Case Report

Non-\textit{Helicobacter pylori} Gastric Intestinal Metaplasia in Children: A Series of Cases and Review of the Literature

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1. Introduction

Gastric Intestinal Metaplasia (GIM) is a finding with unknown frequency and, more importantly, unknown clinical implications in children. We present two cases of pediatric patients with GIM and a review of the literature. We discuss the epidemiology of GIM in patients with \textit{Helicobacter pylori} (HP) gastritis and the potential role of HP gastritis and bile acid reflux in the development of GIM. We discuss histologic risk factors for the progression of GIM to gastric cancer. We also discuss the potential need for the long-term surveillance and natural history studies of GIM in children.

2. Case Report

2.1. Case 1. A 16-year-old female presented to the clinic complaining of progressive worsening of dysphagia to solid foods with sensation of fullness in the chest and sour taste in the mouth but denies heartburn or chest pain. She did not have any weight loss. Her symptoms were unresponsive to PPI therapy started by her primary care physician. There was no family history of gastric cancer. The physical exam was unremarkable and the blood work was normal (including celiac panel, comprehensive metabolic panel, and HP IgG). An esophagram was normal. She underwent esophagogastroduodenoscopy (EGD), which revealed a 4 mm prepyloric nodule (see Figure 1(a)). A rapid urease test for HP was negative. Hematoxylin and eosin-stained sections on the prepyloric nodule biopsy showed antral-type gastric mucosa (see Figures 1(b) and 1(c)). The lamina propria was dissected by a chronic inflammatory cell infiltrate consisting of lymphocytes and plasma cells. Numerous mucin producing cells, characteristic of intestinal epithelium, were identified. The features were those of chronic gastritis with intestinal metaplasia of the complete type. No dysplasia was present. No HP organisms were identified on light microscopy and by immunostain. Patient reports improvement in her symptoms after completing a course of double dose PPI therapy. Repeated EGD 1 year after showed resolution of the gastric erosion and persistence of the prepyloric nodule on EGD. Complete type intestinal metaplasia persisted on histologic evaluation.

2.2. Case 2. An 8-year-old female with generalized but not radiated abdominal pain, described as cramping and
sharp for the last one year, presented to our clinic. She reported intermittent nausea associated with nonbloody and nonbilious vomiting. Vomiting was more frequent at night. There was no family history of gastric cancer. The patient had an unremarkable blood work (including complete blood count, chemistry panel, antibodies to HP, and celiac panel). Computed tomography of the abdomen was significant for mesenteric adenopathy. The EGD showed a prepyloric nodule (Figure 2(a)) and bile-lake. Histological examination showed incomplete intestinal gastric metaplasia with irregular mucin droplets and an absent brush border (see Figures 2(b) and 2(c)). Rapid urease testing for HP, as well as immunostains, was negative. She was treated with double dose PPI therapy. Vomiting improved with cyproheptadine. Repeat EGD 6 months later showed resolution of the prepyloric erosion and continued presence of the prepyloric nodule. Histologic examination showed persistence of incomplete gastric intestinal metaplasia without progression to dysplasia.

3. Discussion

GIM is defined as the replacement of gastric columnar cells by cells of intestinal morphology characterized by the presence of mucin-containing goblet, Paneth, and absorptive cells [1]. The intestinal cells are easily distinguished in the gastric mucosa, because they are not present in healthy gastric mucosa [2]. The histopathologic diagnosis of GIM has been found to have high interobserver agreement [3, 4]. No consensus is available about the optimal number or location of biopsies needed in children [5]. In adults, biopsy mapping of the stomach requires at least 5 biopsy specimens: 2 from the antrum within 2 to 3 cm from the pylorus (1 each from the lesser and greater curvatures); 2 from the corpus approximately 8 cm from the cardia (1 each from the lesser and greater curvatures); and 1 from the incisura angularis [6].

