Vornanigo (vorasidenib) **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.

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

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	OD MEDICAL DI		
MEDICATION	OR MEDICAL DI	SPENSING IN	FORMATION

MEDICATION NAME: DOSE/STRENGTH: FREQUENCY: LENGTH OF QUANTITY: THERAPY/REFILLS: 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 YES (if yes, complete below)	OTHER MEDICATIONS FOR THIS (CONDITION?		
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:		
2. LIST DIAGNOSES:		ICD-10:		
 Astrocytoma or Oligodendroglioma Other diagnosis: 				
3. REQUIRED CLINICAL INFORM. TO SUPPORT A PRIOR AUTHORI	ATION: PLEASE PROVIDE ALL REL ZATION.	EVANT CLINICAL INFORMATION		
Is patient going to be using drug Does patient have a diagnosis of G submit documentation.	in a clinical trial? 🗌 Yes 🗌 No rade 2 astrocytoma or oligodendrogi	ioma? 🗌 Yes 🗌 No Please		
Does tumor have IDH1 or IDH2 mutation? 🗌 Yes 🗌 No Please submit documentation.				
Has patient had at least one(1) surgery, including biopsy, sub-total resection, or gross total resection, prior to requesting Voranigo(vorasidenib)? Yes No Please submit documentation.				
Was surgery for the patient's astrocytoma or oligodendroglioma more than 5 years prior to the patient starting Voranigo(vorasidenib)? 🗌 Yes 🛛 No Please submit documentation.				
Has patient had any prior anticancer therapy, including chemotherapy and/or radiotherapy? Yes No Please submit documentation.				
Does patient have high-risk features including brainstem involvement, either a primary location or by tumor extension? Yes No Please submit documentation.				
Does patient have any clinically relevant functional or neurocognitive deficits due to the tumor? Yes No Please submit documentation.				
Does patient have uncontrolled seizures(defined as persistent seizures interfering with activities of daily life)? Yes No Please submit documentation.				
If patient has uncontrolled seizures, has the patient failed at least 3 lines of antiepileptic drug regimens including at least 2 combination regimen? I Yes I 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

