Locally aggressive giant basal cell carcinoma of the head with focus on surgical treatment– a case report

Lokalnoagresivni gigantski bazocelularni karcinom vlasišta s posebnim osvrtom na kirurško liječenje – prikaz slučaja

Fatima Juković-Bihorac, Hakija Bečulić, Emir Begagić, Rasim Skomorac*

———— Summary –

Introduction: Basal cell carcinoma is the most common non melanoma skin cancer. It accounts for approximately 80% of all skin cancers.

Case report: We presented a 62-year-old patient with a giant, deeply infiltrative, destructive lesion of the head which lasted for 15 years. Microbiological analysis showed contamination, computed tomography (CT) scan showed deep infiltration and bone destruction. The risks of operative treatment were numerous. The treatment was more complicated by infection, infiltrative spreading and the patient's comorbidity. Regardless, we decided on a wide surgical resection with pathohistological evaluation of the resection margins. The patient denied oncological treatment. After 10 years, there were no recurrent tumours.

Conclusion: Aggressive surgical treatment is the treatment of choice for giant basal cell carcinoma. In the case of giant locally aggressive and advanced neoplasms, when surgery is not appropriate or not possible, medical treatment becomes oncological.

Key words: non melanoma skin cancer, basal cell carcinoma, giant skin carcinoma

Uvod: Bazalno-stanični karcinom je najčešći nemelanomski maligni tumor kože. U prosjeku čini 80% svih malignih kožnih tumora.

Prikaz slučaja: Predstavljamo bolesnika u dobi od 62 godine, s gigantskom, dubokoinfiltrativnom, destruktivnom lezijom glave, koja je trajala 15 godina. Mikrobiološke analize pokazale su infekciju, a CT nalaz duboku infiltraciju i destrukciju kosti. Rizici operativnog tretmana bili su brojni. Tretman je dodatno bio kompliciran infekcijom, dubokom infiltracijom i komorbiditetima bolesnika. Odlučili smo se na široku kiruršku resekciju s patohistološkom evaluacijom resekcionih margina. Bolesnik je odbio onkološko liječenje. Nakon 10 godina recidiva tumora nije bilo.

Zaključak: Kirurški tretman je tretman izbora gigantskih, bazalno-staničnih karcinoma. U slučaju neoperabilnih gigantskih, lokalnoagresivnih i uznapredovalih bazalno-staničnih karcinoma, tretman izbora je onkološka terapija.

Ključne riječi: nemelanomski maligni tumor kože, bazocelularni karcinom, gigantski karcinom kože

Med Jad 2023;53(3):219-224

* Cantonal Hospital Zenica, Department of Pathology, Zenica, Bosnia and Herzegovina (Fatima Juković-Bihorac,MD); Cantonal Hospital Zenica, Department of Neurosurgery, Zenica, Bosnia and Herzegovina (PhD Hakija Bečulić, MD; PhD Rasim Skomorac, MD);

University of Zenica, Medical Faculty, Department of General Medicine (Emir Begagić)

Correspondence address/ Adresa za dopisivanje: Fatima Juković-Bihorac, MD, Cantonal hospital Zenica, Department of pathology, Crkvice 67, 72000 Zenica, Bosnia and Herzegovina E-mail: fatima.bihorac@live.com

Received/Primljeno 2022-10-29; Revised/Ispravljeno 2023-06-21; Accepted/Prihvaćeno 2023-08-17

Introduction

There are 2 main types of skin malignancies: melanoma and nonmelanoma skin cancer. Non melanoma skin cancer includes basal cell skin cancer, squamous cell skin cancer and other rare types.¹ Basal cell carcinoma (BCC) is the most common nonmelanoma skin cancer. The incidence of basal cell carcinoma is increasing. It is accounting approximately 80% of all skin cancers.¹ This tumor is most associated with uv radiation. Solar UVB radiation generally is considered to be the most important risk factor for the development of BCC. The other risk factors are light skin color, red or blonde hair color, and a family history of melanoma or sunburn reactions.²

The tumor is slow growing but invasive and locally aggressive. Its metastases are very rare and have been reported in between 0.0028 and 0.55 of all BCC cases.^{1,3} BCC is classified into several basic types of which it is the most common nodular, superficial, nodulocystic, morpheic, metatypical, pigmented, and ulcerative.⁴

