

## **Clinical Policy: Triptorelin Pamoate (Trelstar, Triptodur)**

Reference Number: CP.PHAR.175

Effective Date: 10.01.16

Last Review Date: 11.18

Line of Business: Commercial, HIM\*, Medicaid

[Coding Implications](#)

[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

### **Description**

Triptorelin pamoate (Trelstar<sup>®</sup> and Triptodur<sup>®</sup>) is a gonadotropin-releasing hormone (GnRH) receptor agonist.

### **FDA Approved Indication(s)**

- Trelstar is indicated for the palliative treatment of advanced prostate cancer.
- Triptodur is indicated for the treatment of pediatric patients 2 years and older with central precocious puberty (CPP).

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*\*For Health Insurance Marketplace (HIM), Triptodur is non-formulary and cannot be approved using these criteria; refer to the formulary exception policy, HIM.PA.103.*

### **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 Centene Corporation<sup>®</sup> that Trelstar and Triptodur are **medically necessary** when the following criteria are met:

#### **I. Initial Approval Criteria**

##### **A. Prostate Cancer** (must meet all):

1. Diagnosis of prostate cancer;
2. Request is for Trelstar;
3. Prescribed by or in consultation with an oncologist;
4. Age  $\geq$  18 years;
5. Request meets one of the following:
  - a. Dose does not exceed 3.75 mg per 4 weeks, 11.25 mg per 12 weeks, or 22.5 mg per 24 weeks;
  - a. Dose is supported by practice guidelines or peer-reviewed literature for the relevant off-label use (*prescriber must submit supporting evidence*).

**Approval duration: 12 months**

##### **B. Central Precocious Puberty** (must meet all):

1. Diagnosis of CPP confirmed by all of the following (a through c):
  - a. Elevated basal luteinizing hormone (LH) level  $>$  0.2 - 0.3 mIU/L (dependent on type of assay used) and/or elevated leuprolide-stimulated LH level  $>$  3.3 - 5 IU/L (dependent on type of assay used);

- b. Difference between bone age and chronological age was > 1 year (bone age-chronological age);
- c. Age at onset of secondary sex characteristics is < 8 years if female, or < 9 years if male;
2. Request is for Triptodur;
3. Member meets the following age requirements:
  - a. Female: 2 - 11 years;
  - b. Male: 2 - 12 years;
4. Prescribed by or in consultation with a pediatric endocrinologist;
5. Dose does not exceed 22.5 mg per 24 weeks.

**Approval duration: 12 months**

**C. Other diagnoses/indications**

1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid.

**II. Continued Therapy**

**A. Prostate Cancer** (must meet all):

1. Currently receiving medication via Centene benefit, or documentation supports that member is currently receiving Trelstar for prostate cancer and has received this medication for at least 30 days;
2. Request is for Trelstar;
3. Member is responding positively to therapy;
4. If request is for a dose increase, request meets one of the following (a or b):
  - a. New dose does not exceed 3.75 mg per 4 weeks, 11.25 mg per 12 weeks, or 22.5 mg per 24 weeks;
  - b. New dose is supported by practice guidelines or peer-reviewed literature for the relevant off-label use (*prescriber must submit supporting evidence*).

**Approval duration: 12 months**

**B. Central Precocious Puberty** (must meet all):

1. Currently receiving medication via Centene benefit or member has previously met all initial approval criteria;
2. Request is for Triptodur;
3. Member is responding positively to therapy (e.g., decreased growth velocity, cessation of menses, softening of breast tissue or testes, arrested pubertal progression);
4. Member meets the following age requirement:
  - a. Female: ≤ 11 years;
  - b. Male: ≤ 12 years.
5. If request is for a dose increase, new dose does not exceed Triptodur (IM): 22.5 mg per 24 weeks.

**Approval duration: 12 months**

**C. Other diagnoses/indications (must meet 1 or 2):**

1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.  
**Approval duration: Duration of request or 6 months (whichever is less);** or
2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid.

**III. Diagnoses/Indications for which coverage is NOT authorized:**

- A.** Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policies – CP.CPA.09 for commercial, HIM.PHAR.21 for health insurance marketplace, and CP.PMN.53 for Medicaid or evidence of coverage documents.

