

Characteristics of Upper Gastrointestinal Tract Involvement in Korean Pediatric Crohn's Disease: A Multicenter Study

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Purpose: Crohn's disease (CD) can involve any site of the gastrointestinal tract (GIT). However, the characteristics of upper GIT involvement in CD are unclear, especially in the Eastern pediatric population. This study aimed to estimate the prevalence of upper GIT involvement and identify the clinical features of Korean children with CD.

Methods: This was a retrospective multicenter cohort study that included 52 pediatric patients with CD who underwent esophagogastroduodenoscopy and biopsy. The clinical symptoms and endoscopic and histologic features of the upper GIT were identified according to the presence or absence of upper gastrointestinal symptoms.

Results: Among the 52 patients, upper GIT involvement was noted in 50.0% (26/52). The mean age at CD diagnosis was 14.1±2.1 years. Gastric ulcer was the most common lesion (19.2%) found on upper GIT endoscopy, followed by duodenal ulcers (15.4%). Chronic inflammation was the most common histopathologic feature (75.0%), followed by gastric erosion (17.3%). Granuloma was found in 9.6% of patients. *Helicobacter pylori* infection was identified in 5.8% of patients. Endoscopic and histologic findings were not significantly different, but the mean values of erythrocyte sedimentation rate (60.7±27.1 vs. 43.0±27.6 mm/h, $p=0.037$) and C-reactive protein (16.5±28.2 vs. 6.62±13.4 mg/dL, $p=0.014$) were significantly different between patients with and without upper gastrointestinal CD symptoms.

Conclusion: Upper GIT involvement was relatively common in pediatric patients with CD irrespective of upper gastrointestinal symptoms, and *H. pylori* infection was relatively uncommon. The results of this study should aid the establishment of regional guidelines for upper GIT examination.

Key Words: Crohn disease, Upper gastrointestinal tract, Granuloma, *Helicobacter pylori*, Pediatrics

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INTRODUCTION

Crohn's disease (CD) is a chronic inflammatory bowel disease of unknown cause. It occurs primarily in the terminal ileum, but can involve any area of the gastrointestinal tract (GIT), from the mouth to the anus [1]. To date, much research has been conducted to investigate the involvement of the lower GIT, but the involvement of the upper GIT in CD has not been studied sufficiently.

In studies involving Italian and American adults, the prevalence of upper gastrointestinal (GI) CD is reported to be 16-34% [2,3]. In pediatric patients with CD, studies have reported different findings; for example, the upper GIT was involved in 71% of cases in a study conducted in Canada [4], whereas, in a study conducted in the UK, esophagitis involvement was noted in 72%, gastritis in 92%, and duodenitis in 33% [5]. Moreover, in a study conducted in the US, 36% of patients showed symptoms suggestive of upper GI involvement, and 12% of pediatric patients demonstrated noncaseating granulomas [6]. In a study involving Korean adults who underwent upper GIT biopsy, 59% of cases had upper GI symptoms. Erosive gastritis was the most common endoscopic finding, and was noted in 66% of patients. In another study, approximately 70% of patients had an abnormal histological finding. However, these studies were based on only a few cases, and moreover, the focus was on abnormal findings in the stomach or *Helicobacter pylori* [7]. To date, data has been scarce regarding pediatric patients in Korea. In Asians, CD is more male-predominant (1.67:1 to 2.9:1) compared to in other races. Moreover, CD of an isolated colonic type is most common in Europeans, whereas CD involving both the small and large bowels is most common among Asians [8-10].

Accordingly, this study aimed to examine whether the presence or absence of upper GI symptoms was related to upper GIT lesions in pediatric patients with CD and to investigate whether the relationship would help determine the need for testing. An additional objective of the study was to examine the clinical features, endoscopic and histological findings,

and the prevalence of *H. pylori* in CD with upper GIT involvement to help create a guideline for upper GIT testing in pediatric CD patients in Korea.

