

ForPatients

by Roche

Choroidal neovascularization Myopia

A Study to Evaluate the Efficacy and Safety of Faricimab in Patients With Choroidal Neovascularization Secondary to Pathologic Myopia

Trial Status
Recruiting

Trial Runs In
11 Countries

Trial Identifier
NCT06176352 2023-506707-25-00
CR44829

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 Phase III, multicenter, randomized, double-masked, active comparator-controlled study evaluating the efficacy and safety of faricimab in patients with myopic choroidal neovascularization (CNV). This non-inferiority study will compare 6.0 mg faricimab versus 0.5 mg ranibizumab administered at a pro-re-nata (PRN) dosing regimen after an initial active IVT treatment administration at randomization (Day 1).

Hoffmann-La Roche
Sponsor

Phase 3
Phase

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Eligibility Criteria:

Gender
All

Age
≥18 Years

Healthy Volunteers
No
