Primary Anorectal Amelanotic Melanoma:
The First Case Report from Saudi Arabia

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INTRODUCTION

Anorectal melanomas are exceptionally uncommon. It accounts for less than 1.0% of anorectal malignancies and around 0.3% of all melanomas. 1 Since, most of the cases with anorectal melanomas already have distant metastases at the time of presentation, the prognosis of this disease is poor with a median post-treatment survival time of less than 2 years and the 5-year survival rate of less than 20%. 1,2 Amelanotic melanoma represents about 2-8% of all melanomas, almost 30% of anorectal melanomas are amelanotic, and they are distinct because they contain a little or no-pigmentation. 3,6 However, the exact prevalence of this malignancy may be higher because this type of melanoma is difficult to diagnose. 7,8 Amelanotic melanomas of the anorectum are extremely rare; only 20 cases of amelanotic melanomas of the anorectum have been reported in literature. Amelanotic or hypomelanotic melanomas...
are a subtype of melanomas with no pigmentation and minimally pigmented on clinical examination. Uncommonly, this malignancy lacks melanin pigment, complicating detection. Presenting complaints are non-specific rectal bleeding, pain, itching, or incontinence, mimicking more common disorders, and if malignancy is not suspected these melanomas can be misdiagnosed.\(^3\),\(^4\),\(^8\),\(^9\) Rectal bleeding is the most common symptom in anorectal melanoma, and for this reason this tumor can be misdiagnosed as simple hemorrhoids, which delays the diagnosis.\(^10\),\(^11\)

Amelanotic melanomas are considered as the great masqueraders in dermatology.\(^12\) Several cases of initial misdiagnosis were reported by pathologists, which upon careful investigations and rechecking for the melanoma markers turned out to be amelanotic melanomas.\(^13\) There were reports of amelanotic melanoma cases that were initially misdiagnosed as a basal cell carcinoma,\(^12\) pyogenic granuloma,\(^14\),\(^15\) poorly differentiated adenocarcinoma of the rectum,\(^16\),\(^17\) Bowen’s disease,\(^18\) presenting as contact dermatitis,\(^19\) simulating juvenile xanthogranuloma,\(^20\) and as otitis media.\(^21\) Using the histological criteria alone, differentiation of amelanotic anorectal melanoma from other malignant tumors is difficult, and immunostains are of important help to make the diagnosis accurately.\(^22\) Abnormal expression of immunohistochemical stains such as vimentin, cytokeratin, and c-Kit (CD117) further increases the risk of misdiagnosis.\(^23\)

Due to the aggressive nature and rarity of the disease, there are no separate guidelines for staging of the amelanotic anorectal melanoma tumors, however, anorectal melanomas staging and treatment protocols are also useful for these tumors.\(^9\),\(^24\),\(^25\) Recently, immunotherapy also shows success in treating melanomas in Japan. It has been shown that Nivolumab (OPDIVO), an antibody to programmed death (PD-1) is beneficial in controlling metastasis of amelanotic anorectal melanoma.\(^26\) Surgery in combination with radiotherapy and adjuvant immunotherapy, like interferon, was reported to be a choice of treatment for amelanotic anorectal melanoma if the tumor is not metastasized.\(^27\),\(^28\) Laparoscopic abdominoperineal resection (APR) was also shown to be useful in treating anorectal amelanotic melanoma due to its minimally invasive nature.\(^29\)

**CASE REPORT**

An 84-year-old Saudi Arabian female patient was diagnosed for chronic calculous cholecystitis and had a laparoscopic cholecystectomy 2 years earlier and discharged in stable condition. This time she was admitted to ER with a complaint of anal mass for 3 months. She had a history of diabetes, hypertonse, and hypothyroidism as well. She had no abdominal pain or fever; abdomen was soft and lax with no tenderness or distension. She did not notice any changes in bowel habits, but she noticed fresh blood during defecation. On physical examination there was painful cylindrical mass (7.0×4.0 cm) prolapsing through the anus, which was pale pinkish in color and was thought to be clinically malignant (figure 1).

