



A Rare Case of Primary Mucosal Melanoma of the Palate

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Abstract

Melanoma usually occurs on the skin. Only 1% of all melanomas affect mucosal membrane of the head and neck with 1951 cases reported from 1945 to 2011 in the world [1]. The article describes the case of the 65-year-old patient suffered from the palate melanoma that was irradiated on tumor area. The other therapeutic options for melanoma patients were discussed.

Keywords

Mucosal melanoma, Palate, Radiotherapy

Case Report

A 65-year-old man treated from skin psoriasis was referred to the oncology department by dermatologist with the suspicion of palate melanoma. Physical examination showed painless pigmented lesion present at mucous at the palate durum, palate mole, right buccal mucosa and right upper alveolus (Figure 1a). No enlarged regional lymphatic nodes were detected. Excisional biopsy was performed. Pathological examination proved mucosal lentiginous melanoma. Immunohistochemical analysis revealed that the neoplastic cells were positive for HMB-45, melan-A, S-100 and negative for AE1/AE3, confirming the diagnosis of melanoma.

Computed tomography revealed a thickening of mucosal membrane in the area of the palate durum, especially on the right side (Figure 2a). PET-CT demonstrated active process mainly in the right side of the palate and confirmed absence of any active regional nodes (Figure 2b). Chest X-ray examination excluded metastases in the lungs. The patient refused major surgical treatment, so he underwent up-front conventional 3D RT (X 6 MV) to the tumor area at the dose of 50 Gy in 20 fractions (2,5 Gy per fraction); nodes were not irradiated (Figure 3). The radiation acute side effects were assessed according to Dische scale. The intensity of the skin rash and oral mucositis did not accede 13 points (Figure 1b).

The physical examination performed soon after RT cessation, showed weaker saturation with the black color of the tumor, but no decrease in diameter was described. Two months after RT, the melanoma disappeared from the mucous of the right cheek and alveolus and infiltration was slightly reduced at the periphery of the palate. The extent of the lesion was stable also 3, 4, 5 months after RT, however intensity of the black color was further decreasing (Figure 1c). The melanoma lesion decreased by one third and stabilized one and a half year after RT. Unfortunately, 1 year and 7 months after RT cessation progression of the disease appeared on right alveolus causing two slightly painful, non-healing ulcers. Additionally, new intensive black spots appeared on the palate. Two months later the disease further progressed - a thick tumor was found on the palate, and more black spots on the mucous of the cheek and alveolus developed (Figure 1d). The patient sturdily refused chemotherapy because of a fear of adverse events. The recurrent melanoma was

Introduction

Primary melanomas located in the oral cavity constitute as low as 0,2-8,0% of all melanomas [1] predominantly affected palate and maxillary gingiva [2]. Less frequently oral melanoma (OM) can be found in mucosal epithelium, stria vascularis of inner ears, retina, and uveal tract. Oral melanoma (OM) has been reported in patients aged 20 to 80 years and has a male predilection. The etiology of OM is unknown. Possible risk factors of these tumors may be tobacco use, chronic irritation from ill-fitting dentures as well as factors associated with environmental pollution [3]. It was suggested that mucosal melanomas may arise from both pigment-producing cells and Schwann cells found in the mucous membrane [4]. In case of OM genetic aberrations are less common, than in cutaneous melanoma. The most often occurred aberrations are those, concerning *ckIT* gene (20%), *NRAS* gene (5%) and *BRAF* gene (3%) [5]. Even 15.6-39% of OM patients have *KIT* aberration or more copies of *KIT* gene [6,7]. The clinical presentation of this neoplasm can vary widely if it comes to intensity of black saturation, proliferation rate and metastatic potential [8].

OM is characterized by frequently delayed diagnosis and beginning of subsequent treatment, because most mucosal melanotic lesions are painless in their early stages. These facts worsen prognosis of patients suffered from OM [8]. 5-year survival rate of OM patients amounts 17%, although early recognition and treatment significantly improve the prognosis [9,10].

