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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:		7/17/2024				
		NPI#:	Patient Phone:	Physician Name: Specialty: Physician Office Telephone				
Phy	vsician Office Address:							
Dru	g Name (specify drug)			_				
Quantity: Route of Administration: Diagnosis:		• •	Streng	-				
Con								
Plea 1.	ase check the appropria What is the patient's dia	te answer for each applica agnosis?	ble question.					
	Breast cancer (If chee	cked, go to 2)						
	Central nervous syste	em (CNS) metastases from b	breast cancer (If checked, go to 2)					
	Chordoma (If checke	d, go to 2)						
	Colorectal cancer, inc (If checked, go to 2)	cluding appendiceal adenoca	arcinoma and anal adenocarcinoma					
	Other, please specify	. (If checked, no further que	stions)					
2.	Is this a request for con	tinuation of therapy with the	requested medication?	Y		N		
3.	Is there evidence of una regimen?	acceptable toxicity or disease	e progression while on the current	Y		N		
4.	What is the patient's dia	agnosis?						
	Breast cancer (If cheo	cked, go to 7)						
	Central nervous syste	em (CNS) metastases from b	breast cancer (If checked, go to 5)					
	Chordoma (If checke	d, go to 12)						
	Colorectal cancer, ind (If checked, go to 15)	cluding appendiceal adenoca	arcinoma and anal adenocarcinoma					
5.		attach chart note(s) or test i	r receptor 2 (HER2) status? ACTION results of human epidermal growth					
	HER2-positive (If checked, go to 6)							
	HER2-negative (If ch	ecked, no further questions)						
	Unknown (If checked	, no further questions)						
	Υ.	: Submit supporting docume	entation					
6.	Will the requested medi	cation be given in combinati	on with capecitabine?	Y		N		

7. What is the clinical setting in which the requested medication will be used?

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	No response to preoperative systemic therapy (If checked, go to 8)		
	Recurrent disease (If checked, go to 8)		
	Advanced disease (If checked, go to 8)		
	Metastatic disease (If checked, go to 8)		
	Other, please specify. (If checked, no further questions)		
8.	What is the patient's human epidermal growth factor 2 (HER2) status? ACTION REQUIRED: If HER2 positive, attach chart note(s) or test results of human epidermal growth factor receptor 2 (HER2) status.		
	HER2-positive (If checked, go to 9)		
	HER2-negative (If checked, no further questions)		
	Unknown (If checked, no further questions)		
	ACTION REQUIRED: Submit supporting documentation		
9.	Does the patient have hormone receptor-positive disease? ACTION REQUIRED: If Yes, attach chart note(s) or test results of hormone receptor status.		
	Yes (If checked, go to 10)		
	No (If checked, go to 11)		
	Unknown (If checked, go to 11)		
	ACTION REQUIRED: Submit supporting documentation		
10.	Will the requested medication be given in any of the following regimens?	_	
	In combination with an aromatase inhibitor (e.g., letrozole, anastrazole, or exemestane) (If checked, no further questions)		
	In combination with an aromatase inhibitor (e.g., letrozole, anastrazole, or exemestane) with trastuzumab (Herceptin) (If checked, no further questions)		
	None of the above (If checked, go to 11)		
11.	Will the requested medication be given in combination with capecitabine (Xeloda) or trastuzumab (Herceptin)?	Y 🔲	N 🔲
12.	What is the clinical setting in which the requested medication will be used?		
	Recurrent disease (If checked, go to 13)		
	Other, please specify. (If checked, no further questions)		
13.	What is the patient's epidermal growth factor receptor (EGFR) status? ACTION REQUIRED: If EGFR positive, attach chart note(s) or test results of epidermal growth factor receptor (EGFR) status.		
	EGFR-positive (If checked, go to 14)		
	EGFR-negative (If checked, no further questions)		
	Unknown (If checked, no further questions)		
	ACTION REQUIRED: Submit supporting documentation		
14.	Will the requested medication be given as a single agent?	Y	N 🗌
15.	What is the patient's human epidermal growth factor receptor 2 (HER2) status? ACTION REQUIRED: Attach chart note(s) or test results of human epidermal growth factor receptor 2 (HER2) status.		
	HER2-amplified (If checked, go to 16)		
	Other or unknown, please specify. (If checked, no further questions)		

ACTION REQUIRED: Submit supporting documentation

16.	Does the patient have RAS and BRAF wild-type disease? ACTION REQUIRED: If Yes, attach chart note(s) or test results RAS and BRAF mutation status.			
	Yes (If checked, go to 17)			
	No (If checked, no further questions)			
	Unknown (If checked, no further questions)			
	ACTION REQUIRED: Submit supporting documentation			
17.	Will the requested medication be used in combination with trastuzumab (Herceptin)?	Y	Ν	
18.	Is the patient appropriate for intensive therapy?	Y	Ν	
19.	Will the requested medication be used as subsequent therapy for progression of advanced or metastatic disease?	Y	Ν	

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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