

Lenmeldy™ (atidarsagene autotemcel) Prior Authorization Form

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

Drug Information

Physician billing (HCPCS code: _____) Start Date: _____

Dose: _____ Regimen: _____

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 lifetime)

1. Please indicate the diagnosis and information:

- Metachromatic Leukodystrophy (MLD)
- Other _____

2. How was the diagnosis confirmed? (select one)

- Arylsulfatase A (ARSA) enzyme activity below the normal range in peripheral blood mononuclear cells or fibroblasts **(Please submit results of assay)**
- Molecular genetic testing confirming biallelic pathogenic variants in the ARSA gene of known polymorphisms **(Please submit results of genetic testing)**
 - a. Were novel ARSA variants identified? Yes No
 - i. If yes, did a 24-hour urine collection demonstrate increased urinary excretion of sulfatides? Yes No **(Please submit results)**

3. Does the member have 1 of the following forms of MLD? **(Please submit clinical documentation)**

- Pre-symptomatic late infantile (PSLI) MLD with expected disease onset ≤30 months of age
- Pre-symptomatic early juvenile (PSEJ) MLD with expected disease onset >30 months and <7 years of age
- Early symptomatic early juvenile (ESEJ) MLD with disease onset >30 months and <7 years of age

4. Is Lenmeldy™ prescribed by a geneticist, hematologist/oncologist, neurologist, or other specialist with expertise in the treatment of MLD and the administration of Lenmeldy™? Yes No

5. Does member have a history of prior hematopoietic stem cell transplantation (HSCT)? Yes No

a. If yes, is there evidence of residual cells of donor origin? Yes No

6. Is member clinically stable and eligible to undergo HSCT? Yes No

<p>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. Pharmacy Coverage Guidelines are available at AetnaBetterHealth.com/Oklahoma.</p>	<p>CONFIDENTIALITY NOTICE</p> <p><i>This document, including any attachments, contains information which is confidential or privileged. If you are not the intended recipient, be aware that any disclosure, copying, distribution, or use of the contents of this information is prohibited. If you have received this document in error, please notify the sender immediately by telephone to arrange for the return of the transmitted documents or to verify their destruction.</i></p>
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Criteria**For Authorization (continued):**

7. Does the member have a negative serology test for human immunodeficiency virus 1 & 2 (HIV-1/HIV-2), hepatitis B virus (HBV), hepatitis C virus (HCV), human T-lymphotropic virus 1 & 2 (HTLV-1/HTLV-2), cytomegalovirus (CMV), and mycoplasma prior to apheresis? Yes No
8. For female members of reproductive potential, please answer the following:
- a. Is the member pregnant? Yes No
 - b. Does the member have a negative pregnancy test prior to the start of mobilization, prior to conditioning procedures, and prior to Lenmeldy™ administration? Yes No
9. For all members of reproductive potential, please answer the following:
- a. Does member agree to use an effective method of contraception from the start of mobilization through at least 6 months after administration of Lenmeldy™? Yes No
 - b. Has member been counseled on the potential effects of myeloblastic conditioning on fertility? Yes No
 - c. Is the potential risk of infertility acceptable to the member or member's caregiver? Yes No
10. Has the member been evaluated for and counseled on all warnings and precautions related to Lenmeldy™, including the risk of thrombosis and thromboembolic events, serious infections, and veno-occlusive disease? Yes No
11. Will member be monitored for hematologic malignancies lifelong, with a complete blood count (with differential) performed annually and integration site analysis as warranted for at least 15 years after treatment with Lenmeldy™? Yes No
12. Will Lenmeldy™ be administered at a Lenmeldy™ qualified treatment center? Yes No
- a. Name of facility: _____
 - b. Does the receiving facility have a mechanism in place to track the patient-specific Lenmeldy™ dose from receipt to storage to administration? Yes No

Additional Information: _____

(Page 2 of 2)

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. Failure to complete all pages will result in processing delays.

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