Nodular and superficial type of BCC have a low risk for local recurrences, but morpheic, infiltrating and metatypical type are very destructive and with higher recurrence rate. That division of BCC is of big clinical importance.⁵

Giant basal cell carcinoma (GBCC) is a BCC which is larger than 5cm in diameter. The occurrence rate of GBCC is approximately 0.5–1% out of all BCC types.⁶ Most often BCC are small tumors, mostly localized in the head and neck region.⁷

In the treatment of giant locally aggressive and advanced neoplasms, metastatic or recurrent, when surgery is not appropriate or not possible, medical treatment becomes oncological. The most common treatment is oral hedgehog pathway and check point inhibitors.⁸

There are two ways of hedgehog pathway treatment, which are approved by FDA.^{8,9} Vismodegib is the first hedgehog inhibitor approved by the FDA in 2012. An objective response rate (ORR) was 47.6% for locally advanced BCC and 12 months.^{9,10} metastatic BCC at 30% for Sonidegib is the second oral HHI for the treatment of BCC, approved by the FDA in 2015. It is indicated for the treatment of locally advanced BCC that has recurred after surgery or radiation treatment or for the patients who are not candidates for surgery or radiation therapy. The research revealed an ORR of 56.1% with a median duration of response of 26.1 months and in 93.2% a 2-year survival rate for locally advanced BCC. An ORR of 7.7% was reported for metastatic BCC.¹¹

Immunotherapy of the advanced, non-melanoma skin cancer is gaining importance, because their high mutational burden index is high (MBI). Two anti-PD-1 antibodies, cemiplimab and pembrolizumab are approved.¹²

PD-1 inhibitors produce persistent effects, with improved survival. Patients are capable to stop with therapy while they have benefits from treatment. PD-1 inhibitor therapy is well tolerated and safe among patients specifically with comorbidity.¹³

Several case reports with anti-PD-1 agents¹⁴⁻¹⁶ and anti-CTLA-4 therapy16 revealed good responses in disease. Seven patients advanced received pembrolizumab plus vismodegib and nine patients received only pembrolizumab.¹⁷ The ORRs were 44% and 29% at 18 weeks, respectively. The authors concluded that combination therapy is not much better than monotherapy. The usage of pembrolizumab has been registered in 5 case reports more with complete or partial responses.¹⁴⁻¹⁷

We present the case of locally aggressive and deep penetrating, extremely big basal cell carcinoma, located in the hairy part of head. The treatment options of such highly risked tumors must be carefully considered.

Case report

The Ethics Committee of the Zenica Cantonal Hospital allowed the data used in this case report and the patient gave us informed consent for the data and figures we used in this article also.

A 62-year-old patient arrived by car of the Emergency Medical Service to the Department of Emergency Medical Assistance (UB). On admission to UB he was somnolent, disoriented in time, space and towards persons, gave no answers to questions, could not perform more complicated requests, without pronounced lateralization in neurological findings, febrile (TT 39° C), eupnoic, normotensive, tachycardic (f 100/min). We obtained heteroanamnestic data that the patient had had a "head wound" for 15 years, which was bleeding occasionally. He bandaged the wound and hid it with a scarf. He had not consulted a doctor about this problem earlier.

He has been a diabetic on insulin therapy for many years. In the local finding we found a large ulcerated, deeply infiltrated tumor of the head, about 11cm in diameter, with purulent contents at the bottom, necrotic and gangrenous edges (Figure 1).

At the UB, we cleaned and bandaged the tumor, took wound swabs and made a computed tomography (CT) scan of the head. CT showed extensive and deeply infiltrative lesion that destroys the bone (Figure 2).



Figure 1 Preoperative image of the extensive and ulcerative BCC of the scalp (Department of Neurosurgery Zenica Cantonal Hospital) Slika 1. Preoperativna slika ekstenzivnog i ulcerativnog BCC vlasišta (Odjel Neurokirurgije Kantonalne bolnice Zenica)



Figure 2 CT scan of the head (bone window) (Department of Neurosurgery Zenica Cantonal Hospital) Slika 2. CT snimak glave (prozorzakost) (Odjel Neurokirurgije Kantonalne bolnice Zenica)

After receiving adequate primary care and diagnostic processing at the UB, we transferred the patient to the Intensive Care Unit (ICU).

