

Contents lists available at ScienceDirect

# Saudi Journal of Biological Sciences

journal homepage: www.sciencedirect.com



# Original article

# Microcystic adnexal carcinoma of the scalp treated with surgical resection along with chemoradiation: A case report and review of the literature



Rolina K. Al-Wassia a,\*, Marwah S. Al-Oathmi b

- <sup>a</sup> Department of Radiology, Faculty of Medicine, King Abdulaziz University, Jeddah, Saudi Arabia
- <sup>b</sup> Department of Otolaryngology Head and Neck Surgery, King Abdulaziz University, Jeddah, Saudi Arabia

#### ARTICLE INFO

# Article history: Received 17 April 2021 Revised 31 July 2021 Accepted 1 August 2021 Available online 5 August 2021

Keywords: Microcystic adnexal carcinoma Skin cancer Wide local excision Radiotherapy

#### ABSTRACT

Microcystic adnexal carcinoma (MAC) is an infiltrative rare cutaneous neoplasm for which there are no consensus management guidelines because of the paucity of evidence-based practice; hence, the utility of their management is based only on previously published case reports. We report a case of a scalp lesion that was successfully treated using a combination of surgical resection, chemotherapy, and radiotherapy. © 2021 The Author(s). Published by Elsevier B.V. on behalf of King Saud University. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

# 1. Background

Microcystic adnexal carcinoma (MAC) is an infiltrative rare cutaneous neoplasm for which there are no consensus management guidelines because of the paucity of evidence-based practice; hence, the utility of their management is based only on previously published case reports (Waqas et al., 2017; Tanese et al., 2019). MAC usually presents as a firm dermal papulonodule or flesh-colored plaque on the head or neck. The deeply penetrative growth pattern, coupled with the long latency of MAC, poses a significant diagnostic challenge to histopathologists and surgeons. Tissue invasion may extend beyond the visible clinical margins of the mass seen on the skin surface and may also invade the underlying subcutis or even deeper into soft tissue hence, local recurrence is common. A recent systematic review recommended a deep biopsy specimen that includes subcutaneous fat to help achieve an accurate diagnosis (Worley et al., 2019). Risk factors for MAC are

*E-mail address*: ralwassia@kau.edu.sa (R.K. Al-Wassia). Peer review under responsibility of King Saud University.



Production and hosting by Elsevier

unknown, however ultraviolet light, radiation, immunosuppression, and genetic predisposition have been implicated in the pathogenesis (Abbate et al., 2003; Worley et al., 2019). The most commonly utilized treatment modalities include wide-local excision, Mohs micrographic surgery, and radiation therapy (RT) (Chaudhari et al., 2015).

# 2. Case presentation

A 41-year-old male Filipino nurse, who had no known medical history and was a smoker, presented with a localized 5-cm mass at the vertex of his scalp that had appeared 6 months ago. It was gradually increasing in size. The swelling was first observed after head trauma when the swelling transformed into a vascularized hematoma. The patient had a history of multiple head injuries and keloid formation at 10 years of age. On examination, the lesion showed grayish discoloration, with an irregular surface, obscuring borders, and scattered nodularity over  $7 \times 8$  cm at the vertex. There was associated localized alopecia, ulceration, and necrosis (Fig. 1A). Upon investigation, the patient was found to have multiple episodes of high blood pressure, reaching 182/106 mmHg, with a positive family history of hypertension and paternal lung cancer.

In July 2018, the patient was referred to a plastic surgeon for evaluation of the scalp lesion and further management (Fig. 1A). Complete physical examination including a relevant neurologic assessment, regional lymph node assessment, and baseline imag-

<sup>\*</sup> Corresponding author at: Department of Radiology, Faculty of Medicine King Abdulaziz University, P.O.BOX: 80215, Jeddah 21589, Saudi Arabia.

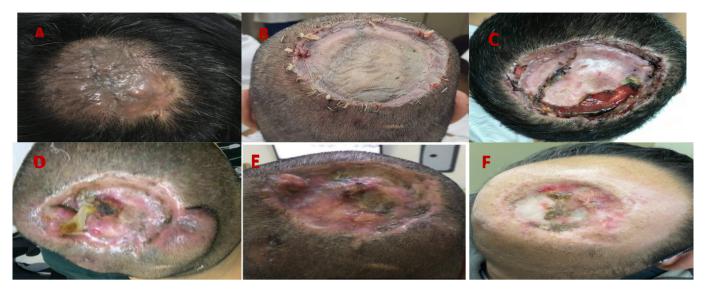


Fig. 1. Microcystic adnexal carcinoma: (A) at presentation (B) after surgical resection and graft (C) graft necrosis (D) before chemoradiotherapy (E) after chemotherapy (F) after radiation.

