Ileocecal Obstruction Due to B-cell Non-Hodgkin Lymphoma

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Abstract

We report a rare case of non-Hodgkin lymphoma presented as an ileocecal mass. The patient was a 77-year-old man with history of symptoms of partial bowel obstruction, intermittent right iliac fossa pain, loss of weight, vomiting and fatigue. Clinical signs included moderate abdominal tenderness with a palpable mass in the right iliac fossa at the physical examination. Colonoscopy revealed an intussusception of the right colon causing a complete stenosis. The patient developed complete bowel obstruction during hospitalization that required emergent surgical intervention. Intraoperatively an ileocecal mass was found measuring 10-12 cm in diameter, causing complete stenosis at its level and bowel dilatation proximally. Multiple nodules were found in the liver and the parietal peritoneum as well. An ileotransverso-anastomosis was performed and biopsies of the nodules were taken. Pathological evaluation revealed a diffuse large B cell non-Hodgkin's lymphoma of the ileocecum and the parietal peritoneum.

Key words: non-Hodgkin lymphoma, ileocecum, bowel obstruction

Background

The latest classification of lymphomas, which was established by the WHO in 2008, had abandoned the terms “Hodgkin” and “non-Hodgkin” using more than 80 different forms of lymphoma in 4 broad groups (1). Despite that the terms “Hodgkin” and “non-Hodgkin” are still used in daily practice.

Primary gastrointestinal lymphoma is very rare, constituting

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Rezumat

Ocluzie ileo-cecalã secundarã unui limfom non-Hodgkin cu celule B


Cuvinte cheie: limfom non-Hodgkin, ileocecal, ocluzie intestinalã

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only about 1-4% of all gastrointestinal malignancies (2). The gastrointestinal (GI) tract is the most common extranodal site of involvement in non-Hodgkin lymphoma (NHL) (3). The stomach is the predominant location for GI lymphomas (50%), whereas intestinal lymphomas are less frequently observed in the small bowel (20-30%) (4, 5). The small intestine is less commonly involved with lymphomas than the large intestine. The ileum is the most common site of small intestinal lymphoma (6).

The most frequent presentation of small bowel tumors is occult gastrointestinal bleeding forming 57% of the cases, while chronic abdominal pain, diarrhea and bowel obstruction account for 15% of them (7). Intussusception is the least frequent presentation, particularly in adults. It is most often seen in infants and children and only 5% of all cases are adults. Adult intussusception is genuinely rare, only 1% of adult bowel obstructions are due to intussusception and about 40% of them are related to malignant lesions (6).

**Case report**

A 77-year-old man presented with symptoms of partial bowel obstruction; vomiting, pain in the right iliac fossa associated with weight loss (about 5 kg in the last month). On physical examination, the patient looked pale and had a palpable mass on the right iliac fossa. Laboratory tests revealed anemia (Hb=9 d/dl; Hct= 27.1%) and anormal tumor marker level (CEA=0.5 ng/mL). An abdominal ultrasound was performed, which detected a mass in the right iliac fossa of about 12 cm in diameter and multiple nodules spreading in the liver measuring up to 1 cm in diameter. To complete the investigations, a total colonoscopy was performed, revealing a stenosis of the right colon, with aspect of a colic intussusception; biopsies were taken from that area as well. Moreover, a chest X-ray was done, but no tumors were discovered in the lungs.

Histopathological examination revealed nonspecific colitis. During hospitalization, the patient developed complete bowel obstruction and we decided to perform the surgery.

Intraoperatively, an ileo-cecal mass was found measuring 10-12 cm in diameter, causing a complete stenosis at its level and bowel dilatation proximally. The parietal peritoneum and the liver were covered with multiple nodules, measuring 1-1.5 cm in diameter. So we decided to perform a side-to-side ileo-transverso-anastomosis. Biopsies from the peritoneal nodules were taken.

Histological examination revealed a high-grade non-Hodgkin’s B-cell lymphoma. To confirm the diagnosis, additional immunohistochemical staining analysis of the peritoneal lesions was performed. We used the following markers for immunohistochemical staining: CD 20+ (Fig. 1), AE1/AE3 (negative, exclude poorly differentiated carcinoma) (Fig. 2), CD 3 (negative on the tumoral cells - positive internal control) (Fig. 3), and Ki67 (present in 90% of tumoral cells).

**Discussion**

Due to the high proportion of lymphoid tissue, the ileo-
The ileocecal region is the most commonly involved site in primary intestinal lymphomas (8, 9). Non-Hodgkin’s B-cell lymphoma is the most common subtype of primary lymphoma of the small intestine, and these patients have been found to have a poor outcome compared to those with lymphoma in other regions of the GI tract (3).

Several studies have shown that surgical resection performed to control the disease locally, to prevent bleeding and/or due to perforation can rarely eliminate the disease when used alone (10-14). Despite this, a surgical resection should always be attempted for localized tumors. On the other hand, the management of extensive GI lymphoma remains a controversy. This lymphoma is commonly diagnosed at an advanced stage, and surgical treatment is only suitable for 30-40% of such patients (3). As a consequence, radiotherapy and adjuvant chemotherapy are essential therapeutic approaches. In case of localized gastric lymphomas radiotherapy is preferred to surgery, but in intestinal lymphomas a combination of surgery and chemotherapy was proven to be superior to any other treatment combination (15).

Regarding the anatomical site, previous studies have shown that ileocecal lymphomas frequently require emergent surgical intervention. Fifty-four percent of ileocecal lymphomas require immediate surgery, while 30% of non-ileocecal lymphomas present with complications (4, 16-19). In our case it was hard to say if it was an ileal or a cecal lymphoma, because this neoplasia includes the area from the distal ileum to the cecum, so it was difficult to precisely designate the primary site of the tumor. Thus, the designation of this region differs among studies, some consider it a part of the small intestine while others a part of the large intestine (20). Regional lymph node involvement is considered as a bad prognosis.

Conclusion

In conclusion, we describe the diagnosis of adult ileocecal obstruction due to B-cell non-Hodgkin lymphoma with detailed histopathological images of this specific case. Obstruction due to B-cell non-Hodgkin lymphoma must be considered in the differential diagnosis of patients with abdominal pain and vomiting. Work-up must include plain abdominal X-ray, ultrasound and CT scan of the abdomen and PET-CT in special cases. Our patient developed a complete bowel obstruction so a CT scan wasn’t necessary, otherwise it could be helpful to determine the cause of the obstruction.

References