

A Case of Secondary Rectal Intramural Hematoma Triggered by Aspiration for Endoscopic Mucosal Resection

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Abstract

Total colonoscopy was performed on an 89-year-old man who was not treated with anticoagulation therapy. He had developed purpura in the rectum below the peritoneal reflection by the aspiration for endoscopic mucosal resection to rectal polyp. The purpura resulted in spontaneous cessation, but he had lower abdominal pain thereafter. He was diagnosed to have intramural hematoma of the rectum from rectosigmoid to above the peritoneal reflection by computed tomography and sigmoidoscopy. It disappeared with conservative therapy. We infer that it was secondary rectal intramural hematoma triggered by excessive straining against purpura.

Keywords: Rectum; Intramural hematoma; Endoscopic mucosal resection; Abdominal pain; Straining

Abbreviations: IH: Intramural Hematoma; TCS: Total Colonoscopy; RB: Rectum below the peritoneal reflection EMR: Endoscopic Mucosal Resection, PT-INR: Prothrombin Time-International Normalized Ratio; APTT: Activated Partial Thromboplastin Time; CT: Computed Tomography; RS: Rectum from Rectosigmoid RA: Rectum above the Peritoneal Reflection

Introduction

Intramural Hematoma (IH) of intestine is a rare condition [1] that may develop from esophagus to rectum. It often occurs in patients with anticoagulation therapy [2-4], hemophilia [5], amyloidosis [6], and Ehlers-Danlos syndrome [7], and occurs rarely without coagulation disorder. We report here a case of secondary rectal IH triggered by aspiration for endoscopic mucosal resection without coagulation disorder.

Case Presentation



Figure 1: Total colonoscopy showed a polyp which was 5mm in size on the rectum below the peritoneal reflection; 39cm from the anus. Purpura was caused by the aspiration for endoscopic mucosal resection.

He was treated with medicines for high blood pressure hypertension, hyperuricemia, iron deficiency anemia, and insomnia. A cecal cancer was diagnosed in him by Total Colonoscopy (TCS) which was performed to check cause of fecal occult blood. We assessed that operation was necessary to cure the lesion. In addition, there was a semi-pedunculated polyp which

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was 5mm size on the rectum below the peritoneal Reflection (Rb) and we tried to perform Endoscopic Mucosal Resection (EMR) for it immediately. However, we interrupted to perform EMR for it because severe purpura was caused by the aspiration for injection to submucosa (Figure 1). We confirmed spontaneous cessation of hemorrhage, and he was admitted to our hospital for observation of the progress on this day. He recognized only slight tenesmus and unpleasant feeling of anus, and his general condition was stable. So that, he was discharged 2 days later.

He repeated straining excessively against these symptoms after TCS. The lower abdominal pain developed from the next day after discharge, and It rapidly got worse. Therefore, he visited our hospital again 4 days after TCS. On physical examination, blood pressure was 134/72mmHg, pulse was 66 beats per minute/regular, and temperature was 37.1 degrees. The auscultatory findings of lungs and heart were normal. The abdomen was soft-flat, bowel sound was normal, and there was no tenderness. In laboratory data, hemoglobin level was 12.0g/dL which was 12.3g/dL before TCS, and there was no appearance of anemia was. The platelet count was 19.2×104/uL, prothrombin time-international normalized

ratio (PT-INR) was 1.02 and Activated Partial Thromboplastin Time (APTT) was 26.9 seconds, all of which were normal.

The abdominal plain Computed Tomography (CT) showed a large highly absorbed mass in the rectum from Rectosigmoid (RS) to above the peritoneal Reflection (Ra) (Figure 2a). There were no definitive findings to suggest active bleeding with contrastenhanced CT (Figure 2b). Sigmoidoscopy demonstrated a large dark-red mass in the rectum from RS to Ra (Figure 3a). A rectal moderate stenosis was caused by the mass, endoscope passed there easily. On the anal side, there was a polyp which was interrupted to perform EMR the previous time. Purpura in the polyp and the mucosa around it was improved (Figure 3b). We diagnosed rectal IH. Since his general condition was stable and IH was examined as spontaneous cessation with CT, we decided to treat conservatively rather than surgically. He was admitted and received conservative therapy which were bowel rest and intravenous hydration. His symptoms improved in several days. We confirmed that IH did not grow with abdominal CT performed 7 days after admission. He was discharged 13 days later.