MATERIALS AND METHODS

Study design and subjects

Data from the medical records of pediatric patients (age < 18 years) with CD who underwent upper GIT endoscopy and biopsy between January 2001 and August 2016 were collected from 4 Korean tertiary hospitals. The 4 medical centers included Gachon University Gil Medical Center located in Incheon, Chungbuk National University Hospital in the central region, Kangwon National University School of Medicine in the north region, and Chung-Ang University Hospital in Seoul, South Korea.

Of 64 subjects, 52 were finally enrolled after meeting the following criteria: (1) patients who underwent biopsy for abnormal gastric lesions and/or normal-looking gastric mucosa and (2) patients who had no known chronic medical illnesses. Our diagnostic criteria for CD has been previously described [8,11].

We retrospectively reviewed the patients' medical records, upper GI endoscopic findings, and histopathological findings from the time of CD diagnosis. The following demographic and clinical information was collected: gender, GI symptoms (diarrhea, lower abdominal pain, weight loss, hematochezia, and poor oral intake), upper GI symptoms (epigastric pain, nausea, vomiting, dyspepsia, burning sensation, epigastric fullness, belching, and melena), perianal lesions (skin tag, fissure, abscess, and fistula), and medications ever used for CD (anti-tumor necrosis factor- α therapy [infliximab, adalimumab], immunosuppressive therapy [steroid], immunomodulators [6-mercaptopurine, azathioprine], 5-aminosalicylic acid [mesalamine, sulfasalazine], and antibiotics). Laboratory findings such as erythrocyte sedimentation rate (ESR) and C-reactive protein (CRP) were identified. Disease locations were evaluated according to the Montreal classification. Disease activity was evaluated with the pediatric Crohn's disease

activity index (PCDAI) and was classified as remission (PCDAI <10), mild activity (PCDAI 10-27.5), moderate activity (PCDAI 30-37.5), and severe activity (PCDAI >40) [12]. The study protocol was approved by the institutional review boards of Gachon University Gil Medical Center (IRB no. 2016-368).

Upper GIT endoscopy, histologic evaluation, and *H. pylori* infection assessment

All upper GI endoscopic evaluations were performed by expert GIT endoscopists of the participating institutions. Biopsy specimens were obtained for

pathologic evaluation from abnormal gastric lesions and normal-looking gastric mucosa at the endoscopists' discretion. Additional biopsies were performed at both the gastric antrum and corpus for the rapid urease test. Standard hematoxylin and eosin staining was performed on biopsy specimens. The diagnosis of *H. pylori* infection was based on either a positive histopathology plus a positive rapid urease test, or a positive culture. Gastric biopsies were obtained for histopathology [13].

The diagnosis of "upper GI involvement of CD" was based on a combination of compatible endo-