ing was performed to assess for local or nodal metastasis and assist in preoperative planning. Computed tomography (CT) brain showed a right-sided, well-defined, ovoid, heterogeneous, subcutaneous lesion measuring  $3\times 2.5\times 2$  cm in the high parietal region. Its average density was approximately 60 HU. The underlying bone appeared intact with no evidence of cortical thickening or erosion. Consequently, the patient was admitted for incisional biopsy, revealing a microcystic adnexal carcinoma with perineural invasion. Tumor cells were positive for CK7 and CK5/6, but negative for CK20 and SMA. The case was discussed with the tumor board, who recommended surgery as the initial management.

In December 2018, CT brain showed interval worsening of the right parietal extracranial soft tissue scalp lesion, with an increase in size (2.3  $\times$  4.5  $\times$  4.8 cm [craniocaudal (CC)  $\times$  anteroposterior (AP)  $\times$  transverse (TR)]), involving subcutaneous tissue and overlying skin. CT chest abdomen pelvis (CAP) and mammograms showed no evidence of metastasis.

On follow-up in February 2019, CT brain showed an interval increase in size of the previously seen subcutaneous, right parietal lesion. The lesion measured 3.6  $\times$  6.1  $\times$  5.1 cm (CC  $\times$  AP  $\times$  TR) with no frank involvement of the underlying bone, and no erosions or sclerosis depressed the lesion. The lesion showed heterogeneous enhancement on post-contrast study. No intracranial extension or brain metastases were noted. A wide local excision was performed with 2-cm safety margins. Intraoperative frozen sections were taken and resulted negative margin except for occipital periosteum and as a result, a neurosurgeon joined surgery for outer and inner table skull drilling exposing the dura in the center of the defect. The specimen was then sent for histopathology examination. Defect reconstruction with a split-thickness skin graft was performed temporary during the same operative session and to be booked for second stage reconstruction. Postoperative pathological examination confirmed that the surgical margins of the excised tumor were tumor-free except for occipital periosteum. The pathological report revealed microcystic adnexal carcinoma with undifferentiated foci, with a maximum tumor dimension of 6.5 cm. perineural invasion, and no lymphovascular invasion was detected. Three weeks later, the patient was admitted for skin graft necrosis in which the debridement was performed intraoperatively.

In June 2019, the scalp tumor recurred in multiple areas around the surgical scar with bilateral cervical lymphadenopathy, where the largest is located at the left level V measuring 1.6 cm on CT scan. The case was discussed with the tumor board once again.

Administration of three cycles of chemotherapy, docetaxel + cispla tin regimen followed by radiation therapy (RT) was recommended. Following chemotherapy, a good response was noted in the scalp lesion along with the cervical lymph nodes based on the CT scan results. However, the patient lost follow-up due to the extensive disease recurrence resulting to delayed RT as the patient returned to his home country for 2 months. Nevertheless, the local control of chemotherapy was preserved and an updated CT scan showed no evidence of gross disease. The patient started on RT course of 66 Gy in 33 fractions using intensity-modulated RT to the primary site along with cervical lymph nodes (Fig. 3). Clinical response of the tumor was observed as early as 6 weeks after completion of RT. Follow-up scans demonstrated a decrease in the size of the previously visualized nodes, with no cervical lymphadenopathy (Fig. 2B). The patient remains asymptomatic and disease free after 18 months of follow-up. Even at the time of writing this manuscript, this was evident on the clinical and radiological imaging that was taken Fig. 4.

# 3. Discussion

Microcystic adnexal carcinoma (MAC) is an uncommon tumor and was first described by Goldstein and colleagues in 1982 (Goldstein et al., 1982). Predominantly, MAC occurs in a population with fair skin and usually in the head and neck region (Chiller et al., 2000; Abbate et al., 2003; Stam et al., 2015; Oyasiji et al., 2018). Due to the rarity, limited exposure, frequent extensive subclinical involvement and slow growing nature of MAC, prognosis and management strategies are not well established. Given these diagnostic challenges, the clinical presentation of MAC can be mistaken for basal cell carcinoma, squamous cell carcinoma, or other variable diagnosis reported in the literature. Hence, skin cancer screening should focus on high-risk anatomical areas, such as the scalp, where detection can be delayed because of hair covering.

