Anorectal malignant melanoma: A case report and treatment options

Abstract
Anorectal malignant melanoma (AMM) is a rare malignant disease with a poor prognosis. This disease is often confused with hemorrhoids. The most common site of malignant melanoma following skin and eye involvement is the anorectal region. This is the most commonly involved site in the gastrointestinal tract. We report the case of a 67-year-old patient with lower gastrointestinal hemorrhage for 4 months and hemorrhoid treatment for 2 months. The imaging revealed no distant metastasis but histopathologically, lymph node metastasis and invasion of surrounding tissues. Laparoscopic abdominoperineal resection (APR) was performed.

Key words: Anorectal diseases, malignant melanoma, surgery.

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Öz

Anahtar Kelimeler: Anorektal hastalıklar, malign melanom, cerrahi.
Introduction
Malignant melanoma is a rare disease, but it has usually a poor diagnosis because it is diagnosed late [1]. It can be mistaken for hemorrhoids. The incidence of malignant melanoma is 0.1-4.6% of all rectal malignancies [3]. The patient's complaints included rectal bleeding, palpable mass, breech pain, tenesmus and pruritis [5]. Its treatment is surgical, with extensive local excision or abdominoperineal resection [6]. In our 67-year-old woman, we performed abdominoperineal resection due to locally advancing of the tumor.

Case report
A 67-year-old female patient was treated with diagnosis of hemorrhoids in another hospital due to lower gastrointestinal hemorrhage, 4 months ago. The patient received medical treatment for 2 months. Physical examination revealed an irregular mass on the anorectal junction. Colonoscopy was planned. The colonoscopy showed a mass located at 3.cm from the anal canal and surrounding the anal canal (Figure 1). Histopathological examination revealed mucosal malignant melanoma (Figure 2). No other features were found in the other system examinations of our patient.

Abdominal computed tomography showed a mass lesion of 2.5 cm in diameter at the thickest site, and lymph nodes in the pararectal region, the largest of which was 3 * 1.5 cm in size. Increased attenuation suggesting infiltration was observed in perirectal fat planes (Figure 3). In addition, multiple diverticula were observed in the sigmoid colon and the descending colon (diverticulosis coli).

Figure 1: Colonoscopic image of anorectal malignant melanoma.

Figure 2: Histopathological imaging of anorectal malignant melanoma.

Figure 3: Computed tomography image of anorectal malignant melanoma.

Following preoperative evaluations, laparoscopic abdominoperineal resection was planned. Invasion of the mass into the posterior wall of the vagina was observed during surgery. The patient was consulted with gynecologist peroperatively. Total excision of the mass was completed including the vagina in the posterior wall. Histopathological examination revealed that the tumor was 5.5 X 4 X 1.2 cm in size, passed through the muscularis propria and showed direct invasion to the adipose tissue and surrounding structures. The patient had uneventful postoperative period and was discharged on the 7th day.

Informed consent was obtained from the patient.

Discussion
Anorectal malignant melanoma is an extremely rare disease of neuroectodermal origin, which is aggressive and has a poor prognosis, and is usually fatal with late diagnosis [1]. Malignant melanoma of the anus and rectum was first reported by Moore in 1857 [2]. Rectal malignant melanomas constitute 0.2-3% of all malignant melanomas and 0.1-4.6% of rectal malignancies [3]. The most common symptom is rectal hemorrhage, so it is confused with hemorrhoidal disease and causes delays in diagnosis. Our patient also had medical treatment for a long time with the diagnosis of hemorrhoids. The most common symptoms of AMM include palpable mass, pain in the breech, changes in bowel habits, pruritis, tenesmus, and anorectal region [1, 5]. In our patient, bleeding and severe pain in the breech area were prominent. The contribution of chemotherapy and radiotherapy is limited. Surgical treatment is still controversial even today [9]. In previous studies, radical resections such as abdominoperineal resection were accepted for treatment, whereas in recent studies there was no significant difference in survival and prognosis with wide local excision (WLE) technique [4]. Thibault et al. [5] recommended WLE...
with a negative surgical margin of at least 1 cm in treatment and recommended APR for palliative or non-WLE-only tumors for obstructing tumors. There is no consensus on the surgical margin in WLE and it should be noted that even a 1 cm negative surgical margin can lead to anal incontinence. Therefore, if negative surgical margin can be achieved without causing anal incontinence, WLE should be performed [6]. However, some studies have suggested that aggressive treatment with APR has a better survival rate, possibly associated with lymphadenectomy, that can control lymphatic spread (especially mesenteric lymph nodes) and that wider negative surgical margins and lower local recurrence rates can be achieved. [4, 8]. Yap et al. [1] reviewed 17 series published and reported that there was no statistically superiority between two techniques in all stages of the disease in their 5-year results. To sum up, treatment of anorectal malignant melanoma should be patient-specific [7].

In conclusion, in light of all these information, we preferred APR because of pathological appearance LAPs in the patient's pararectal region, infiltration in the perirectal fatty area and vaginal invasion. The closeness of the mass to the anal canal played a major role in this decision.

References