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Uncommon features of surgically resected ALK-positive cavitary lung adenocarcinoma: a case report

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Abstract

Some features found on chest computed tomography (CT), such as central tumor location, large pleural effusion, and the absence of a pleural tail, and a patient age of less than 60 years, have been suggested to be useful in predicting anaplastic lymphoma kinase (ALK) rearrangement in patients with non-small cell lung cancer (NSCLC). A 68-year-old female patient with a history of gynecological treatment was found to have a cavitary mass in the right lower lobe on an annual chest roentgenogram. The tumor was located in the peripheral area with a pleural tail showing no pleural effusion. In addition, two pure ground-glass-opacity nodules (p-GGNs) in the right upper lobe of the lung were detected on consecutive chest CT scans. The patient underwent right lower lobectomy, partial resection of the right upper lobe, and hilar mediastinal lymph node dissection for complete resection of each tumor. The pathological diagnosis was invasive mucinous adenocarcinoma with signet-ring cells for the cavitary mass in the right lower lobe and invasive adenocarcinoma for the rest of the p-GGNs; subcarinal lymph node metastasis was also detected. The ALK rearrangement was detected by fluorescence in situ hybridization from the cavitary mass. The patient underwent four cycles of cisplatin and vinorelbine chemotherapy as standard adjuvant chemotherapy for pStage III NSCLC. The ALK fusion gene status of NSCLC with atypical CT features should also be investigated.

Keywords: Anaplastic lymphoma kinase, Cavitary mass, Radiological feature

Background

Several radiological features delineated by chest computed tomography (CT) of non-small cell lung cancer (NSCLC) have been reported to be associated with some utility in detecting the gene mutation status, such as *epidermal growth factor receptor* (EGFR) or *anaplastic lymphoma kinase* (ALK) rearrangement. Certain features of a tumor on chest CT, such as convergence, notch, and a pleural tail, have been reported to have a significant association with an EGFR or ALK fusion gene mutation [1–5]. With regard to the mutation of the ALK fusion gene, the combination of certain tumor features on chest CT, such as a central tumor location, large pleural effusion, and the absence of a pleural tail, and a younger patient age (<60 years old), were shown to have a significant relationship with

We herein report a case of surgically resected lung adenocarcinoma that presented as a cavitary mass harboring *ALK* fusion with rare radiological features.

Case presentation

A 68-year-old female was found to have a cavitary mass on annual chest roentgenography in December 2015. She had a history of treatment for cervical cancer in the uterus in 2001 and had been followed up at our hospital without recurrence. She was a never-smoker and had no significant abnormalities on a physical examination except for an abdominal scar related to the treatment for her cervical cancer.

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ALK rearrangement in patients with NSCLC [6]. In general, larger squamous cell carcinomas often show cavitation, and cavitary masses have been found in a number of adenocarcinoma patients who had a poor prognosis [7].

Chest CT revealed a cavitary mass, approximately 4.3 cm in its maximum dimension, in the basal segment of the right lower lobe of the lung (Fig. 1a). In addition, we noted 2 pure ground-glass-opacity nodules (p-GGNs), 8 and 12 mm in their maximum dimensions. Retrospectively, abdominal CT performed in 2008 occasionally revealed a tiny nodule in the basal segment of the right lower lobe adjacent to the visceral pleura on the diaphragm (Fig. 1b). No significant hilar or mediastinal lymph node swelling was observed. fluorodeoxyglucose positron emission tomography (FDG-PET) showed no abnormal accumulation except in the mass in the right lower lobe. A biopsy of the tumor via fiber optic bronchoscopy could not distinguish the cavitary mass from a metastatic or primary tumor. The two p-GGNs were considered to be multiple primary lung cancers. The patient underwent video-assisted thoracoscopic right lower lobectomy, partial resections for the two p-GGNs in the right upper lobe, and systemic hilar mediastinal lymph node dissection as curative intent resection. An analysis of the intraoperative frozen sections for the three tumors revealed adenocarcinoma.

A pathological examination revealed the cavitary mass in the right lower lobe to be invasive mucinous adenocarcinoma (60% acinar, 30% micropapillary, 10% papillary growth pattern) with signet-ring cells (Fig. 1c). Mediastinal lymph node metastasis was found in the subcarinal lesion. The two p-GGNs were pathologically different from the cavitary mass in the right lower lobe; thus, the pathological diagnosis was multiple primary NSCLC: pT2aN2M0 stage IIIA adenocarcinoma for the cavitary mass in the right lower lobe and pT1aN0M0

stage IA adenocarcinoma for each p-GGN in the right upper lobe. The patient received four cycles of cisplatin and vinorelbine as adjuvant chemotherapy. In addition, ALK rearrangement was detected by FISH from the specimen obtained from the cavitary mass on the right lower lobe. The postoperative course was uneventful, and the patient is currently disease-free at 5 months post operation.

