

Ulcerative Colitis

A Study to Evaluate the Activity, and Safety of Vixarelimab in  
Participants With Moderate to Severe Active Ulcerative Colitis

|                                   |                                   |                                                                     |
|-----------------------------------|-----------------------------------|---------------------------------------------------------------------|
| <b>Trial Status</b><br>Recruiting | <b>Trial Runs In</b><br>1 Country | <b>Trial Identifier</b><br>NCT06693908 2024-516287-29-00<br>GA45735 |
|-----------------------------------|-----------------------------------|---------------------------------------------------------------------|

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 Ic Open-label Study to Evaluate the Pharmacodynamic Effects, Pharmacokinetics, and Safety of Vixarelimab in Patients With Moderate to Severe Active Ulcerative Colitis

Trial Summary:

This study aims to evaluate the pharmacodynamic (PD) effects of vixarelimab in the gut of participants with active moderate to severe ulcerative colitis (UC).

|                                                            |                                 |
|------------------------------------------------------------|---------------------------------|
| <b>Genentech, Inc.</b><br>Sponsor                          | <b>Phase 1/Phase 2</b><br>Phase |
| NCT06693908 2024-516287-29-00 GA45735<br>Trial Identifiers |                                 |

Eligibility Criteria:

|                      |                         |                                 |
|----------------------|-------------------------|---------------------------------|
| <b>Gender</b><br>All | <b>Age</b><br>#18 Years | <b>Healthy Volunteers</b><br>No |
|----------------------|-------------------------|---------------------------------|

Inclusion Criteria:

- Diagnosis of UC established at least 3 months
- Moderately to severely active UC
- Participants must meet criteria for either advanced therapy failure or conventional therapy failure

Exclusion Criteria:

- Diagnosis of Crohn's disease or indeterminate colitis

# ForPatients

*by Roche*

- Suspicion of ischemic colitis, radiation colitis, microscopic colitis or infectious colitis
- Prior colectomy
- Prior treatment with systemic janus kinase (JAK) inhibitors