

# Study of investigational odronextamab in patients with B-cell malignancies



Primary Objective: evaluate the safety and tolerability of odronextamab in patients with CD20+ B-cell malignancies



PATIENTS WITH B-NHL OR CLL (N≈200)

OPEN LABEL INTERVENTION

## Part A

Dose escalation + dose expansion

Odronextamab (IV)

DLBCL post CAR-T therapy, B-NHL, and CLL cohorts

No longer enrolling

## Primary Endpoints

Safety  
Tolerability

DLBCL after failure of CAR-T therapy  
expansion cohort: Objective response rate

## Secondary Endpoints

Pharmacokinetics  
Immunogenicity  
Objective response rate

Progression-free survival  
Overall survival  
Duration of complete response



### FIND OUT MORE

Scan here to find out more about this study at <https://clinicaltrials.gov/ct2/show/NCT02290951>

This information is intended for investigators interested in open clinical trials.  
Odronextamab is an investigational asset and has not been fully evaluated by any regulatory authority.

B-NHL, B-cell non-Hodgkin lymphoma; CAR-T, chimeric antigen receptor T cell; CD, cluster of differentiation; CLL, chronic lymphocytic leukemia; DLBCL, diffuse large B-cell lymphoma; IV, intravenous; N, number of patients; SC, subcutaneous.

# ODRONEXTAMAB

## An Investigational CD20xCD3 Bispecific Antibody

Simultaneously engages CD20 on malignant B cells and CD3 on cytotoxic T cells<sup>1</sup>

### SELECTED INCLUSION CRITERIA<sup>a</sup>



CD20+ B-cell malignancy, with active disease not responsive to prior therapy



ECOG PS 0 or 1



Measurable lesion  $\geq 1.5$  cm documented by CT or MRI scan



Life expectancy of at least 6 months



Patients with B-NHL must have had prior treatment with an anti-CD20 antibody therapy



Adequate bone marrow and organ function

### SELECTED EXCLUSION CRITERIA<sup>a</sup>



Primary CNS lymphoma or CNS pathology



Non-biologic anti-lymphoma chemotherapy or radiotherapy within 28 days prior to first administration of study drug



Infection with HIV or chronic infection with HBV, HCV, or CMV infection



For more information, visit [www.clinicaltrials.gov](http://www.clinicaltrials.gov) or please call [+353 (0)61 533 400 OR 844 REGN-MID].  
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<sup>a</sup>Inclusion/exclusion criteria include a summary of selected criteria. Please review the complete study design on [clinicaltrials.gov](http://clinicaltrials.gov) for complete details.

B-NHL, B-cell non-Hodgkin lymphoma; CD, cluster of differentiation; CMV, cytomegalovirus; CNS, central nervous system; CT, computed tomography; ECOG PS, Eastern Cooperative Oncology Group performance status; HBV, hepatitis B virus; HIV, human immunodeficiency virus; MRI, magnetic resonance imaging.

1. Smith EJ et al. *Sci Rep*. 2015;5:17943.

Current per [clinicaltrials.gov](http://clinicaltrials.gov) as of September 7, 2023.

Odronextamab-EM-0008 November 2023.

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