Endoscopic assessment of children with esophageal atresia: Lack of relationship of esophagitis and esophageal metaplasia to symptomatology

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BACKGROUND: Late complications of esophageal atresia (EA), particularly esophagitis and Barrett’s esophagus, are increasingly being recognized. With the exception of patients with dysphagia associated with esophageal stricture, it is unknown whether patient symptomatology can predict endoscopic findings.

METHODS: Data regarding the digestive symptoms of patients who were referred to the EA multidisciplinary clinic from October 2005 to October 2008, and underwent upper gastrointestinal endoscopic evaluation, were systematically collected. Macroscopic and histological findings were analyzed. Endoscopy was considered normal if no esophagitis, intestinal metaplasia or gastric metaplasia (GM) was discerned.

RESULTS: Sixty-three patients underwent endoscopy. Eighteen had dysphagia related to an esophageal stricture needing dilation and were subsequently excluded from the analysis. Forty-five patients (26 girls) with a median age of 7.3 years (range 0.4 to 17.9 years) were evaluated. Twenty-six patients (58%) were normal at endoscopy, 14 patients (31%) had esophagitis and 16 patients (36%) had GM. No intestinal metaplasia or adenocarcinoma was detected. Six patients with abnormal endoscopy results were asymptomatic. No correlation between digestive symptoms and endoscopy results was found.

CONCLUSION: The present cross-sectional study showed that symptomatology was not predictive of abnormal endoscopy in EA patients. Esophagitis or GM may be discovered, even in the absence of symptoms, suggesting that physicians cannot rely solely on symptomatology to accurately evaluate the extent of these esophageal complications in this population.

Key Words: Barrett’s esophagus; Endoscopic findings; Esophageal atresia; Esophagitis; Gastric metaplasia; Intestinal metaplasia

Since the first successful surgery in 1941 (1), the survival of patients with esophageal atresia (EA) has greatly improved (2-6). Subsequently, the follow-up of these patients has revealed gastrointestinal (GI) complications such as gastroesophageal reflux (GER), esophagitis and their related consequences (7-10), which were unrecognized 60 years previously.

Among these complications, one of the major concerns in the long-term GI follow-up is the increased incidence of gastric metaplasia (GM) of the esophagus (11-13), intestinal metaplasia (8,14) and esophageal adenocarcinoma (15-17), which have all been reported in young adults with EA (14-16). However, the exact incidence and natural history of these complications are unknown in EA patients.

It has not been established whether systematic upper GI endoscopic screening in the follow-up of young patients with EA is recommended. Finding clinical symptoms predictive of
endoscopic appearance would be very helpful in the clinical management of these patients.

Thus, the aims of the present cross-sectional study were to describe esophagitis, Barrett’s esophagus (BE) and metaplastic features (macroscopic and microscopic) of the esophagus in a cohort of children with EA who were proposed a systematic endoscopic examination, and to verify whether a correlation between endoscopic observations and patient symptoms exists.

METHODS

Patients
In October 2005, a specialized EA clinic was created at the Centre Hospitalier Universitaire Sainte-Justine (Montreal, Quebec). Patients of all ages were invited to be followed by the clinic’s multidisciplinary team. At their first visit, all patients were routinely proposed upper GI endoscopy if they were older than two years of age. If they were younger than two years of age, indication for endoscopy was based on symptoms (dysphagia, food impaction and/or hematemesis). The study period extended from October 2005 to October 2008.

Patients were excluded from the study if they had previously undergone gastric tube reconstruction, colonic interposition or esophagogastric dissociation. Patients with dysphagia associated with esophageal stricture needing dilation at endoscopy were not included in the analysis because they did not have esophageal biopsies taken when endoscopic dilation was performed.

Clinical data collection
Digestive symptoms were systematically recorded by asking specific predetermined questions. A data sheet was available at each visit. Questions were appropriate for all ages: patients (or parents) were asked whether they (or their child) experienced regurgitation or ‘food coming up’ (clinical GER), heartburn (pyrosis), trouble swallowing (dysphagia), pain with swallowing (odynophagia), food blockage (food impaction) and coughing at meals.

