ForPatients

by Roche

Multiple Myeloma

A Dose-escalation Study to Evaluate the Safety, Pharmacokinetics, and Activity of XmAb24306 in Combination With Cevostamab in Participants With Relapsed/Refractory Multiple Myeloma

 Trial Status
 Trial Runs In
 Trial Identifier

 Recruiting
 7 Countries
 NCT05646836

 2022-001204-18,2023-505212-38-00
 GO43980

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

Trial Summary:

The purpose of this study is to evaluate the safety, tolerability, pharmacokinetics, and activity of XmAb24306 in combination with cevostamab in participants with relapsed/refractory multiple myeloma (R/R MM) who have received a minimum of three prior treatments, including at least one immunomodulatory drug (IMiD), one proteasome inhibitor (PI), and one anti-CD38 monoclonal antibody.

Genentech, Inc. Sponsor		Phase 1 Phase
NCT05646836 2022-001204-18,2023-505212-38-00 GO43980 Trial Identifiers		
Eligibility Criterio	a:	
Gender All	Age >=18 Years	Healthy Volunteers No