Heterotopic gastric mucosa together with intestinal metaplasia and moderate dysplasia in the gall bladder: report of two clinically unusual cases with literature review

N Xeropotamos, A S Skopelitou, Ch Batsis, A M Kappas

Abstract
We report the clinicopathological findings of two patients with ectopic gastric mucosa within the gall bladder. The first patient, a 78 year old man, was asymptomatic. He was admitted to hospital for a colon adenocarcinoma. Intraoperatively, a firm nodule was palpable in the gall bladder. Histological examination of the resected specimen revealed a body type gastric mucosa in the submucosa, adjacent to which were extensive pyloric gland and intestinal metaplasia with mild to moderate dysplasia. The remaining gall bladder mucosa demonstrated changes of chronic cholecystitis. The second patient was a 62 year old woman with symptoms of chronic cholecystitis. The preoperative diagnosis was consistent with this diagnosis with a “polyp” at the junction of the neck and cystic duct. Cholecystectomy was performed and the histological examination of the resected specimen showed that the “polyp” consisted of heterotopic gastric mucosa with glands of body and fundus type. In the remaining mucosa, chronic cholecystitis was evident. To the best of our knowledge, this is the first report of a clinicopathological presentation of heterotopic gastric mucosa, pyloric gland type, and intestinal metaplasia with dysplastic changes in the gall bladder. As heterotopic tissue may promote carcinogenesis of the gall bladder, close attention should be paid to any occurrence of such lesions in this anatomical region.

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Keywords: heterotopic gastric mucosa; gall bladder; intestinal metaplasia; dysplasia; precancerous lesion

Heterotopia, from the Greek “heteros” (different) and “topos” (location), is defined as the occurrence of normal tissue in an abnormal location. Synonymously, the term “choristoma”, from the Greek “choristos” (separated), has also been used.

Case reports

PATIENT NO 1
A 78 year old man was admitted to our hospital for colon carcinoma. At operation, palpation of the gall bladder revealed an intramural firm nodule. Cholecystectomy was carried out together with colectomy. On gross examination the gall bladder specimen measured 3x1.3x0.8 cm and had a smooth and glistening serosa. After opening the specimen, at the body of the gall bladder, a firm nodular thickening of the wall was noticed, measuring 0.7 cm at its largest diameter. Histological examination revealed that the nodule was within the lamina propria and consisted of heterotopic gastric mucosa, with both body type and pyloric type gastric glands. The overlying mucosa was hyperplastic. In close proximity to this heterotopic mucosa, extensive pyloric gland and...
intestinal metaplasia (fig 1A, B) together with mild to focally moderate dysplastic changes in the metaplastic epithelium were observed (fig 2 A–C). In the remaining mucosa, typical features of chronic cholecystitis were obvious. Cholelithiasis was not found.

PATIENT NO 2
A 62 year old woman presented to our hospital with vague dyspeptic symptoms and intermittent postprandial right upper quadrant abdominal discomfort that was described as dull-aching, radiating sometimes through the back, and associated with nausea. She only rarely experienced vomiting. Physical examination revealed a positive Murphy’s sign. Oral cholangiogram showed abnormal gall bladder function with a filling defect suggesting a polyp at the junction of the neck and cystic duct. A sonogram clearly demonstrated a polyp at the same location. Cholecystectomy was carried out for cholecystitis and gall bladder polyp.

The cholecystectomy specimen measured 8×2×0.9 cm with a smooth gray-whitish glistening serosa. On opening, a smooth velvety green mucosa was found, measuring 0.4 cm in thickness, except at the region of the neck and the orifice of the cystic duct where a polypoid nodule of red-grayish colour was noticed measuring 0.8 cm at its greatest diameter. Gall stones were not observed.

Histological examination revealed that the polypoid lesion consisted of gastric heterotopic mucosa with parietal and columnar mucous secreting cells, almost identical to those observed in the previous case (fig 1B, C). The adjacent gall bladder mucosa showed typical features of chronic cholecystitis.

