



An Unusual Case of Basal Cell Carcinoma with Lung and Endobronchial Metastasis

TaeHwa Kim^{1,2}, Do Hyung Kim³, Seung Eun Lee¹, Min-Young Yang⁴, Yun Seong Kim^{1,2}

¹Department of Internal Medicine, Pusan National University Yangsan Hospital, Pusan National University School of Medicine, Yangsan, ²BioMedical Research Institute for Convergence of Biomedical Science and Technology, Pusan National University Yangsan Hospital, Yangsan, ³Department of Cardiovascular and Thoracic Surgery, Pusan National University Yangsan Hospital, Yangsan, ⁴Department of Dermatology, Pusan National University Hospital, Busan, Korea

Received January 6, 2020
Revised December 21, 2020
Accepted February 1, 2021

Corresponding Author

Yun Seong Kim
Department of Internal Medicine, Pusan National University Yangsan Hospital, Pusan National University School of Medicine, 20 Geumo-ro, Mulgeum-eup, Yangsan 50612, Korea
Tel: +82-55-360-1414
Fax: +82-55-360-1757
E-mail: yskim@pusan.ac.kr
<https://orcid.org/0000-0003-4328-0818>

Recently, some cases of basal cell carcinoma (BCC) with lung metastasis have been reported, but those involving simultaneous tracheal, bronchial, and lung metastases have been rarely reported. Here, we have reported a very unusual case of BCC with metastasis, presenting with lung nodules and endobronchial lesions after two metastasectomies. Since BCC is a slow-growing cancer that rarely metastasizes to distant organs, tumor stage workup including radiological imaging has not been routinely performed in clinical practice. This case showed that BCC can metastasize to the lung, although the currently reported metastasis rate of BCC is extremely low.

Keywords: Basal cell carcinoma, Lung, Neoplasm metastasis

INTRODUCTION

Basal cell carcinoma (BCC) is the most common skin cancer of the head and neck occurring after prolonged sun exposure. It is a slow-growing cancer that rarely metastasizes to other organs. Recently, some cases of BCC with lung and tracheal metastases have been reported. However, cases involving simultaneous lung, endotracheal, and endobronchial metastases have been rarely reported^{1,2}. Here, we have reported a case of facial BCC with endotracheal and endobronchial metastases.

CASE REPORT

A 58-year-old male was diagnosed with BCC on the left cheek 15 years ago at a local hospital and had undergone excision two times. Nine years after the diagnosis, metastasis was detected on chest computed tomography (CT), which showed a nodule measuring 2 cm in the right upper lobe and a nodule measur-

ing 2.5 cm in the left lingular segment. The patient underwent metastasectomy of both the lungs and confirmed the presence of BCC. He received six courses of 5-fluorouracil and cisplatin-based chemotherapy for approximately 1 year. The patient was free of metastatic BCC recurrence for 6 years, until he presented with left upper lobe (LUL) lung metastasis again, for which he underwent video-assisted thoracoscopic surgery. Six years after surgical resection, chest CT, performed as part of a regular checkup, showed an enhancing soft tissue density in the left lateral tracheal wall, luminal narrowing of the proximal portion of the left main bronchus, and anterior arch pathological fracture of the left third rib (Fig. 1A, B). Bronchoscopy and mucosal biopsy revealed a mucosal lesion, a whitish patch, and an elevated polypoid mass in the left main bronchus (Fig. 1C, D). The patient was diagnosed with metastatic BCC recurrence for the third time with endotracheal and bronchial metastases and a pulmonary lesion on the LUL. On January, 2019, the patient underwent tracheal resection and end-to-end anasto-



