

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 requested information:**

A. Will cabozantinib be used a monotherapy? Yes \_\_\_ No \_\_\_

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

**Renal Cell Carcinoma (RCC)**

A. Is diagnosis advanced RCC? Yes \_\_\_ No \_\_\_

B. Will cabozantinib be used in combination with nivolumab for initial treatment of advanced RCC?  
Yes \_\_\_ No \_\_\_

i. Is the diagnosis relapsed or surgically unresectable stage 4 disease? Yes \_\_\_ No \_\_\_

[Please note: Opdivo<sup>®</sup> (nivolumab) requires prior authorization. The Opdivo<sup>®</sup> (nivolumab) prior authorization form (PHARM-64) is available on the OHCA website: <https://oklahoma.gov/ohca/providers/forms/rxforms.html>]

**Hepatocellular Carcinoma (HCC)**

A. Is diagnosis advanced HCC? Yes \_\_\_ No \_\_\_

B. Has the member previously received sorafenib? Yes \_\_\_ No \_\_\_

**Differentiated Thyroid Cancer (DTC)**

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

B. Has disease progressed following prior vascular endothelial growth factor (VEGF)-targeted therapy?  
Yes \_\_\_ No \_\_\_

C. Is disease radioactive iodine-refractory or is member ineligible for radioactive iodine?  
Yes \_\_\_ No \_\_\_

**If diagnosis is not listed above, please indicate diagnosis:** \_\_\_\_\_

**For Continued Authorization:**

1. Date of last dose: \_\_\_\_\_
2. Does member have any evidence of progressive disease while on cabozantinib? Yes \_\_\_ No \_\_\_
3. Has the member experienced adverse drug reactions related to cabozantinib 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.***

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