



Updated: 03/2026
Approved: 04/2026

**Request for Prior Authorization for Testosterone Supplementation
Website Form – www.wv.highmarkhealthoptions.com
Submit request via: Fax - 1-833-547-2030.**

All requests for Testosterone Supplementation require a Prior Authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

Testosterone Supplementation Prior Authorization Criteria:

Coverage may be provided with a diagnosis of hypogonadism and the following criteria is met:

- The member is a male
- The member had or currently has at least two confirmed low testosterone levels according to current practice guidelines or your standard lab reference values
- The member has consistent signs and symptoms of androgen deficiency (e.g. incomplete or delayed sexual development; reduced sexual desire, activity, or spontaneous erections; breast discomfort or gynecomastia; loss of body hair (especially axillary and pubic hair) or reduced need for shaving; very small (< 5 mL) or shrinking testes; inability to father children, low or zero sperm count; height loss, low trauma fracture, low bone mineral density; hot flushes, sweats)
- The member has no contraindications to starting therapy (e.g. breast cancer, prostate cancer, erythrocytosis with hematocrit > 54%, untreated obstructive sleep apnea, severe lower urinary tract symptoms with American Urological Association/International Prostate Symptom Score [IPSS] >21, uncontrolled or poorly controlled heart failure due to risk of increased fluid retention and desire for fertility)
- Baseline PSA, lipids, and hematocrit laboratory levels have been performed
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- Is age-appropriate according to FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature
- Men over age 50 years (or over 40 years who have a family history of prostate cancer or are African-American) have been screened for prostate cancer
- Initial Duration of Approval: 6 months
- Reauthorization criteria:
 - Documentation of improvement in signs and symptoms and tolerance to therapy
 - Documentation that normal serum testosterone concentrations are being achieved
 - Repeat PSA, lipids, and hematocrit laboratory levels have been performed and are within normal range
- Reauthorization Duration of Approval: 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.



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When criteria are not met, the request will be forwarded to a Medical Director for review. The physician reviewer must override criteria when, in their professional judgment, the requested medication is medically necessary.

**TESTOSTERONE SUPPLEMENTATION
PRIOR AUTHORIZATION FORM**

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Highmark Health Options Pharmacy Services. **FAX: (833)-547-2030.**

If needed, you may call to speak to a Pharmacy Services Representative. **PHONE: 1-844-325-6251 Mon – Fri 8 am to 7 pm**

PROVIDER INFORMATION

Requesting Provider:	NPI:
Provider Specialty:	Office Contact:
Office Address:	Office Phone:
	Office Fax:

MEMBER INFORMATION

Member Name:	DOB:
Member ID:	Member weight: Height:

REQUESTED DRUG INFORMATION

Medication:	Strength:
Directions:	Quantity: Refills:
Is the member currently receiving requested medication? <input type="checkbox"/> Yes <input type="checkbox"/> No Date Medication Initiated:	
Is this medication being used for a chronic or long-term condition for which the medication may be necessary for the life of the patient? <input type="checkbox"/> Yes <input type="checkbox"/> No	

Billing Information

This medication will be billed: <input type="checkbox"/> at a pharmacy OR <input type="checkbox"/> medically, JCODE:
Place of Service: <input type="checkbox"/> Hospital <input type="checkbox"/> Provider's office <input type="checkbox"/> Member's home <input type="checkbox"/> Other

Place of Service Information

Name:	NPI:
Address:	Phone:

MEDICAL HISTORY (Complete for ALL requests)

Diagnosis:	ICD Code:
<input type="checkbox"/> Hypogonadism, ICD-10: _____ <ul style="list-style-type: none"> ➢ Is the member 18 years of age or older and a male? <input type="checkbox"/> Yes <input type="checkbox"/> No ➢ Are there <u>two</u> confirmed low testosterone levels according to current practice guidelines or the standard lab reference values? <input type="checkbox"/> Yes <input type="checkbox"/> No ➢ Are there consistent symptoms and signs of androgen deficiency? <input type="checkbox"/> Yes <input type="checkbox"/> No ➢ Does the member have any contraindications to therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No ➢ Have baseline PSA, lipids, and hematocrit been performed before initiation of therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No ➢ Has the member been screened for prostate cancer (if applicable)? <input type="checkbox"/> Yes <input type="checkbox"/> No 	

CURRENT or PREVIOUS THERAPY

Medication Name	Strength/ Frequency	Dates of Therapy	Status (Discontinued & Why/Current)

REAUTHORIZATION

Is the member demonstrating improvement in symptoms and tolerating therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No
Has the member achieved a normal serum testosterone level? <input type="checkbox"/> Yes <input type="checkbox"/> No

SUPPORTING INFORMATION or CLINICAL RATIONALE

Prescribing Provider Signature

Date

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