

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:** _____**Billing Provider 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 chemo-therapy? 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 UnitFax: 1-800-224-4014
Phone: 1-800-522-0114 Option 4**CONFIDENTIALITY NOTICE***This document, including any attachments, contains information which is confidential or privileged. If you are not the intended recipient, be aware that any disclosure, copying, distribution, or use of the contents of this information is prohibited. If you have received this document in error, please notify the sender immediately by telephone to arrange for the return of the transmitted documents or to verify their destruction.*