
Nandkishore Jaiswal¹, Aniket B Payagude²

¹Associate Professor, Department of Surgery, Government Medical College and Hospital, Nagpur.
²Post Graduate Student, Department of Surgery, Government Medical College and Hospital, Nagpur.

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ABSTRACT

Anal canal malignant melanoma is an uncommon malignancy. It is a highly aggressive tumor that tends to spread early in the course and presents with distant metastasis. Due to the rarity of the condition, treatment is yet to be standardized. We report a case of a 47-year-old male patient who presented with the history of mass coming out of anus for 6 months, which was misdiagnosed as a case of hemorrhoids. Abdominoperineal resection was done based on tissue biopsy which suggested it to be a primary anal canal malignant melanoma. The diagnosis was later confirmed by histopathological and immunohistochemistry studies, which was strongly positive for HMB 45 AND Melan-A.

Keywords: Anal canal malignant melanoma, Abdominoperineal resection, HMB 45, Immunohistochemistry, Melan-A.

INTRODUCTION

Primary Anorectal malignant melanoma is a very rare tumor with high malignant potential that tends to spread early with poor overall outcome.¹ It accounts for 1% of all melanomas and 0.5-2% of all melanomas arise in the anorectal region.² ³ After skin and retina, anal canal is the third most frequent site where melanomas are found.⁴ It typically presents in fifth or sixth decades of life, with female preponderance.⁵ To the best of our knowledge, till date only 640 cases have been reported in the literature.⁶ A non-specific presenting symptom of anal melanoma makes early diagnosis difficult. Early diagnosis is crucial so that treatment can be standardized.

CASE REPORT

A 47-year-old male patient presented with a complaint of passing blood in stools associated with mass coming out of anus since 6 months. The patient was in follow up since then misdiagnosed as hemorrhoids by a general practitioner. The patient was then referred to us for further management. Past medical and personal history of the patient was insignificant. On rectal examination, there was a large polypoidal pedunculated friable mass present on the anterior surface of anal canal approx. 1 cm from the anal verge, which was resembling thrombosed hemorrhoids. The biopsy was taken as mass was suspicious for malignancy. Biopsy report suggested it to be a primary malignant melanoma. CECT abdomen [Figure 1A.] demonstrated heterogeneously enhancing intraluminal polypoidal lesion of approx. size 5.7×5×4.1 cm, the length of involvement was approx. 4 cm, fat planes were maintained with surrounding structure with perirectal fat stranding with few necrotic inguinal lymph nodes and no distant metastasis. The patient was planned for surgical intervention. Accordingly, the patient was evaluated with complete blood count, renal function test, liver function test, chest radiograph, echocardiography, pulmonary function test for major surgery. Abdominoperineal resection was done with a permanent colostomy. Gross examination of the specimen showed a black mass of size 5×5×5 cm with an irregular surface which was solid and friable, 1.5 cm from anal verge [Figure 1B]. On microscopic examination [Figure 2A-B], tumor cells were round, polygonal and spindle-shaped with abundant cytoplasm and prominent nucleoli containing coarse black pigment suggestive of malignant melanoma (section from the skin with tumor mass shows epidermis with tumor infiltration) with serosal and lymph node involvement. The diagnosis was further confirmed by immunohistochemistry, strongly positive for Melan-A and HMB 45 [Figure 2C-D]. The patient recovered uneventfully from the surgery and then was managed post-operatively with radiotherapy. Follow up CECT abdomen was done after six months which...
demonstrated hypodense lesion in right lobe of liver with features suggestive of metastasis. The patient is now on chemotherapy.

![Figure 1: CECT ABDOMEN: Heterogeneously enhancing intraluminal polypoidal lesion of approx. size 5.7×5×4.1 cm, the length of involvement was approx. 4 cm (A). Gross examination: showed a black mass of size 5×5×5 cm with an irregular surface which was solid and friable (B).](image)

**DISCUSSION**

Anal canal malignant melanoma is an uncommon malignancy of gastrointestinal tract. It was first reported by Moore DW in 1857[7], since then to best of our knowledge only 640 cases of anorectal melanomas have been reported in the literature of which 375 cases were anal melanoma.[6] Due to its relative rarity, it is difficult to determine the exact Incidence of the disease.[2] Incidence rate of anorectal melanoma is about 2.7 per million per year in the United States.[8] It accounts for 1% of all melanomas and 0.5-2% of all melanomas arise in the anorectal region.[2,3]

Anorectal melanomas arise from the melanocytes found in the intestinal mucosa below the dentate line extending proximally into the rectum, hence also identified as primary anorectal melanomas.[9,10] Unlike cutaneous melanoma, there is no risk factor for the development of mucosal melanoma.[2]

Anorectal melanoma patient usually presents with per rectal bleeding, rectal pain, tenesmus, mass coming out of anus and change in bowel habit’s.[2] Presenting symptoms of anal canal malignant melanoma is non-specific and most of the cases are initially diagnosed and managed as benign conditions, more commonly being internal hemorrhoids. Patients of anal canal malignant melanoma usually present in the later stage of the disease.[11] We hereby report a case of an adult male who presented with bleeding per rectum with an anal mass protruding out and was managed as internal hemorrhoid, which later turned out to be malignant melanoma on tissue biopsy. Approximately 61% of the patients already have lymph node involvement or distant at the time of diagnosis.[12] Our patient was free of distant metastasis at the time of diagnosis, but lymph nodes...
were positive for the malignant cell in pathological specimen. This is important as overall prognosis for anorectal malignant melanoma is poor with overall 5 year survival rate of less than 20%\[^{13}\], the five-year survival rate depends on the extent of the disease and is 32%, 17% and 0% for local, regional and, distant disease, respectively.\[^{11}\]

On microscopic examination presence of melanin in the tumor cell and on immunohistochemistry expression of HMB-45 and Melan-A are strongly suggestive of melanoma\[^{14}\] additionally anal melanoma will stain for S-100, Vimentin, microphthalmia-associated transcription factor proteins, and, Carcinoembryonic antigen.\[^{12}\]

Figure 2: Microscopy examination: Round, polygonal and spindle-shaped tumor cells with abundant cytoplasm and prominent nuclei containing coarse black pigment (A & B) Immunohistochemistry findings: (C) HMB-45 and (D) Melan-A.

Treatment of anorectal melanoma is controversial.\[^{15}\] Abdominoperineal resection [APR] with or without inguinal lymph node dissection OR wide local excision [WLE] are the treatment options available.\[^{3}\] But, due to the relative rarity of the disease and lack of randomized clinical trials consensus on preferred surgical approach has not been reached.\[^{5}\] The difference in Five-year survival rate between patients undergoing APR versus WLE (17% & 19%, respectively) is not significant, Iddings et al.\[^{16,17}\] As overall prognosis of anorectal melanoma is poor, the surgical approach is to be individualized. The postoperative role of adjuvant chemo-radiotherapy has not been well established, but there are studies to support the role of radiotherapy for local control\[^{3,18}\] and, chemotherapy in the management of metastatic disease.\[^{3,19}\] Our patients received postoperative radiotherapy after which patient was followed up for 6 months, Repeat
CECT abdomen was done which suggested liver metastasis and patient is now being advised chemotherapy.

CONCLUSION

Primary Anal canal malignant melanoma is rare and has high malignant potential. Although a rare presentation, diagnosis of anorectal melanoma should be one of the differential diagnoses in the management of benign conditions, particularly hemorrhoids. As there is no consensus regarding management of Anorectal melanoma and its management should be individualized.

REFERENCES


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