An observational study of the epidermal growth factor receptor-tyrosine kinase inhibitor resistance mechanism in epidermal growth factor receptor gene mutation-positive non-small cell lung cancer

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Abstract
Non-small cell lung cancer (NSCLC) patients with epidermal growth factor receptor (EGFR) mutation show a high response to EGFR-tyrosine kinase inhibitor (EGFR-TKI). Clinically, EGFR-positive NSCLC acquires several resistance mechanisms during EGFR-TKI treatment, such as the emergence of a secondary mutation (T790M), MET gene amplification, and transformation to small cell lung cancer. However, the mechanism of resistance to afatinib, a second-generation EGFR-TKI, remains unclear. In this study, we prospectively investigate the mechanism of resistance to afatinib using proteomic analyses.

In total, 35 EGFR-positive NSCLC patients of both sexes and ≥20 years old will be included. NSCLC patients with major obstacles in major organs, such as bone marrow, heart, lung, liver, and kidney, will be excluded. Eligible patients will be administered afatinib or gefitinib until disease progression and proteomic analysis will be performed with biopsy samples before treatment and at disease progression.

The primary outcome is to detect the potential predictive anomalies in proteins that can be candidates for the resistance factor of afatinib. The secondary outcome is to detect gene and protein abnormalities affecting progression-free survival, response rate, and rate of disease control in afatinib therapy.

The protocol was approved by the institutional review boards of Kyoto Prefectural University of Medicine and all the participating hospitals. Written informed consent was obtained from all patients before registration, in accordance with the Declaration of Helsinki.

The results of the study will be disseminated via publications in peer-reviewed journals.

Trial registration number is UMIN000031013.

Abbreviations: DCR = disease control rate, EGFR-TKI = epidermal growth factor receptor-tyrosine kinase inhibitor, NSCLC = non-small cell lung cancer, ORR = objective response rate, PFS = progression-free survival.

Keywords: afatinib, EGFR mutation, EGFR-TKI resistance, NSCLC

1. Introduction
In recent years, treatment advances against non-small cell lung cancer (NSCLC) with driver mutations in genes, such as epidermal growth factor receptor (EGFR), have revealed that patients with driver mutations show greater responses to tyrosine kinase inhibitors (TKIs) compared to chemotherapy. Among EGFR-positive NSCLC patients, treatment with EGFR-TKIs showed significantly longer progression-free survival (PFS) compared to platinum doublet therapy.

Afatinib, a second-generation EGFR-TKI, is an irreversible ErbB family binder and blocker that inhibits EGFR, HER2, and HER4. At present, gefitinib, erlotinib, and afatinib have been approved for EGFR-positive NSCLC patients as a first-line treatment. For the first time, afatinib among EGFR-TKIs showed significantly longer PFS and overall survival compared to platinum doublet therapy in patients with deletion of exon 19 in EGFR. Moreover, afatinib significantly prolonged the PFS compared to gefitinib. On the other hand, strong digestive organ toxicity and dermal toxicity were observed; therefore, afatinib was not recommended in poor performance status and the elderly. However, by adjusting the amount of medicine to reduce these side effects, patients were reported to be able to continue afatinib therapy with a therapeutic effect equivalent to that reported previously in a study including the elderly.
Almost all EGFR-positive NSCLC patients acquire several resistance mechanisms during EGFR-TKIs treatment, such as the emergence of a secondary mutation (T790M), MET gene amplification, and transformation to small cell lung cancer.\textsuperscript{[12,13]} Of these, EGFR-T790M is the most common resistance mechanism and found in approximately 50% of cases of resistance during the administration of first-generation EGFR-TKIs.\textsuperscript{[12,14]} Osimertinib, a third-generation EGFR-TKI, is approved for EGFR-T790M-positive NSCLC patients who experienced disease progression during the first and/or second EGFR-TKI treatment because the AURA-3 study reported that osimertinib significantly prolonged PFS compared to platinum-pemetrexed therapy in EGFR-T790M-positive NSCLC patients after resistance.\textsuperscript{[15]}

Little is known regarding the presence of associations between the different responses to EGFR-TKIs and the emergence of T790M. As EGFR-T790M detection was reported to be associated with the initial EGFR-TKI duration and response rate,\textsuperscript{[16]} afatinib treatment is expected to be associated with the greater emergence of EGFR-T790M in basic and clinical research.

Recently, advances in the proteomic analysis have revealed the anomalies in genes and proteins in increased detail. At this time, we are prospectively investigating the tumor proteome before the different responses to EGFR-TKIs and the emergence of EGFR-T790M in basic and clinical research.

The study is a prospective observational study. Figure 1 depicts a flowchart of the study.

2. Methods and analysis

2.1. Study design

The study is a prospective observational study. Figure 1 depicts a flowchart of the study.

2.2. Study setting

Four hospitals agreed to participate in this study. The study protocol was approved by the institutional review board of each hospital. Written informed consent was obtained from all the patients before registration in accordance with the Declaration of Helsinki. The samples are being analyzed by the National Cancer Center Hospital. At least annual independent monitoring is planned in accordance with the Japanese clinical trial guidelines.

