



**TARGET → RARE → CANCER → KNOWLEDGE**

**TCF-001 TRACK is a rare cancer precision medicine clinical trial, NCT04504604.**

TRACK provides participating rare cancer patients and their physicians with personalized, actionable information to potentially inform treatment, as well as recommendations for on-label, off-label, or clinical trial treatments from an expert panel of rare cancer clinicians and scientists. Simultaneously, TRACK generates critical genomic data to drive a better understanding of often overlooked rare cancers.

**TRACK is currently open to enrollment for:**

Patients with any rare cancer (defined as a solid tumor or lymphoma occurring in less than 6 per 100,000 people per year in the US). TRACK will specifically enroll 100 patients with cholangiocarcinoma.

Patients with cancer of unknown primary.

**How TRACK Works:**

→ Qualifying patients can enroll in TRACK from their home using a remote consenting system, allowing full participation with no requirement to travel or change their treating physician.

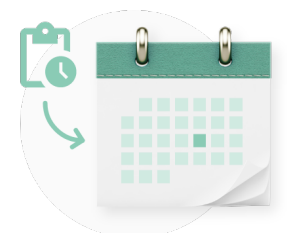


→ Patients enrolled in TRACK receive comprehensive genomic profiling (Foundation One CDx and Foundation One Liquid CDx) at no cost.

→ The TRACK Virtual Molecular Tumor Board, composed of field-leading rare cancer experts, convenes to review the resulting reports and other data, and provides treatment recommendations to the patient and their treating physician.



→ Over the year that follows, the TCF study team collects updated study-related medical information from each patient. In addition, comprehensive genomic profiling of blood is repeated multiple times to identify new alterations which could potentially drive additional treatment recommendations.



**To learn more about TRACK and how to enroll, visit [www.targetcancerfoundation.org/track](http://www.targetcancerfoundation.org/track), or call 617-299-0389.**