

Factor VIII Agents Afstyla, Kovaltry, Nuwiq

CareFirst Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name:		Date:
Patient's ID:	Patient's Date of Birth:	
Physician's Name:		
Specialty:		NPI#:
Physician Office Telephone:	Physician Office Fax:	
Referring Provider Info: 🗖 Same as Re	equesting Provi	der
Name:	NPI#:	
Fax:	Phone:	
Rendering Provider Info: 🗆 Same as Re	eferring Provide	er 🗆 Same as Requesting Provider
Name:	 -	NPI#:
Fax:	Phone:	
		in accordance with FDA-approved labeling, vidence-based practice guidelines.
Patient Weight:	kg	
Patient Height:	cm	
Please indicate the place of service for the	requested drug.	•
		☐ Off Campus Outpatient Hospital
☐ On Campus Outpatient Hospital	□ Office	☐ Pharmacy
Patient Height:Please indicate the place of service for the	cm requested drug.	
- On Campus Outputtent Hospital	= 0,,, ice	= 1 narmacy
What is the ICD-10 code?		

	ception Criteria Questions: What drug is being prescribed?						
	☐ Afstyla ☐ Nuwiq, Skip to Clinical Questions	☐ Kovaltry ☐ Other	, Skip to Clinical Questions				
В.	The preferred products for your patient's he Nuwiq. Can the patient's treatment be switted as Yes – Eloctate, <i>Please obtain Form for Please</i> . Hemlibra, <i>Please obtain Form for Please</i> . Nuwiq, <i>Skip to Clinical Criteria Question Form for Please</i> . Ayntha, <i>Please obtain Form for Please</i> . Ayntha Solofuse, <i>Please obtain Form In No, Continue to Question C</i> .	ched to one of the preferred profeered product and submit preferred product and submit preferred product and subm Questions referred product and submit	oroducts? It for corresponding PA. It for corresponding PA. It for corresponding PA.				
C.	Did the patient have a documented inadequate response, contraindication, or intolerable adverse event to all preferred products (Eloctate, Hemlibra, Xyntha (including Solofuse), and Nuwiq)? **Action Required: If Yes, attach supporting chart note(s). □ Yes □ No If Yes or No, Continue to Clinical Criteria Questions						
	nical Criteria Questions:						
	What is the diagnosis? Hemophilia A, Continue to 2						
	Other, please specify.	, No further questi	ons				
	Will the requested medication be prescribed Yes, <i>Continue to 3</i> No, <i>Continue to 3</i>	by or in consultation with a	hematologist?				
	Is the request for continuation of therapy? Yes, <i>Continue to 8</i> No, <i>Continue to 4</i>						
4.	What is the patient's baseline factor VIII ass	sav level (% activity)?					
	Less than 1% to 5% (moderate or severe dis						
	Greater than 5% (mild disease), Continue to	o 5					
	Has the patient had an insufficient response Yes, <i>No Further Questions</i> No, <i>Continue to 6</i>						
	Is there a clinical reason for not trying desm Yes, <i>Continue to 7</i> No, <i>Continue to 7</i>	opressin first?					
	What is the reason? Please indicate the clini Age less than 2 years, <i>No further questions</i>	cal reason for not trying desi	mopressin first.				
	Pregnancy, No further questions	-4:					
	Fluid/electrolyte imbalance, No further que	STIONS					

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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High risk for cardiovascular or cerebrovascular disease (especially elderly), No juriner questions
Predisposition to thrombus formation, <i>No further questions</i>
Trauma requiring surgery, No further questions
Life-threatening bleed, No further questions
Contraindication or intolerance to desmopressin, <i>No further questions</i>
Severe type 1 von Willebrand disease, No further questions
I Stimate Nasal Spray is unavailable due to backorder/shortage issues (where applicable), No further questions
Other, please specify, No further questions
. Is the patient experiencing benefit from therapy (e.g., reduced frequency or severity of bleeds)?
Yes, No Further Questions
No, No Further Questions

Step Therapy Override: Complete if Applicable for the state of Maryland.		Please Circle	
Is the requested drug being used to treat stage four advanced metastatic cancer?		No	
Is the requested drug's use consistent with the FDA-approved indication or the National Comprehensive Cancer Network Drugs & Biologics Compendium indication for the treatment of stage four advanced metastatic cancer and is supported by peer-reviewed medical literature?	Yes	No	
Is the requested drug being used for an FDA-approved indication OR an indication supported in the compendia of current literature (examples: AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?	Yes	No	
Does the prescribed quantity fall within the manufacturer's published dosing guidelines or within dosing guidelines found in the compendia of current literature (examples: package insert, AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?		No	
Do patient chart notes document the requested drug was ordered with a paid claim at the pharmacy, the pharmacy filled the prescription and delivered to the patient or other documentation that the requested drug was prescribed for the patient in the last 180 days?	Yes	No	
Has the prescriber provided proof documented in the patient chart notes that in their opinion the requested drug is effective for the patient's condition?	Yes	No	

Step Therapy Override: Complete if Applicable for the state of Virginia.		
Is the requested drug being used for an FDA-approved indication or an indication supported in the compendia of current literature (examples: AHFS, Micromedex, current accepted guidelines)?	Yes	No
Does the prescribed dose and quantity fall within the FDA-approved labeling or within dosing guidelines found in the compendia of current literature?	Yes	No
Is the request for a brand drug that has an AB-rated generic equivalent or interchangeable biological product available?	Yes	No
Has the patient had a trial and failure of the AB-rated generic equivalent or interchangeable biological product due to an adverse event (examples: rash, nausea, vomiting, anaphylaxis) that is thought to be due to an inactive ingredient?	Yes	No
Is the preferred drug contraindicated?		No
Is the preferred drug expected to be ineffective based on the known clinical characteristics of the patient and the prescription drug regimen?		No
Has the patient tried the preferred drug while on their current or previous health benefit plan and it was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event?	Yes	No
Is the patient currently receiving a positive therapeutic outcome with the requested drug for their medical condition?	Yes	No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X	
Prescriber or Authorized Signature	Date (mm/dd/yy)