ForPatients
by Roche

Advanced Solid TumorsMetastatic Solid TumorsCancer

Safety and Pharmacokinetics (PK) of Escalating Doses of MTIG7192A as a Single Agent and in Combination With Atezolizumab in Locally Advanced or Metastatic Tumors

Trial Status	Trial Runs In	Trial Identifier
Completed	7 Countries	NCT02794571 2016-000944-33 GO30103

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 Ia/Ib Open-Label, Dose-Escalation Study of the Safety and Pharmacokinetics of Tiragolumab as a Single Agent and in Combination With Atezolizumab and/or Other Anti-Cancer Therapies in Patients With Locally Advanced or Metastatic Tumors

Trial Summary:

This first-in-human open-label, multicenter, dose-escalation and expansion study is designed to evaluate the safety, tolerability, and PK of tiragolumab alone or in combination with atezolizumab and/or other anti-cancer therapies in participants with locally advanced, recurrent, or metastatic incurable tumors for whom standard therapy does not exist, has proven to be ineffective or intolerable, or is considered inappropriate, or for whom a clinical trial of an investigational agent is a recognized standard of care.

Genentech, Inc.	**Phase 1**
Sponsor	Phase

NCT02794571 2016-000944-33 GO30103
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	# 18 Years	No

Inclusion Criteria:

- Adults 18 years of age or older
- Eastern Cooperative Oncology Group (ECOG) performance status of 0 or 1
- Life expectancy at least 12 weeks
- Adequate hematologic and end organ function
- Histologic documentation of locally advanced, recurrent, or metastatic incurable malignancy that has progressed after at least one available standard therapy; or for which standard therapy has proven ineffective, intolerable, or considered inappropriate; or for which a clinical trial of an investigational agent is a recognized standard of care
- Confirmed availability of representative tumor specimens
- Measurable disease according to RECIST Version 1.1

Exclusion Criteria:

- Any anti-cancer therapy, whether investigational or approved, including chemotherapy, hormonal therapy, or radiotherapy, within 3 weeks prior to initiation of study treatment
- Malignancies other than disease under study within 5 years prior to Day 1 of Cycle 1
- Primary central nervous system (CNS) malignancy, or untreated/active CNS metastases
- Leptomeningeal disease
- History of idiopathic pulmonary fibrosis, pneumonitis, organizing pneumonia, or evidence of active pneumonitis on Screening chest computed tomograph (CT) scan
- History of autoimmune disease
- Positive human immunodeficiency virus (HIV) test
- Active hepatitis B or C, or tuberculosis
- Severe infection within 4 weeks prior to randomization
- Prior allogeneic bone marrow or solid organ transplant
- Significant cardiovascular disease
- Known clinically significant liver disease