ForPatients
by Roche

Multiple Sclerosis (MS)

A Study of Ocrelizumab in Participants With Relapsing Remitting Multiple Sclerosis (RRMS) Who Have Had a Suboptimal Response to an Adequate Course of Disease-Modifying Treatment (DMT)

Trial Status	Trial Runs In	Trial Identifier
Completed	17 Countries	NCT02861014 2015-005597-38 MA30005

The source of the below information is the publicly available website ClinicalTrials.gov. It has been summarised and edited into simpler language.

Trial Summary:

The purpose of this prospective, multicenter, open-label, efficacy, and safety study is to assess the efficacy and safety of ocrelizumab in participants with Relapsing Remitting Multiple Sclerosis (RRMS) who have had a suboptimal response to an adequate course of a Disease-Modifying Treatment (DMT). The study will consist of a Screening period (up to 4 weeks), an Open-label treatment period (96 weeks; with last dose administered at Week 72), and a Follow-up period of at least 2 years.

Hoffmann-La Roche	Phase 3
Sponsor	Phase

NCT02861014 2015-005597-38 MA30005
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	>=18 Years & <= 55 Years	No