

Asparlas[®] (calaspargase pegol-mknl) (Intravenous)



Last Review Date: 12/01/2020 Date of Origin: 06/02/2020 Dates Reviewed: 06/2020, 12/2020

I. Length of Authorization

Coverage will be provided for 6 months and may be renewed.

II. Dosing Limits

A. Quantity Limit (max daily dose) [NDC Unit]:

• Asparlas 3,750 units per 5 mL single-use vial: 2 vials every 21 days

B. Max Units (per dose and over time) [HCPCS Unit]:

• 750 billable units (2 vials) per 21 days

III. Initial Approval Criteria¹⁻³

Coverage is provided in the following conditions:

• Patient age is 1 month up to 21 years old; AND

Universal Criteria

- Patient must not have a history of serious hypersensitivity, pancreatitis, severe hepatic impairment, thrombosis, or hemorrhagic events with prior L-asparaginase* therapy; **AND**
- Must be used as a component of multi-agent chemotherapy; AND

Acute Lymphoblastic Leukemia (ALL) † Ф

Preferred therapies and recommendations are determined by review of clinical evidence. NCCN category of recommendation is taken into account as a component of this review. Regimens deemed equally efficacious (i.e., those having the same NCCN categorization) are considered to be therapeutically equivalent.



† FDA Approved Indication(s); **‡** Compendia recommended indication(s); **Φ** Orphan Drug

IV. Renewal Criteria¹⁻³

Coverage can be renewed based upon the following criteria:

- Patient continues to meet universal and indication-specific criteria as identified in section III; AND
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include allergic reactions (including anaphylaxis), thrombosis (including cerebral thrombosis, ischemia, and stroke), coagulopathy/bleeding (including intracranial hemorrhage), severe hepatotoxicity, pancreatitis, etc.; AND

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• Disease stabilization or improvement as evidenced by a complete response [CR] (i.e. morphologic, cytogenetic or molecular complete response CR), complete hematologic response or a partial response by CBC, bone marrow cytogenic analysis, QPCR, or FISH

V. Dosage/Administration

Indication	Dose
All indications	Administer 2,500 units/m ² intravenously given no more frequently than every 21 days.*

VI. Billing Code/Availability Information

HCPCS Code:

• J9118 – Injection, calaspargase pegol-mknl, 10 units: 1 billable unit = 10 units

<u>NDC(s)</u>:

• Asparlas 3,750 units/5 mL single-use vial: 72694-0515-XX

VII. References (STANDARD)

- 1. Asparlas [package insert]. Boston, MA; Servier Pharmaceuticals Inc.; June 2020. Accessed October 2020.
- 2. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium®) for calaspargase pegol-mknl. National Comprehensive Cancer Network, 2020. The NCCN Compendium® is a derivative work of the NCCN Guidelines®. NATIONAL COMPREHENSIVE CANCER NETWORK®, NCCN®, and NCCN GUIDELINES® are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed by Magellan Rx October 2020.
- Silverman LB, Blonquist TM, Hunt SK, et al. Randomized Study of Pegasparagase (SS-PEG) and Calaspargase Pegol (SC-PEG) in Pediatric Patients with Newly Diagnosed Acute Lymphoblastic Leukemia or Lymphoblastic Lymphoma: Results of DFCI ALL Consortium Protocol 11-001. Blood 2016;128:175.



