An Induction Study to Investigate the Efficacy and Safety of Duvakitug in Participants With Moderately to Severely Active Ulcerative Colitis

Status: RECRUITING

Eligibility Criteria

Age: 16 years to 80 years old

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

* Participants aged ≥18 and ≤80 years of age at Screening. Where permitted locally, participants 16 to \<18 years of age who meet the definition of Tanner Stage 5 for development * Confirmed diagnosis of moderately to severely active UC for at least 3 months prior to Baseline * Demonstrated inadequate response, have shown loss of response or intolerance to conventional therapies or advanced therapies

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Exclusion Criteria:

* Participants with Crohn's Disease (CD), indeterminate colitis * Current diagnosis of Ulcerative Proctitis * Participants with surgical bowel resection within the past 3 months prior to Baseline, or a history of \>3 bowel resections * Prior or current high-grade gastrointestinal (GI) dysplasia * Participants on treatment with but not on stable doses of conventional therapies prior to baseline * Participants with prohibited medications or therapies prior to baseline * Participants with previous exposure to anti-TL1A investigational therapy The above information is not intended to contain all considerations relevant to a participant's potential participation in a clinical trial.

Conditions & Interventions

Interventions:
DRUG: Duvakitug, DRUG: Placebo
Conditions:
Ulcerative Colitis

More Information

Contact(s): Trial Transparency email recommended (Toll free for US & Canada) - contact-us@sanofi.com
Principal Investigator:
IRB
Number:
System ID: NCT07184996

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