At the ICU, we connected the patient to vital functions monitoring and observed him intensively. In the laboratory findings, we found hypocalcemia, hypochromic anemia, metabolic ketoacidosis, and the level of glucose in the blood was from 8.5 to 12.5. Microbiological analysis of the wound swab showed contamination with *Staphylococcus aureus*. During the stay at the ICU, we prescribed appropriate antibiotic therapy, according to the antibiogram, and appropriate symptomatic therapy.

After the patient was stabilized, we started the surgery procedure. Following an appropriate positioning of the head, washing and limiting the operative field, we excised the tumor-infiltrated part of the skin. Then we osteoclastically removed the tumor-affected and severely softened bone down to the healthy tissue. We gradually removed the upper layer of the dura mater, which was also thickened and infiltrated by the tumor. After that, we opened the dura mater parasagittally on the left and allowed the spontaneous release of copious purulent content from the subdural space. We additionally washed it with saline and antibiotic solution. After establishing adequate hemostasis, we closed the defect with local transpositional flaps taken from the frontal and occipital regions. We covered the donor defect in that area with epidermal grafts according to Tirsch, which we took from the back of the left upper leg and lower leg. We did not reconstruct the calvaria and dura.

Postoperatively, we referred the patient to the ICU for postoperative treatment. In the further course, the flaps were without local signs of necrosis, as well as the epidermal graft. The patient was vitally stable, subjectively felt well. On the second day postoperatively, he was conscious, oriented, communicative, afebrile, eupnoic and the wounds were properly healed.

On the pathohistological examination we got the diagnosis of a giant basal cell carcinoma – metatypical type, with keratin pearls and infiltrative peripheral border. Locally aggressive, deeply infiltrated, ulcerated mass, resected in its entirety with clear resection margin at the end (Figure 3A-C). Separately delivered dura and bone were tumor-infiltrated with free resection margins at a distance of 5 mm at least.

After 15 days, the patient was discharged from the hospital to home treatment and care. About 30 days after the operation, the patient was ordered for a follow-up examination. During the control examination, the patient was in a normal state of consciousness and neurological findings. In the local findings there were no signs of infection and necrosis (Figure 4).



Figure 3 This picture of a giant basal cell carcinoma demonstrating A) ulcerated surface, B) invasion of the fat by highly infiltrated trabeculae of basaloid cells, with the absence of peripheral palisading (C) and the area of squamous differentiation. Note the formation of keratin pearls (C) (H&E, original magnification ×20)
(Department of Pathology Zenica Cantonal Hospital)
Slika 3. Slika gigantskog bazocelularnog carcinoma koji pokazuje: A) ulceriranu površinu, B) invaziju masnog tkiva visoko infiltrativnim tračcima bazaloidnih stanica, bez izraže neperiferne palisadizacije C) aree skvamozne diferencijacije i keratinske perle (HE, ×20) (Odijel patologije Kantonalne bolnice Zenica)

The patient was recommended further treatment. But, due to his good physical condition, the patient refused oncological treatment. The locoregional status was satisfactory. Ten years after discovering the BCC lesion, the patient died due to a heart attack.



Figure 4 Appearance of the patient 30 days after the operation (Department of Neurosurgery Zenica Cantonal Hospital) Slika 4. Izgled pacijenta 30 dana nakon operacije (Odijel Neurokirurgije Kantonalne bolnice Zenica)

Discussion

Typical BCC lesions are small tumors with indolent behavior, located mostly at the skin which is exposed to the sun and UV light.²

The treatment of the BCC depends on the tumor characteristics which are based on their prognostic factors. The most important prognostic factors are size, site, tumor subtype, status of resection margins and prior treatment. In that regard, BCC are categorized in low and high risk tumor.¹⁸

