Pyloric Gland Adenoma: Case Report

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Abstract

Pyloric gland adenoma (PGA), also called adenoma with gastric differentiation, is a rare neoplasm of the gastric mucosa that can appear as gastric heterotopia in several organs. A 49-year-old woman presented with gastric reflux and chronic elevation of liver enzymes. She had a history of type 2 diabetes mellitus, hypothyroidism and an unspecified allergy treated with deflazacor, and a family history of autoimmune diseases. A liver biopsy showed macro- and microvesicular steatohepatitis. Hepatitis B and virus serum tests were negative. Autoimmune hepatitis was suspected and investigated. As an evaluation for dyspeptic symptoms an upper gastrointestinal endoscopy was performed, showing diffuse gastroduodenitis. A few polyps were found and resected from the gastric fundus; histopathology revealed a pyloric gland adenoma. There is very few clinical data on this tumor type because it is frequently underdiagnosed and reported as dysplasia. Further research is needed on the pathophysiology of this disease.

Key words: Gastric tumors, pyloric gland adenoma, gastric polyps, dyspepsia, upper gastrointestinal endoscopy, Mexico.

Resumen

El adenoma pilórico, llamado también adenoma con diferenciación gástrica, es una neoplasia poco común de la mucosa gástrica que puede aparecer como heterotopia gástrica en diversos órganos. Una mujer de 49 años de edad se presenta con refluo gastroesofágico y elevación crónica de enzimas hepáticas. Cuenta con los antecedentes de diabetes mellitus tipo 2, hipotiroidismo y una alergia no especificada en tratamiento con deflazacort, así como historia familiar de enfermedades autoinmunes. La biopsia hepática mostró estaohepatitis macro y microvesicular. Las pruebas virales para hepatitis B y otros resultaron negativas. Se sospechó hepatitis autoinmune por lo que fue investigada. Debido a los síntomas de refluo gastroesofágico se realizó una endoscopia que mostró gastroduodenitis difusa. Se encontraron algunos pólipos en el fundus gástrico los cuales se resecaron; el examen histopatológico mostró un adenoma pilórico. Hay pocos datos en la literatura debido a que es frecuentemente subdiagnosticado y reportado como displasia. Es necesaria más investigación en la histopatología de esta enfermedad.

Palabras clave: Tumores gástricos, adenoma pilórico, pólipos gástricos, dispepsia, endoscopia digestiva alta, México.

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Introduction

Gastric adenomas are characterized by polypoid projections of dysplastic epithelium. They represent 7% to 10% of gastric polyps and are classified into foveolar and pyloric gland types. This neoplasm was first described by Elster in 1976. Later, Borchard et al. presented two cases of pyloric gland adenoma that showed a transition to a moderately differentiated adenocarcinoma. In 1990, Watanabe included these lesions in the World Health Organization Classification of Gastric Tumors. The most frequent location of these tumors is in the mucosa of the stomach corpus. However, pyloric gland-type adenomas (PGAs) have also been reported at other areas such as the duodenum, gallbladder and bile duct. A case of a 49-year-old woman with a pyloric gland adenoma in the fundic mucosa of the stomach is here presented.

Case report

A 49-year-old woman presented at the gastroenterology service of the Medica Sur Clinic and Foundation in Mexico City for evaluation. She had a history of type 2 diabetes mellitus, hypothyroidism and an unspecified allergy treated with deflazacort, so as family history of autoimmune diseases. She had a history of elevated liver enzymes: AST 321 IU/L, ALT 387 IU/L, LDH 534 IU/L and GGT 179 IU/L. Hepatitis B and C virus infection and autoimmune hepatitis were also ruled out even when serum antinuclear antibodies were positive. After a series of liver biopsies the diagnosis of steatohepatitis (NASH) was established. In February of 2009, she presented again with symptoms of gastroesophageal reflux. Because of her dyspeptic symptoms, an upper GI endoscopy was performed where diffuse gastroduodenitis was found. Incidentally a few polyps from the gastric fundus were resected (Figure 1), and histopathology revealed a PGA (Figure 2). Immunohistochemistry showed the tissue to be positive to mucin core peptide 6 (MUC6) and MUC5AC, whereas it was negative for MUC2.

Discussion

PGA, also called an adenoma with gastric differentiation, is a very rare neoplasm of the gastric mucosa. Since it was first described, there have been cases reported of PGAs arising as gastric heterotopia in the gallbladder, duodenum, pancreatic main duct, rectum and Barrett’s esophagus. These tumors are frequently associated with dysplasia, Michal et al. reported a PGA arising in normal esophageal epithelium; as in the patient here reported, they found no abnormalities in the epithelium other than the pyloric gland adenoma. There is scant clinical data on this type of tumors because they are frequently underdiagnosed and...
Pyloric gland adenomas are more common in the elderly (eighth decade of life). Further evidence was provided by Vieth et al. through an analysis of 90 cases that showed a predominant localization in the corpus mucosa; they also pointed out that PGAs are more frequent in women than in men. Here, we found the tumor in the fundic mucosa of a mature woman.

Regarding the relationship of PGAs with lesions of the gastric mucosa, Abraham et al. have associated intestinal metaplasia and mucosal atrophy with the development of gastric adenomas. In this context, associations of PGAs with autoimmune gastritis have been reported. However, these findings are not present in all cases, and the influence of autoimmune gastritis in the etiology of PGAs has not been confirmed. Our patient, as for other patients with NAFLD, showed ANA-positive serum which does not confirm an autoimmune component. However, further research is needed on the pathophysiology of PGAs and their relationship with other diseases.

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References