

**Aphexda (motixafortide)**  
**Effective 07/01/2025**

<b>Plan</b>	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	<b>Program Type</b>	<input checked="" type="checkbox"/> Prior Authorization
<b>Benefit</b>	<input type="checkbox"/> Pharmacy Benefit <input checked="" type="checkbox"/> Medical Benefit		<input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
<b>Specialty Limitations</b>	N/A		
<b>Contact Information</b>	<b>Medical and Specialty Medications</b>		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
	<b>Non-Specialty Medications</b>		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
<b>Exceptions</b>	N/A		

#### Overview

Aphexda (motixafortide) is a hematopoietic stem cell mobilizer that is indicated in combination with filgrastim (G-CSF) to mobilize hematopoietic stem cells to the peripheral blood for collection and subsequent autologous transplantation in patients with multiple myeloma.

Aphexda is initiated after filgrastim has been administered daily for four days. Aphexda should be administered 10-14 hours before initiating apheresis. A second dose of Aphexda can be administered 10-14 hours before a third apheresis.

#### Coverage Guidelines

Authorization may be granted for members new to the plan within the past 90 days who are currently receiving treatment with the requested medication, excluding when the product is obtained as samples or via manufacturer's patient assistance program

#### OR

Authorization may be granted if the member meets all the following diagnosis-specific criteria:

1. Documented diagnosis of multiple myeloma
2. Member is 18 years of age or older
3. Documentation Aphexda is being used to mobilize hematopoietic stem cells for autologous transplantation
4. Documentation member has been approved by the Plan for autologous hematopoietic stem cell transplant
5. Aphexda will be used in conjunction with a granulocyte colony stimulating factor (G-CSF)
6. Member meets ONE of the following:
  - a. Documentation member has had in inadequate response, adverse reaction, or contraindication to plerixafor
  - b. Prescriber submits documentation of a clinical rationale why plerixafor is not clinically appropriate for the member

### **Continuation of Therapy**

1. Requests for reauthorization must meet initial criteria.

### **Limitations**

1. Initial and reauthorization approvals will be granted for three months.

### **References**

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13. National Comprehensive Cancer Network (NCCN). NCCN Clinical Practice Guidelines in Oncology – Multiple Myeloma v2.2024. November 1, 2023(a). NCCN Web site. [https://www.nccn.org/professionals/physician\\_gls/pdf/myeloma.pdf](https://www.nccn.org/professionals/physician_gls/pdf/myeloma.pdf). Accessed November 1, 2023.



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### Review History

09/11/2024 – Reviewed at September P&T. Effective 11/01/2024.

04/09/2024 – Reviewed and Updated at April P&T. Updated criteria to clarify documentation requirements.  
Effective 07/01/2025.