2.3. Participants

The inclusion criteria are as follows:

(1) Patients with histologically or cytologically confirmed stage IIIb/IV NSCLC and postoperative recurrence.
(2) Chemotherapy-naive patients planning for afatinib or first-generation EGFR-TKI treatment.
(3) Patients whose tissues can be harvested for the study before afatinib or first-generation EGFR-TKI treatment and resistance.
(4) Performance status (ECOG) 0 to 1.
(5) Patients with a life expectancy of at least 3 months.
(6) Patients whose consent has been obtained in the document for this study.
(7) Patients who are ≥20 years of age (at the time of enrollment).
(8) Patients for whom bone marrow, hepatic, and renal functions have all been confirmed as normal within 14 days before enrollment according to the following clinical test standards:
   1) WBC ≥ 3000/mm\(^3\) to ≤12,000/mm\(^3\),
   2) Neutrophil count ≥ 1500/mm\(^3\),
   3) Platelet count ≥ 100,000/mm\(^3\),
   4) Hemoglobin ≥ 9.0 g/dL,
   5) AST, ALT ≤ 100 IU/L,
   6) Total bilirubin ≤ 1.5 mg/dL,
   7) Creatinine ≤ 2.0 mg/dL,
   8) SpO\(_2\) (room air) ≥ 90%, and
   9) Participation in “prospective observational research (LC-SCRUM)” to clarify the clinicopathological and molecular biological characteristics of low-frequency genetic change positive lung cancer, such as presence of the RET fusion gene.

The exclusion criteria are as follows:

(1) Patients with pulmonary disorders, such as idiopathic pulmonary fibrosis, interstitial pneumonia, pneumoniosis, active radiation pneumonitis, and drug-induced pneumonia,
(2) Patients with infectious disorders requiring intravenous injection of antibacterial drugs or antymycotics,
(3) Patients unable to swallow oral medications,
(4) Patients who are pregnant, nursing, or possibly pregnant,
(5) Patients with symptomatic brain metastases,
(6) Patients with active double cancer,
(7) Patients with uncontrollable diabetes mellitus,
(8) Patients with complications of clinical concern (such as uncontrollable cardiac disease, severe cardiac arrhythmia requiring medical treatment, and sustained serious diarrhea), and
(9) Any other patients who are regarded as unsuitable for this study by the investigators.

2.4. Dose and treatment regimens

Afatinib (40 mg) or gefitinib (250 mg) OD tablet is administered orally. Oral administration of these drugs is continued until disease progression or the criteria for discontinuation.

2.5. Rationale for the setting of the number of enrolled subjects

As this study is a prospective observational study for biomarker detection, the number of enrolled subjects is not set. Enrollment of 35 subjects who met the eligibility criteria until March 2020 was planned.

2.6. Proteomic analysis

Samples at diagnosis and upon rebiopsy after resistance are cryopreserved at −80°C after washing with physiological saline. If rebiopsy samples are diagnosed with NSCLC, proteomic analysis will be performed at National Cancer Center Hospital as follows: comprehensive protein expression analysis using mass spectrometry and exhaustive phosphoenzyme activity analysis using PamStation.

2.7. Statistical methods

PFS: survival curve, median (Kaplan–Meier method), confidence interval of the median (Brookmeyer and Crowley method), and standard error of the annual rate (Greenwood method).
ORR: response rate and its two-sided 95% confidence interval (Wilson method). Statistical significance is considered when the lower limit of the estimated confidence interval exceeds a threshold of 35%.

DCR: disease control rate and its two-sided 95% confidence interval (Wilson method).

2.8. Ethics

The trial received ethical approval from the Ethics Committee of Kyoto Prefectural University of Medicine, Kyoto, Japan (number: ERB-C-1106). The trial is subject to the supervision and management of the Ethics Committee.

2.9. Trial status

This study opened to recruitment in May 2018, with a planned last follow-up in March 2020. As of August 2018, 4 subjects have been enrolled.

3. Discussion

Ultimately, EGFR-positive NSCLC patients acquire resistance to EGFR-TKI during therapy. EGFR-T790M, which is found in approximately 50% of patients after resistance, is one of the several resistance mechanisms involved.[12,14] If we detect T790M in rebiopsy samples, EGFR-T790M-positive patients could be administered osimertinib. Several investigations have shown T790M was significantly detected in EGFR-positive NSCLC patients with a longer duration of initial EGFR-TKI and a higher response rate to initial EGFR-TKI. Afatinib differs from first-generation EGFR-TKIs in binding and blocking the ErbB family that inhibits EGFR, HER2, and HER4 irreversibly as shown in basic studies.[6–8] Moreover, afatinib therapy was superior to gefitinib in clinical research with respect to PFS and ORR.[10] The emergence of T790M might show a greater increase during afatinib therapy than during gefitinib therapy, while resistance mechanisms differing from T790M, MET amplification, and transformation to small cell lung cancer might emerge. Therefore, evaluation of the
tumor proteome before treatment and after the acquisition of resistance using proteomic analysis is needed.