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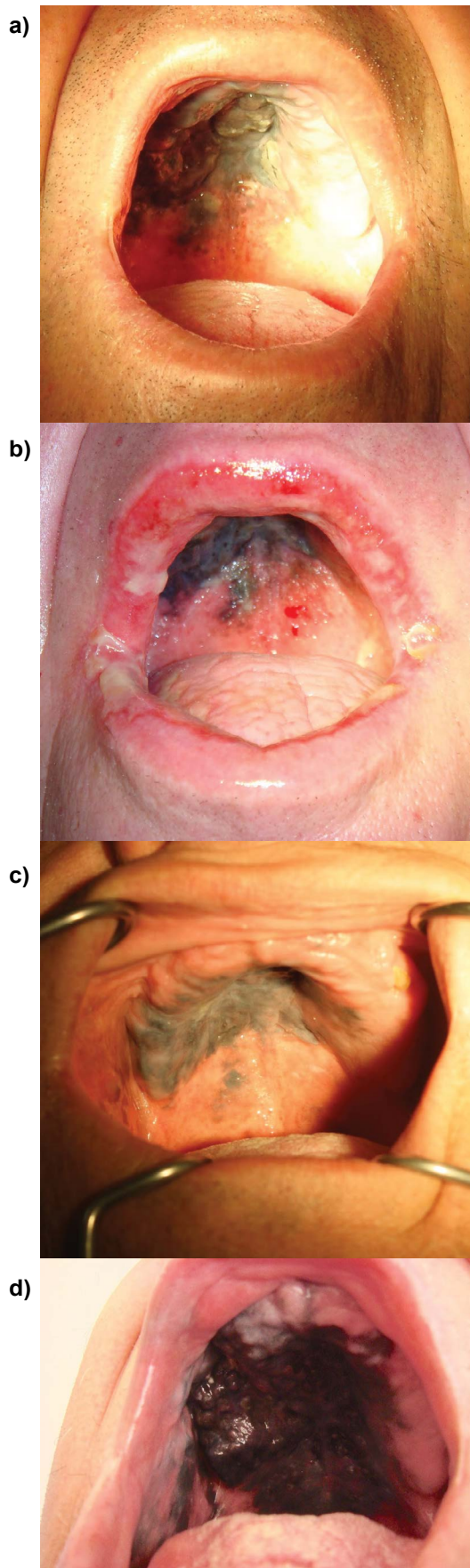


Figure 1: Photographic documentation of the clinical status of the oral melanoma patient.

a: Pre-treatment clinical state of the patient (prior to radiotherapy) - melanoma of the palate; **b:** Acute radiation reaction of the mucous of the oral cavity; **c:** Clinical status of the patient 5 months after radiotherapy cessation. Oral mucous melanoma decreased and stabilized, and was less saturated with the black color; **d:** Clinical status 1 year and 9 month after radiotherapy cessation. Progression of the oral mucosal melanoma: thick, dark tumor is seen on the palate.

slowly, but constantly progressing, which was clinically observed and documented on CT examination. Nearly 3 years after RT cessation, the patient suffered from pain in the tumor area. Chest X-rays revealed suspicious lesions in the right lung (Figure 2c) while head and neck CT revealed enlargement of mucosa infiltration of right alveolus and palate. Tumor exceeded midline and caused osteolysis of right alveolus as well as palate durum (Figure 2d). This time patient signed informed consent to undergo monochemotherapy with dacarbazine and was referred to the regional chemotherapy center located nearest of his home.

Discussion

Mucosal melanoma of the oral cavity is unusual malignancy. There are no specific clinical symptoms of OM, so it may be difficult to distinguish on a clinical basis from solitary oral melanotic macules, labial lentiginos, foreign bodies, or individual pigmentation. The OM cases misdiagnosed with racial pigmentation [11] or pyogenic granulomas [12] were also described. Furthermore it is not easy to differentiate mucosal melanoma from a metastatic melanoma [13]. Additionally the local inflammation may complicate the assessment of infiltrated areas [14]. There are some pathological disturbances that should be differentiated with OM, such as: other forms of pigmented oral diseases, including drug-, disease- or smoking-associated melanosis, oral melanotic macule, Kaposi's sarcoma, melanocytic nevus, melanoacanthoma, amalgam tattoo, vascular blood-related pigments, oral melanotic macule, Peutz-Jeghers syndrome, Addison disease, and drug-induced pigmentation [13].

OM occurs the most commonly in the palate, which is in close anatomic proximity to the most prevalent site for head and neck melanomas - the nasal cavity and maxillo-ethmoid massif [4,10]. The first symptoms of OM may not rivet patient's attention, because there may be some asymptomatic swelling and occasional bleeding. OM may be noticed by the way of treating different illnesses, because most of them are painless in their early stages. The diagnosis of OM is often unfortunately delayed until some symptoms resulting from ulceration, growth, or bleeding reveal. The pain is usually the late manifestation of melanoma infiltration [3].

