ForPatients
by Roche

Multiple Myeloma

A Study to Evaluate the Safety and Pharmacokinetics of XMAB24306 in Combination With Daratumumab in Participants With Relapsed/ Refractory Multiple Myeloma

Trial Status	Trial Runs In	Trial Identifier
Completed	4 Countries	NCT05243342 GO43073

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 Ib, Open-Label, Multicenter, Dose-Escalation Study to Evaluate the Safety and Pharmacokinetics of XMAB24306 in Combination With Daratumumab in Patients With Relapsed/Refractory Multiple Myeloma

Trial Summary:

This study will evaluate the safety, tolerability, pharmacokinetics, and activity of XmAb24306 in combination with a multiple myeloma (MM)-targeting monoclonal antibody capable of inducing antibody-dependent cellular toxicity (ADCC) in participants with relapsed or refractory (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

NCT05243342 GO43073
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	#18 Years	No

Inclusion Criteria:

- Life expectancy of at least 12 weeks
- Measurable disease, as defined by the protocol

- Participants must have received a minimum of 3 prior lines of therapy, including at least one PI, one IMiD, and an anti-CD38 monoclonal antibody
- Best response of stable disease or better with at least one prior anti-CD38 monoclonal antibody containing line of treatment

Exclusion Criteria:

- Any anti-cancer therapy within 3 weeks prior to initiation of study treatment, with exceptions defined by the protocol
- Prior allogeneic stem cell or solid organ transplantation
- Autologous stem cell transplantation within 100 days prior to initiation of study treatment
- Significant cardiovascular disease
- Known clinically significant liver disease
- Active or history of autoimmune disease or immune deficiency
- Known active infection requiring IV anti-microbial therapy within 14 days prior to first study drug administration
- Primary or secondary plasma cell leukemia
- Current CNS involvement by MM
- Other protocol defined inclusion/exclusion criteria may apply