



Breyanzi® (lisocabtagene maraleucel) Prior Authorization Form

Member Name: _____ **Date of Birth:** _____ **Member ID#:** _____

Drug Information

Physician Billing (HCPCS code: _____ **) Start Date (or date of next dose):** _____

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)

- Please include the most recent office visit note or clinical summary from the hospital to support your request.
Is this information attached? Yes ☐ No ☐
- Is the health care facility on the certified list to administer chimeric antigen receptor (CAR) T-cells?
Yes ☐ No ☐
- Is the health care facility trained in the management of cytokine release syndrome (CRS) and neurologic toxicities? Yes ☐ No ☐
- Will the health care facility comply with the Breyanzi® risk evaluation and mitigation strategy (REMS) program requirements? Yes ☐ No ☐
- 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. A patient-specific, clinically significant reason why Kymriah® (tisagenlecleucel) or Yescarta® (axcabtagene) is not appropriate for the member: _____

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Fax completed prior authorization request form to 888-601-8461 or submit Electronic Prior Authorization through CoverMyMeds® or SureScripts. All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at AetnaBetterHealth.com/Oklahoma.

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Member Name: _____ Date of Birth: _____ Member ID#: _____

Criteria

For Authorization (continued):

5. Please indicate the diagnosis and information:

☐ **Chronic Lymphocytic Leukemia (CLL)/Small Lymphocytic Lymphoma (SLL)**

- A. Relapsed or refractory disease after 2 or more lines of systemic therapy? Yes ☐ No ☐
- i. Did therapy include a Burton tyrosine kinase (BTK) inhibitor and a B cell lymphoma-2 (BCL-2) inhibitor? Yes ☐ No ☐
- B. Does member have primary central nervous system (CNS) lymphoma? Yes ☐ No ☐

☐ **Follicular Lymphoma**

- A. Relapsed or refractory disease after 2 or more lines of systemic therapy? Yes ☐ No ☐
- B. Does member have primary central nervous system (CNS) lymphoma? Yes ☐ No ☐
- C. A patient-specific, clinically significant reason why Kymriah® (tisagenlecleucel) or Yescarta® (axcabtagene) is not appropriate for the member: _____
- _____
- _____

☐ **Mantle Cell Lymphoma (MCL)**

- A. Relapsed or refractory disease after 2 or more lines of systemic therapy? Yes ☐ No ☐
- i. Did therapy include a Burton tyrosine kinase (BTK) inhibitor? Yes ☐ No ☐
- B. Does member have primary central nervous system (CNS) lymphoma? Yes ☐ No ☐
- C. A patient-specific, clinically significant reason why Tecartus® (brexucabtagene autoleucel) is not appropriate for the member: _____
- _____
- _____

☐ **Other:** _____

Additional Information: _____

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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.

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