This study reported a 41-year-old male patient, a finding concordant with that of Oyasiji and colleagues in 2018 and Blake and colleagues in 2010 (Blake et al., 2010; Oyasiji et al., 2018). However, this result was in contrast with other studies such as Worley and colleagues (2019) and Barnes and Garcia (2008) (Barnes and Garcia, 2008; Worley et al., 2019). In a recent systematic review conducted on gender distribution, a female predominance was shown with a median age of 61.8 years for the

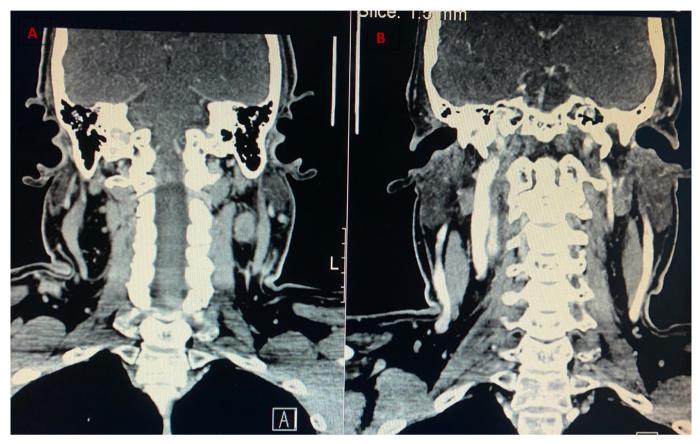


Fig. 2. Coronal view of CT scan (A) before chemotherapy, (B) after chemoradiotherapy.

development of MAC (Worley et al., 2019). The etiology of MAC is unknown nevertheless, possible triggers such as ultraviolet light and radiation have been postulated to their development (Abbate et al., 2003; Worley et al., 2019). Nonetheless, our patient had no exposure to radiation or a history of long hours of sunlight exposure. Because of the widespread infiltration and indistinct depth of invasion of the disease, it may also extend into deeper underlying structures such as muscles, nerves, and bones, leading to deformity (Abbate et al., 2003; Waqas et al., 2017). In 2000, Chiller and colleagues demonstrated that the actual tumor defect can have four-fold increase in size (Chiller et al., 2000). In our case, a clearance margin of 2-cm of the clinically apparent borders of the tumor was observed.

The mainstay treatment modality for MAC is wide local excision along with comprehensive marginal excision while preserving function and cosmesis (Waqas et al., 2017; Worley et al., 2019). As a consequence, this patient underwent complete excision with 2-cm clear surgical margins. Since the tumor was deeply infiltrative, neurosurgery was involved even though no evidence of infiltration to the skull was detected on the imaging. Tissue processing by frozen section to ensure negative peripheral and deep margins and as well as final histopathology were also performed. Nevertheless, the occipital periosteum remained positive owing to the deep infiltrating growth pattern of the tumor. Given the significant soft-tissue defects after tumor resection, reconstruction was performed using a skin graft.

Due to scarcity and diversity of cases treated with radiation therapy (RT) establishing the exact role of RT in MAC would be challenging to interpret. Our experience builds on previously published data reporting its efficacy, summarized in Table 1. The role of adjuvant RT has not been well defined in MAC. However, a number of case series have supported its use in the context of

high risk factors such as close or positive surgical margins, persistent occult tumors at the margins, and perineural invasion. RT may also be beneficial in the definitive setting as a surgical alternative (demonstrated in Table 1). In this case, we administered a dose of 66 Gy in 33 fractions to 2.5 cm beyond the primary tumor location, the borders of surgical bed, cervical lymph nodes bilaterally, and the node at risk to maximize the locoregional control and had no serious adverse events documented. This is in accordance with a recently published systematic review that recommended the use of adjuvant RT (60–66 Gy, 2 Gy per fraction) in the presence of adverse pathologic features (Worley et al., 2019).

In comparison with other case reports, 45 Gy was administered by Waqas and colleagues (2017), 50-60 Gy by (Wang et al., 2017) with 100 % locoregional control, and 55 Gy given by Baxi and colleagues (2010) with 93% locoregional control, despite 69% and 56% having positive margins and peri-neural invasion, respectively. In other published case reports on MAC using upfront definitive RT as a surgical alternative such as those of Gulmen and Pullon (1976), (Schipper et al., 1995), Stein and colleagues (2003), Pugh and colleagues (2012) and Kim and colleagues (2012) only those of Stein and colleagues (2003) and Pugh and colleagues (2012) described positive clinical outcomes. In the recent report of Kim and colleagues (2020), a patient with 12-year history of a histologically diagnosed MAC philtrum nodule presented with worsening numbness of the upper lip and midfacial induration, was treated using RT (70 Gy) targeted to the primary site as well as the nodal area and with concurrent chemotherapy, a progression-free for 6 years from the completion of the treatment was observed (Kim et al., 2020). Moreover, in 2014 Kim and colleagues reported the use of adjuvant RT and demonstrated efficacy in treating scalp lesions (Kim et al., 2014).