Table 1. Baseline Characteristics of Subjects

Characteristic	Total (n=52)	UGI symptoms (n=16)	No UGI symptoms (n=36)	p-value
Male	34 (65.4)	9 (56.3)	25 (69.4)	0.356*
Age at diagnosis of CD (y)	14.1±2.1	14.2±1.7	14.1±2.3	0.773 [§]
Montreal disease location				
L1, terminal ileum	3 (5.8)	0 (0)	3 (8.3)	0.544 [†]
L2, colon	16 (30.8)	7 (43.8)	9 (25)	0.206 [†]
L3, ileocolon	33 (63.5)	9 (56.3)	24 (66.7)	0.472*
L4, upper gastrointestinal only	0 (0)	0 (0)	0 (0)	
Perianal lesion	31 (59.6)	8 (50.0)	23 (63.9)	
Skin tag	11 (21.2)	3 (18.8)	8 (22.2)	0.587 [†]
Fistula	5 (9.6)	2 (12.5)	3 (8.3)	0.505 [†]
Fissure	2 (3.8)	0 (0)	2 (5.6)	0.552 [†]
Abscess	2 (3.8)	0 (0)	2 (5.6)	0.341 [†]
Mixed	11 (21.2)	3 (18.8)	8 (22.2)	0.587 [†]
PCDAI at diagnosis				
Remission	0 (0)	0 (0)	0 (0)	
Mild	8 (15.4)	1 (6.3)	7 (19.4)	0.409 [†]
Moderate	17 (32.7)	3 (18.8)	14 (38.9)	0.153*
Severe	27 (51.9)	12 (75.0)	15 (41.7)	0.056*
HP positive	3 (5.8)	0 (0)	3 (8.3)	0.544 [†]
ESR (mm/h)	48.5±28.4	60.7±27.1	43.0±27.6	0.037 [†]
CRP (mg/dL)	9.7±19.6	16.5±28.2	6.62±13.4	0.014 [§]
Anti-TNF- α therapy	13 (25.0)	4 (25.0)	9 (25.0)	
Infliximab	11 (21.2)	3 (18.8)	8 (22.2)	1.000 [†]
Adalimumab	2 (3.8)	1 (6.3)	1 (2.8)	0.525 [†]
Immunosuppressive therapy	14 (26.9)	3 (18.8)	11 (30.6)	0.506 [†]
5-ASA compounds	43 (82.7)	12 (75.0)	31 (86.1)	
Mesalamine	42 (80.8)	12 (75.0)	30 (83.3)	0.475 [†]
Sulfasalazine	1 (1.9)	0 (0)	1 (2.8)	1.000 [†]
Antibiotic therapy	12 (23.1)	5 (31.3)	7 (19.4)	0.478 [†]
Steroid therapy	34 (65.4)	12 (75.0)	22 (61.1)	0.331*
Surgery	2 (3.8)	1 (6.3)	1 (2.8)	0.525 [†]

Values are presented as number (%) or mean±standard deviation.

UGI: upper gastrointestinal, CD: Crohn's disease, PCDAI: pediatric Crohn's disease activity index, HP: *Helicobacter pylori*, ESR: erythrocyte sedimentation rate, CRP: C-reactive protein, TNF: tumor necrosis factor, 5-ASA: 5-aminosalicylic acid.

*Pearson chi-squared test, [†]Fisher's exact test, [†]Student t-test, [§]Mann-Whitney U-test.

scopic (ulcerations, erosions, strictures, and aphthous lesions) and histologic findings (chronic inflammation, erosion, ulceration, granuloma, and gastric intestinal metaplasia) as described previously [14,15]. Some nonspecific inflammation or inflammatory changes explained by other conditions (reflux esophagitis or *H. pylori* gastritis) were excluded.

After dividing the participants into 2 groups according to the presence or absence of upper GI symptoms, we evaluated the differences in demographics, *H. pylori* infection, treatment, surgery, PCDAI, perianal lesions, laboratory findings (ESR, CRP), and endoscopic and histologic findings.

Statistical analysis

Continuous variables were expressed as means with ranges. Discrete data were expressed as numbers or percentages or both. Demographics, clinical symptoms, and endoscopic and histologic GIT lesions of subjects with and without upper GI CD symptoms were compared. For comparative analyses, the Mann-Whitney U-test and Student's t-test were used for continuous variables, while the chi-squared test or Fisher's exact test were used for categorical variables, where appropriate. All statistical analyses were performed using IBM SPSS for Windows ver. 21.0 (IBM Co., Armonk, NY, USA). $p < 0.05$ was considered statistically significant.

RESULTS

Demographics and disease characteristics

Of the total 64 patients, 2 patients did not undergo upper GI endoscopy and 10 patients were excluded from the study because of inadequate data. A total of 52 patients were enrolled. The subjects included 34 men (65.4%), and the mean age at CD diagnosis was 14.1 ± 2.1 years. According to the Montreal classification, the dominant phenotypes were listed as follows: the sites of disease location were L3 (ileocolon, 33/52, 63.5%), L4 (isolated upper GI, 0/52, 0%), L2 (colon, 16/52, 30.8%), and L1 (terminal ileum, 3/52, 5.8%).

