An aggressive case of anorectal melanoma

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Introduction

All melanomas originate from the melanocyte. They may arise not only from the skin, but also from mucosal epithelium lining of the respiratory, alimentary and genitourinary tracts. Anorectal melanoma accounts for 1–3% of all anal tumors and 0.3% of all melanoma. The majority of cases arise from the mucocutaneous junction (dentate line).1,2

It usually occurs in the 5th or 6th decades of life. The most common presenting complaints are bleeding, anal pain or mass, pruritus, tenesmus and change in bowel habits. Weight loss, anemia and fatigue may indicate metastatic disease. At presentation, 60% of the patients have lymph node involvement and 30% have distant metastases.

This very rare tumor can be classified into three different stages of disease progression: localized disease (I), regional lymph node involvement (II) and distant metastases disease (III), which account for a median survival of 24, 17 and 8 months respectively.3

Surgical resection is the gold standard treatment. Chemotherapy and radiation therapy alone have not been shown to be effective, but may provide some benefit when used as adjuvants. Despite treatment, the prognosis remains poor.4

Case report

This case presents a 55-year-old female patient with a past medical history significant for hemorrhoids and prior depression who was admitted to our medical facility for syncope. As associated symptoms, she reported intermittent rectal bleeding, usually after a bowel movement, and involuntary weight loss over the past 6 months.

Due to family history of colorectal cancer in her father at the age of 47, she had had a total colonoscopy the year before which confirmed just internal hemorrhoids. She also had had CEA and CA 19.9 measurements two months prior to admission and both markers were within normal range.

On physical examination, conjunctiva pallor was noted and digital rectal exam revealed a 5 cm rectal mass palpated 3 cm from the anal verge. Admission laboratory studies were significant for iron-deficiency anemia. Chest X-ray showed multiple bilateral pulmonary cannon ball nodules suggestive of metastasis (Fig. 1).

Flexible sigmoidoscopy was performed. A dark polypoid lesion measuring 5 cm in diameter was identified at the anorectal junction (Fig. 2). Histologically, tumor cells contained apparent brown pigment and were positive for S-100 protein and Melan-A immunohistochemical staining. These findings were consistent with the diagnosis of anorectal melanoma.

Tumor staging included standard head, thoracic, abdominal and pelvic CT imaging. Multiple bilateral pulmonary
metastases of varying sizes were observed on thoracic CT (Fig. 3). In addition, head CT revealed six lesions involving the frontal and temporal lobes bilaterally, some showing necrotic degeneration. The greatest was 3 cm in diameter and presented extensive surrounding edema, causing lateral ventricular displacement (Fig. 4).

She was immediately placed on high dose corticosteroid therapy and prophylactic antiepileptic drugs. On day 5, she developed rapid focal progressive neurological deterioration with confusion, aphasia, right hemiparesis and sphincter control loss. Due to the advanced tumor stage, she was discharged and referred to a palliative care center just one week after admission. She passed away 8 weeks after the initial diagnosis.

**Ethical responsibilities**

**Protection of human and animal subjects.** The authors declare that no experiments were performed on humans or animals for this investigation.

**Confidentiality of Data.** The authors declare that they have followed the protocols of their work center on the publication of patient data and that all the patients included in the study have received sufficient information and have given their informed consent in writing to participate in that study.

**Right to privacy and informed consent.** The authors declare that no patient data appear in this article.

**Conflicts of interest**

The authors have no conflicts of interest to declare.
References