

# OPTIMA HEALTH PLAN

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

**Directions:** The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; fax to **1-800-750-9692**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If information provided is not complete, correct, or legible, authorization may be delayed.**

**Drug Requested:** Orilissa® (elagolix)

**DRUG INFORMATION:** Authorization may be delayed if incomplete.

**Drug Form/Strength:** \_\_\_\_\_

**Dosing Schedule:** \_\_\_\_\_ **Length of Therapy:** \_\_\_\_\_

**Diagnosis:** \_\_\_\_\_ **ICD Code, if applicable:** \_\_\_\_\_

### **Quantity Limits:**

- 150 mg: Maximum of 1 tablet daily; maximum treatment duration of 24 months
- 200 mg: Maximum of 2 tablets daily; maximum treatment duration of 6 months

**\*Total collective approval duration not to exceed 24 months for all GnRH antagonist products\***

**CLINICAL CRITERIA:** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

☐ **Requested Dose: 150 mg, 1 tablet per day**

**Initial Authorization: 6 months**

- ☐ Member is premenopausal
- ☐ Member is 18 years of age or older
- ☐ Medication is being prescribed by or in consultation with a specialist in gynecology or reproductive health
- ☐ Member has a diagnosis of moderate to severe pain associated with endometriosis
- ☐ Diagnosis of endometriosis has been confirmed by direct visualization during surgery and/or histology
- ☐ Member does **NOT** have any contraindications to therapy including osteoporosis, severe hepatic impairment/disease, or concomitant use of hormonal contraceptives
- ☐ Member has history of inadequate response to the following therapies, tried for at least three (3) months each (**must submit chart note documentation of all therapy failures**):
  - ☐ NSAIDs (non-steroidal anti-inflammatory drugs)
  - ☐ Combination (estrogen/progesterone) oral contraceptive
  - ☐ Progestins

**OR**

- ☐ Member has had surgical ablation to prevent recurrence

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**Reauthorization: 18 months.** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

☐ Requested Dose: 150 mg, 1 tablet per day

**Note: Therapy will NOT exceed 24 months per lifetime**

- ☐ Member has improvement in pain associated with endometriosis (e.g., improvement in dysmenorrhea and non-menstrual pelvic pain)
- ☐ Member does **NOT** have any contraindications to therapy including osteoporosis, severe hepatic impairment/disease, or concomitant use of hormonal contraceptives
- ☐ Treatment duration of Orilissa® has not exceeded a total of 24 months.

**CLINICAL CRITERIA:** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

☐ Requested Dose: 200 mg, 2 tablets per day

**Authorization Criteria: Therapy will NOT exceed 6 months per lifetime**

- ☐ Member is premenopausal
- ☐ Member is 18 years of age or older
- ☐ Medication is being prescribed by or in consultation with a specialist in gynecology or reproductive health
- ☐ Member has a diagnosis of moderate to severe pain associated with endometriosis and coexisting condition of dyspareunia
- ☐ Diagnosis of endometriosis has been confirmed by direct visualization during surgery and/or histology
- ☐ Member does **NOT** have any contraindications to therapy including osteoporosis, severe hepatic impairment/disease, or concomitant use of hormonal contraceptives
- ☐ Member has history of inadequate response to the following therapies, tried for at least three (3) months each (**must submit chart note documentation of all therapy failures**):
  - ☐ NSAIDs (non-steroidal anti-inflammatory drugs)
  - ☐ Combination (estrogen/progesterone) oral contraceptive
  - ☐ Progestins

**OR**

- ☐ Member has had surgical ablation to prevent recurrence

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(Please ensure signature page is attached to form.)

Medication being provided by Specialty Pharmacy - PropriumRx

*Not all drugs may be covered under every Plan.*

*If a drug is non-formulary on a Plan, documentation of medical necessity will be required.*

***\*\*Use of samples to initiate therapy does not meet step edit/preauthorization criteria.\*\****

***\*Previous therapies will be verified through pharmacy paid claims or submitted chart notes.\****

Member Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

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

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

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

\*Approved by Pharmacy and Therapeutics Committee: ~~10/17/2018~~; 11/18/2022

REVISED/UPDATED: ~~12/30/2018~~; Reformatted 1/8/2020; 11/30/2022;