

A Real-World Study to Gain Clinical Insights Into Faricimab (FaReal Study)

NCT06680817

Status RECRUITING
Sponsor Hoffmann-La Roche
Enrollment 850 participants

Key Eligibility Criteria

Inclusion (3)

- Patients receiving faricimab according to the local faricimab product label and who have initiated treatment with faricimab at time of the ICF signature date or no more than 3 months prior to the ICF signature date, in diabetic macular edema (DME) or neovascular age-related macular degeneration (nAMD) in at least one eye
- Patients have received at least one faricimab treatment (the first dose) in the study eye
- Patients should have available data for visual acuity (VA) and Central Subfield Thickness (CST) for the Baseline level (i.e. examinations to be performed at the index date or within 4 months prior to it)

Exclusion (6)

- Patient participation in any investigational ophthalmology clinical trial that includes receipt of any ophthalmological investigational drug or procedure within the last 28 days prior to the ICF signature date
 - Concomitant participation in any interventional clinical study
 - Active ocular inflammation and/or suspected/active ocular infection in either eye
 - Patients treated with faricimab who have and are currently participating in patient support programs (PSP) that are Market Research and Patient Support Programs (MAP) including Post Trial Access Programs (PTAP) and Compassionate Use Programs (CUP)
 - Patients with non-ocular sight threatening disease which have an effect on the primary endpoint (e.g., apoplexia)
- ... and 1 more (see full listing online)

Locations (55 total)

LKH-Univ.Klinikum Graz, Graz, Austria
Medizinische Universität Innsbruck, Innsbruck, Austria
Kepler Universitätskliniken GmbH - Med Campus III, Linz, Austria
... and 52 more locations