

# Pharmacy Prior Authorization Form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206

This form applies to:  Commercial (Traditional)  Commercial (Individual/Optimized)  
 Medicaid

This request is:  Urgent (life threatening)  Non-Urgent (standard review)

Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function.

## Hemlibra<sup>®</sup> (emicizumab-kxwh)

### Member

Last Name: \_\_\_\_\_ First Name: \_\_\_\_\_  
 ID #: \_\_\_\_\_ DOB: \_\_\_\_\_ Gender: \_\_\_\_\_  
 Primary Care Physician: \_\_\_\_\_  
 Requesting Provider: \_\_\_\_\_ Prov. Phone: \_\_\_\_\_ Prov. Fax: \_\_\_\_\_  
 Provider Address: \_\_\_\_\_  
 Provider NPI: \_\_\_\_\_ Contact Name: \_\_\_\_\_  
 Provider Signature: \_\_\_\_\_ Date: \_\_\_\_\_

### Product Information

New request  Continuation request

Drug product:  Hemlibra 30 mg/mL **Start date** (or date of next dose): \_\_\_\_\_  
 Hemlibra 60 mg / 0.4 mL **Date of last dose** (if applicable): \_\_\_\_\_  
 Hemlibra 105 mg / 0.7 mL **Dosing frequency:** \_\_\_\_\_  
 Hemlibra 150 mg/mL

### Drug cost information

The wholesale acquisition cost for one dose of Hemlibra is \$373,000.

### Precertification Requirements

**Before this drug is covered, documentation must be submitted to support that the patient meets all of the following requirements:**

1. Diagnosis of Hemophilia A with factor VIII inhibitors (documentation of the presence of inhibitors must be submitted to Priority Health)

**Note:** Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication.

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**Priority Health Precertification Documentation**

**A. What condition is this drug being requested for?**

Hemophilia A with factor VIII inhibitors (documentation of inhibitors must be submitted to Priority Health)

Other – the patient’s condition is: \_\_\_\_\_

Rationale for use: \_\_\_\_\_

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**Additional information**

**Note:** Coverage of Hemlibra is limited to the FDA approved dosing of 3 mg/kg for the first 4 weeks of therapy, then 1.5 mg/kg weekly thereafter. Additionally, when approved, Hemlibra must be obtained from a participating Hemophilia Specialty Pharmacy as noted in Medical Policy 91569.