

SENTARA HEALTH PLANS

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-844-668-1550**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If information provided is not complete, correct, or legible, authorization can be delayed.**

For Medicare Members: Medicare Coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD) and Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. They can be found at: <https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx>. Additional indications may be covered at the discretion of the health plan.

Testosterone Replacement Therapy (TRT) Injectables **(MUST be purchased by Physician's Office)**

Drug Requested: (Check box below that applies)

PREFERRED		
<input type="checkbox"/> testosterone cypionate injection (J1071)	<input type="checkbox"/> testosterone enanthate injection (J3121)	
NON-PREFERRED		
<input type="checkbox"/> Aveed [®] (testosterone undecanoate) IM injection (J3145)	<input type="checkbox"/> Azmiro (testosterone cypionate) IM injection (J1072)	<input type="checkbox"/> Testopel [®] (testosterone pellets) (J1073)

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.

Member Name: _____

Member Sentara #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

NPI #: _____

DRUG INFORMATION: Authorization may be delayed if incomplete.

Drug Form/Strength: _____

Dosing Schedule: _____ Length of Therapy: _____

Diagnosis: _____ ICD Code, if applicable: _____

Weight (if applicable): _____ Date weight obtained: _____

(Continued on next page)

- Standard Review. In checking this box, the timeframe does not jeopardize the life or health of the member or the member's ability to regain maximum function and would not subject the member to severe pain.

NOTE: For the hypogonadism indication, testosterone drugs **CANNOT** be used in conjunction with other erectile dysfunction drugs.

Maximum Dosage:

- Aveed – 3 mL (750 mg) injected intramuscularly, followed by 3 mL (750 mg) injected after 4 weeks, then 3 mL (750 mg) injected every 10 weeks
- Azmiro – 50 to 400 mg administered every two to four weeks as an intramuscular injection
- Testopel – 75 mg implantable pellet; 6 pellets per 90-day supply (6 billable units every 90 days)

CLINICAL CRITERIA: Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Length of Authorization: 12 months

- Member must meet **ONE** of the following:
 - Member has Partial Androgen Insensitivity Syndrome with male gender identity/gender dysphoria or delayed male puberty
 - Member has hypogonadism confirmed by **BOTH** of the following:
 - TWO (2) MORNING (6AM to 11AM)** testosterone levels **obtained on different dates** that are below 300 ng/dL or below the lower limit of normal for the reference range (**attach lab results with reference ranges from the laboratory for both**)
 - First testosterone level: _____
 - Repeat testosterone or free testosterone level: _____
 - Member has at least **one** specific symptom and at least **two** non-specific symptoms [**NOTE: If 'decreased spontaneous erections' is the ONLY symptom documented in chart notes, the request will be denied as testosterone replacement is EXCLUDED from coverage for sexual dysfunction**]:

Specific Symptoms: (≥ 1 of the following)	Non-Specific Symptoms: (≥ 2 of the following)
<input type="checkbox"/> Incomplete or delayed sexual development <input type="checkbox"/> Reduced sexual desire (libido) and activity <input type="checkbox"/> Decreased spontaneous erections* <input type="checkbox"/> Breast discomfort, gynecomastia <input type="checkbox"/> Loss of body hair (axillary, facial, and/or pubic) <input type="checkbox"/> Small testes (<5 mL) or shrinking testes <input type="checkbox"/> Low or zero sperm count <input type="checkbox"/> Height loss, low trauma fracture or low bone mineral density <input type="checkbox"/> Hot flushes, sweats	<input type="checkbox"/> Decreased energy, motivation, initiative and self-confidence <input type="checkbox"/> Depressed mood <input type="checkbox"/> Poor concentration and memory <input type="checkbox"/> Sleep disturbances, increased sleepiness <input type="checkbox"/> Mild anemia (Hgb 10-11 gm/dL) <input type="checkbox"/> Reduced muscle bulk and strength due to cachexia <input type="checkbox"/> Increased body fat, BMI <input type="checkbox"/> Diminished physical or work performance

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AND

- Member has had an inadequate response, contraindication or intolerance to at least a three-month trial with a topical agent e.g., testosterone gel, testosterone patch, testosterone topical solution, testosterone nasal gel (verified by pharmacy paid claims)

_____ (Please document date/name of drug)

AND

- Member has had an inadequate response, contraindication or intolerance to at least a three-month trial of a preferred intramuscular testosterone medication e.g., testosterone cypionate or testosterone enanthate (verified by pharmacy or medical paid claims)

_____ (Please document date/name of drug)

Medication being provided by (check box below that applies):

- Physician's office **OR** Specialty Pharmacy

For urgent reviews: Practitioner should call Sentara Pre-Authorization Department if they believe a standard review would subject the member to adverse health consequences. Sentara's definition of urgent is a lack of treatment that could seriously jeopardize the life or health of the member or the member's ability to regain maximum function.

*****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.*****

****Previous therapies will be verified through pharmacy paid claims or submitted chart notes.****