Abdomen and pelvic computed tomography (CT) showed rectal wall thickening with an eccentric polyloid lesion of soft tissue density arising from the wall of the rectum predominantly from the left side, which measured around 4.24×6.21×7.25 cm (craniocaudal×transverse×anteroposterior) diameters, respectively (figure 2). The mass was situated about 3.9 cm from the anal verge with intraluminal projection and subsequent marked reduction of the anorectal caliber. Perirectal fat planes stranding with presacral lymph node enlargement was seen. The largest lymph node on the left side measured around 1.5×1.8 cm in size (figure 3.B), and left inguinal lymphadenopathy was noted (figure 3.A). Both kidneys showed multiple. The liver was enlarged with diffuse fatty infiltration, no focal lesions were seen. These radiological descriptions suggested the anorectal mass was likely to be malignant and recommended for biopsy correlation. CT of the chest was done, which showed a small (0.77×0.49 cm) nodule in the lower left lobe of the lung that was likely suggestive of metastatic deposit (Figure 3.C). In figure 4 (A and B) contrast enhance axial and coronal magnetic resonance imaging (MRI) of the pelvis with anorectal polyloid mass with marginal and eccentric enhancement is shown. These radiological...
Fig.1: Grossing shows tumor mass extending from the rectum throughout the anus, 4 cm out (yellow arrow), and tumor mass pinkish white color with surface ulceration.

Fig.2: Sagittal (A), and coronal (B) views of abdomen and pelvic CT show cranial caudal extension of anorectal mass (yellow arrow) from the anal verge.

Fig.3: Axial CT of pelvis and chest show rectal wall thickening with an eccentric polypoid lesion of soft tissue density (Fig A & B yellow arrow) arising from the wall of the rectum predominantly from the left side with subsequent marked reduction of the anorectal caliber. Left inguinal lymph node (Fig. B, blue arrow) and presacral lymph node (Fig. A, green arrow) enlargement was also seen. CT of the chest shows a small nodule (Fig C, red arrow) in the lower lobe of the left lung likely representing metastatic deposit.

Fig.4: Contrast enhance axial (A) and coronal MRI of pelvis (B) show anorectal polypoid mass with marginal (yellow arrow) and eccentric (red arrow) enhancement.
descriptions suggested of malignant nature of this tumor. Colonoscopy was also performed to see occult lesion or mucosal changes. Rectal biopsy was obtained from the primary tumor site for histopathological evaluation.

For the histological evaluation, two pieces of soft tissue were received in formalin. The tissues had a solid grayish white surface, covered with ulcerated mucosa measuring 2.5×1.0×1.0 cm. Microscopic examination revealed a piece of skin with hyperplastic squamous epithelium with surface ulceration (figure 5.A). The dermis and underlining tissue were infiltrated by sheets and nests of ovoid and spindle shape cells. The tumor cells had hyperchromatic nuclei with prominent nucleoli and high mitotic activity (Figure 5.B and C). Immunohistochemistry showed immunoreactivity for melanocytic markers such as Human Melanoma Black-45 (HMB-45), and melanoma-associated protein A (Melan-A/MART-1), and also positive for S-100 (figure.6) and negative for P63, CK 5/6, and Pan CK (figure7).

**DISCUSSION**

Melanomas are known to occur on any region of the body, wherever melanocytes present, for example places such as the epidermis, eyes, nasal cavity, oropharynx, vagina, urinary tract, rectum, and anus. The median age for anorectal melanoma cases is 66 years at the time of diagnosis, with a female predominance up to 60%. In

![Fig.5](image-url): (A) H & E stained microscopic pictures showing skin with hyperplastic squamous epithelium with surface ulceration (yellow arrow). Extensive tumor growth beneath the surface squamous epithelium (blue arrow). (B) The dermis and underlining tissue are showing (10x) infiltration by malignant sheets and nests of ovoid and spindle shape cells (black arrows); (C), with prominent and high mitotic figures (yellow arrow). The nuclei are hyper chromatic with some prominent nucleoli (40x).