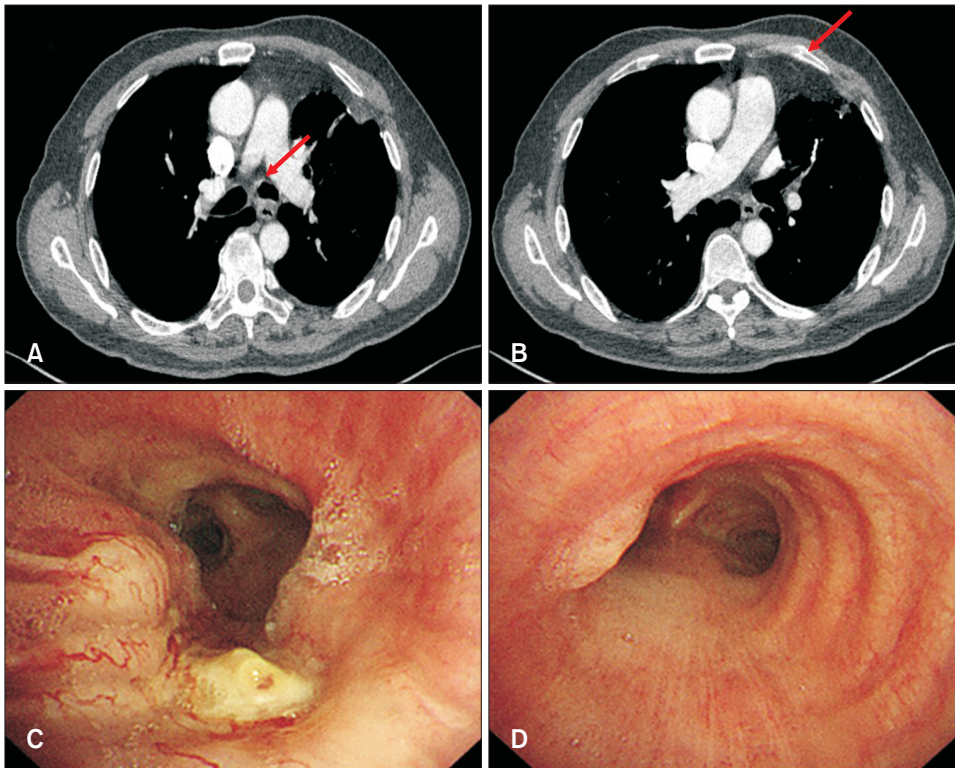


Fig. 1. Chest computed tomography showing the proximal portion of the left main bronchus with luminal narrowing (red arrow) (A) and anterior arch pathological fracture of the left third rib (red arrow) (B). Bronchoscopy showing a mucosal lesion, a whitish patch (C), and an elevated polypoid mass in the left main bronchus (D). We received the patient's consent form about publishing all photographic materials.

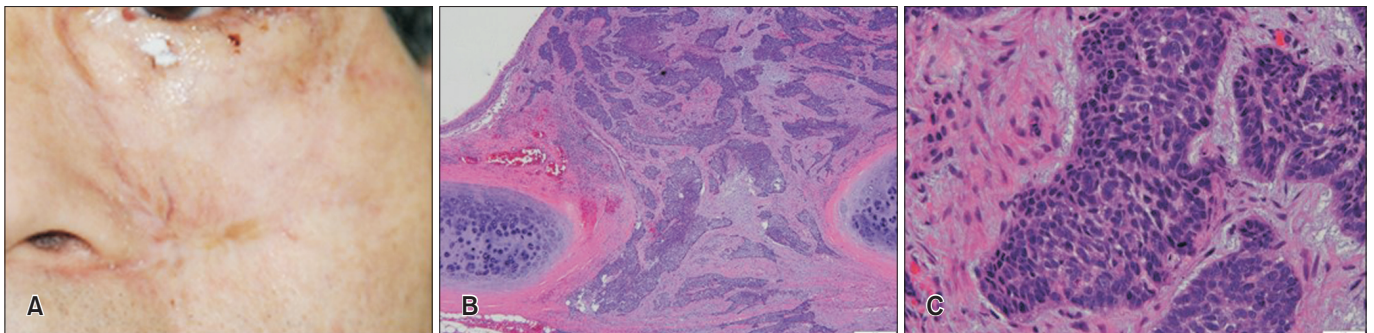


Fig. 2. (A) Recovery of the left side of the face without any side effects after surgery. Tumor cell nests involving the left bronchus invading the peribronchial soft tissue and the tumor in the trachea showing typical features of basal cell carcinoma including peritumoral clefts and basal palisading (B: H&E, $\times 40$; C: H&E, $\times 400$). We received the patient's consent form about publishing all photographic materials.

mosis, LUL sleeve lobectomy, and left third rib resection and reconstruction. Histological examination of all the surgical tissues confirmed the diagnosis of BCC; examination of the skin tissue revealed the same pathological findings (Fig. 2).