The worst impact on tumor prognosis has large, long-standing lesions, infiltrating, morpheic or metatypical pathohistological subtype with an aggressive way of growth in the deeper lesion layers. Also, on the choice of treatments, a great influence is the medical condition of the patient, coexisting disease, and current medication. In addition, experience and the ability of the physician are undoubtedly significant.¹⁸ The treatment of these large-dimensional tumors could be surgical and non-surgical. Surgical treatments are surgical resection, Mohs micrographic resection, radiotherapy and chemotherapy in unresectable tumors.^{18,19}

Tumors located in the head area, especially large tumors, are in themselves classified as high-risk tumors²⁰. High risk tumors require a more careful approach, primarily due to the volume and severity of the resection, but also the completeness of the excision. An aggressive surgical resection with tumor-free resection margins offers the best results and healing. The cure rates are averaging from 90% to 91%.²⁰

In one study, the age of the patient, duration of the lesion and duration of the treatment were not significantly connected with the increased risk of relapse.²¹

Our patient arrived in our hospital's intensive care unit in a serious condition, with deep, infiltrative lesion on the head which had lasted for the last 15 years. The case was complicated by staphylococcal infection of the tumor surface in subdural space, and tumor infiltration of the dura of the upper sagittal sinus.

The risks of operative treatment were reflected in the possibility of intraoperative damage and thrombosis of the upper sagittal sinus, sepsis, damage of intracranial structures, liquefaction and postoperative necrosis of transplanted skin flaps. The case was more difficult due to the insulin-dependent diabetes.

There was a risk of closing the resulting defect with a skin flap due to the possibility of its infection and necrosis. Usually, the flaps are not applied to the infected substrate immediately but after a few days, when there is certainty that the infection has been cured.²² Wide excision of GBCC of the head and neck can consequently have large complex of soft tissue and bone defects and even in brain exposure. How the defect will be reconstructed depends on the extent of the defect. The large and complex defect in the head and neck area can be properly covered by the selection of free, regional, or local flaps.²³ Nevertheless, after extensive removal of the tumor mass, removal of necrotic tissue and washing of the subdural space with an antibiotic solution, we decided to close the defect. The reason for such a decision was the general poor condition of the patient, the possibility of superinfection in an open wound with a more resistant strain, its spread towards the intracranial space, cerebrospinal fluid (CSF) and good blood circulation in the head. Our procedure, considering the postoperative course and rapid recovery of the patient, proved to be justified. After 10 years, there were no recurrent tumors and patient died due to a heart attack.

Metastatic BCC is very rare. It has been described in bone, lungs and axillary lymph nodes.^{24,25} In the case of BCC with pulmonary metastasis, after surgery, sonidegib, a hedgehog signaling inhibitor, was used as a first-line treatment and cemiplimab in the second act.²⁴

In the treatment of giant locally aggressive and advanced neoplasms, metastatic or recurrent after surgery treatment and radiotherapy, then in case surgery is not appropriate or not possible, medical treatment becomes just oncological. The most common treatment is oral hedgehog pathway and check point inhibitors.⁸

Further investigation of immunotherapy, especially of PD-1inhibitors is needed to define its optimal role for patients with this disease.

References

- 1. Mehta KS, Mahajan VK, Chauhan PS, et al. Metastatic Basal cell carcinoma: a biological continuum of Basal cell carcinoma?. Case Rep Dermatol Med. 2012;2012:157187.
- Mott SE, Hunter WJ, Silva E, Huerter CJ. Approach to management of giant basal cell carcinomas. Cutis. 2017;99:356-362.
- 3. Seo SH, Shim WH, Shin DH, Kim YS, Sung HW. Pulmonary metastasis of Basal cell carcinoma. Ann Dermatol. 2011;23:213-216.
- 4. McKee PH, Calonje J, Lazar A, et al, eds. Pathology of the Skin with Clinical Correlations. 4th ed. Vol 2. Philadelphia, PA: Elsevier Mosby; 2011.
- Elder DE. Basal cell carcinoma. In: Elder DE, Elenitsas R, Johnson Jr BL, et al, eds. Lever's Histopathology of the Skin. 10th ed. Philadelphia, PA: Lippincott Williams & Wilkins; 2009;826-832.
- 6. Cantisani C., Rossi R., Nisticò SP. et al. Management of patients with giant basal cell carcinoma during SARS COV2 outbreak in Italy. Transl Biophotonics 2022;\$:e20200009.
- 7. Rubin AI, Chen EH, Ratner D. Basal-cell carcinoma. N Engl J Med. 2005;353:2262-2269.
- Ahmed M, Muradashvili T, Soliman S, Ghaly M. Metastatic insidious super giant basal cell carcinoma. BMJ Case Reports CP 2022;15:e249873.
- Sekulic A, Migden MR, Basset-Seguin N, et al. Longterm safety and efficacy of vismodegib in patients with advanced basal cell carcinoma: final update of the pivotal ERIVANCE BCC study. BMC Cancer. 2017;17:332.
- 10. Sekulic A, Migden MR, Lewis K, et al. Pivotal ERIVANCE basal cell carcinoma (BCC) study: 12-month update of efficacy and safety of vismodegib in advanced BCC. J Am Acad Dermatol 2015;72:1021-1026.
- 11. Lear JT, Migden MR, Lewis KD, et al. Long-term

