Isturisa (osilodrostat) Prior Authorization Request Form

Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

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.

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: <u>PRIMETHERAPEUTICS.COM/NOPP</u>

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

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

Continued on next page.



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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:		
Hyperortisolemia secondary to <u>endogenous</u> Cushing's Syndrome Other diagnosis:ICD-10				
3. REQUIRED CLINICAL INFORMATION: PRIOR AUTHORIZATION.	PLEASE PROVIDE ALL RELEVANT CLINIC	AL INFORMATION TO SUPPORT A		
Clinical Information: <u>Initial Request:</u> Is the prescriber an endocrinologist? Yes No Does the patient have hypercortisolemia? Yes No				
Is the patient's hypercortisolemia due	to endogenous Cushing's Syndrome?	Yes 🗆 No		
Is the endogenous Cushing's Syndrome caused by one of the following?* I Yes I No *Please provide documentation. I an ACTH-dependent (e.g., pituitary corticotrope adenoma, ectopic secretion of ACTH by nonpituitary tumor), I an ACTH-independent (e.g., adrenocortical adenoma, adrenocortical carcinoma, nodular adrenal hyperplasia Select if the patient has tried at least 2 of the listed therapies: *Please provide documentation. I Metyrapone Ketoconazole				
Has the patient failed surgery or are they not a candidate for surgery?* Yes No *Please provide documentation supported by a surgeon or anesthesiologist consult.				
<u>Renewal Request:</u> Is the prescriber an endocrinologist? Yes No Does the patient have a blood cortisol level or urinary free cortisol level at or below the upper limit of normal? Yes No <i>Please provide lab report documentation</i> .				
Was the lab blood cortisol level or urinary free cortisol level drawn more than 30 days prior to the request for renewal? \Box Yes \Box 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?				



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MEMBER'S LAST NAME:

MEMBER'S FIRST NAME:

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: _____ 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