Figure 1  (A) Alcian blue-periodic acid-Schiff (PAS) stain in normal (arrowheads) and metaplastic gall bladder epithelium. The acid mucins of intestinal metaplasia are blue whereas neutral mucins of gastric metaplasia (pyloric type) are red (×320). (B) Heterotopic gastric mucosa of body type in the gall bladder; parietal cells (top) have a large pyramidal shape and plump eosinophilic cytoplasm (haematoxylin-eosin ×480); same area (bottom) stained with Alcian blue-PAS (×480). (C) Polyp of the gall bladder consisting of heterotopic gastric mucosa (haematoxylin-eosin ×180).

Figure 2  (A) Mild and focally moderately dysplastic epithelium (arrow), occurring in intestinal metaplasia (arrowheads) (Alcian blue-periodic acid-Schiff (PAS) stain ×320) (inset: mild dysplasia developing in pyloric type metaplastic epithelium (haematoxylin-eosin ×320). (B) In this field the transition between normal (arrowhead) and metaplastic epithelium (arrow) can be seen (haematoxylin-eosin ×320); mild dysplasia is focally well discerned (inset: haematoxylin-eosin ×850). (C) Intestinal metaplasia of a tubule (arrowheads) in close proximity to the normal gall bladder epithelium (haematoxylin-eosin ×540).
Heterotopic gastric mucosa in the gall bladder

Discussion

Heterotopic gastric mucosa in the gall bladder is extremely unusual. It was first described by Eguchi in 1934. Since then it has been reported in various organs and sites in the gastrointestinal tract, including the tongue, oesophagus, epiglottis, small bowel, vermiform appendix, rectum, and gall bladder. A survey of the world literature revealed 45 reports, including the present cases, and interestingly none shared the broad spectrum of clinical and histological findings of our cases.

There is a wide range in patient age, clinical presentation and symptoms, and roentgenographic and pathological findings. From the six extended reviews, it is apparent that the sex distribution is approximately equal with extended reviews, it is apparent that the sex distribution is approximately equal.

Embryologically, the gall bladder arises from the hepatic primordium. The latter originates from the ventral surface of the foregut in the 2.5 mm embryo, some distance caudal to the developing stomach. Both are in close proximity to the septum transversum during the later stage of development. It seems therefore that heterotopia could result either by entrapment of primitive gastric tissue, by heterotopic differentiation within the primitive gall bladder, by metastatic differentiation of the primitive gall bladder, or by metastatic differentiation of the proximal duodenal bud. Recently, the correct diagnosis of gastric heterotopia was made preoperatively, with detection of gastric mucosa in a Meckel's diverticulum by the use of pertechnetate scintigraphy, associated with H, receptor blocking agent. The differential diagnosis of gastric heterotopia in general includes fixed gall stone or (rarely) a neoplasm, and gastric heterotopia in general includes fixed gall stone or (rarely) a neoplasm, and for the gastric mucosa is the histological presence of fundic glands with both parietal and chief cells as well as pyloric glands, and most investigators have reported that heterotopic gastric mucosa involves all of these components (fundic type). APUD cells have been demonstrated in five cases, thyroid tissue in one, and pancreatic tissue in two. In three cases there was peptic ulceration of the wall of the gall bladder.