The prevalence of GIM in children is largely unknown [7]. Furthermore, endoscopic features of GIM in pediatric patient are poorly defined. A white opaque substance visualized by
Figure 2: Endoscopic and histologic appearance of gastrointestinal metaplasia in case 2. (a) shows the appearance of a prepyloric nodule with an erosion located on the lesser curvature of the antrum. A cold forceps biopsy was taken from a region adjacent to the erosion (arrow). (b) shows a low power image and (c) shows a high power histopathology image of the prepyloric nodule in case 2. The lamina propria is distended by a chronic inflammatory cell infiltrate, irregular columnar cells filled with mucin, and an absence of brush border is noted, signifying an incomplete type.

Magnifying endoscopy with narrow-band imaging (M-NBI) appears to be a useful indicator of the histological diagnosis of GIM [8]. GIM is a common finding on routine endoscopy in adults [9] and is more frequently associated with HP than in children [10]. The frequency of GIM in children related to HP-positive gastritis versus HP-negative gastritis is variable. Shabib et al. [11] reported a frequency of 42% in children with HP-positive gastritis versus 6% in children with HP-negative gastritis. However, Kato et al. [12] documented no difference in the presence of intestinal metaplasia between the study groups of children with and without HP infection. However, no children in a Brazilian cohort of 96 children with HP gastritis were found to have GIM [13].

HP infection causes inflammatory cell infiltration in the gastric mucosa, resulting in atrophy of the foveolar epithelium and long-term mucosal changes such as intestinal metaplasia, which are precursors of gastric cancer [13–15]. HP organisms seem to be the most important member of the gastric microbiota with the highest relative abundance when present, but when it is absent, the stomach has a diverse microbiota [16]. Proteobacteria, Firmicutes, Actinobacteria, Bacteroidetes, and Fusobacteria are the most abundant phyla in both HP-positive and HP-negative patients [16].

Reactive gastropathy represents the second most common cause for the occurrence of age-dependent mucosal alterations [17]. Primary duodenogastric reflux (DGER) could cause gastric mucosal lesions manifested as intestinal metaplasia histologically in children. DGER is probably an independent etiological factor and might play a synergistic role in the pathogenesis of gastric mucosal lesions along with gastric acid and HP infection [18]. Other causes that are associated with GIM in adults include high gastric pH, increased bile acid exposure, smoking [19], and gastric denervation after surgery for benign disease [20].

The diagnosis of intestinal metaplasia can have adverse clinical implications and should be made with caution in a child [21]. The association of GIM with adenoma/dysplasia/carcinoma progression is commonly encountered in adults but is rarely seen in children. Only 10% of gastric cancer cases are found in patients younger than 40 years of age [22]. It is very likely that time plays an important factor the progression of GIM to adenoma. The progression from intestinal metaplasia to gastric adenocarcinoma takes an average of about 7 years in adult studies [23]. Thus, by the time GIM undergoes neoplastic transformation, the patient would become an adult and, therefore, managed in the adult service [5, 10]. The malignant potential of GIM has been shown to vary based on histologic subtype, location, and extent of mucosal involvement [24, 25]. Adults with incomplete GIM subtype versus complete GIM subtype,
diffuse involvement of the antrum and gastric body versus antrum alone, and greater than 20% extension of mucosal involvement between endoscopic sampling had a greater risk of gastric cancer [25, 26]. Family history of gastric cancer on initial evaluation was associated with increased risk of subsequent gastric cancer in adult patients [9, 23, 24].

Treatment, long-term consequences, and surveillance protocols of GIM are not well established in the pediatric population. Contrary to our first case presented, a case report of a 15-year-old with GIM located inside the cryptic antral population. Contrary to our first case presented, a case report protocols of GIM are not well established in the pediatric practice or results in widely varying follow-up frequency or need for consensus on the follow-up of this histopathological finding. At present, GIM is frequently disregarded in clinical practice or results in widely varying follow-up frequency or treatment. These uncertainties require further research in the pediatric population.

Disclosure

An earlier version of this work was presented as a poster at Clinical Vignette Abstracts NASPGHAN 2017.

Conflicts of Interest

The authors have no conflicts of interest to report with regard to this publication.

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