**IV. Appendices/General Information**

*Appendix A: Abbreviation/Acronym Key*

CPP: central precocious puberty

FDA: Food and Drug Administration

NCCN: National Comprehensive Cancer Network

GnRH: gonadotropin-releasing hormone

LH: luteinizing hormone

*Appendix B: Therapeutic Alternatives*

Not applicable.

*Appendix C: Contraindications/Boxed Warnings*

- Contraindication(s):
  - Hypersensitivity to triptorelin or any other component of the product, or other GnRH agonists or GnRH.
  - Pregnancy
- Boxed warning(s): None reported

**V. Dosage and Administration**

Drug Name	Indication	Dosing Regimen	Maximum Dose
Triptorelin pamoate (Trelstar)	Prostate cancer*	IM: 3.75 mg per 4 weeks; 11.25 mg per 12 weeks; 22.5 mg per 24 weeks	See regimen
Triptorelin pamoate (Triptodur)	CPP	IM: 22.5 mg IM every 24 weeks	See regimen

*\*May be used in combination with therapies such as radiation therapy, antiandrogens, glucocorticoids, docetaxel.*

**VI. Product Availability**

Drug Name	Availability
Triptorelin pamoate (Trelstar)	Single-dose vial for reconstitution with Mixject system (kit): 3.75 mg, 11.25 mg, 22.5 mg

Drug Name	Availability
Triptorelin pamoate (Triptodur)	Single-dose vial for reconstitution (kit): 22.5 mg

**VII. References**

1. Trelstar Prescribing Information. Irvine, CA: Allergan USA, Inc.; January 2018. Available at [https://www.allergan.com/assets/pdf/trelstar\\_pi](https://www.allergan.com/assets/pdf/trelstar_pi). Accessed July 30, 2018.
2. Triptodur Prescribing Information. Atlanta, GA: Arbor Pharmaceuticals, LLC; June 2017. Available at <https://www.accessdata.fda.gov>. Accessed July 30, 2018.
3. National Comprehensive Cancer Network Drugs and Biologics Compendium. Triptorelin pamoate. Available at [nccn.org](http://nccn.org). Accessed July 30, 2018.
4. National Comprehensive Cancer Network. Prostate cancer (Version 3.2018). Available at [nccn.org](http://nccn.org). Accessed July 30, 2018.
5. Klein K, Yang J, Aisenberg J, et al. Triptorelin Efficacy and safety of triptorelin 6-month formulation in patients with central precocious puberty. *J Pediatr Endocrinol Metab*. November 2016; 29(11): 1241–1248.
6. Kaplowitz P, Bloch C. Evaluation and referral of children with signs of early puberty. *Pediatrics*. 2016; 137(1): e20153732.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy split from CP.PHAR.118.GnRH Analogs. Max dose added; removed preferencing; staging of advanced prostate cancer restated as stage T3 through T4 or high risk through nodal/metastatic disease per guidelines; approval period extended to 12 months	02.16	02.16
Age removed. Formulations added. NCCN recommended uses added (prostate cancer; doses removed).	01.17	02.17
Age and dosing added for prostate cancer; positive therapeutic response examples added. Prostate cancer FDA/NCCN (categories 1 and 2A) indications listed separately. New drug/indication added: Triptodur/CP. Safety information removed (hypersensitivity).	09.17	11.17
4Q 2018 annual review; policies combined for Centene Medicaid, Commercial (CP.CPA.314; Triptodur only); added HIM; no significant changes; for oncology, summarized NCCN and FDA-approved uses for improved clarity (limited to diagnosis); specialist involvement in care and continuation of care added; no significant changes; references reviewed and updated.	08.07.18	11.18

**Important Reminder**

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program

approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

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**Note:**

**For Medicaid members**, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

**For Health Insurance Marketplace members**, when applicable, this policy applies only when the prescribed agent is on your health plan approved formulary. Request for non-formulary drugs must be reviewed using the non-formulary policy; HIM.PA.103.

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