinreb

An 85 year old man presented with symptoms of dyspepsia and increase in stool frequency of two months duration. Upper endoscopy revealed an ulcer and the biopsy was interpreted as carcinoma with endocrine features. A formal distal gastrectomy was planned, but intraoperatively, because of the patient’s condition, a wedge resection was performed. Histology revealed a neuroendocrine tumour (grade 1), located mainly in the submucosa, which had caused mucosal attenuation and ulceration. Associated with the tumour and at its periphery was heterotopic pancreatic tissue composed of acini, ducts, and endocrine cells. The tumour was strongly positive for chromogranin and focally for synaptophysin. There was no associated chronic atrophic gastritis or G cell hyperplasia. A discrete focus of high grade neuroendocrine carcinoma (grade 3) within the typical (grade 1) neuroendocrine tumour was also present. This case illustrates a grade 1 neuroendocrine carcinoma, with a small grade 3 focus, arising within or intimately associated with heterotopic pancreatic tissue in the stomach. Although the two entities may be separate, their close topographical association favours the possibility of neuroendocrine carcinoma arising from the heterotopic pancreas. Pathologists should be aware of the occurrence of pancreatic heterotopia in the stomach and the association of carcinoma with it.

Pancreatic heterotopia has been defined as “the occurrence of pancreatic tissue in aberrant anatomic sites that lack vascular, neural and anatomic continuity with the pancreas”.¹ In general, pancreatic heterotopia is rare and postmortem frequency rates range from 0.6% to 13.7%.² It is usually asymptomatic and the common gastrointestinal sites of occurrence include the stomach, duodenum, jejunum, Meckel’s diverticulum,³ gallbladder,⁴ and oesophagus.⁵ If symptomatic, abdominal pain, melaena, anaemia, obstruction, and cystic change have all been documented.⁶ Indeed, even carcinomas arising within heterotopic pancreatic tissue have been described.⁷–⁹ To the best of our knowledge, a case of neuroendocrine carcinoma arising in gastric heterotopic pancreas has not been described previously.

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**MATERIALS AND METHODS**

Representative sections from the surgical specimen, which was fixed in 10% neutral buffered formalin, were processed in a routine fashion. In addition to haematoxylin and eosin stains, immunohistochemistry was performed on the formalin fixed, paraffin wax embedded tissue using the liquid streptavidin–biotin complex immunoperoxidase method, with diaminobenzidine as chromogen. Immunohistochemistry was performed with a panel of immunohistochemical markers.

**RESULTS**

**Gross examination**

The surgical specimen was received in three pieces, the largest measuring 6.5 × 1.5 × 1.0 cm. The mucosal aspect appeared slightly protuberant and showed a haemorrhagic focus measuring 1.0 × 0.7 cm, with the remainder of the mucosa appearing normal. The serosal surface was normal. On sectioning the gastric wall, a well circumscribed, pale submucosal mass was noted, underlying the haemorrhagic focus. This mass measured 1.7 × 1.7 × 0.9 cm, and it had a uniform cut surface with no areas of haemorrhage or necrosis. The mass was submitted in toto for histological assessment.
Histological examination
A well demarcated, circumscribed submucosal tumour was identified under low power magnification. The overlying muscularis mucosae was largely intact (fig 1) except for a focal area where it was breached by the tumour. Here, small foci of the tumour lay within the lamina propria. The mucosal surface showed focal ulceration with an acute inflammatory exudate. The deep aspect of the tumour was well confined by the muscularis propria, which was not invaded or breached. At the periphery of the tumour, adjacent to the muscularis propria, was a discrete focus of pancreatic tissue composed mainly of ducts (fig 2). Very occasional acinar cells were evident and a focus of normal appearing islet cells was noted intimately associated with the neuroendocrine tumour, which appeared to be arising from these cells. The entire heterotopic focus was within the tumour mass and confined by the muscularis propria. The tumour was arranged in nests or packets surrounded by delicate capillaries, in addition to trabeculae, pseudo-papillary and pseudo-acinar patterns. Individual tumour cells were uniform with ample eosinophilic cytoplasm, and round to oval nuclei containing dispersed, salt and pepper chromatin (fig 3). There was minimal pleomorphism, and in the most mitotically active areas the mitotic count was three for each 10 high power fields. Necrosis or areas of haemorrhage were not present. These morphological features were typical of a well differentiated gastric neuroendocrine carcinoma (grade 1). A discrete microscopic hypercellular focus at the periphery of the tumour, separate from the heterotopic pancreas, was also present (fig 4). This focus had a more trabecular pattern, with occasional pseudo-acinar areas, and was sharply demarcated from the typical neuroendocrine component. The cells were larger, elongated to spindle shaped, with vesicular nuclei, small but distinct nucleoli, and easily identifiable mitoses. This was construed as a focus of high grade neuroendocrine carcinoma (grade 3). Importantly, the overlying and adjacent gastric epithelium did not show evidence of chronic atrophic gastritis or endocrine cell hyperplasia. This last point was confirmed by the absence of G cell hyperplasia on gastrin immunostaining.

