Appendiceal Crohn’s disease clinically presenting as acute appendicitis

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Abstract

AIM: To determine the incidence of appendiceal Crohn’s disease (CD) and to summarize the characteristic histologic features of appendiceal CD.

METHODS: We reviewed the pathology files of 2179 appendectomy specimens from January 2007 to May 2013. The computer-assisted retrieval search facility was utilized to collect specimens. We selected those cases that were diagnosed as CD or chronic granulomatous inflammation and defined the final diagnosis according to the histologic findings of CD, including transmural lymphocytic inflammation, non-caseating epithelioid granulomas, thickening of the appendiceal wall secondary to hypertrophy of muscularis mucosa, mucosal ulceration with crypt abscesses, mucosal fissures, and fistula formation.

RESULTS: We found 12 cases (7 male and 5 female patients, with an average age of 29.8 years) of appendiceal CD. The incidence of appendiceal CD was 0.55%. The chief complaints were right lower quadrant pain, abdominal pain, lower abdominal pain, and diarrhea. The duration of symptom varied from 2 d to 5 mo. The histologic review revealed appendiceal wall thickening in 11 cases (92%), transmural inflammation in all cases (100%), lymphoid aggregates in all cases (100%), epithelioid granulomas in all cases (100%), mucosal ulceration in 11 cases (92%), crypt abscesses in 5 cases (42%), perforation in 2 cases (17%), muscular hypertrophy in 1 case (8%), neural hyperplasia in 5 cases (42%), and perpendicularly serosal fibrosis in 8 cases (67%).

CONCLUSION: A typical and protracted clinical course, unusual gross features of the appendix and the characteristic histologic features are a clue in the diagnosis of appendiceal CD.

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Key words: Appendix; Appendectomy; Acute appendicitis; Crohn’s disease; Prognosis

Core tip: Appendiceal Crohn’s disease (CD) is relatively rare and is indistinguishable from acute appendicitis. Appendiceal CD shows a favorable clinical outcome with a low recurrence rate. The differential diagnosis includes intestinal tuberculosis, foreign body reaction, diverticulitis of the appendix, sarcoidosis, actinomycosis, and Yersinia infection. Atypical and protracted clinical course, unusual gross features of the appendix and the characteristic histologic features are a clue in the diagnosis of appendiceal CD.


INTRODUCTION

Crohn’s disease (CD) is a chronic inflammatory bowel...
Appendiceal Crohn’s disease is a rare disorder characterized by a transmural inflammatory reaction and non-caseating small granulomas and may involve all parts of the gastrointestinal (GI) tract from the mouth to the anus[1-7]. The most common sites of involvement are the ileum and colon[8]. Appendiceal CD is a rare disease but has been well summarized in the various reports[9-14]. The incidence of appendicitis with granulomatous reaction varies from 0.1% to 2.0%[15]. Since Meyerd ing et al[16] had reported an interesting case of appendiceal CD without demonstrable involvement of the adjacent GI tract in 1953, many additional cases of appendiceal CD have been demonstrated in the literature to date.

The purpose of this retrospective review study was to determine the exact incidence of appendiceal CD in patients who underwent appendectomy and to summarize the common characteristic histologic findings along with a review of the literature.

**MATERIALS AND METHODS**

**Ethics**

The materials used in our study are human appendix tissue samples, which are products of surgical procedures. Our study contains no private information relating to the patients, and so ensures their anonymity. Therefore, our study has no problems in causing any ethical issue or encroachment of human rights.

**Patient tissue**

A retrospective review of 2179 appendectomy specimens from January 2007 to May 2013 was conducted. All patients underwent appendectomy at the Hanyang University Hospital (Seoul, South Korea). The computer-assisted retrieval search facility was utilized to collect appendectomy specimens. Appendices resected for acute appendicitis and those removed as a part of right hemicolectomy and gynecology procedures were collected and reviewed. We selected those cases that were diagnosed as CD or chronic granulomatous inflammation and defined the final diagnosis according to the common histologic findings of CD, including transmural lymphocytic inflammation, non-caseating small epithelioid granulomas, thickening of the appendiceal wall secondary to hypertrophy of muscularis mucosa, mucosal ulceration with crypt abscesses, mucosal fissures, and fistula formation. No evidence of parasitic, fungal and mycobacterial disease, foreign body, or systemic sarcoidosis was found in any patient. The clinical information including age, gender, clinical data, and data about the surgical procedure for each case as well as follow-up data including colonoscopic evaluation was collected. The special staining technique such as Ziehl-Neelsen staining and special molecular technique such as tuberculosis polymerase chain reaction (Tb-PCR) were performed to rule out *Mycobacterium tuberculosis*.

