

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

### Drug Information

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

### Billing Provider Information

Pharmacy NPI: \_\_\_\_\_ Pharmacy Name: \_\_\_\_\_  
Pharmacy Phone: \_\_\_\_\_ Pharmacy Fax: \_\_\_\_\_

### Prescriber Information

Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_  
Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Specialty: \_\_\_\_\_

### Criteria

**\*Page 1 of 2–Please complete and return all pages. Failure to complete all pages will result in processing delays.\*  
For Initial Authorization (Initial approval will be for the duration of 6 months):**

1. Please indicate diagnosis and information:

- Advanced Recurrent/Refractory Ovarian, Fallopian Tube, or Primary Peritoneal Cancer Treatment**
- A. Presence of deleterious or suspected deleterious germline BRCA mutation (*gBRCAm*)? Yes \_\_\_ No \_\_\_
  - B. Was member previously treated with 2 or more lines of prior chemotherapy? Yes \_\_\_ No \_\_\_
    - i. If yes, please provide prior chemotherapy regimens: \_\_\_\_\_
- Maintenance Treatment of Advanced Ovarian, Fallopian Tube, or Primary Peritoneal Cancer**
- A. Is disease in complete or partial response to primary chemotherapy? Yes \_\_\_ No \_\_\_
    - i. Will olaparib be used as a single-agent in deleterious or suspected deleterious *gBRCAm* or somatic BRCA-mutated (*sBRCAm*) disease? Yes \_\_\_ No \_\_\_
    - ii. Will olaparib be used in combination with bevacizumab following a primary therapy regimen that included bevacizumab? Yes \_\_\_ No \_\_\_
  - B. Is disease in complete or partial response to second-line or greater platinum-based chemotherapy? Yes \_\_\_ No \_\_\_
- Breast Cancer**
- A. Is diagnosis metastatic breast cancer? Yes \_\_\_ No \_\_\_
  - B. Has member shown progression on previous chemotherapy in any setting? Yes \_\_\_ No \_\_\_
  - C. Positive test for *gBRCAm*? Yes \_\_\_ No \_\_\_
  - D. 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 \_\_\_
- Pancreatic Cancer**
- A. Is diagnosis metastatic pancreatic adenocarcinoma with known germline BRCA1/BRCA2 mutation? Yes \_\_\_ No \_\_\_
  - B. Will olaparib be used as a single agent for maintenance therapy? Yes \_\_\_ No \_\_\_
  - C. Has member progressed on at least 16 weeks of first-line platinum-based chemotherapy? Yes \_\_\_ No \_\_\_
- Prostate Cancer**
- A. Is diagnosis metastatic castration-resistant prostate cancer? Yes \_\_\_ No \_\_\_
  - B. Has member failed previous first-line therapy? Yes \_\_\_ No \_\_\_
  - C. Will olaparib be used as a single-agent? Yes \_\_\_ No \_\_\_
    - i. If no, will olaparib be used with a gonadotropin-releasing hormone (GnRH) analog? Yes \_\_\_ No \_\_\_
    - ii. If no, does member have a prior history of bilateral orchiectomy? Yes \_\_\_ No \_\_\_
  - D. Is disease positive for a mutation in a homologous recombination gene? Yes \_\_\_ No \_\_\_

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

**Criteria**

**\*Page 2 of 2– Please complete and return all pages. Failure to complete all pages will result in processing delays.\***

**For Initial Authorization, continued:**

1. Please indicate diagnosis and information, continued:

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

**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:* \_\_\_\_\_

Additional Information: \_\_\_\_\_

DRAFT

**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.***

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