EDITÖRE MEKTUP / LETTER TO THE EDITOR

Post-traumatic perforated mesenteric cystic lymphangioma

Travma sonrası perfore mezenterik kistik lenfanjiyom

Mustafa Tuşat1, Mehmet Semih Demirtaş2

1Aksaray University, Faculty of Medicine, Department of Pediatric Surgery, Aksaray, Turkey
2Aksaray University Education and Research Hospital, Department of Pediatrics, Aksaray, Turkey

To the Editor,

Abdominal lymphangiomas are a rare congenital cysts of the mesenteric and/or retroperitoneal lymphatics. They may occur mainly in childhood as a result of proliferation and enlargement of lymphatic vessels due to obstruction of the lymphatic system. Although abdominal lymphangiomas are located most commonly in the small-bowel mesentery, they have been reported as less than 1% of all lymphangiomas. The clinical presentation of the cases is variable, which ranges from asymptomatic abdominal mass to acute abdomen. Although they can be identified by imaging methods such as abdomen ultrasonography, tomography or magnetic resonance imaging, the definitive diagnosis is made by histopathological examination. In this report, a case of mesenteric cystic lymphangioma which was ruptured due to trauma and caused an acute abdomen, is presented as a rare clinical presentation of MCL.

A five-year-old girl presented to pediatric emergency department with complaints of increasing abdominal pain and vomiting, which started after fell on her abdomen. The patient had a history of intermittent abdominal pain in the last one year before admission to hospital. In addition, the child was hospitalized two a year ago due to abdominal pain with a preliminary diagnosis of intussusception and was discharged after spontaneous reduction of the intussusception. The physical examination revealed a palpable mass on the periumbilical region, measuring around 10 cm, mobile, and accompanied by abdominal tenderness with guarding and scarce bowel sounds. Laboratory examination showed erythrocyte sedimentation rate of 89 mm per hour, C reactive protein 118 mg/L, leucocyte elevation of 19.3 x10⁹ /L and hemoglobin 9.1 mg/dl.

Abdominal x-ray revealed sign of ileus (Figure 1). Abdominal CT taken to the patient with a history of trauma showed two cystic lesions compressing the small intestines superiorly and left laterally in the abdominopelvic region and the larger one is 85x95 mm in size. Also, free fluid spread to all abdominal quadrants was observed (Figure 2). The patient then underwent an emergency surgery with preliminary diagnosis of intestinal duplication, mesentery or ovarian cyst perforation. In exploration, there were two cysts, tightly adherent to the jejunum, one intact and the other ruptured in the jejunal mesentery and free fluid with chylose character that spread all the quadrants. The intact cyst was a large 9x8x9 cm in size, yellowish-white looking containing chylous fluid (Figure 3-4). They were located at approximately 45 - 50 cm from the Treitz ligament. Resection of the mass along with the adjacent jejunum was performed (Figure 5), and the continuity of gastrointestinal motility was maintained. Tissue samples were sent for histopathological examination. The patient did not develop any complications in the postoperative period. The patient was discharged after 1 week of treatment in the hospital. The histological analysis of the cysts were reported as MCL. The patient was given detailed information about the current diagnosis and treatment. Signature was obtained for informed consent form from the patient. No
recurrence was observed during the 2 years 2 months follow-up period.

Figure 1. Abdominal X-ray finding of ileus

Figure 2. Axial and coronal section computed tomography: Widespread free fluid and two cystic lesions compressing the small intestines superiority and left laterally, in the abdominopelvic region

Figure 3. Intraoperative view of the non-ruptured MCL with thin-walled chylous fluid-filled

MCLs are an uncommon intraabdominal cystic lesions of the mesenteric lymphatics and compose 1 / 100000-250000 of hospital admissions.

Lymphangiomas which seen in childhood are usually diagnosed 40% in the first year of age and 80% of them are up to 5 years of age. Although MCL is usually asymptomatic, the patient may present with chronic or acute abdominal pain, distension and an palpated abdominal mass. Also MCL may cause acute presentation because of an intestinal obstruction or peritonitis caused by infected cysts, torsion, compression of adjacent organs, intestinal ischemia, malrotation, bleeding into the cyst or as in our case, acute abdomen may develop due to post-traumatic cyst rupture, which is rarely seen. Although abdomen ultrasonography to show its cystic nature,
CT and MRI to show its complex anatomic relationships are highly sensitive. Duplication cyst, ovarian cyst, omentum cyst, peritoneal inclusion cyst, echinococcal cysts and multiloculated acid that causes intra-abdominal cystic appearance should be considered in differential diagnosis. The definitive diagnosis is made by histological examination of tissue.

The treatment for abdominal mesenteric lymphangioma is complete surgical excision, sometimes including resection of the intestinal segment that is closely related with the cyst because of the risk of recurrence and malignant transformation.

As a conclusion, MCLs are extremely rare intra-abdominal cystic masses in children. As they may not lead to any clinical complaints, as in our case, may rarely present with acute abdominal findings due to post-traumatic cyst rupture and should be kept in mind in the differential diagnosis of intra-abdominal cystic masses.

REFERENCES