Uterine metastasis from the rectosigmoid transition cancer: a case report

Nicoli Taiana Henn¹*, Kelen Zanin², Bruna de Oliveira², Daniela Schwingel³ and Julia Pastorello¹

Abstract

Colorectal Cancer is common and it will often present with metastases during the course of the disease and follow-up. Usually metastasis affects liver and lungs, but other organs may also be involved. We report the case of a patient who had cancer in the recto-sigmoid transition in 2002 and had a recurrence 14 years after surgery with metastasis in the uterine body. The patient had the tumor resected and was given adjuvant chemotherapy. It is important to consider this possibility in patients with prior history of neoplasm that presents with abnormal vaginal bleeding.

Keywords: Uterus, metastasis, colon cancer

Introduction

Colorectal cancer includes tumors that affect colon segment or rectum [1]. Usually evolves from adenoma to adenocarcinoma over the years [2]. The segments most affected by this tumor are rectum and sigmoid, being followed by cecal region.

The global estimate in 2012 put the colon cancer and rectum as second most common among women and the third most common among men, and the majority (55%) occur in countries with a high Human Development Index (HDI), unlike its mortality, more common in countries with low HDI [1].

This cancer is closely related to the population way of life, being higher when there is higher consumption of red and processed meats, low intake of fruits and vegetables, physical inactivity, obesity and overweight, and the consumption of alcohol and tobacco habit [1]. Patients with inflammatory bowel disease (Crohn's disease and ulcerative colitis) are also at increased risk of developing this neoplasm [3]. In addition, genetic syndromes increase the risk of colorectal cancer, as Lynch syndrome (hereditary cancer no polypoid) [4] and Familial adenomatous polyposis [5].

The survival rate of colon cancer changes by staging and early diagnosis and treatment. For patients with stage I disease and low-risk stage II or high frequency microsatellite instability, usually surgical resection with removal of lymph nodes in block is the only approach needed [6]. For patients with clinical stage II with high risk factors (T4, poorly differentiated histology, lymphovascular and/or perineural invasion, intestinal obstruction, positive margins or inadequate lymph node resection) after surgery, adjuvant chemotherapy may be considered generally with fluoropyrimidine alone. The patients with stage III, adjuvant therapy is recommended for six months [7] generally with fluoropyrimidine and oxaliplatin.

Some selected patients have benefit from surgery for metastasis when cure is sometimes possible, which is shown especially with liver metastasis [8].

As the progress of therapy for the disease and long-term monitoring of patients, less common sites of metastases are present, as is the case of metastasis in the uterine body. We present the case of a woman 46 years old with metastasis in uterine body from colonic adenocarcinoma.

Case presentation

A 46 years old woman of with recto-sigmoid transition adenocarcinoma history in 2002, submitted to rectosigmoidectomy with pathology staging II (T3N0M0) that received adjuvant treatment with injectable Fluoropyrimidine for six cycles, and later local radiotherapy. She remained in monitoring for five years and then she was discharged. In 2016, she experienced abnormal vaginal bleeding and sought gynecological assistance that requested a pelvic magnetic resonance, which showed small
nodular formation in the fundus of the uterus and proximal segment of the endometrial cavity, exhibiting paramagnetic enhancement. She underwent hysterectomy, pathology and subsequent immunohistochemistry, which showed moderately differentiated adenocarcinoma with positive neoplastic cells in the following antibodies: CK20, CDX2 and CEA, with no positivity for antibodies: CK7, vimentin and Estrogen Receptor. Computed tomography (CT) scan showed no disease at another site and the postoperative value of carcinoembryonic antigen (CEA) was 1.5 ng/mL. She had a recent colonoscopy without evidencing new lesions. It was then decided to start chemotherapy with mFOLFOX6 scheme for the patient for six months. It was done microsatellite instability research which showed immunohistochemistry positivity for MSH-2 and MSH-6 antibodies, and no positivity for MLH-1 and PMS2 antibodies, and because of that, indicated evaluation for BRAF mutation (Figures 1-5).

