Prior Authorization Review Panel

CHC-MCO Policy Submission

A separate copy of this form must accompany each policy submitted for review.
Policies submitted without this form will not be considered for review.

Plan: PA Health & Wellness	**Submission Date: N/A**
Policy Number: PHW.PDL.224	**Effective Date: 01/01/2020** **Revision Date: 07/2020**
Policy Name: Pituitary Suppressive Agents, LHRH	

Type of Submission – Check all that apply:

- ☐ **New Policy**
- ☐ **Revised Policy***
- ✓ **Annual Review - No Revisions**
- ✓ **Statewide PDL -** *Select this box when submitting policies for Statewide PDL implementation and when submitting policies for drug classes included on the Statewide PDL.*

***All revisions to the policy must be highlighted using track changes throughout the document.**

Please provide any changes or clarifying information for the policy below:

Q3 2020 annual review: no changes.

Name of Authorized Individual (Please type or print): **Francis G. Grillo, MD**	**Signature of Authorized Individual:**

Clinical Policy: Pituitary Suppressive Agents, LHRH

Reference Number: PHW.PDL.224
Effective Date: 01/01/2020
Last Review Date: 07/2020

Revision Log

Policy/Criteria
Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of health plans affiliated with PA Health and Wellness® that Luteinizing Hormone-Releasing Hormone Pituitary Suppressive Agents are **medically necessary** when the following criteria are met:

I. Requirements for Prior Authorization of Pituitary Suppressive Agents, LHRH

A. Prescriptions That Require Prior Authorization

All prescriptions for Pituitary Suppressive Agents, LHRH must be prior authorized.

B. Review of Documentation for Medical Necessity

In evaluating a request for prior authorization of a prescription for a Pituitary Suppressive Agent, LHRH, the determination of whether the requested prescription is medically necessary will take into account whether the beneficiary:

1. Is prescribed the Pituitary Suppressive Agent, LHRH for an indication that is included in the U.S. Food and Drug Administration (FDA)-approved package labeling OR a medically accepted indication; **AND**

2. Is age-appropriate according to FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**

3. Is prescribed a dose and duration of therapy that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**

4. Does not have a history of a contraindication to the prescribed medication; **AND**

5. For an adolescent with gender dysphoria, **both** of the following:

    a. Is prescribed the Pituitary Suppressive Agent, LHRH by or in consultation with a pediatric endocrinologist, adolescent medicine specialist, or medical provider with experience and/or training in transgender medicine

b. Is prescribed the Pituitary Suppressive Agent, LHRH in a manner consistent with the current World Professional Association for Transgender Health standards of care for the health of transsexual, transgender, and gender nonconforming people;

AND

6. For an adult with gender dysphoria, **both** of the following:

   a. Is prescribed the Pituitary Suppressive Agent, LHRH by or in consultation with an endocrinologist or medical provider with experience and/or training in transgender medicine
   b. Is prescribed the Pituitary Suppressive Agent, LHRH in a manner consistent with current medical literature;

   AND

7. For a diagnosis of endometriosis, **all** of the following:

   a. Has **one** of the following:

      i. A diagnosis of endometriosis confirmed by laparoscopy
      ii. A diagnosis of endometriosis supported by chart documentation of an adequate work-up that includes the clinical rationale for the diagnosis,

   b. Has a history of **all** of the following:

      i. Therapeutic failure, contraindication, or intolerance of non-steroidal anti- inflammatory drugs
      ii. Therapeutic failure (based on a 3-month trial), contraindication, or intolerance of oral contraceptives,

   c. Is prescribed the Pituitary Suppressive Agent, LHRH by or in consultation with a gynecologist,

   d. For Orilissa (elagolix), if the beneficiary has a history of depression and/or suicidal thoughts or behaviors or is currently receiving treatment for depression and/or suicidal thoughts or behavior, has a behavioral health assessment prior to use;

   AND

8. For a non-preferred Pituitary Suppressive Agent, LHRH, has a history of therapeutic failure, contraindication, or intolerance of the preferred Pituitary Suppressive Agents, LHRH approved or medically accepted for the beneficiary's

indication.

AND

9. If a prescription for a Pituitary Suppressive Agent, LHRH is for a quantity that exceeds the quantity limit, the determination of whether the prescription is medically necessary will also take into account the guidelines set forth in PA.CP.PMN.59 Quantity Limit Override.

NOTE: If the beneficiary does not meet the clinical review guidelines above but, in the professional judgement of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

C. Clinical Review Process

Prior authorization personnel will review the request for prior authorization and apply the clinical guidelines in Section B. above to assess the medical necessity of a prescription for a Pituitary Suppressive Agent, LHRH. If the guidelines in Section B. are met, the reviewer will prior authorize the prescription. If the guidelines are not met, the prior authorization request will be referred to a physician reviewer for a medical necessity determination. Such a request for prior authorization will be approved when, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary.

D. **Approval Duration: 6 months**

E. References

1. Hembree WC, Cohen-Kettenis P, Delemarre-van de Waal HA, et al. Endrocrine Treatment of Transsexual Persons: An Endocrine Society Clinical Practice Guideline. J Clin Endocrinol Metab. 2009;94(9):3132-3154.
2. Schmidt L, Levine R. Psychological Outcomes and Reproductive Issues Among Gender Dysphoric Individuals. Endocrinol Metab Clin N Am. 44(2015)773-785.
3. Coleman E, Bockting W, Botzer M, et al. Standards of care for the health of transsexual, transgender, and gender-nonconforming people, version 7. Int J Transgenderism 2011;13:165-232.
4. Medical Assistance Bulletin Number 99-16-11, Subject: Federal Final Rule, Nondiscrimination in Health Programs and Activities" and Implications for Coverage of Services Related to Gender Transition; http://www.dhs.pa.gov/publications/bulletinsearch/bulletinselected/index.htm?bn= 99-16- 11&o=N&po=OMAP&id=07/18/2016.
5. Schenken RS, Barbieri RL, Eckler K. Endometriosis: Pathogenesis, clinical features, and diagnosis. UpToDate. Accessed August 22, 2019.
6. Schenken RS, Barbieri RL, Eckler K. Endometriosis: Treatment of pelvic

pain. UpToDate. Accessed August 22, 2019.
7. Orilissa [package insert]. North Chicago, IL. AbbVie, Inc. July 2018.

Reviews, Revisions, and Approvals	Date
Policy created	01/01/2020
Q3 2020 annual review: no changes.	07/2020