

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

Prior authorization guidelines for **Antihemophilia Agents** are available on Geisinger Health Plan's website at  
<https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<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 (submit documentation):		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 submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.**

**Prescriber Signature:**

**Date:**

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