

**State of Oklahoma  
SoonerCare  
Zynlonta<sup>®</sup> (Loncastuximab Tesirine-Ipyl)  
Prior Authorization Form**

**Member Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Member ID#:** \_\_\_\_\_

**Drug Information**

Physician billing (HCPCS code: \_\_\_\_\_)  Pharmacy billing (NDC: \_\_\_\_\_)

**Start Date (or date of next dose):** \_\_\_\_\_ **Dose:** \_\_\_\_\_

**Dosing Regimen: Cycles 1 & 2** \_\_\_\_\_ **Subsequent Cycles:** \_\_\_\_\_

**Billing Provider Information**

**Provider NPI:** \_\_\_\_\_ **Provider Name:** \_\_\_\_\_

**Provider Phone:** \_\_\_\_\_ **Provider Fax:** \_\_\_\_\_

**Prescriber Information**

**Prescriber NPI:** \_\_\_\_\_ **Prescriber Name:** \_\_\_\_\_

**Prescriber Phone:** \_\_\_\_\_ **Prescriber Fax:** \_\_\_\_\_ **Specialty:** \_\_\_\_\_

**Criteria**

**For Initial Authorization**

**1. Please indicate the diagnosis and information:**

**Diffuse Large B-cell Lymphoma (DLBCL)**

A. Please select 1 of the following:

- \_\_\_\_\_ DLBCL not otherwise specified
- \_\_\_\_\_ DLBCL arising from low grade lymphoma
- \_\_\_\_\_ High-grade B-cell lymphoma
- \_\_\_\_\_ Other, please specify: \_\_\_\_\_

B. Is disease relapsed or refractory after 2 or more lines of systemic therapy? Yes \_\_\_\_\_ No \_\_\_\_\_

C. Was previous CD19-directed therapy was used? Yes \_\_\_\_\_ No \_\_\_\_\_

i. If yes, does the member have a biopsy that shows CD19 protein expression after completion of the CD19-directed therapy? Yes \_\_\_\_\_ No \_\_\_\_\_

1. If yes, please provide biopsy results.

D. Please provide a patient-specific, clinically significant reason why tafasitamab in combination with lenalidomide is not appropriate for the member:

**If diagnosis is not listed of the above, please indicate diagnosis:** \_\_\_\_\_

**Additional Information:** \_\_\_\_\_

**For Continued Authorization:**

1. Date of last dose: \_\_\_\_\_

2. Does the member have any evidence of progressive disease while on Zynlonta<sup>®</sup>? Yes \_\_\_\_\_ No \_\_\_\_\_

3. Has the member experienced adverse drug reactions related to Zynlonta<sup>®</sup> therapy? Yes \_\_\_\_\_ No \_\_\_\_\_

*If yes, please specify adverse reactions:* \_\_\_\_\_

**Prescriber Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

***I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Failure to complete this form in full will result in processing delays.***

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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