**RESULTS**

Out of these 2179 appendectomy specimens, 12 cases (0.55%) were classified as appendiceal CD. The clinicopathologic characteristics of the appendiceal CD patients are summarized in Tables 1 and 2. Out of these 12 patients, there were 7 male and 5 female patients. The age of patients ranged from 11 to 51 years (average age of 29.8 years). The chief complaints of patients were right lower quadrant (RLQ) pain, abdominal pain, lower abdominal pain, and diarrhea. The duration of symptom with which patients presented varied from 2 d to 5 mo. There was no systemic clinical manifestation such as arthralgia, uveitis, or arthritis. No history of tuberculosis of any organ was found in these patients. There was also no clinical evidence of systemic sarcoidosis. The initial clinical impression was acute appendicitis in all of these 12 patients along with perforation in 2 among these 12 patients. All patients underwent appendectomy. The final pathologic report was CD in all of these 12 cases. All cases showed a negative result for *Mycobacterium tuberculosis* in Ziehl-Neelsen staining and Tb-PCR. The histologic review of these 12 cases revealed appendiceal wall thickening in 11 cases (92%), transmural inflammation in all cases (100%), lymphoid aggregates in all cases (100%), epithelioid granulomas in all cases (100%), mucosal ulceration in 11 cases (92%), crypt abscesses in 5 cases (42%), perforation with abscess formation in 2 cases (17%), muscular hypertrophy in 1 case (8%), neural hyperplasia in 5 cases (42%), and perpendicularly serosal fibrosis in 8 cases (67%). The representative microphotographs are shown in Figure 1. There is no evidence of disease recurrence in these 12 patients to date.

**DISCUSSION**

Crohn first described that CD stops at the ileocecal valve with sparing of the colon and appendix. However, this theory was disproved as patients with CD often have involvement of the colon and appendix[8]. The first isolated appendiceal CD was reported by Meyerd ing et al[16] in 1953. Since Meyerd ing et al[16] had reported a case of CD arising in the appendix, many case reports and some collective reviews have been reported in the literature. The incidence of appendiceal CD is variable[17-21]. Prieto-Nieto *et al* described that approximately 0.2% of patients (10 out of 4468 appendectomies performed during 20 years) had appendiceal CD. In our review, 12 cases (0.55%) out of 2179 appendectomy specimens were revealed as appendiceal CD.

Appendiceal CD is usually found among young patients, however, it can occur at any age[3,12]. Yang *et al* described the age with onset of disease in 14 patients with appendiceal CD, ranged from 10 to 45 years (average age of 21.1 years). Prieto-Nieto *et al* reported the disease onset-age in 10 patients with appendiceal CD, ranged from 10 to 33 years (average age of 29 years). The difference in incidence of disease in males and females has been reported, with male predominance[14,16]. In our study, the age ranged from 11 to 51 years, with an average age of 29.8 years. Among 12 patients, 7 were male, reflecting more male patients with the disease described.
The clinical presentation of appendiceal CD is variable. The most common presenting symptom is acute lower abdominal pain especially in the RLQ, which is very similar to the lower abdominal pain presented in patients with acute appendicitis\(^4,13,22\). Approximately 25% of appendiceal CD patients show chronic abdominal pain in the right lower abdomen\(^13\). The symptoms may be more protracted or recurrent than in the usual case of acute suppurative appendicitis. Appendiceal CD should be suspected when the patients show atypical or protracted clinical course\(^2,13\). In our study, most patients presented with the pain in the RLQ. The initial clinical impression was acute appendicitis in all 12 patients. Most patients had symptoms for two or more days, and 8 patients (67%) presented with these symptoms for over a week.