Perianal lesions were found in 59.6% of the

patients. Of the lesions, the skin tag was the most common (21.2%, 11/52), followed by fistula (9.6%, 5/52). Overall, a mixed-type lesion was noted in 21.2% (11/52) of patients.

According to PCDAI, severe activity was the most common (51.9%, 27/52). Moderate activity was found in 32.7% (17/52) and mild activity in 15.4% (8/52), and remission did not occur in any patients. *H. pylori* test was positive in 3 patients (5.8%).

The mean CRP value at the time of upper GIT endoscopy was 9.7 ± 19.6 mg/dL and the mean ESR level was 48.5 ± 28.4 mm/h (Table 1).

GI symptoms

The most common GI symptom was diarrhea (63.5%, 33/52), followed by lower abdominal pain (61.5%, 32/52) and weight loss (42.3%, 22/52). Hematochezia (21.2%, 11/52) and poor oral intake (17.3%, 9/52) were rare relative to other symptoms.

Upper GI symptoms were noted in 30.8% (16/52). Of those, nausea was the most common symptom (25.0%, 13/52), followed by epigastric pain (19.2%, 10/52) and vomiting (7.7%, 4/52). There were no complaints of epigastric fullness, belching, or burning sensation (Table 2).

Upper GI endoscopic and pathologic findings

The upper GI endoscopic findings were examined

Table 2. Gastrointestinal Symptoms of Pediatric Crohn's Disease (n=52)

Symptom	Case
Diarrhea	33 (63.5)
Low Abdominal pain	32 (61.5)
Weight loss	22 (42.3)
Hematochezia	11 (21.2)
Poor oral intake	9 (17.3)
Upper gastrointestinal symptom	16 (30.8)
Nausea	13 (25.0)
Epigastric pain	10 (19.2)
Vomiting	4 (7.7)
Dyspepsia	1 (1.9)
Melena	1 (1.9)
No symptom	1 (1.9)

Values are presented as number (%).

according to the following sites: esophagus, stomach, and duodenum.

Gastric ulcer was the most common (19.2%) lesion found on upper GIT endoscopy, followed by duodenal ulcers (15.4%). In the group with upper GI symptoms, esophageal ulcer was observed in 12.5% (2/16) of patients, while in the group without upper GI symptoms, esophageal erosion was noted in 11.1% (4/36) of patients. The difference was not statistically significant.

Regarding gastric findings, gastric ulcer was the most common (19.2%, 10/52). In the group with upper GI symptoms, gastric erosion was noted in 25.0% (4/16), while in the group without GI symptoms, gastric ulcer was observed in 19.4% (7/36) of patients. The difference was not statistically significant.

Regarding duodenal findings, duodenal ulcer was the most common (15.4%, 8/52). There was no difference between the groups with and without upper

GI symptoms.

Chronic inflammation was most common histopathologic feature (n=39, 75.0%) followed by gastric erosion (n=9, 17.3%). Chronic inflammation was the most common both in the group with (81.3%, 13/16) and without (72.2%, 26/36) upper GI symptoms. In 5 patients (9.6%), granuloma had previously been identified from biopsy specimens. In addition, erosion, ulceration, gastric intestinal metaplasia, etc. showed non-significant differences based on the presence or absence of upper GI symptoms (Table 3).

Differences between patients with and without upper GI symptoms

In the group with upper GI symptoms, the mean age at CD diagnosis was 14.2±1.7 years and 43.8% (7/16) were female. In the group without upper GI symptoms, the mean age at CD diagnosis was 14.1±2.3 years and 30.6% (11/36) were female. No

Table 3. Endoscopic Findings between Pediatric Crohn’s Disease Patients with and without Upper Gastrointestinal Symptoms

