

Qfitlia (fitusiran)
Prior Authorization Request Form
Caterpillar Prescription Drug Benefit
Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME: _____ **MEMBER'S FIRST NAME:** _____

Instructions: Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

☐ **URGENT**

MEMBER INFORMATION		
LAST NAME:		FIRST NAME:
PHONE NUMBER:		DATE OF BIRTH:
STREET ADDRESS:		
CITY:	STATE:	ZIP CODE:
PATIENT INSURANCE ID NUMBER:		

☐ MALE ☐ FEMALE HEIGHT (IN/CM): _____ WEIGHT (LB/KG): _____ ALLERGIES: _____

IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: [PRIMETHERAPEUTICS.COM/NOPP](https://www.primetherapeutics.com/nopp)

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE): _____
AUTHORIZED REPRESENTATIVE'S PHONE NUMBER: _____

PRESCRIBER INFORMATION	
LAST NAME:	FIRST NAME:
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:
NPI NUMBER:	DEA NUMBER:
PHONE NUMBER:	FAX NUMBER:
STREET ADDRESS:	
CITY:	STATE: ZIP CODE:
REQUESTER (if different than prescriber):	OFFICE CONTACT PERSON:

MEDICATION OR MEDICAL DISPENSING INFORMATION			
MEDICATION NAME:			
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
☐ NEW THERAPY	☐ RENEWAL	IF RENEWAL: DATE THERAPY INITIATED:	
DURATION OF THERAPY (SPECIFIC DATES):			

Continued on next page

Qfitlia (fitusiran)
Prior Authorization Request Form
Caterpillar Prescription Drug Benefit
Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME: _____ MEMBER'S FIRST NAME: _____

1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION?

☐ YES (if yes, complete below) ☐ NO

MEDICATION/THERAPY
(SPECIFY DRUG NAME AND
DOSAGE):

DURATION OF THERAPY
(SPECIFY DATES):

**RESPONSE/REASON FOR
FAILURE/ALLERGY:**

2. LIST DIAGNOSES:

ICD-10:

- ☐ Hemophilia A WITH inhibitors
☐ Hemophilia A WITHOUT inhibitors
☐ Hemophilia B WITH inhibitors
☐ Hemophilia B WITHOUT inhibitors
☐ Other diagnosis: _____ ICD-10 Code(s): _____

3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.

Will the patient be using the drug as a part of the clinical trial? ☐ Yes ☐ No

Hemophilia A WITH inhibitors

Has the patient developed high-titer factor VIII inhibitors (greater than 5 Bethesda units)?

☐ Yes ☐ No

Is the medication being prescribed for the prevention of bleeding episodes (routine prophylaxis)?

☐ Yes ☐ No

Hemophilia B WITH inhibitors

Has the patient developed high-titer factor IX inhibitors (greater than 5 Bethesda units)?

☐ Yes ☐ No

Is the medication being prescribed for the prevention of bleeding episodes (routine prophylaxis)?

☐ Yes ☐ No

Hemophilia A WITHOUT inhibitors

Does the patient have severe hemophilia with endogenous factor VIII levels that are less than 1% of normal (< 0.01 IU/mL)? ☐ Yes ☐ No

Does the patient have moderate hemophilia with endogenous factor VIII levels that are ≥1% and <5% (0.01–0.05 IU/mL)? ☐ Yes ☐ No

Does the patient have mild hemophilia with endogenous factor VIII levels that are ≥5% (≥ 0.05 IU/mL)? ☐ Yes ☐ No

Qfitlia (fitusiran)
Prior Authorization Request Form
Caterpillar Prescription Drug Benefit
Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME: _____ **MEMBER'S FIRST NAME:** _____

Hemophilia B WITHOUT inhibitors

Does the patient have severe hemophilia with endogenous factor IX levels that are less than 1% of normal (< 0.01 IU/mL)? ☐ Yes ☐ No

Does the patient have moderate hemophilia with endogenous factor IX levels that are ≥1% and <5% (0.01–0.05 IU/mL)? ☐ Yes ☐ No

Does the patient have mild hemophilia with endogenous factor IX levels that are ≥5% (≥ 0.05 IU/mL)? ☐ Yes ☐ No

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.

ATTESTATION: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or 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 or Electronic I.D. Verification: _____ **Date:** _____

CONFIDENTIALITY NOTICE: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return FAX) and arrange for the return or destruction of these documents.

FAX THIS FORM TO: 800-424-7640
MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program
Attn: CP-4201
P.O. Box 64811
St. Paul, MN 55164-0811
Phone: 877-228-7909