ForPatients
by Roche

Metastatic Colorectal Cancer

Study To Evaluate Safety, Pharmacokinetics, Pharmacodynamics, And Preliminary Anti-Tumor Activity Of RO7122290 In Combination With Cibisatamab With Obinutuzumab Pre-Treatment

Trial Status	Trial Runs In	Trial Identifier
Completed	5 Countries	NCT04826003 BP42675

The source of the below information is the publicly available website ClinicalTrials.gov. It has been summarised and edited into simpler language.

Trial Summary:

This is an open-label, multicenter, Phase Ib study to determine the maximum tolerated dose (MTD) and/or the recommended Phase 2 dose (RP2D) in the weekly (QW) and/or every 3 weeks (Q3W) regimens, safety, tolerability, PK, immunogenicity, PD profile and to evaluate preliminary anti-tumor activity of RO7122290 in combination with cibisatamab Q3W after pretreatment with obinutuzumab, in participants with previously treated metastatic, microsatellite-stable colorectal adenocarcinoma with high CEACAM5 expression

Hoffmann-La Roche
Sponsor

Phase 1/Phase 2
Phase

NCT04826003 BP42675
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	>=18 Years	No