

# Study of investigational odronextamab + lenalidomide in patients with **relapsed or refractory follicular lymphoma or marginal zone lymphoma**



Primary Objective: evaluate the efficacy and safety of odronextamab + lenalidomide vs rituximab + lenalidomide in patients with R/R FL or R/R MZL



PATIENTS WITH R/R FL OR R/R MZL (N≈325 R/R FL; N≈70 R/R MZL)

## OPEN LABEL INTERVENTION

### Part 1: non-randomized

Odronextamab safety run-in



### Part 2: randomized

R  
1:1

#### Arm 1

Odronextamab (IV)  
+ lenalidomide

#### Arm 2

Rituximab  
+ lenalidomide

### Primary Endpoint

Progression-free survival<sup>a</sup>

### Secondary Endpoints

Complete response rate

Best overall response

Overall survival

#### FIND OUT MORE

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

This information is intended for investigators interested in open clinical trials.

Odronextamab is an investigational agent and has not been evaluated by any regulatory authority.

<sup>a</sup>Based on Independent Central Review (ICR).

FL, follicular lymphoma; IV, intravenous; MZL, marginal zone lymphoma; N, number of patients; R randomized; R/R, relapsed or refractory

# 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>



Histologically confirmed FL Grade 1–3a or MZL (nodal, splenic, or extra nodal)



Refractory disease or relapsed after  $\geq 1$  prior systemic therapy including an anti-CD20 antibody



Measurable disease



ECOG PS  $\leq 2$

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



CNS lymphoma



Histological evidence of transformation to high grade or diffuse large B-cell lymphoma



Prior use of lenalidomide or any CD20xCD3 bispecific antibody within 6 months

For more information, visit [www.clinicaltrials.gov](http://www.clinicaltrials.gov) or please call [+353 (0)61 533 400 OR 844 REGN-MID].  
NCT  
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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.

CD, cluster of differentiation; CNS, central nervous system; ECOG PS, Eastern Cooperative Oncology Group performance status; IPI, International Prognostic Index.

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

<sup>1</sup>Smith EJ et al. *Sci Rep.* 2015;5:17943

Veeva code November 2023.

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