Data regarding OM are scanty. Majority of information about OM comes from case reports, not from randomized clinical trials [2,8,12,13,15]. That is why the clear guidelines for treatment of melanoma in this location are lacking. Therapy of OM is mainly based on surgical excision of the primary tumor [2,8,10,12,13,] which may be supplemented by adjuvant RT [2,15-17]. Adjuvant RT does not appear to affect survival but seems to improve locoregional control thus is recommended by majority of experts in OM patients [10,15]. There are limited data proving efficacy of chemotherapy and immunotherapy [18,19]. Unfortunately the OM is often inoperable, because of a large size of the lesion or localization which is unfavorable for the excision [20].

The described patient did not assent to any surgical treatment, thus was offered up-front RT. The physical examination as well as images did not indicate nodal spread that is why the irradiated fields were limited to the tumor area. The irradiation of the clinically negative neck is controversial [15]. Some authors recommend nodal irradiation because of the high risk of subclinical disease [15,20]. However, it is assumed that about 80% of OM patients have local disease, and only 5-10% - nodal spread at the time of diagnosis [20]. The poor prognosis of even early-staged OM patients results from occult metastases at presentation in the majority of patients [20]. The fractionation schedule varies between the authors from 1,5 Gy to even 13,8 Gy per fraction giving the total dose ranging from 32 to 76 Gy [2,10,15,16]. Taking into consideration the biology of melanoma and palate location 2,5 Gy per fraction was chosen. Radiation therapy is rarely used as a primary modality in OM patients - in patients with inoperable tumour, in medically compromised patients and in the older population [21]. Although cutaneous melanomas present weak radio sensitivity, patients may occasionally have good response to RT, especially in the case of early melanomas or in melanomas *in situ* [22]. The data from literature indicate that in some cases the definitive RT

