

Cancer

Study Evaluating the Safety, Pharmacokinetics (PK), Pharmacodynamics (PD), and Therapeutic Activity of Selicrelumab (RO7009789) With Vanucizumab or Bevacizumab in Participants With Metastatic Solid Tumors

Trial Status
Completed

Trial Runs In
7 Countries

Trial Identifier
NCT02665416 2015-003480-11
BP29889

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

Trial Summary:

This open-label, two-part study is designed to assess the safety, PK, PD, and therapeutic activity of Selicrelumab in combination with vanucizumab or bevacizumab in participants with metastatic solid tumors not amenable to standard treatment. Part I (dose escalation) is designed to establish the maximum tolerated dose (MTD) of Selicrelumab in this combination. Part II (expansion) is intended to characterize the safety and tolerability of Selicrelumab in combination with bevacizumab among indication-specific cohorts and to confirm the recommended dose.

Hoffmann-La Roche
Sponsor

Phase 1
Phase

NCT02665416 2015-003480-11 BP29889
Trial Identifiers

Eligibility Criteria:

Gender
All

Age
>= 18 Years

Healthy Volunteers
No