The characteristic feature of heterotopic gastric mucosa is the histological presence of fundic glands with both parietal and chief cells as well as pyloric glands, and most investigators have reported that heterotopic gastric mucosa involves all of these components (fundic type). APUD cells have been demonstrated in five cases, thyroid tissue in one, and pancreatic tissue in two. In three cases there was peptic ulceration of the wall of the gall bladder. Embryologically, the gall bladder arises from the hepatic primordium. The latter originates from the ventral surface of the foregut in the 2.5 mm embryo, some distance caudal to the developing stomach. Both are in close proximity to the septum transversum during the later stage of development. It seems therefore that heterotopia could result either by entrapment of primitive gastric tissue, by heterotopic differentiation within the primitive gall bladder, or by metastatic differentiation of the primitive gall bladder. Recently, the correct diagnosis of gastric heterotopia was made preoperatively, with detection of gastric mucosa in a Meckel's diverticulum by the use of pertechnetate scintigraphy, associated with H, receptor blocking agent. The differential diagnosis of gastric heterotopia in general includes fixed gall stone or (rarely) a neoplasm, and for the gastric mucosa is the histological presence of fundic glands with both parietal and chief cells as well as pyloric glands, and most investigators have reported that heterotopic gastric mucosa involves all of these components (fundic type). APUD cells have been demonstrated in five cases, thyroid tissue in one, and pancreatic tissue in two. In three cases there was peptic ulceration of the wall of the gall bladder. Embryologically, the gall bladder arises from the hepatic primordium. The latter originates from the ventral surface of the foregut in the 2.5 mm embryo, some distance caudal to the developing stomach. Both are in close proximity to the septum transversum during the later stage of development. It seems therefore that heterotopia could result either by entrapment of primitive gastric tissue, by heterotopic differentiation within the primitive gall bladder, or by metastatic differentiation of the primitive gall bladder. Recently, the correct diagnosis of gastric heterotopia was made preoperatively, with detection of gastric mucosa in a Meckel's diverticulum by the use of pertechnetate scintigraphy, associated with H, receptor blocking agent. The differential diagnosis of gastric heterotopia in general includes fixed gall stone or (rarely) a neoplasm, and for the gastric mucosa is the histological presence of fundic glands with both parietal and chief cells as well as pyloric glands, and most investigators have reported that heterotopic gastric mucosa involves all of these components (fundic type). APUD cells have been demonstrated in five cases, thyroid tissue in one, and pancreatic tissue in two. In three cases there was peptic ulceration of the wall of the gall bladder.
results are expected as it is well known that p53 immunoreactivity is detected in approximately 47–50% of colorectal carcinomas, tends to increase with tumour stage, and seems to be a late event in colon carcinogenesis. This may suggest that epithelium is in keeping with the results of incidental found fourth lesions should be undertaken to better understand its pathogenesis.

In contrast, the histological changes observed in our cases could be explained by the multidirectional differentiation of gall bladder epithelium during inflammation. Metaplasia could progress to dysplasia and hence to adenocarcinoma via a multistep process which may result from accumulation of genetic abnormalities. It is interesting that the opposite has also been observed—that is, most gall bladder adenocarcinomas in the series studied by Kushima and colleagues displayed gastric and/or intestinal type differentiation, both immunohistochemically and morphologically—suggesting that gall bladder adenocarcinomas develop and progress under induction of gastric and intestinal differentiation. Numerous studies on this subject have appeared in the literature. We believe that further investigations of the molecular biology of gall bladder precancerous lesions should be undertaken to better understand its pathogenesis.

To the best of our knowledge, this is the fourth incidentally found case in the literature, and the first demonstrating a wide spectrum of epithelial changes, including metaplasia and dysplasia.

Our second patient was not symptomatic until admission to hospital. This is curious given the role of acid secreting parietal cells in the production of symptoms. In terms of a developmental abnormality during organogenesis—that is, entrapment of gastric mucosa in the gall bladder—one could expect a clinically active lesion, secreting acid and producing symptoms. If it is the case, why did this lesion remain silent for so many years? A reduced number of parietal cells could be a possible explanation, as has previously been reported. In our case, a paucity of parietal cells was not observed. This histological finding, in relation to the non-specific clinical signs and late symptoms, was challenging and led us to hypothesise that this ectopic gastric mucosa could represent a heteroplastic or metaplastic process, due to a developmental error of multipotential/multidirectional mesenchymal cells, rather than a congenital defect.

In conclusion, we have presented the first report of asymptomatic heterotopic gastric mucosa in the gall bladder accompanied by pyloric and intestinal metaplasia together with dysplasia, and the first clinically silent, probably of metaplastic origin, gastric mucosa in the gall bladder, presenting as a polyp. Both conditions emphasise the role of inflammation in the sequence metaplasia → hyperplasia → dysplasia → carcinogenesis of the gall bladder. Furthermore, we propose that the term choristoma, in its etymological sense, could also be applied to lesions derived through a heteroplastic process, and not only to congenital tissue development error.  


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