

Beleodaq

CareFirst Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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| Patient's Name: | Date: |
|--|---|
| Patient's ID: | Patient's Date of Birth: |
| Physician's Name: | |
| Specialty: | NPI#: |
| Physician Office Telephone: | Physician Office Fax: |
| Referring Provider Info: ☐ Same as Re | uesting Provider |
| Name: | NPI#: |
| Fax: | Phone: |
| Rendering Provider Info: 🗆 Same as Re | erring Provider 🗆 Same as Requesting Provider |
| Name: | |
| | |
| Fax: | Phone: |
| Fax: Approvals may be subject | Phone: o dosing limits in accordance with FDA-approved labeling, ndia, and/or evidence-based practice guidelines. |
| Fax: Approvals may be subject accepted comp | Phone: o dosing limits in accordance with FDA-approved labeling, ndia, and/or evidence-based practice guidelines. |
| Fax: | Phone: o dosing limits in accordance with FDA-approved labeling, ndia, and/or evidence-based practice guidelineskg |
| Fax: Approvals may be subject accepted comp Required Demographic Information: Patient Weight: | Phone: o dosing limits in accordance with FDA-approved labeling, ndia, and/or evidence-based practice guidelines. kgcm |
| Fax: | Phone: o dosing limits in accordance with FDA-approved labeling, ndia, and/or evidence-based practice guidelines. kgcm |

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. CareFirst Beleodaq SGM 1701-A - 07/2025.

Criteria Questions: 1. What is the diagnosis? ☐ Peripheral T-cell lymphoma (PTCL) [including the following subtypes: anaplastic large cell lymphoma, peripheral T-cell lymphoma not otherwise specified, angioimmunoblastic T-cell lymphoma, enteropathy associated T-cell lymphoma, monomorphic epitheliotropic intestinal T-cell lymphoma, nodal peripheral T-cell lymphoma with TFH phenotype, or follicular T-cell lymphoma], Continue to 2 ☐ Adult T-cell leukemia/lymphoma (ATLL), Continue to 2 ☐ Extranodal NK/T-cell lymphoma, Continue to 2 ☐ Hepatosplenic T-cell lymphoma, Continue to 2 ☐ Breast implant associated anaplastic large cell lymphoma (ALCL), Continue to 2 ☐ Other, please specify. ________, Continue to 2 2. Is this a request for continuation of therapy with the requested medication? ☐ Yes, Continue to 3 ☐ No, Continue to 4 3. Is there evidence of unacceptable toxicity or disease progression while on the current regimen? ☐ Yes, No Further Questions ☐ No, *No Further Questions* 4. What is the diagnosis? ☐ Peripheral T-cell lymphoma (PTCL) [including the following subtypes: anaplastic large cell lymphoma, peripheral T-cell lymphoma not otherwise specified, angioimmunoblastic T-cell lymphoma, enteropathy associated T-cell lymphoma, monomorphic epitheliotropic intestinal T-cell lymphoma, nodal peripheral T-cell lymphoma with TFH phenotype, or follicular T-cell lymphoma], Continue to 5 ☐ Adult T-cell leukemia/lymphoma (ATLL), Continue to 8 ☐ Extranodal NK/T-cell lymphoma, Continue to 10 ☐ Hepatosplenic T-cell lymphoma, Continue to 14 ☐ Breast implant-associated anaplastic large cell lymphoma (ALCL), Continue to 16 5. Will the requested medication be used as a single agent? ☐ Yes, Continue to 6 □ No, Continue to 6 6. Is the disease relapsed or refractory? ☐ Yes, No Further Questions □ No, Continue to 7 7. Is the requested medication being used for palliative intent? ☐ Yes, No Further Questions ☐ No, *No Further Questions* 8. Will the requested medication be used as a single agent?

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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☐ Yes, Continue to 9☐ No, Continue to 9☐

| Prescriber or Authorized Signature | Date (mm/dd/yy) |
|--|---------------------------------------|
| X | |
| I attest that this information is accurate and true, and the information is available for review if requested by CVS | |
| ☐ Subsequent treatment, No Further Questions | |
| 17. What is the place in therapy in which the requested medic First-line treatment, <i>No Further Questions</i> | ation will be used? |
| 16. Will the requested medication be used as a single agent? ☐ Yes, Continue to 17 ☐ No, Continue to 17 | |
| 15. How many previous lines of chemotherapy has the patient lines, <i>No Further Questions</i> | t received? |
| 14. Will the requested medication be used as a single agent? ☐ Yes, Continue to 15 ☐ No, Continue to 15 | |
| 13. Does the patient have a contraindication to asparaginase-by Tes, <i>No Further Questions</i> ☐ No, <i>No Further Questions</i> | pased therapy (e.g., pegaspargase)? |
| 12. Has the patient had an inadequate response to asparaginas ☐ Yes, <i>No Further Questions</i> ☐ No, <i>Continue to 13</i> | e-based therapy (e.g., pegaspargase)? |
| 11. Is the disease relapsed or refractory? ☐ Yes, Continue to 12 ☐ No, Continue to 12 | |
| 10. Will the requested medication be used as a single agent? ☐ Yes, Continue to 11 ☐ No, Continue to 11 | |
| 9. What is the place in therapy in which the requested medical ☐ First-line treatment, <i>No Further Questions</i> ☐ Subsequent treatment, <i>No Further Questions</i> | tion will be used? |

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