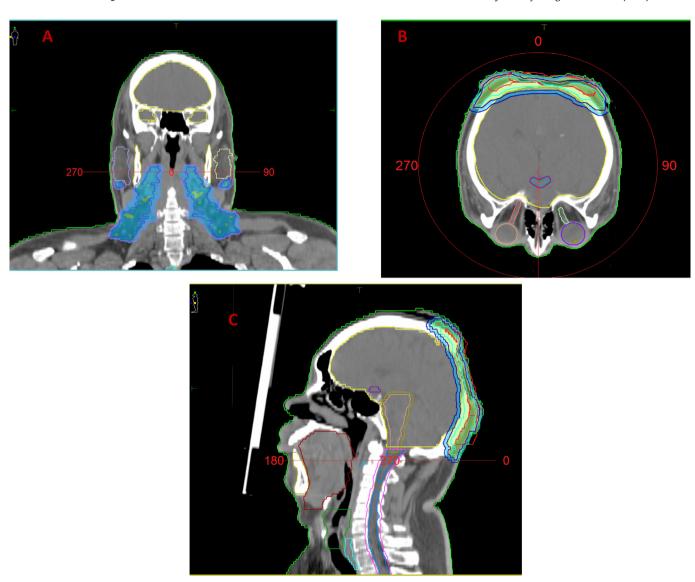


Fig. 3. Radiation plan for treatment of microcystic adnexal carcinoma primary tumor and cervical lymph nodes in (A) coronal, (B), axial and (C) sagittal view.

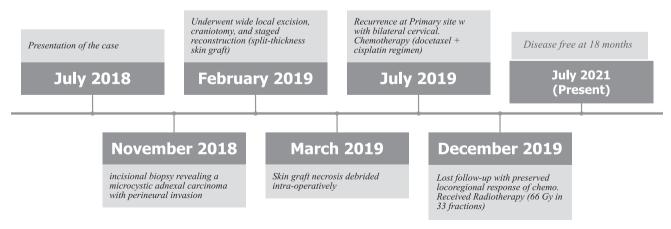


Fig. 4. Timeline of the case.

In addition to RT, we incorporated sequential chemotherapy as it could potentially act as a radiosensitizer to improve local control of the treatment. Although the time period between the last

chemotherapy dose and the start of radiation was 2 months, local control of chemotherapy was still preserved, clinically and radiologically. Nonetheless, as summarized in Table 2, the role of