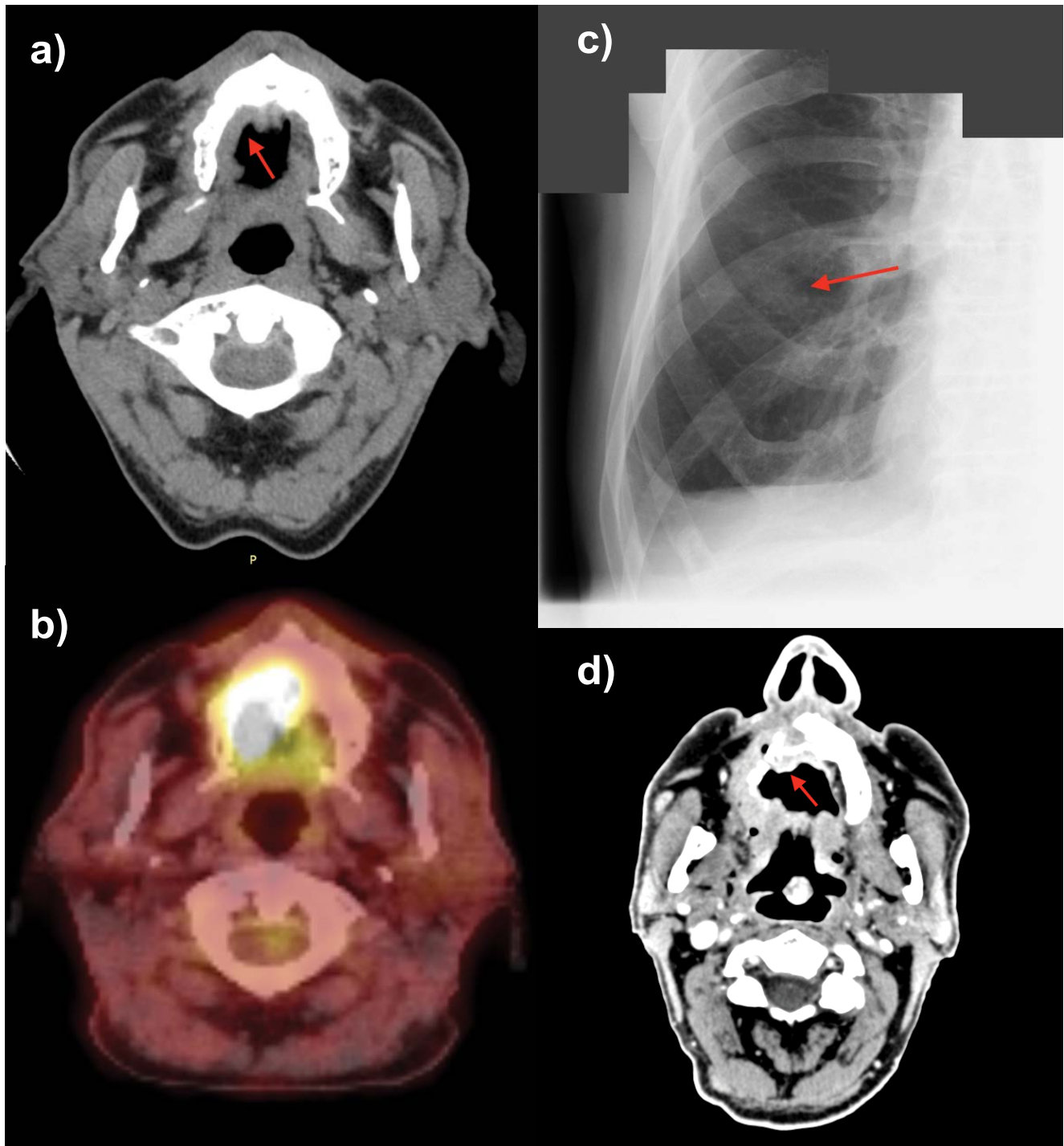


Figure 2: Imaging studies (CT, PET-CT, RTG) of oral melanoma patient in pre-treatment period and during the treatment.

a: CT scan of patient's head performed prior to the beginning of the radiotherapy. Red arrow indicates mucosal melanoma of the palate; **b:** PET-CT scan. ¹⁸F-FDG intake by oral mucosal melanoma indicates increased metabolic activity mainly in the right palate; **c:** Chest RTG. Metastatic lesion in the right lung indicated by red arrow; **d:** CT scan of patient's head, 2 years and 9 months after radiotherapy cessation. Red arrow indicates osteolysis of maxilla (local progression of the disease).

may result in survival rates comparable to that received by surgery, however, the efficacy of definitive RT in III and IV stage of the disease is poor reviewed in [2,15,16]. The possible late complications of the RT as mucosal ulcer, radiation necrosis or bone exposure are another limiting factors of definitive RT [15,16]. There are single reports that definitive particle radiation therapy promises to provide high rates of local control for OM inoperable patients. Unfortunately, overall survival rate does not extend significantly after particle radiotherapy treatment [23].

To date, separate small studies demonstrated, that adjuvant chemotherapy and immunotherapy may prevent distant metastases formation in the subset of OM patients improving also survival rates [19,23-25]. Recent molecular evidence suggests that proto-oncogene

KIT aberrations determined in advanced head and neck melanoma patients may represent a potential diagnostic value and serve as a therapeutic target for tyrosine kinase inhibitors also in an adjuvant setting of OM patients [1].

Recurrences of the disease may occur even 10-15 years after primary therapy of OM patients. Distant metastases to the brain, liver, lungs, and bones are frequently observed as well. Unfortunately, most patients die because of the progressive disease in 2 years [24,25]. Systemic therapy is the management of choice in case of recurrent or metastatic OM. Dacarbazine treatment of cutaneous melanoma patients increased overall survival by 5,6-7,8 months, however in OM patients dacarbazine alone was not effective [24]. Combination of dacarbazine with interleukin-2 (IL-2) is unsatisfactory as well [24,25]. Increase of

