

Lynparza[®] (Olaparib) Prior Authorization Form

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:

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 disease human epidermal growth factor receptor 2 (HER2)-negative? Yes ___ No ___
 - B. Is disease high-risk early breast cancer previously treated with neoadjuvant or adjuvant chemotherapy?
Yes ___ No ___
 - i. Will olaparib be used in the adjuvant setting? Yes ___ No ___
 - ii. Positive test for *gBRCAm*? Yes ___ No ___
 - C. Is diagnosis metastatic breast cancer? Yes ___ No ___
 - i. Has member shown progression on previous chemotherapy? Yes ___ No ___
 - ii. Is disease hormone receptor (HR)-positive? Yes ___ No ___
 - 1. 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 ___

Page 1 of 2

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Fax completed prior authorization request form to **888-601-8461** or submit Electronic Prior Authorization through CoverMyMeds[®] or SureScripts.
All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at **AetnaBetterHealth.com/Oklahoma**.

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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:

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 ____

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: _____

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. Failure to complete all pages will result in processing delays.

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