

**Tibsovo® (Ivosidenib) Prior Authorization Form**

Member Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Member ID#: \_\_\_\_\_

**Drug Information**

Pharmacy Billing (NDC: \_\_\_\_\_) Start Date (or date of next dose): \_\_\_\_\_

Dose: \_\_\_\_\_ Regimen: \_\_\_\_\_

**Pharmacy Information**

Pharmacy NPI: \_\_\_\_\_ Pharmacy Name: \_\_\_\_\_

Pharmacy Phone: \_\_\_\_\_ Pharmacy Fax: \_\_\_\_\_

**Prescriber Information**

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_

Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

**Criteria**

**For Initial Authorization:**

**1. Please indicate the diagnosis and information:**

**Acute Myeloid Leukemia (AML)**

A. Is AML newly-diagnosed? Yes \_\_\_ No \_\_\_

i. If member is younger than 75 years of age, are they unable to tolerate intensive induction chemotherapy? Yes \_\_\_ No \_\_\_

ii. Has an IDH1 mutation been detected? Yes \_\_\_ No \_\_\_

iii. Will Tibsovo® (ivosidenib) be used as a single-agent or in combination with azacitidine? Yes \_\_\_ No \_\_\_

B. Is AML relapsed or refractory? Yes \_\_\_ No \_\_\_

i. Will Tibsovo® (ivosidenib) be used as a single-agent? Yes \_\_\_ No \_\_\_

ii. Has an IDH1 mutation been detected? Yes \_\_\_ No \_\_\_

**Cholangiocarcinoma**

A. Is diagnosis locally advanced or metastatic cholangiocarcinoma? Yes \_\_\_ No \_\_\_

B. Has an IDH1 mutation been detected? Yes \_\_\_ No \_\_\_

C. Has the member received prior treatment for this diagnosis? Yes \_\_\_ No \_\_\_

**Myelodysplastic Syndromes (MDS)**

A. Is diagnosis relapsed or refractory MDS? Yes \_\_\_ No \_\_\_

B. Is there presence of isocitrate dehydrogenase-1 (IDH1) mutation, as detected by an FDA-approved test? Yes \_\_\_ No \_\_\_

**If answer is none of the above, please indicate diagnosis:** \_\_\_\_\_

Additional Information: \_\_\_\_\_

**For Continued Authorization:**

1. Date of last dose: \_\_\_\_\_

2. Does member have any evidence of progressive disease while on ivosidenib? Yes \_\_\_ No \_\_\_

3. Has the member experienced adverse drug reactions related to ivosidenib therapy? Yes \_\_\_ No \_\_\_

If yes, please specify adverse reactions: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

*I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Failure to complete this form in full will result in processing delays.*

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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