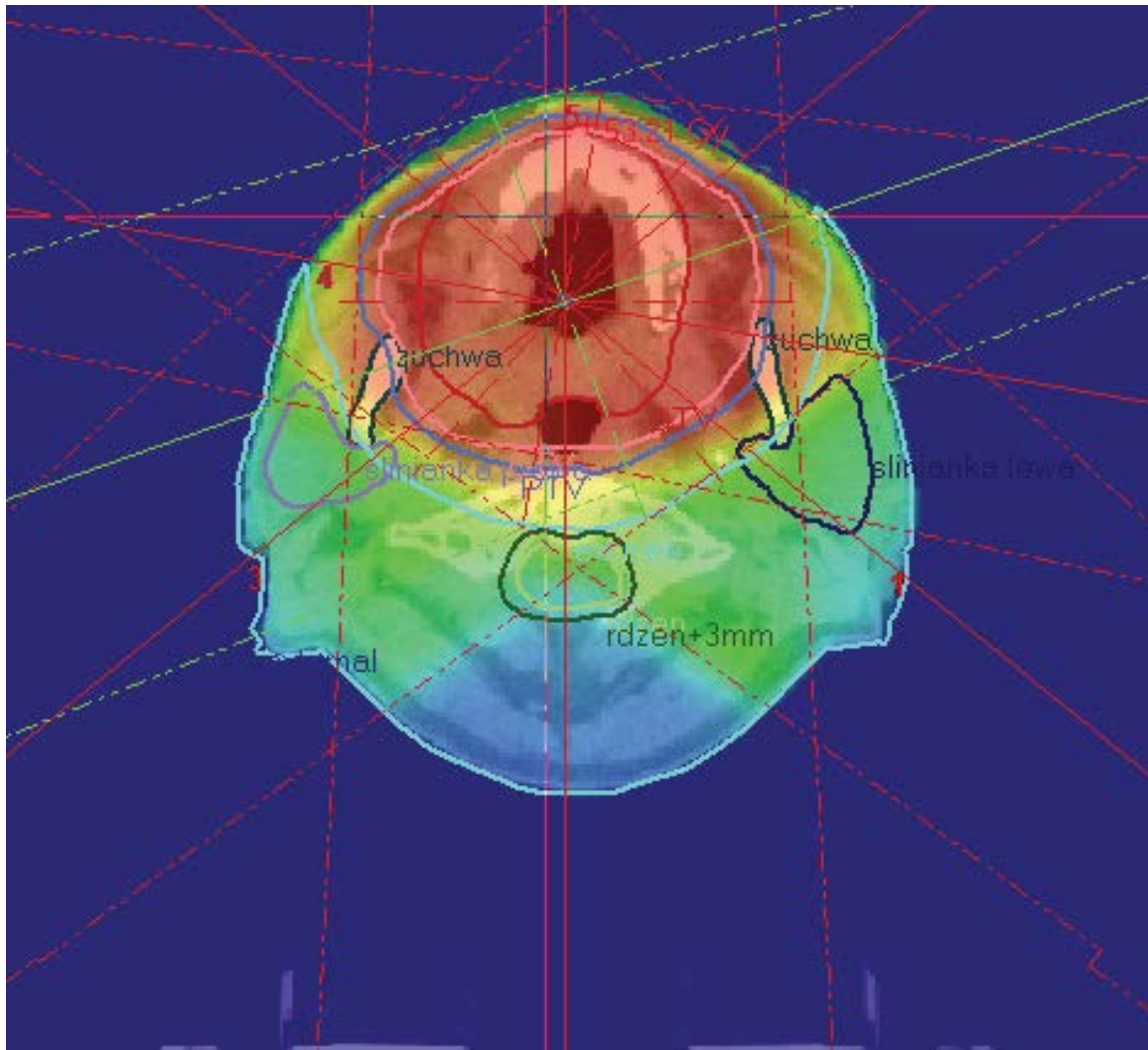


Figure 3: Oral mucous melanoma patient underwent up-front conventional megavoltage 3D RT (X 6 MV) to the tumor area at the dose of 50 Gy in 20 fractions (2,5 Gy per fraction); nodes were not irradiated. Red isodoses indicate prescribed high dose of radiation.

survival of melanoma patients occurred with the introduction of targeted therapies. Administering of ipilimumab for advanced melanoma patients increased overall survival from 6.4 months (melanoma peptide vaccine group) to 10.1 months [26,27]. This human IgG1 monoclonal antibody (MoAb), by inhibiting the activation of CTLA-4, enhances T-cell proliferation and T-lymphocytes' dependent antitumor response [26,27]. Based on results of clinical trials, in 2011, the Food and Drug Agency (FDA) approved ipilimumab for treatment of patients with inoperable or metastatic melanoma [26,27]. Unfortunately, at the time when the patient was treated due to progressive disease, there was no administrative possibility to qualify him to ipilimumab, which was only registered for cutaneous melanoma patients. Nowadays, in Poland, targeted therapy is refunded both for cutaneous and oral melanoma patients and is offered for them.

The review of literature indicates that to improve treatment efficacy of OM patients both early detection [28] as well as a lifelong follow-up [20] are of paramount importance in the management of this aggressive disease.

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