

**OPTIMA HEALTH COMMUNITY CARE
AND
OPTIMA FAMILY CARE
(MEDICAID)**

PHARMACY/MEDICAL 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 the information is not complete, correct, or legible, the authorization can be delayed.**

Drug Requested: **Orilissa[®]** (lusutrombopag)

DRUG INFORMATION: Complete information below or authorization will be delayed

Drug Form/Strength: _____

Dosing Schedule: _____ **Length of Therapy:** _____

Diagnosis: _____ **ICD Code, if applicable:** _____

CLINICAL CRITERIA: Check below **ALL** that apply. **ALL** criteria **must** be met. **ALL** documentation including labs or chart notes (if required) **must** be submitted or request will be denied.

Initial Authorization Approval – 6 months

1. The member has a confirmed diagnosis of endometriosis. Yes No

AND

2. Is the member 18 years old or older? Yes No

AND

3. The member has failed an adequate trial of the following therapies: Yes No

- non-steroidal anti-inflammatory drugs (NSAIDs),

AND

- hormonal contraceptives (including oral or transdermal formulations, vaginal ring, or intrauterine device),

AND

- gonadotropin-releasing hormone (GnRH) agonist (e.g., nafarelin [Synarel[®]], leuprolide [Lupron[®]], goserelin [Zoladex[®]])

AND

4. Elagolix is prescribed by or in consultation with an obstetrics/gynecology or reproductive specialist Yes No

AND

5. Pregnancy is excluded prior to initiating treatment. Yes No

AND

(continued on next page)

6. The member will use effective non-hormonal contraception during treatment with elagolix and 1 week after stopping therapy. Yes No

AND

7. The member does not have osteoporosis as evident by a Z score > -1.5 at spine and femur (total hip). Yes No

AND

8. The member does not have severe hepatic impairment (Child-Pugh C). Yes No

AND

9. The member is not on concomitant strong organic anion transport polypeptide (OATP) 181 inhibitor (e.g., cyclosporine, gemfibrozil). Yes No

Renewal Authorization Approval – 18 months. To receive approval for the 150mg tablet for a maximum duration of 24 months, the following questions must be checked. No Renewal for the 200mg tablet.

10. The member continues to meet the initial criteria. Yes No

AND

11. The member is considered to have clinically meaningful response to treatment. Yes No

Medication being provided by a Specialty Pharmacy – PropriumRx

****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.

Patient Name: _____

Member Optima #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

*REVISED/UPDATED: 2/25/2019