



## Leuprolide Acetate Prior Authorization Request

Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. **Please complete the information requested on the form below and fax this form along with supporting clinical documentation to Priority Partners, toll-free at 1-866-212-4756 to initiate the review process.** If you have questions regarding the prior authorization please contact Priority Partners at 888-819-1043 Option 4.

**Patient's Name:** \_\_\_\_\_ **Date:** \_\_\_\_\_  
**Patient's ID:** \_\_\_\_\_ **Patient's Date of Birth:** \_\_\_\_\_  
**Physician's Name:** \_\_\_\_\_  
**Specialty:** \_\_\_\_\_ **NPI#:** \_\_\_\_\_  
**Physician Office Telephone:** \_\_\_\_\_ **Physician Office Fax:** \_\_\_\_\_

**Referring Provider Info:**  Same as Requesting Provider

**Name:** \_\_\_\_\_ **NPI#:** \_\_\_\_\_  
**Fax:** \_\_\_\_\_ **Phone:** \_\_\_\_\_

**Rendering Provider Info:**  Same as Referring Provider  Same as Requesting Provider

**Name:** \_\_\_\_\_ **NPI#:** \_\_\_\_\_  
**Fax:** \_\_\_\_\_ **Phone:** \_\_\_\_\_

*Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.*

**Required Demographic Information:**

*Patient Weight:* \_\_\_\_\_ kg

*Patient Height:* \_\_\_\_\_ cm

*Please indicate the place of service for the requested drug:*

- Ambulatory Surgical       Home       Off Campus Outpatient Hospital  
 On Campus Outpatient Hospital       Office

**Drug Information:**

*Strength/Measure* \_\_\_\_\_ *Units*  ml  Gm  mg  ea  Un

*Directions(sig)* \_\_\_\_\_ *Route of administration* \_\_\_\_\_

*Dosing frequency* \_\_\_\_\_

**Send completed form to: Priority Partners Fax: 1-866-212-4756**

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Leuprolide Acetate SGM – 10/2021.

**Priority Partners • 7231 Parkway Drive Suite 100 • Hanover, MD 21076**

**Phone: 888-819-1043 • Fax: 1-866-212-4756 • www.jhhc.com**

**Criteria Questions:**

1. What is the requested drug being used for?
  - Ovulation induction (e.g., intrauterine insemination [IUI])
  - Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT])
  - Central precocious puberty (CPP) (including use as a stimulation test to confirm diagnosis of CPP)
  - Prostate cancer
  - Treatment of advancing puberty and growth failure
  - Recurrent salivary gland tumors
  - Other
2. What is the ICD-10 code? \_\_\_\_\_

**Complete the following section based on the patient's diagnosis, if applicable.**

**Section A: Central Precocious Puberty**

3. Is leuprolide being requested for use as a stimulation test to confirm diagnosis of CPP?  
*If Yes, no further questions*  Yes  No
4. Is the patient currently receiving the prescribed therapy for central precocious puberty through a paid pharmacy or medical benefit?  Yes  No *If no, skip to #6*
5. Is the patient experiencing signs of treatment failure such as clinical pubertal progression, lack of growth deceleration and continued excessive bone age advancement?  Yes  No *No further questions*
6. Has the patient been evaluated for intracranial tumor(s) by appropriate lab tests and diagnostic imaging, such as computed tomography (CT scan), magnetic resonance imaging (MRI), or ultrasound?  Yes  No
7. Has the diagnosis of central precocious puberty been confirmed by a pubertal response to a GnRH (gonadotropin-releasing hormone) agonist test **or** a pubertal level of a third-generation LH (luteinizing hormone) assay? **Action Required: If yes, collect laboratory report or medical record of pubertal response to a GnRH agonist test or a pubertal level of a third-generation LH assay.**  Yes  No
8. Does the assessment of bone age versus chronological age support the diagnosis of central precocious puberty?  
 Yes  No
9. How old was the patient **AT THE ONSET** of secondary sexual characteristics? \_\_\_\_\_ years

**Section B: Prostate Cancer, and Recurrent Salivary Gland Tumors**

10. Is the patient currently receiving treatment with the requested medication?  Yes  No  
**For recurrent salivary gland tumors requests: If No, skip to #14**  
**For Prostate Cancer: If No, no further questions.**
11. Has the patient experienced clinical benefit while receiving the requested drug?  Yes  No
12. Has the patient experienced an unacceptable toxicity while receiving the requested drug?  
 Yes  No *No further questions*
13. *If the diagnosis is recurrent salivary gland tumors*, is the tumor androgen receptor positive?  Yes  No

**Section C: Advancing puberty and growth failure**

14. Is the patient also requesting or is currently receiving growth hormone?  Yes  No

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Section D: Infertility; Ovulation induction (e.g., intrauterine insemination [IUI]), Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT])

15. Is coverage for the drug being requested for a procedure that has been approved by the patient's medical benefit plan?  Yes  No *If No, skip to #18*
16. Has the medical authorization number been provided?  
 Yes, please indicate: \_\_\_\_\_  
 No, *skip to #18*  
 Not applicable, patient's medical benefit plan does not require precertification for the requested procedure
17. Please indicate the type of procedure that has been approved by the medical benefit plan: *No further questions*  
 Ovulation induction (e.g., intrauterine insemination [IUI])  
 Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT], intracytoplasmic sperm injection [ICSI])  
 Mature oocyte cryopreservation  
 Embryo cryopreservation  
 Preimplantation genetic diagnosis
18. What is the type of procedure the patient will be undergoing?  
 Ovulation induction (e.g., intrauterine insemination [IUI])  
 Assisted reproductive technology (e.g., in vitro fertilization [IVF], frozen embryo transfer, gamete intrafallopian transfer [GIFT], zygote intrafallopian transfer [ZIFT], intracytoplasmic sperm injection [ICSI])  
 Other
19. What is the intent of therapy?  
 Inhibition of premature luteinizing hormone (LH) surge  
 Ovulation trigger  
 Other

*I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by Priority Partners.*

X \_\_\_\_\_

Prescriber or Authorized Signature

Date (mm/dd/yy)

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