	Total (n=52)	UGI symptoms (n=16)	No UGI symptoms (n=36)	p-value
Endoscopic findings				
Esophagus				
Ulcer	2 (3.8)	2 (12.5)	0 (0)	0.090*
Erosion	4 (7.7)	0 (0)	4 (11.1)	0.299*
Stomach				
Erosions	9 (17.3)	4 (25.0)	5 (13.9)	0.431*
Ulcers	10 (19.2)	3 (18.8)	7 (19.4)	1.000*
Aphthous	1 (1.9)	1 (6.3)	0 (0)	0.308*
Stricture	1 (1.9)	0 (0)	1 (2.8)	1.000*
Duodenum				
Erosions	3 (5.8)	1 (6.3)	2 (5.6)	0.578*
Ulcers	8 (15.4)	4 (25.0)	4 (11.1)	0.231*
Aphthous	2 (3.8)	1 (6.3)	1 (2.8)	0.525*
Stricture	2 (3.8)	1 (6.3)	1 (2.8)	0.525*
Histologic findings				
Chronic inflammation	39 (75.0)	13 (81.3)	26 (72.2)	0.730*
Erosion	9 (17.3)	4 (25.0)	5 (13.9)	0.431*
Ulceration	4 (7.7)	3 (18.8)	1 (2.8)	0.081*
Granuloma	5 (9.6)	3 (18.8)	2 (5.6)	0.163*
Gastric intestinal metaplasia	3 (5.8)	2 (12.5)	1 (2.8)	0.221*
Normal	5 (9.6)	0 (0)	5 (13.9)	1.000*

Values are presented as number (%).

UGI: upper gastrointestinal.

*Fisher’s exact test.

between-group differences were statistically significant.

When the association between upper GI symptoms and *H. pylori* infection, CD medications, PCDAI, perianal lesions, etc., was analyzed, no variables showed an association with upper GI symptoms except for the mean values of ESR (60.7 ± 27.1 vs. 43.0 ± 27.6 mm/h, $p=0.037$) and CRP (16.5 ± 28.2 vs. 6.62 ± 13.4 mg/dL, $p=0.014$) (Table 1).

DISCUSSION

Upper GI symptoms were not correlated with upper GI lesions, and *H. pylori* infections were relatively uncommon in Korean pediatric CD. However, the extent of inflammation suggested the presence of upper GI symptoms. These results can aid the establishment of guidelines for upper GIT examination.

According to studies involving adults in Western countries, the prevalence of upper GI CD ranges from 16-34% [2,3]. However, the correlation between upper GI symptoms and true endoscopically and pathologically proven disease has not been fully established, even in pediatric patients. In the present study, upper GI involvement in pediatric CD was 50.0% (26/52), which is higher compared to results in Western countries. We believe the differences are due to dissimilar subject baseline characteristics such as age, race, and disease severity, and differences in diagnostic criteria.

A previous study involving adult Korean patients with CD reported that 59.6% demonstrated upper GI symptoms [7]. In the present study, 16 (30.8%) patients showed upper GI symptoms, which is consistent with other studies, and there was no significant difference based on the presence or absence of symptoms. Concerning individual symptoms, nausea (25.0%) was the most common, followed by epigastric pain (19.2%), vomiting (7.7%), dyspepsia (1.9%), and melena (1.9%).

Regarding endoscopic findings, gastric ulcer was the most common (19.2%) lesion found on upper GIT endoscopy, followed by duodenal ulcers (15.4%). In the presence of upper GI symptoms, gastric ero-

sion (25.0%) and duodenal ulcer (25.0%) were the most common, and gastric ulcer (18.8%) was the most common in the absence of upper GI symptoms. The difference in endoscopic findings based on the presence and absence of upper GI symptoms was not statistically significant. A study involving adults found gastric erosion in 9.6%, duodenal ulcer in 5.3%, duodenal erythema or erosion in 5.1% [2]. There were fewer abnormal findings in that study compared to the present one [2]. It is speculated that the discrepancy may be due to inter-observer differences and the presence or absence of overlapping findings.

