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Rectal mature teratoma: A case report

Liu JL *et al.* Rectal mature teratoma

Jia-Li Liu, Ping-Liang Sun

Abstract**BACKGROUND**

Rectal mature teratoma is rare and has been reported as a case report in this study. Herein, clinical presentation, magnetic resonance imaging findings, and immunohistochemistry showed a pelvic rectal mature teratoma. The case report and the surgical treatment procedure have been discussed below.

CASE SUMMARY

A 29-year-old Chinese female showed up with over a one-month history of perianal mass that emerged after defecation. Physical examination indicated that the mass was 4 cm × 3 cm × 3 cm. The intraoperative procedure involved ligation of the sigmoid colon 10 cm above the upper edge of the tumor, followed by ligation of the rectum 3.5 cm above the upper edge of the tumor, and subsequent complete removal of the mass. The histopathology confirmed the mature teratoma.

CONCLUSION

The tumor can be completely removed using surgery to prevent its recurrence.

Key Words: Rectal; Mature teratoma; Therapy; Case report

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Core Tip: Herein, a rectal mature teratoma patient is reported. However, only a few similar cases have been reported. Currently, it is difficult to diagnose mature rectal teratoma using a computed tomography scan. However, complete removal of the tumor using surgery can prevent its recurrence.

INTRODUCTION

Teratoma is a tumor caused by pluripotent cells, especially the embryonic stem or seed cells in the gonad or embryonic part of the body. It occurs in the midline or on both sides of the body. It often originates from the Hensen's node, (the location of pluripotent stem cells. Teratoma also occurs in the sacral region, where pluripotent cells are also located^[1]. Teratoma is mostly benign with low malignant potential, but it can also develop into a malignancy^[2]. Rectal teratoma is rare, and there are few reports worldwide. Mature teratoma is a benign tumor (dermoid cyst) and accounts for over 95% of teratomas. Mature teratoma mostly occurs in women of childbearing age, and sometimes in young girls and postmenopausal women. It rarely occurs in males ^[3]. This study aimed to review the diagnosis and treatment of rectal teratoma and to determine the clinical characteristics associated with this rare tumor.

CASE PRESENTATION

Chief complaints

A 29-year-old female, G1P0, with over one-month history of a perianal mass out after defecation, ¹ was hospitalized in the First Affiliated Hospital of Guangxi Chinese Medicine University.

History of present illness

She reported a one-month medical history of perianal mass out after defecation and also complained about the anal bulge. The patient had not used contraceptives, was not injured, had no pain, chills, or fever, and no difficulty during defecation.

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History of past illness

The patient had not past illness.

Personal and family history

The patient had a history of artificial abortion and no family history of Rectal mature teratoma. The condition was diagnosed as a rectal mass (nature to be investigated).

Physical examination

The mass was 4 cm × 3 cm × 3 cm inside the anus with a dentate line distance of about 6 cm and was smooth upon palliation, and non-tender mass was seen outside the anus.

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Laboratory examinations

Hematological examinations, including serum electrolyte levels, human chorionic gonadotropin, comprehensive metabolic panel, and complete blood count, were normal.

Imaging examinations

Electronic colonoscopy: Rectal mass (nature to be investigated) (Figure 1).

The computed tomography (CT) scan revealed: (1) A 6.3 cm × 4.7 cm × 5.1 cm round mass, flaky low-density shadow and calcification on center, enhanced scanning lesions with circular mild enhancement, non-enhancement on center, and clear boundary on pelvic(unclear if this is a teratoma); and (2) Double-sided adnexal area low-density shadow(cyst) (Figure 2).

A rectal mass resection was performed *via* laparoscopic under anesthesia to alleviate the patient's symptoms.

FINAL DIAGNOSIS

The condition was diagnosed as mature rectal teratoma based on the above physical examinations and imaging data.

TREATMENT

Surgical procedure

A rectal mass resection was conducted *via* laparoscopy under anesthesia. Intraoperative ligation was conducted on the sigmoid colon 10 cm above the upper edge of the tumor and on the rectal 3.5 cm above the upper edge of the tumor, followed by complete removal of the mass. Full hemostasis, sigmoid colon and rectal suture repair, placement of a negative pressure drainage tube in the anus and abdominal cavity, and layer-by-layer suture repair of the incision was then conducted (Figure 3).

Pathological examination

In the intestinal section, two connected tumors, about 6 cm × 5 cm × 4 cm mass and 2 cm × 2 cm × 2 cm mass, were seen in the intestinal mucosa and intestinal serosal layer, respectively. In the microscopic view, skin and appendages, glands, fat, bone tissue, bone marrow tissue, and brain tissue indicated mature teratoma. No tumor tissue was seen at the two ends (upper and lower margins) after the examination. Six lymph nodes were found, and no tumor metastasis was identified (0/6). Therefore, the condition was diagnosed as mature teratoma (Figure 4).

