



MASSACHUSETTS

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CAR T-Cell Therapy Services for Non-Hodgkin Lymphoma (Lisocabtagene Maraleucel) Prior Authorization Request Form #941

Medical Policy #066 Chimeric Antigen Receptor Therapy for Leukemia and Lymphoma

CLINICAL DOCUMENTATION

- Clinical documentation that supports the medical necessity criteria for CAR T-Cell Therapy Services for Non-Hodgkin Lymphoma (Lisocabtagene Maraleucel) must be submitted.
- If the patient does not meet all the criteria listed below, please submit a letter of medical necessity with a request for [Clinical Exception \(Individual Consideration\)](#) explaining why an exception is justified.

Requesting Prior Authorization Using Authorization Manager

Providers will need to use [Authorization Manager](#) to submit initial authorization requests for services. Authorization Manager, available 24/7, is the quickest way to review authorization requirements, request authorizations, submit clinical documentation, check existing case status, and view/print the decision letter. For commercial members, the requests must meet medical policy guidelines.

To ensure the request is processed accurately and quickly:

- Enter the facility's NPI or provider ID for where services are being performed.
- Enter the appropriate surgeon's NPI or provider ID as the servicing provider, *not* the billing group.

Authorization Manager Resources

- Refer to our [Authorization Manager](#) page for tips, guides, and video demonstrations.

Complete Prior Authorization Request Form for CAR T-Cell Therapy Services for Non-Hodgkin Lymphoma (Lisocabtagene Maraleucel) ([941](#)) using [Authorization Manager](#)

For out of network providers: Requests should still be faxed to 888-973-0726.

Patient Information	
Patient Name:	Today's Date:
BCBSMA ID#:	Date of Treatment:
Date of Birth:	Place of Service: Outpatient <input type="checkbox"/> Inpatient <input type="checkbox"/>

Physician Information	Facility Information
Name:	Name:
Address:	Address:
Phone #:	Phone #:
Fax#:	Fax#:
NPI#:	NPI#:

Please check off if the patient is enrolled in a Clinical Trial.

Clinical Trial #	<input type="checkbox"/>
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Please check off if the patient has the following diagnosis and HAS RELAPSED^c or is REFRACTORY^c :	
Histologically confirmed diagnosis of: Diffuse large B-cell lymphoma not otherwise specified (including diffuse large B-cell lymphoma arising from indolent lymphoma)	<input type="checkbox"/>
High-grade B-cell lymphoma, OR	<input type="checkbox"/>
Primary mediastinal large B-cell lymphoma, OR	<input type="checkbox"/>
Follicular lymphoma grade 3B.	<input type="checkbox"/>

^c Relapsed or refractory disease is defined as progression after 2 or more lines of systemic therapy (which may or may not include therapy supported by autologous cell transplant).

Please check off that the patient meets ALL the following criteria:	
1. Histologically confirmed diagnosis of large B-cell lymphoma, including diffuse large B-cell lymphoma not otherwise specified (including diffuse large B-cell lymphoma arising from indolent lymphoma), high-grade B-cell lymphoma, primary mediastinal large B-cell lymphoma, and follicular lymphoma grade 3B; AND	<input type="checkbox"/>
2. Meets at least one of the following: a. Primary refractory or relapsed disease ^c within 12 months of first-line chemo-immunotherapy that included an anti-CD20 monoclonal antibody and anthracycline-containing regimen; OR b. Primary refractory or relapsed disease within 12 months of first-line chemo-immunotherapy that included an anti-CD20 monoclonal antibody and anthracycline-containing regimen and are not eligible for hematopoietic stem cell transplantation due to comorbidities or age; OR c. Relapsed or refractory disease as defined as progression after ≥2 lines of systemic therapy including anti-CD20 monoclonal antibody for CD20-positive tumor and anthracycline-containing chemotherapy i. When the individual has histological transformation of follicular lymphoma or marginal zone lymphoma to diffuse large B-cell lymphoma: prior chemotherapy for follicular lymphoma or marginal zone lymphoma and ≥2 chemo-immunotherapy regimens for the transformed disease; AND	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
3. At least 18 years of age at the time of infusion; AND	<input type="checkbox"/>
4. Have adequate organ and bone marrow function as determined by the treating oncologist/hematologist; AND	<input type="checkbox"/>
5. Have not received prior CD19-directed chimeric antigen receptor T-cell therapy treatment, any other cell therapy, or any gene therapy or are being considered for treatment with any other cell therapy or any gene therapy; AND	<input type="checkbox"/>
6. Do not have primary central nervous system lymphoma.	<input type="checkbox"/>

^c Relapsed or refractory disease is defined as progression after 2 or more lines of systemic therapy (which may or may not include therapy supported by autologous cell transplant).

CPT CODES/ HCPCS CODES/ ICD CODES

HCPCS codes:	Code Description	
C9076	Lisocabtagene maraleucel, up to 110 million autologous anti-cd19 car-positive viable t cells, including leukapheresis and dose preparation procedures, per therapeutic dose	<input type="checkbox"/>
C9399	Unclassified drugs or biologicals	<input type="checkbox"/>
J3490	Unclassified drugs	<input type="checkbox"/>
J3590	Unclassified biologics	<input type="checkbox"/>

J9999	Not otherwise classified, antineoplastic drugs	<input type="checkbox"/>
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Providers should enter the relevant diagnosis code(s) below:

Code	Description	
		<input type="checkbox"/>
		<input type="checkbox"/>

Providers should enter other relevant code(s) below:

Code	Description	
		<input type="checkbox"/>
		<input type="checkbox"/>