

Real World Evidence in China: Faricimab Use in Diabetic Macular Edema, Retinal Vein Occlusion, and Neovascular Age-Related Macular Degeneration (The Farseeing Study)

NCT06439576

Status RECRUITING
Sponsor Hoffmann-La Roche
Enrollment 1,000 participants

Key Eligibility Criteria

Inclusion (5)

- Have signed the informed consent
- Female and male Chinese patients, who had been diagnosed with nAMD, DME, or RVO by CFP, OCT, FFA, ICGA, or OCTA
- e50years old for patients with nAMD, e18 years old both for patients with DME and RVO, at the time of signing informed consent (or the first administration of faricimab, whichever occurs first)
- Patients for whom the decision to receive treatment with faricimab is made prior to and independent from study participation
- Patients have received at least one faricimab treatment (the first dose) in the study eye

Exclusion (6)

- Patients not receiving treatment for nAMD/DME/RVO with faricimab according to the standard of care and in line with the current summary of product characteristics (SPC) / labeling in China
- Active ocular inflammation or suspected / active ocular infection in either eye
- Received any other anti-VEGF treatment after faricimab
- Received any steroid treatment within 6 months (180 days) before the first faricimab treatment in the study eye
- Any participation in any other clinical trials currently
- ... and 1 more (see full listing online)

Locations (41 total)

Joint Shantou International Eye Center, Shantou, Guangdong, China
The First Affiliated Hospital of Henan UN of Science and Technology, Luoyang, Henan, China
Taihe Hospital of Shiyan, Shiyan, Hubei, China
... and 38 more locations

<https://clinicaltrials.gov/study/NCT06439576>

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