

Fax completed form to: (855) 840-1678 If this is an URGENT request, please call (800) 882-4462

(800.88.CIGNA)

Hemlibra (emicizumab-kxwh)

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on		
Specialty:	Specialty: * DEA, NPI or TIN:		this form are completed.*		
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:	* Date of Birth:	
Office Fax:			* Patient Street Address:		
Office Street Address:			City: St	tate:	Zip:
City:	State:	Zip:	Patient Phone:	Patient Phone:	
Urgency: Standard Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)					
Medication Requested: Hemlibra 30mg/ml vial Hemlibra 60 mg/0.4ml vial Hemlibra 105mg/0.7ml vial Hemlibra 150mg/1ml vial					
ICD10: Dose and Quantity: Duration of therapy: Frequency of administration: What is your patient's current weight? Is this a new start or continuation of therapy? If your patient has already begun treatment with drug samples of Hemlibra, please choose "new start of therapy". In new start of therapy (if continued therapy) Has your patient had a good response to therapy with this drug (such as reduction in frequency of bleeding episodes)?					
Where will this medication be obtained? Accredo Specialty Pharmacy** Prescriber's office stock (billing on a medical claim form) Characterization of the control of the cont					red specialty pharmacy
Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient?					
What is your patient's diag		ciency)	other (please specify)):	
Clinical Information **This drug requires supportive documentation (chart notes, lab and test results, etc). ** **Supportive documentation for all answers must be attached with this request** Is there documentation that your patient has one of the following? factor XIII inhibitors mild or moderate hemophilia (defined as factor VIII level of 1% to less than 40%) severe hemophilia defined as pre-treatment factor VIII level less than 1% none of the above (if mild/moderate) Which of the following applies to your patient? Please provide documentation. 1 or more episodes of bleeding into the central nervous system or other serious, life-threatening bleed 1 or more episodes of bleeding into large joint (ankles, knees, hips, elbows, shoulders) and age 3 years or younger 2 or more episodes of bleeding into large joints (ankles, knees, hips, elbows, shoulders) presence of joint disease documented by physical examination and plain radiographs of the affected joints none of the above Is Hemlibra being used for routine prophylaxis to prevent or reduce the frequency of bleeding episodes? Yes No (if no) Please specify the use for which Hemlibra is being prescribed.					
(if no) Please specify the us	e for which Hem	ilibra is being prescri	lbed.		

Additional pertinent information (including prior therapy, disease stage, performance status, and names/doses/admin schedule of any agents to be used concurrently):
Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or
insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.
Prescriber Signature: Date: Date:
Save Time! Submit Online at: www.covermymeds.com/main/prior-authorization-forms/cigna/ or via SureScripts in your EHR.
Our standard response time for prescription drug coverage requests is 5 business days. If your request is urgent, it is important that you call us to expedite the request. View our Prescription Drug List and Coverage Policies online at cigna.com.

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