VIII. References (ENHANCED)

- 1e. Referenced with permission from the NCCN Drugs & Biologics Compendium (NCCN Compendium[®]) for Acute Lymphoblastic Leukemia, Version 1.2020. National Comprehensive Cancer Network, 2020. The NCCN Compendium[®] is a derivative work of the NCCN Guidelines[®]. NATIONAL COMPREHENSIVE CANCER NETWORK[®], NCCN[®], and NCCN GUIDELINES[®] are trademarks owned by the National Comprehensive Cancer Network, Inc. To view the most recent and complete version of the Compendium, go online to NCCN.org. Accessed by Magellan Rx October 2020.
- 2e. Angiolillo AL, Schore RJ, Devidas M, et al. Pharmacokinetic and pharmacodynamic properties of calaspargase pegol Escherichia coli L-asparaginase in the treatment of patients with acute lymphoblastic leukemia: results from Children's Oncology Group Study AALL07P4. J Clin Oncol. 2014;32(34):3874–3882. doi:10.1200/JCO.2014.55.5763.
- 3e. Magellan Health, Magellan Rx Management. Asparlas Clinical Literature Review Analysis. Last updated October 2020. Accessed October 2020.

ICD-10	ICD-10 Description			
C83.50	Lymphoblastic (diffuse) lymphoma, unspecified site			
C83.51	Lymphoblastic (diffuse) lymphoma, lymph nodes of head, face, and neck			
C83.52	Lymphoblastic (diffuse) lymphoma, intrathoracic lymph nodes			
C83.53	Lymphoblastic (diffuse) lymphoma, intra-abdominal lymph nodes			
C83.54	Lymphoblastic (diffuse) lymphoma, lymph nodes of axilla and upper limb			
C83.55	Lymphoblastic (diffuse) lymphoma, lymph nodes of inguinal region and lower limb			
C83.56	Lymphoblastic (diffuse) lymphoma, intrapelvic lymph nodes			
C85.57	Lymphoblastic (diffuse) lymphoma, spleen			
C83.58	Lymphoblastic (diffuse) lymphoma, lymph nodes of multiple sites			
C83.59	Lymphoblastic (diffuse) lymphoma, extranodal and solid organ sites			
C91.00	Acute lymphoblastic leukemia not having achieved remission			
C91.01	Acute lymphoblastic leukemia, in remission			
C91.02	Acute lymphoblastic leukemia, in relapse			

Appendix 1 – Covered Diagnosis Codes

Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD), Local Coverage Determinations (LCDs), and Local Coverage Articles (LCAs) may exist and compliance with these policies is required where applicable. They can be found at: <u>http://www.cms.gov/medicare-coverage-</u>



<u>database/search/advanced-search.aspx</u>. Additional indications may be covered at the discretion of the health plan.

	Medicare Part B Administrative Contractor (MAC) Jurisdictions							
Jurisdiction	Applicable State/US Territory	Contractor						
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC						
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC						
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)						
6	MN, WI, IL	National Government Services, Inc. (NGS)						
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.						
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)						
N (9)	FL, PR, VI	First Coast Service Options, Inc.						
J (10)	TN, GA, AL	Palmetto GBA, LLC						
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA, LLC						
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.						
К (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)						
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Appendix 3 – CLINICAL LITERATURE REVIEW

OS = overall survival; PFS = progression-free survival; ORR = objective response rate; CR = complete response; PR = partial response; DoR = duration of response; TTP = time to progression; FFS = failure-free survival; EFS = event-free survival; PFR = progression free rate

Acute lymphoblastic leukemia (ALL)

Regimen	NCCN Category	FDA Approved	Trial Design	Comparator	Primary End-Point	Line of Therapy	Conclusion
Calaspargase pegol-mknl + multi-agent chemotherapy	2A	Yes	<u>Phase 2,</u> randomized	Pegaspargase + multi-agent chemotherapy	Toxicity Pharmacokinetics	Induction therapy (newly diagnosed)	• During remission induction, calaspargase pegol led to a more sustained serum asparaginase activity without excess toxicity or significant difference in the proportion of patients with low end-induction minimal residual disease compared to pegaspargase.
Calaspargase pegol-mknl + multi-agent chemotherapy	2A	Yes	AALL07P4, randomized, open-label, multi-center	Pegaspargase + multi-agent chemotherapy		Induction therapy (newly diagnosed)	 Calaspargase pegol given as 2,500 IU/m² achieves a significantly longer period of asparaginase activity above defined thresholds and asparagine depletion compared with pegaspargase.