Anorectal melanoma and seborrheic dermatitis—A case report

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SUMMARY

Introduction Anorectal melanoma (ARM) is a rare and aggressive neoplasm with predisposition for early infiltration, distant spread, and unfavorable prognosis. It has been speculated that Malassezia yeasts could possibly have an impact on skin carcinogenesis and development of melanoma, especially in patients with seborrheic dermatitis (SD), due to production of aryl hydrocarbon receptor (AhR) agonists. Case Outline A 52-year-old man with intensive SD complained of a four-month-long rectal bleeding, tenesmus, pain, and difficulty during defecation. On examination, a rectal tumor was detected and histopathology of tumor tissue revealed ARM with positive protein S100, melanoma antigen HMB45 and melan-A expression. After the diagnosis was established, abdominoperineal resection of the anus and rectum was performed, since the tumor was large, obstructive, and the anal sphincter was invaded. Conclusion Because of the possible impact of intensive SD to the cross-link between Malassezia yeasts AhR agonists and skin carcinogenesis, we discussed on this matter and reviewed the literature data regarding ARM. In addition to “pathogenic” and “non-pathogenic” Malassezia subtypes based on AhR agonist production, future studies on Malassezia metabolites, their carcinogenic effect in the skin and development of melanoma are needed. If the cross-link between Malassezia AhR agonists and skin carcinogenesis exists, timely prevention of ARM could be done with Malassezia eradication, especially in patients with severe SD. Keywords: anorectal melanoma; cancerogenesis; Malassezia; aryl hydrocarbon receptor; seborrheic dermatitis

INTRODUCTION

Anorectal melanoma (ARM) is a rare and aggressive neoplasm with predisposition for early infiltration, distant spread, and unfavorable prognosis, which represents 0.05–4.6% of all anal malignancies [1, 2]. Although being behind cutaneous and retinal, ARM, the third most common melanoma, accounts for 0.4–1.6% of all melanomas [2, 3]. ARM originates from melanocytes, the carcinogenic changed pigmented dendritic-like cells, which are present in all three mucosal zones of the anal canal [4]. Although solar ultraviolet radiation (UVR) is the most important carcinogen for melanocytes, not all melanomas can be attributed to ambient UVR, since some tumors arise on sun protected localizations, such as colon and anal region [5].

The literature data indicate possible link between melanoma, specific products of Malassezia yeasts, and aryl hydrocarbon receptors (AhR) [6, 7, 8]. Malassezia (Pityrosporum) species are lipophilic yeasts that are recognized as skin commensals capable of producing an extensive array of bioactive indolic compounds, such as malassezin, indolo[3,2-b]carbazole (ICZ), and indirubin, especially in pathogenic condition, e.g. seborrheic dermatitis (SD) [7]. Recent data show that the presence of different Malassezia species, the yeast density, and different enzyme production, could be important for Malassezia role in pathogenesis of different conditions and in different patient groups, especially those mediated by activation of AhR [7–10]. AhR are widely expressed in humans and are involved in various signaling pathways critical to normal cell homeostasis. Dysregulation of these physiological processes is known to contribute to skin tumor induction, by ‘initiation’, ‘promotion’, and ‘progression’, which represent different stages of the carcinogenic process. AhR are expressed in multiple tumor types, suggesting their pro-oncogenic role. It has been proposed that compounds produced by Malassezia yeasts are responsible for alterations in melanocytic function [7, 11]. Malassezia yeasts may produce enzymes important for melanoma progression making melanoma more sensitive to AhR-activation simultaneously. Therefore, AhR could play a central role in the ecological ‘cross-talk’ between Malassezia yeasts and development of melanoma in the upper part of the epithelium/epidermis.

CASE REPORT

A 52-year-old man with a four-month history of rectal bleeding, bowel obstruction symp-
Anamnestic data revealed untreated extensive long-standing SD. A huge anorectal tumor in the region of the dentate line was detected by clinical inspection (Figure 1). Biopsies demonstrated that it was a malignant melanoma, as protein S100, melanoma antigen HMB-45, and Melan-A expression were found. On computed tomography (CT) scanning, a circumferential anorectal thickening 3.2 cm above the anus, 7.2 cm in length and 1.8 cm in thickness, narrowing the rectal lumen, was detected, associated with several adjacent enlarged perirectal lymph nodes (Figure 2). CT scans of the thorax and abdomen showed no signs of distant metastases. In the inguinal region, no enlarged lymph nodes were detected.

