

**MYFEMBREE**  
**(relugolix, estradiol, and norethindrone acetate)**

## **Pre - PA Allowance**

None

---

## **Prior-Approval Requirements**

**Age** 18 years of age and older

**Gender** Female

### **Diagnoses**

Patient must have **ONE** of the following:

1. Heavy menstrual bleeding associated with uterine leiomyomas (fibroids)
2. Moderate to severe pain associated with endometriosis

**AND ALL** of the following:

1. Patient is premenopausal
2. Pregnancy has been excluded
3. Medication is being prescribed by or in consultation with an obstetrician-gynecologist (OB-GYN)
4. Patient has **NOT** already used Myfembree or Oriahnn cumulatively for 24 months
5. **NOT** used in combination with Oriahnn
6. Patient does **NOT** have current, or history of thrombotic or thromboembolic disorders **AND** patient is not at increased risk for these events (e.g., women over 35 years of age who smoke or women with uncontrolled hypertension)
7. **NO** known liver impairment or disease (e.g., clinically significant elevated transaminases > 2-3 times upper limit of normal, fibrosis F1-F4, etc.)
8. **NO** known osteoporosis
9. Prescriber agrees to monitor for suicidal ideation and mood disorders

## **Prior - Approval Limits**

**Quantity** 84 tablets per 84 days

**Duration** 12 months

---

## **Prior – Approval *Renewal* Requirements**

**Age** 18 years of age and older

**Gender** Female



**BlueCross  
BlueShield**

Federal Employee Program.

**MYFEMBREE**  
**(relugolix, estradiol, and norethindrone acetate)**

**Diagnoses**

Patient must have **ONE** of the following:

1. Heavy menstrual bleeding associated with uterine leiomyomas (fibroids)
2. Moderate to severe pain associated with endometriosis

**AND ALL** of the following:

1. Documented improvement in patient's condition
2. Medication is being prescribed by or in consultation with an obstetrician-gynecologist (OB-GYN)
3. Patient has **NOT** already used Myfembree or Oriahnn cumulatively for 24 months
4. **NOT** used in combination with Oriahnn
5. Patient does **NOT** have current, or history of thrombotic or thromboembolic disorders **AND** patient is not at increased risk for these events (e.g., women over 35 years of age who smoke or women with uncontrolled hypertension)
6. **NO** known liver impairment or disease (e.g., clinically significant elevated transaminases > 2-3 times upper limit of normal, fibrosis F1-F4, etc.)
7. **NO** known osteoporosis
8. Prescriber agrees to monitor for suicidal ideation and mood disorders

**Prior - Approval *Renewal* Limits**

**Quantity**      84 tablets per 84 days  
**Duration**      12 months – **One renewal ONLY**