Preclinical and clinical studies have informed the development of increasingly effective cancer therapies. However, in the majority of cases patients subsequently develop resistance to the therapies that previously worked.

This collaborative team comprises cancer biologists, physician scientists with expertise in clinical oncology, and mathematical modelers. Using patient samples of two cancers as test cases (acute myeloid leukemia and non-small cell lung cancer), they are investigating the dynamics of therapeutic response and resistance in patients. These models will change in response to treatment and tumor evolution, allowing investigators to computationally test millions of possible treatment regimens and select the most promising results for examination in cell culture, mouse models, and eventually in clinical trials. This research will help scientists understand the emergence by cancer cells of resistance to therapies and to test new treatments to overcome that resistance.

The team reported the following progress:

April 2019

- The team is analyzing longitudinal samples obtained from leukemia patients with minimal residual disease (MRD). Team members’ observations from this set of samples include the discovery of a subset of mutations associated with a higher risk of developing metastatic disease, present in both the pre-therapy and the MRD samples, and a set of mutations that expand in response to chemotherapy and may be involved in resistance.
- The team is also studying drug resistance in both lung cancer and leukemia to gain an understanding of the time frame in which resistance can develop, as well as investigating combination therapies to overcome single-agent resistance.

January 2018

- Began enrolling EGFR mutant non-small cell lung cancer patients in clinical trial of first-line gefitinib + EGFR816.
- Performing single-cell RNA sequencing on pre-treatment and on-treatment pleural effusions and core biopsies.
- Continued genomic studies of minimal residual disease in AML patients.
- Began a focus on the statistics and dynamics of mutational diversity of normal and precancerous blood.
Team Progress Updates

- Initiated analysis of an initial set of whole genome sequence data from seven AML samples at diagnosis and four metastases each from two autopsies of non-small cell lung cancer patients.

**September 2017**

- For studies of AML, the team has pursued the molecular characterization of response and resistance via bone marrow sampling, pre-therapy, after best response, and at the time of relapse.
- 31 AML cases have been accrued with banked pre-treatment and clinical remissions specimens collected.
- Early data reveal that mutations present at diagnosis tend to also be present in MRD, and to be variably represented in differentiated mononuclear cells in the remission biospecimens.
- The team undertook genome sequencing from a limited number of non-small cell lung cancer cases for which serial biopsies and autopsy tissues were available.
- A computational study has examined rearrangement junctions in AML looking for nearby DNA binding factor motifs.

**November 2016**

- Multiple samples from different times and tissues have been assembled from 11 individuals to examine the evolution of resistance in non-small cell lung cancer patients.
- Initiated studies of the T-cell repertoire in 250 elderly patients with AML at the time of diagnosis.
- Work to date suggests that genetic mechanisms of resistance evolves during treatment and an understanding of that process in patients may point to innovative therapeutic strategies.