ForPatients

by Roche

Neovascular Age-related Macular Degeneration

A Study Of The Efficacy, Safety, And Pharmacokinetics Of The Port Delivery System With Ranibizumab In Chinese Patients With Neovascular Age-Related Macular Degeneration

Trial Status	Trial Runs In	Trial Identifier
Recruiting	1 Countries	NCT05562947 YR42983

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 study will evaluate the efficacy, safety, and pharmacokinetics of ranibizumab 100 mg/mL delivered Q24W via the PDS implant compared with ranibizumab 0.5 mg delivered as a Q4W intravitreal injection in Chinese patients with nAMD.

Hoffmann-La Roche Sponsor	Phase 3 Phase	
NCT05562947 YR42983 Trial Identifiers		
Eligibility Criteria:		
Gender All	Age >=50 Years & <= 80 Years	Healthy Volunteers