ForPatients
by Roche

Breast Cancer

A Study to Evaluate the Safety and Efficacy of Ipatasertib in Combination With Atezolizumab and Paclitaxel or Nab-Paclitaxel in Participants With Locally Advanced or Metastatic Triple-Negative Breast Cancer

Trial Status	Trial Runs In	Trial Identifier
Completed	5 Countries	NCT03800836 CO40151

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 is a study consisting of four cohorts in this setting. In Cohort 1, the safety and efficacy of ipatasertib (ipat) in combination with atezolizumab (atezo) and paclitaxel (pac) or nab-paclitaxel will be evaluated for participants with locally advanced or metastatic triple-negative breast cancer (TNBC) who have not previously received chemotherapy. In Cohort 2, ipatasertib and atezolizumab (with no chemotherapy), will be administered to participants with locally advanced or metastatic TNBC. In Cohort 3, the safety and efficacy of neoadjuvant ipatasertib, atezolizumab, doxorubicin and cyclophosphamide (AC) (Ipat + Atezo + AC) followed by Ipat + Atezo + Pac will be evaluated in participants with locally advanced Type 2-4 (T2-4) TNBC. In Cohort 4, the safety and efficacy of Ipat + Atezo + Pac will be evaluated in participants with PD-L1 (Programmed Death-Ligand-1) positive locally advanced or metastatic TNBC that is not amenable to resection and who have not previously received chemotherapy in the advanced setting.

Hoffmann-La Roche
Sponsor

Phase 1
Phase

NCT03800836 CO40151
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	>=18 Years	No