ForPatients
by Roche

Parkinson's Disease (PD)

A Study to Evaluate the Efficacy and Safety of Intravenous (IV) Prasinezumab in Participants With Early-Stage Parkinson's Disease

Trial Status	Trial Runs In	Trial Identifier
Not yet recruiting		NCT07174310 2025-522683-32-00 BN44715

The information is taken directly from public registry websites such as ClinicalTrials.gov, EuClinicalTrials.eu, ISRCTN.com, etc., and has not been edited.

Official Title:

A Phase III, Randomized, Double-Blind, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of Intravenous Prasinezumab in Participants With Early-Stage Parkinson's Disease

Trial Summary:

The purpose of this study is to evaluate the efficacy, safety, and pharmacokinetics (PK) of prasinezumab compared with placebo in participants with early-stage Parkinson's disease (PD) on stable symptomatic monotherapy with levodopa.

Hoffmann-La Roche
Sponsor

Phase 3
Phase

NCT07174310 2025-522683-32-00 BN44715
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	#50 Years & # 85 Years	No

Inclusion Criteria:

- Body weight within 40-110 kilograms (kg) (88-242 pounds [lbs]) and a body mass index within the range 18-34 kg/m2
- Diagnosis of idiopathic PD based on Movement Disorder Society (MDS) criteria
- Has received monotherapy treatment
- An MDS-UPDRS Part IV score of 0 at screening and prior to randomization
- Hoehn and Yahr (H&Y) Stage 1 or 2 off medication at screening and prior to randomization
- Agreement to adhere to the contraception requirements

Exclusion Criteria:

- Pregnant or breastfeeding, or intention of becoming pregnant during the study or within the time frame in which contraception is required
- Medical history indicating a parkinsonian syndrome other than idiopathic PD
- Diagnosis of a significant neurologic disease other than PD
- Chronic uncontrolled hypertension