According to a meta-analysis conducted on studies with adult patients with CD in Western countries, nonspecific gastric inflammation was the most common histological finding (32%) [2]. Gastric granuloma was noted in 7.9% and focal gastritis in 30.9% [2]. Regarding inflammatory findings by site, 84% occurred in the stomach, 28.2% in the duodenum, and 23.2% in the gastric granuloma [2]. In the present study, the most common histological finding in all pediatric patients with CD, regardless of the anatomical site, was chronic inflammation (75.0%), followed by erosion (17.3%). Granuloma was observed in 9.6% of patients, which was higher than in adults. Moreover, there was no association between the presence or absence of upper GI symptoms and histological findings (Table 3).

In Western countries, one study reported that in small bowel or colonic surgical specimens of patients with CD, perianal fistula increased significantly in the presence of noncaseating granuloma [16]. Other studies reported that gastric granuloma was found in 5-83% of gastric biopsy specimens from patients with CD and suggested an association between gastric noncaseating granuloma and perianal abscess/fistula [17,18]. In the present study of pediatric patients, gastric granuloma occurred in 9.6% (5 patients) and perianal abscess/fistula co-occurred in none of the 5 patients with gastric granuloma. We believe this finding shows that in pediatric patients with CD, unlike in adult patients, there is no association between gastric granuloma and perianal ab-

cess/fistula. This finding should be clarified with studies including a larger number of cases.

H. pylori infections are acquired early in life by young children and adolescents. In one study involving South Korea children, the prevalence of *H. pylori* infection was 22% [19]. Another study reported that *H. pylori* infections were found in 7.4% of South Korean children with recurrent abdominal pain [20]. A previous study reported a lower *H. pylori* infection rate in patients with CD than patients without CD [21]. Additionally, another study based on a meta-analysis showed a significant negative association between *H. pylori* infection and irritable bowel syndrome (IBD) and suggested a possible protective effect of *H. pylori* infection against IBD [22]. In the present study on pediatric patients with CD, *H. pylori* infection was identified in 5.8% of patients, which was lower than in pediatric patients without CD. We believe this finding shows that in pediatric patients with CD, there is negative association between *H. pylori* infection and pediatric patients with CD in South Korea. This finding should be clarified with studies including a larger number of cases.

CD is commonly complicated by perianal manifestations. In studies of adult patients with CD in Western countries, the reported incidence of perianal CD varied from 3.8% [23] to 80% [24]. In a study involving children [25], perianal disease co-occurred in 21% of pediatric patients with CD and was also more common among black people (26%) compared to white people (20%, $p=0.017$). In the present study, 59.6% (31/52) of the pediatric patients showed more than one perianal lesion such as skin tag, fissure, fistula, and abscess. It seems that racial characteristics are important, suggesting that if an Asian pediatric patient with CD presents with a perianal lesion, more attention should be paid to the CD.

According to previous research, increased CRP in adult patients with CD increases the risk of CD-related hospitalization and CD-related intestinal resection. CRP levels >1 mg/dL have been found to be correlated with granulomatous CD in pediatric patients in the United States [26-29]. In the present study, we found that ESR and CRP significantly in-

creased in the presence of upper GI symptoms. CRP testing was conducted at slightly different time points across different studies, but care should be taken if CRP and ESR are elevated in a patient with CD even when there is no difference in PCDAI or histological or endoscopic findings.

According to the European Society of Pediatric Gastroenterology, Hepatology and Nutrition (ESPGHN) guidelines, upper GIT endoscopy and ileocolonoscopy are recommended in all patients with pediatric-onset inflammatory bowel disease [30]. However, there is no such guideline in Korea. Although endoscopically abnormal findings are rare in patients with upper GI symptoms, endoscopic or histologic abnormalities are relatively common according to the findings of routinely performed endoscopy. These results suggest that upper GIT endoscopy should be performed on pediatric patients with CD in South Korea.

To our knowledge, this is the first retrospective multicenter study in Korea aimed at evaluating the prevalence of upper GIT involvement in pediatric patients with CD, irrespective of upper GI symptoms. In the present series, 50% of the patients showed upper GI CD involvement, a higher value than expected.