After the diagnosis was established, abdominoperineal resection of the rectum was performed, since the tumor was large, obstructive, and the anal sphincter was invaded (Figure 3). The patient recovered well from the operation and was dismissed on the ninth postoperative day.

Histology analysis confirmed malignant metastatic epithelioid melanoma. The tumor showed invasion into the muscular layer. Only one of 14 removed perirectal lymph nodes was histologically positive on paraffin embedded sections. Melanoma was classified as stage 3. No adjuvant therapy was prescribed by the oncologist.

Eleven months after the initial surgery treatment was performed, the patient began to feel pain in the perineal region. Clinical examination showed palpable tumor mass in the perineal area. The 7.5 × 6.2 cm tumor was confirmed by magnetic resonance imaging and associated with enlarged lymph nodes in the inguinal region. The abdominal magnetic resonance imaging showed no signs of distant metastases. A second surgical procedure was carried out, showing recurrence of ARM in the small pelvis, with no infiltration of the prostate, urinary bladder or pelvic bones. The tumor was removed, the patient recovered well from the operation and was released on the tenth postoperative day and directly forwarded to a dermatologist to initiate treatment of SD and to an oncologist to initiate chemotherapy. Ten months after the second operation, the patient was disease-free, with SD in remission.

**DISCUSSION**

ARM is a rare disease with exceedingly poor prognosis, worse than cutaneous melanoma, with a median survival of one year and with five-year survival rate of 10% [12]. ARM could be located in the anal canal or at the anal verge (2/3), or in the distal rectum (1/3), and often presents with non-specific local symptoms such as rectal bleeding (55%), rectal masses (34%), anal pain (13%), and change in bowel habits (63%) [13, 14]. However, women are more likely than men to be diagnosed with ARM (male to female ratio is 0.75) [15], and lesions are usually non-specific, amelanotic, and commonly mistaken for benign conditions such as hemorrhoids or rectal polyposis, which prolongs the establishment of the diagnosis [16, 17].

Apart from UVR, the relevant impact of other factors on carcinogenesis in melanoma remains controversial. Exposure to solar UVR is the most important independent carcinogen for melanoma, but additional factors related to the host or local environment may contribute substantially to the pathogenesis of melanoma [18]. The host dependent factors and the polymorphisms of genes encoding for the biosynthesis of detoxifying enzymes are important for melanoma cases in different patient groups [19]. Therefore, UVR-independent anatomical distribution of
melanoma is in line with the current hypothesis that carcinogenic factors acting in loco independently or in addition to UVR, may play a distinctive role in the pathogenesis of ARM. In this setting, the important locally modifying factor in human skin carcinogenesis could be lipophilic yeasts of the genus *Malassezia*, previously known as *Pityrosporum*. The genus is characterized by the number of different species production of bioactive enzymes and production of certain metabolites [9, 18]. This is further in line with the observed coincidence of ARM and *Malassezia* colonization areas that are commonly found in some relatively UVR-protected areas, which are also locations of ARM [23]. Epidemiological observations that indirectly link *Malassezia* with melanoma come from the study of patients with Parkinson’s disease in whom SD and skin carcinomas appeared in higher rates than expected by chance, despite the fact that the most of other malignancies are found with reduced prevalence in this group of patients [9, 19]. The core mechanism by which *Malassezia* yeasts could probably mediate the carcinogenesis acts through the production of locally acting potent indolic AhR ligands [7]. The AhR belong to orphan receptor group that in adults mediate metabolism of diverse chemical and physical environmental damages to the cell, including UVR [21, 22]. Recent evidence indicates that AhR ligands are able to modify differentiation of keratinocytes in vitro and accelerate terminal skin development [23]. Moreover, AhR have pluripotent functions in adult human skin, e.g. they regulate many aspects of immune response, the cell cycle, and also part of the UVR induced cell alterations [21, 22]. The relationship of rapidly metabolized AhR ligands to carcinogenesis, including those naturally-occurring, like ICZ and indirubin, is more complex because they usually occur as mixtures affecting complex molecular interactions with potentially variable biological outcomes [7].

