

Solid TumorsHealthy Volunteers

## A Clinical Trial to Investigate the Clinical Drug-Drug Interaction of Divarasib With Probe Substrates of P-Glycoprotein and Breast Cancer Resistance Protein in Healthy Participants

**Trial Status**  
Completed

**Trial Runs In**  
1 Country

**Trial Identifier**  
NCT06677957 GP45712

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 I, Open-Label, Two-Period, One-Sequence, Crossover Study in Healthy Subjects to Evaluate the Clinical Drug-Drug Interaction of Divarasib With Probe Substrates of P-Glycoprotein (Digoxin) and Breast Cancer Resistance Protein (Rosuvastatin)

### Trial Summary:

This is a Phase 1, open-label, two-period, one-sequence, crossover drug-drug interaction study to assess the P-gp and BCRP inhibition potential of divarasib using digoxin and rosuvastatin as probe substrates, respectively, in healthy participants.

**Genentech, Inc.**  
Sponsor

**Phase 1**  
Phase

**NCT06677957 GP45712**  
Trial Identifiers

### Eligibility Criteria:

**Gender**  
All

**Age**  
#18 Years & # 60 Years

**Healthy Volunteers**  
Accepts Healthy Volunteers

### Inclusion Criteria:

- Males or females of non-childbearing potential
- Within body mass index (BMI) range of 18.0 to 32.0 kg/m<sup>2</sup>, inclusive

### Exclusion Criteria:

# ForPatients

*by Roche*

- Significant history or clinical manifestation of any metabolic, allergic, dermatological, hepatic, renal, hematological, pulmonary, cardiovascular, gastrointestinal (GI), neurological, or psychiatric disorder
- History of significant hypersensitivity, intolerance, or allergy to any drug compound, food, or other substance, unless approved by the investigator
- Poor peripheral venous access