

Pancreatic Ductal Adenocarcinoma

A Study to Evaluate the Safety and Activity of GDC-7035 With Anti-Cancer Therapies in Participants With KRAS G12D-Mutated Unresectable Advanced or Metastatic Colorectal Cancer or Pancreatic Ductal Adenocarcinoma

Trial Status
Not yet recruiting

Trial Runs In

Trial Identifier
NCT07859085 2026-525460-17-00
GO46205

The information is taken directly from public registry websites such as ClinicalTrials.gov, EuClinicalTrials.eu, ISRCTN.com, etc., and has not been edited.

Official Title:

A Phase Ib, Multicenter, Randomized Study Evaluating the Safety and Activity of GDC-7035 With Anti-Cancer Therapies as First-Line Treatment in Patients With KRAS G12D-Mutated Unresectable Advanced or Metastatic Colorectal Cancer or Pancreatic Ductal Adenocarcinoma

Trial Summary:

The primary purpose of this study is to evaluate the safety of GDC-7035 as a single agent and in combination with other anti-cancer therapies in participants with KRAS G12D-mutated metastatic colorectal cancer (CRC) or pancreatic ductal adenocarcinoma (PDAC). The study consists of a safety run-in period followed by a randomized/expansion period.

Genentech, Inc.
Sponsor

Phase 1
Phase

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Trial Identifiers

Eligibility Criteria:

Gender
All

Age
#18 Years

Healthy Volunteers
No

Inclusion Criteria:

ForPatients

by Roche

- Confirmed presence of the Kirsten rat sarcoma viral oncogene homolog (KRAS) G12D mutation as documented through central laboratory testing or prior local laboratory testing
- Agreement to adhere to the contraception requirements described in the protocol for participants of childbearing potential and participants who produce sperm

Exclusion Criteria:

- Malabsorption syndrome or other conditions that would interfere with enteral absorption
- Known and untreated, or active central nervous system (CNS) metastases
- Known clinically significant liver disease
- History of or active clinically significant cardiovascular dysfunction

Other protocol-defined inclusion/exclusion criteria may apply.