

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

**Drug Information**

Physician billing (HCPCS code: \_\_\_\_\_) Start Date: \_\_\_\_\_

**Billing Provider Information**

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

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

**Prescriber Information**

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

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

**Criteria**

**For Authorization:** (approvals will be for 1 dose per member per lifetime)

1. Please include the most recent office visit note or clinical summary from the hospital to support your request. Is this information attached? Yes \_\_\_ No \_\_\_
2. Is the health care facility on the certified list to administer chimeric antigen receptor (CAR) T-cells? Yes \_\_\_ No \_\_\_
3. Is the health care facility trained in the management of cytokine release syndrome (CRS) and neurologic toxicities? Yes \_\_\_ No \_\_\_
4. Will the health care facility comply with the Breyanzi® risk evaluation and mitigation strategy (REMS) program requirements? Yes \_\_\_ No \_\_\_
5. Please indicate the diagnosis and information:

**Large B-cell Lymphoma**

A. Please provide additional information regarding previous therapies member has tried and failed:

\_\_\_\_\_

B. Does the member have any of the following?

- Refractory disease to frontline chemoimmunotherapy.
- Relapse within 12 months of frontline chemoimmunotherapy.
- Relapse after frontline chemoimmunotherapy and is not eligible for hematopoietic stem cell transplantation (HSCT) due to comorbidity or age.
- Relapsed or refractory disease after 2 or more lines of systemic therapy.

C. Does member have primary central nervous system (CNS) lymphoma? Yes \_\_\_ No \_\_\_

D. Please provide a patient-specific, clinically significant reason why Kymriah® (tisagenlecleucel) or Yescarta® (axicabtagene) is not appropriate for the member: \_\_\_\_\_

\_\_\_\_\_

**If answer is none of the above, please indicate diagnosis:** \_\_\_\_\_

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

\_\_\_\_\_

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. Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full and attach requested clinical notes 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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