A Study to Assess Adverse Events and Change in Disease Activity in Participants 12 Years of Age or Older With Locally Advanced or Metastatic Solid Tumors That Harbor MET Amplification Receiving Intravenously Infused Telisotuzumab Adizutecan

NCT07196644

Status	RECRUITING
Phase	Phase 2
Sponsor	AbbVie
Enrollment	125 participants

Key Eligibility Criteria

Inclusion (5)

- Locally advanced/metastatic solid tumors with documented MET amplification via Local next generation sequencing (NGS) or Central NGS via FoundationOne Companion Diagnostic (F1CDx).
- Eastern Cooperative Oncology Group (ECOG) Performance Status of 0 to 1
- Measurable disease at baseline per Response Evaluation Criteria in Solid Tumors (RECIST) version 1.1 or Response Assessment in Neuro-Oncology (RANO) criteria as appropriate to tumor type.
- Received prior systemic therapy appropriate for their tumor type and stage of disease and who have no satisfactory alternative therapy for advanced solid tumors that would be expected to provide a substantial survival benefit for their tumor type.
- If participant has central nervous system (CNS) metastasis, these should be clinically asymptomatic or radiologically stable (i.e., without evidence of progression after definitive treatment

Exclusion (2)

- Current, historical, or suspected (non-infectious) interstitial lung disease (ILD)/pneumonitis that required steroids.
- Any major, life-threatening conditions and life expectancy should be at least 12 weeks.

Locations (26 total)

City of Hope National Medical Center /ID# 275613, Duarte, California, United States
Valkyrie Clinical Trials /ID# 275547, Los Angeles, California, United States
Yale University School of Medicine /ID# 275978, New Haven, Connecticut, United States
... and 23 more locations