Peritoneal carcinomatosis of the cutaneous malignant melanoma: A report of very unusual case and review of the literature

Kutanöz malign melanomanın peritoneal karsinomatozisi: Çok nadir görülen bir olgu sunumu ve literatür taraması

Özge Kömürcü Karuserci1, Seyhun Sucu1, Betül Öğüt2, Suna Erkılıç2, Özcan Balat1
1 Gaziantep University Faculty of Medicine, Department of Obstetrics and Gynecology, Gaziantep, Turkey
2 Gaziantep University Faculty of Medicine, Department of Pathology, Gaziantep, Turkey

Abstract
Cutaneous malignant melanoma is a relatively rare and aggressive malignancy that arises from melanocytes of the skin, which is able to spread quickly and widely. It accounts for approximately 3% of all malignancies reported in Europe. It often demonstrates unpredictable metastatic behaviour, but usually spreads to lymph node, lung, liver, brain and bone. The pelvis are a rare site of metastasis and peritoneal carcinomatosis caused by cutaneous malignant melanoma is a rare condition. In this article, we present a case of metastatic cutaneous malignant melanoma presenting as clinical findings such as peritoneal carcinomatosis due to advanced stage ovarian cancer.

Keywords: Peritoneal carcinomatosis, cutaneous malignant melanoma, ovary.

Öz

Anahtar Sözcükler: Peritoneal karsinomatozis, kutanöz malign melanoma, over.

Introduction
Cutaneous malignant melanoma is a destructive malignancy initiating from melanocytes of the skin with a wide metastasis profile. It accounts for approximately 3% of all malignancies reported in Europe. The tumor generally spreads from skin to the lymph nodes (42-59%), lungs (18-36%), liver (14-20%), brain (12-20%), and skeletal bones (11-17%) (1,2). Less frequent sites include the eye, dura, pleura, duodenum, uterine cervix, and vagina. Nevertheless, peritoneal carcinomatosis of a malignant melanoma has been reported limited edition.

We report an exceptional case of peritoneal carcinomatosis originating from malignant cutaneous melanoma metastasis to the adnex, omentum, and uterus that was initially considered to be ovarian carcinoma. A review of the recent literature is also provided.

Case Report
A 43-year old woman was referred to our clinic with lower abdominal pain and distension with weight loss over six months and reduced appetite. Examination revealed tenderness and fullness in the abdomen and pelvis. Transvaginal ultrasonography revealed a 125x78 mm solid mass indistinguishable from the uterus and common ascites in the pelvis. Computerized tomography (CT) revealed solid lesions, the largest 10 cm in size, in front of the uterus and in the left paracervical space, together with peritoneal implants, ascites and bilateral multiple inguinal lymph nodes. No abnormal findings were determined at colonoscopy or upper endoscopy.
Frozen section biopsy of the ovaries was reported as malignant, but it was not clear whether this was primary or metastasis. Total abdominal hysterectomy, bilateral salpingo-oophorectomy, omentectomy, and peritonectomy was performed.

Macroscopically, the right salpingo-oophorectomy specimen was the largest tumor nodule at 15 cm in diameter. Histological examination revealed a malignant tumor within the peritoneum, myometrium, bilateral ovaries, and fallopian tubes, but not the endometrium or uterine cervix. Immunohistochemically, tumor cells stained with S100, HMB45, MART-1, and SOX10 and no reaction was determined with cytokeratin, CD34, desmin, actin, calretinin, ER, PR, or inhibin-A. The histopathological diagnosis was melanoma (Figure 1). Ascites fluid cytopathology was also reported as malignant.

Physical examination revealed a suspected, irregular nevus-like lesion in her back. The patient was transferred to the department of plastic surgery for skin excisional biopsy. Histopathological diagnosis was superficial spreading melanoma and V600 mutations in the serine–threonine protein kinase B-RAF (BRAF) were observed in the tumor cells. The patient was afterwards transferred to the department of oncology for Vemurafenib (PLX4032) treatment which frequently provides tumor regression in patients with BRAF V600–mutation and improved survival time (3).

Discussion
Cutaneous malignant melanoma metastasis to the peritoneum is a rare condition, only three cases of which have previously been reported (4-6). To the best of our knowledge, this is the fourth case of a malignant melanoma with peritoneal carcinomatosis mimicking advanced ovarian cancer. Clinical characteristics, age ranges and origins were very similar in all cases, but two (4,6) were reported as recurrence of malignant melanoma, and one (5) was identified after exploratory laparotomy for peritoneal carcinomatosis, as our case.

In conclusion, although very rare, metastatic malignant melanomas may cause pelvic spread and peritoneal carcinomatosis, mimicking advanced stage ovarian cancer, especially in young women.

References