

Tibsovo (ivosidenib)
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.

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

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:
REQUESTOR (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):			

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:**

<input type="checkbox"/> Acute myeloid leukemia(AML) <input type="checkbox"/> Cholangiocarcinoma <input type="checkbox"/> Other diagnosis: _____ ICD-10 _____	
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3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.

Clinical Information:
Is drug going to be used in conjunction with a clinical trial? Yes No

Does patient have an IDH1 mutation? Yes No *Please submit documentation of mutation.*
Does patient have an ECOG score 0 or 1? Yes No *Please submit documentation.*
Does patient have an ECOG score 0 to 2? Yes No *Please submit documentation.*
Does patient have an ECOG score 2 or greater? Yes No *Please submit documentation.*
Has patient been previously treated with an IDH1 inhibitor such as but not limited to Idhifa(enasidenib)? Yes No *Please submit documentation.*

For Newly Diagnosis of Acute Myeloid Leukemia(AML), please aslo answer the following:

Is patient 75 years of age or older? Yes No
Is patient 18 to 74 years of age inclusive? Yes No
If patient is 18 to 74 years of age, does the patient have at least one comorbid condition below that precludes use of intensive induction chemotherapy? Yes No *Please submit documentation.*
 Baseline ECOG ≥ 2
 Severe cardiac or pulmonary disease, such as congestive heart failure with an EF $\leq 50\%$, chronic stable angina, or FEV1 $\leq 65\%$
 Hepatic impairment with bilirubin >1.5 times the limit of normal
 Creatine Clearance $<45\text{mL/min}$

Has patient been previously treated with azacitidine or decitabine for myelodysplastic syndrome
Will Tibsovo(ivosidenib) be used as monotherapy? Yes No
Will Tibsovo(ivosidenib) be used in combination with azacitidine? Yes No

For Relapsed or Refractory Acute Myeloid Leukemia(AML), please also answer the following

Has the patient had at least one prior chemotherapy treatment? Yes No *Please submit documentation.*

Is patient ineligible for chemotherapy? Yes No *Documentation why patient is ineligible for chemotherapy is required.*

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For Diagnosis of Cholangiocarcinoma, please also answer the following:

Does the patient have nonresectable or metastatic cholangiocarcinoma? Yes No *Please submit documentation.*

Has the patient's disease progressed following at least 1 or 2 prior regimens? Yes No *Please submit documentation.*

Has the patient's disease progressed following on 3 or more regimens? Yes No *Please submit documentation.*

Was one of the prior regimens gemcitabine or 5-FU containing regimens? Yes No *Please submit documentation.*

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