ForPatients
by Roche

Schizophrenia

A study of patients with Schizophrenia (FlashLyte)

A Study of RO4917838 in Participants With Persistent, Predominant Negative Symptoms of Schizophrenia (FlashLyte)

Trial Status	Trial Runs In	Trial Identifier
Completed	1 Countries	NCT01192867 2010-020370-42 (EudraCT Number) NN25310

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 multi-center, randomized, double-blind, parallel-group, placebo-controlled study will evaluate the efficacy and safety of RO4917838 in participants with persistent, predominant negative symptoms of schizophrenia. Participants, on stable treatment with antipsychotics, will be randomized to receive daily oral doses of RO4917838 or matching placebo for 56 weeks (treatment period 1 of 24 weeks and treatment period 2 of 32 weeks), followed by an optional treatment extension for up to 3 years. After 52 weeks, participants who were originally randomized to an active treatment will be randomly assigned to receive either placebo or continue on the originally assigned study treatment for 4 weeks washout period (Week 52 to Week 56) for the assessment of potential withdrawal effects in a blinded manner using participants staying on active treatment as a control. Participants initially randomized to placebo will remain on placebo. After 56 weeks, participants who were switched to placebo in the washout period will return to their blinded, active treatment arm.

Sponsor	Phase
F. Hoffmann-La Roche Ltd	**Phase 3**

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Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	18 Years and older	No