



Please complete ALL information below and fax your request to 1-888-671-5285

Lynparza® Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:	Specialty:	
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:	Dosage Form:	
<input type="checkbox"/> Check if generic substitution is acceptable			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below:					
<input type="checkbox"/> Advanced ovarian cancer					
<input type="checkbox"/> Breast cancer					
<input type="checkbox"/> Epithelial ovarian cancer					
<input type="checkbox"/> Fallopian tube cancer					
<input type="checkbox"/> Primary peritoneal cancer					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Prescriber's Specialty:					
Is Lynparza prescribed by or in consultation with an oncologist? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For advanced ovarian cancer, answer the following:					
Is there presence of deleterious or suspected deleterious germline BRCA-mutations as detected by an FDA-approved test or Clinical Laboratory Improvement Amendments-approved facility? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient had trial and failure, contraindication, or intolerance to three or more prior lines of chemotherapy (e.g., paclitaxel with cisplatin)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For breast cancer, answer the following:					
Does the patient have metastatic disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is there presence of deleterious or suspected deleterious germline BRCA-mutations as detected by an FDA-approved test or Clinical Laboratory Improvement Amendments-approved facility? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have human epidermal growth factor receptor 2 (HER2)-negative disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient been previously treated with chemotherapy (e.g., anthracycline, taxane) in the neoadjuvant, adjuvant, or metastatic setting? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have hormone-receptor (HR)-positive disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Has the patient been treated with prior endocrine therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
If "no" to the above question, is the patient considered to be an inappropriate candidate for endocrine therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For epithelial ovarian cancer, fallopian tube cancer, or primary peritoneal cancer, answer the following:					
Does the patient have recurrent disease? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Will Lynparza be used for maintenance treatment in patients who are in complete or partial response to platinum -based chemotherapy (e.g., cisplatin, carboplatin)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Reauthorization:					
If this is a reauthorization request, answer the following question:					
Does the patient show evidence of progressive disease while on Lynparza therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					

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Office use only: Lynparza_FSP_2018Jun-W



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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.