efficacy and safety of sonidegib in patients with locally advanced and metastatic basal cell carcinoma: 30month analysis of the randomized phase 2 BOLT study. J Eur Acad Dermatol Venereol 2018;32:372–81

- 12. Ascierto PA, Schadendorf D. Update in the treatment of non-melanoma skin cancers: the use of PD-1 inhibitors in basal cell carcinoma and cutaneous squamous-cell carcinoma. J Immunother Cancer 2022;10:e005082.
- 13. In GK, Nallagangula A, Choi JS, et al. Clinical activity of PD-1 inhibition in the treatment of locally advanced or metastatic basal cell carcinoma. J Immunother Cancer 2022;10:e004839.
- 14. Ikeda S, Goodman AM, Cohen PR. Metastatic basal cell carcinoma with amplification of PD-L1: exceptional response to anti-PD1 therapy. NPJ Genom Med 2016;1:16037.
- 15. Lipson EJ, Lilo MT, Ogurtsova A. et al. Basal cell carcinoma: PD-L1/PD-1 checkpoint expression and tumor regression after PD-1 blockade. J Immunother Cancer 2017;5:23.
- 16. Cannon JGD, Russell JS, Kim J, Chang ALS. A case of metastatic basal cell carcinoma treated with continuous PD-1 inhibitor exposure even after subsequent initiation of radiotherapy and surgery. JAAD Case Rep 2018;4:248–50.
- 17. Chang ALS, Tran DC, Cannon JGD, et al. Pembrolizumab for advanced basal cell carcinoma: an investigator-initiated, proof of-concept study. J Am Acad Dermatol 2019;80:564–6.

- 18. Telfer NR, Colver GB, Morton CA; British Association of Dermatologists. Guidelines for the management of basal cell carcinoma. Br J Dermatol 2008;159:35-48.
- Peris K, Fargnoli MC, Garbe C. et al. Diagnosis and treatment of basal cell carcinoma: European consensusbased interdisciplinary guidelines. Eur J Cancer 2019;118:10-34.
- 20. Dubin N, Kopf AW. Multivariate risk score for recurrence of cutaneous basal cell carcinomas. Arch Dermatol 1983;119:373-377.
- 21. Silverman MK, Kopf AW, Bart RS, Grin CM, Levenstein MS. Recurrence rates of treated basal cell carcinomas. Part 3: Surgical excision. J Dermatol Surg Oncol 1992;18:471-476.
- 22. Faruk Konjihodžić: Udžbenik hirurgije, Sarajevo: NIR, 2001; 335-6.
- 23. Arslan H, Güzel MZ, Cnar C.Treatment of giant basal cell carcinomas of the head and neck with aggressive resection and complex reconstruction. J Caniofac Surg 2012;23: 1634–1637.
- 24. Fordham SA, Shao EX, Banney L, Azer M, Dettrick A. Management of basal cell carcinoma with pulmonary metastasis. BMJ Case Rep 2023;16:e251700.
- 25. Li R, Lee G, Huang M, El-Sherief A. Rare basal cell metastasis of a basal-squamous skin collision tumour to the lung and axillary lymph node. BMJ Case Rep 2019;12:e231487.