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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: Physician Office Address:			<pre>_ Date: _ Patient Date Of Birth:</pre>		9/9/2024				
		NPI#:	Patient Phone:	Physician Name: Specialty:					
				Phys	sician C	Office	Telephone:		
Drug Name (specify drug)									
Quantity: Route of Administration:		Frequency:	Strer	igth:					
Dia	gnosis:		_ ICD Code:						
Con									
	ase check the appropriat What is the diagnosis?	te answer for each applica	ble question.						
1.	Ū	ancer (If checked, go to 2)							
	Anaplastic thyroid cancer (If checked, go to 8)								
	Thyroid cancer (If checked, go to 13)								
	Other, please specify. (If checked, no further questions)								
2.	Is the patient currently r	eceiving treatment with the r	requested medication?	Y		N			
3.	Is there evidence of una regimen?	acceptable toxicity or disease	e progression while on the current	Y		Ν			
4.	What is the clinical setti	ng in which the requested m	edication will be used?						
	Recurrent disease (If checked, go to 5)								
	Advanced disease (If	checked, go to 5)							
	Metastatic disease (If	checked, go to 5)							
	Other, please specify	. (If checked, no further ques	stions)						
5.	Will the requested medi	cation be used as a single a	gent?	Y		N			
6.	Does the patient have a REQUIRED: If Yes, atta	rearranged during transfect ach chart note(s) or test resu	ion (RET) gene fusion? ACTION Its for RET gene fusion.						
	Yes (If checked, go to	o 7)							
	No (If checked, no fur	rther questions)							
	Unknown (If checked	, no further questions)							
7.	Has the patient experier positive-targeted regime		n therapy with a RET rearrangement	Y		Ν			
8.	Is the patient currently r	eceiving treatment with the r	requested medication?	Y		N			

9.	Is there evidence of unacceptable toxicity or disease progression on the current regimen?	Y		N	
10.	What is the clinical setting in which the requested medication will be used? Stage IV disease (If checked, go to 11)				
	Other, please specify. (If checked, no further questions)				
11.	Does the patient have a rearranged during transfection (RET) gene fusion? ACTION REQUIRED: If Yes, attach chart note(s) or test results for RET gene fusion. Yes (If checked, go to 12)				
	No (If checked, no further questions)				
	Unknown (If checked, no further questions)				
12.	Will the requested medication be used as a single agent?	Y		N	
13.	Is the patient currently receiving treatment with the requested medication?	Y		N	
14.	Is there evidence of unacceptable toxicity or disease progression on the current regimen?	Y		N	
15.	Which of the following applies to the patient's disease? Follicular thyroid cancer (If checked, go to 16)				
	Oncocytic thyroid cancer (If checked, go to 16)				
	Papillary thyroid cancer (If checked, go to 16)				
	Other, please specify. (If checked, no further questions)				
16.	What is the clinical setting in which the requested medication will be used? Advanced disease (If checked, go to 17)				
	Metastatic disease (If checked, go to 17)				
	Other, please specify. (If checked, no further questions)				
17.	Is the disease amenable to radioactive iodine therapy (RAI)?	Y		N	
18.	What is the patient's age (in years)?				
	Less than 12 years old (If checked, no further questions)				
	Greater than or equal to 12 years old (If checked, go to 19)				
19.	Does the patient have a rearranged during transfection (RET) gene fusion? ACTION REQUIRED: If Yes, attach chart note(s) or test results for RET gene fusion.				
	Yes (If checked, no further questions)				
	No (If checked, no further questions)				
	Unknown (If checked, no further questions)				

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.

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