Letter to the Editor

A rare case of acute abdomen in an adult: spontaneous Meckel’s diverticulum perforation

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Meckel’s diverticulum (MD) is the most common congenital anomaly of the gastrointestinal system. It is located approximately 60–80 cm proximal to the cecum, on the antimesenteric face of the small bowel. The average length is 4–6 cm and it is a real diverticulum. It occurs if the omphalomesenteric tract does not close between the 5th and 7th weeks of intrauterine life [1]. Meckel’s diverticulum was first described in the 16th century by Fabricus Hildanus. It was embryologically described and named by the German Johann Friedrich Meckel in 1809 [2, 3]. The rule of 2s is a useful memory aid. MD occurs in 2% of the population, is 2 inches (5 cm) long, 2 feet (60 cm) from the ileocecal valve, two thirds have ectopic mucosa, 2 types of ectopic tissue are commonly present (mostly gastric and pancreatic) and 2% become symptomatic [4–6].

Patients usually have nonspecific complaints such as nausea, vomiting, abdominal pain, and loss of appetite [7]. These findings were exacerbated after perforation and abdominal symptoms may progress. Many complications occur such as massive gastrointestinal hemorrhage [8], cecal volvulus [9], ileus [10], subphrenic abscess [11], small bowel torsion due to MD [12], and perforation, as in our case and in the literature. The mortality rate is reported to be between 1.6% and 7.7% [13]. Mortality usually occurs in delayed cases and with intestinal gangrene [7]. Here we present our case, which was diagnosed as spontaneous MD perforation due to acute abdomen, in light of the literature.

A 27-year-old male patient presented with abdominal pain, anorexia, nausea and fever to the emergency department. The patient was examined in emergency service. On physical examination, rebound tenderness and guarding was positive. The patient had an acute abdomen. There was nothing in his anamnesis. Digital rectal examination was performed. The rectum was empty and without hemorrhage. In the patient’s blood tests, white blood count (WBC) was 26.52 10⁹/UL, glucose 131 mg/dl, lactic dehydrogenase 350 U/l, creatine kinase 2742 U/l, C-reactive protein 4.89 mg/dl, sodium 135 mmol/l. The patient’s complete urinalysis included 7 erythrocytes/HPF and 3 leukocytes/HPF. Nothing was detected in the direct abdominal X-ray (Figure 1). Abdominal ultrasonography report of the patient: "Mesenteric echo is heterogeneous
There were fibrin plates together. No additional pathology was found in exploration. We decided to perform resection and anastomosis. The abdomen was washed with plenty of physiological saline solution. One drain was placed in the pelvis. After the operation the patient was referred to an infectious diseases specialist for antibiotic therapy selection. Antibiotic therapy was administered. On the third postoperative day the drain was shortened because the drain was clear. On the fourth postoperative day oral intake was started as regime 1. On the postoperative fifth day the drain was pulled and the oral intake resumed as normal. The WBC value fell to within normal limits 6 days after the operation. The patient was discharged on the 7th day. Sutures were taken out on postoperative day 15. Blood levels were completely normal. Pathological examination revealed MD.

Meckel's diverticulum is a congenital anomaly commonly seen in the gastrointestinal tract. The rate in the population is approximately 2% (1–3%) [14]. It occurs as the result of the omphalomesenteric tract not closing [1]. It is 3 times more frequent in men [15].

In intrauterine life in the early weeks, the fetus takes nutrients from the yolk sac via the vitelline duct. This tube disappears in the 7th week of preg-
nancy. If not, MD occurs [1]. Meckel's diverticulum is frequently located 40–60 cm proximal to the ileocecal valve on the antimesenteric face of the ileum. The size of the diverticulum varies within 1–10 cm, often 6 cm [16].

Hemorrhage is a frequent complication. Bleeding in the form of fresh blood in the child may be confused with invagination. In adults, melena can be seen with abdominal pain [17]. There are cases reported in the literature that cause massive gastrointestinal bleeding [8]. Bleeding is mainly due to the ectopic tissue of the diverticula. The most common is stomach tissue followed by pancreatic tissue. Tissue enzyme damage of the small intestine is responsible for this [18]. Diverticulitis is a common complication [19].

Occasionally, perforations associated with a fish follicle [19, 20], due to a blunt abdominal trauma [21] or button cell [22, 23] and spontaneous [24, 25] cases are found in the literature. Resection is recommended for patients with complicated MD during the operation, and resection for incidental cases has been discussed. The operative technique can vary depending on the patient's condition and because of perforation during the operation. The presence of symptoms is very important. The surgeon must ensure that the diverticular structure is completely removed in the operation and should not narrow the passage.

Schlicke and Johnston advised elective diverticulectomy for cases in the small intestines during laparotomy [26]. However, Cullen et al. recommend resection for all coincidentally detected MDs in patients below the age of 80 [27]. Park et al. recommended resection in symptomatic cases. Asymptomatic and uncomplicated diverticula suggest simple diverticulectomy [15].

Contrary to these ideas, Soltero and Bill reported that they rarely operated for prophylactic resection except for definite acute abdomen [28].

As a result, segmental small bowel resection or diverticulectomy may be performed to include the diverticulum as a surgical treatment. The surgical technique should be decided intraoperatively according to the patient's pathology. It should not be forgotten that there may be a complication due to MD in clinically unexplained cases.

Conflict of interest

The authors declare no conflict of interest.

References


