

# ForPatients

by Roche

Solid Tumors

## A Study to Assess the Safety and Tolerability of Atezolizumab in Combination With Other Immune-Modulating Therapies in Participants With Locally Advanced or Metastatic Solid Tumors

**Trial Status**  
Completed

**Trial Runs In**  
2 Countries

**Trial Identifier**  
NCT02174172 2014-000812-33  
GO29322

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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 global, multicenter, open-label study will evaluate the safety and tolerability of atezolizumab in combination with other immune-modulating therapies in the treatment of selected advanced or metastatic malignancies. The atezolizumab plus ipilimumab arm (Arm A) will focus primarily on participants with advanced or metastatic non-small cell lung cancer (NSCLC). The atezolizumab plus interferon alfa-2b arm (Arm B), plus pegylated interferon alfa-2a (PEG-interferon alfa-2a, Arm C), and atezolizumab plus PEG-interferon Alfa-2a plus bevacizumab (Arm D) will enroll participants with advanced or metastatic renal cell carcinoma (RCC), metastatic NSCLC and melanoma. The atezolizumab plus obinutuzumab (Arm E) will enroll participants with recurrent and/or metastatic (R/M) head and neck squamous cell carcinoma (HNSCC). Atezolizumab will be administered as intravenous (IV) infusion every 3 weeks (q3w).

**Hoffmann-La Roche**  
Sponsor

**Phase 1**  
Phase

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**NCT02174172 2014-000812-33 GO29322**  
Trial Identifiers

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### *Eligibility Criteria:*

**Gender**  
All

**Age**  
≥18 Years

**Healthy Volunteers**  
No

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