The typical neuroendocrine areas were strongly immunoreactive for chromogranin and focally for synaptophysin, CD56, and AE1/3. The high grade area showed a similar immunoprofile. The ducts in the heterotopic pancreatic tissue stained with epithelial markers, whereas the non-neoplastic endocrine cells were positive for chromogranin and focally positive for synaptophysin. The MIB-1 labelling index in both the classic grade 1 areas and the grade 3 focus was 5–10%.
Take home messages

- We present the case of an 85 year old man who had a grade 1 neuroendocrine carcinoma, with a small grade 3 focus, arising within or intimately associated with heterotopic pancreatic tissue in the stomach.
- Although the two entities may be separate, their close topographical association favours the possibility of neuroendocrine carcinoma arising from the heterotopic pancreas.
- Practising pathologists should be aware of the occurrence of pancreatic heterotopia in the stomach and the association of carcinoma with it.

The surrounding non-neoplastic tissue showed a labelling index of less than 1%. All other markers were negative.

**DISCUSSION**

Several theories have been advanced to explain the pathogenesis and occurrence of pancreatic heterotopia. The most tenable and plausible implicates migration and/or rests of embryonic pancreatic tissue buds from the developing pancreas during embryogenesis. This theory is most likely when one considers gastrointestinal pancreatic heterotopia: however, pancreatic tissue occurring in more exotic, anatomically remote sites, such as the thyroid gland or fallopian tube, may have a different pathogenesis. The possibility of origin from teratomas has been advanced in these instances.

Heterotopic pancreas in the stomach is usually located within 5 cm of the pylorus, may occur on either anterior or posterior gastric walls, and is more common along the greater curve.\(^1\) The submucosal and subserosal localisation has also been encountered.\(^2\) The submucosal lesions may be sessile, polypoid, or nipple-like, where the duct associated with the heterotopic pancreas may communicate with the gastric lumen.\(^3\)

Gaspar-Fuentes and colleagues modified von Heinrich’s classification of pancreatic heterotopia.\(^12\) Type I was total pancreatic heterotopia with all pancreatic cell types present, type II is composed of pancreatic ducts only (the so-called canalicular variety), type III comprises acinar tissue only (exocrine pancreas), and type IV is made up of islet cells only (endocrine pancreas). The histological classification of the heterotopic pancreas in our case is type I because all elements were present, although the ducal component dominated.

As mentioned earlier, gastric heterotopic pancreas is most frequently asymptomatic, but when symptoms do arise they do so because of gastric mucosal erosion, so that abdominal pain, melena, anaemia, and nausea are the usual symptoms. Unusually, obstruction of the pyloric outflow tract and mucous retention have been described.

**REFERENCES**

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