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This fax machine is located in a secure location as required by HIPAA regulations. Fax complete signed and dated forms to CVS/Caremark at . Please contact CVS/Caremark at 1-888-413-2723 with questions regarding the prior authorization process. When conditions are met, we will authorize the coverage of the medication.

Patient Name: Patient ID: Patient Group No:			_ Date: _ Patient Date Of Birth: Patient Phone:		9/6/2024 Physician Name:			
	<u>.</u>	NPI#:		Spec	Specialty: Physician Office Telephone			
Phy	ysician Office Address:							
Dru	ig Name (specify drug)			_				
	antity:							
			Expected Length of Therapy:ICD Code:					
Cor								
		te answer for each applica	ble question.					
1.	What is the patient's dia Non-small cell lung ca cancer) (If checked, g	•	tases from non-small cell lung					
	Pancreatic cancer (If							
	Chordoma (If checked	d, go to 2)						
	Renal cell carcinoma	(If checked, go to 2)						
	Other, please specify.	. (If checked, no further ques	stions)					
2.	Is the patient currently re	eceiving treatment with the r	requested medication?	Y		N		
3.	Is the patient currently re	eceiving treatment with the r	requested medication?	Y		N		
4.	Is there evidence of una regimen?	cceptable toxicity or disease	e progression while on the current	Υ		N		
5.	Is the disease T790M ne	egative?		Y		N		
6.	Is there evidence of eith current regimen?	er unacceptable toxicity or c	disease progression while on the					
	Yes, unacceptable to	xicity (If checked, no further	questions)					
	Yes, disease progress	sion (If checked, no further o	questions)					
	No (If checked, no fur	ther questions)						
7.	Is there evidence of una regimen?	cceptable toxicity or disease	e progression while on the current	Y		N		
8.	What is the patient's dia	gnosis?						
	Non-small cell lung ca cancer) (If checked, g	ancer (including brain metas jo to 9)	tases from non-small cell lung					
	Pancreatic cancer (If	checked, go to 13)						
	Chardoma (If checked	d go to 15)						

	Renal cell carcinoma (If checked, go to 17)				
9.	What is the clinical setting in which the requested drug will be used?				
	Recurrent disease (If checked, go to 10)				
	Advanced disease (If checked, go to 10)				
	Metastatic disease (If checked, go to 10)				
	Other, please specify. (If checked, no further questions)				
10.	Does the patient have sensitizing epidermal growth factor receptor (EGFR) mutation-positive disease? ACTION REQUIRED: If Yes, attach chart note(s) or test results of EGFR mutation.				
	Yes (If checked, go to 11)				
	No (If checked, no further questions)				
	Unknown (If checked, no further questions)				
	ACTION REQUIRED: Submit supporting documentation				
11.	Will the requested drug be used as a single agent?	Y		N	
12.	Will the requested drug be used in combination with ramucirumab or bevacizumab?	Y		N	
13.	What is the clinical setting in which the requested drug will be used? Locally advanced disease (If checked, go to 14)				
	Unresectable disease (If checked, go to 14)				
	Recurrent disease (If checked, go to 14)				
	Metastatic disease (If checked, go to 14)				
	Other, please specify. (If checked, no further questions)				
14.	Will the requested drug be used in combination with gemcitabine?	Υ		N	
15.	What is the clinical setting in which the requested drug will be used?				
	Recurrent disease (If checked, go to 16)				
	Other, please specify. (If checked, no further questions)				
16.	Will the requested drug be used as a single agent?	Y		N	
17.	What is the clinical setting in which the requested drug will be used? Relapsed disease (If checked, go to 18)		П		
	Stage IV disease (If checked, go to 18)		П		
	Other, please specify. (If checked, no further questions)				
18.	Does the disease have non-clear cell histology?	Y		N	
19.	Will the requested drug be used as a single agent?	Y		N	
20.	Will the requested drug be used in combination with bevacizumab?	Υ		N	

Γ

I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.

Prescriber (Or Authorized) Signature and Date

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