

# ForPatients

by Roche

Breast Cancer Er-Positive Breast Cancer Breast Neoplasms

## A Study of Multiple Immunotherapy-Based Treatment Combinations in Hormone Receptor (HR)-Positive Human Epidermal Growth Factor Receptor 2 (HER2)-Negative Breast Cancer

**Trial Status**  
Completed

**Trial Runs In**  
3 Countries

**Trial Identifier**  
NCT03280563 2017-000335-14  
CO39611

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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:***

This study is designed to evaluate the efficacy, safety, and pharmacokinetics of several immunotherapy-based combination treatments in participants with inoperable locally advanced or metastatic HR-positive, HER2-negative breast cancer who have progressed during or following treatment with a cyclin-dependent kinase (CDK) 4/6 inhibitor in the first- or second-line setting, such as palbociclib, ribociclib, or abemaciclib. The study will be performed in two stages. During Stage 1, participants will be randomized to fulvestrant (control) or an atezolizumab-containing doublet or triplet combination. Those who experience disease progression, loss of clinical benefit, or unacceptable toxicity may be eligible to receive a new triplet combination treatment in Stage 2 until loss of clinical benefit or unacceptable toxicity. New treatment arms may be added and/or existing treatment arms may be closed during the course of the study on the basis of ongoing clinical efficacy and safety as well as the current treatments available.

**Hoffmann-La Roche**  
Sponsor

**Phase 1/Phase 2**  
Phase

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**NCT03280563 2017-000335-14 CO39611**  
Trial Identifiers

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### ***Eligibility Criteria:***

**Gender**  
Female

**Age**  
>= 18 Years

**Healthy Volunteers**  
No

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