Discussion

Previous reports have shown that some CT features are significantly associated with the ALK fusion gene [2–5]. Yamamoto et al. reported that three CT features, namely a central tumor location, large pleural effusion, and the absence of pleural tail, with a patient age of less than 60 years, were good predictors of the presence of the ALK fusion gene in cases of NSCLC [6]. In addition, Yamamoto et al. attempted to predict the presence of the ALK fusion gene using those three CT features and the age of patients and revealed a sensitivity and specificity of 89.4 and 100.0%, respectively, for the presence of the ALK fusion gene in certain cohorts of patients. However, the equation failed to demonstrate the presence of the ALK fusion gene in our patient in the present report. Retrospectively, a tiny nodule was occasionally found in the basal segment adjacent to the visceral pleura (the peripheral area of the right lung on an abdominal CT taken 8 years prior as a follow-up of cervical cancer of the uterus; Fig. 1b). The present case therefore differed with respect to the predictive factors for the presence of the ALK gene fusion in the cavity formation in the tumor, an initial tumor location in a peripheral site

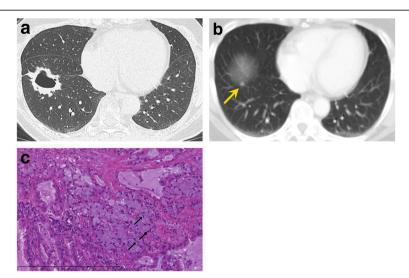


Fig 1 a Chest cross-sectional computed tomography (CT) showing a cavitary mass in the S6 segment in the peripheral area with a pleural tail and without pleural effusion. **b** An abdominal CT scan taken in 2008 occasionally showed a tiny nodule on the basal segment of the *right* lower lobe adjacent to the visceral pleura on the diaphragm (*arrow*). **c** The results of a histological examination. Invasive mucinous adenocarcinoma with signet-ring cells in an S6 tumor (hematoxylin and eosin staining, original magnification ×400)

without pleural tail formation, and a patient age over 60 years old.

With regard to the pathological features of this case, the permanent specimen showed invasive mucinous adenocarcinoma with signet-ring cells (Fig. 1c). Signet-ring cells are a rare feature of primary lung adenocarcinoma [7]; the presence of signet-ring-cell elements is significantly more frequent in ALK+ lung cancer [8]. Given that the ALK status of the patient was positive, these findings were consistent with those of previous reports.

Conclusions

We report a case of NSCLC harboring *ALK* rearrangement with uncommon radiological features. The *ALK* gene status of NSCLC with atypical CT features should also be investigated.

Abbreviations

ALK: Anaplastic lymphoma kinase; CT: Computed tomography; EGFR: Epidermal growth factor receptor; FDG-PET: Fluorodeoxyglucose positron emission tomography; NSCLC: Non-small cell lung cancer; p-GGNs: Pure ground-glass-opacity nodules

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Authors' contributions

ST wrote the manuscript. ME and SS participated in the surgery. KT analyzed the pathologic specimens. MY supervised the writing of the manuscript. RT, KN, FH, TS, MT and YI assisted in drafting the manuscript. All authors significantly contributed to this study. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

Consent for publication

Informed consent was obtained from the patient for publication of this case report and the accompanying images.

Ethics approval and consent to participate

This study was carried out in accordance with the principles of the Declaration of Helsinki.

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References

- Hasegawa M, Sakai F, Ishikawa R, Kimura F, Ishida H, Kobayashi K, et al. CT features of epidermal growth factor receptor-mutated adenocarcinoma of the lung: comparison with nonmutated adenocarcinoma. J Thorac Oncol. 2016;11(6):819–26.
- Kim TJ, Lee CT, Jheon SH, Park JS, Chung JH. Radiologic characteristics of surgically resected non-small cell lung cancer with ALK rearrangement or EGFR mutations. Ann Thorac Surg. 2016;101(2):473–80.
- Halpenny DF, Riely GJ, Hayes S, Yu H, Zheng J, Moskowitz CS, et al. Are there imaging characteristics associated with lung adenocarcinomas harboring ALK rearrangements? Lung Cancer. 2014;86(2):190–4.

- Choi CM, Kim MY, Hwang HJ, Lee JB, Kim WS. Advanced adenocarcinoma of the lung: comparison of CT characteristics of patients with anaplastic lymphoma kinase gene rearrangement and those with epidermal growth factor receptor mutation. Radiology. 2015;275(1):272–9.
- Rizzo S, Petrella F, Buscarino V, De Maria F, Raimondi S, Barberis M, et al. CT radiogenomic characterization of EGFR, K-RAS, and ALK mutations in nonsmall cell lung cancer. Eur Radiol. 2016;26(1):32–42.
- Yamamoto S, Korn RL, Oklu R, Migdal C, Gotway MB, Weiss GJ, et al. ALK molecular phenotype in non-small cell lung cancer: CT radiogenomic characterization. Radiology. 2014;272(2):568–76.
- Vallonthaiel AG, Jain D, Madan K, Arava S. Pulmonary adenocarcinoma with signet ring features: detailed cytomorphologic analysis. Diagn Cytopathol. 2016;44(7):607–11.
- Rodig SJ, Mino-Kenudson M, Dacic S, Yeap BY, Shaw A, Barletta JA, et al. Unique clinicopathologic features characterize ALK-rearranged lung adenocarcinoma in the western population. Clin Cancer Res. 2009;15(16):5216–23.

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