

Opfolda (miglustat)
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:
<input type="checkbox"/> NEW THERAPY	<input type="checkbox"/> RENEWAL	IF RENEWAL: DATE THERAPY INITIATED:	
DURATION OF THERAPY (SPECIFIC DATES):			

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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:

☐ Pompe disease

☐ Other diagnosis: _____ ICD-10 Code(s):

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

Is patient going to be using drug in combination with a clinical trial? ☐ Yes ☐ No

Does patient weighs > 40 kg? ☐ Yes ☐ No Please submit documentation.

Does patient have a sitting FVC \geq 30% of the predicted value prior to starting Opfolda? ☐ Yes ☐ No Please submit documentation.

Will Opfolda(miglustat) be used in combination with Pombiliti (cipaglucosidase alfa-atga)? ☐ Yes ☐ No

Did patient fail to demonstrate an improvement in objective measures after receiving ONE of the following for at least one year (Note: Examples of objective measures include forced vital capacity (FVC) and six-minute walk test (6MWT))? ☐ Yes ☐ No Please submit documentation.

☐ Lumizyme (alglucosidase alfa) intravenous infusion, OR

☐ Nexviazyme (avalglucosidase alfa-ngpt) intravenous infusion

Does patient have late-onset acid alpha-glucosidase deficiency (late-onset Pompe disease) with diagnosis established by ONE of the following? ☐ Yes ☐ No Please submit documentation.

☐ Patient has a laboratory test demonstrating deficient acid alpha-glucosidase activity in blood, fibroblasts, or muscle tissue, OR

☐ Patient has a molecular genetic test demonstrating biallelic pathogenic or likely pathogenic acid alpha-glucosidase (GAA) gene variants

Is Opfolda prescribed by or in consultation with a geneticist, neurologist, a metabolic disorder sub-specialist, or a physician who specializes in the treatment of lysosomal storage disorders? ☐ Yes ☐ No

Will Opfolda(miglustat) be used in combination with any another enzyme replacement medication or Miplyffa(arimoclomol)? ☐ Yes ☐ No Please submit documentation.

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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