

ForPatients

by Roche

Chronic Lymphocytic Leukemia

A Study to Evaluate the Effect of Venetoclax on Participants Receiving a Covalent Bruton's Tyrosine Kinase Inhibitor (cBTKi) for First-Line Chronic Lymphocytic Leukemia (1L CLL) to Achieve Deep Durable Remissions to Allow Off-Treatment Period

Trial Status
Recruiting

Trial Runs In
1 Countries

Trial Identifier
NCT06524375 ML45219

The source of the below information is the publicly available website [ClinicalTrials.gov](https://clinicaltrials.gov). It has been summarised and edited into simpler language.

Trial Summary:

The main purpose of the study is to evaluate if adding venetoclax to participants receiving cBTKi for the 1L CLL can achieve deep durable remissions of undetectable measurable residual disease [uMRD < or 10^{-4} in peripheral blood (PB)] by end of combination treatment (EOCT) to allow off-treatment period. The acronym BRAVE stands for Btki Responders to Achieve deep remission (or off-treatment periods) with VEnetoclax.

Genentech, Inc.
Sponsor

Phase 2
Phase

NCT06524375 ML45219
Trial Identifiers

Eligibility Criteria:

Gender
All

Age
>=18 Years

Healthy Volunteers
No