Appendiceal CD usually shows an enlarged appendix with marked thickening of the appendiceal wall and fibrous adhesion to the periappendiceal soft tissue\(^2,22,23\). Microscopically, the histologic features are characterized by transmural chronic inflammation with marked fibrous thickening of the wall, lymphoid aggregates, small non-caseating granulomas, ulcerative mucosal change, crypt abscesses, muscular hypertrophy, and neural hyperplasia\(^13,24-26\). In our study, the features were similar to the

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**Figure 1** Appendiceal Crohn’s disease. A: The appendix with Crohn’s disease shows transmural inflammation with markedly thickened wall; B: There is a prominent lymphoid hyperplasia in the mucosa and serosa; C: The mucosa shows many small non-caseating granulomas; D: The serosa shows creeping fat with perpendicular thick fibrous bands.

**Table 1** Summary of the patients with appendiceal Crohn’s disease

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Sex</th>
<th>Age (yr)</th>
<th>c/c</th>
<th>SD</th>
<th>Clinical impressions</th>
<th>AFB</th>
<th>Tb-PCR</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>30</td>
<td>RLQ pain</td>
<td>None</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>2</td>
<td>M</td>
<td>38</td>
<td>RLQ pain</td>
<td>14 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>3</td>
<td>M</td>
<td>26</td>
<td>RLQ pain</td>
<td>None</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>28</td>
<td>RLQ pain</td>
<td>7 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>25</td>
<td>RLQ pain</td>
<td>14 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>6</td>
<td>F</td>
<td>29</td>
<td>RLQ pain</td>
<td>7 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>7</td>
<td>M</td>
<td>51</td>
<td>RLQ pain</td>
<td>5 mo</td>
<td>Acute appendicitis, perforation</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>8</td>
<td>F</td>
<td>49</td>
<td>RLQ pain</td>
<td>8 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>9</td>
<td>M</td>
<td>30</td>
<td>Abdominal pain</td>
<td>10 d</td>
<td>Acute appendicitis, perforation</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>10</td>
<td>M</td>
<td>23</td>
<td>Lower abdominal pain</td>
<td>2 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>11</td>
<td>M</td>
<td>18</td>
<td>Lower abdominal pain, diarrhea</td>
<td>3 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>12</td>
<td>F</td>
<td>11</td>
<td>RLQ pain</td>
<td>7 d</td>
<td>Acute appendicitis</td>
<td>Negative</td>
<td>Negative</td>
</tr>
</tbody>
</table>

F: Female; M: Male; c/c: Chief complaint; RLQ: Right lower quadrant; SD: Symptom duration; AFB: Acid-fast bacillus; Tb-PCR: Tuberculosis polymerase chain reaction.
The incidence of appendiceal CD is variable. The purpose of this study was to determine the exact incidence of appendiceal CD in patients who underwent appendectomy and to summarize the common characteristic histologic findings along with a review of the literature.

Innovations and breakthroughs
The histologic features are characterized by transmural chronic inflammation, with marked fibrous thickening of the wall, lymphoid aggregates, small non-caseating granulomas, ulcerative mucosal change, crypt abscesses, muscular hypertrophy, and neural hyperplasia. In this study, the features were similar to the previously described histologic characteristics. However, the authors found that appendiceal CD had the characteristic perpendicularly fibrous band formation in 8 out of 12 cases.

Applications
With the characteristic clinical presentation and the typical pathologic findings, the clinicians and pathologists can consider the possibility of appendiceal CD. Atypical and protracted clinical course, unusual gross features of the appendix and the characteristic histologic features are a clue in the diagnosis of appendiceal CD.

Terminology
CD is a chronic inflammatory bowel disorder characterized by a transmural inflammatory reaction and non-caseating small granulomas and may involve all parts of the gastrointestinal tract from the mouth to the anus.

Peer review
The authors described appendiceal CD clinically presenting as acute appendicitis. This is an interesting review and CD in appendix is a rare condition. Whenever it is encountered, the surgeon must know what to do and be aware of its prognosis. This paper will lead surgeons to this condition.

