

Study Assessing Activity of Intravenous (IV) Etentamig Monotherapy Versus Standard Available Therapies in Adult Participants With Relapsed or Refractory Multiple Myeloma

Status: RECRUITING

Eligibility Criteria

Age: 18 years and over

This study is NOT accepting healthy

Healthy Volunteers: volunteers

Inclusion Criteria:

* Eastern Cooperative Oncology Group (ECOG) performance of ≤ 2 . * Diagnosis of relapsed/refractory (R/R) multiple myeloma (MM) during or after the participant's last treatment as stated in the protocol. * Must have measurable disease with at least 1 of the following assessed within 28 days of enrollment: * Serum M-protein ≥ 0.5 g/dL (≥ 5 g/L). * Urine M-protein ≥ 200 mg/24 hours. * In participants without measurable serum or urine M protein, serum free light chain (FLC) ≥ 100 mg/L (10 mg/dL) (involved light chain) and an abnormal serum kappa lambda ratio. * Must have received at least 2 or more lines of therapy, including a proteasome inhibitor (PI), an immunomodulatory imide (IMiD), and an anti-CD38 monoclonal antibody (mAb). * Must be eligible to receive the Investigator's choice standard available therapy (SAT) based on approved prescribing information, previous MM treatment history, and institutional guidelines.

Exclusion Criteria:

* Clinically significant (per Investigator's judgment) drug or alcohol abuse within the last 6 months. * Clinically significant conditions such as but not limited to the following: neurologic, psychiatric, endocrine, metabolic, immunologic, cardiovascular, pulmonary, or hepatic disease within the last 6 months that would adversely affect the participant's participation in the study. * Central nervous system involvement of MM. * Has received B-cell maturation antigen (BCMA)-targeted therapy.

Conditions & Interventions

Interventions:

DRUG: Etentamig, DRUG: Carfilzomib, DRUG: Pomalidomide, DRUG: Elotuzumab, DRUG: Selinexor, DRUG: Bortezomib, DRUG: Dexamethasone

Conditions:

Multiple Myeloma

Keywords:

Cervino, Multiple Myeloma, Etentamig, Carfilzomib, Pomalidomide, Elotuzumab, Selinexor, Bortezomib, Dexamethasone

More Information

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Principal Investigator:

Phase: PHASE3

IRB

Number:

System ID: NCT06158841

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