DISCUSSION

BCC is the most common skin cancer worldwide as well as in Korea with predilection site of the head and neck¹. It is a slow-growing cancer that rarely metastasizes to other

organs³⁻⁵. Recently, some cases of BCC with lung metastasis have been reported, but cases involving simultaneous lung, endotracheal, and endobronchial metastases have been rarely reported^{1,2}. This report pertains to a very rare case of BCC with lung, endobronchial, and bronchial metastases after two metastasectomies.

In a previous study, the median age of onset of primary BCC was 45 years, and the average period between the diagnosis of the primary lesion and development of metastasis was approximately 10 years⁶. In our patient, lung metastasis was

found 6 years after primary tumor diagnosis. It took 13 years for the duration between primary tumor onset and simultaneous metastasis to lung, tracheal and bronchial portions.

The optimal treatment for metastatic BCC is complete surgical resection, if possible, as it confers favorable survival rates and recurrence-free time. Radiation and chemotherapy are therapeutic choices for metastatic BCC. Our patient underwent surgery and adjuvant chemotherapy. The patient underwent two additional surgeries for lung, tracheal, and bronchial metastases and is being closely followed up.

Although the patient with previous history BCC develops suspicious lesions as cancer in internal organs, it may be reasonable to suspect other primary or secondary cancer. Therefore, the diagnosis should be confirmed histologically for that suspicious lesions. Since BCC is a slow-growing cancer that rarely metastasizes, initial work-up with chest CT and bronchoscopy usually do not reveal BCC. Although survival is short after metastasis, longer survival is possible without recurrence for years after metastatic cancer treatment, as seen in our patient. Therefore, detailed examination and evaluation are needed for discrimination when finding lesions suspected of metastasis in patients with a history of BCC.

CONFLICTS OF INTEREST

The authors have nothing to disclose.

FUNDING SOURCE

This study was supported by a 2020 research grant from Pu-

san National University Yangsan Hospital.

ORCID

TaeHwa Kim, <https://orcid.org/0000-0003-3722-0261>

Do Hyung Kim, <https://orcid.org/0000-0002-8774-3397>

Seung Eun Lee, <https://orcid.org/0000-0002-4266-7722>

Min-Young Yang, <https://orcid.org/0000-0001-8994-8401>

Yun Seong Kim, <https://orcid.org/0000-0003-4328-0818>

REFERENCES

1. Verma S, Sahni S, Zeb S, Esposito MJ, Talwar A. A rare case of endobronchial and lung metastasis in basal cell carcinoma. *JAAPA* 2014;27:27-28.
2. Lattes R, Kessler RW. Metastasizing basal-cell epithelioma of the skin; report of two cases. *Cancer* 1951;4:866-878.
3. Aldhaban S, Marc S, Eshki M, Girod A, Boissonet H, Chapelier A, et al. Giant basal cell carcinoma with regional lymph node and distant lung metastasis. *Eur J Dermatol* 2011;21:972-975.
4. Motegi S, Tamura A, Tanaka S, Nagai K, Ishikawa O. Aggressive basal cell carcinoma with pulmonary metastases. *Eur J Dermatol* 2006;16:585-586.
5. Copcu E, Aktas A. Simultaneous two organ metastases of the giant basal cell carcinoma of the skin. *Int Semin Surg Oncol* 2005;2:1.
6. Von Domarus H, Stevens PJ. Metastatic basal cell carcinoma. Report of five cases and review of 170 cases in the literature. *J Am Acad Dermatol* 1984;10:1043-1060.