Figure 1. H&E-2.5x-adenocarcinoma with foci of necrosis infiltrating the myometrium.

Figure 2. Immunohistochemistry 2.5x-Vimentin-absence of marking.

Figure 3. Immunohistochemistry 2.5x-Estrogen receptor-absence of marking.

Figure 4. Immunohistochemistry 2.5x-CK20-presence of marking.

Figure 5. Immunohistochemistry 2.5x-CDX2-presence of marking.
Discussion
If the tumor were mucinous, it might be very difficult to distinguish between colonic origin or the female genital tract [9-12]. In the immunohistochemistry, it could have presented CK7 positive and CK20 negative even if it was from colonic origin [13], but our patient presented the most common profile in colon cancer (CK7 negative and CK20 positive).

Usually colonic cancer metastases are concentrated in the liver, lymph nodes and lungs, but other sites may also be involved. When the female genital tract is affected, usually it occurs in the ovaries [14]. Metastasis to the uterine body is infrequently diagnosed. The metastasis could be in the cervix uterine, as also previously described [15,16].

The oligometastatic disease can be characterized by the presence of metastasis in two or three sites, occasionally five or sometimes more lesions, particularly in viscera. In these patients, we can use local therapies to improve their outcomes. When metastatic disease is confined to an organ, a potential curative treatment exist [17].

A case of metastasis to the uterine body of a sigmoid cancer was reported by Tokoro Y [18], but in this case metastasis was synchronous to the initial tumor and the diagnosis was also made only with the help of immunohistochemistry, as originally also thought to be of endometrial origin. After resection of both lesions was indicated adjuvant therapy with Xelox scheme for six months.

In a series of 63 reported cases of metastasis to the uterine body by extragenital primary neoplasms, the colon was the second most common primary site, it lost only to breast. Remember also that when the uterus is affected, usually occurs by direct extension to adjacent organs or peritoneal dissemination of ovaries carcinomas implants [14].

The patient in this case report had her primary tumor in 2002, 14 years later she had recurrence. CT scans showed no lesions elsewhere, so we opted to extrapolate postresection data to liver and lung metastases and, following the same idea than Tokoro Y, we also indicated adjuvant chemotherapy scheme based in fluoropyrimidine and oxaliplatin and the patient is tolerating well.

In addition, based on the results of immunohistochemistry in which MLH-1 and PMS2 (proteins of DNA repair) are impaired, we follow the Guidelines for Genetics evaluation and Lynch Syndrome management [19] and request search for BRAF mutation. Considering the 2002 approach, if it had the availability and knowledge of the microsatellites instability test, this patient probably would have not received chemotherapy at the time.

Conclusion
We should pay attention to patients with previous long term history of cancer if they show abnormal vaginal bleeding complaint, we must remember that there is a probability that we are facing a possible metastasis, as well as uterine primary malignancy or benign causes.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions

<table>
<thead>
<tr>
<th>Authors’ contributions</th>
<th>NTH</th>
<th>KZ</th>
<th>BO</th>
<th>DS</th>
<th>JP</th>
</tr>
</thead>
<tbody>
<tr>
<td>Research concept and design</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
</tr>
<tr>
<td>Collection and/or assembly of data</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>--</td>
</tr>
<tr>
<td>Data analysis and interpretation</td>
<td>✓</td>
<td>--</td>
<td>--</td>
<td>✓</td>
<td>--</td>
</tr>
<tr>
<td>Writing the article</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Critical revision of the article</td>
<td>✓</td>
<td>--</td>
<td>--</td>
<td>--</td>
<td>--</td>
</tr>
<tr>
<td>Final approval of article</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
</tr>
</tbody>
</table>

Acknowledgements
We thank the patient who allowed the publication of her case.

Publication history
Editor: Stefano Fratoni, St.Eugenio Hospital of Rome, Italy.
Received: 12-Mar-2017 Final Revised: 15-Apr-2017
Accepted: 05-May-2017 Published: 19-May-2017

References


Citation: