

Non Hodgkin Lymphoma (NHL)

## A Dose Escalation Study Evaluating the Safety and Tolerability of GDC-0032 in Participants With Locally Advanced or Metastatic Solid Tumors or Non-Hodgkin's Lymphoma (NHL) and in Combination With Endocrine Therapy in Locally Advanced or Metastatic Hormone Receptor-Positive Breast Cancer

**Trial Status**

Active, not recruiting

**Trial Runs In**

4 Countries

**Trial Identifier**

NCT01296555 GO00886  
PMT4979g

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 is an open-label, multicenter, Phase I/II study to assess the safety, tolerability, and pharmacokinetics of GDC-0032. The Phase I portion will be divided into two stages. During Stage 1, GDC-0032 will be administered every day orally and at escalating doses in participants with locally advanced or metastatic solid tumors. During Stage 2, GDC-0032 will be administered alone or as combination therapy within indication-specific cohorts. In Phase II of the study, the efficacy and safety of the combination GDC-0032 and fulvestrant will be evaluated in post-menopausal female participants with locally advanced or metastatic human epidermal growth factor receptor 2 (HER2)-negative, hormone receptor-positive breast cancer.

**Genentech, Inc.**

Sponsor

**Phase 1**

Phase

**NCT01296555 GO00886 PMT4979g**

Trial Identifiers

### *Eligibility Criteria:*

**Gender**

All

**Age**

**>= 18 Years**

**Healthy Volunteers**

No