



Lytgobi® (Futibatinib) 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:

Unresectable, locally advanced, or metastatic intrahepatic cholangiocarcinoma

a. Was member previously treated with least 1 prior therapy? Yes ___ No ___

b. Is tumor positive for fibroblast growth factor receptor 2 (FGFR2) gene fusion or rearrangement?
Yes ___ No ___

If diagnosis is not listed 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 futibatinib therapy? Yes ___ No ___

3. Has member experienced any adverse drug reactions related to futibatinib therapy? Yes ___ No ___

If yes, please specify adverse reactions: _____

Additional information: _____

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 do not send in chart notes. Specific information will be requested if necessary.

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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