**Table 1**Case reports of MAC in the head and neck treated with radiation therapy.

Author (date)	Age/Sex	Primary tumor site	Primary treatment	TTR (month)	Nature of recurrence	Salvage treatment	Outcome
Definitive Radiotherapy							
Kim et al. (2020)	73/M	Philtrum	RT to primary (70 Gy) + lymph node (56– 63 Gy) + chemo; declined surgery	NA	NA	NA	No tumor progression at 72 mo
Pugh et al. (2012)	53/F	Upper lip	RT (63.6 Gy)	48	Primary site	RT (64 Gy)	Disease free at 15 mo
Stein et al. (2003)	76/F	Right nasal dorsum	RT (58 Gy)	6	Primary site extend to right medial cheek + left nasal dorsum	Total rhinectomy, partial septectomy /turbinectomy	Disease free at 14 mo
Schipper et al. (1995)	65/M	Tongue	RT (NR): refused surgery	NA	NA	NA	No change in tumor size. Disease free at 21 mo
Gulmen and Pullon (1976)	35/F	Left lower lip + submental lymph node	RT to primary (60 Gy) + lymph node (50 Gy)	NA	NA	NA	Disease free at 6 mo
<u>Adjuvant Radiotherapy</u>							
Brent et al. (2018)	NS	Orbit	complete orbital exenteration with positive margin + RT (66 Gy)	NA	NA	NA	Mass reduced in size with remnant post-RT scarring.  Disease free at 12 years
Waqas et al. (2017)	59/F	Scalp	Surgical excision with positive margins + RT (45 Gy)	NA	NA	NA	Disease free at 3 years
Waqas et al. (2017)	53/M	Left temporal scalp	Surgical excision with positive margins + RT (45 Gy)	NA	NA	NA	Disease free at 3 years
Chaudhari et al. (2015)	14/M	Right medial upper lip + submandibular lymph node	Surgical excision + sentinel lymph node biopsy > RT + chemo (NR)	NA	NA	NA	NR
Kim et al. (2014)	56/F	Scalp	Surgical excision with positive margins, PNI, and periosteal involvement > Wider excision + RT (NR)	NA	NA	NA	Disease free at 3 years
Pugh et al. (2012)	60/F	Chin	MMS with NR muscle and PNI > surgical excision with reconstruction of lower lip + RT (66 Gy)	NA	NA	NA	Disease free at 30 mo
Pugh et al. (2012)	63/F	Right cheek	Surgical excision with positive margin and PNI + RT (60 Gy)	NA	NA	NA	Disease free at 26 mo
Baxi et al. (2010)	14 patients, median age 71 years	Head and neck region	Surgical excision (56% PNI, 69% positive margins) + RT (median dose 55 Gy)		(35 Gy) > progress	sed after 2 years > surgical ex	rate of 93%. One pt with local recurrence (CN V salvaged > R scision + RT(45 Gy). One pt with ipsilateral cervical nodal on > RT (60 Gy). Both diseased free
Ong et al. (2004)	89/F	Right eyebrow	Surgical excision with positive margin (declined further excision) + RT	NA	NA	NA	Disease free at 6 mo
Kirkland et al. (1997)	55/F	Nasal septum	Surgical excision with positive margins + RT (55 Gy)	NA	NA	NA	Disease free at 6 mo
Yuh et al. (1991)	51/M	Left lower lip	Surgical excision with positive margins + RT (57.5 Gy)	36	Primary site with invasion of mandible	Surgical excision + RT	Disease free at 18 mo

Table 1 (continued)

Author (date)	Age/Sex	Primary tumor site	Primary treatment	TTR (month)	Nature of recurrence	Salvage treatment	Outcome
Birkby et al. (1989)	51/M	Left lower lip	Surgical excision with positive margins + RT (57.5 Gy)	36	Primary site + ipsilateral mandible invasion	MMS + partial hemi- mandibulectomy + RT (61.2 Gy)	Disease free at 18 mo
Salvage Radiotherapy							
Haga et al. (2019)	78/F	Philtrum	Surgical excision	Unknown Primary site	Unknown Primary site	RT (60 Gy)	Recurrence at primary site and ala of the nose. Chemo (S-1 monotherapy) at 6 years for recurrence. NP at 15 mo.
King et al. (2018)	3 patients	NR	Surgical excision	NR	NR	Surgical excision + RT	2 had progressive disease with 1 developing fatal metastases to skin, lymph node, and lung.
Mamic et al. (2018)	74/F	Left upper lip	Surgical excision	36	Primary site	Surgical excision with positive margin	Surgical excision + RT (58 Gy). Disease free at 18 mo
Gomez- Maestra (2009)	75/F	Right eyebrow	MMS	24	Right supraorbital nerve	Superior orbitotomy + RT (61.2 Gy)	Local recurrence 24 mo after RT > Exenteration. Developed mesencephalon and cavernous sinus metastases 21 mo later. Managed with supportive care.
Clement et al. (2005)	59/M	Left temporal	Surgical excision with positive margin and PNI	8	Primary site	MMS	Recurrence at 4 mo > surgical excisions + RT (57.6 Gy). Third recurrence over left cheek treated with RT (45 Gy). Disease free at 71 mo.
Clement et al. (2005)	83/M	Right canthus	Surgical excision	7	Right lateral pper eye lid	Multiple surgical excisions with positive margin + RT (60 Gy)	Disease free at 47 mo.
Sebastien et al. (1993)	57/F	Chin	MMS	60	Primary site	MMS	MMS + adjuvant RT (55 Gy) for local recurrence 48 mo later. Disease free at 16 mo.
Carroll et al. (2000)	86/M	Left upper forehead	MMS	5	Satellite deposits (3 new distant nodules in scalp from primary)	MMS	3 months later developed 5 satellite deposits on scalp > RT (60 Gy, 6 megavolt scalp) > developed left postauricular nodal recurrence 3 mo later. Underwent MMS but found to have lymph node infiltration. Expired shortly after from metastatic small cell lymphoma.
Bier-Laning et al. (1995)	55/M	Left posterior scalp	Multiple surgical excisions	10	Primary site with dura involvement (craniectomy)	Debulking only + RT (54 Gy).	Progression at 46 mo. Disease free at 10 mo.
Bier-Laning et al. (1995)	85/M	Right cheek	Multiple surgical excisions	NR	Primary site + right CN Vb	MMS with positive margin > surgical excision (right cheek, anterior wall of maxilla, infraorbital nerve)	Right lower eyelid involvement at 13 mo salvaged with RT (60 Gy). disease fee at 7 mo.
Bier-Laning et al. (1995)	46	Right upper and medial cheek	Surgical excision	96	Primary site	RT (60 Gy)	Field edge recurrence at 18 mo, treated with reirradiation (45 Gy). Recurrence at 19 mo requiring multiple reirradiation + resections + chemo. Remained with suspicious lesion on lip.

