

Member Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Member ID#: \_\_\_\_\_

**Drug Information**

Pharmacy billing (NDC: \_\_\_\_\_) Start Date (or date of next dose): \_\_\_\_\_

Dose: \_\_\_\_\_ Regimen: \_\_\_\_\_

**Billing Provider Information**

Provider NPI: \_\_\_\_\_ Provider Name: \_\_\_\_\_

Provider Phone: \_\_\_\_\_ Provider Fax: \_\_\_\_\_

**Prescriber Information**

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_

Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

**Criteria**

For Initial Authorization (Initial approval will be for the duration of 6 months):

1. Please indicate diagnosis and information:

**Advanced Recurrent/Refractory Ovarian Cancer Treatment**

A. Presence of deleterious or suspected deleterious germline BRCA mutation (gBRCAm)? Yes \_\_\_ No \_\_\_

B. Member previously treated with three or more lines of prior chemotherapy? Yes \_\_\_ No \_\_\_

i. If yes, please provide prior chemotherapy regimens: \_\_\_\_\_

**Maintenance Treatment of Advanced Ovarian Cancer**

A. Is member in complete or partial response to first-line platinum based chemotherapy? Yes \_\_\_ No \_\_\_

i. If yes, presence of deleterious or suspected deleterious germline BRCA mutation (gBRCAm)? Yes \_\_\_ No \_\_\_

B. Is member in complete or partial response to second-line or greater platinum based chemotherapy?

Yes \_\_\_ No \_\_\_

**Breast Cancer**

A. Metastatic breast cancer? Yes \_\_\_ No \_\_\_

B. Has member shown progression on previous chemotherapy in any setting? Yes \_\_\_ No \_\_\_

C. Human epidermal growth factor receptor 2 (HER2)-status? Positive \_\_\_ Negative \_\_\_

D. Positive test for germline BRCA-mutation? Yes \_\_\_ No \_\_\_

E. Hormone receptor (HR)-positive? Yes \_\_\_ No \_\_\_

i. If yes, has member failed prior endocrine therapy or considered to not be a candidate for endocrine therapy? Yes \_\_\_ No \_\_\_

**Epithelial Ovarian, Fallopian Tube, or Primary Peritoneal Cancer**

A. Will olaparib be used for maintenance treatment in recurrent disease where member had a complete or partial response to platinum-based chemotherapy? Yes \_\_\_ No \_\_\_

B. Has member completed platinum-based therapy in the past 8 weeks? Yes \_\_\_ No \_\_\_

**Other, please provide diagnosis:** \_\_\_\_\_

For Continued Authorization:

1. Date of last dose: \_\_\_\_\_

2. Does member have any evidence of progressive disease while on olaparib? Yes \_\_\_ No \_\_\_

3. Has member experienced adverse drug reactions related to olaparib therapy? Yes \_\_\_ No \_\_\_

If yes, please specify adverse reactions: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

**I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge.**

*Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.*

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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