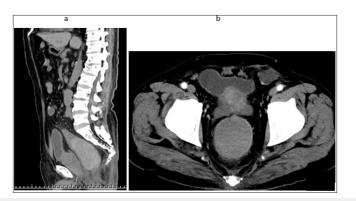


Figure 2:

(a): Sagittal section of abdominal plain computed tomography showed a large highly absorbed mass (white arrows) which was 9*9*5cm in size in the rectum from rectosigmoid to above the peritoneal reflection.

(b): Axial section of abdominal contrast-enhanced computed tomography showed no definitive findings in it (white arrow heads).

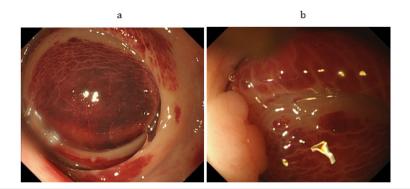


Figure 3:

(a): Sigmoidoscopy showed a large dark-red mass in the rectum above the peritoneal reflection.

(b): On the anal side, sigmoidoscopy showed a polyp (white arrows) which was interrupted to perform endoscopic mucosal resection in the previous time. Purpura in the polyp and the mucosa around it was improved.

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About two months later, sigmoidoscopy demonstrated that rectal IH disappeared (Figure 4). Cecal cancer operation was performed and examined as stage I/radical excision by histological diagnosis. Now, four years later, rectal IH and cecal cancer have not relapsed.

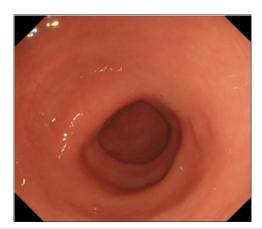


Figure 4: Sigmoidoscopy demonstrated that the rectal mass disappeared about two months later.

Discussion

IH of intestine often occurs in small intestine and duodenum, whereas colonic IH is rare [8,9]. Its typical symptoms are abdominal pain and nausea/vomiting [1,9]. It is diagnosed by endoscopy, abdominal ultrasonography, and CT [1,3,10]. In addition, there is a report that endoscopic ultrasonography was useful for diagnosis of colonic IH [11]. However, it is often difficult to distinguish IH from neoplasm [12]. In this case, we confirmed that there was no rectal IH by TCS several days earlier. It was relatively easy to diagnose rectal mass with IH from his history and sigmoidoscopy/CT findings. The standard treatment for IH of intestine is conservative therapy if general condition is stable [3,4,10,13], but we should consider operation when it is accompanied by intestinal obstruction and peritonitis [1,2].

The major cause of IH is abdominal blunt trauma [3,8,10]. In addition, it is often caused by coagulation disorder [1-5]. However, this case did not have these factors. There is a report of a colonic IH associated with colon cancer [14], and our case had a cecal cancer. However, we think that it was unrelated among both for our case because there was distance from each other, and it was stage I/radical excision lesion. There were some cases of colonic IH which were caused by endoscopic procedures such as polypectomy [11,15], and tattooing [16]. Because severe purpura was caused by aspiration for EMR several days before rectal IH occurred, we thought that there was a relationship between both in this case. However, purpura by aspiration was located in Rb whereas IH was located in RS-Ra. In addition, purpura improved when IH appeared. Therefore, it was thought that aspiration did not influence occurrence of IH directly. A non-traumatic rectal IH caused by straining was reported [2], and we guessed that the mechanism in our case was also the same. In summary, purpura by aspiration for EMR gave him some symptoms such as tenesmus and unpleasant feeling of anus, and he repeated straining excessively against them. The straining increased abdominal pressure and rectal muscular

contraction. They gave rectum severe stress, and it led to secondary rectal IH. We infer that a case like this is extremely rare, but it is important to recognize that excessive straining may cause rectal IH. For preventing rectal IH, we should perform instruction and prescribe medicine to avoid excessive straining for patients with some conditions, such as colitis, constipation, and rectal purpura caused by aspiration.

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Kazu Hamada wrote the manuscript; Masashi Okuro and Ken Kawaura collected data for review; all authors read and approved the final version of the manuscript. We thank all the investigators, staff, and participants associated with this report for helpful discussions during manuscript development.

Conflict of Interest

The authors declare no conflict of interest.

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