The present study had the following limitations. First, it was based on a retrospective design and second, the study sample was small. Third, we could not review the pathology again and simply check the pathology reports.

In conclusion, upper GI symptoms may not be correlated with upper GIT lesions. *H. pylori* infection was relatively uncommon in Korean pediatric patients with CD. However, the extent of inflammation suggests the presence of upper GI symptoms. These results should aid the establishment of regional guidelines for upper GIT examination.

REFERENCES

1. Van Limbergen J, Russell RK, Drummond HE, Aldhous MC, Round NK, Nimmo ER, et al. Definition of phenotypic characteristics of childhood-onset inflammatory bowel disease. *Gastroenterology* 2008;135:1114-22.
2. Diaz L, Hernandez-Oquet RE, Deshpande AR, Moshiree

- B. Upper gastrointestinal involvement in Crohn disease: histopathologic and endoscopic findings. *South Med J* 2015;108:695-700.
3. Annunziata ML, Caviglia R, Papparella LG, Cicala M. Upper gastrointestinal involvement of Crohn's disease: a prospective study on the role of upper endoscopy in the diagnostic work-up. *Dig Dis Sci* 2012;57:1618-23.
 4. Lenaerts C, Roy CC, Vaillancourt M, Weber AM, Morin CL, Seidman E. High incidence of upper gastrointestinal tract involvement in children with Crohn disease. *Pediatrics* 1989;83:777-81.
 5. Tobin JM, Sinha B, Ramani P, Saleh AR, Murphy MS. Upper gastrointestinal mucosal disease in pediatric Crohn disease and ulcerative colitis: a blinded, controlled study. *J Pediatr Gastroenterol Nutr* 2001;32:443-8.
 6. Ammourey RF, Pfefferkorn MD. Significance of esophageal Crohn disease in children. *J Pediatr Gastroenterol Nutr* 2011;52:291-4.
 7. So H, Ye BD, Park YS, Kim J, Kim JS, Moon W, et al. Gastric lesions in patients with Crohn's disease in Korea: a multicenter study. *Intest Res* 2016;14:60-8.
 8. Yang SK, Yun S, Kim JH, Park JY, Kim HY, Kim YH, et al. Epidemiology of inflammatory bowel disease in the Songpa-Kangdong district, Seoul, Korea, 1986-2005: a KASID study. *Inflamm Bowel Dis* 2008;14:542-9.
 9. Thia KT, Loftus EV Jr, Sandborn WJ, Yang SK. An update on the epidemiology of inflammatory bowel disease in Asia. *Am J Gastroenterol* 2008;103:3167-82.
 10. Park SH, Yang SK, Park SK, Kim JW, Yang DH, Jung KW, et al. Long-term prognosis of Crohn's disease and its temporal change between 1981 and 2012: a hospital-based cohort study from Korea. *Inflamm Bowel Dis* 2014;20:488-94.
 11. Loftus EV Jr, Silverstein MD, Sandborn WJ, Tremaine WJ, Harmsen WS, Zinsmeister AR. Crohn's disease in Olmsted County, Minnesota, 1940-1993: incidence, prevalence, and survival. *Gastroenterology* 1998;114:1161-8.
 12. Hyams J, Markowitz J, Otley A, Rosh J, Mack D, Bousvaros A, et al. Evaluation of the pediatric Crohn disease activity index: a prospective multicenter experience. *J Pediatr Gastroenterol Nutr* 2005;41:416-21.
 13. Koletzko S, Jones NL, Goodman KJ, Gold B, Rowland M, Cadranel S, et al. Evidence-based guidelines from ESPGHAN and NASPGHAN for *Helicobacter pylori* infection in children. *J Pediatr Gastroenterol Nutr* 2011;53:230-43.