Abbreviations: F - female, M - male, MMS - Mohs - micrographic surgery, NA - not applicable, NR- not reported, PNI - perineural invasion, RT - radiation therapy, TTR - time to recurrence., NP - No progression

case reports of MAC of the head and neck treated with chemotherapy

Author (date)	Age in years/ Sex	Age in Primary site years/ Sex	Primary treatment	Nature of recurrence	Salvage treatment Chemotherapy	Chemotherapy	Outcome
Kim et al. (2020) 73/M	73/M	Philtrum	RT to primary (70 Gy) + lymph node NA (56–63 Gy) + chemo; declined surgery	NA	NA	4 cycles of concomitant weekly disease free at 72 mo carboplatin and paclitaxel	disease free at 72 mo
Haga et al. (2019) 78/F	78/F	Philtrum	Surgical excision	Primary site + ala of nose	RT > recurrence resulting in chemo	Oral S-1 monotherapy, combination drug (tegafur/ gimeracil/oteracil)	PR at 8 weeks. NP at 15 mo.
Chaudhari et al. (2015)	14/M	Right medial upper lip +submandibular lymph node	Surgical excision + sentinel lymph node biopsy > RT + chemo (NR)	NA	NA	V.	NR
BierLaning et al. (1995)	46/F	Right upper lip and medial cheek	Surgical excision	Primary site + lymph node	Multiple RT + surgical excisions + chemo	Single course of cisplatin + 5- fluorouracil	Recurrence at 8 mo resulting in multiple surgical excision + RT. Remained with suspicious lesion on lip.

Abbreviations: F - female, M - male, NA - not applicable, NR - not reported, RT - radiation therapy, TP - tumor progression, PR - partial response, NP - No progression

chemotherapy remains unclear and further studies are necessary to definitively determine this role (Chaudhari et al., 2016). In 1995, Bier-Laning and colleagues reported a 46-year-old woman presented with right upper and medical cheek lesion who was initially misdiagnosed (Bier-Laning et al., 1995). This patient underwent multiple extensive surgical resections due to recurrence and metastasis. Additionally, variant courses of radiation therapy was administered to the primary site and cervical lymph node bilaterally, and as well as received a single course of cisplatin and 5-fluorouracil. Even though with chemotherapy, it remained resistant and failed to eradicate the tumor (Bier-Laning et al., 1995). In 2015, Chaudhari and colleagues used chemoradiation on the involved lymph node of a previously resected primary lip lesion, however details of chemoradiation and clinical outcomes have not been reported (Chaudhari et al., 2015). In 2019, Haga and colleagues reported a MAC case of a locally infiltrated recurrent ulcerative tumor on the philtrum and ala of the nose which was previously resected and irradiated with 60 Gy, and successfully treated with combination chemotherapy oral S-1 (tegafur/ gimeracil/oteracil) with partial response at 8-weeks and no progression at 15-months follow-up (Haga et al., 2019). As previously mentioned, Kim and colleagues (2020) used RT (70 Gy) to the primary site and nodal area, with concurrent chemotherapy (four cycles of concomitant weekly carboplatin and paclitaxel) and achieved 6 years of progression-free survival (Kim et al., 2020).