REFERENCES

11. Agha FP, Chahremani GG, Panella JS, Kaufman MW. Appendicitis as the initial manifestation of Crohn’s disease.

**Table 2** Summary of histologic features of appendiceal Crohn’s disease

<table>
<thead>
<tr>
<th>Histologic features</th>
<th>Number of cases</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wall thickening</td>
<td>11/12</td>
<td>92</td>
</tr>
<tr>
<td>Transmural inflammation</td>
<td>12/12</td>
<td>100</td>
</tr>
<tr>
<td>Lymphoid aggregates</td>
<td>12/12</td>
<td>100</td>
</tr>
<tr>
<td>Epithelioid granulomas</td>
<td>12/12</td>
<td>100</td>
</tr>
<tr>
<td>Mucosal ulceration</td>
<td>11/12</td>
<td>92</td>
</tr>
<tr>
<td>Crypt abscess</td>
<td>5/12</td>
<td>42</td>
</tr>
<tr>
<td>Perforation</td>
<td>2/12</td>
<td>17</td>
</tr>
<tr>
<td>Muscular hypertrophy</td>
<td>1/12</td>
<td>8</td>
</tr>
<tr>
<td>Neural hyperplasia</td>
<td>5/12</td>
<td>42</td>
</tr>
<tr>
<td>Perpendicular serosal fibrosis</td>
<td>8/12</td>
<td>67</td>
</tr>
</tbody>
</table>

COMMENTS

Background
Appendiceal Crohn’s disease (CD) is a rare disease. Since MeyeRd et al reported an interesting case of appendiceal CD without demonstrable involvement of the adjacent gastrointestinal tract in 1953, many additional cases of appendiceal CD have been demonstrated in the literature to date.

Research frontiers
The incidence of appendicitis with granulomatous reaction varies from 0.1% to 2.0%. The incidence of appendiceal CD is variable. The purpose of this study was to determine the exact incidence of appendiceal CD in patients who underwent appendectomy and to summarize the common characteristic histologic findings along with a review of the literature.

Table 2 Summary of histologic features of appendiceal Crohn’s disease

The differential diagnosis includes intestinal tuberculosis, foreign body reaction, diverticulitis of the appendix, sarcoidosis, actinomycosis, and *Yersinia* infection

Appendiceal tuberculosis results in the formation of epithelioid granulomas, however, the granulomas in tuberculosis are larger with a central caseous necrosis and less discrete than those in Crohn’s disease. If a foreign body is present, histologic examination should reveal the offending material and diverticulitis disease may be excluded via careful examination. Intestinal sarcoidosis is extremely rare and does not occur as an isolated finding. Actinomycosis also results in a vague granulomatous tissue reaction, however, actinomycosis shows neutrophilic abscess formation with floating bacterial colonies (sulphur granules). *Yersinia* infection results in necrotizing granulomatous reaction in the appendiceal mucosa or submucosa and shows microabscess formation.

The treatment of choice for appendiceal CD is appendectomy. Appendiceal CD shows lower recurrence rate compared with CD arising in other parts of the intestine. The prognosis of appendiceal CD seems to be much better than that of CD arising in the small or large bowel.

In conclusion, we described the incidence of appendiceal CD in patients who underwent appendectomy and summarized the common characteristic histologic findings along with a review of the literature. Atypical and protracted clinical course, unusual gross features of the appendix and the characteristic features are a clue in the diagnosis of appendiceal CD.
Radiologic features and prognosis. AJR Am J Roentgenol 1987; 149: 515-518 [PMID: 3497553 DOI: 10.2214/ajr.149.3.515]


16 Meyering EV, Bertram HF. Nonspecific granulomatous inflammation (Crohn’s disease) of the appendix; a case report. Surgery 1953; 34: 891-894 [PMID: 13113513]


21 Warren S, Sommers SC. Cicatrizing enteritis as a pathologic entity; analysis of 120 cases. Am J Pathol 1948; 24: 475-501 [PMID: 18859355]


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