

ANTIHEMOPHILIA AGENTS PRIOR AUTHORIZATION FORM *(form effective 1/8/2024)*

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

1. Request is for HEMLIBRA (emicizumab):

- Has a diagnosis of severe congenital hemophilia A
- Has a diagnosis of congenital hemophilia A with inhibitors
- Has a diagnosis of congenital hemophilia A and a history of at least 1 spontaneous joint bleed or other serious bleeding event

2. Request is for a BYPASSING AGENT (eg, FEIBA NF, NovoSeven, Sevenfact):

- Has hemophilia A with inhibitors AND:
 - Is using the requested medication for episodic/on-demand treatment OR intermittent/periodic prophylaxis
 - Is using the requested medication for routine prophylaxis AND:

Failed to achieve clinical goals with Hemlibra (emicizumab)
 Has a medical reason why Hemlibra (emicizumab) 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 medication 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 medications 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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