ForPatients
by Roche

Von Willebrand disease (VWD)

An Observational Study of Participants With Type 3 Von Willebrand Disease on Prophylactic Standard-of-Care Treatment

Trial Status	Trial Runs In	Trial Identifier
Recruiting	3 Countries	NCT06883240 WP45335

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 non-interventional study (NIS) is designed to collect information on the effectiveness and safety of treatment received in routine clinical care, as well as measure the health-related quality of life (HRQoL) of participants with Type 3 von Willebrand disease (VWD) receiving prophylactic therapy per local standard of care (SOC) over an observation period of at least 24 weeks.

Hoffmann-La Roche Sponsor	**N/A** Phase

NCT06883240 WP45335
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	>=2 Years	No