

Prior Authorization Form



**Note:** Please provide as much information as possible on this form. Missing data may cause processing delays for requested authorization(s). Attach additional sheets to this form if necessary.

Please fax the completed PA form and any additional informational sheets to Nirvanahealth at the following fax number: +1(866) 871-8565

Patient Information	Prescriber Information
Patient Name: _____	Prescriber Name: _____
Health Plan Name: _____	Prescriber Address: _____
Patient Insurance Id: _____	_____
Patient Date of Birth: _____	Prescriber Phone: (     ) _____
Patient Phone: _____	Prescriber Fax: (     ) _____
	Prescriber Specialty: _____
	Prescriber DEA: _____
	Prescriber NPI: _____

Medication & Medical Information	
Requested Drug(s) & Strength(s):	<input type="checkbox"/> Zejula 100 mg capsule
Requested Daily Quantity Limit – Amount:	
Requested Daily Quantity Limit – Days:	
Requested Quantity Limit Over Time – Amount:	
Requested Quantity Limit Over Time – Days:	
Requested Quantity Per Rx – Amount:	
Expected Length of Therapy:	
Directions:	
Diagnosis and Diagnosis Codes (ICD-10 Standard Codes):	
List drugs used previously to treat the same condition:	
Additional clinical information or history. Please include any relevant test results and/or medical record notes:	

### Questionnaire

Q1: I, as the provider or designated representative of the provider, certify and attest that the information provided is complete and accurate and that, upon request, I shall provide any information to RxAdvance that RxAdvance determines is reasonably necessary to verify my responses. (Check only one that apply)

Yes

No

Q2: Is the member currently treated with this medication? (Check only one that apply)

Yes (please list start date of therapy (month/day/year)) \_\_\_\_\_

(\*Required)

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No

Q3: What is the member's diagnosis? (Check only one that apply)

- Advanced epithelial ovarian cancer
- Advanced fallopian tube cancer
- Advanced primary peritoneal cancer
- Recurrent epithelial ovarian cancer
- Recurrent fallopian tube cancer
- Recurrent primary peritoneal cancer
- Other (please specify the member's diagnosis and provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q4: What is the member's diagnosis? (Check only one that apply)

- Advanced epithelial ovarian cancer
- Advanced fallopian tube cancer
- Advanced primary peritoneal cancer
- Recurrent epithelial ovarian cancer
- Recurrent fallopian tube cancer
- Recurrent primary peritoneal cancer
- Other (please specify the member's diagnosis and provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q5: Will the requested drug be used for maintenance treatment in patients who are in a complete or partial response to first-line platinum-based chemotherapy (e.g., cisplatin, carboplatin)? (Check only one that apply)

- Yes
- No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q6: Will the requested drug be used for maintenance treatment in patients who are in a complete or partial response to platinum-based chemotherapy (e.g., cisplatin, carboplatin)? (Check only one that apply)

- Yes
- No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q7: Has the member been treated with three or more prior chemotherapy regimens? (Check only one that apply)

- Yes (please specify treatment details) \_\_\_\_\_ (\*Required)
- No (please provide clinical rationale for the request) \_\_\_\_\_ (\*Required)

Q8: Is the member's cancer associated with homologous recombination deficiency (HRD) positive status defined by deleterious or suspected deleterious BRCA mutation? (Check only one that apply)

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Yes (please specify documentation of lab test, lab values and date of lab test)  
\_\_\_\_\_ (\*Required)

No

Q9: Is the member's cancer associated with homologous recombination deficiency (HRD) positive status defined by genomic instability? (Check only one that apply)

Yes (please specify) \_\_\_\_\_ (\*Required)

No (please provide clinical rationale for the request) \_\_\_\_\_  
(\*Required)

Q10: Is the member's cancer associated with homologous recombination deficiency (HRD) positive status defined as the cancer has progressed more than 6 months after response to the last platinum-based chemotherapy (e.g., cisplatin, carboplatin)? (Check only one that apply)

Yes (please specify) \_\_\_\_\_ (\*Required)

No (please provide clinical rationale for the request) \_\_\_\_\_  
(\*Required)

Q11: Is the medication prescribed by or in consultation with an oncologist? (Check only one that apply)

Yes

No (please provide clinical rationale for the request) \_\_\_\_\_  
(\*Required)

<b>Attestation:</b> I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, Insurer, Medical Group or its designated representatives may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.	
Signature of Prescriber or Authorized Representative:	Date:
Print Authorized Representative Name:	