

Medical Drug Clinical Criteria

Subject: Aphexda (motixafortide)

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Overview

This document addresses the use of Aphexda (motixafortide), a chemokine receptor type 4 inhibitor which impairs binding of hematopoietic stem cells within the bone marrow microenvironment. Aphexda is approved in combination with filgrastim to mobilize hematopoietic stem cells to the peripheral blood for subsequent autologous transplantation in individuals with multiple myeloma.

The National Comprehensive Cancer Network (NCCN) guideline on Hematopoietic Cell Transplantation recommends the use of Aphexda in combination with filgrastim (or biosimilar) or tbo-filgrastim as a hematopoietic cell mobilization regimen for autologous donors undergoing transplantation for multiple myeloma.

Clinical Criteria

When a drug is being reviewed for coverage under a member's medical benefit plan or is otherwise subject to clinical review (including prior authorization), the following criteria will be used to determine whether the drug meets any applicable medical necessity requirements for the intended/prescribed purpose.

Aphexda (motixafortide)

Requests for Aphexda (motixafortide) may be approved if the following criteria are met:

- I. Individual is 18 years or older; **AND**
- II. Individual has a diagnosis of multiple myeloma; **AND**
- III. Aphexda is being used to mobilize autologous hematopoietic stem cells; **AND**
- IV. Individual is using in combination with filgrastim, filgrastim biosimilar, or tbo-filgrastim (NCCN 2A); **AND**
- V. After stem cell mobilization and collection, a subsequent autologous hematopoietic stem cell transplant is anticipated; **AND**
- VI. The total number of Aphexda injections does not exceed two doses total; one dose prior to first and third apheresis.

Requests for Aphexda (motixafortide) may not be approved for the following:

- I. More than one treatment cycle*; **OR**
- II. Individual is using as a mobilizing agent for an allogeneic stem cell donor (NCCN); **OR**
- III. Individual is using as a mobilizer of leukemic cells; **OR**
- IV. When the above criteria are not met or for all other indications.

Approval duration: *One treatment cycle (includes two doses total; one dose prior to first and third apheresis)

Coding

The following codes for treatments and procedures applicable to this document are included below for informational purposes. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member

HCPCS

Injection, motixafortide, 0.25 mg [Aphexda]

C90.00-C90.02

Multiple Myeloma

Z52.011

Autologous donor, stem cells

Z94.84

Stem cells transplant status

Reviewed: 02/21/2025

- 02/21/2025 – Annual Review: No Changes. Coding Reviewed: Added ICD-10-CM Z52.011 and Z94.84.
- 02/23/2024 - Annual Review: No changes. Coding Reviewed: No Changes. Effective 4/1/2024 Added HCPCS J2277. Added ICD-10-CM C90.00-C90.02. Removed J3490, J3590, J9999.
- 11/17/2023 - Select Review: Create new clinical criteria for Aphexda. Coding Reviewed: Added HCPCS J3490, J3590, J9999. All diagnoses pend.

1. DailyMed. Package inserts. U.S. National Library of Medicine, National Institutes of Health website. <http://dailymed.nlm.nih.gov/dailymed/about.cfm>. Accessed: January 23, 2024.
2. DrugPoints® System [electronic version]. Truven Health Analytics, Greenwood Village, CO. Updated periodically.
3. Lexi-Comp ONLINE™ with AHFS™, Hudson, Ohio: Lexi-Comp, Inc. Updated periodically.
4. Crees ZD, Rettig MP, Jayasinghe RG, et al. Motixafortide and G-CSF to mobilize hematopoietic stem cells for autologous transplantation in multiple myeloma: a randomized phase 3 trial. *Nat Med.* 2023;29(4):869-879.
5. NCCN Clinical Practice Guidelines in Oncology™. © 2024 National Comprehensive Cancer Network, Inc. For additional information visit the NCCN website: <http://www.nccn.org/index.asp>. Accessed on January 5, 2024.
 - a. Hematopoietic Cell Transplantation (HCT). V3.2023. Revised October 9, 2023.

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