PA Request Criteria

9.

What is the requested regimen?





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Patient Name: Patient ID: Patient Group No:		Date: Patient Date Of Birt Patient Phone:		6/13/2025 Physician Name: Specialty:			
Phy	ysician Office Address:			Phys	sicián C	ffice	Telephone
_	tient ID: tient Group No: NPI#: ysician Office Address: ug Name (specify drug) antity: ute of Administration: Patient Date Of Birth: Patient Phone: Patient Date Of Birth: Patient Phone:						
				nath:			
Route of Administration:							
Dia	gnosis:		ICD Code:				
Cor							
Ple 1.	What is the diagnosis?		·				
	·		intolleal calicel (il checked, go to 2)				
	-						
	•	,					
	Other, please specify	. (If checked, no further q	uestions)		Ш		
2.	Is the patient currently re	eceiving treatment with th	ne requested medication?	Υ		N	
3.	Is there evidence of una	cceptable toxicity while o	on the current regimen?	Υ		N	
4.	Is there evidence of dise	ease progression while or	n the current regimen?	Υ		N	
5.	· ·						
	Epithelial ovarian, fall	opian tube, or primary pe	ritoneal cancer (If checked, go to 6)				
	Uterine leiomyosarco	ma (If checked, go to 12)					
	Prostate cancer (If ch	ecked, go to 16)					
6.	Is the requested medica	tion being used as mainte	enance treatment?	Υ		N	
7.			platinum-based (e.g., cisplatin,	Y		N	
8.	What clinical setting will	the requested medication	n be used?				
	Advanced (Stage II-I\	/) disease (If checked, go	o to 9)				
	Recurrent disease (If	checked, go to 10)					
	Other, please specify	. (If checked, no further q	uestions)				

Single agent (If checked, no further questions)		
In combination with bevacizumab (Avastin) (If checked, no further questions)		
Other, please specify. (If checked, no further questions)		
What is the requested regimen?	_	
Single agent (If checked, go to 11)	Ш	
Other, please specify. (If checked, no further questions)		
Does the patient have a deleterious or suspected deleterious germline BRCA mutation? ACTION REQUIRED: If Yes, attach chart note(s) or test results confirming BRCA mutation status.	ı	
Yes (If checked, no further questions)		
No (If checked, no further questions)		
Unknown (If checked, no further questions)		
ACTION REQUIRED: Submit supporting documentation		
Does the patient have BRCA2-altered uterine leiomyosarcoma? ACTION REQUIRED: If Yes, attach chart note(s) or test results confirming BRCA2 mutation status. Yes (If checked, go to 13)		
No (If checked, no further questions)		
Unknown (If checked, no further questions) ACTION REQUIRED: Submit supporting documentation		
What is the place in therapy in which the requested medication will be used? First-line treatment (If checked, no further questions)		
Subsequent treatment (If checked, go to 14)		
Will the requested medication be used as a single agent?	Y 🗀	N 🗆
What is the clinical setting in which the requested medication will be used? Advanced disease (If checked, no further questions)	П	
Recurrent disease (If checked, no further questions)		
Metastatic disease (If checked, no further questions)		
Inoperable disease (If checked, no further questions)		
Other, please specify. (If checked, no further questions)		
What is the clinical setting in which the requested medication will be used?		
Metastatic disease (If checked, go to 17)		
Other, please specify. (If checked, no further questions)		
Is the disease castration-resistant?	Υ 🔲	N 🗆
Does the disease have BRCA1 or BRCA2 mutation? ACTION REQURED: If Yes, attach chart note(s) or test results confirming BRCA mutation status.		
Yes (If checked, go to 19)		
No (If checked, no further questions)		
Unknown (If checked, no further questions)		

	ACTION REQUIRED: Submit supporting documentation			
19.	Has the patient had a bilateral orchiectomy?	Υ	N	
20.	Will the requested medication be used in combination with a luteinizing hormone-releasing hormone (LHRH) agonist (e.g., goserelin, leuprolide) or antagonist (e.g., degarelix, relugolix)?	Y	N	
21.	Will the requested medication be used in combination with abiraterone and concurrent steroids (prednisone or methylprednisolone)?	Y	N	
and tr	st that the medication requested is medically necessary for this patient. I further attest that the informatio ue, and that the documentation supporting this information is available for review if requested by the claim sponsor, or, if applicable a state or federal regulatory agency.			

Prescriber (Or Authorized) Signature and Date

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