ForPatients
by Roche

Solid Tumors

A Safety and Efficacy Extension Study of Pertuzumab in Patients With Solid Tumors Previously Enrolled in a Hoffmann-La Roche-Sponsored Pertuzumab Clinical Trial

Trial Status	Trial Runs In	Trial Identifier
Active, not recruiting	15 Countries	NCT02320435 2014-002048-42,2023-505102-42-00 MO29406

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 is a single-arm, multi-center, open-label extension study designed to provide continued pertuzumab therapy to patients receiving pertuzumab as an investigational medicinal product (IMP) in a Roche-sponsored global study and who continue to receive pertuzumab at the end of the Parent study, as well as to collect long-term safety and efficacy data of pertuzumab therapy. Patients with solid tumors who have not experienced progressive disease in the Parent study and, in the investigator's opinion, may potentially benefit from continued pertuzumab treatment, will continue to receive pertuzumab until disease progression, unacceptable toxicity, investigator/patient decision, patient non-compliance, patient death, patient request to withdraw, or study termination by the Sponsor, whichever occurs first.

Hoffmann-La Roche Sponsor	**Phase 3** Phase

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Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	>=18 Years	No