

# Checklist for Prior Authorization Submission

Your patient's health plan will likely require prior authorization (PA) before it approves Triptodur (triptorelin). The checklist below provides basic guidance on what may be needed to obtain a PA decision. It's important to note that PA requirements will vary among insurers. We encourage health providers to review PA guidelines on the insurer's website or to contact the insurer's customer service to confirm requirements, forms, and contacts.

Use of this checklist does not guarantee coverage or that a health plan will provide reimbursement for Triptodur and is not intended to be a substitute for or to influence the independent medical judgment of the physician.

## Completed PA Request Form\*

If required, complete and submit the PA request form to the insurer. PA forms can be obtained through the insurer's website or by contacting the insurer's customer service department.

## Clinical Documentation

- Letter of Medical Necessity
- Office visit notes, progress report notes, and/or clinical notes
- Any laboratory findings such as:
  - Gonadotropin-Releasing Hormone (GnRH) levels: Luteinizing Hormone (LH) and Follicle-Stimulating Hormone (FSH)
  - Sex Steroid Levels: Estradiol, Testosterone
  - Measurement of height and bone age
- Other

## If the information below is not part of the PA request form, it may be beneficial to provide to the insurer:

- Patient name, date of birth, insurance policy number/member ID and other relevant information
- Physician and facility information (e.g., name, provider ID number and tax ID number)
- Relevant procedure and HCPCS codes for products/services to be provided/performed
- Relevant information regarding the treatment decision:
  - Triptodur prescribing information and NDC 24338-150-20
  - Peer-reviewed journal articles

For expedited requests, adequate information should be provided to support the urgent nature of the request.

\*Specific prior authorization forms may need to be completed for select products or therapeutic areas. Always verify that the correct form has been completed. HCPCS, Healthcare Common Procedure Coding System; NDC, National Drug Code.

Triptodur<sup>®</sup> is manufactured by Debiopharm Research & Manufacturing SA on behalf of Azurity Pharmaceuticals, Inc., and its applicable affiliates. Triptodur<sup>®</sup> is a registered trademark of Debiopharm International SA.

© 2023 Azurity Pharmaceuticals, Inc. All Rights Reserved. All brand names referenced herein are subject to trademark protection, and are the property of their respective owners. PP-TRIP-US-0727