A Phase II Study of Osimertinib in Combination With Selumetinib in EGFR Inhibitor–Naïve Advanced EGFR-Mutant Lung Cancer

Principal Investigator: Pasi Janne, MD

Condition: Non–small cell lung cancer

Institution: Dana-Farber Cancer Institute

This phase II clinical trial is studying a combination of 2 targeted therapies as a possible treatment for non–small cell lung cancer (NSCLC) with an *EGFR* mutation. Osimertinib alone has been shown to benefit some patients who have received prior treatment for their *EGFR*-mutant NSCLC. Selumetinib is to be administered orally intermittently (4 days on, 3 days off), and osimertinib is to be administered orally on a daily basis. The main purpose of the study is to evaluate information on any potential side effects that this drug combination may cause and collect data on how NSCLC responds to the combination of drugs. Specific tyrosine kinase inhibitors have been approved by the FDA for first-line treatment of patients with NSCLC with an *EGFR* mutation.

Primary Objective:

- Best objective response (time frame: 2 years)

Secondary Objectives:

- Progression-free survival (time frame: 2 years)
- Overall survival (time frame: 2 years)
- Tolerability (time frame: 2 years)
- Toxicity (time frame: 2 years)

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NCCN ORP studies typically explore new avenues of clinical investigation and seek answers to important cancer-related questions. All studies are approved and funded through a scientific peer-review process and are overseen by the ORP.

This feature highlights an NCCN study funded through the grant mechanism.

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For more information on specific trials, including patient selection criteria, use the contact information listed with each study.

For more information on the NCCN ORP, including a complete detailing of the clinical studies currently underway at NCCN Member Institutions, go to www.nccn.org/education-research/nccn-oncology-research-program/orp-main-page.