 14. De Felice KM, Katzka DA, Raffals LE. Crohn's disease of the esophagus: clinical features and treatment outcomes in the biologic era. *Inflamm Bowel Dis* 2015;21:2106-13.
 15. Sonnenberg A, Melton SD, Genta RM. Frequent occurrence of gastritis and duodenitis in patients with inflammatory bowel disease. *Inflamm Bowel Dis* 2011;17:39-44.
 16. Denoya P, Canedo J, Berho M, Allende DS, Bennett AE, Rosen L, et al. Granulomas in Crohn's disease: does progression through the bowel layers affect presentation or predict recurrence? *Colorectal Dis* 2011;13:1142-7.
 17. Wagtmans MJ, van Hogezaand RA, Griffioen G, Verspaget HW, Lamers CB. Crohn's disease of the upper gastrointestinal tract. *Neth J Med* 1997;50:S2-7.
 18. Kefalas CH. Gastroduodenal Crohn's disease. *Proc (Bayl Univ Med Cent)* 2003;16:147-51.
 19. Malaty HM, Kim JG, Kim SD, Graham DY. Prevalence of *Helicobacter pylori* infection in Korean children: inverse relation to socioeconomic status despite a uniformly high prevalence in adults. *Am J Epidemiol* 1996;143:257-62.
 20. Jang KM, Choe BH, Choe JY, Hong SJ, Park HJ, Chu MA, et al. Changing prevalence of *Helicobacter pylori* infections in Korean children with recurrent abdominal pain. *Pediatr Gastroenterol Hepatol Nutr* 2015;18:10-6.
 21. El-Omar E, Penman I, Cruikshank G, Dover S, Banerjee S, Williams C, et al. Low prevalence of *Helicobacter pylori* in inflammatory bowel disease: association with sulphasalazine. *Gut* 1994;35:1385-8.
 22. Rokkas T, Gisbert JP, Niv Y, O'Morain C. The association between *Helicobacter pylori* infection and inflammatory bowel disease based on meta-analysis. *United European Gastroenterol J* 2015;3:539-50.
 23. Sangwan YP, Schoetz DJ Jr, Murray JJ, Roberts PL, Collier JA. Perianal Crohn's disease. Results of local surgical treatment. *Dis Colon Rectum* 1996;39:529-35.
 24. McClane SJ, Rombeau JL. Anorectal Crohn's disease. *Surg Clin North Am* 2001;81:169-83, ix.
 25. Adler J, Dong S, Eder SJ, Dombkowski KJ; ImproveCareNow Pediatric IBD Learning Health System. Perianal Crohn disease in a large multicenter pediatric collaborative. *J Pediatr Gastroenterol Nutr* 2017;64:e117-24.
 26. Vaiopoulou A, Gazouli M, Papadopoulou A, Anagnostopoulos AK, Karamanolis G, Theodoropoulos GE, et al. Serum protein profiling of adults and children with Crohn disease. *J Pediatr Gastroenterol Nutr* 2015;60:42-7.
 27. Kellermayer R, Mir SA, Nagy-Szakal D, Cox SB, Dowd SE, Kaplan JL, et al. Microbiota separation and C-re-

- active protein elevation in treatment-naïve pediatric granulomatous Crohn disease. *J Pediatr Gastroenterol Nutr* 2012;55:243-50.
28. Motil KJ, Grand RJ, Maletskos CJ, Young VR. The effect of disease, drug, and diet on whole body protein metabolism in adolescents with Crohn disease and growth failure. *J Pediatr* 1982;101:345-51.
29. Oh K, Oh EH, Baek S, Song EM, Kim GU, Seo M, et al. Elevated C-reactive protein level during clinical remission can predict poor outcomes in patients with Crohn's disease. *PLoS One* 2017;12:e0179266.
30. Levine A, Koletzko S, Turner D, Escher JC, Cucchiara S, de Ridder L, et al. ESPGHAN revised porto criteria for the diagnosis of inflammatory bowel disease in children and adolescents. *J Pediatr Gastroenterol Nutr* 2014;58:795-806.