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References

[1] Mok TS, Wu YL, Thongprasert S, et al. Gefitinib or carboplatin-paclitaxel in pulmonary adenocarcinoma. N Engl J Med 2009;361:947–57. doi: 10.1056/NEJMoA0810699.

[2] Mitsudomi T, Morita S, Yatabe Y, et al. Gefitinib versus cisplatin plus docetaxel in patients with non-small-cell lung cancer harbouring mutations of the epidermal growth factor receptor (WJTOG3405): an open-label, randomised phase 3 trial. Lancet Oncol 2010;11:121–8. doi: 10.1016/S1470-2045(09)70364-X.

[3] Inoue A, Kobayashi K, Maemondo M, et al. Updated overall survival results from a randomized phase III trial comparing gefitinib with carboplatin-paclitaxel for chemo-naive non-small-cell lung cancer with sensitive EGFR gene mutations (NEJ002). Ann Oncol 2013;24:54–9. doi: 10.1093/annonc/mdt214.

[4] Zhou C, Wu YL, Chen G, et al. Erlotinib versus chemotherapy as first-line treatment for patients with advanced EGFR mutation-positive non-small-cell lung cancer (OPTIMAL, CTONG-0802): a multicentre, open-label, randomised, phase 3 study. Lancet Oncol 2011;12:735–42. doi: 10.1016/S1470-2045(11)70184-X.

[5] Rosell R, Carcereny E, Gervais R, et al. Erlotinib versus standard chemotherapy as first-line treatment for European patients with advanced EGFR mutation-positive non-small-cell lung cancer (EUR TAC): a multicentre, open-label, randomised phase 3 trial. Lancet Oncol 2012;13:239–46. doi: 10.1016/S1470-2045(11)70393-X.

[6] Yang JC, Shih JY, Su WC, et al. Afatinib for patients with lung adenocarcinoma and epidermal growth factor receptor mutations (LUX-Lung 2): a phase 2 trial. Lancet Oncol 2012;13:539–48. doi: 10.1016/S1470-2045(12)70086-4.

[7] Sequist LV, Yang JC, Yamamoto N, et al. Phase III study of afatinib or cisplatin plus gemcitabine for first-line treatment of Asian patients with advanced non-small-cell lung cancer harbouring EGFR mutations (LUX-Lung 6): an open-label, randomised phase 3 trial. Lancet Oncol 2014;15:213–22. doi: 10.1016/S1470-2045(13)70604-1.

[8] Wu YL, Zhou C, Hu CP, et al. Afatinib versus cisplatin plus gemcitabine for first-line treatment of Asian patients with advanced non-small-cell lung cancer harbouring EGFR mutations (LUX-Lung 6): a phase 3 trial. Clin Cancer Res 2013;19:5327–34. doi: 10.11200/JCO.2012.44.2806.

[9] Yang JC, Wu YL, Schuler M, et al. Afatinib versus cisplatin-based chemotherapy for EGFR mutation-positive lung adenocarcinoma (LUX-3 and LUX-Lung 6): analysis of overall survival data from two randomised, phase 3 trials. Lancet Oncol 2015;16:141–51. doi: 10.1016/S1470-2045(14)7173-8.

[10] Park K, Tan EH, O’Byrne K, et al. Afatinib versus gefitinib as first-line treatment of patients with EGFR mutation-positive non-small-cell lung cancer (LUX-Lung 7): a phase 2B, open-label, randomised controlled trial. Lancet Oncol 2016;17:577–89. doi: 10.1016/S1470-2045(16)30033-X.

[11] Wind S, Schnell D, Ebner T, et al. Clinical pharmacokinetics and pharmacodynamics of afatinib. Clin Pharmacokinet 2017;56:235–50. doi: 10.1007/s40262-016-0440-1.

[12] Sequist LV, Waltman BA, Dias-Santagata D, et al. Genotypic and histological evolution of lung cancers acquiring resistance to EGFR inhibitors. Sci Transl Med 2011;3:75ra26. doi: 10.1126/scitranslmed.3002003.

[13] Yano S, Wang W, Li Q, et al. Hepatocyte growth factor induces gefitinib resistance of lung adenocarcinoma with epidermal growth factor receptor-activating mutations. Cancer Res 2008;68:9479–87. doi: 10.1158/0008-5472.CAN-08-1643.

[14] Wu SG, Liu YN, Tsai MF, et al. The mechanism of acquired resistance to irreversible EGFR tyrosine kinase inhibitor-afatinib in lung adenocarcinoma patients. Oncotarget 2016;7:12404–6. doi: 10.18632/oncotarget.11267.

[15] Oya Y, Yoshida T, Kuroda H, et al. Association between EGFR T790M status and progression patterns during initial EGFR-TKI treatment in patients harboring EGFR mutation. Clin Lung Cancer 2017;18:698–705.e2. doi: 10.1016/j.cllc.2017.05.004.