

ANTIHEMOPHILIA AGENTS PRIOR AUTHORIZATION FORM *(form effective 1/9/2023)*

Prior authorization guidelines for **Antihemophilia Agents** are available on the DHS Pharmacy Services website at:

<https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>.

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	Total # of pgs: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	State license #:
LTC facility contact/phone:			Street address:	
Beneficiary name:			City/State/Zip:	
Beneficiary ID#:		DOB:	Phone:	Fax:

CLINICAL INFORMATION.

Drug #1 requested:	Strength & package size:		
Directions:	Quantity:	Refills:	
Drug #2 requested:	Strength & package size:		
Directions:	Quantity:	Duration:	
Diagnosis <i>(submit documentation)</i> :			Dx code <i>(required)</i> :
Is the medication prescribed by a hematologist or hemophilia treatment center practitioner? <input type="checkbox"/> Yes <input type="checkbox"/> No			

Complete the section(s) below applicable to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

INITIAL REQUESTS

- Request is for HEMLIBRA (emicizumab):**
 - Has a diagnosis of severe hemophilia A
 - Has a diagnosis of severe hemophilia A with inhibitors
 - Has a diagnosis of hemophilia A and a history of at least 1 spontaneous joint bleed or other serious bleeding event
- Request is for a BYPASSING AGENT (eg, FEIBA NF, NovoSeven, Sevenfact) *(check all that apply)*:**
 - Has hemophilia A with inhibitors:
 - Is using the requested medication for episodic/on-demand treatment OR intermittent/periodic prophylaxis
 - Is using the requested medication for routine prophylaxis
 - Failed to achieve clinical goals with Hemlibra

- Has a medical reason why Hemlibra cannot be used
- Has been using the requested bypassing agent for routine prophylaxis within the past 90 days
- Has hemophilia B with inhibitors
- Has acquired hemophilia
- Has congenital factor VII deficiency
- Has Glanzmann's thrombasthenia

3. Request is for a non-preferred FACTOR VIII, FACTOR IX, or VWF:

- Both of the following:
 - Has been using the requested product within the past 90 days
 - Has a medical reason to continue using the requested medication
- Failed to achieve clinical goals with or has a contraindication or an intolerance to the preferred FVIII, FIX, or FVIII/VWF products with the same half-life (standard v. extended half-life), if applicable. Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.
- Has a diagnosis for which no preferred Antihemophilia Agents are appropriate. Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.

RENEWAL REQUESTS

- Experienced a positive clinical response since starting the requested medication

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION

Prescriber Signature:	Date:
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