Endoscopic data collection
Macroscopic data were collected prospectively. Reports of macroscopic esophagitis were noted according to the Savary-Miller classification or the presence of endoscopically visible breaks in the esophageal mucosa at or immediately above the gastroesophageal junction (18). The BE macroscopic aspect was defined as velvety-red tongue extending up in the esophagus from the proximal gastric folds at the gastroesophageal junction (Figure 1). BE endoscopic aspect was differentiated from hiatal hernia or gastric pull-up by carefully delimiting the gastroesophageal junction, identified as the proximal margin of the gastric folds (Figure 2). Each patient systematically underwent at least one esophageal biopsy. When a BE macroscopic aspect was seen, at least one supplementary biopsy was systematically taken from the segment in question. Care was taken to accurately biopsy the esophageal mucosa and not the gastric mucosa. Biopsies were taken above the proximal gastric folds, at least 1 cm away from the gastroesophageal junction. Such caution was needed mostly in patients with long-gap atresia, where gastric pull-up was often noticed.

A total of eight different pediatric endoscopists evaluated the patients. A specialized endoscopic nurse was present for each endoscopy.

Both macroscopic esophagitis and BE had to be confirmed by histology to be considered for analysis in the present study.

Pathology data
Esophagitis was defined as the elongation of papillae and basal hyperplasia. Biopsies from endoscopically suspected esophageal metaplasia that showed a columnar epithelium were ‘flagged’ as BE. As proposed by the Montreal definition and classification of GER (18), endoscopically suspected esophageal
metaplasia, both cardiac- and intestinal-type metaplasia should be included in the definition of BE (18,19). Metaplasia type was defined as BE, gastric GM positive (GM+) (Figure 3) or BE, specialized intestinal metaplasia positive (19-22). In biopsies for which metaplasia was suspected, Alcian blue dye was used to highlight goblet cells and readily distinguish intestinal metaplasia. One pathologist (DBDS) systematically reviewed all the biopsies that were previously analyzed by another pathologist who was a member of the department of pathology. Discrepancies in analyses were resolved by discussion, with final diagnosis established by consensus between the two participating pathologists.

**pH monitoring**
When a pH study was performed, the reflux index (23) was used as the criterion for acid reflux: for children younger than one year of age, the percentage of time at pH<4 for more than 10%, and for older than one year of age, the percentage of time for more than 5% were considered to be abnormal.

**Statistical analysis**
Endoscopy was considered normal if there was no evidence of macroscopic and microscopic esophagitis, no BE/GM+, and no BE/specialized intestinal metaplasia positive. Symptoms were analyzed according to endoscopic findings; differences between groups were analyzed by the $\chi^2$ test, with P<0.05 considered to be statistically significant.

The study was approved by the local institutional board.

**RESULTS**
Sixty-three patients underwent endoscopy. No patient refused the evaluation. Eighteen patients needed dilation for esophageal stricture and were subsequently excluded from the analysis (Figure 4). The study group consisted of 45 patients (26 girls) with a median age of 7.3 years at the time of endoscopy (range five months to 17 years, 11 months). Thirty-seven patients (82%) had type C atresia (24), 10 (22%) had long-gap atresia, and 11 patients (24%) had VACTERL (vertebral, anal, esophageal, renal, limb defects) association (Online Mendelian Inheritance in Man 192350). Twenty patients (44%) underwent a fundoplication procedure before the endoscopy and 10 (22%) had a history of esophageal stricture needing dilation.

Twenty-six patients (58%) were normal at endoscopy. Fourteen patients (31%) had histologically proven esophagitis, and 10 (36%) had GM (BE/GM+) (Figure 3). In the latter patient group, the mean age was 9.8 years (range 3.4 to 13.2 years), and five patients had long-gap atresia. No intestinal metaplasia, dysplasia or adenocarcinoma was found. As reported in Figure 5, both esophagitis and GM were encountered in patients four years of age and older.

pH monitoring was performed in 24 of the 63 patients. Of the 14 patients with esophagitis, six underwent pH monitoring, which revealed abnormal results in all patients who were not on antacid medication (ie, proton-pump inhibitors). Of the 16 patients with GM, seven underwent pH monitoring, with abnormal results in only three of these seven.

Twenty-eight patients (62%) were symptomatic. The symptoms reported were dysphagia in 40%, food impaction in 36%, coughing during meals in 27%, regurgitation in 20%, pyrosis in 13% and odynophagia in 2%.

Of 17 completely asymptomatic patients, six had abnormal endoscopy results. Of these six patients, four had both esophagitis and GM. In contrast, of the 28 symptomatic patients, 15 had normal endoscopic findings.