Despite the aggressive presentation, we treat each case considering age, health status, with exhaust treatment option to reach best local control and survival rate with utilizing available modality and treat the patient with best option. This particular case presented had experiences that warrants additional information on the limited resources on the management strategies of MAC. Also, this may pose as a model treatment modality to most likely similar cases. Nevertheless, close clinical follow-up and periodic selfexamination are important to monitor locoregional, distant metastasis, as well as side effects in cancer treatment. Although the side effects of such treatment have significant impact on survivors' quality of life, morbidity, and mortality. The long-term sequelae of the multiple modalities were discussed with and agreed by the patient prior treatment. Starting from locoregional side effects of alopecia, pigmentation, defect, and neuropathic pain to generalized latent side effects include musculoskeletal and neuromuscular dysfunction, oropharyngeal issues (dental caries and dysphagia) psychosocial, hypothyroidism, increased risk of other cancers, stroke, osteoporosis, and infertility.

Photoprotection, smoking cessation, and possible normal changes or alterations in local sensation during healing have been and need to be discussed with patients.

# 4. Conclusion

Microcystic adnexal carcinoma (MAC) is a rare tumor with diverse histological patterns and a tendency for locoregional and distant metastasis. Surgical excision followed by adjuvant radiation therapy (RT) and chemotherapy offers excellent local–regional control for patients with skin adnexal carcinomas and risk factors for local regional recurrence.

Funding

No funding source was available for this study.

Availability of data and materials

Please contact author for data requests.

Consent for publication

Written informed consent was obtained from the patient for the publication of this case report and accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

# **Declaration of Competing Interest**

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

### References

- Abbate, M., Zeitouni, N.C., Seyler, M., Hicks, W., Loree, T. and Cheney, R.T., 2003. Clinical course, risk factors, and treatment of microcystic adnexal carcinoma: a short series report. Dermatol Surg 29(10): 1035-1038. https://doi.org/: 10.1046/ i.1524-4725.2003.29296.x.
- Barnes, J., Garcia, C., 2008. Primary cutaneous adenoid cystic carcinoma: a case report and review of the literature. Cutis 81 (3), 243–246.
- Baxi, S., Deb, S., Weedon, D., Baumann, K. and Poulsen, M., 2010. Microcystic adnexal carcinoma of the skin: the role of adjuvant radiotherapy. J Med Imaging Radiat Oncol 54(5): 477-482. https://doi.org/: 10.1111/j.1754-9485.2010.02200.x.
- Bier-Laning, C.M., Hom, D.B., Gapany, M., Manivel, J.C. and Duvall, A.J., 3rd, 1995. Microcystic adnexal carcinoma: management options based on long-term follow-up. Laryngoscope 105(11): 1197-1201. https://doi.org/: 10.1288/00005537-199511000-00011.
- Blake, P.W., Bradford, P.T., Devesa, S.S. and Toro, J.R., 2010. Cutaneous appendageal carcinoma incidence and survival patterns in the United States: a populationbased study. Arch Dermatol 146(6): 625-632. https://doi.org/: 10.1001/ archdermatol.2010.105.
- Chaudhari, S., Blanchard, L., Chaudhari, P.J. and Lythgoe, T., 2015. The first reported case of metastatic microcystic adnexal carcinoma in an adolescent. Dermatol Surg 41(2): 286-288. https://doi.org/: 10.1097/dss.0000000000000258.
- Chaudhari, S.P., Mortazie, M.B., Blattner, C.M., Garelik, J., Wolff, M., Daulat, J. and Chaudhari, P.J., 2016. Treatments for microcystic adnexal carcinoma–A review. J Dermatolog Treat 27(3): 278-284. https://doi.org/: 10.3109/09546634.2015.1089351.
- Chiller, K., Passaro, D., Scheuller, M., Singer, M., McCalmont, T. and Grekin, R.C., 2000. Microcystic adnexal carcinoma: forty-eight cases, their treatment, and their outcome. Arch Dermatol 136(11): 1355-1359. https://doi.org/: 10.1001/ archderm.136.11.1355.
- Goldstein, D.J., Barr, R.J. and Santa Cruz, D.J., 1982. Microcystic adnexal carcinoma: a distinct clinicopathologic entity. Cancer 50(3): 566-572. https://doi.org/: 10.1002/1097-0142(19820801)50:3<566::aid-cncr2820500330>3.0.co;2-q.
- Gulmen, S. and Pullon, P.A., 1976. Sweat gland carcinoma of the lips. Oral Surg Oral Med Oral Pathol 41(5): 643-649. https://doi.org/: 10.1016/0030-4220(76) 90317-0.
- Haga, T., Fujimura, T. and Aiba, S., 2019. Successful treatment of microcystic adnexal carcinoma on the philtrum with oral S-1 monotherapy. J Dermatol 46(6): e208-e209. https://doi.org/: 10.1111/1346-8138.14770.