According to literature data, the presence of AhR ligands from *Malassezia* spp. in the skin could modify epidermal carcinogenesis and induce melanoma by (i) affecting the immune state of the skin; (ii) perturbing the homeostasis between cell cycle progression and apoptosis; (iii) increasing expression of matrix metalloproteinase-1 (MMP-1) in keratinocytes with consecutively enhanced tumor invasion; (iv) affecting survival of the initiated tumor cells; (v) modulating UVR induced carcinogenesis of the epidermis [7, 8]. Within the above framework, a key component of the proposed association to be dissected in future studies will be the behavior of *Malassezia* yeasts under UVR. The increased prevalence of SD among patients with parkinsonism also highlights the impact of the patients’ genetic profile, which is important in SD pathogenesis [9, 24].

Although there are various treatment strategies for ARM, surgical treatment remains the main therapeutic modality, with radical abdominoperineal resection (APR) and conservative wide local excision (WLE) as the most common [25]. The necessity of APR in the treatment of ARM beyond WLE remains controversial. Because of the low incidence of this disease, treatment guidelines are based on small retrospective studies. Ross with colleagues showed a 29% recurrence rate in primary ARM after APR, and a 58% recurrence rate after WLE [26]. In a study by Thibault et al. [13], 37 patients had curative resection with no significant survival difference between WLE and APR when comparing disease stage and five-year survival. Before 1997, approximately 70% of all patients underwent APR, while after 1997 almost 90% of patients underwent WLE [14]. The benefits of WLE are obvious, and include fast recovery time from a significantly less invasive procedure, minimal impact on bowel function, and avoidance of permanent colostomy [13]. WLE with or without adjuvant therapy seems to offer good locoregional control without reducing the survival rate, and may be a therapeutic modality of choice for patients with small, superficial ARM [14, 27]. APR should be offered to patients with obstructive bowel symptoms, locally advanced and invasive ARM, or as a salvage therapy following recurrence [27]. Moreover, the role of adjuvant chemotherapy and immunotherapy in the treatment of ARM has not been established, particularly in cases involving metastatic disease. Some patients with metastatic melanomas have been shown to respond to therapy with interferon-α and interleukin-2, suggesting that metastatic melanoma is susceptible to immune assault [28]. However, ARM has always been considered to be a radioresistant tumor [27], as radiation therapy provides no benefit, except occasionally for palliative care in cases with unresectable tumors [28].

In conclusion, besides surgery, the optimal treatment strategy for ARM requires multidisciplinary approach, taking into consideration the possible link between *Malassezia* yeasts and skin carcinogenesis, especially in patients with severe SD. Based on AhR agonist production, future studies are needed to determine a cross-link between ARM and *Malassezia* yeasts. If a cross-link between agonists (malassezin, ICZ, or indirubin) and carcinogenesis exists in patients with severe SD, timely prevention of melanoma and ARM could possibly be done with *Malassezia* eradication.

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Аноректални меланом и себореични дерматитис – приказ случаја

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КРАТАК САДРЖАЈ
Увод Аноректални меланом (АРМ) ретка је и агресивна неоплазма са израженом способношћу ране инфилтрације, удалеженог ширења и лоше прогнозе. Познато је да кваснице рода Malassezia имају способност продукције протеина, који представљају агонисте за арил хидрокарбон рецепторе (AhR), што се доводи у везу са њиховим могућим карциногеним потенцијалом и способношћу учешћа у развоју меланома коже, посебно код пацијената са себореичним дерматитисом (СД).

Приказ болесника У раду је приказан пацијент стар 52 године, мушкарац са израженим СД и симптомима аналног крварења, грчева, бола и отежане дефекције током четири месеца. Дијагностикован је тумор ануса, а патохистолошко испитање потврдило сумњу да се ради о АРМ са положним налазом антигена S100, меланома антигена HMB45 и меланоцитног антигена melan-A (MART1). По постављању патохистопатолошке дијагнозе начињена је абдомино-перинална ресекција ануса и ректума с обзиром на величину тумора, његову опструктивну природу и инвазију аналног сфинктера.

Закључак Због могуће повезаности СД и продуката гљива рода Malassezia са процесом карциногенезе у кожи и последичним развојем АРМ, у раду је дискуторан приказани случај уз преглед литературе. Подела на “патогене” и “непатогене” Malassezia врсте може да се доведе у везу са њиховим могућим потенцијалом да продукују агонисте AhR рецептора, због чега су неопходна даља испитивања. Уколико се потврди веза између ових агониста и канцерогенезе коже код пацијената са израженим СД, могуће је развијати стратегију превенције меланома правовременом ерадикацијом гљива рода Malassezia.

Кључне речи: аноректални меланома; канцерогенеза; Malassezia;aryl hydrocarbon receptor; себореични дерматитис

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