No association was found between the symptoms and endoscopic findings analyzed either as a whole (abnormal endoscopy) (Table 1) or separately (esophagitis or GM).

**DISCUSSION**
The present study was one of the largest pediatric endoscopic series of patients with EA (11-14,25). Our study was based on systematic endoscopic evaluation of all EA patients, with the intention of specifically screening for BE and esophagitis. Mucosal abnormalities at endoscopy were observed in 19 of 45 patients (42%) in our cross-sectional study. Symptoms were reported by 62% of patients, but none could be identified as being statistically associated with an abnormal endoscopic finding. Furthermore, six asymptomatic patients had abnormal endoscopy results, reflecting the difficulty in adequately deciding which patients need to be investigated.
Patients with strictures needing dilation were excluded from the analysis because the majority of them did not have biopsies at endoscopy – most gastroenterologists at our institution prefer not to perform an esophageal biopsy when performing a dilation. Furthermore, these patients had obvious symptoms requiring immediate attention. If patients with strictures had been included in the analysis, dysphagia would have been statistically associated with abnormal endoscopic findings but only due to cases of significant strictures requiring dilation.

In patients followed for EA in recent years, our attention has been focused on significant complications of GER, such as BE and esophageal adenocarcinoma. BE with intestinal metaplasia has been focused on significant complications of GER, such as BE and esophageal adenocarcinoma. BE with intestinal metaplasia has been associated with significant complications of GER, such as BE and esophageal adenocarcinoma. BE with intestinal metaplasia has been statistically associated with abnormal endoscopic findings but only due to cases of significant strictures requiring dilation.

In patients followed for EA in recent years, our attention has been focused on significant complications of GER, such as BE and esophageal adenocarcinoma. BE with intestinal metaplasia has been identified as the proximal margin of the gastric mucosal folds (Figures 1 and 2), as defined by Prague C & M criteria (32). Moreover, only six of the 16 subjects with GM had long-gap atresia. Therefore, we believe that the specimens diagnosed as GM were truly GM and not erroneously diagnosed gastric mucosa.

Finding specific symptoms associated with abnormal endoscopy would have been helpful. In their long-term follow-up of adults with EA, Taylor et al (33) investigated endoscopic abnormalities only in symptomatic patients and reported a 58% incidence of esophagitis and an 11% incidence of intestinal metaplasia. Based on our study, esophageal mucosal abnormalities can be observed in EA patients at endoscopy despite the absence of symptoms (six of 17 patients [35%]), making the recommendation of endoscopic assessment based solely on symptomatology inappropriate.

On the other hand, our investigation shows that some symptomatic patients were completely normal at endoscopy. Symptoms such as dysphagia, odynophagia or food impaction can be related to esophageal dysmotility in this population of patients, as demonstrated by manometric studies (12,34).

A limitation of the present study is the total number of biopsies taken because it did not strictly follow the guidelines for BE surveillance (18,20). Therefore, we acknowledge that some GM or intestinal metaplasia may have been missed and that our results are most likely an underestimation of the true number of these conditions.

Another limitation is the sample size; therefore, it is possible that no association between symptoms and endoscopic findings could be statistically demonstrated because of an insufficient number of patients. However, Locke et al (35) noted that in a non-EA population of 1011 adults, symptoms were only weakly predictive of findings at endoscopy. In infants, the presence of esophagitis is difficult to predict on the basis of symptoms. Chadwick et al (36) reported a poor correlation between symptoms and histological changes. Salvatore et al (37) demonstrated that questionnaires are poorly predictive of GER disease severity because they do not correlate with esophageal acid exposure, measured by pH-metry, and with esophagitis, evaluated by the histological analysis of esophageal biopsies.

**CONCLUSION**

Our study indicates that symptoms are not predictive of abnormal endoscopy in patients with EA. Furthermore, even in the absence of symptoms, abnormal findings may still be discovered. Many of these patients present with GM for which no exact modalities exist with regard to its follow-up and surveillance. Considering the high prevalence of GER...
in this population, the systematic endoscopic follow-up and evaluation of esophageal biopsies in EA patients to accurately evaluate complications, response to treatment and outcomes are warranted. Close collaboration between gastroenterologists and pathologists should be established, lesions should be described according to standardized landmarks and biopsy samples should be taken with special care.

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