OUTCOME AND FOLLOW-UP

Postoperatively, the patient was discharged after healing. She returned for a follow-up in August 2018. On examination, there was evident wound healing and no tumor

recurrence. Additionally, the patient was free of discomfort, pain, and fecal incontinence.

The colonoscopy and CT scan revealed a rectal mass, 6 cm × 5 cm × 4 cm in intestinal mucosa and 2 cm × 2 cm × 2 cm in the intestinal serosal layer, which was diagnosed as mature rectal teratoma. Laparoscopic tumor resection was conducted to remove the tumor. No tumor metastasis was found six months after successful 1-mo treatment. The teratoma was located in the rectal wall, which is close to the pelvic cavity. The teratoma volume increases and breaks into the intestinal wall, ad bulging to the posterior wall of the rectum. The teratoma then comes out of the anus and can only be returned by hand.

DISCUSSION

Clinical reports of teratoma are common in the sacrococcygeal, appendix, ovary, testis, retroperitoneum, mediastinum, *etc.* Several studies have shown that the incidence of teratoma may be related to various factors, such as genetic, environmental, and gene-level regulation^[4,5]. Teratoma can be divided into benign and malignant transformations based on the degree of tissue differentiation. Teratoma incidence is about 1:35000-1:40000^[6] and mostly occurs in women (the ratio of male to female is about 1:2-4) with few occurrences in children and postmenopausal women^[7,8]. Although mostly reported in the ovary and testis, it also occurs in the midline of the mediastinum, appendix, sacrococcygeal, pineal body, mediastinum, posterior peritoneal cavity, omentum, uterine rectum, vagina, and cervix^[9-12]. Immature teratomas occur in adolescents. Most malignancies transform into cancer (squamous cell carcinoma). About 1%-2% of teratoma cases are malignant and are common in young women (the average age of onset, 11 to 19 years) with poor prognosis^[13-15]. CT images of mature teratoma reveal calcification, adipose tissue, bone, tooth, and Obviously cyst^[16,17].

CT scan is sensitive to calcification and fat, common and quick, combined with enhanced scan can evaluate the soft tissue composition well, but lacks specificity for differentiating between tumor types. While MRI has a higher resolution of fat, soft tissue, which helps to determine the retrorectal tumors and their relationships of

surrounding structures and cystic degeneration, but shows poorly for calcification^[18,19]. To some extent, MRI is more accurate than CT to estimate the possible complications such as torsion, rupture, and malignant transformation.

Badmos reported that laparoscopic surgery can enlarge the field of view, reducing the incision and intraoperative blood loss^[20]. Lee *et al*^[21] also reported that laparoscopic surgery could significantly reduce the body's inflammatory response compared to open surgery. Tharintorn reported a case of complicated duodenal mature teratoma, which was resected *via* laparoscopic surgery^[22]. Herein, the mature cystic teratoma was identified, and the patient was discharged after the operation. No recurrence occurred after six months of follow-up. Laparoscopic pelvic and teratogenic teratoma surgery is widely used because of the minimally invasive advantages. Laparoscopic surgery completely removes the tumor without damaging adjacent tissues and organs, avoiding the rupture of the tumor and preventing leakage of the teratoma, thus inhibiting malignant transformation, recurrence, and metastasis^[23,24].

Murdock and Abbas^[25] reported that an anorectal cystic teratoma ⁵transabdominal approach is necessary, which can be done laparoscopically safely and successfully, even for a large lesion.

Wang *et al*^[26] ²reported that generally not recommended to use preoperative biopsy of retrorectal tumors because of the risk of infection or tumor seeding in the pelvis. As such, a definitive diagnosis is best to obtain by following complete resection of the tumor. Resection of retrorectal teratoma is generally regarded as appropriate because of the malignant potential.

Aiken *et al*^[27] reported that the diagnosis can be made with endoscopy alone by the presence of hair over the mass.

Nam and Kim^[28] reported that the mass was removed by polypectomy because of patient's lesion was a pedunculate polyp measuring approximately 4cm located approximately 15cm from the anal, endoscopic resection was performed to make a diagnosis. Endoscopic resection is indicated for a pedunculate polyp that measures < 4 cm. If the diagnosis is unclear or malignancy cannot be excluded, surgical resection is

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preferable. The summaries of reported cases of rectal mature teratoma are shown in Table 1.

CONCLUSION

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