![Fig.6](image-url): (A) Immunostaining for S-100 shows positivity in tumor cells; (B). Immunostaining for HMB-45 shows positivity in tumor cells. (C) Melan A- is positive in tumor cells (green arrow). The epidermis is negative (Red arrow).
a retrospective analysis of 61 cases of anorectal malignant melanomas during 2005-2014, Tariq and colleagues have reported that around 21% of cases were completely amelanotic.\textsuperscript{31} Examination by CT was used to detect metastasis of this tumor to distant organs such as liver, lungs, and kidneys. Also, the most common sites for metastases are inguinal and pre-sacral lymph nodes as we have observed in this case.\textsuperscript{32-34} Khaled and co-workers have reported a case of amelanotic anorectal malignant melanoma with satellite lesions or an unusual metastatic deposit in the vulva.\textsuperscript{35} Amelanotic melanomas also were reported on body parts such as in the middle ear,\textsuperscript{21} on the upper arm and the shoulder,\textsuperscript{36} in the esophagus,\textsuperscript{37} in duodenum,\textsuperscript{38} on perineum and perianal sites,\textsuperscript{39} on penis,\textsuperscript{40} on female urethra,\textsuperscript{41} and in vagina, breasts, and vulva.\textsuperscript{42-44} Further, amelanotic forms of acral melanomas, hallux (subungual amelanotic melanoma), subungual acrolentiginous melanoma, conjunctival melanoma, irido-ciliary ring melanoma, melanoma of the nasal cavity, melanoma of the tongue, and intracranial amelanotic melanoma involving brain were also reported previously.\textsuperscript{45–54}

Distinction of amelanotic anorectal melanoma from lymphoma, undifferentiated carcinoma, and gastrointestinal stromal tumor (GIST) is difficult on the basis of histological criteria alone. HMB-45 and Melan-A are useful in the differential diagnosis between granular cell tumor and malignant melanoma.\textsuperscript{55} Abnormal expression of immunohistochemical stains such as vimentin, cytokeratin, and c-Kit (CD117) further increases the risk of misdiagnosis.\textsuperscript{31} However, activating c-Kit mutations have been associated with the pathogenesis of malignant melanoma, which are commonly found in leukemia and gastrointestinal stromal tumors.\textsuperscript{56} There are few reports of amelanotic melanomas with S-100 negative and c-Kit positive, one such case that confused clinicians, was from Japan that was difficult to diagnose between amelanotic melanomas and GIST of the anorectum.\textsuperscript{23} In their case the authors reported that the anorectal tumor was positive for vimentin and c-Kit, and negative for S-100, however, an excisional biopsy of the inguinal lymph node showed positive for S-100, Melan-A, HMB-45, and tyrosinase. And upon repeating the immuno-histochemistry of the anorectal tumor, this was positive for Melan-A, HMB-45, and tyrosinase but negative for S-100. Immunological markers are very important to make the diagnosis accurately. We have shown the prevalence of immuno-markers that were used in the histopathology for differential diagnosis of anorectal melanomas in table 1. As shown, S100 and HMB45 are positive in all cases tested, and cytokeratin is negative in all cases of anorectal melanomas. Immuno-staining with melanoma markers such as protein S-100, melanoma antigen HMB-45, and vimentin are important for final diagnosis in amelanotic melanoma, and most of the times it is positive. Whereas, positive staining for carcinoembryonic antigen (CEA), cytokeratin, epithelial membrane antigen (EMA or MUC1) is suggestive of adenocarcinoma.\textsuperscript{72,77} Canales and Blesa\textsuperscript{22} have reported that anorectal melanomas are strongly positive for vimentin, CD117 (C-kit), S-100 protein, and S-100,
Melan-A; however, HMB-45 stain was focally positive. Stains for cytokeratin, smooth muscle actin, desmin, h-caldesmon, chromogranin, synaptophysin, and CD34 were negative. The Ki67 proliferation index was 60%. In another case studied an anorectal melanoma was misdiagnosed as a GIST because it was positive for CD56, S100, and CD117; however, further confirmation with HMB-45 and Melan-A led to the correct diagnosis.

CONCLUSION

This is the first case of amelanotic anorectal melanoma reported from Saudi Arabia. Amelanotic anorectal melanomas spread by lymphatics to regional nodes, and hematogenously to the lungs and to other organs. Thickness of the tumor appears to be correlated with the clinical outcome in anorectal melanoma. Thickness of 4.0 mm or more is more prone to relapse, metastasis, and low disease-free survival rate. Thus, this case showed little promise of survival as the tumor was significantly large and she was discharged on her own will.

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ETHICAL APPROVAL

There is nothing to be declared.

CONFLICT OF INTEREST

The authors declare no conflict of interest related to this work.

REFERENCES


5. Jaimes N, Braun RP, Thomas L, Marghoob AA.

Table 1: Prevalence of Immuno-Markers in Anorectal Melanomas

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