- Kim, D.W., Lee, G., Lam, M.B., Harris, E.J., Lam, A.C., Thomas, T., Chau, N.G. and Tishler, R.B., 2020. Microcystic Adnexal Carcinoma of the Face Treated With Definitive Chemoradiation: A Case Report and Review of the Literature. Adv Radiat Oncol 5(2): 301-310. https://doi.org/: 10.1016/j.adro.2019.11.008.
- Kim, L.H., Teston, L., Sasani, S. and Henderson, C., 2014. Microcystic adnexal carcinoma: successful management of a large scalp lesion. J Plast Surg Hand Surg 48(2): 158-160. https://doi.org/: 10.3109/2000656x.2013.777218.
- Oyasiji, T., Tan, W., Kane, J., 3rd, Skitzki, J., Francescutti, V., Salerno, K. and Khushalani, N.I., 2018. Malignant adnexal tumors of the skin: a single institution experience. World J Surg Oncol 16(1): 99. https://doi.org/: 10.1186/s12957-018-1401-y.
- Pugh, T.J., Lee, N.Y., Pacheco, T. and Raben, D., 2012. Microcystic adnexal carcinoma of the face treated with radiation therapy: a case report and review of the literature. Head Neck 34(7): 1045-1050. https://doi.org/: 10.1002/hed.21690.
- Schipper, J.H., Holecek, B.U. and Sievers, K.W., 1995. A tumour derived from Ebner's glands: microcystic adnexal carcinoma of the tongue. J Laryngol Otol 109(12): 1211-1214. https://doi.org/: 10.1017/s0022215100132475.
- Stam, H., van de Wiel, B.A., Klop, W.M.C., Zupan-Kajcovski, B., Janssens, S., Karakullukcu, M.B., van der Noort, V. and Lohuis, P., 2015. Skin adnexal carcinoma of the head and neck: a retrospective study in a tertiary referral center. Eur Arch Otorhinolaryngol 272(4): 1001-1010. https://doi.org/: 10.1007/s00405-014-3324-8.
- Stein, J.M., Ormsby, A., Esclamado, R. and Bailin, P., 2003. The effect of radiation therapy on microcystic adnexal carcinoma: a case report. Head Neck 25(3): 251-254. https://doi.org/: 10.1002/hed.10167.
- Tanese, K., Nakamura, Y., Hirai, I. and Funakoshi, T., 2019. Updates on the Systemic Treatment of Advanced Non-melanoma Skin Cancer. Front Med (Lausanne) 6: 160. https://doi.org/: 10.3389/fmed.2019.00160.
- Wang, L.S., Handorf, E.A., Wu, H., Liu, J.C., Perlis, C.S. and Galloway, T.J., 2017. Surgery and Adjuvant Radiation for High-risk Skin Adnexal Carcinoma of the Head and Neck. Am J Clin Oncol 40(4): 429-432. https://doi.org/: 10.1097/coc.00000000000000178.
- Waqas, O., Faisal, M., Haider, I., Amjad, A., Jamshed, A. and Hussain, R., 2017. Retrospective study of rare cutaneous malignant adnexal tumors of the head and neck in a tertiary care cancer hospital: a case series. J Med Case Rep 11(1): 67. https://doi.org/: 10.1186/s13256-017-1212-8.
- Worley, B., Owen, J.L., Barker, C.A., Behshad, R., Bichakjian, C.K., Bolotin, D., Bordeaux, J.S., Bradshaw, S., Cartee, T.V., Chandra, S., Cho, N., Choi, J., Council, M. L., Eisen, D.B., Golda, N., Huang, C.C., Ibrahim, S.F., Jiang, S.I.B., Kim, J., Lacutoure, M., Lawrence, N., Lee, E.H., Leitenberger, J.J., Maher, I.A., Mann, M., Minkis, K., Mittal, B., Nehal, K.S., Neuhaus, I., Ozog, D.M., Petersen, B., Samie, F., Shin, T.M., Sobanko, J.F., Somani, A.K., Stebbins, W.G., Thomas, J.R., Thomas, V., Tse, D., Waldman, A., Xu, Y.G., Yu, S.S., Zeitouni, N.C., Ramsay, T., Poon, E. and Alam, M., 2019. Evidence-Based Clinical Practice Guidelines for Microcystic Adnexal Carcinoma: Informed by a Systematic Review. JAMA Dermatol 155(9): 1059-1068. https://doi.org/: 10.1001/jamadermatol.2019.1251.