

**PITUITARY SUPPRESSIVE AGENTS, LHRH PRIOR AUTHORIZATION FORM** (form effective 1/3/2022)

Prior authorization guidelines for **Pituitary Suppressive Agents, LHRH and Quantity Limits/Daily Dose Limits** are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	Total # of pgs: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	State license #:
LTC facility contact/phone:			Street address:	
Beneficiary name:			Suite #:	City/State/Zip:
Beneficiary ID#:	DOB:	Phone:	Fax:	

**CLINICAL INFORMATION**

Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.

Drug requested:	Strength:	
Directions/frequency:	Quantity:	Refills:
Diagnosis (submit documentation):	Dx code ( <i>required</i> ):	
<p><b>For a non-preferred Pituitary Suppressive Agent, LHRH:</b> Does the beneficiary have a history of trial and failure of or contraindication or intolerance to the preferred drugs in this class approved or medically accepted for treatment of the beneficiary's condition? Refer to <a href="https://papdl.com/preferred-drug-list">https://papdl.com/preferred-drug-list</a> for a list of preferred and non-preferred drugs in this class.</p>	<input type="checkbox"/> Yes – Submit documentation. <input type="checkbox"/> No	

**Complete the section(s) below applicable to the beneficiary and this request and SUBMIT DOCUMENTATION for each item.**

- For the treatment of CENTRAL PRECOCIOUS PUBERTY:**
  - Is prescribed the medication by or in consultation with a pediatric endocrinologist
  - Is female:
    - Is ≤11 years of age
    - Experienced onset of secondary sexual characteristics earlier than 8 years of age
  - Is male:
    - Is ≤12 years of age
    - Experienced onset of secondary sexual characteristics earlier than 9 years of age
- For the treatment of GENDER DYSPHORIA:**
  - Is prescribed the medication by or in consultation with an adult or pediatric endocrinologist or other provider with experience/training in transgender medicine
  - Is prescribed the medication in a manner consistent with current WPATH standards of care or other medical literature
- For the treatment of ENDOMETRIOSIS:**
  - Is prescribed the medication by or in consultation with a gynecologist
  - Diagnosis confirmed by laparoscopy
  - Diagnosis supported by chart documentation of adequate work-up that includes the clinical rationale for the diagnosis

<input type="checkbox"/> Tried and failed NSAIDs or has a contraindication or intolerance to NSAIDs
<input type="checkbox"/> Failed a 3-month trial of oral contraceptives or has a contraindication or intolerance to oral contraceptives
<input type="checkbox"/> <b>For PRESERVATION OF OVARIAN FUNCTION:</b>
<input type="checkbox"/> Is receiving cancer treatment that is associated with premature ovarian failure based on NCCN guidelines or peer-reviewed medical literature
<input type="checkbox"/> <b>For MYFEMBREE (relugolix/estradiol/norethindrone), ORIAHNN (elagolix/estradiol/norethindrone + elagolix), and ORILISSA (elagolix):</b>
<input type="checkbox"/> Has a history of depression and/or suicidal thoughts or behaviors OR is receiving treatment for depression and/or suicidal thoughts or behaviors
<input type="checkbox"/> Had a behavioral health assessment prior to use of the requested medication
<input type="checkbox"/> <b>For MYFEMBREE (relugolix/estradiol/norethindrone) and ORIAHNN (elagolix/estradiol/norethindrone + elagolix):</b>
<input type="checkbox"/> Is being treated for <b>HEAVY MENSTRUAL BLEEDING ASSOCIATED WITH UTERINE LEIOMYOMAS (FIBROIDS)</b>
<input type="checkbox"/> Is pre-menopausal
<input type="checkbox"/> Tried and failed a 3-month trial of or has a contraindication or intolerance to contraceptives

Please submit to PromptPA <https://ghp.promptpa.com> OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

<b>Prescriber Signature:</b>	<b>Date:</b>
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